

I have a list of web resources:

- The Ehlers-Danlos Society (formerly ednf.org) <https://www.ehlers-danlos.com/>
- EDS Awareness <https://www.chronicpainpartners.com/>
- Ehlers Danlos International Registry <http://www.edsregistry.org/>
- Ehlers-Danlos Syndrome Network C.A.R.E.S. Inc. <http://www.ehlersdanlosnetwork.org/>

And also online support:

- Inspire <https://www.inspire.com/>
- EDS Today~Advocates <https://themighty.com/partner/edstoday/>
- Rare Connect <https://www.rareconnect.org/en/community/ehlers-danlos-syndrome>
- Strength/flexibility/health/EDS <http://strengthflexibilityhealtheds.com/>

Check them out – very helpful information is available!