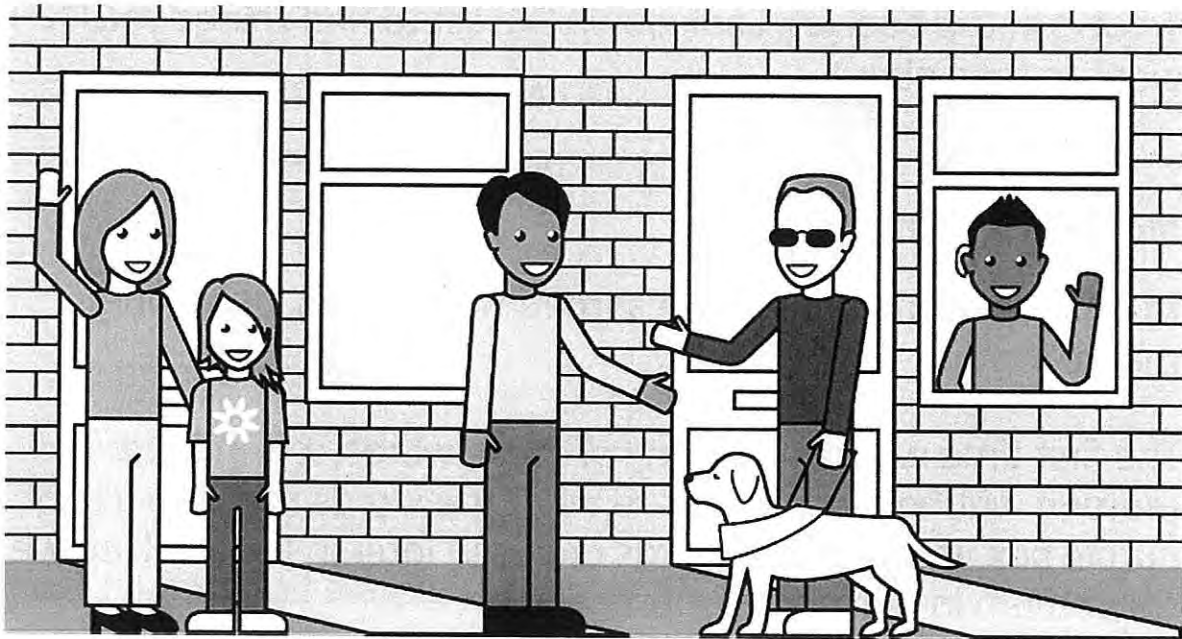


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Consultation about changing Disability Living Allowance to a new benefit

Tell us what you think

December 2010



Easy Read

DWP

Department for
Work and Pensions

Questions

Question 1

What things stop disabled people joining in with other people and living full, active and independent lives?

lack of confidence people looking at
you differently and more scared of going
out on your own.

Question 2

What parts of Disability Living Allowance do you think we should keep?

all of it. it helps to get you lots
of things

Question 3

What extra things do disabled people need to spend money on?

more Heating, more transport, more household items. extra care for household chores + Shaves it back feet

Question 4

The new benefit will have 2 amounts for each of the 2 parts of the benefit. Do you think this will make the benefit easier to understand and also easier for us to run?

Do you think just having the 2 amounts for each part will cause any problems?

No

Question 5

Do you think some health conditions or disabilities should allow people to get an amount of the benefit automatically?

Or do you think that all claims should be based on the needs of the person asking for the benefit?

based on the needs

Question 6

How can we make sure that disabled people who most need the new benefit can get it?

What activities or actions are the most important to live an independent life?

by asking what they need

Question 7

How can we make sure that the new benefit takes into account the way a person's health condition can change?

by ~~changing~~ giving us phone numbers
to the person or family so it
could be changed right away

Question 8

When a person makes a claim to the new benefit, should we take account of any aids or adaptations that they use?

What aids and adaptations should we take into account?

Should we only take aids and adaptations into account if the person already uses them? Or should we take aids and adaptations into account that a person could use and get hold of easily?

all aids + adaptations + ~~but~~
should be taken in to account
right away + in the future
if they need more

Question 9

How could we make the way a person asks for benefit better.
For example

- How could we make the claim form easier to fill in?
- How could we tell people about the new benefit so that they know what the benefit is for and who is likely to get the benefit?

make it short + simple
by making Doctors + hospitals
aware so people can get it
with one leaflet

Question 10

Who are the best people to tell us about the needs of the person asking for benefit?

What information will we need to make it clear what the person can and cannot do?

ask the person or family
hospital or doctor

Question 11

An important part of the new benefit may be talking face to face with an independent person about how well you can do the things you need to do to take part in everyday life.

What good things and bad things may this bring?

Is there any time when it would not be right to say that a person had to meet an independent person face to face, either in the person's own home or somewhere else?

face to face person wood be nice
but make ~~Shure~~ sure the they know
about the persons disability or do
reverse

Question 12

What should we use to decide how often we should look at a claim again and check it?

Should the way we look at a claim again depend on the needs of the person and their health condition or disability?

*Should be on the health and the
disability of the person ~~to~~ and how
often ~~to~~ it changes*

Question 13

The new benefit will be easier for people to understand, so we will expect people to tell us when things change in their lives.

How can we get people to tell us about the changes in their lives?

by phone or letter or meeting

Question 14

What types of help and advice are people who will ask for the new benefit likely to need?

Would it help if we told people to get help and advice and where to get it from?

leaflet ~~to~~ from Doctor or hospital
yes to get help + advice

Question 15

How do disabled people pay for their aids and adaptations at the moment?

Should disabled people be allowed to use the new benefit to pay for a one-off cost?

Some aids + adaptations should be free
Some at low cost

Question 16

What are the main differences we should think about when we are dealing with claims for children instead of adults?

children claims should be made through
hospital or Doctor or ~~the~~ social workers

Question 17

How important or useful has Disability Living Allowance been in getting people to use other services or to get other benefits?

What can we do to make things better?

At the moment people who get Disability Living Allowance automatically get help from other benefits and services, like the Blue Badge scheme and the Warm Front scheme.

What would it mean to disabled people if they did not automatically get help from these other benefits or services?

all people that get disability should
get leaflets on extra help they
can get depending on their age

Question 18

What information about the disabled person could we share with other services or government departments to stop the disabled person having to tell lots of people the same thing?

*make them aware of the Disability
and what it does*

Question 19

How would our ideas for the new benefit affect different equality groups? For example, the equality groups looking at disability, age, race, gender, **sexual orientation** and religion and belief.

Sexual orientation

This is about whether a person is

- heterosexual – sexually attracted to people of the other sex.
- lesbian – a woman who is sexually attracted to women.
- gay – a man who is sexually attracted to men.
- bisexual – sexually attracted to men and women.
- asexual – not sexually attracted to men or women

Should not matter what a person is

Question 20

Is there anything else you would like to tell us about our plans?

*to do your homework on the people
they are dealing with*