Acquired cystic kidney disease: A benign yet potentially fatal condition

Vikrampal Bhatti, Abhilash Koratala, Ashutosh M. Shukla

ABSTRACT

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CASE REPORT

A 48-year-old African-American male with end stage renal disease (ESRD) on hemodialysis for 14 years developed acute abdominal pain during his outpatient dialysis session and was sent to the emergency room. The patient was found to be hypotensive with a blood pressure of 93/72 mmHg requiring fluid resuscitation, which later deteriorated further requiring vasopressors. He was found to have acute anemia with a hemoglobin level of 7.5 g/dL. Abdominal computed tomography (CT) scan revealed large hemoperitoneum with a right perinephric sentinel clot, bilateral multiple renal cysts and features suspicious of bleeding originating from the upper anterior pole of right kidney (Figure 1). The largest of the cysts measured ~3.2 cm in diameter. Renal angiogram showed active ongoing extravasation from multiple branches of the right renal artery (Figure 2). Selective right renal artery embolization was unsuccessful. An emergent laparotomy revealed a torn right renal capsule with subcapsular hemorrhage and a 1.4 cm ragged defect in the inferior portion of the kidney. He underwent right nephrectomy and pathology was negative for malignancy. It essentially showed end stage kidney with arteriolonephrosclerosis, tubular atrophy, interstitial fibrosis, thyroidization, cysts and chronic interstitial inflammation with associated intra-parenchymal and perirenal hemorrhage. Patient refused blood transfusion because of religious reasons and unfortunately died of hemorrhagic shock after the surgery.

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Received: 01 March 2017  
Accepted: 13 July 2017  
Published: 01 August 2017

DISCUSSION

Acquired cystic kidney disease (ACKD) in patients with advanced chronic kidney disease (CKD) and end
stage renal disease, unlike other forms of cystic renal diseases, is largely considered a benign pathology with no clear recommendations for follow-up care. Our case represents a rare but fatal complication of ACKD. Definition of ACKD usually requires three or more cysts in each kidney in a patient with CKD or end stage renal disease who does not have a hereditary cause of cystic disease such as autosomal-dominant polycystic kidney disease or tuberous sclerosis [1]. It is estimated that hemorrhagic cysts are seen in approximately 50% of the patients with ACKD and perinephric hematomas in about 13% of patients [2], which can generally be managed conservatively. Severe bleeding can be fatal and may require interventions such as transcatheter embolization or surgery. More studies are needed to determine the benefit of periodic screening in patients with ACKD to prevent potentially fatal complications.

Without clear guidelines for follow-up and majority of the studies on cystic renal diseases excluding patients with ACKD, the true incidence of complications in these patients is not known. We believe an observational cohort study on the lines of Bosniak classification, in patients with advanced CKD and end stage renal disease should be considered such that natural history of this condition, including potential complications e.g. malignancy [3] and bleeding can be better predicted.

CONCLUSION

Acquired cystic kidney disease (ACKD) is characterized by development of numerous cysts in both the kidneys in individuals without history of hereditary cystic disease. Hemorrhagic cysts can sometimes lead to severe bleeding that can be life-threatening. High index of suspicion is required for this condition when end stage renal disease patients present with abdominal pain and drop in hemoglobin. Until clear guidelines are established, it would be prudent to monitor ACKD patients with periodic renal imaging.

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Keywords: Acquired cystic kidney disease, Hemoperitoneum, Renal disease

How to cite this article

Bhatti V, Koratala A, Shukla AM. Acquired cystic kidney disease: A benign yet potentially fatal condition. Int J Case Rep Images 2017;8(8):561–563.

doi:10.5348/ijcri-201719-CL-10129

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Author Contributions

Vikrampal Bhatti – Substantial contributions to conception and design, Acquisition of data, Analysis and interpretation of data, Drafting the article, Revising it critically for important intellectual content, Final approval of the version to be published

Abhilash Koratala – Analysis and interpretation of data, Revising it critically for important intellectual content, Final approval of the version to be published

Ashutosh M. Shukla – Analysis and interpretation of data, Revising it critically for important intellectual content, Final approval of the version to be published

Guarantor

The corresponding author is the guarantor of submission.

Conflict of Interest

Authors declare no conflict of interest.

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REFERENCES

1. Levine E. Acquired cystic kidney disease. Radiol Clin North Am 1996 Sep;34(5):947–64.
2. Choyke PL. Acquired cystic kidney disease. Eur Radiol 2000;10(11):1716–21.
3. Bhatti V, Vangapalli A, Bhattacharya D, Koratala A. Renal cell carcinoma of the native kidney in a renal transplant recipient. Int J Case Rep Images 2017;8(2):165–7.
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