Date: \_\_\_\_\_

## Genetics

| 1. Trisomy 21 Down Syndrome occurs due to  | A. Oogenesis           |
|--|------------------------|
| 2. This pattern of inheritance requires both parents to be a carrier for the child to be affected              | B. Down syndrome       |
| 3. When looking at the family history, there is at least one affected person in each generation                | C. DNA recombination   |
| 4. Formation of male gamete in the seminiferous tubules  | D. Non-disjunction     |
| 5. Formation of female gamete, ovum, in the ovaries  | E. Spermatogenesis     |
| 6. This is an example of a congenital abnormality caused by both genetic and environmental factors play a role | F. Spina bifida        |
| 7. Common chromosomal disorder   | G. Autosomal dominant  |
| 8. A single gene disorder  | H. Achondroplasia      |
| 9. Exchange of genetic material between two chromosomes  | I. Mitosis             |
| 10. This process results in two identical daughter cells   | J. Autosomal Recessive |