

The Pediatric and Adult Intercontinental Registry on Chronic ITP (PARC-ITP)

The PARC-ITP is a worldwide multi-center registry designed to collect data prospectively of children and adults with newly diagnosed idiopathic thrombocytopenic purpura (ITP). Since chronic ITP of children and adults shows similarities a common database is warranted to coordinate scientific activities in the field of ITP.

Clinical data, bleeding symptoms and management will be observed continuously by long-term follow-up over several years with the goal to recruit as many patients as possible.

You can read the protocol and download all the documents at address www.unibas.ch/itpbase1