















BEAVER CREEK

BRECKENRIDGE

KEYSTONE

HEAVENLY

NORTHSTAR

KIRKWOOD

Acute Emergencies in Rheumatology

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Acute Rheumatological Emergencies

- The Acute Hot joint
- Inflammatory back pain...
- Systemic lupus erythematosus(SLE)
- Inflammatory muscle problems
- Adults onset Stills Disease

The Acute Hot joint!





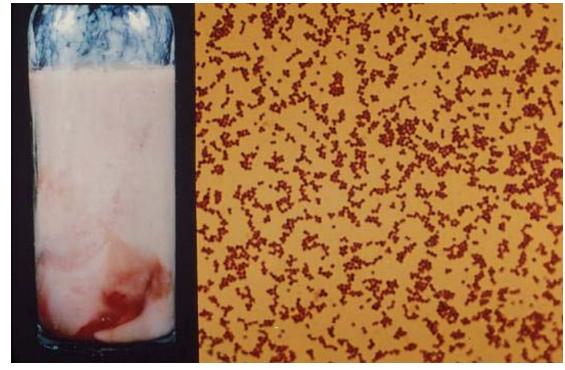




Should we aspirate this?







What is the differential Diagnosis?

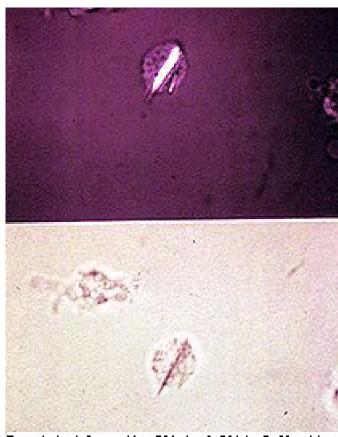
- Acute crystal arthritis
- Urate crystals- gout
 - Calcium pyrophosphate
 - - Pseudo gout
 - Hydroxyapatite-
 - Milwaukee shoulder
- Septic arthritis
- Osteomyelitis
- New presentation of inflammatory arthritis



How should it be diagnosed?

Do we need this?

 Do you know where to send it?



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Calcified hyaline and fibrocartilage with linear and spotty appearances and well-preserved joint space

Challenges with the Acute Hot Joint

- Missing the acute septic arthritis
- Missing the osteomyelitis.
- Crystals not obtained from the joint.

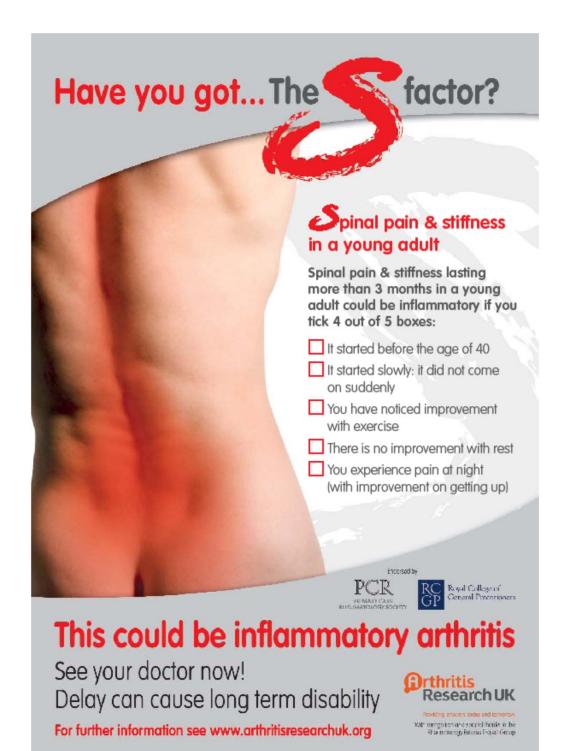
Challenges with Acute Hot Joints

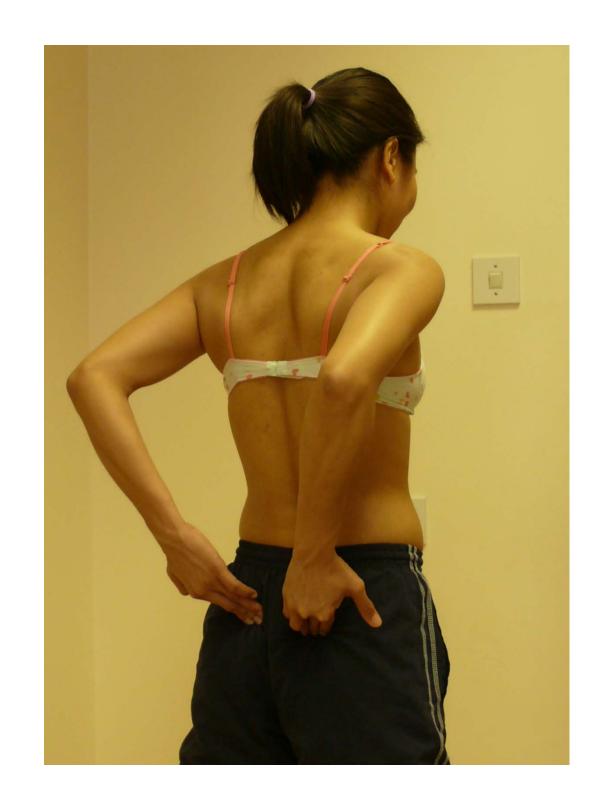
- Diagnosis
- ??



Back Pain?



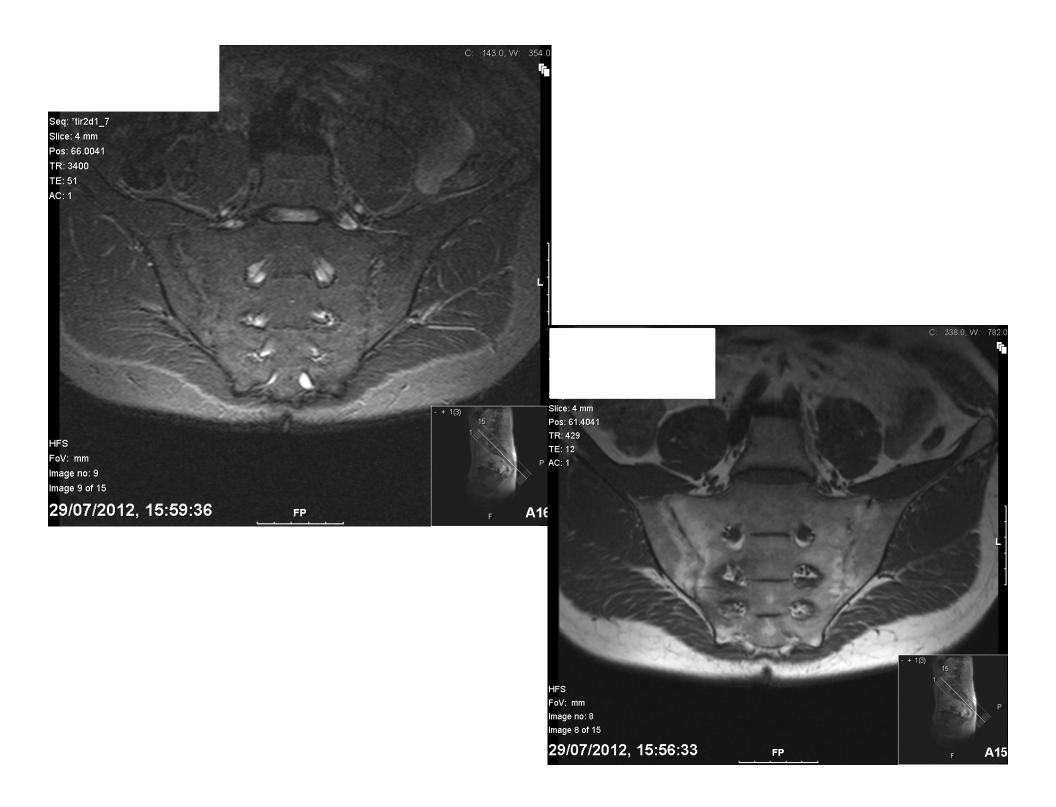




inflammatory back pain

ASAS Criteria (Sieper J et al Ann Rheum Dis 2009;68:784-8): Back pain of more than 3 months duration is inflammatory if:

- Age at onset less than 40 years
- Insidious onset
- Improvement with exercise
- No improvement with rest
- Pain at night (with improvement on getting up)
 The criteria are fulfilled if at least 4 of 5 parameters are present

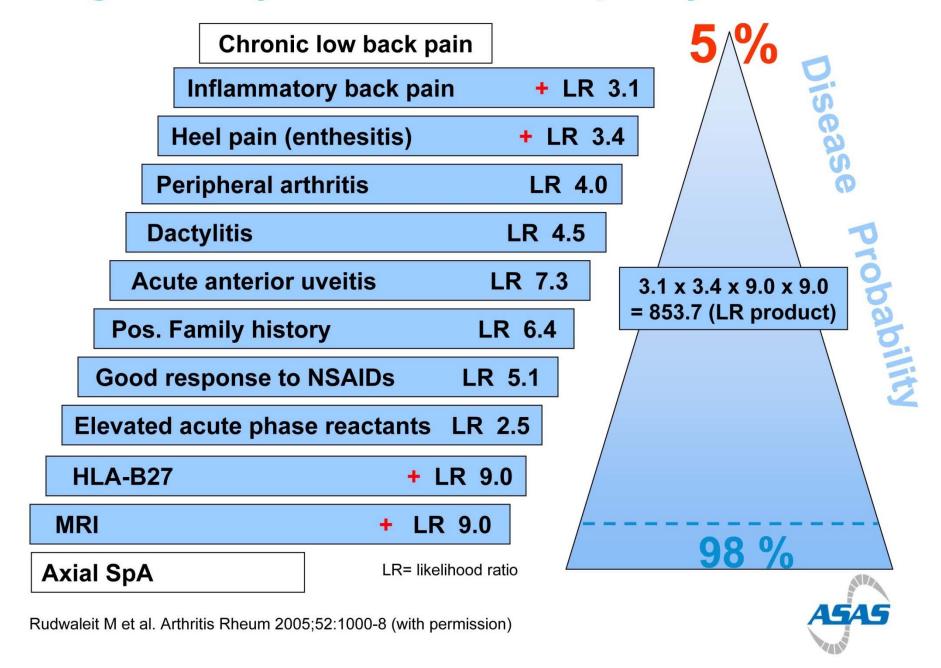


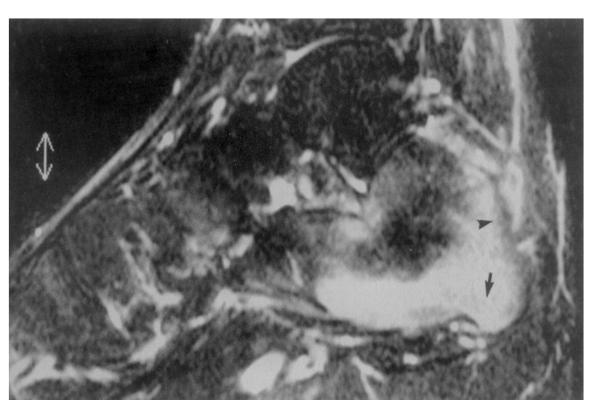




and 50% of people with AS have other associated problems.....

Diagnostic Pyramide for Axial Spondyloarthritis

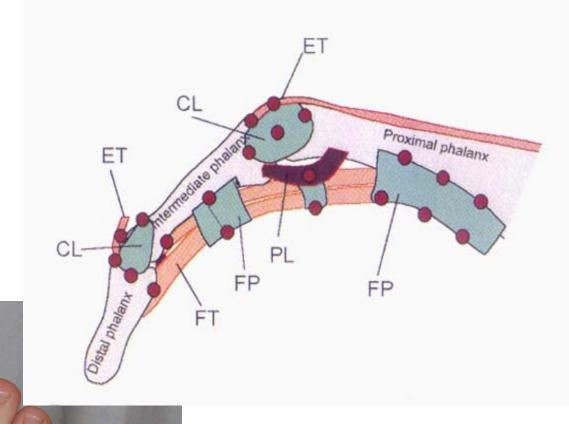










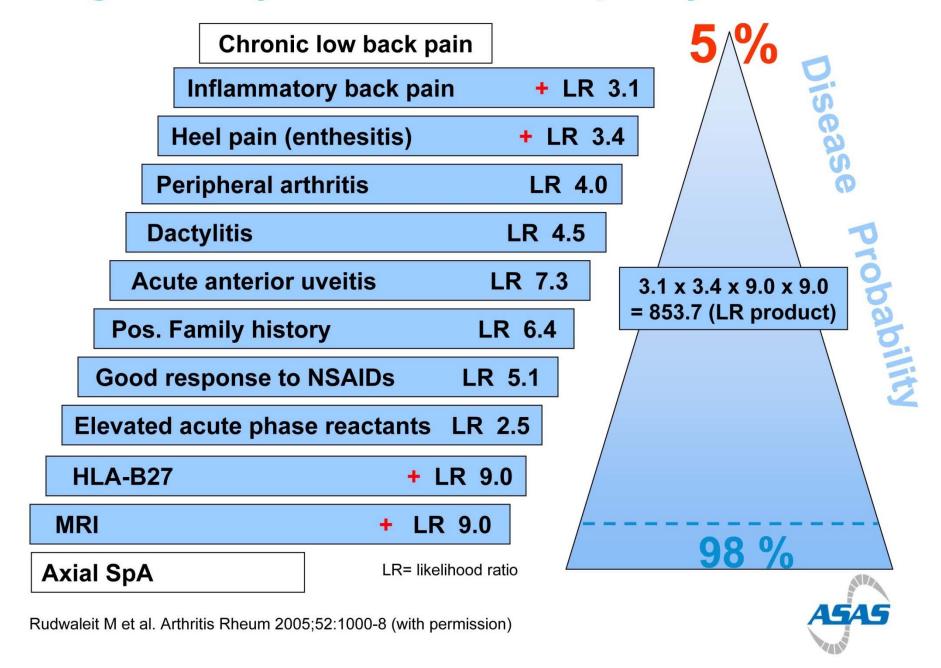








Diagnostic Pyramide for Axial Spondyloarthritis

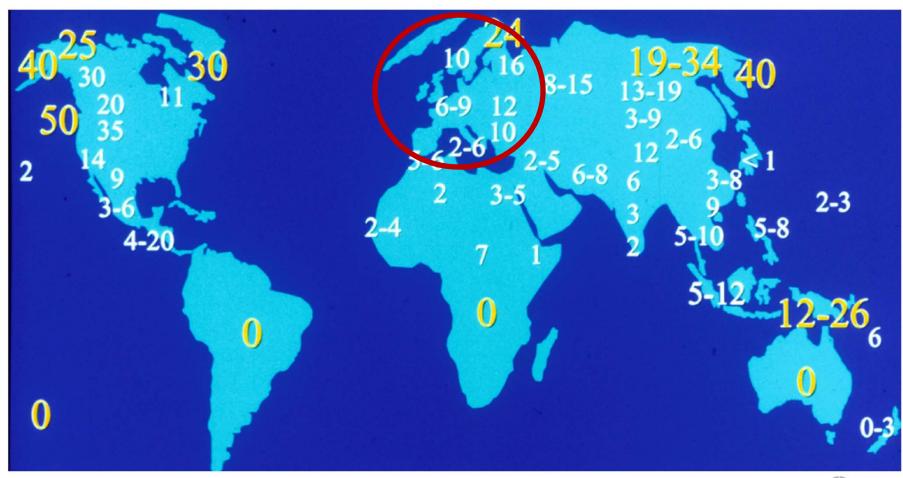


Spondyloarthritides

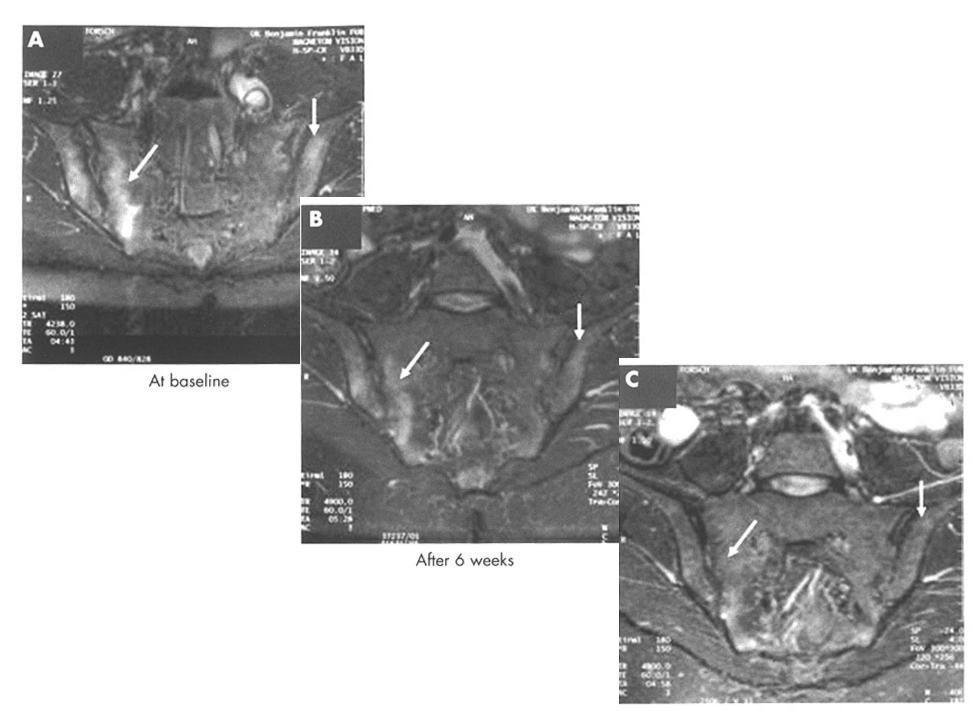
- This group have a similar prevelence to RA
- They share common clinical lesions.
- Inheritance of HLA-B27 is common to all the SpA. The prevalence of these disorders relate to HLA-B27.
- Diagnosis of Ankylosing spondylitis is often delayed
- Identification of inflammatory back pain is very important in determining the diagnosis.
- Use of Anti- tumour necrosis factor biologic drugs has revolutionised the treatment of severe AS.



Percentage Prevalence of HLA-B27 in Indigenous Populations of the World

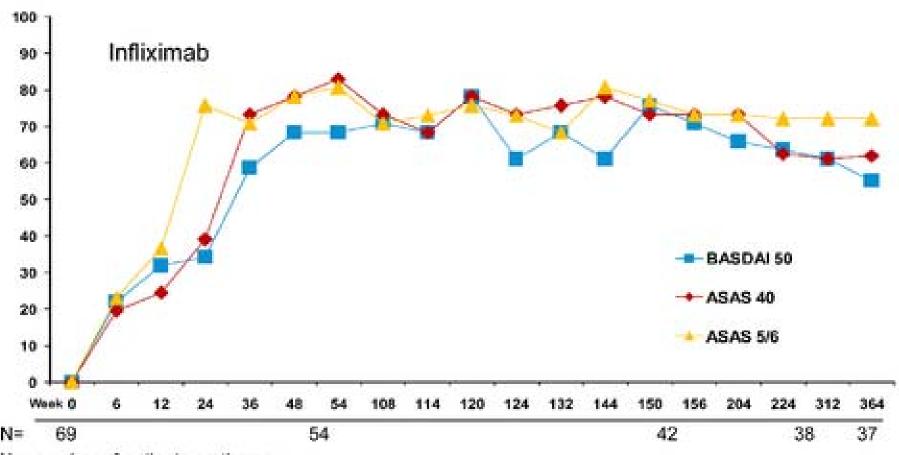






After 24 weeks

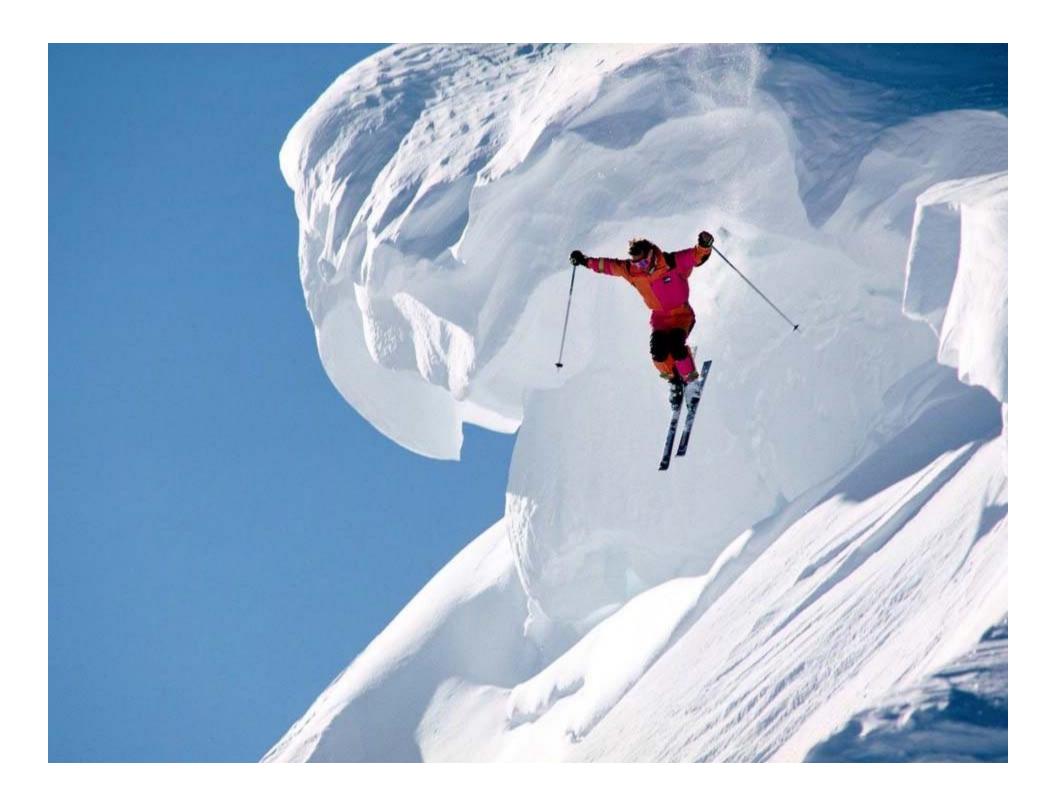
Longterm Clinical Efficacy of TNFα-Blocker in AS Results over 7 years



N = number of patients on therapy

Braun J et al. Lancet 2002;359:1187-93 Braun J et al. Ann Rheum Dis 2008;67:340-5 Baraliakos X et al. EULAR 2008, Paris, FRI0290





Connective tissue diseases. Acute presentations



Case Study AT

- 24yr old vegetarian Asian female
- presented via emergency dept.
- Vomiting, 5kg wt loss, night sweats
- Productive cough & pleurisy
- Initial Diagnosis Pulmonary Tuberculosis
- Investigation. Hb 7.8g/dl(low).blood white cells 3.5(low)
- ESR 105mm/hr C reactive protein <4(n)

.

Reviewed 24hrs later

- Facial rash, alopecia, febrile, chest X ray clear
- Diagnosis Probable SLE
- ? Why??

AT (2)

- Further investigations.ANA 1/2560 homogenous(very high)
- DNA binding .>300;(HIGH)
- IgG 26.7;IgA 6.9 IgM 2.4;g/I(raised)
- C3 0.3 C4 0.05;g/IComplements(-low)
- Coombs (+) creatinine 105umol/L 24hr urinary protein 2.35 grams/24hrs

- Renal Biopsy WHO Class IV (Diffuse Proliferative Glomurulonephritis)
- Treatment Intravenous Methyl Prednisolone 1gram X3
- Dramatic recovery!

Recognizing SLE

"Typical bloods"

- Anaemia
- Lymphopaenia
- Normal/low platelets
- High ESR
- Low C3 and C4
- Autoantibodies



Is it a connective tissue disease?

- Raynauds syndrome of recent onset.
- Non specific inflammation with no infection
- Screen for autoantibodies positive.(ANA/DNA)
- Raised ESR with normal C reactive protein.
- Low white cell count with low lymphocyte count and low platelets.
- Rashes.
- Multisystem disease.

I can't get out of my chair!

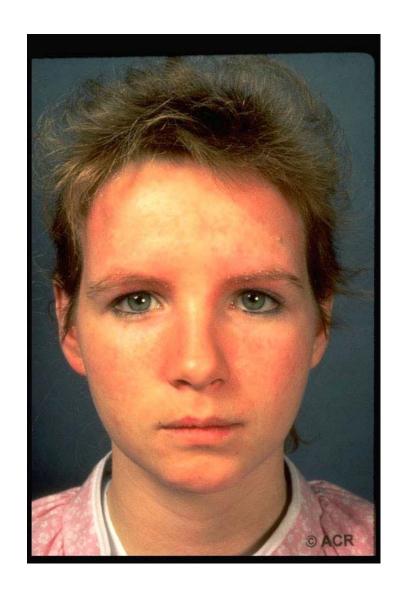


Mrs AU age 35 presented to the dept unable to get out of a chair with severe pain and weakness.

- Facial rash
- Rash on the hands and fingers
- Shortness of breath
- Weakness and muscle pain for 4 weeks
- ?Raynauds syndrome
- Generally unwell.
- Raised ESR, Raised muscle enzymes.
- Raised anti nuclear antibody 1/2500

Dermatomyositis - facial rashes

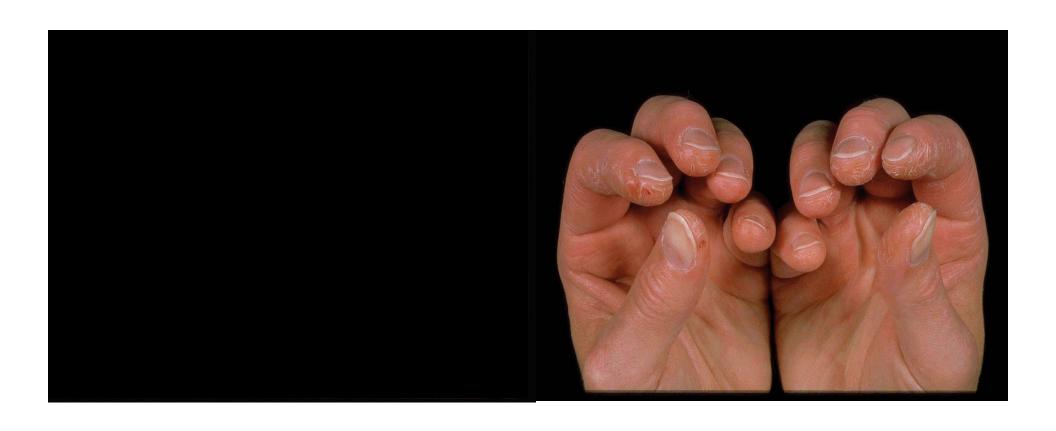




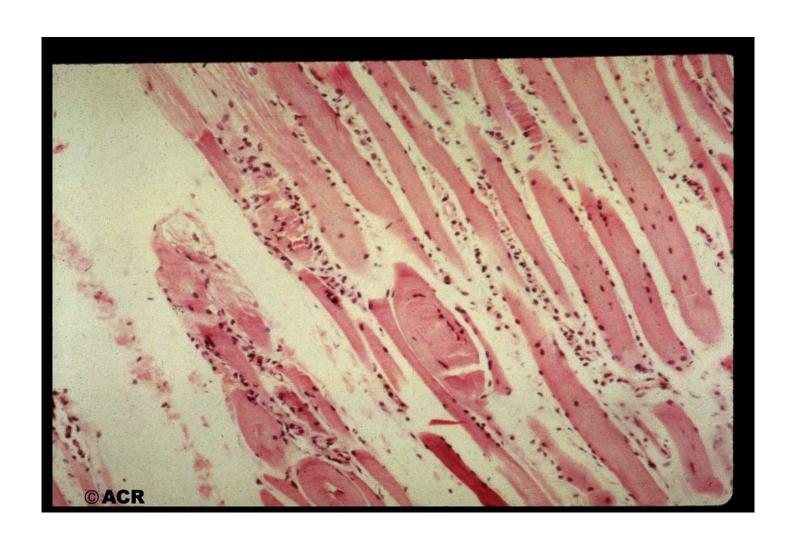
Dermatomyositis - Gottron's papules

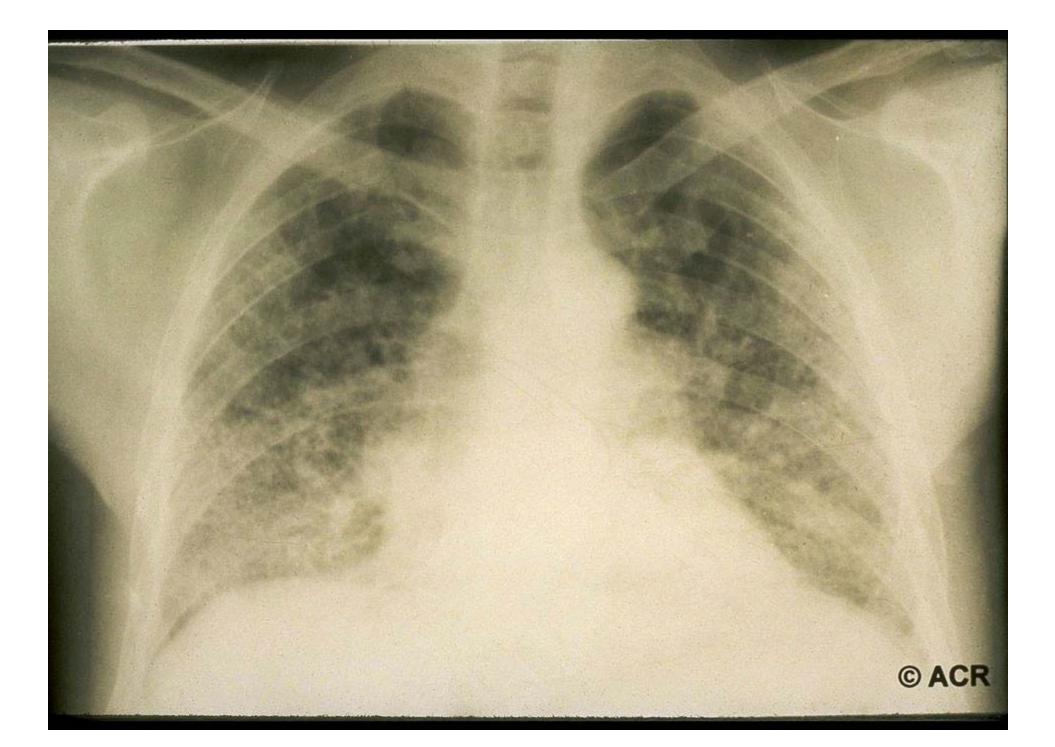


Anti JO 1- mechanic's hands



Acute Myositis





Dermatomyositis

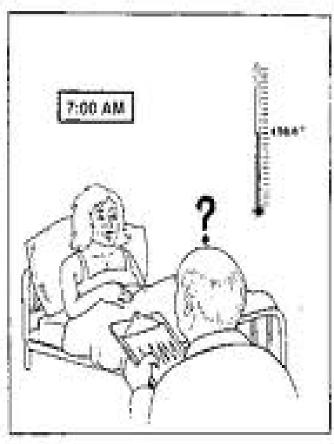
- Rash
- Abnormal muscle enzymes
- Abnormal muscle biopsy and electrical studies and MRI of the thighs.
- Autoantibodies- positive ANA and Extractable nuclear antibodies:--
- Treatment steroids and cyclophosphamide
- ,intravenous Immunoglobulin.
- Rituximab???

Myositis and anti synthetase syndrome

- Mechanic hands
- Myositis
- Anti Jo1 antibodies
- Other antibodies positive –anti trna synthetase
- Manual identification.-Ku etc

Periodic fevers!





Patient with Fever and acute arthritis miss LR

- Pyrexia often Periodic, at night.
- Rose coloured rash on the body and face
- Acutely swollen joints
- General Malaise and weight loss over 2 weeks.

- Abnormal liver function tests.
- Raised ESR

Adult onset Still's disease.



- Fever, rash +/-arthritis,
 pharyngitis, serositis
- Raised inflammatory markers
- Negative RF/ANA
- High ferritin>20,000
- (normal less than 150)

Still's Disease Rash

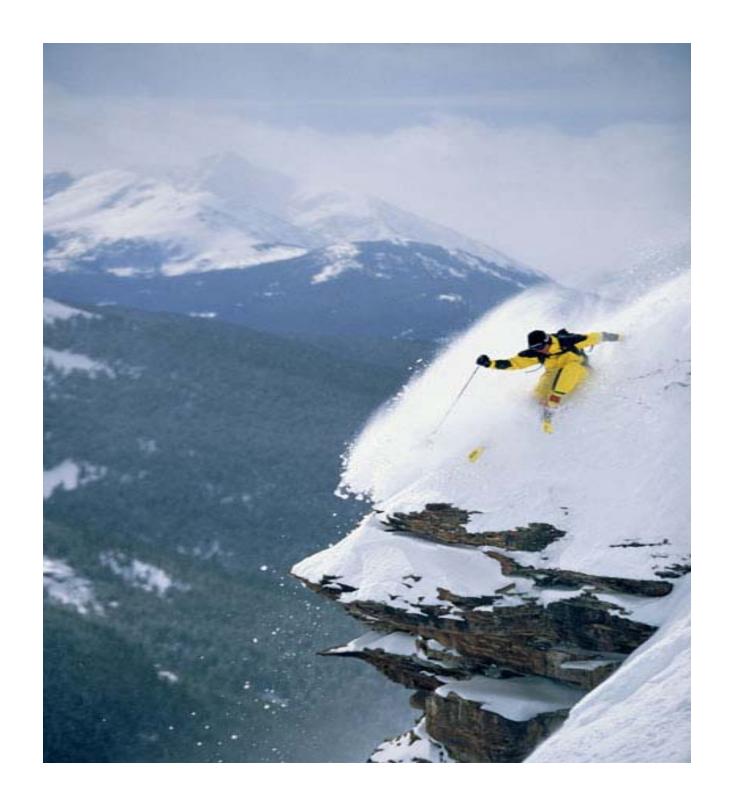


Treatment

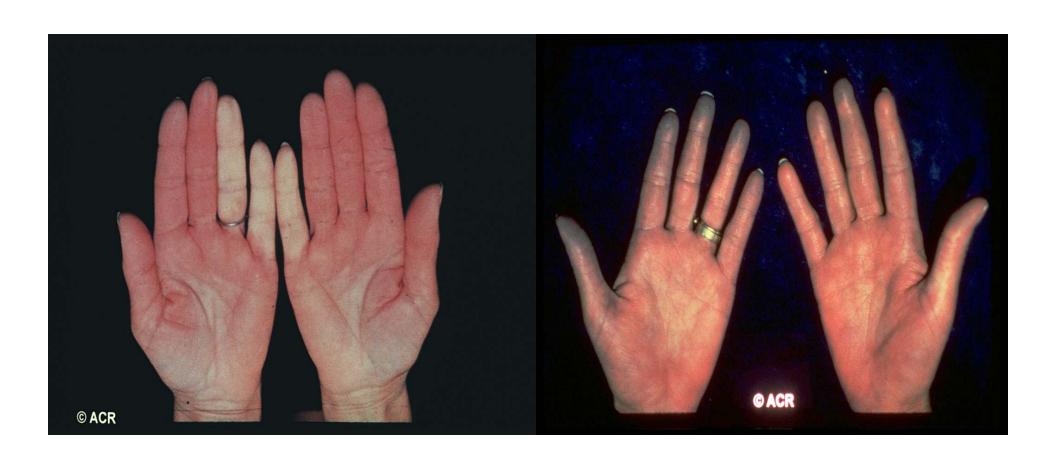
- Anti inflammatories
- Steroids
- Biological therapy- Tocilizumab-Anti IL-6

Northwick park hospital Harrow





Raynauds



Small vessel ischaemia

