

We are conducting a comprehensive study on Coronin 1A deficiency, a rare autosomal recessive immunodeficiency characterized by defects in T-cell development and function. Our goal is to expand the current knowledge of this condition by gathering and analysing more cases.

If you have patients diagnosed with Coronin 1A deficiency and are interested in contributing to this study, we would greatly appreciate your collaboration. Your participation will help us better understand the disease, its clinical spectrum, and genotype-phenotype correlations, ultimately leading to improved patient care and outcomes.

Please contact us at [Dr. Despina Moshous; Despina.moshous@inserm.fr , Dr. Reem Elfeky; reem.elfeky@gosh.nhs.uk] to discuss how you can contribute to this important research.

Thank you for your valuable support.