



I created this piece to bring visibility to an invisible illness; as someone who lives with endometriosis I know first hand how little is known about this chronic condition both in and out of medical spheres, as well as the amount of misinformation about the condition. It affects one in ten people assigned female at birth and as of today has no cure (getting surgery every few years and having to live on painkillers is not a cure). I urge my peers as well as medical professionals to educate themselves on endometriosis from reputable sources that don't spread incorrect information, I suggest following Instagram accounts made by people who live with this condition to start.

