

Open Peer Review on Qeios

Lelis syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. <u>Lelis</u> <u>syndrome</u>. ORPHA:140936

Lelis syndrome is characterised by the association of ectodermal dysplasia (hypotrichosis and hypohidrosis) with acanthosis nigricans.

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