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# Galloway-Mowat Syndrome

National Cancer Institute

## Source

National Cancer Institute. *Galloway-Mowat Syndrome*. NCI Thesaurus. Code C132195.

An autosomal recessive neurodegenerative condition caused by mutation(s) in the WDR73 gene, encoding WD repeat-containing protein 73. It is characterized by microcephaly and severely delayed psychomotor development.