

Genetic Mind Reader Card Games

<b>Concept</b>	<b>Answer</b>
<b>1. A chromosome map of a species that shows the specific physical location of its genes and/or markers on each chromosome</b>	<b>1. Physical map</b>
<b>2. A chromosome map of a species that shows the position of its known genes and/or markers relative to each other</b>	<b>2. Genetic map</b>
<b>3. Number of DNA base pairs in the bacterium <i>Escherichia coli</i></b>	<b>3. 4.6 million</b>
<b>4. Number of DNA base pairs in bakers' yeast (<i>Saccharomyces cerevisiae</i>)</b>	<b>4. 12 million</b>

<b>5. Number of DNA base pairs in the roundworm (<i>Caenorhabditis elegans</i>)</b>	<b>5. 97 million</b>
<b>6. Number of DNA base pairs in the fruit fly (<i>Drosophila melanogaster</i>)</b>	<b>6. 165 million</b>
<b>7. Number of DNA base pairs in the mouse (<i>Mus musculus</i>)</b>	<b>7. 3 billion</b>
<b>8. Number of chromosomes in a normal human cell</b>	<b>8. 23 pairs of chromosomes</b>

<b>9. Segments of DNA; most contain information for making a specific protein</b>	<b>9. Genes</b>
<b>10. The form in which genes are passed from parent to offspring</b>	<b>10. A chromosome</b>
<b>11. A large complex molecule made up of one or more chains of amino acids. This molecule can perform a wide variety of activities in the cell</b>	<b>11. A protein</b>
<b>12. The names of the bases that, as part of nucleotides, make up the ladder of DNA</b>	<b>12. (A) Adenine, (G) Guanine, (C) Cytosine, and (T) Thymine</b>

<b>13. A blood disorder caused by a single base pair change in one of the genes that code for hemoglobin, the blood protein that carries oxygen</b>	<b>13. Sickle cell anemia</b>
<b>14. Number of DNA base pairs in barley (<i>Hordeum vulgare</i>)</b>	<b>14. 4.8 billion</b>
<b>15. Complex diseases such as diabetes and Alzheimer's</b>	<b>15. Disorders that arise from mutations in multiple genes</b>
<b>16. This contains 6 feet of DNA</b>	<b>16. The nucleus of the human cell</b>

<b>17. These cells contain single sets of chromosomes</b>	<b>17. Egg and sperm cells</b>
<b>18. Knowing the DNA sequence of a gene</b>	<b>18. Reveals the amino acid sequence of the protein that the gene encodes</b>
<b>19. Using healthy genes to treat genetic diseases</b>	<b>19. Gene therapy</b>
<b>20. Knowing an individual's DNA sequence</b>	<b>20. Can predict the risk of developing a condition like heart disease, diabetes, or prostate cancer later in life</b>

<b>21. A person who predicts the likelihood that a couple's hidden mutations will be passed on to future generations</b>	<b>21. Genetic counselor</b>
<b>22. A person who uses altered genes to increase the nutritional value of various foods and dairy products</b>	<b>22. A genetic engineer</b>
<b>23. Huntington's disease, cystic fibrosis, and sickle cell anemia</b>	<b>23. Just a few of the 5,000 hereditary diseases that have been identified</b>
<b>24. Can make a protein malfunction, which may cause disease</b>	<b>24. Mutations of DNA</b>

<b>25. The number of genes in human DNA</b>	<b>25. 25,000</b>
<b>26. The number of base pairs in human DNA</b>	<b>26. About 3 billion</b>
<b>27. One-letter variations in human DNA sequence that contribute to differences among individuals</b>	<b>27. Single-nucleotide polymorphisms or SNP's</b>
<b>28. Tailoring a drug to an individual patient, whose response to that drug can be predicted by genetic fingerprinting</b>	<b>28. One possible future of Pharmacogenomics</b>

<p><b>29. Technologies that allow the detection of thousands of genes as they are turned on or off in different types of cells and in response to different stimuli</b></p>	<p><b>29. Microarrays and DNA chips</b></p>
<p><b>30. An organism's entire set of DNA</b></p>	<p><b>30. Genome</b></p>