



Evaluation and Management of Polyarthrititis?

June 26, 2026

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Disclosures

- Clinical Trial Support
 - Fate Therapeutics
 - GSK
 - Sanofi
 - Astra Zeneca

Case 1

- **CC:** “my back hurts”
- **HPI:** 29 yo male with 4 mos of bilateral buttock and lower back pain. Dull, achy but can be sharp with coughing, sneezing, bending; radiates to mid thigh; 4 hours am stiffness. Pain improves with hot showers, walking.
- **ROS:** No ankle pain, eye redness, STD, rash, bloody diarrhea
- **SH:** No tob/etoh/drugs
- **Labs:** CBC, CMP normal, ESR: 40 mm/hr (<30 mm/hr)
- **Plain films of L spine and pelvis:** ‘degenerative changes of sacroiliac joints’

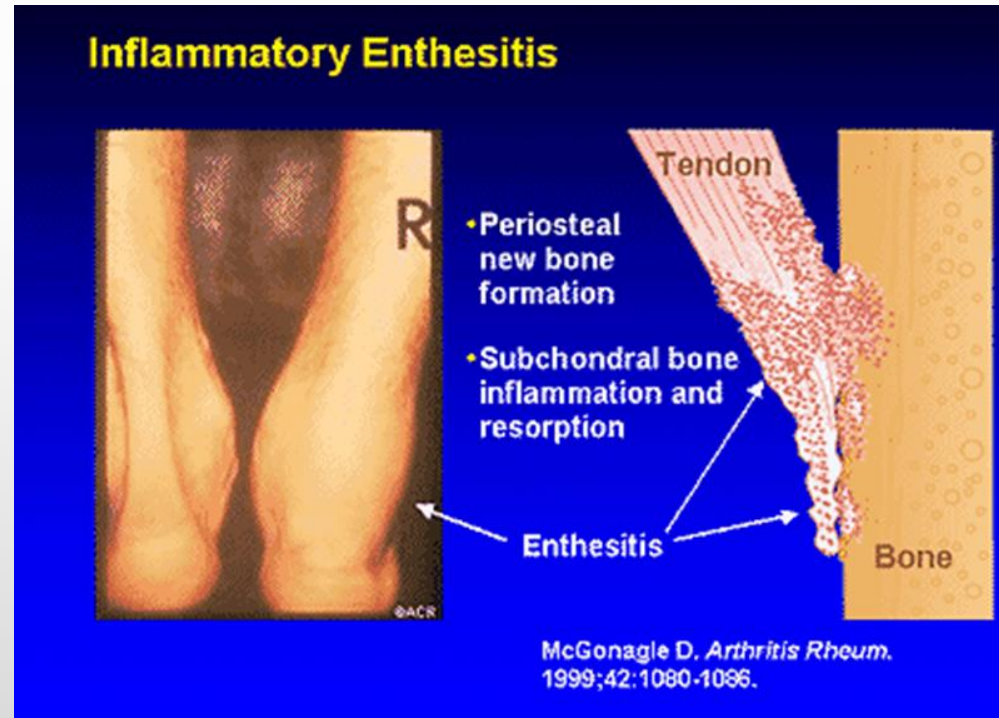


What is the most likely cause of this man's symptoms?

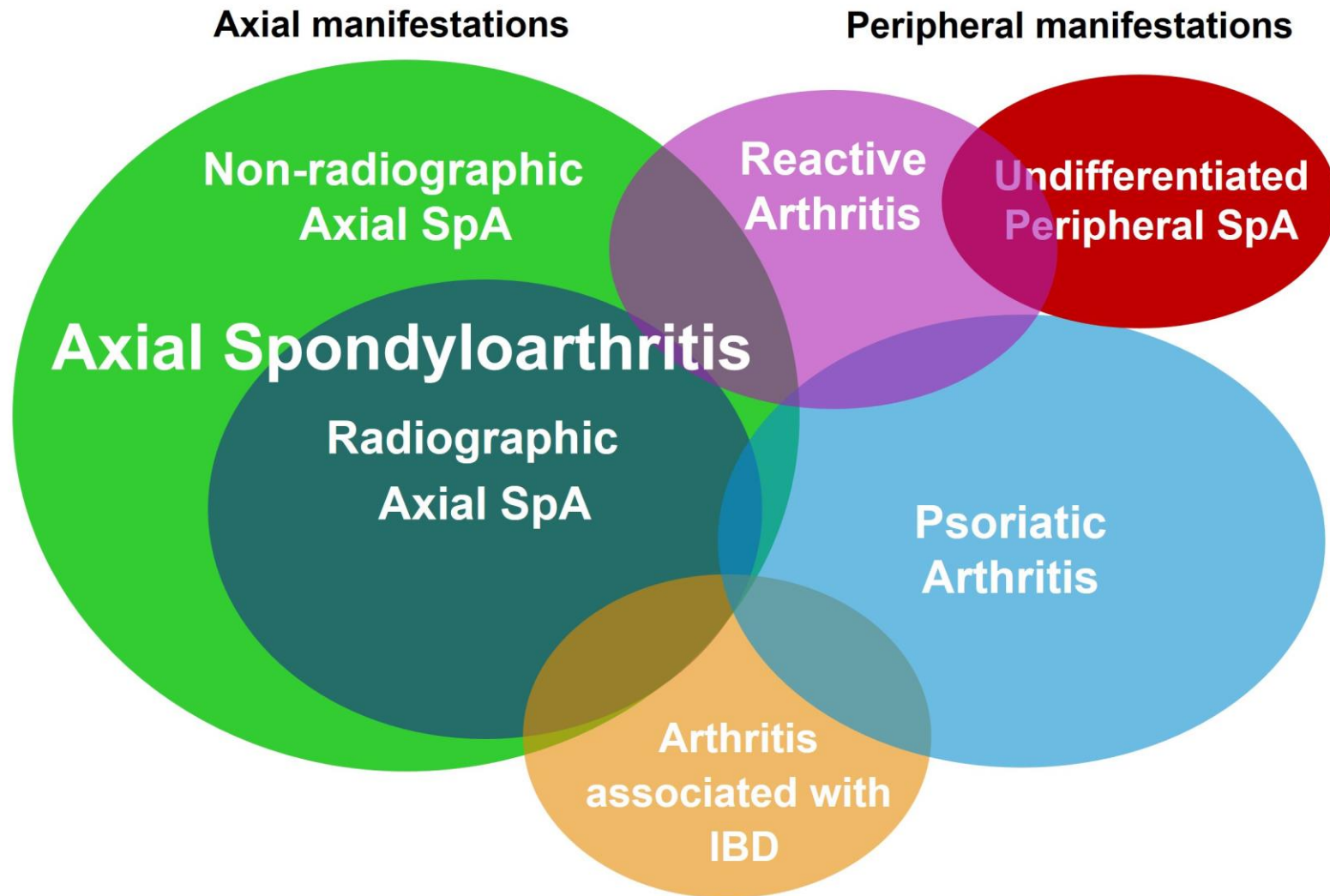
1. Spondylolisthesis
2. Spondylosis
3. Spondylolysis
4. Scoliosis
- ✓ 5. Inflammatory back pain

Axial Spondyloarthritis: General Points

- Highly heterogeneous presentation
 - Inflammatory low back pain
 - Sacroileitis
 - Dactylitis
 - Enthesitis



Spondyloarthritides (SpA)



ASAS Inflammatory Back Pain Criteria by Experts (Chronic Back Pain; n=648)

- age at onset < 40 years
- insidious onset
- improvement with exercise
- no improvement with rest
- pain at night (with improvement upon getting up)

Sensitivity: 79.6%; Specificity: 72.4%

Inflammatory back pain present if at least 4 out of 5 parameters are fulfilled.

ASAS Classification Criteria for Spondyloarthritis (SpA)

In patients with ≥ 3 months back pain and age at onset < 45 years

In patients with peripheral symptoms ONLY

Sacroiliitis on imaging plus ≥ 1 SpA feature

OR

HLA-B27 plus ≥ 2 other SpA features

Arthritis or enthesitis or dactylitis plus

SpA features

- inflammatory back pain (IBP)
- arthritis
- enthesitis (heel)
- uveitis
- dactylitis
- psoriasis
- Crohn's/colitis
- good response to NSAIDs
- family history for SpA
- HLA-B27
- elevated CRP

≥ 1 SpA feature

- uveitis
- psoriasis
- Crohn's/colitis
- preceding infection
- HLA-B27
- sacroiliitis on imaging

OR

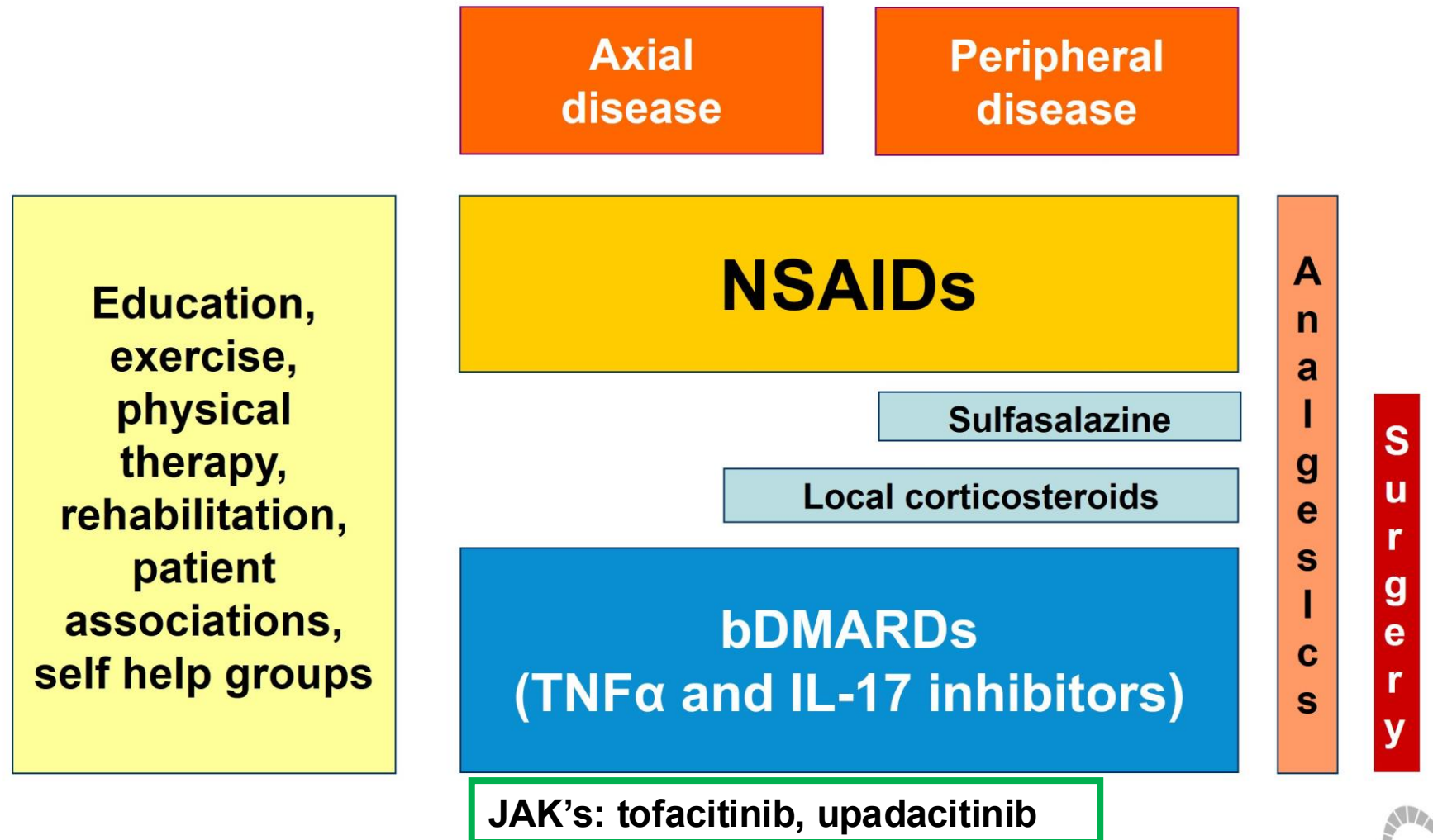
≥ 2 other SpA features

- arthritis
- enthesitis
- dactylitis
- IBP ever
- family history for SpA

Sensitivity: 79.5%, Specificity: 83.3%; n=975



ASAS-EULAR Recommendations for the Management of Axial Spondyloarthritis



Case #2

- CC: finger pain, swelling
- HPI: 84 yo female with known OA complains of chronic aching and swelling of DIPs – gradually over past year; episodes of warmth and tenderness that comes and goes, but no pain in feet, knees or swelling elsewhere.
- PMH: hypertension, DMII, no psoriasis, crohns
- Meds:
 - hydrochlorothiazide 25 mg daily
 - metformin 500 mg bid
- Exam:
 - Moderate heberden and bouchard nodes, moderate mal-alignment, but with no MCP synovitis
 - soft tissue swelling in DIPs of L hand



What is the best explanation for these symptoms?

- A. Erosive osteoarthritis
- ✓ B. Elderly onset gout
- C. Calcium pyrophosphate deposition disease
- D. Rheumatoid arthritis
- E. Amyloid arthritis

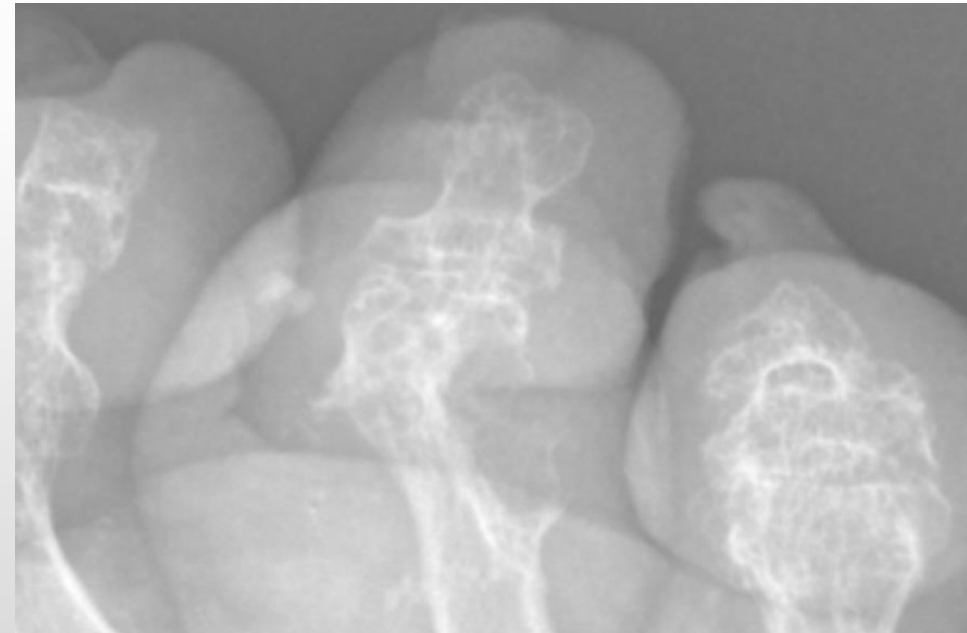
Gout:

- Common in men >40 yrs but **rising in women**
 - ↑prevalence (~4%)
 - chronic disease, aging, ↓GFR
 - ↑obesity
 - medications
- Uric acid often **deceptively low/normal during attacks** (our pt: 11.3 mg/dL), adjusting treatments can **precipitate attacks**
- Phases:
 - acute
 - chronic tophaceous →
 - intercritical period



Elderly Onset Gout (EOG)

- subacute/chronic polyarticular hand involvement
- localization of tophi on heberden nodes
- increased female:male ratio
- association with drugs → decrease renal urate excretion (diuretics and low-dose aspirin)



Treatment Principles

- Terminate the acute attack
 - High dose NSAIDs
 - Corticosteroids (PO or intra-articular with arthrocentesis)
 - Colchicine, if early
 - FDA indication 2009
 - 1.2 mg then 0.6 an hour later then qd-tid
- Treat hyper-uricemia to limit progression
 - Lower serum uric acid
 - Xanthine oxidase inhibitors
 - Allopurinol
 - Febuxostat (2009)
 - Pegloticase (2011)

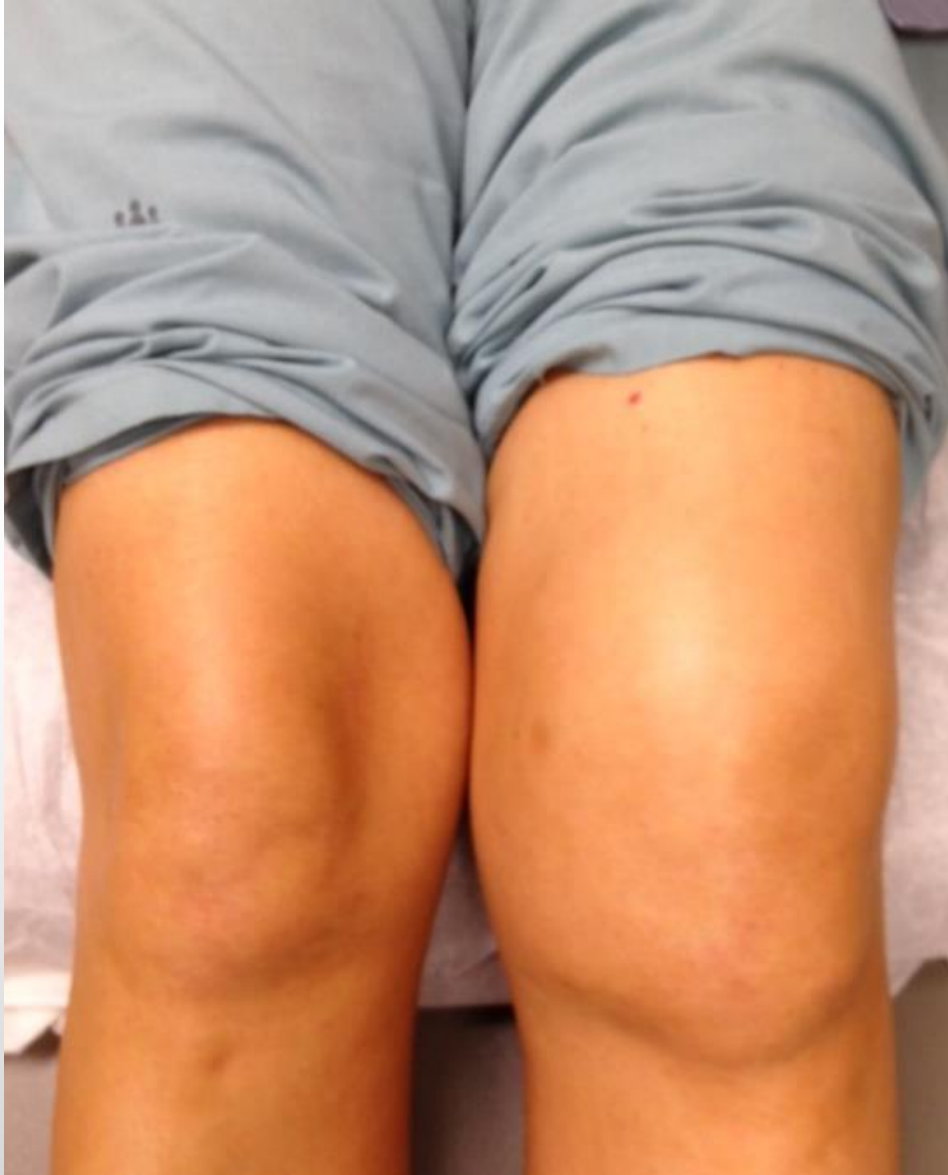
Gout: Need to know...

- Gout prevalence is on the rise (acute, chronic, EOG)
- Risk factors are traditional (diet) - and more recently recognized
- Serum uric acid is not always elevated during attacks
- Newer approaches to treatment:
 - Colchicine dosing
 - Start allopurinol in low dose; titrate gradually
 - Febuxostat if allopurinol intolerant, ckd
 - Pegloticase for severe tophaceous gout



Case #3

- A 74 year old woman presents with 2 days of spontaneous L knee swelling and moderate pain with weight bearing. Symptoms are worsening, though improved with OTC ibuprofen. She reports mild intermittent low grade fevers and occasional chills. No prior episodes.
- PMH:
 - Psoriasis
 - Diabetes mellitus with CKD
- SH:
 - 1 ppd tobacco x 20 yrs
 - Moderate alcohol x yrs




Exam:

- T: 100.2, rr: 16, hr: 98, bp: 110/70
- Gen: mild distress with L knee exam
- Msk: mod pain elicited with passive ROM testing in all fields; mild erythema and tenderness with moderate effusion in suprapatellar bursa
- WBC: 14K
- CRP: 79.6
- ESR: 64



What is the most likely cause of her swelling?

- A. Psoriatic arthritis
- B. Gout
- C. Osteoarthritis
-  D. Calcium pyrophosphate deposition
- E. Polymyalgia rheumatica

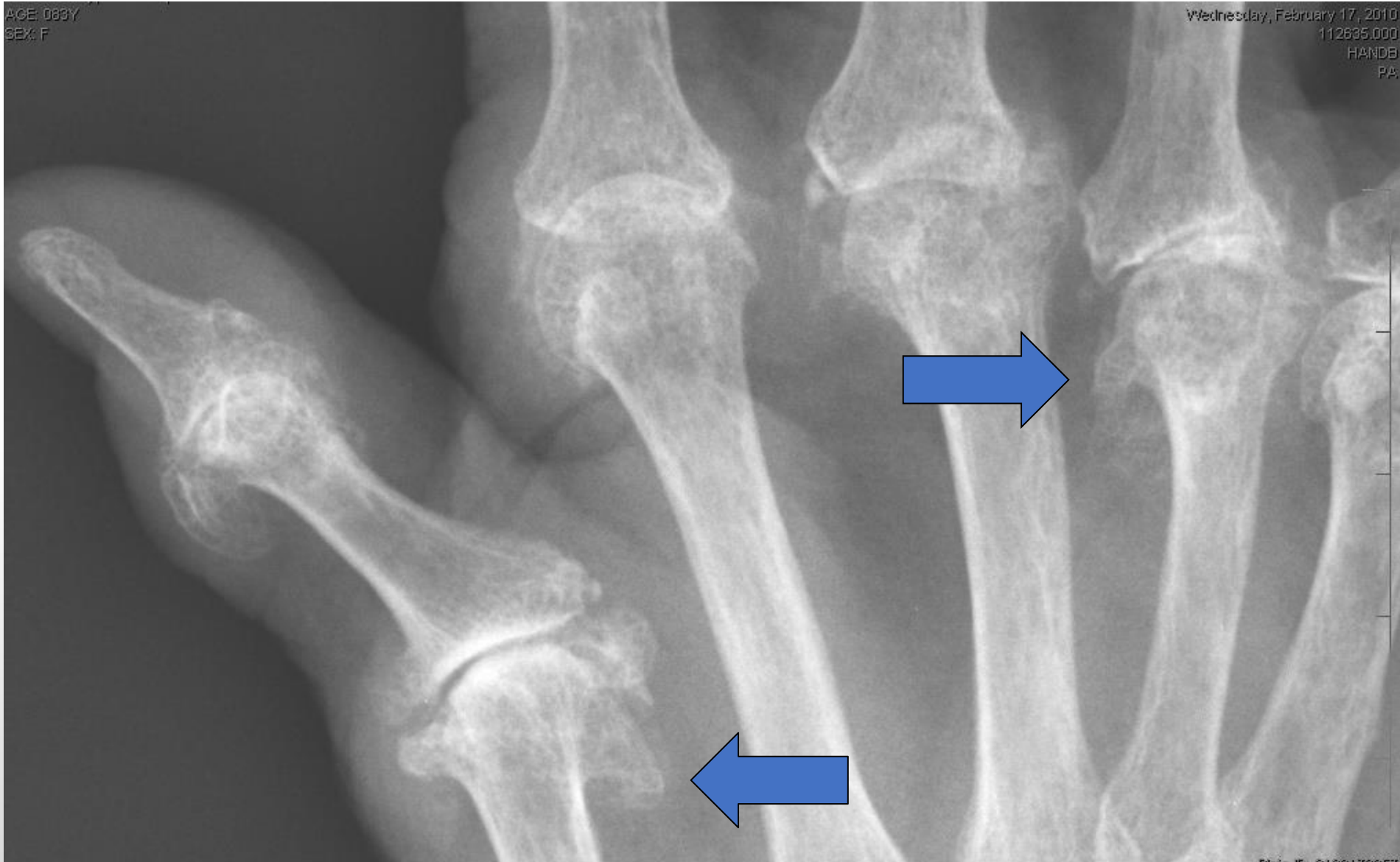


chondrocalcinosis



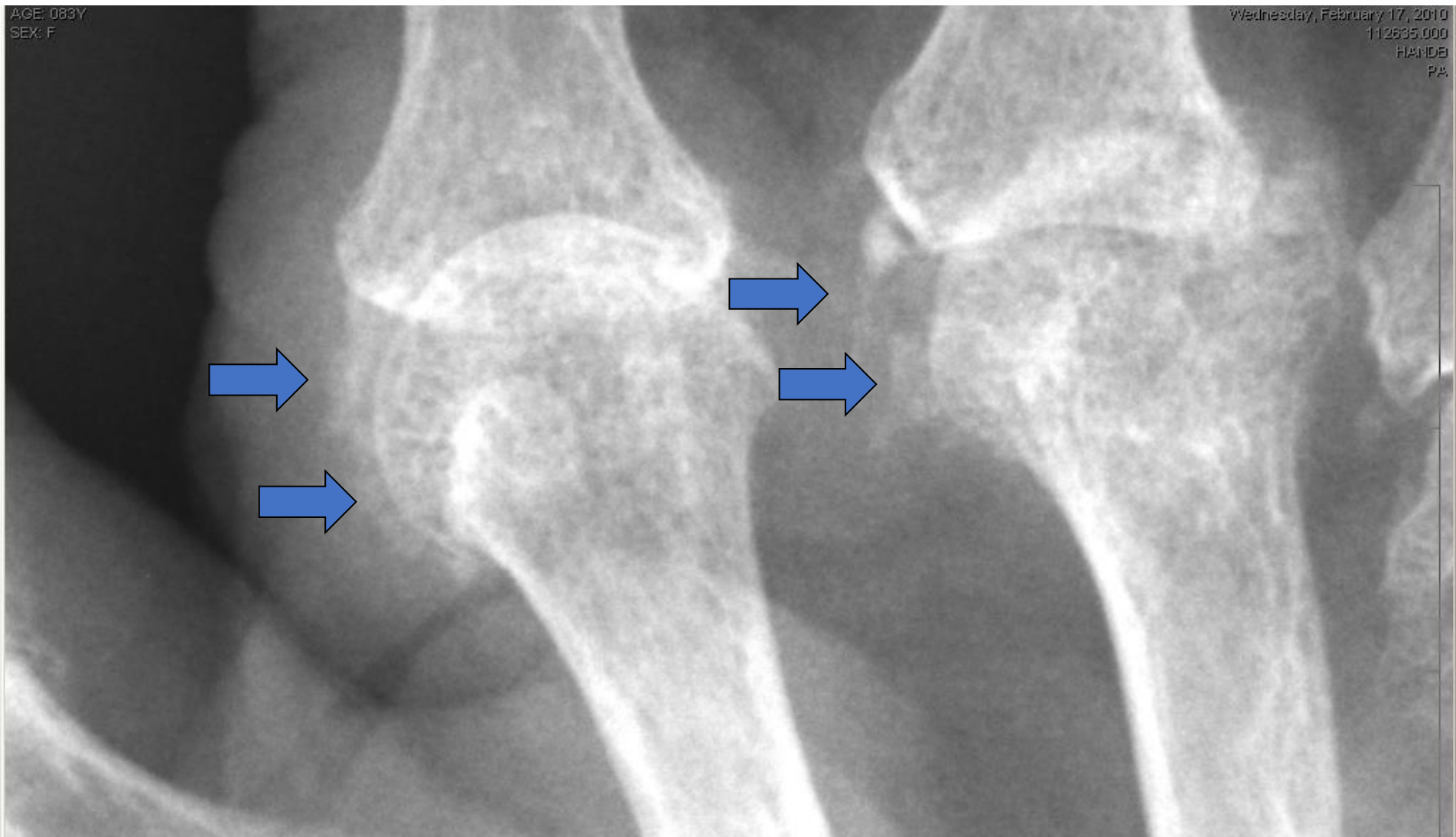
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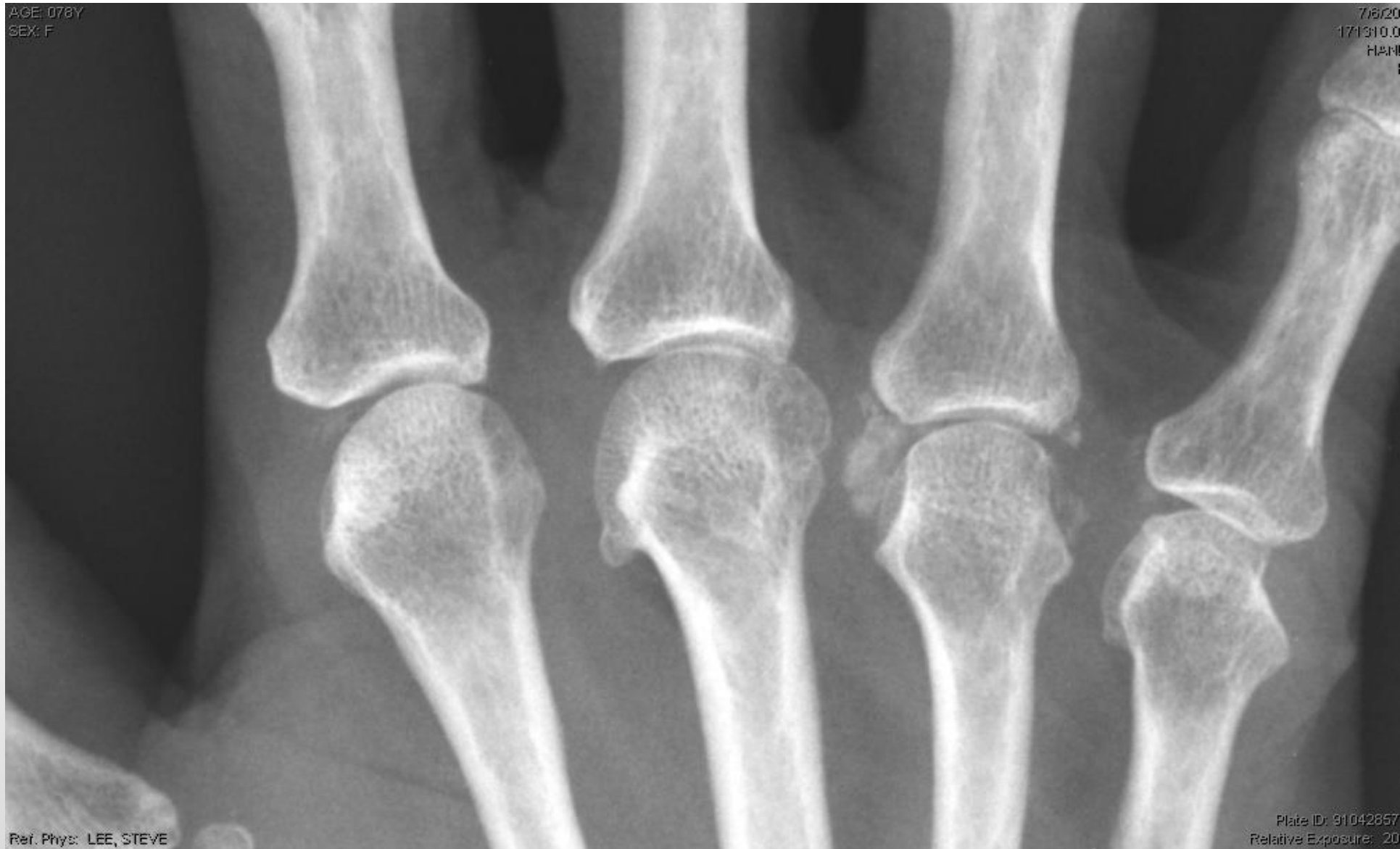
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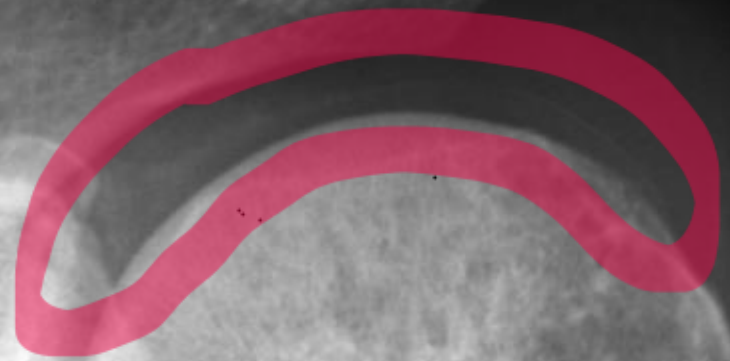
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Metacarpophalangeal osteophytosis and chondrocalcinosis

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Need to know: CPPD

- Common cause of acute inflammatory arthritis; mimics gout, RA, septic and OA
- Variable presentations; prevalence and RF's
 - Aging
 - Endocrinopathies
 - Hemochromatosis
- Recognize chondrocalcinosis on xrays
- Treatments:
 - Acutely:
 - Ice, arthrocentesis and steroid injection
 - NSAIDs or colchicine
 - Long term:
 - Low dose colchicine
 - Methotrexate/hydroxychloroquine



Case 5


- HPI: 32 year man presents with one month of gradually worsening stiffness and aching in fingers that improves with use and OTC ibuprofen. Moderate morning stiffness in low back x 1 hour. Significant limitation in daily function.

- SH: moderate etoh; no drugs/tobacco
- FH: mother with psoriasis
- Labs:
- Exam.....

WBC'S AUTO	4.0 - 11.0 x1000/mcL	5.9
RBC AUTO	4.70 - 6.10 Mill/mcL	4.59 (L)
HGB	14.0 - 18.0 g/dL	14.4
HCT AUTO	42.0 - 52.0 %	42.3
MCV	80.0 - 94.0 fL	92.0
MCH	27.0 - 35.0 pg/cell	31.4
MCHC	32.0 - 37.0 g/dL	34.2
RDW, BLOOD	11.5 - 14.5 %	13.4
PLT'S AUTO	130 - 400 x1000/mcL	197
CRP	<=7.4 mg/L	1.4
ESR	0 - 15 mm/Hr	9
ALB	3.3 - 4.8 g/dL	3.9
ALT	<=63 U/L	41



What is the most likely cause of this patient's symptoms?

- A. Multicentric reticulohistiocytosis
- B. Sarcoid arthritis
- C. Osteoarthritis
- D. Rheumatoid Arthritis
-  E. Psoriatic arthritis

Psoriatic Arthritis Domains



- Peripheral arthritis
- Dactylitis/Enthesitis
- Axial disease
- Skin domain
 - Nail disease
- Arthritis mutilans



Psoriatic Arthritis Treatments

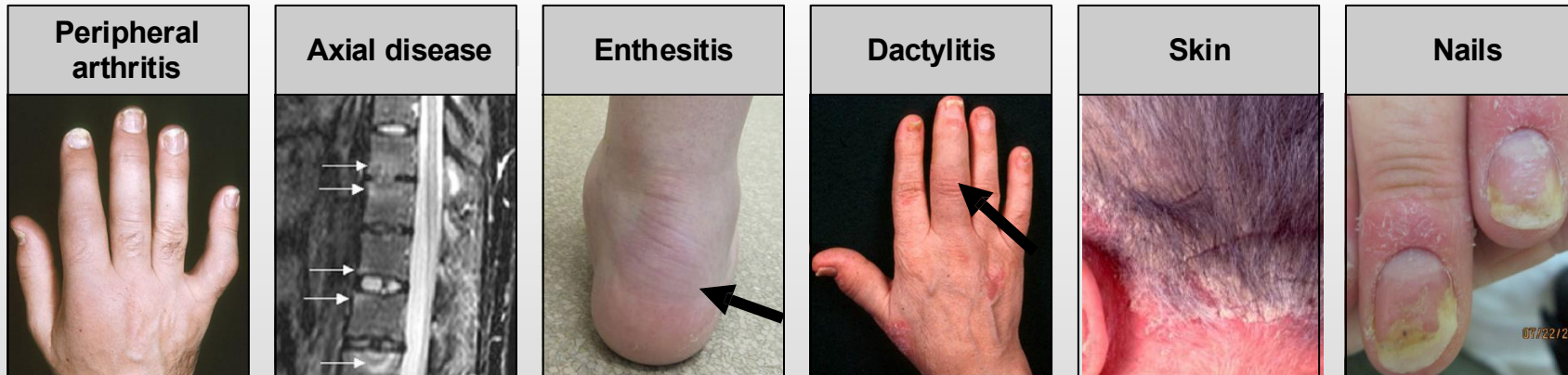
- NSAIDs
- Steroids (topical, IA)
- DMARDs
 - Leflunomide
 - Sulfasalazine
 - Methotrexate
 - Apremilast/Otezla (PDE-4)
- Interleukin inhibitors:
 - Ustekinumab/Stelara (IL-12/23) and **biosimilars**
 - Guselkumab/Tremfya (IL-23)
 - Risankizumab/Skyrizi (IL-23)

 - Secukinumab/Cosentyx (IL-17)
 - Ixekizumab/Taltz (IL-17)
 - Bimekizumab/Bimzelx (IL17A,F)
- T cell costimulation inhibitor:
 - Abatacept/Orencia
- Tumor necrosis factor inhibitors
 - Etanercept/Enbrel
 - Infliximab/Remicade
 - Adalimumab/Humira
 - Certolizumab/Cimzia
 - Golimumab/Simponi
 - **Biosimilar** infliximab and adalimumab
- JAK's, TYK
 - Tofacitinib/Xeljanz (JAK)
 - Upadacitinib/Rinvoq (JAK)
 - Deucravacitinib/Sotyktu (TYK2)



Case 5: Psoriatic arthritis key points

- Another multi-domain condition
- Recognition and refer
- Newer therapies depending on domains
 - Traditional dmards (skin and mild disease)
 - Biologics (TNF's in severe disease first line)



Final Case

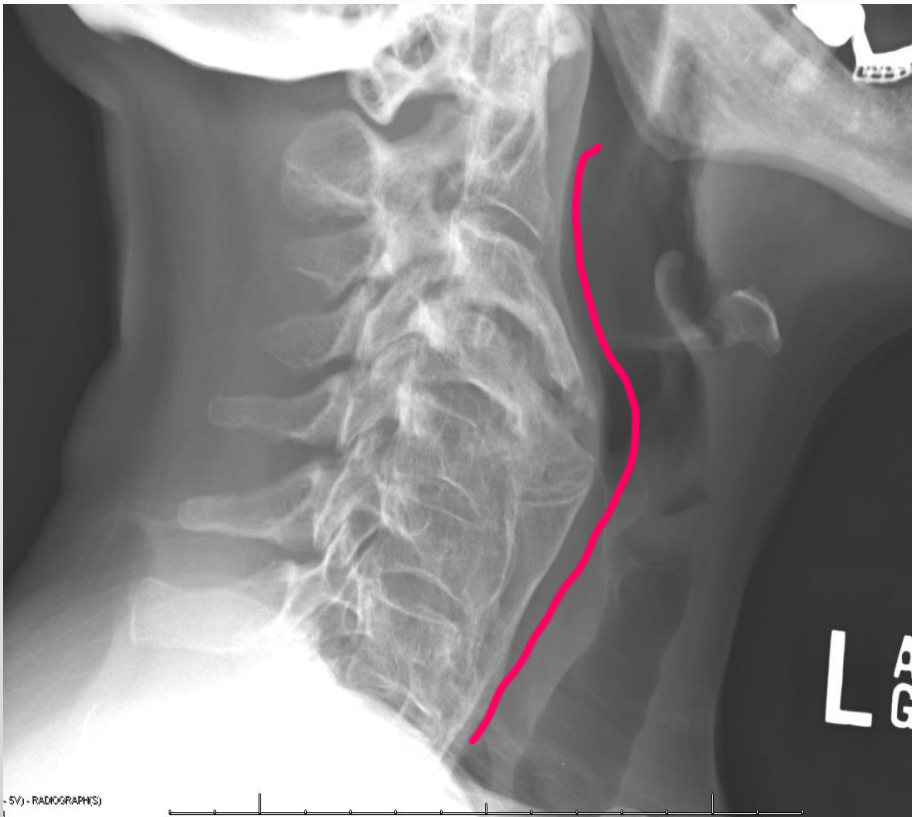
- 64 yo man with longstanding DM presents with 6 months of worsening chronic neck stiffness



Diffuse Idiopathic Skeletal Hyperostosis (DISH)

“Candle wax pattern”:

1. presence of continuous ossification/calcification along anterolateral aspects of 4 contiguous vertebrae
2. preservation of disks without degeneration
3. absence of facet joint fusion or sacroiliac erosions/fusion



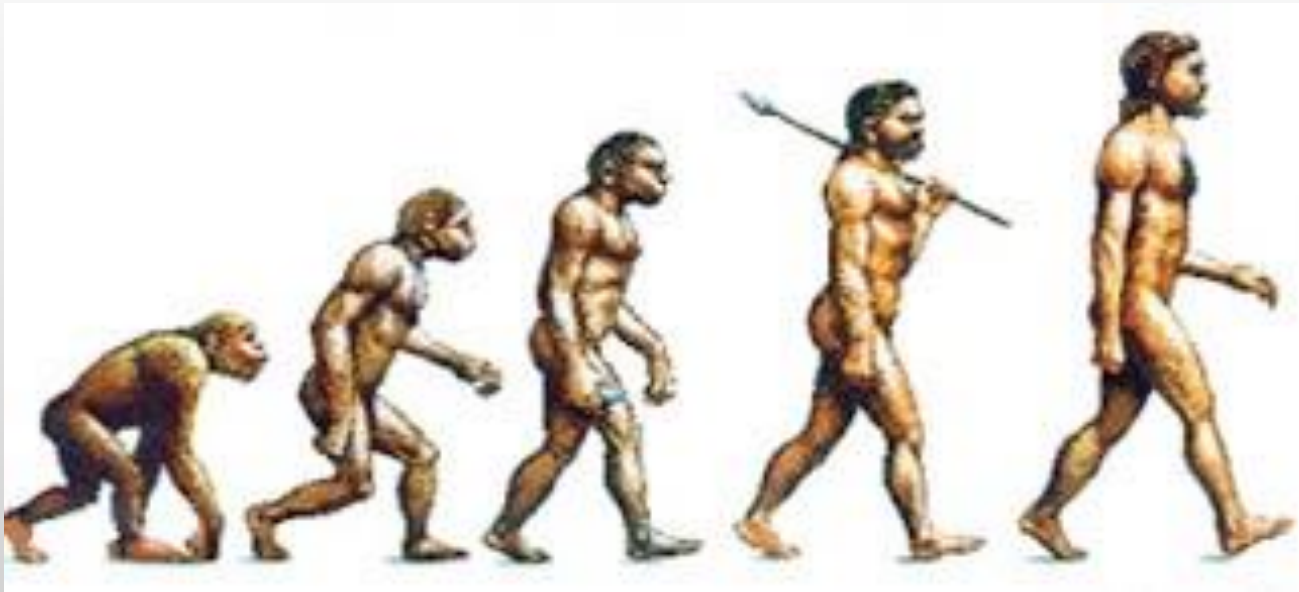
Axial Spondyloarthritis

- Marginal syndesmophytes

Diffuse Idiopathic Skeletal Hyperostosis (DISH)

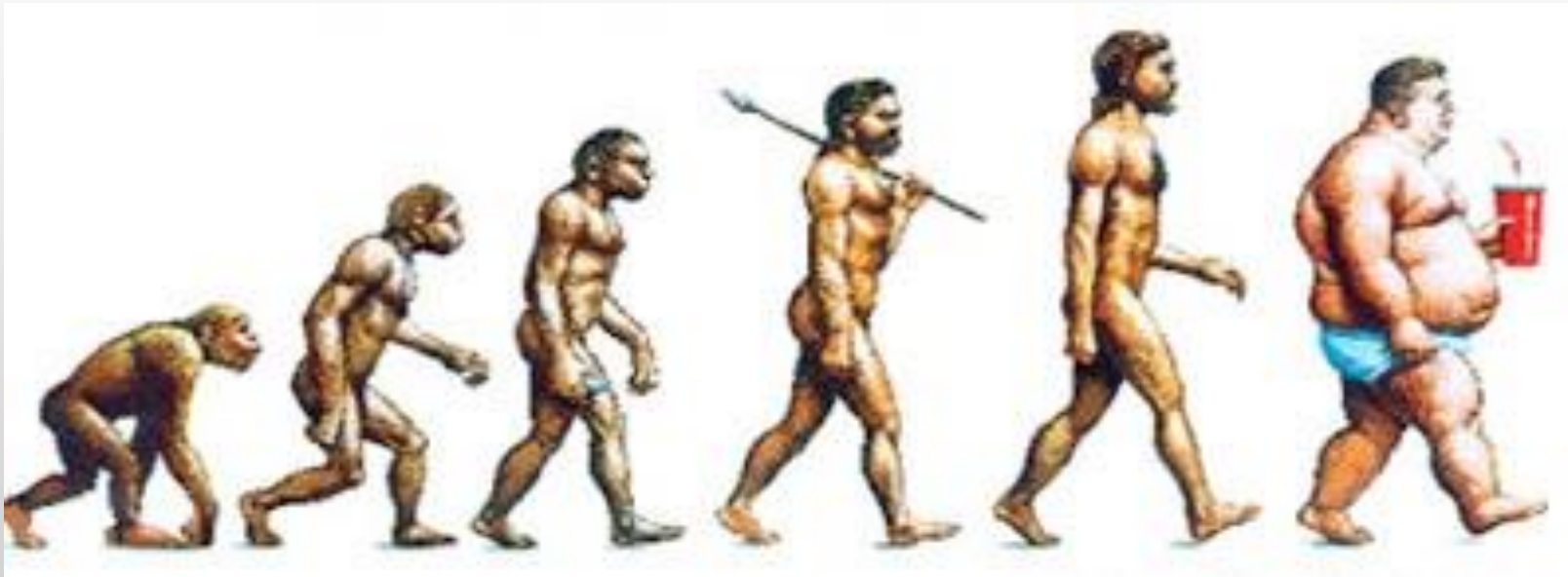
- Commonly mistaken for Spondyloarthritis
- Age of onset >50
- Genetic associations
- Endocrinopathies (DMII, PTH)
- Ossification of ALL, extraspinal ligaments
- No role for immunomodulators
- Analgesia, NSAIDs, PT, surgical options PRN

OA contributing factors



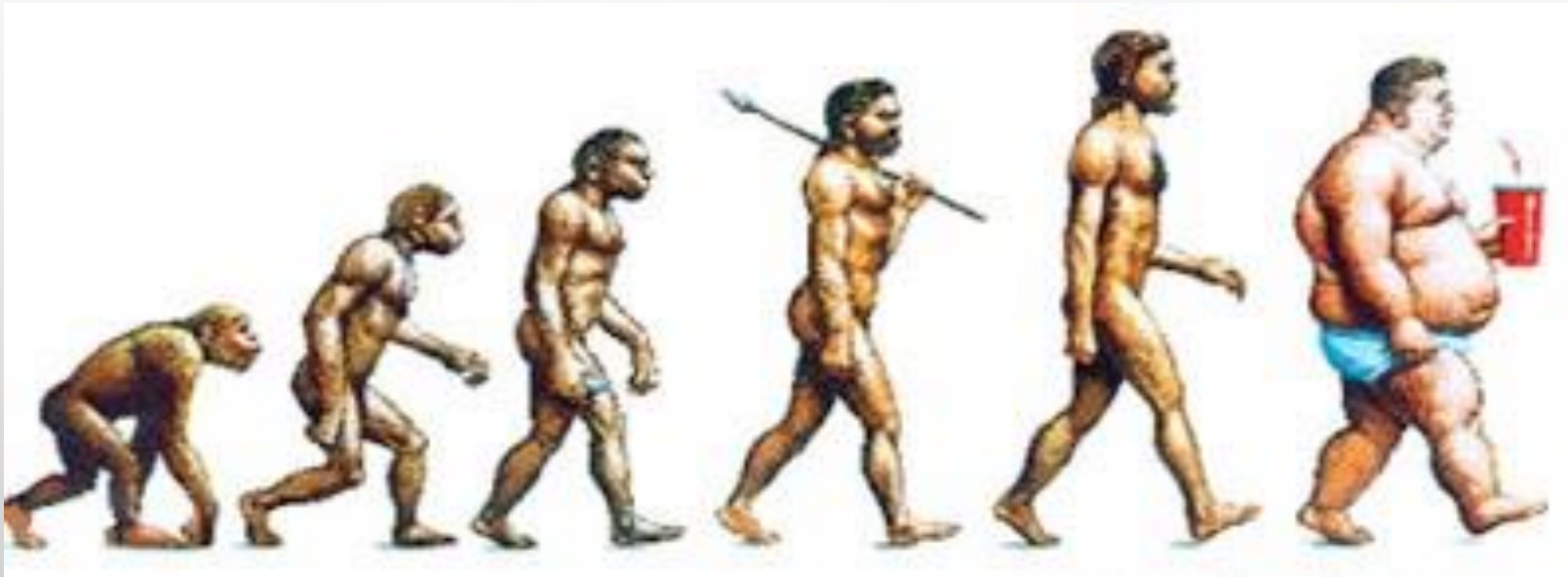
OA contributing factors

- Obesity



OA contributing factors

- Obesity
- Prior trauma and injury
- Menopause: estrogen?
- Family history



OA

- Pathophysiology
 - Periosteal bone remodeling (bone morphogenetic proteins) - subchondral microfractures
 - Cartilage degradation (matrix metalloproteinases)
- Exam: bony enlargement, decreased ROM, peri-articular muscle weakness, crepitus
- Joints distribution:
 - DIPs, PIPs, CMCs
 - Cervical, lumbar spine
 - 1st MTPs, knees, hips
 - **NOT shoulders, wrists or MCP's**



OA treatment

- Non-pharmacologic:
 - Wt loss, PT, muscle strengthening, aerobic exercise
 - Education
- Pharmacologic:
 - NSAIDs
 - Analgesia (tylenol, tramadol, opiates)
 - Anti-depressants (nortriptyline, duloxetine)
 - Anti-epileptics (fibromyalgia)
 - GLP-1?
- Injections of steroids or hyaluronic acid or GLP-1?

Summary: evaluation of polyarthrititis

- Spondyloarthritis – numerous domains, including LBP
- Elderly onset gout – traditional and even DIP involvement
- CPPD – can appear similar to knee OA, even RA
- Psoriatic arthritis – DIP involvement among other domains
- DISH – a form of axial OA