WS 7-3: Human Genetic Disorders

| Name | <u>Key</u> | |
|------|---|--------------------------------|
| | ine provided, write the letter of the term from the list an be used once, more than once, or not at all. | that matches each description. |
| c_ | 1. disorder that causes a rapid breakdown of the nervous system beginning at age 2 or 3 | a. albinism |
| _ | , , , | b. cystic fibrosis |
| F | 2. process that takes place when a c'some pair fail to separate correctly during meiosis | c. Tay-Sachs disease |
| H | _3. form of trisomy in which there is an extra copy of c'some 21 | d. phenylketonuria |
| Α | 4. disorder that results in lack of pigment in hair or skin | e. Huntington disease |
| ^_ | | f. nondisjunction |
| B | 5. disorder caused by a recessive allele on c'some 7 that results in fluid buildup in the lungs6. Condition that exists when an individual is born with cells that contain 3 copies of a c'some | g. trisomy |
| | | h. Down syndrome |
| G | | i. neurofibromatosis |
| D | 7. genetic disorder known as PKU, for which newborn infants in PA are tested | j. fragile-X syndrome |
| | | k. deletion |
| K | _8. process that occurs during meiosis when pieces of c'somes break off and are lost | |
| E | 9. disorder for which symptoms typically don't appear until late 30s or 40s | |
| c_ | _10. nervous system disorder that is most prevalent in Jewish and French Canadian populations | |
| D | _11. metabolic disease that if untreated can damage the nervous system | |
| | 12. disease in which a skin spots may develop into tumors | |
| J | 13. disease that occurs mostly in males; when part of the X c'come may be deleted | |