



NATIONAL
ORGANISATION
— FOR —
FASD

FASD Policy Matters

What's new?

1



Department
of Health &
Social Care

New from
the
Department
of Health

Fetal Alcohol Spectrum Disorder

Health Needs Assessment for England

Published 9 September 2021

“The government
recognises the
importance
of FASD”

What is a needs assessment?

This identifies the needs of:

- People with FASD
- Their carers and families
- People at risk of alcohol-exposed pregnancies

This means,
no one can
tell you FASD
doesn't matter
ever again!



It highlights priorities for improving healthcare.

1

We need to understand how many people have FASD.
We need better studies.



2

Professionals in health and social care, education, justice and industry need to understand how to support people with FASD **throughout their lives.**



This can help you fight for services when you need them.

That last bit is important. The Department of Health officially recognises that “FASD is a lifelong condition”

Priority needs (continued)

3

Health professionals need to be more aware of FASD.

They need better training about FASD and understand how it's relevant to their work.



They also need to explore their own perspectives on alcohol.

"The neuro-developmental problems associated with FASD can create great difficulties for individuals in their childhood that persist throughout life. Impaired ability to learn, remember, make judgements and forward plan make day-to-day life very challenging."

"There is no 'mild' FASD."

Priority needs (continued)

4

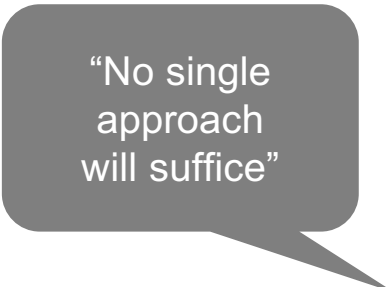
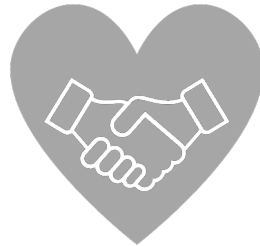
Services need to be organized better to be more accessible for people with FASD and their families.



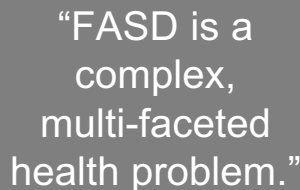
Families can show their local policy makers this and use it to fight for better services.

5

Better ways to support people with FASD are needed.



“No single approach will suffice”



“FASD is a complex, multi-faceted health problem.”

The FASD Health
Needs Assessment
says your input
mattered!

“The voices of people with
lived experience are
important.”



There is a whole section about how important meetings with the Deputy Chief Medical Officer and others were, and how important it was to share our views and research. National FASD was proud to work with other members of the FASD UK Alliance to make this happen.

So what does this mean?

Times are changing!



If someone is denying you access to diagnosis or support, you can tell them doing nothing about FASD is no longer an option.

Tell them the Department of Health and Social Care has identified priority areas for improving care for people with FASD.

Give them this link:

<https://tinyurl.com/FASDNeedsAssessment>

Tell them a NICE Quality Standard on FASD is also coming. It's time to get serious about FASD.



Any questions?

**You can find us at
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