

Purpose of the exam

The exam is designed to evaluate the knowledge, diagnostic reasoning, and clinical judgment skills expected of the certified rheumatologist in the broad domain of the discipline. The ability to make appropriate diagnostic and management decisions that have important consequences for patients will be assessed. The exam may require recognition of common as well as rare clinical problems for which patients may consult a certified rheumatologist.

Exam content

Exam content is determined by a pre-established blueprint, or table of specifications. The blueprint is developed by ABIM and is reviewed annually and updated as needed for currency. Trainees, training program directors, and certified practitioners in the discipline are surveyed periodically to provide feedback and inform the blueprinting process.

The primary medical content categories of the blueprint are shown below, with the percentage assigned to each for a typical exam:

Medical Content Category	% of Exam
Basic and Clinical Sciences	7%
Crystal-induced Arthropathies	5%
Infections and Related Arthritides	6%
Metabolic Bone Disease	5.5%
Osteoarthritis and Related Disorders	5%
Rheumatoid Arthritis	13%
Spondyloarthritis	6.5%
Other Rheumatic and Connective Tissue Disorders (ORCT)	16.5%
Lupus Erythematosus	9%
Nonarticular and Regional Musculoskeletal Disorders	7%
Nonrheumatic Systemic Disorders	9%
Vasculitides	8.5%
Miscellaneous Topics	2%
	100%

Exam questions in the content areas above may also address clinical topics in geriatrics, pediatrics, pharmacology and topics in general internal medicine that are important to the practice of rheumatology.

ABIM is committed to working toward health equity and believes that board-certified physicians should have an understanding of health care disparities. Therefore, health equity content that is clinically important to each discipline will be included in assessments, and the use of gender, race, and ethnicity identifiers will be re-evaluated.

Exam format

The exam is composed of up to 240 single-best-answer multiple-choice questions, of which approximately 40 are new questions that do not count in the examinee's score. Most questions describe patient scenarios and ask about the work done (that is, tasks performed) by physicians in the course of practice:

- Making a diagnosis
- Ordering and interpreting results of tests
- Recommending treatment or other patient care
- Assessing risk, determining prognosis, and applying principles from epidemiologic studies
- Understanding the underlying pathophysiology of disease and basic science knowledge applicable to patient care

Clinical information presented may include patient photographs, radiographs, micrographs, DXA scans, electrocardiograms, angiograms, and other media to illustrate relevant patient findings. <u>Learn more information on how exams are developed.</u>

A tutorial including examples of ABIM exam question format can be found at <u>http://www.abim.org/certification/exam-information/rheumatology/exam-tutorial.aspx</u>.

The blueprint can be expanded for additional detail as shown below. Each of the medical content categories is listed there, and below each major category are the content subsections and specific topics that *may* appear in the exam. <u>Please note:</u> actual exam content may vary.



Anatomy, biology, and structure of musculoskeletal tissues	<2%
Joints and ligaments, intervertebral discs, synovium,	
and cartilage	
Connective tissue cells, matrix components, and macromolecul	es
Bone	
Muscles, tendons, and bursae	
Blood vessels	
Nerves	
Immunology	2.5%
Anatomy and cellular elements of the immune system	
Lymphoid organs: gross and microscopic anatomy	
and function	
Organization of immune system: innate and adaptive re	esponses
Specific cell types: ontogeny, structure, phenotype, fund	ction,
and activation markers and cell membrane receptors	
Immune and inflammatory mechanisms	
Antigens: types, structure, processing, presentation,	
and elimination	
Components and regulation of innate immune system	
Major histocompatibility complex: structure, function,	
and nomenclature	
B-cell receptors and immunoglobulins: structure, function	on,
antigen binding, signaling, genetic basis, and effector	function
T-cell receptors: structure, function, antigen binding, sig	gnaling,
and genetic basis	
Receptor-ligand interactions, adhesion molecules, comp	olement
receptors, Fc receptors, and signal transduction	
Complement and kinin systems: structure, function, and	d regulation
Acute-phase reactants and enzymatic defenses	
Cellular interactions, immune regulation, and immunomodulat	ion
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	
Mechanisms of immune tolerance	



Immune responses

IgE-mediated: acute- and late-phase reactions Immunoglobulin-mediated: opsonization, complement fixation, and antibody-dependent cellular cytotoxicity Immune complex-mediated: physiochemical properties and clearance of immune complexes Cell-mediated: cells and effector mechanisms in cellular cytotoxicity and granuloma formation 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 Cellular and molecular mediators Proteases and collagenases

Research principles in basic and clinical investigation

<2%

Design of experimental protocols, clinical trials, and

outcomes research

Outcome assessment techniques: scales, guestionnaires,

performance-based and capacity-based measurements,

health status, disease activity, and functional assessment

Controls, validity, reliability, and responsiveness

Other design of experimental protocols, clinical trials, and outcomes research

Principles of epidemiology and health services research

Prevalence and incidence

Measurement of disease frequency

Application of epidemiologic data

Data analysis, biostatistics, meta-analysis, and medical informatics

Principles of quality assessment and improvement

Ethical and legal issues

Bioethics of basic research and clinical trials

Patient rights and confidentiality

Laboratory and research techniques

Serologic: enzyme-linked immunosorbent assay (ELISA),

radioimmunoassay (RIA), radial immunodiffusion (RID),



nephelometry, immunoblots, protein electrophoresis, and circulating immune complex assays Cellular: lymphocyte proliferation, flow cytometry Histochemistry and immunofluorescence of biopsied tissues Molecular: Northern, Southern, and Western blotting, polymerase chain reaction, genetic mapping techniques, gene sequencing, and gene expression analysis Monoclonal antibody production Transgenic and gene knockout animals Principles of genetic and proteomic analysis: genetic epidemiology, gene transcription, and protein expression analysis **Clinical analysis** Synovial fluid analysis Appropriate use and interpretation of serologic, chemical, biochemical, and microbiologic laboratory tests Diagnostic imaging techniques Plain radiographs: in the assessment of normal and diseased joints, bones, and periarticular structures and prosthetic joints Computed tomography, magnetic resonance imaging, radionuclide scanning, bone densitometry, and arteriography: principles of imaging of joints, bones, and periarticular structures and tissues Ultrasonography: principles of imaging of joints and periarticular structures and tissues Electromyograms and nerve conduction studies: indications for and interpretation of results Biopsy and pathology: diagnostic interpretation of pathologic specimens of specific tissues Pharmacology: dosing, pharmacokinetics, metabolism, mechanisms of action, adverse effects, and drug interactions 2.5% Nonsteroidal anti-inflammatory drugs Glucocorticoids: topical, intra-articular, and systemic Systemic antirheumatic drugs Conventional synthetic disease-modifying antirheumatic drugs (DMARDs)



Targeted synthetic DMARDs Biologics Urate-lowering therapy Agents for bone health Opioid and nonopioid analgesics Colchicine Plasma exchange Vaccines Intravenous immunoglobulin (IVIG) Vasodilator medications Anti-fibrotic agents

Crystal-induced Arthropathies

Spine

Gout 2.5% Primary gout Asymptomatic hyperuricemia Acute gout Intercritical periods Tophaceous gout Conditions associated with gout Lead intoxication Secondary gout Calcium pyrophosphate dihydrate deposition (CPPD) <2% Familial Secondary to primary metabolic disorders **Idiopathic CPPD** <2% Basic calcium phosphate crystal deposition **Infections and Related Arthritides** 6% of Exam Infections 5% Bacterial (nongonococcal and gonococcal) Native joint Prosthetic joint

B

5% of Exam

	Bone	
	Soft tissue	
	Mycobacterial	
	Spirochetal (syphilis, Lyme disease)	
	Viral (human immunodeficiency virus [HIV], hepatitis B virus,	
	hepatitis C virus, parvovirus, chikungunya virus, and others)	
	Fungal	
	Parasitic	
	Whipple disease	
Related	arthritides	<2%
	Acute rheumatic fever and poststreptococcal arthritis	
	Arthritis associated with bacterial endocarditis	
	Postimmunization arthritis	

Metabolic Bone Disease	5.5% of Exam
Low bone mass	<2%
Osteoporosis	2.5%
Primary	
Postmenopausal	
Male	
Secondary	
Other causes of bone loss	<2%
Paget disease of bone	<2%
Bone disease related to renal disease	<2%
Osteomalacia	<2%

Osteoarthritis and Related Disorders	5% of Exam
Osteoarthritis	3%
Diffuse idiopathic skeletal hyperostosis (DISH)	<2%
Hypertrophic osteoarthropathy	<2%
Malignant and nonmalignant tumors of bones, tendons, and joints	<2%
Benign tumors	
Malignant tumors	
Osteonecrosis	<2%



Rheumatoid Arthritis	13% of Exam
Seropositive rheumatoid arthritis	8%
Early disease	
Established disease	
Feltys syndrome	
Seronegative inflammatory polyarthritis	<2%
Complications of established disease	4%
Extra-articular manifestations	
Cardiovascular disease: atherosclerotic cardiovascular	
disease and congestive heart failure	
Malignancy	
Vasculitis	
Immunologic considerations	

Spondyloarthritis	6.5% of Exam
Axial spondyloarthritis	2%
Ankylosing spondylitis	<2%
Skeletal manifestations	
Extra-articular manifestations	
Nonradiographic axial spondyloarthritis	<2%
Skeletal manifestations	
Extra-articular manifestations	
Reactive arthritis	<2%
Skeletal manifestations	
Extra-articular manifestations	
Arthritis associated with inflammatory bowel disease (IBD)	<2%
Skeletal manifestations	
Extra-articular manifestations	
Psoriatic arthritis	2%
Skeletal manifestations	
Extra-articular manifestations	
Arthritis associated with other skin diseases	<2%
SAPHO syndrome (synovitis, acne, pustulosis,	
hyperostosis, and osteitis)	
Peripheral spondyloarthritis	<2%
Skeletal manifestations	
Extra-articular manifestations	



Othe	er Rheumatic and Connective Tissue Disorders (ORCT)	16.5% of Exam
	Paynaud nhonomonon	~ 29/
		NZ /0
	Fillidiy	
	Secondary Drimorily fibrosing rhoumatic dispasses	20/
		3%
	Systemic scierosis	
	Skin	
	Gastrointestinai	
	Pulmonary	
	Renal	
	Scieroderma mímics	
	Eosinophilic fasciitis	
	Retroperitoneal fibrosis (Ormond disease)	20/
	Myopathies	3%
	Idiopathic inflammatory myopathies	
	Metabolic myopathies	
	Medication-associated	
	Critical illness-associated	
	Sjögren syndrome	<2%
	Primary antiphospholipid antibody syndrome	<2%
	Skin-associated rheumatic diseases	<2%
	Erythema nodosum	
	Other forms of panniculitis	
	Multicentric reticulohistiocytosis	
	Fever-associated rheumatic disorders	<2%
	Autoinflammatory disorders	
	Adult-onset Still disease (AOSD)	
	Hemophagocytic lymphohistiocytosis and macrophage	
	activation syndrome (HLH/MAS)	
	Joint-associated rheumatic diseases	<2%
	Polymyalgia rheumatica (PMR)	
	Remitting seronegative symmetric synovitis	
	with pitting edema (RS3PE)	
	Palindromic rheumatism	
	Miscellaneous rheumatic disorders	2%
	Autoimmune hearing loss	



Autoimmune eye disease
IgG4-related disease
Relapsing polychondritis
Overlap syndromes
Undifferentiated connective tissue disease
Mixed connective tissue disease
Autoimmune encephalitis
VEXAS
Pediatric disorders
Juvenile idiopathic arthritis (JIA)
Childhood disease
Complications in adulthood
Kawasaki disease (KD)
Juvenile dermatomyositis (JDM)
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	9% of Exam
Drug-induced	<2%
Cutaneous	<2%
Isolated	
In systemic disease	
Systemic	7.5%
Renal	
Immune-mediated glomerular and tubular dis	sease
Antiphospholipid antibody syndrome and mic	croangiopathies
Renal insufficiency and hypertension	
Urologic complications	
Neurologic	
Central nervous system: inflammatory, vaso-	occlusive,
microangiopathies, and others	
Spinal cord	
Peripheral nerves	
Neuromyelitis optica	



2%

Pulmonary

Pneumonitis

Thromboembolism

Pulmonary hypertension

Cardiovascular

Myocardial disease

Valvular disease

Accelerated atherosclerosis

Serositis

Pleuritis

Pericarditis

Peritonitis

Hematologic

Autoimmune cytopenias

Hemolytic uremic syndrome (HUS) and thrombotic

thrombocytopenic purpura (TTP)

Autoimmune clotting factor deficiencies (overlap with

antiphospholipid antibody syndrome)

Musculoskeletal

Joints, tendons, and ligaments

- Muscle disease
- Lupus in pregnancy

Neonatal lupus

Vasculitis

Antiphospholipid antibody syndrome (APS)

Clinical features excluding pregnancy

Pregnancy

Catastrophic APS

Nonarticular and Regional Musculoskeletal Disorders

Diffuse pain syndromes

Fibromyalgia

Complex regional pain syndrome (reflex sympathetic dystrophy)

Medication-induced diffuse pain

7% of Exam

<2%



Regional musculoskeletal disorders

Axial syndromes Back pain Neck pain Thoracic outlet syndrome Shoulder disorders Joint Soft tissue 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 Leg disorders **Neuropathies** Axial disorders Peripheral disorders **Entrapment neuropathies** Mononeuritis multiplex Polyneuropathy Small fiber neuropathy

Nonrheumatic Systemic Disorders

<2%

9% of Exam

<2%



Hereditary, congenital, and inborn errors of metabolism

Osteogenesis imperfecta	
Ehlers-Danlos syndromes including hypermobility	
Mucopolysaccharidoses	
Osteochondrodysplasias	
Multiple epiphyseal dysplasia	
Spondyloepiphyseal dysplasia	
Inborn errors of metabolism affecting connective tissue	
Homocystinuria	
Ochronosis	
Storage disorders	
Immunodeficiencies	<2%
Immunoglobulin A (IgA) deficiency	
Complement component deficiencies	
Common variable immunodeficiency	
Metabolic-associated rheumatic disorders	2.5%
Diabetes mellitus	
Acromegaly	
Thyroid disease	
Cushing disease	
Parathyroid disease	
Renal failure and dialysis	
Hematologic and oncologic malignancy-associated	
rheumatic disorders	3%
Amyloidosis	
Primary	
Secondary	
Hereditary	
Lymphoma	
Myelodysplastic syndromes	
Leukemia	
Solid tumors	
Plasma cell dyscrasias	
Hemoglobinopathies	
Sickle cell	
Hemophilias	



Arthritic and rheumatic disorders

Hemochromatosis Myositis ossificans progressiva Wilson disease Sarcoidosis Scurvy Pancreatic disease Primary biliary cholangitis Cystic fibrosis Graft-versus-host disease Celiac disease Drug-associated Environmental agent-associated Neurologic <2% Amyotrophic lateral sclerosis (ALS) Neuropathic arthropathy

Vasculitides	8.5% of Exam
Large-vessel vasculitis	<2%
Takayasu arteritis	
Giant cell arteritis	
Medium-vessel vasculitis	<2%
Polyarteritis nodosa	
Small-vessel vasculitis	2.5%
Antineutrophil cytoplasmic antibody (ANCA)-associated vasculitis	5
Granulomatosis with polyangiitis	
Microscopic polyangiitis	
Eosinophilic granulomatosis with	
polyangiitis (Churg-Strauss)	
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	<2%
Behçet disease	
Cogan syndrome	
Single-organ vasculitis	<2%
Cutaneous leukocytoclastic angiitis	
Cutaneous arteritis	
Primary central nervous system angiitis	
Isolated aortitis	
Vasculitis associated with probable etiology	<2%
Hepatitis C virus-associated cryoglobulinemic vasculitis	
Hepatitis B virus-associated vasculitis	
Syphilis-associated aortitis	
Drug-induced vasculitis	
Drug-induced ANCA-associated vasculitis	
Drug-induced immune complex vasculitis	
Other drug-induced vasculitis	
Cancer-associated vasculitis	
Vasculitis mimickers	<2%
Buerger disease (thromboangiitis obliterans)	
Cholesterol emboli	
Fibromuscular dysplasia	
Segmented arterial mediolysis	
Warfarin necrosis	
Reversible cerebral vasoconstriction syndrome	
Moyamoya disease	
Atrial myxoma	
Endocarditis	
Calciphylaxis	
Amyloid angiopathy	

Miscellaneous Topics

Arthrocentesis and injections

Anatomy Precautions Potential sequelae

<2%

2% of Exam



General clinical care

Rehabilitation in rheumatic diseases	
Exercise	
Therapeutic modalities	
Thermal modalities	
Adaptive equipment and assistive devices	
Footwear and orthotics	
Functional status and disability determination	
Pain management	
Physiology of pain	
Opioid contract	
Psychosocial aspects of rheumatic diseases	
Psychological and emotional factors including sexuality	
Economic and vocational issues	
Perioperative management of rheumatic diseases	
Nutrition	
Complementary and alternative practices	
Treatment adherence	<2%
Barriers	
Health literacy	
Professionalism and ethical behavior	<2%
Ethical dilemmas	
Professionalism	
Communication	
Interpersonal communication skills	
Use of medical interpreters	

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<2%