

Chapter 1: Human Genome

1.1. INTRODUCING GENES AND GENOME

- **Genetics** – study of heredity, traits and their variations.
 - study of how traits are transmitted.
- **Heredity**: transmission of traits and biological information.
- **Genes**: basic unit of heredity.
 - inherit cell's production of proteins.
 - consist of long DNA.
 - protein-encoding genes are scattered among 3.2 billion DNA bases in each set of chromosomes.
- **Proteins** – control the characteristics of every individual.
- **DNA (Deoxyribonucleic Acid)** – harbors information in sequence of nitrogenous bases.
- **Genome** – made up of complete set of genetic instructions characteristic of each cell organism.
 - all of our cells contain two sets copies of genome.
- **Genetics** – scientific field that analyzes genomes.
 - reveals how closely related are different species.

1.2 LEVELS OF GENETICS AND GENOMICS

- **DNA (Deoxyribonucleic Acid)**
 - Shape: double helix
 - Consist of: phosphate sugar and phosphates
 - Function: provide information to manufacture specific proteins



- 4 types of nitrogenous bases:
 - Adenine (A) and Thymine (T)
 - Cytosine (C) and Guanine (G)
 - Each set of these complementary bases is a code for a particular amino acid.
- **Amino Acid** – building blocks of protein.
 - align and link like-into bonds forming protein.

1.3. THREE PROCESSES OF ACQUIRED GENETIC INFORMATION

- **DNA replication** – (1) chains of double helix untwist and separate.
 - (2) each part builds a new partner chain from free DNA bases.
- **Transcription** – (3) copies the sequence of one strand of DNA molecule into mRNA.
- **Translation** – (4) each 3 DNA bases in a row attract another type of RNA, that functions as a connector, bringing in a particular amino acid.
 - (5) Amino acids align and link to form protein.

1.2 THE HUMAN GENOME

- **20,125 protein-encoding genes** comprise the genome.
 - Accounts: 1.5 percent of human genome, 8.5 percent of known genetic diseases.
- **Alleles** ("gene variants") – some proteins-encoding genes may vary in DNA base sequence from parent to parent that is passed to the next generation.
- **Mutations** ("changed gene") – alteration or change in genes, causing a disease.

1.3. INHERITED DISEASES

- **Cystic Fibrosis** – inherited disease
 - Change of a "C" to "T" inserts amino acid rather than the amino acid glutamic which results in protein becoming abnormal.
 - **Cystic Fibrosis Transmembrane**

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Mikaeel D. Shabazz Muhammad



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On Military Science: A Guide to Understanding the Meaning of F.O.I. (Volume 2) Mikaeel D. Shabazz
Muhammad,2011-12-16 Collection of essays letters and class outlines based on military science and the teachings of the Most Hon Elijah Muhammad as represented by the Hon Louis Farrakhan of the Nation of Islam Forthcoming Books Rose Army,1998-06 *Whitaker's Cumulative Book List* ,1962 *Books of the Month* ,1960 *The Human Genome in Health and Disease* Tore Samuelsson,2019-02-07 The human genome is a linear sequence of roughly 3 billion bases and information regarding this genome is accumulating at an astonishing rate Inspired by these advances *The Human Genome in Health and Disease A Story of Four Letters* explores the intimate link between sequence information and biological function A range of sequence based functional units of the genome are discussed and illustrated with inherited disorders and cancer In addition the book considers valuable medical applications related to human genome sequencing such as gene therapy methods and the identification of causative mutations in rare genetic disorders The primary audiences of the book are students of genetics biology medicine molecular biology and bioinformatics Richly illustrated with review questions provided for each chapter the book helps students without previous studies of genetics and molecular biology It may also be of benefit for advanced non academics which in the era of personal genomics want to learn more about their genome Key selling features Molecular sequence perspective explaining the relationship between DNA sequence motifs and biological function Aids in understanding the functional impact of mutations and genetic variants Material presented at basic level making it accessible to students without previous studies of genetics and molecular biology Richly illustrated with questions provided to each chapter **The Human Genome** R. Scott Hawley,Catherine A. Mori,1999 *The Human Genome A User s Guide* conveys both the essence and the excitement of modern human genetics Incorporating all of researchers latest discoveries the authors ground their work in the discussion of a major function of the human gene that of sex determination and development This focus opens the discussion to the interactions between science and society Hawley and Mori take care to examine the process of genetic analysis and to explore relevant topics such as the genetics of cancer behavior and personality AIDS mental illness cloning and gene therapy The reader gains sophisticated insight into human heredity beyond the misconceptions of folklore **Human Genome** ,2010 **The Book of Man** Walter Fred Bodmer,1995 This astonishing work is the first history and analysis of the Human Genome Project the massive international effort aimed at mapping each of the three billion molecules that make up human DNA The authors chronicle the startling progress of this mammoth undertaking and examine the moral questions that arise from such research **Encyclopedia of the Human Genome** ,2005 **Human Genome Project, Chromosome Number 01** Human Genome Project,2002 Human Genome Project, Build 34, Chromosome Number 01 Human Genome Project,2004

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