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521

Dear Mr [REDACTED]

I am a 59 year old woman who is now campaigning for Social Justice for people with Autism, Aspergers, ADHD, AOD and other related disorders.

I through personal experience know what the lack of support from an Education System that is full of nepotism and Hypocrisy can do.

In the last two years
I have been a member of
the two Artistic Societies
in my area of Stoke-on
Trent.

These conditions are after
missed by psychologists
as I now know myself.

I have been asked to
lead this, your Disability
Living Allowance reform.

I am very concerned
about how a panel
will be able to diagnose
these conditions when
clearly so called experts
are unable to.

It does not say that
because we can not see

2

a disability it does not exist.

I have not answered the questions because as someone who does not receive any benefits I do not know a lot about them.

I only know what the lack of care does to people.

I feel the whole system is wrong with people (eg) who have children who are "smatic" (like both my brothers are) get vast sums of money when they both smoke.

I do feel there are people using DLA to top up their income.

I am a carer for my

daughter and son who both look normal but having two older children I know they both struggle to cope. My aunt and mother both get carer's allowance.

They are quite worried it will stop as they both need extra support and both pay carer workers.

Myself and my husband both help out with my mother and aunt.

My aunt is unable to walk very well and can not leave the house without she is taken in a car.

My aunt has had a review and they tried to

3

cut her case and also take away her blue badge which is essential for the few trips she takes.

luckily she was allowed to keep it on appeal.

Yours Sincerely


(Campaigner for Social
Justice for people on the
Autistic Spectrum)