

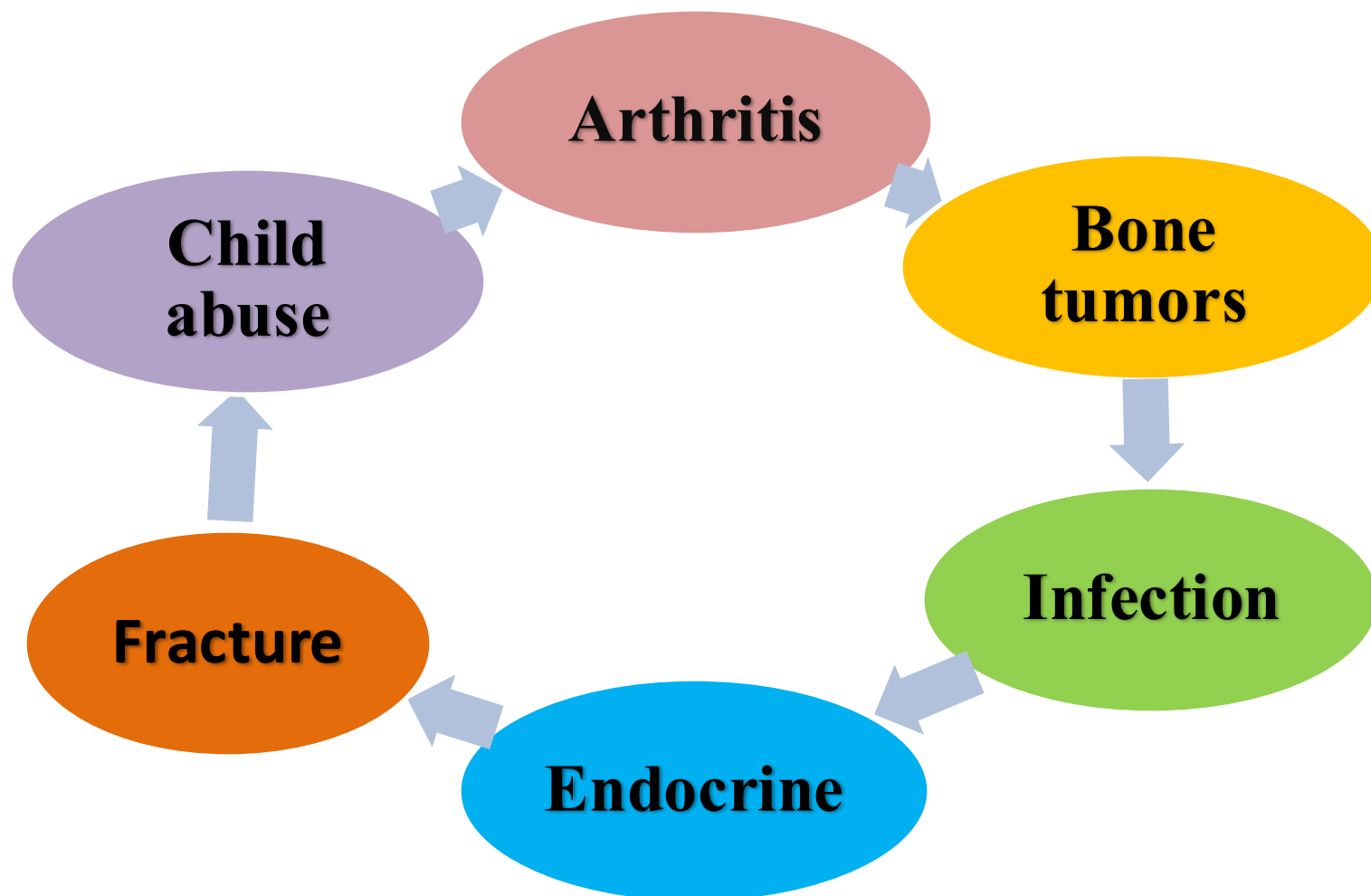


How To Manage G.P. Clinic 12/

MSK Radiology

O. M. Albtoush

Outline



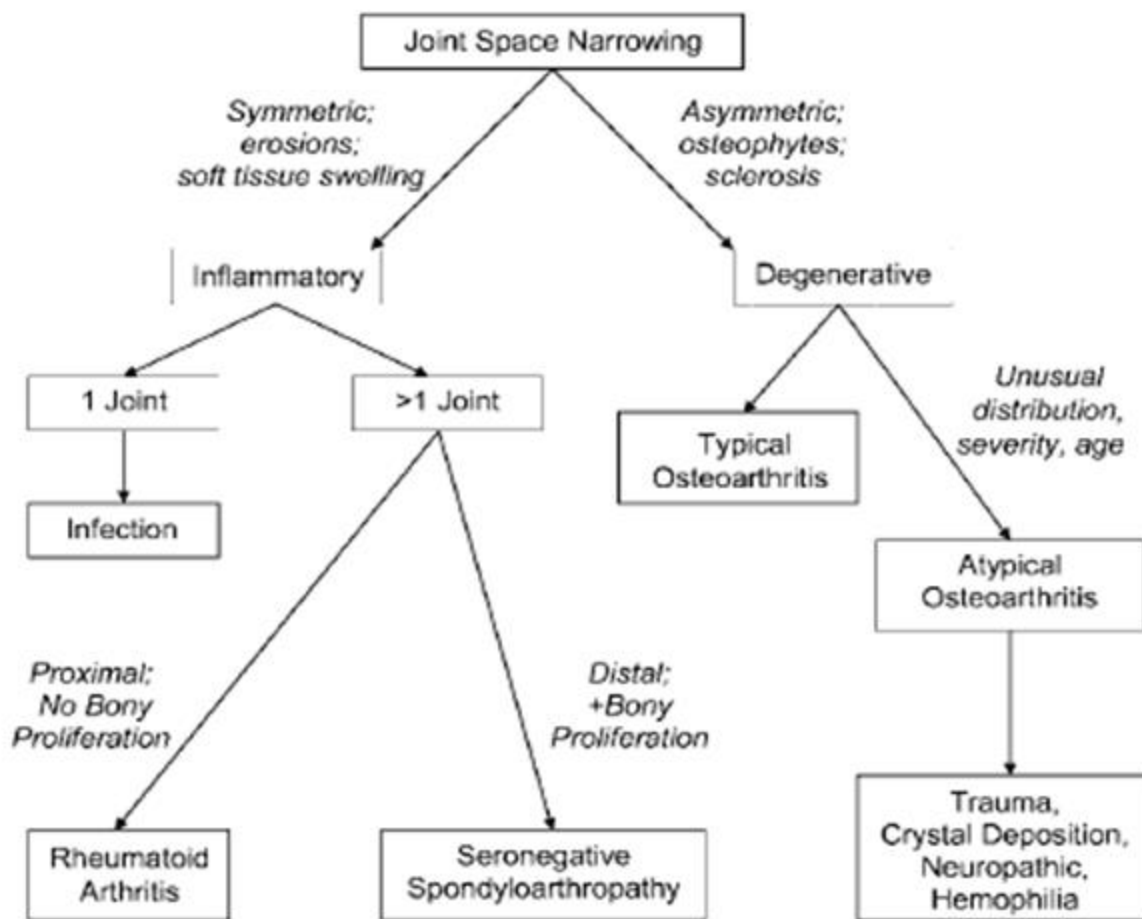
Radiological approach to arthritis

Arthritis

General classification:

1- Inflammatory

2- Degenerative



Inflammatory arthritis

Single versus Multiple

Single joint involvement concern  infection.

Multiple joints inflammation in **proximal** distribution in the hands or feet **without** bone proliferation suggests **Rheumatoid arthritis**.

Multiple joint inflammation in **distal** distribution in the hands or feet **with** bone proliferation suggests **seronegative spondyloarthropathy**.

Signs of inflammatory arthritis

1- Uniform joint space narrowing

2- Soft-tissue swelling

3- Periarticular osteopenia

4- Bone erosion

Rheumatoid Arthritis

Multiple joints involvement.

Proximal distribution of the hand or feet

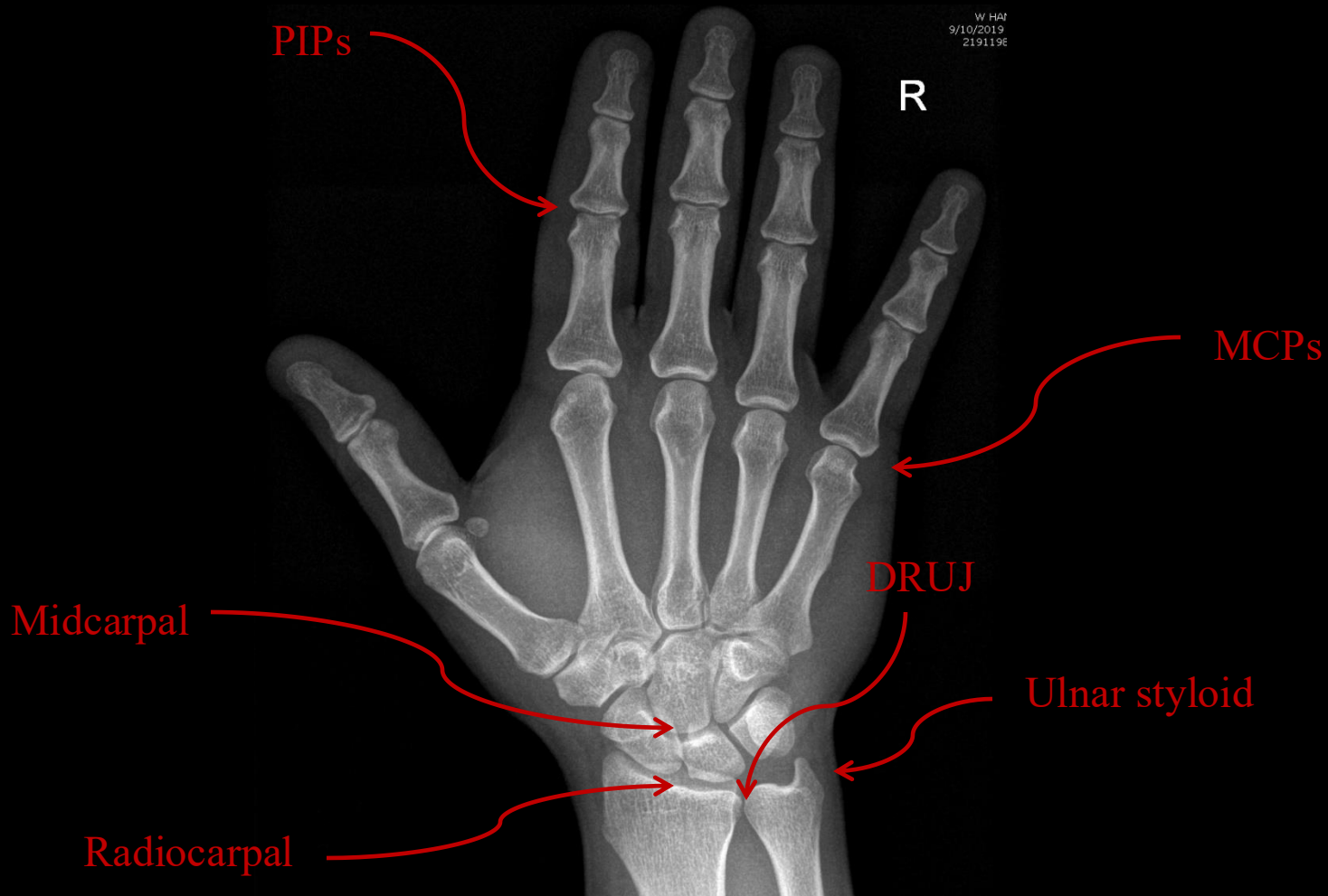
Lack of bone proliferation.

Most common in women aged 30–60 years.

Proximal distribution.

Bilateral involvemnt.

Distribution of Rheumatoid Arthritis



Rheumatoid Arthritis

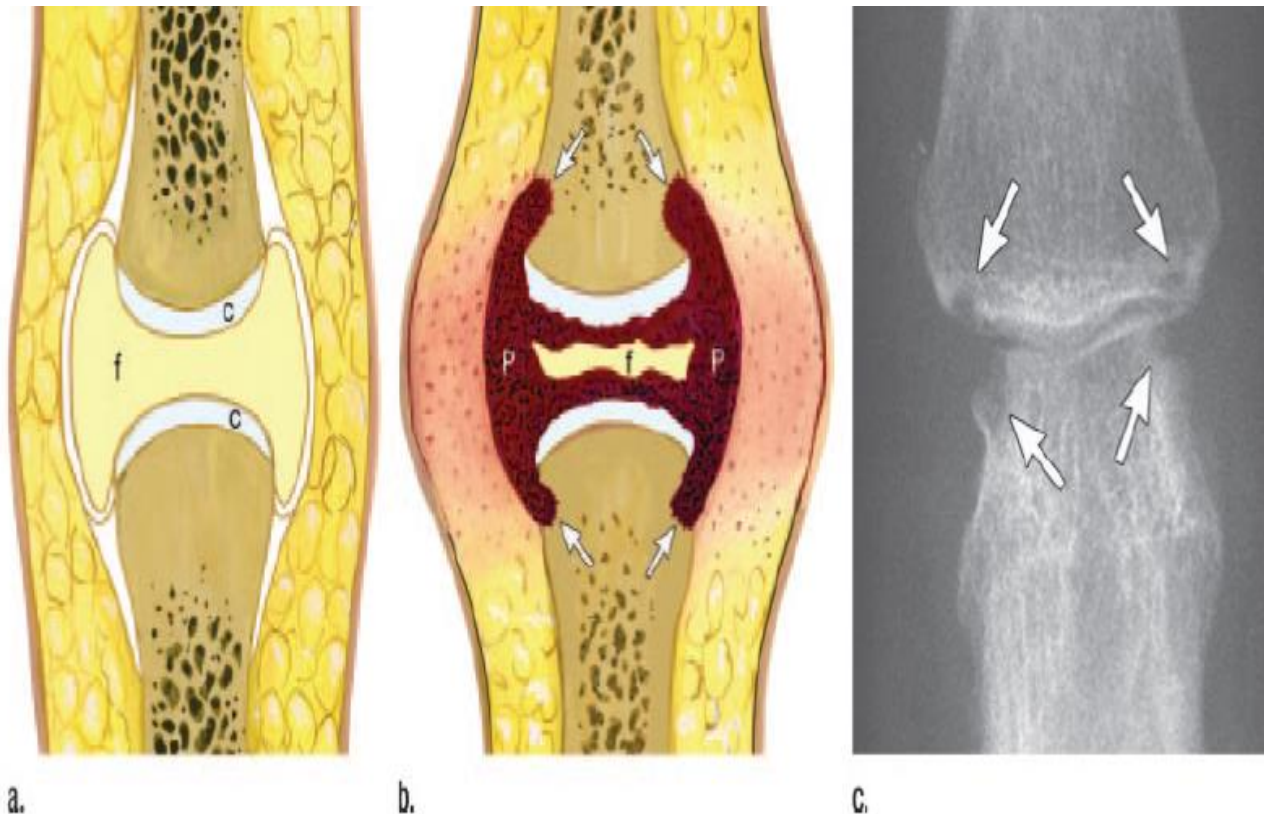
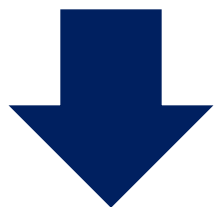


Figure 3: (a) Illustration of synovial joint shows joint fluid (*f*) and articular cartilage (*c*). (b) Illustration and (c) radiograph show inflammatory arthritis, synovitis, and pannus (*P*) causing cartilage destruction. Marginal erosions (arrows) are seen where subchondral bone plate is exposed to intraarticular synovitis. *f* = Fluid.

Normal Radiograph

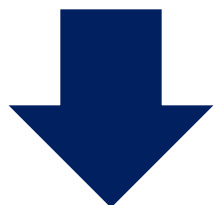


Joint narrowing
Erosions
Osteopenia



**MCP, DRUJ,
Radiocarpal,
Midcarpal**

Subluxation



PIPs.



Joint space narrowing:

PIP

MTC

Midcarpal

Radiocarpal

DRUJ

**Erosions of the ulnar styloid
process (arrowhead).**



**Extension
and flexion
radiographs
show:**

**widening of
Atlantoaxial
interval
(arrowheads)**



Seronegative spondyloarthropathies

**Multiple distal joint involvement in the hands and feet
with bone proliferation**

**Include: psoriatic arthritis, reactive arthritis and
ankylosing spondylitis.**

Ankylosing spondylitis

Idiopathic inflammatory arthritis.

96% of patients are HLA-B27 positive.

Men affected three times more frequent than women

Between 20 - 40 years

Involves the axial skeleton

Ankylosing spondylitis

Spine involvement is characterized by osteitis, syndesmophyte formation, facet inflammation, and eventual facet joint and vertebral body fusion

Early radiographic findings are erosions at the anterior margins of the vertebral body at the discovertebral junction. These focal areas of osteitis become increasingly sclerotic, a finding termed the “**shiny corner sign**”

Thin and slender syndesmophytes are generally evident, representing ossification of the outer layer of the anulus fibrosus

As the syndesmophytes thicken and become continuous, “**bamboo spine**”

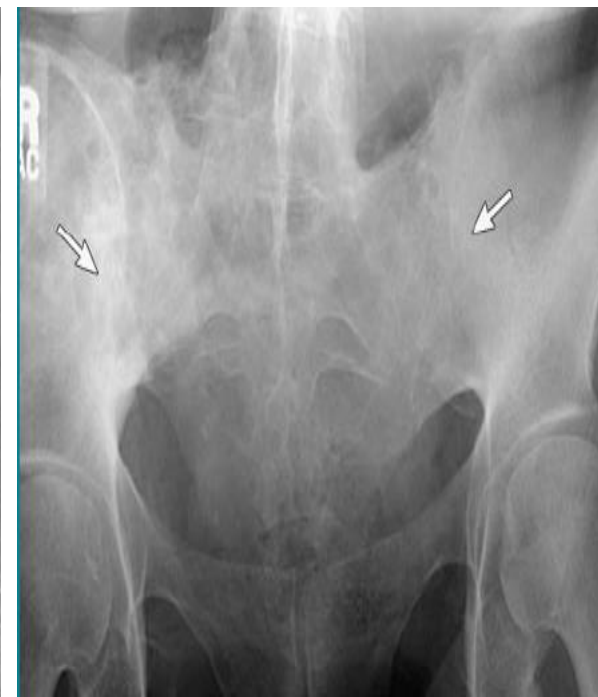
Ankylosing spondylitis

Ossification of the posterior interspinous ligament produces a dense radiopaque line, designated the “**dagger sign**”.

The combination of the fused facets and ossification of the interspinous ligaments produces the “**trolleytrack sign**”.

Bilateral and symmetrical sacroiliac joint disease and usually precedes spinal involvement.

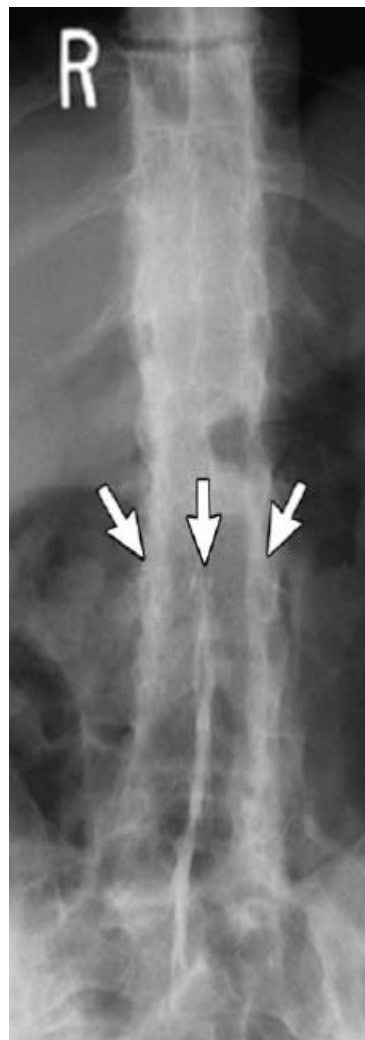
Sacroiliitis (Normal, Early and Late)



**AP lumbar spine
radiograph
shows:**

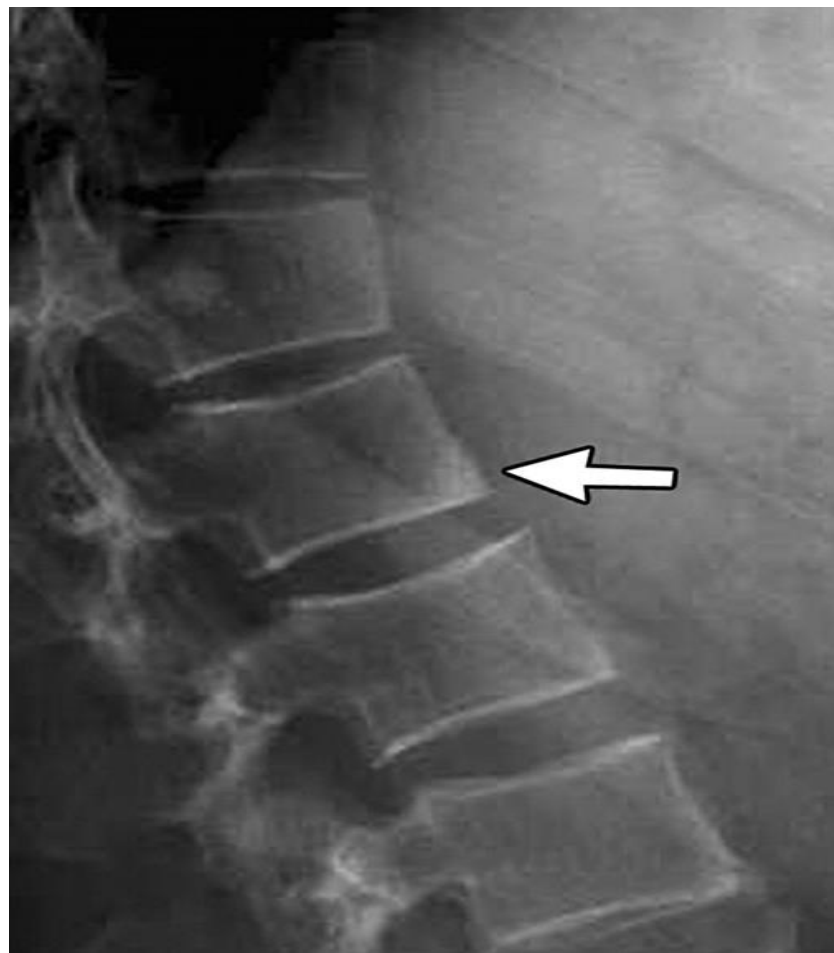
**Bridging
syndesmophytes
(bamboo spine)**





**Trolleytrack
sign**

Dagger sign



Shiny corner sign

Septic arthritis

Usually staph. or strep through hematogenous spread.

Radiographic features:

Uniform joint space narrowing (initially it maybe widened due the presence of effusion).

Periarticular osteopenia.

Soft-tissue swelling.

Bone erosions (not seen acutely!!)

Clinical data and physical examination can aid in the diagnosis

Septic arthritis: Joint
Space narrowing
(arrows), osteopenia,
soft-tissue swelling,
and bone erosion
(arrowhead)



Degenerative Joint Disease

Osteoarthritis

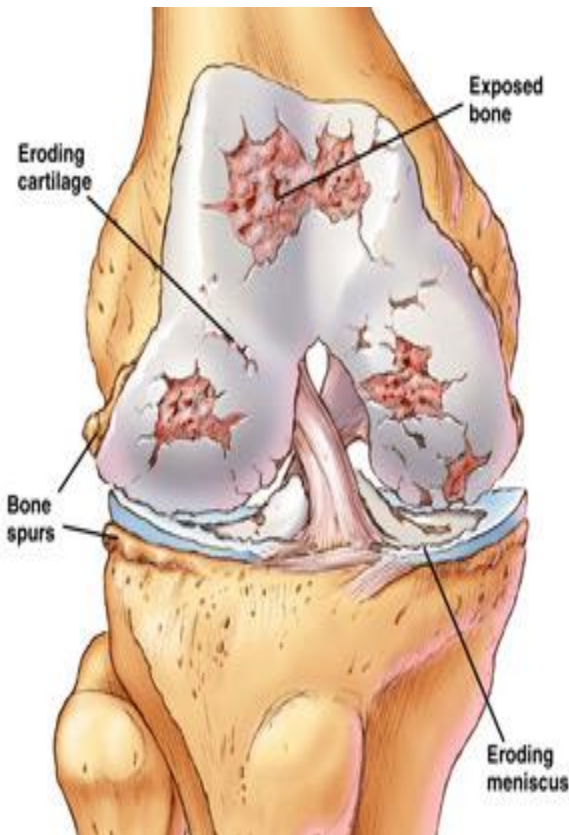
Characterized by Osteophytes, bone sclerosis, and subchondral cysts (geodes).

Absence of inflammatory features such as erosions, uniform Joint space narrowing.

Involves specific joints at a particular age (distribution).

When atypical in location, early age, or unusual radiographic appearance; other causes for cartilage destruction should be considered, such as trauma, crystal deposition, neuropathic joint, and hemophilia.

Knee osteoarthritis





Inflammatory arthritis

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Osteoarthritis

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Unusual location, secondary cause: FAI



Unusual location, secondary cause: Traumatic

Diagnosis clue:

Young age

Atypical joint

Unilaterality



Neuropathic joint

Characteristic findings: **Sclerosis, fragmentation, and subluxation** are obvious in the later stages of this process, the early changes of a neuropathic joint will often appear non-specific.

The primary clue is the distribution of radiographic changes

Midfoot is characteristic with diabetes mellitus, where findings of joint space narrowing, bone sclerosis, and osteophytes are seen.

A coexisting **arterial calcifications** further suggest the diagnosis of neuropathic joint, and correlation with patient history

Joint space narrowing, sclerosis, subchondral cyst, and osteophyte formation. Note; plantar navicular tilt and pes planus



Gout

Erosions are frequently near a joint but not specifically marginal and they have sclerotic margins “**punched-out**”.

Periarticular osteopenia is absent.

Gouty tophus deposition.

Most common site for gout involvement is the first MTPJ.

Soft-tissue swelling from bursitis, such as olecranon bursitis

Multiple
punched-out
sclerotic
erosions
(arrows), with
soft-tissue
swelling



Radiological approach to Bone tumors

Bone lesion

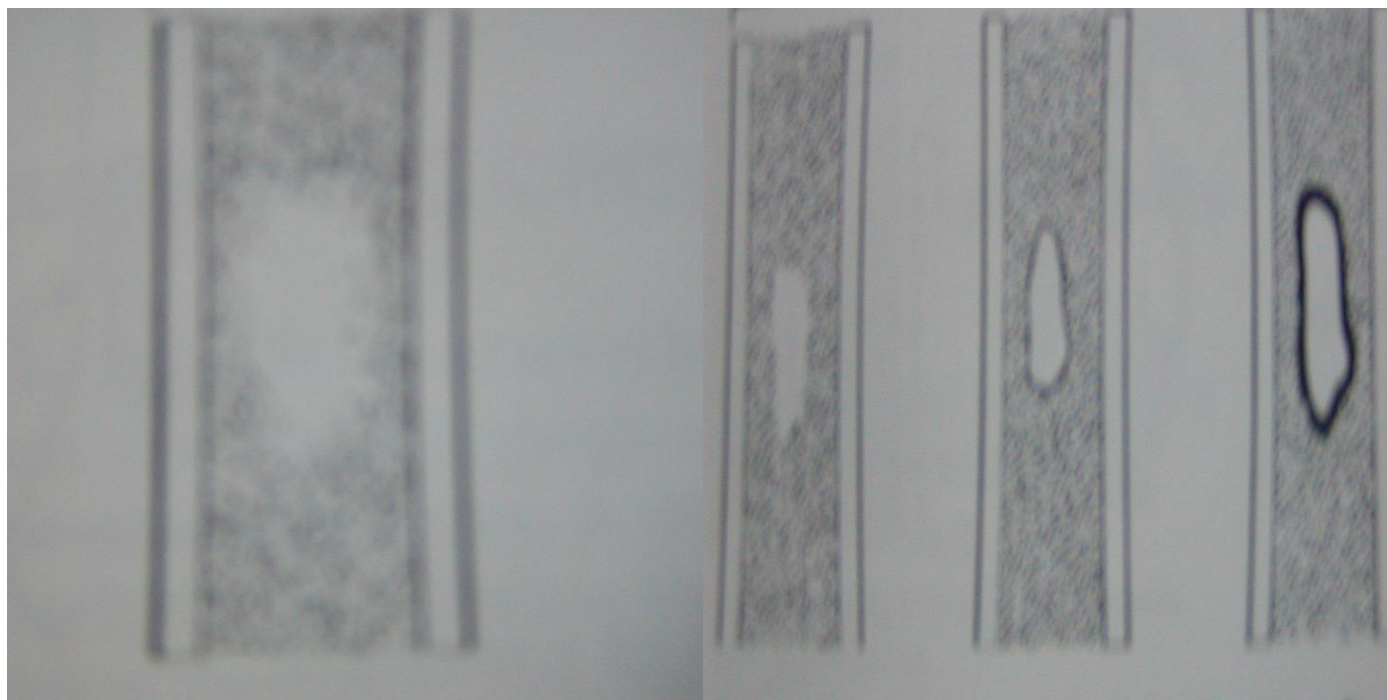
Identification then determine if aggressive or not.

Border definition (well- or ill-defined/ wide and narrow zone of transition) of the lesion

The presence of a **sclerotic rim** is a sign of a benign nature indicate non- or slowly-growing.

An ill-defined border/ wide transition zone indicates that the lesion is very **rapidly** growing.

If the lesion is well-defined but has no sclerotic rim, it may be benign or aggressive.



Aggressive lesions

An aggressive lesion may represent a malignant bone tumor or infection (osteomyelitis).

Signs that indicate malignancy:

I- Endosteal **cortical** erosion (internal scalloping).

II- **Periosteal** reactions: Codman's triangle, sun burst (hair-on-end), and interrupted periosteal reaction.

III- Evidence of **soft tissue** extension.

Osteosarcoma

Defined as a malignant mesenchymal tumor in which the cancerous cells produce osseous matrix.

75% in younger than age 20 in adolescence.

About 50% arise in the metaphysis around the knee, either in the distal femur or proximal tibia -skeletal growth-.

In older age it is usually secondary, due malignant transformation of a relatively benign tumor or other condition i.e. radiotherapy.

Osteosarcoma

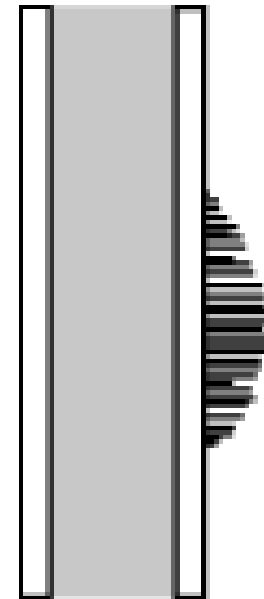
Large, ill-defined, destructive,
mixed lytic and sclerotic mass.

Frequently breaks through the
cortex and elevates the
periosteum, resulting in periosteal
bone formation.

The triangular shadow between the
cortex and raised ends of
periosteum is known as
“Codman’s triangle” and is
characteristic, though not
diagnostic feature.



Osteosarcoma



**Hair-on-end periosteal
reaction**

Ewing`s Sarcoma

Most common site: diaphysis of long bones and appears purely lytic.

Clinically: pain, swelling, fever and leucocytosis.

Plain X-ray: permeative, poorly margined destructive lesion, endosteal erosion, cortical disruption, periosteal reaction (maybe interrupted).

Associated with large adjacent extra-osseous mass (direct extension).

Ewing's Sarcoma



Ewing's Sarcoma



A

Common Benign bone tumors

Common Benign Tumors

- Symptomatic (mostly)
 - Osteoid osteoma
- Asymptomatic (mostly)
 - Fibrous cortical Defect
 - Non ossifying Fibroma “NOF”
 - Osteochondroma (exstosis)

Osteoid Osteoma

Benign bone tumor less than **2 cm** in greatest dimension and usually occur in patients in their teens and twenties.

75% of patients are **under age 25**.

Cortical lesions with a predilection for femur or tibia **50%**.

Painful: The pain is caused by excess prostaglandin E₂ which is produced by the proliferating osteoblasts.

Characteristically occurs at **night** and is dramatically relieved by **aspirin**.

Osteoid Osteoma

Hip X-Ray shows:
fusiforme cortical
thickening (blue
arrow) with a small
rounded lucent center
representing nidus
(red arrow)



Fibrous Cortical Defect and Non-ossifying Fibroma

Common, incidental.

Believed to be developmental defects rather than neoplastic process.

The vast majority arise in the diaphysis of the distal femur and proximal tibia, and almost one half are bilateral or multiple.

FCD are small and when larger 5 or 6 cm called NOF.

Fibrous Cortical Defect and Non-ossifying Fibroma



Radiological approach to Bone Infection

Acute Osteomyelitis

Plain X-ray:

Latent period of 10 days.

Deep soft tissue swelling.

Ill-defined area of bone destruction.

Solid periosteal reaction

Bone destruction becomes more prominent with time

Acute Osteomyelitis



Radiological approach to Metabolic and Endocrine Disorders

Rickets

Hyperparathyroidism

Rickets

Premature infants and usually develops between 6 and 12 months of age.

Classic radiographic signs include:

Osteopenia.

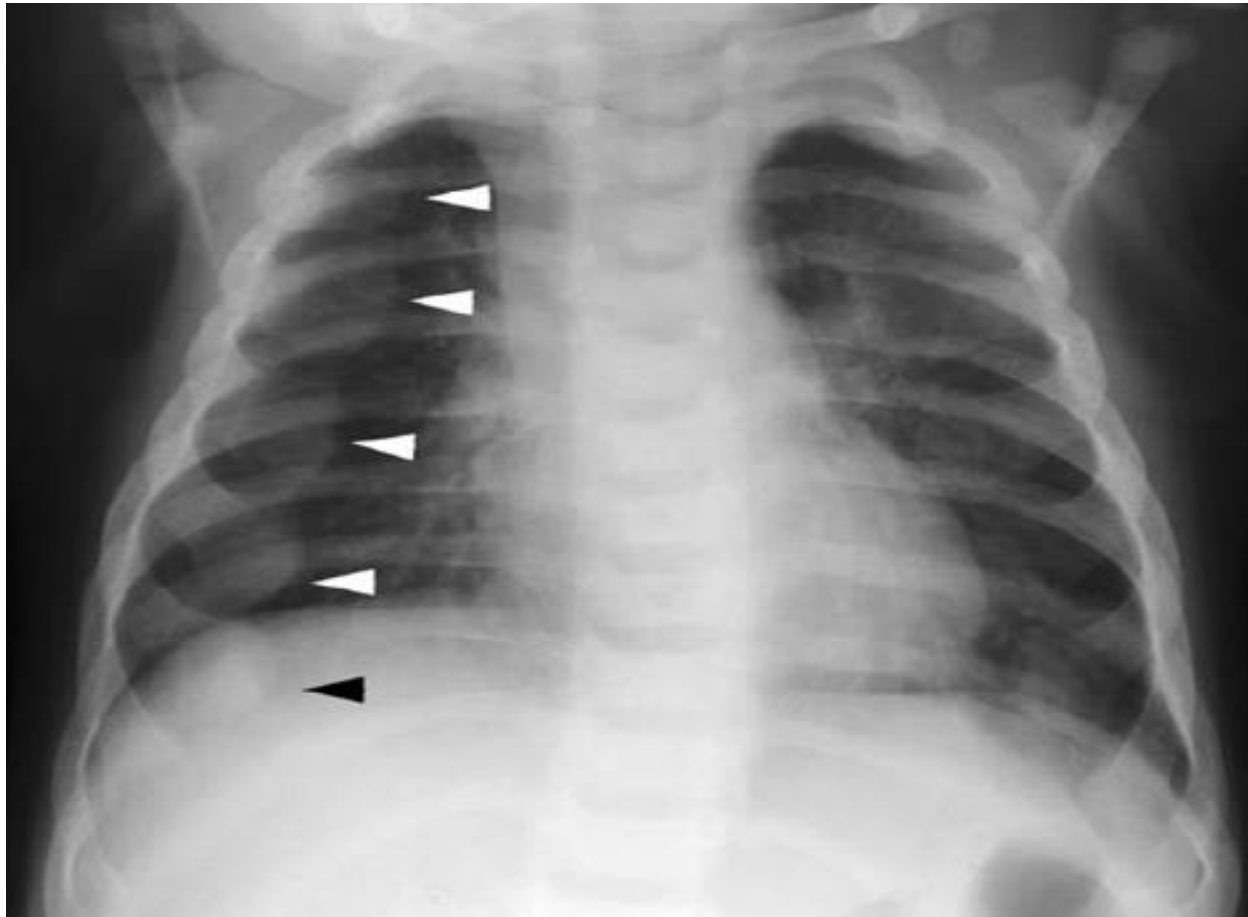
Cupping and fraying of metaphyseal ends of bone.

Delayed appearance of epiphyseal ossification centers which have blurred margins.

Excessive osteoid tissue in the sternal ends of ribs producing characteristic beading (rachitic rosary).



Rachitic rosary



Hyperparathyroidism

Excessive secretion of parathyroid primary or secondary (more common and most often due to chronic renal failure).

Classic radiographic signs include: Generalized osteopenia, brown tumors, salt-and-pepper skull, peptic ulcer, pancreatitis and gallstones.

Secondary HPT associated with sclerosis (including rugger-jersey spine), soft-tissue calcification, renal stones

Salt and pepper skull and brown tumor





**Looser's
zone and
Rugger
jersey
spine**

Incomplete stress fractures.

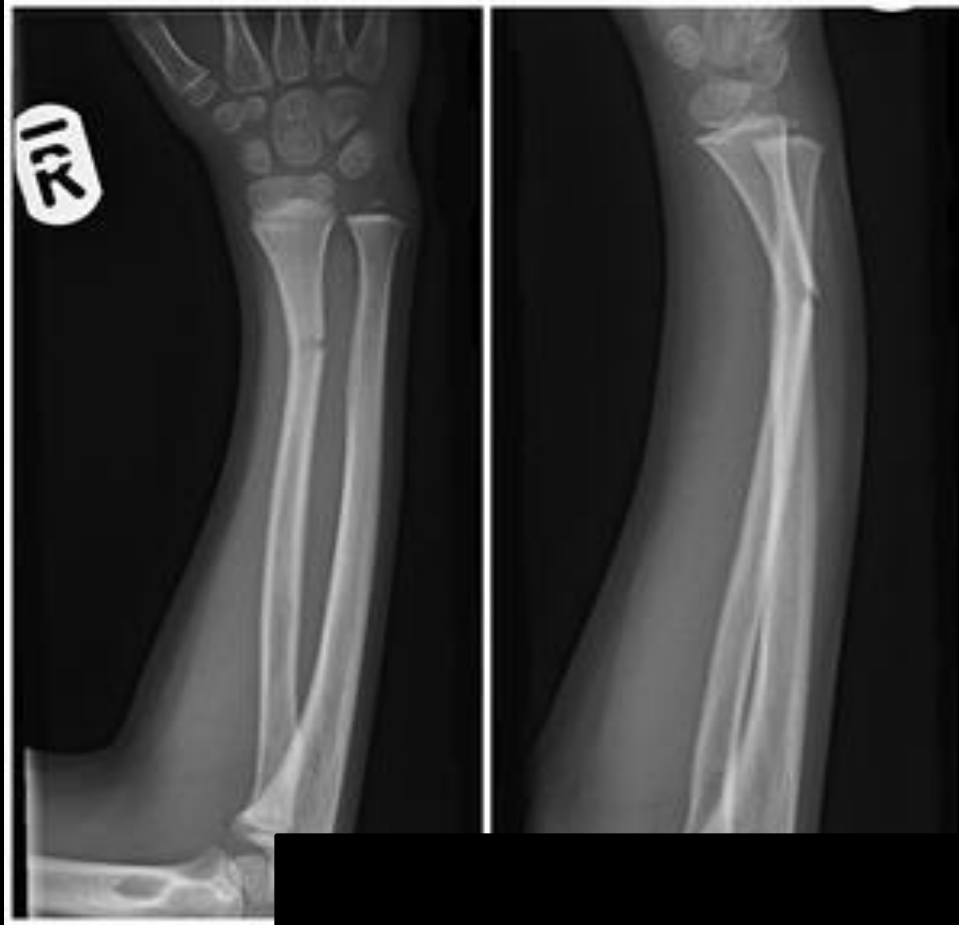
Radiological approach to fracture

- Fracture can occur at any site.
- Fractures could be classified into:
 - ➔ Displaced **vs** non-displaced
 - ➔ Partial **vs** complete
 - ➔ Comminuted **vs** non-comminuted
 - ➔ **Traumatic vs pathological**
 - ➔ Extension through the growth plate at younger age group “Salter Harris”

Displaced vs non-displaced



Partial vs complete



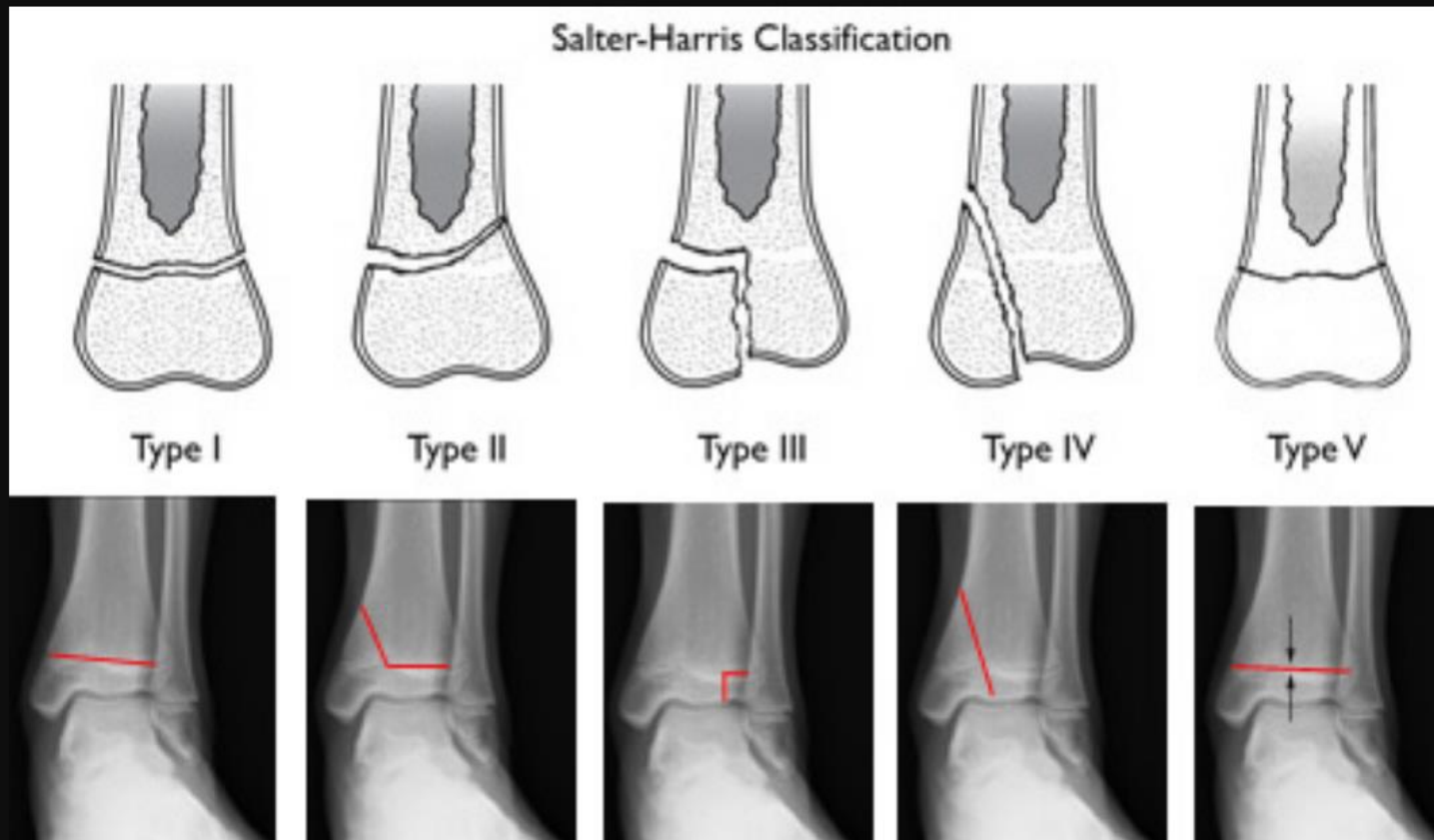
Comminuted vs non-comminuted



Traumatic vs pathological



Growth plate extension “Salter Harris”



- SH I (Slipped or Straight through): fracture plane passes through the growth plate without involving the bones Salter
- SH II (Above): commonest; metaphysis + epiphyseal plate; corner sign (metaphyseal chip)
- SH III (Low): epiphyseal plate + epiphysis
- SH IV (Together): Metaphysis + Epiphyseal plate + Epiphysis
- SH V (Rammed): Crush injuries to the epiphyseal plate; rarest; often missed diagnosis.

Radiological approach to child abuse

Child abuse

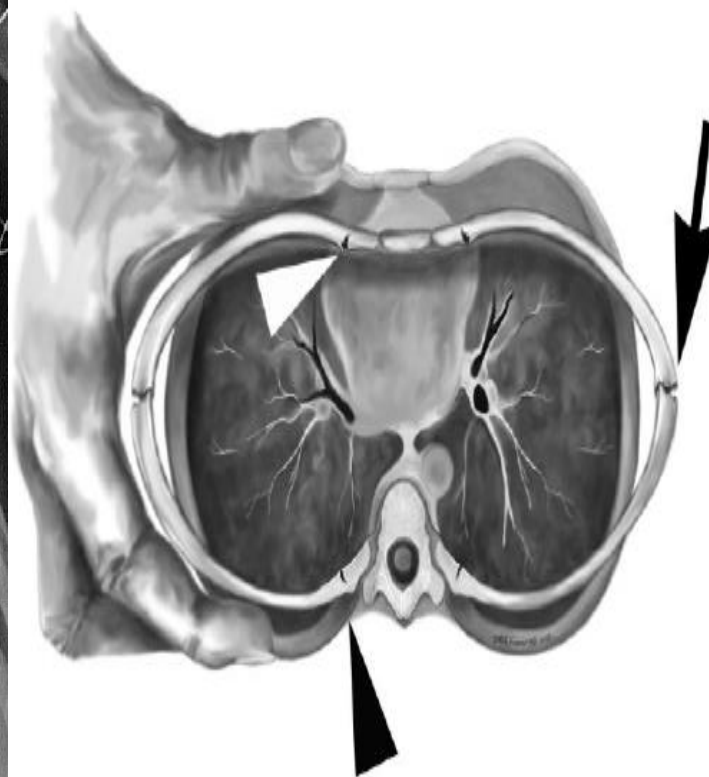
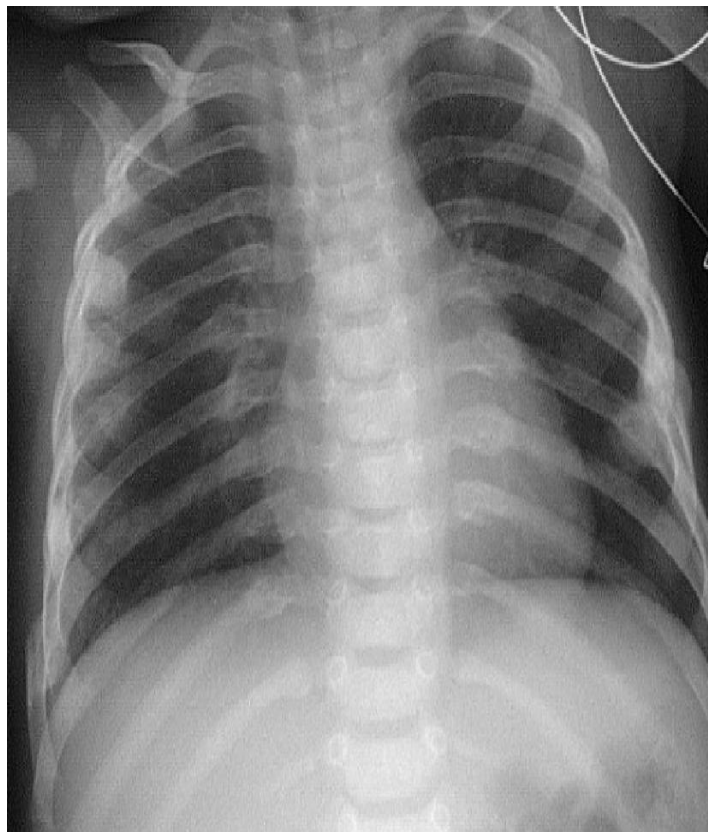
Should be considered in children with a **suspicious** clinical history and supporting examination findings.

Skeletal trauma is the most commonly seen injury in non-accidental injury.

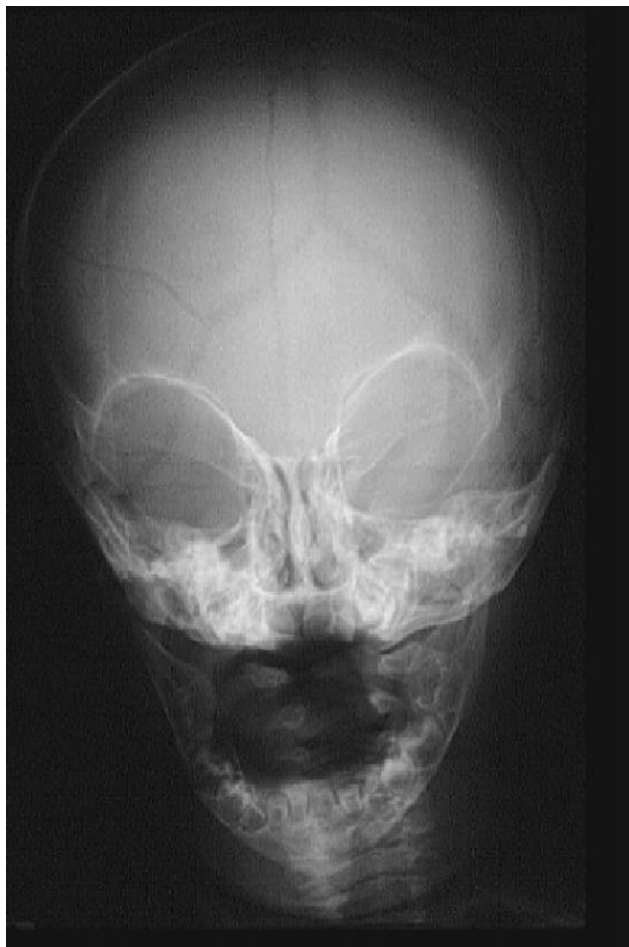
The presence of **multiple** fractures with varying of healing **ages** is characteristic.

Classical: Acromial fx, Metaphyseal corner fx, skull fx,
Bilateral posterior rib fx, Spiral fx of proximal humerus

Child abuse



Child abuse



**Medially
displaced
Ulna and
radius
relative to
humerus**



