

《简明内科学》

图书基本信息

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内容概要

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书籍目录

SECTION	Introduction to Molecular Medicine	1 Changes to Medical Practice in the Age of Genomics
SECTION	Decision-making in Clinical Medicine	2 Evidence-based Medicine
SECTION	Cardiovascular Disease	3 Structure and Function of the Normal Heart and Blood Vessels
3.1	Heart	3.2
Circulation	4 Evaluation of the Patient with Cardiovascular Disease	4.1 Symptoms
examination	4.2 Physical	4.3 Examination of the heart
5 Diagnostic Tests and Procedures	5.1 Electrocardiogram	5.2 Echocardiography
6 Coronary Heart Disease	6.1 Epidemiology of atherosclerosis	6.2 Coronary atherosclerosis
synonyms (ACS)	6.3 Stable angina pectoris (SAP)	6.4 Acute coronary
6.5 Acute myocardial infarction	6.6 Percutaneous coronary intervention	7 Heart Failure
8 Cardiac Arrhythmias	8.1 Definition	8.2 Symptoms of cardiac arrhythmias
arrhythmias	8.3 Investigation of	8.4 Bradycardias
8.5 Tachycardias	8.6 Extrasystoles	8.7 Atrial fibrillation and atrial flutter
8.8 Treatment of cardiac arrhythmias	9 Valvular Heart Disease	9.1 Mitral stenosis
9.2 Mitral regurgitation	9.3 Aortic stenosis	9.4 Aortic regurgitation
9.5 Tricuspid stenosis	9.6 Tricuspid regurgitation	9.7 Pulmonary valve disease
10 Myocarditis and Cardiomyopathy	10.1 Myocarditis	10.2
Cardiomyopathy	11 Hypertension	11.1 Definitions and classification
Pathophysiology	11.2 Etiology	11.3
11.4 Symptoms	11.5 Measuring blood pressure	11.6 Subclinical organ damage
11.7 Identifying secondary causes of hypertension	11.8 Diagnosis	11.9 Treatment
11.10 Patients follow-up	12 Syncope	13 Acute Bronchitis
SECTION	14 Pneumonia	14.1
Pneumococcal pneumonia	14.2 Pneumonia, fungal	14.3 Pneumonia, viral
Tuberculosis	14.4 Bronchiectasis	15 Bronchiectasis
16.1 Primary pulmonary tuberculosis	16.2 Postprimary pulmonary tuberculosis	17 Chronic
Obstructive Pulmonary Disease	18 Bronchial Asthma	19 Pulmonary Embolism
20 Pulmonary Hypertension	20.1 Secondary pulmonary artery hypertension (SPAH)	20.2 Pulmonary hypertension (PPH)
Interstitial Lung Disease	22 Disease of Pleura	23 Pneumothorax
24 Primary Bronchogenic Carcinoma	25 Respiratory Failure	SECTION
Renal Disease	26 Structure and Function of the Kidney	27 Water and
Electrolyte Metabolism	28 Approach to the Patient with Renal Disease	29 Acute Renal Failure
30 Chronic Kidney Disease	31 Glomerular Diseases	32 Major Nonglomerular Disorders
33 Vascular Disorders of Kidney	SECTION	Gastrointestinal and Liver Disease
34 Structure and Function of the Gut	35	Symptomatology of Gastrointestinal Disease
36 Upper Gastrointestinal Endoscopy	37 Peptic Ulcer Disease	38
Cirrhosis	39 Crohn's Disease	SECTION
Endocrine and Metabolic Disorders	40 Diabetes Mellitus	41
Thyroid Disease	41.1 Hypothyroidism	41.2 Thyrotoxicosis
43 Anemias	42 Dyslipidaemia	SECTION
43.1 Iron deficiency anemia	43.2 Folic acid and vitamin B12 deficiency anemia	Hematology
44 Aplastic anemia	44.1 The acute leukemias	44.2 Chronic myelogenous leukemia
45 Disorders of Hemostasis	46 Blood Transfusions	SECTION
47 Rheumatic Diseases	48 Rheumatic Disease	49 Systemic Lupus Erythematosus

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章节摘录

插图：Given the great effort needed to define allelic variants contributing to complex disease, it is reasonable to ask whether such a large investment of resources is warranted. To be able to answer in the affirmative, it is necessary to demonstrate that benefits will accrue to everyday medical practice and patient health. Understanding genetic factors that contribute to disease could help establish a more rational basis for many aspects of patient care by providing deep insights into molecular pathogenesis and through improved molecular diagnostic tools that allow individually tailored preventive and/or therapeutic regimens. Better Understanding of Molecular Mechanisms of Disease Despite the extraordinary advances in our understanding of the functions of cells and organ systems in states of health and disease, it is somewhat humbling that fewer than 5000 human genes have been functionally characterized, many in only a cursory fashion. Clearly, it is difficult to provide full descriptions of the ways in which disease processes perturb cellular function in the absence of a comprehensive catalogue of genes that are either affected by these disease processes or are involved in the response to disease. The Human Genome Project provides such a catalogue, giving a complete description of the DNA and protein sequences of all of these genes.

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编辑推荐

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精彩短评

- 1、很不错，不过内容太简易了，很多内容都省略掉了。我是学消化的，结果拿到书后发现消化内科部分好少。这一点有点不好
- 2、rt！！
- 3、图书这种商品，没什么太多说的，不错。
- 4、但是书没翻两下就要散架了
- 5、还没有怎么读刚刚收到,看了几眼 还可以吧

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