## **Genetic Disorders Presentation**

<b>Genetic Disorder</b> :	Due:	
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You will present on a genetic disorder using PowerPoint. Your PowerPoint should serve as a visual <u>aid</u> to your oral presentation. The PowerPoint should not contain long sentences, paragraphs, or irrelevant images and information; it should be neat, organized and concise. Remember the PowerPoint slides should be used as a <u>supplement</u> to your presentation not be the main part of the presentation. You must show that you <u>know</u> and <u>understand</u> the information being presented and not merely read off the slides. You may include images and diagrams that can enhance your oral presentation and may also include a relevant video on the disorder. All of these aspects are part of your grade. This project will be graded as a test grade.

Your presentation must be written in your own words and include each of the following:

- 1. The name of your disease.
- 2. Describe the genetics of the disorder. For example:
  - a. Is it a gene mutation, a chromosomal mutation, or a result of non-disjunction?
  - b. State the chromosome number (autosome) that is affected or state if it is sex-linked (on a sex chromosome).
- 3. How many people are affected by the disorder?
- 4. Phenotypic characteristic: What are the phenotypic characteristics of someone with this disorder? If there are no phenotypic features, state that there are none.
- 5. What are the symptoms and complications?
- 6. What is the life expectancy of an individual with the disorder?
- 7. Are current treatments? If yes, how effective are they? If there are no current treatments include a description of what type of research is being done to find a treatment.
- 8. What are the support groups that exist for this disorder? List at least one support organization and describe what support they provide.
- 9. Bibliography. A minimum of three sources are required from the library (catalog/databases, e-books, news articles, and/or journals).

## **Recommended Sources**

Gulliver Library: CREDO Reference

National Institutes of Health: <a href="http://www.nih.gov/">http://www.nih.gov/</a> Genetics Home Reference: <a href="http://ghr.nlm.nih.gov/">http://ghr.nlm.nih.gov/</a>

Mayo Clinic: http://www.mayoclinic.com/

Duke Center for Human Genetics: http://www.chg.duke.edu/index.html

Merck Manual: http://www.merckmanuals.com/home/index.html

## Genetic Disorders List

Achondroplasia

Marfan Syndrome

Alkaptonuria Neurofibromatosis

Cystic Fibrosis Oculocutaneous Albinism

(Type I or II)

Ehlers—Danlos Syndrome
Classical (Type I or II)
Osteogenesis Imperfecta

Fragile X Syndrome Progeria

Hemophilia A Sickle Cell Disease

Huntington's Disease Tay-Sachs Disease

Klinefelter Syndrome Trisomy 21

Lou Gehrig's Disease (ALS)

Trisomy X

Color Blindness Turner's Syndrome