

Open Peer Review on Qeios

## Kabuki syndrome

**INSERM** 

## Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. <u>Kabuki syndrome</u>. ORPHA:2322

Kabuki syndrome (KS) is a multiple congenital anomaly syndrome characterized by typical facial features, skeletal anomalies, mild to moderate intellectual disability and postnatal growth deficiency.

Qeios ID: M8K9OS · https://doi.org/10.32388/M8K9OS