

Global CRO, Local Expertise

YOUR TRUSTED PARTNER FOR RARE DISEASES STUDIES

In partnership with leading medical professionals, patient advocates, and cutting-edge technology, we are committed to uncovering solutions for those affected by rare diseases. Backed by years of pioneering work, GCT has forged an unparalleled path in understanding and addressing the unique challenges posed by rare diseases. From innovative patient recruitment strategies and optimization of study endpoints for robust results to a deep understanding of rare disease regulations, - GCT provides services that honor the unique aspects of each condition, aiming to bring transformative solutions to individuals and families facing these challenges. Our CRO's global reach ensures trials are conducted across diverse geographic regions, maximizing patient representation and the generalizability of findings.

From our portfolio

Phase	Indication
III	Trombocytopenic purpura
II	Immune Thrombocytopenic Purpura (ITP)
1/11	Fabry
III	Congenital fibrinogen deficiency
III	Phenylketonuria
III	CDKL5 deficiency disorder (CDD)
II II	Bullous Pemphigoid
II	Controlled Bullous Pemphigoid
I/II	Peripheral T-Cell Lymphoma
III	PCDH19-related epilepsy
Observational	Mitochondrial Neurogastrointestinal Encephalomyopathy
II	Idiopathic Pulmonary Fibrosis (IPF)
11/111	Mucopolysaccharidosis Type II
II	Dravet Syndrome
I/II	Kabuki Syndrome
II	Primary Mitochondrial Myopathy
III	Primary Generalized Tonic-Clonic Seizures
III	Lennox-Gastaut Syndrome (LGS)