Case Report

Hypophosphatemic vitamin D resistant rickets in an adult: A rare case report

Vidiyala Harika1, Abhilash Mishra2, Sourya Acharya3*, Samarth Shukla4

1,2Resident, 3,4Professor, 1,2Dept. of Internal Medicine, 4Department of Pathology, Acharya Vinoba Bhave Rural Hospital and Jawaharlal Nehru Medical College, DMIMS University, Wardha, Maharashtra, India

*Corresponding author:
Email: souryaacharya74@gmail.com

Abstract
Hypophosphatemic Vitamin D resistant rickets is a rare disease, which is characterized by lack response to vitamin D at the level of receptors leading to rickets. It is also associated with disturbed metabolism and activity of vitamin D and/or disturbed phosphate metabolism. We present a case of a 19 year old male presented with symptoms and signs of Vitamin D deficiency and diagnosed as Vitamin D resistant rickets.

Keywords: Vitamin D, Phosphate, Rickets.

Received: 13th November, 2017 Accepted: 15th December, 2017

Introduction
Hypophosphatemic Vitamin D resistant rickets is a condition characterized by, features of rickets that is resistant to normal therapeutic doses of vitamin D. the treatment usually requires high doses of vitamin D. This rare condition was first described by Albright, Butler and Boonberg in 1927.

For normal calcium homeostasis, calcitriol or 1, 25-dihydroxy cholecalciferol, the active form of vitamin D is crucial. Vitamin D receptors play an important role for down regulation of calcitriol on the target tissues. This rare autosomal recessive genetic disease in humans, is caused by heterogeneous mutations in vitamin D receptor gene. The cardinal clinical presentation is characterized by features of rickets along with typical biochemical abnormalities in form of, hypocalcaemia, elevated serum calcitriol, and secondary hyperparathyroidism. Some patients can present with associated dermatological abnormalities like alopecia and dermal cyst.

Case Report
A 19 year old male presented to us with complains of back pain, groin pain and lower limb weakness since 3 years duration. The pain was intermittent and dull aching type. The lower limb weakness was gradual in onset aggravated by exercise and relieved slightly on rest. There was no history of fever, night sweats, morning stiffness, small joints swelling, paresthesias and urinary incontinence. On examination vitals were stable mild pallor was present systemic examination was normal. Musculoskeletal examination revealed short stature, bowed legs and scoliosis. (Fig. 1, 2)

Fig. 1: Showing short stature

Fig. 2: Scoliosis and Bowed legs

Investigations revealed, Normal complete blood counts. KFT, LFT was normal. X ray
Thoraco-lumbar spine revealed looser zones, decreased mineralization, reduced disc spaces. (Fig. 3) Serum calcium 6.4 mg/dl, alkaline phosphatase was 1100 IU, Parathyroid hormone: 865.5 pg/ml, serum inorganic phosphate: 1.4 mg%, whereas the values of 1, 25-dihydroxyvitamin D was high (185 μmol/L), suggesting end organ resistance to vitamin D.

![Fig. 3: X-ray Thoraco-lumbar spine showing looser zones, decreased mineralization, reduced disc spaces, and scoliosis](image)

The patient was treated with Vitamin D 50,000 IU/week for 8 weeks with tablet sodium and potassium phosphate 250 mg twice a day.

**Discussion**

Hypophosphatemic rickets is a rare autosomal recessive disorder. It usually manifests in pediatric age group. There is no established guidelines regarding indications of treatment of this disease in adult patients. The aim of the therapy is to decrease the pain, which usually occur because of microfractures and/or osteomalacia of bones. All symptomatic cases should receive treatment. Surgical interventions like corrective osteotomy, may be planned to be temporarily promote bone mineralization.

The conventional treatment is based on oral phosphate salts, usually given twice daily, and active vitamin D metabolites. The goal of management is to improve the symptoms, and not to normalize serum phosphate levels. Careful monitoring of plasma calcium, PTH, creatinine, and 24-h urinary calcium excretion is required. Over treatment usually leads to tertiary hyperparathyroidism, hypercalciuria with nephrocalcinosis and renal insufficiency, resulting from excess calcitriol. Tertiary hyperparathyroidism responds to adjunctive therapy by 1,25-dihydroxyvitamin D, even if surgery is the likely modality of management.

**References**


