Open Peer Review on Qeios

Fibrodysplasia ossificans progressiva

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Fibrodysplasia ossificans progressiva. ORPHA:337

Fibrodysplasia ossificans progressiva (FOP) is a severely disabling heritable disorder of connective tissue characterized by congenital malformations of the great toes and progressive heterotopic ossification that forms qualitatively normal bone in characteristic extraskeletal sites.