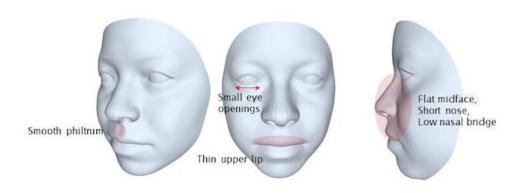


Fetal alcohol syndrome: Facial modeling study explores technology to aid diagnosis

June 8 2023, by Tinashe Ernest Muzvidzwa Mutsvangwa, Bernhard Egger and Felix Atuhaire



Facial regions associated with foetal alcohol syndrome on a normal face. Credit: Tinashe Mutsvangwa

Fetal alcohol syndrome is a lifelong condition <u>caused</u> by exposing an unborn baby to alcohol. It's a pattern of mental, <u>physical</u> and behavioral symptoms seen in some people whose mothers consumed alcohol during pregnancy. Not all prenatal alcohol exposure results in the syndrome; it



is the most severe form of a range of effects called fetal alcohol spectrum disorders.

South Africa has the <u>highest reported rates</u> of fetal alcohol spectrum disorders in the world: 111.1 per 1,000 population. The disorders may affect <u>seven million</u> people in the country. The number could be higher because of under-diagnosis.

Fetal alcohol syndrome can't be reversed. But confirmed diagnosis can have benefits. It can lead to <u>early intervention</u> and therapy (physical, occupational, and speech, among others), and a better <u>understanding</u> from parents and teachers. Diagnosis can also ensure that adults are eligible for social services support.

Clinicians use a range of methods to <u>diagnose fetal alcohol syndrome</u>, including assessing abnormal growth and brain function. A key part of the process is looking at the individual's facial features. Typical <u>features</u> are small eye openings, a thin upper lip, and a smooth area between the nose and upper lip.

But visual examination of the facial features can be subjective and often depends on the clinician's experience and expertise. Another challenge arises in <u>low-resource settings</u> when there aren't many doctors specially trained to do this.

A more objective and standard way to detect fetal alcohol syndrome early would therefore be useful.

One method that's being used to aid diagnosis is <u>three-dimensional (3D)</u> <u>surfaces</u> produced by devices that scan the face. The technology is costly and complex. Two-dimensional (2D) images are easier to get—it can be done with a <u>digital camera</u> or smartphone—but are not accurate enough for diagnosis.



Our study sought to explore whether it was possible to use normal 2D <u>face images</u> to approximate 3D surfaces of the face. We showed that it was. Our method involved using 3D models that can change their shape based on a variety of real human faces, combined with 3D facial analysis technology.

We argue in our paper that our findings show the technology can improve early detection, intervention and treatment for people affected by fetal alcohol syndrome, particularly in low-resource settings.

We hope to contribute to the global effort to prevent and manage the lifelong consequences of the syndrome and disorders.

How it would work

We constructed a <u>flexible 3D model</u> that can alter its shape based on a variety of real human faces. The changes are guided by statistical patterns learned from a <u>dataset of high-quality 3D scans</u> from 98 individuals. This international open-source dataset was carefully curated to represent different demographic groups.

We didn't have access to image data of individuals affected by fetal alcohol syndrome. We therefore used 2D and 3D images of individuals without this condition to develop and validate our approach. We nevertheless reasoned that our method should work equally well for any scenario where the model and the <u>test subjects</u> are closely matched.

We then set out to develop and validate a machine learning algorithm for predicting 3D faces of unseen subjects, from their 2D face images only, using our 3D model.

This was a pioneering step in our research, where we aimed to create a "smart" tool that could bring flat images to life in three dimensions. The



results of the study were encouraging.

Our 3D-from-2D prediction algorithm performed well in three ways:

- capturing facial variations
- representing unique features
- summarizing information of faces from 2D images.

Since we had actual 3D face scans to use for comparison, we were able to calculate the average difference between these scans and the face shapes predicted by our model. This allowed us to measure the error in our fitting, which we found to be in <u>line with other studies</u>.

We particularly focused on specific regions of the face: the eyes, midface, upper lip, and philtrum (the groove between the nose and the top lip). These regions provide crucial information for clinicians when examining the facial markers of fetal alcohol syndrome.

We could accurately predict these facial regions, and concluded from this that our method could form the foundation of an image-based diagnostic tool for fetal alcohol syndrome.

Our study also showed that the quality of our predictions was independent of skin tone. This is a crucial finding. Certain 3D scanning technologies have been known to struggle with accurately capturing darker skin tones. This issue is being addressed. Nevertheless, our findings gave us confidence that there was additional potential for use of our approach in diverse populations.

We did identify some limitations. Access to 3D data of individuals with



fetal <u>alcohol</u> syndrome remains a challenge. Future research could focus on reducing reconstruction errors to acceptable clinical standards by collecting and analyzing larger datasets, including data from underrepresented populations.

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