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CLINICAL VIGNETTE

A Case of Congenital Benign Renal Glycosuria Likely Caused by Genetic SGLT2 Mutation

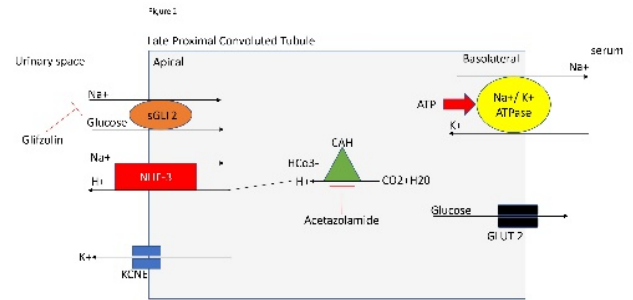
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A 51 year old female with chronic yeast infections and multiple urinary tract infections was referred to nephrology for glycosuria. Her fasting blood sugars have been slowly increasing towards 100mg/dL. She has prehypertension and history of paroxysmal supraventricular tachycardia, well controlled on beta-blockers. She had been evaluated in the past because both her mother and sister had renal glycosuria, and was advised it was benign without need for further follow up. Serum BMP, phosphorus, and urinary pH were all within normal limits. Her fasting and non-fasting blood sugars were between 90 and 120 mg/dL and never exceeded the tubular maximum of 180 mg/dL expected before glycosuria manifests. A random urine glucose level was 160 mg/dL, which was greater than the serum glucose level.

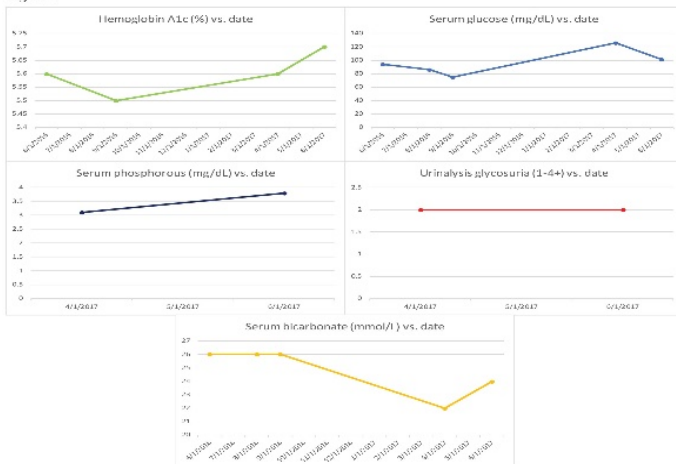
She had multiple urinary tract infections consistent with the presentation usually seen with patients on gliflozins, but without acute kidney injury or chronic kidney disease, as is sometimes a risk in patients on SGLT-2 antagonists. Despite the glycosuria, the patient’s prediabetes has worsened. Figure 1 presents a graphical trend of glycosuria and electrolytes. Figure 2 depicts the mechanism of SGLT 2 function in the renal proximal convoluted tubule.



Diabetes management is experiencing a wave of innovation with the addition of the gliflozin class of anti-hyperglycemic agents. These drugs are currently used in conjunction with metformin, thiazolidinediones, gliptins, and sulfonylureas to help control blood sugar without the need for injectable agents (like insulin and incretin mimetics).^{1,2} Given the increased microvascular³ and cardiovascular risks with diabetes⁴, the decrease in cardiovascular events among patients treated with gliflozins is encouraging.⁵⁻⁸ The new agents seen are not without risks. These include lactic acidosis, urinary tract infections, and acute kidney and chronic kidney disease. These risks were recently reiterated by the Food and Drug Administration (FDA).⁹

The gliflozins are antagonists of SGLT2 (sodium glucose linked transporter). These transporters are a family of channels that couple sodium movement across its chemical gradient to power glucose symport into cellular cytoplasm against its concentration gradient. They are expressed in the latter part of the proximal convoluted tubules (PCT) where they are involved in glucose reabsorption from filtered plasma. Gliflozins inhibit this target to increase glucose excretion and help lower serum glucose levels in diabetics. The gliflozins produce isolated glycosuria, whereas other proximal tubular dysfunction syndromes cause aminoaciduria, bicarbonaturia, and phosphaturia. The bicarbonaturia noted with PCT dysfunction is due to proximal renal tubular acidosis (type II). Fanconi’s syndrome usually results in the former acidosis as well as aminoaciduria and concomitant phosphaturia with hypophosphatemia due to global PCT dysfunction. Conditions like multiple myeloma and medications such as topiramate,

Figure 1



acetazolamide, and tenofivir can also cause proximal convoluted tubular dysfunction.¹⁰ This is in opposition to the urinary profile of glifzolins, which cause glycosuria alone. Table 1 lists medication induced, genetic, and acquired causes of proximal convoluted tubular dysfunction.

Table 1: Causes of Proximal convoluted tubule dysfunction	
Genetic	pattern
hereditary cystinosis	Fanconi's
tyrosinemia	Fanconi's
galactosemia	Fanconi's
glycogen storage diseases	Fanconi's
hereditary fructose intolerance	Fanconi's
Dent's disease	Fanconi's
Medications	pattern
acetazolamide	isolated type II RTA
topiramate	isolated type II RTA
tenofivir	varies, isolated glycosuria up to full fanconi's syndrome
Glifzolin class	isolated glycosuria
Expired tetracyclines	Fanconi's
Acquired disease	pattern
Multiple myeloma/Monoclonal gammopathy	RTA, proteinuria
Lead poisoning	Fanconi's
Sjogren's syndrome	Fanconi's
Pleuritis and other auto immune disease	Fanconi's

Conclusions

This patient has isolated glycosuria without hypophosphatemia, acidosis (especially a renal tubular acidosis of proximal type), or Fanconi's syndrome that is generally seen in global proximal convoluted tubular dysfunction. Glycosuria may be protective given her comorbid pre-hypertension and prediabetes. It is important to distinguish benign renal glycosuria from a medication effect, Fanconi's syndrome, and proximal renal tubular acidosis due to myeloma. The patient is considering genetic testing to define if SGLT2 or another closely related protein is the causative agent of her glycosuria, though her clinical course will be quite benign with a favorable long term renal prognosis.

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Figure legends

Figure 1 - Graph of RUA measurement of glycosuria as measured on routine urinalysis vs. date, serum glucose levels in mg/dL vs. date, serum Hco3 mmol/L vs. date, serum phosphorous mg/dL vs. date, and hemoglobin a1c (%) vs. date

Figure 2 - Schematic of proximal convoluted tubule function in its three segments ATP=adenosine triphosphate, CAH=Carbonic anhydrase inhibitor, CO2=carbon dioxide, GLUT2= glucose channel basolateral membrane, H2O=water, H+=proton, Hco3=bicarbonate, K+=potassium, KCNE=voltage gated potassium channel, Na+=sodium, NHE-3= sodium proton antiporter , SGLT2= sodium glucose linked transporter 2.

Table 1 - Causes of genetic, acquired, and medication induced proximal convoluted tubular dysfunction. See attached figure legend.

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