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That Matters 

Full Lesson:

# Genetic Mutations



Learning Standard(s)	Essential Question(s) / Thought(s)
<ul style="list-style-type: none"> <li>Students will know that genetic information can be altered by mutations and that those mutations can be harmful, neutral, or beneficial.</li> </ul>	<ul style="list-style-type: none"> <li>How do mutations occur?</li> <li>Are all mutations bad?</li> <li>How can the same genetic mutation be beneficial in some situation yet harmful in others?</li> </ul>

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**Genetic Mutations**

Student Name: \_\_\_\_\_ Date: \_\_\_\_\_

Teacher Name: \_\_\_\_\_

Directions: Read the passage and answer the questions below.

1. A mutation is a change in the DNA sequence. It can be caused by errors during DNA replication or by external factors like radiation. Mutations can be harmful, neutral, or beneficial.
2. Some mutations can lead to the development of new traits. For example, a mutation in a flower's DNA can change its color from white to red.
3. A single change in one letter of the DNA code can change the instructions for making a protein. This can affect the function of the protein and the organism's health.

Answer the following questions:

1. What is a mutation?
2. How can a mutation be harmful?
3. How can a mutation be beneficial?

# Mutations And Genetic Conditions Unit 07 Lesson

**Lauren Gardner**



## **Mutations And Genetic Conditions Unit 07 Lesson :**

### **Hidden Secrets and Lessons From the Crystal Structures of Integral Membrane Proteins Channels, Pumps and Receptors** Mario Díaz, Garth L. Nicolson, 2019-03-07      **Thymic Stromal Alterations and Genetic Disorders of Immune System** Claudio Pignata, Ana E. Sousa, 2015-12-07

The pathogenic mechanisms underlying primary T cell disorders are mainly related to molecular alterations of genes whose expression is intrinsic to hematopoietic cells. However, since the differentiation process requires a crosstalk among thymocytes and the thymic microenvironment, molecular alterations of genes involved in the differentiation and functionality of the stromal component of the thymus may lead to a severe T cell defect or failure of central tolerance as well. The first example of severe combined immunodeficiency SCID not related to an intrinsic alteration of the hematopoietic cell but rather of the thymic epithelial component is the Nude SCID phenotype inherited as an autosomal recessive disorder whose hallmarks are the T cell defect and the absence of the thymus. The clinical and immunological phenotype is the human equivalent of the murine Nude SCID syndrome which represents the first spontaneous SCID identified in nude mice in 1966. For over 3 decades, studies of immune system in these mice enormously contributed to the overall knowledge of cell mediated immunity in the assumption that the athymia of these mice was solely responsible for the T cell immunological defect. This syndrome is due to mutations of the transcription factor FOXP1 belonging to the forkhead box gene family which is mainly expressed in the thymus and skin epithelial cells where it plays a critical role in differentiation and survival. An alteration of the thymic structure is also a feature of the DiGeorge syndrome DGS which has been long considered the human counterpart of the nude mice phenotype. This syndrome is frequently associated to a deletion of the 22q11 region which contains approximately 30 genes including the TBX1 gene which is responsible for most of the clinical features of DGS in humans and mice. In this syndrome, common manifestations are cardiac malformations, speech delay, hypoparathyroidism, and immunodeficiency, even though the immunological hallmarks of the T cell defect in DiGeorge syndrome are profoundly different from those reported in human Nude SCID. The divergence of the phenotype among these 2 entities raised the possibility that the FOXP1 transcription factor represents the real key stromal molecule implicated in directing the hematopoietic stem cell toward a proper T cell fate. Thymic stromal component of the primary lymphoid organ is also required to negatively select the autoreactive clones, a process driven by the expression of tissue specific antigens TSA by medullary thymic epithelial cells mTECs. The expression of genes encoding TSA antigens is mediated by autoimmune regulator AIRE gene encoding a transcription factor expressed in mTECs. Molecular alterations of this gene are associated to autoimmune polyendocrinopathy, candidiasis, ectodermal dystrophy (APECED), a rare autosomal disorder which may be considered the prototype of an autoimmune disease due to the failure of central tolerance homeostasis. All these experiments of nature led to unravel novel pathogenic mechanisms underlying inherited disorders of immune system and of note to clarify the pivotal role of epithelial cells in the maturation and education process of T cell.

precursors NJN ... Secondary Teacher Manual, Grades 7-12 ,1988 **Health: Teacher ed** Linda Brower Meeks,1990

Ecology and Evolution of Infectious Diseases Benjamin Roche,Hélène Broutin,Frédéric Simard,2018 Provides an up to date authoritative and challenging review of the ecology and evolution of infectious diseases focusing on low income countries for effective public health applications and outcomes **Paller and Mancini - Hurwitz Clinical Pediatric**

**Dermatology E-Book** Amy S Paller,Anthony J. Mancini,2020-12-08 Written by two leaders in the field of pediatric dermatology this classic text provides both detailed content for the specialist and easily accessible information for the non dermatologist and less experienced clinician Paller and Mancini Hurwitz Clinical Pediatric Dermatology 6th Edition comprehensively covers the full range of skin disorders in children offering authoritative practical guidance on diagnosis and treatment in a single volume This award winning evidence based text has been fully revised and updated and is an essential resource for anyone who sees children with skin disorders Features new content outline boxes for faster navigation hundreds more clinical images and authors tips for the diagnosis and treatment of pediatric skin diseases Contains updated evidence based guidance and the latest drug developments and disease classifications Provides a careful balance of narrative text useful tables and 1 300 high quality clinical photographs helping you recognize virtually any skin condition you re likely to see Includes a greatly expanded discussion of atopic dermatitis and psoriasis and new therapeutic approaches for treating genetic disorders and systemic diseases such as ichthyoses and rheumatologic disorders Discusses new tests for subclassifying disease such as the myositis specific antibodies of juvenile dermatomyositis genotyping and immunophenotypes of inflammatory skin disorders Contains new and updated tables on psoriasis co morbidities genetic syndrome classifications acne therapies pediatric histiocytoses PHACE syndrome criteria HSV therapies and juvenile dermatomyositis Features updated sections on infections exanthems vascular disorders dermatoses and genodermatoses Discusses hot topics such as the use of stem cell and cell therapy as well as recombinant protein for treating epidermolysis bullosa the resurgence of measles congenital Zika virus infections and much more *Cumulated Index Medicus* ,1997

**Advanced Genetic Analysis** Philip Meneely,Matthew R. Willmann,2009-01-15 Advanced Genetic Analysis explores the question How can the principles of genetics be used as analytical tools to solve biological problems Drawing on the latest experimental tools including microarrays RNAi and bioinformatics approaches it provides a state of the art review of the field but in a truly student friendly manner **NJN Instructional Resource Manual, K-12** ,1990 **Index Medicus** ,2001

Ion Channels and Genetic Diseases Society of General Physiologists. Symposium,1995 *Study Guide [to] Microbiology, Principles & Applications, Third Edition [by] Jacquelyn G. Black* Jacquelyn G. Black,George Allan Wistreich,Black,1996

**Adventures** Mick Gammidge,Ben Wetz,2005-08 Adventures builds students confidence through the use of clear grammar presentations varied practice exercises and guided skills work The highly motivating topics have been carefully chosen to appeal to teenagers The clear syllabus offers an analytical approach to grammar as well as extensive and varied practice

Strong emphasis is placed on skills development and study skills There is a systematic approach to vocabulary building The variety of challenging and enjoyable activities motivate students to participate fully in class The optional World of English sections exploit functional and everyday language cross cultural topics and authentic pop songs     Molecular Basis of Cardiovascular Disease Kenneth R. Chien, 2004 The 2nd Edition of this heralded companion to Braunwald's Heart Disease explores the molecular mechanisms of cardiology and the scientific advances that are changing the practice of cardiology today International experts discuss the role of genetics in cardiovascular disease the molecular basis of ischemic disease thrombosis and hypertension genetic mapping approaches to inherited disorders biologically targeted agents for hypertension and heart failure and much more Abundant figures and tables illustrate key concepts Addresses most common cardiovascular problems from a molecular standpoint Explores developing treatments for cardiovascular problems based on genetics Provides references to Braunwald's Heart Disease 6th Edition Examines today's cutting edge advances in molecular cardiology and the future of gene therapy Examines the implications of cellular cholesterol metabolism in health and disease Delivers up to date information on understanding the origin of inherited disease     **The Education Index** ,1991

*Practical Genetic Counselling* Peter S. Harper, 1988     **NJN Instructional Resource Manual** ,1990     Genetics Abstracts ,2000     **Developmental Disabilities in Infancy and Childhood: The spectrum of developmental disabilities** Arnold J. Capute, Pasquale J. Accardo, 1996     Pathophysiology Kathryn L. McCance, Sue E. Huether, 2006 This fully updated textbook provides pathophysiology students with a complete knowledge base that includes the latest data from research on environmental and genetic factors that influence the disease process

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