The Genetic Information Nondiscrimination Act

The Genetic Information Nondiscrimination Act (GINA) is a Federal law that prohibits discrimination in health coverage and employment based on genetic information.

This Federal law generally makes it illegal for health insurance companies, group health plans, and most employers to discriminate against you based on your genetic information. All health insurance companies and group health plans must follow this law by May 21, 2010. All employers with 15 or more employees must follow this law as of November 21, 2009.

This law will generally protect you in the following ways:

• Health insurance companies and group health plans may not request your genetic information that is obtained from research.

• Health insurance companies or health plan administrators engaged in research may not use the information obtained to discriminate against you.

• Health insurance companies and group health plans may not use your genetic information when making decisions regarding your eligibility or premiums.

• Employers with 15 or more employees may not use your genetic information from research when making a decision to hire, promote, or fire you or when setting the terms of your employment.

Limitations (The following are not all inclusive, but the main limitations of GINA):

• The law does not exclude life insurance companies from using genetic information to make decisions.

• The law does not protect an individual if they already have a disease. It only protects an individual that has a genetic predisposition to a disease.

• Be aware that this new Federal law does not protect you against genetic discrimination by companies that sell life insurance, disability insurance, or long-term care insurance.

For more detailed information from the Department of Health and Human Services (HHS) of GINA see: [http://www.genome.gov/Pages/PolicyEthics/GeneticDiscrimination/GINAInfoDoc.pdf](http://www.genome.gov/Pages/PolicyEthics/GeneticDiscrimination/GINAInfoDoc.pdf)

Just as in any research project, participation in genetic research is voluntary. It is your choice whether or not to participate or withdraw at any time.