Brachyolmia

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Brachyolmia. ORPHA:1293

Brachyolmia is a rare, clinically and genetically heterogeneous group of bone disorders characterized by short trunk, mild short stature, scoliosis and generalized platyspondyly without significant abnormalities in the long bones.