

## Genetic Testing Discussion

1. Would you want the Alzheimer's or Parkinson's Disease information from 23andMe? Would you want the results of an incidental finding in another genetic test? What kind of genetic information do you want?
2. Who should you tell about this information? Who do you not want to know about this? Who should your physician tell about this? What were the reasons these people were on these lists?
3. What are the responsibilities of genetic testing companies? To whom do they have a responsibility?
4. Are the policies on genetic testing of children in conflict? In what ways are they in what ways not?
5. In these policies on genetic testing, who are the stakeholders? How are their needs, desires, and responsibilities balanced?