

# Introduction

McKittrick Wheelock Syndrome (MWS) is a rare disorder characterized by distal colorectal tumors, most commonly benign secretory villous adenoma leading to secretory diarrhea with electrolyte depletion syndrome. Patients present with volume depletion, severe electrolyte derangement, specifically hyponatremia and hypokalemia, along with acute kidney injury (AKI). We present a rare case of an elderly woman with severe electrolyte derangement in the setting of MWS.

# **Case Description**

71-year-old woman with history of 4 months of watery diarrhea, fatigue, and anorexia presented after a syncopal episode. Labs notable for Na 114 mEq/L, K 2.2 mEq/L, WBC 23.5K, and Cr 2.91 mg/dL. Stool electrolytes resulted in stool osmotic gap 48 mOsm/kg consistent with secretory diarrhea. Of note, patient was hospitalized 3 times in the past 2 months for hyponatremia, hypokalemia, and AKI requiring temporary dialysis secondary to profuse diarrhea.

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# A Rare Case of McKittrick Wheelock Syndrome

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A colonoscopy revealed a large rectal polypoid lesion with pathology consistent

Villous adenomas, normally a benign with a tubulovillous adenoma (Figure A). condition, can present with a life-She had aggressive electrolyte and fluid threatening electrolyte derangements and repletion with a robot volume depletion which makes the ability assisted abdominoperineal resection. to diagnose and adequately treat MWS Subsequently, noted to have resolution of critical. Patients typically have multiple her symptoms and complete electrolyte admissions with watery or mucous correction upon follow up (Table 1). diarrhea, nausea, and vomiting. Labs significant for hyponatremia, hypokalemia, AKI, and leukocytosis. The tumors are large and often past the splenic flexure and low in the rectum, Table 1. Electrolytes: Comparison of initial admission therefore flexible sigmoidoscopy can be vs. post-surgical resection reliably used rather than colonoscopy, which often delays diagnosis due to patients' inability to prep. Treatment includes aggressive fluid and electrolyte repletion until tumor can be surgically resected. Few case reports suggest using indomethacin or octreotide as a bridge to surgery or as medical management for patients who are not surgical candidates. However, patients who are managed medically have a mortality rate up to ~61-100%. Surgical management definitively resolves symptoms, although minimally invasive options are being explored.

Electrolytes	Initial Admission	Post-surgical resection
Na+	114	137
K+	2.2	4.8
CI-	79	106
Creatinine	2.91	1.06

### Discussion

#### References

1. Challis, Benjamin G et al. "The McKittrick-Wheelock syndrome: a rare cause of curable diabetes." Endocrinology, diabetes & metabolism case reportsvol. 2016 (2016): 160013.

2. Mois, Emil Ioan et al. "McKittrick-Wheelock syndrome: a rare case report of acute renal failure." Clujul medical (1957) vol. 89,2 (2016): 301-3.

3. Orchard, M R et al. "A systematic review of McKittrick-Wheelock syndrome." Annals of the Royal College of Surgeons of England, vol. 100,8 1-7. 16 Oct. 2018



A high index of suspicion and a systematic approach is critical to diagnose and provide life-saving treatment for MWS patients.

Figure A. Rectal villous adenoma.

