

# MSK I: Arthritis/Rheum Cheat Sheet

by ksellybelly via cheatography.com/19318/cs/2449/

#### Osteoarthritis

# Definition

Progressive loss of articular cartilage with reactive changes in the bone, results in pain and joint destruction (most common arthropathy in adults)

#### Clinical Features

Decreased ROM, joint crepitus, pain worsening througout the day

# Common sites

DIP joint (Heberden's nodes + PIP joint (Bouchard's nodes), and wrist/hip/knee/spine

#### **Imaging**

Asymmetric narrowing or joint spaces, subchondral sclerosis, cysts, marginal osteophytes

# Treatment

Weight reduction, physical actibity, acetaminophen, NSAIDs, intra-articular steroids. Total joint replacement in advanced cases.

# Rheumatoid Arthritis

# Definition

A chronic autoimmune disease with synovitis affecting multiple joints and other systemic extra-articular manifestations. Joint destruction ultimately occurs.

# Typical patient

F>>M, 40-60yo at onset (juvenile in pts <16yo)

# Clinical Features

Morning stiffness, symmetric, subcutaneous nodules, RF level >95th percentile, arthritis of hand joints, softtissue swelling (DIP joints spared)

# Extra-articular manifestations

Osteoporosis, changes in skin/lungs/kidneys/eyes/liver/bloodstream/hear t

# Rheumatoid Arthritis (cont)

Lab Studies

Elevated ESR + CRP, RF and anti-CCP positive in up to 80%

#### Treatment

PT + NSAIDS + DMARDS (MTX, corticosteroids, biologics) and reconstructive surgery for severe cases.

#### Gout

#### Definition

A systemic disease of altered purine metabolism and subsequent sodium urate crystal precipitation into synovial fluid

# Typical patient

M>>W (until menopause, then 1:1)

Clinical Features of Initial Attack

Metatarsal phalangeal joint of the great toe (podagra)

Symptoms around involved joint

Pain, swelling, redness, exquisite tenderness

Substance that may form adjacent to the joint, diagnostic!

Tophi (chalky deposits of uric acid)

# Lab Studies

Joint fluid shows rod-shaped, negatively birefringent urate crystals seen. Serum uric acid level often > 8 mg/dL

# Lifestyle Modifications

Elevation, dietary modifications (avoid purines and EtOH)

# Pharmacotherapy

NSAIDs (indomethacin), corticosteroid injections, colchicine in between attacks

# Polymyalgia Rheumatica (PMR)

#### Definition

Syndrome with pain and stiffness in the neck/shoulder/pelvic girdles and is accompanied by constitutional symptoms (fever, fatigue, weight loss, depression)

#### Etiology

Unknown (F>>M)

Associated with

Temporal arteritis (30% of cases)

# Clinical features

Stiffness, worse after rest and in the morning. MSK symptoms are bilateral, proximal, symmetrical

#### Must r/o...

Giant Cell Arteritis (scalp tenderness, jaw claudication, headache, temporal artery tenderness-->can lead to vision loss)

# Lab Studies

ESR elevated (>50mm/hr)

# Treatment

Low-dose corticosteroids (higher doses if GCA), might need to be on for 2 years

# Sjogren's Syndrome

# Definition

An autoimmune disease that destroys the salivary and lacrimal glands (exocrine glands)

May be a secondary complication to preexisting disorders like...

RA, SKE, polymyositis, scleroderma

Classic Patient

Middle-aged females

# Clinical Features

*Mucus membranes* most affected. Parotid glands might be enlarged.



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# Sjogren's Syndrome (cont)

Characteristic features of primary Sjogren's

Dry mouth (xerostomia) + dry eyes (seropthalmia or keratoconjunctivitis)

#### Lab Studies

RF in 70% of cases, ANA in 60%, anti-Ro Abs in 60%, anti-La in 40%. Schirmer's tear test wetting of <5mm of filter paper in lower eyelid in 5min = + for decreased secretions

How to confirm lymphocytic infiltrate and gland fibrosis

Lip bx

#### Treatment

Mainly symptomatic management, goal of keeping mycosal surfaces moist. Can give artificial tears and saliva, increased oral fluid intake, and lubricants for eyes/vagina. Pilocarpine may increase saliva flow

# Childhood-onset idiopathic arthritis

#### Definition

Characterized by chronic synovitis and extra-articular manifestations (fever, rash, weight loss)

# Typical Patient

F>>M, at age 1-3yo (males older 8-12yo)

# Forms of arthritis

Pauciarticular (50%) polyarticular (35%), systemic (15%)

# Systemic (JRA) clinical features

Spiking fevers, myalgias, salmon-pink maculopapular rashin evening. Hepatosplenomegaly, lymphadenopathy, leukocytosis, pericarditis, myocarditis

# Pauciarticular clinical features

4 or less medium to large joints. Also at risk for asymptommatic uveitis (can lead to blindness if +ANA)

# Childhood-onset idiopathic arthritis (cont)

Polyarticular clinical features

Resembles adult RA, symmetri, 5+ joints. Systemic sx: low-grade fever, fatigue, rheumatoid nodules, anemia.

# Treatment

PT + NSAIDs

# **Psoriatic Arthritis**

# Definition

An inflammatory arthritis with skin involvement usually preceding joint disease

#### Clinical features

Symmetric, hands and feet. Affects few joints. Pitting of nails and onycholysis. "Sausage-finger" appearance

# Lab Studies

ESR elevated, hyperuricemia if severe skin involvement, "pencil in a cup" deformity on x-ray

#### Treatment

NSAIDs, MTX, reconstructive surgery as last resort

# **Pseudogout**

# Definition

Intra-articular deposition of calcium pyrophosphate dehydrate (CPPD) in peripheral joints

Acute presentations mimic...

Gout (may be recurrent and abrubt)

# Clinical features

Painful inflammation (when crystals shed into joint), most common in knees/wrist/elbow

# Lab Studies

Rhomboid-shaped CPPD crystals, negatively birefringement. Will see chondrocalcinosis in radiographs (fine, linear calcifications)

# Treatment

NSAIDS, colchicine, and intra-articular steroid injections

# **Polymyosotis**

#### Definition

An inflammatory disease of striated muscle affecting the proximal limbs/neck/pharynx (skin can also be

affected=dermatomyositis)

# Etiology

Unknown, but strong association with occult malignancy

Classic patient

F>>M

# Clinical Features

Insidious painless proximal muscle weakness, dysphagia, skin rash (malar or heliotrope), polyarthralgias, muscle atrophy

#### Lab Studies

CPK and aldolase elevated. Muscle bx will show myopathic inflammatory changes

#### Treatment

High-dose steroids, MTX, or azathioprine

# Scleroderma (Systemic Sclerosis)

# Definitions

Characterized by deposition of collagen in the skin, and also lungs, kidney, heart stomach. Unknown etiology.

Classic Patient

F>>M, 30-50yo

# Clinical Features in general

95% of patients have skin involvement, starts with swelling of fingers and hands, may spread to trunk and face. R

Clinical Features: Limited

Mostly affects skin of face, neck, distal elbows and knees. Causes pulmonary hypertension later in disease.

Clinical Features: Diffuse

Affects the skin plus the heart, lungs, GI tract, kidneys



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# Scleroderma (Systemic Sclerosis) (cont)

CREST Syndrome (associated with limited scleroderma

Calcinosis, Raynaud's, esophageal dysfunction, sclerodactyly, telangiectasias

# Initial presentation

Skin changes, polyarthralgias, esophageal dysfunction

#### Lab Studies

+ANA in 90% of pts, +anticentromere Ab assoc. w/ limited scleroderma, watch for HTN

#### Treatment

No cure. Treat specific manifestations of disease (ie PPIs for GERD, ACEis for renal dz, aboid triggers, and immunosuppressives for pulmonary HTN)

# Septic (Infectious) Arthritis

#### Definition

The hematogenous spread of bacteremia infection (osteomyelitis) caused by diagnostic or therapeutic procedure (injection) or infection elsewhere (cellulitis, bursitis)

# Classic joint

Single joint, usually knee (can also be hip, shoulder, ankle)

# Most common pathogen

Staph. aureus

Pathogen in sexually-active young adults

Neisseria gonorrhea

# Clinical Presentation

Acute swelling, fever, joint warmth and effusion, tenderness to palpation, increased pain w/ minimal ROM

# Lab Studies

Collect/aspirate synovial fluid, many will also have a positive blood culture.

# Septic (Infectious) Arthritis (cont)

#### Treatmen

Aggressive IV Abx followed by oral antibiotics, sometimes arthrotomy and arthrocentesis are required.

# Reactive Arthritis (Reiter Syndrome)

#### Definition

A seronegative arthritis that has a tetrad of: urethritis, conjunctivitis, oligoarthritis, and mucosal ulcers (leading cause of nontraumatic monoarthritis)

Can be seen as a sequele to...

STDs or gastroenteritis

# Clinical Features

Asymmetric arthritis in large joints below the waist (knee, ankle), mucocutaneous lesions (balanitis, stomatitis), urethritis, conjunctivitis

#### Typical patient

M>>F after STDs (1:1 ratio after enteric infx)

#### Lab Studies

Up to 80% HLA-B27+, synovial fluid negative culture

# Treatment

PT + NSAIDs. Abx can reduce chance of developing disorder, but they don't alleviate sx of the reactive arthritis

# SLE (Systemic Lupus Erythematous)

# Definition

An autoimmune disorder characterized by inflammation, plus +ANA, and involvement of multiple organs

# Classic Patient

Women of childbearing age, and more in African-American women

# SLE (Systemic Lupus Erythematous) (cont)

# Clinical Features

(4+ criteria including high ANA): malar rash, discoid rash, photosensitivity, oral ulcers, arthritis, serositis, renal dz, ANA, hematologic/immunologic/neurologic disorders

#### What must be ruled out?

Drug-induced lupus/lupus-like syndrome (from INH, hydralazine, quinidine). Usually pt. will have positive antihistone Abs

#### Lab Studies

Get CBC BUN, Cr, U/A, ESR, serum complement (C3 or C4), anti-Smith antibodies to mark for progression. ANA is present 99% of the time, but low titers have a low predictive value.

#### Treatment

Exercise + sun protection, NSAIDs, Antimalarials (hydroxychloroquine), corticosteroids, MTX

# **Polyarteritis Nodosa**

# Definition

Small/medium artery inflammation involving the skin, kidney, peripheral nerves, muscles, and gut

# Classic patient

M>>F, 40-60yo, Hep B pts

# Etiology

Unknown (concurrent in Hep B pts. 30% of the time)

# Clinical Features

Fever, anorexia, weight loss, abdominal pain, peripheral neuropathy, arthralgias, arthritis, skin lesions. If renal involvement: HTN, edema, oliguria, uremia (if renal involvement.

# Lab Studies

Vessel bx or angiography to diagnose (might also see ANCA, elevated ESR and CRP)



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# Polyarteritis Nodosa (cont)

# Treatment

High-dose corticosteroids, cytotoxic drugs, immunotherapy. Might need to also treat for Hep B. Treat HTN if present

# Fibromyalgia Syndrome

# Definition

A central pain disorder; cause and pathogenesis are poorly understood

Can occur concurrently with...

RA, SLE, Sjogren's

# Clinicla Features

Nonarticular MSK aches, fatigue, sleep disturbance, multiple tender points on exam, anxiety, depression, headaches, irritable bowel syndrome, dysmenorrhea, paresthesias

# Lab Studies

Diagnosis of exclusion, must r/o hormonal and vitamin disturbances. Sometimes abnormalities of T-cell subsets

# Treatment

SSRIs, SSNRIs, RCAs. Lyrica is only FDA-approved drug to specifically treat fibromyalgia. Aerobic exercise, stress reduction, and sleep assistance are helpful.



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