Gerstmann-Straussler-Scheinker syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Gerstmann-Straussler-Scheinker syndrome. ORPHA:356

Gerstmann-Straussler-Scheinker syndrome (GSSS) is a particular and rare form of human transmissible spongiform encephalopathy (TSE) due to a defective gene encoding the prion protein (PRNP gene) and marked by particular multicentric amyloid plaques in the brain.