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ල් ^න Components of Innate Immunity	,	Robbins and Cotran PATHOLOGIC
P Cellular Receptors for	Kumar, Vinay, MBBS, MD, FRCPath; Abbas, Abul K., MBBS; Aster, Jon C., MD, PhD;	
Damaged Cells, and	Robbins and Cotran Pathologic Basis of Disease, Chapter 6, 185-264	
⊘ Toll-Like Receptors.	The immune system is vital for survival, because it protects us from infectious pathogens that	-
€ ⁹ NOD-Like Receptors		Robbins and Cotran
and the Inflammasome.	Examples of disorders caused by immune responses include <i>allergies</i> and reactions against an	Pathologic Basis of Disease
Other Receptors for	individual's own tissues and cells <i>(autoimmunity)</i> .	Ninth Edition
	This chapter is devoted to diseases caused by too little immunity or too much immunologic	Kumar, Vinay, MBBS, MD, FRCPath; Abbas,
Immunity		Abul K., MBBS; Aster, Jon C., MD, PhD
Adaptive Immunity	of the important features of normal immune responses, to provide a foundation for	Copyright © 2015, 2010, 2004, 1999, 1994, 1989, 1984, 1979, 1974 by Saunders, an imprint
P Cells of the Immune System	understanding the abnormalities that give rise to immunologic diseases.	of Elsevier Inc.
P Lymphocyte Diversity	The Normal Immune Personance	Launch Bookshelf >
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₀⁰ B Lymphocytes	The classic definition of <i>immunity</i> is protection from infectious pathogens, and the normal	
o [©] Dendritic Cells ↓	immune response is best understood in this context. The mechanisms of defense against	
	microbes fall into two broad categories ($Fig. 6-1$). <i>Innate immunity</i> (also called natural, or	
	 The Normal Immune Response Innate Immunity Components of Innate Immunity Cellular Receptors for Microbes, Products of Damaged Cells, and Foreign Substances Toll-Like Receptors. NOD-Like Receptors and the Inflammasome. Other Receptors for Microbial Products. Reactions of Innate Immunity Adaptive Immunity Cells of the Immune System Lymphocyte Diversity T Lymphocytes B Lymphocytes 	 Che Normal Immune Response Chanate Immunity Centrate Receptors for Microbes, Products of Damaged Cells, and Foreign Substances Celluita Receptors for Microbes, Products of Damaged Cells, and Foreign Substances Colluita Receptors for Microbes and the Inflammasone. Cother Receptors for Microbia Products. Mother Receptors for Microbia Products. Microbia Microbia Products. Microbia Products. Microbia Microbia Products. Microbia Products. Microbia Products. Microbia Microbia Microbia Products. Microbia Microbia Microbia Products. Microbia Products.<!--</td-->

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Figure 2-23 Mechanisms of apoptosis. The two pathways of apoptosis differ in their induction and regulation, and both culminate in the activation of caspases. In the mitochondrial pathway, proteins of the BCL2 family, which regulate mitochondrial permeability, become imbalanced and leakage of various substances from mitochondria leads to caspase activation. In death receptor pathway, signals from plasma membrane receptors lead to the assembly of adaptor proteins into a "death-including signaling complex," which activates caspases, and the end result is the same.

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Pathology literally translates to the study of *suffering* (Greek *pathos* = suffering, *logos* = study); more prosaically, the term *pathology* is invoked to represent the study of *disease*. Germane to this opening chapter, Virchow coined the term *cellular pathology* to emphasize the basic tenet that all diseases originate at the cellular level. Thus, modern pathology is basically the study of *cellular* abnormalities. Therefore, diseases and the underlying mechanisms are best understood in the context of *normal* cellular structure and function.

irable) to condense the vast and fascinating field of cell biology into a its of biology are likely quite familiar with many of the broader concepts insequently, rather than attempting a comprehensive review, our goal is and highlight some recent advances that are relevant to the pathologic ed throughout the text. We hope this chapter will be useful to review key hey apply to the areas of Pathology that are covered from **Chapter 2**

大聲朗讀

The sequencing of the human genome represented a landmark achievement of biomedical science. Published in draft form in 2001 and more completely detailed in 2003, the information has already led to remarkable advances in science and medicine. Since then there has been an exponential decrease in to cost of sequencing and an exponential increase in data accrual; this new information, now literally at

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The Systemic Vasculitides

John H. Stone

Definition

The vasculitides are a heterogeneous group of disorders linked by the common finding of destructive inflammation within blood vessel walls. The most current nomenclature scheme¹ identifies at least 27 different forms of primary vasculitis (Table 270-1). The major forms of vasculitis are discussed in this chapter.

TABLE 270-1

NAMES FOR VASCULITIDES ADOPTED BY THE 2012 INTERNATIONAL CHAPEL HILL CONSENSUS CONFERENCE ON THE NOMENCLATURE OF VASCULITIDES.¹

LA	ARGE-VESSEL VASCULITIS
	Takayasu arteritis Giant cell arteritis
M	EDIUM-VESSEL VASCULITIS
	Polyarteritis nodosa Kawasaki disease Buerger disease*
SN	AALL-VESSEL VASCULITIS
	Antineutrophil cytoplasmic antibody (ANCA)-associated vasculitis Microscopic polyangiitis Granulomatosis with polyangiitis (formerly Wegener granulomatosis) Eosinophilic granulomatosis with polyangiitis (Churg-Strauss syndrome Immune complex small-vessel vasculitis Antiglomerular basement membrane disease Cryoglobulinemic vasculitis Immunoglobulin (Ig)A vasculitis (Henoch-Schönlein purpura) Hypocomplementemic urticarial vasculitis
VA	ARIABLE-VESSEL VASCULITIS
	Behçet syndrome Cogan syndrome
SI	NGLE-ORGAN VASCULITIS
	Cutaneous leukocytoclastic angiitis Cutaneous arteritis Primary central nervous system vasculitis Isolated aortitis

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LARGE-VESSEL VASCULITIS		
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新增附註		新組
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	Takayasu arteritis	TABLE NAME S
	Giant cell arteritis	LARG
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H. Stone

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Polyarteritis nodosa Kawasaki disease Buerger disease*
SMALL-VESSEL VASCULITIS
Antineutrophil cytoplasmic antibody (ANCA)-associated vasculitis Microscopic polyangiitis Granulomatosis with polyangiitis (formerly Wegener granulomatosis) Eosinophilic granulomatosis with polyangiitis (Churg-Strauss syndrome) Immune complex small-vessel vasculitis

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	scheme

The Systemic Vasculitides

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	ayasu arteritis nt cell arteritis
Giai	nt cell arteritis
MEDIU	UM-VESSEL VASCULITIS
Poly	yarteritis nodosa
Kav	vasaki disease
Bue	erger disease*
SMAL	L-VESSEL VASCULITIS
Ant	ineutrophil cytoplasmic antibody (ANCA)-associated vasculitis
M	licroscopic polyangiitis
G	ranulomatosis with polyangiitis (formerly Wegener granulomatosis)
	osinophilic granulomatosis with polyangiitis (Churg-Strauss syndrom
	nune complex small-vessel vasculitis

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