Chapter 1 project: Exploration questions

DUE) 11:00am Wed., January 23

Student: \_\_\_\_\_

Partner (if any): \_\_\_\_\_

Part I: Regarding SNP rs34637584

- 1) On which human chromosome is this SNP located and at what position?
- 2) Does rs34637584 occur within a gene or between genes? If it occurs within a gene, is it within an exon or an intron?
- 3) Describe the Alleles that occur through variation at this site?
- 4) List the genes found within approximately 500kb on either side of the SNP
- 5) Has this site or any of the nearby genes been associated with PD previously?
- 6) What evidence did you find to support the identification of one or more of the genes in this region as a candidate for PD-associated gene?

## Part II: Regarding SREBF1 gene

- 7) How long is the entire SREBF1 gene?
- 8) When you click on the CDS link in a GenBank entry, only short segments of the gene sequence are highlighted. What do these highlighted segments represent?
- 9) How long is the spliced mRNA for SREBF1? What fraction of the gene is thrown away in the form of spliced-out introns?
- 10) What accounts for the difference between the sequence segments that are highlighted when you click on the mRNA link versus when you click on the CDS link?
- 11) How long is the SREBF1 protein in amino acids? What are the first 10 amino acids in the protein sequence?
- 12) What is known about the function of this gene?
- 13) Besides its hypothetical associate with PD, what are two other known connections of SREBF1 to disease?