Open Peer Review on Qeios

Rombo syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. <u>Rombo</u> <u>syndrome</u>. ORPHA:3110

Rombo syndrome is characterized by vermiculate atrophoderma, milia, hypotrichosis, trichoepitheliomas, peripheral vasodilation with cyanosis and basal cell carcinomas.