

Open Peer Review on Qeios

Familial dysautonomia

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. <u>Familial</u> <u>dysautonomia</u>. ORPHA:1764

Hereditary sensory and autonomic neuropathy, type 3 (HSAN3) is an inherited disorder characterized by sensory dysfunction and severe impairment of the autonomic nervous system activity, resulting in multisystem dysfunction.

Qeios ID: 99M625 · https://doi.org/10.32388/99M625