A Report of an Indian Boy with a Delayed Diagnosis of Pseudochondroplasia

ABSTRACT
The mutations in the Cartilage Oligomeric Matrix Protein (COMP) gene are associated two common and allelic bony dysplasias: Pseudochondroplasia (PSACH) and Multiple epiphyseal dysplasias-1 (MED-1). The characteristic radiological features of both have been well established in the literature, with areas of overlap between the two in certain forms of mild PSACH and severe MED. MED is also a genotypically and a phenotypically heterogeneous disease. Here, we emphasise the salient radiological features which aid in the diagnosis of PSACH and COMP MED; which may enable a targeted molecular analysis.

Key words: Pseudochondroplasia, Multiple Epiphyseal Dysplasia, COMP gene

INTRODUCTION
Pseudochondroplasia (PSACH) and Multiple Epiphyseal Dysplasia type 1 (MED1) are autosomal dominant skeletal dysplasias which are caused by mutations in the COMP gene. PSACH manifests around 2 years of age and it is characterised by a moderate to severe short stature, deformities of the lower limbs, vertebral anomalies and early onset progressive osteoarthritis. MED 1, on the other hand, is characterised by an onset in mid or late childhood, a mild to moderate short stature and a normal spine. The radiological changes progress and they change or evolve with increasing age and if they are seen later, it may be difficult to differentiate between mild PSACH and severe COMP MED. The COMP gene mutations, though they are available as clinical tests, are expensive. So, we attempted to highlight the salient features which could help the clinicians in making the decision of COMP testing.

CASE REPORT
A 9-years old male child presented with a short stature and bony deformities. He was born of a non – consanguineous parentage. The antenatal and the perinatal periods were uneventful. The child’s growth was normal till the age of two years. Thereafter, he started showing a decrease in the growth velocity and he developed lower limb deformities. The child achieved his developmental (motor, fine motor, language, personal and social) milestones normally and he presently studies in class seventh and is doing average in studies.

On examination, he was found to have a severe short stature with a height of 92cm (< - 3 SD scores) with disproportionately short limbs (US/LS= 1.24). He had micromelia and his arm span was 82 cm. He had widening at all the joints, a bilateral genu varus deformity, a waddling gait and short stubby fingers. He did not have any dysmorphic facial features, contractures or kyphoscoliosis. He had a normal dentition and a normal IQ (IQ-110, was assessed by DASIll). His systemic examination was normal.

Radiography revealed a marked flaring of the metaphyses at all the joints [Table/Fig-1 and 2]. The epiphyses were fragmented, deformed with delay in ossification. The carpal bones were deformed and there was a markedly delayed carpal bone age [Table/Fig-3]. The spine showed undulations of the superior and the inferior plates of the multiple dorsolumbar vertebrae, which resembled Schmorl’s nodes and anterior tonguing [Table/Fig-4 and 5]. An X-ray of the pelvis showed bilateral iliac flaring, coxa vara and a tri radiate acetabulum [Table/Fig-6].

[Table/Fig-1]: Radiography revealed marked flaring of the metaphyses at all the joints, epiphyses were fragmented, deformed with delay in ossification.

[Table/Fig-2]: Small, late appearing epiphysis and flared, ragged metaphyses
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suspected. His mutational analysis revealed a novel de novo missense mutation in the exon 14 of the COMP gene [c.1554C>G (p.Asp518Glu)].

**DISCUSSION**

PSACH and MED are a group of osteochondrodysplasias with variable phenotypes and heterogeneous genotypes. PSACH has a single genetic locus (19p13.11) with single gene (COMP) involvement, whereas MED has multiple genetic loci (19p13.11, 1p33-p33.2, 20q13.3, 5q32-q33.1, 2p24-p23,6q13) with multiple gene involvements (COMP, COL9A2, COL9A3, SLCE26A2, MATN3, COL9A1). PSACH has frequency of 4 per million population, while MED has high frequency of 9-16 per million [1,2,3]. PSACH and MED -1, 3, 5, 6] are inherited in an autosomal dominant manner, except MED4, which is inherited in an autosomal recessive form. COMP-MED is the commonest of all the six MEDs and it is allelic to PSACH. The COMP mutations are responsible for these two allelic disorders [4,5].

COMP is a modular protein which comprises the amino-terminal coiled-coil oligomerization domain, 4 type II (EGF-like) domains, 8 type III Calmodulin like repeats (CLR) and a C-terminal globular domain (CTD); which are expressed in cartilage, tendons and ligament [6]. A majority of the COMP mutations (85 %) are clustered in the CLR repeats [7]. The COMP gene is comprised of 19 exons, with the maximum number of missense mutations clustering in the exons 13 and 14 [8]. Exon 14 is comprised of the CLR and the CTD domains. We found a missense mutation (c.1554C>G) in exon 14 in the CLR domain of chromosome 19, which had led to a protein change (p.Asp5218Glu). The calmodulin like repeats are highly conserved and so any change which involves the aspartic acid residue will lead to functional and conformational changes in the COMP gene product. This mutation was absent in the parents, sibs and in the ethnic controls. Thus, it was a de novo pathogenic mutation, which was later substantiated by doing an in silico analysis.

PSACH and COMP-MED represents the two ends of a clinical spectrum, with many cases falling in between the two extremes. It is not unusual for PSACH to get diagnosed as late as our present case (9 years). Hence, a diagnostic dilemma occurred, as this is the period of presentation of MED-1. It is difficult to distinguish mild PSACH from very severe MED, thus delaying the correct radiological diagnosis. There are certain radiological signs which have been designated as COMPY [Table/Fig-7] that can guide the clinicians in the COMP mutation testing [9].

We tabulated the radiological findings of our case and compared them with the characteristic radiological findings of PSACH and COMP-MED [Table/Fig-8]. The following radiological findings strongly favoured the diagnosis of PSACH: an anterior tongue, biconvex vertebrae, platyspondyly (Spine); short tubular bones

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<td>Very small capital femoral epiphysis</td>
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<td>Brachydactyly</td>
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<th>Spine</th>
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<td>Indentations of superior and inferior ring epiphyseal regions (early age)</td>
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[Table/Fig-7]: Radiographic MED with “COMPY” changes (suggesting COMP mutations)

The skull radiograph was normal, with a J shaped sella. His bone age (4 – 5 years) was markedly delayed, as was assessed by the Greulich and Pyle method. His baseline investigations were within normal limits. His thyroid function tests were normal. Based on his history and radiographic examination, a working diagnosis of PSACH, MED and Spondylometaphysal dysplasia was

[Table/Fig-3]: The carpal bones were deformed and there was a markedly delayed carpal bone age

[Table/Fig-4 & 5]: Spine showed undulations of the superior and inferior plates of multiple dorsolumbar vertebrae resembling Schmorl’s nodes and anterior tonguing

[Table/Fig-6]: X-ray pelvis showed bilateral iliac flaring, coxa vara and tri radiate acetabulum
with a frayed metaphysis and a fragmented epiphysis (All the large joints); a tri radiate appearance of the acetabulum with a short and beaked femoral neck (Pelvis). The findings which were characteristic of COMP-MED were- vertebral end plate irregularities with Schmorl bodies (Spine) and a delay in the appearance of the carpal/tarsal bones. However, a generalised epiphyseal ossification delay and deformed carpal/tarsal bones were the common radiological features which were seen in both. Thus, our case demonstrated major radiological findings which were suggestive of PSACH, which shared a few radiological features with COMP-MED. So, tabulating and comparing the radiological features helped in predicting the correct phenotype and in targeting the molecular study to the COMP gene rather than to other genetic loci in MED.

To conclude, this is a case of PSACH with an overlap of some features of COMP MED. It shared two unusual features of COMP MED- vertebral end plate irregularities with Schmorl bodies and a delay in the appearance of the carpal and the tarsal bones. Though the features were classical of PSACH, the diagnosis was delayed till 9 years of age. There are psychological and treatment implications of making a late diagnosis and so, we emphasise an early diagnosis in the prototype patients and highlight certain radiological clues in milder PSACH and COMP MED, to direct the COMP gene testing to reach a correct molecular diagnosis.

**REFERENCES**


