



The **Endo** Educational
Organization of Canada

Resource Library: Extrapelvic Not Rare

“Extrapelvic Not Rare, was created in 2018 by Dr. Wendy Bingham, DPT. It is a non-profit organization focused on grassroots fundraising, independent research and advancement of awareness and education. Our target audience includes healthcare disciplines and general public about endometriosis as a disease with *bodywide potential*. A disease not confined to the female reproductive organs and tissues.” -Extrapelvic Not Rare Website

We support this resource because Extrapelvic Not Rare has done an amazing job at articulating details about endometriosis that are factual, up-to-date and inclusive. In particular, the “Encyclopedia of Extrapelvic Disease” featured on their website offers some important insights into endometriosis that we often don’t see; endometriosis is a systemic disease. Extrapelvic Not Rare can be found on Instagram, Facebook, Twitter, YouTube and the web.

Link to Extrapelvic Not Rare Website
<https://extrapelvicnotrare.org>