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(FRACP) examination. Written by a team of physicians based at Flinders Medical Centre, and covering the key components of the FRACP basic training syllabus, this guide presents over 500 multiple-choice questions on all major topics covered in the examination. It provides coverage

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of rapidly evolving topics such as healthcare in an ageing population, disparity in indigenous health outcomes, advances in molecular science and genetics, and the complexity of care arising from multiple chronic illnesses. Questions echo the written examination, including those on

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both 'Basic Sciences' and 'Clinical Practice'. Many of the questions are similar to those in the actual examination; others are designed to 'teach' particularly important issues or to draw attention to contemporary topics. Each question has an answer that fully explains the

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correct and incorrect responses. This study aid also includes:

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practice and AP® test preparation; it also highlights careers and research opportunities in biological sciences. Transfusion Medicine, Apheresis, and Hemostasis: Review Questions and Case Studies is the collaborative effort that spanned a time period of 2 years and included 50 experts, many whom

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are national leaders in their respected fields. It also represents the passion and privilege we feel to teach the next generation of physicians in Transfusion Medicine and Apheresis. The main goal for this book is to help the readers build a solid foundation of both basic and advanced conceptual knowledge

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to prepare for the American Board of Pathology (ABP) certification exam in Transfusion Medicine. This book is not intended to be a substitute for textbooks, original research or review articles, and/or clinical training. Further, since the field of medicine, both from a scientific and regulatory

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perspective, rapidly changes, the readers are advised to continuously update their knowledge by attending national meetings and reading clinical journals. To equip the readers with the basic knowledge in critical reading and data analysis, which is an essential skill in daily medical practice, a novel

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chapter titled "Data Interpretation in Laboratory Medicine was included in this book. In this chapter, the readers are asked to make logical conclusions based on the given data and/or statistical results. Moreover, there is also a chapter on "Practical Calculations in Transfusion Medicine,

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Apheresis, and Hemostasis to help consolidate all the necessary formulas commonly used in daily practice for easy reference. These chapters are unique to our book and will not be found in any other currently on the market. All of the questions in this book were originally created by the authors

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of each chapter. Each question can either be standalone or part of a case scenario representing challenge cases in Transfusion Medicine, Apheresis, and Hemostasis. These questions often represent both rare and common clinical scenarios that the authors have seen during their clinical practice. Each

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question is then followed by 5 possible answers, with only one being correct (or the best answer). After the question, there is a conceptual explanation followed by a more factual explanation of the right and wrong answers. We gave the individual authors the freedom to choose how

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they explained the wrong answer choices. Some authors chose to be more direct (e.g. Answer A is incorrect because...), while other authors chose a more conversational style (e.g. Human resources (answer A) includes staffing, selection, orientation, training, and competency assessment of

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employees). This format is designed to help the student linking the conceptual and factual knowledge together to form a solid foundation for use in clinical practice. At the end of each chapter, there is a list of articles and textbooks that will prove useful to the motivated student who wishes to become an

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expert in the field. Another special feature to our textbook is the presence of a pre-test and post-test, which are provided to help the readers with self-assessment. As stated above, the main focus of this book is to help the readers preparing for the ABP certification exam in Transfusion

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Medicine. However, due to the interdisciplinary nature of the field of Transfusion Medicine, Apheresis, and Hemostasis, we believe that this book is also beneficial to and can be used by all clinicians involved in the management of complex transfusion, apheresis, and hemostasis issues,

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such as hematologists, anesthesiologists, surgeons, and critical care physicians. We further believe that it is a helpful guide for these specialists to prepare for their own specialty certification exam, when the topics are related to Transfusion Medicine, Apheresis, and Hemostasis.

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The major histocompatibility

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complex (MHC) is a highly polymorphic and diverse multigene locus in all jawed vertebrate species that has an integral role in adaptive/innate immune systems, transplantation, and infectious

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and autoimmune diseases. The MHC supra-locus in mammalian vertebrates is usually partitioned into three distinct regions, known as classes I, II, and III, which, to varying extents, can be found conserved in

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nonmammalian jawed vertebrates, such as bony fish, amphibians, and bird lineages. The MHC gene region is characterized particularly by the expression of class I and class II glycoproteins that bind peptides

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derived from intracellular or extracellular antigens to circulating T-cells. While this expressed antigenic specificity remains the predominant interest with respect to MHC function and polymorphism in a population, a

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broader concept has emerged that examines the MHC as a multifunctional polymorphic controller that facilitates and regulates genome diversity with a much greater array of functions and effects than just MHC-

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restricted antigen recognition. This volume of 19 reprints presented by various experts and collected from the Special Issue of Cells on "MHC in Health and Disease" covers a broad range of topics on the genomic

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diversity of the MHC regulatory system in various vertebrate species, including MHC class I, II, and III genes; innate and adaptive immunity; neurology; transplantation; haplotypes; infectious and autoimmune

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diseases; fecundity;
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these articles demonstrate the
immense complexity and
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brief description of the disorder followed by a discussion that includes tables detailing the laboratory evaluation of specific disorders, diagnosis, baseline

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*tests to exclude
diagnostic
possibilities, and
clinical indications
that warrant further
screening and special
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experimentalists detail
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biology, and immunology, they describe step-by-step methods designed to question how MHC-binding peptides are generated, to test how peptides are delivered to MHC

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Furthermore, a book with concise overview of the field and highlights of clinical applications will certainly help those trainees, including pathology

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residents, molecular
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you understand and can
apply important concepts
of the sciences basic to**

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**the practice of
medicine, with special
emphasis on principles
and mechanisms
underlying health,
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mastery of not only the sciences that provide a foundation for the safe and competent practice of medicine in the present, but also the scientific principles

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cytoplasm and in the endoplasmic reticulum) and at the cell surface (between T cells and MHC molecules). To investigate such a diverse array, it is

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**necessary that
biochemical, cell
biology, and
immunological techniques
all be employed. The
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how MHC-binding peptides are generated, to test how peptides are delivered to MHC molecules, to analyze MHC peptide-binding patterns, and to assay

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of research in extending the questions that can be asked and answered by the application of these current methods. For editorial assistance, I would like to thank

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book, and can be purchased separately. It contains animations and videos with voiceover narration, as well as the figures from the text for presentation purposes.

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**Immunologic privilege of the
cornea and anterior chamber has,
in most cases, been believed to be
a sufficient guarantee for survival
of the donor cornea. If this failed,**

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it was thought that adjunctive immunomodulating agents would compensate. But the question of how to care for better tolerance and safer long-term survival of corneal grafts after withdrawal of a potent immunomodulating

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shield still remained unanswered.

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well as experienced keratoplasty surgeons. These experts conclude that HLA matching is, in fact, a valuable adjunct for long-term survival of corneal grafts not only in those at high risk but notably also in patients at normal risk.

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This volume opens new pathways towards the successful application of HLA matching to keratoplasty surgeons, experimental and clinical immunologists and cornea bank officers, at the same time acting as an aid to

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information is ever more important as the limits of transplantation continue to be reduced, including the recent success of bone marrow transplantation between unrelated but closely matched individuals.

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In addition, the need to transfuse platelets in the face of immunologic barriers continues to challenge transfusion services. Thus, the serologic information summarized in this volume is essential for optimal patient care.

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At the same time, recombinant DNA technology has led to a revolution in our understanding of many aspects of basic biology. Among the advances has been the initial characterization of the structure of some HLA loci. While

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this will ultimately improve clinical services, constant reference to serologic data is essential so that the powerful new techniques can be applied in the most effective ways. The timing of the First Red Cross International

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Histocompatibility Workshop is fortunate as it brings together experts from around the world to address the state of the art. We are all grateful to Dr. John Lee and his colleagues for organizing the workshop, and for bringing

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**together in this volume the
material to be presented in
Beijing during October 17-23,
1990. Leon W. Hoyer, M.D.
Laposata's Laboratory Medicine
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**Molecular Biology of the Cell
Major Histocompatibility
Complex (MHC) in Health and
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Disease

Heart transplantation remains one of the major scientific achievements of twentieth century medicine. During the past four decades, it has evolved from an unproven

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experimental surgical technique to the most effective form of therapy for refractory end-stage heart disease. It has captured the public's imagination and expanded our understanding of fundamental immunologic mechanisms that are responsible for cellular and

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***humorally-mediated immunity.
Despite its successes, many clinical
and scientific problems remain. One
or more bouts of acute cellular or
humoral (vascular) rejection will
occur in over 75% of transplant
recipients despite current
immunosuppressive strategies.***

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Further, rejection directly results in approximately 20% of post-transplant deaths and is believed to play a major role in the development of late allograft dysfunction and coronary vasculopathy. This book by international experts in the fields of

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transplantation medicine, immunobiology and cardiac imaging provides the reader with an up-to-date, concise summary of the latest developments in the diagnosis and treatment of acute cardiac rejection. It is axiomatic that a more complete understanding of

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the pathogenic processes involved in rejection will ultimately lead to its prevention. This volume will be useful to transplant cardiologists, cardiovascular surgeons, cardiac pathologists and transplant scientists who seek to prolong the lifespan and improve the quality of

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*life of their transplant recipients.
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review of medical microbiology and immunology Review of Medical Microbiology and Immunology is a succinct, high-yield review of the medically important aspects of microbiology and immunology. It covers both the basic and clinical aspects of bacteriology, virology,

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mycology, parasitology, and immunology and also discusses important infectious diseases using an organ system approach. The book emphasizes the real-world clinical application of microbiology and immunology to infectious diseases and offers a unique mix of

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these questions for successfully certifying this exam.

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communities of plants and animals
Autoimmune disease
disproportionately affects more
women than men. Rheumatoid
arthritis (RA) and systemic lupus
erythematosus (SLE) are two
prototypic female-predominant

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autoimmune diseases. Despite different clinical manifestations, similarities exist. Both are characterized by a dysregulated immune response to self-tissues. For both diseases, the strongest genetic risk factors are linked to human

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leukocyte antigen (HLA) alleles. Pregnancy affects their course and may also contribute to risk. This dissertation investigates mother-child HLA relationships and risk of developing RA or SLE. Chapter 1 provides background on RA and SLE

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and a literature review outlining the rationale for the dissertation. The chapter states the hypotheses of the project and the specific aims addressed in each chapter. Chapter 2 addresses the question of mother-child histocompatibility and risk of

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RA or SLE. The study tests the hypothesis that having any children who are compatible from the perspective of the mother increases the risk of developing RA or SLE. Results show that having any histocompatible children at HLA-B

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and DPB1 increase risk of RA in the mother. Increased structural similarity as quantified by a sequence similarity matching score at DPB1 was associated with decreased risk of SLE. Mother-child compatibility at minor histocompatibility antigen ZAPHIR

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was likewise associated with decreased risk of SLE but compatibility at LB-WNK1 increased risk for mothers who were also HLA-compatible at HLA-A with their children. This is the first study investigating the relationship between minor

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histocompatibility antigen compatibility and risk of SLE. This study supports the hypothesis that mother-child HLA relationships are associated with risk of RA and SLE. Chapter 3 addresses the question of whether having children who carry

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HLA alleles associated with increased risk of SLE contributes to its development among mothers. The alleles investigated include DRB1*03:01, *15:01, *08:01. In addition, the study tests the hypothesis that molecular mimicry by

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means of exposure to DRB1*04:01 that codes for a similar amino acid sequence to the Epstein-Barr virus, a known SLE risk factor, increases risk among mothers. There was no association between SLE risk-associated alleles. However, increased

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risk of SLE was associated with having at least one child who carried the DRB1*04:01 allele among women negative for the allele but with at least one of the predisposing alleles. This study supports the hypothesis that a child's alleles inherited from the father

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influence a mother's subsequent risk of SLE. Chapter 4 investigates the association between having children who carry alleles encoding amino acids (AA) associated with RA including the "shared epitope", the sequence DERA A at AA positions

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70-74, AA valine, lysine, alanine at positions 11, 71, 74 of the HLA-DRB1 molecule, aspartic acid at position 9 of HLA-B and phenylalanine at position 8 of DPB1. The study finds increased risk of RA among mothers who had any child

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with SE, DERAA, valine, lysine and alanine encoding alleles, independent of maternal carrier status. Among non-carrier mothers, increased risk of RA was associated with having children who carried DERAA. Chapter 5 summarizes the key findings and

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results from each study and the conclusions of this work. Future directions are also addressed.

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report concludes that human reproductive cloning would be dangerous for the woman, fetus, and newborn, and is likely to fail. The study panel did not address the issue of whether human reproductive cloning, even if it were

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found to be medically safe, would be "or would not be" acceptable to individuals or society.

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