

RAB18 gene

RAB18, member RAS oncogene family

Normal Function

The *RAB18* gene provides instructions for producing the RAB18 protein, which functions as a GTPase. Often referred to as molecular switches, GTPases can be turned on and off. They are turned on (active) when they are attached (bound) to a molecule called GTP and are turned off (inactive) when they are bound to another molecule called GDP. When active, RAB18 is involved in a process called vesicle trafficking, which moves proteins and other molecules within cells in sac-like structures called vesicles. RAB18 regulates the movement of substances between compartments in cells and the storage and release of fats (lipids) by structures called lipid droplets. The protein also appears to play a role in a process called autophagy, which helps clear unneeded materials from cells. RAB18 is important for the organization of a cell structure called the endoplasmic reticulum, which is involved in protein processing and transport.

Health Conditions Related to Genetic Changes

RAB18 deficiency

At least five mutations in the *RAB18* gene have been found to cause Warburg micro syndrome, which is the most severe of the disorders caused by RAB18 deficiency. Warburg micro syndrome is characterized by multiple eye abnormalities, vision impairment, severe intellectual disability, and a reduction of the hormones that direct sexual development (hypogonadotropic hypogonadism).

The *RAB18* gene mutations that cause Warburg micro syndrome eliminate the function of the RAB18 protein. It is unclear how a shortage (deficiency) of RAB18 activity leads to eye problems, brain abnormalities, and other features of Warburg micro syndrome.

Other Names for This Gene

- RAB18 small GTPase
- RAB18LI1
- ras-related protein Rab-18 isoform 1
- ras-related protein Rab-18 isoform 2

- ras-related protein Rab-18 isoform 3
- ras-related protein Rab-18 isoform 4
- ras-related protein Rab-18 isoform 5
- WARBM3

Additional Information & Resources

Tests Listed in the Genetic Testing Registry

- Tests of RAB18 ([https://www.ncbi.nlm.nih.gov/gtr/all/tests/?term=22931\[geneid\]](https://www.ncbi.nlm.nih.gov/gtr/all/tests/?term=22931[geneid]))

Scientific Articles on PubMed

- PubMed (<https://pubmed.ncbi.nlm.nih.gov/?term=%28RAB18%5BTIAB%5D%29+OR+%28RAB18+small+GTPase%5BTIAB%5D%29+AND+%28%28Genes%5BMH%5D%29+OR+%28Genetic+Phenomena%5BMH%5D%29%29+AND+english%5Bla%5D+AND+human%5Bmh%5D+AND+%22last+2880+days%22%5Bdp%5D%29>)

Catalog of Genes and Diseases from OMIM

- RAS-ASSOCIATED PROTEIN RAB18; RAB18 (<https://omim.org/entry/602207>)

Gene and Variant Databases

- NCBI Gene (<https://www.ncbi.nlm.nih.gov/gene/22931>)
- ClinVar ([https://www.ncbi.nlm.nih.gov/clinvar?term=RAB18\[gene\]](https://www.ncbi.nlm.nih.gov/clinvar?term=RAB18[gene]))

References

- Bem D, Yoshimura S, Nunes-Bastos R, Bond FC, Kurian MA, Rahman F, Handley MT, Hadzhiev Y, Masood I, Straatman-Iwanowska AA, Cullinane AR, McNeill A, Pasha SS, Kirby GA, Foster K, Ahmed Z, Morton JE, Williams D, Graham JM, Dobyns WB, Burglen L, Ainsworth JR, Gissen P, Muller F, Maher ER, Barr FA, Aligianis IA. Loss-of-function mutations in RAB18 cause Warburg micro syndrome. *Am J Hum Genet.* 2011 Apr 8;88(4):499-507. doi: 10.1016/j.ajhg.2011.03.012. Citation on PubMed (<https://pubmed.ncbi.nlm.nih.gov/21473985>) or Free article on PubMed Central (<https://www.ncbi.nlm.nih.gov/pmc/articles/PMC3071920/>)
- Feldmann A, Bekbulat F, Huesmann H, Ulbrich S, Tatzelt J, Behl C, Kern A. The RAB GTPase RAB18 modulates macroautophagy and proteostasis. *Biochem Biophys Res Commun.* 2017 May 6;486(3):738-743. doi: 10.1016/j.bbrc.2017.03.112. Epub 2017 Mar 22. Citation on PubMed (<https://pubmed.ncbi.nlm.nih.gov/28342870>)
- Gerondopoulos A, Bastos RN, Yoshimura S, Anderson R, Carpanini S, Aligianis I,

Handley MT, Barr FA. Rab18 and a Rab18 GEF complex are required for normal ERstructure. J Cell Biol. 2014 Jun 9;205(5):707-20. doi: 10.1083/jcb.201403026. Epub 2014 Jun 2. Citation on PubMed (<https://pubmed.ncbi.nlm.nih.gov/24891604>) or Free article on PubMed Central (<https://www.ncbi.nlm.nih.gov/pmc/articles/PMC4050724/>)

- Handley MT, Carpanini SM, Mali GR, Sidjanin DJ, Aligianis IA, Jackson IJ, FitzPatrick DR. Warburg Micro syndrome is caused by RAB18 deficiency ordysregulation. Open Biol. 2015 Jun;5(6):150047. doi: 10.1098/rsob.150047. Citation on PubMed (<https://pubmed.ncbi.nlm.nih.gov/26063829>) or Free article on PubMed Central (<https://www.ncbi.nlm.nih.gov/pmc/articles/PMC4632505/>)

Genomic Location

The *RAB18* gene is found on chromosome 10 (<https://medlineplus.gov/genetics/chromosome/10/>).

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