

Incidental Finding of Rare Mucinous Carcinoma of Renal Pelvis in Post Nephrectomy Specimen of Pyonephrotic Non-Functioning Kidney

Saurabh Kumar Negi ^{a++*}, Sandip Desai ^{a++},
Gaurav Faujdar ^{a++}, Ram Dayal Sahu ^{a#}
and Nachiket Vyas ^{at}

^a Department of Urology, SMS Medical College, Jaipur, Rajasthan, India.

Authors' contributions

This work was carried out in collaboration among all authors. All authors read and approved the final manuscript.

Article Information

Open Peer Review History:

This journal follows the Advanced Open Peer Review policy. Identity of the Reviewers, Editor(s) and additional Reviewers, bpeer review comments, different versions of the manuscript, comments of the editors, etc are available here: <https://www.sdiarticle5.com/review-history/109578>

Case Report

Received: 14/09/2023
Accepted: 18/11/2023
Published: 20/11/2023

ABSTRACT

Primary mucinous adenocarcinoma of renal pelvis is very rare and difficult to diagnose preoperatively due to lack of specific symptoms and radiological findings. We are reporting a case of 52-year old man diagnosed with pyonephrosis with renal stone who underwent percutaneous nephrostomy (PCN) initially later on open simple nephrectomy was diagnosed primary mucinous adenocarcinoma of the renal pelvis in histopathology. Postoperative serum CEA and CA19.9 levels were normal. By assessing a literature review we recommend that careful history taking, tumor markers and CT scans may improve the diagnostic accuracy.

⁺⁺Mch Urology Resident;

[#]Asst Professor Urology;

[†]Professor Urology;

^{*}Corresponding author: Email: negisaurabh375@gmail.com;

Keywords: Mucinous carcinoma; nephrectomy; radiological findings.

1. INTRODUCTION

“The transitional cell carcinoma of renal pelvis is a common subtype, which accounts 90% of cases” [1]. “Renal pelvis adenocarcinoma accounts for less than 1% cases which is subclassified as tubulovillous (71.5%), mucinous (21.5%) and papillary non-intestinal (7.0%)” [2,3]. “Primary mucinous adenocarcinoma of renal pelvis is rare and often discovered accidentally by nephrectomy” [4]. “First reported in 1960 and till date fewer than 100 cases have been reported” [5,6]. “It is related to chronic irritation such as stone, infection, inflammation and obstruction” [7]. “It is difficult to diagnose before surgery without characteristic symptoms or specific radiological features. Also, because of its rarity, no standard treatment protocols has been proposed. We are reporting a case of mucinous adenocarcinoma of renal pelvis presented with feature of calculus and pyonephrosis” [8].

2. CASE PRESENTATION

Fifty two (52) years old male patient presented to our hospital with Leftt flank pain and fever for 20 days. On examination vital stable, Hemoglobin/Total Leucocyte Count/Serum Creatinine were 9.3g/dl /13k/ μ l and 1.01lakh/ μ l, Ultrasonography abdomen- s/o pyonephrosis with multiple renal stone with perinehric collection. CT urography (Fig. 1)- LT kidney multiple calculus with grossly dilated pelvicalyceal system (PCS) with mild perinephric collection, no contrast excretion. RT kidney normal. Patient underwent percutaneous nephrostomy (PCN) and 500 ml purulent fluid was drained. PCN kept for 2 weeks blater on underwent DTPA scan bwchich showed non-functioning bsame kidney. We performed an open simple nephrectomy via flank approach (Fig. 2). He was diagnosed as renal pelvis primary mucinous adenocarcinoma in histopathology report (Fig. 3).

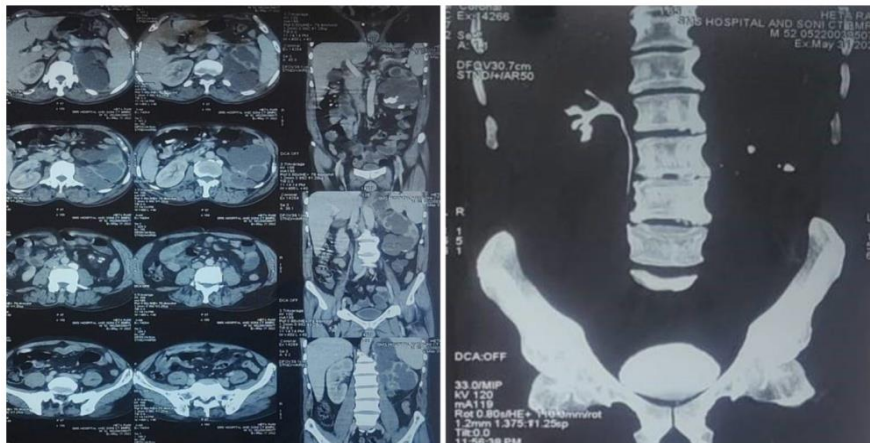


Fig. 1. CECT Abdomen & Urography -Left dilated pcs with calculus & perinephric collection with non excretion of contrast

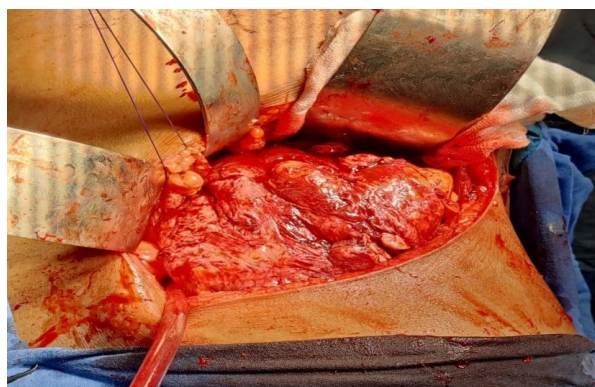


Fig. 2. Intra-operative image of nephrectomy

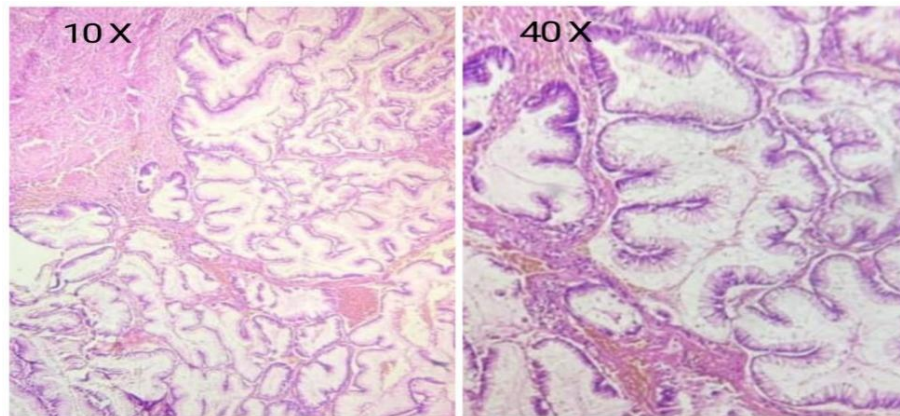


Fig. 3. HPE image -well differentiated mucinous adenocarcinoma of renal pelvis stage- pT3NxMX (AJCC2017)

2.1 Follow Up

Post operative patient underwent serum CEA and CA19.9 levels which were normal levels. Patient refused for chemotherapy and presently on regular follow up.

3. DISCUSSION

“Mucinous adenocarcinoma of renal pelvis is rare disease, discovered mainly in Asian population. Most adenocarcinomas are of high grade and invasive at presentation. No proper specific protocols has been proposed for renal mucinous adenocarcinoma. Early radical surgery appears to be the best option however the misdiagnosis with benign renal tumor, cyst or pyonephrosis may lead to delayed treatment and serious consequences” [9]. “CEA and CA19-9 may act as independent markers for prognosis and recurrence, however there can be normal in some cases” [10]. The images of CT and MRI are not specific and almost confirmed accidentally in pathological specimen. The standard of care is radical nephroureterectomy. The role of adjuvant chemotherapy and radiotherapy is controversial. These tumour are aggressive and has a poor prognosis. Early diagnosis is an important with preoperative CEA levels, CT scan and high level of suspicion help in diagnosis and treatment.

4. CONCLUSION

Primary mucinous adenocarcinoma of the renal pelvis is difficult to diagnose preoperatively. Thus, the patient usually have prolonged stone impaction with associated hydronephrosis or pyonephrosis. We should keep high suspicion. Early operation is the most effective therapy.

ETHICAL APPROVAL

As per international standards or university standards written ethical approval has been collected and preserved by the author(s).

CONSENT

As per international standards or university standards, patient(s) written consent has been collected and preserved by the author(s).

COMPETING INTERESTS

Authors have declared that no competing interests exist.

REFERENCES

1. Julian W, Kana A, Lee W. Primary mucinous adenocarcinoma of renal pelvis in solitary pelvic kidney. *Urology*. 1993; 41:292e4
2. Han DS, Yuk SM, Youn CS, et al. Primary mucinous cystadenocarcinoma of the renal pelvis misdiagnosed as ureteropelvic junction stenosis with renal pelvis stone: A case report and literature review. *World J Surg Oncol*. 2015;13:324.
3. Lai C, Teng XD. Primary enteric-type mucinous adenocarcinoma of the renal pelvis masquerading as cystic renal cell carcinoma: A case report and review of the literature. *Pathol Res Pract*. 2016;212:842-8.
4. Gupta P, Agarwal D, Shruti S, Chandra M. Mucinous adenocarcinoma of renal pelvis in a young male: A diagnostic challenge. *German Medical Science : GMS E-journal*. 2020;18:Doc11.

- DOI: 10.3205/000287
PMID: 33299389; PMCID: PMC7705116.
5. Hasebe M, Serizawa S, Chino S. On a case of papillary cystadenocarcinoma following malignant degeneration of a papillary adenoma in the kidney pelvis. *Yokohama Med Bull.* 1960;11:491-500.
 6. Shah VB, Amonkar GP, Deshpande JR, et al. Mucinous adenocarcinoma of the renal pelvis with pseudomyxoma peritonei. *Indian J Pathol Microbiol.* 2008;5.
 7. Joshi K, Jain K, Mathur S, Mehrotra G. Mucinous adenocarcinoma of the renal pelvis. *Postgrad Med J.* 1980;56: 442e4.
 8. Li H, Xie F, Zhao C, Yi Z, Chen J, Zu X. Primary mucinous adenocarcinoma of the renal pelvis misdiagnosed as calculous pyonephrosis: A case report and literature review. *Translational Andrology and Urology.* 2020 Apr;9(2):781.
 9. Yadav R, Kataria K, Balasundaram P, et al. Mucinous cystadenocarcinoma arising in an ectopic kidney simulating a retroperitoneal dermoid cyst: A rare tumour presenting as a diagnostic dilemma, *Malays. J. Pathol.* 2013;35:95–98.
 10. Raphael V, Sailo S, Bhuyan A, et al., Mucinous adenocarcinoma of the renal pelvis with adenocarcinoma in situ of the ureter, *Urol. Ann.* 2011;3:164–166.

© 2023 Negi et al.; This is an Open Access article distributed under the terms of the Creative Commons Attribution License (<http://creativecommons.org/licenses/by/4.0>), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

Peer-review history:
The peer review history for this paper can be accessed here:
<https://www.sdiarticle5.com/review-history/109578>