CDKN2A gene
cyclin dependent kinase inhibitor 2A

Normal Function

The \textit{CDKN2A} gene provides instructions for making several proteins. The most well-studied are the p16(INK4A) and the p14(ARF) proteins. Both function as tumor suppressors, which means they keep cells from growing and dividing too rapidly or in an uncontrolled way. Both proteins are also involved in stopping cell division in older cells (senescence).

The p16(INK4A) protein attaches (binds) to two other proteins called CDK4 and CDK6. These proteins help regulate the cell cycle, which is the cell's way of replicating itself in an organized, step-by-step fashion. CDK4 and CDK6 normally stimulate the cell to continue through the cycle and divide. However, binding of p16(INK4A) blocks CDK4's or CDK6's ability to stimulate cell cycle progression. In this way, p16(INK4A) controls cell division. Cells begin to produce p16(INK4A) when they are no longer able to undergo cell division.

The p14(ARF) protein protects a different protein called p53 from being broken down. The p53 protein is an important tumor suppressor that is essential for regulating cell division, senescence, and self-destruction (apoptosis). By protecting p53, p14(ARF) also helps prevent tumor formation. The p14(ARF) and p53 proteins are often made in cells that are unable to undergo cell division.

Health Conditions Related to Genetic Changes

Head and neck squamous cell carcinoma

Mutations in the \textit{CDKN2A} gene are found in up to one-quarter of head and neck squamous cell carcinomas (HNSCC). This type of cancerous tumor occurs in the moist lining of the mouth, nose, and throat. \textit{CDKN2A} gene mutations associated with this condition are acquired during a person's lifetime and are found only in tumor cells; these changes are known as somatic mutations. Most of these mutations lead to production of little or no functional p16(INK4A) protein. Without p16(INK4A) to regulate cell growth and division (proliferation), cells can continue to grow and divide without control, which can lead to tumor formation.

A different type of alteration involving the \textit{CDKN2A} gene can result in reduced amounts or an absence of the p16(INK4A) or p14(ARF) protein. This alteration, known as promoter hypermethylation, turns off the production of p16(INK4A) or p14(ARF). Without one of these tumor suppressors, cells can grow and divide unchecked, leading to the development of cancer.
Lung cancer

Melanoma

Mutations in the CDKN2A gene are also associated with melanoma, a type of skin cancer that begins in pigment-producing cells called melanocytes. CDKN2A gene mutations are found in up to 40 percent of familial cases of melanoma, in which multiple family members develop the cancer. These mutations, classified as germline mutations, are typically inherited and are present in essentially all of the body’s cells. The CDKN2A gene mutations found in melanoma result in a nonfunctional p16(INK4A) protein. In many cases, a second, somatic mutation occurs in the normal copy of the gene in melanocytes. In about half of melanomas, part or all of the CDKN2A gene is missing (deleted). In many other cases, the CDKN2A gene has a mutation or is turned off (inactive). Somatic mutations in other genes involved in cell growth are also needed for a melanoma to develop. Together, the germline and somatic mutations impair the function of proteins that regulate division and senescence, leading to uncontrolled cell growth and the formation of a melanoma.

Individuals with a CDKN2A gene mutation tend to develop melanoma at an earlier age than those without a mutation in the gene. They also tend to develop other cancers during their lifetime, particularly cancers of the pancreas or lung.

Other cancers

Germline mutations affecting the CDKN2A gene are associated with other cancers, including breast cancer and pancreatic cancer. In some families, CDKN2A gene mutations are associated with development of only one type of cancer. In other families, mutations can lead to a cancer predisposition syndrome, which increases the risk of developing multiple types of cancer. CDKN2A gene mutations involved in cancer impair production of functional p16(INK4A) or, less commonly, p14(ARF), which can result in uncontrolled cell growth and tumor formation.

Somatic CDKN2A gene mutations have been found in some people with brain tumors and in children with a blood cancer called acute lymphoblastic leukemia.
Chromosomal Location

Cytogenetic Location: 9p21.3, which is the short (p) arm of chromosome 9 at position 21.3

Molecular Location: base pairs 21,967,752 to 21,995,324 on chromosome 9 (Homo sapiens Updated Annotation Release 109.20190607, GRCh38.p13) (NCBI)

Credit: Genome Decoration Page/NCBI

Other Names for This Gene

- ARF
- CDK4 inhibitor p16-INK4
- CDK4I
- CDKN2
- cell cycle negative regulator beta
- CMM2
- cyclin-dependent kinase 4 inhibitor A
- cyclin-dependent kinase inhibitor 2A
- cyclin-dependent kinase inhibitor 2A (melanoma, p16, inhibits CDK4)
- cyclin-dependent kinase inhibitor 2A isoform p12
- cyclin-dependent kinase inhibitor 2A isoform p14ARF
- cyclin-dependent kinase inhibitor 2A isoform p16gamma
- cyclin-dependent kinase inhibitor 2A isoform p16INK4a
- INK4
- INK4A
- MLM
- MTS-1
- MTS1
• multiple tumor suppressor 1
• P14
• P14ARF
• P16
• P16-INK4A
• P16INK4
• P16INK4A
• P19
• P19ARF
• TP16

Additional Information & Resources

Educational Resources
• Molecular Cell Biology (fourth edition, 2000): Loss-of-Function Mutations in Tumor-Suppressor Genes Are Oncogenic
  https://www.ncbi.nlm.nih.gov/books/NBK21662/#_A7100_

Scientific Articles on PubMed
• PubMed
  https://www.ncbi.nlm.nih.gov/pubmed?term=%28%28CDKN2A%5BTI%5D%29+OR+%28cyclin-dependent+kinase+inhibitor+2A%5BTI%5D%29%29+AND+%28%28Genes%5BMH%5D+OR+%28Genetic+Phenomena%5BMH%5D%29+AND+english%5Bla%5D+AND+human%5Bmh%5D+AND+%22last+720+days%22%5Bdp%5D

Catalog of Genes and Diseases from OMIM
• CYCLIN-DEPENDENT KINASE INHIBITOR 2A
  http://omim.org/entry/600160
• MELANOMA-ASTROCYTOMA SYNDROME
  http://omim.org/entry/155755
• MELANOMA-PANCREATIC CANCER SYNDROME
  http://omim.org/entry/606719
• PANCREATIC CANCER
  http://omim.org/entry/260350
Research Resources

- Atlas of Genetics and Cytogenetics in Oncology and Haematology
  http://atlasgeneticsoncology.org/Genes/CDKN2aID146.html
- ClinVar
- HGNC Gene Symbol Report
- Monarch Initiative
  https://monarchinitiative.org/gene/NCBIGene:1029
- NCBI Gene
- UniProt: ARF_HUMAN
  https://www.uniprot.org/uniprot/Q8N726
- UniProt: CDN2A_HUMAN
  https://www.uniprot.org/uniprot/P42771

Sources for This Summary

- Baker DJ, Childs BG, Durik M, Wijers ME, Sieben CJ, Zhong J, Saltness RA, Jeganathan KB,
  Verzosa GC, Pezeshki A, Khazaie K, Miller JD, van Deursen JM. Naturally occurring p16(Ink4a)-
nature16932. Epub 2016 Feb 3.
  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/26840489
  Free article on PubMed Central: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC4845101/

- Cánepa ET, Scassa ME, Ceruti JM, Marazita MC, Carcagno AL, Sirkin PF, Ogara MF. INK4
  proteins, a family of mammalian CDK inhibitors with novel biological functions. IUBMB Life. 2007
  Jul;59(7):419-26. Review.
  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/17654117

- Goldstein AM, Chan M, Harland M, Gillanders EM, Hayward NK, Avril MF, Azizi E, Bianchi-Scarra
  DE, Ghiorzo P, Gruis NA, Hansson J, Hogg D, Holland EA, Kanetsky PA, Kefford RF, Landi MT,
  Lang J, Leachman SA, Mackie RM, Magnusson V, Mann GJ, Niendorf K, Newton Bishop J, Palmer
  JM, Puig S, Puig-Butille JA, de Snoo FA, Stark M, Tsao H, Tucker MA, Whitaker L, Yakobson
  E; Melanoma Genetics Consortium (GenoMEL). High-risk melanoma susceptibility genes and
  pancreatic cancer, neural system tumors, and uveal melanoma across GenoMEL. Cancer Res.
  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/17047042

- He S, Sharpless NE. Senescence in Health and Disease. Cell. 2017 Jun 1;169(6):1000-1011. doi:
  10.1016/j.cell.2017.05.015. Review.
  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/28575665
  Free article on PubMed Central: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC5643029/
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  Free article on PubMed Central: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC3715072/

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