

About the questions

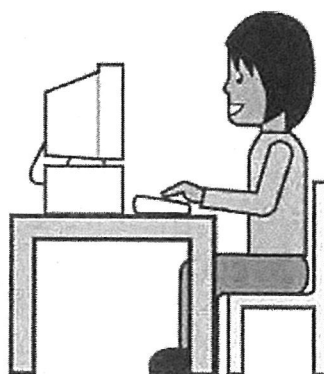
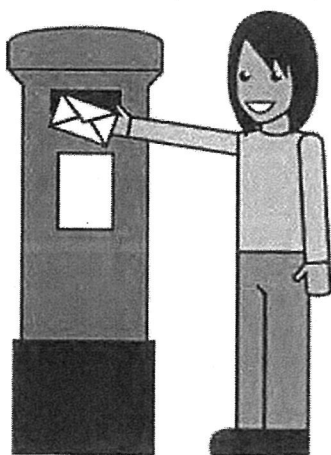
On the next few pages there are questions that we would like you to give us your answers to.

This consultation starts on 6 December 2010 and will end on 14 February 2011.

Send your replies to

DLA Reform Team
1st Floor
Caxton House
Tothill Street
London
SW1H 9NA

Or you can email your reply to
consultation.dlareform@dwp.gsi.gov.uk



And finally, thank you

Thank you for taking the time to look at this booklet and taking part in the consultation.

Questions

Question 1

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

I think by staff stopping them from joining in with other people. Too many carers not letting them live independently.

Question 2

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

We should keep to the lowest rate because that's all I get. I use it for going out and getting clothes and food. At least it gets me out the flat a bit.

Question 3

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

Clothes and help with getting around on public transport, for rail travel - they need a lot of help with that. Not everybody feels comfortable getting out on their own.

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?

I don't think it will be better because it will confuse people.

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?

I think people should get it ~~automatically~~ based on the individual themselves.

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?

You can't really be sure but you can send letters out, some people forget. most important activities to get your own money, food themselves instead of being stopped.

Question 7

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

Go to ~~go~~ the doctors every month to be checked up, because you can get people that fiddle it and get money when you didn't need it.

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?

I think people with aids should get more money because they can't do things for themselves.

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?

To make it easier for people to
fill out so they can do it for
themselves.

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?

citizens advice are the best to ask
about benefits.

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?

I think it's good for the person to get involved with Independent Advocacy because they help me out and can help people more unable than me. Leaves a voice for disabled people & learning disabilities.

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?

see an independent doctor for assessment
on the health of the person.

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?

free phone and to be able to live
independently.

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?

To get advice from the C.A.B.
for the person that's disabled to
go to their doctor.

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?

Should be monthly installments.

Question 16

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

who is looking after the child - their parents

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?

no it should be looked at sepearte from the doctor to see if you need the blue badge because somebody abuse it and use the blue badge.

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?

They should share it with other agencies about their disabilities and proof of a letter from the doctor.

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

no

Question 20

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

Please can I be invited to attend
meetings my address is

my name is