Chest X-Ray as the First Pointer in Various Skeletal Dysplasia and Related Disorders

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Abstract

Keywords
► radiograph
► CXR
► dysplasia
► mucopolysaccharidosis
► spondylocostal dysostosis

Chest X-ray (CXR) is the most commonly used imaging modality. It is commonly used for respiratory or cardiac ailments; however, it is also used routinely as a part of skeletal surveys. In the case of suspected skeletal dysplasia, the viewer is alerted regarding the presence of some skeletal abnormality. But in case of a routine CXR performed for some other reason, it is not uncommon to miss subtle pointers of skeletal dysplasia. Sometimes routine CXR is the first pointer to alert a radiologist toward some generalized skeletal anomaly and therefore, initiate its proper evaluation by the skeletal survey.

Introduction

Skeletal dysplasia comprises a group of disorders involving abnormalities in the bones and cartilages. According to the current nosology of skeletal dysplasias,1 there are 461 diseases and 42 classes of disorders. The skeletal survey is the most important radiological investigation in the assessment of dysplasias. Chest X-ray (CXR) forms an integral part of the skeletal survey. While assessing chest radiographs, a systematic approach is to be followed. Chest radiographs can reveal characteristic findings in the bones (including clavicles, ribs, vertebrae, sternum, scapula, humeral head) in a large number of skeletal dysplasias. In an unsuspected case, CXR can even provide the first diagnostic clue to skeletal dysplasia. This article will review the chest radiograph findings in various common and clinically important skeletal dysplasias.

Imaging Features in Specific Skeletal Dysplasias

Since a detailed discussion of all dysplasias is beyond the scope of this article, we shall have an imaging feature-based discussion, based on the information a chest radiograph (►Fig. 1) can provide.

An assessment of the bone density should be the first step. A generalized decrease in bone density points toward a metabolic bone disease or a specific group of bone dysplasia.
(e.g., osteogenesis imperfecta). On the contrary, an increased bone density can be indicative of sclerosing bone dysplasia, besides other diagnostic considerations.

**Normal Bone Density**

**Abnormality in the Ribs**

*Mucopolysaccharidoses (MPS):* Defect in mucopolysaccharide degradation pathway leads to its progressive accumulation in the body, predominantly in central nervous system, bones, and heart.

Prominent imaging finding on a chest radiograph (CXR) includes paddle shaped/spatulated appearance of ribs (Fig. 2). Inferior beaking of vertebrae in MPS type I and central beaking in MPS type 4 is evident on a lateral radiograph. Other imaging findings include platyspondyly and widening of disk spaces, varus deformity of humerus with mildly hypoplastic epiphysis and proximal humeral notching, small scapula with spaces, varus deformity of humerus with mildly hypoplastic epiphysis and metaphyseal flaring. Characteristic vertebral changes are evident on a lateral radiograph and include platyspondyly and anterior tongue-like projections. Other imaging findings in the skeletal survey are listed in Table 1.4

*Hereditary multiple exostoses:* It is caused by an abnormality in bone remodeling of epiphyseal growth plate cartilage. CXR may show bony outgrowth having continuity with underlying ribs (sessile form may not be very apparent; Fig. 5). Similar exostoses may be evident on other visible bones as well.

*Encondromatosis:* In this disorder, multiple enchondromas are associated with faulty cartilaginous development and abnormal formation of intraosseous cartilaginous foci. Enchondromas may be seen on CXR as expansive lytic rib lesions, with/without ring or arc-like calcific foci within (Fig. 6). When no calcific focus is apparent, it closely mimics fibrous dysplasia.

*Fibrous dysplasia:* It is characterized by the localized developmental defect of osteoblast with its replacement by fibrous tissue and immature woven bone. On CXR, the lesions are seen as smooth expansive lytic ground glass lesion with “rind sign” (Fig. 7). Polyostotic fibrous dysplasia may be associated with deformity of the long bones as well.

*Spondylocostal dysostosis:* Various types of spondylolocostal dysostosis are described. The rib anomalies on CXR include absence, abnormal fusion, abnormal bital ribs, etc., always associated with variable segmentation anomalies involving ≥ 10 thoracic vertebrae (Fig. 8).

*Nonaccidental injury:* Its hallmark is evidence of repeated injury clinically and radiologically as a consequence of child abuse in infants/young children. Sinister findings include multiple posterior rib fractures of different ages. Other imaging clues on a CXR include scapular fracture (Fig. 9), sternal fracture (difficult to detect on CXR), and the classic metaphyseal lesion or bucket-handle type of fracture involving the proximal/distal humeri. Other imaging findings are enlisted in Table 1.

*Dysplasias with short ribs:* Normally, the anterior ends of the true ribs are seen anteriorly for about half the hemithoracic diameter. When the ribs are not seen in the anterior part/stops right after the lateral ends, a short-rib dysplasia may be looked for. Important entities in this group are short-rib polydactyly syndromes (SRPS), Jeune’s asphyxiating thoracic dystrophy (JATD), and Ellis Van Creveld syndrome (EVC). On a molecular basis, they all fall in the spectrum of ciliary disorders (ciliopathy). CXR gives a clue to diagnosis; however, imaging differentiation between them is not always possible based on a CXR alone and may need a complete skeletal survey. SRPS has polydactyly and a small narrow thorax with short ribs. JATD shows a typical narrow “bell-shaped” thorax and elevated “handle-bar” clavicles (Fig. 10). EVC clinically presents...
<table>
<thead>
<tr>
<th>Disease entity</th>
<th>Findings in CXR</th>
<th>Findings in other radiographs</th>
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<tbody>
<tr>
<td><strong>Mucopolysaccharidoses</strong></td>
<td>Paddle shaped/spatulated appearance of ribs (tapered posteriorly and widened anteriorly) Thick and short clavicles Thoracolumbar kyphosis/gibbus Vertebral bodies malformation • Inferior beaking (middle beaking in morquio) • Posterior scalloping Varus deformity of humerus with mildly hypoplastic epiphysis Proximal humeral notching Short sternum Small scapula with glenoid cavities flattening</td>
<td>Macrocephaly Thickening of cortical bone Abnormal J- shaped sella turcica Lack of pneumatization of paranasal sinuses Observe mandibular angle Prognathism Atlantoaxial instability Cervical kyphosis Anteroinferior/ central beaking Anisospondyly Rounded iliac wings Inferior tapering of ilium, Hip dysplasia Poorly developed acetabulum Underdeveloped proximal femoral epiphysis Coxa valgus Genu valgum Irregular hypoplastic tarsal and carpal bones Proximally pointed metatarsals and metacarpals Bullet shaped phalanges</td>
</tr>
<tr>
<td><strong>Pseudoachondroplasia</strong></td>
<td>Costa-transverse and costo-chondral junction widening Shortening of long bones (proximal &gt; distal) Irregular and fragmented epiphysis Metaphyseal flaring Central anterior tongue appearance (pathognomic): Anterior part of vertebral body has tongue like protrusion (in lateral radiograph) Platyspondyly: At older age Disc space widening</td>
<td>Odontoid dysplasia Squared pelvis with broad iliac wings Narrow saccroscatic notch Poorly formed acetabulum with horizontal roofs Shortening of long bones (proximal &gt; distal) Irregular and fragmented epiphysis Metaphyseal flaring Medial beaking of femoral neck (characteristic) Skull, facial bones and interpeduncular distance are normal (vs achondroplasia)</td>
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<tr>
<td><strong>Non accidental injury</strong></td>
<td>Multiple rib fractures, posterior especially Classic metaphyseal lesions- bucket handle, corner fracture (humerus) Sternal fractures Scapular fractures Spinous process fractures Clavicular fractures Costochondral fractures of humerus Epiphyseal fractures Vertebral body fractures Note: Fractures of different ages are highly suspicious</td>
<td>Complex and linear skull fracture Classic metaphyseal lesions- (bucket handle, corner fracture other long bones) Fractures of long bones Epiphyseal fractures and separations Subperiosteal new bone formation Digital fractures</td>
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<td><strong>Campomelic dysplasia</strong></td>
<td>Hypoplastic vertebral bodies (thoracic vertebral pedicle absence) Hypoplastic scapulae 11 pair of ribs instead of 12 Narrow thorax (may be bell shaped) Scoliosis/kyphoscoliosis Short bowed humerus</td>
<td>Narrow iliac wings Poor pubis ossification Short bowed femur</td>
</tr>
<tr>
<td><strong>Spondyloepiphyseal dysplasia congenita</strong></td>
<td>Platspondyly (with maintained bone density) Pear shaped and bulbous vertebrae may be seen Kyphoscoliosis Intervertebral disc space is narrowed Delayed ossification of humeral epiphysis Bowing of humerus, short bone, early onset arthritis can be seen</td>
<td>Flat face Micrognathia Delayed ossification of pubis Bowing of femur can be seen</td>
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<tr>
<td><strong>Achondroplasia</strong></td>
<td>Marked symmetrical shortening of long bones particularly humerus Normal epiphysis with relative splaying and flaring involving metaphysis, may give ball and socket appearance Kyphoscoliosis Posterior scalloping of vertebral bodies Narrowing of spinal canal Height of discs nearly equal to vertebral body Short and thick pedicle Ulna often shorter than radius</td>
<td>Large cranial vault and small skull base Foramen magnum narrowing Short hand phalanges with trident-hand appearance Caudal narrowing of lumbar interpediculate distance Notch-like sacroiliac groove Decreased acetabular angles Small and square iliac wings (tombstone-shaped) Champagne glass shaped pelvic inlet Long fibulae</td>
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with postaxial polydactyly, disproportionate dwarfism (acromesomelia), congenital heart defects (ASD most common), nail, and teeth changes. Apart from short ribs, one typical imaging finding is the outward bowing of the humerus along with short forearm bones (►Fig. 11).

**Caffey’s disease:** It is a self-limiting disease usually presenting before 5 months of birth. Pathological hallmark is periosteal new bone formation with associated cortical thickening. The common clinical manifestation is classic triad including hyperirritability, soft tissue swelling, and hard mass over affected bones (most commonly involving mandible, clavicle, and ribs symmetrically). The imaging findings in CXR in Caffey’s disease include thickened cortex and widened clavicle, cortical hyperostosis involving lateral ribs, unilateral involvement of scapula common (may be confused with malignancy), thick cortex involving diaphysis of tubular bones, and sparing epiphysis (ulnar involvement more common; ►Fig. 12). Interoosseous bridging can be seen in ribs and between radius and ulna. Vertebrae are spared. The close differential is child abuse, however symmetrical involvement of clavicle and ribs is seen in Caffey’s disease.

Other diseases having abnormal shapes of ribs include neurofibromatosis 1 (twisted ribbon ribs) and Melnick-Needles syndrome (wavy ribs with diaphyseal constrictions and curvature in long bones), among others (►Fig. 13).

**Abnormality in the Thoracic Spine**

**Campomelic dysplasia:** It is diagnosed antenatally commonly with key features including narrow thorax, bowed femur and

### Table 1 (Continued)

<table>
<thead>
<tr>
<th>Disease entity</th>
<th>Findings in CXR</th>
<th>Findings in other radiographs</th>
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<tbody>
<tr>
<td><strong>Metaphyseal chondrodysplasia</strong></td>
<td>Metaphyseal broadening and irregularities with preserved epiphysis</td>
<td>Wide-open sutures and patent fontanelles, Wormian bones, Depressed nasal bridge, Hypertelorism, Hypoplastic mid-facial region, Prognathic mandible, High arched palate, Retention of deciduous teeth, Delayed eruption of permanent dentition, Short/absent radius may be seen, Wide pubic symphysis, Hypoplastic iliac wings, Widened sacroiliac joints, Large femoral neck with coxa vara, Lengthening of the second metacarpal, Hypoplastic pointed phalanges may be seen.</td>
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<tr>
<td><strong>Clidocranial dysostosis</strong></td>
<td>Anomalous clavicular development (partly/completely absent)</td>
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<td></td>
<td>Short and oblique ribs, may be supernumerary</td>
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<td>Delayed mineralization, hemivertebrae, spina bifida occulta and biconvex bodies may be found in spine (upper thoracic predominantly)</td>
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<td></td>
<td>Small, winged and elevated scapula</td>
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<td>Narrow/cone shaped chest (frequent)</td>
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<td></td>
<td>Kyphoscoliosis</td>
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<td></td>
<td>Shoulder dislocation may be seen</td>
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<tr>
<td><strong>Osteogenesis imperfecta</strong></td>
<td>Osteopenia</td>
<td>Osteopenia, fractures (long bone diaphyses, apophyses and spine -most common site)</td>
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<td></td>
<td>Hyperplastic callus, Pseudoarthrosis, Deformities, Ossification of interosseous membrane, popcorn calcification, dense metaphyseal bands</td>
<td>Deformities, Prominent occipital region (Darth Vader appearance), Cranial vault flattening with transverse infolding of base (Tam O’Shanter skull), diaphyseal bending or angulation, Wormian bones</td>
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<td></td>
<td>Platyspondylodycodfish vertebra Spondylylosis and spondylolisthesis, kyphoscoliosis can be seen</td>
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**Fig. 3 (A,B)** Mucolipidosis. CXR (A) shows uniformly widened ribs without increased density, pelvis and lower limb radiograph (B) shows widened undermodeled femora with periosteal thickening (arrows), note comma shaped iliac bones (dotted arrows).

**Fig. 4 (A,B)** Pseudoachondroplasia. CXR (A) shows striking finding in the form of widening of costovertebral junctions (arrows), lateral radiograph of DL spine (B) shows anterior tongue like projection in the vertebral bodies (arrows).
tibia, and hypoplastic scapulae. Imaging findings include the absence of thoracic vertebral pedicles. Other important features include hypoplastic scapulae, occasional short bowed upper limbs, and 11 pairs of ribs (Fig. 14). Other imaging features are listed in Table 1.

Spondyloepiphyseal dysplasia congenita (SEDC): This is a form of collagenopathy, having clinical features secondary to multiple organ affection. CXR shows universal platyspondyly (more evident on a lateral view) with maintained bone density, kyphoscoliosis, and delayed ossification of humeral epiphysis. Secondary to the epiphyseal abnormality, early degenerative changes may be evident at multiple joints. Other imaging features are listed in Table 1.

Spondyloepiphyseal dysplasia tarda (SEDT): Many forms exist. Diagnostic imaging findings on a CXR include platyspondyly with posterior “heaping” of vertebral bodies (Fig. 15). Visualized epiphyses show irregularity and early degenerative changes.

Chondrodysplasia punctata (CDP): It has an abnormal epiphyseal osseous nucleus and has two main forms, rhizomelic type (lethal) and Conradi–Hünermann variety (milder type). Imaging findings on a CXR include stippling of vertebral bodies (Fig. 16), coronal clefting (lateral radiograph), kyphoscoliosis, and stippled epiphysis of the humerus. Short humeri are evident in the rhizomelic form.

Spondylometaphyseal dysplasias (SMD): On a frontal CXR, the most prominent abnormalities lie in the long bone metaphyses (discussed later).

Long Bone (Humerus/Ulna) Abnormal

Achondroplasia: It is the most common cause of congenital dwarfism (rhizomelic) resulting from disturbance in FGF3 gene. On a CXR, the most striking abnormality is a shortening of humeri (rhizomelic shortening). The proximal...
humeral metaphyses are flared, and the inferior angle of scapula are square in shape (Fig. 17). On lateral CXR, several other imaging findings may be appreciated, namely, posterior or scalloping of vertebral bodies, kyphoscoliosis, and short and thick pedicles. Other characteristic imaging findings are discussed in Table 1.

Metaphyseal chondrodysplasia (MCD): Several forms exist (e.g., Schmid, McKusick, and Jansen type). The common imaging finding evident on a frontal CXR is irregularity and broadening of proximal humeral metaphyses (Fig. 18). It is important to differentiate them from metabolic bone disease (rickets) and spondylometaphyseal group of disorders. While rickets show reduced bone density, widening of growth plate, and widening of anterior rib ends, MCDs usually have normal bone density. A differentiation of MCD from SMD requires evaluation of a complete skeletal survey although a lateral CXR can also provide a clue about the shape and size of vertebrae. Differentiation between various types requires the analysis of patterns of metaphyseal involvement. Characteristic imaging findings are discussed in Table 1.

SMD: It is a heterogeneous group of disorders consisting of several different varieties; SMD is manifested on radiographs as a combination of metaphyseal and spinal abnormalities. The spinal manifestations vary, ranging from universal platyspondyly and medially located rounded pedicles (in Kozlowski subtype) to rounded vertebral bodies (in Sutcliffe subtype) (Fig. 19).

SEDC and SEDT: These entities are discussed above in the section “Abnormality in the Thoracic Spine.”

Non-accidental injury: This entity is mentioned above in the section “Abnormality in the Ribs.”

Multiple epiphyseal dysplasia (MED): On a CXR, notable imaging findings of MED include delayed ossification of epiphysis with irregular flattened epiphysis, early osteoarthritic changes, and metaphyseal widening.

Metatropic dysplasia: This rare dysplasia is evidenced on a frontal CXR with rhizomelic shortening with prominent metaphyseal flaring (resembling dumbbell appearance) of the humeri (Fig. 20). Other features in chest radiograph include marked platyspondyly and kyphoscoliosis.

Diastrophic dysplasia: Clinical clues in this dysplasia include a deformed “Hitchhiker’s thumb” and cauliflower ear. On a frontal CXR, imaging findings include shortened long bones with metaphyseal flaring.

Pseudoachondroplasia, MPS, EVC, HME, SEDC, and CDP also show abnormality in the long bones (discussed earlier).

Clavicle Abnormal

Cleidocranial dysostosis: It is characterized by defective ossification of membranous and enchondral bones (Fig. 21). Diagnostic imaging finding is an anomalous
clavicular development (partly/completely absent). Other features in CXR include supernumerary ribs, short or oblique ribs, hemivertebrae, spina bifida occulta, biconvex bodies, and kyphoscoliosis (upper thoracic predominantly). Other features are listed in Table 1.29

Nonaccidental injury, MPS, JATD, and Caffey’s disease also show abnormality in clavicles (discussed above).

Soft Tissue Abnormality

Fibrodysplasia ossificans progressiva: It involves progressive ossification of soft tissue (striated muscle, ligaments, tendon, fascia, and aponeuroses) with associated congenital micro-

dactyly of great toe, and hallux valgus. CXR may show sheet-like soft tissue ossification in the lateral chest wall, axilla, and neck (Fig. 22).30 Secondary to these heterotopic ossifications, variable spinal deformities may ensue, eventually leading to restrictive lung disease and respiratory failure.30

Calcinosis universalis: It is characterized by the diffusion of calcium deposition (extensive sheet-like) in skin, subcutaneous tissue, tendons, or muscles.31
Reduced Bone Density

Abnormality in Ribs and/or Thoracic Spine

*Achondrogenesis*: This lethal dysplasia is characterized by decreased bone mineralization, micromelia, short trunk, and macrocranium. On a CXR, the crucial imaging finding is poor ossification of vertebral bodies and only pedicles visualized. Other features in chest radiograph include shortened long bones (as a result of multiple intrauterine fractures) and thin ribs with/without fracture (fracture in Type 1A).
Osteogenesis imperfecta (OI): It is characterized by reduced bone density with increased fragility due to abnormal type I collagen. It has different types of varying severity. Imaging findings include diffuse osteopenia, platyspondyly/codfish vertebra, thin ribs, and multiple fractures with/without hypertrophic callus formation (►Fig. 23). Secondary to multiple repeated fractures, several deformities may be evident. Ossification of interosseous membrane is a typical imaging finding in type V OI. Popcorn calcification and dense metaphyseal bands can be seen. Other characteristic imaging findings are discussed in ►Table 1.

Abnormality in Long Bones

Achondrogenesis: It is explained in "Reduced Bone Density" section.

OI: It is mentioned above
Increased Bone Density

There are few skeletal dysplasias which present with diffuse increased bone density; however, increased bone density can be a normal finding in a neonatal CXR and needs to be meticulously evaluated.

Abnormality in the Ribs/Spine

Osteopetrosis: It is caused by defective osteoclastic resorption leading to abnormal bone maturation with the formation of dense and brittle bones. Severe cases are diagnosed in infancy with clinical features including anemia, hepatosplenomegaly, bleeding episodes, fractures, and failure to thrive. Diagnostic imaging pointers are sclerotic ribs with obliteration of medullary cavity, sclerotic clavicle, and humeri with “bone within bone” appearance, sandwich vertebrae on a lateral CXR, and evidence of repeated fractures (►Fig. 24).

Pyknodysostosis: This often comes as an imaging differential of osteopetrosis. It is characterized by osteosclerosis; however, medullary canal of long bones is preserved (versus osteopetrosis). Dense vertebral bodies with sparing of the transverse processes and a typical “spool shape” are described. Other important associated findings are acro-osteolysis, wide open sutures, obtuse mandibular angle, etc. (►Fig. 25).

Table 2 Role of CXR in skeletal dysplasia diagnosis

<table>
<thead>
<tr>
<th>Conditions where CXR alone may be sufficient</th>
<th>Conditions where CXR plays an important adjunct role</th>
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<tbody>
<tr>
<td>• Spondylocostal dysostosis</td>
<td>• Osteopetrosis</td>
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<tr>
<td>• Short rib dysplasias</td>
<td>• Craniodiaphyseal dysplasias</td>
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<tr>
<td>• Ellis–van Crevel syndrome</td>
<td>• Achondroplasia</td>
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<tr>
<td>• Cleidocranial dysostosis</td>
<td>• Pseudoachondroplasia</td>
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<tr>
<td>• Osteogenesis imperfecta</td>
<td>• Mucopolysaccharidosis</td>
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<tr>
<td>• Fibrodysplasia ossificans progressiva</td>
<td>• Polyostotic fibrous dysplasia</td>
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<tr>
<td>• Polyostotic fibrous dysplasia</td>
<td>• Enchondromatosis</td>
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<tr>
<td>• Hereditary multiple exostoses</td>
<td>• Hereditary multiple exostoses</td>
</tr>
<tr>
<td></td>
<td>• Epiphyseal dysplasias</td>
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<td>• Metaphyseal dysplasias</td>
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Abbreviation: CXR, chest X-ray.

Table 3 Conditions where CXR is diagnostic at birth

| JATD | SRPS | Achondroplasia | Osteogenesis imperfecta |

Abbreviations: CXR, chest X-ray; JATD, Jeune’s asphyxiating thoracic dystrophy; SRPS, short-rib polydactyly syndromes.

Long Bone (Humerus/Ulna) Abnormal

Several other sclerosing bone dysplasias may present with sclerosis of long bones, namely, osteopathia striata, melorheostosis, progressive diaphyseal dysplasias, and craniodiaphyseal dysplasias (►Fig. 25). While some (melorheostosis) may have characteristic imaging findings on a single CXR, most of them require a full skeletal survey and clinical details to reach a diagnosis.

Fig. 24 Osteopetrosis with rickets. Note the increased densities involving all ribs, humerus, and vertebrae, obliterated medullary cavities. Rachitic changes are evident in the form of metaphyseal widening and irregularity.

Fig. 25 (A,B,C) Craniodiaphyseal dysplasia in a 15 year boy, having no organomegaly or anemia. CXR (A) shows diffuse osteosclerosis (arrows) involving all visualized bones, mimicking osteopetrosis. However, lateral skull radiograph (B) reveals extreme bone sclerosis of facial and calvarial bones (arrows), hand radiograph (C) show diaphyseal sclerosis involving the phalanges and metacarpals (arrows); however, the medullary cavity (bent arrow) is maintained.
The Role of Chest X-Ray Alone in Making the Diagnosis

Most often, CXR is useful as a part of the whole skeletal survey. However, there are a few conditions where the diagnosis of the specific skeletal dysplasia/close differential can be reasonably accurately made on a CXR alone. These are listed in Table 2. There are few dysplasias where CXR is diagnostic at birth as listed in Table 3.

Conclusion

Skeletal dysplasia is not a single-film imaging diagnosis. It requires evaluation of the complete skeletal survey, along with appropriate clinical details. However, often the CXR may be the first imaging clue to an unsuspecting case, being evaluated for unrelated causes. Hence, radiologists should be aware of the various subtle and overt imaging findings of different dysplasias on a CXR. Segment-wise approach toward skeletal abnormalities in a CXR is given in Fig. 26.

Availability of Data and Material

Data available on request.

Contribution of Authors

R.J. contributed to data collection, literature review, and analysis and drafted the manuscript and critical revision. P.N. contributed to data collection, literature review, and analysis and drafted the manuscript and critical revision. N.G. contributed to data collection, literature review, analysis, and critical revision. P.S. contributed to data collection, literature review, analysis, and critical revision. M.K. contributed to data collection, literature review, analysis, and drafted the manuscript and critical revision. A.K.G. contributed to data collection, literature review, analysis, and critical revision. M.J. contributed to data collection, literature review, and analysis and drafted the manuscript and critical revision.

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Conflict of Interest

None declared.

Declaration of Patient Consent

The authors certify that they have obtained all appropriate patient consent forms. In the form, the patient(s) has/have given his/her/their consent for his/her/their images and other clinical information to be reported in the journal. The patients understand that their names and initials will not be published and due efforts will be made to conceal their identity but anonymity cannot be guaranteed.

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