Familial articular hypermobility syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Familial articular hypermobility syndrome. ORPHA:2295

Ehlers-Danlos syndrome, type 11, is characterised by generalized joint hypermobility often complicated by dislocation of major joints, particularly the shoulder but in some cases the kneecap. Congenital hip dislocation has also been frequently reported. The syndrome has been described in several families. It is transmitted as an autosomal dominant trait, with high penetrance.