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Mesomelic dysplasia, Kantaputra type

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. <u>Mesomelic</u> <u>dysplasia, Kantaputra type</u>. ORPHA:1836

Mesomelic dysplasia Kantaputra type (MDK) is a rare skeletal disease characterized by symmetric shortening of the middle segments of limbs and short stature.

Qeios ID: E1YQIS · https://doi.org/10.32388/E1YQIS