

SUMMARY OF NARRATIVE SYNTHESIS OF THE HEALTH EFFECTS OF POTENTIAL DIETARY FIBRE COMPONENTS

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1. Introduction

The Food Standards Agency asked the Scientific Advisory Committee on Nutrition (SACN) to consider which components should be included in the definition of dietary fibre and to review the available evidence on potential dietary fibre components.

To enable SACN to assess the available evidence the Food Standards Agency with SACN input commissioned Human Nutrition Research (HNR), Cambridge (via competitive tender) to undertake a comprehensive review of the health effects of potential dietary fibre components following the guidelines outlined in the SACN framework for the evaluation of evidence (2002)

(www.sacn.gov.uk/reports_position_statements/reports/risk_assessment.html)

This paper was used as a background paper to enable SACN to produce a position statement the "SACN statement on Dietary Fibre" (www.sacn.gov.uk)

A summary of the evidence reviewed is presented in appendices A-E of this document.

2. Method

A narrative synthesis of the health effects of potential dietary fibre components was produced using the SACN Framework for the Evaluation of Evidence (2002) www.sacn.gov.uk/reports_position_statements/reports/risk_assessment.html

2.1 Literature

The literature review was based on a structured search of the electronic bibliographic database PubMed, and also the reference lists from relevant primary studies, meta analyses and review articles, including the WHO/FAO Scientific Update on Carbohydrate and Human Nutrition (2007), the CRC Handbook of Dietary Fibre in Human Nutrition, the Institute of Medicine of the National Academy of Sciences, Report on the Dietary Reference Intakes for Macronutrients, the WCRF Report on Food, Nutrition, Physical Activity and the Prevention of Cancer (2007) and research registers, such as The Cochrane Controlled Trials Register. The types of study that were included for the narrative synthesis were randomised control trials, acute experimental studies, prospective cohort and cross sectional studies.

2.2 Search terms and inclusion/exclusion criteria

2.2.1 Obesity and metabolic disease

The following criteria were used to help identify journal articles to be included in the review. Articles were included if:

- They were carried out in human subjects and reported in English
- Fibre was quantified and measured using the AOAC method, Asp method, Englyst method, Southgate method, or Neutral Detergent Fibre for insoluble fibre.
- For effects on body weight: studies reported either body weight as an outcome or another shorter-term outcome with a plausible link to body weight (including appetite, energy intake/expenditure hormones linked to satiety)
- For effects on metabolic health: studies reported type 2 diabetes incidence or another shorter-term outcome with a plausible link to this endpoint (including fasting or postprandial glucose/insulin, insulin sensitivity, glucose tolerance)

Exclusion criteria were as follows:

- 'Crude fibre' was used to define fibre
- Multiple dietary/lifestyle factors were changed such that the effect of fibre could not be separated from that of other dietary/lifestyle factors
- For weight and diabetes outcomes (but not shorter-term metabolic outcomes) crosssectional observational studies were excluded because of the potential for reversecausality
- Studies including subjects who were diabetic at the point of inclusion in the study (as treatment of diabetes was not considered in the report)
- Blood pressure and cholesterol/triacylglycerol outcomes were excluded from the obesity section as these were for considered in the context of cardiovascular

A PubMed search was performed using the following terms:

- Fiber OR Fibre OR Non-starch polysaccharide OR Resistant starch OR Fructo(-) oligosaccharide OR Galacto(-)oligosaccharide OR Polydextrose OR Inulin OR Lignin

AND

- Weight OR Obesity OR Obese OR Overweight OR BMI OR Hunger OR Satiety OR Fullness OR Appetite OR Insulin OR Glucose OR Diabetes OR Metabolic

7545 articles were identified; 595 were potentially relevant and abstracts examined in further detail; 242 full papers were obtained for consideration; 159 of these were relevant for inclusion. A further 42 papers were identified for inclusion from scrutiny of reference lists of journal articles and book chapters.

2.2.2 Cardiovascular disease

The following criteria were used to help identify journal articles to be included in the review. Articles were included if:

- They were carried out in human subjects and reported in English
- The fibre content was measured using the method of the Association of Official Analytical Chemists (AOAC), the Englyst method, Southgate method (or Neutral Detergent Fibre (NDF) in the case of insoluble fibre sources)
- They reported blood lipid fractions (i.e. total cholesterol, low density lipoprotein cholesterol, high density lipoprotein cholesterol, and/or triacylglycerol) or blood pressure.
- They specified the amount and the type of dietary fibre employed*

* Observational studies were included if the quantity of total dietary fibre could be determined from whole grain intake. If for example, dietary fibre (g/d) was tabulated according to quartile or quintiles of whole grain intake and met the other predetermined inclusion criteria, an observational study examining the association between dietary fibre intake and cardiovascular disease was included.

Articles were excluded if:

- Fibre content was defined as 'crude fibre'.
- The intervention was of a short duration (< 3 weeks)*
- Participants were taking lipid-lowering drugs or antihypertensive medication**
- Participants achieved significant weight loss (pre Vs post treatment)***
- A dietary portfolio (combination diet) was the intervention, e.g. the prescribed diet included plant sterols, soy protein, viscous fibres, and nuts etc.
- If the study reported C-Reactive Protein (CRP) as an intermediate marker of cardiovascular disease. It was considered that raised CRP levels were not a suitable indicator of disease related to atherosclerosis, especially in an epidemiological setting. Raised CRP measures may reflect inflammation in the body associated with infection and disease independent of cardiovascular disease.

* In proposing to authorise the use on food labels of health claims on the association between oat products and reduced risk of coronary heart disease, the Food and Drug Administration (1994) considered whether the intervention studies that it evaluated were of long enough duration to ensure stabilization of blood lipids (greater than or equal to 3 weeks duration).

** Studies were included if only a small number of the participants were reported to be taking medication to treat hyperlipidemia or to treat hypertension.or where participants were taking a stable dose of lipid-lowering or antihypertensive drugs the study has been included in the review.

In some instances a meta-analysis was taken as the starting point from which to search for relevant scientific literature. This was the case for several types of soluble fibre. The meta-analysis of Brown et al (1999) included 67 controlled trials and quantified the cholesterol-lowering effect of oat products, psyllium, pectin and guar gum. Studies published from 1966 to June 1996 were included in this meta-analysis. Relevant journal articles published subsequent to these dates were considered in relation to this review.

A literature search was also carried out on studies relating concerned with the relating blood pressure to fibre intake published subsequent to the meta-analysis of Streppel et al (2005).

A PubMed search was performed using the following terms:

- Fiber OR Fibre OR Oligosaccharide OR Resistant starch OR Fructo(-)oligosaccharide OR Galacto(-)oligosaccharide OR Polydextrose OR Inulin OR Lignin
- AND
- Cholesterol OR Cardiovascular disease OR Blood lipids OR Blood pressure OR Hypertension OR Guar gum OR Psyllium OR Pectin

2.2.3 Colorectal cancer

The following criteria were used to help identify journal articles to be included in the review. Articles were included if:

- They referred to human subjects and reported in English
- They specified the amount and the type of dietary fibre investigated
- The fibre content was measured using the method of the Association of Official Analytical Chemists (AOAC), the Englyst method, Southgate method (or Neutral Detergent Fibre (NDF) in the case of insoluble fibre sources)
- They reported incident colon, rectal or colorectal cancer, and/or incidence or recurrence of colorectal adenoma.

Articles were excluded if:

- Effect estimates for fibre as a main exposure were not given
- Data were not prospective
- The amount of fibre consumed per day was not given
- A combination diet was the intervention, e.g. the prescribed diet included increased fibre and reduced fat content
- Fibre content was defined as 'crude fibre'
- More than a single type of fibre was incorporated into the fibre supplement

Search Terms/Strategy

A PubMed search was performed using the following terms:

Fiber OR fibre OR resistant starch

AND

colorectal cancer OR colon cancer OR colorectal adenoma

Reference lists of appropriate meta-analyses such as the WCRF 2007 report on Food, Nutrition, Physical Activity, and the Prevention of Cancer, the Pooling Project of Prospective Studies of Diet and Cancer and the Cochrane Review on dietary fibre for the prevention of colorectal adenomas and carcinomas were scrutinised for additional relevant papers. Initial searches in PubMed: 197 papers, of which 22 were included. Searching of reference lists led to a further 4 papers. Searches using 'colorectal adenoma' instead of 'colorectal cancer' produced no additional papers. 26 prospective investigations involving 17 cohort studies (table 1C) and 2 RCTs with single interventions published between 1989 and 2007 were reviewed (table 2C). Studies using aggregate data, such as ecological studies, were not reviewed as the effects of an individual's dietary intake of fibre on their risk of colorectal cancer are difficult to determine using group level data on average intakes and disease rates.

2.2.4 Colonic function

The following criteria were used to help identify journal articles to be included in the review. Articles were included if:

- They referred to human studies and were in English
- They specified the amount and the type of dietary fibre employed
- The fibre content was measured using the method of the Association of Official Analytical Chemists (AOAC), the Englyst method, Southgate method (or Neutral Detergent Fibre (NDF) in the case of insoluble fibre sources)
- They reported faecal output in g/d, collected from at least a 24 hour collection of all stools passed.

Articles were excluded if:

- The faecal measure was something other than daily faecal output in g/d, including measures of 'stool' weight (generally obtained where only one stool was collected.)
- Fibre content was defined as crude fibre.

Search Terms/Strategy

A PubMed search was performed using the following terms:

- Fiber OR Fibre OR Resistant starch OR Fructo(-)oligosaccharide OR Polydextrose OR Inulin OR Lignin

AND

- Colon OR Bowel OR Faecal OR fecal

Additionally, a manual search was made using reference lists of original research articles and review articles to help identify any further relevant articles.

2.2.5 Prebiotics

The following criteria were used to help identify journal articles to be included in the review. Articles were included if:

They refer to candidate materials which fulfilled the prebiotic requirement criteria set out by (Gibson et al, 2004):

- resists gastric acidity, hydrolysis by mammalian enzymes and gastrointestinal absorption
- is fermented by intestinal microflora
- selectively stimulates the growth and/or activity of intestinal bacteria associated with health and well-being.
- They were reported in the English language
- They were in human subjects, including infants

Articles were excluded if:

- They did not fully meet the three prebiotic criteria

A PubMed search was performed using the following terms:

 Prebiotic OR Bifidobacteria OR Lactobacilli OR Inulin OR Fructo(-)oligosaccharide OR Galacto(-)oligosaccharide OR Lactulose

Excluded terms:

- Non-selective saccharolytic fermentation OR Proteolysis OR Upper gut absorption OR Clinical situations

Reference lists of recent books on prebiotics were also searched for articles fulfilling the above search criteria.

2.3 Presentation of data

The data were extracted from papers and tabulated in accordance with the SACN framework. Where the term fibre is used in the review it reflects the terminology used by the author of the paper and does not necessarily reflect SACNs opinion.

3. Results

Results for all sections are presented in the tables the appendices A-E.

3.1 Obesity and metabolic health

Studies investigating fibre intake and weight related outcomes, type II diabetes and metabolic risk factors are listed in Tables 1A-14A. Metabolic risk factors include measures of insulin sensitivity and glucose response.

Evidence from four of the five prospective cohort studies identified suggest that increasing amounts of total fibre (as determined by AOAC method) in the diet are associated with lower body weight and waist circumference (Table 1A). There is no evidence for an association of fibre intake with weight change in children.

Prospective studies investigating total fibre intake and the incidence of type 2 diabetes are detailed in Table 2A. In terms of total fibre, the evidence is inconsistent. For cereal fibre, nine of the ten studies observed that increasing intake significantly reduced the risk of type 2 diabetes, but overall, the evidence does not support a positive association for fibre from fruit, vegetables or legumes on the outcomes investigated. Two of three studies observed an association with insoluble fibre intakes, but no association with soluble fibre (Meyer *et al* 2000; Montonen *et al* 2003). Seven of the nine cross sectional studies (table 3A) reported an association between fibre intake and various measures of insulin sensitivity and glucose tolerance.

Intervention studies have investigated the relationship between fibre supplements and food sources of fibre on weight and related outcomes such as energy intake and excretion (Tables 4A-8A). Although different forms of fibre have been administered, in general those studies that have provided, fibre as part of *ad libitum* diet or in the context of healthy eating regimens have not shown conclusive evidence that fibre intake in general is related to weight related outcomes. However, supplementation appears to be efficacious in overweight and obese subjects when dietary advice is given. The balance of evidence suggests that fibre supplementation with certain fibres in sufficient amounts is more likely to be efficacious in assisting weight loss as an adjunct to a weight-reducing diet in overweight and obese subjects (Table 6A). It should be noted that, due to the variation in study duration, and fibre types and

amounts, it is difficult to draw conclusions about the overall role of fibre intake in relation to the more important outcome of long term weight loss. There is no evidence to suggest an effect of fibre supplementation on weight control in children (Table 9A); this is consistent with the evidence from observational studies.

Studies evaluating the effect of fibre supplementation on metabolic risk factors such as insulin sensitivity and glucose tolerance are listed in Tables 10A-13A. There is heterogeneity in the study designs, and fibre types and amounts, but overall, beneficial effects of supplementation have been reported when more sensitive measures, such as the euglycaemic hyperinsulinaemic clamp have been used, with fewer effects seen in those studies using fasting measures. This suggests that any effect size is likely to be small. Furthermore, improvements in metabolic risk factors have generally been observed in subjects at higher metabolic risk, with very little evidence to suggest that insulin sensitivity or glucose tolerance can be further improved in healthy subjects. A wide range of supplements has been used in these studies, with few direct comparisons between types of fibre, making it difficult to draw conclusions about health benefits of different components. These data suggest that there may be some beneficial effect of increasing total fibre/non starch polysaccharide intake in reducing metabolic disease risk, independent of weight.

In terms of insulin sensitivity and glucose tolerance, only a small number of studies have investigated the effects of resistant starch (Table 11A), isolated oligosaccharides (Table 12A), or polydextrose (Table 13 A). The four studies which have assessed the effect of resistant starch suggest that it could have a positive effect on insulin sensitivity and glucose tolerance; however, further research is required to confirm this relationship. There is insufficient evidence to show an association between oligosaccharides, inulin, or polydextrose on weight outcomes or metabolic profiles.

Table 14A lists acute experimental feeding studies, which have investigated immediate postprandial effects associated with the consumption of fibre and subsequent glucose and insulin responses. There is a trend towards reductions in post-prandial glucose and insulin levels with a number of different sources of fibre, especially resistant starch, oat bran and guar gum. Some studies have reported transient increases in satiety with specific fibres, but there is much less evidence that these fibres produce decreases in food intake at later meals. The limited number of studies, together with the diversity of study design and outcome measures makes it difficult to draw clear conclusions on the acute effects of different types of fibre and whether any effects may translate into long term health benefits.

3.2 Cardiovascular disease

Studies investigating fibre and cardiovascular disease risk are listed in Tables 1B-18B. The majority of studies have focussed on lipid outcomes (cholesterol and triglyceride levels) with only a few measuring the effect of fibre intake on blood pressure or disease endpoints.

Prospective studies investigating cardiovascular disease risk are detailed in Table 1B: of the six studies five measured disease end points. Two studies reported that total fibre (Jensen et al 2004; Pietinen et al 1996b) and cereal fibre intake (Pietinen et al 1996b; Wolk et al 1999) decreased the risk of cardiovascular disease. Four cross-sectional studies were identified that evaluated the effect of fibre consumption on lipid outcomes (Table 1B). Of these two observed that increasing total fibre intake was related to lower cholesterol levels (McKeown

et al 2002; Newby et al 2007). Newby et al (2007) also noted an association with cereal fibre. In terms of observational studies, the findings are inconsistent, thus making it difficult to draw any firm conclusions about fibre intake and cardiovascular disease.

RCT's which have investigated the effect of supplemention with various components on lipid profiles are listed in Tables 6B-16B. In terms of oat products: the 13 of the 16 studies demonstrated that consumption of soluble fibre had significant effects on lowering total- and LDL-C levels (Tables 5B and 6B). Psyllium intake was also associated with lower total or LDL-cholesterol levels in three out of five trials (Table 7B and 8B). Data from a meta-analysis of 67 RCT's looking at the outcome of different types of fibre on the levels of blood lipids, which included 17 on oat products, 17 on psyllium, 7 on pectin and 18 on guar gum was also considered (Brown et al 1999). Taken together with the studies presented in the tables, the evidence suggests that soluble fibre, in particular that from oats, psyllium, pectin and guar gum may be effective in lowering total cholesterol and LDL cholesterol.

Three of the intervention studies (Behall et al 2004a; Behall et al 2004b; Keenan et al 2007) which investigated barley products were identified (Table 10B) demonstrated that fibre from barley could be effective in lowering total and LDL cholesterol concentrations in hypercholesterolemic subjects.

The findings from trials investigating isolated polysaccharides, wheat, fibre supplement mixtures, legumes, resistant starch and oligosaccharides and inulin are inconsistent (Tables 11B-16B), therefore there is insufficient evidence to conclude whether these different components are associated with lipid outcomes. In terms of legumes, three out of the five studies (Cobiac et al 1990; Fruhbeck et al 1997; Pittaway et al 2006; Anderson et al 1990) suggested that fibre from this source may have a cholesterol lowering effect (Table 14B), However, due to limitations in study design, it is difficult to draw any firm conclusions.

Studies investigating the effect of fibre intake on blood pressure are listed in Tables 3B, 4B, 17B and 18B. There is insufficient evidence from the five observational (prospective and cross sectional) and four intervention studies to demonstrate that dietary fibre has any effect on blood pressure. These studies were considered in addition to a meta-analysis of 24 trials investigating fibre supplementation with blood pressure (Streppel et al 2005), which overall did not find any evidence for an association.

3.3 Colorectal cancer

Studies which have evaluated the effect of fibre intake and the risk of colorectal cancer and adenoma are listed in Tables 1C-7C. The majority of the data were derived from prospective cohort studies with only two RCT's reporting on the effect of fibre supplementation.

For colorectal cancer the evidence was variable, with eight out of 17 cohort studies finding no statistically significant association after multiple adjustment for confounding factors between AOAC determined dietary fibre and colorectal cancer (Table 1C). Of these, the EPIC Europe study was one of the largest studies, and utilised the techniques of increased dietary heterogeneity and calibration to reduce measurement error. This study showed a strong protective effect of dietary fibre.

Nine studies have investigated the link between fruit, vegetable and grain sources of fibre and colorectal cancer (Bingham et al 2003; Bingham et al 2005; Fuchs *et al* 1999; Giovannucci *et al* 1994; Lin *et al* 2005; Michels *et al* 2005; Nomura *et al* 2007; Schatzkin *et al* 2007; Terry *et al* 2001; Wakai *et al* 2007; Willett *et al.*, 1990) (Table 6C). and most of these reported no significant effect. Two of the studies which involved an increased intake of vegetable fibre, and one study investigating grain fibre, reported a reduced the risk of colorectal cancer. Of the two cohort studies which assessed the intake of soluble and insoluble fibre in relation to the risk of colorectal cancer (Table 7C), one found an association with insoluble fibre.

It should be noted that all but one of the cohort studies (Pietinen et al 1999) on dietary fibre and colorectal cancer measured total fibre, as determined by the AOAC method. This method does not allow separation into the different components, therefore the individual fibre components cannot be directly associated with risk of colorectal cancer. In addition, there are no reliable biomarkers of fibre intake, and it is possible that measurement error in dietary assessment leading to misclassification of exposure may have affected the results of the observational studies.

Table 3C details five cohort studies investigating the risk of colorectal adenoma with total fibre intake. Of these, two studies reported an association with adenoma occurrence (Giovanucci et al 1992; Peters et al 2003). Three studies have distinguished between fibre from fruit, vegetable and grain sources and risk of colorectal adenoma (Giovannucci et al 1992; Jacobs et al 2002; Platz et al 1997) (Table 4C). Two of these studies showed a decreased risk of adenoma with fruit fibre intake and only one study found an association with vegetable and grain fibre. Platz et al (1997) reported that soluble fibre intake was associated with a decreased risk of adenoma; however, this was the only study which quantified soluble and insoluble fibre in the diet (Table 5C).

Table 2C lists two RCTs (The Wheat Bran Fibre Trial (WBFT) and European Cancer Prevention Organisation Intervention Study (ECPOIS)) which evaluated fibre supplementation and colorectal adenoma reoccurrence (Alberts *et al* 2000; Bonithon-Kopp *et al* 2000). These both investigated wheat bran and psyllium supplements, respectively, and found no evidence of an effect on subsequent adenoma risk. Therefore there is insufficient evidence from trials that dietary fibre components can reduce the occurrence of colorectal adenoma.

The findings suggest that increased overall fibre intake may reduce cancer risk; however further evidence is required to confirm this observation. On balance, due to the paucity of data and inconsistent findings, there is not enough evidence to conclude whether specific forms of fibre intake have an association with the risk of colorectal cancer or adenoma.

3.4 Colonic function

Studies which have evaluated the effect of intake of different types of 'dietary fibre' and colonic function are listed in Tables 1D-4E.

There are very few observational studies where diet has been assessed accurately and faecal collections made for sufficient periods to evaluate the effect of dietary fibre on faecal output in free-living individuals, so that intake of 'dietary fibre' that may contribute to faecal output can be examined. These are listed in table 1D.Colonic function as determined by faecal weight has been demonstrated to relate to NSP intake (Davies et al 1986; Cummings et al 1992; Birkett et al 1997) but not intakes of starch or resistant starch (Birkett et al 1997).

Several small intervention studies have investigated the effect of 'fibre' components including fibre from grains and vegetables, isolated polysaccharides, resistant starch and oligosaccharides on faecal weight. Results are presented in tables 2D-4D.

The mean increase in daily faecal weight was greater for components such as wheat bran (5.4g/g) followed by fruit and vegetables (4.1g/g), gums such as psyllium (4g/g), soya products (2.5g/g) and pectin least of all (1.2g/g) (Cummings 2001). It should be noted that many of these studies were only on a small number of subjects and were insufficiently powered.

Fibre type source	or	Number of studies	Mean increase in daily faecal weight g/g Dietary Fibre fed (SEM)	Comments
Wheat		41	5.4 (0.7)	Mainly bran. Raw 7.2 g/g; cooked 4.9 g/g
Fruit vegetables	and	28	4.1 (0.7)	Carrot, cabbage, peas, apple, potato, banana, prunes, mixed sources
Gums mucilages	and	27	3.7 (0.5)	Psyllium/ispaghula 4.0 g/g (n=14); gum Arabic 4.0 g/g (n=9); tragacanth, sterculia, bassara, xylan, agar
Cellulose		7	3.5 (0.8)	Also carboxymethylcellulose, 4.9 g/g (n=3), methylcellulose 8.9 g/g (n=4)
Oats		4	3.4 (1.1)	Oat bran or oats
Corn		5	3.3 (0.3)	Corn meal or bran
Legumes		17	2.2 (0.3)	Soya products 2.5 g/g (n=11)
Pectin		11	1.2 (0.3)	Degree of methoxylation not important

Table II: Summary of studies on effect of dietary fibre on faecal weight (from Cummings,2001)

Intervention studies have investigated the effect of fibre from various grains including wheat, rye and barley. Only one study investigating wheat was identified (a wheat fibre supplement; Fibrotein) (Vuksan et al 1999) and this increased faecal bulk significantly compared to the negative control group ($P \le 0.01$). Based on the two studies (Grasen et al 2000;McIntosh et al 2003) in table 2D, rye would appear to have a significant effect on faecal weight and transit time. Three intervention studies investigating barley were identified (Bird et al 2008, Li et al 2003, Lupton et al 1993) and demonstrated that barley significantly increased faecal weight. A number of studies investigated various new fibre supplements made from plant sources (see table 2D iv) including sugar beet (Castaglia-Delavind et al 1998), potato (Cherbut et al 1997) and flax (Dahl et al 2005). Overall these plant sources have a significant effect on stool weight. Fours small intervention studies (Chen et al 2006; Chaplin et al 2000; Daly et al 1993 and Robinson et al 2001) investigated isolated polysaccharides in the form of powders and gels. Overall these did not provide evidence of stool bulking.

Seven intervention studies on resistant starch and colonic function were reviewed by, Cummings et al (2001). These described 11 treatments where the effects of resistant starch on colonic function were studied. Several of these investigated the same resistant starch product, namely Hylon VII, either RS_2 or RS_3 . Four studies (Jenkins et al 1998; Behall et al 2002; Muir et al 2004; Grubbens et al 2001) were considered in addition to those reported by Cummings et al. (see table 3D). Overall, resistant starch has a modest effect on faecal weight (Cummings et al 2001; Jenkins et al 1998; Behall et al 2002; Muir et al 2004; Grubbens et al 2001) and this could not be considered of a size which would make it appropriate for consideration as a dietary fibre according to this criterion

Reference	Source of RS	Number of subjects	Amount of RS fed (g)	Increase in daily faecal weight (g/g RS fed)
Cummings et al (1996)	Potato RS2	9	26.8	1.6
Cummings et al (1996)	Banana RS2	8	30.0	1.7
Cummings et al (1996)	Wheat RS2	9	17.4	2.5
Cummings et al (1996)	Maize RS2	8	19.0	2.7
Heijnen et al (1998)	Hylon VIIRS2	23	32	1.4
Heijnen et al (1998)	Hylon VIIRS3	23	32	2.2
Hylla et al (1998)	Hylon VII	12	55	0.8
Silvester et al (1997)	Mixed potato RS2 and Hylon VII	8	40	0.9
Phillips et al (1995)	Mixed food sources	11	39	1.8
Tomlin and Read (1990)	Cornflakes	8	10	-1.9
Van Munster et al (1994)	Hylon VII RS2	14	28	1.0

Fable I: Effects of resistant starch on faecal	weight	(Cummings 20	01)
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Ten studies on oligosaccharides and inulin are summarised in table 4D. Overall oligosaccharides (Alles et al 1999; Bouhnik et al 2007; Brighenti et al 1999;; Chen et al 2001;Molis et al 1996; Ten Bruggencate 2006) and inulin (Causey et al 2000; Den Hond et al 2000; Sairanen et al 2007) had very little effect on faecal weight or transit time when these materials are fed and hence an effect on faecal weight is not a criterion that is satisfied by these materials to enable them to be considered dietary fibre.

In addition, several studies reported no significant change in SCFA concentration (Alles et a 1999; Causey et al 2000) and two studies (Chen et al 2001; Scholtens et al 2006) reported a decrease in butyrate concentration. Four studies investigated diets with mixed fibre sources from fruit, vegetables and/or grains. All four studies reported an increase in faecal weight when fibre was increased in the diet.

3.5 Prebiotics

Studies which have evaluated the effect of prebiotics on the gut microflora in adults and infants are detailed in tables 1E and 2E. Studies investigating prebiotics were of short durations (often days or weeks) and on a very small number of human subjects. Overall studies demonstrate that lactulose, fructo-oligosaccharides and galacto-oligosaccharides significantly increased bifidobacteria but as yet there is no convincing evidence that as potential fibre components they confer any specific health benefit to the human host.

4. Interpretation of results

In formulating a recommendation SACN considered the strength of the available evidence from both intervention and observational studies. Consideration was given to the sample size, dose, duration and significance of the studies presented.

There were concerns regarding the quality of some studies presented in the report and where there appear to be conflicting results in outcomes, this may be accounted for by differences in study quality. Furthermore, studies have often poorly characterised the fibre components under investigation as in the use of AOAC method to determine fibre or the use of generic terms such as soluble or insoluble fibre. Therefore, making it difficult to distinguish which specific components could be contributing to any associations observed or readily compare the findings of different studies.

In addition consideration was given to whether the studies demonstrated a clear physiological effect on human health.

The data presented in the appended tables will contribute to the cardio-metabolic health and colorectal health sections of the SACN Carbohydrate and Health report once the committee have finished their deliberation.

5. SACN recommendation

SACN consider that a material can be considered as dietary fibre if it is resistant to digestion and absorption in the small intestine and has a demonstrable physiological effect potentially associated with health benefits in the body, such as increasing stool bulk, decreasing intestinal transit time or decreasing post prandial glycaemia. Evidence only of increased fermentation in the gut should not be included under this definition, since although this has a direct effect on the microflora, it must also be shown to have a demonstrable benefit to the host to be considered as dietary fibre.

6.0 Carbohydrate definitions

Monosaccharides- single sugar molecules e.g. glucose, fructose galactose.

Disaccharides- consist of two monosaccharides linked together e.g. sucrose, lactose, maltose, trehalose.

Oligosaccharides- short chain carbohydrates comprising of 3-9 monomers joined together by glycosidic linkages.

Polysaccharides- long chain carbohydrates comprising of ≥ 10 monomers.

Sugars- monosaccharides, disaccharides and polyols (sugar alcohols).

Alpha glucans- oligosaccharides with an alpha linkage between monomers.

Inulin, Fructooligosaccharides (FOS)- non- α -glucan oligosaccharides, known as fructans, and are the storage component of artichokes and chicory.

Polydextrose -non- α–glucan oligosaccharide.

Galactooligosaccharides (GOS)/ milk oligosaccharides- oligosaccharides found in milk, which principally contain galactose.

Non starch polysaccharide (NSP)- non- α -glucan polysaccharides that are mainly found in the plant cell walls and consist of a large umber of monosaccharides. This includes cellulose, hemicellulose, pectin, arabinoxylans, plant gums, β -glucans.

Pectin- an NSP which is common to all cell walls.

Gums/mucilages- an NSP which is which is chemically related to the cell wall, but is not strictly a cell wall component. Plant gums are sticky exudates which are formed at the site of injury e.g. guar gum.

Starch- the storage carbohydrate of plants, such as cereals, root vegetables and legumes, and consists of only glucose molecules.

Resistant starch- the sum of starch and products of starch digestion (such as maltose, maltotriose and α -limit dextrins) that are not absorbed by the small bowel.

Prebiotic- a non-digestible food component that stimulates the growth and/ or activity of the bacteria in the bowel.

Lignin- non-carbohydrate component associated with plant walls.

Englyst method- specifically determines NSP using an enzymatic-chemical method, which can be modified to yield soluble and insoluble fractions.

American Association of Analytical Chemists (AOAC)/ enzymatic-gravimetric method-

Determines total, soluble and insoluble residue containing carbohydrate and non-carbohydrate material in unknown proportions by measuring total residue weight and subtracting ash and protein content.

Soluble fibre- relates to NSP components which can be rendered soluble by changing the pH conditions. These generally undergo significant fermentation, and viscous forms of these may also slow rates of glucose and lipid absorption from the small intestine^A. Examples of soluble fibres include pectin, beta-glucan (from oats and barley) and psyllium.

Insoluble fibre- NSP components that tend to undergo slow and incomplete fermentation and can have a greater effect on bowel habit^A. Insoluble fibres are found in vegetables and wholegrain products.

Available and unavailable carbohydrate- relates to carbohydrates which are available to the body for metabolism (starch/soluble sugars), or those which are not (cellulose and hemicellulose) and pass to the colon. It should be noted that carbohydrate reaching the colon is still able to provide energy through fermentation and, therefore the definition is not exact.

Glycaemic carbohydrates- carbohydrates which provide glucose following digestion and absorption in the small intestine e.g. mono- and disaccharides, some oligosaccharides (maltodextrins), rapid and slowly digested starch.

Non-glycaemic carbohydrates- carbohydrates which pass to the large intestine e.g. other oligosaccharides, resistant starch, NSP.

^AThe division between soluble and insoluble fibre is extremely pH dependent. Also, a large proportion of insoluble fibres are completely fermented and not all soluble fibre have effects on glucose and lipid absorption. Therefore, WHO have considered these definitions to be less useful when characterising fibre components.

Appendix A: Obesity and metabolic Health

Table 1 A: Prospective studies of fibre intake and body weight/composition

Table 2A: Prospective studies of fibre intake and type 2 diabetes incidence

Table 3A: Cross sectional studies of fibre intake and metabolic risk factors

Table 4A: Highly-controlled intervention studies of fibre intake and body weight/weight-related outcomes

Table 5A: Effect of fibre on body weight in 'healthy-eating' dietary interventions

Table 6A: Effect of fibre supplementation of habitual diets on body

 Table 7A: Effect of fibre supplementation of weight-reducing diets on body

Table 8A: Effect of fibre supplementation in maintenance of weight-loss

Table 9A: Effect of fibre supplementation on body weight in children

Table 10A: Effect of fibre supplementation on metabolic risk factors

Table 11A: Effect of resistant starch on weight and metabolic outcomes

Table 12A: Effect of oligosaccharides and inulin on weight and metabolic outcomes

Table 13A: Effect of polydextrose on weight and metabolic outcomes

Table 14A: Acute experimental studies

Reference Cohort	Subjects N (M/F) Age Follow-up	Determination of fibre	Fibre intake	Adjusted result/OR/RR	P/Ptrend ^A	Factors adjusted for
Iqbal et al (2006) Danish 1936 cohort (n=284) & MONICA1 study	1,762 (862/900) 30-60 years	AOAC-defined total dietary fibre by 7 day weighed record (baseline)	Mean ± SD (g/d)	Mean change in body weight (kg) ($\beta \pm SE$) per 1 kg/d increase in fibre intake		Baseline BMI, age, physical activity, smoking, education, cohort, dietary volume, energy intake
(n=1478)	5 years follow- up		M: 18.8 ± 7.3 F: 13.9 ± 5.5	8.6 ± 6.9 -22.3 ± 13.4	0.17 0.10	
Koh-Banerjee et al (2003) Health Professionals Follow-up Study	16,587 (M) 40-75 years 9 year follow- up	AOAC-defined total dietary fibre by FFQ	Baseline / follow-up mean intake \pm SD (g/d): 40-49y: 18.8 \pm 7.3 / 21.3 \pm 7.1 50-59y: 21.1 \pm 6.7 / 22.7 \pm 7.3 60-75y: 22.3 \pm 7.4 / 24.2 \pm 7.6	Mean waist change ± SE (cm) with 12 g/d increase in total fibre intake: -0.23 ± 0.09	0.008	Age, baseline waist, baseline and change in BMI, baseline and change in energy intake, change in smoking status, baseline and change in physical activity, baseline and change in alcohol intake
Koh-Banerjee et al (2004) Health Professionals Follow-up Study	27,082 (M) 40-75 years 8 year follow- up	AOAC-defined total dietary fibre by FFQ	Tertile median change in intake (g/d): Total fibre Q1: -5.2 Q2: +0.5 Q3: +8.5 Cereal fibre Q1: -2.2 Q2: +1.0 Q3: +5.1 Fruit fibre Q1: -2.2 Q2: +0.2 Q3: +3.7 Vegetable fibre Q1: -3.2 Q2: 0.0 Q3: +4.1	Mean weight change \pm SE (kg) 1.40 \pm 0.20 1.04 \pm 0.20 0.39 \pm 0.20 1.30 \pm 0.27 1.15 \pm 0.26 0.91 \pm 0.26 1.59 \pm 0.27 0.96 \pm 0.26 0.64 \pm 0.26 1.08 \pm 0.27 1.26 \pm 0.26 1.12 \pm 0.26	<0.0001 0.0004 <0.0001 0.8	Age, baseline weight, baseline smoking status, baseline and change in other dietary factors (including other fibre types), baseline and change in physical activity

Table 1A: Prospective studies of fibre intake and body weight/composition

^A When data is reported across tertiles, quartiles or quintiles (up to Q3, Q4, or Q5) the values represent Ptrend BMI- body mass index; FFQ Food frequency questionnaire

Reference Cohort	Subjects N (M/F) Age Follow-up	Determination of fibre	Fibre intake	Adjusted result/OR/RR	P/Ptrend ^A	Factors adjusted for
Lindstrom et al (2006a) Finnish Diabetes Prevention Study Cohort	500 40-64 years 3 year follow- up from 3 year active intervention period	AOAC-defined total dietary fibre by 3 day food record (mean of intervention years 1-3)	Quartile range (g/4.18MJ) Q1: <10.85 Q2: 10.85-13.00 Q3: 13.00-15.55 Q4: >15.55	Mean weight change year 1 to 3 (kg) (LOCF) -0.4 -1.6 -2.5 -3.0	0.001	Intervention group, gender, age, VLCD-use, baseline weight, baseline & follow-up physical activity, baseline fibre intake (NOT fat intake)
	penod		Quartile range (g/4.18MJ) Q1: <10.85 Q2: 10.85-13.00 Q3: 13.00-15.55 Q4: >15.55	Mean waist change year 1 to 3 (cm) (LOCF) -1.6 -2.2 -2.5 -2.9	0.033	
Liu et al (2003) Nurses Health Study	74,091 (F) 38-63 years 2-4 years follow-up	AOAC-defined total dietary fibre by FFQ	Quintile Q1 Q2 Q3 Q4 Q5	Mean ± SE weight change (kg) 1.73 ± 0.03 1.50 ± 0.03 1.37 ± 0.02 1.34 ± 0.02 0.97 ± 0.02	<0.0001	Age, baseline BMI; changes in exercise, smoking, HRT, alcohol, caffeine, total energy, SFA, PUFA, MUFA, transFA and protein
Ludwig et al (1999) CARDIA Study	1,302 to 1,602 depending on outcome 18-30 years 10 year follow- up	AOAC-defined total dietary fibre by FFQ	Quintile median (g/4.18MJ) White (n=1,602) Q1: 5.2 Q5: 12.3 Black (n=1,307) Q1: 5.2 Q5: 12.3	Mean year 10 body weight (kg) 79.2 75.6 84.2 80.6	<0.001 0.001	Baseline weight, gender, age, field centre, education, energy intake, physical activity, smoking, alcohol, vitamin supplement use
			Quintile median (g/4.18MJ) White (n=1,598) Q1: 5.2 Q5: 12.3 African-American (n=1,302) Q1: 5.2 Q5: 12.3	Mean year 10 waist- to-hip ratio 0.813 0.801 0.809 0.799	0.004	

^A When data is reported across tertiles, quartiles or quintiles (up to Q3, Q4, or Q5) the values represent Ptrend BMI- body mass index; FFQ Food frequency questionnaire; SFA- saturated fatty acids; PUFA- polyunsaturated fatty acids; transFA- trans fatty acids

STUDIES IN CH	STUDIES IN CHILDREN							
Reference Cohort	Subjects N (M/F) Age Follow-up	Determination of fibre	Fibre intake	Adjusted result/OR/RR	P	Factors adjusted for		
Berkey et al (2000) Growing Up Today Study	10,769 (4,620/6,149) 9-14 years 1 year follow- up	AOAC-defined total dietary fibre by FFQ	Baseline mean age-specific intake ± SD (g/d)	Mean annual change in BMI ($\beta \pm SE$) per 1 g/d increase in fibre intake in year before baseline BMI		Race, menarche history, annual height growth, baseline BMI, age, Tanner stage, energy & fat intake, Physical activity measures		
			M9y: 16.9 ± 4.2 M14y: 17.8 ± 4.4	-0.0059 ± 0.0045	0.186			
			F9y: 16.4 ± 4.1 F14y: 16.8± 4.6	0.0023 ± 0.00415	0.577			
Newby et al (2003) North Dakota WIC Program	1,379 (690/689) 2-5 years 1 year follow up	AOAC-defined total dietary fibre by FFQ	Baseline mean intake ± SD (g/d) M: 11.9 ± 3.0 F: 11.9 ± 3.1	Mean annual change in weight (kg) ($\beta \pm$ SE) per 1 g/d increase in baseline fibre intake 0.001 ± 0.02	0.96	Age, gender, energy intake, ethnicity, residence, poverty level, maternal education, birth weight		

FFQ Food frequency questionnaire, BMI- body mass index

Reference Cohort	Subjects N (M/F) Age Follow-up	Determination of fibre	Fibre intake	Adjusted result/OR/RR (95%CI)	P/Ptrend ^A	Factors adjusted for
Colditz et al (1992)	84,360 (F)	Southgate-defined total dietary fibre	Energy-adjusted total fibre intake			Age, BMI, alcohol, family history of diabetes, prior weight change & time
Nulses health Study	702 00303	bying	BMI<29			period
	30-55 years		Q1 Q2	1.00 (ref) 0.89	0.60	
	6 years follow-		Q3	0.82		
	up		Q4	1.19		
			BMI≥29	0.75 (0.50-1.15)		
			Q1	1.00 (ref)	0.97	
			Q2	1.21		
			04	1.21		
			Q5	1.08 (0.78-1.48)		
Hodge et al (2004)	31,641	AOAC-defined total dietary fibre by	Increase in intake (g/d)			Age, country of birth, physical activity, 5-year
Melbourne	40-69 years	FFQ	Total fibre: +20	1.02 (0.81-1.30)	0.46	weight change, education, family history of diabetes
Study	4 years follow-		Fruit fibre: +10	0.97 (0.81-1.16)	0.96	energy intake, BMI,
	up		Vegetable fibre: +5	1.00 (0.86-1.17)	0.67	waist:hip ratio
			Legume fibre: +1	1.01 (0.96-1.06)	0.65	
				1.03 (0.91-1.16)		
Hu et al (2001)	84,941 (F)	AOAC-defined total dietary fibre by	Cereal fibre intake			Age, time, family history of diabetes, menopausal
Nurses Health Study	3,300 cases	FFQ	Q1 Q5	1.00 (ref) 0.6 (0.5-0.7)	<0.001	status, HRT, smoking, BMI, physical activity, alcohol,
	34-59 years			(estimated from		PUFA:SFA, transFA, glycaemic load
	16 years follow-up			figure)		

Table 2A: Prospective studies of fibre intake and type 2 diabetes incidence

^A When data is reported across tertiles, quartiles or quintiles (up to Q3, Q4, or Q5) the values represent Ptrend

BMI- body mass index; FFQ Food frequency questionnaire; SFA- saturated fatty acids; PUFA- polyunsaturated fatty acids; transFA- trans fatty acids

Reference Cohort	Subjects N (M/F) Age Follow-up	Determination of fibre	Fibre intake	Adjusted result/OR/RR (95%CI)	P/Ptrend ^A	Factors adjusted for
Krishnan et al (2007)	40,078 (F) 1,938 cases	AOAC-defined total dietary fibre by FFQ	Quintile median energy- adjusted cereal fibre intake (g/d)			Age, BMI, energy intake, family history diabetes, physical activity, smoking,
Health Study	21-69 years 8 years follow- up		Q1: 1.7 Q2: 2.7 Q3: 3.7 Q4: 4.9 Q5: 7.6	1.00 (ref) 0.91 (0.78-1.05) 0.89 (0.76-1.04) 0.83 (0.70-0.96) 0.82 (0.70-0.96)	0.01	total fat intake
Lindstrom et al (2006a) Finnish Diabetes Prevention Study Cohort	500 114 cases 40-64 years 4 years follow- up from 3 year active intervention period)	AOAC-defined total dietary fibre by 3 day record (mean of intervention years 1-3)	Quartile range energy- adjusted fibre intake (g/d) Q1: <10.85 Q2: 10.85-13 Q3: 13-15.55 Q4: >15.55	1.00 (ref) 0.50 (0.28-0.89) 0.71 (0.40-1.23) 0.38 (0.19-0.77)	<0.05 ^B	Intervention group, gender, age, VLCD-use, baseline weight, baseline & follow-up physical activity, baseline fibre intake, weight change
Lindstrom et al (2006b) Finnish Diabetes Prevention Study Cohort	406 69 cases 40-64 years 3 years follow- up from 3 year active intervention period	AOAC-defined total dietary fibre by 3 day record (mean of intervention years 1-3)	Energy-adjusted fibre intake (g/d) <15 by year 3 ≥15 by year 3	1.00 (ref) 0.97 (0.63-1.51)	>0.05	Also meeting fat, SFA & physical activity goals

^A When data is reported across tertiles, quartiles or quintiles (up to Q3, Q4, or Q5) the values represent Ptrend
 ^B Ptrend for this maximally adjusted model was not stated, however p-trend for a similar model was 0.04. The authors stated that the result was not affected by the additional adjustment.

BMI- body mass index; FFQ Food frequency questionnaire; SFA- saturated fatty acids; VLCD

Reference Cohort	Subjects N (M/F) Age	Determination of fibre	Fibre intake	Adjusted result/OR/RR	Ptrend	Factors adjusted for
Montonen et al (2003) Finnish Mobile Clinic Health Examination Survey	4,316 (2,286/2,030) 156 cases 40-69 years	AOAC-defined total dietary fibre by diet history interview	Quartile range intake (g/d) Total fibre Q1: 2.6-19.2 Q2: 19.3-25.3 Q3: 25.4-33.1 Q4: 33 2-118	1.00 (ref) 0.70 (0.46-1.07) 0.67 (0.40-1.11) 0.51 (0.26-1.00)	0.04	Age, gender, geographic area, energy intake, fruit, berry & vegetable intake
	10 years follow-up		Soluble fibre Q1: 0.53-4.5 Q2: 4.6-5.8 Q3: 5.9-7.3 Q4: 7.4-22.7	1.00 (ref) 0.50 (0.31-0.81) 0.74 (0.44-1.25) 0.57 (0.29-1.12)	0.21	
			Insoluble fibre Q1: 1.1-8.7 Q2: 8.8-12.0 Q3: 12.1-16.5 Q4: 16.6-69.3	1.00 (ref) 0.75 (0.50-1.12) 0.72 (0.45-1.17) 0.47 (0.25-0.91)	0.03	
			Cellulose Q1: 0.48-3.2 Q2: 3.3-4.2 Q3: 4.3-5.3 Q4: 5.4-15.2	1.00 (ref) 0.53 (0.32-0.85) 0.67 (0.39-1.14) 0.60 (0.29-1.21)	0.19	
			Lignin Q1: 0.48-2.3 Q2: 2.4-3.1 Q3: 3.2-4.1 Q4: 4.2-14.5	1.00 (ref) 0.79 (0.52-1.20) 0.69 (0.42-1.15) 0.68 (0.36-1.30)	0.16	
			Cereal fibre Q1: 0.47-12.0 Q2: 12.1-17.3 Q3: 17.4-24.4 Q4: 24.5-111	1.00 (ref) 0.81 (0.54-1.21) 0.74 (0.46-1.18) 0.39 (0.20-0.77)	0.01	
			Fruit fibre Q1: 0-0.99 Q2: 1.0-2.0 Q3: 2.1-3.3 Q4: 3.4-36.8	1.00 (ref) 0.68 (0.39-1.20) 0.79 (0.39-1.60) 0.92 (0.40-2.13)	0.87	
			Vegetable fibre Q1: 0.11-3.7 Q2: 3.8-5.0 Q3: 5.1-6.7 Q4: 6.8-26.5	1.00 (ref) 1.02 (0.58-1.79) 0.89 (0.43-1.85) 1.19 (0.46-3.04)	0.86	

Reference Cohort	Subjects N (M/F) Age Follow-up	Determination of fibre	Fibre intake	Adjusted result/OR/RR (95%CI)	Ptrend	Factors adjusted for
Salmeron et al (1997a)	65,173 (F) 915 cases	AOAC-defined total dietary fibre by FFQ	Quintile median energy- adjusted fibre intake (g/d)			Energy, age, BMI, alcohol, smoking, physical activity, family history of diabetes
Nurses' Health		_	Total fibre			
Study	40-65 years		Q1: 11.8 O2: 14.7	1.00 (ref) 1.01 (0.83-1.24)	0.02	
	6 year follow-		03: 17.0	0.90(0.73-1.11)		
	up		Q4: 19.6	0.91 (0.74-1.13)		
			Q5: 24.1	0.78 (0.62-0.98)		
			Cereal fibre			
			Q1: 2.0	1.00 (ref)	0.001	
			Q2: 2.9	1.01 (0.83-1.23)		
			Q3: 3.7	0.85 (0.69-1.04)		
			Q4: 4.9	0.82 (0.66-1.01)		
			Q5: 7.5	0.72 (0.58-0.90		
			Fruit fibre			
			Q1: 1.4	1.00 (ref)	0.39	
			Q2: 2.6	0.87 (0.70-1.07)		
			Q3: 3.7	0.95 (0.77-1.18)		
			Q4: 5.1	0.94 (0.76-1.16)		
			Q5: 7.6	0.87 (0.70-1.08)		
			Vegetable fibre			
			Q1: 3.4	1.00 (ref)	0.54	
			Q2: 4.8	1.40 (1.13-1.73)		
			Q3: 5.9	1.23 (0.99-1.53)		
			Q4: 7.2	1.29 (1.04-1.61)		
			Q5: 9.6	1.17 (0.93-1.46)		
1						

BMI- body mass index; FFQ Food frequency questionnaire

Reference Cohort	Subjects N (M/F) Age	Determination of fibre	Fibre intake	Adjusted result/OR/RR	P/Ptrend	Factors adjusted for
	Follow-up			(95%CI)		
Salmeron et al	42,759 (M)	AOAC-defined total	Quintile median energy-			Energy, age, BMI, alcohol,
(1997b)		dietary fibre by	adjusted fibre intake (g/d)			smoking, physical activity,
	523 cases	FFQ				family history of diabetes
Health Professionals'			Total fibre			
Follow-up Study	40-75 years		Q1: 13.4	1.00 (ref)	0.70	
			Q2: 17.1	0.98 (0.75-1.29)		
	6 year follow-		Q3: 20.0	1.08 (0.83-1.42)		
	up		Q4: 23.5	0.87 (0.65-1.17)		
			Q5: 29.7	0.98 (0.73-1.33)		
			Cereal fibre			
			$01^{\circ} 25$	1 00 (ref)	0.007	
			02: 3.8	1.14 (0.89-1.46)	0.007	
			03: 5.0	0.95 (0.73-1.25)		
			04: 6.8	0.91 (0.69-1.20)		
			Q5: 10.2	0.70 (0.51-0.96		
			Fruit fibre			
			Q1: 1.2	1.00 (ref)	0.68	
			Q2: 2.8	1.01 (0.76-1.34)		
			Q3: 3.8	0.89 (0.67-1.19)		
			Q4: 5.3	1.14 (0.86-1.51)		
			Q5: 8.3	1.01 (0.76-1.36)		
			Vegetable fibre			
			Q1: 3.5	1.00 (ref)	0.65	
			Q2: 4.9	1.12 (0.85-1.49)		
			Q3: 6.3	1.22 (0.93-1.61)		
			Q4: 7.9	1.10 (0.83-1.46)		
			Q5: 11.3	1.12 (0.84-1.49)		
1						

BMI- body mass index; FFQ Food frequency questionnaire

Reference Cohort	Subjects N (M/F) Age Follow-up	Determination of fibre	Fibre intake	Adjusted result/OR/RR (95%CI)	P/Ptrend	Factors adjusted for
Schulze et al (2004)	91,249 (F)	AOAC-defined total	Quintile median energy-			Age, BMI, energy intake,
		dietary fibre by	adjusted fibre intake (g/d)			alcohol, family history of
Nurses Health Study	741 cases	FFQ	Tatal fibra			diabetes, physical activity,
11	24-44 years			1.00 (rof)	0.80	smoking, HRT, oral
			$02^{\circ} 15.4$	0.94 (0.76-1.17)	0.00	high blood pressure or
	8 years follow-		03: 17.7	0.87 (0.68-1.11)		cholesterol, glyycaemic
	up		Q4: 20.2	0.84 (0.65-1.10)		load, magnesium, caffeine;
			Q5: 24.9	1.00 (0.75-1.34)		sub-types also adjusted for total fibre.
			Cereal fibre			
			Q1: 3.1	1.00 (ref)	0.004	
			Q2: 4.2	0.85 (0.69-1.05)		
			Q3: 5.2	0.87 (0.69-1.08)		
			Q4: 6.4	0.82 (0.65-1.04)		
			Q5: 8.8	0.64 (0.48-0.86)		
			Fruit fibre			
			Q1: 1.1	1.00 (ref)	0.04	
			Q2: 2.0	0.93 (0.75-1.15)		
			Q3: 2.9	0.80 (0.63-1.00)		
			Q4: 4.1	0.77 (0.60-0.98)		
			Q5: 6.2	0.79 (0.60-1.02)		
			Vegetable fibre			
			01.34	1 00 (ref)	0 192	
			02: 4.8	0.97(0.77-1.22)	01152	
			Q3: 6.1	1.01 (0.80-1.28)		
			Q4: 7.6	1.19 (0.94-1.51)		
			Q5: 10.4	1.12 (0.87-1.46)		

BMI- body mass index; FFQ Food frequency questionnaire

Reference	Subjects	Determination of	Fibre intake	Adjusted	Ptrend	Factors adjusted for
Cohort	N (M/F) Age Follow-up	fibre		result/OR/RR (95%CI)		
Schulze et al (2007)	25,067	AOAC-defined total	Quintile median energy-			Energy, age, sex,
EDIC Detedam Study	(9,702/1,536)	dietary fibre by	adjusted fibre intake (g/d)			education, sports activity,
EPIC-POISUAIII Sludy	844 cases	FFQ	Total fibre			activity, smoking, alcohol,
			Q1: 15.8	1.00 (ref)	0.11	BMI, waist, Mg, PUFA:SFA,
	35-65 years		Q2: 18.8	1.10 (0.88-1.36)		MUFA:SFA, CHO, other
	9 years follow-		Q4: 23.4	0.96 (0.75-1.22)		types of fibre
	up		Q5: 27.9	0.86 (0.65-1.14)		
			Soluble fibre			
			Q1: 5.3	1.00 (ref)	0.45	
			Q2: 6.4	0.85 (0.66-1.08)		
			Q3: 7.2 Q4: 8 1	0.99 (0.75-1.30)		
			Q5: 9.6	0.83 (0.57-1.22		
			Incoluble fibro			
			01: 10.3	1.00 (ref)	0.62	
			Q2: 12.3	1.11 (0.87-1.42)		
			Q3: 13.7	0.94 (0.70-1.26)		
			Q4: 15.4	0.99 (0.71-1.38)		
			Q3. 10. 1	0.00 (0.02 11.10)		
			Cereal fibre	1.00 (0.00	
			Q1: 6.6 Q2: 9.0	1.00 (ref) 0.86 (0.70-1.06)	0.02	
			Q3: 10.8	0.94 (0.76-1.16)		
			Q4: 12.8	0.85 (0.68-1.06)		
			Q5: 16.6	0.72 (0.56-0.93)		
			Fruit fibre		0.22	
			Q1: 0.2	1.00 (ref)		
			Q2: 1.0 Q3: 1.5	1 01 (0 81-1 26)		
			Q4: 2.8	0.93 (0.74-1.16)		
			Q5: 4.7	0.89 (0.70-1.13)		
			Vegetable fibre		0.66	
			Q1: 0.7	1.00 (ref)		
			Q2: 1.3	0.87 (0.70-1.08)		
			Q3. 1.7 Q4: 2.3	0.86 (0.68-1.07)		
			Q5: 3.4	0.93 (0.74-1.17)		

FFQ Food frequency questionnaire; BMI- body mass index; SFA- saturated fatty acids; PUFA- polyunsaturated fatty acids, MUFA- monounsaturated fatty acids, CHO- carbohydrate

Reference Cohort	Subjects N (M/F) Age	Determination of fibre	Fibre intake	Adjusted result/OR/RR	Р	Factors adjusted for
	Follow-up			(95%CI)		
Stevens et al (2002)	12,251	AOAC-defined total	Quintile median energy-			Age, BMI, gender, field
Atherosclerosis Risk	1 447 cases	FFO	aujustea fibre filtake (g/a)			smoking physical activity
in Communities	1,11, 60565		Total fibre			Sinoking, physical activity
(ARIC) Study	9 years follow-		White			
	up		Q1: 11.2	1.00 (ref)	0.915	
			Q5: 27.5	0.999 (0.987-1.012)		
			African-American			
			Q1: 10.2	1.00 (ref)	0.849	
			Q5: 26.1	0.998 (0.980-1.017)		
			Coroal fibro			
			White			
			01	1.00 (ref)	0.006	
			05	0.956 (0.925-0.987)	01000	
			African-American	, , , , , , , , , , , , , , , , , , ,		
			Q1	1.00 (ref)	0.525	
			Q5	0.982 (0.927-1.039)		
			Fruit fibre			
			01	1.00 (rof)	0.841	
			05	1.00(10) 1.002(0.983-1.021)	0.041	
			African-American	1.002 (0.903 1.021)		
			01	1.00 (ref)	0.479	
			Q5	1.009 (0.985-1.033)		
			Legume fibre			
			White	1.00 (0 774	
				1.00 (ret)	0.774	
			QJ African-American	1.001 (0.928-1.028)		
				1.00 (ref)	0 366	
			05	0.961(0.882-1.047)	0.000	

FFQ Food frequency questionnaire; BMI- body mass index

GESTATIONAL DIABETES							
Reference Cohort	Subjects N (M/F) Age Follow-up	Determination of fibre	Fibre intake	Adjusted result/OR/RR (95%CI)	Ptrend	Factors adjusted for	
Zhang et al (2006)	13,110 (F)	AOAC-defined total dietary fibre by	Quintile median energy- adjusted fibre intake (g/d)			Age, parity, BMI, ethnicity, smoking, family history of	
II	8 years follow- up		Total fibre Q1: 12.4 Q2: 15.4 Q3: 17.5 Q4: 20.1 Q5: 24.8	1.00 (ref) 0.97 (0.78-1.20) 1.00 (0.80-1.24) 0.84 (0.66-1.07) 0.67 (0.51-0.90)	0.005	activity, total E, other nutrients, separate fibre sources	
			Cereal fibre Q1: 2.9 Q2: 4.1 Q3: 5.1 Q4: 6.3 Q5: 8.9	1.00 (ref) 1.04 (0.84-1.30) 0.83 (0.65-1.04) 0.89 (0.71-1.15) 0.76 (0.59-0.98)	0.02		
			Fruit fibre Q1: 1.1 Q2: 2.0 Q3: 2.9 Q4: 4.1 Q5: 6.2	1.00 (ref) 0.75 (0.60-0.94) 0.86 (0.69-1.07) 0.82 (0.65-1.04) 0.67 (0.51-0.87)	0.02		
			Vegetable fibre Q1: 3.3 Q2: 4.8 Q3: 6.1 Q4: 7.7 Q5: 10.6	1.00 (ref) 1.06 (0.86-1.32) 1.04 (0.83-1.29) 1.01 (0.80-1.28) 0.87 (0.67-1.13)	0.24		

FFQ Food frequency questionnaire; BMI- body mass index

Reference Cohort	Subjects N (M/F) Age Follow-up	Determination of fibre	Fibre intake	Adjusted result/OR/RR (95%CI)	P/Ptrend ^A	Factors adjusted for
Boeing et al (2000)	1,773 (745/1,028)	AOAC-defined total dietary fibre by		Odds ratio (95%CI) Q1-5 total fibre		Age, sex, obesity, physical activity level, education,
	35-64 years	Mean ± SD (g/d) M: 27.1 ± 9.4 F: 24.2 ± 7.7	High (3 rd tertile) HbA1c	1.00 (ref) 1.05 (0.72-1.55) 0.83 (0.57-1.23) 0.97 (0.66-1.43) 0.93 (0.62-1.37)	0.595	SHOKING
Feskens et al (1994)	389 (M)	AOAC-defined total		Correlation coefficient		Age, BMI, physical activity,
Zutphen Elderly Study	70-89 years	diet history interview (previous	OGTT AUC Insulin	-0.12	<0.05	of coronary heart disease
,		2-4 weeks)	Fasting C-peptide	-0.11	<0.05	
		Mean ± SD (g/d)	HOMA-IR	-0.15 (unadjusted)	<0.01	
		25.4 ± 7.3	β-cell mass	-0.14 (unadjusted)	<0.01	
Lau et al (2005)	5,675	AOAC-defined total dietary fibre by		HOMA-IR Ratio (95%CI)		Age, gender, smoking, physical activity, energy
Danish Inter99 Study	30-60 years	FFQ	Change in HOMA-IR with 10g/d increase in intake	0.97 (0.96-0.99)	0.001	intake, BMI, waist circumference
Liese et al (2005)	979	AOAC-defined total		β ± SE		Age, gender,
Insulin Resistance Atherosclerosis	(442/337) 33% IGT	FFQ	FSIVGTT SI	0.1250 ± 0.0306	<0.001	history of diabetes, current
Study	67% NGT	Mean ± SD (g/d)	Acute insulin response	0.0317 ± 0.0393	0.420	expenditure, energy intake
	40-69 years	16.9 ± 7.9	Glucose disposal	0.0933 ± 0.0452	0.039	
			Fasting insulin	-0.0815 ± 0.0361	0.024	

Table 3A: Cross sectional studies of fibre intake and metabolic risk factors

^A When data is reported across tertiles, quartiles or quintiles (up to Q3, Q4, or Q5) the values represent Ptrend.

FFQ Food frequency questionnaire; IGT- impaired glucose tolerance; NGT- normal glucose tolerance; HOMA- IR homeostasis model assessment of insulin resistance; OGTT- oral glucose tolerance test; AUC- area under the curve; FSIVGTT- frequently-sampled intravenous glucose tolerance test; BMI- body mass index;

Reference Cohort	Subjects N (M/F) Age	Determination of fibre	Fibre intake	Adjusted result/OR/RR	P/Ptrend ^A	Factors adjusted for
	Follow-up			(95%CI)		
Lovejoy & DiGirolamo (1992)	45 (11/34) 22 lean 23 obese	AOAC-defined total dietary fibre by FFQ Mean ± SD (g/d)	FSIVGTT SI	Partial R ² = 0.18 Data not shown	0.007 NSD	Multiple regression model with total energy intake, % energy from fat & CHO BMI
		Lean: 15.8 ± 7.9 Obese: 10.8 ± 1				
McKeown et al	2,834	AOAC-defined total	Quintile median (g/d)	Mean HOMA-IR		Age, gender, BMI, WHR,
(2004)	(1,290/1,544)	dietary fibre by	Tabal films	(95%CI)		smoking, E intake,
Framingham	26.92 40250	FFQ		70(6972)		alconol, %E SFA, %E
Offspring Cohort	20-02 years		02.149	(0.8-7.3)	< 0.001	physical activity BP
Chispining Conort			03: 17.4	6.7 (6.5-7.0)	<0.001	treatment
			04: 20.1	6.7 (6.5-7.0)		
			Q5: 25.5	6.4 (6.1-6.1)		
			Cereal fibre			
			Q1: 2.6	6.8 (6.5-7.0)		
			Q2: 3.7	6.9 (6.7-7.2)	0.02	
			Q3: 4.6	6.8 (6.6-7.0)		
			Q4: 5.8	6.6(6.4-6.9)		
			Q5: 6.0 Fruit fibro	0.5 (0.3-0.8)		
				70(67-72)		
			02:17	6.8 (6.5-7.0)	< 0.001	
			03: 2.8	6.8 (6.5-7.0)	101001	
			Q4: 4.2	6.6 (6.4-6.8)		
			Q5: 5.8	6.5 (6.2-6.7)		
			Vegetable fibre			
			Q1: 2.4	6.7 (6.4-6.9)		
			Q2: 3.7	6.9 (6.6-7.2)	0.64	
			Q3: 4.8	6.7 (6.4-6.9)		
			Q4: 6.1	6.8 (6.5-7.0)		
			US: 8.4	0.0 (0.7-2.0)		
			$01 \cdot 0.23$	68(65-70)		
			02: 0.69	6.8 (6.6-7 1)	0.58	
			03: 1.0	6.8 (6.5-7.0)		
			Q4: 1.4	6.7 (6.5-6.9)		
			Q5: 2.5	6.7 (6.5-7.0)		

^A When data is reported across tertiles, quartiles or quintiles (up to Q3, Q4, or Q5) the values represent Ptrend.

Reference Cohort	Subjects N (M/F) Age	Determination of fibre	Fibre intake	Adjusted result/OR/RR	Ptrend	Factors adjusted for
	Follow-up			(95%CI)		
McKeown et al (2004) continued				Odds Ratio metabolic syndrome (95%CI)		Age, gender, BMI, WHR, smoking, Energy intake, alcohol, %E SFA,
Framingham Offspring Cohort				1.00 (ref) 0.81 (0.61-1.09) 0.88 (0.65-1.19) 0.81 (0.59-1.07) 0.73 (0.51-1.03)	0.11	%Energy PUFA, vitamin use, physical activity, Blood pressure treatment
				1.00 (ref) 0.87 (0.65-1.16) 0.88 (0.66-1.18) 0.74 (0.54-1.00) 0.62 (0.45-0.86)	0.002	
				1.00 (ref) 1.07 (0.80-1.43) 0.74 (0.55-1.01) 0.89 (0.65-1.21) 0.88 (0.64-1.22)	0.36	
				1.00 (ref) 1.08 (0.81-1.45) 1.04 (0.77-1.40) 1.00 (0.74-1.36) 1.15 (0.84-1.57)	0.51	
				1.00 (ref) 0.91 (0.68-1.23) 0.90 (0.67-1.20) 1.00 (0.75-1.34) 0.96 (0.72-1.29)	0.98	

FFQ Food frequency questionnaire; BMI- body mass index; SFA- saturated fatty acids; PUFA- polyunsaturated fatty acids; CHO- carbohydrate; FSIVGTT- frequently-sampled intravenous glucose tolerance test; HOMA- IR homeostasis model assessment of insulin resistance; WHR- waist hip ratio

Reference Cohort	Subjects N (M/F) Age Follow-up	Determination of fibre	Fibre intake	Adjusted result/OR/RR (95%CI)	P/Ptrend	Factors adjusted for
Mooy et al (1998) Hoorn Study	2,226 (1,013/1,213) 50-74 years 1,878 NGT 239 IGT 109 newly diagnosed diabetics	AOAC-defined total dietary fibre by FFQ Mean ± SD (g/d) M: 28 ± 9 F: 26 ± 7 All: 27 ± 8	Fasting insulin	% change with +2SD fibre intake M: -1.5 F: -6.0 All: -3.0	NSD P<0.05 NSD	Age, family history diabetes, BMI, WHR, physical activity, alcohol, smoking, energy intake
Newby et al (2007) Baltimore Longitudinal Study of Ageing	455 to 1,324 depending on outcome 27-88 years	AOAC-defined total dietary fibre by FFQ Quintile median cereal fibre intake (g/d) Q1: 2.2 Q5: 9.5	Fasting glucose Mean ± SEM mmol/l n=1324 OGTT 2h glucose Mean ± SEM mmol/l n=882	Q1-5 cereal fibre intake 5.55 ± 0.05 5.48 ± 0.05 5.53 ± 0.05 5.52 ± 0.05 5.52 ± 0.05 8.05 ± 0.21 7.94 ± 0.20 7.72 ± 0.19 7.55 ± 0.20 6.48 ± 0.21	0.95	Age, gender, total energy, decade of visit, BMI, race, education, supplement use, smoking, % energy from SFA & alcohol
			Fasting insulin Mean ± SEM pmol/l n=460 OGTT 2h insulin Mean ± SEM pmol/l n=455	68.9 ± 4.0 72.2 ± 3.8 73.0 ± 3.7 71.3 ± 3.8 73.0 ± 4.0 438 ± 38.8 477 ± 36.2 404 ± 35.8 356 ± 36.3 413 ± 38.2	0.68	

^A When data is reported across tertiles, quartiles or quintiles (up to Q3, Q4, or Q5) the values represent Ptrend.

IGT- impaired glucose tolerance; NGT- normal glucose tolerance; OGTT- oral glucose tolerance test; FFQ Food frequency questionnaire; BMIbody mass index; SFA- saturated fatty acids; NSD- no significant difference

Reference Cohort	Subjects N (M/F) Age Follow-up	Determination of fibre	Fibre intake	Adjusted result/OR/RR (95%CI)	Р	Factors adjusted for
Ylonen et al (2003)	552 (248/304)	Enzymatic method of Asp et al 1983-	HOMA-IR	Adjusted $\beta \pm SE$	0.012	Age, gender, BMI, waist: hip ratio, education,
Botnia Dietary		total dietary fibre		Total fibre		physical activity, blood
Study	20-70 years	by 3 day diet record	HOMA-IR	0.17 ± 0.07	0.024	pressure, blood lipids
				Insoluble fibre		
		Median intake (IQR) (g/d)	HOMA-IR	-0.15 ± 0.07	0.049	
				Soluble fibre		
		M:		-0.14 ± 0.07		
		21.6 (17.6-25.7)				
		F:				
		17.1 (14.2-20.6)				

HOMA-IR- homeostasis model assessment of insulin resistance; BMI- body mass index

Reference	Subjects N (M/F) Age Health	Study duration & design	Intervention	Fibre dose	Outcome	Fibre group	Control group	P
Barroso Aranda et al (2002)	19 (8/11) 34 years (mean) Healthy	2 x 3 days 3 day run- in & 4 day washout	Psyllium & chitosan supplement or placebo	300 mg/d psyllium 2100 mg/d chitosan	Mean faecal fat excretion (g/d)	9.30	5.82	0.002
Jenkins et al (1993)	43 (15/28) 29-70 years Hyper- lipidaemic	2 x 4 months 2 month run-in and 2 month washout Crossover	Controlled diet high in soluble (barley, legumes, oat bran, psyllium-enriched cereal) or insoluble (wheat bran cereal & bread, crackers) fibre; run-in & washout on NCEP step 2 diet	Mean fibre \pm SE (g/d) soluble / insoluble diet: Total fibre: 49.8 \pm 1.7 / 58.3 \pm 2.1 Soluble fibre: 16.1 \pm 0.5 / 10.2 \pm 0.4 Insoluble fibre: 33.6 \pm 1.2 / 47.9 \pm 1.8	Mean ± SE Weight change (g/wk)	Soluble diet: 29 ± 16	Insoluble diet: 62 ± 19	0.058
Rolls et al (1999)	33 (F) 18-45 years n=16 lean n=17 obese	2 x 1 week Crossover	Portion of each meal manipulated (low or high energy density) and consumed in full, rest of intake ad libitum	Low & high ED foods: identical energy and macronutrients Mean fibre intake from compulsory foods \pm SEM (g/d): Lean: LED 25 \pm 1.0 HED 11 \pm 0.4 Obese: LED 34 \pm 1.6 HED 15 \pm 0.7	Mean energy intake ± SEM (MJ/d) Total (lean) Total (obese) Side dishes (lean) Side dishes (obese) Snacks (lean) Snacks (obese)	8.03 ± 0.37 9.75 ± 0.42 2957 ± 213 3751 ± 237 0.99 ± 0.19 0.60 ± 0.13	8.47 ± 0.37 10.52 ± 0.38 3.51 ± 0.18 4.47 ± 0.24 0.90 ± 0.21 0.68 ± 0.18	NSD <0.025 <0.025 <0.025 NSD NSD
Ryttig et al (1990)	19 (10/9) 18-40 years Healthy	2 x 2 weeks 2 weeks run-in Crossover	Pectin or placebo supplement	7g/d soluble fibre from pectin	Mean ± SE Weight (kg) Lean mass (kg) Faecal energy excretion Basal metabolic rate (W) 24h energy expenditure (MJ)	64.6 ± 2.0 47.0 ± 2.0 2.32 ± 0.25 81 ± 3.7 10.71 ± 0.09	64.1 ± 2.0 46.5 ± 2.0 2.30 ± 0.19 82 ± 3.3 10.84 ± 0.36	NSD NSD NSD NSD

Table 4A: Highly-controlled intervention studies of fibre intake and body weight/weight-related outcomes

NCEP- National Cholesterol Education Program; NSD- no significant difference

Reference	Subjects N (M/F) Age Health	Study duration & design	Intervention	Fibre dose	Outcome	Fibre group	Control group	P
Southgate et al (1976)	5 (3/2) 25-67 years	2 x 1 week Crossover	Controlled diet with high or low fibre biscuits	Bran biscuits: 13.2 g/100g total fibre (9.2 non-cellulosic polysaccharide, 2.7 cellulose, 1.2 lignin) Mean increase in fibre intake ± SD (g/d): 13.8 ± 4.1	Energy excretion (MJ/d)	+0.08 to 0.40		Not stated
Stevens et al (1987)	12 (F) 22-38 years Healthy	4 x 1 week Crossover	Psyllium or wheat bran or psyllium + wheat bran or control crackers before meals, then ad libitum intake of pre- weighed low fibre meals	Psyllium (P): 7g/d insoluble + 17g/d soluble fibre Wheat bran (B): 21g/d insoluble + 1g/d soluble fibre Combination: 15g/d NDR + 7g/d soluble fibre Control: 3g/d NDR + 1g/d soluble fibre	Mean energy intake (MJ/d) Energy excretion (MJ/d)	Psyllium: 8.60 Bran: 9.02 P+B: 8.78 P: 0.67 ± 0.09 B: 0.67 ± 0.07 P+B: 0.66 ± 0.10	9.26 0.40 ± 0.10	<0.05 NSD <0.05 NSD relative to intake
Weinreich et al (1977)	25 19-29 years	2 week control period, 5 weeks intervention Within- subject	Controlled diet ± wheat bran	24g/d wheat bran added to food	Mean weight change (kg)	-0.4	0	Not stated
Wisker et al (1988)	6 (F) 23-27 years Healthy	2 x 3 weeks 4 weeks washout Crossover	Controlled diet with low or high fibre cereal products; constant weight maintained	High fibre (HF): 48.3 g/d (91% NSP, 9% lignin) Low fibre (LF): 19.7 g/d (98% NSP, 2% lignin) Carbohydrate intake HF>LF (314 ± 13 / 262 ± 12 g/d)	Mean total energy excretion ± SE (MJ/d) Mean energy intake ± SD (MJ/d) Difference in metabolisable energy (MJ/d) HF-LF	1.71 ± 0.42 9.80 ± 1.22 +297	1.06 ± 0.19 8.85 ± 1.25	<0.001 Not stated

NSD- no significant difference; NSP- non starch polysaccharide

Reference	Subjects N (M/F) Age Health	Study duration & design	Intervention	Fibre dose	Outcome	Fibre group	Control group	P
Cicero et al (2007)	141 (70/71) 50-70 years Overweight	6 months 4 week run- in Parallel	4 week run-in: all given AHA dietary advice, then guar gum or psyllium powder before meals or control - no placebo	3.5 g/d guar gum (G) or psyllium (P)	Mean changes: BMI ± SD (kg/m²)	G: -1.8 P: -1.1 (both P<0.01)	-0.1 (NSD)	<0.01 NSD
Frost et al (2004)	55 (49/6) 30-70 years CHD patients	12 weeks Parallel	Individual nutrition advice: low glycaemic index (GI) or general healthy-eating Only differences between diets were fibre & sucrose intakes	Mean fibre intake \pm SE (g/d) Southgate Low GI: 27 \pm 2 Control: 21 \pm 2 (P=0.0334) Englyst Low GI: 20 \pm 1 Control: 15 \pm 1 (P=0.0059) Mean sucrose intake \pm SE (g/d) Low GI: 37 \pm 5 Control: 27 \pm 2 (P=0.0029)	Mean changes: Weight (kg) Waist:Hip ratio	-1.4 -0.01	-2.1 -0.01	NSD NSD

Table 5A: Effect of fibre on body weight in `healthy-eating' dietary interventions

CHD- coronary heart disease; HOMA-IR- homeostasis model assessment of insulin resistance; HOMA- β - an index of pancreatic β -cell function ; AHA- American Heart Association; NSD- no significant difference
Reference	Subjects N (M/F) Age Health	Study duration & design	Intervention	Fibre dose	Outcome	Fibre group	Control group	Ρ
Howard et al (2006) Women's Health Initiative	46,808 (F) 50-79 years	7.5 years Parallel	Group & individual counselling sessions to promote behaviour change to reduce fat & increase fruit, vegetables & grains; Control group given Dietary Guidelines for Americans	Mean fibre intake \pm SD (g/d) Baseline: 14.4 \pm 6.0 Intervention: 16.9 \pm 7.1 Control: 14.4 \pm 6.1 (P<0.001)	Mean weight loss ± SD (kg)	0.8 ± 10.1	0.1 ± 10.1	<0.001
Mackay & Ball (1992)	39 (22/17) 28-66 years	3 x 6 weeks 4 week run- in Crossover	Low fat diet run-in, then + high (HF) or low fibre (LF) oat bran or beans (quantity to match soluble fibre content of HF oat bran)	HF bran: 55g/d (5.4% β- glucan) Beans: 80g/d LF bran: 55g/d (3.5% β- glucan)	Mean weight ± SD (kg) Mean energy intake ± SD (MJ/d)	HF bran: 76.6 ± 14.0 Beans: 76.9 ± 14.1 HF bran: 7.84 ± 2.31 Beans: 7.16 ± 1.67	<i>LF bran:</i> 76.7 ± 13.5 <i>LF bran:</i> 7.61 ± 1.79	NSD NSD NSD NSD
Reyna- Villasmil et al (2007)	38 (M) 55-72 years Overweight	8 weeks Parallel	AHA step II diet with bread $\pm \beta$ -glucan	6g/d β-glucan	Mean changes: Weight ± SE (kg)	-5.8	-3.8	<0.002
Tuomilehto et al (1980)	32 (F)	4 months Parallel	Guar gum (n=10) or placebo (n=11) or no treatment (n=11)	15 g/d guar gum	Mean weight ± SE (kg)	-2.5 (P<0.0005)	-0.4 (NSD) Untreated: -0.5 (NSD)	Not given

AHA- American Heart Association; NSD- no significant difference

Reference	Subjects N (M/F) Age Health	Study duration & design	Intervention	Fibre dose	Outcome	Fibre group	Control group	Ρ
Davy et al (2002)	36 (M) 50-75 years Overweight and obese	12 weeks Parallel	Wholegrain oat or wheat cereals substituted into diet	Both 14 g/d fibre; similar energy and macronutrient contents Oat products: 5.5g/d β-glucan	Mean (SD) Weight (Kg) BMI (Kg/m ²)	Oat 92.1 (± 3.0) 29.8 (± 0.8)	Wheat 93.2 (± 3.0 29.5 (± 0.8	*
Effertz et al (1991)	30 (1/29)	12 week	Isocaloric crackers 3x/d ± soy polysaccharide	Soy polysaccharide crackers 20.3 g/d	Waist circumference (cm) Mean weight change (kg)	-0.04	+0.76	Not stated NSD
	18+ years Overweight and obese	2 week run-in Parallel		fibre (83% insoluble, 17% soluble) placebo crackers 0.7g/d fibre	Mean change in energy intake (MJ)	-1.36	-0.84	NSD
Eliasson et al (1992)	63 (39/24) 18-75 years	3 months Parallel	Fibre (n=32) or placebo (n=31) tablet	7 g/d beet, barley & citrus fibre	Mean weight change (kg)	-0.8 (P=0.05)	-0.2 (NSD)	0.04
Evans & Miller (1975)	11 (4/7)	2 x 2 week	Methylcellulose or guar gum supplement	2 x 15.5g/d methylcellulose granules (64% methylcellulose) 2 x 16g/d guar supplement (56% guar gum)	Mean energy intake as % baseline ± SE Mean weight change ± SE (kg/wk)	Supplements similar results combined: Obese: 75 ± 8 Non-obese: 96 ± 3 Obese: -1.7 ± 0.4 Non-obese: -0.1 ± 0	rly efficacious so	<0.05 NSD <0.01 NSD
Howarth et al (2003)	11 (4/7) 23-46 years	2 x 3 week 4 week washout	Fermentable (pectin, β- glucan) or non-fermentable (methylcellulose) fibre supplement before meals	30g/d fibre	Mean energy intake ± SE (MJ/d)	Non-fermentable: 7.7 ± 0.5	Fermentable 8.2 ± 0.8	NSD
	Healthy	Crossover			Mean weight change (kg)	-0.3 (NSD)	-0.1 (NSD)	NSD
					fat (%)	+0.13 (NSD)	+0.13 (NSD)	NSD

Table 6A: Effect of fibre supplementation of habitual diets on body weight

Authors stated that there were significant changes over time (P<0.05), but the differences between groups are significant was not reported.

BMI= body mass index; NSD- no significant difference

Reference	Subjects N (M/F) Age Health	Study duration & design	Intervention	Fibre dose	Outcome	Fibre group	Control group	Р
Lo & Cole (1990)	18 (10/8) 27-60 years Healthy	2 x 6 weeks 3 week washout	Cereal & bread ± soy fibre substituted into diet	15g/d soy cotyledon fibre 79% NSP (Englyst): 21% soluble, 58% insoluble	Weight	No change	No change	NSD
Marett & Slavin (2004)	54 (28/26)	6 month	Larch (n=19) or tamarack (n=19) arabinogalactan supplement or placebo	8.4g/d arabinogalactan	Weight Mean change in	No change	No change -0.51 ($P < 0.01$)	NSD
(2004)	18-55 years		(n=17)		fasting glucose (mmol/l)	T: -0.33 (P<0.05)	0.51 (1 < 0.01)	NSD
	neatiny				Mean change in fasting insulin (pmol/l)	L: -1.04 (NSD) T: +0.48 (NSD)	-0.76 (NSD)	NSD
Pelkman et al (2007)	29 (F)	3 x 1 week 1 week	Alginate-pectin beverage or placebo	1.0 or 2.8g alginate + pectin	Mean energy intake ± SD (MJ/d)	1.0g: 10.86 ± 0.46	11.37 ± 0.46	NSD
	Overweight and obese	Crossover				2.8g: 10.85 ± 0.46		
Ross et al	5 (M)	1 week	Gum arabic supplement	25 g/d Gum arabic	Dietary intakes	-	-	NSD
(1905)	30-55 years Healthy	then 3 week intervention			Median faecal fat excretion (range) (mmol/24h)	18.6 (8.4-46.6)	15.2 (9.7-24.1)	
Rigaud et al (1987)	20 (10/10)	2 x 4 week 2 week run-in	Fibre or placebo tablets	7.3g/d total fibre	Mean energy intake (95%CI)	8.19 (7.35-9.03)	8.24 (7.54-8.95)	NSD
	21-30 years Healthy	Crossover			Mean faecal energy (95%CI) (MJ/d)	0.72 (0.68-0.77)	0.64 (0.57-0.72)	<0.05
Vajifdar et al (2000)	114 (91/23)	6 months	Fibre or placebo powder supplement 2x/d after meals	9.2 g/d fibre (5.8 g/d soluble, 3.4	Mean weight change (kg)	-1.0 (P=0.002)	-0.8 (P=0.002)	NSD
	35-84 years	Parallel		g/d insoluble)	Mean change in waist (cm)	-1.3 (P=0.03)	0	NSD
	Chronic IHD							
Walsh et al (1984)	20 (F) Obese	8 weeks Parallel	Glucomannan supplement vs placebo	1g/d glucomannan	Mean weight change ± SE (kg)	-2.5 ± 0.7	+0.7 ± 0.7	<0.005

NSD- no significant difference; NSP non starch polysaccharide

Reference	Subjects N (M/F)	Study duration	Intervention	Fibre dose	Outcome	Fibre group	Control group	Р
Astrup et al (1990)	Age Health 22 Obese	& design 2 x 2 weeks Crossover	VLCD formula (M: 1.95 MJ/d, F: 1.62 MJ/d) ± fibre	30 g/d birch fibre (98.5% cellulose)	Weight loss (kg)	- 38.0	- 44.5	NSD P<0.01
Birkevedt et al (2000)	53 (F) 18-67 years Overweight	24 weeks Parallel	5.0 MJ/d diet with 15 g/d dietary fibre + fibre supplement or placebo	6g/d 8 weeks, then 4g/d cereal & citrus fibre (85% insoluble, 15% soluble)	Mean weight change ± SD (kg)	-	-	NSD
Birkevedt et al (2005)	167 30-60 years Overweight	5 weeks Parallel	5.0 MJ/d diet + fibre tablet or placebo	 (1) glucomannan (4320 mg/d) + guar (900 mg/d) + alginate (900 mg/d) (2) glucomannan (1240 mg/d) (3) glucomannan (420 mg/d) + guar (420 mg/d) 	Mean weight change ± SD (kg)	(1) -4.4 ± 2.0 (2) -3.8 ± 0.9 (3) -4.1 ± 0.6 NSD between supplements	-2.7 ± 1.3 -2.5 ± 0.5 -2.1 ± 0.5	<0.001 <0.01 <0.01
Duncan et al (1960)	57 (7/50) 38-78 years Obese	8 weeks 4 week run-in Parallel	Methylcellulose supplement or placebo	4.5g/d methylcellulose	Mean weight change ± SD (kg)	-0.3 ± 1.9	-0.1 ± 1.2	NSD
Hylander & Rossner (1983)	110 (8/102) 15-72 years Obese	2 weeks Parallel	5.86 MJ/d dietary advice + wheat bran (n=44) or psyllium granules (n=43) before meals or control - no placebo (n=23)	19.8 g/d wheat bran or psyllium	Mean weight change ± SD (kg)	Bran: -4.6 ± 2.3 Psyllium: -4.2 ± 3.2	-4.6 ± 2.7	NSD NSD

Table 7A: Effect of fibre supplementation of weight-reducing diets on body weight

VLCD- very low calorie diet; NSD- no significant difference

Reference	Subjects N (M/F) Age Health	Study duration & design	Intervention	Fibre dose	Outcome	Fibre group	Control group	P
Kovacs et al (2001)	28 (M) 19-56 years	3 x 2 weeks	Low energy meals ± fibre for breakfast, lunch and snack; dinner <i>ad libitum</i>	7.5 g/d guar gum	Mean weight change ± SD (kg)	-2.1 ± 0.3	-1.6 ± 0.2	NSD
	Overweight	run-in			Mean energy intake snacks ± SD (MJ/d)	0.8 ± 0.2	1.0 ± 0.2	<0.01
					Mean energy intake dinner ± SD (MJ/d)	3.2 ± 0.2	3.4 ± 0.2	NSD
Rossner et	54 (F)	2 months	5.86 MJ/d diet with 5 g/d fibre + fibre tablet or placebo	1.68 g/d cereal & citrus fibre	Mean weight	-7.0	-6.0	<0.05
	18-60 years	Parallel			(kg)	(P<0.01)	(P<0.01)	
	Obese							
	41 (F)	3 months	6.70 MJ/d diet with 7 g/d	2.16 g/d vegetable, cereal &	Mean weight	-6.2	-4.1	<0.05
	18-60 years	Parallel			(kg)	(P<0.001)	(P<0.001)	
	Obese							
Rossner et al (1988)	62 (F)	12 weeks	6.7 MJ/d diet + fibre tablet or placebo	6 g/d vegetable, cereal & citrus fibre	Mean weight change (range)	-4.1 (-11.4, 0.2)	-4.4 (-16.9, 0.6)	NSD
		i di dilei				(1 < 0.05)	(1 < 0.05)	NCD
	Obese				circumference	(P<0.05)	(P<0.05)	NSD
Ryttig et al (1989)	97 (F) 18-55 years	27 weeks Parallel	11 week 5 MJ/d diet with 26g/d fibre + fibre supplement (n=62) or	6g/d fibre (predominantly insoluble)	Mean weight change ± SE (kg)	-3.8 ± 0.5	-2.8 ± 0.9	<0.05
	Overweight		placebo (n=35); then 16 week 6.72 MJ/d diet with 34g/d fibre + fibre supplement or placebo		Change in Waist:Hip	Decrease (P<0.01)	Decrease (P<0.01)	NSD
Rigaud et al	52	6 months	25-30% hypoenergetic diet +	7g/d beet, barley and citrus	Mean weight	-5.5 ± 0.7	-3.0 ± 0.5	0.005
(1990)	(11/41)	Parallel	fibre tablet or placebo	fibre	change ± SE (kg)			
	16-60 years							
	Overweight				Mean energy intake ± SE (MJ/d)	8.35 ± 0.46	9.17 ± 0.65	<0.01

NSD- no significant difference

Reference	Subjects N (M/F) Age Health	Study duration & design	Intervention	Fibre dose	Outcome	Fibre group	Control group	Р
Saltzman et al (2001a; 2001b)	41 (20/21) 19-30 and 64-78 BMI 20.5- 33.9	6 weeks 2 week run-in	Run-in on controlled low soluble fibre diet & energy requirement established; 6 weeks 4.18 MJI/d reduction: macronutrient matched diets with oat or low fibre wheat products	45 g oats/4.18 MJ Mean fibre intake \pm SD (g/d): Total Oat 16.3 \pm 6.9 Control 12.5 \pm 5.1 Soluble Oat 7.2 \pm 3.5 Control 3.5 \pm 1.4	Mean changes: Weight ± SD (kg)	-3.9 ± 1.6	-4.0 ± 1.1	NSD
Solum et al (1987)	60 (F) 30-60 years Overweight	12 weeks Parallel	5.0 MJ diet with 25 g/d dietary fibre + fibre tablet or placebo	6 g/d cereal & citrus fibre	Mean weight change (95%CI) (kg)	-8.5 (-7.5,-9.5) (P<0.01)	-6.4 (-4.8,-8.0) (P<0.01)	<0.01
Yudkin (1959)	20 Overweight	6 weeks	Low carbohydrate diet with unlimited fat and protein ± fibre tablet – no placebo	10g/d methylcellulose	Mean weight change ± SE (kg)	-4.4 ± 0.7	-2.8 ± 0.5	0.05
Valle-Jones (1980)	53 (13/40) 19-65 years Overweight	6 weeks Parallel	4.18 MJl/d diet ± fibre supplement	20g/d fibre supplement (60% vegetable gums: sterculia & guar gum)	Median weight change (kg)	-3.6	-1.8	<0.005

NSD- no significant difference

Table 8A: Effect of fibre	supplementation in	maintenance of w	eiaht-loss
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Reference	Subjects N (M/F) Age Health	Study duration & design	Intervention	Fibre dose	Outcome	Fibre group	Control group	Ρ
Cairella et al (1995)	30 (8/22) Obese	85 days Parallel	15 days VLCD 10 days re-introduction food 60 days fibre tablet or placebo	6 g/d vegetable, citrus & cereal fibre	Mean BMI ± SD (kg/m²)	32.3 ± 4.0	32.9 ± 3.8	<0.01
Pasman et al (1997a)	31 (F) 41 years (mean) Obese	2 month VLCD, then 14 months Parallel	2 month VLCD then fibre (n=20) or control - no placebo (n=11) Fibre group subdivided by compliance A: >80% (n=10) B: 50-80% (n=10)	20g/d guar gum	Mean weight regain ± SD (%)	A: 65 ± 65 B: 123 ± 63	61 ± 66	0.07
Pasman et al (1997b)	17 (F) 39 years Overweight & obese	2 x 1 week Crossover	3 months after 10 kg weight loss on VLCD Energy intake of 4 or 6 MJ/d prescribed ± fibre supplement	20 g/d guar gum	Mean energy intake ± SEM (MJ/d)	5.4 ± 0.24	6.7 ± 0.39	<0.05

VLCD- very low calorie diet

Reference	Subjects N (M/F) Age Health	Study duration & design	Intervention	Fibre dose	Outcome	Fibre group	Control group	Ρ
Gropper et al (1987)	8 (3/5) 6-12 years Overweight	2 x 4 weeks Crossover	2.1 MJ/d less than habitual diet + fibre supplement or placebo	Fibre tablet (corn bran, wheat bran, oat flakes, corn germ meal) 15g/d fibre	Mean weight loss (g)	336	33	NSD
Pena et al (1989)	80 (40/40) 10-15 Overweight	4 weeks Parallel	High or low fibre diet ± physical activity	Total dietary fibre 20 ± 5 vs 3-6g/d	Weight loss high vs low fibre	-	-	NSD
Vido et al (1993)	60 8-14 years Overweight	2 months Parallel	Glucomannan or placebo supplement	2g/d Glucomannan	% overweight mean ± SD (%)	49.5 ± 18.3 to 46.1 ± 18.7 (P=0.01)	43.9 ± 19.7 to 41.7 ± 18.0 (P=0.05)	NSD

Table 9A: Effect of fibre supplementation on body weight in children

NSD- no significant difference

Reference	Subjects N (M/F) Age Health	Study duration & design	Intervention	Fibre dose	Outcome	Fibre group	Control group	P
Aller et al (2004)	53 (19/34)	3 months Parallel	Fibre-enriched diet or control diet	30.5 g/d total fibre Control: 10.4 g/d total fibre	Change in mean ± SE			
	18-70 years				Fasting glucose (mmol/l)	-0.7 (P<0.05)	+0.1 (NSD)	NSD
	Treatiny				Fasting insulin (pmol/I)	+3.9 (NSD)	+2.1 (NSD)	NSD
Birkevedt et al (2000)	53 (F) 18-67 years Overweight	24 weeks Parallel	5.0 MJ/d diet with 15 g/d dietary fibre + fibre supplement or placebo	6g/d 8 weeks, then 4g/d cereal & citrus fibre (85% insoluble, 15% soluble)	Change in fasting glucose	-	-	NSD
Cicero et al (2007)	141 (70/71) 50-70 years	6 months 4 week run- in	4 week run-in: all given AHA dietary advice, then guar gum or psyllium powder before meals or	3.5 g/d guar gum (G) or psyllium (P)	Fasting glucose ± SD (mmol/l)	G: -0.4 P: -1.3 (both P<0.01)	-0.7 (P<0.01)	NSD <0.01
	Overweight	Parallel	control - no placebo		Fasting insulin ± SD (mmol/I)	G: -18.8 P: -35.4 (both P<0.01)	+2.1 (NSD)	<0.01 <0.01
					HOMA-IR ± SD	G: -0.8 P: -2.9 (both P<0.01)	-0.3 (NSD)	NSD <0.01
Chearskul et al (2006)	28 (F)	3 menstrual cycles	Controlled diet provided: low fibre for 1 cycle then	High fibre: 25-30 g/d fibre Low fibre: 8-10	Mean ± SE			
	18-20 years	Within-	high fibre for 2 successive cycles	g/d fibre	Fasting glucose (mmol/l)	5.20 ± 0.07	5.26 ± 0.10	NSD
	пеаітлу	SUDJECT			Fasting insulin (pmol/l)	60.91 ± 7.15	62.99 ± 7.15	NSD

Table 10A: Effect of fibre supplementation on metabolic risk factors

NSD- no significant difference

Reference	Subjects N (M/F) Age Health	Study duration & design	Intervention	Fibre dose	Outcome	Fibre group	Control group	Ρ
Davy et al	36 (M)	12 weeks	Wholegrain oat or wheat	Both 14 g/d fibre;	Mean change in:	Oat diet:	Wheat diet:	
(2002)	50-75 years	Parallel	diet	macronutrient	IVGTT SI	0.0	-0.4	NSD
	Overweight and obese			Oat products: 5.5g/d β-glucan	IVGTT SG (min)	-0.0011	+0.0036	0.03
					Acute insulin response (pmol/ml/min)	-55.7	-4.5	NSD
					Fasting glucose (mmol/l)	+0.2	0	NSD
					Fasting insulin (pmol/l)	-5.4	+3.5	NSD
Eliasson et al (1992)	63 (39/24)	3 months Parallel	Fibre (n=32) or placebo (n=31) tablet	7 g/d beet, barley & citrus fibre	Mean change in fasting insulin (µU/mI)	28.5 (P<0.05)	+8.3 (NSD)	NSD
	10-75 years				Mean change in HbA1c (%)	-0.8 (P<0.0005)	-1.1 (P<0.0005)	NSD
Frost et al (2004)	55 (49/6)	12 weeks Parallel	Individual nutrition advice: low glycaemic index (GI) or general	Mean fibre intake ± SE (g/d)	Fasting glucose (mmol/l)	+0.01	-0.24	NSD
	30-70 years		healthy-eating	Southgate Low GI: 27 ± 2	Fasting insulin (pmol/l)	+4.6	-8.53	NSD
			between diets were fibre	(P=0.0334)	HOMA-Sensitivity	-9.2	-6.76	NSD
				Englyst Low GI: 20 ± 1	ΗΟΜΑ-β	+3.08	+4.2	NSD
				Control: 15 ± 1 (P=0.0059)	HbA1c (%)	+0.04	+0.02	NSD
				Mean sucrose intake \pm SE (g/d) Low GI: 37 \pm 5 Control: 27 \pm 2 (P=0.0029)				

CHD- coronary heart disease; HOMA-IR- homeostasis model assessment of insulin resistance; HOMA-β- an index of pancreatic β-cell function; IVGTT- SI : intravenous glucose tolerance test- insulin sensitivity IVGTT- SG: intravenous glucose tolerance test—glucose effectiveness; NSD- no significant difference

Reference	Subjects N (M/F)	Study duration &	Intervention	Fibre dose	Outcome	Fibre group	Control group	Р
	Age Health	design						
Garcia et al (2007)	11 (4/7)	2 x 6 week	Arabinoxylan supplement as bread	15g/d arabinoxylan	Change in mean ± geometric SE			
	48-70 years	washout	rolls & powder		Fasting glucose (mmol/l)	0	+0.3	0.029
	IGT				Fasting insulin (pmol/l)	-8.0	+10.0	NSD
					MTT AUC glucose	< Placebo		0.005
					MTT AUC insulin	< Placebo		0.003
					MTT AUC ghrelin	< Placebo		<0.001
Hanai et al (1997)	38 (20/18)	2 x 6 month	Corn bran supplementation then	2 x 5g/d hemicellulose	Mean ± SE			
	35-68 years	Crossover	control period - no placebo		<u>Fasting glucose</u> (mmol/l): Obese IGT	5.78 ± 0.93	7.11 ± 0.79	<0.05
	20 obese IGT 8 lean IGT				Non-obese IGT Non-obese NGT	7.11 ± 0.79 4.23 ± 7.1	7.67 ± 0.93 4.76 ± 0.53	P<0.05 NSD
	10 lean NGT				<u>Glucose response to</u> OGTT: Obese IGT	Decrease		<0.05
					Non-obese IGT Non-obese NGT	Decrease No change		<0.05 NSD
					Fasting insulin (pmol/l):			
					Obese IGT Non-obese IGT Non-obese NGT	74.2 ± 13.5 33.72 ± 12.0 40.44 ± 12.0 (Decreases in	91.0 ± 23.6 47.22 ± 20.2 46.92 ± 12.0	NSD NSD NSD
					Inculin recoonce to	hyperinsulinaemic subjects)		
					OGTT: Obese IGT Non-obese IGT	Decrease No change		<0.05 NSD
					Non-obese NGT	No change		NSD
					HbA1c (%): Obese IGT Non-obese IGT Non-obese NGT	6.8 ± 0.3 Data not given Data not given	6.3 ± 0.2 - -	<0.05 NSD NSD

IGT- impaired glucose tolerance, NGT- normal glucose tolerance; MTT- meal tolerance test; AUC- area under the curve; OGTT- oral glucose tolerance test; NSD- no significant difference

Reference	Subjects N (M/F) Age Health	Study duration & design	Intervention	Fibre dose	Outcome	Fibre group	Control group	P
Jenkins et al (1977)	3 (M) Healthy	3 week control period then 6 week intervention	Metabolically controlled diet (control) then same diet + pectin supplement	36g/d pectin	Mean ± SE Fasting glucose (mmol/l)	4.51 ± 0.28	4.97 ± 0.33	NSD
					Fasting insulin (pmol/I)	24 ± 36	30 ± 6	NSD
Jenkins et al (1993)	43 (15/28) 29-70 years Hyper- lipidaemic	2 x 4 months 2 month run- in and 2 month washout Crossover	Controlled diet high in soluble (barley, legumes, oat bran, psyllium-enriched cereal) or insoluble (wheat bran cereal & bread, crackers) fibre; run-in & washout on NCEP step 2 diet	Mean fibre \pm SE (g/d) soluble / insoluble diet: Total fibre: 49.8 \pm 1.7 / 58.3 \pm 2.1 Soluble fibre: 16.1 \pm 0.5 / 10.2 \pm 0.4 Insoluble fibre: 33.6 \pm 1.2 / 47.9 \pm 1.8	Day-profile glucose (mmol/l) Day-profile insulin (pmol/l)	5.61 ± 0.22 224 ± 20	5.78 ± 0.11 251 ± 35	NSD NSD
Keogh et al (2003)	18 (M) 18-65 years Overweight	2 x 4 weeks 4 weeks washout Crossover	Identical provided diets + enriched barley fibre or glucose (control)	8.1-11.9g/d β -glucan from enriched barley fibre Mean NSP intake \pm SD (Englyst; g/d) Barley: 35.8 \pm 4.8 Control: 28.7 \pm 2.6 (P<0.001)	Mean ± SE Fasting glucose (mmol/l) OGTT AUC glucose	5.23 ± 0.11 Data on figure	5.28 ± 0.10 Data on figure	NSD NSD
Kestin et al (1990)	24 (M) 25-65 years	3 x 4 weeks 3 weeks run- in Crossover	Run-in on low fibre diet, then supplementation with fibre from wheat or rice or oat bran	Low fibre run-in diet: 11.2 g/d total fibre, then +11.8g/d fibre Soluble: Insoluble fibre ratio: Wheat bran 0.26 Rice bran 0.32 Oat bran 1.20	Fasting glucose Fasting insulin MTT glucose response MTT insulin response	Data not given	Data not given	NSD NSD NSD NSD
Li et al (2003)	10 (F) 20 (mean) Healthy Lean	2 x 4 weeks Crossover	Barley or control (100% rice) diet	Barley: carbohydrate 30% from barley 70% rice	Mean ± SD Fasting glucose (mmol/l) HbA1c (%) OGTT AUC glucose	4.94 ± 0.77 5.1 ± 0.8 12.39 ± 2.61	5.44 ± 0.77 5.4 ± 0.7 12.89 ± 1.72	NSD NSD NSD

Reference	Subjects N (M/F) Age Health	Study duration & design	Intervention	Fibre dose	Outcome	Fibre group	Control group	Ρ
Maki et al	60	12 weeks	NCEP dietary advice + 3		Mean change \pm SE			
(2007)	(33/17) 40+ years	Parallel	servings/d provided foods: β-glucan- enriched or control		Fasting glucose (mmol/l)	+0.08 ± 0.16	+0.11 ± 0.09	NSD
	Overweight and obese				OGTT AUC glucose (mmol/l/hr)	+8 ± 37	+38 ± 33	NSD
					OGTT peak glucose (mmol/l)	-0.21 ± 0.29	$+0.19 \pm 0.20$	NSD
					Fasting insulin (pmol/l)	+2.8 ± 4.9	+1.4 ± 12.5	NSD
					OGTT AUC insulin ((pmol/l)h)	-3209 ± 2507	-278 ± 4487	0.034
					OGTT peak insulin (pmol/l)	-41.0 ± 35.4	+0.7 ± 63.9	0.037
Munoz et al (1979)	15 (M)	30 days	Controlled low fibre diet	26 g/d fibre	OGTT AUC glucose			
	19-54 years	Crossover	bread	% fibre in supplements: wheat bran 50.8 white wheat 44.1 corn bran 92.1 soy hulls 86.7	Wheat bran White wheat Corn bran Soy hull Apple Carrot	Data on figure	Data on figure	NSD NSD <0.05 <0.05 <0.05 <0.05
				apple powder 25.6 carrot powder 31.0	OGTT AUC insulin All			NSD
Landin et al (1992)	25 (M)	2 x 6 week	Granulated guar gum or placebo before meals	30g/d granulated guar gum	Mean ± SD			
	52 years (mean)	2 week run- in & 2 week washout			Fasting insulin (pmol/l)	57 ± 14	57 ± 14	NSD
	Healthy				Fasting glucose (mmol/l)	4.5 ± 0.5	4.8 ± 0.4	<0.001
					Glucose disposal (mg/kgLBM/min)	15.0 ± 2.4	13.9 ± 2.8	<0.01

OGTT; oral glucose tolerance test; NCEP- National Cholesterol Education Program; MTT- meal tolerance test; AUC- area under the curve; OGTToral glucose tolerance test; NSD- no significant difference

Reference	Subjects N (M/F) Age Health	Study duration & design	Intervention	Fibre dose	Outcome	Fibre group	Control group	Р
Pereira et al (2002)	11 (5/6)	2 x 6 weeks 6-9 weeks	Controlled whole (WG) or refined grain (RG) diet	Fibre intakes whole / refined grain diet (g/d):	Mean ± SE Hunger	WG <rg< td=""><td></td><td>0.08</td></rg<>		0.08
	Overweight and obese	washout		Total fibre: 28.0 / 17.8 Insoluble fibre: 19.7 /	Fasting insulin (2,4,6 wk) (pmol/I)	141 ± 3.9	156 ± 3.9	<0.01
				10.8 Soluble fibre: 7.7 /	HOMA-IR (2,4,6 wk)	5.4 ± 0.18	6.2 ± 0.18	<0.01
				6.7 [(Difference in clamp insulin sensitivity (95%CI) WG-RG	0.7 (0.03-1.44) x10 ⁻⁵		<0.01
Reyna- Villasmil et	38 (M)	8 weeks	AHA step II diet with bread $\pm \beta$ -glucan	6g/d β-glucan	Fasting glucose ± SE (mmol/I)	-0.3	-0.1	NSD
al (2007)	55-72 years	Parallel						
	Overweight							
(2004)	42 (31/11)	5 weeks Parallel	High fibre, high carbohydrate, low fat dietary advice ± wheat	10.5g/d wheat fibre powder for 1st week, then 21g/d	Change in mean fasting glucose (95%CI) (mmol/l)	-0.4 (P<0.05)	+0.1 (NSD)	Not given
	18-70 years	i ululei	fibre supplement - no placebo					
Saltzman et al (2001a;	41 (20/21)	6 weeks	Run-in on controlled low soluble fibre diet &	45 g oats/4.18 MJ	Fasting glucose ± SD (mmol/l)	-0.0005 ± 0.9	-0.21 ± 0.29	NSD
20016)	19-30 and 64-78	2 week run- in	energy requirement established; 6 weeks 4.18 MJI/d reduction:	SD (g/d):	Fasting insulin ± SD (pmol/l)	-9.3 ± 46.7	-28.7 ± 26.5	NSD
	BMI 20.5-		macronutrient matched diets with oat or low	Total Oat 16.3 ± 6.9	HOMA-IR	-0.5 ± 2.1	-1.1 ± 1.2	NSD
	33.9		fibre wheat products	Control 12.5 \pm 5.1				
				Soluble Oat 7.2 ± 3.5 Control 3.5 ± 1.4				
Villaume et al (1984)	5 (M)	3 week control period	Metabolically controlled diet (control) then same	20g/d wheat bran supplement (11g	Mean ± SD			
	21 years (mean)	then 7 week intervention	diet + wheat bran supplement	fibre)	Mean 30 min MTT glucose (mmol/l) Day 0	7 38 + 0 11		
	Healthy MTT-	Within- subject	Controlled diet: 32g/d fibre (Southgate)		Day 24 Day 48	5.00 ± 1.11 Data not given		<0.05 NSD
					60 min MTT insulin (pmol/l)			
					Day 0 Day 24	Data not given 61.8 ± 8.9		NSD
					Day 48	45.5 ± 12.3		<0.02

Reference	Subjects N (M/F) Age Health	Study duration & design	Intervention	Fibre dose	Outcome	Fibre group	Control group	P
Vuksan et al (2000)	11 (5/6)	2 x 3 weeks 8 weeks run-	Controlled matched NCEP step 2 diet enriched with konjac-	15 g/4.18 MJ total fibre from konjac- mannan or wheat	Mean change ± SE Fasting glucose	-13.0 ± 2.48	-9.6 ± 4.27	NSD
	45-65 years	in & 2 weeks washout	mannan (KJM) or wheat bran (WB) fibre	bran	(mmol/l)	(P<0.05)	(NSD)	NCD
	101	Crossover	provided	Baseline 24.2 \pm 11.0 KJM 34.7 \pm 8.4	(pmol/l)	(NSD)	-3.0 ± 9.67 (NSD)	NSD
				WB 33.4 ± 9.6	Fructosamine	-5.6 ± 1.46 (P<0.05)	-0.4 ± 1.3 (NSD)	0.0013
Weickert et al (2006)	17 (F)	3 days	3 macronutrient- matched portions per	Oat fibre product: 70% cellulose, 25%	Mean ± SE			
	53 years (mean)	Crossover	day of oat-fibre enriched or control bread as only food	hemicellulose, 3-5% lignin, 0.2% β-glucan, 0.1% fat, 0.25%	Fasting glucose (mmol/l)	4.91 ± 0.10	4.96 ± 0.10	NSD
	Overweight and obese		Energy provided = 1.5 x Resting Energy Expenditure	protein	Fasting insulin (pmol/l)	29.7 ± 4.3	32.3 ± 4.9	NSD
					Fasting C-peptide (nmol/l)	0.65 ± 0.05	0.68 ± 0.06	NSD
					Adiponectin (µg/ml)	15.4 ± 1.7	16.3 ± 1.9	NSD
					Ghrelin (pg/ml)	163.7 ± 17.5	175.9 ± 18.8	NSD
					Respiratory quotient	0.87 ± 0.02	0.86 ± 0.01	NSD
					Resting energy Expenditure (MJ/d)	4.61 ±0.19	4.66 ± 0.20	NSD
					Glucose disposal (mg/min/kg)	6.56 ± 0.32	6.07 ± 0.27	0.043
					Steady state insulin (mU/ml)	183.7 ± 5.2	193.7 ± 4.9	NSD
					Insulin action (mg/min/kg/mU/l)	3.61 ± 0.20	3.21 ± 0.22	0.023
					Posthepatic insulin clearance (I/min)	1.03 ± 0.04	0.98 ± 0.04	NSD

AHA- American Heart Association; NCEP- National Cholesterol Education Program; HOMA-IR- homeostasis model assessment of insulin resistance MTT- meal tolerance test; NSD- no significant difference

Reference	Subjects N (M/F) Age Health	Study duration & design	Intervention	Fibre dose	Outcome	Fibre group	Control group	Ρ
Noakes et al (1996)	23 (13/10) 44-64 years Overweight + risk factor	3 x 4 weeks Crossover	High-amylose starch or oat bran or low-amylose starch products incorporated into low fat (15%E), low fibre (<10g/d) background diet	 (a) High-amylose starch: Fibre (g/d) M: 48 ± 7, F: 33 ± 3 RS products 33% RS (b) Oat bran: Fibre (g/d) M: 42 ± 9, F: 30 ± 6 Low RS (c) Low-amylose starch: Fibre (g/d) M: 21 ± 3, F: 16 ± 3 Low RS 	Mean ± SD Weight Fasting glucose (mmol/I) Fasting insulin	(a) 5.67 ± 0.64 (b) 5.73 ± 0.57 (a) 79 ± 50 (b) 80 ± 50	(c) 5.58 ± 0.65 (c) 77 ± 57	NSD NSD <0.01 NSD NSD
Park et al ^a (2004)	25 (F) 26-57 years Overweight and obese	3 weeks Parallel	Resistant corn starch (n=12) or corn starch as controls (n=13) supplement	Both supplements 40g/d starch	Fasting glucose Fasting insulin	↓ (P<0.05) No change	No change No change	Not Stated Not stated
de Roos et al (1995)	24 (M) 20-27 years Healthy	3 x 1 week Crossover	RS2 or RS3 or control (glucose) supplement	Both RS supplements: 32 g/d RS Placebo: 4 g/d RS All: 2g/d fibre	Energetic compensation for supplement (%) C-peptide excretion	RS_2 : -5% (increase in energy intake) RS_3 : 15% RS_2 : 4.39 ± 1.46 RS_3 : 3.74 ± 1.42	24% 4.71 ± 1.73	Not stated NSD 0.0001

Table 11A: Effect of resistant starch on weight and metabolic outcomes

^A Park et al (2004) glucose and insulin measurements only shown graphically. Fasting serum glucose was significantly different compared to baseline following resistant starch supplementation (p<0.05), but not following the control starch intervention. NSD- non significant difference; RS2- high amylase corn starch; RS3- extruded and retrograde high amylose corn starch

Reference	Subjects N (M/F)	Study duration	Intervention	Fibre dose	Outcome	Fibre group	Control group	Ρ
	Age Health	& design						
Robertson et al (2005)	10 (4/6)	2 x 4 week	RS_2 or placebo supplement	50 g/d Hi-Maize 260 (30 g RS ₂ + 20 g RDS)	Mean ± SD			
	24-61 years	4 week washout		Placebo: 20g/d	Weight (kg)	71.0 ± 3.88	70.6 ± 3.74	NSD
	Healthy	Crossover		(0 g RS ₂ + 20 g RDS)	Lean body mass (kg)	52.5 ± 3.08	51.4 ± 3.03	0.003
					HOMA-Sensitivity	76.75 ± 6.72	77.4 ± 5.55	NSD
					ΗΟΜΑ-β	138 ± 8.83	128 ± 9.21	NSD
					Fasting glucose (mmol/l)	5.06 ± 0.14	5.04 ± 0.12	NSD
					AUC glucose (mmol/300min/l)	1830 ± 28.9	1890 ± 27.7	NSD
					Fasting insulin (pmol/l)	84 ± 17.4	80 ± 16.0	NSD
					AUC insulin (pmol/300min/l)	55200 ± 10500	63000 ± 10600	0.024
					C-peptide:insulin AUC	7.48 ± 0.734	6.08 ± 0.523	0.027
					Oral insulin sensitivity (x10 ⁻³)	1.82 ± 0.36	1.36 ± 0.19	0.050
					Adipose tissue glucose uptake (µmol/100ml)	141.0 ± 59.3	54.4 ± 62.5	0.007
					Skeletal muscle glucose clearance	43.9% > control		0.013
					Clamp insulin sensitivity (x10 ⁻² M/I)	9.7 ± 1.09	8.5 ± 0.87	0.027

RS2- high amylose corn starch; RDS- rapidly digestible starch; AUC- area under the curve; HOMA- homeostasis model assessment; HOMA-βan index of pancreatic β–cell function; NSD- no significant difference

Reference	Subjects N (M/F) Age Health	Study duration & design	Intervention	Fibre dose	Outcome	Fibre group	Control group	P
Brighenti et al (1999)	12 (M) 23 years Healthy	2 x 4 weeks 4 week washout Healthy	50 g/d cereal ± inulin substituted for habitual breakfast	9 g/d inulin/FOS mixture	Mean weight ± SD	73.3 ± 4.3	78.4 ± 4.2	NSD
Cani et al (2006)	10 (5/5) 21-39 years Healthy	2 x 2 weeks	FOS or placebo at breakfast and dinner	16 g/d FOS	Energy intake	8.94 ± 0.92	9.44 ± 0.70	0.05
Castiglia- Delavaud et al (1998)	9 (M) 22 years (mean) Healthy	3 x 4 weeks Crossover	Provided diet ± sugar beet fibre (SBF) or inulin	Control diet: 22g/d NSP Supplemented with 50g/d sugar beet fibre or inulin (amount built up over 1st 2 weeks)	Mean faecal energy excretion (MJ/d) Metabolisability Energy expenditure (MJ/d)	Inulin: 0.707 SBF: 0.734 Inulin: 0.891 SBF: 0.889 Inulin: 9.90 SBF: 10.04	0.59 0.900 9.84	<0.01 <0.001 <0.01 <0.001 NSD <0.05
Daubioul et al (2005)	7 (M) 37-66 years NASH patients	2 x 8 week 5 week washout Crossover	FOS or placebo at breakfast & dinner	16 g/d FOS	Change in mean ± SE Energy intake Weight Body composition Fasting glucose (mmol/I) Fasting insulin (pmol/I) Fasting C-peptide (pmol/I)	No change No change -0.7 -16.0 -133	No change No change No change +0.6 +9.7 +219	NSD NSD NSD NSD NSD

Table 12A: Effect of oligosaccharides and inulin on weight and metabolic outcomes

FOS- fructo-oligosaccharides; NSD- no significant difference

Reference	Subjects N (M/F)	Study duration & design	Intervention	Fibre dose	Outcome	Fibre group	Control group	Р
	Health	a acsign						
Ellegard et	10	3 x 3 days	Matched meals with inulin or	17g/d inulin or FOS	Mean (95%CI)	Inulin:	FOS:	
ai (1997)	30-71 years	Crossover			Energy excretion relative to control (kJ/d)	+245 (190-307)	+230 (217-315)	<0.05
	lleostomy patients				Recovery (% of ingested)	88 (76-100)	89 (64-114)	
Giacco et al	27	2 x 2	FOS or placebo	10.6 g/d FOS	Mean ± SD			
(2004)	46 years	1 month	avoidance of foods	supplement	Weight	No change	No change	NSD
	(mean)	run-in			Fasting glucose (mmol/l)	5.44 ± 1.00	5.38 ± 0.83	NSD
					Fasting insulin (pmol/l)	51.7 ± 15.1	50.2 ± 11.0	NSD
Jackson et al	54	8 weeks	Inulin or placebo	10 g/d inulin	Mean ± SD			
(1999)	35-65 years	Parallel	(matodextrine) in beverage		Weight	No change	No change	NSD
	Healthy				Fasting glucose (mmol/l)	4.84 ± 0.51	4.99 ± 0.49	NSD
					Fasting insulin (pmol/l)	37.5 ± 18.3	44.9 ± 27.5	NSD
Luo et al	12 (M)	2 x 4	FOS or sucrose	20 g/d FOS	Mean ± SE			
(1990)	19-32 years	Dweeks	supplemented cookies		Weight	No change	No change	NSD
	Healthy	washout			Fasting glucose (mmol/l)	4.90 ± 0.11	4.86 ± 0.09	NSD
		Crossover			Fasting insulin (pmol/l)	57 ± 5	56 ± 3	NSD
					Clamp steady state: Glucose Insulin C-peptide Glucagon			NSD NSD NSD NSD

FOS- fructo-oligosaccharides; NSD- no significant difference

Reference	Subjects N (M/F) Age Health	Study duration & design	Intervention	Fibre dose	Outcome	Fibre group	Control group	P
Schaafsma et al (1998)	30 (M)	2 x 3 week	Fermented milk drink ± FOS with meals	3×125 ml/d fermented milk drink $\pm 2.5\%$ FOS	Weight	No change	No change	NSD
	33-64 years Healthy	1 week washout		with habitual meals	Fasting glucose (mmol/l)	5.47	5.37	NSD
	11	Crossover		Dee filme	Maan watalat	1 ((1 0 2 2)	1 2 (1 0 2 2)	NCD
(2006)	(5/6)	2 x 2 weeks	based on calculated TEE) ± pea-fibre and FOS	Mean (95%CI) (g/d): 18.3 (16.8-19.9)	change (95%CI)	-1.6 (1.0-2.3)	-1.3 (1.0-2.3)	NSD
	25-30 years	6 weeks		(control: 0)				
		washout						
	Healthy	Character		FOS Mean (95%CI)				
		Crossover		(y/u): 9 8 (8 9-10 7)				
				(control: 0)				
STUDIES I		EN	1					1
Abrams et al	97	1 year	FOS + inulin or placebo	8 g/d FOS + inulin mix	Mean ± SD			
(2007)	(49/48)		(maltodextrin) in calcium-					
	0.40	1 year	fortified juice/milk at		BMI Z-score	0.25 ± 0.045	0.38 ± 0.044	0.048
	9-13 years	rollow-up	breaktast		BMI (ka/m^2)	19 52 + 0 15	20.03 + 0.15	0.016
	Healthy	Parallel				19.52 ± 0.15	20.05 ± 0.15	0.010
	,				Weight	47.7 ± 0.4	49.0 ± 0.4	0.048
					Fat mass	11.24 ± 0.25	12.07 ± 0.25	0.022
					Follow-up BMI difference (kg/m ²)		+0.68 ± 0.36	0.061

FOS- fructo-oligosaccharides; BMI-body mass index; TEE- total energy expenditure; NSD- no significant difference

Reference	Subjects N (M/F) Age Health	Study duration & design	Intervention	Fibre dose	Outcome	Fibre group	Control group	Ρ
Schwab et al (2006)	66 (29/37) 30-65 years	12 weeks	Nutrition counselling (dietary goals for prevention of CVD & T2D) + sugar beet pectin (SBP) or polydextrose (PDX) or	16g/d sugar beet pectin or polydextrose	Change in mean weight (kg)	PDX: -1.0 (P=0.007) SBP: -0.3 (NSD)	-0.9 (0.007)	NSD NSD
	Non-obese (10 T2D)		control drink		Change in mean HbA1c (%)	PDX: +0.2 SBP: +0.1 (both P<0.05)	0.0	NSD NSD
					Change in mean fasting glucose (mmol/l)	PDX: +0.1 SBP: +0.1 (both NSD)	+0.2 (0.007)	NSD NSD

Table 13A: Effect of polydextrose on weight and metabolic outcomes

T2D- type 2 diabetes; CVD- cardiovascular disease; NSD- non significant difference

Table 14A: Acute experimental studies

Reference	Subjects	Durati	Intervention	Glucose	Insulin	Other
	N (M/F) Age	on				
Whoat	Health					
Bonini ot al	8 (5/3)	4	High (M:17.4: E:14.0g) or low (M:3.5: E:2.8g) whoat			thunger at 120 & 180 min post-
(1995)	8 (3/3) 28-41	hours	fibre matched meals	▼ AUC (P<0.03)		\checkmark hunger at 120 & 180 min post- consumption (P<0.03)
(1990)	Healthy	nours				
Delargy et al	12 (M)	14	High (20g, 50:50 insoluble: soluble) or low (3g) fibre			NSD appetite or later energy intake (#)
(1995)	Healthy	hours	cereals			
Fontvieille et al	12 (8/4)	3	Bran-enriched (4.2g fibre) or control (1.3g fibre) toast	NSD AUC (#)	NSD AUC (#)	
(1988)	24 (mean)	hours				
Grimos &			Wholemest or control white bread			A amount consumed to reach point of
Gordon (1978)	Healthy		wholemear of control white bread			fullness (P<0.05)
Hallfrisch et al	24 (12/12)	3	OGTT + 0.08 / 0.17 / 0.33g/kg body weight insoluble	NSD AUC (#);	NSD AUC (#);	NSD glucagon responses (#)
(2002)	41 (mean)	hours	fibre (Z-trim: oat, wheat, corn, rice, soy and pea)	\downarrow peak with \uparrow	delayed peak	
	Healthy			dose (P=0.04)	with ↑ dose	
					(P<0.0001)	
Hamberg et al	8 (5/3)	2	OGII \pm 30g pea fibre (PF) / 36g wheat bran (WB) / 22g	Ψ AUC with PF	NSD AUC (#);	
(1989	Z3-30 Healthy	nours	Sugar beet libre (SBF)	NSD ALC with	Ψ at 30 mm with PE only (P<0.05)	
	Treaterry			WB/SBF (#)	11 Only (1 < 0.05)	
Jeffrys (1974)	6 (4/2)	2	Glucose ± unprocessed wheat bran / bagasse / wood	Bran ↓ AUC		
	20-25 years	hours	cellulose (0.2g/kg body weight)	(P<0.01);		
	Healthy			Others ↑ peak		
				(P<0.01)		
Jenkins et al	13(6/7)	2	High / low fibre bread ($10.2/2.8g$), pasta ($7.3/2.0g$), rice	NSD AUC (#)	NSD AUC (#)	
(1901)	SU (mean) Healthy	nours	(3.3/1.4g)			
McIntosh et al	28 (M)	1 hour	High rye fibre (RF) or wheat fibre meal (WF) or control	↓ fed (1hr) –	↓ fed (1hr) –	
(2003)	40-65		low fibre meal	fasted change RF	fasted change RF	
	Overweight			& WF (P<0.0001)	& WF(P<0.0005)	
Molnar et al	10 (4/6)	3	OGTT \pm 15 g unprocessed wheat bran (21% cellulose,	↓ 30 min	↓ 30 min	
(1985)	12 (mean)	hours	26% hemicellulose, 3% pectin, 4% lignin)	(P<0.02)	(P<0.02)	
Porikos &	50 (M)	1 hour	High $(3.3a)$ or control low $(0.2a)$ wheat fibre bread meal			y hunger (immediate post-
Hagamen	18-25	1 noui	(3.3g) of control low (0.2g) wheat the bread theat			consumption $P < 0.005$: 30 min $P = 0.08$)
(1986)	Healthy					\downarrow later energy intake (P<0.05 in total
	,					sample; obese subjects only P<0.025)
Samra &	16 (M)	1.5	High (33g insoluble) or low (1g) fibre cereal	NSD AUC (#)		ψ energy intake at 75 min (P<0.05)
Anderson	20-35	hours				NSD appetite AUC (#)
(2007)	Healthy	15	High (33g incoluble) or low (1g) fibre coreal			NSD alucasa talaransa ta sat sasand
	20-35	hours		NSD AUC (P=0.4)		meal (#)
	Healthy	nours				

Reference	Subjects N (M/F) Age	Durati on	Intervention	Glucose	Insulin	Other
	Health					
Weickert et al (2005)	14 (F) 24 (mean) Healthy	2 hours	Fibre-enriched bread: 10.5g wheat fibre (WF; 94.5% insoluble) / 10.6g oat fibre (OF; 93% insoluble) or un- enriched control bread	NSD AUC (P>0.15)	↑ early (45 min) AUC with OF (P=0.027), NSD WF (P>0.15); NSD 180 min AUC (P>0.15)	Earlier GIP peak with OF (P<0.01)
Wolover et al	77 (M)	2	High (36.7g fibro, 35.8g insoluble) or low (0.5g fibro)	JL AUC & poak		
(2004)	18-75 Healthy	hours	fibre cereal	(both P<0.001)	(P=0.015) & peak (P=0.012)	
Other grain	S		•			
Behall et al (2005)	10 (F) 28-58 Overweight	3 hours	Oat / barley meals (flour & flakes) or glucose (control)	Ψ AUC & peak with all (P<0.05)	Ψ AUC & peak with barley meals only (P<0.05); NSD with oats (#)	NSD glucagon (P>0.71) or leptin (P>0.80) AUC
Bourdon et al (1999)	12 (M) 21-42 Healthy	6 hours	High barley (A: naturally high β -glucan, B: enriched with β -glucan; both 15.7g fibre, 5g β -glucan) or low (4.6g) fibre meals	NSD AUC & peak (#)	NSD AUC with both (#); ψ peak with B (P<0.05) but NSD with A (#)	NSD CCK AUC (#)
Braaten et al (1994)	11 (7/4) 52 (mean) Healthy	3 hours	Wheat farina + oat gum / oat bran / wheat farina (control) breads (insoluble/soluble fibre (g): 0.6/9.5, 10.8/9.2, 0.7/0.8)	 ↓ AUC & peak with both (P<0.05); NSD between oat breads (#) 		
Dahl et al (2005)	11 Overweight	2 hours	White bread ± flax fibre	\downarrow AUC & peak (both P<0.05)		
Karlstrom et al (1988)		4 hours	Cereal (19g fibre) / legume (15.4g fibre) / mixed (18.4g fibre) / control low (5.7g fibre) fibre meal	NSD AUC (#)	 ↓ AUC legume only (P<0.05), NSD AUC other meals (#) 	
O'Dea et al (1980)	6 (M) 20-28 Healthy	4 hours	Brown or white rice meals	NSD AUC (#)	NSD AUC (#)	
Turnbull & Thomas (1995)	17 (F) 23 (mean) Healthy	24 hours	MTT ± 20g plantago granules / placebo / water			↑ fullness at 1 hour vs water & placebo (P>0.05), NSD at 2 hours (P=0.88) & 3 hours (0.35) NSD later energy intake (#)

Reference	Subjects N (M/F) Age Health	Durati on	Intervention	Glucose	Insulin	Other
Wood et al (1994)	9 (4/5) 32 (mean) Healthy	3 hours	OGTT ± oat gum: 1.8g / 3.6g / 7.2g	↓ AUC (P<0.05) & peak (P<0.001) dose-response with \uparrow dose	↓ AUC (P<0.05) & peak (P<0.01) dose-response with \uparrow dose	
	11 (6/5) 38 (mean) Healthy	3 hours	OGTT ± oat gum: 7.2g low / high viscosity	NSD AUC & peak (#)	NSD AUC & peak (#)	
Fruit						
Bolton et al (1981)	Not given 20-43 Healthy	3 hours	Whole grapes (1.4% fibre) or grape juice (0% fibre)	NSD AUC (#); ↓ rate increase (P<0.005)	↑ AUC (P<0.02)	↑ satiety (P<0.05)
			Whole oranges (2.5% fibre) or orange juice (0% fibre)	NSD AUC (#); ↓ reactive late hypoglycaemia (P<0.05)	↓ AUC (P<0.02) & peak (P<0.05)	↑ satiety (P<0.05)
Haber et al (1977)	10 (5/5) 22-40 Healthy	3 hours	Whole apples (2.9% fibre) or apple juice (0% fibre)	↓ reactive late hypoglycaemia (P<0.05)	↓ AUC (P<0.01) & peak (P<0.005)	↑ satiety (P<0.05)
Kay (1978)	10 (F) 20-22 Healthy	3 hours	Whole oranges (8g pectin) or orange juice (0.2g pectin)	NSD AUC (#); ↑ level at 180 min P<0.01)		ψ hunger during 3rd hour (#)

Reference	Subjects N (M/F) Age Health	Durati on	Intervention	Glucose	Insulin	Other
Vegetables						
Moorhead et al (2006)	36 (F) 21-40 Healthy	3.5 hours	Whole carrots (4.4g fibre) or carrot nutrients (0g fibre)			↑ satiety (P<0.05) ↓ later energy intake (P<0.05)
Onyechi et al (1998)	15 (12/3) 24-38 Healthy	2.5 hours	MTT ± vegetable flours: detarium (legume) or cissus (shrub); Both 10-11g total NSP	Ψ AUC (detarium: P<0.001; cissus: P<0.0005)	NSD AUC (pooled P=0.4)	
Raben et al (1994)	10 (M) 20-50 Healthy	6 hours	Pea fibre (4.7g/4.18MJ) or low fibre (1.7g/4.18MJ) meal	NSD AUC (P=0.48)		\downarrow DIT (P<0.05) NSD suppression fat oxidation (P=0.11)
Legumes						
Bourdon et al (2001)	8 (M) 21-45 Healthy	6 hours	MTT \pm bean flakes (11.8g fibre, 3.2g insoluble)	NSD AUC (#)	NSD AUC (#)	↑ CCK AUC (P<0.05)
Burley et al (1993)	18 (9/9) Healthy	4.5 hours	High (11g fibre: 33% chitin 64% insoluble β -glucan cell wall material) or low (3g) fibre lunch			\downarrow hunger at 4 hrs only (P<0.04) \downarrow energy intakes at 4.5 hrs (P<0.001)
Tredger et al (1981)	6 (4/2) 18-21 Healthy	3 hours	MTT ± 20g sugar beet pulp	NSD AUC (#)	NSD AUC (#)	
Turnbull et al (1993)	13 (F) 25 (mean) Healthy	3 hours	Mycoprotein (16.8g fibre) or control (10.1g fibre) meals			
Mixed sour	ces					
Burton-Freeman et al (2002)	15 (7/8)	6 hours	High (20g/4.18MJ) or low (7g/4.18MJ) fibre mixed meal	↓ AUC (P<0.005)	NSD AUC (#)	↓ hunger (in women only; P<0.05) ↑ CCK AUC (P<0.0002)
Fraser et al (1983)	25 (F) Pregnant	24 hours	High (51.4g) or low (12.4g) fibre mixed meals	Ψ mean at 29 (P<0.001) but not at 35 (#) weeks gestation	Ψ mean at 35 P<0.0001) but not at 29 (#) weeks gestation	
Scalfi et al (1987)	7 (M) 30 (mean)	6 hours	High (26g, 9.7g cellulose) or low (8g, 1.4g cellulose) fibre meal	↓ AUC (P<0.05)	↓ AUC (P<0.05)	↓ DIT (P<0.05)
	Healthy		Low (8g, 1.4g cellulose) fibre meal \pm 6g glucomannan	↓ AUC (P<0.05)	↓ AUC (P<0.05)	↓ DIT (P<0.01)
Sparti et al (2000)	14 (7/7) 20-30 Healthy	24 hours	High (60g) or low (3g) fibre mixed meals			↑ fullness post-meals (#) ↓ carbohydrate oxidation post-meals (P<0.05), but NSD 24 hr substrate oxidation (#)
Isolated po	lysaccharid	e sourc	ces	1	1	
Bergmann et al (1992)	12 (3/12) 18-65 Healthy	6 hours	Psyllium or placebo immediately before MTT			

Reference	Subjects N (M/F) Age Health	Durati on	Intervention	Glucose	Insulin	Other
Burley et al (1987)	20 (F) Healthy	2.5 hours	MTT ± guar gum (12.5 or 3.0g fibre)			NSD fullness or later energy intakes (#)
Di Lorenzo et al (1988)	9 (1/8) 24-53 Healthy	2 hours	MTT + 10g pectin (P) or 10g methylcellulose (M)			↓ rate gastric emptying P vs M (P<0.001) ↑ satiety (P<0.001) & time to next meal (P<0.05) P vs M
Durrant & Royston (1978)	13 Overweight		0.84 MJ preload \pm 1 g methylcellulose			NSD hunger (#)
Ebihara et al (1981)	7 (M) 22-32 Overweight	3 hours	OGTT ± 5g konjac-mannan fibre	NSD AUC (#); ↑ 180 min (P<0.01)	↓ AUC (P<0.05) & peak (P<0.05)	
Edwards et al (1987)	16 (12/4) 18-25 Healthy	2.5 hours	OGTT ± 2.5g various viscous polysaccharides (guar (G), xanthan (X), locust bean gum (LBG), meyproydn (M), X/LBG, X/M)		↓ AUC X (P<0.01), G (P<0.001) & X/LBG (P<0.01)	No correlation between <i>in vitro</i> - determined viscosity & AUC
Ellis et al (1985)	9 (4/5) 23-54 Healthy	2 hours	Bread ± guar: 5% (3.2g), 10% (5.9g), 15% (7.7g)			↑ satiety at 0-60 min post- consumption with 10 & 15% (P<0.05), at 120 min with 15% only (P<0.05), NSD at any time with 5% (#)
Fairchild et al (1996)	10 (3/7) 22 (mean) Healthy	4 hours	Guar-enriched (5.5g soluble, 3.6g insoluble fibre) or control wheatflakes (1.0g soluble, 3.0g insoluble fibre)	↓ AUC to 60 and 120 (P<0.05) but not 240 min (#)	↓ AUC to 60, 120 and 240 min (P<0.05)	NSD AUC C-peptide (#)
French & Read (1994)	8 (M) 22-30 Healthy	1 hour	High/low fat MTT ± 12g guar gum			 ↓ rate gastric emptying with guar for both high & low fat meals (P<0.05) ↑ time to return of hunger with guar for high fat meal only (P<0.05)
Frost et al (2003)	10 (4/6) 34 (mean) Healthy	4 hours	High/low fat MTT ± 1.7g psyllium	Ψ AUC for high fat meal (P=0.004), NSD low fat meal (#)	NSD AUC (#)	NSD GLP-1 AUC (#)
Geleva et al (2003)	33 20-75 Healthy		LMTT \pm 5g solubilised cellulose gel (74.5% cellulose, 23% lignin, 1.5% hemicellulose)	NSD AUC (P=0.22) & peak (P=0.51)		↓ CCK AUC (P=0.08) & peak (P=0.01)
Greundel et al (2006)	20 (9/11) 22-62 Healthy	6 hours	Liquid meals enriched with 0, 5, 10 or 20g carob fibre (68.4% insoluble)	NSD AUC (#)	NSD AUC (#)	 ↓ ghrelin response (10g: P=0.021, 20g P=0.046) ↓ glucose oxidation (P<0.001) ↓ triglycerides (P<0.001) ↑ energy expenditure (P<0.001)
Groop et al (1986)	10 (3/7) 40-55 Healthy	7 hours	5g guar gum or placebo before meals	NSD AUC (#)	NSD AUC (#)	↑ AUC C-peptide & GIP (both P<0.05) NSD glucagon (#)
Heini et al (1998)	25 (F) 46 (mean) Healthy		LMTT \pm 8g partially hydrolysed guar gum	NSD AUC (#)	NSD AUC (#)	↑ CCK peak (P<0.01)
Hoad et al (2004)	12 (3/9) 19-29 Healthy	4 hours	LMTT ± 3g guar / weakly-gelling alginate (WGA) / strongly-gelling alginate (SGA)			NSD gastric emptying (P=0.51) ↑ fullness at same gastric volume (all P<0.05) ↓ hunger at 4 hrs with SGA (P=0.041), guar & WGA NSD (#)

Reference	Subjects N (M/F) Age Health	Durati on	Intervention	Glucose	Insulin	Other
Innami et al (2005)	7 (M) 21-25 Healthy	2 hours	OGTT ± 6g or 15g Jew's mellow leaves powder (viscous)	↓ peak (P<0.05)	NSD peak (#)	
Jarjis et al (1984)	16 (11/5) 20-30 Healthy	2 hours	OGTT ± 2.5g or 14.5g guar gum	↓ 30 & 60 min (both P<0.05)	↓ 30 (P<0.01) & 60 (P<0.05) min	NSD gastric emptying
			OGTT ± 3.5g or 7g psyllium	NSD AUC (#)	NSD AUC (#)	
Jenkins et al (1977)	13 (11/2) 19-33	2 hours	LMTT ± 14.5g guar flour	↓ 30 min (P<0.05)	NSD any time point (#)	
	Healthy		MTT ± 10g pectin	↓ 15 min (P<0.01)	↓ 15, 30 (P<0.01) & 45 (P<0.05) min	
			MTT \pm 14.5g guar flour + 10g pectin	↓ 15 (P<0.002) & 30 min (P<0.01)	Ψ all time points (P<0.05)	
Krotkiewski (1984)	9 (F) 30-57 Obese	2 hours	MTT ± 10g guar gum	↓ 30 min (P<0.05)		No further suppression by guar gum after 8 weeks supplementation when excluding outliers
Lavin et al (1995)	10 (M) 21-32 Healthy	3 hours	OGTT ± 5g guar gum	↓ AUC (P<0.004) & peak (P<0.05)	↓ AUC (P<0.0001) & peak (P<0.003)	 ↓ hunger (P<0.002) & ↑ satiety (P<0.029) & fullness (P<0.003) over 3 hrs post-consumption NSD gastric emptying (#)
Leclere et al (1994)	6 (3/3) 21-39 Healthy		Oral glucose (G) or starch (S) + high or low (control) viscosity guar gum	G: ↑ 120-180 min (P<0.05); S: ↓ 45 min (P<0.05)	G: ↓ 45-90 min (P<0.05); S: ↓ 45-75 min (P<0.05)	
Leeds et al (1978)	5 (F) 21-25 Healthy	2 hours	OGTT ± 12g guar gum			No H ₂ production after either meal meaning glucose was not malabsorbed (indicating prevention of glucose absorption by guar is not the mechanism by which it causes \downarrow postprandial glycaemia)
Levitt et al (1980)	12 (M) Healthy	4 hours	MTT \pm 5g guar + 5g pectin	↓ AUC (P<0.05)	NSD AUC (#)	↑ AUC glucagon (P<0.05) NSD AUC GIP (#)
Maki et al (2007a)	31 (6/25) 18-64 Healthy	3 hours	MTT with 8g cellulose (control) or 4g high-viscosity hydroxypropylmethycellulose (HV-HPMC) + 4g cellulose or 8g HV-HPMC	\downarrow AUC (P<0.05) & peak (P<0.001) both doses	\downarrow AUC (P<0.05) & peak (P<0.01) both doses	
Maki et al (2007b)	49 (24/25) 18-64 Healthy	2 hours	OGTT + 1 or 2g high-viscosity (HV) hydroxypropyl- methylcellulose (HPMC) or 2g ultra-high-viscosity (UHV) HPMC or 4g medium-viscosity (MV) HPMC	NSD AUC (#); \downarrow peak UHV only (P<0.001)	Ψ AUC (P<0.001) & peak (#) with all formulations	
Mattes (2007)	25 (10/15) 27 (mean) Healthy	5 hours	High fibre (4.5g from guar + alginate) or placebo meal replacement bar			NSD hunger (#) NSD later energy intake (#)
Morgan et al	6 (M)	3	MTT ± 10g galactomannan (guar gum)	↓ AUC (P<0.02)	↓ AUC (P<0.02)	↓ AUC GIP (P<0.05)
(1990)	20 (mean)	hours	MTT ± 5g glucomannan (konjac-mannan)	NSD AUC (#)	↑ AUC (P<0.02)	NSD AUC GIP (#)
	Healthy		MTT ± 10g sugar beet fibre		NSD AUC (#)	\uparrow AUC GIP (P<0.05)
1	1		MIII ± 10g soy cotyleaon fibre	INSD AUC (#)	个 AUC (P<0.02)	NSD AUC GIP (#)

Reference	Subjects N (M/F) Age	Durati on	Intervention	ntervention Glucose Ins		Other
Rigaud et al (1998)	Health 14 (7/7) 18-50 Healthy	3 hours	7.4g psyllium or placebo preload to MTT	7.4g psyllium or placebo preload to MTT ψ AUC (P<0.05)		NSD gastric emptying (#) ↓ hunger (P<0.05) ↓ later energy intake (total: P<0.05, snacks: P<0.02, dinner; NSD #)
Sanaka et al (2007)	10 (M) 21-33 Healthy	2 hours	MTT ± 2.6g pectin or 2.5g agar	NSD AUC (#)		
Sels et al (1992)	11 (5/6) 18-34 Healthy	2 hours	TT ± 8.1g guar NSD AUC (#)			NSD C-peptide AUC (#)
Sierra et al	10 (F)	2	OGTT ± 10.5g guar gum	NSD AUC (#)	↓ AUC (P<0.05)	
(2001)	30-48 Healthy	hours	OGTT ± 10.5g psyllium	↓ AUC (P<0.05)	↓ AUC (P<0.05)	
Tiwary et al (1997)	74 (49/25) 18-53 Healthy		LMTT ± 5, 10, 15 or 20g pectin			 ↑ satiety (P<0.001 at all time points), NSD between doses (#) ↑ satiety after 2nd fixed meal (P≤0.014 at all time points)
Tomlin (1995a; 1995b)	17 (10/7) 21-31 Healthy	26 hours	Liquid fibre (8.5g ethyl hydroxy ethyl cellulose) or placebo drink			↓ hunger at 0.5-1 hour (P<0.01) ↑ time to next meal (P≤0.01) NSD next day energy intake (P=0.13)
Van den Ven (1994)	24 (F) 24-40 Healthy	24 hours	5 / 10g guar gum or placebo preload 1 hr before meal			\downarrow energy intake at both doses (preload + meal; P<0.001) NSD 24 hr energy intake (#)
Wilmshurst & Crawley (1980)	7 36 (mean) Healthy	8 hours	MTT ± 2g guar gum			\downarrow rate gastric emptying (P<0.05) Delayed return of hunger (P<0.01)
Wolf et al (2003)	30 (13/17) 18-75 Healthy	3 hours	MTT ± 5g guar gum	↓ AUC (P<0.01)		
Wolever et al (1991)	9 (4/5) Healthy	1.5 hours	Branflakes (15.8g fibre) ± psyllium: 5, 10, 15, 20% (17.1, 21.3, 23.4, 24.3g fibre)	↓ AUC dose- response (P<0.002)		No effect of psyllium before cereal (#)
Resistant sta	rch					
Behall & Hallfrisch (2002)	25 (13/12) 23-58 Healthy	3 hours	30-70% amylose bread (2.0-13.4g RS) (10% increments)	\downarrow AUC with \uparrow % amylose (P<0.0001)	\downarrow AUC with \uparrow % amylose (P<0.0001)	
Behall et al (2006)	20 (F) 43 (mean) 10 normal wt 10 overweight	4 hours	MTT: glucose control & muffins with 0.9/3.4/6.5 g RS (at levels of 0.3/0.9/3.7g β -glucan)	$↓$ AUC with \uparrow RS (P<0.05)	↓ AUC with ↑ RS (P<0.0001)	
Brighenti et al (2006)	10 (8/2) 40 (mean) Healthy	8.5 hours	High (13g) or control low (1g) RS breakfast, followed by standardised lunch at 5 hrs	↓ 30 min peak (P<0.03)	↓ 1 hr peak (P<0.02) & 2 hr P<0.05)	\downarrow glucose response to lunch (P<0.05 at 4 hr post-lunch) NSD insulin response to lunch (#)
Granfeldt et al (1995)	9 (4/5) Healthy	3 hours	45g starch from high amylose corn (16g RS) or dent corn arepas (control; 1.8g RS)	↓ at 30, 45 & 70 min (P<0.05)	↓ at 30 & 45 min (P<0.05)	
Heijenen et al (1995)	10 (M) 20-26 Healthy	5 hours	50g raw (54% RS2) or pregelatinised potato starch (control; 100% digestible)	↓ peak (P<0.0001)	↓ peak (P<0.01)	↓ DIT (P<0.05)

Reference	Subjects	Durati	Intervention	Glucose	Insulin	Other
	Health					
Leeman et al (2005)	13 (3/10) 19-32 years Healthy	2 hours	50g starch from boiled & cooled (5.2% RS) or freshly boiled (control; 3.3% RS) potatoes	↓ 1 hr AUC (P<0.05); NSD 2 hr AUC (#)	NSD 1 hr AUC (#); ↓ 2hr AUC (P<0.05)	
Liljeberg et al (1999)	10 (4/6) 22-57 Healthy	7 hours	50g starch from white bread + raw potato starch (9.7g RS) or white bread (control; 0.1g RS) for breakfast, followed by standardised lunch at 4 hrs	NSD AUC (GI 92 vs 100; #)	NSD AUC (II 92 vs 100; #)	NSD glucose response to lunch (#) NSD insulin response to lunch (#)
Meance et al (1999)	8 (4/4) 19-43 Healthy		Non-extruded (5.4% RS) or extruded (control; 1.1%) corn flour-based porridge			NSD satiety scores (AUC marginally higher extruded vs non-extruded; P=0.06)
Raben et al (1994a)	10 (M) Healthy		50g raw (54% RS2) or pregelatinised potato starch (control; 100% digestible)	↓ AUC (P=0.0015)	↓ AUC (P=0.0008)	 ↓ AUC GIP (P=0.03) & GLP-1 (P=0.06) ↑ satiety (P=0.03) & fullness (P=0.012)
Ranganathan et al (1994)	7 (M) 23-26 Healthy	12 hours	RMR measured then 30g cellulose / pectin / lintner (RS) consumed	NSD AUC (#)	NSD AUC (#)	
Robertson et al (2003)	10 (4/6) 23-65 Healthy	24 hours	Low fibre meals + 100g high amylose maize starch (60g RS + 40g digestible starch) or 40g waxy maize starch (40g digestible starch) then fibre-free MTT	Ψ AUC following MTT (P=0.037)	Ψ AUC following MTT (P=0.038)	\uparrow C-peptide: insulin AUC (#) NSD HOMA-IR (#) / HOMA-β (#)
Tagliabue et al (1995)	15 (M) 20-31 Healthy	5 hours	50g raw (54% RS2) or pregelatinised potato starch (control; 100% digestible)			 ↓ DIT (P=0.008) ↑ fat oxidation (P=0.013) ↓ carbohydrate oxidation (<0.0005)
Yamada et al (2005)	20 (9/11) 50 (mean) 12 hyper- glycaemic, 8 normal	2 hours	Bread with 6g RS or control bread (0g RS)	NSD AUC (#)	↓ AUC (P<0.05)	Subdivision of group: \forall AUC insulin in high fasting glucose group (P<0.05) but NSD in normal group (#)
Oligosacchai	rides/inulin					
Archer et al (2004)	33 (M) 37-64 Healthy	4.5 hours	Sausage patty + 24g inulin / lupin kernal fibre (LKF)			Ψ AUC satiety inulin vs LKF (P<0.05)
van Dokkum et al (1999)	12 (M) 23 (mean) Healthy		50g glucose + 5g inulin / FOS / GOS / nothing added	NSD (#)	NSD (#)	
Polydextrose	2					
King et al (2005)	16 (8/8) 30 (mean) Healthy	1.5 hours	Yogurt + 25g polydextrose (P) / 12.5g polydextrose + 12.5g xylitol (PX) / 25g xylitol (X) / 10g sucrose (control)			 ↑ fullness following XP only (P=0.003) ↓ energy intake following P only after accounting for differing energy content of yoqhurts (P<0.001)

AUC	Area Under the Curve
CCK	Cholecystokinin
DIT	Diet-Induced Thermogenesis
GI	Glycaemic Index
GIP	Gastric Inhibitory Polypeptide
GLP-1	Glucagon-Like Peptide-1
HOMA-IR	Homeostasis Model Assessment of Insulin Resistance
ΗΟΜΑ-β	Homeostasis Model Assessment of beta-cell function
II	Insulinaemic index
MTT	Meal Tolerance Test
NSD	No significant difference
OGTT	Oral Glucose Tolerance Test
RMR	Resting Metabolic Rate
RS	Resistant Starch
#	P value not given

Appendix B: list of cardiovascular disease tables

Table 1B: Prospective studies investigating the association of fibre and cardiovascular disease risk Table 2B: Cross-sectional studies investigating the association of fibre with serum lipid outcomes Table 3B: Prospective studies investigating the association of fibre with blood pressure Table 4B: Cross-sectional studies investigating the association of fibre with blood pressure Table 5B: Intervention trials investigating the effect of oat products on cholesterol levels Table 6B: Effect of oat intake on lipid outcomes in weight loss studies Table 7B: Intervention trials investigating the effect of psyllium on cholesterol levels Table 8B: Effect of psyllium supplementation on lipid outcomes in weight loss studies Table 9B: The effect of pectin on cholesterol levels Table 10B: Intervention trials investigating the effect of fibre from barley on cholesterol levels Table 11B: Intervention trials investigating the effect of isolated polysaccharides on cholesterol levels Table 12B: Intervention trials investigating the effect of fibre from wheat on cholesterol levels Table 13B: Intervention trials investigating the effect of fibre supplement mixtures on cholesterol levels Table 14B: Intervention trials investigating the effect of legumes on cholesterol levels Table 15B: Intervention trials investigating the effect of resistant starch on cholesterol levels Table 16B: Intervention trials investigating the effect of oligosaccharides and inulin on cholesterol levels Table 17B: Intervention studies investigating the effect of dietary fibre on blood pressure Table 18B: Effect of fibre supplementation weight studies on blood pressure in loss

Table 1B: Prospective studies investigating the association of fibre and cardiovascular disease risk

Reference	Subject Population	Measure of Exposure	Fibre intake/level	Adjusted Relative Risk or Odds Ratio	P trend	Factors adjusted for in analysis
Jensen et al (2004) Health Professionals Follow-Up Study	42,850 men 1818 cases of coronary heart disease 40-75 years 14 y follow-up	Total dietary fibre according to quintile of mean whole-grain intake by FFQ	Quintile mean fibre intake (g/d) Total fibre Q1: 17 Q2: 19 Q3: 21 Q4: 22 Q5: 26	Hazard Ratio 1.00 (ref) 0.97 (0.84, 1.11) 0.94 (0.82, 1.09) 0.86 (0.74, 1.01) 0.82 (0.70, 0.96)	0.01	Age, energy intake, smoking, alcohol intake, physical activity, family history of myocardial infarction, vitamin E use, intakes of fats (saturated, polyunsaturated, and trans fats), fruit, vegetables and fish.
Liu et al (2002a) Women's Health Study	38,480 women 570 cases of cardiovascular disease 46-64 years 6 y follow-up	AOAC defined total dietary fibre by FFQ	Quintile mean fibre intake (g/d) Total fibre Q1: 12.5 Q2: 15.7 Q3: 18.2 Q4: 21.1 Q5: 26.3	1.00 (ref) 0.81 (0.61, 1.07) 0.85 (0.64, 1.12) 0.78 (0.57, 1.05) 0.79 (0.58, 1.09)	0.17	Age, smoking, exercise, alcohol intake, use of postmenopausal hormones, body mass index, use of multivitamin or vitamin C supplements, history of hypertension, high cholesterol or diabetes mellitus, parental history of myocardial infarction before age 60, energy intake.
			Cereal fibre Q1: 3.0 Q2: 3.8 Q3: 4.4 Q4: 5.0 Q5: 6.5	1.00 (ref) 1.00 (0.75, 1.33) 1.09 (0.82, 1.45) 1.08 (0.81, 1.43) 1.11 (0.84, 1.46)	0.38	
			Vegetable fibre Q1: 5.9 Q2: 6.4 Q3: 6.8 Q4: 7.2 Q5: 8.0	1.00 (ref) 0.89 (0.67, 1.18) 1.10 (0.83, 1.44) 1.07 (0.81, 1.41) 0.96 (0.72, 1.28)	0.78	
			Fruit fibre Q1: 2.5 Q2: 3.5 Q3: 4.2 Q4: 4.9 Q5: 6.0	1.00 (ref) 0.94 (0.72, 1.24) 1.10 (0.85, 1.44) 0.80 (0.60, 1.06) 0.82 (0.61, 1.09)	0.09	

Reference	Subject Population	Measure of Exposure	Fibre intake/level	Adjusted Relative Risk or Odds Ratio	P trend	Factors adjusted for in analysis
Liu et al (2002a) continued			Soluble fibre Q1: 3.7 Q2: 4.8 Q3: 5.6 Q4: 6.5 Q5: 8.6	1.00 (ref) 0.99 (0.75, 1.32) 1.02 (0.77, 1.35) 0.99 (0.74, 1.31) 0.90 (0.68, 1.21)	0.50	
			Insoluble fibre Q1: 9.5 Q2: 12.3 Q3: 14.2 Q4: 16.5 Q5: 21.8	1.00 (ref) 0.92 (0.62, 1.03) 0.75 (0.56, 1.00) 0.78 (0.59, 1.00) 0.78 (0.57, 1.06)	0.09	
Liu et al (2002b) Women's Health Study	38,480 women Myocardial Infarction 46-64 years 6 y follow-up	AOAC defined total dietary fibre by FFQ	Quintile mean fibre intake (g/d) Total fibre Q1: 12.5 Q2: 15.7 Q3: 18.2 Q4: 21.1 Q5: 26.3	1.00 (ref) 0.70 (0.44, 1.15) 0.66 (0.40, 1.09) 0.58 (0.33, 0.99) 0.68 (0.39, 1.22)	0.13	Age, smoking, exercise, alcohol intake, use of postmenopausal hormones, body mass index, use of multivitamin or vitamin C supplements, history of hypertension, high cholesterol or diabetes mellitus, parental history of myocardial infarction before age 60, energy intake.
			Cereal fibre Q1: 3.0 Q2: 3.8 Q3: 4.4 Q4: 5.0 Q5: 6.5	1.00 (ref) 0.93 (0.58, 1.50) 0.54 (0.31, 0.96) 0.95 (0.59, 1.53) 0.91 (0.56, 1.47)	0.74	
			Vegetable fibre Q1: 5.9 Q2: 6.4 Q3: 6.8 Q4: 7.2 Q5: 8.0	1.00 (ref) 0.87 (0.51, 1.49) 1.39 (0.86, 2.66) 1.21 (0.73, 2.00) 0.89 (0.52, 1.53)	0.87	
			Fruit fibre Q1: 2.5 Q2: 3.5 Q3: 4.2 Q4: 4.9 Q5: 6.0	1.00 (ref) 1.88 (1.14, 3.11) 1.50 (0.89, 2.54) 1.23 (0.71, 2.12) 1.11 (0.62, 1.96)	0.63	

Reference	Subject Population	Measure of Exposure	Fibre intake/level	Adjusted Relative Risk or Odds Ratio	P trend	Factors adjusted for in analysis
Liu et al (2002b) continued			Soluble fibre Q1: 3.7 Q2: 4.8 Q3: 5.6 Q4: 6.5 Q5: 8.6	1.00 (ref) 0.93 (0.57, 1.49) 0.79 (0.47, 1.33) 0.77 (0.45, 1.33) 0.83 (0.47, 1.48)	0.40	
			Insoluble fibre Q1: 9.5 Q2: 12.3 Q3: 14.2 Q4: 16.5 Q5: 21.8	1.00 (ref) 1.04 (0.65, 1.67) 0.79 (0.47, 1.31) 0.71 (0.41, 1.21) 0.74 (0.42, 1.30)	0.12	
Liu et al (2000) Nurses' Health Study	75, 521 women 352 cases of ischemic stroke 38-63 years 12 y follow-up	Total dietary fibre according to quintile of whole- grain servings (median/d) by FFQ	Quintile mean fibre intake (g/d) Total fibre Q1: 14 Q2: 15 Q3: 16 Q4: 18 Q5: 20	1.00 (ref) 0.72 (0.53, 1.00) 0.78 (0.58, 1.08) 0.60 (0.43, 0.86) 0.69 (0.50, 0.98)	0.08	Age, body mass index, physical activity, smoking, alcohol intake, parental history of myocardial infarction at <60y, aspirin use, menopausal status, self-reported hypertension, self- reported high blood cholesterol level, multivitamin use, vitamin E use, saturated fat intake, trans fatty acids, energy intake.
Ludwig et al (1999) Cardia study	2506 men and women Cholesterol levels (mean) 18-30 years 10 y follow-up	AOAC defined total dietary fibre by FFQ	Quintile median fibre intake (g/4184KJ/d) Q1: <5.9 Q5: >10.5	LDL-cholesterol (mmol/l) Adjusted means White Men & Women Q1: 2.93 Q5: 2.81 Black Men & Women Q1: 2.82 Q5: 2.72 HDL-cholesterol (mmol/l) White Men & Women Q1: 1.21 Q5: 1.27 Black Men & Women Q1: 1.34 Q5: 1.36	0.06 0.20 0.005 0.28	Baseline value of respective risk factor (except fibrinogen), age, sex, Cardia field centre, education, energy intake, physical activity, smoking, alcohol, vitamin supplement use.

Reference	Subject Population	Measure of	Fibre intake/level	Adjusted Relative	P trend	Factors adjusted for in
Ludwig et al (1999) Continued	Population			Triglyceride (mmol/l) White Men & Women Q1: 1.0 Q5: 0.91 Black Men & Women Q1: 0.79	0.05	
				Fibrinogen		
				White Men & Women Q1: 7.76 Q5: 7.29	0.005	
				Black Men & Women Q1: 8.06 Q5: 7.91	0.80	

Pietinen et al (1996b)	21,930 men	Englyst defined	Total fibre	1 00 (ref)		Age, treatment group, smoking,
Alpha-Tocopherol Beta-Carotene Cancer Prevention Study	581 cases of coronary heart disease	FFQ	Q1: 10:1 Q2: 20.7 Q3: 24.3 Q4: 28.3 Q5: 34.8 Soluble fibre	1.00 (161) 0.91 (0.72, 1.15) 0.83 (0.64, 1.06) 0.72 (0.55, 0.93) 0.73 (0.56, 0.95) 1.00 (ref)	0.004	pressure, energy intake, alcohol intake, saturated fatty acid intake, education, physical activity, beta-carotene intake, vitamin C intake, and vitamin E use.
			Q2: 4.7 Q3: 5.4 Q4: 6.2 Q5: 7.4	0.73 (0.58, 0.93) 0.77 (0.61, 0.98) 0.54 (0.41, 0.71) 0.61 (0.46, 0.79)	0.003	
			Insoluble fibre Q1: 12.2 Q2: 15.9 Q3: 18.9 Q4: 22.3 Q5: 27.7	1.00 (ref) 0.98 (0.77, 1.24) 0.86 (0.67, 1.10) 0.74 (0.57, 0.97) 0.75 (0.58, 0.98)	0.01	
			Cereal fibre Q1: 8.8 Q2: 12.8 Q3: 16.0 Q4: 19.9 Q5: 26.3	1.00 (ref) 0.92 (0.72, 1.17) 0.92 (0.72, 1.17) 0.83 (0.64, 1.07) 0.74 (0.57, 0.96)	0.01	
			Vegetable fibre Q1: 2.9 Q2: 3.9 Q3: 4.7 Q4: 5.6 Q5: 7.1	1.00 (ref) 0.86 (0.68, 1.09) 0.89 (0.70, 1.14) 0.66 (0.49, 0.87) 0.88 (0.66, 1.99)	0.08	
			Fruit fibre Q1: 0.7 Q2: 1.5 Q3: 2.4 Q4: 3.4 Q5: 5.3	1.00 (ref) 1.09 (0.84, 1.41) 1.39 (1.06, 1.84) 1.26 (0.92, 1.73) 1.16 (0.80, 1.67)	0.77	
Reference	Subject Population	Measure of Exposure	Fibre intake/level	Adjusted Relative Risk or Odds Ratio	P trend	Factors adjusted for in analysis
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Wolk et al (1999)	68, 782 women	AOAC defined total	Quintile median fibre			Age, study period, body mass
Nurses' Health Study	591 cases of coronary	dietaly libre by FFQ	illake (g/u)			status, aspirin use, multivitamin
	heart disease		Total fibre			supplement use, vitamin E use,
			Q1: 11.5	1.00 (ref)	0.07	physical activity, hypertension,
	37-64 years		Q2: 14.3	0.98 (0.77, 1.24)		parental history of myocardial
			Q3: 16.4	0.92 (0.71, 1.18)		<60 y, alcohol intake,
	10 y follow-up		Q4: 18.8	0.87 (0.66, 1.15)		carbohydrate intake, energy
			Q5: 22.9	0.77 (0.57, 1.04)		Intake.
			Cereal fibre			also adjusted for the other
			01: 2.2	1.00 (ref)	< 0.001	sources of fibre, folate, vitamins
			Q2: 3.1	1.06 (0.85, 1.34)		B6, C, and E, beta carotene and
			Q3: 3.8	0.71 (0.55, 0.93)		magnesium intake.
			Q4: 4.9	0.76 (0.58, 0.99)		
			Q5: 7.7	0.66 (0.49, 0.88)		
			Vegetable fibre			
			Q1: 3.6	1.00 (ref)	0.63	
			Q2: 4.9	1.05 (0.80, 1.37)		
			Q3: 5.9	1.11 (0.82, 1.49)		
			Q4: 7.1	1.10 (0.79, 1.52)		
			Q5: 9.5	1.13 (0.77, 1.64)		
			Q5: 9.5	1.13 (0.77, 1.64)		

Table 2B: Cross-sectional studies investigating the association of fibre with serum lipid outcomes

Reference	Subject Population	Measure of Exposure	Fibre intake/level	Adjusted Relative Risk or Odds Ratio	P trend	Factors adjusted for in analysis
Lairon et al (2005) SU.VI.MAX study	2,532 men 3,429 women 35-60 years Cholesterol (>6.2 mmol/l Vs \leq 6.2 mmol/l)	AOAC defined total dietary fibre by 24-h recalls	Mean (g/d) Men: 21.9 Women: 17.9	1.00 (ref) 0.84 (0.71, 1.01) 0.85 (0.70, 1.02) 0.83 (0.68, 1.02) 0.68 (0.53, 0.87)	0.15	Age, sex, energy intake, tobacco use, carbohydrate intake, saturated fatty acid intake, alcohol intake, leisure- time exercise, intervention supplement.
McKeown et al (2002)	1338 men 1603 women Cholesterol levels 54 years (mean)	Total dietary fibre according to quintile of whole-grain servings/wk (median) by FFQ	Total fibre Q1: 13.8 Q2: 15.0 Q3: 17.3 Q4: 19.0 Q5: 21.2	Total Cholesterol (mmol/l) Q1: 5.20 Q2: 5.14 Q3: 5.24 Q4: 5.18 Q5: 5.09	0.06	Age, sex, energy intake, treatment of hypertension, smoking, alcohol intake, multivitamin use, estrogen use, physical activity, body mass index, % of polyunsaturated fatty acids, intakes of meat, fish, fruit & vegetables.
				HDL-C (mmol/l) Q1: 1.20 Q2: 1.21 Q3: 1.21 Q4: 1.18 Q5: 1.23	0.16	
				LDL-C (mmol/l) Q1: 3.16 Q2: 3.13 Q3: 3.21 Q4: 3.16 Q5: 3.04	0.02	
				TAG (mmol/l) Q1: 1.69 Q2: 1.61 Q3: 1.64 Q4: 1.68 Q5: 1.58	0.07	

Total-C- total cholesterol; HDL-C- high density lipoprotein cholesterol; LDL-C low density lipoprotein cholesterol; TAG- triacylglycerol

Reference	Subject Population	Measure of Exposure	Fibre intake/level	Adjusted Relative Risk or Odds Ratio	P trend	Factors adjusted for in analysis
Newby et al (2007) Baltimore	1516 men Cholesterol levels	Total dietary fibre according to quintile of whole-grain intake (median/d) by 7d	Total fibre Q1: 13.9 Q5: 24.7	Total-C (mmol/l) Q1: 5.71 Q5: 5.49	0.02	Age, sex, energy intake, decade of visit, race, education, vitamin supplement use, smoking, % energy from saturated fat alcohol
Aging	27-88 years	food diaries		HDL-C (mmol/l) Q1: 1.27 Q5: 1.22	0.07	body mass index, lipid-lowering medication, hypercholesterolemia.
				LDL-C (mmol/l) Q1: 3.16 Q5: 2.96	0.04	
				TAG (mmol/l) Q1: 1.23 Q5: 1.16	0.22	
			Cereal fibre Q1: 14.5 Q5: 25.9	Total-C (mmol/l) Q1: 5.73 Q5: 5.44	0.005	
				HDL-C (mmol/l) Q1: 1.23 Q5: 1.25	0.59	
				LDL-C (mmol/l) Q1: 3.13 Q5: 2.99	0.07	
				TAG (mmol/l) Q1: 1.24 Q5: 1.15	0.12	

Total-C- total cholesterol; HDL-C- high density lipoprotein cholesterol; LDL-C low density lipoprotein cholesterol; TAG- triacylglycerol

Reference	Subject Population	Measure of Exposure	Fibre intake/level	Adjusted Relative Risk or Odds Ratio	P trend	Factors adjusted for in analysis
Ascherio et al (1996)	41,541 women	AOAC defined total dietary fibre by	Total dietary fibre (g/d)			Age, BMI & alcohol intake.
Nurses' Health Study	2526 cases of self- reported hypertension	FFQ	Category <10.0 10.0-14.9	1.0 (ref) 1.04 (0.85, 1.28)	0.75	
	38-63 years 4 y follow up		15.0-19.9 20.0-24.9 ≥25.0	0.99 (0.81, 1.22) 1.02 (0.82, 1.27) 1.01 (0.80, 1.29)		
Ludwig et al (1999)	2,731 men & women	AOAC defined total dietary fibre by	Quintile median fibre intake (g/4184KJ/d)	Mean SBP (mmHg) White Men & Women		Age, sex, Cardia field centre, education, energy intake,
Cardia study	18-30 years	FFQ		Q1: 109.1 Q5: 106.9	0.01	physical activity, smoking, alcohol, vitamin supplement
	10 y follow-up		Q1: <5.9 Q5: >10.5	Black Men & Women Q1: 111.6 Q5: 111.5	0.77	use.
				Mean DBP (mmHg) White Men & Women Q1: 72.4 Q5: 69.7	<0.001	
				Black Men & Women Q1: 74.0 Q5: 73.3	0.70	

Table 3B: Prospective studies investigating the association of fibre with blood pressure

SBP- systolic blood pressure; DBP- diastolic blood pressure

Reference	Subject Population	Measure of Exposure	Fibre intake/level	Adjusted Relative Risk or Odds Ratio	P trend	Factors adjusted for in analysis
Lairon et al (2005) SU.VI.MAX study	2,532 men 3,429 women 35-60 years Hypertension	AOAC defined total dietary fibre by 24-h recalls	Mean (g/d) Men: 21.9 Women: 17.9	Total fibre 1.00 (ref) 0.88 (0.73, 1.07) 0.74 (0.60, 0.91) 0.83 (0.66, 1.04) 0.71 (0.54, 0.93)	0.02	Age, sex, energy intake, tobacco use, carbohydrate intake, saturated fatty acid intake, alcohol intake, leisure- time exercise, intervention supplement.
	(defined as systolic BP >140 mm Hg, diastolic BP >90 mm Hg, or use of antihypertensive medication)			Insoluble fibre 1.00 (ref) 0.84 (0.69, 1.01) 0.77 (0.62, 0.95) 0.78 (0.62, 0.98) 0.68 (0.52, 0.89)	0.01	
McKeown et al (2002)	2480 men & women Blood pressure outcomes 54 years (mean)	Total dietary fibre according to quintile of whole-grain servings/wk (median) by FFQ	Total fibre Q1: 13.8 Q2: 15.0 Q3: 17.3 Q4: 19.0 Q5: 21.2	Mean SBP (mmHg) Q1: 124.4 Q2: 123.2 Q3: 123.3 Q4: 122.5 Q5: 123.1	0.38	Age, sex, energy intake, treatment of hypertension, smoking, alcohol intake, multivitamin use, estrogen use, physical activity, body mass index, % of polyunsaturated fatty acids, intakes of meat,
				Mean DBP (mmHg) Q1: 75.6 Q2: 74.4 Q3: 74.6 Q4: 74.7 Q5: 73.8	0.19	fish, fruit & vegetables.
Newby et al (2007) Baltimore Longitudinal Study of Aging	1516 men Blood pressure outcomes 27-88 years	Total dietary fibre according to quintile of whole-grain intake (median/d) by 7d food diaries	Total fibre Q1: 13.9 Q5: 24.7	Mean SBP (mmHg) Q1: 129.2 Q5: 128.3 Mean DBP (mmHg) Q1: 79.8 Q5: 79.2	0.79 0.42	Age, sex, energy intake, decade of visit, race, education, vitamin supplement use, smoking, % energy from saturated fat, alcohol, body mass index, blood pressure-lowering medication, hypertension.

Table 4B: Cross-sectional studies investigating the association of fibre with blood pressure

SBP- systolic blood pressure; DBP- diastolic blood pressure

Reference	Subjects N (M/F) Age Health	Study duration & design	Intervention	Fibre dose	Outcome (mmol/l)	Fibre group	Control group	P
Noakes et al (1996)	23 (10/13) 51y (mean) Hyper-TAG	3 x 4 weeks Crossover No washout period	High-amylose diet Diet high in oat bran Low-amylose diet	87g/d oat bran (women) 121g/d oat bran (men)	Mean ± SD Total-C HDL-C LDL-C TAG	$\begin{array}{c} 6.04 \pm 0.85 \\ 0.88 \pm 0.16 \\ 3.98 \pm 0.69 \\ 2.61 \pm 0.75^* \end{array}$	High- amyloseLow- rabic 6.06 ± 0.71 6.12 ± 0.86 0.91 ± 0.21 0.93 ± 0.23 3.77 ± 0.74 3.74 ± 0.86 3.05 ± 0.87 3.23 ± 1.10	*P<0.02
Pick et al (1996)	8 (M) 45y (mean) Type II DM	2 x 12 weeks Crossover No washout period	Individualized diet plans constructed + Oat bran concentrate bread or White bread	40g/d oat bran concentrate (mean)	Mean ± SE Total-C HDL-C LDL-C TAG	$\begin{array}{c} 4.56 \pm 0.11 \\ 1.04 \pm 0.05 \\ 2.59 \pm 0.12 \\ 2.03 \pm 0.16 \end{array}$	$5.30 \pm 0.11 \\ 0.96 \pm 0.05 \\ 3.36 \pm 0.12 \\ 2.14 \pm 0.16$	P < 0.01 NS P < 0.01 NS
Gerhardt et al (1998)	44 (23/21) 51 y (mean) Mod Hyper- chol	6 weeks Parallel	Low fat diet + Rice starch Rice bran or Oat bran	84g/d of each product added to usual low fat diet	Mean ± SD Total-C HDL-C LDL-C TAG	$6.25 \pm 0.58 \\ 1.33 \pm 0.33 \\ 4.19 \pm 0.50 \\ 1.61 \pm 1.03$	Rice starchRice bran 7.20 ± 0.78 6.34 ± 0.76 1.30 ± 0.33 1.43 ± 0.28 5.02 ± 0.56 4.26 ± 0.74 2.04 ± 0.87 1.41 ± 0.97	NS NS NS NS
Onning et al (1999)	52 (M) 62 y (mean) Mod Hyper- chol	2 x 5 weeks Crossover 5 week washout	Oat milk Rice milk 0.25ldrunk 3 x daily. Substitution of test beverages for usual beverages	Oat milk: 3.8g/d ß-glucan Rice milk:0.15g/d ß- glucan	Mean ± SD Total-C HDL-C LDL-C TAG	$6.25 \pm 0.67 \\ 1.37 \pm 0.33 \\ 4.14 \pm 0.56 \\ 1.67 \pm 0.67$	$6.58 \pm 0.80 \\ 1.39 \pm 0.34 \\ 4.38 \pm 0.82 \\ 1.85 \pm 0.92$	P < 0.005 NS P < 0.036 NS
Lovegrove et al (2000)	62 (31/31) 35-70y Mod Hyper- chol	8 weeks Parallel	Habitual diet + Oat bran concentrate or Wheat bran (control)	20g/d of each product consumed with low-fat yoghurt or low-fat milk Oat bran: 3g/d ß-glucan	Mean ± SD Total-C HDL-C LDL-C TAG	$6.3 \pm 1.1 \\ 1.4 \pm 0.5 \\ 4.2 \pm 0.8 \\ 1.5 \pm 0.8$	$\begin{array}{c} 6.5 \pm 0.8 \\ 1.5 \pm 0.7 \\ 4.3 \pm 0.8 \\ 1.7 \pm 0.8 \end{array}$	NS NS NS NS
Davy et al (2002)	36 (M) 50-75y Overweight	12 weeks Parallel	Habitual diet + Oat cereal or Wheat cereal	136g/d oat cereal (5.5g/d ß- glucan). Both cereals provided 14g/d dietary fibre.	Mean ± SE Total-C HDL-C LDL-C TAG	$5.15 \pm 0.21 \\ 0.86 \pm 0.05 \\ 3.49 \pm 0.14 \\ 1.71 \pm 0.18$	$5.22 \pm 0.18 \\ 0.85 \pm 0.05 \\ 3.57 \pm 0.14 \\ 1.83 \pm 0.17$	NS NS P = 0.02 NS

Table 5B: Intervention trials investigating the effect of oat products on cholesterol levels

Reference	Subjects N (M/F) Age Health	Study duration & design	Intervention	Fibre dose	Outcome (mmol/l)	Fibre group	Control group	P
Keenan et al (2002)	18 44 y (mean) Hypertensive	6 weeks Parallel (pilot study)	Habitual diet + Oat cereal Low fibre cereal	137g/d oat cereal (5.52 g/d ß- glucan)	Mean ± SE Total-C HDL-C LDL-C TAG	$\begin{array}{l} 4.75 \pm 0.15 \\ 1.08 \pm 0.10 \\ 2.96 \pm 0.20 \\ 2.10 \pm 0.67 \end{array}$	5.10 ± 0.17 1.06 ± 0.08 3.38 ± 0.16 1.70 ± 0.24	P < 0.05 NS P < 0.05 NS
Pins et al (2002)	88 (45/43) 48 y (mean) Hypertensive	12 weeks Parallel	Habitual diet + Oat cereal Low fibre cereal	137g/d oat cereal (11.7g/d fibre, 5.52 g/d ß- glucan)	Mean ± SE Total-C HDL-C LDL-C TAG	$\begin{array}{c} 4.68 \pm 0.14 \\ 1.16 \pm 0.04 \\ 3.04 \pm 0.11 \\ 1.95 \pm 0.07 \end{array}$	$5.36 \pm 0.17 \\ 1.12 \pm 0.04 \\ 3.42 \pm 0.12 \\ 2.08 \pm 0.08$	NS NS P < 0.05 NS
Kerckhoffs et al (2003a)	48 (21/27) 18-65y Mild Hyper- chol	3 week control period then 4 week intervention Parallel	ß-glucan bread & cookies or Control bread & cookies	ß-glucan bread & cookies: 5.9g/d ß-glucan (mean)	Mean ± SE Total-C HDL-C LDL-C TAG	$5.85 \pm 0.18 \\ 1.47 \pm 0.08 \\ 3.86 \pm 0.17 \\ 1.13 \pm 0.14$	$\begin{array}{c} 6.04 \pm 0.15 \\ 1.41 \pm 0.09 \\ 4.11 \pm 0.16 \\ 1.12 \pm 0.10 \end{array}$	NS NS NS NS
Kerckhoffs et al (2003b)	25 (10/15) 18-65y Mild Hyper- chol	2 x 2 weeks Crossover 1 week washout	Orange juice enriched with ß-glucan from oats Orange juice with wheat fibre (control)	Orange juice + β- glucan: 5.0g/d β- glucan (mean)	Mean ± SE Total-C HDL-C LDL-C TAG	$5.36 \pm 0.13 \\ 1.28 \pm 0.08 \\ 3.50 \pm 0.14 \\ 1.24 \pm 0.12$	$5.58 \pm 0.13 \\ 1.25 \pm 0.07 \\ 3.77 \pm 0.14 \\ 1.22 \pm 0.09$	P < 0.01 NS P < 0.01 NS
Karmally et al (2005)	152 Hispanic Americans (49/103) 30-70y Mod Hyper- chol	5 week run- in period (NCEP Step 1 diets) then 6 week intervention Parallel	NCEP Step 1 diet + Oat cereal or Corn cereal (control)	90g/d oat cereal:3g/d soluble fibre Corn cereal: 0g/d soluble fibre	Mean ± SD Total-C HDL-C LDL-C TAG	5.15 ± 0.65 0.93 ± 0.26 3.43 ± 0.56 1.71 ± 0.71	$5.21 \pm 0.70 \\ 0.97 \pm 0.23 \\ 3.49 \pm 0.58 \\ 1.63 \pm 0.75$	P = 0.0003 NS P = 0.0007 NS

Reference	Subjects N (M/F) Age Health	Study duration & design	Intervention	Fibre dose	Outcome (mmol/l)	Fibre group	Control group	Р
Chen et al (2006)	110 (44/66) 30-65y	12 weeks Parallel	Habitual diet + Oat bran cereals or Low fibre (wheat	144g/d oat cereal: 8g/d ß- glucan	Mean change (95%CI)			
	, Healthy		cereal)		Total-C	-0.06 (-0.23,0.11)	0 (-0.14, 0.14)	NS
					HDL-C	-0.01 (-0.06, 0.04) -0.05	0.04 (-0.02, 0.09) -0.02	NS
					TAG	(-0.19, 0.09) -0.01 (-0.11, 0.10)	(-0.14, 0.10) -0.05 (-0.20, 0.10)	NS
Naumann et al (2006)	47 (18/29) 18-70y Healthy	3 week run- in period consuming rice starch (placebo) then 5 week intervention	Oat drink enriched with β-glucan or Rice starch (placebo)	500mL oat drink: 5g/d ß-glucan	Mean ± SD Total-C HDL-C LDL-C TAG	5.75 ± 0.58 1.27 ± 0.32 3.67 ± 0.61 1.88 ± 0.95	$5.67 \pm 0.66 \\ 1.40 \pm 0.31 \\ 3.54 \pm 0.71 \\ 1.58 \pm 0.57$	
Queenan et al (2007)	75 (25/50) 22-65y Hyperchol	Parallel 6 weeks Parallel	Oat bran concentrate or placebo (dextrose monohydrate) provided as a powder to be mixed with beverages	12g/d Oat bran concentrate: (6g/d ß-glucan)	Mean change (SE) Total-C HDL-C LDL-C TAG	-0.3 ± 0.1 -0.02 ± 0.02 -0.3 ± 0.1 0.09 ± 0.1	-0.1 ± 0.08 -0.009 ± 0.02 -0.04 ± 0.08 -0.2 ± 0.1	NS NS P = 0.026 P = 0.030
Theuwissen et al (2007)	42 (20/22) 18-65y Mild Hyper- chol	3 x 4 week Crossover 2 week washout	Habitual diet + 81 Muesli with wheat fibre (control) ii) Muesli with ß- glucan or iii) Muesli with ß- glucan & plant stanols	100g/d Muesli with ß-glucan: 5g/d ß-glucan. 100g/d Muesli with ß-glucan & plant stanols: 5g/d ß-glucan & 1.5g/d plant stanols	Mean ± SD Total-C HDL-C LDL-C TAG	6.26 ± 0.96^{-1} 1.45 ± 0.49^{-1} $4.09 \pm 0.91^{*1}$ 1.66 ± 1.36^{-1}	iii) Muesli ß- Control glucan & stanols $6.01 \pm 0.92 \ 6.49 \pm 1.03$ $1.45 \pm 0.52 \ 1.46 \pm 0.46$ $3.89 \pm 0.90 \ 4.31 \pm 0.96$ $1.56 \pm 0.94 \ 1.52 \pm 0.84$	ii) Muesli ß- glucan Vs Control ^P = 0.015 *P = 0.013

Table 6B: Effect of oat intake on lipid outcomes in weight loss studies

Reference	Subjects N (M/F) Age Health	Study duration & design	Intervention	Fibre dose	Outcome (mmol/l)	Fibre group	Control group	P
Saltzman et al (2001)	43 (20/23) 18-75y Overweight/ Obese	2 weeks control weight maintenance diet then 6 week hypocaloric diet (- 4.2MJ/d) Parallel	Hypocaloric diet + oats or Hypocaloric diet without the addition of oats (control)	Hypocaloric diet: 45g/d oats/4.2MJ dietary energy	Mean change (SD) Total-C HDL-C LDL-C TAG	-0.87 ± 0.47 0.09 ± 0.13 -0.6 ± 0.41 -0.36 ± 0.36	-0.34 ± 0.5 -0.04 ± 0.14 -0.2 ± 0.41 -0.22 ± 0.23	P < 0.05 NS P < 0.05 NS
Reyna- Villasmil et al (2007)	38 (M) 59.8 y (mean) Hyperchol	8 weeks Parallel	AHA Step II diet + Oat bread or Wheat bread (control)	Oat bread: 6g/d β-glucan	Mean ± SE Total-C HDL-C LDL-C TAG	5.05 0.11 1.29 0.05 3.14 0.11 1.24 0.08	5.27 0.17 1.08 0.06 3.46 0.14 1.35 0.08	NS P < 0.001 P < 0.04 NS

Reference	Subjects N (M/F) Age Health	Study duration &	Intervention	Fibre dose	Outcome (mmol/l)	Fibre group	Control group	P
Davidson et al (1998)	196 21-80 y Mild Hyper- chol	8 week run- in period (NCEP Step 1 diets) then 24 week intervention Parallel	NCEP Step 1 diet + Randomisation to one of four treatment arms: Control or one of three doses of psyllium provided in food products	81 control ii) psyllium 3.4g/d iii) psyllium 6.8g/d iv) psyllium 10.2g/d	Mean ± SE Total-C HDL-C LDL-C TAG Total-C HDL-C LDL-C TAG	$\begin{array}{c} 3.4g/d\\ 6.15 \pm 0.11\\ 1.25 \pm 0.04\\ 4.17 \pm 0.08\\ 1.63 \pm 0.11\\ 6.8g/d\\ 6.27 \pm 0.12\\ 1.22 \pm 0.04\\ 4.25 \pm 0.10\\ 1.73 \pm 0.14 \end{array}$	$\begin{array}{l} 6.27 \pm 0.11 \\ 1.44 \pm 0.04 \\ 4.39 \pm 0.08 \\ 1.36 \pm 0.11 \end{array}$	NS difference found between the 3 treatment groups
					Total-C HDL-C LDL-C TAG	$10.2g/d 6.14 \pm 0.12^{1} 1.29 \pm 0.04 4.17 \pm 0.10^{*} 1.51 \pm 0.13$		10.2g/d Vs control ^P = 0.02 *P = 0.03
Anderson et al (2000)	248 (130/118) 21-70y Hyperchol	8 week run- in period (AHA Step 1 diets) then 26 week intervention Parallel	AHA Step 1 diet + Psyllium or cellulose (control) provided as powders to be mixed with beverages.	Psyllium 10.2g/d	Mean ± SE Total-C HDL-C LDL-C TAG	5.08 ± 0.05 1.29 ± 0.02 3.86 ± 0.04 3.2 ± 2.3	$5.98 \pm 0.11 \\ 1.20 \pm 0.03 \\ 4.11 \pm 0.10 \\ 1.51 \pm 0.11$	P = 0.001 NS P = 0.001 NS
Wolever et al (2002)	62 (34/28) 63 y (mean) Type II DM	24 weeks Parallel	Low fibre breakfast cereal High fibre breakfast cereal or MUFA diet (cereal forbidden)	Low fibre: 23.1g/d total fibre. High fibre: 50.3g/d total fibre 8.5g/d psyllium MUFA: 23.5g/d total fibre	Mean ± SD Total-C HDL-C TAG	5.06 ± 0.14 0.92 ± 0.03 3.16 ± 0.49	Low-fibreMUFA 5.00 ± 0.25 5.88 ± 0.21 1.03 ± 0.07 1.24 ± 0.09 2.59 ± 0.55 1.95 ± 0.18	Diet x time interaction NS 0.003 NS
Sola et al (2007)	28 (M) 61 y (mean) Ischemic heart disease	2 x 8 weeks Crossover Controlled low-fat diet + treatment 8 week washout	Plantago ovata husk (soluble fibre) Plantago ovata seeds (control/insoluble fibre)	10.5g/d P. Ovata husk 10.5g/d P. Ovata seeds	Mean ± SD Total-C HDL-C LDL-C TAG	5.06 ± 0.67 1.15 ± 0.27 3.26 ± 0.67 1.45 ± 0.83	5.09 ± 0.65 1.06 ± 0.23 3.35 ± 0.61 1.50 ± 0.88	NS P = 0.006 NS NS

Table 7B: Intervention trials investigating the effect of psyllium on cholesterol levels

Table 8B: Effect	of psyllium	supplementation	on lipid outcome	s in weight loss studies
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Reference	Subjects N (M/F) Age Health	Study duration & design	Intervention	Fibre dose	Outcome (mmol/l)	Fibre group	Control group	Ρ
Anderson et al (1999a) Metabolic Ward	29 (M) 30-70y Type II DM & mild hyperchol	2 week diet stabalization then 8 weeks treatment Parallel	Psyllium or cellulose (control) provided as powders to be mixed with beverages.	10.2g/d psyllium	Mean % change (SE) Total-C HDL-C LDL-C TAG	-2.1 ± 2.3 0.6 ± 3.1 -4.7 ± 4.3 6.5 ± 6.8	6.9 ± 2.4 2.0 2.2 8.3 5.3 13.7 7.3	P < 0.05 NS NS NS
Anderson et al (1999b) Outpatient setting					Mean % change (SE) Total-C HDL-C LDL-C TAG	-2.3 ± 2.2 -0.9 ± 3.0 -4.9 ± 2.4 -7.0 ± 13.3	$2.8 \pm 2.3 \\ 8.8 \pm 2.3 \\ 2.8 \pm 3.4 \\ -0.4 \pm 5.3$	NS NS NS NS

Table 9B: The effect of pectin on cholesterol levels

Reference	Subjects	Study	Intervention	Fibre dose	Outcome	Fibre group	Control group	Р
	N (M/F)	duration &			(mmol/l)			
	Age Health	design						
Schwab et al	66 (29/37)	12 weeks	Control	SBP: 16g/d total	Mean ± SD		Control PDX	
(2006)			Polydextrose (PDX)	fibre (76% soluble)				
	30-65y	Parallel	Sugar beet pectin (SBP)	PDX: 16g/d total	Total-C	5.54 ± 0.68	5.32 ± 1.03 5.27 ± 0.82	NS
				fibre	HDL-C	1.19 ± 0.25	$1.17 \pm 0.31 1.32 \pm 0.35$	NS
	Abnormal	Nutrition	Fibre enrichment		LDL-C	3.65 ± 0.77	3.28 ± 0.79 3.29 ± 0.72	NS
	glucose	counselling	consumed as a drink (4		TAG	1.60 ± 0.25	$1.96 \pm 1.46 1.46 \pm 0.71$	NS
	metabolism	provided	dl/d) with meals					

Reference	Subjects N (M/F) Age Health	Study duration & design	Intervention	Fibre dose	Outcome (mmol/l)	Fibre group	Control group	P
Behall et al (2004)	25 (7/18) 47 y (mean) Mild hyperchol	2 weeks run- in period (AHA Step 1 diet) then 5 weeks on each treatment Latin-square	AHA Step 1 diet + Control Med B-glucan High B-glucan	Control: 0g/d Med: 3g/d ß- glucan High: 6g/d ß- glucan	Mean ± SE Total-C HDL-C LDL-C TAG	MedHigh β -glucan5.17 \pm 0.13*5.12 \pm 0.33*1.22 \pm 0.061.22 \pm 0.063.57 \pm 0.13*3.50 \pm 0.13*1.90 \pm 0.222.03 \pm 0.23	$5.44 \pm 0.13 \\ 1.22 \pm 0.06 \\ 3.82 \pm 0.13 \\ 2.02 \pm 0.22$	NS difference between Med & High groups Med & High Vs Control *P < 0.05
Behall et al (2004)	18 (M) 45 y (mean) Mild hyperchol	2 weeks run- in period (AHA Step 1 diet) then 5 weeks on each treatment Latin-square	AHA Step 1 diet + Control Med ß-glucan High ß-glucan	Control: 0g/d Med: 3g/d B- glucan High: 6g/d B- glucan	Mean ± SE Total-C HDL-C LDL-C TAG	MedHigh β -glucanHigh β -glucan 5.18 ± 0.20 $4.85 \pm 0.20^*$ 1.01 ± 0.04 1.04 ± 0.04 3.28 ± 0.20 $3.00 \pm 0.20^*$ 1.94 ± 0.21 1.82 ± 0.21	5.24 ± 0.20 1.06 ± 0.04 3.28 ± 0.21 2.01 ± 0.21	High Vs Control & Med *P <0.05
Keenan et al (2007)	155 (75/80) 55 y (mean) Hyperchol	4 week run- in period (low fat diet) then 6 week intervention 5-arm parallel	Control High molecular weight (HMW) barley ß- glucan (BBG) at 2 doses Low molecular weight (LMW) barley ß- glucan at 2 doses. Provided as breakfast cereal and juice beverage	Control 0g/d BBG HMW 5g/d BBG LMW 5g/d BBG HMW 3g/d BBG LMW 3g/d BBG	Mean ± SD Total-C HDL-C LDL-C TAG Total-C HDL-C LDL-C TAG	HMW 5g/d LMW 5g/d 5.35 ± 0.65 5.50 ± 0.53 1.35 ± 0.33 1.29 ± 0.33 3.43 ± 0.30 3.49 ± 0.33 $1.51 \pm 0.54^*$ 1.65 ± 0.71 HMW 3g/d LMW 3g/d 5.58 ± 0.56 5.69 ± 0.52 1.23 ± 0.29 1.32 ± 0.41 3.61 ± 0.53 3.65 ± 0.39 1.72 ± 0.63 1.61 ± 0.56	Control 6.01 ± 0.70 1.30 ± 0.36 3.92 ± 0.63 1.79 ± 0.73	Compared to control Total-C & LDL-C in all groups (P < 0.05) NS difference between treatment groups *Compared to control & treatment groups P < 0.05

Table 10B: Intervention trials investigating the effect of fibre from barley on cholesterol levels

Table 11B: Intervention trials investigating the effect of isolated polysaccharides on cholesterol levels

Reference	Subjects N (M/F) Age Health	Study duration & design	Intervention	Fibre dose	Outcome (mmol/l)	Fibre group	Control group	Р
Konjac-mai	nnan							
Arvill et al (1995)	63 (M) 47 y (mean) Healthy	2 x 4 weeks Crossover 2 week washout	Habitual diet + glucomannan or cornstarch (placebo) provided as capsules to be taken ½ hour before meals.	3.9g/d glucomannan	Mean ± SD Total-C HDL-C LDL-C TAG	$6.51 \pm 0.85 \\ 1.14 \pm 0.26 \\ 4.28 \pm 0.69 \\ 2.37 \pm 1.59$	$6.72 \pm 0.67 \\ 1.25 \pm 0.39 \\ 4.54 \pm 0.71 \\ 2.88 \pm 2.32$	P = 0.0001 NS P 0.007 P = 0.026
Vuskan et al (2000)	11 (5/6) 55 y (mean) Insulin resistance syndrome	8 week run-in period (NCEP Step 2 diet) then 3 week intervention Crossover 2 week washout	NCEP Step 2 diet enriched with glucomannan or wheat (control)	8-13g/d glucomannan	Mean % change (SE) Total-C HDL-C LDL-C TAG	-19 ± 2.7 1.2 ± 2.2 -29 ± 3.4 10.1 ± 9.9	-6.3 ± 3.4 -9.6 ± 2.2 -6.6 ± 5.0 12.1 ± 14	P = 0.0038 NS P = 0.0017 NS
Wood et al (2007)	29 (M) 38 y (mean) Overweight	12 weeks Parallel	Carbohydrate restricted diet + 6 capsules containing glucomannan or maltodextrin (placebo) taken daily before meals	3.0g/d glucomannan	Mean ± SD Total-C HDL-C LDL-C TAG	$\begin{array}{c} 4.18 \pm 0.63 \\ 1.17 \pm 0.32 \\ 2.61 \pm 0.72 \\ 0.86 \pm 0.39 \end{array}$	$\begin{array}{c} 4.30 \pm 0.92 \\ 1.23 \pm 0.35 \\ 2.72 \pm 0.95 \\ 0.77 \pm 0.35 \end{array}$	NS NS NS NS

NCEP- National Cholesterol Education Program NS- non significant; Total-C- total cholesterol; HDL-C- high density lipoprotein cholesterol; LDL- low density lipoprotein cholesterol; TAG- triacylglycerol

Reference	Subjects N (M/F) Age Health	Study duration & design	Intervention	Fibre dose	Outcome (mmol/l)	Fibre group	Control group	P
Gums								
Blake et al (1997)	11 (M) 44 y (mean) Mod hyperchol	3 x 2 weeks Crossover 4 week washout	Habitual diet + bread products containing guar gum flour or control flour	Guar bread: 17.2g/d NSP (mean) & 13.9g/d guar gum (mean). Control: 16.2g/d NSP	Mean ± SE Total-C HDL-C LDL-C TAG	$5.89 \pm 0.15 \\ 1.23 \pm 0.05 \\ 3.81 \pm 0.12 \\ 1.85 \pm 0.25$	$\begin{array}{c} 6.53 \pm 0.16 \\ 1.31 \pm 0.06 \\ 4.31 \pm 0.14 \\ 1.86 \pm 0.32 \end{array}$	P < 0.001 NS P < 0.001 NS
Maki et al (1999)	154 (102/52) 21-75y Hyperchol	8 week run-in period (NCEP Step 1 diet) then 6 intervention Parallel	NCEP Step 1 diet + Placebo or hydroxypropylmethylcellulose (HPMC) provided as 3 different doses as powder form to be mixed with beverage	HPMC: 2.5g/d 5g/d 7.5g/d	Mean ± SE Total-C HDL-C LDL-C TAG (log ¹⁰) Total-C HDL-C TAG (log ¹⁰) Total-C HDL-C LDL-C TAG (log ¹⁰)	HPMC 2.5g/d 6.09 ± 0.12 1.20 ± 0.05 4.11 ± 0.10 1.85 HPMC 5g/d $5.63 \pm 0.11^*$ 1.19 ± 0.06 $3.68 \pm 0.10^*$ 1.85 HPMC 7.5g/d $5.76 \pm 0.12^*$ 1.28 ± 0.05 $3.85 \pm 0.09^*$ 1.68	5.98 0.11 1.27 0.05 4.04 0.09 1.68	*Compared to placebo P < 0.05

Hyperchol- hypercholesterolemia; NCEP- National Cholesterol Education Program NS- non significant; Total-C- total cholesterol; HDL-Chigh density lipoprotein cholesterol; LDL- low density lipoprotein cholesterol; TAG- triacylglycerol

Reference	Subjects N (M/F) Age Health	Study duration & design	Intervention	Fibre dose	Outcome (mmol/l)	Fibre group	Control group	P
Arabinogala	ctan (AG)							
Robinson et al (2001)	20 Age (not reported) Healthy	7d control period followed by 2 x 3 week intervention Crossover No washout	Control: 0g/d AG Followed by AG intervention incorporated into sweetened beverage	AG 15g/d AG 30g/d	Mean ± SE Total-C HDL-C LDL-C TAG (log ¹⁰) Total-C HDL-C LDL-C TAG (log ¹⁰)	AG $15g/d$ 5.09 ± 0.08 1.35 ± 0.05 3.16 ± 0.08 2.03 ± 0.26 AG $30g/d$ 5.15 ± 0.08 1.26 ± 0.05 3.08 ± 0.08 2.10 ± 0.26		AG had no significant effect on cholesterol levels
Marett et al (2004)	54 29 y (mean) Healthy	24 weeks Parallel	Habitual diet + Rice starch (placebo) Larch AG Tamarack AG provided in foods & beverages	Rice Starch 8.4g/d Larch AG: 8.4g/d Tamarack AG: 8.4g/d	Mean ± SD Total-C HDL-C LDL-C TAG Total-C HDL-C LDL-C TAG	Larch AG 4.52 ± 0.79 1.24 ± 0.25 2.55 ± 0.76 1.15 ± 0.71 Tamarack AG 4.86 ± 0.98 1.35 ± 0.29 2.98 ± 0.92 1.18 ± 0.53	Control 4.32 ± 0.92 1.30 ± 0.22 2.55 ± 0.76 1.01 ± 0.55	AG had no significant effect on cholesterol levels
Arabinoxyla	n (AX)							
Lu et al (2004)	15 (6/9) 60 y (mean) Type II DM	2 x 5 weeks Crossover No washout	Habitual diet + majority of starchy foods replaced with control or AX-rich food products	AX: 14-17g/d NSP	Mean ± SE Total-C HDL-C LDL-C TAG	5.31 ± 0.23 1.04 ± 0.04 3.32 ± 0.21 1.86 ± 0.29	5.33 ± 0.24 1.04 ± 0.05 3.35 ± 0.26 2.06 ± 0.39	NS NS NS NS
Carob pulp				1				
Zunft et al (2003)	58 (25/33) 34-70y Hyperchol	6 weeks Parallel	Bread (2 servings) & fruit bar (1 serving) with or without carob pulp	Carob pulp 15g/d	Mean ± SD Total-C HDL-C LDL-C TAG ¹	$6.86 \pm 0.96 \\ 1.46 \pm 0.47 \\ 4.03 \pm 0.75 \\ 1.91 \\ (0.99, 3.71)$	7.23 ± 0.96 1.48 ± 0.34 4.47 ± 0.78 1.59 (1.16, 2.20)	P=0.001 NS P=0.010 NS NS

¹ Values given as geometric mean and geometric SD Hyperchol- hypercholesterolemia, Type II DM- type 2 diabetes mellitus; NS- non significant;Total-C- total cholesterol; HDL-C- high density lipoprotein cholesterol; LDL- low density lipoprotein cholesterol; TAG- triacylglycerol

Reference	Subjects N (M/F) Age Health	Study duration & design	Intervention	Fibre dose	Outcome (mmol/l)	Fibre group	Control group	P
Stasse- Wolthius et al (1980)	62 (40/22) 18-28 Healthy	2.5 weeks run-in (low- fibre diet) then 5 week intervention 4-arm Parallel	Controlled diet: All foodstuffs provided. Low-fibre High-fibre (fruit & vegetables) Citrus Pectin Wheat bran	Low fibre: 15g/d total fibre High-fibre: 41g/d total fibre Citrus Pectin: 24g/d total fibre; 9g/d citrus pectin Wheat bran: 38g/d bran	Mean change (SD) Total-C HDL-C Total-C HDL-C	High-fibre -0.17 \pm 0.63* 0.01 \pm 0.15 Citrus Pectin -0.34 \pm 0.34* 0.02 \pm 0.18	0.10 ± 0.34 0.01 ± 0.12	*Compared to control P < 0.05 ^Compared
					Total-C HDL-C	Wheat bran 0.34 ± 0.41^ 0.07 ± 0.16		to high-fibre & citrus pectin P < 0.05
Jenkins et al (1999a) Metabolic Study	24 (16/8) 57y (mean) Hyperchol	3 x 4 weeks Crossover 2 week washout	Identical metabolic diets + low-wheat fibre bread (control) or high-wheat fibre bread at 2 particle sizes (PS) (medium or ultra-fine)	Low-fibre: 6g/d total fibre Medium (PS): 25g/d total fibre Ultra-fine (PS): 25g/d total fibre	Mean ± SE Total-C HDL-C LDL-C TAG Total-C HDL-C LDL-C TAG	Medium PS 5.88 ± 0.17 1.01 ± 0.06 4.02 ± 0.12 $2.01 \pm 0.17*$ Ultra-fine PS 5.95 ± 0.16 1.03 ± 0.07 4.07 0.11 $2.06 \pm 0.20*$	5.92 ± 0.19 0.99 ± 0.05 $3.91 \ 0.16$ 2.41 ± 0.21	*Compared to control P < 0.005
Jenkins et al (1999b) <i>Ad libitum</i> study	24 (12/12) 36y (mean) Normolipidemic	3 x 2 weeks Crossover 2 week washout	Habitual diet + low- wheat fibre breakfast cereal (control) or high-wheat fibre cereal at 2 particle sizes (PS) (coarse or medium)	Low-fibre: 2.4g/d total fibre Coarse: 21.5g/d total fibre Medium: 21.5g/d total fibre	Mean ± SE Total-C HDL-C LDL-C TAG Total-C HDL-C LDL-C TAG	Coarse PS 4.85 ± 0.18 1.20 ± 0.07 2.99 ± 0.16 1.45 ± 0.16 Medium PS 4.78 ± 0.18 1.20 ± 0.06 2.97 ± 0.16 1.36 ± 0.16	$4.80 \pm 0.19 \\ 1.21 \pm 0.06 \\ 2.97 \pm 0.17 \\ 1.37 \pm 0.17$	PS had no significant effect on cholesterol levels

Table 12B: Intervention trials investigating the effect of fibre from wheat on cholesterol levels

Hyperchol- hypercholesterolemia, Type II DM- type 2 diabetes mellitus; NS- non significant;Total-C- total cholesterol; HDL-C- high density lipoprotein cholesterol; TAG- triacylglycerol

Reference	Subjects N (M/F) Age Health	Study duration & design	Intervention	Fibre dose	Outcome (mmol/l)	Fibre group	Control group	P
Jenkins et al (2002)	23 (16/7) 63y (mean) Type II DM	2 x 12 weeks Crossover 8 week washout	Habitual diet + Low-wheat fibre or High-wheat fibre provided as cereals & breads	Low-fibre: Additional 4g/d total fibre High-fibre: Additional 19g/d cereal fibre	Mean ± SE Total-C HDL-C LDL-C TAG	$\begin{array}{r} 4.97 \pm 0.17 \\ 1.23 \pm 0.05 \\ 3.00 \pm 0.15 \\ 1.63 \pm 0.20 \end{array}$	$4.87 \pm 0.15 \\ 1.17 \pm 0.03 \\ 2.99 \pm 0.14 \\ 1.53 \pm 0.18$	NS NS NS NS
Lampe et al (1991)	34 (18/16) 27y (mean) Healthy	6 x 3 weeks Crossover	Six controlled formula diets supplemented with Low-fibre Wheat bran (WB) Vegetable fibre (VF) Sugar beet fibre (SBF)	Low-fibre: 0g/d total fibre WB: 10g/d total fibre WB: 30g/d total fibre VF: 10g/d total fibre VF: 30g/d total fibre SBF: 30g/d total fibre	Mean change (SE) ¹ Total-C HDL-C LDL-C TAG Total-C HDL-C LDL-C TAG Total-C HDL-C LDL-C TAG	WB: $30g/d$ -0.05 ± 0.10 -0.06 ± 0.03 0.06 ± 0.08 -0.12 ± 0.06 VF: $30g/d$ -0.24 ± 0.10** -0.03 ± 0.03 -0.07 ± 0.08 -0.30 ± 0.06 SBF: $30g/d$ -0.70 ± 0.16** -0.09 ± 0.03 -0.46 ± 0.14 -0.27 ± 0.08	$\begin{array}{c} -0.13 \pm 0.10 \\ -0.06 \pm 0.03 \\ -0.02 \pm 0.08 \\ -0.12 \pm 0.06 \end{array}$	Soluble fibre (VF & SBF) proved more effective in ↓ serum lipids (Total-C & LDL-C) than insoluble fibre (WB) **Changes significantly greater compared to WB

¹Owing to the number of diets and the inappropriateness of making all possible comparisons, only a few comparisons between diets on changes in serum lipids were made.

Type II DM- type 2 diabetes mellitus; NS- non significant; Total-C- total cholesterol; HDL-C- high density lipoprotein cholesterol; LDL- low density lipoprotein cholesterol; TAG- triacylglycerol

Table 13B: Intervention trials investigating the effect of fibre supplement mixtures on cholesterol	levels
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Reference	Subjects N (M/F)	Study duration &	Intervention	Fibre dose	Outcome (mmol/l)	Fibre group	Control group	Р
Aller et al (2003)	Age Health 53 (19/34) 18-70 y Healthy	design 12 weeks Parallel	81 diets controlled diets: High-fibre Low-fibre	High fibre: 4.11g/d soluble fibre; 25.08g/d insoluble fibre Low-fibre: 1.97g/d soluble fibre; 8.13g/d	Mean ± SD Total-C HDL-C LDL-C TAG	4.9 ± 0.9 1.41 ± 0.4 3.1 ± 0.8 1.02 ± 0.3	5.0 ± 0.9 1.76 ± 0.3 2.9 ± 0.9 1.27 ± 0.8	P < 0.05 NS P < 0.05 NS
Davidson et al (1998)	85 (59/26) 21-75 y Hyperchol	8 week run- in period (AHA Step 1 diet) then 12 week intervention 4-arm parallel	AHA Step 1 diet + control or gum rabic & pectin mixture (4:1 ratio) provided in apple juice (240ml/d)	Control: 0g/d Mixture: 5g/d Mixture: 9g/d Mixture: 15g/d	Mean ± SE Total-C HDL-C LDL-C TAG Total-C HDL-C LDL-C TAG Total-C HDL-C LDL-C TAG	5g/d 6.46 ± 0.15 1.46 ± 0.10 4.31 ± 0.14 1.78 0.15 9g/d 6.38 ± 0.17 1.47 ± 0.07 4.19 ± 0.16 2.22 ± 0.20 15g/d 6.66 ± 0.20 1.21 ± 0.07 4.42 ± 0.14 2.23 ± 0.20	$\begin{array}{c} 6.88 \pm 0.26 \\ 1.55 \pm 0.08 \\ 4.58 \pm 0.25 \\ 1.81 \pm 0.17 \end{array}$	NS differences between any groups
Knopp et al (1999)	125 (104/21) 18-70 y Mild hyperchol	9 week run- in period (NCEP Step 1 diet) followed by 15 week intervention	NCEP Step 1 diet + Placebo or fibre mixture (guar gum, pectin, soy fibre, pea fibre & corn bran) provided as powder to be mixed with beverage	Fibre mixture: 20g/d	Mean change (SD) Total-C HDL-C LDL-C TAG	-0.44 ± 0.45 -0.02 ± 0.14 -0.41 ± 0.36 0.01 ± 0.43	-0.03 ± 0.47 -0.02 ± 0.16 0.00 ± 0.41 -0.03 ± 0.50	P < 0.001 NS P < 0.001 NS

Hyperchol- hypercholesterolemia NCEP- National Cholesterol Education Program; AHA-American Heart Association; NS- non significant; Total-C- total cholesterol; HDL-C- high density lipoprotein cholesterol; LDL- low density lipoprotein cholesterol; TAG- triacylglycerol

Reference	Subjects N (M/F) Age Health	Study duration & design	Intervention	Fibre dose	Outcome (mmol/l)	Fibre group	Control group	P
Knopp et al (1999) Extension phase	85 26-69y	36 weeks	All subjects received fibre mixture.	Fibre mixture: 20g/d	Mean change (SD) from baseline Total-C LDL-C	Fibre throughout -0.30 ± 0.68 -0.43 ± 0.50	Placebo to fibre -0.23 ± 0.71 -0.32 ± 0.64	Reductions smaller for those in placebo/fibre
Solum et al (1987)	60 (F) 30-60y Overweight	12 weeks Parallel	Calorie restricted diet (1200kcal/d) with fibre tablet mixture (cereal & citrus fruit) or placebo taken with water	Fibre mixture: 6g/d	Mean (95% CI) Total-C TAG	5.6 (4.7, 6.4) 1.15 (0.9, 1.4)	5.7 (4.1, 6.5) 1.18 (0.93, 1.41)	NS NS

NS- non significant; Total-C- total cholesterol; LDL- low density lipoprotein cholesterol; TAG- triacylglycerol

Reference	Subjects N (M/F) Age Health	Study duration & design	Intervention	Fibre dose	Outcome (mmol/l)	Fibre group	Control group	P
Cobiac et al (1990)	20 (M) 29-65y Mild hyperchol	81 x 4 wee ks Crossover	Substitution of one daily meal with canned spaghetti (control) or baked beans	440g/d baked beans: 19.9g/d NSP	Mean ± SE Total-C HDL-C LDL-C TAG	6.29 0.20 1.22 0.05 4.61 0.19 1.26 0.08	6.32 0.15 1.26 0.05 4.63 0.12 1.25 0.08	NS NS NS NS
Fruhbeck et al (1997)	40 (M) 18-21 y Mild-Mod hyperchol	4 weeks Parallel	Controlled low-fat diet (20% total energy as fat) + control flour (group A) cooked field bean flour (groups B & C) or raw field bean flour (group D) to be mixed with soups, yoghurt and cereal.	90g/d of each type of flour	Adjusted Mean (analysis of covariance) Total-C HDL-C LDL-C TAG Total-C HDL-C LDL-C TAG Total-C HDL-C LDL-C TAG	Group B 5.7^ 1.17* 3.87 1.43 Group C 5.35* 1.17* 3.67* 1.11* Group D 5.28* 1.18* 3.65* 1.01*	Group A 5.81 1.02 4.03 1.67	*Compared to Group A P < 0.001 ^Compared to Group A P < 0.05
Pittaway et al (2006)	47 (19/28) 53y (mean) Healthy	81 x 5 wee ks Crossover	Chickpea- supplemented diet or wheat-supplemented diet (control).	140g/d chickpeas	Mean (95% CI) Total-C HDL-C LDL-C TAG	5.75 (5.40, 6.11) 1.39 (1.27, 1.51) 3.71 (3.41, 4.01) 1.46 (1.15, 1.76)	5.98 (5.62, 6.63) 1.41 (1.29, 1.53) 3.89 (3.58, 4.20) 1.5 (1.28, 1.72)	P < 0.05 NS P < 0.05 NS

Table 14B: Intervention trials investigating the effect of legumes on cholesterol levels

Reference	Subjects N (M/F) Age Health	Study duration & design	Intervention	Fibre dose	Outcome (mmol/l)	Fibre group		Р
Anderson et al (1990)	24 (M) 64y (mean) Hyperchol	81 weeks Parallel	7d control period containing no beans followed by one of three bean diets.	A: 120g/d beans (single dose) B: 120g/d beans (divided dose) C: 162g/d beans (divided dose)	Mean ± SE Total-C HDL-C LDL-C TAG	Group AGroup 7.03 ± 0.50 $6.41 \pm 0.96 \pm 0.05$ $1.05 \pm 0.15 \pm 0.12 \pm 0.47$ $4.40 \pm 0.18 \pm 0.24$ 2.18 ± 0.24 2.18 ± 0.24	B Group C 0.30 7.09 ± 0.53 0.09 0.98 ± 0.11 0.31 4.73 ± 0.54 0.35^* 2.92 ± 0.49	* P < 0.05
Insoluble L	egumes				•			•
Stephen et al (1995)	9 (M) 19-38 v	81 x 3 weeks	Diet low in NSP + same diet with lentils incorporated into	130g/d lentils: 11.8g/d NSP	Mean ± SE (range)	Lentil period	Control period	
	Healthy	Crossover	baked products		Total-C	5.1 0.2 (4.1-5.9)	5.2 0.2 (4.1-5.8)	NS
	,				HDL-C	1.4 0.1 (1.0-2.0)	1.4 0.1 (1.0-2.1)	NS
					LDL-C	3.2 0.2 (2.4-3.9)	3.2 0.1 (2.6-3.8)	NS
					TAG	1.28 0.14 (0.57-1.98)	1.16 0.11 (0.57-1.75)	NS

Reference	Subjects N (M/F)	Study duration &	Intervention	Fibre dose	Outcome (mmol/l)	Fibre group	Control grou	dı	Р
	Age Health	design							
Heijnen et al (1996)	57 (27/33) 24y (mean) Healthy	81 x 3 wee ks Latin-square	Habitual diet + supplements containing glucose (control) raw high-amylose cornstarch (RS ₂) or retrograded high- amylose cornstarch (RS ₃)	RS₂: 30g/d RS₃: 30g/d	Mean ± SE Total-C HDL-C LDL-C TAG Total-C HDL-C LDL-C TAG	RS_{2} 4.61 ± 0.13 1.45 ± 0.05 2.71 ± 0.10 0.98 ± 0.05 RS_{3} 4.61 ± 0.13 1.45 ± 0.04 2.69 ± 0.10 1.03 ± 0.08	4.69 ± 0.14 1.47 ± 0.04 2.72 ± 0.11 1.09 ± 0.07		NS difference between groups
Noakes et al (1996)	23 (10/13) 51y (mean) Hyper-TAG	81 x 4 wee ks Crossover No washout period	High-amylose diet (33% resistant starch) Diet high in oat bran Low-amylose diet	16.5g/d high- amylose (women) 24.4g/d high- amylose (men)	Mean ± SD Total-C HDL-C LDL-C TAG	High-amylose 6.06 ± 0.71 0.91 ± 0.21 3.77 ± 0.74 3.05 ± 0.87	Oat-bran 5.94 0.81 0.92 0.20 3.80 0.83 2.72 0.76*	Low- amylose 6.12 0.86 0.93 0.23 3.74 0.86 3.23 1.10	*P<0.02

Table 15B: Intervention trials investigating the effect of resistant starch on cholesterol levels

Reference	Subjects N (M/F) Age Health	Study duration & design	Intervention	Fibre dose	Outcome (mmol/l)	Fibre group	Control grou	пр	P
Luo et al (1996)	12 (M) 24y (mean) Healthy	2 x 4 weeks Crossover 2 week washout	FOS or sucrose incorporated into cookies eaten as snacks	20g/d FOS	Mean ± SE Total-C HDL-C TAG	3.96 ± 0.22 0.97 ± 0.05 0.83 ± 0.16	3.91 ± 0.17 1.05 ± 0.06 0.72 ± 0.05		NS NS NS
Pedersen et al (1997)	64 (F) 20-36 y Healthy	81 x 4 wee ks Crossover No washout period	Usual spreads replaced with low-fat inulin or control spread	14g/d Inulin	Mean ± SD Total-C HDL-C LDL-C TAG	4.24 ± 0.75 1.38 ± 0.30 2.38 ± 0.67 0.97 ± 0.39	$\begin{array}{c} 4.25 \pm 0.63 \\ 1.37 \pm 0.30 \\ 2.39 \pm 0.56 \\ 0.98 \pm 0.42 \end{array}$		NS NS NS NS
Davidson et al (1998)	21 30-75 y Mod hyperchol	2 x 6 weeks Crossover 6 week washout	NCEP Step 1 diet + food products with or without inulin	18g/d Inulin	Mean ± SE Total-C HDL-C LDL-C TAG	6.07 ± 0.17 1.35 ± 0.07 3.98 ± 0.12 1.61 ± 0.08	$6.28 \pm 0.12 \\ 1.39 \pm 0.07 \\ 4.10 \pm 0.10 \\ 1.70 \pm 0.17$		P < 0.05 NS P < 0.05 NS
Schaafsma et al (1998)	30 (M) 48y (mean) Healthy	2 x 3 weeks Crossover 1 week washout	Test milk containing FOS or a reference product (control milk) consumed daily with main meals	9.4g/d FOS	Mean Total-C HDL-C LDL-C TAG	5.17 1.21 3.33 1.52	5.41 1.23 3.52 1.57	SE of difference between means 0.058 0.026 0.059 0.105	P < 0.001 NS P < 0.005 NS
Brighenti et al (1999)	12 (M) 23y (mean) Healthy	4 weeks Sequential No washout	50g/d breakfast cereal (control) followed by same cereal containing inulin	9g/d Inulin	Mean ± SE Total-C HDL-C LDL-C TAG	$3.89 \pm 0.19 \\ 1.30 \pm 0.08 \\ 2.31 \pm 0.21 \\ 0.61 \pm 0.05$	$\begin{array}{c} 4.10 \pm 0.23 \\ 1.23 \pm 0.06 \\ 2.48 \pm 0.20 \\ 0.84 \pm 0.07 \end{array}$		NS NS NS P < 0.05
Jackson et al (1999)	27 52y (mean) Mod hyper- TAG	8 weeks Parallel	Inulin and placebo (maltodextrin) provided as powder to be added to beverages	10g/d Inulin	Mean ± SD Total-C HDL-C LDL-C TAG	5.90 ± 0.97 1.31 ± 0.33 4.00 ± 0.85 1.29 ± 0.35	$6.46 \pm 0.91 \\ 1.31 \pm 0.39 \\ 4.43 \pm 1.08 \\ 1.59 \pm 0.58$		NS NS NS P < 0.05

Table 16B: Intervention trials investigating the effect of oligosaccharides and inulin on cholesterol levels

Hyperchol- hypercholesterolemia NCEP- National Cholesterol Education Program; FOS- fructo-oligosaccharide; Total-C- total cholesterol; HDL-C- high density lipoprotein cholesterol; LDL- low density lipoprotein cholesterol; TAG- triacylglycerol; NS- non significant

Reference	Subjects N (M/F) Age Health	Study duration & design	Intervention	Fibre dose	Outcome (mmol/l)	Fibre group	Control group	P
van Dokkum et al (1999)	12 (M) 23y (mean) Healthy	4 x 3 weeks Latin Square	Controlled basal diet supplemented with inulin, FOS, GOS or placebo	15g/d Inulin 15g/d FOS 15g/d GOS	Mean ± SD Total-C HDL-C LDL-C TAG Total-C HDL-C LDL-C TAG Total-C HDL-C LDL-C LDL-C TAG	Inulin 4.51 ± 0.56 1.16 ± 0.22 2.81 ± 0.50 1.31 ± 0.58 FOS 4.51 ± 0.62 1.14 ± 0.20 2.82 ± 0.61 1.29 ± 0.51 GOS 4.58 ± 0.78 1.11 ± 0.20 2.87 ± 0.67 1.46 ± 0.66	$4.56 \pm 0.62 \\ 1.14 \pm 0.22 \\ 2.82 \pm 0.51 \\ 1.40 \pm 0.68$	NS differences between any groups
Kruse et al (1999)	8 26-53 y Healthy	8d run-in consuming 'Western' diet (45% energy as fat, 40% carbohydrate) followed by low-fat diet for 64d	Reduced fat diet (30% energy as fat, 55% as carbohydrate) with inulin incorporated into yoghurt	22-34g/d depending on energy needs	Mean ± SD Total-C HDL-C LDL-C TAG	5.0 ± 0.5 1.2 ± 0.2 3.4 ± 0.4 1.2 ± 0.3	5.3 ± 0.8 1.1 ± 0.2 3.6 ± 0.6 1.2 ± 0.2	NS NS NS NS
Causey et al (2000)	12 (M) 27-49 y Mod hyper- TAG	81 x 3 wee ks Crossover	Controlled diets (NCEP Step 1 diet) + control ice cream or ice cream containing inulin	20g/d Inulin	Mean ± SD Total-C HDL-C LDL-C TAG	5.73 ± 0.82 0.92 ± 0.20 3.82 ± 0.71 2.75 ± 1.83	5.93 ± 0.96 0.96 ± 0.26 3.90 ± 0.83 3.20 ± 2.19	NS NS NS P = 0.05
Luo et al (2000)	10 (6/4) 57y (mean) Type II DM	2 x 4 weeks Crossover 2 week washout	FOS or placebo provided as powder to be mixed with beverages or yoghurt	20g/d FOS	Mean ± SE Total-C HDL-C LDL-C TAG	5.13 ± 0.27 1.02 ± 0.08 3.85 ± 0.23 1.33 ± 0.16	5.15 ± 0.24 1.01 ± 0.06 3.85 ± 0.20 1.42 ± 0.12	NS NS NS NS

Letexier et al (2003)8 (4/4)81 x 3 wee ksHigh carbohydrate (55% of total energy) plus inulin or placebo (maltodextrin) consumed before breakfast and evening meals10g/d Inulin	Mean \pm SE4.35 \pm 0.304.12 \pm 0.32NSHDL-C1.31 \pm 0.101.20 \pm 0.11NSLDL-C2.90 \pm 0.222.77 \pm 0.21NSTAG0.77 \pm 0.080.92 \pm 0.10P < 0	0.05
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Hyperchol- hypercholesterolemia NCEP- National Cholesterol Education Program; FOS- fructo-oligosaccharide; GOS- galacto oligosaccharides; Total-C- total cholesterol; HDL-C- high density lipoprotein cholesterol; LDL- low density lipoprotein cholesterol; TAG-triacylglycerol; NS- non significant.

Table 17B: Intervention studies investigating the effect of dietary fibre on blood pressure

Reference	Subjects N (M/F) Age Health	Study duration & design	Intervention	Fibre dose	Outcome (mmol/l)	Fibre group	Control group	P
Lu et al (2004)	15 (6/9) 60 y (mean) Type II DM	81 x 5 wee ks Crossover	Habitual diet + majority of starchy foods replaced with control or AX-rich food products	AX: 14-17g/d NSP	Mean ± SE Systolic BP Diastolic BP	136 ± 4 75 ± 2	139 ± 5 77 ± 2	NS NS
		No washout						

Type II DM- type 2 diabetes; AX- Arabinoxylan, NSP- non starch polysaccharide, BP- blood pressure, NS- non significant.

Table 18B: Effect of fibre supplementation on blood pressure in weight loss studies

Reference	Subjects N (M/F) Age Health	Study duration & design	Intervention	Fibre dose	Outcome (mmol/l)	Fibre group	Control group	Р
Wood et al (2007)	29 (M) 38 y (mean) Overweight	12 weeks Parallel	Carbohydrate restricted diet + 6 capsules containing glucomannan or maltodextrin (placebo) taken daily before meals	3.0g/d glucomannan	Mean ± SD Systolic BP Diastolic BP	119.9 ± 7.7 84.3 ± 7.4	113.7 ± 9.4 77.7 ± 4.6	P < 0.05 NS
Behall et al (2006)	25 (7/18) 38-53 Healthy	2 week run- in (AHA Step 1 diet) followed by 5 week on each treatment	AHA Step 1 diet + whole grain foods containing ß-glucan from barley	0g/d ß-glucan 3g/d ß-glucan 6g/d ß-glucan	Mean ± SE Systolic BP Diastolic BP Systolic BP Diastolic BP	$3g/d \beta$ -glucan 108.7 ± 2.4 65.8 ± 1.7 6g/d β -glucan 114.0 ± 2.4 66.1 ± 1.7	110.2 ± 2.4 65.3 ± 1.7	NS difference between groups
Schwab et al (2006)	66 (29/37) 30-65y Abnormal glucose metabolism	12 weeks Parallel Nutrition counselling provided	Control Polydextrose (PDX) Sugar beet pectin (SBP) Fibre enrichment consumed as a drink (4 dl/d) with meals	SBP: 16g/d total fibre (76% soluble) PDX: 16g/d total fibre	Mean ± SD Systolic BP Diastolic BP Systolic BP Diastolic BP	SBP 129 ± 15 80 ± 7 PDX 135 ± 18 84 ± 9	Control 134 ± 14 81 ± 9	NS difference between groups

AHA- Amercian heart association; NSP- non starch polysaccharide, BP- blood pressure, NS- non significant.

Appendix C: list of colorectal cancer tables

Table 1C: Prospective studies investigating the association of fibre with colorectal cancer

Table 2C: Randomised controlled trials investigating the association of fibre with colorectal adenoma

Table 3C: Prospective studies investigating the association of fibre with colorectal adenoma

Table 4C: Prospective studies investigating the association of fibre from fruits, vegetables and grains with colorectal adenoma

Table 5C: Prospective studies investigating the association of soluble or insoluble fibre with colorectal adenoma

Table 6C: Prospective studies investigating the association of fibre from fruits, vegetables and grains with colorectal cancer

Table7C: Prospective studies investigating the association of soluble or insoluble fibre with colorectal cancer

Reference	Subject population	Measure of exposure	Fiber intake/level	Adjusted Relative Risk or Odds Ratio	P trend	Factors adjusted for in analysis
Bingham et al (2003) Prospective cohort EPIC	519,978 men and women 1,065 case of incident CRC	AOAC and Englyst defined dietary fibre by FFQ	g/day Q1 = 12.8 (M) 12.6 (F) Q2 = 18.0 (M) 17.5 (F) Q3 = 22.0 (M) 20.9 (F) Q4 = 26.5 (M) 24.7 (F)	1.00 0.94 (0.78-1.13) 0.77 (0.63-0.95) 0.76 (0.61-0.95)	0.005	Stratified by centre: age, sex, weight, height, non-fat energy, energy from fat.
Europe	4.5 y follow up		Q5 = 35.6 (M) 31.9 (F) Colon Cancer Q1 = 12.8 (M) 12.6 (F) Q2 = 18.0 (M) 17.5 (F) Q3 = 22.0 (M) 20.9 (F) Q4 = 26.5 (M) 24.7 (F) Q5 = 35.6 (M) 31.9 (F)	0.75 (0.59-0.95) 1.00 0.95 (0.75-1.19) 0.75 (0.58-0.96) 0.71 (0.55-0.94) 0.72 (0.54-0.97)	0.006	
Bingham et al (2005) Prospective cohort EPIC Europe	519,978 men and women 1,826 case of incident CRC 6.2 y follow up	AOAC and Englyst defined dietary fibre by FFQ	g/day (mean values from 24hr recall calibration) Q1 = 18.2 (M) 15.9 (F) Q2 = 21.0 (M) 17.8 (F) Q3 = 23.2 (M) 19.4 (F) Q4 = 25.6 (M) 21.3 (F) Q5 = 30.1 (M) 24.3 (F) Colon Cancer Q1 = 18.2 (M) 15.9 (F) Q2 = 21.0 (M) 17.8 (F) Q3 = 23.2 (M) 19.4 (F) Q4 = 25.6 (M) 21.3 (F) Q5 = 30.1 (M) 24.3 (F)	1.00 0.93 (0.80-1.08) 0.82 (0.69-0.97) 0.79 (0.66-0.96) 0.79 (0.63-0.99) 1.00 0.89 (0.74-1.07) 0.72 (0.59-0.89) 0.70 (0.55-0.88) 0.77 (0.58-1.02)	0.01	Stratified by centre: age, sex, weight, height, non-fat energy, energy from fat, folate intake, red and processed meat intake, physical activity, alcohol consumption, smoking and education.
Fuchs et al (1999) Prospective cohort NHS USA	88,757 women 787 cases of incident CRC 16 y follow up	AOAC defined total dietary fibre by FFQ	g/day Q1 = 9.8 Q2 = 13.1 Q3 = 15.9 Q4 = 19.1 Q5 = 24.9 Colon cancer Q1 = 9.8	1.00 0.90 (0.71-1.13) 0.96 (0.75-1.21) 0.93 (0.72-1.19) 0.95 (0.73-1.25) 1.00	0.59	Age, smoking, BMI, exercise, aspirin, family history, endoscopy, history of adenoma, red meat, alcohol, folate, methionine, calcium, vitamin D.

Table 1C: Prospective studies investigating the association of fibre with colorectal cancer

	Q2 = 13.1	1.04 (0.78-1.39)	
	Q3 = 15.9	1.26 (0.94-1.68)	
	Q4 = 19.1	1.06 (0.77-1.45)	
	Q5 = 24.9	1.04 (0.74-1.46)	

CRC- colorectal cancer

Reference	Subject population	Measure of exposure	Fiber intake/level	Adjusted Relative Risk or Odds Ratio	P trend	Factors adjusted for in analysis
Gaard et al (1996)	50,535 young men and women	AOAC defined total dietary fibre by FFO	g/day standardized to average caloric intake			Age, attained age, BMI, energy intake, height, smoking.
Prospective cohort	143 cases of incident colon cancer		Men Q1 <13.5 Q2 = 13.6-15.6	Men 1.00 0.85 (0.43-1.37)	0.6	
Norwegian National Health	11.4 y follow up		Q3 = 15.7-17.8 Q4 >17.9	0.93 (0.30-1.09) 0.82 (0.46-1.46)		
Screen Norway			Women Q1 <8.5 Q2 = 8.6-9.8	Women 1.00 1.73 (0.73-4.13)	0.12	
,			Q3 = 9.9-11.2 Q4 >11.3	2.42 (1.06-5.51) 2.10 (0.90-4.87)		
Giovannucci et al (1994)	47,949 men	AOAC defined total dietary fibre by	g/day	1.00	0.12	Age, total energy, previous polyps, previous endoscopic screening,
Prospective cohort	cancer	rrų	Q1 = 14.2 Q2 = 18.3 Q3 = 21.7	1.00 0.63 (0.39-1.01) 0.59 (0.36-0.97)	0.12	total pack-years of cigarette smoking, aspirin use, and intake of red meat,
HPFS	6 y follow up		Q4 = 25.6 Q5 = 32.8	1.19 (0.78-1.82) 1.08 (0.68-1.70)		methionine and alcohol
USA						
Heilbrun et al (1989)	102 male cases of colon cancer	AOAC defined total dietary fibre by 24	g/day	1.00 (rof)	0.35	Age
Nested case- control study	361 male controls	in recail	$Q_2 = 13.10-14.79$ $Q_3 = 10.40-13.09$	0.89 0.86	0.55	
Japan-Hawaii Cancer Study	16 y follow up in cohort		Q4 = 7.50-10.39 Q5 <7.50	0.77 1.40		
USA						
Higginbotham et al (2004)	38,451 women	AOAC defined total dietary fibre by	g/day			Age, BMI, history of oral contraceptive use, postmenopausal hormone use,
Prospective cohort	7.9 y follow up	FFQ	Q1 <12.5 Q2 12.5<17.0 Q3 17.0<20.0	1.00 0.99 (0.61-1.62) 0.57 (0.32-1.00)	0.5	family history of CRC, smoking, alcohol use, physical activity, NSAID use, total energy, energy-adjusted total fat,
			Q4 20.0<23.1	0.96 (0.58-1.59)		energy-adjusted folate, energy-

Women's		Q5 >23.1	0.79 (0.45-1.38)	adjusted calcium, and energy-adjusted
Health Study				vitamin D
USA				

CRC- colorectal cancer; NSAID- non-steroidal anti-inflammatory drugs

Reference	Subject population	Measure of exposure	Fiber intake/level	Adjusted Relative Risk or Odds Ratio	P trend	Factors adjusted for in analysis
Lin et al (2005) Prospective cohort WHS USA	36,976 women 223 cases of incident CRC 10 y follow up	AOAC defined total dietary fibre by FFQ	g/day Q1 = 12 Q2 = 16 Q3 = 18 Q4 = 21 Q5 = 26	1.00 0.90 (0.59-1.38) 0.62 (0.39-0.98) 0.84 (0.54-1.31) 0.75 (0.47-1.18)	0.11	Age, treatment assignment, BMI, family history of CRC, history of colon polyps, physical activity, smoking status, baseline aspirin use, red meat intake, alcohol consumption, total energy intake, menopausal status, baseline post-menopausal HT use, folate intake and multivitamin use
Mai et al (2003) Prospective cohort Breast Cancer Detection Demonstration Project Follow- up Study USA	45,491 women 487 cases of incident CRC 8.5 y follow up	AOAC defined total dietary fibre by FFQ	g/1000kcal /day Q1 = 4.9 Q2 = 7.2 Q3 = 8.8 Q4 = 10.7 Q5 = 14.8	1.00 0.90 (0.67-1.19) 0.67 (0.49-0.91) 1.00 (0.76-1.33) 0.94 (0.70-1.26)	Not stated	Non-steroidal anti-inflammatory drugs, smoking, alcohol, calcium, vitamin D, red meat, height, BMI, education
McCullough et al (2003) Prospective cohort Cancer Prevention Study II Nutrition Cohort Study USA	62,609 men 70,554 women 298 male and 210 female cases of incident CRC 5 y follow up	AOAC defined total dietary fibre by FFQ	g/day Men Q1 <9.3 Q2 9.3-11.3 Q3 11.4-13.5 Q4 13.6-16.6 Q5 >16.6 Women Q1 <8.0 Q2 8.0-9.8 Q3 9.9-11.7 Q4 11.8-14.4 Q5 >14.4	Men 1.00 0.63 (0.44-0.92) 0.64 (0.43-0.96) 0.84 (0.55-1.28) 1.01 (0.62-1.65) Women 1.00 1.41 (0.90-2.19) 1.04 (0.62-1.75) 1.79 (1.07-2.99) 1.09 (0.58-2.05)	0.594 0.65	Age, exercise METs, aspirin, smoking, family history of colorectal cancer, BMI, education, energy, multivitamin use, total calcium, red meat intake, HRT use, dietary folate

CRC- colorectal cancer

Reference	Subject population	Measure of exposure	Fiber intake/level	Adjusted Relative Risk or Odds Ratio	P trend	Factors adjusted for in analysis
Michels et al (2005) Prospective cohort Nurses' Health Study & Health Professionals Follow-up Study USA	76,947 women & 47,279 men 1596 cases of incident CRC 1.8 million person- years follow up	AOAC defined total dietary fibre by FFQ	g/1,000 kcal/day at baseline Men Q1 = 6.5 Q2 = 8.6 Q3 = 10.1 Q4 = 11.9 Q5 = 16.1 Women Q1 = 6.5 Q2 = 8.4 Q3 = 9.8 Q4 = 11.4 Q5 = 14.9 Colon cancer Men Q1 < 8.0 Q2 = $8.0-10.0$ Q3 = $10.0-12.0$ Q4 = $12.0-14.0$ Q5 > 14.0 Women Q1 < 8.0 Q2 = $8.0-10.0$ Q3 = $10.0-12.0$ Q4 = $12.0-14.0$ Q5 > 14.0	or Odds Ratio Men 1.00 0.79 (0.61- 1.01) 0.71 (0.54- 0.93) 0.85 (0.65- 1.12) 0.79 (0.59- 1.07) Women 1.00 1.01 (0.81- 1.26) 1.11 (0.89- 1.38) 1.17 (0.93- 1.46) 0.90 (0.70- 1.17) Men 1.00 0.85 (0.63- 1.15) 0.78 (0.56- 1.09) 0.94 (0.65-	0.36 0.57 0.76 0.63	Age, time period, family history of CRC, history of colonoscopy, height, BMI, physical activity, regular aspirin use, duration of aspirin use, pack-yrs of early-onset smoking, multivitamin supplement use, fat energy, non-fat energy, alcohol consumption, dietary folate, red meat consumption, processed meat, glycemic load, calcium, methionine, menopausal status and postmenopausal hormone use
				0.85 (0.56- 1.30)		

Women 1.00
1.18 (0.91- 1.53) 1.24 (0.93-
1.64) 1.18 (0.86- 1.63)
0.95 (0.65- 1.39)

CRC- colorectal cancer; FFQ- food frequency questionnaire

Reference	Subject population	Measure of exposure	Fiber intake/level	Adjusted Relative Risk or Odds Ratio	P trend	Factors adjusted for in analysis
Nomura et al	85,903 men	AOAC defined total	g/1,000 kcal/day			Ethnicity, time since study entry, age,
(2007)	105,108	dietary fibre by				family history of colorectal cancer,
	women	FFQ	Men	Men		history of colorectal polyp, pack-years
Prospective			Q1 = 6.1	1.00	0.002	of cigarette smoking, BMI, hours of
cohort	1,138 male		Q2 = 8.4	0.75 (0.62;0.91)		vigorous activity, aspirin use,
	and 972		Q3 = 10.3	0.73 (0.60;0.89)		multivitamin use, and replacement
Multiethnic	female cases		Q4 = 12.5	0.77 (0.62;0.96)		hormone use (women)
Cohort Study	of incident CRC		Q5 = 16.5	0.62 (0.48;0.79)		
USA			Women	Women		
	7 y follow up		Q1 = 7.5	1.00	0.245	
			Q2 = 10.0	1.07 (0.86;1.34)		
			Q3 = 12.1	0.84 (0.66;1.07)		
			Q4 = 14.5	0.97 (0.76;1.24)		
			Q5 = 18.6	0.88 (0.67;1.14)		
			Colon cancer			
			Men	Men		
			Q1 = 6.1	1.00	0.031	
			Q2 = 8.4	0.76 (0.60-0.96)		
			Q3 = 10.3	0.78 (0.61-0.99)		
			Q4 = 12.5	0.88 (0.69-1.14)		
			Q5 = 16.5	0.64 (0.48-0.86)		
			Women	Women		
			Q1 = 7.5	1.00	0.361	
			$Q^2 = 10.0$	1.16 (0.90-1.50)		
			Q3 = 12.1	0.88 (0.67-1.16)		
			Q4 = 14.5	1.02 (0.77-1.34)		
			Q5 = 18.6	0.92 (0.68-1.25)		

CRC- colorectal cancer ; FFQ- food frequency questionnaire
Otani et al (2006) Prospective cohort	78,326 men and women	AOAC defined total	g/day			
Japan Public Health Center Study Japan	522 cases of incident CRC 5.8 y follow up	FFQ	Men Q1 = 6.4 Q2 = 9.1 Q3 = 11.2 Q4 = 13.6 Q5 = 18.7	Men 1.00 0.90 (0.63-1.3) 0.70 (0.47-1.1) 0.88 (0.58-1.3) 0.85 (0.53-1.4)	0.48	Age, alcohol, smoking, BMI, physical exercise, folate, calcium, vitamin D, red meat intake, family history of CRC, study areas
			Women Q1 = 8.3 Q2 = 11.2 Q3 = 13.3 Q4 = 15.6 Q5 = 20.0	Women 1.00 0.61 (0.35-1.0) 0.62 (0.36-1.1) 0.77 (0.44-1.3) 0.58 (0.31-1.10)	0.21	
			Colon cancer Men Q1 = 6.4 Q2 = 9.1 Q3 = 11.2 Q4 = 13.6 Q5 = 18.7	Men 1.00 0.84 (0.55-1.3) 0.63 (0.38-1.0) 0.77 (0.46-1.3) 0.80 (0.45-1.4)	0.39	
			Women Q1 = 8.3 Q2 = 11.2 Q3 = 13.3 Q4 = 15.6 Q5 = 20.0	Women 1.00 0.67 (0.36-1.3) 0.60 (0.31-1.2) 0.80 (0.42-1.5) 0.48 (0.23-1.0)	0.12	
Park et al (2005) 13 Pooled prospective cohort studies Europe and North America	725,628 men and women 8081 cases of incident CRC 6-20 y follow up (7,328,414 py)	AOAC and Englyst defined total dietary fibre by FFQ	g/day <10 10-<15 15-<20 20-<25 25-<30 ≥30 Colon cancer	1.18 (1.05-1.31) 1.00 (ref) 1.02 (0.95-1.10) 1.01 (0.92-1.10) 0.99 (0.87-1.12) 1.00 (0.85-1.17)	0.68	Age, BMI, height, education, physical activity, family history of colorectal cancer, HRT use, oral contraceptive use, NSAID use, multivitamin use, smoking, alcohol intake, folate intake, red meat intake, milk intake and total energy

10-<15 15-<20 20-<25 25-<30 ≥30	1.00 (ref) 1.04 (0.95-1.13) 1.03 (0.93-1.15) 1.08 (0.93-1.25) 1.04 (0.86-1.26)	

Reference	Subject population	Measure of exposure	Fiber intake/level	Adjusted Relative Risk or Odds Ratio	P trend	Factors adjusted for in analysis
Pietinen et al (1999) Prospective cohort Alpha Tocopherol Beta Carotene (ATBC) trial Finland	29,133 male smokers 185 cases of incident colon cancer 8 y follow up	Englyst defined total dietary fibre by FFQ	g/day Q1 = 16 Q2 = 22.2 Q3 = 26.9 Q4 = 34.1	1.00 1.0 (0.7-1.6) 1.0 (0.7-1.5) 1.0 (0.6-1.5)	0.79	Smoking years, BMI, alcohol, education, physical activity at work, and calcium intake
Schatzkin et al (2007) Prospective cohort NIH-AARP USA	291,988 men 197,623 women 2974 cases of incident CRC 5 y follow up	AOAC defined total dietary fibre by FFQ	g/1,000 kcal/day Q1 = 6.6 Q2 = 8.6 Q3 = 10.3 Q4 = 12.3 Q5 = 15.9 Colon cancer Q1 = 6.6 Q2 = 8.6 Q3 = 10.3 Q4 = 12.3 Q5 = 15.9	1.00 0.92 (0.82-1.03) 0.93 (0.82-1.06) 0.90 (0.78-1.04) 0.99 (0.85-1.15) 1.00 0.90 (0.79-1.03) 0.93 (0.80-1.08) 0.87 (0.74-1.02) 0.96 (0.80-1.15)	0.96	Sex, physical activity, smoking, menopausal HRT use, and intakes of red meat, dietary calcium, dietary folate, and total energy
Sellers et al (1998) Prospective cohort Iowa Women's Heath Study USA	35,216 women 241 cases of incident colon cancer 10 y follow up	AOAC defined total dietary fibre by FFQ	g/day Without family history T1 <16.17 T2 = 16.18-22.59 T3 >22.60 With family history T1 <16.17 T2 = 16.18-22.59 T3 >22.60	1.0 0.8 (0.5-1.1) 0.8 (0.5-1.2) 1.0 1.2 (0.7-2.3) 1.2 (0.6-2.6)	0.3	Age, total energy, history of rectal colon polyps
Steinmetz KA 1994 Prospective cohort Iowa Women's Heath Study	35216 women 212 cases of incident colon cancer	AOAC defined total dietary fibre by FFQ	g/day Q1 <14.5 Q2 = 14.5-18.9 Q3 = 19.0-24.7 Q4 >24.7	1.00 1.14 (0.79-1.65) 0.95 (0.63-1.43) 0.80 (0.49-1.31)		Age, energy

	5 y follow up			
USA				

Reference	Subject population	Measure of exposure	Fiber intake/level	Adjusted Relative Risk or Odds Ratio	P trend	Factors adjusted for in analysis
Terry et al (2001) Prospective cohort Swedish Mammography Cohort Sweden	61 463 women 460 cases of incident CRC 9.6 y follow up	AOAC defined total dietary fibre by FFQ	g/day standardized to average caloric intake Q1 = 12.3 Q2 = 15.6 Q3 = 18.1 Q4 = 21.8	1.00 0.96 (0.73-1.28) 1.05 (0.79-1.40) 0.96 (0.70-1.33)	0.98	Age, BMI, educational level, intake of energy, and quartiles of alcohol, red meat, total fat, folic acid, vitamin D, vitamin C, and calcium
Wakai et al (2007) Prospective cohort Japan Multi-Centered Study Japan	43,115 men and women 443 cases of incident CRC 7.6 y follow up	AOAC defined total dietary fibre by FFQ	g/day Men Q1 = 6.7 Q2 = 9.4 Q3 = 11.3 Q4 = 13.4 Women Q1 = 7.4 Q2 = 9.8 Q3 = 11.5 Q4 = 13.4 Colon cancer Men Q1 = 6.7 Q2 = 9.4	Men 1.00 1.12 (0.77-1.62) 0.62 (0.40-0.96) 0.69 (0.43-1.11) Women 1.00 0.73 (0.47-1.14) 0.84 (0.54-1.33) 0.75 (0.46-1.25) Men 1.00 1.06 (0.66-1.69)	0.023 0.41 0.004	Age, sex, area, educational level, fam hist of CRC, alcohol consumption, smoking, BMI, daily walking habits, exercise, sedentary work, beef intake, pork intake, energy intake, and energy- adjusted intakes of folate, calcium, and vitamin D
Willett et al (1990) Prospective cohort NHS USA	88,751 women 150 cases of incident CRC 6 y follow up	AOAC defined total dietary fibre by FFQ	Q2 = 9.4 Q3 = 11.3 Q4 = 13.4 Women Q1 = 7.4 Q2 = 9.8 Q3 = 11.5 Q4 = 13.4 g/day standardized to average caloric intake Q1 <11.6 Q2 = 11.6-14.3 Q3 = 14.4-17.2 Q4 = 17.3-21.1	1.00 (0.00-1.09) 0.43 (0.24-0.76) 0.52 (0.28-0.96) Women 1.00 0.72 (0.44-1.18) 0.71 (0.42-1.20) 0.64 (0.36-1.13) 1.00 0.74 (0.43-1.28) 1.16 (0.72-1.88) 0.69 (0.40-1.17)	0.15	Age and energy

		Q5 >21.3	0.90 (0.54-1.49)	
	c			

Reference	Subject population and Design	Time period	Intervention	No. with ≥1 recurrent adenoma/total no. (%)	Adjusted Odds Ratio	P trend	Factors adjusted for in analysis
Alberts et al (2000) Wheat Bran Fiber Trial	1429 (1303 completed) men and women with previous adenoma parallel	34 months	2 g/day Wheat Bran Fibre 12.5 g/day Wheat Bran fibre	299/584 (51.2) 338/719 (47.0)	0.88 (0.70-1.11)	0.28	Randomization scheme
Bonithon-Kopp et al (2000) European Cancer Prevention Organisation (ECP) Intervention Study	665 (552 completed) men and women with previous adenoma parallel	36 months	placebo 3.5 g/day ispaghula husk	36/178 (20.2) 58/198 (29.3)	1.67 (1.01-2.76)	0.042	Age, sex, adenoma history, and number and location of adenomas inclusion

Table 2C: Randomised controlled trials investigating the association of fibre with colorectal adenoma

Reference	Subject population	Measure of exposure	Fiber intake/level	Adjusted Relative Risk or Odds Ratio	P trend	Factors adjusted for in analysis
Fuchs et al (1999) Prospective cohort Nurses Health Study USA	27,530 women with sigmoidoscopy 1012 cases of adenoma 16 y follow up	AOAC defined total dietary fibre by FFQ	g/day Q1 = 9.8 Q2 = 13.1 Q3 = 15.9 Q4 = 19.1 Q5 = 24.9	1.00 0.98 (0.79-1.21) 1.07 (0.86-1.33) 0.95 (0.76-1.20) 0.91 (0.71-1.16)	0.36	Age, smoking, BMI, exercise, aspirin, family history, endoscopy, history of adenoma, red meat, alcohol, folate, methionine, calcium, vitamin D
Giovannucci et al (1992) Prospective cohort HPFS USA	7284 men with colonoscopy 170 cases of colorectal adenoma	AOAC defined total dietary fibre by FFQ	g/day Q1 <16.6 Q2 = 16.6-20.0 Q3 = 20.1-23.5 Q4 = 23.6-28.3 Q5 >28.3	1.00 1.04 (0.67;1.61) 0.75 (0.47;1.20) 0.64 (0.40;1.01) 0.36 (0.22;0.60)	<0.001	Age, total energy intake, and family history of colon cancer
Jacobs et al (2002) Prospective cohort Wheat Bran Fiber Trial; baseline analysis USA	1304 Men and women 638 cases of recurrent colorectal adenoma 3 y follow up colonoscopy	AOAC defined total dietary fibre by FFQ	g/day Q1 = 10.1 Q2 = 15.3 Q3 = 20.0 Q4 = 27.7	1.00 0.79 (0.56;1.12) 0.76 (0.54;1.08) 0.83 (0.57;1.19)	0.31	History of polyps prior to baseline colonoscopy, age at randomization, sex, number of colonoscopies, aspirin use, total calcium intake at baseline, and number of baseline adenomas
Jacobs et al (2002) Prospective cohort Wheat Bran Fiber Trial; analysis by adherence to intervention USA	1208 men and women 575 cases of recurrent colorectal adenoma 3 y follow up colonoscopy	AOAC defined total dietary fibre by FFQ	g/day Q1 = 4.9-17.8 Q2 = 17.9-23.7 Q3 = 23.8-30.3 Q4 = 30.4-66.3	1.00 0.87 (0.62;1.23) 0.80 (0.56;1.14) 0.98 (0.68;1.42)	0.82	History of polyps prior to baseline colonoscopy, age at randomization, gender, number of colonscopies, total calcium intake at baseline, and number of baseline adenomas

Table 3C: Prospective studies investigating the association of fibre with colorectal adenoma

FFQ- food frequency questionnaire

Reference	Subject population	Measure of exposure	Fiber intake/level	Adjusted Relative Risk or Odds Ratio	P trend	Factors adjusted for in analysis
Peters et al (2003) Nested case-control study PLCO Cancer Screening Trial, screening arm USA	3591 cases with ≥1 polyp 33971 controls	AOAC defined total dietary fibre by FFQ	g/day Q1 <15.4 Q2 = 15.4-19.7 Q3 = 19.8-24.2 Q4 = 24.3-30.5 Q5 >30.6	1.00 0.91 (0.81-1.01) 0.85 (0.76-0.96) 0.79 (0.69-0.90) 0.73 (0.62-0.86)	0.002	Age, sex, centre, energy intake, ethnicity, education, smoking, alcohol use, use of aspirin and ibuprofen separately, physical activity, BMI, red meat intake, calcium intake, and folate intake
Platz et al (1997) Prospective cohort HPFS USA	16,448 men 1017 cases of adenomatous polyps of the distal colon 8 y follow up	AOAC defined total dietary fibre by FFQ	g/day Q1 = 11.6 Q2 = 16.1 Q3 = 19.7 Q4 = 24.2 Q5 = 32.2	1.00 0.89 (0.67-1.17) 0.97 (0.72-1.30) 0.86 (0.62-1.21) 0.88 (0.59-1.31)	0.1	Age at endoscopy, history of endoscopy prior to 1986, family history of CRC, total daily energy intake BMI, pack- years smoked, daily alcohol consumption, multivitamin use, physical activity, regular aspirin use, and intake of red meat, folate, and methionine
Robertson et al (2005) Prospective cohort Pooling of the Antioxidant Polyp Prevention Study and Calcium Polyp Prevention Study USA	1520 men and women 540 cases of colorectal adenoma recurrence 4 yr follow up colonoscopy	AOAC defined total dietary fibre by FFQ	g/day standardized to 2,000 kcal/day diet Q1 = 7.92 Q2 not given Q3 not given Q4 = 25.75	1.00 1.10 (0.91, 1.33) 0.90 (0.74, 1.10) 0.85 (0.69-1.05)	0.07	Age, sex, clinical center, treatment category, study, and duration of observation

Table 4C: Prospective studies investigating the association of fibre from fruits, vegetables and grains with colorectal adenoma

Reference	Subject population	Measure of exposure	Fiber intake/level	Adjusted Relative Risk or Odds Ratio	P trend	Factors adjusted for in analysis
Giovannucci et al (1992)	7284 men with colonoscopy	AOAC defined dietary fibre by	g/day			Age, total energy intake, and family history of colon cancer
Prospective cohort Health Professions Follow up Study USA	170 cases of colorectal adenoma	rrų	Q1 <1.9 Q2 = $1.9-3.2$ Q3 = $3.3-4.8$ Q4 = $4.9-7.1$ Q5 >7.1	1.00 0.69 (0.43-1.10) 0.67 (0.42-1.08) 0.63 (0.41-0.97) 0.53 (0.28-1.02)	0.02	
			Fibre from Vegetables Q1 <3.7 Q2 = 3.7-5.1 Q3 = 5.2-6.8 Q4 = 6.9-9.3 Q5 >9.3	1.00 0.99 (0.64-1.53) 0.76 (0.47-1.23) 0.65 (0.39-1.08) 0.53 (0.30-0.91)	0.003	
			Fibre from Grains Q1 <2.8 Q2 = 2.8-4.6 Q3 = 4.8-7.0 Q4 = 7.1-10.5 Q5 >10.6	1.00 0.79 (0.50-1.23) 0.78 (0.49-1.23) 0.73 (0.46-1.17) 0.44 (0.26-0.76)	<0.001	
Jacobs et al (2002) Prospective cohort Wheat Bran Fiber Trial; baseline analysis	1304 Men and women 638 cases of recurrent colorectal adenoma	AOAC defined dietary fibre by FFQ	g/day Fibre from Fruits Q1 = 1.5 Q2 = 3.7 Q3 = 6.2 Q4 = 11.5	1.00 0.89 (0.63-1.25) 0.86 (0.60-1.21) 0.92 (0.64-1.32)	0.62	History of polyps prior to baseline colonoscopy, age at randomization, sex, number of colonoscopies, aspirin use, total calcium intake at baseline, and number of baseline adenomas
USA	3 y follow up colonoscopy		Fibre from Vegetables Q1 = 1.5 Q2 = 2.7 Q3 = 3.4 Q4 = 6.8	1.00 0.98 (0.70-1.38) 0.86 (0.60-1.20) 1.34 (0.94-1.91)	0.21	

	Fibre from Grains Q1 = 1.4 Q2 = 2.8 Q3 = 4.4 Q4 = 8.8	1.00 0.92 (0.65-1.29) 0.88 (0.62-1.24) 0.84 (0.59-1.19)	0.31	
	Q+ - 0.0	0.04 (0.05 1.15)		

FFQ- food frequency questionnaire

Reference	Subject population	Measure of exposure	Fiber intake/level	Adjusted Relative Risk or Odds Ratio	P trend	Factors adjusted for in analysis
Platz et al (1997) Prospective cohort Health Professions	16,448 men 1017 cases of adenomatous polyps of the	AOAC defined dietary fibre by FFQ	g/day Fibre from Fruits Q1 = 1.3 Q2 = 2.5	1.00 0.94 (0.72-1.24)	0.03	Age at endoscopy, history of endoscopy prior to 1986, family history of CRC, total daily energy intake BMI, pack- years smoked, daily alcohol consumption, multivitamin use,
USA	8 y follow up		$Q_3 = 3.9$ $Q_4 = 5.4$ $Q_5 = 8.4$	1.06 (0.81-1.39) 0.73 (0.54-0.98) 0.81 (0.59-1.11)		and intake of red meat, folate, and methionine
			Q1 = 3.2 Q2 = 4.7 Q3 = 6.1 Q4 = 7.9 Q5 = 11.5 Fibre form Ordina	1.00 0.95 (0.72-1.26) 1.14 (0.86-1.50) 1.07 (0.80-1.43) 0.93 (0.67-1.30)	0.33	
			$\begin{array}{l} \text{Fibre from Grains} \\ \text{Q1} = 2.1 \\ \text{Q2} = 3.7 \\ \text{Q3} = 5.2 \\ \text{Q4} = 7.1 \\ \text{Q5} = 10.6 \end{array}$	1.00 1.12 (0.85-1.48) 1.11 (0.83-1.49) 1.31 (0.97-1.78) 1.20 (0.86-1.66)	0.76	

Reference	Subject population	Measure of exposure	Fiber intake/level	Adjusted Relative Risk or Odds Ratio	P trend	Factors adjusted for in analysis
Platz et al (1997)	16,448 men	AOAC defined dietary fibre by	g/day			Age at endoscopy, history of endoscopy prior to 1986, family history of CRC,
Prospective cohort Health Professions Follow up Study USA	1017 cases of adenomatous polyps of the distal colon 8 y follow up	FFQ	Insoluble fibre Q1 = 8.5 Q2 = 12 Q3 = 14.9 Q4 = 18.5 Q5 = 25.0	1.00 1.02 (0.77-1.35) 1.08 (0.80-1.46) 1.24 (0.90-1.73) 1.14 (0.77-1.69)	0.59	total daily energy intake BMI, pack- years smoked, daily alcohol consumption, multivitamin use, physical activity, regular aspirin use, and intake of red meat, folate, and methionine
			Soluble fibre Q1 = 3.4 Q2 = 4.7 Q3 = 5.8 Q4 = 7.1 Q5 = 9.4	1.00 0.91 (0.69-1.19) 1.01 (0.76-1.35) 0.78 (0.56-1.08) 0.69 (0.46-1.03)	0.007	

Table 5C: Prospective studies investigating the association of soluble or insoluble fibre with colorectal adenoma

Reference	Subject population	Measure of exposure	Fibre intake/level	Adjusted Relative Risk or Odds Ratio	P trend	Factors adjusted for in analysis
Bingham et al (2003)	519,978 men and	AOAC and Englyst	g/day			Stratified by centre: age, sex, weight,
Prospective cohort	women	fibre by FFO	Fibre from Fruits			neight, non-fat energy, energy from fat
	1.065 case of	nore by rr q	01 = 2.21	1.00	0.17	
EPIC	incident CRC		$Q^2 = 3.41$	0.69 (0.57-0.85)		
			Q3 = 4.29	0.76 (0.63-0.92)		
Europe	4.5 y follow up		Q4 = 5.36	0.82 (0.66-0.99)		
			Q5 = 7.76	0.78 (0.64-0.97)		
			Fibre from Vegetables			
			Q1 = 2.83	1.00	0.52	
			Q2 = 3.77	0.94 (0.77-1.15)		
			Q3 = 4.42	0.95 (0.77-1.16)		
			Q4 = 5.11	1.00 (0.81-1.24)		
			Q5 = 6.48	0.88 (0.70-1.11)		
			Fibre from Grains			
			Q1 = 4.72	1.00	0.06	
			Q2 = 6.61	0.89 (0.74-1.08)		
			Q3 = 7.93	0.85 (0.69-1.03)		
			Q4 = 9.35	0.88 (0.71-1.08)		
			Q5 = 12.05	0.78 (0.62-0.98)		
Bingham et al (2005)	519,978 men and	AOAC and Englyst	g/day			Stratified by centre: age, sex, weight,
	women	defined dietary				height, non-fat energy, energy from
Prospective cohort	1.026 anal of	fibre by FFQ	Fibre from Fruits	1.00	0.42	fat, folate intake, red and processed
EDIC	1,020 Case OI		QI = 2.7 (M) 2.8 (F) Q2 = 3.1 (M) 3.4 (F)	1.00	0.42	consumption smoking and education
			$O_2 = 3.1 (M) 3.4 (T)$ $O_3 = 3.8 (M) 3.9 (F)$	0.78 (0.67-0.92)		consumption, smoking and education.
Europe	6.2 v follow up		O4 = 4.3 (M) 4.4 (F)	0.90 (0.76-1.06)		
	- , F		Q5 = 5.3 (M) 5.4 (F)	0.81 (0.68-0.97)		
			Fibre from vegetables $O_1 = 2.7 (M) - 2.8 (E)$	1.00	0.52	
			QI = 2.7 (M) 2.8 (F) Q2 = 3.1 (M) 3.4 (F)	1.00 0.99 (0.84-1.17)	0.52	
			$Q_2 = 3.1 (M) 3.4 (T)$ $Q_3 = 3.8 (M) 3.9 (F)$	0.97 (0.82-1.16)		
			Q4 = 4.3 (M) 4.4 (F)	0.96 (0.80-1.16)		
			Q5 = 5.3 (M) 5.4 (F)	0.94 (0.76-1.16)		
			Fibre from Grains			
			Q1 = 6.6 (M) 4.9 (F)	1.00	0.44	

Table 6C: Prospective studies investigating the association of fibre from fruits, vegetables and grains with colorectal cancer

Q5 = 13.1 (M) 9.2 (F) 0.93 (0.76-1.15)
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Reference	Subject population	Measure of exposure	Fibre intake/level	Adjusted Relative Risk or Odds Ratio	P trend	Factors adjusted for in analysis
Fuchs et al (1999) Prospective cohort Nurses Health Study	88,757 women 787 cases of incident CRC 16 y follow up	AOAC defined dietary fibre by FFQ	g/day Fibre from Fruits Q1 = 0.8 Q2 = 1.9 Q3 = 3.1 Q4 = 4.5 Q5 = 7.2	1.00 0.94 (0.74-1.18) 1.03 (0.82-1.29) 0.88 (0.69-1.12) 0.86 (0.67-1.10)	0.16	Age, smoking, BMI, exercise, aspirin, family history, endoscopy, history of adenoma, red meat, alcohol, folate, methionine, calcium, vitamin D.
USA			Fibre from Vegetables Q1 = 2.7 Q2 = 4.0 Q3 = 5.3 Q4 = 6.8 Q5 = 10.0 Fibre from Grains	1.00 0.98 (0.78-1.23) 0.98 (0.78-1.25) 1.13 (0.89-1.43) 1.35 (1.05-1.72)	0.004	
			Q1 = 1.0 Q2 = 1.8 Q3 = 2.5 Q4 = 3.3 Q5 = 4.8	1.00 0.90 (0.71-1.13) 0.95 (0.75-1.19) 0.99 (0.79-1.24) 1.00 (0.79-1.27)	0.69	
Giovannucci et al (1994) Prospective cohort Health Professions Follow up	47,949 men 205 cases of incident colon cancer 6 y follow up	AOAC defined dietary fibre by FFQ	g/day Fibre from Fruits Q1 = 1.2 Q2 = 2.6 Q3 = 4.1 Q4 = 5.8 Q5 = 9.2	1.00 1.6 (1.01-2.55) 2.03 (1.29-3.20) 1.61 (0.98-2.81) 1.66 (0.98-2.81)	0.08	Age, total energy, previous polyps, previous endoscopic screening, parental history of colorectal cancer, total pack-years of cigarette smoking, aspirin use, and intake of red meat, methionine and alcohol
Study USA			Fibre from Vegetables Q1 = 2.8 Q2 = 4.5 Q3 = 6.0 Q4 = 7.9 Q5 = 11.7 Fibre from Grains Q1 = 2.3 Q2 = 4.4	1.00 1.13 (0.74-1.73) 1.26 (0.82-1.94) 0.86 (0.52-1.40) 1.17 (0.72-1.90) 1.00 1.27 (0.82 1.06)	0.91	

Q3 = 6.6 $1.22 (0.77-1.93)$ $Q4 = 9.4$ $1.63 (1.04-2.57)$ $Q5 = 15.3$ $1.28 (0.78-2.09)$	
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Reference	Subject population	Measure of exposure	Fibre intake/level	Adjusted Relative Risk or Odds Ratio	P trend	Factors adjusted for in analysis
Lin et al (2005)	36,976 women	AOAC defined	g/day			Age, treatment assignment, BMI,
Due en estive se heut	222 anone of	dietary fibre by	Fibus fuero Fuuite			family history of CRC, history of colon
Prospective conort	incident CRC	FFQ	FIDE FROM FRUILS $01 = 2.5$	1.00	0.65	status baseline aspirin use red meat
WHS			$Q_1 = 2.5$ $Q_2 = 3.5$	0.78 (0.51-1.19)	0.05	intake, alcohol consumption, total
	10 y follow up		Q3 = 4.2	0.95 (0.63-1.43)		energy intake, menopausal status,
USA			Q4 = 4.9	0.67 (0.44-1.04)		baseline post-menopausal HT use,
			Q5 = 6.0	1.0 (0.67-1.49)		folate intake and multivitamin use.
			Fibre from Vegetables			
			01 = 5.9	1.00	0.66	
			$Q^2 = 6.4$	0.90 (0.57-1.41)		
			Q3 = 6.8	1.16 (0.75-1.78)		
			Q4 = 7.3	1.46 (0.97-2.20)		
			Q5 = 8.0	1.00 (0.65-1.56)		
			Fibre from Grains			
			Q1 = 3.1	1.00	0.66	
			Q2 = 3.9	1.0 (0.67-1.49)		
			Q3 = 4.4	0.56 (0.35-0.90)		
			Q4 = 5.0 O5 = 6.1	0.95 (0.03-1.42)		
			25 - 0.1	0.57 (0.00 1.42)		

Reference	Subject population	Measure of exposure	Fibre intake/level	Adjusted Relative Risk or Odds Ratio	P trend	Factors adjusted for in analysis
Michels et al (2005) Prospective cohort Nurses' Health Study & Health Professionals Follow-up	76,947 women & 47,279 men 1596 cases of incident CRC 1.8 million person-years follow up	AOAC defined dietary fibre by FFQ	g/1,000 kcal/day at baseline Fibre from Fruits Men Q1 = 1.4 Q2 = 2.8 Q3 = 4.1 Q4 = 5.6 Q5 = 9.3	Men 1.00 1.02 (0.79-1.33) 0.91 (0.70-1.19) 1.08 (0.82-1.41) 0.92 (0.68-1.13)	0.62	Age, time period, family history of CRC, history of colonoscopy, height, BMI, physical activity, regular aspirin use, duration of aspirin use, pack-yrs of early-onset smoking, multivitamin supplement use, fat energy, non-fat energy, alcohol consumption, dietary folate, red meat consumption, processed meat, glycemic load, calcium, methionine, menopausal status and postmenopausal hormone
Study USA			Women Q1 = 1.4 Q2 = 2.5 Q3 = 3.5 Q4 = 4.7 Q5 = 7.3	Women 1.00 0.94 (0.75-1.16) 1.05 (0.84-1.31) 0.85 (0.67-1.07) 0.88 (0.68-1.13)	0.20	use
			Fibre from Vegetables Men Q1 = 3.6 Q2 = 5.3 Q3 = 6.6 Q4 = 8.3 Q5 = 12.2	Men 1.00 1.04 (0.82 -1.32) 0.84 (0.65-1.08) 0.93 (0.72-1.20) 1.09 (0.83-1.42)	0.57	
			Women Q1 = 3.6 Q2 = 5.0 Q3 = 6.0 Q4 = 7.2 Q5 = 10.0	Women 1.00 1.12 (0.90-1.39) 1.07 (0.85-1.34) 1.26 (1.01-1.59) 1.20 (0.94-1.56)	0.11	
			Fibre from Grains Men Q1 = 2.8 Q2 = 4.3 Q3 = 5.6 Q4 = 7.3 Q5 = 11.45	Men 1.00 0.81(0.63-1.04) 0.98 (0.76-1.25) 0.89 (0.69-1.16) 0.79 (0.60-1.05)	0.19	
			Women	Women		

	Q1 = 2.3 Q2 = 3.3 Q3 = 4.1 Q4 = 5.2	1.00 0.81 (0.66-1.00) 0.88 (0.71-1.08) 0.86 (0.69-1.07)	0.63	
	Q5 = 8.0	0.89 (0.71-1.12)		

Reference	Subject population	Measure of exposure	Fibre intake/level	Adjusted Relative Risk or Odds Ratio	P trend	Factors adjusted for in analysis
Nomura et al (2007) Prospective cohort Multiethnic Cohort Study	85,903 men 105,108 women 1,138 male and 972 female cases of incident CRC 7 y follow up	AOAC defined dietary fibre by FFQ	g/1,000 kcal/day Fibre from Fruit Men Q1 = 0.9 Q2 = 2.5 Q3 = 4.3 Q4 = 6.9 Q5 = 12.6	Men 1.00 0.86 (0.71-1.05) 0.76 (0.62-0.93) 0.78 (0.63-0.96) 0.78 (0.63-0.97)	0.076	Ethnicity, time since study entry, age. Family history of colorectal cancer, history of colorectal polyp, pack-years of cigarette smoking, BMI, hours of vigorous activity, aspirin use, multivitamin use, and replacement hormone use (women)
USA			Women Q1 = 1.2 Q2 = 3.0 Q3 = 5.0 Q4 = 7.8 Q5 = 14.0	Women 1.00 0.84 (0.66-1.05) 0.81 (0.64-1.02) 1.01 (0.80-1.27) 0.82 (0.64-1.05)	0.479	
			Fibre from Vegetables Men Q1 = 3 Q2 = 5.1 Q3 = 7.2 Q4 = 10.4 Q5 = 18.4	Men 1.00 0.90 (0.74-1.08) 0.83 (0.69-1.01) 0.91 (0.75-1.10) 0.78 (0.62-0.97)	0.052	
			Women Q1 = 3.0 Q2 = 5.0 Q3 = 7.0 Q4 = 10.0 Q5 = 17.2	Women 1.00 0.91 (0.74-1.14) 0.85 (0.68-1.05) 0.92 (0.74-1.14) 0.95 (0.75-1.20)	0.767	
			Fibre from Grains Men Q1 = 2.8 Q2 = 4.7 Q3 = 6.7 Q4 = 9.5 Q5 = 15.6	Men 1.00 0.84 (0.69-1.02) 0.86 (0.70-1.05) 0.92 (0.75-1.13) 0.86 (0.69-1.07)	0.479	
			Women Q1 = 2.4	Women 1.00	0.675	

	Q2 = 4.2 Q3 = 6.0 Q4 = 8.5 Q5 = 14.0	1.06 (0.84-1.33) 1.17 (0.93-1.46) 1.02 (0.80-1.28) 1.0 (0.78-1.27)	

Reference	Subject population	Measure of exposure	Fibre intake/level	Adjusted Relative Risk or Odds Ratio	P trend	Factors adjusted for in analysis
Schatzkin et al (2007) Prospective	291,988 men 197,623 women 2974 cases of incident CRC	AOAC defined dietary fibre by FFQ	g/1,000 kcal/day Fibre from fruit Q1 = 0.5 Q2 = 1.2	1.00	0.14	Sex, physical activity, smoking, menopausal HRT use, and intakes of red meat, dietary calcium, dietary folate, and total energy
NIH-AARP	5 y follow up		$Q_2 = 1.2$ $Q_3 = 2.0$ $Q_4 = 2.9$ $Q_5 = 4.8$	0.91 (0.81-1.03) 0.90 (0.80-1.02) 1.08 (0.95-1.23)		
			Fibre from Vegetables Q1 = 1.7 Q2 = 2.5 Q3 = 3.2 Q4 = 4.2 Q5 = 6.0	1.00 0.91 (0.81-1.02) 0.93 (0.83-1.04) 0.92 (0.81-1.03) 1.01 (0.89-1.15)	0.7	
			Fibre from Grains Q1 = 1.7 Q2 = 2.5 Q3 = 3.2 Q4 = 4.0 Q5 = 5.7	1.00 1.03 (0.92, 1.15) 0.94 (0.83, 1.05) 0.94 (0.83, 1.06) 0.86 (0.76, 0.98)	0.01	
Terry et al (2001)	61 463 women 460 cases of incident CRC	AOAC defined dietary fibre by FFQ	g/day standardized to average caloric intake			Age, BMI, educational level, intake of energy, and quartiles of alcohol, red meat, total fat, folic acid, vitamin D,
Prospective cohort Swedish Mammography Cohort	9.6 y follow up		Fibre from Fruits Q1 = 0.8 Q2 = 2.0 Q3 = 3.1 Q4 = 5.2	1.00 0.92 (0.70-1.20) 0.97 (0.72-1.31) 0.97 (0.69-1.38)	0.93	vitamin C, and calcium
Sweden			Fibre from Vegetables Q1 = 0.6 Q2 = 1.0 Q3 = 1.5 Q4 = 2.5	1.00 0.89 (0.68-1.18) 1.08 (0.81-1.44) 1.17 (0.85-1.61)	0.22	
			Fibre from Grains Q1 = 5.4 Q2 = 8.3 Q3 = 10.4	1.00 0.80 (0.60;1.07) 1.20 (0.93;1.56)	0.82	

		Q4 = 13.6	0.91 (0.69;1.20)	

Reference	Subject population	Measure of exposure	Fibre intake/level	Adjusted Relative Risk or Odds Ratio	P trend	Factors adjusted for in analysis
Wakai et al (2007) Prospective cohort Japan Multi-	43,115 men and women 443 cases of incident CRC 7.6 y follow up	AOAC defined dietary fibre by FFQ	g/day Fibre from Fruit Q1 = 0.4 Q2 = 1.0 Q3 = 1.7 Q4 = 2.2	1.00 1.05 (0.78-1.40) 1.23 (0.92-1.64) 1.06 (0.78-1.43)	0.55	Age, sex, area, educational level, family history of CRC, alcohol consumption, smoking, BMI, daily walking habits, exercise, sedentary work, beef intake, pork intake, energy intake, and energy-adjusted intakes of folate, calcium, and vitamin D
Study			Fibre from Vegetables $Q1 = 2.0$	1.00	0.65	
Japan			$Q^2 = 3.1$ $Q^3 = 4.0$ $Q^4 = 5.1$	0.81 (0.60-1.08) 0.86 (0.64-1.16) 0.89 (0.65-1.24)		
Willett et al (1990)	88,751 women	AOAC defined dietary fibre by	g/day standardized to average caloric intake			Age and energy
Prospective	150 cases of incident CRC	FFQ	Fibre from Fruit			
cohort Nurses Health Study	6 y follow up		Q1 <0.8 Q2 = 0.8-1.6 Q3 = 1.7-2.6 Q4 = 2.7-4.0 Q5 >4.1	1.00 0.94 (0.57-1.55) 0.87 (0.53-1.44) 0.81 (0.49-1.34) 0.62 (0.37-1.05)	0.12	
USA			Fibre from Vegetables Q1 <3.8 Q2 = 3.8-5.1 Q3 = 5.2-6.8 Q4 = 6.9-9.3 Q5 >9.4	1.00 1.02 (0.61-1.71) 1.03 (0.62-1.72) 0.78 (0.43-1.34) 1.07 (0.65-1.76)	0.87	
			Fibre from Grains Q1 <2.8 Q2 = 2.8-3.9 Q3 = 4.0-5.2 Q4 = 5.3-7.0 Q5 >7.1	1.00 0.72 (0.43-1.21) 0.58 (0.34-1.00) 1.04 (0.65-1.64) 0.74 (0.43-1.21)	0.62	

Reference	Subject population	Measure of exposure	Fibre intake/level	Adjusted Relative Risk or Odds Ratio	P trend	Factors adjusted for in analysis
Pietinen et al (1999)	29,133 male smokers	Englyst defined dietary fibre by	g/day			Smoking years, BMI, alcohol, education, physical activity at work,
Prospective cohort		FFQ	Insoluble fibre			and calcium intake
	185 cases of		Q1 = 12.2	1.0	0.73	
Alpha Tocopherol Beta	incident colon		Q2 = 17.2	1.2 (0.8-1.8)		
Carotene (ATBC) trial	cancer		Q3 = 21.0	1.1 (0.7-1.6)		
			Q4 = 27.1	1.0 (0.6-1.5)		
Finland	8 y follow up					
			Soluble fibre		0.01	
			Q1 = 3.7	1.0	0.91	
			$Q_2 = 5.0$	1.0 (0.7-1.6)		
			$Q_3 = 5.9$	1.0(0.6-1.5) 1.1(0.7,1.6)		
			Q4 = 7.5	1.1 (0.7-1.0)		
Wakai et al (2007)	43,115 men and	AOAC defined	g/dav			Age, sex, area, educational level,
	women	dietary fibre by	5, 1			family history of CRC, alcohol
Prospective cohort		FFQ	Insoluble fibre			consumption, smoking, BMI, daily
	443 cases of		Q1 = 5.3	1.00	0.041	walking habits, exercise, sedentary
Japan Multi-Centered	incident CRC		Q2 = 7.0	1.06 (0.80-1.40)		work, beef intake, pork intake, energy
Study			Q3 = 8.2	0.77 (0.56-1.05)		intake, and energy-adjusted intakes of
	7.6 y follow up		Q4 = 9.6	0.77 (0.55-1.08)		folate, calcium, and vitamin D
Japan						
			Soluble fibre	1 00		
			Q1 = 1.2	1.00	0.022	
			$Q_2 = 1.7$	0.85(0.64-1.14)		
			$Q_3 = 2.1$	0.70(0.55-1.04)		
			$Q^{4} = 2.0$	0.07 (0.47-0.95)		
	1	1			1	1

Table 7C: Prospective studies investigating the association of soluble or insoluble fibre with colorectal cancer

Appendix D: Colonic Function

 $\label{eq:table_$

Table 2D: Intervention trials investigating the effect of fibre from grain,

vegetable or isolated polysaccharide intake on colonic function

Table 3D: Intervention trials investigating the effect of resistant starch andcolonic function

Table 4D: Intervention trials investigating the effect of oligosaccharides andcolonic function

Table 5D: Intervention trials investigating the effect of mixed diets

Reference	Subjects N (M / F)	Duration of study	Duration of faecal collection	Dietary assessment	Fibre intake (mean ± SD g/d)	Faecal wt (mean ± SD g/d)	Correlation Coefficients	Comments
Birkett et al (1997)	53 (16 Men / 37 Women) 18-67 years Mean 29 ± 13 years	7d	3d	7d weighed	Total NSP: 14 ± 7 Insol: 8 ± 5 Sol: 6 ± 2 RS: 5 ± 2	Mean 127 ± 70 Range 41-340	NSP r= 0.49 AOAC fibre r=0.54 NSP and RS r= 0.44	Faecal weight related to NSP intake, but not starch or RS alone. Individuals consuming more NSP had faster transit times
Davies et al (1986)	51 17 (10 men / 7 women) in each of omnivores, vegetarians and vegans 35, 34 and 31 years (mean)	7d	7d	7d weighed	Total DF (Southgate)	Omnivores 153 ± 79 Range 54-415 Vegetarians 168 ± 56 Range 81-265 Vegans 225 ± 91 Range 129- 499	Total fibre r= 0.96 Cereal fibre r= 0.93 vegetable fibre r= 0.87 Fruit fibre r=0.78	Dietary Fibre correlated with faecal weight, especially cereal fibre (see correlation coefficients) Dietary Fibre significantly increased mean transit time decreased, stool frequency increased and the stools became softer.
Cummings et al (1992)	220 healthy UK adults		weeks		NSP quantified	Median daily stool weight 106 104 men 99 women	NSP r=0.84	NSP content showed a significant correlation between fibre intake and mean daily stool weight (r=0.84)

Table 1D: Observational studies investigating the association of `dietary fibre' and colonic function

Insol: Insoluble; Sol: Soluble; RS: resistant starch; NSP: non-starch polysaccharide

Table 2D: Intervention trials investigating the effect of fibre from grain, vegetable or isolated polysaccharide intake on colonic function

Year	Subjects	Fibre source	Duration of study	Dose (g/d)	How given	Diets	Duration of faecal collection	Faecal weight	Increase g/g fibre	Comments
Grains		1				•	•			
(i) Whea	at fibre									
Vuksan (1999)	24 (12 men/ 12 women) 21-60 years Mean 31 ± 2y	Wheat supplement - Fibrotein is produced by amylolytic digestion of wheat in making ethanol as	14 d periods Crossover design	21 fibrotein, 21 AACC wheat bran (positive control) 1.7 low-fibre control supplement consisting of crushed corn flakes (Negative	Flake form, on cereal or yogurt	Usual diets	4d	Control: 165.6 ± 16 (SE) +Wheat: 216.7 ± 19 +Fibrotein: 239.5 ± 19	Wheat 2.4 Fibrotein 3.5	The test supplement and the positive control (wheat bran) increased faecal bulk significantly compare to negative control (P≤0.01)
(ii) Rve		luel)		controly		1				
Grasten (2000)	17 (9 women/ 8 men) 28-51 years Mean 40.6y	Rye	4 wk periods Crossover design	13.5 increase in DF	Rye bread, versus wheat bread	Usual diets replaced with rye or wheat breads	5d	Females Control: $151 \pm 63 (SD)$ +Fibre: 203 ± 58 Males: Control: 198 $\pm 61 (SD)$ +Fibre: 335 ± 921	F: 3.9 M: 10.19 Both: 6.8	Faecal weight was significantly greater during rye bread period than wheat bread period in both women and men ($P < 0.05$) Mean intestinal transit time significantly shorter during rye bread period in men ($P < 0.05$)
McIntosh (2003)	28 (Men) 40-65years	Rye vs Wheat	4 wk periods Crossover design	Low fibre 19 Rye 32 Wheat 32	Incorporated into a variety of grain products	Usual diets	Not given "collected end of periods"	Con: 203 ± 18 (SEM) + Rye 278 ± 16 + Wheat 257 ± 21	Rye: 5.8 Wheat: 4.2	Both high-fibre rye and wheat foods increased faecal output by $33-36\%$ ($P = 0.004$)

(iii) Barl	ey									
Bird et al (2008)	18 (10 men / 8 women) 31-66 years Mean 55.9±2.0y	Barley vs wheat	4 wk periods Crossover design	low fibre 21.4, wheat 32.4, barley 44.6	Incorporated into a variety of grain products	Usual diets	48h	Control: 150 ± 19 (SEM) + Wheat 187 ± 25 + Barley 200 ± 22	Wheat: 3.4 Barley: 2.3	High amylose barley variety (Himalaya 292, novel hull- less) - more RS than normal barley Faecal output greater for Himalaya 292 than refined cereal diets (p<0.05) 33% higher faecal weight 91% higher butyrate excretion
Li (2003)	10 (Women) Mean 20.4 ± 1.3y	Barley	4 wk periods Crossover design	9.4 increase	Incorporated into foods	Meals provided	4 wk	Control: 95 ± 26 (SD) +Fibre 140 ± 24	4.8	Barley intake significantly increased stool volume compared to control (p<0.05).
Lupton et al (1993)	22 adults 20-64 years	Barley Bran flour vs cellulose	5 d control 7 d test Crossover design	20 cellulose, 30 barley bran flour (21 fibre)	NCEP Step 1 diet	NCEP Step 1 diet	5d	Control: 150.7 ± 12 (SD) +Fibre: 147.5 ± 1.2 Control: 123.6 ± 8 (SD) +Fibre: 172.2 ± 12	Cellulose: 0.2 Barley bran: 2.3	Significant increase in faecal weight (p=0.0001) The group that consumed barley bran flour significantly decreased transit time by 8.02 hours from baseline
(iv) Vari	ous fibre ty	pes				1	1	•		
Cherbut et al (1997)	9 adults 24-48 years	Potato Fibre, Maize fibre	1 month periods Crossover design	15 DF (22g source)		Control 15g sucrose. Subjects given ingredient s to prepare food at home	7d	Potato Control: 79.3 ± 9.7 (SEM) Range 44- 115 +Fibre: 115.7 ± 9.1 Range 59-141 Maize Control: 72.7 ± 7.4 Range 23-93 +Fibre:	Potato: 2.4 Maize 2.4	Increase in faecal weight compared to control (p=0.011) Significant increase in faecal weight (p=0.007)

							108.4 ± 5.4 Range 89-142		
Castaglia- Delavand et al (1998)	9 (Men) Mean age 21.5y	Sugar beet fibre Chicory inulin	28 d periods Crossover design	50	Controlled diets	8d	Control: 129 +sugar beet fibre: 202 Chicory inulin: 204 Pooled SEM =16	1.5	Increase in faecal weight with sugar beet fibre (p<0.05) Increase in faecal weight with chicory inulin (p<0.01)

Dahl et al (2005)	26 adults (7 men, 19 women) 18-60 years	Flax fiber (14 subjects) vs psyllium (12 subjects)	3 wk control 2 wk test Parallel design	Flax: 15 DF intended - 9.0 taken Psyllium: 15 DF intended - 10.4 g taken		Usual diets, but assigned breakfasts	3 wk control, 2 wk test	Flax Control: 162.1 ± 89.1 (SD) +Fibre: 188.6 ± 101.6 Psyllium Control: 124.9 ± 92.0 (SD) +Fibre: 175.3 ± 138.7	Flax: 2.9 Psyllium: 4.8	Significant increase in faecal weight Flax (p<0.05) Psyllium (P<0.005)
Jenkins (2000)	25 (13 men/ 12 women) 22-57 years Mean 37 ± 2y	Cocoa bran	2 wk periods Plus 2 week washout Crossover design	19.4	Incorporated into breakfast cereal and low fibre breakfast cereal	Usual diets	4d	Control: 135 ± 10 (SE) +Fibre: 191 ± 16	2.9	Included 19 subjects who had had AACC wheat bran (+ve control) 1-2 y before Mean faecal output was significantly higher for cocoa- bran than for low- fibre cereal (P<.001) but faecal weight was not significantly different (p=0.21)
Johnson et al (2006)	38 (Men) 24-64 years Mean 41 ± 2y	Lupin kernel	28 d periods Crossover design	22.2 increase in DF	Lupin fibre (LK fibre in various foods)	Usual diets + products provided	3d	Control: 172 ± 11 (SE) + lupin Fibre: 208 ± 14	1.6	Significant increase in faecal output (P=0.02) Significant increase in butyrate concentration by 16% (p=0.006) 40% Increase in butyrate output (p=0.002)

Spiller et al (2003)	13 (7 Men / 6 women) 27-65 years	Sun-dried raisins	3 wk periods Crossover design	8.4 (60-80g/kg given as DF in raisins)	Usual diets	4d	Control: 132 ± 68 (SD) +Fibre: 177 ± 78	4.1	Exact fibre content of raisins not given - raisins given as food hence contained other components
									Intestinal transit time decreased from 42 h on the baseline diet to 28 h on sun-dried raisins ($P < 0.05$). sun-dried raisins increased faecal wet weight ($P < 0.05$),
									Increase in butyrate excretion (p=0.02)

(v)Isolated	polysaccha	rides								
Chen (2006)	8 (1Man/ 7women) 21-54 years Mean 35 ± 5y	Konjac glucomannan	3 wk periods Crossover design	4.5	In water 3 x 1.5g/d	Diets provided 7d menu of low fibre Chinese food.	7d	Control: 132 ± 26.5 (SE) +Fibre: 171.8 ± 29.3	8.8	Significantly increased faecal weight (p<0.05)
Chaplin et al (2000)	16 (6men / 10 women) 22-36 years	Ispaghula (psyllium)	12 wks 2 wk pre- treatment 8 wk treatment 2 wk post- treatment	3.5 fybogel (2.94 NSP)		Usual diets	One 24h stool collection each week for 8 wks	Pre-treatment: 121 Range 93-153 +End treatment: 147 Range 116-183	8.8	Not continuous collection Difference in faecal weight not significant (p>0.05)
Daly et al (1993)	18 (Men) 18-34 years	Xanthan gum	10 d periods Crossover design	15 DF		Not given	10d	Control: 190.3 ± 13.6 (SE) +Fibre: 242.8 ± 15.8	3.5	Significant increases in faecal weight (P < 0.001) Increased SCFA production

									(p=0.049)
Robinson et al (2001)	22 (11 men / 11	Arabingalact an (LAREX	3 wk periods	15 or 30 AG	Usual diet AG powder	5d	Control: 136.9 ± 7.8 (SE)	15g: 0.5	No effect on stool weight (p=0.37)
	women)	from Larch	Crossover design		in drinks		+15a AG'	30a.	No significant
			deorgin				130.1 ± 7.8	0.3	changes in SCFA Total SCFA
							+30g AG:		(p=0.41)
							126.9 ± 7.8		Proprionate
									(p=0.31)

Table 3D: Intervention trials investigating the effect of resistant starch and colonic function

Reference	Subjects	Duration of study	Fibre source	Dose (g/d)	How given	Diets	Duration of faecal collection	Faecal weight (g)	Increase g/g fibre	Comments
Behall et al (2002)	24 (Men) 28-58 years Mean 41y	14 wk periods 4 wks controlled Crossover design	High amylose cornstarch	RS intake: Control 1 Test 30	Added to baked products	Usual 10wk, controlled 4 wk of each period	7d	Control: 246 ± 14.5 (SE) +RS: 269.4 ± 14.5	0.8	Study investigated mineral balances, with resistant starch on colonic function. No significant relationships or p values relating to colonic function reported.
Grubben et al (2001)	24 (13Men / 11women) 25-75 years Patients with adenomas removed	4 wk periods, after 4wk maltodextrin for all Crossover design	Hylon VII	RS₂: 45g	Powder 3 x 15g/d	Usual	2d	2 day weights Control: 303 ± 43 (SE) +RS: 331 ± 43 vs control Control: 316 ± 30 Change -36 ± 41 (280)	0.3	Adenoma patients, not healthy controls; usual diet had higher fibre than representative Australian population. No significant increase in faecal weight (p=0.235)
Jenkins et al (1998)	24 (12men / 12 women) 22-53 years Mean 33 ± 2y	2 wk periods Crossover Design	RS ₂ , RS ₃ or wheat bran (positive control)	Wheat bran: 30 (TDF) RS₂: 30 (TDF) 21.5 (RS) RS₃: 30 (TDF) 27.9 (RS)	In cereals and muffins	Usual	4d	Control: 163 ± 23 (SE) +Wheat bran: 258 ± 22 +RS2: 187 ± 24 +RS3 182 ± 23	Wheat bran: 3.2 TDF RS ₂ : 1.1 RS ₃ : 0.7	The wheat bran supplement increased faecal bulk compared with the low-fibre control (p<0.001) with the mean for both resistant starches also being greater than the low-fibre control (p=0.013).
Muir et al (2004)	20 (11 men / 9	3 wk periods	Hi-maize with and	19.8 RS 11.6/11.2g	In foods	Other foods	5d	Control: 131 ± 64 (SD)	WB: 2.6	
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	women)		without wheat	DF from WB		provided		+Wheat bran:	RS: 2.2	Significant increase in faecal weight (p<0.001) for wheat bran plus RS diet.
	Mean 42 ± 2v		bran					161 ± 67		
	_,							+WB+RS: 204 ± 84		

TDF: total dietary fibre; RS: resistant starch

Table 4D: Intervention trials investigating the effect of oligosaccharides and colonic function

Reference	Subjects	Fibre source	Duration of study	Dose	How given	Diets	Duration of faecal collection	Faecal weight (g)	Increase g/g fibre	Results/Comments
Alles et al (1999)	40 (22 men / 18 women) 3 groups mean ages: 37.8 ± 17.6 years 36.5 ± 17.6 years 42.9 ± 14.8 years	Trans-galacto- oligosaccharides (Elix'or)	3 wk periods Parallel design Placebo, low, or high doses	7.5g or 15g TOS	In fruit juice	Controll ed - 21 menus for 3 wk	21d	Control: $147 \pm 11 (SE)$ +Placebo: 139 ± 14 Control: 113 ± 12 +7.5g TOS: 127 ± 14 Control: 146 ± 22 +15g TOS: 142 ± 18	7.5gTOS: 1.9 15g TOS: - 0.3	Author reported that TOS did not significantly affect bowel habits, stool composition or the concentration of SCFA. No P values reported
Bouhnik et al (2007)	12 (6 men / 6 women) Mean 69 ± 2y Elderly	Short chain Fructo- oligosaccharides (sc-FOS)	2 wk periods, crossover design	8g	2 x 4g doses, one each breakfast and dinner	Usual	2d	Control: 155.4 ± 20.9 (SE) +FOS: 137.7 ± 17.3	-2.2	Faecal bifidobacteria counts significantly increased during sc-FOS period (p<0.05) Faecal weight decreased No change in transit time, no p value reported

Brighenti et al (1999)	12 (men) Mean 23.3 ± 0.5 years	FOS/Inulin mixture (Fibruline)	4 wk periods All control first, test second	9g	In breakfast cereal	Usual	3d Subjects weighed own stools	Control: 159.3 ± 17.6 (SE) +FOS: 155.6 ± 13.8	-0.4	No significant change in SCFA No significant change in faecal weight. no p values given
Causey et al (2000)	12 (Men) 27-49 years	Inulin	3 wk each diet Cross over design with no washout period	20g	In ice cream	Controll ed diet, rotating menu	5d	Control: 150.3 ± 54.0 (SD) +Inulin: 164.3 ± 56.8	0.72	Transit time did not differ significantly ($p=0.33$) but decreased in inulin phase. No significant change in SCFA (Total SCFA $p=0.13$, butyrate p= 0.12, propionate $p=0.14$) No significant change in faecal weight ($P=0.20$)

Chen et al (2001)	7 (Men) Mean 75.2 ± 4.0 years Constipated elderly (age 60+) men	Isomalto- oligosaccharide (IO)	4 wk periods Control first Test second	10g active compone nt	In afternoon dessert	Constan t (nursing home)	5d	Control: 47.7 ± 81.1 (SE) +isomalto- oligosacc: 81.1 ± 1.5	3.7	Faecal weight increased significantly with IO supplement. p values not reported Significantly increased faecal proprionate concentration (p<0.05) but not butyrate concentration (p value not reported).
Den Hond et al (2000)	6 (1 man / 5 women) 20-49 years Mean 28.5 ± 11.1 years	Inulin (Raftilene)	1 wk periods 1 wk washout	15g	Combined with meals	Usual, but constan t advised	3d	Control: 91 ± 107 (SE) +Inulin: 113 ± 22	1.5	Significant increase in stool frequency with inulin (p=0.02) but not faecal weight (p=0.28)
Molis et al (1996)	6 (3 men / 3 women) 20-27 years	FOS (Actilight)	Control 6d Test 11d Crossover design With 1 week washout	20.1g FOS	Powder with meals	Controll ed diet	2d	Control: 210 ± 12.6 (SE) +FOS: 222 ± 29.7	0.6	No significant difference in faecal weight No p values reported

Scholtens et al (2006)	12 (6 men/ 6 women) 18-35 years	FOS (Raftilene) or maltodextrin (Control)	2 wk periods, crossover design	25-30g dependent on body weight	As powder in sachets	Usual	3d	Control: 174 ± 19.3 (SE) +FOS: 225 ± 19.3	1-7-2.0	Not stated how many subjects on each dose - hence range of effect 1.9-2.0 g/g (divided by 25 and by 30) Total stool output was higher in the FOS period, (p=0.097) Increased acetate production and decreased butyrate production.
Sairanen et al (2007)	66 (22 men/ 44 women) 3 groups mean ages: 41.3 (22-59)y 37.0 (22-54)y 41.4 (23-60)y	Inulin (Raftilene)	3 wk periods Parallel design Control Probiotic Probiotic + inulin	12g	In milk	Usual	5d	5d results Control: 829 ± 385 (SD) +Milk: 935 ± 432 Control: 735 ± 428 (SD) +Probiotic: 781 ± 441 Control: 746 ± 372 (SD) +Probiotic+inul in: 861 ± 340	1.9	Results given per 5 days; inulin is with probiotic, not alone No significant increase in faecal weight (p=0.566)
Ten Bruggencate (2006)	34 (Men) 29 for faecal data 18-55 years	FOS (Raftilose)	2 wk periods - 1 week between, crossover Design	20g FOS	In lemonade: dose divided into 3 portions, morning, afternoon, evening	Usual	24h	Control: 203.7 ± 28.3 (SE) +FOS: 268.0 ± 29.4	3.2	n=29 for faecal data. 24h collection only

Table 5D: Intervention trials inve	stigating the effect of mixed diets
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Reference	Subjects	Duration of study	Fibre source	Dose (g/d)	Diets	Duration of faecal collection	Faecal weight	Increase g/g fibre	Comments
Haack et al (1998)	9 (Men) Mean 22 ± 1y	28d periods, in order of increasing fibre	Mixed diets: low, medium and high fibre	Low 15.7 Medium 30.2 High 41.8	Provided Mixed fruit, veg, grains and legumes	18d	Low 109 ± 15 (SEM) Medium Fibre 156 ± 12 High Fibre 195 ± 17	Med 3.2 High 3.3	Increase faecal weight (no P value reported)
Hovey et al (2003)	12 (4 men/ 8 women) 31 years mean	7d periods	Mixed diets	Average DF intake 83.4	Usual then mixed diet of wholegrains - linseeds, sunflower, sesame, wheat grains, beans, chickpeas prepared, whole or ground.	3d	Usual 125 ± 46 (SD) Range 40-184 Whole grain 258 ± 123.2 Range 106-562 Ground grain 170.5 ± 63 Range 42-288	Whole 1.6 Ground 0.5	Consumption of intact seeds compared to ground seed significantly increased faecal weight (p=0.005)
Jenkins et al (2001)	10 (8 men / 2 women) 24-60 years Mean 38 ± 4y	14d periods Crossover	Mixed diets: high starch high vegetable	Control 25.1 High starch 45.4 High vegetable 149.6	Provided Starch diet: grains, legumes, bread etc + some fruit and veg. Veg diet: only fruit and veg, no grains or legumes	3d	Control 172 ± 28 (SEM) High starch 279 ± 27 High vegetable 906 ± 130	High starch 5.3 High vegetable 5.9	Significant increase in faecal weight (P=0.000).

Nagengast	22	4wk low,	Test mixed diet	Low fibre usual 20.3	Usual (7d diam()	3d	Low usual	3.5 (addition	Increased faceal
et al (1993)	12 (6 men/ 6	IUWK LESL	fruit and	High fibre usual 31.1			100 ± 12 (3LM)	of mixed fibre)	weight (p<0.05)
	women) on test		vegetables vs usual				Test high fibre		
	10 (5 men / 5		low fibre control vs				156 ± 11		No effect on transit
	women) on high		usual high fibre						time
	fibre usual						High fibre usual		
							initial 174 ± 22		
							final 163 ± 18		

Appendix E: Prebiotic Studies

Table 1E: Relevant studies with oligosaccharides attempting to confirm a prebiotic effect

Table 2E: Studies with prebiotics in infants

Table 1E: Relevant studies with oligosaccharides attempting to confirm a prebiotic effect

Reference	Subjects	Substrate	Dose	Duration	Results
Lactulose					
Ballongue et al (1997)	12 humans	Lactulose	2 x 10g/d	4 weeks	Bifidobacteria, streptococci and lactobacilli significantly increased ($P < 0.01$ for all) compared to placebo whilst bacteroides, clostridia, coliforms and eubacteria significantly decreased ($P < 0.01$ for all)
Bouhnik et al (2004a)	65 volunteers	Lactulose	a) 20 g/day then b) (10-30 g/day) then c) (10-30 g/day)	4w a) 1 week + b) 1 week + c) 2 weeks	An increase in faecal bifidobacteria counts ($P = 0.04$) and beta-galactosidase activity ($P < 0.001$) was observed
Bouhnik et al (2004b)	16 healthy volunteers	Lactulose	5g/d	бw	Lactulose ingestion led to a significant increase in faecal bifidobacteria counts when compared to placebo $(P = 0.03)$
Terada et al (1992)	8 humans	Lactulose	3g/d	14d	Bifidobacteria significantly increased (P<0.001) while clostridia incl. <i>Cl.perfringens</i> (P<0.05), bacteroides (P<0.01), streptococci (P<0.05) and Enterobacteriaceae decreased (P<0.05 at 7d, not significant at 14d)
Tuohy et al (2002)	30 adults	Lactulose	10g/d	3w	Significantly increased bifidobacteria on Raffinose- Bifidobacterium agar (P=0.03). Non significant increase on Beerens' agar.
Fructo-oligosacch	narides (FOS)				
Bouhnik et al (1996)	20 adults	FOS	12.5g/d	12d	Significant increase in bifidobacteria ($P < 0.01$) by about 10 times was demonstrated on selective agars
Bouhnik et al (1999)	40 adults	FOS	0, 2.5, 5, 10 and 20g/d	7d	Selective agars showed that bifidobacteria were most increased by 10g (P = 0.02) and 20g (P = 0.01) doses of FOS and increases at these doses where significantly higher compared to 2.5g and placebo (P < 0.05). The optimum dose of oligofructose was found to be 10g/d
Bouhnik et al (2007)	12 elderly persons	scFOS	8g/d	4w	Bifidobacterial counts were significantly increased (P < 0.05)
Buddington et al (1996)	12 adults	Neosugar	4g/d (in controlled diet)	25d	Significant increase in anaerobes (P<0.0003), and

					bifidobacteria ($P < 0.03$) and non significant increase in enterobacteria.
de Preter et al (2008)	19 adults	FOS enriched inulin	10g/d	4w	Total faecal bifidobacteria increased significantly after intake of FOS ensriched inulin ($p < 0.001$)

Gibson et al (1995)	8 adults	FOS and inulin	15g/d (in controlled diet)	15d	FOS and inulin both significantly increased bifidobacteria (P<0.01 and P=0.0002 respectively). FOS significantly reduced bacteroides (P<0.01), clostridia (P<0.05) and fusobacteria (P<0.01).
Guigoz et al (2002)	19 elderly persons	FOS	8g/d	3w	Increase in faecal bifidobacteria of approximately 2.8 log cfu/g of faeces compared to baseline (P < 0.001)
Harmsen et al (2002)	14 adult volunteers	Inulin	9g/d	2w	Quantification of all bacteria, bifidobacteria, the <i>Eubacterium rectale–Clostridium coccoides</i> (Erec) group, <i>Bacteroides</i> , and eubacteria were counted with FISH probes. A significant increase in bifidobacteria (P < 0.05) and a significant decrease in Erec group was observed (P < 0.05).
Hidaka et al (1986)	23 Senile adults	Neosugar	8g/d	14d	Significantly increased bifidobacteria (P<0.01)
Hidaka et al (1986)	2 adults	Neosugar	8g/d	2 months	Increase in bifidobacteria, Reduction in SCFA and Putrefaction [DN: Only 2 subjects]
Kleesen et al (1997)	10 senile adults	Inulin	20g/d, then 40g/d	8d, then 11d	Significant increase in bifidobacteria (P<0.05). For 40g/d, significant reduction in enterococci (P<0.01) and non-significant decreases in bacteroides and enterobacteria
Kolida et al (2007)	30 adults	Inulin	5 and 8g/d	2w	Bifidobacterial levels increased significantly upon ingestion of both the low ($P < 0.05$) and high inulin dose ($P = 0.05$) compared to placebo.
Kruse et al (1999)	8 persons	Inulin	up to 34g/d (based on individual energy requirements)	2 months	FISH revealed a significant increase in bifidobacteria from 9.8 to 11.0 \log_{10} cells/g dry faeces. The effect lasted for the whole 2 months period that the volunteers received the prebiotic. [DN Paper states "significant increase", but no p-value given. However, graph with error bars indicates statistical significance]
Kleesen et al (2007)	45 adult volunteers	Jerusalem artichoke or chicory inulin in snack bars	7.7g/d then 15.4g/d	7d then 14d	Significant increase in bifidobacteria (p<0.05) and significant reductions in <i>Bacteroides/Prevotella</i> (P<0.05) i and the <i>Clostridium histolyticum/C. lituseburense</i> group (p<0.05)
Menne et al (2000)	8 persons	Chicory inulin hydrolysate	8g/d	2w	Selective agars showed anincrease in faecal bifidobacteria ($P < 0.01$).
Mitsouka et al (1987)	23 adults	FOS	8g/d	2 weeks	Increase in faecal bifidobacteria by about 10x

					(P<0.005) and decrease in stool pH
Rao (2001)	8 young volunteers	FOS	5g/d	3w	By means of selective agars, an increase in faecal bifidobacteria was observed ($P < 0.001$)

Tuohy et al (2001a)	10 adults	HP-inulin	8g/d	2w	A small but statistically significant increase in bifidobacteria after 7 days (P<0.05) and non significant increase after 14 days		
Tuohy et al (2001b)	31 adults	Biscuits containing FOS and partially hydrolysed guar gum (PHGG)	6.6g FOS 3.4g PHGG	3w	FISH revealed a significant increase in faecal bifidobacteria compared to placebo (P = 2.5×10^{-5})		
Williams et al (1994)	10 adults	Neosugar	4g/d	14d	Significantly increased bifidobacteria (P < 0.05)		
Galacto-oligosaccharides (GOS)							
Bouhnik et al (1997)	8 adults	Trans-GOS	10g/d	21d	Significantly increased bifidobacteria (P < 0.05), significant reduction in breath H_2 (P < 0.01)		
Depeint et al (2008)	30 adults	Trans-GOS (BiMuno)	7.5	7d	Significant increase in bifidobacteria compared to pre-treatment ($P < 0.05$)		
Ito et al (1990)	12 men	Trans-GOS (Oligomate)	0, 2.5, 5 then 10g/d	1 week for each dose	Bifidobacteria increased with dose. For $10g/d$, significant increase in bifidobacteria (P < 0.001) and lactobacilli (P < 0.05)		
Ito et al (1993)	12 men	Trans-GOS	15g/d	6d	Significant increase in bifidobacteria (P < 0.01) and lactobacilli (P < 0.05), significantly lower bacteroides (P < 0.05). Significant decrease in indole (P < 0.01), p-cresol (P < 0.05), NH ₃ (P < 0.01), propionate (P < 0.05), valerate (P < 0.05), isovalerate (P < 0.01) and isobutyrate (P < 0.01).		
Tanaka et al (1983)	5 men	Trans-GOS	3g/d, then 10g/d	1 week for each dose	Non-significant increase in bifidobacteria and significant decrease in bacteroides ($P < 0.05$).		
Teuri et al (1998)	6 adults	Trans-GOS	15g/d	14d	Significant increase in total bacterial count on media for lactic acid bacteria (P=0.03). No change in bifidobacteria.		

Table 2E: Studies with prebiotics in infants

Reference	Test oligosaccharide	Study design	Dose	Evidence of prebiotic efficacy
Bakker-Zierikzee et al (2006)	GOS & FOS; Bifidobacterium animalis	3 groups of 19 healthy, formula fed infants, 63 breast fed (ref. group) randomised, double blind parallel, from birth to 16 wk	6g/l prebiotic/formula; 6×10 ¹⁰ viable <i>B. animalis</i> /l formula; standard formula	Similar metabolic activity of the flora in GOS/FOS group as breast fed (#), <i>B. animalis</i> group similar to standard formula (#)
Ben et al (2004)	GOS	69 healthy term infants fed GOS, parallel study, 59 fed formula, 124 mixed; 6 month intervention	2.4g/l prebiotic/formula; formula; mixed (breast fed & prebiotic formula)	Significant increases in bifidobacteria, lactobacilli & stool frequency in prebiotic and mixed groups (all P<0.05) but not the standard formula group (P≥0.05)
Boehm et al (2002)	GOS & FOS	19 Preterm infants on prebiotic, 19 maltodextrin placebo, 12 fortified breast milk parallel study, 28d intervention	10g/L prebiotic/formula (90% GOS)	Significantly higher bifidobacteria compared with placebo group (P=0.0008); significantly higher stool frequency as compared with placebo (P=0.0079)
Costalos et al (2008)	GOS & long chain FOS	Healthy bottle fed infants, randomised, double blind, parallel followed up to 6 wk of age	4g/l prebiotic/formula or standard formula (no prebiotic)	Significant decrease in clostridia (FISH) (#), trend of increased bifidobacteria (P=0.2) and <i>E. coli</i> (P=0.3), higher stool frequency and softer stools compared with control group
Haarman & Knol (2006) Haarman & Knol (2005)	GOS & FOS	2 groups of 10 healthy, formula fed infants 28-90d age, parallel study	8g/l prebiotic/formula (90% GOS); breast fed control group	Real time PCR analysis, similar flora composition between formula and breast fed infants (P≥0.05 for most lactobacillus strains)
Kim et al (2007) Native inulin		14, 12, 6 wk formula fed healthy infants, 6 wk intervention (3 wk inulin, 3 wk without)	0.25g/kg/d native inulin	Inulin significantly increased lactobacilli and bifidobacteria (both P<0.05), stool frequency not affected (#)
Mihatsch et al (2006)	GOS & FOS	20 preterm infants on enteral nutrition, assigned into 2 groups, placebo controlled, double blind, 14d supplementation	10g/l prebiotic/formula or standard formula	Significant reduction in gastrointestinal transit time (P=0.037) and trend towards higher stool frequency (P=0.059); well tolerated

Moro et al (2005)	GOS & FOS	Healthy formula fed infants, 28d feeding period	8g/l prebiotic/formula; maltodextrin control	Significantly higher bifidobacteria with prebiotic compared with control (P<0.001)
Savino et al (2006)	GOS & FOS	199 formula fed infants with colic, 96 prebiotic, aged >4 months, 103 standard formula parallel randomised, 2 wk	8g/l prebiotic/formula (90% GOS), formula & simethicone (6mg/kg)	Significant reduction in colic episodes (as diagnosed by ≥ 3 episodes of full-forced crying lasting ≥ 3 hrs on $\geq 3d$ per week) after 7 and 14d as compared to standard formula (P<0.0001 at both time points)
Scholtens et al (2006)	GOS & FOS	35 formula fed infants in weaning, aged 4-6 months, double blind, randomised, 6 wk supplementation	4.5g/d prebiotic in weaning food or weaning food (no prebiotic)	Significant increase in bifidobacteria % (FISH) with prebiotic compared with control (P=0.026)
Ziegler et al (2007)	GOS & polydextrose, lactulose	226 healthy formula fed term infants, assigned to treatment grps of 76 parallel design, followed up to 120d age	4g/l or 8g/l prebiotic/formula	Normal growth and stool characteristics similar to breast fed (P≥0.005)

7. Reference section

All references assessed in the HNR report are listed in the following section

References

Abrams SA, Griffin IJ, Hawthorne KM, Liang L, Gunn SK, Darlington G & Ellis KJ (2005) A combination of prebiotic short- and long-chain inulin-type fructans enhances calcium absorption and bone mineralization in young adolescents *Am J Clin Nutr* **82**, 471-476.

Alberts DS, Martinez ME, Roe DJ, Guillen-Rodriguez JM, Marshall JR, van Leeuwen JB et al. (2000) Lack of effect of a high-fiber cereal supplement on the recurrence of colorectal adenomas. Phoenix Colon Cancer Prevention Physicians' Network *N Engl J Med* **342** (16), 1156-1162.

Aller R, de Luis DA, Izaola O, La Calle F, del Olmo L, Fernandez L, Arranz T & Hernandez JM (2004) Effect of soluble fiber intake in lipid and glucose levels in healthy subjects: a randomised clinical trial *Diabetes Res Clin Pract* **65**, 7-11.

Alles MS, Hartemink R, Meyboom S, Harryvan JL, Van Laere KM, Nagengast FM, Hautvast JG. (1999) Effect of transgalactooligosaccharides on the composition of the human intestinal microflora and on putative risk markers for colon cancer. *Am J Clin Nutr* **69**, 980-91.

American Association of Cereal Chemists (2001) The Definition of Dietary Fiber. Report of the Dietary Fiber Definition Committee to the Board of Directors of AACC International *Cereal Foods World* **46**, 112-126.

Amos AF, McCarty DJ & Zimmet P (1997) The rising global burden of diabetes and its complications: estimates and projections to the year 2010 *Diabet Med* **14 Suppl 5**, S1-85

Anderson JW, Allgood LD, Turner J, Oeltgen PR & Daggy BP (1999) Effects of psyllium on glucose and serum lipid responses in men with type 2 diabetes and hypercholesterolemia *Am J Clin Nutr* **70**, 466-73.

Anderson JW, Davidson MH, Blonde L, Brown WV, Howard WJ, Ginsberg H, Allgood LD & Weingand KW (2000) Long-term cholesterol-lowering effects of psyllium as an adjunct to diet therapy in the treatment of hypercholesterolemia *Am J Clin Nutr* **71**, 1433-8.

Anderson JW, Gustafson NJ, Spencer DB, Tietyen J & Bryant CA (1990) Serum lipid response of hypercholesterolemic men to single and divided doses of canned beans *Am J Clin Nutr* **51**, 1013-9.

Andrieux C & Szylit O (1992) Effects of galacto-oligosaccharides (TOS) on bacterial enzyme activities and metabolite production in rats associated with a human faecal flora *Proc Nutr Soc* **51**, 7A.

Archer BJ, Johnson SK, Devereux HM & Baxter AL (2004) Effect of fat replacement by inulin or lupin-kernel fibre on sausage patty acceptability, post-meal perceptions of satiety and food intake in men Br J Nutr 91(4), 591-9.

Arvill A & Bodin L (1995) Effect of short-term ingestion of konjac glucomannan on serum cholesterol in healthy men *Am J Clin Nutr* **61**, 585-9.

Asano T, McLeod RS. (2002) Dietary fibre for the prevention of colorectal adenomas and carcinomas *Cochrane Database Syst Rev* (2), CD003430.

Ascherio A, Hennekens C, Willett WC, Sacks F, Rosner B, Manson J, Witteman J & Stampfer MJ (1996) Prospective study of nutritional factors, blood pressure, and hypertension among US women *Hypertension* 27, 1065-72.

Association of Official Analytic Chemists (AOAC) (1990) Official methods of analysis. 15[Sec.985.29]. Arlington, VA, AOAC.

Astrup A, Vrist E & Quaade F (1990) Dietary fibre added to very low calorie diet reduces hunger and alleviates constipation *Int J Obes* **14**(2), 105-12.

Bach Knudsen KE & Hessov I (1995) Recovery of inulin from Jerusalem artichoke (*Helianthus tuberosus* L.) in the small intestine of man *Br J Nutr* **74**, 101-13.

Bakker-Zierikzee AM, Tol EA, Kroes H, Alles MS, Kok FJ & Bindels JG (2006) Faecal SIgA secretion in infants fed on pre- or probiotic infant formula *Ped Allergy Immunol* **17**, 134-140, 2006.

Ballongue J, Schumann C & Quignon P (1997) Effects of lactulose and lactitol on colonic microflora and enzymatic activity *Scand J Gastroenterol* **32**, 41-44.

Barcenilla A, Pryde SE, Martin JC, Duncan SH, Stewart CS, Henderson C & Flint HJ (2000) Plylogenetic relationships of butyrate-producing bacteria from the human gut. *Appl Env Microbiol* **66**, 1654-1661.

Barroso Aranda J, Contreras F, Bagchi D & Preuss HG (2002) Efficacy of a novel chitosan formulation on fecal fat excretion: a double-blind, crossover, placebo-controlled study *J Med* **33**(1-4), 209-25.

Behall KM, Howe JC, Anderson RA (2002) Apparent mineral retention is similar in control and hyperinsulinemic men after consumption of high amylose cornstarch *J Nutr* **132**, 1886-91.

Behall KM & Hallfrisch J (2002) Plasma glucose and insulin reduction after consumption of breads varying in amylose content *Eur J Clin Nutr* **56**(9), 913-20.

Behall KM, Scholfield DJ & Hallfrisch J (2004) Diets containing barley significantly reduce lipids in mildly hypercholesterolemic men and women *Am J Clin Nutr* **80**, 1185-93.

Behall KM, Scholfield DJ & Hallfrisch J (2004) Lipids significantly reduced by diets containing barley in moderately hypercholesterolemic men *J Am Coll Nutr* **23**, 55-62.

Behall KM, Scholfield DJ & Hallfrisch J (2006) Whole-grain diets reduce blood pressure in mildly hypercholesterolemic men and women *J Am Diet Assoc* **106**, 1445-9.

Behall KM, Scholfield DJ, Hallfrisch JG & Liljeberg-Elmstahl HG (2006) Consumption of both resistant starch and beta-glucan improves postprandial plasma glucose and insulin in women *Diab Care* **29**(5), 976-81.

Ben XM, Zhou XY, Zhao WH, Yu WL, Pan W, Zhang WL, Wu SM, Van Beusekom CM & Schaafsma A (2004) Supplementation of milk formula with galacto-oligosaccharides improves intestinal micro-flora and fermentation in term infants *Chin Med J* **117**, 1268-1270.

Benini L, Castellani G, Brighenti F, Heaton KW, Brentegani MT, Casiraghi MC, Sembenini C, Pellegrini N, Fioretta A, Minniti G & et al. (1995) Gastric emptying of a solid meal is accelerated by the removal of dietary fibre naturally present in food *Gut* **36**(6), 825-30.

Benno Y, Sawada K & Mitsuoka T (1984) The intestinal microflora of infants: composition of fecal flora in breast-fed and bottle-fed infants *Microbiol Immunol* **28**, 975-986.

Bergmann JF, Chassany O, Petit A, Triki R, Caulin C & Segrestaa JM (1992) Correlation between echographic gastric emptying and appetite: influence of psyllium *Gut* **33**(8), 1042-3.

Berkey CS, Rockett HR, Field AE, Gillman MW, Frazier AL, Camargo CA Jr & Colditz GA (2000) Activity, dietary intake, and weight changes in a longitudinal study of preadolescent and adolescent boys and girls *Pediatrics* **105**(4), E56.

Bezkorovainy A (2001) Probiotics: determinants of survival and growth in the gut Am J Clin Nutr 73, 3998-4058.

Bingham SA, Day NE, Luben R, Ferrari P, Slimani N, Norat T et al (2003) Dietary fibre in food and protection against colorectal cancer in the European Prospective Investigation into Cancer and Nutrition (EPIC): an observational study *Lancet* **361** (9368), 1496-1501.

Bingham SA, Norat T, Moskal A, Ferrari P, Slimani N, Clavel-Chapelon F et al (2005) Is the association with fiber from foods in colorectal cancer confounded by folate intake? *Cancer Epidemiol Biomarkers Prev* **14** (6), 1552-1556.

Bird AR, Vuaran MS, King RA, Noakes M, Keogh J, Morell MK & Topping DL (2008) Wholegrain foods made from a novel high-amylose barley variety (Himalaya 292) improve indices of bowel health in human subjects *Br J Nutr* **99**, 1032-40.

Birkett AM, Jones GP, de Silva AM, Young GP & Muir JG (1997) Dietary intake and faecal excretion of carbohydrate by Australians: importance of achieving stool weights greater than 150 g to improve faecal markers relevant to colon cancer risk *Eur J Clin Nutr* **51**, 625-32.

Birketvedt GS, Aaseth J, Florholmen JR & Ryttig K (2000) Long-term effect of fibre supplement and reduced energy intake on body weight and blood lipids in overweight subjects *Acta Medica* (Hradec Kralove) **43**(4), 129-32.

Birketvedt GS, Shimshi M, Erling T & Florholmen J (2005) Experiences with three different fiber supplements in weight reduction *Med Sci Monit* **11**(1), PI5-8.

Blake DE, Hamblett CJ, Frost PG, Judd PA & Ellis PR (1997) Wheat bread supplemented with depolymerized guar gum reduces the plasma cholesterol concentration in hypercholesterolemic human subjects *Am J Clin Nutr* **65**, 107-13.

Boehm G, Lidestri M, Casetta P, Jelinek J, Negretti F, Stahl B & Marini A (2002) Supplementation of a bovine milk formula with an oligosaccharides mixture increases counts of faecal bifidobacteria in preterm infants *Arch Dis Children* **86**, F178-182.

Boeing H, Weisgerber UM, Jeckel A, Rose HJ & Kroke A (2000) Association between glycated hemoglobin and diet and other lifestyle factors in a nondiabetic population: cross-sectional evaluation of data from the Potsdam cohort of the European Prospective Investigation into Cancer and Nutrition Study *Am J Clin Nutr* **71**(5), 1115-22.

Bolton RP, Heaton KW & Burroughs LF (1981) The role of dietary fiber in satiety, glucose, and insulin: studies with fruit and fruit juice *Am J Clin Nutr* **34**(2), 211-7.

Bonithon-Kopp C, Kronborg O, Giacosa A, Rath U & Faivre J. (2000) Calcium and fibre supplementation in prevention of colorectal adenoma recurrence: a randomised intervention trial. European Cancer Prevention Organisation Study Group *Lancet* **356** (9238), 1300-1306.

Bouhnik Y, Achour L, Paineau D, Riottot M, Attar A & Bornet F (2007) Four-week short chain fructooligosaccharides ingestion leads to increasing fecal bifidobacteria and cholesterol excretion in healthy elderly volunteers *Nutr J* **6**, 42-46.

Bouhnik Y, Attar A, Joly FA, Riottot M, Dyard F & Flourié B (2004b) Lactulose ingestion increases faecal bifidobacterial counts: a randomised double-blind study in healthy humans *Eur J Clin Nutr* **58**, 462-466.

Bouhnik Y, Flourie B, D'Agay-Abensour L, Pochart P, Gramet G, Durand M & Rambaud JC (1997) Administration of *trans*galacto-oligosaccharides increases fecal bifidobacteria and modifies colonic fermentation metabolism in healthy humans *J Nutr* **127**, 444-448.

Bouhnik Y, Flourie B, Riottot M, Bisetti N, Gailing M, Guibert A, Bornet F and Rambaud, JC (1996) Effects of fructooligosaccharide ingestion on fecal bifidobacteria and selected metabolic indexes of colon carcinogenesis in humans *Nutr Cancer* **26**, 21-29.

Bouhnik Y, Neut C, Raskine L, Michel C, Riottot M, Andrieux C, Guillemot F, Dyard F & Flourié B. (2004a) Prospective, randomised, parallel-group trial to evaluate the effects of lactulose and polyethylene glycol-4000 on colonic flora in chronic idiopathic constipation *Alimentary Pharmacol Ther* **19**, 889-899.

Bouhnik Y, Vahedi K, Achour L, Attar A, Salfati J, Pochart P, Marteau P, Flourie B, Bornet F & Rambaud JC (1999) Short-chain fructo-oligosaccharide administration dose- dependently increases fecal bifidobacteria in healthy humans *J Nutr* **129**, 113-116.

Bourdon I, Yokoyama W, Davis P, Hudson C, Backus R, Richter D, Knuckles B & Schneeman BO (1999) Postprandial lipid, glucose, insulin, and cholecystokinin responses in men fed barley pasta enriched with betaglucan *Am J Clin Nutr* **69**(1), 55-63.

Bourdon I, Olson B, Backus R, Richter BD, Davis PA & Schneeman BO (2001) Beans, as a source of dietary fiber, increase cholecystokinin and apolipoprotein b48 response to test meals in men *J Nutr* **131**(5), 1485-90.

Braaten JT, Scott FW, Wood PJ, Riedel KD, Wolynetz MS, Brule D & Collins MW (1994) High beta-glucan oat bran and oat gum reduce postprandial blood glucose and insulin in subjects with and without type 2 diabetes *Diabet Med* **11**(3), 312-8.

Brighenti F, Casiraghi MC, Canzi E & Ferrari A (1999) Effect of consumption of a ready-to-eat breakfast cereal containing inulin on the intestinal milieu and blood lipids in healthy male volunteers *Eur J Clin Nutr* **53**, 726-33.

Brighenti F, Benini L, Del Rio D, Casiraghi C, Pellegrini N, Scazzina F, Jenkins DJ & Vantini I (2006) Colonic fermentation of indigestible carbohydrates contributes to the second-meal effect *Am J Clin Nutr* **83**(4), 817-22.

Brown L, Rosner B, Willett WW & Sacks FM (1999) Cholesterol-lowering effects of dietary fiber: a metaanalysis Am J Clin Nutr **69**, 30-42.

Buddington RK, Williams CH, Chen SC & Witherly SA (1996) Dietary supplement of neosugar alters the fecal flora and decreases activities of some reductive enzymes in human subjects *Am J Clin Nutr* **63**, 709-716.

Burley VJ, Leeds AR & Blundell JE (1987) The effect of high and low-fibre breakfasts on hunger, satiety and food intake in a subsequent meal *Int J Obes* **11** Suppl 1, 87-93.

Burley VJ, Paul AW & Blundell JE (1993) Influence of a high-fibre food (myco-protein) on appetite: effects on satiation (within meals) and satiety (following meals) *Eur J Clin Nutr* **47**(6), 409-18.

Burton-Freeman B, Davis PA & Schneeman BO (2002) Plasma cholecystokinin is associated with subjective measures of satiety in women *Am J Clin Nutr* **76**(3), 659-67.

Cairella G, Cairella M & Marchini G (1995) Effect of dietary fibre on weight correction after modified fasting *Eur J Clin Nutr* **49** Suppl 3, S325-7.

Campbell J, Fahey GC Jr & Wolf BW (1997) Selected indigestible oligosaccharides affect large bowel mass, cecal and fecal short-chain fatty acids, pH, and microflora in rats *J Nutr* **127**, 130-136.

Cani PD, Joly E, Horsmans Y & Delzenne NM (2006) Oligofructose promotes satiety in healthy human: a pilot study *Eur J Clin Nutr* **60**(5), 567-72.

Castiglia-Delavaud C, Verdier E, Besle JM, Vernet J, Boirie Y, Beaufrere B, De Baynast R & Vermorel M (1998) Net energy value of non-starch polysaccharide isolates (sugarbeet fibre and commercial inulin) and their impact on nutrient digestive utilization in healthy human subjects *Br J Nutr* **80**, 343-52.

Causey JL, Feirtag JM, Gallaher DD, Tungland BC & Slavin JL (2000) Effects of Dietary Inulin on Serum Lipids, Blood Glucose and the Gastrointestinal Environment in Hypercholesterolemic men *Nutr Res* **20**, 191-201.

Chaplin MF, Chaudhury S, Dettmar PW, Sykes J, Shaw AD & Davies GJ (2000) Effect of ispaghula husk on the faecal output of bile acids in healthy volunteers *J Steroid Biochem Mol Biol* **72**, 283-92.

Chearskul S, Supingklud N, Nitithamyong A & Sirichakwal P (2006) Assessment of hormonal and metabolic effects of dietary fiber in young Thai women *J Med Assoc Thai* **89**(7), 997-1003.

Chen HC, Chang CC, Mau WJ & Yen LS (2002) Evaluation of N-acetylchitooligosaccharides as the main carbon sources for the growth of intestinal bacteria *FEMS Microbiology Letters* **209**, 53-56.

Chen HL, Cheng HC, Liu YJ, Liu SY & Wu WT (2006) Konjac acts as a natural laxative by increasing stool bulk and improving colonic ecology in healthy adults *Nutrition* **22**, 1112-9.

Chen HL, Lu YH, Lin JJ & Ko LY (2001) Effects of isomalto-oligosaccharides on bowel functions and indicators of nutritional status in constipated elderly men *J Am Coll Nutr* **20**, 44-9.

Chen J, He J, Wildman RP, Reynolds K, Streiffer RH & Whelton PK (2006) A randomised controlled trial of dietary fiber intake on serum lipids *Eur J Clin Nutr* **60**, 62-8.

Cherbut C, Aube AC, Mekki N, Dubois C, Lairon D & Barry JL (1997) Digestive and metabolic effects of potato and maize fibres in human subjects *Br J Nutr* **77**, 33-46.

Cherbut C, Michel C & Lecannu G (2003) The prebiotic characteristics of fructooligosaccharides are necessary for reduction of TNBS-induced colitis in rats *J Nutr* **133**, 21-27.

Cicero AF, Derosa G, Manca M, Bove M, Borghi C & Gaddi AV (2007) Different effect of psyllium and guar dietary supplementation on blood pressure control in hypertensive overweight patients: a six-month, randomised clinical trial *Clin Exp Hypertens* **29**(6), 383-94.

Cobiac L, McArthur R & Nestel PJ (1990) Can eating baked beans lower plasma cholesterol? *Eur J Clin Nutr* 44, 819-22.

Colditz GA, Manson JE, Stampfer MJ, Rosner B, Willett WC & Speizer FE (1992) Diet and risk of clinical diabetes in women *Am J Clin Nutr* **55**(5), 1018-23.

Collins MD & Gibson GR (1999) Probiotics, prebiotics, and synbiotics: approaches for modulating the microbial ecology of the gut *Am J Clin Nutr* **69**, 1052S-1057S.

Conly JM, Stein K, Worobetz L & Rutledge-Harding S (1994) The contribution of vitamin K2 (menaquinones) produced by the intestinal microflora to human nutritional requirements for vitamin K. *Amer J Gastroenter* **89**, 915-923.

Costalos C, Kapiki A, Apostolou M & Papathoma E (2008) The effect of a prebiotic supplemented formula on growth and stool microbiology of term infants *Early Human Development* **84**, 45-49.

Coudray C & Fairweather-Tait SJ (1998) Do oligosaccharides affect the intestinal absorption of calcium in humans *Am J Clin Nutr* **68**, 921-923.

Crittenden RG & Playne MJ (1996) Production, properties and applications of food-grade oligosaccharides *Trends in Food Science and Technology* **7**, 353-361.

Crittenden RG (1999) Prebiotics. In *Probiotics: A critical review*, pp. 141-156 [G. Tannock, editor] Wymondham: Horizon Scientific Press.

Cummings JH (1995) Short chain fatty acids. In *Human colonic bacteria: Role in nutrition, physiology and pathology*, pp. 101-130 [Gibson G.R. and Macfarlane G.T. editors]. Boca Raton: CRC Press.

Cummings JH & Macfarlane GT (1991) The control and consequences of bacterial fermentation in the human colon *J Appl Bacteriol* **70**, 443-459.

Cummings JH, Bingham SA, Heaton KW & Eastwood MA (1992) Fecal weight, colon cancer risk, and dietary intake of nonstarch polysaccharides (dietary fiber) *Gastroenterol* **103**, 1783-9.

Cummings JH (1985) The effect of dietary fiber on fecal weight and composition. in: CRC Handbook of Dietary Fiber ain Human Nutrition. 1st edition. Edit GA Spiller. *CRC Press*, Boca Raton.

Cummings JH (2001) The effect of dietary fiber on fecal weight and composition. Chapter 4.4 in: CRC Handbook of Dietary Fiber ain Human Nutrition. 3rd edition. Edit GA Spiller. *CRC Press*, Boca Raton.

Cummings JH, Christie S & Cole TJ (2001) A study of fructo oligosaccharides in the prevention of travellers' diarrhoea *Alimentary Pharmacol Ther* **15**, 1139-1145.

D'Sousa A, Rajkumar C, Cooke J & Bulpitt CJ (2002) Probiotics in prevention of antibiotic associated diarrhoea: meta-analysis. *Brit Med J* 324, 1361-1364.

Dahl WJ, Lockert EA, Cammer AL & Whiting SJ (2005) Effects of flax fiber on laxation and glycemic response in healthy volunteers *J Med Food* **8**, 508-11.

Daly J, Tomlin J & Read NW (1993) The effect of feeding xanthan gum on colonic function in man: correlation with in vitro determinants of bacterial breakdown *Br J Nutr* **69**, 897-902.

Davidson MH, Dugan LD, Stocki J, Dicklin MR, Maki KC, Coletta F, Cotter R, McLeod M & Hoersten K (1998) A low-viscosity soluble-fiber fruit juice supplement fails to lower cholesterol in hypercholesterolemic men and women *J Nutr* **128**, 1927-32.

Davidson MH, Maki KC, Kong JC, Dugan LD, Torri SA, Hall HA, Drennan KB, Anderson SM, Fulgoni VL, Saldanha LG & Olson BH (1998) Long-term effects of consuming foods containing psyllium seed husk on serum lipids in subjects with hypercholesterolemia *Am J Clin Nutr* **67**, 367-76.

Davidson MH, Maki KC, Synecki C, Torri SA & Drennan KB (1998) Effects of dietary inulin on serum lipids in men and women with hypercholesterolemia *Nutrition Res* **18**, 503-517.

Davies GJ, Crowder M & Dickerson JW (1985) Dietary fibre intakes of individuals with different eating patterns *Hum Nutr Appl Nutr* **39**, 139-48.

Davies GJ, Crowder M, Reid B & Dickerson JW (1986) Bowel function measurements of individuals with different eating patterns *Gut* 27, 164-9.

Davy BM, Davy KP, Ho RC, Beske SD, Davrath LR & Melby CL (2002) High-fiber oat cereal compared with wheat cereal consumption favorably alters LDL-cholesterol subclass and particle numbers in middle-aged and older men *Am J Clin Nutr* **76**, 351-8.

de Preter V, Vanhoutte T, Huys G, Swings J, Rutgeerts P & Verbeke K (2008) Baseline microbiota activity and initial bifidobacteria counts influence responses to prebiotic dosing in healthy subjects *Alimentary Pharmacology Therapy* **15**, 504-513.

de Roos N, Heijnen ML, de Graaf C, Woestenenk G & Hobbel E (1995) Resistant starch has little effect on appetite, food intake and insulin secretion of healthy young men *Eur J Clin Nutr* **49**(7), 532-41.

de Vrese M, Stegelmann A, Richter B, Fenselau S, Laue C & Schrezenmeir J (2001) Probiotics - compensation for lactose insufficiency *Am J Clin Nutr* **73**, 421S-429S.

Delargy HJ, Burley VJ, O'Sullivan KR, Fletcher RJ & Blundell JE (1995) Effects of different soluble: insoluble fibre ratios at breakfast on 24-h pattern of dietary intake and satiety *Eur J Clin Nutr* **49**(10), 754-66.

Delzenne N, Aertssens J, Verplaetse H, Roccaro M & Roberfroid M (1995) Effect of fermentable fructooligosaccharides on mineral, nitrogen and energy digestive balance in the rat *Life Science* **57**, 1579-1587.

Den Hond E, Geypens B & Ghoos Y (2000) Effect of High Performance Chicory Inulin on Constipation Nutr Res 20, 731-36.

Department of Health (1991) Dietary Reference Values for Food Energy and Nutrients for the United Kingdom. Report of the panel of Dietary Reference Values of the Committee on Medical Aspects of Food Policy. Report on Health and Social Subjects **41**. HMSO: London.

Department of Health (2008) Health Survey for England, 2006, Volume 1: Cardiovascular diseases & risk factord in adults

Depeint F, Tzortzis G, Vulevic J, I'anson, K & Gibson GR (2008) Prebiotic evaluation of a novel galactooligosaccharide mixture produced by the enzymatic activity of *Bifidobacterium bifidum* NCIMB 41171, in healthy humans: a randomised, double-blind, crossover, placebo-controlled intervention study *Am J Clin Nutr* **87**, 785-791.

Daubioul CA, Horsmans Y, Lambert P, Danse E & Delzenne NM (2005) Effects of oligofructose on glucose and lipid metabolism in patients with nonalcoholic steatohepatitis: results of a pilot study *Eur J Clin Nutr* **59**(5), 723-6.

Di Lorenzo C, Williams CM, Hajnal F & Valenzuela JE (1988) Pectin delays gastric emptying and increases satiety in obese subjects *Gastroenterol* **95**(5), 1211-5.

Djouzi Z & Andrieux C (1997) Compared effects of three oligosaccharides on metabolism of intestinal microflora in rats inoculated with a human faecal flora. *Brit J Nutr* **78**, 313-324.

Djouzi Z, Andrieux C, Pelenc V, Somarriba S, Popot F, Paul F, Monsan P & Szylit O (1995) Degradation and fermentation of α -gluco-oligosaccharides by bacterial strains from human colon: in vitro and in vivo studies in gnotobiotic rats *J Appl Bacteriol* **79**, 117-127.

Drasar BS & Barrow PA (1985) Intestinal microbiology. Van Nostrand Reinhold Co Limited, UK.

Duncan LJ, Rose K & Meiklejohn AP (1960) Phenmetrazine hydrochloride and methylcellulose in the treatment of "refractory" obesity *Lancet* 1(7137), 1262-5.

Dunne C, O'Mahony L, Murphy L, Thornton G, Morrissey D, O'Halloran S, Feeney M, Flynn S, Fitzgerald G, Daly C, Kiley B, O'Sullivan GC, Shanahan F & Collins K (2001) In vitro selection criteria for probiotic bacteria of human origin: correlation with in vivo findings *Am J Clin Nutr* **73**, 386S-392S.

Durrant ML & Royston P (1978) The effect of preloads of varying energy density and methyl cellulose on hunger, appetite and salivation [proceedings] *Proc Nutr Soc* **37**(3), 87A.

Ebihara K, Masuhara R & Kiriyama S (1981) Effect of konjac mannan, a water soluble dietary fibre on plasma glucose and insulin responses in young men undergone glucose tolerance test *Nutr Rep Int* **23**(4), 577.

Eckburg PB, Bik EM, Bernstein, CN Purdom E, Dethlefsen L, Sargent M, Gill SR, Nelson KE, & Relman DA (2005) Diversity of the human intestinal microbial flora. *Science* **308**, 1635-1638.

Edwards CA, Blackburn NA, Craigen L, Davison P, Tomlin J, Sugden K, Johnson IT & Read NW (1987) Viscosity of food gums determined in vitro related to their hypoglycemic actions *Am J Clin Nutr* **46**(1), 72-7.

Effertz ME, Denman P & Slavin JL (1991) The effect of soy polysaccharide on body weight, serum lipids, blood glucose and fecal parameters in moderately obese adults *Nutrition Research* **11**(8), 849-859.

Elia M & Cummings JH (2007) Physiological aspects of energy metabolism and gastrointestinal effects of carbohydrates *Eur J Clin Nutr* **61**(S1), S40-S74.

Eliasson K, Ryttig KR, Hylander B & Rossner S (1992) A dietary fibre supplement in the treatment of mild hypertension. A randomised, double-blind placebo-controlled trial *J Hypertens* **10**(2), 195-9.

Ellegärd L, Andersson H & Bosaeus I (1997) Inulin and oligofructose do not influence the absorption of cholesterol, or the excretion of cholesterol, Ca, Mg, Zn, Fe or bile acids but increases energy excretion in ileostomy subjects. *Eur J Clin Nutr* **51**, 1-5.

Ellis PR, Apling EC, Leeds AR, Peterson DB & Jepson EM (1985) Guar bread and satiety: effects of an acceptable new product in overweight diabetic patients and normal subjects *J Plant Foods* **6**, 253-262.

Evans E & Miller DS (1975) Bulking agents in the treatment of obesity Nutr Metab 18(4), 199-203.

Fairchild RM, Ellis PR, Byrne AJ, Luzio SD & Mir MA (1996) A new breakfast cereal containing guar gum reduces postprandial plasma glucose and insulin concentrations in normal-weight human subjects Br J Nutr **76**(1), 63-73.

Feskens EJ, Loeber JG & Kromhout D (1994) Diet and physical activity as determinants of hyperinsulinemia: the Zutphen Elderly Study *Am J Epidemiol* **140**(4), 350-60.

Figdor SK & Rennhart HH (1981) Caloric utilisation and disposition of $[^{14}C]$ polydextrose in man *J Agricul Food Chem* **29**, 1181-1189.

Fontvieille AM, Bornet F, Rizkalla SW, Le Francois P, Pichard P, Desplanque N, Chevalier A, Letanoux M, Verel A, Tchobroutsky G et al (1988) In vitro and in vivo digestibility and metabolic effects of 3 wheat-flour products (white bread, french toast (rusk) and french toast bran-enriched) in normal subjects *Diabet Metab* **14**(2), 92-6.

Food and Agriculture Organisation (1998) Carbohydrates in human nutrition. Report of the Expert Panel meeting Rome April 1997 Rome. FAO, Rome.

Food and Agriculture Organisation (1980) Carbohydrates in Human Nutrition. Report of an expert meeting Geneva September 1979. FAO, Rome.

Fraser RB, Ford FA & Milner RD (1983) A controlled trial of a high dietary fibre intake in pregnancy--effects on plasma glucose and insulin levels *Diabetologia* **25**(3), 238-41.

French SJ & Read NW (1994) Effect of guar gum on hunger and satiety after meals of differing fat content: relationship with gastric emptying *Am J Clin Nutr* **59**(1), 87-91.

Frost GS, Brynes AE, Bovill-Taylor C & Dornhorst A (2004) A prospective randomised trial to determine the efficacy of a low glycaemic index diet given in addition to healthy eating and weight loss advice in patients with coronary heart disease *Eur J Clin Nutr* **58**(1), 121-7.

Frost GS, Brynes AE, Dhillo WS, Bloom SR & McBurney MI (2003) The effects of fiber enrichment of pasta and fat content on gastric emptying, GLP-1, glucose, and insulin responses to a meal *Eur J Clin Nutr* **57**(2), 293-8.

Fruhbeck G, Monreal I & Santidrian S (1997) Hormonal implications of the hypocholesterolemic effect of intake of field beans (Vicia faba L.) by young men with hypercholesterolemia *Am J Clin Nutr* **66**, 1452-60.

Fuchs CS, Giovannucci EL, Colditz GA, Hunter DJ, Stampfer MJ, Rosner B et al (1999) Dietary fiber and the risk of colorectal cancer and adenoma in women *N Engl J Med* **340** (3),169-176.

Fujiti K, Ogata Y & Hara K (1995) Effect of 4 G- -Galactosylsucrose (lactosucrose) on intestinal flora. *Seito Gigutsu Kenkyukai-shi* **43**, 83-91.

Fuller R & Gibson GR (1997) Modification of the intestinal microflora using probiotics and prebiotics *Scand J Gastroenterol* **32**, 28-31.

Fuller R (1989) Probiotics in man and animals J Appl Bacteriol 66, 365-378.

Fuller R (1992) The effect of probiotics on the gut micro-ecology of farm animals. In *The lactic acid bacteria in health and disease, vol. 1.*, pp. 171-192. [B.J.B. Wood, editor] Cambridge: Elsevier Science Publishers LTD.

Fuller R [editor] (1997) Probiotics 2: Applications and practical aspects. London: Chapman & Hall.

Furrie E, Macfarlane S, Kennedy A, Cummings JH, Walsh SV, O'Neil DA & Macfarlane GT (2005) Synbiotic therapy (*Bifidobacterium longum*/Synergy 1) initiates resolution of inflammation in patients with active ulcerative colitis: a randomised controlled pilot trial *Gut* **54**, 242-249.

Gaard M, Tretli S & Loken EB (1996) Dietary factors and risk of colon cancer: a prospective study of 50,535 young Norwegian men and women *Eur J Cancer Prev* **5** (6), 445-454.

Garcia AL, Otto B, Reich SC, Weickert MO, Steiniger J, Machowetz A, Rudovich NN, Mohlig M, Katz N, Speth M, Meuser F, Doerfer J, Zunft HJ, Pfeiffer AH & Koebnick C (2007) Arabinoxylan consumption decreases postprandial serum glucose, serum insulin and plasma total ghrelin response in subjects with impaired glucose tolerance *Eur J Clin Nutr* **61**(3), 334-41.

Geleva D, Thomas W, Gannon MC & Keenan JM (2003) A solubilized cellulose fiber decreases peak postprandial cholecystokinin concentrations after a liquid mixed meal in hypercholesterolemic men and women J *Nutr* **133**(7), 2194-203.

Gerhardt AL & Gallo NB (1998) Full-fat rice bran and oat bran similarly reduce hypercholesterolemia in humans J Nutr 128, 865-9.

Giacco R, Clemente G, Luongo D, Lasorella G, Fiume I, Brouns F, Bornet F, Patti L, Cipriano P, Rivellese AA & Riccardi G (2004) Effects of short-chain fructo-oligosaccharides on glucose and lipid metabolism in mild hypercholesterolaemic individuals *Clin Nutr* **23**(3), 331-40.

Gibson & Roberfroid MB [editors] (2008) A handbook of prebiotics. Taylor & Francis, Boca Raton.

Gibson GR & Collins MD (1999) Concept of balanced colonic microbiota, prebiotics, and synbiotics. In *Probiotics, other nutritional factors, and intestinal microflora, vol.* 42., pp. 139-152. [L.A. Hanson and R.H. Yolken, editors] Ltd, Philadelphia: Lippincott-Raven Publinshers.

Gibson GR & Macfarlane GT [editors] (1995) Human colonic bacteria: Role in nutrition, physiology and pathology. Boca Raton: CRC Press.

Gibson GR & McCartney AL (1998) Modification of the gut flora by dietary means *Biochem Soc Transactions* **26**, 222-228.

Gibson GR & R.A. Rastall [editors] (2006) *Prebiotics: Development and application*. Chichester: John Wiley & Sons Ltd.

Gibson GR & Roberfroid MB (1995) Dietary modulation of the human colonic microflora introducing the concept of probiotics *J Nutr* **125**, 1401-1412.

Gibson GR & Roberfroid MB [editors] (1999) Colonic microbiota, nutrition and health. Dodrecht: Kluwer Academic Publishers.

Gibson GR & Wang X (1994) Selective enrichment of bifidobacteria from human gut contents by oligofructose using continuous culture *FEMS Microbiol Letters* **118**, 121-128.

Gibson GR & Wang X (1994) Bifidogenic properties of different types of fructo-oligosaccharides *Food Microbiology* **11**, 491-498.

Gibson GR, Beatty ER, Wang X & Cummings JH (1995) Selective stimulation of bifidobacteria in the human colon by oligofructose and inulin *Gastroenterol* **108**, 975-982.

Gibson GR, Probert HM, van Loo JAE, Rastall RA & Roberfroid MB (2004) Dietary modulation of the human colonic microbiota: Updating the concept of prebiotics *Nutr Res Rev* **17**, 259-275.

Gibson GR, Willis CL & Van Loo J (1994) Non digestible oligosaccharides and bifidobacteria - implications for health *Int Sugar J* 96, 381-387.

Giovannucci E, Rimm EB, Stampfer MJ, Colditz GA, Ascherio A & Willett WC (1994) Intake of fat, meat, and fiber in relation to risk of colon cancer in men *Cancer Res* **54**(9), 2390-2397.

Giovannucci E, Stampfer MJ, Colditz G, Rimm EB & Willett WC (1992) Relationship of diet to risk of colorectal adenoma in men *J Natl Cancer Inst* **84**(2), 91-98.

Goldin BR (1998) Health benefits of probiotics Brit J Nutr 80, S203-S207.

Granfeldt Y, Drews A & Bjorck I (1995) Arepas made from high amylose corn flour produce favorably low glucose and insulin responses in healthy humans *J Nutr* **125**(3), 459-65.

Greundel S, Garcia AL, Otto B, Mueller C, Steiniger J, Weickert MO, Speth M, Katz N & Koebnick C (2006) Carob pulp preparation rich in insoluble dietary fibre and polyphenols enhances lipid oxidation and lowers postprandial acylated ghrelin in humans *J Nutr* **136**, 1533-1538.

Gråsten SM, Juntunen KS, Poutanen KS, Gylling HK, Miettinen TA, Mykkänen HM. (2000) Rye bread improves bowel function and decreases the concentrations of some compounds that are putative colon cancer risk markers in middle-aged women and men *J Nutr* **130**, 2215-21.

Grimes DS & Gordon C (1978) Satiety value of wholemeal and white bread Lancet 2(8080), 106.

Grubben MJ, van den Braak CC, Essenberg M, Olthof M, Tangerman A, Katan MB & Nagengast FM (2001) Effect of resistant starch on potential biomarkers for colonic cancer risk in patients with colonic adenomas: a controlled trial *Dig Dis Sci* **46**, 750-6.

Groop PH, Groop L, Totterman KJ & Fyhrquist F (1986) Relationship between changes in GIP concentrations and changes in insulin and C-peptide concentrations after guar gum therapy *Scand J Clin Lab Invest* **46**(6), 505-10.

Guarner F & Malagelada JR (2003) Gut flora in health and disease Lancet 361, 512-519.

Guarner F & Schaafsma GJ (1998) Probiotics. Int J Food Microbiol 39, 237-238.

Guarner F (2006) Enteric flora in health and disease Digestion 73, 5-12.

Guigoz Y, Rochat F, Perruisseau-Carrier G, Rochat I & Schriffin EJ (2002) Effects of oligosaccharide on the fecal flora and non-specific immune system in elderly people *Nutr Rev* 22, 13-25.

Haack VS, Chesters JG, Vollendorf NW, Story JA & Marlett JA (1998) Increasing amounts of dietary fiber provided by foods normalizes physiologic response of the large bowel without altering calcium balance or fecal steroid excretion *Am J Clin Nutr* **68**, 615-22.

Haarman M & Knol J (2005) Quantitative real-time PCR assays to identify and quantify fecal *Bifidobacterium* species in infants receiving a prebiotic infant formula. *Appl Env Microbiol* **71**, 2318-2324.

Haarman M & Knol J (2006) Quantitative real-time PCR analysis of fecal *Lactobacillus* species in infants receiving a prebiotic infant formula *Appl Env Microbiol* **72**, 2359-2365.

Haber GB, Heaton KW, Murphy D & Burroughs LF (1977) Depletion and disruption of dietary fibre. Effects on satiety, plasma-glucose, and serum-insulin *Lancet* **2**(8040), 679-82.

Hallfrisch J, Scholfield DJ & Behall KM (2002) Glucose and insulin responses to a new zero-energy fiber source *J Am Coll Nutr* **21**(5), 410-5.

Hamberg O, Rumessen JJ & Gudmand-Hoyer E (1989) Inhibition of starch absorption by dietary fibre. A comparative study of wheat bran, sugar-beet fibre, and pea fibre *Scand J Gastroenterol* **24**(1), 103-9.

Hanai H, Ikuma M, Sato Y, Iida T, Hosoda Y, Matsushita I, Nogaki A, Yamada M & Kaneko E (1997) Longterm effects of water-soluble corn bran hemicellulose on glucose tolerance in obese and non-obese patients: improved insulin sensitivity and glucose metabolism in obese subjects *Biosci Biotechnol Biochem* **61**(8), 1358-61.

Hara H, Li S, Sasaki M, Maruyama T, Terada A, Ogata Y, Fujita K, Ishigami H, Hara K, Fujimori I & Mitsuoka T (1994) Effective dose of lactosucrose on fecal flora and fecal metabolites of humans *Bifidobacteria Microflora* **13**, 51-63.

Harmsen HJ, Raangs GC, Franks A, Wildeboer-Veloo AC & Welling GW (2002) The effect of the prebiotic inulin and the probiotic *Bifidobacterium longum* on the fecal microflora of healthy volunteers measured by FISH and DGGE *Microbial Ecol Health Dis* **14**, 219-225.

Harmsen HJM, Wildeboer-Veloo ACM, Grijpstra J, Knol J, Degener JE & Welling GW (2000) Development of 16S rRNA-based probes for the *Coriobacterium* group and the *Atopobium* cluster and their application for enumeration of *Coriobacteriaceae* in human faeces from volunteers of different age groups *Appl Env Microbiol* **66**, 4523-4527.

Hayakawa K, Mizutani J, Wada K, Masai T, Yoshihara I & Mitsuoka T (1990) Effects of soybean oligosaccharides on human faecal flora *Microbial Ecol Health Dis* **3**, 293-303.

Heijnen ML, Deurenberg P, van Amelsvoort JM & Beynen AC (1995) Replacement of digestible by resistant starch lowers diet-induced thermogenesis in healthy men *Br J Nutr* **73**(3), 423-32.

Heijnen ML, van Amelsvoort JM, Deurenberg P & Beynen AC (1996) Neither raw nor retrograded resistant starch lowers fasting serum cholesterol concentrations in healthy normolipidemic subjects Am J Clin Nutr **64**, 312-8.

Heilbrun LK, Nomura A, Hankin JH & Stemmermann GN (1989) Diet and Colorectal-Cancer with Special Reference to Fiber Intake *Int J Cancer* 44(1), 1-6.

Heini AF, Lara-Castro C, Schneider H, Kirk KA, Considine RV & Weinsier RL (1998) Effect of hydrolyzed guar fiber on fasting and postprandial satiety and satiety hormones: a double-blind, placebo-controlled trial during controlled weight loss *Int J Obes* **22**(9), 906-9.

Hidaka H, Eida T, Takizawa T, Tokunaga T & Tashiro Y (1986) Effects of fructooligosaccharides on intestinal flora and human health *Bifidobacteria Microflora* **5**, 37-50.

Higginbotham S, Zhang ZF, Lee IM, Cook NR, Giovannucci E, Buring JE et al (2004) Dietary glycemic load and risk of colorectal cancer in the Women's Health Study *J Natl Cancer Inst* **96** (3), 229-233.

Hoad CL, Rayment P, Spiller RC, Marciani L, de Celis Alonso B, Traynor C, Mela DJ, Peters HPF & Gowland PA (2004) In vivo imaging of intragastric gelation and its effect on satiety in humans *J Nutr* **134**: 2293-2300

Hodge AM, English DR, O'Dea K & Giles GG (2004) Glycemic index and dietary fiber and the risk of type 2 diabetes *Diab Care* **27**(11), 2701-6.

Holzapfel WH, Haberer P, Snel J, Schillinger U & Huis isn't Veld JH (1998) Overview of gut flora and probiotics *Int J Food Microbiol* **41**, 85-101.

Hovey AL, Jones GP, Devereux HM & Walker KZ (2003) Whole cereal and legume seeds increase faecal short chain fatty acids compared to ground seeds *Asia Pac J Clin Nutr* **12**, 477-82.

Howard BV, Manson JE, Stefanick ML, Beresford SA, Frank G, Jones B, Rodabough RJ, Snetselaar L, Thomson C, Tinker L, Vitolins M & Prentice R (2006) Low-fat dietary pattern and weight change over 7 years: the Women's Health Initiative Dietary Modification Trial *JAMA* **295**(1), 39-49.

Howarth NC, Saltzman E, McCrory MA, Greenberg AS, Dwyer J, Ausman L, Kramer DG & Roberts SB (2003) Fermentable and nonfermentable fiber supplements did not alter hunger, satiety or body weight in a pilot study of men and women consuming self-selected diets *J Nutr* **133**(10), 3141-4.

Hu FB, Manson JE, Stampfer MJ, Colditz G, Liu S, Solomon CG & Willett WC (2001) Diet, lifestyle, and the risk of type 2 diabetes mellitus in women *N Engl J Med* **345**(11), 790-7.

Hylander B & Rossner S (1983) Effects of dietary fiber intake before meals on weight loss and hunger in a weight-reducing club *Acta Med Scand* **213**(3), 217-20.

Imamura L, Hisamitsu K & Kobashi K (1994) Purification and characterization of β-fructofuranosidase from *Bifidobacterium infantis Biol Pharmacol Bull* **17**, 596-602.

Innami S, Ishida H, Nakamura K, Kondo M, Tabata K, Koguchi T, Shimizu J & Furusho T (2005) Jew's mellow leaves (Corchorus olitorius) suppress elevation of postprandial blood glucose levels in rats and humans *Int J Vitam Nutr Res* 75(1), 39-46.

Institute of Medicine of the National Academies (2002) *Reference Intakes for Energy, Carbohydrate, Fiber, Fat, Fatty Acids, Cholesterol, Proteins and Amino Acids.* Washington, DC.

Iqbal SI, Helge JW & Heitmann BL (2006) Do energy density and dietary fiber influence subsequent 5-year weight changes in adult men and women? *Obesity* **14**(1), 106-14.

Ito M, Deguchi Y, Miyamori A, Matsumoto K, Kikuchi H, Matsumoto K, Kobayashi Y, Yajima T & Kan T (1990) Effects of administration of galactooligosaccharides on the human fecal microflora, stool weight and abdominal sensation *Microb Ecol Health Dis* **3**, 285-292.

Ito M, Kimura M, Deguchi Y, Miyamori-Watabe A, Yajima T & Kan T (1993) Effects of *trans*galactosylated disaccharides on the human intestinal microflora and their metabolism *J Nutr Sci Vitaminol* **39**, 279-288.

Jackson KG, Taylor GR, Clohessy AM & Williams CM (1999) The effect of the daily intake of inulin on fasting lipid, insulin and glucose concentrations in middle-aged men and women *Br J Nutr* **82**, 23-30.

Jacobs ET, Giuliano AR, Roe DJ, Guillen-Rodriguez JM, Alberts DS & Martinez ME (2002) Baseline dietary fiber intake and colorectal adenoma recurrence in the wheat bran fiber randomised trial *J Natl Cancer Inst* **94** (21), 1620-1625.

Jacobs ET, Giuliano AR, Roe DJ, Guillen-Rodriguez JM, Hess LM, Alberts DS et al (2002) Intake of supplemental and total fiber and risk of colorectal adenoma recurrence in the wheat bran fiber trial *Cancer Epidemiol Biomarkers Prev* **11** (9), 906-914.

Jarjis HA, Blackburn NA, Redfern JS & Read NW (1984) The effect of ispaghula (Fybogel and Metamucil) and guar gum on glucose tolerance in man *Br J Nutr* **51**(3), 371-8.

Jaskari J, Kontula P, Siitonen A, Jousimies-Somer H, Mattila-Sandholm T & Poutanen K (1998) Oat β -glucan and xylan hydrolysates as selective substrates for *Bifidobacterium* and *Lactobacillus* strains *Appl Microbiol Biotechnol* **49**, 175-181.

Jeffrys DB (1974) The efffect of dietary fibre on the response to orally administered glucose *Proc Nutr Soc* 33, 11A.

Jenkins DJ, Leeds AR, Gassull MA, Cochet B & Alberti GM (1977) Decrease in postprandial insulin and glucose concentrations by guar and pectin *Ann Intern Med* **86**(1), 20-3.

Jenkins DJ, Leeds AR, Houston H, Hinks L, Alberti KG & Cummings JH (1977) Carbohydrate tolerance in man after six weeks of pectin administration *Proc Nutr Soc* **36**(2), 60A.

Jenkins DJ, Wolever TM, Taylor RH, Barker HM, Fielden H & Gassull MA (1981) Lack of effect of refining on the glycemic response to cereals *Diabetes Care* **4**(5), 509-13.

Jenkins DJ, Wolever TM, Rao AV, Hegele RA, Mitchell SJ, Ransom TP, Boctor DL, Spadafora PJ, Jenkins AL, Mehling C et al (1993) Effect on blood lipids of very high intakes of fiber in diets low in saturated fat and cholesterol *N Engl J Med* **329**(1), 21-6.

Jenkins DJ, Vuksan V, Kendall CW, Würsch P, Jeffcoat R, Waring S, Mehling CC, Vidgen E, Augustin LS & Wong E (1998) Physiological effects of resistant starches on fecal bulk, short chain fatty acids, blood lipids and glycemic index *J Am Coll Nutr* **17**, 609-16.

Jenkins DJ, Kendall CW, Vuksan V, Augustin LS, Mehling C, Parker T, Vidgen E, Lee B, Faulkner D, Seyler H, Josse R, Leiter LA, Connelly PW & Fulgoni V 3rd (1999) Effect of wheat bran on serum lipids: influence of particle size and wheat protein *J Am Coll Nutr* **18**, 159-65.

Jenkins DJ, Kendall CW, Vuksan V, Vidgen E, Wong E, Augustin LS, Fulgoni V 3rd (2000) Effect of cocoa bran on low-density lipoprotein oxidation and fecal bulking *Arch Intern Med* **160**, 2374-9.

Jenkins DJ, Kendall CW, Popovich DG, Vidgen E, Mehling CC, Vuksan V, Ransom TP, Rao AV, Rosenberg-Zand R, Tariq N, Corey P, Jones PJ, Raeini M, Story JA, Furumoto EJ, Illingworth DR, Pappu AS & Connelly PW (2001) Effect of a very-high-fiber vegetable, fruit, and nut diet on serum lipids and colonic function *Metabolism* **50**, 494-503.

Jenkins DJ, Kendall CW, Augustin LS, Martini MC, Axelsen M, Faulkner D, Vidgen E, Parker T, Lau H, Connelly PW, Teitel J, Singer W, Vandenbroucke AC, Leiter LA & Josse RG (2002) Effect of wheat bran on glycemic control and risk factors for cardiovascular disease in type 2 diabetes *Diab Care* **25**, 1522-8.

Jensen MK, Koh-Banerjee P, Hu FB, Franz M, Sampson L, Gronbaek M & Rimm EB (2004) Intakes of whole grains, bran, and germ and the risk of coronary heart disease in men *Am J Clin Nutr* **80**, 1492-9.

Jie Z, Bang-Yao L, Ming-Jie X, Hai-Wei L, Zu-Kang Z, Ting-Song W & Craig, SA (2000) Studies on the effects of poydextrose intake on physiologic functions in Chinese people *Am J Clin Nutr* **72**, 1503-1509.

Johnson SK, Chua V, Hall RS & Baxter AL (2006) Lupin kernel fibre foods improve bowel function and beneficially modify some putative faecal risk factors for colon cancer in men *Br J Nutr* **95**, 372-8.

Kanyshkova TG, Buneva VN & Nevinskii GA (2002) Biological functions of human milk and its components *Upsekhi Sovremennoi Biologii* **122**, 259-271.

Karlstrom B, Vessby B, Asp NG & Ytterfors A (1988) Effects of four meals with different kinds of dietary fibre on glucose metabolism in healthy subjects and non-insulin-dependent diabetic patients *Eur J Clin Nutr* **42**(6), 519-26.

Karmally W, Montez MG, Palmas W, Martinez W, Branstetter A, Ramakrishnan R, Holleran SF, Haffner SM & Ginsberg HN (2005) Cholesterol-lowering benefits of oat-containing cereal in Hispanic americans *J Am Diet Assoc* **105**, 967-70.

Kay RM (1978) Food form, postprandial glycemia, and satiety Am J Clin Nutr 31, 738-741.

Keenan JM, Goulson M, Shamliyan T, Knutson N, Kolberg L & Curry L (2007) The effects of concentrated barley beta-glucan on blood lipids in a population of hypercholesterolaemic men and women *Br J Nutr* **97**, 1162-8.

Keenan JM, Pins JJ, Frazel C, Moran A & Turnquist L (2002) Oat ingestion reduces systolic and diastolic blood pressure in patients with mild or borderline hypertension: a pilot trial *J Fam Pract* **51**, 369.

Keogh GF, Cooper GJ, Mulvey TB, McArdle BH, Coles GD, Monro JA & Poppitt SD (2003) Randomised controlled crossover study of the effect of a highly beta-glucan-enriched barley on cardiovascular disease risk factors in mildly hypercholesterolemic men *Am J Clin Nutr* **78**(4), 711-8.

Kerckhoffs DA, Hornstra G & Mensink RP (2003) Cholesterol-lowering effect of beta-glucan from oat bran in mildly hypercholesterolemic subjects may decrease when beta-glucan is incorporated into bread and cookies *Am J Clin Nutr* **78**, 221-7.

Kestin M, Moss R, Clifton PM & Nestel PJ (1990) Comparative effects of three cereal brans on plasma lipids, blood pressure, and glucose metabolism in mildly hypercholesterolemic men *Am J Clin Nutr* **52**(4), 661-6.

Kim SH, Lee da H & Meyer D (2007) Supplementation of baby formula with native inulin has a prebiotic effect in formula-fed babies *Asia Pac J Clin Nutr* **16**, 172-177.

King NA, Craig SAS, Pepper T and Blundell JE (2005) Evaluation of the independent and combined effects of xyliton and polydextrose consumed as a snack on hunger and energy intake over 10d *Brit J Nutr* **93**: 911-915.

Kleessen B, Schwarz S, Boehm A, Fuhrmann H, Richter A, Henle T & Krueger M (2007) Jerusalem artichoke and chicory inulin in bakery products affects faecal microbiota of healthy volunteers *Br J Nutr* **98**, 540-549.

Kleessen B, Sykura B, Zunft HJ & Blaut M (1997) Effects of inulin and lactose on faecal microflora, microbial activity and bowel habit in elderly constipated persons. *Am J Clin Nutr* **65**, 1397-1402.

Knopp RH, Superko HR, Davidson M, Insull W, Dujovne CA, Kwiterovich PO, Zavoral JH, Graham K, O'Connor RR & Edelman DA (1999) Long-term blood cholesterol-lowering effects of a dietary fiber supplement *Am J Prev Med* **17**, 18-23.

Knotula P, Jaskari J, Nollet L, De Smet I, von Wright A, Poutanen K & Mattila-Sandholm T (1998) The colonization of a simulator of the human intestinal microbial ecosystem by a probiotic strain fed on a fermented oat bran product: effects on the gastrointestinal microbiota *Appl Microbiol Biotechnol* **50**, 246-252.

Koh-Banerjee P, Chu NF, Spiegelman D, Rosner B, Colditz G, Willett W & Rimm E (2003) Prospective study of the association of changes in dietary intake, physical activity, alcohol consumption, and smoking with 9-y gain in waist circumference among 16 587 US men *Am J Clin Nutr* **78**(4), 719-27.

Koh-Banerjee P, Franz M, Sampson L, Liu S, Jacobs DR, Jr., Spiegelman D, Willett W & Rimm E (2004) Changes in whole-grain, bran, and cereal fiber consumption in relation to 8-y weight gain among men *Am J Clin Nutr* **80**(5), 1237-45.

Kohmoto T, Fukui F, Takaku H & Mitsuoka T (1991) Dose-response test of isomaltooligosaccharides for increasing fecal bifidobacteria *Agric Biol Chem* **55**, 2157-2159.

Kohmoto T, Fukui F, Takaku H, Machida Y, Arai M & Mitsuoka T (1988) Effect of isomalto-oligosaccharides on human fecal flora *Bifidobacteria Microflora* **7**, 61-69.

Kolida S, Meyer D & Gibson GR (2007) A double-blind placebo-controlled study to establish the bifidogenic dose of inulin in healthy humans *Eur J Clin Nutr* **61**, 1189-1195.

Kovacs EM, Westerterp-Plantenga MS, Saris WH, Goossens I, Geurten P & Brouns F (2001) The effect of addition of modified guar gum to a low-energy semisolid meal on appetite and body weight loss *Int J Obes* **25**(3), 307-15.

Krishnan S, Rosenberg L, Singer M, Hu FB, Djousse L, Cupples LA & Palmer JR (2007) Glycemic index, glycemic load, and cereal fiber intake and risk of type 2 diabetes in US black women *Arch Intern Med* **167**(21), 2304-9.

Krotkiewski M (1984) Effect of guar gum on body-weight, hunger ratings and metabolism in obese subjects Br J *Nutr* **52**(1), 97-105.

Kruse HP, Kleessen B & Blaut M 1999) Effects of inulin on faecal bifidobacteria in human subjects. *Brit J Nutr* **82**, 375-382.

Kullen MJ & Klaenhammer T (1999) Genetic modification of lactobacilli and bifidobacteria. In: *Probiotics: A critical review*, pp. 65-83 [G Tannock, editor] Wymondham: Horizon Scientific Press.

Lairon D, Arnault N, Bertrais S, Planells R, Clero E, Hercberg S & Boutron-Ruault MC (2005) Dietary fiber intake and risk factors for cardiovascular disease in French adults *Am J Clin Nutr* **82**, 1185-94.

Lambert J & Hull R (1996) Upper gastrointestinal disease and probiotics Asia Pac J Clin Nutr 5, 31-35.

Lampe JW, Slavin JL, Baglien KS, Thompson WO, Duane WC & Zavoral JH (1991) Serum lipid and fecal bile acid changes with cereal, vegetable, and sugar-beet fiber feeding *Am J Clin Nutr* **53**, 1235-41.

Lampe JW, Slavin JL, Melcher EA & Potter JD (1992) Effects of cereal and vegetable fiber feeding on potential risk factors for colon cancer *Cancer Epidemiol Biomarkers Prev* **1**, 207-11.

Landin K, Holm G, Tengborn L & Smith U (1992) Guar gum improves insulin sensitivity, blood lipids, blood pressure, and fibrinolysis in healthy men *Am J Clin Nutr* **56**(6), 1061-5.

Langendijk P, Schut F, Jansen G, Raangs GC, Kamphuis GR, Wilkinson MH & Welling GW (1995) Quantitative fluorescent in situ hybridization of *Bifidobacterium* spp. with genus-specific 16S rRNA-targeted probes and its application in faecal samples *Appl Env Microbiol* **61**, 3069-3075.

Lau C, Faerch K, Glumer C, Tetens I, Pedersen O, Carstensen B, Jorgensen T & Borch-Johnsen K (2005) Dietary glycemic index, glycemic load, fiber, simple sugars, and insulin resistance: the Inter99 study *Diab Care* **28**(6), 1397-403.

Lavin JH & Read NW (1995) The effect on hunger and satiety of slowing the absorption of glucose: relationship with gastric emptying and postprandial blood glucose and insulin responses *Appetite* **25**(1), 89-96.

Leclere CJ, Champ M, Boillot J, Guille G, Lecannu G, Molis C, Bornet F, Krempf M, Delort-Laval J & Galmiche JP (1994) Role of viscous guar gums in lowering the glycemic response after a solid meal *Am J Clin Nutr* **59**(4), 914-21.

Leeds AR (1987) Treatment of obesity with dietary fibre: present position and potential developments *Scand J Gastroenterol Suppl* **129**, 156-8.

Leeman M, Ostman E & Bjorck I (2005) Vinegar dressing and cold storage of potatoes lowers postprandial glycaemic and insulinaemic responses in healthy subjects *Eur J Clin Nutr* **59**(11), 1266-71.

Letexier D, Diraison F & Beylot M (2003) Addition of inulin to a moderately high-carbohydrate diet reduces hepatic lipogenesis and plasma triacylglycerol concentrations in humans *Am J Clin Nutr* **77**, 559-64.

Levitt NS, Vinik AI, Sive AA, Child PT & Jackson WP (1980) The effect of dietary fiber on glucose and hormone responses to a mixed meal in normal subjects and in diabetic subjects with and without autonomic neuropathy *Diab Care* **3**(4), 515-9.

Lewis S, Burmeister S & Brazier J (2005) Effect of the prebiotic oligofructose on relapse of *Clostridium difficile*-associated diarrhea: a randomised, controlled study *Clin Gastroenterol Hepatol* **3**, 442-448.

Li J, Kaneko T, Qin LQ, Wang J & Wang Y (2003) Effects of barley intake on glucose tolerance, lipid metabolism, and bowel function in women *Nutrition* **19**(11-12), 926-9.

Liese AD, Schulz M, Fang F, Wolever TM, D'Agostino RB, Jr., Sparks KC & Mayer-Davis EJ (2005) Dietary glycemic index and glycemic load, carbohydrate and fiber intake, and measures of insulin sensitivity, secretion, and adiposity in the Insulin Resistance Atherosclerosis Study *Diab Care* **28**(12), 2832-8.

Liljeberg HG, Akerberg AK & Bjorck IM (1999) Effect of the glycemic index and content of indigestible carbohydrates of cereal-based breakfast meals on glucose tolerance at lunch in healthy subjects *Am J Clin Nutr* **69**(4), 647-55.

Lin J, Zhang SM, Cook NR, Rexrode KM, Liu S, Manson JE et al (2005) Dietary intakes of fruit, vegetables, and fiber, and risk of colorectal cancer in a prospective cohort of women (United States) *Cancer Causes Control* **16** (3), 225-233.

Lindstrom J, Peltonen M, Eriksson JG, Louheranta A, Fogelholm M, Uusitupa M & Tuomilehto J (2006a) Highfibre, low-fat diet predicts long-term weight loss and decreased type 2 diabetes risk: the Finnish Diabetes Prevention Study *Diabetologia* **49**(5), 912-20.

Lindstrom J, Ilanne-Parikka P, Peltonen M, Aunola S, Eriksson JG, Hemio K, Hamalainen H, Harkonen P, Keinanen-Kiukaanniemi S, Laakso M, Louheranta A, Mannelin M, Paturi M, Sundvall J, Valle TT, Uusitupa M & Tuomilehto J (2006b) Sustained reduction in the incidence of type 2 diabetes by lifestyle intervention: followup of the Finnish Diabetes Prevention Study *Lancet* **368**(9548), 1673-9.

Liu S, Manson JE, Stampfer MJ, Rexrode KM, Hu FB, Rimm EB & Willett WC (2000) Whole grain consumption and risk of ischemic stroke in women: A prospective study *JAMA* **284**, 1534-40.

Liu S, Buring JE, Sesso HD, Rimm EB, Willett WC & Manson JE (2002) A prospective study of dietary fiber intake and risk of cardiovascular disease among women *J Am Coll Cardiol* **39**, 49-56.

Liu S, Willett WC, Manson JE, Hu FB, Rosner B & Colditz G (2003) Relation between changes in intakes of dietary fiber and grain products and changes in weight and development of obesity among middle-aged women *Am J Clin Nutr* **78**(5), 920-7.

Lo GS & Cole TG (1990) Soy cotyledon fiber products reduce plasma lipids Atherosclerosis 82(1-2), 59-67.

Lovegrove JA, Clohessy A, Milon H & Williams CM (2000) Modest doses of beta-glucan do not reduce concentrations of potentially atherogenic lipoproteins *Am J Clin Nutr* **72**, 49-55.

Lovejoy J & DiGirolamo M (1992) Habitual dietary intake and insulin sensitivity in lean and obese adults Am J Clin Nutr 55(6), 1174-9.

Lu ZX, Walker KZ, Muir JG & O'Dea K (2004) Arabinoxylan fibre improves metabolic control in people with Type II diabetes *Eur J Clin Nutr* **58**, 621-8.

Ludwig DS, Pereira MA, Kroenke CH, Hilner JE, Van Horn L, Slattery ML & Jacobs DR Jr (1999) Dietary fiber, weight gain, and cardiovascular disease risk factors in young adults *JAMA* **282**, 1539-46.

Luo J, Rizkalla SW, Alamowitch C, Boussairi A, Blayo A, Barry JL, Laffitte A, Guyon F, Bornet FR & Slama G (1996) Chronic consumption of short-chain fructooligosaccharides by healthy subjects decreased basal hepatic glucose production but had no effect on insulin-stimulated glucose metabolism *Am J Clin Nutr* **63**, 939-45.

Luo J, Van Yperselle M, Rizkalla SW, Rossi F, Bornet FR & Slama G (2000) Chronic consumption of shortchain fructooligosaccharides does not affect basal hepatic glucose production or insulin resistance in type 2 diabetics *J Nutr* **130**, 1572-7.

Lupton JR, Morin JL, Robinson MC. (1993) Barley bran flour accelerates gastrointestinal transit time. J Am Diet Assoc. 93, 881-5.

Macfarlane GT & McBain AJ (1999) The human colonic microbiota. In *Colonic microbiota, nutrition and health,* pp. 1-25 [GR Gibson and MB Roberfroid editors]. Dordrecht: Kluwer Academic Publishers.

Macfarlane GT, Gibson GR & Cummings JH (1992) Comparison of fermentation reactions in different regions of the colon *J Appl Bacteriol* **72**, 57-64.

Macfarlane GT, Macfarlane S & Gibson GR (1998) Validation of a three-stage compound continuous culture system for investigating the effect of retention time on the ecology and metabolism of bacteria in the human colonic microbiota. *Microbial Ecol* **35**, 180-187.

Macfarlane S, Macfarlane GT & Cummings JH (2006) Review article: prebiotics in the gastrointestinal tract *Alimentary Pharmacol Ther* **24**, 701-714.

MacGillivray PC, Finlay HVL & Binns TB (1959) Use of lactulose to create a preponderance of lactobacilli in the intestine of bottle-fed infants *J Med* **4**, 182-189.

Mackay S & Ball MJ (1992) Do beans and oat bran add to the effectiveness of a low-fat diet? *Eur J Clin Nutr* **46**(9), 641-8.

Mai V, Flood A, Peters U, Lacey JV, Jr., Schairer C & Schatzkin A (2003) Dietary fibre and risk of colorectal cancer in the Breast Cancer Detection Demonstration Project (BCDDP) follow-up cohort *Int J Epidemiol* **32** (2), 234-239.

Maki KC, Davidson MH, Torri S, Ingram KA, O'Mullane J, Daggy BP & Albrecht HH (2000) High-molecularweight hydroxypropylmethylcellulose taken with or between meals is hypocholesterolemic in adult men *J Nutr* **130**, 1705-10.

Maki KC, Galant R, Samuel P, Tesser J, Witchger MS, Ribaya-Mercado JD, Blumberg JB & Geohas J (2007) Effects of consuming foods containing oat beta-glucan on blood pressure, carbohydrate metabolism and biomarkers of oxidative stress in men and women with elevated blood pressure *Eur J Clin Nutr* **61**(6), 786-95.

Maki KC, Carson ML, Miller MP, Turowski M, Bell M, Wilder DM & Reeves MS (2007) High-viscosity hydroxypropylmethylcellulose blunts postprandial glucose and insulin responses *Diab Care* **30**(5), 1039-43.

Maki KC, Carson ML, Miller MP, Turowski M, Bell M, Wilder DM, Rains TM & Reeves MS (2008) Hydroxypropylmethylcellulose and methylcellulose consumption reduce postprandial insulinemia in overweight and obese men and women *J Nutr* **138**(2), 292-6.

Marett R & Slavin JL (2004) No long-term benefits of supplementation with arabinogalactan on serum lipids and glucose *J Am Diet Assoc* **104**, 636-9.

Marteau P, Cuillerier E, Meance S, Gerhardt MF, Myara A, Bouvier M, Bouley C, Tondu F, Bommelaer G, Grimaud JC (2002) *Bifidobacterium animalis* strain DN-173 010 shortens the colonic transit time in healthy women: a double-blind, randomised, controlled study *Alimentary Pharmacol Ther* **16**, 587-593.

Marteau P, de Vrese M, Cellier CJ & Schrezenmeir J (2001) Protection from gastrointestinal diseases with the use of probiotics *Am J Clin Nutr* **73**, 430S-436S.

Mattes RD (2007) Effects of a combination fiber system on appetite and energy intake in overweight humans *Physiol Behav* **90**(5), 705-11.

McCartney AL (2002) Application of molecular biological methods for studying probiotics and the gut flora *Brit J Nutr* **88**, S29-S37.

McCullough ML, Robertson AS, Chao A, Jacobs EJ, Stampfer MJ, Jacobs DR et al (2003) A prospective study of whole grains, fruits, vegetables and colon cancer risk *Cancer Causes Control* **14** (10), 959-970.

McIntosh GH, Noakes M, Royle PJ & Foster PR (2003) Whole-grain rye and wheat foods and markers of bowel health in overweight middle-aged men *Am J Clin Nutr* **77**(4), 967-74.

McKeown NM, Meigs JB, Liu S, Wilson PW & Jacques PF (2002) Whole-grain intake is favorably associated with metabolic risk factors for type 2 diabetes and cardiovascular disease in the Framingham Offspring Study *Am J Clin Nutr* **76**, 390-8.

Meance S, Achour L & Briend A (1999) Comparison of starch digestibility of a blended food prepared with and without extrusion cooking *Eur J Clin Nutr* **53**(11), 844-8.

Menne E, Guggenbuhl N & Roberfroid M (2000) Fn-type inulin hydrolysate has a prebiotic effects in humans J *Nutr* **130**, 1197-1199.

Metchnikoff E (1907) The prolongation of life. London: William Heineman.

Michels KB, Fuchs CS, Giovannucci E, Colditz GA, Hunter DJ, Stampfer MJ et al. (2005) Fiber intake and incidence of colorectal cancer among 76,947 women and 47,279 men *Cancer Epidemiol Biomarkers Prev* **14**(4), 842-849.

Meyer KA, Kushi LH, Jacobs DR, Jr., Slavin J, Sellers TA & Folsom AR (2000) Carbohydrates, dietary fiber, and incident type 2 diabetes in older women *Am J Clin Nutr* **71**(4), 921-30.

Mihatsch WA, Hoegel J & Pohlandt F (2006) Prebiotic oligosaccharides reduce stool viscosity and accelerate gastrointestinal transport in preterm infants *Acta Paediatrics* **95**, 843-848.

Mitsuoka T, Hidaka H & Eida T (1987) Effect of fructooligosaccharides on intestinal microflora *Die Nahrung* **31**, 427-436.

Molis C, Flourié B, Ouarne F, Gailing MF, Lartigue S, Guibert A, Bornet F, Galmiche JP (1996) Digestion, excretion, and energy value of fructooligosaccharides in healthy humans *Am J Clin Nutr* **64**, 324-8.

Molly K, Vande Woestyne M & Verstraete W (1993) Development of a 5-step multi-chamber reactor as a simulation of the human intestinal microbial ecosystem *Appl Microbiol Biotechnol* **139**, 254-258.

Molnar D, Dober I & Soltesz G (1985) The effect of unprocessed wheat bran on blood glucose and plasma immunoreactive insulin levels during oral glucose tolerance test in obese children *Acta Paediatr Hung* 26(1), 75-7.

Montonen J, Knekt P, Jarvinen R, Aromaa A & Reunanen A (2003) Whole-grain and fiber intake and the incidence of type 2 diabetes *Am J Clin Nutr* **77**(3), 622-9.

Moore WEC & Holdeman LV (1974) Human fecal flora: the normal flora of 20 Japanese-Hawaiiens Appl Env Microbiol 27, 961-979.

Moorhead S, Welch RW, Barbara M, Livingstone E, McCourt M, Burns AA & Dunne A (2006) The effects of the fibre content and physical structure of carrots on satiety and subsequent intakes when eaten as part of a mixed meal *Br J Nutr* **96**(3), 587-95.

Mooy JM, Grootenhuis PA, de Vries H, Bouter LM, Kostense PJ & Heine RJ (1998) Determinants of specific serum insulin concentrations in a general Caucasian population aged 50 to 74 years (the Hoorn Study) *Diabet Med* **15**(1), 45-52.

Morgan LM, Tredger JA, Wright J & Marks V (1990) The effect of soluble- and insoluble-fibre supplementation on post-prandial glucose tolerance, insulin and gastric inhibitory polypeptide secretion in healthy subjects Br J Nutr **64**(1), 103-10.

MorishitaY & Konishi Y (1994) Effects of high dietary cellulose on the large intestinal microflora and shortchain fatty acids in rats *Letters Appl Microbiol* **19**, 433-435.

Moro, G.E., Stahl B, Fanaro S, Jelinek J, Boehm G & Coppa GV (2005) Dietary prebiotic oligosaccharides are detectable in the faeces of formula-fed infants *Acta Paediatrics* **94**, 27-30.

Mountzouris KC, McCartney AL & Gibson GR (2002) Intestinal microflora of human infants and current trends for its nutritional modulation *Br J Nutr* **87**, 405-420.

Muir JG, Yeow EG, Keogh J, Pizzey C, Bird AR, Sharpe K, O'Dea K & Macrae FA (2004) Combining wheat bran with resistant starch has more beneficial effects on fecal indexes than does wheat bran alone *Am J Clin Nutr* **79**, 1020-8.

Munoz JM, Sandstead HH, Jacob RA, Logan GM Jr, Reck SJ, Klevay LM, Dintzis FR, Inglett GE & Shuey WC (1979) Effects of some cereal brans and textured vegetable protein on plasma lipids *Am J Clin Nutr* **32**(3), 580-592.

Mykkanen H, Laiho K & Salminen S (1998) Variations in faecal bacterial enzyme activities and associations with bowel function and diet in elderly subjects *J Appl Microbiol* **85**, 37-41.

Nagengast FM, van den Ban G, Ploemen JP, Leenen R, Zock PL, Katan MB, Hectors MP, de Haan AF & van Tongeren JH (1993) The effect of a natural high-fibre diet on faecal and biliary bile acids, faecal pH and whole-gut transit time in man. A controlled study *Eur J Clin Nutr* **47**, 631-9.

Naidu AS, Bidlack WR & Clemens RA (1999) Probiotic spectra of lactic acid bacteria. Crit Rev Food Sci Nutr **38**, 13-126.

Nakajima Y & Nishio K (1993) Isomaltulose. Oligosaccharides. Production, properties and applications *Jap Tech Rev* **3**,107-117.

National Audit Office (2001) Tackling Obesity in England. London: The Stationery Office

Naumann E, van Rees AB, Onning G, Oste R, Wydra M & Mensink RP (2006) Beta-glucan incorporated into a fruit drink effectively lowers serum LDL-cholesterol concentrations *Am J Clin Nutr* **83**, 601-5.

Newby PK, Maras J, Bakun P, Muller D, Ferrucci L & Tucker KL (2007) Intake of whole grains, refined grains, and cereal fiber measured with 7-d diet records and associations with risk factors for chronic disease *Am J Clin Nutr* **86**, 1745-53.

Newby PK, Peterson KE, Berkey CS, Leppert J, Willett WC & Colditz GA (2003) Dietary composition and weight change among low-income preschool children *Arch Pediatr Adolesc Med* **157**(8), 759-64.

Nishida C, Nocito, FM & Mann J (2007) Joint FAO/WHO Scientific update on carbohydrates in Human Nutrition *Eur J Clin Nutr* **61**, Suppl 1 S1-S137.

Noakes M, Clifton PM, Nestel PJ, Le Leu R & McIntosh G (1996) Effect of high-amylose starch and oat bran on metabolic variables and bowel function in subjects with hypertriglyceridemia *Am J Clin Nutr* **64**, 944-51.

Nobaek S, Johansson, ML, Molin G, Ahrné S & Eppsson B (2000) Alteration of intestinal microflora is associated with reduction in abdominal bloating and pain in patients with irritable bowel syndrome Am J *Gastroenterol* **95**, 1231-1238.

Nomura AM, Hankin JH, Henderson BE, Wilkens LR, Murphy SP, Pike MC et al (2007) Dietary fiber and colorectal cancer risk: the multiethnic cohort study *Cancer Causes Control* **18**(7), 753-764.

O'Dea K, Nestel PJ & Antonoff L (1980) Physical factors influencing postprandial glucose and insulin responses to starch *Am J Clin Nutr* **33**(4), 760-5.

O'Sullivan DJ (2000) Methods for analysis of the intestinal microflora. Curr Iss Intestinal Microbiol 1, 39-50.

Okazaki M, Fujikawa S & Matsumoto N (1990) Effects of xylooligosaccharide on growth of bifidobacteria *J Jap Soc Nutr Food Sci* **43**, 395-401.

Oku T (1994) Special physiological functions of newly developed mono- and oligosaccharides. In: *Functional foods: Designer foods, pharma foods, nutraceuticals,* pp. 202-217 [I. Goldberg, I editor]. London: Chapman & Hall.

Olano-Martin E, Gibson GR & Rastall RA (2002) Comparison of the *in vitro* bifidogenic properties of pectins and pectic-oligosaccharides *J Appl Microbiol* **93**, 505-511.

Onning G, Wallmark A, Persson M, Akesson B, Elmstahl S & Oste R (1999) Consumption of oat milk for 5 weeks lowers serum cholesterol and LDL cholesterol in free-living men with moderate hypercholesterolemia *Ann Nutr Metab* **43**, 301-9.

Onyechi UA, Judd PA & Ellis PR (1998) African plant foods rich in non-starch polysaccharides reduce postprandial blood glucose and insulin concentrations in healthy human subjects *Br J Nutr* **80**(5), 419-28.

Otani T, Iwasaki M, Ishihara J, Sasazuki S, Inoue M, Tsugane S. (2006) Dietary fiber intake and subsequent risk of colorectal cancer: the Japan Public Health Center-based prospective study *Int J Cancer* **119**(6), 1475-1480.

Park OJ, Kang NE, Chang MJ & Kim WK (2004) Resistant starch supplementation influences blood lipid concentrations and glucose control in overweight subjects *J Nutr Sci Vitaminol (Tokyo)* **50**(2), 93-9.

Park Y, Hunter DJ, Spiegelman D, Bergkvist L, Berrino F, van den Brandt PA et al. (2005) Dietary fiber intake and risk of colorectal cancer: a pooled analysis of prospective cohort studies *JAMA* **294**(22), 2849-2857.

Parker RB (1974) Probiotics, the other half of the antibiotic story. Anals of Nutrition and Health 29, 4-8.

Pasman WJ, Saris WH, Wauters MA & Westerterp-Plantenga MS (1997) Effect of one week of fibre supplementation on hunger and satiety ratings and energy intake *Appetite* **29**(1), 77-87.

Pasman WJ, Westerterp-Plantenga MS, Muls E, Vansant G, van Ree J & Saris WH (1997) The effectiveness of long-term fibre supplementation on weight maintenance in weight-reduced women *Int J Obes* **21**(7), 548-55.

Pedersen A, Sandstrom B & Van Amelsvoort JM (1997) The effect of ingestion of inulin on blood lipids and gastrointestinal symptoms in healthy females *Br J Nutr* **78**, 215-22.

Pelkman CL, Navia JL, Miller AE & Pohle RJ (2007) Novel calcium-gelled, alginate-pectin beverage reduced energy intake in nondieting overweight and obese women: interactions with dietary restraint status *Am J Clin Nutr* **86**(6), 1595-602.

Pereira MA, Jacobs DR, Jr., Pins JJ, Raatz SK, Gross MD, Slavin JL & Seaquist ER (2002) Effect of whole grains on insulin sensitivity in overweight hyperinsulinemic adults *Am J Clin Nutr* **75**(5), 848-55.

Peters U, Sinha R, Chatterjee N, Subar AF, Ziegler RG, Kulldorff M et al. (2003); Dietary fibre and colorectal adenoma in a colorectal cancer early detection programme. *Lancet* **361** (9368), 1491-1495.

Petschow BW & Talbott RD (1991) Response of *Bifidobacterium* species to growth promoters in human and cow milk. *Pediatric Research* 29, 208-213.

Pick ME, Hawrysh ZJ, Gee MI, Toth E, Garg ML & Hardin RT (1996) Oat bran concentrate bread products improve long-term control of diabetes: a pilot study *J Am Diet Assoc* **96**, 1254-61.

Pietinen P, Malila N, Virtanen M, Hartman TJ, Tangrea JA, Albanes D et al. (1999) Diet and risk of colorectal cancer in a cohort of Finnish men *Cancer Causes Control* **10**(5), 387-396.

Pietinen P, Rimm EB, Korhonen P, Hartman AM, Willett WC, Albanes D & Virtamo J (1996) Intake of dietary fiber and risk of coronary heart disease in a cohort of Finnish men. The Alpha-Tocopherol, Beta-Carotene Cancer Prevention Study *Circulation* **94**, 2720-7.

Pins JJ, Geleva D, Keenan JM, Frazel C, O'Connor PJ & Cherney LM (2002) Do whole-grain oat cereals reduce the need for antihypertensive medications and improve blood pressure control? *J Fam Pract* **51**, 353-9.

Pittaway JK, Ahuja KD, Cehun M, Chronopoulos A, Robertson IK, Nestel PJ & Ball MJ (2006) Dietary supplementation with chickpeas for at least 5 weeks results in small but significant reductions in serum total and low-density lipoprotein cholesterols in adult women and men *Ann Nutr Metab* **50**, 512-8.

Platz EA, Giovannucci E, Rimm EB, Rockett HR, Stampfer MJ, Colditz GA et al. (1997) Dietary fiber and distal colorectal adenoma in men. *Cancer Epidemiol Biomarkers Prev* **6** (9), 661-670.

Playne MJ & Crittenden R (1996) Commercially available oligosaccharides. *Bull Int Dairy Foundation* **313**, 10-22.

Porikos K & Hagamen S (1986) Is fiber satiating? Effects of a high fiber preload on subsequent food intake of normal-weight and obese young men *Appetite* **7**(2), 153-62.

Probert HM & Gibson GR (2002) Investigating the prebiotic and gas-generating effects of selected carbohydrates on the human colonic microflora *Letters Appl Microbiol* **35**, 473-480.

Probert HM, Apajalahti JHA, Routenden N, Stowell J & Gibson GR (2004) Polydextrose and lactitol fermentation by colonic bacteria in a three-stage continuous culture system. *Appl Env Microbiol* **70**, 4505-4511.

Prosky L, Asp NG, Schweizer TF, DeVries JW & Furda I (1988) Determination of Insoluble, Soluble, and Total Dietary Fiber in Foods and Food-Products - Interlaboratory Study *J AOAC* **71**(5), 1017-1023.

Queenan KM, Stewart ML, Smith KN, Thomas W, Fulcher RG & Slavin JL (2007) Concentrated oat betaglucan, a fermentable fiber, lowers serum cholesterol in hypercholesterolemic adults in a randomised controlled trial *Nutr J* **6**, 6. Raben A, Christensen NJ, Madsen J, Holst JJ & Astrup A (1994) Decreased postprandial thermogenesis and fat oxidation but increased fullness after a high-fiber meal compared with a low-fiber meal *Am J Clin Nutr* **59**(6), 1386-94.

Raben A, Tagliabue A, Christensen NJ, Madsen J, Holst JJ & Astrup A (1994) Resistant starch: the effect on postprandial glycemia, hormonal response, and satiety *Am J Clin Nutr* **60**(4), 544-51.

Ranganathan S, Champ M, Pechard C, Blanchard P, Nguyen M, Colonna P & Krempf M (1994) Comparative study of the acute effects of resistant starch and dietary fibers on metabolic indexes in men *Am J Clin Nutr* **59**(4), 879-83.

Rao VA (2001) The prebiotic properties of oligofructose at low intake levels. Nutrition Research 6, 843-848.

Reyna-Villasmil N, Bermudez-Pirela V, Mengual-Moreno E, Arias N, Cano-Ponce C, Leal-Gonzalez E, Souki A, Inglett GE, Israili ZH, Hernandez-Hernandez R, Valasco M & Arraiz N (2007) Oat-derived beta-glucan significantly improves HDLC and diminishes LDLC and non-HDL cholesterol in overweight individuals with mild hypercholesterolemia *Am J Ther* **14**, 203-12.

Rigaud D, Ryttig KR, Leeds AR, Bard D & Apfelbaum M (1987) Effects of a moderate dietary fibre supplement on hunger rating, energy input and faecal energy output in young, healthy volunteers. A randomised, doubleblind, cross-over trial *Int J Obes* **11** Suppl 1, 73-8.

Rigaud D, Ryttig KR, Angel LA & Apfelbaum M (1990) Overweight treated with energy restriction and a dietary fibre supplement: a 6-month randomised, double-blind, placebo-controlled trial *Int J Obes* **14**(9), 763-9.

Rigaud D, Paycha F, Meulemans A, Merrouche M & Mignon M (1998) Effect of psyllium on gastric emptying, hunger feeling and food intake in normal volunteers: a double blind study *Eur J Clin Nutr* **52**(4), 239-45.

Roberfroid MB, Cumps J & Devogelaer JP (2002) Dietary chicory inulin increases whole-body bone mineral density in growing male rats *J Nutr* **132**, 3599-3602.

Robertson DJ, Sandler RS, Haile R, Tosteson TD, Greenberg ER, Grau M et al (2005) Fat, fiber, meat and the risk of colorectal adenomas *Am J Gastroenterol* **100** (12), 2789-2795.

Robertson MD, Currie JM, Morgan LM, Jewell DP & Frayn KN (2003) Prior short-term consumption of resistant starch enhances postprandial insulin sensitivity in healthy subjects *Diabetologia* **46**(5), 659-65.

Robertson MD, Bickerton AS, Dennis AL, Vidal H & Frayn KN (2005) Insulin-sensitizing effects of dietary resistant starch and effects on skeletal muscle and adipose tissue metabolism *Am J Clin Nutr* **82**(3), 559-67.

Robinson RR, Feirtag J & Slavin JL (2001) Effects of dietary arabinogalactan on gastrointestinal and blood parameters in healthy human subjects *J Am Coll Nutr* **20**, 279-85.

Rolls BJ, Bell EA, Castellanos VH, Chow M, Pelkman CL & Thorwart ML (1999) Energy density but not fat content of foods affected energy intake in lean and obese women *Am J Clin Nutr* **69**(5), 863-71.

Ross AH, Eastwood MA, Brydon WG, Anderson JR & Anderson DM (1983) A study of the effects of dietary gum arabic in humans *Am J Clin Nutr* **37**(3), 368-75.

Rossner S, Andersson IL & Ryttig K (1988) Effects of a dietary fibre supplement to a weight reduction programme on blood pressure. A randomised, double-blind, placebo-controlled study *Acta Med Scand* **223**(4), 353-7.

Rossner S, von Zweigbergk D, Ohlin A & Ryttig K (1987) Weight reduction with dietary fibre supplements. Results of two double-blind randomised studies *Acta Med Scand* **222**(1), 83-8.

Rowland IR & Tanaka R (1993) The effects of *trans*galactosylated oligosaccharides on gut flora metabolism in rats associated with a human faecal microflora *J Appl Bacteriol* **74**, 667-674.

Rycroft CE, Jones MR, Gibson GR & Rastall RA (2001) Fermentation properties of gentio-oligosaccharides *Letters Appl Microbiol* **32**, 156-161.

Ryttig KR, Tellnes G, Haegh L, Boe E & Fagerthun H (1989) A dietary fibre supplement and weight maintenance after weight reduction: a randomised, double-blind, placebo-controlled long-term trial *Int J Obes* **13**(2), 165-71.

Ryttig KR, Lammert O, Nielsen E, Garby L & Poulsen K (1990) The effect of a soluble dietary fibre supplement on 24-hour energy expenditure during a standardized physical activity programme *Int J Obes* **14**(5), 451-5.

Sabovic M, Lavre S & Keber I (2004) Supplementation of wheat fibre can improve risk profile in patients with dysmetabolic cardiovascular syndrome *Eur J Cardiovasc Prev Rehabil* **11**(2), 144-8.

Sairanen U, Piirainen L, Gråsten S, Tompuri T, Mättö J, Saarela M & Korpela R (2007) The effect of probiotic fermented milk and inulin on the functions and microecology of the intestine *J Dairy Res* **74**, 367-73.

Saito Y, Takano T & Rowland I (1992) Effects of soybean oligosaccharides on the human gut microflora in in vitro culture *Microb Ecol Health Dis* **5**, 105-110.

Salmeron J, Manson JE, Stampfer MJ, Colditz GA, Wing AL & Willett WC (1997a) Dietary fiber, glycemic load, and risk of non-insulin-dependent diabetes mellitus in women *JAMA* **277**(6), 472-7.

Salmeron J, Ascherio A, Rimm EB, Colditz GA, Spiegelman D, Jenkins DJ, Stampfer MJ, Wing AL & Willett WC (1997b) Dietary fiber, glycemic load, and risk of NIDDM in men *Diab Care* **20**(4), 545-50.

Salminen S, Bouley C, Boutron-Ruault MC, Cummings JH, Franck A, Gibson GR, Isolauri I, Moreau MC, Roberfroid M & Rowland IR (1998) Functional food science and gastrointestinal physiology and function *Brit J Nutr* **80**, S147-S171.

Saltzman E, Das SK, Lichtenstein AH, Dallal GE, Corrales A, Schaefer EJ, Greenberg AS & Roberts SB (2001) An oat-containing hypocaloric diet reduces systolic blood pressure and improves lipid profile beyond effects of weight loss in men and women *J Nutr* **131**(5), 1465-70.

Saltzman E, Moriguti JC, Das SK, Corrales A, Fuss P, Greenberg AS & Roberts SB (2001) Effects of a cereal rich in soluble fiber on body composition and dietary compliance during consumption of a hypocaloric diet JAm *Coll Nutr* **20**(1), 50-7.

Samra RA & Anderson GH (2007) Insoluble cereal fiber reduces appetite and short-term food intake and glycemic response to food consumed 75 min later by healthy men *Am J Clin Nutr* **86**(4), 972-9.

Sanaka M, Yamamoto T, Anjiki H, Nagasawa K & Kuyama Y (2007) Effects of agar and pectin on gastric emptying and post-prandial glycaemic profiles in healthy human volunteers *Clin Exp Pharmacol Physiol* **34**(11), 1151-5.

Sanders ME, Gibson GR, Gill HS & Guarner F (2007) Probiotics: Their potential to impact human health *CAST Issue Paper* **36**, 1-20.

Savino F, Palumeri E, Castagno E, Cresi F, Dalmasso P, Cavallo F & Oggero R (2006) Reduction of crying episodes owing to infantile colic: A randomised controlled study on the efficacy of a new infant formula *Eur J Clin Nutr* **60**, 1304-1310.

Scalfi L, Coltorti A, D'Arrigo E, Carandente V, Mazzacano C, Di Palo M & Contaldo F (1987) Effect of dietary fibre on postprandial thermogenesis *Int J Obes* **11**(S1), 95-9.
Schaafsma G, Meuling WJ, van Dokkum W & Bouley C (1998) Effects of a milk product, fermented by Lactobacillus acidophilus and with fructo-oligosaccharides added, on blood lipids in male volunteers *Eur J Clin Nutr* **52**, 436-40.

Schatzkin A, Mouw T, Park Y, Subar AF, Kipnis V, Hollenbeck A et al (2007) Dietary fiber and whole-grain consumption in relation to colorectal cancer in the NIH-AARP Diet and Health Study *Am J Clin Nutr* **85** (5), 1353-1360.

Scholtens PA, Alles MS, Willemsen LE, van den Braak C, Bindels JG, Boehm G & Govers MJ (2006) Dietary fructo-oligosaccharides in healthy adults do not negatively affect faecal cytotoxicity: a randomised, doubleblind, placebo-controlled crossover trial *Br J Nutr* **95**, 1143-9.

Scholtens PA, Alles MS, Bindels JG, van der Linde EG, Tolboom JJ & Knol J (2006) Bifidogenic effects of solid weaning foods with added prebiotic oligosaccharides: A randomised controlled clinical trial *J Ped Gastroenterol Nutr* **42**, 553-559, 2006.

Schulze MB, Liu S, Rimm EB, Manson JE, Willett WC & Hu FB (2004) Glycemic index, glycemic load, and dietary fiber intake and incidence of type 2 diabetes in younger and middle-aged women *Am J Clin Nutr* **80**(2), 348-56.

Schulze MB, Schulz M, Heidemann C, Schienkiewitz A, Hoffmann K & Boeing H (2007) Fiber and magnesium intake and incidence of type 2 diabetes: a prospective study and meta-analysis *Arch Intern Med* **167**(9), 956-65.

Schwab U, Louheranta A, Torronen A & Uusitupa M (2006) Impact of sugar beet pectin and polydextrose on fasting and postprandial glycemia and fasting concentrations of serum total and lipoprotein lipids in middle-aged subjects with abnormal glucose metabolism *Eur J Clin Nutr* **60**, 1073-80.

Sellers TA, Bazyk AE, Bostick RM, Kushi LH, Olson JE, Anderson KE et al (1998) Diet and risk of colon cancer in a large prospective study of older women: an analysis stratified on family history (Iowa, United States) *Cancer Causes Control* **9** (4), 357-367.

Sierra M, Garcia JJ, Fernandez N, Diez MJ, Calle AP & Sahagun AM (2001) Effects of ispaghula husk and guar gum on postprandial glucose and insulin concentrations in healthy subjects *Eur J Clin Nutr* **55**(4), 235-43.

Smith EA & Macfarlane GT (1996) Enumeration of human colonic bacteria producing phenolic and indolic compounds: effects of pH, carbohydrate availability and retention on dissimilatory aromatic amino acid metabolism *J Appl Bacteriol* **81**, 288-302.

Sola R, Godas G, Ribalta J, Vallve JC, Girona J, Anguera A, Ostos M, Recalde D, Salazar J, Caslake M, Martin-Lujan F, Salas-Salvado J & Masana L (2007) Effects of soluble fiber (Plantago ovata husk) on plasma lipids, lipoproteins, and apolipoproteins in men with ischemic heart disease *Am J Clin Nutr* **85**, 1157-63.

Solum TT, Ryttig KR, Solum E & Larsen S (1987) The influence of a high-fibre diet on body weight, serum lipids and blood pressure in slightly overweight persons. A randomised, double-blind, placebo-controlled investigation with diet and fibre tablets (DumoVital) *Int J Obes* **11**(S1), 67-71.

Southgate DA, Branch WJ, Hill MJ, Drasar BS, Walters RL, Davies PS & Baird IM (1976) Metabolic responses to dietary supplements of bran *Metabolism* **25**(10), 1129-35.

Sparti A, Milon H, Di Vetta V, Schneiter P, Tappy L, Jequier E & Schutz Y (2000) Effect of diets high or low in unavailable and slowly digestible carbohydrates on the pattern of 24-h substrate oxidation and feelings of hunger in humans *Am J Clin Nutr* **72**(6), 1461-8.

Spiller GA, Story JA, Furumoto EJ, Chezem JC & Spiller M (2003) Effect of tartaric acid and dietary fibre from sun-dried raisins on colonic function and on bile acid and volatile fatty acid excretion in healthy adults *Br J Nutr* **90**, 803-7.

Stark PL & Lee A (1982) The microbial ecology of the large bowel of breast-fed and formula-fed infants during the first year of life *J Med Microbiol* **15**, 189-203.

Stasse-Wolthuis M, Albers HF, van Jeveren JG, Wil de Jong J, Hautvast JG, Hermus RJ, Katan MB, Brydon WG & Eastwood MA (1980) Influence of dietary fiber from vegetables and fruits, bran or citrus pectin on serum lipids, fecal lipids, and colonic function *Am J Clin Nutr* **33**, 1745-56.

Steer T, Carpenter H, Tuohy KM & Gibson GR (2001) Perspectives on the role of the human gut microflora in health and disease and its modulation by pro- and prebiotics *Nutr Res Rev* **13**, 229-254.

Steinmetz KA, Kushi LH, Bostick RM, Folsom AR, Potter JD. (1994) Vegetables, Fruit, and Colon-Cancer in the Iowa Womens Health Study *Am J Epidemiol* **139**(1), 1-15.

Stephen AM, Dahl WJ, Sieber GM, van Blaricom JA & Morgan DR (1995) Effect of green lentils on colonic function, nitrogen balance, and serum lipids in healthy human subjects *Am J Clin Nutr* **62**, 1261-7.

Stevens J, Ahn K, Juhaeri, Houston D, Steffan L & Couper D (2002) Dietary fiber intake and glycemic index and incidence of diabetes in African-American and white adults: the ARIC study *Diabet Care* **25**(10), 1715-21.

Stevens J, Levitsky DA, VanSoest PJ, Robertson JB, Kalkwarf HJ & Roe DA (1987) Effect of psyllium gum and wheat bran on spontaneous energy intake *Am J Clin Nutr* **46**(5), 812-7.

Streppel MT, Arends LR, van 't Veer P, Grobbee DE & Geleijnse JM (2005) Dietary fiber and blood pressure: a meta-analysis of randomised placebo-controlled trials *Arch Intern Med* **165**, 150-6.

Sunvold GD, Titgemeyer EC, Bourquin LD, Fahey GC Jr & Garleb KA (1995) Alteration of the fiber and lipid components of a defined-formula diet: effects on stool characteristics, nutrient digestibility, mineral balance, and energy metabolism in humans *Am J Clin Nutr* **62**, 1252-60.

Suzuki K, Endo Y, Uehara M, Yamada H, Goto S, Imamura M & Shioza S (1985) Effect of lactose, lactulose and sorbital on mineral utilisation and intestinal flora *J Jap Soc Nutr Food Sci* **38**, 39-42.

Szilagyi A (2002) Review article: lactose a potential prebiotic Alimentary Pharmacol Ther 16, 1591-1602.

Tagliabue A, Raben A, Heijnen ML, Deurenberg P, Pasquali E & Astrup A (1995) The effect of raw potato starch on energy expenditure and substrate oxidation *Am J Clin Nutr* **61**(5), 1070-5.

Tahiri M, Tressol JC, Arnaud Y, Bornet FRJ, Bouteloup-Demange C, Feillet-Coudray C, Brandolini M, Ducros, V, Pepin D, Brouns F, Roussel AM, Rayssiguier Y & Coudray C (2003) Effect of short-chain fructooligosaccharides on intestinal calcium absorption and calcium status in postmenopausal women: a stable-isotope study *Am J Clin Nutr* **77**, 449-457.

Tamura Z (1983) Nutriology of bifidobacteria Bifidobacteria Microflora 2, 3-16.

Tanaka R, Takayama H, Morotomi M, Kuroshima T, Ueyama S, Matsumoto K, Kuroda A & Mutai M (1983) Effects of administration of TOS and *Bifidobacterium breve* 4006 on the human fecal flora *Bifidobacteria Microflora* **2**, 17-24.

Tannock GW (1999) A fresh look at the intestinal microflora. In *Probiotics: A critical review*, pp. 5-12 [GW Tannock editor]. Wymondham: Horizon Scientific Press.

Ten Bruggencate SJ, Bovee-Oudenhoven IM, Lettink-Wissink ML, Katan MB & van der Meer R (2006) Dietary fructooligosaccharides affect intestinal barrier function in healthy men *J Nutr* **136**, 70-4.

Terada A, Hara H, Kataoka M & Mitsuoka T (1992) Effect of lactulose on the composition and metabolic activity of the human faecal flora. *Microbial Ecol Health Dis* **5**, 43-50.

Terry P, Hu FB, Hansen H & Wolk A (2001) Prospective study of major dietary patterns and colorectal cancer risk in women *Am J Epidemiol* **154** (12), 1143-1149.

Teuri U, Korpela R, Saxelin M, Montonen L & Salminen S (1998) Increased fecal frequency and gastrointestinal symptoms following ingestion of galacto-oligosaccharide-containing yogurt *J Nutr Sci Vitaminol* **44**, 465-471.

Theuwissen E & Mensink RP (2007) Simultaneous intake of beta-glucan and plant stanol esters affects lipid metabolism in slightly hypercholesterolemic subjects *J Nutr* **137**, 583-8.

Tiwary CM, Ward JA & Jackson BA (1997) Effect of pectin on satiety in healthy US Army adults *J Am Coll Nutr* **16**(5), 423-8.

Tomlin J (1995) The effect of liquid fibre on feeding behaviour Eur J Clin Nutr 49 (S3), S246-9.

Tomlin J (1995) The effect of the gel-forming liquid fibre on feeding behaviour in man Br J Nutr 74(3), 427-36.

Tredger J, Sheard C & Marks V (1981) Blood glucose and insulin levels in normal subjects following a meal with and without added sugar beet pulp *Diabet Metab* **7**(3), 169-72.

Tsukahara T, Koyama H, Okada M & Ushida K (2002) Stimulation of butyrate production by gluconic acid in batch culture of pig cecal digesta and identification of butyrate-producing bacteria. *J Nutr* **132**, 2229-2234.

Tuohy KM, Finlay RK, Wynne AG & Gibson GR (2001a) A human volunteer study on the prebiotic effects of HP-Inulin – gut bacteria enumerated using fluorescent *in situ* hybridisation (FISH) *Anaerobe* **7**, 113-118.

Tuohy KM, Kolida S, Lustenberger A & Gibson GR (2001b) The prebiotic effects of biscuits containing partially hydrolyzed guar gum and fructooligosaccharides – A human volunteer study *Br J Nutr* **86**, 341-348.

Tuohy KM, Ziemer CJ, Klinder A, Knobel Y, Pool-Zobel BL & Gibson GR (2002) A human volunteer study to determine the prebiotic effects of lactulose powder on human colonic bacteria *Microb Ecol Health Dis* **14**, 165-173.

Tuomilehto J, Voutilainen E, Huttunen J, Vinni S & Homan K (1980) Effect of guar gum on body weight and serum lipids in hypercholesterolemic females *Acta Med Scand* **208**(1-2), 45-8.

Turnbull WH & Thomas HG (1995) The effect of a Plantago ovata seed containing preparation on appetite variables, nutrient and energy intake *Int J Obes Relat Metab Disord* **19**(5), 338-42.

Tzortzis G, Goulas AK & Gibson GR (2005a) Synthesis of prebiotic galactooligosaccharides using whole cells of a novel strain, *Bifidobacterium bifidum* NCIMB 41171 *Appl Microbiol Biotechnol* **68**, 412-416.

Tzortzis G, Goulas AK, Gee JM & Gibson GR (2005b) A novel galactooligosaccharide mixture increases the bifidobacterial population numbers in a continuous in vitro fermentation system and in the proximal colon of pigs in vivo. *Journal of Nutrition* **135**, 1726-1731.

United Nations Food and Agriculture Organization (UNFAO) (2001) Health and nutritional properties of probiotics in food including powder milk with live lactic acid bacteria, http://ftp.fao.org/es/esn/food/probio_report_en.pdf (4 October 2006)

United Nations Food and Agriculture Organization (UNFAO) (2002) Guidelines for the evaluation of probiotics in Food. http://ftp.fao.org/es/esn/food/wgreport2.pdf (4 October 2006)

USDA Agricultural Research Service (2005) What we eat in America *NHANES* 2001/02: Usual nutrient intakes from food compared to Dietary Reference Intakes.

Vajifdar BU, Goyal VS, Lokhandwala YY, Mhamunkar SR, Mahadik SP, Gawad AK, Halankar SA & Kulkarni HL (2000) Is dietary fiber beneficial in chronic ischemic heart disease? *J Assoc Physicians India* **48**(9), 871-6.

Valette P, Pelenc V, Djouzi Z, Andrieux C, Paul F, Monsan P & Szylit O (1993) Bioavailability of new synthesised glucooligosaccharides in the intestinal tract of gnotobiotic rats *J Sci Food Agric* **62**, 121-127.

Valle-Jones JC (1980) The evaluation of a new appetite-reducing agent (Prefil) in the management of obesity *Br J Clin Pract* **34**(3), 72-4.

van de Ven ML, Westerterp-Plantenga MS, Wouters L & Saris WH (1994) Effects of liquid preloads with different fructose/fibre concentrations on subsequent food intake and ratings of hunger in women *Appetite* 23(2), 139-46.

van den Heuvel EG, Schoterman MH & Muijs T (2000) *Trans*galactooligosaccharides stimulate calcium absorption in postmenopausal women *J Nutr* **130**, 2938-2942.

van der Waaij D (1999) Microbial ecology of the intestinal microflora: influence of interactions with the host organism. In *Probiotics, other nutritional factors, and intestinal microflora, vol 42,* pp 1-16 [Hanson LA & Yolken RH editors] Philadelphia: /Lippincott-Raven Publishers.

van Dokkum W, Wezendonk B, Srikumar TS & van den Heuvel EG (1999) Effect of nondigestible oligosaccharides on large-bowel functions, blood lipid concentrations and glucose absorption in young healthy male subjects *Eur J Clin Nutr* **53**(1), 1-7.

van Laere KMJ, Hartemink R, Beldman G, Pitson S, Dijkema C, Schols, HA & Voragen AGJ (1999) Transglycosidase activity of *Bifidobacterium adolescentis* DSM 20083 α -galactosidase *Appl Microbiol Biotechnol* **52**, 681-688.

van Loo JAE, Coussement P, De Leenheer L, Hoebregs H & Smits G (1995) On the presence of inulin and oligofructose as natural ingredients in the Western diet. *CRC Crit Rev Food Sci Nutr* **35**, 525-552.

Videla S (1999) Deranged luminal pH homeostasis in experimental colitis can be restored by a prebiotic *Gastroenterol* **116**, A942.

Videla S, Vilaseca J, Antolin, M, Garcia-Lafuente A, Guarner F, Crespo E, Casalots J, Salas A & Malagelada JR (2001) Dietary inulin improves distal colitis induced by dextran sodium sulfate in the rat *Am J Gastroenterol* **96**, 1486-1493.

Villaume C, Beck B, Gariot P, Desalme A & Debry G (1984) Long-term evolution of the effect of bran ingestion on meal-induced glucose and insulin responses in healthy man *Am J Clin Nutr* **40**(5), 1023-1026.

Vuksan V, Jenkins DJ, Vidgen E, Ransom TP, Ng MK, Culhane CT & O'Connor D (1999) A novel source of wheat fiber and protein: effects on fecal bulk and serum lipids *Am J Clin Nutr* **69**, 226-30.

Vuksan V, Sievenpiper JL, Owen R, Swilley JA, Spadafora P, Jenkins DJ, Vidgen E, Brighenti F, Josse RG, Leiter LA, Xu Z & Novokmet R (2000) Beneficial effects of viscous dietary fiber from Konjac-mannan in subjects with the insulin resistance syndrome: results of a controlled metabolic trial *Diab Care* 23(1), 9-14.

Wada K, Watabe J, Mizutani J, Tomoda M, Suzuki H & Saitoh Y (1992) Effects of soybean oligosaccharides in a beverage on human fecal flora and metabolites *J Agric Chem Soc Jap* **66**, 127-135.

Wakai K, Date C, Fukui M, Tamakoshi K, Watanabe Y, Hayakawa N et al (2007) Dietary fiber and risk of colorectal cancer in the Japan collaborative cohort study. *Cancer Epidemiol Biomarkers Prev* **16**(4), 668-675.

Walsh DE, Yaghoubian V & Behforooz A (1984) Effect of glucomannan on obese patients: a clinical study *Int J Obes* **8**(4), 289-93.

Wang X & Gibson GR (1993) Effects of the in vitro fermentation of oligofructose and inulin by bacteria growing in the human large intestine *J Appl Bacteriol* **75**, 373-380.

Weickert MO, Mohlig M, Koebnick C, Holst JJ, Namsolleck P, Ristow M, Osterhoff M, Rochlitz H, Rudovich N, Spranger J & Pfeiffer AF (2005) Impact of cereal fibre on glucose-regulating factors *Diabetologia* **48**(11), 2343-53.

Weickert MO, Mohlig M, Schofl C, Arafat AM, Otto B, Viehoff H, Koebnick C, Kohl A, Spranger J & Pfeiffer AF (2006) Cereal fiber improves whole-body insulin sensitivity in overweight and obese women *Diabetes Care* **29**(4), 775-80.

Weinreich J, Pedersen O & Dinesen K (1977) Role of bran in normals: serum levels of cholesterol, triglyceride, calcium and total 3-alphahydroxycholanic acid and intestinal transit time *Acta Med Scand* **202**, 125-130.

Welters CF, Heineman E, Thunnissen FB, van den Bogaard AE, Soeters PB & Baeten CG (2002) Effect of dietary inulin supplementation on inflammation of pouch mucosa in patients with an ileal pouch-anal anastomosis *Dis Colon Rectum* **45**, 621-627.

Whelan K, Efthymiou L, Judd PA, Preedy VR & Taylor MA (2006) Appetite during consumption of enteral formula as a sole source of nutrition: the effect of supplementing pea-fibre and fructo-oligosaccharides Br J Nutr **96**(2), 350-6.

White LA, Newman MC, Comwell GL & Lindemann MD (2002) Brewers dried yeast as a source of mannan oligosaccharides for weaning pigs *J Anim Sci* **80**, 2619-2628.

Willett WC, Stampfer MJ, Colditz GA, Rosner BA & Speizer FE (1990) Relation of meat, fat, and fiber intake to the risk of colon cancer in a prospective study among women *N Engl J Med* **323**(24), 1664-1672.

Williams CH, Witherly SA & Buddington RK (1994) Influence of dietary neosugar on selected bacterial groups of the human faecal microbiota *Microb Ecol Health Dis* **7**, 91-97.

Wilmshurst P & Crawley JC (1980) The measurement of gastric transit time in obese subjects using 24Na and the effects of energy content and guar gum on gastric emptying and satiety Br J Nutr 44(1), 1-6.

Wisker E, Maltz A & Feldheim W (1988) Metabolizable energy of diets low or high in dietary fiber from cereals when eaten by humans *J Nutr* **118**(8), 945-52.

Wolever TM, Vuksan V, Eshuis H, Spadafora P, Peterson RD, Chao ES, Storey ML & Jenkins DJ (1991) Effect of method of administration of psyllium on glycemic response and carbohydrate digestibility *J Am Coll Nutr* **10**(4), 364-71.

Wolever TM, Campbell JE, Geleva D & Anderson GH (2004) High-fiber cereal reduces postprandial insulin responses in hyperinsulinemic but not normoinsulinemic subjects *Diabetes Care* 27(6), 1281-5.

Wolever TM, Schrade KB, Vogt JA, Tsihlias EB & McBurney MI (2002) Do colonic short-chain fatty acids contribute to the long-term adaptation of blood lipids in subjects with type 2 diabetes consuming a high-fiber diet? *Am J Clin Nutr* **75**, 1023-30.

Wolf BW, Wolever TM, Lai CS, Bolognesi C, Radmard R, Maharry KS, Garleb KA, Hertzler SR & Firkins JL (2003) Effects of a beverage containing an enzymatically induced-viscosity dietary fiber, with or without fructose, on the postprandial glycemic response to a high glycemic index food in humans *Eur J Clin Nutr* **57**(9), 1120-7.

Wolk A, Manson JE, Stampfer MJ, Colditz GA, Hu FB, Speizer FE, Hennekens CH & Willett WC (1999) Long-term intake of dietary fiber and decreased risk of coronary heart disease among women *JAMA* **281**, 1998-2004.

Wood PJ, Braaten JT, Scott FW, Riedel KD, Wolynetz MS & Collins MW (1994) Effect of dose and modification of viscous properties of oat gum on plasma glucose and insulin following an oral glucose load Br J *Nutr* **72**(5), 731-43.

Wood PJ, Arrigoni E, Miller SS & Amado R (2002) Fermentability of oat and wheat fractions enriched in β -glucan using human faecal inoculation *Cereal Chem* **79**, 445-454.

Wood RJ, Fernandez ML, Sharman MJ, Silvestre R, Greene CM, Zern TL, Shrestha S, Judelson DA, Gomez AL, Kraemer WJ & Volek JS (2007) Effects of a carbohydrate-restricted diet with and without supplemental soluble fiber on plasma low-density lipoprotein cholesterol and other clinical markers of cardiovascular risk *Metabolism* **56**, 58-67.

World Cancer Research Fund/ American Institute of Cancer Research. (2007) Food Nutrition, Physical Activity, and the Prevention of Cancer: a Global Perspective Washington DC: AICR.

Wullt M, Hagslätt MLJ & Odenholt I (2003) *Lactobacillus plantarum* 299v for the treatment of recurrent *Clostridium difficile*-associated diarrhoea: a double-blind, placebo-controlled trial *Scand J Infect Dis* **35**, 365-367.

Yamada Y, Hosoya S, Nishimura S, Tanaka T, Kajimoto Y, Nishimura A & Kajimoto O (2005) Effect of bread containing resistant starch on postprandial blood glucose levels in humans *Biosci Biotechnol Biochem* **69**(3), 559-66.

Yamano T, Lino H, Takada M, Blum S, Rochat F & Fukushima Y (2006) Improvement of the human intestinal flora by ingestion of the probiotic strain *Lactobacillus johnsonii* La1 *Br J Nutr* **95**, 303-312.

Ylonen K, Saloranta C, Kronberg-Kippila C, Groop L, Aro A & Virtanen SM (2003) Associations of dietary fiber with glucose metabolism in nondiabetic relatives of subjects with type 2 diabetes: the Botnia Dietary Study *Diabetes Care* **26**(7), 1979-85.

Yudkin J (1959) The causes and cure of obesity Lancet 2(7112), 1135-8.

Zhang C, Liu S, Solomon CG & Hu FB (2006) Dietary fiber intake, dietary glycemic load, and the risk for gestational diabetes mellitus *Diab Care* **29**(10), 2223-30.

Ziegler, E., Vanderhoof JA, Petschow B, Mitmesser SH, Stolz SI, Harris CL & Berseth CL (2007) Term infants fed formula supplemented with selected blends of prebiotics grow normally and have soft stools similar to those reported for breast-fed infants *J Ped Gastroenterol Nutr* **44**, 359-364.

Zunft HJ, Luder W, Harde A, Haber B, Graubaum HJ, Koebnick C & Grunwald J (2003) Carob pulp preparation rich in insoluble fibre lowers total and LDL cholesterol in hypercholesterolemic patients *Eur J Nutr* **42**, 235-42.