KID syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. KID syndrome. ORPHA:477

Keratitis (and hystrix-like) ichthyosis deafness (KID/HID) syndrome is a rare congenital ectodermal disorder characterized by vascularizing keratitis, hyperkeratotic skin lesions and hearing loss.