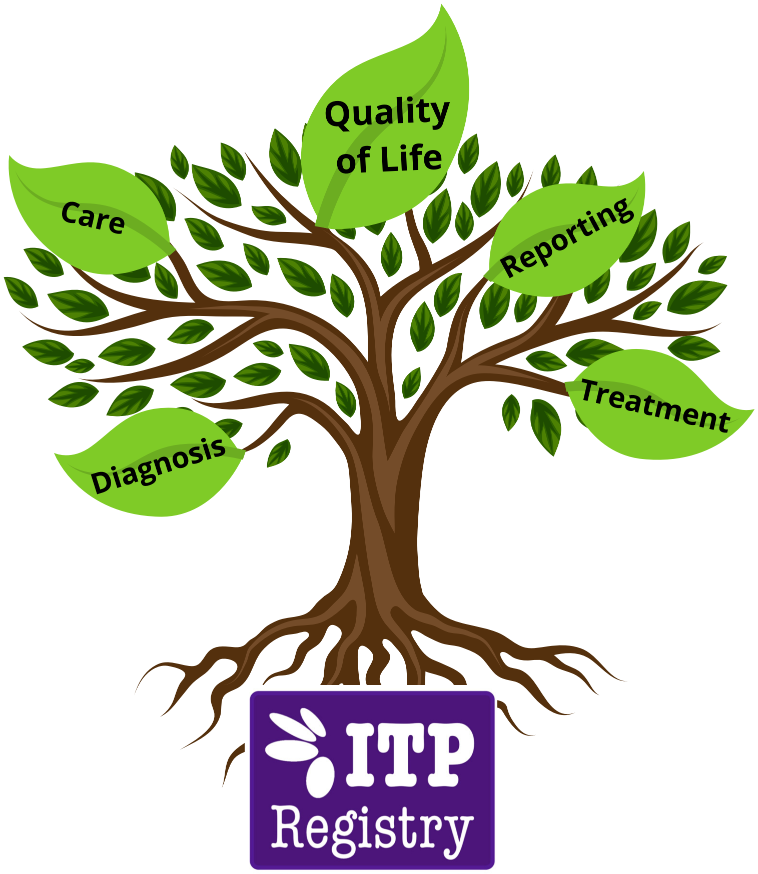
What is the ITP Natural History Study Registry?



The ITP Natural History Study Registry is *an international patient-consented registry of individuals with ITP.*

The registry aims *to collect data on the natural progression of ITP,* enabling PDSA to gather data on ITP diagnosis, treatment, management of care, quality of life, and clinician reporting.

Participation in the ITP Natural History Study is free and voluntary, and participants may withdraw at any time.

Enroll at: **ITPstudy.org**

WHY PARTICIPATE?

Participation is vital given the rarity of ITP - every patient experience is a unique and invaluable part of the natural history of ITP. **Participation in the registry will allow for:**

- Greater Understanding of ITP
- Better Standards of Care
- Understanding of Impact on Quality of Life
- Better Treatments & Clinical Trials
- Cutting-Edge Research
- Patient Voice to be Heard

Platelet Disorder Support Association
Empowering ITP Patients

For further information or to join, please contact: PDSA at 440-746-9003 or email Jennifer DiRaimo, at *jdiraimo@pdsa.org*. The ITP Natural History Study Registry is supported by the National Organization for Rare Disorders (NORD).