Genetic conditions:



Healthcare provider card

Summary of family health history

I am concerned about my family history of (check all that apply):

Heart disease or heart attack Breast cancer Stroke Ovarian cancer Sudden death/ruptured vessels Endometrial (uterine) cancer Bleeding/clotting disorders Colon cancer High blood pressure Other cancer: ___ High cholesterol Mental health: _____ Diabetes/sugar disease Alzheimer's disease/dementia Obesity Mental retardation/developmental delay Asthma Speech delay Hearing loss as child/young adult Birth defects Vision loss as child/young adult Miscarriage/stillbirth

Identify family members with each condition checked, including age of diagnosis, current age or age at death and cause of death (use extra pages if needed).

Relationship	Condition	Age of onset	Current age	Age, cause of death
Example:				
	High blood			
Brother	pressure	35		
	High blood			
Mother	pressure	40	45	65, stroke

Be sure to include information about your children, siblings, mother and father, and remember to include relatives from both sides of the family.

After you complete this paper, bring it to your primary care provider.



Resources for the provider

Recognizing family risk (genetic red flags)

Family history of known genetic disorder

Multiple affected family members with same or related disorders

Earlier age at onset of disease than expected

- o Breast, ovarian and endometrial cancer <50 years (premenopausal)
- o Colon and prostate cancer <50 years
- o Stroke and non-insulin-dependent diabetes <50 years
- o Dementia < 60 years
- o Coronary artery disease <55 years in males and <65 years in females

Sudden cardiac death in a person who seemed healthy

Multifocal or bilateral occurrence in paired organs

Ethnic predisposition to certain genetic disorders

General guidelines for risk stratification

High risk:

- 1. Premature disease in a first-degree relative (sibling, parent or child)
- 2. Premature disease in a second-degree relative (CAD only)
- 3. Two affected first-degree relatives
- 4. One first-degree relative with late or unknown disease onset and an affected second-degree relative with premature disease from the same lineage
- 5. Two second-degree maternal or paternal relatives with at least one having premature onset of disease
- 6. Three or more affected maternal or paternal relatives
- 7. Presence of a "moderate risk" family history on both sides of the pedigree

Moderate risk:

- 1. One first-degree relative with late or unknown onset of disease
- 2. Two second-degree relatives from the same lineage with late or unknown disease onset

Average risk:

- 1. No affected relatives
- 2. Only one affected second-degree relative from one or both sides of the pedigree
- 3. No known family history
- 4. Adopted person with unknown family history

Scheuner et al., Am J Med Genet 1997; 71:315-324







Family history website resources

- 1. CDC cdc.gov/genomics/famhistory/famhist.htm
- 2. AAFP Genomics CME aafp.org
- 3. U.S. Surgeon General hhs.gov/familyhistory
- 4. Genetic Alliance Genetic Alliance.org/fhh
- 5. Recent literature GeneticAlliance.org/fhh.literature

Have questions? Contact us at familyhistory@geisinger.edu or 570-214-6065.











