Woodhouse-Sakati syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Woodhouse-Sakati syndrome. ORPHA:3464

Woodhouse-Sakati syndrome is a multisystemic disorder characterized by hypogonadism, alopecia, diabetes mellitus, intellectual deficit and extrapyramidal signs with choreoathetoid movements and dystonia.