



Department  
of Health

# The Adult Social Care Outcomes Framework 2013/14

Handbook of Definitions

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# Handbook of Definitions

**Prepared by the Department of Health**

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# Introduction

The Adult Social Care Outcomes Framework for 2013/14 was published on 22 November 2012. This technical handbook of definitions should be read in conjunction with the framework document.<sup>1</sup> Both the framework and this handbook have been developed by the Department of Health (DH), the Association of Directors of Adult Social Services (ADASS), and the Local Government Association (LGA).

The Adult Social Care Outcomes Framework (ASCOF) is used both locally and nationally to set priorities for care and support, measure progress and strengthen transparency and accountability.

The purpose of the ASCOF is three-fold:

- Locally, the ASCOF supports councils to improve the quality of care and support. By providing robust, nationally comparable information on the outcomes and experiences of local people, the ASCOF supports meaningful comparisons between councils, helping to identify priorities for local improvement and stimulating the sharing of learning and best practice;
- The ASCOF fosters greater transparency in the delivery of adult social care, supporting local people to hold their council to account for the quality of the services they provide. A key mechanism for this is through councils' local accounts, where the ASCOF is already being used as a robust evidence base to support councils' reporting of their progress and priorities to local people; and,
- Nationally, the ASCOF measures the performance of the adult social care system as a whole and its success in delivering high-quality, personalised care and support. The framework will support Ministers in discharging their accountability to the public and Parliament for the adult social care system and will inform and support national policy development.

The Government will not seek to performance manage councils in relation to any of the measures set out in this framework. Instead, the ASCOF will inform and support improvement led by the sector itself, underpinned by strengthened transparency and local accountability.

The Care and Support White Paper, published in July 2012, set out the Government's vision for a reformed care and support system, building on the 2010 Vision for Adult Social Care, and Transparency in Outcomes: a framework for quality in adult social care<sup>2</sup>. The ASCOF for 2013/14 will support councils to rise to the challenge of delivering key White Paper priorities by providing a clear focus for local priority setting and improvement and by strengthening the accountability of councils to local people.

This handbook sets out the detailed definition of each ASCOF measure with worked examples, to support consistency in reporting and interpretation of the measures. The intended audience for this handbook is local authorities, members of the public and other stakeholders with an interest in social care outcomes, such as health and wellbeing boards and the voluntary sector.

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<sup>1</sup> <http://www.dh.gov.uk/health/2012/11/ascof1314/>

<sup>2</sup> [http://webarchive.nationalarchives.gov.uk/+www.dh.gov.uk/en/Aboutus/Features/dh\\_121664](http://webarchive.nationalarchives.gov.uk/+www.dh.gov.uk/en/Aboutus/Features/dh_121664)

Through Joint Strategic Needs Assessments (JSNAs), health and wellbeing boards will identify the current and future health and care needs of the local population, building a robust evidence base of local needs and also looking at local assets available. From this, boards will develop Joint Health and Wellbeing Strategies (JHWSs), to drive local services by setting the framework for NHS, public health and social care commissioning, and delivering improved outcomes for local communities.

Health and wellbeing boards will have an interest in where the NHS, Public Health and Adult Social Care Outcomes Frameworks overlap locally, and the measures in the frameworks can help inform their local priorities (as well as other local evidence), and be used to transparently demonstrate their progress in improving outcomes to their community. However, the measures from the outcomes frameworks are not intended to overshadow local evidence to inform JSNAs and JHWSs. Where the NHS, Public Health and Adult Social Care Outcomes Frameworks (and future Child Health Outcomes Strategy) come together, local partners will be able to see how well they are delivering integrated services for their communities, especially around specific health and care issues.

Performance against the ASCOF, at both the national and individual council level, will be published annually by the Health and Social Care Information Centre, and the Department will also release an annual commentary on the national picture.

Measures will be broken down by certain equality characteristics, to show how the outcomes achieved by people and their experiences of care and support, might differ between groups.

# Development of the Adult Social Care Outcomes Framework

The ASCOF was first published in March 2011 and is updated annually in partnership with local government. For 2013/14, the ASCOF has been strengthened with the addition of new measures, reflecting the Department's and local government's key priorities for social care. In the development of new measures, we remain mindful of the reporting burden on councils and the need to retain a focus on measuring the success of the adult social care system in delivering high-quality care and support that is of use both nationally and locally.

There have been three key drivers for change to the ASCOF for 2013/14: the Care and Support White Paper, the 'zero-based review' of adult social care data collections from local authorities, and increasing alignment with the NHS and Public Health Outcomes Frameworks.

## The Care and Support White Paper

This year, the ASCOF has been augmented to better reflect key White Paper priorities for care and support. Although many of the themes of the White Paper were already reflected in the framework, it has been strengthened in some areas – including the promotion of care and support which is integrated and joins up around the needs of the people who use them, and the need to tackle loneliness and social isolation in our communities. It also reflects the Government's objective to improve the quality of life for people with dementia.

## The Zero-Based Review

Since November 2010, the Health and Social Care Information Centre has led a 'zero-based review' of adult social care collections from local authorities. The objective of the review is to ensure that the information we collect nationally reflects the changing face of adult social care – keeping pace with the transformation of care towards more personalised, preventative forms of support. The review has actively supported the development of the ASCOF for 2013/14 and beyond, and when the proposals are fully implemented they will support several new measures. Where new ASCOF measures rely on changes to collections after 2013/14, measures will be implemented in full when the supporting data are available. Placeholders have been used to indicate the nature of the proposed measures.

## Alignment of the Outcomes Frameworks

The Department has committed to improve alignment between the ASCOF and the Public Health and NHS Outcomes Frameworks, reflecting the joint contribution of health and social care to improving outcomes. The 2013/14 ASCOF and NHS Outcomes Framework and the technical refresh of the Public Health Outcomes Framework include an increased number of shared and complementary measures and placeholders.<sup>3</sup> In November, the Department also published, *'Improving health and care: the role of the Outcomes Frameworks.'*<sup>4</sup> This document sets out how the three frameworks work together to support improvement in outcomes for people who use health and care services and the public.

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<sup>3</sup> A measure is shared when the same measure appears in more than one framework. A measure is complementary when a similar measure addressing the same issue features in more than one of the frameworks.

<sup>4</sup> <http://www.dh.gov.uk/health/2012/11/health-care-of/>



## Future Developments

The ASCOF is a key mechanism by which the Government sets national priorities for adult social care, and measures national progress against those priorities. The Department will continue to use national performance against the ASCOF to inform national policy-making and will work with local government to ensure the continued focus of the framework on measuring the success of the adult social care system in delivering high-quality care and support.

The Department has commissioned a number of pieces of research to support the development of the ASCOF, including work on generating a social care-related quality of life 'value added' measure (described in more detail below) and further work by the Personal Social Services Research Unit on promoting the use of Adult Social Care Survey data locally. The objective of this work is to demonstrate the purpose and value of surveys locally and further information will be available at [www.maxproject.org.uk](http://www.maxproject.org.uk) later in the spring.

# Changes to the Adult Social Care Outcomes Framework for 2013/14

This section outlines the changes introduced to the ASCOF for 2013/14. Where placeholders have been added, or the implementation of new measures deferred until 2014/15, this reflects ongoing development work and the need to secure underpinning data sources before the measure can be implemented.

## New measures for 2013/14

A new measure of social isolation for users of care and support and carers has been added to Domain 1, in response to the key White Paper commitment to address loneliness and social isolation (measure shared with the Public Health Outcomes Framework 1.18)

## Changes deferred to 2014/15

The Department has committed to revise measure 1C (*Proportion of people using social care who receive self-directed support, and those receiving direct payments*) from 2014/15, to better reflect local authorities' progress in delivering personalised services. This revision is dependent on the full proposals of the zero-based review being implemented.

A new measure of the effectiveness of reablement care in supporting people to maintain their independence has been added to Domain 2, for implementation in 2014/15. This measure will provide evidence of a good outcome in delaying dependency or supporting recovery – short-term support that results in no further need for care. The inclusion of this measure is dependent on the full proposals of the zero-based review being implemented.

## New placeholders for 2013/14

A new placeholder measure has been added to Domain 2, to support interpretation of the new measure of effectiveness of reablement services when implemented. This is intended to support a more rounded view of the success of short-term support in supporting people to recover their independence.

A new placeholder measure on the effectiveness of post-diagnosis care in sustaining independence and improving quality of life for people with dementia has also been added to Domain 2. This is a priority area for the ASCOF and will promote joined up working across adult social care and the NHS (measure shared with the NHS Outcomes Framework 2.6ii).

A new placeholder measure on improving people's experience of integrated care (measure shared with the NHS Outcomes Framework 4.9) has been added to Domain 3, in response to the Care and Support White Paper, which restated the Department's commitment to a clear, ambitious and measurable goal to drive further improvements in people's experiences of integrated care.

A new placeholder measure has been added to Domain 4 on the proportion of completed safeguarding referrals where people report they feel safe, reflecting the Care and Support White Paper's statement that a high-quality service must be one which keeps people safe from harm.

# Using the Handbook of Definitions

The handbook sets out the following information for each measure:

Detail	Description
Title	Identifier (1A, 1B, etc) and name of the measure as it appears in the ASCOF
Domain/Outcome	The Domain of the ASCOF in which the measure appears and the associated outcome statement within the domain
Rationale	A brief description of the rationale for the inclusion of the measure
Definition/interpretation	Guidance on the definition of the measure, including the definition of related terms and any notes on interpretation
Alignment	Whether the measure is shared or complementary with a measure in the Public Health or NHS Outcomes Frameworks
Risk adjustment	Comments on factors that could affect the comparability of the measure, for example age distribution of the local population, and possible adjustments to support more meaningful comparisons between areas
Formula	The detail of how the measure will be calculated, with a formula and precise definitions of each component (for example, the source table of a data collection or question in a survey)
Worked example	An example of how this formula would be applied to a particular set of data to yield the measure
Disaggregation available	A list of client groups and equality groups by which the measure can be disaggregated, to identify outcomes for different groups and highlight any equality issues
Frequency of collection	How frequently the data will be disseminated – biennially, annually or more frequently
Data source	The data collection or survey from which the measure is drawn – in some cases this may combine data from more than one source
Return format	Whether the measure will be presented as a percentage or as a number
Decimal places	Number of decimal places used in the presentation of the measure
Longer-term development options	Potential improvements or alternatives to current measures to be explored for future iterations of the ASCOF
Further guidance	Where to find further guidance relating to the data collections underpinning the measure

## Risk adjustment

The aim of risk adjustment of the measures is to improve meaningful comparisons between different local authorities by controlling for factors which are not within the power of local authorities to influence. An example of this is measure 2A (permanent admissions to residential and nursing care homes, per 100,000 population). This measure is presented as two separate measures, one covering those aged 18 to 64 and the other covering those aged 65 and over, to reflect that the likelihood of admissions to residential and nursing care is related to the age of the client.

However, risk adjustment can also make measures more difficult to understand and interpret. As a result, risk adjustment should only be applied when the improvement in the comparability of the measure is significant enough to outweigh the additional complexity in understanding a risk-

adjusted measure. Where risk adjustment is not thought to be appropriate, the current practice of comparing councils with similar authorities can be used for benchmarking purposes.

This handbook sets out suggested factors which could be explored for the risk adjustment of measures. Decisions on whether risk adjustment is justified and should be applied should be made on a case-by-case basis.

# The Adult Social Care Outcomes Framework for 2013/14

## Domain 1 – Enhancing quality of life for people with care and support needs

(1A) Social care-related quality of life	
<b>Domain / Outcome</b>	1. Enhancing quality of life for people with care and support needs ( <i>Overarching Measure</i> )
<b>Rationale</b>	This measure gives an overarching view of the quality of life of users of social care. It is based on the outcome domains of social care-related quality of life identified in the Adult Social Care Outcomes Toolkit (ASCOT) developed by the Personal Social Services Research Unit ( <a href="http://www.pssru.ac.uk/ascot">www.pssru.ac.uk/ascot</a> ) <sup>5</sup>
<b>Definition / Interpretation</b>	<p>This measure is an average quality of life score based on responses to the Adult Social Care Survey. It is a composite measure using responses to survey questions covering the eight domains identified in the ASCOT.<sup>6</sup></p> <p>(control, dignity, personal care, food and nutrition, safety, occupation, social participation and accommodation). The relevant questions are listed below:</p> <ul style="list-style-type: none"> <li>▪ <i>Control</i> - Q3a: Which of the following statements best describes how much control you have over your daily life?</li> <li>▪ <i>Personal care</i> - Q4a: Thinking about keeping clean and presentable in appearance, which of the following statements best describes your situation?</li> <li>▪ <i>Food and Nutrition</i> - Q5a: Thinking about the food and drink you get, which of the following statements best describes your situation?</li> <li>▪ <i>Accommodation</i> - Q6a: Which of the following statements best describes how clean and comfortable your home/care home is?</li> <li>▪ <i>Safety</i> - Q7a: Which of the following statements best describes how safe you feel?</li> <li>▪ <i>Social participation</i> - Q8a: Thinking about how much contact you've had with people you like, which of the following statements best describes your social situation?</li> <li>▪ <i>Occupation</i> - Q9a: Which of the following statements best describes how you spend your time?</li> <li>▪ <i>Dignity</i> - Q11: Which of these statements best describes how the way you are helped and treated makes you think and feel about yourself?</li> </ul> <p>Each of the questions has four possible answers, which are equated with having:</p> <ul style="list-style-type: none"> <li>▪ no unmet needs in a specific life area or domain (the ideal state);</li> <li>▪ needs adequately met;</li> <li>▪ some needs met, and;</li> <li>▪ no needs met.</li> </ul> <p>Responses to the questions indicate whether the individual has unmet needs in any of the eight areas. The measure gives an overall score based on respondents' self-reported quality of life across the eight questions. All eight questions are given equal weight.</p> <p><b>Interpretation</b></p> <p>Guidance on the interpretation of this measure is presented in Appendix 4 to this document. The measure gives an overall indication of reported outcomes for individuals – it</p>

<sup>5</sup> The „ASCOT“ (Adult Social Care Outcomes Toolkit) measure (1A) is designed to capture information about an individual's social care-related quality of life (SCRQoL). The ASCOT is also the source for the questions in the Adult Social Care Survey. Users wishing to make commercial use of any of the ASCOT materials should contact the ASCOT team ([ascot@kent.ac.uk](mailto:ascot@kent.ac.uk)), who will then be put into contact with Kent Innovation and Enterprise, as people need to register to use the ASCOT. Also see <http://www.pssru.ac.uk/ascot/>.

<sup>6</sup> The „ASCOT“ (Adult Social Care Outcomes Toolkit) measure (1A) is designed to capture information about an individual's social care-related quality of life (SCRQoL).

	does not, at present, identify the contribution of councils' adult social care services towards those outcomes (see longer-term development options below).					
<b>Alignment</b>	This measure is complementary with Measure 2 (health-related quality of life for people with long-term conditions) in the NHS Outcomes Framework ( <a href="https://www.wp.dh.gov.uk/publications/files/2012/11/121109-Technical-Appendix.pdf">https://www.wp.dh.gov.uk/publications/files/2012/11/121109-Technical-Appendix.pdf</a> page 29). Health-related quality of life is measured using the EQ5D tool.					
<b>Risk adjustment</b>	<p>A range of factors will be considered to adjust the measure to improve comparability between councils. Some example are:</p> <ul style="list-style-type: none"> <li>▪ Age of users</li> <li>▪ Needs of users</li> <li>▪ Client groups of users</li> </ul> <p>See the section on longer term developments for information about progress in this area.</p>					
<b>Formula</b>	$\left( \frac{X}{Y} \right)$ <p>Where:</p> <p><b>X:</b> Each respondent is assigned a score based on their answers to questions 3a to 9a and 11. Higher scores are assigned to better outcomes. Scores are assigned as follows:</p> <ul style="list-style-type: none"> <li>▪ No needs met (the last answer option for each question) = 0</li> <li>▪ Some needs met (3rd answer option) = 1</li> <li>▪ Needs adequately met (2nd answer option) = 2</li> <li>▪ No unmet needs (1st answer option) = 3</li> </ul> <p>The numerator is then the sum of the scores for all respondents who have answered questions 3a to 9a and 11.</p> <p>The responses of respondents who were sent the version of the questionnaire for people with a learning disability will be treated in the same way, as this questionnaire has been designed to be equivalent to the non-learning disabilities version.</p> <p><b>Y:</b> The number of respondents who answered questions 3a to 9a and 11.</p> <p>For both the numerator (X) and denominator (Y), weighted data should be used to calculate the measure. The data from the survey will be weighted by the Health and Social Care Information Centre to take account of the stratified sampling technique that has been used when conducting the survey. The weights are automatically calculated within the survey data return along with the ASCOF outcome measures. Further details of how to use the weights when analysing the survey data are available in Appendix I of the guidance for the 2012-13 Adult Social Care Survey (<a href="http://www.ic.nhs.uk/CHttpHandler.ashx?id=10142&amp;p=0">http://www.ic.nhs.uk/CHttpHandler.ashx?id=10142&amp;p=0</a>).</p> <p><b>Exclusions</b></p> <p>Any respondents who failed to answer all of the questions from 3a to 9a and question 11, are excluded from the calculation of the measure. For example, a respondent who answered questions 3a to 8a and 11 but did not answer 9a would be excluded from the calculation.</p>					
<b>Worked example</b>	<p>The table below represents the responses of 145 users who answered questions 3a to 9a and 11. The data has been weighted to reflect the stratified sampling technique used when conducting the survey.</p> <table style="margin-left: auto; margin-right: auto;"> <thead> <tr> <th>No unmet needs</th> <th>Needs adequately</th> <th>Some needs</th> <th>No needs met</th> <th>Total</th> </tr> </thead> </table>	No unmet needs	Needs adequately	Some needs	No needs met	Total
No unmet needs	Needs adequately	Some needs	No needs met	Total		

		met	met			
	Control (Q3a)	20	58	48	19	145
	Personal Care (Q4a)	65	65	15	0	145
	Food and Nutrition (Q5a)	78	65	2	0	145
	Accommodation (Q6a)	45	36	55	9	145
	Safety (Q7a)	30	75	35	5	145
	Social Participation (Q8a)	36	46	36	27	145
	Occupation (Q9a)	28	51	46	20	145
	Dignity (Q11)	33	47	47	18	145
	Total	335	443	284	98	
	<p>Scores are assigned as follows:</p> <ul style="list-style-type: none"> <li>▪ No unmet needs (1st answer option) = 3</li> <li>▪ Needs adequately met (2nd answer option) = 2</li> <li>▪ Some needs met (3rd answer option) = 1</li> <li>▪ No needs met (the last answer option for each question) = 0</li> </ul> <p>Higher scores are assigned to better outcomes, so the higher the overall score the better the average social care-related quality of life.</p> <p>The numerator for the measure is [(335*3)+ (443*2)+(284*1)+(98*0)]=2,175</p> <p>The denominator for the measure is 145</p> <p>Therefore the measure value is 2175/145 which equals 15.0</p>					
<b>Disaggregation available</b>	<b>Equalities:</b> Age, Gender, Ethnicity**, Religion*, Sexual orientation*					
	<b>Client groups:</b> Physical Disability (all ages)**, Learning Disability (all ages)**, Mental Health (all ages), Substance Misuse (all ages)**, Vulnerable People (all ages)**					
<b>Frequency of collection</b>	Annual	<b>Data source</b>		Adult Social Care Survey (ASCS)		
<b>Return format</b>	Numeric	<b>Decimal places</b>		One		
<b>Longer-term development options</b>	<p>The social care-related quality of life measure tells us about outcomes for social care users but does not isolate the impact that care and support services have on those outcomes. The Department has commissioned research from the Quality and Outcomes of Person Centred Care Policy Research Unit to identify a way of generating a social care-related quality of life 'value added' measure, which would allow us to identify the impact of adult social care on people's quality of life. This research is due to report its findings in summer 2014. If successful, this will allow us to develop a new or additional measure for the ASCOF.</p>					
<b>Further guidance</b>	<p>Guidance for 2012/13 onwards can be found via the user survey guidance page at <a href="http://www.ic.nhs.uk/article/2213/User-survey-guidance-2012-13">http://www.ic.nhs.uk/article/2213/User-survey-guidance-2012-13</a>.</p>					

\*\* This information is not published as part of the adult social care outcomes returns; however it is part of the publication of the data source.

\* Although the underlying survey results will in theory be disaggregated by religion and sexual orientation, in practice there are likely to be significant gaps in the data on these characteristics at least in the short/medium term. This reflects the content of records held locally by councils.

<b>(1B) The proportion of people who use services who have control over their daily life</b>	
<b>Domain / Outcome</b>	1. Enhancing quality of life for people with care and support needs. <i>People manage their own support as much as they wish, so that they are in control of what, how and when support is delivered to match their needs.</i>
<b>Rationale</b>	<p>A key objective of the drive to make care and support more personalised is that support more closely matches the needs and wishes of the individual, putting users of services in control of their care and support. Therefore, asking users of care and support about the extent to which they feel in control of their daily lives is one means of measuring whether this outcome is being achieved.</p> <p>This measures one component of the overarching measure 1A – social care-related quality of life. A preference study conducted by RAND<sup>7</sup> found that members of the public gave this domain the highest weight of the eight included, i.e. of all the domains included in the overarching measure, this is the one that is considered by the public to be the most important.</p>
<b>Definition / Interpretation</b>	<p>The relevant question drawn from the Adult Social Care Survey is Question 3a: ‘Which of the following statements best describes how much control you have over your daily life?’, to which the following answers are possible:</p> <ul style="list-style-type: none"> <li>▪ I have as much control over my daily life as I want</li> <li>▪ I have adequate control over my daily life</li> <li>▪ I have some control over my daily life but not enough</li> <li>▪ I have no control over my daily life</li> </ul> <p>The measure is defined by determining the percentage of all those responding who identify no needs in this area or no needs with help – i.e. by choosing the answer ‘I have as much control over my daily life as I want’ or ‘I have adequate control over my daily life’. These have been chosen to focus the measure on those individuals achieving the best outcomes, to allow for better use in benchmarking.</p> <p><b>Interpretation</b></p> <p>The measure gives an overall indication of the reported outcome for individuals – it does not, at present, identify the specific contribution of councils’ adult social care towards the outcome (see longer-term development below).</p>
<b>Alignment</b>	ASCOF measure only
<b>Risk adjustment</b>	<p>A range of factors will be considered to adjust the measure to improve comparability between councils. Some examples are:</p> <ul style="list-style-type: none"> <li>▪ Age of users</li> <li>▪ Needs of users</li> <li>▪ Client groups of users</li> </ul>
<b>Formula</b>	$\left(\frac{X}{Y}\right) * 100$ <p>Where:</p> <p><b>X:</b> In response to Question 3a, those individuals who selected the response ‘I have as much control over my daily life as I want’ and ‘I have adequate control over my daily life’.</p>

<sup>7</sup> Burge, P et al (2010) How do the public value different social care outcomes? Estimation of preference weights for ASCOT



	<p>The responses of respondents who were sent the version of the questionnaire for people with a learning disability will be treated in the same way, as this questionnaire has been designed to be equivalent to the non-learning disabilities version.</p> <p><b>Y:</b> All those that respond to the question.</p> <p>For both the numerator (X) and denominator (Y), weighted data should be used to calculate the measure. The data from the survey will be weighted by the Health and Social Care Information Centre to take account of the stratified sampling technique that has been used when conducting the survey. The weights are automatically calculated within the survey data return along with the ASCOF outcome measures. Further details of how to use the weights when analysing the survey data are available in Appendix I of the guidance for the 2012-13 Adult Social Care Survey (<a href="http://www.ic.nhs.uk/CHttpHandler.ashx?id=10142&amp;p=0">http://www.ic.nhs.uk/CHttpHandler.ashx?id=10142&amp;p=0</a>).</p>		
<b>Worked example</b>	<p>The number of users who said ‘I have as much control over my daily life as I want or ‘I have adequate control over my daily life’” was 156.</p> <p>In total the number of users who responded to the questions was 210.</p> <p>(Data weighted to reflect the stratified sampling technique that has been used when conducting the survey)</p> <p>The measure value is <math>[(156/210)*100] = 74.3\%</math></p>		
<b>Disaggregation available</b>	<b>Equalities:</b> Age, Gender, Ethnicity **, Religion *, Sexual orientation*		
	<b>Client groups:</b> Physical Disability (all ages)**, Learning Disability (all ages)**, Mental Health (all ages)**, Substance Misuse (all ages)**, Vulnerable People (all ages)**		
<b>Frequency of collection</b>	Annual	<b>Data source</b>	Adult Social Care Survey (ASCS)
<b>Return format</b>	Percentage	<b>Decimal places</b>	One
<b>Longer-term development options</b>	Develop a ‘value-added’ measure which quantifies the contribution of social services to quality of life as described for Measure 1A.		
<b>Further guidance</b>	Guidance for 2012/13 onwards can be found via the user survey guidance page at <a href="http://www.ic.nhs.uk/article/2213/User-survey-guidance-2012-13">http://www.ic.nhs.uk/article/2213/User-survey-guidance-2012-13</a>		

\*\* This information is not published as part of the adult social care outcomes returns; however it is part of the publication of the data source.

Although the underlying survey results will in theory be disaggregated by religion and sexual orientation, in practice there are likely to be significant gaps in the data on these characteristics at least in the short/medium term. This reflects the content of records held locally by councils.

<b>(1C) Proportion of people using social care who receive self-directed support, and those receiving direct payments</b>	
<b>Domain / Outcome</b>	1. Enhancing quality of life for people with care and support needs. <i>People manage their own support as much as they wish, so that they are in control of what, how and when support is delivered to match their needs.</i>
<b>Rationale</b>	<p>Research has indicated that personal budgets have a positive effect in terms of impact on well-being, increased choice and control, cost implications and improving outcomes.<sup>8</sup> Studies have shown that direct payments make people happier with the services they receive and are the purest form of personalisation.<sup>9</sup></p> <p>The Department remains committed to the 2013 objective for personal budgets; meaning everyone eligible for long-term community-based care should be provided with a personal budget, preferably as a Direct Payment, by April 2013. As there are recognised limitations in the current measure meaning that 100% provision of personal budgets is not possible nationally, the April 2013 objective was set at 70% of people receiving a personal budget nationally. The April 2013 objective, along with the proposals in the Care and Support White Paper and draft Bill to place personal budgets into legislation as part of the care and support plan, will ensure that personalised care becomes standard practice. This measure reflects the success of councils in delivering personalised services, through self-directed support, including direct payments.</p> <p><b>Measure 1C from 2014/15</b></p> <p>There are recognised limitations to the current measure, for example its scope includes some services and users of care and support for whom self-directed support may not be appropriate, and so it currently does not reflect the true extent of the provision of self-directed support and direct payments to those who are eligible.</p> <p>To address these issues, implementation of the proposals of the zero-based review will result in a strengthened measure 1C, by limiting its scope to people who only receive long-term support, for whom self-directed support is most relevant, to better reflect councils' progress in delivering personalised services. The existing measure will also be replaced by two measures: one which focuses on users; and another measure will be introduced which focuses on carers. Each will have a sub-measure for users/carers in receipt of direct payments, showing progress made on personalisation for users and carers separately. The final change is for this measure to be based on 'snapshot' rather than full-year data, to better reflect the progress made on personalisation at the end of the year.</p> <p><b>For 2013/14, the current definition of this measure remains.</b></p>
<b>Definition / Interpretation</b>	<p>This is a two-part measure which reflects both the proportion of people using services who receive self-directed support (1C part 1), and the proportion who receive a direct payment either through a personal budget or other means (1C part 2).</p> <p>1C part 1 is presented as the number of adults, older people and carers receiving self-directed support in the year to 31st March as a percentage of all clients receiving community based services and carers receiving carer specific services.<sup>10</sup></p> <p>To be counted as receiving self-directed support, the person (adult, older person or carer) must either:</p> <ul style="list-style-type: none"> <li>• be in receipt of a direct payment; or</li> </ul>

<sup>8</sup> Quoting; C Glendinning et al, The national evaluation of the Individual Budgets pilot programme (IBSEN (Individual Budgets Evaluation Network); Social Policy Research Unit, University of York, 2008); Individual Budgets: Impacts and outcomes for carers, (2009, IBSEN; Social Policy Research Unit, University of York); Choice and competition in public services: a guide for policy makers (2010, OFT/Frontier Economics)

<sup>9</sup> Quoting; Choice and competition in public services: a guide for policy makers (2010, OFT/Frontier Economics)

<sup>10</sup> For the purposes of this measure the following age brackets are used:

Adult: aged 18-64

Older person: aged 65 and over

Carer: aged 16 or over but caring for an adult aged 18 or over

	<ul style="list-style-type: none"> <li>• have in place a personal budget which meets all the following criteria: <ol style="list-style-type: none"> <li>1. The person (or their representative) has been informed about a clear, upfront allocation of funding, enabling them to plan their support arrangements; and</li> <li>2. There is an agreed support plan making clear what outcomes are to be achieved with the funding; and</li> <li>3. The person (or their representative) can use the funding in ways and at times of their choosing.</li> </ol> </li> </ul> <p>Councils will need to evidence that these criteria are met, for example through local monitoring of outcomes and satisfaction, as outlined in paragraph 69 of <i>Transforming social care (LAC (DH) (2008) 1)</i>.</p> <p><b>1C Part 1:</b></p> <p>The data collections will record for each category:</p> <p>i) people who have been through a self-directed support planning process:</p> <ul style="list-style-type: none"> <li>• people receiving a personal budget in the form of a direct payment for all or some of the package</li> <li>• people receiving a personal budget (based on the above definition) and who do not receive a direct payment</li> </ul> <p>ii) of people who have not been through a self-directed support planning process:</p> <ul style="list-style-type: none"> <li>• people receiving an existing or new direct payment (they may also be receiving other services).</li> </ul> <p><b>1C Part 2:</b></p> <p>Those receiving direct payments. The denominator remains the same (i.e. all adults and carers receiving community-based services), but the numerator captures only those from part 1 with direct payments.</p> <p><b>Interpretation</b></p> <p>There are established issues with the data definitions in relation to this measure, which means that care must be taken when interpreting the information for analysis and benchmarking.</p> <p>The denominator of the current measure is based upon a definition of people receiving community-based services which includes some individuals for whom self-directed support may not be appropriate, for instance those receiving some one-off, short-term or universal services such as equipment and reablement. This means the overall proportion does not reflect the true extent of the provision of self-directed support to those who are eligible and it is not possible to reach 100%. The Department has been working with the sector to address this issue, and recently agreed that for the April 2013 objective, a milestone of 70% is realistic.</p>
<b>Alignment</b>	ASCOF measure only
<b>Risk adjustment</b>	Risk adjustment does not seem appropriate for this measure since the objective is that self directed support is offered to all users regardless of ages, client group etc.
<b>Formula</b>	$\left(\frac{X}{Y}\right) * 100$ <p>Where, for <b>1C part 1</b> (receiving self-directed support):</p> <p><b>X:</b> The number of users and carers receiving self-directed support in the year to 31st</p>

	<p>March. Source: RAP Tables SD1 and SD3<sup>11</sup></p> <p><b>Y:</b> Clients receiving community-based services (aged 18 or over) and carers (caring for someone aged 18 or over) receiving carer-specific services in the year to 31<sup>st</sup> March. Source: RAP Table P2f<sup>12</sup> and Table C2<sup>13</sup></p> <p>For <b>1C part 2</b> (direct payments):</p> <p><b>X:</b> The number of users and carers receiving direct-payments in the year to 31st March. Source: RAP Tables SD1 and SD3<sup>14</sup></p> <p><b>Y:</b> Clients receiving community-based services (aged 18 or over) and carers aged (caring for someone aged 18 or over) receiving carer specific services in the year to 31<sup>st</sup> March. Source: RAP Table P2f and Table C2</p>		
<b>Worked example</b>	<p><b>1C part 1</b></p> <p>Suppose the total number of people who received self directed support (existing/new direct payment or personal budget) in the year 2011/12 to March 31 = 600, and the total number of people receiving community-based services and carers receiving carer-specific services = 2,000</p> <p>The measure value is <math>[(600/2000) \times 100] = 30.0\%</math></p> <p><b>1C part 2</b></p> <p>If the total number of people receiving a direct payment (whether part of a self directed process or not) is 195.</p> <p>Then the measure value is <math>[(195/2000) \times 100] = 9.8\%</math></p>		
<b>Disaggregation available</b>	<p><b>Equalities:</b> Age</p> <p><b>Client groups:</b> Physical Disabilities (18-64 and 65+**), Learning Disabilities (18-64 and 65+**), Mental Health (18-64, 65+), Substance Misuse (18-64, 65+**), Other vulnerable people (18-64, 65+**), Older People (65+), Carers (18-64, 65-74 and 75+)**</p>		
<b>Frequency of collection</b>	Annual	<b>Data source</b>	Referrals, Assessments and Packages of care (RAP) – table, page and cell references given here are based on the 2012/13 proforma
<b>Return format</b>	Percentage	<b>Decimal places</b>	One
<b>Longer-term development options</b>	<p>As above, this measure will be revised to focus only on those for whom self-directed support and direct payments are appropriate, which is not possible from the current data collections. This will give a better representation of the progress of the personalisation agenda and enable fairer benchmarking between councils.</p> <p>The development of the data collections required is being taken forward as part of the 'zero-based review' of social care data. Revisions to this measure are contingent on the implementation of the proposals of the zero-based review.</p>		
<b>Further guidance</b>	<p>Guidance for 2012/13 onwards can be found via the social care collection page at <a href="http://www.ic.nhs.uk/socialcare/collections">http://www.ic.nhs.uk/socialcare/collections</a> by clicking on the year.</p>		

<sup>11</sup> RAP SD1 Row 10 Column 5 (Total 18 and over) plus RAP SD3 Row 6 Column 5 (Total all ages)

<sup>12</sup> (Number of clients receiving community-based services during the period, provided or commissioned by the CASSR, by age group, primary client type and components of service) pages 1 and 3, row 11 (Total of above), column 1 (Total of clients)

<sup>13</sup> (Number of carers receiving different types of services as an outcome of assessment or review by age group of carer, and age group and primary client type of the person cared for by the carer) page 1, row 5 (all ages), column 1 (services including respite for the carer and/or other carer specific services)

<sup>14</sup> RAP SD1 Row 10 sum of columns 1, 2 and 4 (Total 18 and over) plus RAP SD3 Row 6 sum of columns 1, 2 and 4 (Total all ages)

\*\* This information is not published as part of the adult social care outcomes returns; however it is part of the publication of the data source.

(1D) Carer-reported quality of life	
<b>Domain / Outcome</b>	1. Enhancing quality of life for people with care and support needs. <i>Carers can balance their caring roles and maintain their desired quality of life.</i>
<b>Rationale</b>	This measure gives an overarching view of the quality of life of carers based on outcomes identified through research by the Personal Social Services Research Unit. This is the only current measure related to quality of life for carers available, and supports a number of the most important outcomes identified by carers themselves to which adult social care contributes.
<b>Definition / Interpretation</b>	<p><b>This measure will be deferred in 2013/14.</b></p> <p>This is a composite measure which combines individual responses to six questions measuring different outcomes related to overall quality of life. These outcomes are mapped to six domains (occupation, control, personal care, safety, social participation and encouragement and support). This is an overarching outcome measure for carers, similar to the equivalent for people who use services (1A – social care-related quality of life).</p> <p>The six questions, drawn from the Carers Survey, are:</p> <ul style="list-style-type: none"> <li>▪ <i>Occupation</i> – Q7: Which of the following statements best describes how you spend your time?</li> <li>▪ <i>Control</i> - Q8: Which of the following statements best describes how much control you have over your daily life?</li> <li>▪ <i>Personal care</i> - Q9: Thinking about how much time you have to look after yourself – in terms of getting enough sleep or eating well – which statement best describes your present situation?</li> <li>▪ <i>Safety</i> – Q10: Thinking about your personal safety, which of the statements best describes your present situation?</li> <li>▪ <i>Social participation</i> - Q11: Thinking about how much social contact you've had with people you like, which of the following statements best describes your social situation?</li> <li>▪ <i>Encouragement and support</i> - Q12: Thinking about encouragement and support in your caring role, which of the following statements best describes your present situation?</li> </ul> <p>Each of the questions has three possible answers, which are equated with having:</p> <ul style="list-style-type: none"> <li>▪ no unmet needs in a specific life area or domain (the ideal state);</li> <li>▪ some needs met, and;</li> <li>▪ no needs met.</li> </ul> <p>Responses to the questions indicate whether the carer has unmet needs in any of the six areas. The measure gives an overall score based on respondents' self-reported quality of life across the six questions. All six questions are given equal weight.</p> <p><b>Interpretation</b></p> <p>The measure gives an overall indication of the reported outcomes for carers – it does not, at present, identify the specific contribution of councils' adult social care services towards those outcomes.</p>
<b>Alignment</b>	This measure is complementary with Measure 2.4 (health-related quality of life for carers) in the NHS Outcomes Framework ( <a href="https://www.wp.dh.gov.uk/publications/files/2012/11/121109-Technical-Appendix.pdf">https://www.wp.dh.gov.uk/publications/files/2012/11/121109-Technical-Appendix.pdf</a> page 38). Health related quality of life is measured using the EQ5D tool.
<b>Risk adjustment</b>	<p>A range of factors may be considered to adjust the measure to improve comparability between councils. Some example are:</p> <ul style="list-style-type: none"> <li>▪ The intensity of the caring role</li> <li>▪ Age of carer</li> <li>▪ Characteristics of the cared for person</li> </ul>

<p><b>Formula</b></p>	$\left(\frac{X}{Y}\right)$ <p>Where:</p> <p><b>X:</b> Each respondent is assigned a score based on their answers to the six questions above. Each of the questions has three answers. Higher scores are assigned to better outcomes, Scores are assigned to answers as follows:</p> <ul style="list-style-type: none"> <li>▪ No needs met (the last answer option for each question) = 0</li> <li>▪ Some needs met (2nd answer option) = 1</li> <li>▪ No unmet needs (1st answer option) = 2</li> </ul> <p>The numerator is then a sum of the scores for all respondents who have answered all six questions.</p> <p><b>Y:</b> The number of respondents who answered all six questions.</p> <p><b>Exclusions</b></p> <p>Any respondents who failed to answer any of the six questions above are excluded from the calculation of the measure.</p>																																													
<p><b>Worked example</b></p>	<p>The table below represents the responses of 105 carers who answered all six questions.</p> <table border="1" data-bbox="352 902 1150 1272"> <thead> <tr> <th></th> <th>No unmet needs</th> <th>Some needs met</th> <th>No needs met</th> <th>Total</th> </tr> </thead> <tbody> <tr> <td>Occupation</td> <td>45</td> <td>45</td> <td>15</td> <td>105</td> </tr> <tr> <td>Control</td> <td>33</td> <td>52</td> <td>20</td> <td>105</td> </tr> <tr> <td>Personal Care</td> <td>65</td> <td>38</td> <td>2</td> <td>105</td> </tr> <tr> <td>Safety</td> <td>85</td> <td>20</td> <td>0</td> <td>105</td> </tr> <tr> <td>Social</td> <td></td> <td></td> <td></td> <td></td> </tr> <tr> <td>Participation</td> <td>58</td> <td>35</td> <td>12</td> <td>105</td> </tr> <tr> <td>Encouragement and Support</td> <td>22</td> <td>36</td> <td>47</td> <td>105</td> </tr> <tr> <td>Total</td> <td>308</td> <td>226</td> <td>96</td> <td></td> </tr> </tbody> </table> <p>Scores are assigned as follows:</p> <ul style="list-style-type: none"> <li>▪ No needs met (the last answer option for each question) = 0</li> <li>▪ Some needs met (2nd answer option) = 1</li> <li>▪ No unmet needs (1st answer option) = 2</li> </ul> <p>Higher scores are assigned to better outcomes so the higher the overall score the better the average social care related quality of life.</p> <p>The numerator for the measure is [(308*2)+ (226*1)+(96*0)]=842</p> <p>The denominator for the measure is 105.</p> <p>Therefore the measure value is 842/105 which equals 8.0.</p>		No unmet needs	Some needs met	No needs met	Total	Occupation	45	45	15	105	Control	33	52	20	105	Personal Care	65	38	2	105	Safety	85	20	0	105	Social					Participation	58	35	12	105	Encouragement and Support	22	36	47	105	Total	308	226	96	
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\*\* This information is not published as part of the adult social care outcomes returns; however it is part of the publication of the data source.

Although the underlying survey results will in theory be disaggregated by religion and sexual orientation, in practice there are likely to be significant gaps in the data on these characteristics at least in the short/medium term. This reflects the content of records held locally by councils.

<b>Frequency of collection</b>	Biennial (Carers Survey to be next conducted in 2014/15)	<b>Data source</b>	Carers Survey
<b>Return format</b>	Numeric	<b>Decimal places</b>	One
<b>Longer-term development options</b>	<p>There remains potential for moving to an annual Carers Survey if burden can be reduced significantly, subject to the agreement of local government.</p> <p>The research project to develop a value added measure for social care-related quality of life for users will also investigate whether it is possible to develop a value added measure for carer-reported quality of life. However, a value added measure for carers' quality of life is likely to be a longer-term piece of development beyond 2014.</p>		
<b>Further guidance</b>	<p>Guidance for 2012/13 onwards can be found via the user survey guidance page at <a href="http://www.ic.nhs.uk/article/2213/User-survey-guidance-2012-13">http://www.ic.nhs.uk/article/2213/User-survey-guidance-2012-13</a></p>		

### (1E) Proportion of adults with a learning disability in paid employment

<b>Domain / Outcome</b>	<p>1. Enhancing quality of life for people with care and support needs.</p> <p><i>People are able to find employment when they want, maintain a family and social life and contribute to community life, and avoid loneliness or isolation.</i></p>
<b>Rationale</b>	<p>The measure is intended to improve the employment outcomes for adults with a learning disability, reducing the risk of social exclusion. There is a strong link between employment and enhanced quality of life, including evidenced benefits for health and wellbeing<sup>15</sup> and financial benefits<sup>16</sup>.</p>
<b>Definition / Interpretation</b>	<p>The measure shows the proportion of all adults with a learning disability who are known to the council, who are recorded as being in paid employment. The information would have to be captured or confirmed within the reporting period 1 April 2013 to 31 March 2014.</p> <p>The definition of individuals '<b>known to the council</b>' is restricted to those adults with a learning disability (with a primary client group of LD) who have been assessed or reviewed by the council during the year (irrespective of whether or not they receive a service) or who should have been reviewed but were not.</p> <p>The measure is focused on 'paid' employment, to be clear that voluntary work is to be excluded for the purposes of this measure. Paid employment is measured using the following two categories:</p> <ul style="list-style-type: none"> <li>▪ Working as a paid employee or self-employed (16 or more hours per week); and,</li> <li>▪ Working as a paid employee or self-employed (up to 16 hours per week).</li> </ul> <p>A '<b>paid employee</b>' is one who works for a company, community or voluntary organisation, council or other organisation and is earning at or above the National Minimum Wage. This includes those who are working in supported employment (i.e. those receiving support from a specialist agency to maintain their job) who are earning at or above the National Minimum Wage.</p> <p>'<b>Self-employed</b>' is defined as those who work for themselves and generally pay their National Insurance themselves. This should also include those who are unpaid family workers (i.e. those who do unpaid work for a business they own or for a business a relative owns).</p> <p>The measure will not require collection of any further employment status (e.g. unpaid voluntary work); though councils may choose to provide this in addition to support their own benchmarking.</p>

<sup>15</sup> Vigna, E., Beyer, S. and Kerr, M. (2011) The role of supported employment agencies in promoting the health of people with learning disabilities. Cardiff: Welsh Centre for Learning Disabilities.

<sup>16</sup> Beyer, S. (2008) *An evaluation of the outcomes in supported employment in North Lanarkshire*. North Lanarkshire Social Work Service

<p><b>Alignment</b></p>	<p>This measure is complementary with Measure 1.6 (improved functional ability, and ability to work, in people with long-term conditions) in the Public Health Outcomes Framework (<a href="http://www.dh.gov.uk/prod_consum_dh/groups/dh_digitalassets/@dh/@en/documents/digitalasset/dh_132558.pdf">http://www.dh.gov.uk/prod_consum_dh/groups/dh_digitalassets/@dh/@en/documents/digitalasset/dh_132558.pdf</a> page 14) and Measure 2.2 (employment of people with long-term conditions) in the NHS Outcomes Framework (<a href="https://www.wp.dh.gov.uk/publications/files/2012/11/121109-Technical-Appendix.pdf">https://www.wp.dh.gov.uk/publications/files/2012/11/121109-Technical-Appendix.pdf</a> page 33).</p> <p>Although the Public Health Outcomes Framework and the ASCOF both include measures connected with employment for people with a learning disability and people with mental health problems, the Public Health Outcomes Framework measures the gap between the employment rate for those groups and the overall employment rate. This reflects the approach taken in the NHS Outcomes Framework for a complementary measure on employment of people with long term conditions. Although aligning the ASCOF with the other two frameworks was considered, in developing the framework with local government, it was agreed that this would not support local interpretation and benchmarking, and so the ASCOF will retain the current approach, with changes made to reflect the updated NMDS-MH dataset. Furthermore, although the Public Health Outcomes Framework uses the same data sources for rates of employment for these group as the ASCOF, the NHSOF uses the Labour Force Survey. This source cannot be used for the ASCOF because it does not provide robust results at the local authority level.</p>
<p><b>Formula</b></p>	$\left(\frac{X}{Y}\right) * 100$ <p><b>Where:</b></p> <p><b>X:</b> All people within the denominator, who are in employment. The numerator should include those recorded as in paid employment irrespective of whether the information was recorded in an assessment or review. However, the information would have to have been captured within the current financial year. <i>Source: Table L1, ASC-CAR<sup>17</sup></i></p> <p><b>Y:</b> Number of working-age learning-disabled clients known to CASSRs during the period. This includes:</p> <ul style="list-style-type: none"> <li>▪ those who are assessed or reviewed in the financial year and have received a service;</li> <li>▪ those who are assessed or reviewed in the financial year and have not received a service, and;</li> <li>▪ those who should have been reviewed in the financial year but were not.</li> </ul> <p>In other words, this is a count of eligible adults with learning disabilities (aged 18-64), without duplication, who are assessed, or reviewed (regardless of whether they have received a service), or should have been reviewed during the year; i.e. they are recorded within an A or P table within the RAP return.</p> <p><i>Source: Table L1, ASC-CAR<sup>18</sup></i></p>
<p><b>Worked example</b></p>	<p>Adults with learning disabilities known to CASSRs = 722</p> <p>Of those adults with learning disabilities known to CASSRs, those who are recorded as being in paid employment within the current financial year = 134</p> <p>The measure value = <math>(134/722) \times 100 = 18.6\%</math></p>
<p><b>Disaggregati</b></p>	<p><b>Equalities:</b> Gender</p>

<sup>17</sup> (Number of working age learning disabled clients known to CASSRs during year to 31<sup>st</sup> March, by service type and gender and by employment status at the time of their latest assessment or review) sum of rows 1 to 5 column 9

<sup>18</sup> (Number of working age learning disabled clients known to CASSRs during year to 31<sup>st</sup> March, by service type and gender and by employment status at the time of their latest assessment or review) row 9 column 9.



<b>on available</b>	<b>Client groups:</b> Learning disability (18-64)		
<b>Frequency of collection</b>	Annual	<b>Data source</b>	Adult Social Care Combined Activity Return (ASC-CAR) – table, page and cell references given here are based on the 2012/13 proforma.
<b>Return format</b>	Percentage	<b>Decimal places</b>	One
<b>Longer-term development options</b>	The 2012/13 ASCOF included a placeholder on employment for all people with a disability, which would replace the existing measure for adults in contact with secondary mental health services and for adults with a learning disability. This placeholder was to combine all adults into a single measure for employment status. This placeholder has been removed for the 2013/14 ASCOF, as, in consultation with local authorities, it was agreed that it would be too onerous to collect additional information on people with physical disabilities in employment in order to populate the proposed measure.		
<b>Further guidance</b>	Guidance for 2012/13 onwards can be found via the social care collection page at <a href="http://www.ic.nhs.uk/socialcare/usersurveys">http://www.ic.nhs.uk/socialcare/usersurveys</a> by clicking on the year.		

### (1F) Proportion of adults in contact with secondary mental health services in paid employment

<b>Domain / Outcome</b>	1. Enhancing quality of life for people with care and support needs. <i>People are able to find employment when they want, maintain a family and social life and contribute to community life and avoid loneliness or isolation.</i>
<b>Rationale</b>	The measure is of improved employment outcomes for adults with mental health problems, reducing their risk of social exclusion and discrimination. Supporting someone to become and remain employed is a key part of the recovery process <sup>19</sup> . Employment outcomes are a predictor of quality of life, and are indicative of whether care and support is personalised. Employment is a wider determinant of health and social inequalities.
<b>Definition/ Interpretation</b>	<p>The measure shows the percentage of adults receiving secondary mental health services in paid employment at the time of their most recent assessment, formal review or other multi-disciplinary care planning meeting.</p> <p>Adults <b>‘in contact with secondary mental health services’</b> is defined as those aged 18 to 69 who are receiving secondary mental health services and who are on the Care Programme Approach (CPA).</p> <p>The measure is focused on ‘paid’ employment, to be clear that voluntary work is to be excluded for the purposes of this measure. Employment status is recorded using the following categories:</p> <ul style="list-style-type: none"> <li>01 Employed</li> <li>02 Unemployed and Seeking Work</li> <li>03 Students who are undertaking full (at least 16 hours per week) or part-time (less than 16 hours per week) education or training and who are not working or actively seeking work</li> <li>04 Long-term sick or disabled, those who are receiving Incapacity Benefit, Income Support or both; or Employment and Support Allowance</li> <li>05 Homemaker looking after the family or home and who are not working or actively seeking work</li> </ul>

<sup>19</sup> Waddell, G. & Burton, A. (2006). *Is Work Good for your Health and Well-being?* London: TSO

	<p>06 Not receiving benefits and who are not working or actively seeking work          07 Unpaid voluntary work who are not working or actively seeking work          08 Retired          ZZ Not Stated (person asked but declined to provide a response)</p> <p>Further development work will explore those on the CPA who it may be appropriate to exclude from the measure – for example those who are detained under the Mental Health Act for a significant portion of the year.</p> <p>From 2012-13 the data source (Mental health minimum data set) has moved to a monthly collection. Therefore the definition of the measure has been amended slightly in consultation with stakeholders. Details are given below.</p> <p><i>Interpretation</i></p> <p>Interpretation of the measure should take into account the above point regarding scope, and the likelihood that some people in contact with secondary mental health services are being supported in paid employment by the council, but are not captured within the current definition. Additional local data may be available to support analysis.</p>
<p><b>Alignment</b></p>	<p>This measure is complementary with Measure 1.8 (employment for those with a long-term health condition including those with a learning difficulty/disability or mental illness) in the Public Health Outcomes Framework (<a href="http://www.dh.gov.uk/prod_consum_dh/groups/dh_digitalassets/@dh/@en/documents/digitalasset/dh_132558.pdf">http://www.dh.gov.uk/prod_consum_dh/groups/dh_digitalassets/@dh/@en/documents/digitalasset/dh_132558.pdf</a> page 14) and Measure 2.5 (employment of people with mental illness) in the NHS Outcomes Framework (<a href="https://www.wp.dh.gov.uk/publications/files/2012/11/121109-Technical-Appendix.pdf">https://www.wp.dh.gov.uk/publications/files/2012/11/121109-Technical-Appendix.pdf</a> page 40).</p> <p>Although the Public Health Outcomes Framework and the ASCOF both include measures connected with employment for people with a learning disability and people with mental health problems, the Public Health Outcomes Framework measures the gap between the employment rate for those groups and the overall employment rate. This reflects the approach taken in the NHS Outcomes Framework for a complementary measure on employment of people with long term conditions. Although aligning the ASCOF with the other two frameworks was considered, in developing the framework with local government, it was agreed that this would not support local interpretation and benchmarking, and so the ASCOF will retain the current approach, with changes made to reflect the updated NMDS-MH dataset. Furthermore, although the Public Health Outcomes Framework uses the same data sources for rates of employment for these groups as the ASCOF, the NHSOF uses the Labour Force Survey. This source cannot be used for the ASCOF because it does not provide robust results at the local authority level.</p>
<p><b>Formula</b></p>	<p><math>\left(\frac{X}{Y}\right) * 100</math> is calculated each month. The twelve monthly figures are summed and then divided by 12 to derive an average.</p> <p>Where:</p> <p><b>X:</b> Number of working age adults (18-69 years) who are receiving secondary mental health services and who are on the Care Programme Approach recorded as being in employment. The most recent record of employment status for the person during the previous twelve months is used.  <i>Source: Mental Health Minimum Data Set v4.1</i></p> <p><b>Y:</b> Number of working age adults (18-69 years) who have received secondary mental health services and who were on the Care Programme Approach at the end of the month  <i>Source: Mental Health Minimum Data Set v4.1</i>          Where X and Y are measured at the end of each month</p>
<p><b>Worked example</b></p>	<p>In January, adults receiving secondary mental health services = 964</p> <p>In January, adults receiving secondary mental health services in paid employment = 196</p>

	The measure value for January = $(196/964) \times 100 = 20.3\%$		
	This is calculated for each of the twelve months, then the monthly figures are averaged.		
Disaggregation available	Equalities: Gender		
	Client groups: Mental health (18-69)		
Frequency of collection	Annual report based on monthly collection	Data source	Mental Health Minimum Data Set v4.1 (MHMDS)
Return format	Percentage	Decimal places	One
Longer-term development options	<p>The 2012/13 ASCOF included a placeholder on employment for all people with a disability, which would replace the existing measures for adults in contact with secondary mental health services and for adults with a learning disability. The placeholder was to combine all adults into a single measure for employment status. This placeholder has been removed for the 2013/14 ASCOF, as, in consultation with local authorities, it was agreed that it would be too onerous to collect additional information on people with physical disabilities in employment in order to populate the proposed measure.</p> <p>A time-series will be maintained for this measure. Further detail will be provided in the Handbook of Definitions for 2014/15.</p>		
Further guidance	<p>Guidance for 2012/13 onwards can be found via the social care collection page at <a href="http://www.ic.nhs.uk/socialcare/collections">http://www.ic.nhs.uk/socialcare/collections</a> by clicking on the year.</p> <p>Guidance and information relating to the Minimum Mental Health Dataset can be found at <a href="http://www.ic.nhs.uk/mentalhealth">http://www.ic.nhs.uk/mentalhealth</a>.</p> <p>The Community and Mental Health team at the Health and Social Care Information Centre now publish data from the Mental Health Minimum Data Set on a monthly basis. The most recently published data are for June 2013 and are available at: <a href="http://www.hscic.gov.uk/catalogue/PUB11729">http://www.hscic.gov.uk/catalogue/PUB11729</a>. By mapping this CCG level data to CASSR, local authorities will be able to monitor their outcomes for 1F and 1H throughout the year with reasonable accuracy. Mapping data are available at: <a href="http://www.england.nhs.uk/wp-content/uploads/2013/03/ccg-Isa.csv">http://www.england.nhs.uk/wp-content/uploads/2013/03/ccg-Isa.csv</a>. Please send any queries about the Mental Health Minimum Data Set publication to <a href="mailto:MHMDS@hscic.gov.uk">MHMDS@hscic.gov.uk</a>.</p>		

### (1G) Proportion of adults with a learning disability who live in their own home or with their family

Domain / Outcome	<p>1. Enhancing quality of life for people with care and support needs.</p> <p><i>People are able to find employment when they want, maintain a family and social life and contribute to community life and avoid loneliness or isolation.</i></p>
Rationale	<p>The measure is intended to improve outcomes for adults with a learning disability by demonstrating the proportion in stable and appropriate accommodation. The nature of accommodation for people with a learning disability has a strong impact on their safety and overall quality of life and the risk of social exclusion.</p>
Definition / Interpretation	<p>The measure shows the proportion of all adults with a learning disability who are known to the council, who are recorded as living in their own home or with their family. The information would have to be captured or confirmed within the reporting period 1 April 2013 to 31 March 2014.</p> <p>The definition of individuals '<b>known to the council</b>' is currently restricted to those adults with a learning disability (with a primary client group of LD) who have been assessed or reviewed by the council during the year (irrespective of whether or not they receive a service) or who</p>

	<p>should have been reviewed but were not.</p> <p><b>‘Living on their own or with their family’</b> is intended to describe arrangements where the individual has security of tenure in their usual accommodation, for instance because they own the residence or are part of a household whose head holds such security. This has the same definition as ‘living independently, with or without support’ in Measure 1H (see below), however different wording is used to capture the emphasis on avoiding residential care homes.</p> <p>Situations included within the scope of ‘living on their own or with their family’:</p> <ul style="list-style-type: none"> <li>▪ Owner occupier or shared ownership scheme;</li> <li>▪ Tenant (including local authority, arm’s-length management organisation, registered social landlord, housing association);</li> <li>▪ Tenant – private landlord</li> <li>▪ Settled mainstream housing with family/friends (including flat-sharing);</li> <li>▪ Supported accommodation/supported lodgings/supported group home (i.e. accommodation supported by staff or resident caretaker);</li> <li>▪ Adult Placement Scheme</li> <li>▪ Approved premises for offenders released from prison or under probation supervision (e.g. probation hostel);</li> <li>▪ Sheltered housing/extra care housing/other sheltered housing; and,</li> <li>▪ Mobile accommodation for Gypsy/Roma and Traveller communities.</li> </ul> <p>The following circumstances are <u>not</u> included within the scope of ‘living on their own or with their family’:</p> <ul style="list-style-type: none"> <li>▪ Rough sleeper/squatting;</li> <li>▪ Night shelter/emergency hostel/direct access hostel (temporary accommodation accepting self-referrals);</li> <li>▪ Refuge;</li> <li>▪ Placed in temporary accommodation by council (including homelessness resettlement);</li> <li>▪ Staying with family/friends as a short-term guest;</li> <li>▪ Acute/long-stay healthcare residential facility or hospital (e.g. NHS independent general hospital/clinic, long-stay hospital, specialist rehabilitation/recovery hospital);</li> <li>▪ Registered care home</li> <li>▪ Registered nursing home;</li> <li>▪ Prison/Young Offenders Institution/detention centre; and,</li> <li>▪ Other temporary accommodation.</li> </ul>
<b>Alignment</b>	<p>This measure is shared with Measure 1.6i (people with a learning disability in settled accommodation) in the Public Health Outcomes Framework  <a href="http://www.dh.gov.uk/prod_consum_dh/groups/dh_digitalassets/@dh/@en/documents/digitalasset/dh_132558.pdf">http://www.dh.gov.uk/prod_consum_dh/groups/dh_digitalassets/@dh/@en/documents/digitalasset/dh_132558.pdf</a> page 12).</p>
<b>Risk adjustment</b>	<p>It is not clear whether any factors should be considered for risk adjustment for this measure.</p>
<b>Formula</b>	$\left(\frac{X}{Y}\right) * 100$ <p>Where:</p> <p><b>X:</b> All people within the denominator who are living in their own home or with their family. The numerator should include those living in their own home or with their family irrespective of whether they have had a review during the year, but the information would have to be captured within the current financial year.  <i>Source: Table L2, ASC-CAR<sup>20</sup></i></p> <p><b>Y:</b> Number of working-age (aged 18-64) learning disabled clients known to councils. This</p>

<sup>20</sup> (Number of working age learning disabled clients known to CASSRs during year to 31<sup>st</sup> March, by gender and by accommodation status at the time of their latest assessment or review) sum of rows 12 to 20 column 3

	<p>includes:</p> <ul style="list-style-type: none"> <li>▪ those who are assessed or reviewed in the financial year and have received a service;</li> <li>▪ those who are assessed or reviewed in the financial year and have not received a service; and,</li> <li>▪ those who should have been reviewed in the financial year but were not.</li> </ul> <p>Source: Table L2, ASC-CAR<sup>21</sup></p> <p>In other words, this is a count of eligible adults with a learning disability, without duplication (aged 18-64) who are assessed, or reviewed (regardless of whether they have received a service), or should have been reviewed during the year; i.e. they are recorded within an A or P table within the RAP return.</p>		
<b>Worked example</b>	<p>Adults with a learning disability known to CASSRs = 722</p> <p>Of those adults with a learning disability known to CASSRs, those who are recorded as living in their own home or with their family within the current financial year = 455</p> <p>The measure value = <math>(455/722) \times 100 = 63.0\%</math></p>		
<b>Disaggregation available</b>	<b>Equalities:</b> Gender		
	<b>Client groups:</b> Learning disability (18-64)		
<b>Frequency of collection</b>	Annual	<b>Data source</b>	Adult Social Care Combined Activity Return (ASC-CAR) – table, page and cell references given here are based on the 2012/13 proforma.
<b>Return format</b>	Percentage	<b>Decimal places</b>	One
<b>Further guidance</b>	Guidance for 2012/13 onwards can be found via the social care collection page at <a href="http://www.ic.nhs.uk/socialcare/collections">http://www.ic.nhs.uk/socialcare/collections</a> by clicking on the year.		

### (1H) Proportion of adults in contact with secondary mental health services living independently, with or without support

<b>Domain / Outcome</b>	<p>1. Enhancing quality of life for people with care and support needs.</p> <p><i>People are able to find employment when they want, maintain a family and social life and contribute to community life and avoid loneliness or isolation.</i></p>
<b>Rationale</b>	<p>The measure is intended to improve outcomes for adults with mental health problems by demonstrating the proportion in stable and appropriate accommodation. This is closely linked to improving their safety and reducing their risk of social exclusion.</p>
<b>Definition / Interpretation</b>	<p>The measure shows the percentage of adults receiving secondary mental health services living independently at the time of their most recent assessment, formal review or other multi-disciplinary care planning meeting.</p> <p>Adults <b>‘in contact with secondary mental health services’</b> is defined as those aged 18 to 69 who are receiving secondary mental health services and who are on the Care Programme Approach (CPA).</p> <p><b>‘Living independently, with or without support’</b> refers to accommodation arrangements where the occupier has security of tenure or appropriate stability of residence in their <i>usual</i></p>

<sup>21</sup> (Number of working age learning disabled clients known to CASSRs during year to 31<sup>st</sup> March, by gender and by accommodation status at the time of their latest assessment or review) row 22 column 3

	<p>accommodation in the medium-to-long-term, or is part of a household whose head holds such security of tenure/residence. These accommodation arrangements are recorded as settled accommodation in the Mental Health Minimum Data Set. This has the same definition as 'living on their own or with their family' in Measure 1G (see above); however different wording is used to capture the emphasis on general independence.</p> <p>Accommodation arrangements that are precarious, or where the person has no or low security of tenure/residence in their usual accommodation and so may be required to leave at very short notice, are excluded from the definition of 'living independently, with or without support'. These accommodation arrangements are recorded as non-settled accommodation in the MHMDS.</p> <p>Accommodation types that represent settled or non-settled accommodation for the purpose of this measure are presented in Appendix 5 to this document.</p> <p>Further development work will explore those on the CPA who it may be appropriate to exclude from the measure – for example those who are detained under the Mental Health Act for a significant portion of the year.</p> <p>From 2012-13 the data source (Mental health minimum data set) has moved to a monthly collection. Therefore the definition of the measure has been amended slightly in consultation with stakeholders. Details are given below.</p> <p><i>Interpretation</i></p> <p>Interpretation of the measure should take into account the point above regarding scope, and the likelihood that some people in contact with mental health services are being supported in accommodation by the council, but are not captured within the current definition because they are not on the CPA. Additional local data may be available to support analysis.</p>
<b>Alignment</b>	<p>This measure is shared with Measure 1.6ii (adults receiving mental health services in settled accommodation) in the Public Health Outcomes Framework (<a href="http://www.dh.gov.uk/prod_consum_dh/groups/dh_digitalassets/@dh/@en/documents/digitalasset/dh_132558.pdf">http://www.dh.gov.uk/prod_consum_dh/groups/dh_digitalassets/@dh/@en/documents/digitalasset/dh_132558.pdf</a> page 12).</p>
<b>Risk adjustment</b>	<p>It is not clear whether any factors should be considered for risk adjustment for this measure.</p>
<b>Formula</b>	<p><math>\left(\frac{X}{Y}\right) * 100</math> is calculated each month. The twelve monthly figures are summed and then divided by 12 to derive an average.</p> <p>Where:</p> <p><b>X:</b> Number of adults aged 18-69 who are receiving secondary mental health services on the Care Programme Approach recorded as living independently (with or without support). The most recent record of whether or not the person is in settled accommodation during the previous twelve months is used. <i>Source: Mental Health Minimum Data Set v4.1</i></p> <p><b>Y:</b> Number of adults aged 18-69 who have received secondary mental health services and who were on the Care Programme Approach at the end of the month. <i>Source: Mental Health Minimum Data Set v4.1</i></p> <p>Where X and Y are measured at the end of each month,</p>
<b>Worked example</b>	<p>In January, adults receiving secondary mental health services = 964</p> <p>In January, adults receiving secondary mental health services living independently = 655</p> <p>The measure value in January = <math>(655/964) \times 100 = 67.9\%</math></p> <p>This is calculated for each of the twelve months, then the monthly figures are averaged.</p>

<b>Disaggregation available</b>	<b>Equalities:</b> Gender		
	<b>Client groups:</b> Mental health (18-69)		
<b>Frequency of collection</b>	Annual report based on monthly collections	<b>Data source</b>	Mental Health Minimum Data Set v4.1 (MHMDS)
<b>Return format</b>	Percentage	<b>Decimal places</b>	One
<b>Longer-term development</b>	<p>The 2012/13 ASCOF included a placeholder on employment for all people with a disability, which would replace the existing measures for adults in contact with secondary mental health services and for adults with a learning disability. The placeholder was to combine all adults into a single measure for employment status. This placeholder has been removed for the 2013/14 ASCOF, as, in consultation with local authorities, it was agreed that it would be too onerous to collect additional information on people with physical disabilities in employment in order to populate the proposed measure.</p> <p>A time-series will be maintained for this measure. Further detail will be provided in the Handbook of Definitions for 2014/15.</p>		
<b>Further guidance</b>	<p>Guidance for 2012/13 onwards can be found via the social care collection page at <a href="http://www.ic.nhs.uk/socialcare/collections">http://www.ic.nhs.uk/socialcare/collections</a> by clicking on the year.</p> <p>Guidance and information relating to the Minimum Metal Health Dataset can be found at <a href="http://www.ic.nhs.uk/mentalhealth">http://www.ic.nhs.uk/mentalhealth</a>.</p> <p>The Community and Mental Health team at the Health and Social Care Information Centre now publish data from the Mental Health Minimum Data Set on a monthly basis. The most recently published data are for June 2013 and are available at: <a href="http://www.hscic.gov.uk/catalogue/PUB11729">http://www.hscic.gov.uk/catalogue/PUB11729</a>. By mapping this CCG level data to CASSR, local authorities will be able to monitor their outcomes for 1F and 1H throughout the year with reasonable accuracy. Mapping data are available at: <a href="http://www.england.nhs.uk/wp-content/uploads/2013/03/ccg-isoa.csv">http://www.england.nhs.uk/wp-content/uploads/2013/03/ccg-isoa.csv</a>. Please send any queries about the Mental Health Minimum Data Set publication to <a href="mailto:MHMDS@hscic.gov.uk">MHMDS@hscic.gov.uk</a>.</p>		

#### New measure for 2013/14

#### (1I) Proportion of people who use services and their carers, who reported that they had as much social contact as they would like.

<b>Domain / Outcome</b>	1. Enhancing quality of life for people with care and support needs.
<b>Rationale</b>	There is a clear link between loneliness and poor mental and physical health. A key element of the Government's vision for social care is to tackle loneliness and social isolation, supporting people to remain connected to their communities and to develop and maintain connections to their friends and family. This measure will draw on self-reported levels of social contact as an indicator of social isolation for both users of social care and carers.
<b>Definition / interpretation</b>	<p>The relevant question drawn from the Adult Social Care Survey is question 8a – "Thinking about how much contact you've had with people you like, which of the following statements best describes your social situation?"</p> <ul style="list-style-type: none"> <li>- I have as much social contact as I want with people I like</li> <li>- I have adequate social contact with people</li> <li>- I have some social contact with people, but not enough</li> <li>- I have little social contact with people and feel socially isolated</li> </ul> <p>The relevant question drawn from the Carers Survey is question 11 – "By thinking about social contact you've had with people you like, which statement best describes your present social situation?"</p> <ul style="list-style-type: none"> <li>- I have as much social contact as I want with people I like</li> </ul>

	<ul style="list-style-type: none"> <li>- I have some social contact with people but not enough</li> <li>- I have little social contact and I feel socially isolated</li> </ul> <p>The measure is defined by determining the percentage of users responding “I have as much contact as I want with people I like” and carers choosing “I have as much contact as I want with people I like”. Measures for users and carers will be presented separately. These responses have been chosen to focus the measure on individuals achieving the best outcomes, to allow for better use in benchmarking.</p> <p><b>Interpretation</b></p> <p>The measure gives an overall indication of the reported outcomes for individuals – it does not at present identify the specific contribution of councils’ adult social care towards the outcome (see longer term development below).</p>
<b>Alignment</b>	<p>This measure is shared with Measure 1.18 (social connectedness placeholder) in the Public Health Outcomes Framework  <a href="http://www.dh.gov.uk/prod_consum_dh/groups/dh_digitalassets/@dh/@en/documents/digitalasset/dh_132558.pdf">http://www.dh.gov.uk/prod_consum_dh/groups/dh_digitalassets/@dh/@en/documents/digitalasset/dh_132558.pdf</a> page 25)</p>
<b>Risk adjustment</b>	<p>There are a range of factors which are likely to have an impact on this measure – for example the severity of needs of users or the amount of care provided by carers.</p>
<b>Formula</b>	$\left(\frac{X}{Y}\right) * 100$ <p>Where, for <b>11 part 1</b> (users):</p> <p><b>X:</b> In response to Question 8a of the ASCS, those individuals who selected the response “I have as much social contact as I want with people I like”.</p> <p>Those respondents who were sent the version of the questionnaire for people with learning disabilities will be treated in the same way, as this questionnaire has been designed to be equivalent to the non-learning disabilities version.</p> <p><b>Y:</b> All those that responded to the question.</p> <p>For both the numerator (X) and denominator (Y), weighted data should be used to calculate the measure. The data from the survey will be weighted by the Health and Social Care Information Centre to take account of the stratified sampling technique that has been used when conducting the survey. The weights are automatically calculated within the survey data return along with the ASCOF outcome measures. Further details of how to use the weights when analysing the survey data are available in Appendix I of the guidance for the 2012-13 Adult Social Care Survey.</p> <p>For <b>11 part 2</b> (carers)</p> <p><b>X:</b> The sum of all those who in response to question 11 of the Carers Survey, selected the response “I have as much social contact as I want with people I like”.</p> <p>In years where the Carers Survey has not been completed (since this is initially a biennial collection), the most recent value for the carers element should be carried over and counted in the second of these parts in the numerator. In these years, only the changes in the service user element (drawn from the ASCS) will be identifiable.</p> <p><b>Y:</b> The sum of all those that responded to the above question of the Carers Survey.</p> <p><b>Exclusions</b></p> <p>People who select the response “I’ve never tried to find information or advice” for the ASCS or “I have not tried to find information or advice in the last 12 months” for the Carers Survey will</p>



	not be counted in either the numerator or the denominator.		
<b>Worked example</b>	<p><b>11 part 1 – users</b></p> <p>The number of users who said “I have as much social contact as I want with people I like” was 242. In total the number of users who responded to the question was 548. (Data weighted to reflect the stratified sampling technique that has been used when conducting the survey). The indicator value is <math>[(X/Y)*100] = 44.2\%</math></p> <p><b>11 part 2 – carers</b></p> <p>The number of carers who said “I have as much social contact as I want” was 197. In total the number of carers who responded to the question was 420. (Data weighted to reflect the stratified sampling technique that has been used when conducting the survey). The indicator value is <math>[(X/Y)*100] = 46.9\%</math></p>		
<b>Disaggregation available</b>	<b>Equalities:</b> Age, Gender, Ethnicity**, Religion*, Sexual orientation*		
	<b>Client groups:</b> Physical Disability (all ages)** , Learning Disability (all ages)** , Mental Health (all ages), Substance Misuse (all ages)** , Vulnerable People (all ages)**		
<b>Frequency of collection</b>	Annual for social care users  Biennial for Carers (Carers Survey to be next conducted in 2014/15)	<b>Data source</b>	<i>Adult Social Care Survey</i>  <i>Carers Survey</i>
<b>Return format</b>	Percentage	<b>Decimal places</b>	One
<b>Longer-term development options</b>	<p>Initially, this measure will focus on social care users and carers, rather than the broader population. However, the problems of loneliness and social isolation are not limited to these groups, and all parts of the health and care system have a role to play in preventing and reducing social isolation and loneliness in the broader population. Work will continue to pursue the development of a population based measure of loneliness, with a view to including this in both the ASCOF and the Public Health Outcomes Framework in future years.</p> <p>Develop a ‘value-added’ measure which quantifies the contribution of social services to quality of life as described for Measure 1A.</p>		
<b>Further guidance</b>	Guidance for 2012/13 onwards can be found via the user survey guidance page at <a href="http://www.ic.nhs.uk/socialcare/usersurveys">http://www.ic.nhs.uk/socialcare/usersurveys</a>		

\* Although the underlying survey results will in theory be disaggregated by religion and sexual orientation, in practice there are likely to be significant gaps in the data on these characteristics at least in the short/medium term. This reflects the content of records held locally by councils.

\*\* This information is not published as part of the adult social care outcomes returns; however it is part of the publication of the data source.

## Domain 2 – Delaying and reducing the need for care and support

<b>(2A) Permanent admissions to residential and nursing care homes, per 100,000 population</b>	
<b>Domain / Outcome</b>	2. Delaying and reducing the need for care and support. ( <i>Overarching measure</i> )
<b>Rationale</b>	Avoiding permanent placements in residential and nursing care homes is a good measure of delaying dependency, and the inclusion of this measure in the framework supports local health and social care services to work together to reduce avoidable admissions. Research suggests that, where possible, people prefer to stay in their own home rather than move into residential care.
<b>Definition / interpretation</b>	<p>This is a two part-measure reflecting the number of admissions of younger adults (part 1) and older people (part 2) to residential and nursing care homes relative to the population size of each group. The measure compares council records with ONS population estimates.</p> <p>People counted as a permanent admission should include:</p> <ul style="list-style-type: none"> <li>▪ Residents where the local authority makes any contribution to the costs of care, no matter how trivial the amount and irrespective of how the balance of these costs are met;</li> <li>▪ Supported residents in: <ul style="list-style-type: none"> <li>○ Local authority-staffed care homes for residential care;</li> <li>○ Independent sector care homes for residential care;</li> <li>○ Registered care homes for nursing care; and,</li> <li>○ Residential or nursing care which is of a permanent nature and where the intention is that the spell of care should not be ended by a set date.</li> </ul> </li> </ul> <p>For people classified as permanent residents, the care home would be regarded as their normal place of residence.</p> <p>Where a person who is normally resident in a care home is temporarily absent at 31 March 2013 (e.g. through temporary hospitalisation) and the local authority is still providing financial support for that placement, the person should be included in the numerator.</p> <p>Trial periods in residential or nursing care homes where the intention is that the stay will become permanent should be counted as permanent.</p> <p>Whether a resident or admission is counted as permanent or temporary depends on the intention of the authority making the placement.</p> <p><b>Interpretation</b></p> <p>Analysis shows that older people have a higher rate of permanent admissions than younger adults. Using a two-part measure means that we can separate age as a factor in the level of admissions and focus on the contribution of services to reducing admissions. It will also help highlight, both nationally and locally, the separate issues that relate to the rates of permanent admissions for younger adults and for older people.</p>
<b>Alignment</b>	ASCOF measure only

<b>Risk adjustment</b>	Analysis identified age as a factor that influenced the rate of admissions. Instead of applying risk adjustment, the measure has been expressed separately for those aged 18-64 years, and those aged 65 years and over. There were no other influencing factors identified from the data available.
<b>Formula</b>	$\left(\frac{X}{Y}\right) \times 100,000$ <p>Where:</p> <p><b>for 2A part 1 (younger adults):</b></p> <p><b>X:</b> Number of council-supported permanent admissions of younger adults (aged 18-64) to residential and nursing care during the year (excluding transfers between residential and nursing care).  <i>Source: Table S3, ASC-CAR<sup>22</sup></i></p> <p><b>Y:</b> Size of younger adult population (aged 18-64) in area (ONS mid year population estimates).  <i>Source: Office of National Statistics</i></p> <p><b>For 2A part 2 (older people):</b></p> <p><b>X:</b> Number of council-supported permanent admissions of older people (aged 65 and over) to residential and nursing care during the year (excluding transfers between residential and nursing care).  <i>Source: Table S3, ASC-CAR<sup>23</sup></i></p> <p><b>Y:</b> Size of older people population (aged 65 and over) in area (ONS mid year population estimates).  <i>Source: Office of National Statistics</i></p> <p><b>Exclusions</b>  People funding their own residence in a care home with no support from the council are excluded.</p>
<b>Worked example</b>	<p><b>2A Part 1 (younger adults)</b>  Suppose the number of permanent admissions to residential or nursing care for younger adults (aged 18-64) during the year was 26. Suppose the population of younger adults in the area is 153,471</p> <p>The measure value is <math>[(26/153,471) * 100,000] = 16.9</math></p> <p><b>2A Part 2 (older people)</b>  Suppose the number of permanent admissions to residential or nursing care for older people (aged 65 and over) during the year was 312. Suppose the population of older people in the area is 43,384</p> <p>The measure value is <math>[(312/43,384) * 100,000] = 719.2</math></p>
<b>Disaggregation Available</b>	<b>Equalities:</b> Age (18-64, 65 and over)

<sup>22</sup> (Number of LA supported permanent admissions to residential and nursing care during 1 April to 31 March (excluding admissions to group homes) by type of residence, primary client type and age group), page 1, row 14, sum of columns 1 to 3

<sup>23</sup> (Number of LA supported permanent admissions to residential and nursing care during 1 April to 31 March (excluding admissions to group homes) by type of residence, primary client type and age group), page 1, row 15, sum of columns 1 to 3

	<b>Client groups:</b> Physical Disabilities (18-64), Mental Health (18-64), Learning Disabilities (18-64), Substance misuse and other vulnerable people (as one group)**, Not allocated by client group**		
<b>Frequency of collection</b>	Annual	<b>Data source</b>	<i>Adult Social Care Combined Activity Return (ASC-CAR) – table, page and cell references given here are based on the 2012/13. Office of National Statistics</i>
<b>Return format</b>	Rate per 100,000 population	<b>Decimal places</b>	One
<b>Longer-term development options</b>	The data collections required for the current measure have been reviewed as part of the 'zero-based review'. Work is ongoing to understand whether and how the new ZBR collections, which are yet to be finalised, will impact on this measure. If the new collections are implemented for 2014-15 this will be made clear in the handbook for 2014-15 measures.		
<b>Further guidance</b>	Guidance for 2012/13 onwards can be found via the social care collection page at <a href="http://www.ic.nhs.uk/socialcare/usersurveys">http://www.ic.nhs.uk/socialcare/usersurveys</a> by clicking on the year.		

### **(2B) Proportion of older people (65 and over) who were still at home 91 days after discharge from hospital into reablement/rehabilitation services**

<b>Domain / Outcome</b>	2. Delaying and reducing the need for care and support. <i>When people develop care needs, the support they receive takes place in the most appropriate setting and enables them to regain their independence.</i>
<b>Rationale</b>	<p>There is strong evidence that reablement services lead to improved outcomes and value for money across the health and social care sectors. Reablement seeks to support people and maximise their level of independence, in order to minimise their need to ongoing support and dependence on public services.</p> <p>This measures the benefit to individuals from reablement, intermediate care and rehabilitation following a hospital episode, by determining whether an individual remains living at home 91 days following discharge – the key outcome for many people using reablement services. It captures the joint work of social services and health staff and services commissioned by joint teams, as well as adult social care reablement.</p>
<b>Definition / interpretation</b>	<p>This is a two-part measure which reflects both the effectiveness of reablement services (part 1), and the coverage of the service (part 2).</p> <p><b>2B Part 1:</b></p> <p>The proportion of older people aged 65 and over discharged from hospital to their own home or to a residential or nursing care home or extra care housing for rehabilitation, with a clear intention that they will move on/back to their own home (including a place in extra care housing or an adult placement scheme setting), who are at home or in extra care housing or an adult placement scheme setting 91 days after the date of their discharge from hospital.</p> <p>Those who are in hospital or in a registered care home (other than for a brief episode of respite care from which they are expected to return home) at the three month date and those who have died within the three months are not reported in the numerator.</p> <p>The collection of the denominator will be between 1 October 2013 and 31 December 2013, with a 91-day follow-up for each case included in the denominator to populate the numerator i.e. the numerator will be collected from 1 January 2014 to 31 March 2014.</p> <p><b>2B Part 2:</b></p>

\*\* This information is not published as part of the adult social care outcomes returns; however it is part of the publication of the data source.

	<p>The proportion of older people aged 65 and over offered reablement services following discharge from hospital.</p> <p>This measure will take the denominator from part 1 as its numerator (the number of older people offered reablement services). The denominator will be the total number of older people discharged from hospitals based on Hospital Episode Statistics (HES).</p> <p>The collection of the numerator and the denominator will be from 1 October 2013 to 31 December 2013.</p> <p><b>Interpretation</b></p> <p>The rationale for a two-part measure is to capture the volume of reablement offered as well as the success of the reablement service offered. This will prevent areas scoring well on the measure while offering reablement services to only a very small number of people.</p> <p>The measure includes social care-only placements. Therefore, those that were assessed just on social care needs would now be included in the data collection.</p> <p>In the future it may also be possible to expand the measure to include individuals assessed only on health needs, on the basis that this is a measure of joint working and is due to be replicated in the NHS Outcomes Framework once it comes into use in 2013/14. In addition, even in circumstances where there has been an assessment conducted by the NHS not including social care needs, social care may still be involved in delivering the service to the individual.</p>
<b>Alignment</b>	<p>This measure is shared with Measure 3.6i (the proportion of older people aged 65 and over who were still at home 91 days after discharge into rehabilitation) in the NHS Outcomes Framework (<a href="https://www.wp.dh.gov.uk/publications/files/2012/11/121109-Technical-Appendix.pdf">https://www.wp.dh.gov.uk/publications/files/2012/11/121109-Technical-Appendix.pdf</a> page 61).</p>
<b>Risk adjustment</b>	<p>None.</p>
<b>Formula</b>	$\left(\frac{X}{Y}\right) \times 100$ <p>Where, for <b>2B part 1</b> (proportion of successful reablement):</p> <p><b>X:</b> Number of older people discharged from acute or community hospitals to their own home or to a residential or nursing care home or extra care housing for rehabilitation, with a clear intention that they will move on/back to their own home (including a place in extra care housing or an adult placement scheme setting), who are at home or in extra care housing or an adult placement scheme setting 91 days after the date of their discharge from hospital. This should only include the outcome for those cases referred to in the denominator.</p> <p><i>Source: Table I1, ASC-CAR<sup>24</sup></i></p> <p><b>Y:</b> Number of older people discharged from acute or community hospitals from hospital to their own home or to a residential or nursing care home or extra care housing for rehabilitation, with a clear intention that they will move on/back to their own home (including a place in extra care housing or an adult placement scheme setting). <i>Source: Table I1, ASC-CAR<sup>25</sup></i></p>

<sup>24</sup> (Number of clients aged 65 and over achieving independence through rehabilitation during 1 October to 31 December by age group and gender), row 1 column 9

<sup>25</sup> (Number of clients aged 65 and over discharged to rehabilitation during 1 October to 31 December by age group and gender), row 2 column 9

	<p>For <b>2B part 2</b> (coverage of reablement services):</p> <p><b>X:</b> Number of older people discharged from acute or community hospitals from hospital to their own home or to a residential or nursing care home or extra care housing for rehabilitation, with a clear intention that they will move on/back to their own home (including a place in extra care housing or an adult placement scheme setting). <i>Source: Table 11, ASC-CAR<sup>2</sup></i></p> <p><b>Y:</b> Total number of people, aged 65 and over, discharged alive from hospitals in England between 1 October 2013 and 31 December 2013. This includes all specialities and zero-length stays. Data for geographical areas is based on usual residence of patient. <i>Source: Hospital Episode Statistics</i></p> <p>HES data for the full calendar year of 2012 will be provided to local authorities via the HSCIC website. The data will be broken down by month as well as by local authority, and could be used as a proxy for 2013 data to estimate monthly measure values.</p>		
<p><b>Worked example</b></p>	<p><b>2B Part 1</b></p> <p>Suppose the number of people aged 65+ on discharge and who were discharged and benefited from intermediate care/ rehabilitation still living at home 3 months after discharge = 217.</p> <p>And if the number of people discharged from hospital aged 65+ and entering into joint 'intermediate care' or a 'rehabilitation service' = 306.</p> <p>Therefore the percentage achieving independence = <math>(217 / 306) \times 100 = 70.9\%</math></p> <p><b>2B Part 2</b></p> <p>If the number of people discharged from hospital aged 65+ and entering into joint 'intermediate care' or a 'rehabilitation service' = 306 (using same figure as above).</p> <p>And if the total number of people aged 65+ discharged from hospital = 6,857.</p> <p>Then, the proportion offered reablement services = <math>(306/6,857) \times 100 = 4.5\%</math></p>		
<p><b>Disaggregation available</b></p>	<p><b>Equalities:</b> Age (65-74, 75-84, 85+), Gender</p> <p><b>Client groups:</b> Older people (65+)</p>		
<p><b>Frequency of collection</b></p>	<p>Annual</p>	<p><b>Data source</b></p>	<p><i>Adult Social Care Combined Activity Return (ASC-CAR) – table, page and cell references given here are based on the 2012/13 proforma Hospital Episode Statistics</i></p>
<p><b>Return format</b></p>	<p>Percentage</p>	<p><b>Decimal places</b></p>	<p>One</p>
<p><b>Longer-term development options</b></p>	<p>Over time, we will aim to measure the success of all those offered a reablement service, rather than restricting measurement to those discharged from hospital only.</p>		
<p><b>Further guidance</b></p>	<p>Guidance for 2012/13 onwards can be found via the social care collection page at <a href="http://www.ic.nhs.uk/socialcare/collections">http://www.ic.nhs.uk/socialcare/collections</a> by clicking on the year.</p> <p>Guidance for HES data can be found at: <a href="http://www.hesonline.nhs.uk">http://www.hesonline.nhs.uk</a></p>		

**New measure for 2014/15**

**(2D) The outcome of short-term services: sequel to service**

<p><b>Domain / Outcome</b></p>	<p>2. Delaying and reducing the need for care and support.  <i>Earlier diagnosis, intervention and reablement means that people and their carers are less dependent on intensive services.</i></p>
<p><b>Rationale</b></p>	<p>This measure will reflect the proportion of those people who received short-term services during the year, where no further request was made for ongoing support. Since the aim of short-term services is to reable people and promote their independence, this measure will provide evidence of a good outcome in delaying dependency or supporting recovery – short-term support that results in no further need for services.</p>
<p><b>Definition / interpretation</b></p>	<p>In this context, short-term support is defined as ‘short-term support which is designed to maximise independence’, and therefore will exclude carer contingency and emergency support. This prevents the inclusion of short-term support services which are not reablement services.</p> <p>This measure will be included when the proposals of the zero-based review are fully implemented.</p> <p>Once implemented, this measure should be viewed in the context of a second new measure in this domain, 2E – the effectiveness of reablement services - to understand whether there are any unintended consequences of the decision to provide no further services. Measure 2E is still to be developed.</p> <p>Further work is needed to develop the detailed definition for this measure which is subject to the development of the new data collections. However the definition is likely to be along the lines of the following:</p> <p>Percentage of those that received a short term service during the year where the sequel was no ongoing support = numerator divided by denominator</p> <p><b>Where:</b></p> <p>Numerator: Those with a sequel to short-term service where no services were provided (none offered/individual not eligible). Those where no services were provided but where the individual went on to purchase services because they fund their own care should be subtracted from this total.</p> <p>Denominator: Number of people who had a short-term support to maximise independence. Those with a sequel of either early cessation due to a life event, or those with no services provided (individual declined), should be subtracted from this total.</p> <p>The numerator and denominator would be summed for new and existing clients.</p> <p><b>Comment</b></p> <p>The definition above would include those that fund their own care in the measure – but would ensure that where the intervention was not successful they were identified and excluded from the numerator. This approach would be reliant on councils properly recording those that leave short term services but go on to purchase their own care.</p>

<b>Alignment</b>	ASCOF measure only
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**New placeholder for 2013/14****(2E) The effectiveness of reablement services**

<b>Domain / Outcome</b>	2. Delaying and reducing the need for care and support <i>Earlier diagnosis, intervention and reablement means that people and their carers are less dependent on intensive services.</i>
<b>Rationale</b>	This placeholder signals intent to measure the effectiveness of short-term services, to be viewed in the context of Measure 2D, to understand whether there are any unintended consequences of the decision to provide no further services, Measures 2D and 2E, once implemented alongside the current measure of outcomes from reablement/rehabilitation services (Measure 2B), will provide a more comprehensive view of the effectiveness of reablement care and support.
<b>Definition / interpretation</b>	Under development.
<b>Alignment</b>	ASCOF measure only

**(2C) Delayed transfers of care from hospital, and those which are attributable to adult social care per 100,000 population**

<b>Domain / Outcome</b>	2. Delaying and reducing the need for care and support. <i>When people develop care needs, the support they receive takes place in the most appropriate setting, and enables them to regain their independence.</i>
<b>Rationale</b>	This measures the impact of hospital services (acute, mental health and non-acute) and community-based care in facilitating timely and appropriate transfer from all hospitals for all adults. This indicates the ability of the whole system to ensure appropriate transfer from hospital for the entire adult population. It is an important marker of the effective joint working of local partners, and is a measure of the effectiveness of the interface between health and social care services. Minimising delayed transfers of care and enabling people to live independently at home is one of the desired outcomes of social care.
<b>Definition / interpretation</b>	This is a two-part measure that reflects both the overall number of delayed transfers of care (2C part 1) and, as a subset, the number of these delays which are attributable to social care services (2C part 2).  A delayed transfer of care occurs when a patient is ready for transfer from a hospital bed, but is still occupying such a bed.  A patient is ready for transfer when: (a) a clinical decision has been made that the patient is ready for transfer AND (b) a multi-disciplinary team decision has been made that the patient is ready for transfer AND (c) the patient is safe to discharge/transfer.  Set out below is a table showing UNIFY2 definitions for the attribution of different reasons for delay:



		<i>Attributable to NHS</i>	<i>Attributable to Social Care</i>	<i>Attributable to both</i>
	A. Awaiting completion of assessment	✓	✓	✓
	B. Awaiting public funding	✓	✓	✓
	C. Awaiting further non-acute (including community and mental health) NHS care (including intermediate care, rehabilitation services etc)	✓	x	x
	Di). Awaiting residential home placement or availability	✓	✓	x
	Dii). Awaiting nursing home placement or availability	✓	✓	✓
	E. Awaiting care package in own home	✓	✓	✓
	F. Awaiting community equipment and adaptations	✓	✓	✓
	G. Patient or family choice	✓	✓	x
	H. Disputes	✓	✓	x
	I. Housing – patients not covered by NHS and Community Care Act	✓	x	x
	<b>Interpretation</b>			
	Using a two-part measure means that we can maintain a focus on joint working, while balancing this with a measure that focuses more closely on the specific contribution of social care services.			
<b>Alignment</b>	ASCOF measure only			
<b>Risk adjustment</b>	Risk adjustment does not seem appropriate for this measure since the objective is that delayed transfers of care are minimised. The factors affecting whether this is achieved should largely be within the control of local health and care services.			
<b>Formula</b>	$\left(\frac{X}{Y}\right) \times 100,000$ <p>Where, for <b>2C part 1</b> (total delayed transfers):</p> <p><b>X:</b> The average number of delayed transfers of care (for those aged 18 and over) on a particular day taken over the year. This is the average of the 12 monthly snapshots collected in the monthly Situation Report (SitRep). Source: <i>Unify2</i></p> <p><b>Y:</b> Size of adult population in area (aged 18 and over) Source: <i>ONS mid year population estimates</i><sup>26</sup></p> <p>For <b>2C part 2</b> (delayed transfers attributable to social care):</p> <p><b>X:</b> The average number of delayed transfers of care (for those aged 18 and over) on a particular day taken over the year, that are attributable to social care or jointly to social care and the NHS. This is the average of the 12 monthly snapshots. Source: <i>UNIFY2</i></p> <p><b>Y:</b> Size of adult population in area (aged 18 and over) Source: <i>ONS mid year population estimates</i><sup>27</sup></p>			

<sup>26</sup> If a population estimate does not exist for the current year then the previous year's estimate will be used.

<sup>27</sup> If a population estimate does not exist for the current year then the previous year's estimate will be used.

<b>Worked example</b>	<p>Suppose the total number of delayed discharges from the 12 monthly snapshots is 812.</p> <p>Divide this by 12 for a monthly figure.</p> <p>And if the ONS mid-year population estimate = 570,562</p> <p>Therefore the average rate of delayed transfers is calculated as:</p> $((812 / 12) / 570,562) * 100,000$ <p>= <b>11.9</b></p> <p>If the total number of delays attributable to social care or jointly to social care and the NHS is 271, the average rate of delayed transfers of care attributable to social care or social care and the NHS jointly is calculated as:</p> $((271 / 12) / 570,562) * 100,000$ <p>= <b>4.0</b></p>		
<b>Disaggregation available</b>	<p><b>Equalities:</b> Age (18+)</p> <p><b>Client groups:</b> Adults aged 18+</p>		
<b>Frequency of collection</b>	Annual	<b>Data source</b>	UNIFY2 (DH) Office of National Statistics
<b>Return format</b>	Numeric	<b>Decimal places</b>	One
<b>Longer-term development options</b>	None identified		
<b>Further guidance</b>	<p>Guidance for 2012/13 onwards can be found via the social care collection page at <a href="http://www.ic.nhs.uk/services/social-care/social-care-collections">http://www.ic.nhs.uk/services/social-care/social-care-collections</a> by clicking on the year.</p> <p>Guidance on UNIFY2 can be found at: <a href="http://transparency.dh.gov.uk/2012/06/21/dtoc-information/">http://transparency.dh.gov.uk/2012/06/21/dtoc-information/</a></p> <p>Delayed discharges data can be found at: <a href="http://www.dh.gov.uk/en/Publicationsandstatistics/Statistics/Perfomancedataandstatistics/AcuteandNon-AcuteDelayedTransfersofCare/index.htm">http://www.dh.gov.uk/en/Publicationsandstatistics/Statistics/Perfomancedataandstatistics/AcuteandNon-AcuteDelayedTransfersofCare/index.htm</a></p>		

***New placeholder for 2013/14***

***(2F) Dementia – a measure of the effectiveness of post-diagnosis care in sustaining independence and improving quality of life***

<b>Domain / Outcome</b>	<p>2. Delaying and reducing the need for care and support.</p> <p><i>When people develop care needs, the support they receive takes place in the most appropriate setting and enables them to regain their independence.</i></p>
<b>Rationale</b>	<p>The Care and Support White Paper reinforced the Prime Minister's 'Challenge on Dementia,' which sets out a renewed ambition to go 'further and faster', building on progress made through the National Dementia Strategy to secure greater improvements in dementia care. The placeholder signals the intent to develop a measure to assess the impact of this challenge, which will focus on the effectiveness of post-diagnosis care in sustaining independence and improving quality of life.</p>

<b>Alignment</b>	This measure is shared with Measure 2.6ii (effectiveness of post-diagnosis care in sustaining independence for people with dementia) in the NHS Outcomes Framework ( <a href="https://www.wp.dh.gov.uk/publications/files/2012/11/121109-Technical-Appendix.pdf">https://www.wp.dh.gov.uk/publications/files/2012/11/121109-Technical-Appendix.pdf</a> page 42).
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## Domain 3 – Ensuring that people have a positive experience of care and support

<b>(3A) Overall satisfaction of people who use services with their care and support</b>	
<b>Domain / Outcome</b>	3. Ensuring people have a positive experience of care and support. <i>People who use social care and their carers are satisfied with their experience of care and support services.</i> <i>(Overarching measure)</i>
<b>Rationale</b>	This measures the satisfaction with services of people using adult social care, which is directly linked to a positive experience of care and support. Analysis of surveys suggests that reported satisfaction with services is a good predictor of people’s overall experiences of services.
<b>Definition / interpretation</b>	<p>The relevant question drawn from the Adult Social Care Survey is Question 1: “Overall, how satisfied or dissatisfied are you with the care and support services you receive?”, to which the following answers are possible:</p> <ul style="list-style-type: none"> <li>▪ I am extremely satisfied</li> <li>▪ I am very satisfied</li> <li>▪ I am quite satisfied</li> <li>▪ I am neither satisfied nor dissatisfied</li> <li>▪ I am quite dissatisfied</li> <li>▪ I am very dissatisfied</li> <li>▪ I am extremely dissatisfied</li> </ul> <p>The relevant question drawn from the Easy Read Adult Social Care questionnaire is Question 1: “How happy are you with the way staff help you?”, to which the following answers are possible:</p> <ul style="list-style-type: none"> <li>▪ I am very happy with the way staff help me, it’s really good</li> <li>▪ I am quite happy with the way staff help me</li> <li>▪ The way staff help me is OK</li> <li>▪ I do not think the way staff help me is that good</li> <li>▪ I think the way staff help me is really bad</li> </ul> <p>The measure is defined by determining the percentage of all those responding who identify strong satisfaction – i.e. by choosing the answer “I am extremely satisfied” or the answer “I am very satisfied”, and of those responding to the Easy Read questionnaire, who choose the answer “I am very happy with the way staff help me, it’s really good”.</p>
<b>Alignment</b>	ASCOF only measure
<b>Risk adjustment</b>	While this question asks directly about services, it is potentially subject to influence of exogenous factors. For example a previous study of home care users suggested that better perceptions of home care were related to, amongst other things, receiving less than ten hours home care (a proxy for need) and receiving help from others. Further analysis will be required to explore this and establish whether risk adjustment should be applied.

<b>Formula</b>	$\left(\frac{X}{Y}\right) \times 100$ <p>Where:</p> <p><b>X:</b> In response to Question 1, those individuals who selected the response “I am extremely satisfied” or “I am very satisfied”, and those who select the response “I am very happy with the way staff help me, it’s really good”, in response to Question 1 of the Easy Read questionnaire.</p> <p><b>Y:</b> All those that responded to the question.</p> <p>For both the numerator (X) and denominator (Y), weighted data should be used to calculate the measure. The data from the survey will be weighted by the HSCIC to take account of the stratified sampling technique that has been used when conducting the survey. The weights are automatically calculated within the survey data return along with the ASCOF outcome measures. Further details of how to use the weights when analysing the survey data are available in Appendix I of the guidance for the 2012-13 Adult Social Care Survey.</p>		
<b>Worked example</b>	<p>The number of users who said “I am extremely satisfied” or “I am very satisfied” was 217 and the number of users who said “I am very happy with the way staff help me, it’s really good”, in response to Question 1 of the Easy Read questionnaire was 30.</p> <p>In total the number of users who responded to the question (including the easy read questionnaire) was 398.</p> <p>(Data weighted to reflect the stratified sampling technique that has been used when conducting the survey.)</p> <p>The measure value is <math>[(217 + 30)/398] \times 100 = 62.1\%</math></p>		
<b>Disaggregation available</b>	<b>Equalities:</b> Age, Gender, Ethnicity**, Religion*, Sexual orientation*		
	<b>Client groups:</b> Physical Disability (all ages**), Learning Disability (all ages**), Mental health (all ages)**, Substance Misuse (all ages)**, Vulnerable People (all ages)**		
<b>Frequency of collection</b>	Annual	<b>Data source</b>	<i>Adult Social Care Survey</i>
<b>Return format</b>	Percentage	<b>Decimal places</b>	One
<b>Longer-term development options</b>	None identified		

\*\* This information is not published as part of the adult social care outcomes returns, however it is part of the publication of the data source

Although the underlying survey results will in theory be disaggregated by religion and sexual orientation, in practice there are likely to be significant gaps in the data on these characteristics at least in the short/medium term. This reflects the content of records held locally by councils.

<b>Further guidance</b>	Guidance for 2012/13 onwards can be found via the user survey guidance page at <a href="http://www.ic.nhs.uk/socialcare/usersurveys">http://www.ic.nhs.uk/socialcare/usersurveys</a>
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<b>(3B) Overall satisfaction of carers with social services</b>	
<b>Domain / Outcome</b>	3. Ensuring people have a positive experience of care and support. <i>People who use social care and their carers are satisfied with their experience of care and support services.</i> <i>(Overarching measure)</i>
<b>Rationale</b>	This measures the satisfaction with services of carers of people using adult social care, which is directly linked to a positive experience of care and support. Analysis of user surveys suggests that reported satisfaction with services is a good predictor of the overall experience of services and quality.
<b>Definition / interpretation</b>	<p><b>This measure will be deferred in 2013/14</b></p> <p>The relevant question drawn from the Carers Survey is question 4: “Overall, how satisfied or dissatisfied are you with the support or services you and the person you care for have received from Social Services in the last 12 months?”, to which the following answers are possible:</p> <ul style="list-style-type: none"> <li>▪ We haven’t received any support or services from Social Services in the last 12 months</li> <li>▪ I am extremely satisfied</li> <li>▪ I am very satisfied</li> <li>▪ I am quite satisfied</li> <li>▪ I am neither satisfied nor dissatisfied</li> <li>▪ I am quite dissatisfied</li> <li>▪ I am very dissatisfied</li> <li>▪ I am extremely dissatisfied</li> </ul> <p>The measure is defined by determining the percentage of all those responding who identify strong satisfaction – i.e. by choosing the answer “I am extremely satisfied” or the answer “I am very satisfied”.</p>
<b>Alignment</b>	ASCOF measure only
<b>Risk adjustment</b>	While this question asks directly about services, it is potentially subject to influence of exogenous factors. For example a previous study of home care users suggested that better perceptions of home care were related to receiving less than 10 hours home care (a proxy for need) and receiving help from others . Further analysis will be required to explore this and establish whether risk adjustment should be applied.

<b>Formula</b>	$\left(\frac{X}{Y}\right) \times 100$ <p><b>Where:</b></p> <p><b>X:</b> In response to the question above, those individuals who selected the response “I am extremely satisfied” or “I am very satisfied”.</p> <p><b>Y:</b> All those that responded to the question.</p> <p><b>Exclusions</b></p> <p>People who select the response “We haven’t received any support or services from Social Services in the last 12 months” will not be counted in either the numerator or the denominator.</p>		
	<b>Worked example</b>	<p>The number of carers who said “I am extremely satisfied” or “I am very satisfied” was 112.</p> <p>In total the number of carers who responded to the question was 160 but 7 gave a response of “We haven’t received any support or services from Social Services in the last 12 months”.</p> <p>The measure value is <math>[(112/(160-7))*100] = 73.2\%</math></p>	
<b>Disaggregation available</b>	<b>Equalities:</b> Age, Gender, Ethnicity**, Religion*, Sexual orientation		
	<b>Client groups:</b> Carers		
<b>Frequency of collection</b>	Biennial (Carers Survey to be next conducted in 2014/15)	<b>Data source</b>	<i>Carers Survey</i>
<b>Return format</b>	Percentage	<b>Decimal places</b>	One
<b>Longer-term development options</b>	There remains potential for moving to an annual collection if burden can be reduced significantly, subject to the agreement of local government.		
<b>Further guidance</b>	Guidance for 2012/13 onwards can be found via the user survey guidance page at <a href="http://www.ic.nhs.uk/socialcare/usersurveys">http://www.ic.nhs.uk/socialcare/usersurveys</a> .		

**New placeholder for 2013/14****(3E) Improving people’s experience of integrated care**

<b>Domain / Outcome</b>	3. Ensuring that people have a positive experience of care and support. <i>People who use social care and their carers are satisfied with their experience of care and support services.</i>
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\*\* This information is not published as part of the adult social care outcomes returns, however it is part of the publication of the data source.

Although the underlying survey results will in theory be disaggregated by religion and sexual orientation, in practice there are likely to be significant gaps in the data on these characteristics at least in the short/medium term. This reflects the content of records held locally by councils.

<b>Rationale</b>	<p>As highlighted by the NHS Future Forum, 'integration is a vitally important aspect of the experience of health and social care for millions of people. It has perhaps the greatest relevance for the most vulnerable and those with the most complex and long-term needs'. However, at present, no direct measurement of people's experience of integrated care exists.</p> <p>The Care and Support White Paper restated the Department's commitment to a clear, ambitious and measurable goal to drive further improvements in people's experience of integrated care.</p>
<b>Alignment</b>	<p>This placeholder is shared with measure 4.9 (people's experience of integrated care) in the NHS Outcomes Framework (<a href="https://www.wp.dh.gov.uk/publications/files/2012/11/121109-Technical-Appendix.pdf">https://www.wp.dh.gov.uk/publications/files/2012/11/121109-Technical-Appendix.pdf</a> page 86).</p>

**(3C) The proportion of carers who report that they have been included or consulted in discussion about the person they care for**

<b>Domain / Outcome</b>	<p>3. Ensuring people have a positive experience of care and support. <i>Carers feel that they are respected as equal partners throughout the care process.</i></p>
<b>Rationale</b>	<p>Carers should be respected as equal partners in service design for those individuals for whom they care – this improves outcomes both for the cared for person and the carer, reducing the chance of breakdown in care. This measure reflects the experience of carers in how they have been consulted by both the NHS and social care.</p>
<b>Definition / interpretation</b>	<p><b>This measure will be deferred in 2013/14.</b></p> <p>The relevant question drawn from the Carers Survey is Q15: "In the last 12 months, do you feel you have been involved or consulted as much as you wanted to be, in discussions about the support or services provided to the person you care for?", to which the following answers are possible:</p> <ul style="list-style-type: none"> <li>▪ There have been no discussions that I am aware of, in the last 12 months</li> <li>▪ I always felt involved or consulted</li> <li>▪ I usually felt involved or consulted</li> <li>▪ I sometimes felt involved or consulted</li> <li>▪ I never felt involved or consulted</li> </ul> <p>The measure is defined by determining the percentage of all those responding who choose the answer "I always felt involved or consulted" and "I usually felt involved or consulted".</p>
<b>Alignment</b>	<p>ASCOF measure only</p>
<b>Risk adjustment</b>	<p>None</p>

<b>Formula</b>	$\left(\frac{X}{Y}\right) \times 100$ <p>Where:</p> <p><b>X:</b> In response to the above question, all those individuals who selected the response "I always felt involved or consulted" and "I usually felt involved or consulted".</p> <p><b>Y:</b> All those that responded to the question.</p> <p><i>Exclusions</i> People who select the response "There have been no discussions that I am aware of, in the last 12 months" will not be counted in either the numerator or the denominator.</p>		
<b>Worked example</b>	<p>The number of carers who said "I always felt involved or consulted" (and "I usually felt involved or consulted") was 129.</p> <p>In total the number of carers who responded to the question was 160 with 7 giving a response of "There have been no discussion that I am aware of, in the last 12 months".</p> <p>The measure value is <math>[(129/(160-7))*100] = 84.3\%</math></p>		
<b>Disaggregation available</b>	<p><b>Equalities:</b> Age, Gender, Ethnicity, Religion*, Sexual Orientation *</p> <p><b>Client groups:</b> Carers</p>		
<b>Frequency of collection</b>	Biennial (Carers Survey to be next conducted in 2014/15)	<b>Data source</b>	<i>Carers Survey</i>
<b>Return format</b>	Percentage	<b>Decimal places</b>	One
<b>Longer-term development options</b>	There remains potential for moving to an annual collection if burden can be reduced significantly, subject to the agreement of local government.		
<b>Further guidance</b>	Guidance for 2012/13 onwards can be found via the user survey guidance page at <a href="http://www.ic.nhs.uk/services/social-care/social-care-collections/user-surveys">http://www.ic.nhs.uk/services/social-care/social-care-collections/user-surveys</a> .		

### (3D) The proportion of people who use services and carers who find it easy to find information about services

<b>Domain / Outcome</b>	3. Ensuring people have a positive experience of care and support. <i>People know what choices are available to them locally, what they are entitled to, and who to contact when they need help.</i>
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\* Although the underlying survey results will in theory be disaggregated by religion and sexual orientation, in practice there are likely to be significant gaps in the data on these characteristics at least in the short/medium term. This reflects the content of records held locally by councils.



<p><b>Rationale</b></p>	<p>This measure reflects social services users' and carers' experience of access to information and advice about social care in the past year. Information is a core universal service and a key factor in early intervention and reducing dependency.</p> <p>Improved and/or more information benefits carers and the people they support by helping them to have greater choice and control over their lives. This may help to sustain caring relationships through, for example, reduction in stress, improved welfare and physical health improvements. These benefits accrue only where information is accessed that would not otherwise have been accessed, or in those cases where the same information is obtained more easily.</p>
<p><b>Definition / interpretation</b></p>	<p>This measure is in two parts and uses questions in the Adult Social Care Survey and Carers Survey.</p> <p>The question from the Adult Social Care Survey is Question 12: "In the past year, have you generally found it easy or difficult to find information and advice about support, services or benefits?", to which the following answers are possible:</p> <ul style="list-style-type: none"> <li>▪ Very easy to find</li> <li>▪ Fairly easy to find</li> <li>▪ Fairly difficult to find</li> <li>▪ Very difficult to find</li> <li>▪ I've never tried to find information or advice</li> </ul> <p>This portion of the measure is defined by determining the percentage of all those responding who select the response "very easy to find" and "fairly easy to find".</p> <p>The relevant question drawn from the Carers Survey is Question 13 : "In the last 12 months, have you found it easy or difficult to find information and advice about support, services or benefits? Please include information and advice from different sources, such as voluntary organisations and private agencies as well as Social Services". The following answers are possible:</p> <ul style="list-style-type: none"> <li>▪ I have not tried to find information or advice in the last 12 months</li> <li>▪ Very easy to find</li> <li>▪ Fairly easy to find</li> <li>▪ Fairly difficult to find</li> <li>▪ Very difficult to find</li> </ul> <p>This portion of the measure is defined by determining the percentage of all those responding who select the response "very easy to find" and "fairly easy to find".</p> <p>The measure is then defined by determining the average percentage across the two surveys of all those responding who select the response "very easy to find" and "fairly easy to find".</p> <p>The Adult Social Care Survey will be annual whereas the Carers Survey will, at least initially, be biennial.</p>
<p><b>Alignment</b></p>	<p>ASCOF only measure</p>
<p><b>Risk adjustment</b></p>	<p>None</p>
<p><b>Formula</b></p>	$\left( \frac{X}{Y} \right) * 100$ <p>Where:</p> <p>For <b>3D part 1</b> (users):</p>

	<p><b>X:</b> In response to Question 12 of the ASCS, those individuals who selected the response “very easy to find” and “fairly easy to find”.</p> <p>Those respondents who were sent the version of the questionnaire for people with learning disabilities will be treated in the same way, as this questionnaire has been designed to be equivalent to the non-learning disabilities version.</p> <p><b>Y:</b> All those that responded to the question.</p> <p>For both the numerator (X) and denominator (Y), weighted data should be used to calculate the measure. The data from the survey will be weighted by the Health and Social Care Information Centre to take account of the stratified sampling technique that has been used when conducting the survey. The weights are automatically calculated within the survey data return along with the ASCOF outcome measures. Further details of how to use the weights when analysing the survey data are available in Appendix I of the guidance for the 2012-13 Adult Social Care Survey.</p> <p>Where:</p> <p>For <b>3D part 2</b> (carers):</p> <p><b>X:</b> The sum of all those who in response to the above question of the Carers Survey, selected the response “very easy to find” and “fairly easy to find”.</p> <p>In years where the Carers Survey has not been completed (since this is initially a biennial collection), the most recent value for the carers element should be carried over and counted in the second of these parts in the numerator. In these years, only the changes in the service user element (drawn from the ASCS) will be identifiable.</p> <p><b>Y:</b> The sum of all those that responded to the above question of the Carers Survey.</p> <p><b>Exclusions</b></p> <p>People who select the response “I’ve never tried to find information or advice” for the ASCS or “I have not tried to find information or advice in the last 12 months” for the Carers Survey will not be counted in either the numerator or the denominator.</p>
<p><b>Worked example</b></p>	<p><b>3D Part 1</b> (users)</p> <p>The number of respondents to the Adult Social Care Survey who select the response “Very easy to find” or “fairly easy to find” was 191.</p> <p>In total the number of users who responded to the question was 350 of whom 8 gave a response of “I’ve never tried to find information or advice”.</p> <p>The score for the ASCS is <math>[(191/(350-8))*100] = 55.8\%</math></p> <p>(Data weighted to reflect the stratified sampling technique that has been used when conducting the survey)</p> <p><b>3D Part 2</b> (carers)</p> <p>The number of respondents to the Carers Survey who select the responses “very easy to find” or “fairly easy to find” was 93.</p> <p>In total the number of users who responded to the question was 220 of whom 8 gave a response of “I have not tried to find information or advice in the last 12 months”.</p> <p>The score for the Carers Survey is <math>[(93/(220-8))*100] = 43.9\%</math></p>

<b>Disaggregation available</b>	<b>Equalities:</b> Age, Gender, Ethnicity**, Religion*, Sexual orientation*		
	<b>Client groups:</b> Physical Disability (all ages)** , Learning Disability (all ages)** , Mental Health (all ages)** , Substance Misuse (all ages)** , Vulnerable people (all ages)** , Carers		
<b>Frequency of collection</b>	Annual (ASCS) Biennial (Carers Survey to be next conducted In 2014/15)	<b>Data source</b>	<i>Adult Social Care Survey Carers Survey</i>
<b>Return format</b>	Percentage	<b>Decimal places</b>	One
<b>Longer-term development options</b>	<p>This measure does not include self-funders or people with low-level services that may have been directed to voluntary organisations. In the future, we will look at the feasibility of putting in place a broader measure to capture outcomes for these groups.</p> <p>There remains potential for moving to an annual carers collection if burden can be reduced significantly, subject to the agreement of local government.</p>		
<b>Further guidance</b>	Guidance for 2012/13 onwards can be found via the user survey guidance page at <a href="http://www.ic.nhs.uk/socialcare/usersurveys">http://www.ic.nhs.uk/socialcare/usersurveys</a>		

## Domain 4 – Safeguarding adults whose circumstances make them vulnerable and protecting from avoidable harm

### (4A) The proportion of people who use services who feel safe

<b>Domain / Outcome</b>	4. Safeguarding people whose circumstances make them vulnerable and protecting from avoidable harm. ( <i>Overarching measure</i> )
<b>Rationale</b>	<p>This measures one component of the overarching 'social care-related quality of life' measure. It provides an overarching measure for this domain.</p> <p>Safety is fundamental to the wellbeing and independence of people using social care, and the wider population. Feeling safe is a vital part of users' experience of their care and support. There are legal requirements about safety in the context of service quality, including CQC's essential standards for registered services. There is also a vital role of being safe in the quality of the individual's experience.</p>

\* Although the underlying survey results will in theory be disaggregated by religion and sexual orientation, in practice there are likely to be significant gaps in the data on these characteristics at least in the short/medium term. This reflects the content of records held locally by councils.

\*\* This information is not published as part of the adult social care outcomes returns; however it is part of the publication of the data source.

<p><b>Definition / interpretation</b></p>	<p>The relevant question drawn from the Adult Social Care Survey is Question 7a: “Which of the following statements best describes how safe you feel?”, to which the following answers are possible:</p> <ul style="list-style-type: none"> <li>▪ I feel as safe as I want</li> <li>▪ Generally I feel adequately safe, but not as safe as I would like</li> <li>▪ I feel less than adequately safe</li> <li>▪ I don't feel at all safe</li> </ul> <p>The measure is defined by determining the percentage of all those responding who choose the answer “I feel as safe as I want”.</p> <p>Those respondents who were sent the version of the questionnaire for people with learning disabilities will be treated in the same way, as this questionnaire has been designed to be equivalent to the non-learning disabilities version.</p> <p><i>Interpretation</i></p> <p>The measure gives an overall indication of a reported outcome for individuals – it does not, at present, identify the specific contribution of councils’ adult social care towards to feeling safe (see measure 4B below).</p> <p>While the measure will measure those choosing the most positive response - "I feel as safe as I want" - it will be important locally to analyse responses on safeguarding in the context of the distribution of answers across all four possible responses. For example, if a council has a relatively high proportion of respondents selecting "I feel as safe as I want" (i.e. scores highly on the measure) but also has a relatively high proportion of respondents selecting "I don't feel at all safe", this could reflect gaps in safeguarding services.</p>
<p><b>Alignment</b></p>	<p>This measure is complementary to measure 1.19 (older people's perception of community safety placeholder) in the Public Health Outcomes Framework (<a href="http://www.dh.gov.uk/prod_consum_dh/groups/dh_digitalassets/@dh/@en/documents/digitalasset/dh_132558.pdf">http://www.dh.gov.uk/prod_consum_dh/groups/dh_digitalassets/@dh/@en/documents/digitalasset/dh_132558.pdf</a> page 25).</p>
<p><b>Risk adjustment</b></p>	<p>A range of factors will be considered to adjust the measure to improve comparability between councils. Some example are:</p> <ul style="list-style-type: none"> <li>▪ Age of users</li> <li>▪ Needs of users</li> <li>▪ Client groups of users</li> </ul>
<p><b>Formula</b></p>	$\left( \frac{X}{Y} \right) * 100$ <p>Where:</p> <p><b>X:</b> In response to Question 7a, those individuals who selected the response “I feel as safe as I want”.</p> <p>Those respondents who were sent the version of the questionnaire for people with learning disabilities will be treated in the same way, as this questionnaire has been designed to be equivalent to the non-learning disabilities version.</p> <p><b>Y:</b> All those that responded to the question.</p> <p>For both the numerator (X) and denominator (Y), weighted data should be used to calculate the measure. The data from the survey will be weighted by the Health and Social Care Information Centre to take account of the stratified sampling technique that has been used when conducting the survey. The weights are automatically calculated within the survey data return along with the ASCOF outcome measures. Further details of how to use the</p>

	weights when analysing the survey data are available in Appendix I of the guidance for the 2012-13 Adult Social Care Survey.		
<b>Worked example</b>	<p>The number of users who said “I feel as safe as I want” was 214.</p> <p>In total the number of users who responded to the question was 345.</p> <p>(Data weighted to reflect the stratified sampling technique that has been used when conducting the survey)</p> <p>The measure value is <math>[(214/345)*100] = 62.0\%</math></p>		
<b>Disaggregation available</b>	<b>Equalities:</b> Age, Gender, Ethnicity**, Religion*, Sexual Orientation*		
	<b>Client groups:</b> Physical Disability (all ages)** , Learning Disability (all ages)** , Mental Health (all ages)** , Substance misuse (all ages)** , Vulnerable People (all ages)**		
<b>Frequency of collection</b>	Annual	<b>Data source</b>	<i>Adult Social Care Survey</i>
<b>Return format</b>	Percentage	<b>Decimal places</b>	One
<b>Longer-term development options</b>	Develop a broader 'value-added' measure which quantifies the contribution of social services to people feeling safe.		
<b>Further guidance</b>	We will consider whether and how the development of a broader 'value-added' measure for measure 1A, which quantifies the contribution of social services to social care related quality of life, can or should be applied to this measure.		

#### **(4B) The proportion of people who use services who say that those services have made them feel safe and secure**

<b>Domain / Outcome</b>	<p>4. Safeguarding people whose circumstances make them vulnerable and protecting from avoidable harm</p> <p><i>Everyone enjoys physical safety and feels secure.</i></p> <p><i>People are free from physical and emotional abuse, harassment, neglect and self-harm.</i></p> <p><i>People are protected as far as possible from avoidable harm, disease and injury.</i></p> <p><i>People are supported to plan ahead and have the freedom to manage risks in the way that they wish.</i></p>
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\* Although the underlying survey results will in theory be disaggregated by religion and sexual orientation, in practice there are likely to be significant gaps in the data on these characteristics at least in the short/medium term. This reflects the content of records held locally by councils.

\*\* This information is not published as part of the adult social care outcomes returns, however it is part of the publication of the data source.

<b>Rationale</b>	<p>Safety is fundamental to the wellbeing and independence of people using social care, and the wider population. Feeling safe is a vital part of users' experience and their care and support. There are legal requirements about safety in the context of service quality, including CQC essential standards for registered services.</p> <p>This measure supports measure 4A by reflecting the extent to which users of care services feel that their care and support has contributed to making them feel safe and secure. As such, it goes some way to separate the role of care and support in helping people to feel safe from the influence of other factors, such as crime levels and socio-economic factors.</p>
<b>Definition / interpretation</b>	<p>The relevant question drawn from the Adult Social Care Survey is Question 7b: "Do care and support services help you in feeling safe?" To which the following answers are possible:</p> <ul style="list-style-type: none"> <li>▪ Yes</li> <li>▪ No</li> </ul> <p>Those respondents who were sent the version of the questionnaire for people with learning disabilities will be treated in the same way, as this questionnaire has been designed to be equivalent to the non-learning disabilities version.</p> <p><i>Interpretation</i></p> <p>Whilst the overarching measure (4A) indicates a higher-level individual perspective on feeling safe, this measure complements this with a specific response on the impact of services on this outcome.</p>
<b>Alignment</b>	ASCOF measure only
<b>Risk adjustment</b>	While this question asks directly about services, it is potentially subject to influence of exogenous factors, for example the characteristics of users. Further analysis will be required to explore this and establish whether risk adjustment should be applied.
<b>Formula</b>	$\left( \frac{X}{Y} \right) * 100$ <p>Where:</p> <p><b>X:</b> In response to Question 7b, those individuals who selected the response "yes".</p> <p>Those respondents who were sent the version of the questionnaire for people with learning disabilities will be treated in the same way, as this questionnaire has been designed to be equivalent to the non-learning disabilities version.</p> <p><b>Y:</b> All those that responded to the question.</p> <p>For both the numerator (X) and denominator (Y), weighted data should be used to calculate the measure. The data from the survey will be weighted by the Health and Social Care Information Centre to take account of the stratified sampling technique that has been used when conducting the survey. The weights are automatically calculated within the survey data return along with the ASCOF outcome measures. Further details of how to use the weights when analysing the survey data are available in Appendix I of the guidance for the 2012-13 Adult Social Care Survey.</p>

<b>Worked example</b>	<p>The number of users who said services had helped them feel safe was 197.</p> <p>In total the number of users who responded to the question was 345.</p> <p>(Data weighted to reflect the stratified sampling technique that has been used when conducting the survey.)</p> <p>The measure value is <math>[(197/345)*100] = 57.1\%</math></p>		
<b>Disaggregation available</b>	<b>Equalities:</b> Age, Gender, Ethnicity **, Religion*, Sexual orientation *		
	<b>Client groups:</b> Physical Disability (all ages)*, Learning Disability (all ages)** , Mental health (all ages)** , Substance Misuse (all ages)** , Vulnerable People (all ages)**		
<b>Frequency of collection</b>	Annual	<b>Data source</b>	<i>Adult Social Care Survey</i>
<b>Return format</b>	Percentage	<b>Decimal places</b>	One
<b>Longer-term development options</b>	Develop a broader 'value-added' measure which quantifies the contribution of social services to people feeling safe.		
<b>Further guidance</b>	Guidance for 2012/13 onwards can be found via the user survey guidance page at <a href="http://www.ic.nhs.uk/article/2213/User-survey-guidance-2012-13">http://www.ic.nhs.uk/article/2213/User-survey-guidance-2012-13</a> .		

#### ***New placeholder for 2013/14***

#### ***(4C) Proportion of completed safeguarding referrals where people report that they feel safe***

<b>Domain / Outcome</b>	<p>4. Safeguarding adults whose circumstances make them vulnerable and protecting from avoidable harm.</p> <p><i>Everyone enjoys physical safety and feels secure</i>  <i>People are free from physical and emotional abuse, harassment, neglect and self-harm</i>  <i>People are protected as far as possible from avoidable harm, disease and injuries</i>  <i>People are supported to plan ahead and have the freedom to manage risks the way that they wish</i></p>
<b>Rationale</b>	<p>A high-quality service must be one which keeps people safe from harm and the area of safeguarding is one of the core priorities of adult social care. This area remains one of the critical developmental priorities for the future of the ASCOF, and as part of the zero-based review, work has been taken forward to develop a potential measure of the outcomes of safeguarding interventions.</p> <p>This placeholder signals the Department's intention to measure the proportion of completed safeguarding referrals where service users reported they felt safe. To develop this measure, further work and piloting is required. Should a pilot be successful, this measure will be included in a future ASCOF.</p>
<b>Definition / interpretation</b>	Under development.

\*\* This information is not published as part of the adult social care outcomes returns; however it is part of the publication of the data source.

Although the underlying survey results will in theory be disaggregated by religion and sexual orientation, in practice there are likely to be significant gaps in the data on these characteristics at least in the short/medium term. This reflects the content of records held locally by councils.

<b>Alignment</b>	ASCOF measure only



# Appendix 1: The Adult Social Care Outcomes Framework 2013/14 at a glance

## Adult Social Care Outcomes Framework 2013/14

### At a glance

<b>1</b>	<b>Enhancing quality of life for people with care and support needs</b>
<p><b>Overarching measure</b></p> <p>1A. Social care-related quality of life * (NHSOF 2)</p> <p><b>Outcome measures</b></p> <p><b>People manage their own support as much as they wish, so that are in control of what, how and when support is delivered to match their needs.</b></p> <p>1B. Proportion of people who use services who have control over their daily life  <b>To be revised from 2014/15:</b> 1C. Proportion of people using social care who receive self-directed support, and those receiving direct payments</p> <p><b>Carers can balance their caring roles and maintain their desired quality of life.</b></p> <p>1D. Carer-reported quality of life * (NHSOF 2.4)</p> <p><b>People are able to find employment when they want, maintain a family and social life and contribute to community life, and avoid loneliness or isolation.</b></p> <p>1E. Proportion of adults with a learning disability in paid employment *** (PHOF 1.8, NHSOF 2.2)</p> <p>1F. Proportion of adults in contact with secondary mental health services in paid employment *** (PHOF 1.8, NHSOF 2.5)</p> <p>1G. Proportion of adults with a learning disability who live in their own home or with their family ** (PHOF 1.6)</p> <p>1H. Proportion of adults in contact with secondary mental health services living independently, with or without support ** (PHOF 1.6)</p> <p><b>New measure for 2013/14:</b>  <b>1I. Proportion of people who use services and their carers, who reported that they had as much social contact as they would like. ** (PHOF 1.18)</b></p>	
<b>3</b>	<b>Ensuring that people have a positive experience of care and support</b>
<p><b>Overarching measure</b></p> <p><b>People who use social care and their carers are satisfied with their experience of care and support services.</b></p> <p>3A. Overall satisfaction of people who use services with their care and support</p> <p>3B. Overall satisfaction of carers with social services  <b>New placeholder 3E: Improving people's experience of integrated care ** (NHS OF 4.9)</b></p> <p><b>Outcome measures</b></p> <p><b>Carers feel that they are respected as equal partners throughout the care process.</b></p> <p>3C. The proportion of carers who report that they have been included or consulted in discussions about the person they care for</p> <p><b>People know what choices are available to them locally, what they are entitled to, and who to contact when they need help.</b></p> <p>3D. The proportion of people who use services and carers who find it easy to find information about support</p> <p><b>People, including those involved in making decisions on social care, respect the dignity of the individual and ensure support is sensitive to the circumstances of each individual.</b></p> <p><i>This information can be taken from the Adult Social Care Survey and used for analysis at the local level.</i></p>	

<b>2</b>	<b>Delaying and reducing the need for care and support</b>
<p><b>Overarching measures</b></p> <p>2A. Permanent admissions to residential and nursing care homes, per 100,000 population</p> <p><b>Outcome measures</b></p> <p><b>Everybody has the opportunity to have the best health and wellbeing throughout their life, and can access support and information to help them manage their care needs.</b></p> <p><b>Earlier diagnosis, intervention and reablement means that people and their carers are less dependent on intensive services.</b></p> <p>2B. Proportion of older people (65 and over) who were still at home 91 days after discharge from hospital into reablement/rehabilitation services ** (NHSOF 3.6)</p> <p><b>New measure for 2014/15: 2D. The outcomes of short-term services: sequel to service.</b>  <b>New placeholder 2E: Effectiveness of reablement services</b></p> <p><b>When people develop care needs, the support they receive takes place in the most appropriate setting, and enables them to regain their independence.</b></p> <p>2C. Delayed transfers of care from hospital, and those which are attributable to adult social care</p> <p><b>New placeholder 2F: Dementia - a measure of the effectiveness of post-diagnosis care in sustaining independence and improving quality of life. ** (NHSOF 2.6ii)</b></p>	

<b>4</b>	<b>Safeguarding adults whose circumstances make them vulnerable and protecting from avoidable harm</b>
<p><b>Overarching measure</b></p> <p>4A. The proportion of people who use services who feel safe * (PHOF 1.19)</p> <p><b>Outcome measures</b></p> <p><b>Everyone enjoys physical safety and feels secure.</b>  <b>People are free from physical and emotional abuse, harassment, neglect and self-harm.</b>  <b>People are protected as far as possible from avoidable harm, disease and injuries.</b>  <b>People are supported to plan ahead and have the freedom to manage risks the way that they wish.</b></p> <p>4B. The proportion of people who use services who say that those services have made them feel safe and secure</p> <p><b>New placeholder 4C: Proportion of completed safeguarding referrals where people report they feel safe</b></p>	
<p><b>Aligning across the Health and Care System</b></p> <p>* Indicator complementary</p> <p>** Indicator shared</p> <p>*** Indicator complementary with the Public Health Outcomes Framework and the NHS Outcomes framework</p> <p><b>Shared indicators:</b> The same indicator is included in each outcomes framework, reflecting a shared role in making progress</p> <p><b>Complementary indicators:</b> A similar indicator is included in each outcomes framework and these look at the same issue</p>	

# Appendix 2: The Public Health Outcomes Framework 2013-16 at a glance

## VISION

To improve and protect the nation's health and wellbeing and improve the health of the poorest fastest

### Outcome measures

- Outcome 1) Increased healthy life expectancy, i.e. taking account of the health quality as well as the length of life
- Outcome 2) Reduced differences in life expectancy and healthy life expectancy between communities (through greater improvements in more disadvantaged communities)

### Alignment across the Health and Care System

- \* Indicator shared with the NHS Outcomes Framework.
- \*\* Complementary to indicators in the NHS Outcomes Framework
- † Indicator shared with the Adult Social Care Outcomes Framework
- †† Complementary to indicators in the Adult Social Care Outcomes Framework
- Indicators in italics are placeholders, pending development or identification*

## Public Health Outcomes Framework 2013-2016 At a glance (Autumn 2012)

### 1 Improving the wider determinants of health

#### Objective

Improvements against wider factors which affect health and wellbeing and health inequalities

#### Indicators

- 1.1 Children in poverty
- 1.2 *School readiness (Placeholder)*
- 1.3 Pupil absence
- 1.4 First time entrants to the youth justice system
- 1.5 16-18 year olds not in education, employment or training
- 1.6 Adults with a learning disability / in contact with secondary mental health services, who live in stable and appropriate accommodation<sup>†</sup> (ASCOF 1G and 1H)
- 1.7 *People in prison who have a mental illness or a significant mental illness (Placeholder)*
- 1.8 Employment for those with long-term health conditions including adults with a learning disability or who are in contact with secondary mental health services<sup>†</sup> (i-NHSOF 2.2)<sup>††</sup> (ii-ASCOF 1E)<sup>†††</sup> (iii-NHSOF 2.5)<sup>††</sup> (iii-ASCOF 1F)
- 1.9 Sickness absence rate
- 1.10 Killed and seriously injured casualties on England's roads
- 1.11 *Domestic abuse (Placeholder)*
- 1.12 Violent crime (including sexual violence)
- 1.13 Re-offending levels
- 1.14 The percentage of the population affected by noise
- 1.15 Statutory homelessness
- 1.16 Utilisation of outdoor space for exercise / health reasons
- 1.17 *Fuel poverty (Placeholder)*
- 1.18 *Social isolation (Placeholder)*<sup>†</sup> (ASCOF 1I)
- 1.19 *Older people's perception of community safety (Placeholder)*<sup>††</sup> (ASCOF 4A)

### 2 Health improvement

#### Objective

People are helped to live healthy lifestyles, make healthy choices and reduce health inequalities

#### Indicators

- 2.1 Low birth weight of term babies
- 2.2 Breastfeeding
- 2.3 Smoking status at time of delivery
- 2.4 Under 18 conceptions
- 2.5 *Child development at 2 – 2.1/2 years (Placeholder)*
- 2.6 Excess weight in 4-5 and 10-11 year olds
- 2.7 Hospital admissions caused by unintentional and deliberate injuries in under 18s
- 2.8 Emotional well-being of looked after children
- 2.9 *Smoking prevalence – 15 year olds (Placeholder)*
- 2.10 *Self-harm (Placeholder)*
- 2.11 Diet
- 2.12 Excess weight in adults
- 2.13 Proportion of physically active and inactive adults
- 2.14 Smoking prevalence – adults (over 18s)
- 2.15 Successful completion of drug treatment
- 2.16 People entering prison with substance dependence issues who are previously not known to community treatment
- 2.17 Recorded diabetes
- 2.18 *Alcohol-related admissions to hospital (Placeholder)*
- 2.19 Cancer diagnosed at stage 1 and 2
- 2.20 Cancer screening coverage
- 2.21 Access to non-cancer screening programmes
- 2.22 Take up of the NHS Health Check programme – by those eligible
- 2.23 Self-reported well-being
- 2.24 Injuries due to falls in people aged 65 and over

### 3 Health protection

#### Objective

The population's health is protected from major incidents and other threats, whilst reducing health inequalities

#### Indicators

- 3.1 Fraction of mortality attributable to particulate air pollution
- 3.2 Chlamydia diagnoses (15-24 year olds)
- 3.3 Population vaccination coverage
- 3.4 People presenting with HIV at a late stage of infection
- 3.5 Treatment completion for TB
- 3.6 Public sector organisations with board approved sustainable development management plan
- 3.7 *Comprehensive, agreed inter-agency plans for responding to public health incidents and emergencies (Placeholder)*

### 4

#### Healthcare public health and preventing premature mortality

#### Objective

Reduced numbers of people living with preventable ill health and people dying prematurely, whilst reducing the gap between communities

#### Indicators

- 4.1 Infant mortality\* (NHSOF 1.6i)
- 4.2 Tooth decay in children aged 5
- 4.3 Mortality rate from causes considered preventable\*\* (NHSOF 1a)
- 4.4 Under 75 mortality rate from all cardiovascular diseases (including heart disease and stroke)\* (NHSOF 1.1)
- 4.5 Under 75 mortality rate from cancer\* (NHSOF 1.4i)
- 4.6 Under 75 mortality rate from liver disease\* (NHSOF 1.3)
- 4.7 Under 75 mortality rate from respiratory diseases\* (NHSOF 1.2)
- 4.8 Mortality rate from infectious and parasitic diseases
- 4.9 Excess under 75 mortality rate in adults with serious mental illness\* (NHSOF 1.5)
- 4.10 Suicide rate
- 4.11 Emergency readmissions within 30 days of discharge from hospital\* (NHSOF 3b)
- 4.12 Preventable sight loss
- 4.13 *Health-related quality of life for older people (Placeholder)*
- 4.14 Hip fractures in people aged 65 and over
- 4.15 Excess winter deaths
- 4.16 Estimated diagnosis rate for people with dementia\* (NHSOF 2.6i)

# Appendix 3: The NHS Outcomes Framework 2013/14 at a glance

<b>1</b>	<b>Preventing people from dying prematurely</b>
<b>Overarching indicators</b>	
1a Potential Years of Life Lost (PYLL) from causes considered amenable to healthcare i Adults ii Children and young people 1b Life expectancy at 75 i Males ii Females	
<b>Improvement areas</b>	
<b>Reducing premature mortality from the major causes of death</b> 1.1 Under 75 mortality rate from cardiovascular disease* (PHOF 4.4) 1.2 Under 75 mortality rate from respiratory disease* (PHOF 4.7) 1.3 Under 75 mortality rate from liver disease* (PHOF 4.6) 1.4 Under 75 mortality rate from cancer* (PHOF 4.5) i One- and ii Five-year survival from all cancers iii One- and iv Five-year survival from breast, lung and colorectal cancer	
<b>Reducing premature death in people with serious mental illness</b> 1.5 Excess under 75 mortality rate in adults with serious mental illness* (PHOF 4.9)	
<b>Reducing deaths in babies and young children</b> 1.6 i Infant mortality* (PHOF 4.1) ii Neonatal mortality and stillbirths iii Five year survival from all cancers in children	
<b>Reducing premature death in people with a learning disability</b> 1.7 Excess under 60 mortality rate in adults with a learning disability	

<b>3</b>	<b>Helping people to recover from episodes of ill health or following injury</b>
<b>Overarching indicators</b>	
3a Emergency admissions for acute conditions that should not usually require hospital admission 3b Emergency readmissions within 30 days of discharge from hospital* (PHOF 4.11)	
<b>Improvement areas</b>	
<b>Improving outcomes from planned treatments</b> 3.1 Total health gain as assessed by patients for elective procedures i Hip replacement ii Knee replacement iii Groin hernia iv Varicose veins v Psychological therapies	
<b>Preventing lower respiratory tract infections (LRTI) in children from becoming serious</b> 3.2 Emergency admissions for children with LRTI	
<b>Improving recovery from injuries and trauma</b> 3.3 Proportion of people who recover from major trauma	
<b>Improving recovery from stroke</b> 3.4 Proportion of stroke patients reporting an improvement in activity/lifestyle on the Modified Rankin Scale at 6 months	
<b>Improving recovery from fragility fractures</b> 3.5 Proportion of patients recovering to their previous levels of mobility/walking ability at: 30 and ii 120 days	
<b>Helping older people to recover their independence after illness or injury</b> 3.6 i Proportion of older people (65 and over) who were still at home 91 days after discharge from hospital into reablement/rehabilitation service*** (ASCOF 2B) ii Proportion offered rehabilitation following discharge from acute or community hospital	

<b>4</b>	<b>Ensuring that people have a positive experience of care</b>
<b>Overarching indicators</b>	
4a Patient experience of primary care i GP services ii GP Out of Hours services iii NHS Dental Services 4b Patient experience of hospital care 4c Friends and family test	
<b>Improvement areas</b>	
<b>Improving people's experience of outpatient care</b> 4.1 Patient experience of outpatient services	
<b>Improving hospitals' responsiveness to personal needs</b> 4.2 Responsiveness to in-patients' personal needs	
<b>Improving people's experience of accident and emergency services</b> 4.3 Patient experience of A&E services	
<b>Improving access to primary care services</b> 4.4 Access to i GP services and ii NHS dental services	
<b>Improving women and their families' experience of maternity services</b> 4.5 Women's experience of maternity services	
<b>Improving the experience of care for people at the end of their lives</b> 4.6 Bereaved carers' views on the quality of care in the last 3 months of life	
<b>Improving experience of healthcare for people with mental illness</b> 4.7 Patient experience of community mental health services	
<b>Improving children and young people's experience of healthcare</b> 4.8 An indicator is under development	
<b>Improving people's experience of integrated care</b> 4.9 An indicator is under development*** (ASCOF 3E)	

<b>2</b>	<b>Enhancing quality of life for people with long-term conditions</b>
<b>Overarching indicator</b>	
2 Health-related quality of life for people with long-term conditions** (ASCOF 1A)	
<b>Improvement areas</b>	
<b>Ensuring people feel supported to manage their condition</b> 2.1 Proportion of people feeling supported to manage their condition**	
<b>Improving functional ability in people with long-term conditions</b> 2.2 Employment of people with long-term conditions** (ASCOF 1E PHOF 1.8)	
<b>Reducing time spent in hospital by people with long-term conditions</b> 2.3 i Unplanned hospitalisation for chronic ambulatory care sensitive conditions (adults) ii Unplanned hospitalisation for asthma, diabetes and epilepsy in under 19s	
<b>Enhancing quality of life for carers</b> 2.4 Health-related quality of life for carers** (ASCOF 1D)	
<b>Enhancing quality of life for people with mental illness</b> 2.5 Employment of people with mental illness*** (ASCOF 1F & PHOF 1.8)	
<b>Enhancing quality of life for people with dementia</b> 2.6 i Estimated diagnosis rate for people with dementia* (PHOF 4.16) ii A measure of the effectiveness of post-diagnosis care in sustaining independence and improving quality of life*** (ASCOF 2F)	

<b>NHS Outcomes Framework 2013/14 at a glance</b>	
<b>Alignment across the Health and Social Care System</b>	
* Indicator shared with Public Health Outcomes Framework (PHOF)	
** Indicator complementary with Adult Social Care Outcomes Framework (ASCOF)	
*** Indicator shared with Adult Social Care Outcomes Framework	
**** Indicator complementary with Adult Social Care Outcomes Framework and Public Health Outcomes Framework	
***** Indicators in italics are placeholders, pending development or identification	

<b>5</b>	<b>Treating and caring for people in a safe environment and protect them from avoidable harm</b>
<b>Overarching indicators</b>	
5a Patient safety incidents reported 5b Safety incidents involving severe harm or death 5c Hospital deaths attributable to problems in care	
<b>Improvement areas</b>	
<b>Reducing the incidence of avoidable harm</b> 5.1 Incidence of hospital-related venous thromboembolism (VTE) 5.2 Incidence of healthcare associated infection (HCAI) i MRSA ii C. difficile 5.3 Incidence of newly-acquired category 2, 3 and 4 pressure ulcers 5.4 Incidence of medication errors causing serious harm	
<b>Improving the safety of maternity services</b> 5.5 Admission of full-term babies to neonatal care	
<b>Delivering safe care to children in acute settings</b> 5.6 Incidence of harm to children due to 'failure to monitor'	

## Appendix 4: Interpretation of social care-related quality of life measure

The social care related quality of life score for an individual is a composite measure using responses to questions from the ASCS covering eight domains (control, dignity, personal care, food and drink, safety, occupation, social participation and accommodation). The ASCOF measure provides a social care related quality of life score averaged across each of the users who responded to the Adult Social Care Survey (ASCS) in an authority i.e. it's an average quality of life score for those that responded to the ASCS.

The score will be influenced by a range of factors, one of which is the services provided by the authority. Some of the other factors that are likely to have had an influence are the needs of individuals, age and whether people receive informal care. Therefore, in its current form this measure does not solely reflect the impact of social care services but does capture people's experience in aspects of life relevant to social care.

The social care related quality of life measure tells us about outcomes for social care users but does not isolate the impact that care and support services have on those outcomes. The Department has commissioned research from the Quality and Outcomes of Person Centred Care Policy Research Unit to identify a way of generating a social care related quality of life 'value added' measure, which would allow us to identify the impact of adult social care on people's quality of life. This research is due to report its findings in summer 2014. If successful, this will allow us to develop a new or additional measure for the ASCOF.

### **How can the measure be used?**

If using the measure for benchmarking, then it is important that comparisons are made with authorities that have similar characteristics. A starting point might be the standard comparator groups as otherwise comparisons can be misleading.

At a local level the score for each of the questions that measures outcomes across the eight domains could be investigated. Comparing this to a national average or similar councils would help understand whether scores on any of the individual domains are better or worse than would be expected.

Also at a local level it may be useful to look at the distribution of scores of individuals on the social care related quality of life measure. This would help understand whether most people's scores are around the average or are distributed widely. This analysis could be repeated by service user characteristics such as primary client group, or services being used.

When the survey is repeated, time series comparisons can be made and a change in the level of the measure should be investigated. Reasons for the change in the level of the measure may be a change in the impact of service but could also be related to changes in the needs of the local population etc.

However when making comparisons it's important to remember that the results are estimates from survey data and so there will be a degree of uncertainty which will be greater as the results are broken down further and therefore based on fewer service users. The level of uncertainty is commonly represented by a confidence interval which gives a range around the estimate in which you can be reasonably confident that the true figure lies.

If you would like more information on calculating confidence intervals please see the links under "Helping you make better use of the results from User Surveys" on the following page of the Information Centre website <http://www.ic.nhs.uk/socialcare/usersurveys> .

## Appendix 5: Accommodation types that represent settled or non-settled accommodation for the purpose of measure 1H, 'Proportion of adults in contact with secondary mental health services living independently, with or without support'.

MHMDS Code	Accommodation type	Settled accommodation status
	1 = Settled accommodation 0 = Non-settled accommodation	
<b>Mainstream Housing (MA00)</b>		
MA01	Owner occupier	1
MA02	Settled mainstream housing with family/friends	1
MA03	Shared ownership scheme e.g. Social Homebuy Scheme (tenant purchase percentage of home value from landlord)	1
MA04	Tenant – Local Authority/Arms Length Management Organisation/Registered Landlord	1
MA05	Tenant – Housing Association	1
MA06	Tenant – private landlord	1
MA09	Other mainstream housing	1
<b>Homeless (HM00)</b>		
HM01	Rough sleeper	0
HM02	Squatting	0
HM03	Night shelter/emergency hostel/Direct access hostel (temporary accommodation accepting self referrals, no waiting list and relatively frequent vacancies)	0
HM04	Sofa surfing (sleeps on different friends floor each night)	0
HM05	Placed in temporary accommodation by Local Authority (including Homelessness resettlement service) e.g. Bed and Breakfast	0

accommodation

HM06 Staying with friends/family as a short term guest 0

HM07 Other homeless 0

**Accommodation with mental health care support (MH00)**

MH01 Supported accommodation (accommodation supported by staff or resident caretaker) 1

MH02 Supported lodgings (lodgings supported by staff or resident caretaker) 1

MH03 Supported group home (supported by staff or resident caretaker) 1

MH04 Mental Health Registered Care Home 0

MH09 Other accommodation with mental health care and support 1

**Acute/long stay healthcare residential facility/hospital (HS00)**

HS01 NHS acute psychiatric ward 0

HS02 Independent hospital/clinic 0

HS03 Specialist rehabilitation/recovery 0

HS04 Secure psychiatric unit 0

HS05 Other NHS facilities/hospital 0

HS09 Acute/long stay healthcare residential facility/hospital 0

**Accommodation with other (not specialist mental health) care support (CH00)**

CH01 Foyer – accommodation for young people aged 16-25 who are homeless or in housing need 1

CH02 Refuge 0

CH03 Non-Mental Health Registered Care Home 0

CH09 Other accommodation with care and support (not specialist mental health) 1

**Accommodation with criminal justice support (CJ00)**

Appendix 5: Accommodation types that represent settled or non-settled accommodation for the purpose of measure 1H, 'Proportion of adults in contact with secondary mental health services living independently, with or without support'.

CJ01	Bail/Probation hostel	1
CJ02	Prison	0
CJ03	Young Offenders Institution	0
CJ04	Detention Centre	0
CJ09	Other accommodation with criminal justice support such as ex-offender support	1

**Sheltered Housing (accommodation with a scheme manager or warden living on the premises or nearby, contactable by an alarm system if necessary) (SH00)**

SH01	Sheltered housing for older persons	1
SH02	Extra care sheltered housing (also known as 'very sheltered housing'. For people who are less able to manage on their own, but who do need an extra level of care. Services offered vary between schemes, but meals and some personal care are often provided.)	1
SH03	Nursing Home	0
SH09	Other sheltered housing	1

**Mobile accommodation**

ML00	Mobile accommodation (for Gypsy/Roma and Traveller community)	1
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**Other codes**

OC96	Not elsewhere classified
OC97	Not specified
OC98	Not applicable
OC99	Not applicable