RHEUMATOLOGY OVERVIEW

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What is Rheumatology?

- Medical science devoted to autoimmune diseases, particularly connective tissue disease, and certain musculoskeletal disorders
- "rheuma" "a substance that flows"
- "rheumatism" emphasis that arthritis could be a systemic disorder

Rheumatology Classification

(shortened list)

- Systemic connective tissue disease
- Vasculitis
- Seronegative spondyloarthropathies
- Arthritis associated with infection
- Inflammatory Myopathy
- Rheumatic disorders associated with metabolic, endocrine, and hematologic disease

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Anatomy of a Joint PERIOSTEUM SUBCHONDRAL BONE PLATE SYNOVIUM ARTICULAR CARTILAGE

Diagnostic Approach

- · Articular vs non-articular
- Mechanical vs inflammatory
- Poly- vs oligo- vs monoarticular
- Acute vs chronic
- · Localized vs systemic

Inflammatory vs Non-inflammatory

- Erythema
- Warmth
- Pain
- Swelling
- -
- Prolonged stiffness
- Systemic symptoms
- · Laboratory abnormalities

- Mechanical pain (worse with activity)
- Improves with rest
- Stiffness after brief periods of rest (not prolonged)
- Absence of systemic signs

Diagnostic Approach • Articular vs non-articular · Mechanical vs inflammatory • Poly- vs oligo- vs monoarticular · Acute vs chronic • Localized vs systemic **Differential Diagnosis** • Monoarticular inflammatory - trauma, hemarthrosis, spondyloarthropathy - Septic arthritis, crystal induced Oligoarticular - Spondyloarthropathy, crystal induced, infection related Polyarticular - RA, SLE, crystal induced, infectious Physical Exam • Inspection • Palpation • Maneuvers

Rheum Diseases You Will Encounter

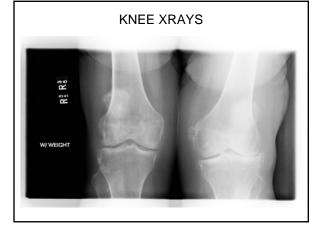
- Osteoarthritis
- Rheumatoid Arthritis
- Seronegative spondyloarthropathy
- · Crystal induced arthritis
- Systemic lupus erythematosus and related connective tissue diseases
- Vasculitis
- Idiopathic Inflammatory Myopathy



OSTEOARTHRITIS

- Most common form of arthritis
- > 50 years of age
- Risk factors: age, obesity, occupation, history of trauma
- Most common sites: hands, feet, knees, hips, AC joints, and facet joints of the cervical and lumbosacral spine
- PAIN (mechanical type), stiffness (< 30 minutes), loss of function are presenting features
- No systemic involvement
- DIP/PIP involvement; spares the wrists (Heberden's/ Bouchard's)
- Non-inflammatory synovial fluid

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RHEUMATOID ARTHRITIS

- Chronic (>6 wks), inflammatory
- Female > Male
- AM stiffness lasting at least 1 hr
- Soft- tissue swelling in at least 3 joint areas simultaneously Including wrist, MCP, or PIP joint Symmetric
- Swan neck/Boutonniere/ulnar deviation , erosive
- Rheumatoid nodules, extra-articular manifestations
- Positive rheumatoid factor , anti-CCP



RHEUMATOID ARTHRITIS

RHEUMATOID ARTHRITIS



Seronegative Spondyloarthropathy

- Seronegative
- Oligoarticular, assymetric
- Chronic, inflammatory
- Sacroiliac involvement
- Enthesopathy
- Spinal involvement (inflammatory)
- HLA B27

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Seronegative Spondyloarthropathies

- Ankylosing spondylitis
- IBD associated arthropathy
- Psoriatic arthritis
- · Reactive arthritis
- Undifferentiated spondyloarthropathy

Dactylitis (Sausage Toes)





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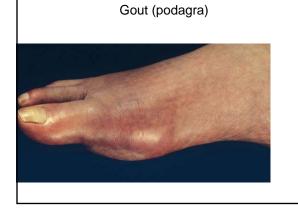




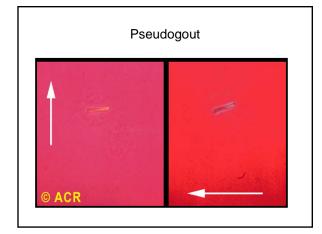
GOUT

- · Recurrent, episodic inflammatory arthritis
- Peak of pain: 24 hours; subside in 3-10 days
- 75 % of initial attacks in 1st MTP joint (podagra)
- Usually monoarticular, may be polyarticular
- Hyperuricemia may or may not be present
- Predisposing factors and associated conditions: surgery, medications (DIURETICS, low dose aspirin, cyclosporine A), alcohol ingestion, hypertension, renal insufficiency, hyperlipidemia

Gout (diagnosis) A B C



Tophaceous gout



Connective Tissue Diseases

- Systemic Lupus Erythematous
- Sjögren's Syndrome Sicca symptoms, +SSA/SSB
- <u>Scleroderma</u>
- Mixed Connective Tissue Disease features of SLE and Scl, +RNP
- Overlap/ Undifferentiated Connective Tissue Disease

Systemic Lupus Erythematosus

- Malar Rash
- Discoid Rash
- Serositis
- Oral ulcers
- Arthritis
- Photosensitivity
- Blood disorder
- Renal disorder
- ANA*
- Immunologic abnormalities
 - (anti-Smith antibody, anti-double stranded DNA, anti-phospholipid antibodies)
- Neurologic symptoms



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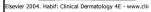
Scleroderma

- Sclero = thickened, derma = skin
- Systemic : Limited or Diffuse
 - Limited = CREST (calcinosis, Raynaud's, Esophageal dysmotility, Sclerodactyly, Telangiectasias)
 - Diffuse = scleroderma proximal to MCPs
 - Pulmonary (ILD, pHTN), Renal involvement (renal crisis)
- Localized
- Overlap syndrome











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Vasculitis

- Inflammation & necrosis of blood vessel
- Perforation & hemorrhage, thrombosis, ischemia
- Large vessel
 - Takayasu, Giant Cell Arteritis
- Medium vessel
 - Polyarteritis nodosa, Kawasaki's
- Small
 - Henoch-Schonlein purpura, Wegener's granulomatosis, Microscopic polyangiitis, Churg-Strauss
 - ANCA -antineutrophil cytoplasmic antibodies*



Giant Cell Arteritis

- Patients >50 y/o
- Cranial symptoms—superficial HA, scalp tenderness, jaw claudication, blindness
- Polymyalgia rheumatica—pain and stiffness of proximal joints
- Fever, systemic symptoms
- · Elevated ESR and CRP
- Diagnosis: Biopsy of temporal artery

Idiopathic Inflammatory Myopathy

- Polymyositis
- Dermatomyositis
- Inclusion body myositis
- · Proximal muscle weakness
 - (muscle pain not a typical symptom)
- Elevated muscle enzymes: CK, Aldolase, LDH
- Diagnosis: biopsy
- Lung involvement, increased risk for malignancy



Case 1	
 67 y/o man with DM, HTN, hyperlipidemia, h/o arthritis (unclear what kind)and EF of 25%. He came with SOB and signs of CHF. Treated with IV Lasix -> he did well, SOB improved, lungs are now clear 	
Pt is ready for discharge but now complains of severe right knee pain	
 Examination: his right knee is flexed and looks much bigger than the left; you touch it and it feels warm; you try to move it but patient is in too much pain and you cannot flex or extend it further. Also, you look at the vital signs and the last temperature checked at 6 am showed 37.8 °C 	
and the last temperature checked at 6 am showed 37.8 °C	
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What are your thoughts?	
Crystal induced arthritis?	
2. Infection?	
Acute trauma?	
4. Hemorrhage?	
	1
What test needs to be done?	
ARTHROCENTESIS OF RIGHT KNEE	

		Synovial	Fluid Analy	/sis	
C	ell Count				
C	rystal analy	ysis			
G	ram Stain	and culture			
_		To a	T.,	T	
		Normal	Non- inflammatory	Inflammatory	Septic
-	WBC	0-200	200-2000	2000 – 50 K	> 50 K
Ī	% PMN	25%	50%	75%	> 90%
L					
			Case 2		
	Scenario	: Loyola Rheumat	ology Clinic		
	CC: joint	pain			
•	she attrib	uted the pain to th	mplains of bilateral he he fact that she was	taking too many n	otes in
	minimal r	elief at this point.	kles and right wrist. She denies swelling	but has trouble to	
	engagem	ent ring off. Also	with oral ulcers for t	he last 6 months	
•			s, weight loss, Rayna		since her
		_			
 PE: +2 oral ulcers, malar rash, mild synovitis of right wrist, 2nd and 3rd PIPs bilaterally 					
			Case 3		
•	Scenario	: Hines Arthritis Cl	linic		
•	CC: joint	pain			
•	HPI: 29 y	/o Hispanic man o	complains of pain in oints include: right sl	his joints x 7 years	(since he
	ankle. He	states that pain i	s worse when he way years. Despite his j	akes up. He believe	es his left
	to work o	ut 5 x week. He fe	eels that being active	e helps with his syr	nptoms.
•	ROS: low least 2 ho		his military time; mo	rning stiffness that	lasts at
			und his left ankle with	h doorooca BOM	a that icint
•	r ⊑. very	ımıu sweiling arot	and monent ankle with	ii ueciease KOM II	ı uıat JUIIII

Case 4

- Scenario: Loyola Gen Med Clinic
- CC: headaches
- HPI: 74 y/o male with 3 weeks headache. Has pain at temporal areas, which are also painful with combing hair. Has had low grade fevers for last 2 weeks and loss of energy. For the last two days, has noticed decreased vision in his right eye.
- PE: afebrile. Decreased temporal artery pulse. Temporal artery tender to touch.