

Open Peer Review on Qeios

Osteomesopyknosis

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Osteomesopyknosis. ORPHA:2777

Osteomesopyknosis is a very rare benign bone disorder characterized by bone dysplasia manifested by patchy sclerosis of the axial skeleton and increased bone mineral content.

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