

Open Peer Review on Qeios

Hennekam syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. <u>Hennekam</u> <u>syndrome</u>. ORPHA:2136

Hennekam syndrome is characterised by the association of lymphoedema, intestinal lymphangiectasia, intellectual deficit and facial dysmorphism.

Qeios ID: BDZLTM · https://doi.org/10.32388/BDZLTM