6q16 microdeletion syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. 6q16 microdeletion syndrome. ORPHA:171829

Deletion 6q16 syndrome is a Prader-Willi like syndrome characterized by obesity, hyperphagia, hypotonia, small hands and feet, eye/vision anomalies, and global developmental delay.