Coexistence of vitamin-D deficiency rickets and iron deficiency anemia has been described in the past. Occasionally, a more serious hematological condition, such as myelofibrosis, can occur in association with vitamin-D deficiency. We present a toddler with rickets due to vitamin-D deficiency and associated myelofibrosis which responded to oral cholecalciferol therapy.

A 16-month-old girl presented with failure to thrive and progressive lower limb deformities noticed since she was 6 months old. She was born of a nonconsanguineous union at term, had a birth weight of 2.5 kg and was appropriately weaned. She had delayed motor development and on enquiry, there was no past history of seizures, diarrhea, polyuria, jaundice, or blood transfusions. She was a playful child who weighed 7 kg (−2.93 standard deviation [SD]) with a length of 67 cm (−4.25 SD). Pallor, umbilical hernia, and features of rickets along with anterolateral bowing of bilateral lower limbs and generalized hypotonia were observed. Additionally, a soft hepatomegaly of 2 cm (span, 10 cm) and a firm splenomegaly of 7 cm along the long axis was found; rest of the systemic examination was unremarkable. The possible differential diagnoses considered were rickets in an infant with a storage disorder. Characteristically, a bone marrow aspiration results in a dry tap with biopsy demonstrating hypocellular bone marrow with increased reticulin deposition. Although, in our child, a bone marrow biopsy could not be performed, the presence of vitamin-D deficiency, characteristic peripheral blood picture, normal reticulocyte index, and the rapid improvement of bicytopenia and organomegaly on oral cholecalciferol therapy favored the diagnosis of vitamin-D deficiency–associated myelofibrosis.

In the pediatric population, myelofibrosis commonly occurs secondary to neoplasms, such as leukemia and lymphoma, or to nonneoplastic conditions, such as vitamin-D deficiency, systemic lupus erythematosus, Fanconi’s anemia, and hyperparathyroidism; primary myelofibrosis is rare.
Children with myelofibrosis associated with vitamin-D deficiency typically present in infancy. Hyperparathyroidism resulting from vitamin-D deficiency is postulated to be responsible for the myelofibrosis. Increased parathormone levels lead to decreased erythropoietin receptors on erythroid progenitor cells and increased medullary fibrosis. The hepatosplenomegaly results from subsequent extramedullary hematopoiesis. Another possible pathophysiologic feature of vitamin-D deficiency induced myelofibrosis could involve reduced levels of active vitamin-D metabolites which are responsible for the maturation of the monocyte and megakaryocyte precursors leading to reduced degradation and increased deposition of collagen. Along with cellular differentiation, vitamin D also plays a role in cellular proliferation; children with vitamin D–induced myelofibrosis have lower concentrations of CD34+ cells in bone marrow but increased levels in peripheral blood and splenic aspirates both of which return to normal with treatment of rickets.

Previous published case reports mainly showed a partial or complete improvement of clinical, radiological, and hematologic parameters with vitamin-D therapy, irrespective of the regimen used. However, persistent pancytopenia and death due to septic shock have also been reported.

Vitamin-D supplementation up to 1 year of age is routinely recommended. Also, secondary myelofibrosis must be kept in mind for any child presenting with rickets and hepatosplenomegaly. A high index of suspicion can obviate the need for extensive workup and support a more conservative approach toward management.

Conflict of Interest
None declared.

References
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