IMMUNOLOGY: Mary K. Crow, MD (Immunology); Bruce N. Cronstein, MD (Inflammation)

- Anatomy and cellular elements of the immune system
 - Lymphoid organs: gross and microscopic anatomy and function
 - Organization of immune system: innate and adaptive responses
 - o Specific cell types: ontogeny, structure, phenotype, function, and activation markers and cell membrane receptors
- Immune and inflammatory mechanisms
 - o Antigens: types, structure, processing, presentation, and elimination
 - Components and regulation of innate immune system
 - o Major histocompatibility complex: structure, function, and nomenclature
 - B-cell receptors and immunoglobulins: structure, function, antigen binding, signaling, genetic basis, and effector function
 - o T-cell receptors: structure, function, antigen binding, signaling, and genetic basis
 - o Receptor-ligand interactions, adhesion molecules, complement receptors, Fc receptors, and signal transduction
 - o Complement and kinin systems: structure, function, and regulation
 - Acute-phase reactants and enzymatic defenses
- Cellular interactions, immune regulation, and immunomodulation
 - Activating and inhibitory immune receptors
 - Cellular activation, suppression, and regulation of each cell type
 - Origin, structure, effect, site of action, metabolism, and regulation of cytokines, chemokines, and other inflammatory mediators
 - o Mechanisms of immune tolerance
- Immune responses
 - o IgE-mediated: acute- and late-phase reactions
 - o Immunoglobulin-mediated: opsonization, complement fixation, and antibody-dependent cellular cytotoxicity
 - o Immune complex-mediated: physiochemical properties and clearance of immune complexes
 - o Cell-mediated: cells and effector mechanisms in cellular cytotoxicity and granuloma formation
 - o Mucosal immunity: interactions between gut and bronchus-associated lymphoid tissue and secretory IgA
 - Natural killer cells, lymphokine-activated killer cells, and graft-versus-host reaction
 - Autoantibodies
- Tissue destruction and repair
 - o Cellular and molecular mediators

o Proteases and collagenases

CRYSTAL INDUCED ARTHROPATHIES: Theodore R. Fields, MD

- Gout
 - o Primary gout
 - Asymptomatic hyperuricemia
 - Acute gout
 - Intercritical periods
 - Tophaceous gout
 - Conditions associated with gout
 - Lead intoxication
 - Secondary gout
 - Lesch-Nyhan syndrome
 - Other secondary types of gout
- Calcium pyrophosphate dihydrate deposition (CPPD)
 - o Familial
 - o Secondary to primary metabolic disorders
 - o Idiopathic CPPD
- Basic calcium phosphate crystal deposition

INFECTIONS AND RELATED ARTHRITIDES: Leonard H. Calabrese, DO

- Infections
 - o Bacterial (nongonococcal and gonococcal)
 - Native joint
 - Prosthetic joint
 - Spine
 - Bone
 - Soft tissue
 - o Mycobacterial
 - o Spirochetal (syphilis, Lyme disease)

- Viral (human immunodeficiency virus [HIV], hepatitis B virus, hepatitis C virus, parvovirus, chikungunya virus, and others)
- o Fungal
- o Parasitic
- o Whipple's disease
- Related arthritides
 - o Acute rheumatic fever and poststreptococcal arthritis
 - o Arthritis associated with bacterial endocarditis
 - o Postimmunization arthritis

METABOLIC BONE DISEASE: Nancy E. Lane, MD

- Low bone mass
- Osteoporosis
 - o Primary
 - Postmenopausal
 - Male
 - o Secondary
 - Medication-induced
- Other causes of bone loss
- Paget's disease of bone
- Bone disease related to renal disease
- Osteomalacia

OSTEOARTHRITIS & RELATED DISORDERS: Marc C. Hochberg, MD, MPH

- Osteoarthritis
- Diffuse idiopathic skeletal hyperostosis (DISH)
- Malignant and nonmalignant tumors of bones, tendons, and joints
 - o Benign tumors
 - Malignant tumors
- Osteonecrosis

RHEUMATOID ARTHRITIS: Stephen A. Paget, MD

- Seropositive rheumatoid arthritis
 - o Early disease
 - o Established disease
 - o Late disease
- Seronegative inflammatory polyarthritis
- Complications of established disease
 - Extra-articular manifestations
 - o Cardiovascular disease: atherosclerotic cardiovascular disease and congestive heart failure
 - Malignancy
 - Vasculitis
 - o Immunologic considerations

SERONEGATIVE SPONDYLOARTHROPATHIES: John J. Cush, MD

- Ankylosing spondylitis
 - Skeletal manifestations
 - Extra-articular manifestations
- Reactive arthritis
 - Skeletal manifestations
 - Extra-articular manifestations
- Arthropathy associated with inflammatory bowel disease (IBD)
 - Skeletal manifestations
 - o Extra-articular manifestations
- Psoriatic arthropathy
 - Skeletal manifestations
 - Extra-articular manifestations
- Arthritis associated with other skin diseases
 - SAPHO syndrome (synovitis, acne, pustulosis, hyperostosis, and osteitis)
- Undifferentiated spondyloarthropathies
 - Skeletal manifestations
 - Extra-articular manifestations

OTHER RHEUMATIC AND CONNECTIVE TISSUE DISORDERS: Edward J. Parrish, MD

- Raynaud's phenomenon
- Primarily fibrosing rheumatic diseases
 - o Systemic sclerosis
 - Skin
 - Gastrointestinal
 - Cardiac
 - Pulmonary
 - Renal
 - Scleroderma mimics
 - Scleromyxedema
 - Nephrogenic fibrosis
 - Scleredema
 - o Eosinophilic fasciitis
 - o Retroperitoneal fibrosis (Ormond disease)
- Primarily myopathic rheumatic diseases
 - o Polymyositis
 - o Dermatomyositis
 - Inclusion body myositis
 - o Metabolic myopathies
 - Medication-associated
 - Critical illness-associated
- Sjögren's syndrome
- Primary antiphospholipid antibody syndrome
- Primarily skin-associated rheumatic diseases
 - o Erythema nodosum
 - o Other forms of panniculitis
 - o Multicentric reticulohistiocytosis
- Fever-associated rheumatic disorders
 - o Autoinflammatory disorders
 - Adult-onset Still's disease (AOSD)

o Hemophagocytic lymphohistiocytosis and macrophage activation syndrome (HLH/MAS)

• Primarily joint-associated rheumatic diseases

- o Polymyalgia rheumatica (PMR)
- Remitting seronegative symmetric synovitis with pitting edema (RS3PE)
- o Palindromic rheumatism

• Miscellaneous rheumatic disorders

- Autoimmune hearing loss
- o Autoimmune eye disease
- o IgG4-related disease
- Relapsing polychondritis
- Overlap syndromes
- Undifferentiated connective tissue disease
- Mixed connective tissue disease

Pediatric disorders

- o Juvenile idiopathic arthritis (JIA)
 - Childhood disease
 - Complications in adulthood
- Kawasaki disease (KD)
- o Juvenile dermatomyositis (JDM)
- o Juvenile localized scleroderma (JLS)
- Pediatric joint disorders seen in adulthood
 - Developmental dysplasia of the hip (DDH)
 - Slipped capital femoral epiphysis (SCFE)
 - Legg-Calvé-Perthes disease

LUPUS ERYTHEMATOSUS: Michelle A. Petri, MD, MPH

- Drug-induced
- Cutaneous
 - Isolated
 - o In systemic disease

Renal

- o Immune-mediated glomerular and tubular disease
- o Antiphospholipid antibody syndrome and microangiopathies
- o Renal insufficiency and hypertension
- o Urologic complications

Neurologic

- o Central nervous system: inflammatory, vaso-occlusive, microangiopathies, and others
- o Spinal cord
- o Peripheral nerves
- o Neuromyelitis optica
- Affective disorders

Pulmonary

- o Pneumonitis
- o Thromboembolism
- o Pulmonary hypertension
- o Pneumonia

Cardiovascular

- Myocardial disease
- Valvular disease
- Accelerated atherosclerosis

Serositis

Hematologic

- o Autoimmune cytopenias
- o Hemolytic uremic syndrome (HUS) and thrombotic thrombocytopenic purpura (TTP)
- o Autoimmune clotting factor deficiencies (overlap with antiphospholipid antibody syndrome)

Musculoskeletal

- o Joints, tendons, and ligaments
- o Muscle disease

• Lupus in pregnancy

Neonatal lupus

Vasculitis

• Antiphospholipid antibody syndrome (APS)

- o Clinical features excluding pregnancy
- o Pregnancy
- o Catastrophic APS

NONARTICULAR AND REGIONAL MUSCULOSKELETAL DISORDERS: Terence W. Starz, MD

- Diffuse pain syndromes
 - o Fibromyalgia
 - o Complex regional pain syndrome (reflex sympathetic dystrophy)
 - o Medication-induced diffuse pain
- Regional musculoskeletal disorders
 - Axial syndromes
 - Back pain
 - Neck pain
 - Thoracic outlet syndrome
 - o Shoulder disorders
 - Joint
 - Soft tissue
 - o Elbow disorders
 - Joint
 - Soft tissue
 - Wrist and hand disorders
 - Joint
 - Soft tissue
 - Hip disorders
 - Joint
 - Soft tissue
 - Knee disorders
 - Joint
 - Soft tissue
 - Ankle and foot disorders

- Joint
- Soft tissue
- o Leg disorders

Neuropathies

- o Axial disorders
- o Peripheral disorders
 - Entrapment neuropathies
 - Mononeuritis multiplex
 - Polyneuropathy
 - Small fiber neuropathy

NONRHEUMATIC SYSTEMIC DISORDERS: Karen B. Onel, MD

- Hereditary, congenital, and inborn errors of metabolism associated with rheumatic syndromes
 - Disorders of connective tissue
 - Marfan syndrome
 - Osteogenesis imperfecta
 - Ehlers-Danlos syndromes
 - Pseudoxanthoma elasticum
 - Hypermobility syndrome
 - o Mucopolysaccharidoses
 - o Osteochondrodysplasias
 - Multiple epiphyseal dysplasia
 - Spondyloepiphyseal dysplasia
 - o Inborn errors of metabolism affecting connective tissue
 - Homocystinuria
 - Ochronosis
 - Storage disorders

Immunodeficiencies

- o Immunoglobulin A (IgA) deficiency
- o Complement component deficiencies
- o Severe combined immunodeficiency (SCID) and adenosine deaminase (ADA) deficiency

- o Purine nucleoside phosphorylase (PNP) deficiency
- o Common variable immunodeficiency

NONRHEUMATIC SYSTEMIC DISORDERS: C. Ronald MacKenzie, MD

- Metabolic-associated rheumatic disorders
 - o Diabetes mellitus
 - o Acromegaly
 - o Thyroid disease
 - o Cushing's disease
 - o Parathyroid disease
 - o Renal failure and dialysis
- Hematologic and oncologic malignancy-associated rheumatic disorders
 - o Amyloidosis
 - Primary
 - Secondary
 - Hereditary
 - o Lymphoma
 - o Myelodysplastic syndromes
 - o Leukemia
 - Solid tumors
 - o Plasma cell dyscrasias
 - o Hemoglobinopathies
 - Sickle cell
 - o Hemophilias

VASCULITIDES: Robert F. Spiera, MD

- Large-vessel vasculitis
 - o Takayasu's arteritis
 - o Giant cell arteritis
- Medium-vessel vasculitis
 - o Polyarteritis nodosa

Small-vessel vasculitis

- o Antineutrophil cytoplasmic antibody (ANCA)-associated vasculitis
 - Granulomatosis with polyangiitis (Wegener's)
 - Microscopic polyangiitis
 - Eosinophilic granulomatosis with polyangiitis (Churg-Strauss)
- o Immune complex small-vessel vasculitis
 - Anti-glomerular basement membrane disease
 - Cryoglobulinemic vasculitis
 - IgA vasculitis (Henoch-Schönlein purpura)
 - Hypocomplementemic urticarial vasculitis (anti-C1q vasculitis)

Variable-vessel vasculitis

- o Behçet's disease
- o Cogan's syndrome

• Single-organ vasculitis

- o Cutaneous leukocytoclastic angiitis
- Cutaneous arteritis
- o Primary central nervous system angiitis
- Isolated aortitis

Vasculitis associated with probable etiology

- o Hepatitis C virus-associated cryoglobulinemic vasculitis
- o Hepatitis B virus-associated vasculitis
- o Syphilis-associated aortitis
- o Drug-induced vasculitis
 - Drug-induced ANCA-associated vasculitis
 - Drug-induced immune complex vasculitis
 - Other drug-induced vasculitis
- Cancer-associated vasculitis

Vasculitis mimickers

- o Buerger's disease (thromboangiitis obliterans)
- Cholesterol emboli
- o Fibromuscular dysplasia

- o Segmented arterial mediolysis
- o Warfarin necrosis
- o Reversible cerebral vasoconstriction syndrome
- o Moyamoya disease
- o Atrial myxoma
- o Ergotism
- o Endocarditis
- o Calciphylaxis