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Feingold syndrome type 1

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. <u>Feingold</u> <u>syndrome type 1</u>. ORPHA:391641

Feingold syndrome type 1 (FS1) is a rare inherited malformation syndrome characterized by microcephaly, short stature and numerous digital anomalies.

Qeios ID: TRGP30 · https://doi.org/10.32388/TRGP30