New onset hyperglycemia attributed to renal cell carcinoma

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Summary
A 61-year-old male was admitted from the outpatient setting for treatment of severe hyperglycemia. Five months earlier, his hemoglobin A1c had been 5 mmol/mol. At presentation, hemoglobin A1c was 11.3 mmol/mol and he required insulin therapy at discharge. Later magnetic resonance imaging (MRI) identified bilateral renal masses, previously seen on ultrasound during workup for chronic kidney disease, as being suspicious for renal cell carcinoma (RCC). He underwent partial nephrectomy and cryoablation with pathology showing papillary type RCC. Hyperglycemia resolved after resection and insulin therapy was discontinued, requiring only an oral hypoglycemic. Hyperglycemia as a paraneoplastic syndrome related to RCC is rare. The cause of this acute hyperglycemia is not understood, though previously suggested mechanisms include ectopic glucagon production, autoimmune causes and interleukin-6 (IL-6) mediated pathways. Severe, new-onset hyperglycemia in the absence of common causes and with a renal mass on imaging may represent an uncommon paraneoplastic syndrome secondary to RCC.

Keywords: Hyperglycemia, malignancy, paraneoplastic syndrome, renal cell carcinoma

1. Introduction
Renal cell carcinoma (RCC) has an incidence of approximately 65,000 cases annually in the United States with approximately 15,000 deaths a year (1). Over time, mortality rates have been steadily decreasing along with decreasing tumor size at the time of initial finding, likely due to an increased rate of incidental detection on abdominal imaging (2). Currently, most RCC is diagnosed while the disease is still localized, with 65% of cases diagnosed while still confined to the kidney. While earlier stages of RCC have a 5-year survival of approximately 90%, stages III and IV have a 5-year survival rate of 59-70% and median survival of 1 year respectively. It is estimated that approximately 20% of patients with RCC will have manifestations of paraneoplastic syndromes and in a significant number of patients; this may be the presenting complaint leading to a diagnostic workup and detection of RCC (3). The most common paraneoplastic syndromes associated with RCC are anemia, erythrocytosis, thrombocytosis, fever, cachexia, AA amyloidosis, hepatic abnormalities (Stauffer's syndrome), and polymyalgia rheumatica. Only a handful of cases have reported hyperglycemia secondary to RCC paraneoplastic syndrome with one case reported in the past decade. Here we describe the case of a patient whose presentation with severe hyperglycemia without previous history of diabetes, contributes to the diagnosis of renal cell carcinoma.

2. Case Report
A 61-year-old male with a history of essential hypertension, hyperlipidemia, coronary artery disease with past stenting and chronic kidney disease (CKD) was sent for admission from his nephrologist's office in October of 2016 when found to have serum blood glucose of 667 mg/dL. Per hospital records, the patient had not had a hemoglobin A1c over 5.5 mmol/mol since the beginning of his records 8 years ago with no diabetes medication during that time. A month prior to admission, the patient was started on metformin 500 mg twice daily by another provider and he reported he had run out of medication 10 days ago. At presentation,
he complained of fatigue, polyuria, polydipsia, blurred vision and lightheadedness. Further review revealed the patient had a renal ultrasound a month before this admission while establishing care with a nephrologist for management of his CKD. At that time, bilateral renal masses included a $3.8 \times 2.9 \times 3.7$ cm hypoechoic, exophytic mass on the right kidney and 1.6 cm $1.3 \times 1.4$ cm hypoechoic solid mass on the left kidney. Suggestion for urologic consultation and further imaging to rule out malignancy had been made, though these had not yet been completed.

On physical exam, the patient was afebrile with heart rate of 89 beats per minute and blood pressure of 104/74 mmHg and saturating at 96% on room air. He was not in distress, oral mucosa was dry and he was alert, awake and oriented. Lab values were significant for sodium of 129 mEq/L, potassium 4.8 mEq/L, bicarbonate of 22 mEq/L, blood urea nitrogen of 36 mEq/L, serum creatinine of 2.3 mEq/L and serum glucose of 667 mg/dL. Calculated serum osmolality was 308 mOsm/kg, liver panel was normal, white blood cell count of $6.3 \times 10^3$ cells/µL and hemoglobin of $11.3 \times 10^6$ cells/µL. Serum pH was 7.39 with no detectible beta-hydroxybutyrate and hemoglobin A1c was 11.3 mmol/mol and otherwise ranging from 4.5-5.5 mmol/mol over the last eight years.

During admission, computed tomography (CT) abdomen displayed a right indeterminate $3.6 \times 3.7$ cm exophytic inferior cortical renal mass. Additionally, there was a $1.0 \times 0.7$ cm left mid-pole lesion. Upon urologic evaluation, the CT scan findings were thought to be low risk for malignancy and reimaging in 6 months was suggested. The patient was discharged on insulin, requiring up to 43 units per day, with instruction to follow up with an ultrasound of the abdomen. Repeat ultrasound six months later revealed an enlarging right renal mass measured as $4.2 \times 4.0$ cm in diameter and left to 1.3 cm and patient was scheduled for MRI to further characterize the mass. MRI re-demonstrated the right sided mass that had increased to 4.5 cm with diffuse post contrast enhancement (Figure 1) as well as left sided hypo-intense mass measuring 1.2 cm with post contrast enhancement (Figure 2) concerning for renal cell carcinoma. The patient underwent right-sided partial nephrectomy with pathology revealing papillary renal cell carcinoma, type 1, and grade 2. Immunohistochemical stains showed tumor cells positive for CK7 and vimentin and negative for CD117, consistent with a diagnosis of RCC (Figure 3). Left sided renal mass was treated with cryoablation and interventional radiologic (IR) embolization. The patient recovered well from his surgery and was discharged.

Four months after discharge, patient's hemoglobin A1c had trended down to 4.5 mmol/mol and he was switched from insulin to glipizide 2.5 mg. The patient was contacted after discharge and informed consent was obtained for publication of this study.

3. Discussion

In this case, a patient with no former history of diabetes and with a hemoglobin A1c of 5.5 mmol/mol just five months prior to this hospitalization presented with severe hyperglycemia. During this five-month
period, the patient was found to have new, bilateral renal masses. The acute elevation of hemoglobin A1c to 11.3 mmol/mol is an atypical presentation of new onset diabetes in the absence of steroid use, severe pancreatitis, pancreatic cancer or pancreatectomy. Insulin therapy was required for 6 months with up to 43 units of long acting insulin per day at peak usage. Insulin requirements did appear to fluctuate rather dramatically, eventually requiring only glipizide. Following nephrectomy, the patient was able to remain on low dose oral hypoglycemic therapy as his hemoglobin A1c fell to 4.5 mmol/mol. This presentation and resolution of hyperglycemia following right-sided partial nephrectomy with embolization and cryoablation of the left sided mass is strongly suggestive of RCC as cause of the hyperglycemia.

A review of the literature revealed 7 previously reported cases of RCC with hyperglycemia in the English literature that were available for review (4-10). Years of publication ranged from 1981 to 2016. Available patient demographics and case details are included below (Table 1). One of the seven patients presented in diabetic ketoacidosis (DKA) (5). In all cases, there was improvement in glycemic control following nephrectomy. Both clear cell type and papillary type of RCC have been documented as causing these hyperglycemic states, however most case reports have not noted the histologic type of RCC. Furthermore, there are not enough cases reported to determine strength of correlation between RCC type and this hyperglycemic syndrome.

While the mechanism of hyperglycemia as a paraneoplastic syndrome in RCC has not yet been elucidated, a few possible mechanisms have been presented. One case noted increased levels of glucagon, which trended down after nephrectomy and presents the possibility of an endocrine mechanism of hyperglycemia (6). Another case detected elevation of anti-glutamic acid dehydroxylase (GAD) and anti-islet cell antibodies, which also normalized following nephrectomy and suggested a possible insulin deprivation mechanism due to pancreatic dysfunction on the level of the beta-islet cell (5). However, a review of the few other cases is not consistent with these previously noted findings. For instance, in the case of Harada et al, anti-GAD was tested for but was negative (9). For Callewaert and colleagues, glucagon, growth hormone, insulin like growth factor, cortisol and adrenocorticotropic hormone levels were all reported as normal (4). It may be that multiple mechanisms account for these few cases of significant hyperglycemia. More recent theories point to the effects of IL-6, which plays a role in other RCC paraneoplastic syndromes as well. In a study comparing cancer vs non-cancer patients, there was significantly decreased uptake of glucose in cancer patients with elevated IL-6 levels compared to those without elevated IL-6 (11). In other studies, injection of recombinant human IL-6 increased plasma glucose levels in a dose-dependent manner (12) and decreased insulin secretion from beta-islet cells (13).

<table>
<thead>
<tr>
<th>Author</th>
<th>Year</th>
<th>Synopsis</th>
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<tbody>
<tr>
<td>Pavelic (6)</td>
<td>1981</td>
<td>59 yo F, no history given with left flank pain. Left kidney mass 25 × 15 × 8 cm. Detected high levels of serum glucagon and insulin, which trend down after surgery, fluctuating blood sugars.</td>
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<td>Palgon (8)</td>
<td>1986</td>
<td>67 yo F with no previous IDDM, presents with AMS and serum glucose 650 mg/dL and requires 50 U insulin per day. Eight cm diameter mass, no metastasis, clear cell type, glucagon and somatostatin stains negative. Right radical nephrectomy with no insulin required after resection.</td>
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<td>Jobe (10)</td>
<td>1993</td>
<td>66 yo M, with no previous IDDM, presents in DKA with serum glucose of 847 mg/dL and requires up to 80 U/day of insulin. Right renal mass 9 × 10 × 11.5 cm, no metastasis, and clear cell type. Right nephrectomy with no insulin required after removal. Discharged on 2.5 mg of glyburide.</td>
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<tr>
<td>Callewaert (4)</td>
<td>1999</td>
<td>35 yo F with previous IDDM requires 600 U/day of insulin at presentation. Bilateral multifocal papillary renal cell carcinoma, grade 2, with no metastasis. Bilateral radical nephrectomy and returns to premorbid insulin use of 80 U/day.</td>
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<td>Macaulay (7)</td>
<td>2002</td>
<td>69 yo M with previous non-IDDM (on gliclazide) with hyperglycemia and elevated alkaline phosphatase requiring insulin (amount not specified). Six cm, right, necrotic mass, grade 2 with renal vein involvement, no metastasis. Left radical nephrectomy with return to gliclazide after nephrectomy.</td>
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<tr>
<td>Elias (5)</td>
<td>2005</td>
<td>52 yo F with no previous IDDM admitted with DKA. Elevated anti-GAD and islet cell antibody, which normalized after nephrectomy. Negative stain for glucagon, growth hormone and insulin. Insulin amounts prior and following tumor removal not specified though noted as decreasing.</td>
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<td>Harada (9)</td>
<td>2016</td>
<td>68 yo F with no previous IDDM, presents with serum glucose 353 mg/dL and A1c of 11.7 mmol/mol, elevated alkaline phosphatase requires 43 U/day insulin. Type of RCC not specified, grade II. Right nephrectomy and required 750 mg metformin daily at 2 month follow up.</td>
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Also, Blay and his colleagues studied the role of IL-6 levels in multiple paraneoplastic syndromes in patients with metastatic RCC and found that 90% of these patients had elevated IL-6 levels (14). Previous works by his group also showed expression of IL-6 from RCC cells in vitro as well. However, if 90% of patients with RCC have elevated levels of IL-6, the question of why so few cases of severe, new-onset hyperglycemia exist in RCC patients remains.

Hyperglycemia as a paraneoplastic syndrome in RCC is rare. The few reported incidences often show acute and severe hyperglycemia, which resolves with nephrectomy, as seen in the present case. These cases collectively indicate RCC as the most plausible cause of hyperglycemia. Further study is needed to determine the mechanism of hyperglycemia in RCC. While it is obvious that every episode of new onset hyperglycemia does not warrant a workup for RCC, new onset of severe hyperglycemia with common causes excluded and the presence of other signs of RCC might indicate a paraneoplastic etiology.

References


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