

What you should know about Hereditary Melanoma

Most melanoma is not inherited; only 10% of all melanomas are thought to be hereditary. Some cases of hereditary melanomas (approximately 20-57%) are associated with a mutation in the *CDKN2* gene (also called p16). 72-81% of families with hereditary melanomas and pancreatic cancer will have a *CDKN2* mutation that can be identified. Other as yet unidentified genes account for other hereditary melanoma cases.

The risk for cancer associated with *CDKN2* mutations

- Mutations in the *CDKN2* gene are associated with up to a 76% risk for melanoma by age 80.
- Individuals with a *CDKN2* mutation who have had melanoma have an increased risk for developing a second melanoma.
- *CDKN2* mutations are also associated with an increased risk (up to 17% by age 75) for pancreatic cancer in some families.

The risks to family members

CDKN2 mutations are inherited in an autosomal dominant fashion. The children, brothers, sisters, and parents of individuals with *CDKN2* mutations have a 50% (1 in 2) risk to have the mutation. Individuals with a *CDKN2* mutation may develop one cancer, more than one cancer, or none at all.

Managing the risk

Increased Surveillance

- Regular self-examinations of the skin
- Clinical skin examinations every 6-12 months, beginning in childhood
- Baseline photography of the entire body by a dermatologist

Preventative surgery

- Biopsy and/or removal of suspicious moles

Lifestyle modifications

- Limit sun exposure and avoid tanning beds
- Wear protective clothing
- Apply sunscreen with SPF of 15 or higher at regular intervals
- Avoid smoking, as smoking significantly increases the risk for pancreatic cancer in individuals with *CDKN2* mutations.