Feingold syndrome type 2

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Feingold syndrome type 2. ORPHA:391646

Feingold syndrome type 2 (FS2) is a rare inherited malformation syndrome characterized by skeletal abnormalities and mild intellectual disabilities similar to those seen in Feingold syndrome type 1 (FS1; see this term) but that lacks the manifestations of gastrointestinal atresia and short palpebral fissures.