Calcium Tartrate Tetrahydrate, Case Report of a Novel Human Kidney Stone

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Abstract

Background: Calcium tartrate tetrahydrate has been reported as the main mineral in urinary stones in rats that have significant tartrate in their diet, but in humans, there has been only one mention of calcium tartrate stones in the form of bladder stone, and that case was in Africa.

Case Presentation: Patient is a 34-year-old Caucasian male who presented with typical symptoms of nephrolithiasis. CT abd/pelvis (renal stone protocol) revealed a 2 cm nonobstructing stone of the right renal pelvis. Patient underwent an uncomplicated right percutaneous nephrolithotomy and was noted to be stone free after surgery. Stone analysis was difficult with regard to determining composition, but was finally identified as calcium tartrate tetrahydrate.

Conclusion: This was an unusual case, as this is the first recorded case of a calcium tartrate tetrahydrate outside of Africa. This type of stone had only been mainly described in rat models with *dl*-bitartrate in their diet. Our patient was an otherwise healthy, relatively muscular individual with no obvious source for this stone other than a vitamin and amino acid supplement that he takes regularly that contains L-carnitine (as tartrate) and choline (as bitartrate and citrate). The prevalence of this stone type is presently unknown, as stone analysis laboratories have not had the ability to recognize it. Although a connection between the supplement and stone formation is conjecture at this time, we believe this necessitates further investigation.

Keywords: urolithiasis, calcium tartrate tetrahydrate, dietary supplement

Introduction and Background

UROLITHIASIS IS A significant clinical issue in both adults and children and its prevalence has been increasing in the United States and across the globe over the last few decades. Overall prevalence varies between studies, but may range from 7% to 13%¹ in North America to 15% or more.² The most common type of urinary stones are made of calcium, specifically calcium oxalate,³ and this is particularly true in the younger age groups. The increased prevalence of stones is certainly multifactorial and may be associated with some lifestyle choices. These choices include poor dietary habits, such as increased consumption of animal protein (and decreased fruits and vegetables), and poor fluid intake that often consists of high sugar beverages.^{1,2,4} Certain supple-

ments and vitamins may also play a role in increasing the rate of kidney stones.

In this study, we present a case of urolithiasis that revealed a stone composed of calcium tartrate tetrahydrate, which could have some association with the use of drink supplements containing tartrate. This composition of stone had previously been described mainly in rat models. An association of urolithiasis and renal failure with a purified diet consisting of *dl*-choline bitartrate was noted in rats by Klurfeld⁵ and Newland et al.⁶ The composition of the stones in these rats was not known until Le Bail et al.⁷ did the crystal determination, confirming the infrared (IR) spectrum published by Newland et al. and identifying the compound as racemic calcium tartrate tetrahydrate, form II.

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Presentation of Case

The patient is a 34-year-old male who initially presented to our clinic for evaluation of recurrent passage of kidney stones. He was initially seen in July 2017 and described the passage of three stones that he was aware of, in the 2 to 3 months leading up to his presentation. During his initial encounter, he described typical symptoms of urolithiasis with primarily right-sided flank pain and intermittent episodes of gross hematuria. Baseline labs were relatively unremarkable (white blood cell [WBC] count 8.2, Hb 12.0 and Cr 1.13). Urinalysis demonstrated 2+ blood, 1+ leukocyte esterase, negative nitrites, and 3 to 9 red blood cells and WBCs. Urine culture was negative. Physical examination on presentation was unremarkable.

CT abdomen/pelvis without contrast (renal calculus protocol) was done and demonstrated a 2 cm nonobstructing stone in the right renal pelvis (Fig. 1). The average Hounsfield unit value was noted to be 825. The patient was scheduled to undergo right percutaneous nephrolithotomy (PCNL) within 2 weeks of presentation.

FIG. 1. Preoperative CT abdomen/pelvis (renal stone protocol).

Surgery

The PCNL procedure was uneventful. A lower pole access was established through biplanar fluoroscopy, access was established on the first try, and the tract was then dilated to 30F using the single step balloon dilation system. Lithotripsy was then achieved using an ultrasonic lithotripter. The stone was noted to be hard, but broke without difficulty. A 10F Cope loop nephrostomy tube was left in the upper pole as a nephrostomy tube and a 5F ureteral catheter was left down the ureter to help secure the access. The patient underwent a repeat CT without contrast (renal calculus protocol) the next morning. CT demonstrated the nephrostomy tube in appropriate position and no evidence of residual stone. Hospital course was uncomplicated, right antegrade nephrostogram was done with the contrast seen to be freely flowing down the ureter and the bladder, the nephrostomy tube was removed per protocol, and the patient was discharged on postoperative day 1.

Postoperative Follow-Up

UroRisk profile and a future renal ultrasonography (US) were ordered. The renal US was done at 3 weeks was unremarkable,

FIG. 2. Photo of stone specimen on mm paper (top), with micro-CT image slice (bottom). Micro-CT slice includes standard specimen composed of pure COM (thick arrow), and apparent CaOx inclusions in the specimen are indicated with thin arrow. Arrowheads indicate bulk of specimen, which was verified by infrared spectroscopy to be calcium tartrate. CaOx, calcium oxalate; COM, calcium oxalate monohydrate.

FIG. 3. FT-IR spectrum from the stone specimen. FT-IR, Fourier transform infrared.

without evidence of obstruction or shadowing to suggest the possible recurrence of stone. Patient was stone free and asymptomatic 10 weeks postoperatively.

Stone Analysis

The initial stone analysis reported the stone was a 100% metabolite, composed strictly of Ciprofloxacin, but there was no reason to believe that this was an accurate analysis. The patient was briefly placed on Levofloxacin 2 days before and 5 days after surgery as prophylaxis, but he had no previous history of Ciprofloxacin use. This unusual stone composition resulted in sending the specimen to the stone analysis laboratory at Indiana University, where micro-CT imaging revealed a majority stone material with X-ray attenuation significantly less compared with calcium oxalate (Fig. 2). The IR spectrum of the specimen did not match any known spectrum, and a consult was sought with the laboratory at Tenon Hospital in Paris. That laboratory identified the spectrum as being from calcium tartrate tetrahydrate (Fig. 3).

History/Exposure

After the finding of calcium tartrate in the stone, a thorough history was obtained. The patient reported no previous history of stones before the events described, no family history of stone formation, and no history of urinary tract infection. With regard to stone formation, he admitted to substantial intake of added salt in his daily diet. The patient worked as a contractor, but only with nonlead-based paints, and there was no apparent exposure to heavy metals or agricultural products. The patient did report drinking “large quantities ...4 or more scoops” of a vitamin and amino acid supplement energy drink/powder every day. Examination of the active ingredients of the product showed the presence of L-carnitine (as tartrate) and choline (as bitartrate and citrate). Each serving (one scoop) contained 10 mg of L-carnitine and

500 mg choline, and, as such, our patient was drinking 40+ mg and 2000+ mg per day, respectively, with thus an apparent intake of tartrate in the range of grams per day. Currently, there is no FDA-established daily value for intake of any of these substances.

UroRisk profile (24-hour urine study) was largely unrevealing, and the patient had an appropriate total volume of urine, 2.97 L, and a pH of 6.5. Specifically, there was no evidence of hyperuricemia (555 mg/day), hypocitraturia (484 mg/day), hyperoxaluria (36 mg/day), or hypercalciuria (218 mg/day). Only a mild elevation of urine sodium (216 mEq/day) was present.

Discussion

Calcium tartrate has been reported previously in only a single person, with data contained only in the tables of an article describing the composition of stones from the West African nation of Burkina Faso.⁸ There are no data in that article on the single male who had a calcium tartrate stone. Before this, calcium tartrate stones had been demonstrated only in rats, in association with tartrate in the diet.^{5–7} The IR spectrum found for the patient in our study (Fig. 3) is strikingly similar to that reported by Newland et al. in their rat study.⁶ The composition of the stone in rats is conclusively calcium tartrate tetrahydrate,⁷ which fits with the presence of tartrate in the diets that caused such stones.

Our patient was a relatively muscular and fit younger male with no significant comorbidities, no significant family history, no known metabolic disorders, and no medications that would be associated with stone formation. As mentioned in his history, he also had no significant chemical exposures, other than the dietary supplement that he took. This supplement contains L-carnitine (as tartrate) and choline (as bitartrate and citrate). Although this is conjecture at this point, as calcium readily binds tartrate, we believe it is reasonable that our patient was ingesting the offending agent, and it was the tartrate in his diet that led to his stones.

Conclusion

We report a 34-year-old male with kidney stones composed mainly of calcium tartrate. The patient reported consuming a large amount of tartrate as part of a dietary supplement, which thus could be the cause of his stones. We include the IR spectrum of this patient's stone specimen to facilitate the identification of this unique form of stone in other laboratories.

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Disclosure Statement

No competing financial interests exist.

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Abbreviations Used

CT = computed tomography
IR = infrared
PCNL = percutaneous nephrolithotomy
US = ultrasonography
WBC = white blood cell

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