

WS 7-3: Human Genetic Disorders

Name _____ Key _____

On the line provided, write the letter of the term from the list that matches each description. Some can be used once, more than once, or not at all.

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| ___ C ___ 1. disorder that causes a rapid breakdown of the nervous system beginning at age 2 or 3 | a. albinism |
| ___ F ___ 2. process that takes place when a c'some pair fail to separate correctly during meiosis | b. cystic fibrosis |
| ___ H ___ 3. form of trisomy in which there is an extra copy of c'some 21 | c. Tay-Sachs disease |
| ___ A ___ 4. disorder that results in lack of pigment in hair or skin | d. phenylketonuria |
| ___ B ___ 5. disorder caused by a recessive allele on c'some 7 that results in fluid buildup in the lungs | e. Huntington disease |
| ___ G ___ 6. Condition that exists when an individual is born with cells that contain 3 copies of a c'some | f. nondisjunction |
| ___ D ___ 7. genetic disorder known as PKU, for which newborn infants in PA are tested | g. trisomy |
| ___ K ___ 8. process that occurs during meiosis when pieces of c'somes break off and are lost | h. Down syndrome |
| ___ E ___ 9. disorder for which symptoms typically don't appear until late 30s or 40s | i. neurofibromatosis |
| ___ C ___ 10. nervous system disorder that is most prevalent in Jewish and French Canadian populations | j. fragile-X syndrome |
| ___ D ___ 11. metabolic disease that if untreated can damage the nervous system | k. deletion |
| ___ I ___ 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 | |