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**Case report  
An unusual case of short stature – Morquio’s syndrome**

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**Abstract:**

Mucopolysaccharidosis (MPS) are rare genetic conditions in the group of lysosomal storage disorders, characterised by constellation of symptomsand signs due to accumulation of lysosomes by undegraded glycosaminoglycans(GAGs) resulting in cellular and organ dysfunction. Morquio’s syndrome or MPS type IV is characterised by skeletal manifestation, short stature and normal intellectual. There is paucity of clinical reports on mucopolysaccharidosis across the world and there are very few case reports of Morquio’s syndrome in Indian literature and hence this case is reported for its varied presentation and its rarity .Early clinical suspicion by a comprehensive examination of the patient for starting enzyme replacement therapy(ERT) before the onset of irreversible organ damage and a multidisciplinary approach of the patient are the mainstay of treatment for this progressive condition.

**Keywords:** Mucopolysaccharidosis, Morquio’s syndrome, short stature