Severe Neonatal Encephalopathy Due to MECP2 Mutations

National Cancer Institute

Source

National Cancer Institute. Severe Neonatal Encephalopathy Due to MECP2 Mutations. NCI Thesaurus. Code C132293.

An X-linked recessive condition caused by mutation(s) in the MECP2 gene, encoding methyl-CpG-binding protein 2. It is characterized by severe neonatal encephalopathy.