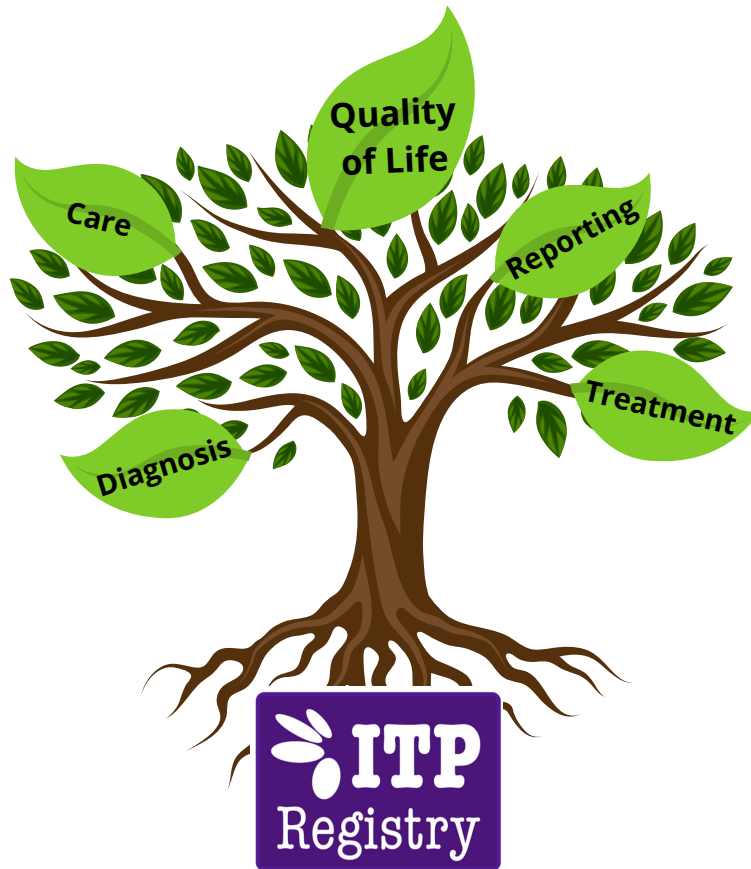


What is the ITP Natural History Study Registry?



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



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