The DiG FASD Study

Visit https://digfasd.org to enroll or learn more.

Questions? Contact us!

844-378-0002
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Find us on Facebook:
FASD Research at Indiana University
www.facebook.com/FASDresearch

Look for us on Twitter:
@FASDResearch

Everyone with FASD is different.

We think genetics plays a role in these differences.

The Dissecting the Genetic Contributions to Fetal Alcohol Spectrum Disorders (DiG FASD) study is researching how genetics explains why people with prenatal alcohol exposure or FASD are different from each other.

Who is Eligible?

- Anyone who had prenatal alcohol exposure or has an FASD diagnosis (parents can sign up their kids).
- You don’t have to know anything about your birth history.
- Adopted people and their families are welcome!
- Anyone who speaks English can participate, from anywhere in the world!

What Happens in the Study?

- You fill out some forms at home, when it is convenient for you.
- You take pictures of your face.
- You provide a saliva (spit) sample for DNA using a kit we send you.
- You will receive a $50 gift card if you complete the study.
- We keep your information, pictures, and DNA secure and confidential and use them for research purposes only.
- We will send you a newsletter with updates on the study’s findings.

Enroll by December 31, 2019 and NOFAS will provide an additional $50 when you complete the study.