Open Peer Review on Qeios

Muenke Syndrome

National Cancer Institute

Source

National Cancer Institute. <u>Muenke Syndrome</u>, NCI Thesaurus. Code C84904.

A rare autosomal dominant inherited disorder caused by mutations in the FGFR3 gene. It is characterized by premature fusion of cranial bones, resulting in head shape abnormalities, flattened cheekbones, and wide-set eyes.