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D. J. Weatherall

The Red Blood Cell Douglas MacN. Surgenor, 1974 The Red Blood Cell ...

Red Blood Cell Aggregation Oguz Baskurt, Björn Neu, Herbert J.

Meiselman, 2011-09-28 Red blood cells in humans—and most other mammals—have a tendency to form aggregates with a characteristic face-to-face morphology, similar to a stack of coins. Known as rouleaux, these aggregates are a normally occurring phenomenon and have a major impact on blood rheology. What is the underlying mechanism that produces this pattern? Does this really happen in blood circulation? And do these rouleaux formations have a useful function? The first book to offer a comprehensive review of the subject, *Red Blood Cell Aggregation* tackles these and other questions related to red blood cell (RBC) aggregates. The book covers basic, clinical, and physiological aspects of this important biophysical phenomenon and integrates these areas with concepts in bioengineering. It brings together state-of-the-art research on the determinants, mechanisms, and measurement and effects of RBC aggregation as well as on variations and comparative aspects. After an introductory overview, the book outlines factors and conditions that affect RBC aggregation. It presents the two hypotheses—the bridging model and the depletion model—that provide potential mechanisms for the adhesive forces that lead to the regular packing of the cells in rouleaux formations. The book also reviews the methods used to quantify RBC aggregation in vitro, focusing on their importance in clinical practice. Chapters discuss the

effect of RBC aggregation on the in vitro rheology of blood as well as on tube flow. The book also looks at what happens in the circulation when red blood cells aggregate and examines variations due to physiological and pathophysiological challenges. The concluding chapter explores the formation of red blood cell aggregates in other mammals. Written by leading researchers in the field, this is an invaluable resource for basic science, medical, and clinical researchers; graduate students; and clinicians interested in mammalian red blood cells.

Red Blood Cell Membranes Peter Agre, 2020-08-12 This book is devoted to the red blood cell membrane, its structure and function, and abnormalities in disease states. It presents a well-documented and well-illustrated comprehensive picture of clinical manifestations of red blood cell disorders.

Red Blood Cells at the Mount of Truth: Highlights of the 22nd Meeting of the European Red Cell Research Society Anna Bogdanova, Lars Kaestner, 2021-01-06

The Morphology of Human Blood Cells Lemuel W. Diggs, Dorothy Sturm, Ann Bell, 1985

The Red Blood Cell Douglas MacN. Surgenor, 1974

Red Blood Cell Aging Antonio De Flora, Mauro Magnani, 2013-03-08 The mammalian erythrocyte is a very suitable model for the study of aging at the cellular and molecular level. It is not only a matter of apparent simplicity in terms of biochemistry, biophysics and physiology but more likely this cell

offers a great possibility for elucidating some basic problems in the process of aging. In fact, nowadays, it is possible to follow individual cells all along their life span in circulation, it is possible to obtain these cells when young, middle aged or old and it is possible to obtain cells from individuals of defined ages and transfuse them into compatible recipients to investigate the role of the environment where the cell lives, and finally it is possible to easily manipulate the red cell content in terms of enzymatic activities and/or metabolic properties to investigate the possible effect of these manipulations on cell survival. This book, *Red Blood Cell Aging*, is based on a symposium held in Urbino, Italy, at the end of 1990 and examines the impact of age on the membrane, metabolism, structural and enzymatic proteins of mammalian erythrocytes. The various contributions to this symposium not only described those processes of aging which affect the cell but also provided a nearly complete picture of the event{s} and mechanism{s} that every day permits to recognize among 25 trillion circulating red cells {in an average adult} that 1 percent that have reached the end of their 120 day life span in circulation.

Red Blood Cell count S K Bajaj, 2019-08-31 Red Blood Cell count

Red Blood Cell Diameters Cecil Price-Jones, 1933

Advances in Red Blood Cell Biology D. J. Weatherall, 1982-01-01

Focus on Sickle Cell Research Ralph L. Plasmar, 2004 Sickle cell anaemia is an inherited blood disorder, characterised primarily by chronic anaemia and

periodic episodes of pain and occurring in approximately 1 in every 400 African-American infants born in the United States each year. Individuals of Mediterranean, Arabian, Caribbean, South and Central American, and East Indian ancestry can also be affected. The underlying problem involves haemoglobin, a component of the red cells in the blood. The haemoglobin molecules in each red blood cell carry oxygen from the lungs to the body organs and tissues and bring back carbon dioxide to the lungs. In sickle cell anaemia, the haemoglobin is defective. After the haemoglobin molecules give up their oxygen, some of them may cluster together and form long, rod-like structures. These structures cause the red blood cells to become stiff and to assume a sickle shape. Unlike normal red cells, which are usually smooth and donut-shaped, the sickled red cells cannot squeeze through small blood vessels. Instead, they stack up and cause blockages that deprive the organs and tissue of oxygen-carrying blood. This process produces the periodic episodes of pain and ultimately can damage the tissues and vital organs and lead to other serious medical problems. Unlike normal red blood cells, which last about 120 days in the bloodstream, sickled red cells die after only about 10 to 20 days. Because they cannot be replaced fast enough, the blood is chronically short of red blood cells, a condition called anaemia. Sickle cell anaemia is caused by an error in the gene that tells the body how to make haemoglobin. The defective gene tells the body to make the abnormal haemoglobin that results in deformed red blood cells. This book gathers the

latest research in this important field.

The Red Blood Cell Charles William Bishop, Douglas MacN. Surgenor, 1964

Morphology of Human Blood Cells Ann Bell, 2005

Blood Groups and Red Cell Antigens Laura Dean, 2005

Red Blood Cells of Domestic Mammals N. S. Agar, P. G. Board, 1983-01-01 The membrane of the red blood cell; Glycolipids of the red blood cells. Erythrocyte deformability; Red cell life-span; The blood groups of domestic mammals; Hemoglobins and globin genes; Hemoglobin switching; Red cell metabolism; Glutathione metabolism in erythrocytes; Erythrocyte amino acid and nucleoside transport; Cation transport in red blood cells; Erythrocyte organic phosphates and hemoglobin functions in domestic mammals; Postnatal changes in energy metabolism of mammalian red cells. Storing and preserving erythrocytes; Electrophoretic variation of erythrocyte enzymes of domesticated mammals.

Corpuscles Marcel Bessis, 1964

The Morphology of Human Blood Cells Lemuel W. Diggs, Dorothy Sturm, Ann Bell, 1970

Pictured Glossary in Biology Prof. Amal Attia El-Morsy Ibrahim, 2017-01-01 The glossary continues to be a valuable guidance tool for biological students those studying biology either in High Schools or Science Colleges as well as scientific researchers. Everything you need for learning biological terminology is right in your hands. The language of biology is rigorous. It

is among the great tools of the mind for a better understanding and more accurate network between all biologists of the life sciences. The lists of prefixes, suffixes and terms arranged alphabetically, which lets students look terms up even if they are not sure about their exact spellings. It provides comprehensive coverage of biology, and biochemistry entries on key scientists. This glossary will contain 8000 scientific words expressing all biology branches (Zoology, Botany & Microbiology). The number of the glossary in this book is more than that found in Oxford Dictionary.

Imaging Life Lawrence R. Griffing, 2023-03-20 Hands-on resource to understand and successfully process biological image data In *Imaging Life: Image Acquisition and Analysis in Biology and Medicine*, distinguished biologist Dr. Lawrence R. Griffing delivers a comprehensive and accessible exploration of scientific imaging, including but not limited to the different scientific imaging technologies, image processing, and analysis. The author discusses technical features, challenges, and solutions of the various imaging modalities to obtain the best possible image. Divided into three sections, the book opens with the basics such as the various image media, their representation and evaluation. It explains in exceptional detail pre- and postprocessing of an image. The last section concludes with common microscopic and biomedical imaging modalities in light of technical limitations and solutions to achieve the best possible image acquisition of the specimen. *Imaging Life: Image Acquisition and Analysis in Biology and*

Medicine is written specifically for readers with limited mathematical and programming backgrounds and includes tutorials on image processing in relevant chapters. It also contains exercises in the use of popular, open-source software. A thorough introduction to imaging methods, technical features, challenges, and solutions to successfully capture biological images Offers tutorials on image processing using open-source software in relevant chapter Discusses details of acquisition needs and image media covering pixels, pixel values, contrast, tonal range, and image formats In-depth presentation of microscopic and biomedical imaging modalities Perfect for professionals and students in the biological sciences and engineering, Imaging Life: Image Acquisition and Analysis in Biology and Medicine is an ideal resource for research labs, biotech companies, and equipment vendors.

Dacie and Lewis Practical Haematology E-Book Barbara J. Bain, Imelda Bates, Mike A Laffan, 2016-08-11 For more than 65 years, this best-selling text by Drs. Barbara J. Bain, Imelda Bates, and Mike A. Laffan has been the worldwide standard in laboratory haematology. The 12th Edition of Dacie and Lewis Practical Haematology continues the tradition of excellence with thorough coverage of all of the techniques used in the investigation of patients with blood disorders, including the latest technologies as well as traditional manual methods of measurement. You'll find expert discussions of the principles of each test, possible causes of error, and the interpretation and clinical significance of the findings. A unique section on haematology in

under-resourced laboratories. Ideal as a laboratory reference or as a comprehensive exam study tool. Each templated, easy-to-follow chapter has been completely updated, featuring new information on haematological diagnosis, molecular testing, blood transfusion- and much more. Complete coverage of the latest advances in the field. An expanded section on coagulation now covers testing for new anticoagulants and includes clinical applications of the tests.

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