

Open Peer Review on Qeios

Fuhrmann syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. <u>Fuhrmann</u> <u>syndrome</u>. ORPHA:2854

Fuhrmann syndrome is mainly characterized by bowing of the femora, aplasia or hypoplasia of the fibulae and poly-, oligo-, and syndactyly.

Qeios ID: 7IRMPR · https://doi.org/10.32388/7IRMPR