



An observational case study of Morquio syndrome in a pediatric patient – a rare disease

*Amisha Bavarva * & Dr. Apexa G. Vyas*

Department of Kaumarbhritya, Institute of Teaching and Research in Ayurveda, Jamnagar

1. Dr. Amisha A. Bavarva- 3rd year MD Scholar, Department of Kaumarbhritya - Institute of Teaching & Research in Ayurveda, Jamnagar.
2. Dr. Apexa G. Vyas – Reader, Department of Kaumarbhritya - Institute of Teaching & Research in Ayurveda, Jamnagar.

Email: serviceheb@gmail.com

Abstract:

Mucopolysaccharidosis type IV (MPS IV), also known as Morquio syndrome, is a progressive condition that mainly affects the skeleton. The first signs and symptoms of MPS IV usually become apparent during early childhood. Affected individuals develop various skeletal abnormalities including short stature, knock knees, abnormalities of the ribs, chest, spine, hips and wrists. People with MPS IV often have restricted movement in certain joints. Here, we present a case of Morquio syndrome. A 14-year-old girl presented with multiple skeletal abnormalities. The physical appearance was characterized by short stature, protuberant chest, scoliosis, deformed knees and bowing of both lower extremities.

KEY WORDS: Kubjata, Morquio syndrome, MPS IV A, Mucopolysaccharidosis.

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