Title
Nutritional Rickets Presenting as Chronic Episodic Extremity Pain in a 9-year-old with Autism

Permalink
https://escholarship.org/uc/item/6fz5j6qv

Journal
Clinical Practice and Cases in Emergency Medicine, 2(3)

Authors
Tripathi, Noopur
Shankar, Roopa K.
Baghdassarian, Aline

Publication Date
2018

DOI
10.5811/cpcem.2018.2.37206

Copyright Information
Copyright 2018 by the author(s). This work is made available under the terms of a Creative Commons Attribution License, available at https://creativecommons.org/licenses/by/4.0/

Peer reviewed
**Nutritional Rickets Presenting as Chronic Episodic Extremity Pain in a 9-year-old with Autism**

Noopur Tripathi, BA*
Roopa Kanakatti Shankar, MBBS, MS†
Aline Baghdassarian, MD, MPH‡

*Virginia Commonwealth University School of Medicine, Richmond, Virginia
†Children's Hospital of Richmond at VCU, Department of Pediatrics, Division of Pediatric Endocrinology, Richmond, Virginia
‡Virginia Commonwealth University School of Medicine, Department of Emergency Medicine, Division of Pediatric Emergency Medicine, Richmond, Virginia

Section Editor: Rick A. McPheeters, DO
Submission history: Submitted December 6, 2017; Revision received February 15, 2018; Accepted February 27, 2018
Electronically published July 16, 2018
Full text available through open access at http://escholarship.org/uc/uciem_cpcem
DOI: 10.5811/cpcem.2018.2.37206

Rickets due to vitamin D deficiency, typically presenting as bowed legs in toddlers, is uncommon in the modern era. We describe the case of a nine-year-old girl with autism and developmental delay who was evaluated for chronic intermittent extremity pain for more than one year prior to referral to the emergency department for hypocalcemia and increased alkaline phosphatase, which eventually led to the diagnosis of rickets confirmed by radiographic and laboratory findings. This report highlights the importance of the patient's history of developmental delay and autism in the evaluation and approach to limb pain, and discusses the appropriate diagnostic approach. [Clin Pract Cases Emerg Med. 2018;2(3):251–254.]

INTRODUCTION

Nutritional rickets is a metabolic bone disease marked by failure of bone mineralization and architectural disruption at the growth plate. Calcipenic rickets is due to a deficiency in calcium; it can occur either due to inadequate intake or metabolism of vitamin D, or inadequate intake or absorption of calcium in the setting of normal vitamin D levels. This commonly presents at an early age with delayed closure of the fontanelle, frontoparietal bossing of the skull, bowing of the legs, widening of the wrists, and craniotabes. Rickets can also cause developmental delay, delayed achievement of motor milestones, hypocalcemic seizures in the first year of life, and neurologic as well as cardiac pathology. While essentially eradicated in the developed world due to vitamin D supplementation of newborns, there are still reports of nutritional rickets worldwide. This case highlights the importance of considering rickets in the differential diagnosis, especially if there is history of developmental delay and autism.

CASE REPORT

A nine-year-old Hispanic female with a past medical history of autism and global developmental delay presented to our emergency department (ED) complaining of a one-year history of pain in her extremities. The pain initially started in the right leg causing her to limp, trip, and fall. She was evaluated by her primary care physician and referred to a physical medicine and rehabilitation clinic that prescribed supramalleolar/ankle foot orthosis (SMAFO). The leg pain resolved, but she developed episodic pain in her bilateral upper extremities a month later. Initially manifesting as pain in her left arm, it was managed with nonsteroidal anti-inflammatory medications; then as this resolved she developed pain in her right arm. The episodic chronic pain in her extremities prompted laboratory evaluation and eventually her referral to the ED because of an elevated alkaline phosphatase (1,847 international units/liter [L]) and low serum calcium (6.4 milligrams [mg]/ deci-liter [dL]). Her past medical history was significant for autism and developmental delay. She did not have a family history of frequent fractures, bone pathology, or calcium problems.

On exam, she was non-verbal but followed commands and was comfortable with no acute distress. Her weight and height were less than the third percentile for age with minimal subcutaneous fat but normal body mass index (twelfth percentile). She had angular deformity and diffuse tenderness in the right and left arms and proximal forearms. She was able to bear weight but...
Nutritional Rickets Presenting as Chronic Episodic Extremity Pain in a 9-year-old with Autism

Tripathi et al.

had lower extremity pain and difficulty with ambulation. The rest of her physical exam was normal. We noted no spine tenderness, brachydactyly or other dysmorphic features.

Initial laboratory findings were remarkable for hypocalcemia and elevated alkaline phosphatase (Table). Radiographs of her extremities revealed multiple healed and healing fractures (Image 1), initially raising concern for non-accidental trauma. Further review of films with radiology revealed generalized bony demineralization, widened growth plates and metaphyseal fraying and flaring consistent with the diagnosis of rickets (Image 2). No vertebral compression fractures were noted, nor was rachitic rosary noted on chest radiographs. An elevated intact parathyroid hormone level (PTH) and extremely low serum 25-hydroxyvitamin D (25-OH vitamin D) concentration (Table) confirmed a diagnosis of severe hypocalcemic rickets due to vitamin D deficiency.

Endocrinology was consulted and elicited on history that she was a picky eater, only eating rice, fries and potato chips. She drank homemade green juices and smoothies but had limited dairy intake. On further review of systems, the family had not noticed any muscle spasms, seizures or twitching. Physical exam by the endocrinologist revealed widened wrists and ankles and rachitic rosary-prominent costochondral junctions of the ribs. She had no symptoms of neuromuscular irritability (negative Chvostek sign) despite ionized calcium of only 0.93 millimoles/L, highlighting the chronicity of the presentation.

The patient received calcium carbonate and calcitriol therapy followed by gradual supplementation of vitamin D3. She experienced significant improvement of pain and gait issues one month after initiation of calcitriol, calcium carbonate, and vitamin D3 supplementation along with orthotics. She was followed by endocrinology and was taken off calcitriol with normalization of calcium and serum 25-OH vitamin D concentrations in a month. One year after the diagnosis, calcium supplementation was also stopped with complete normalization of her calcium (10.1 mg/dL), PTH (38.6 picogram/mL), alkaline phosphatase (355 units/L) and vitamin D (48.7 nanogram/mL). All fractures were well healed except for a malaligned right humerus with no functional disability. She had improved dietary calcium intake with supplemental nutritional shakes and remained on 2,000 IU of vitamin D3 daily.

**DISCUSSION**

Rickets is a metabolic bone disease marked by failure of bone mineralization in the growth plates. The disease had been

---

**CPC-EM Capsule**

What do we already know about this clinical entity?
Nutritional rickets commonly presents at an early age with delayed closure of the fontanelle, frontoparietal bossing of the skull, bowing of the legs, widening of the wrists, and craniotabes.

What makes this presentation of disease reportable?
Late presentation without overt and classical signs and symptoms. The initial differential focused on non-accidental trauma (NAT) and did not entertain rickets.

What is the major learning point?
While NAT should remain high on the differential diagnosis, it’s important to remain mindful of the increased risk of nutritional rickets in patients with autism and developmental delay.

How might this improve emergency medicine practice?
With early diagnosis and management in the Emergency Department, Vitamin D deficiency causing rickets can be easily treated and serious complications avoided.
Bone pain was the first symptom on presentation in pediatric patients with acute lymphoblastic leukemia. Bone pain was also the initial complaint in 67.5% of patients with primary lymphoma. When children present with chronic bone pain to the ED, clinical suspicion is generally high for physical and sexual abuse, musculoskeletal abnormalities, hematologic processes, or malignancy. Children with disabilities are at higher risk for physical/emotional abuse than those without disabilities. Hence, we entertained this diagnosis in the pediatric ED while the workup was ongoing and the Child Protective Team was consulted.

With its decreasing incidence in the developed world, nutritional rickets due to vitamin D deficiency was not high on the initial differential. However, children with autism must be evaluated on different premises. In fact, children with autism and developmental delay are five times more likely to have feeding difficulty, or specific feeding behaviors, and are particularly susceptible to calcium deficiencies. Food selection in children with autism is often based on color and texture preferences, in addition to aversions to dairy products or foods rich in protein. Deficiencies have been observed in vitamins A, B6, B12, C, D, and K. It has also been observed that children with autism have significantly lower vitamin D levels compared to those without autism and that children with vitamin D-dependent rickets have autistic markers, which have resolved after supplementation with vitamin D.

CONCLUSION

Although the differential diagnosis of a child presenting with bone pain includes an extensive list of conditions ranging from benign growing pains to malignancies, in children with developmental delay, failure to thrive, or other musculoskeletal abnormalities, non-accidental trauma, and nutritional rickets should be high on the differential diagnosis. It is important for physicians to remain mindful of the increased risk of nutritional rickets in patients with autism and developmental delay. Vitamin D deficiency causing rickets can be easily treated and serious complications avoided with early diagnosis and management.
ACKNOWLEDGMENT

The authors would like to thank Dr. Joo Y. Cho, MD, Assistant Professor of Pediatric Radiology at Children's Hospital of Richmond at VCU, for the radiographs.

Documented patient informed consent and/or Institutional Review Board approval has been obtained and filed for publication of this case report.

Address for Correspondence: Aline Baghdassarian, MD, MPH, Virginia Commonwealth University School of Medicine, 1250 East Marshall Street, 2nd floor, Suite 500, PO Box 980401 Richmond, VA 23298-0401. Email: Aline.baghdassarian@vcuhealth.org

Conflicts of Interest: By the CPC-EM article submission agreement, all authors are required to disclose all affiliations, funding sources and financial or management relationships that could be perceived as potential sources of bias. The authors disclosed none.

Copyright: © 2018 Tripathi et al. This is an open access article distributed in accordance with the terms of the Creative Commons Attribution (CC BY 4.0) License. See: http://creativecommons.org/licenses/by/4.0/

REFERENCES

1. Singleton R, Lescher R, Gessner BD, et al. Rickets and vitamin D deficiency in Alaska native children. J Pediatr Endocrinol Metab. 2015;28(7-8):815-23.
2. Dietz, WH. Vitamin D Expert Panel Meeting October 11 - 12, 2001. Atlanta, Georgia. Final Report. 1-42.
3. Severe Malnutrition Among Young Children — Georgia, January 1997-June 1999. Available at: https://www.cdc.gov/mmwr/preview/mmwrhtml/mm5012a3.htm. Accessed on June 8, 2017.
4. Arvind R. 100 Clinical Cases in Pediatrics. 4th edition. New Delhi: Jaypee Brothers Medical Publishers Ltd, 2016.
5. Tietz NW. Textbook of Clinical Chemistry. Third edition, edited by Burtis and Ashwood. Philadelphia, PA, WB Saunders Company, Philadelphia, 1999.
6. Tiosano D, Hochberg Z. Hypophosphatemia: the common denominator of all rickets. J Bone Miner Metab. 2009;27(4):392-401.
7. Jonsson OG, Sartain P, Ducore JM, et al. Bone pain as an initial symptom of childhood acute lymphoblastic leukemia: association with nearly normal hematologic indexes. J Pediatr. 1990;117(2):233-7.
8. Horsman J, Thomas J. Primary bone lymphoma: a retrospective analysis. J Oncol. 2006. Available at: https://www.spandidos-publications.com/ijo/28/6/1571/download. Accessed on June 12, 2017.
9. Mandell DS, Walrath CM, Manteuffel B, et al. The prevalence and correlates of abuse among children with autism served in comprehensive community-based mental health settings. Child Abuse Negl. 2005;29(12):1359-72.
10. Horne BSV, Moffitt KB, Canfield MA, et al. Maltreatment of children under age 2 with specific birth defects: a population-based study. Pediatrics. 2015;136(6):e1504-12.
11. Sharp WG, Berry RC, McCracken C, et al. Feeding problems and nutrient intake in children with autism spectrum disorders: a meta-analysis and comprehensive review of the literature. J Autism Dev Disord. 2013;43(9):2159-73.
12. Shmaya Y, Ellat-Adar S, Leitner Y, et al. Nutritional deficiencies and overweight prevalence among children with autism spectrum disorder. Res Dev Disabil. 2015;38:1-6.
13. Bener A, Khattab AO, Al-Dabbagh MM. Is high prevalence of Vitamin D deficiency evidence for autism disorder?: In a highly endogamous population. J Pediatr Neurosci. 2014;9(3):227-33.
14. Roberts AL, Koenen KC, Lyall K, et al. Association of autistic traits in adulthood with childhood abuse, interpersonal victimization, and posttraumatic stress. Child Abuse Negl. 2015;45:135-42.
15. McCarville MB. The child with bone pain:malignancies and mimickers. Cancer Imaging. 2009;(Special issue A):S115-21.