LUCY RAYMON

Dr Lucy Raymond is Reader in Neurogenetics at the University of Cambridge and Honorary Consultant in Medical Genetics at Addenbrooke's Hospital, Cambridge, UK. Her research interest is understanding the genetic basis of intellectual disability and has identified a number of new disease genes over the years which have been rapidly translated into clinical service. She is a leader in a number of large collaborative efforts to identify the remaining causes of rare disease genes where intellectual disability, epilepsy or neurological phenotypes predominate. She is currently assistant director of the UK NIHR Rare Diseases Bioresource.