



Imaging Features of Urachal Cancer: A Case Report

Xiaoyan Chen^{1*}, Chunsong Kang¹ and Mingxia Zhang²

¹ Department of Ultrasound, Shanxi Academy of Medical Sciences, Shanxi DAYI Hospital, Taiyuan, China, ² Department of Clinical Laboratory Medicine, Shanxi Academy of Medical Sciences, Shanxi DAYI Hospital, Taiyuan, China

Urachal adenocarcinoma originates from the space of Retzius. It is a rare but aggressive neoplasm. Typical imaging findings of urachal cancer are difficult to find; this report provides an ultrasonographic (US) and computed tomographic (CT) description of the above. We present a case of a 45-year-old male patient presenting with painless hematuria of 1 week's duration and display the US and CT images. Imaging shows: (1) a solid, ill-defined, irregularly shaped mass invading the bladder wall located between the dome of the bladder and the abdominal wall, and which underwent significant enhancement; (2) hypo-echoic, heterogeneous echopattern with some calcifications and (3) patchy and short-line blood flow signals within the mass. Final histopathology confirmed urachal adenocarcinoma.

OPEN ACCESS

Edited by:

Cicero Matthew R. Habito, Massachusetts General Hospital, Harvard Medical School, United States

Reviewed by:

Yanwei Miao, Dalian Medical University, China Johanna Patricia A. Canal, University of the Philippines Manila, Philippines

*Correspondence:

Xiaoyan Chen xiaoguoguo119@163.com

Specialty section:

This article was submitted to Cancer Imaging and Image-directed Interventions, a section of the journal Frontiers in Oncology

> Received: 28 August 2019 Accepted: 04 November 2019 Published: 26 November 2019

Citation:

Chen X, Kang C and Zhang M (2019) Imaging Features of Urachal Cancer: A Case Report. Front. Oncol. 9:1274. doi: 10.3389/fonc.2019.01274

Keywords: urachus, adenocarcinoma, bladder, ultrasonography, computed tomography

INTRODUCTION

The urachus is a tubular structure extending from the dome of the bladder to the allantois in early embryonic development. After birth, it becomes a fibrous cord located in the loose connective tissue between the transverse fascia and the peritoneum (i.e., within the space of Retzius) (1-4). Failure of complete luminal obliteration has been described in up to one-third of adults and can rarely lead to various anomalies including cysts, fistulae, diverticula, or, rarely, malignant transformations (2, 5).

Based on the statistics of the last decade, the case reports of urachal cancer are still rare each year since the seminal studies by Begg in the 1930s (6). Urachal carcinoma accounts for \sim 0.1% of all malignant tumors and 0.35–0.7% of all bladder tumors (7). It is usually undiscovered for a long time because the early stages are often asymptomatic. It may be identified incidentally during an imaging examination performed for other reasons or when symptoms appear at advanced stages (4, 8–10). Imaging examinations, such as ultrasonography (US), computed tomography (CT), and magnetic resonance imaging (MRI) play an important role in the workup of urachal tumors (11).

CASE PRESENTATION

A 45-year-old man presented to our hospital with 1 week of painless hematuria with no obvious cause. He was previously healthy with no known diseases. Physical examination showed no obvious abnormalities.

Urinalysis was positive for red blood cells, 814×10^6 /L and white blood cells, 3 oun⁶/L. Tumor marker determination showed carcinoembryonic antigen (CEA) 2.8×10^{-3} ng/L (normal range: $0-5 \times 10^{-3}$ ng/L), and prostate specific antigen (PSA) 0.48×10^{-3} ng/L (normal range: $0-4 \times 10^{-3}$ ng/L).

US examination showed a solid mass in the midline of the lower abdomen, invading the antero-inferior bladder wall. It measured $6.3 \times 4.9 \times 3.4$ cm, with ill-defined, irregularly

1

shaped protrusions outside the bladder extending to the umbilicus. Internal echogenicity was heterogeneous, and patchy calcifications and a small amount of point-like and short-line blood flow were observed within the tumor. No enlarged lymph nodes were noted in the abdominal cavity or retroperitoneum. We diagnosed an umbilical tumor (**Figure 1**).

The diagnosis on contrast-enhanced abdominopelvic CT was also umbilical tumor. CT confirmed that there was an intraabdominal mass with calcification, invasion of the adjacent bladder dome and peritoneum, and which underwent significant enhancement (**Figure 2**).

The patient underwent surgery to remove the mass. The urachus, umbilicus, and part of the bladder dome 2.0 cm from the tumor were also resected. Intraoperative findings showed an \sim 5.0 × 5.5 × 5.0 cm mixed cystic and solid mass invading the dome of the bladder and adherent to the peritoneum gross pathology showed a 6.0 × 5.5 × 3.5 cm mass with a grayish-white cut surface, coated in a jelly-like substance. The tumor was friable. Histopathology of the tumor showed that part of the bladder mucosa was included in the tumor, without vascular and neurological invasion recidivism (**Figure 3A**). According to the Mayo staging system (3), the pathological diagnosis was urachal carcinoma (grade II mucinous adenocarcinoma)

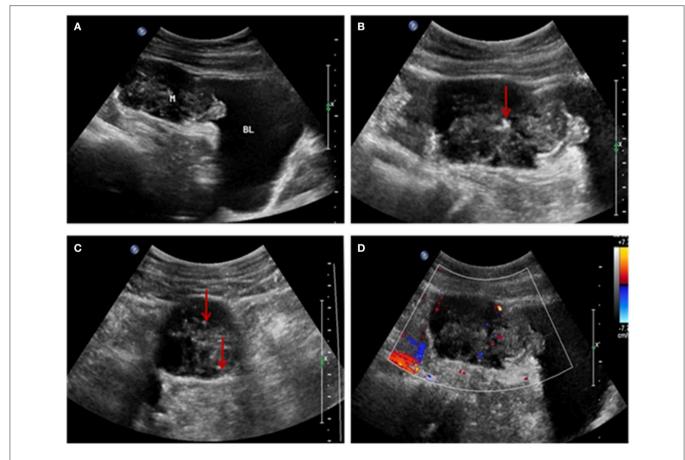
with calcification (**Figure 3B**). The patient received docetaxel combined with tegafur chemotherapy after surgery.

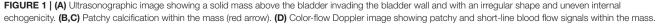
DISCUSSION

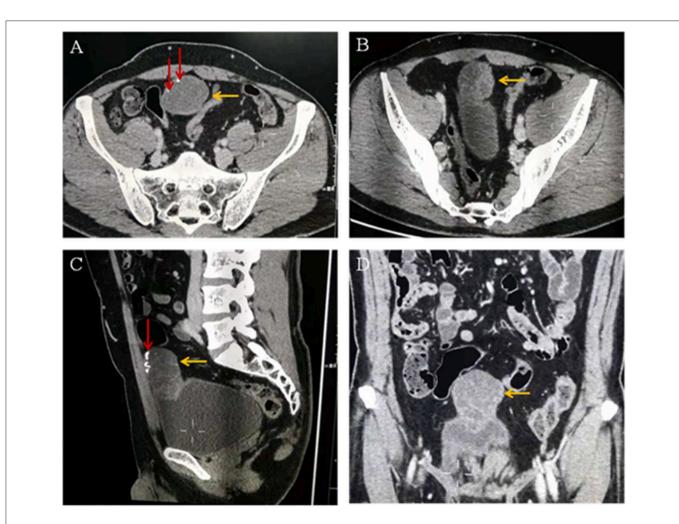
Urachal cancers can originate from any of the various layers of the urachus tract and arise mainly from epithelial cells; 94% of umbilical urachal cancers are adenocarcinomas, the diagnosis of our patient (6). The tumor invaded and actually eroded the bladder wall, causing painless hematuria, similar to a bladder cancer (12). Because of the non-specific clinical signs of this case, preoperative diagnosis depended on imaging.

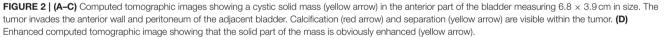
Urachal cancer usually appears as an intra-abdominal mass adjacent to the bladder dome, extending into the space of Retzius. It can be cystic, solid, or mixed (11). US and CT can provide a general impression of the lesion, including the location, size, and invasiveness of the mass (8, 9, 13).

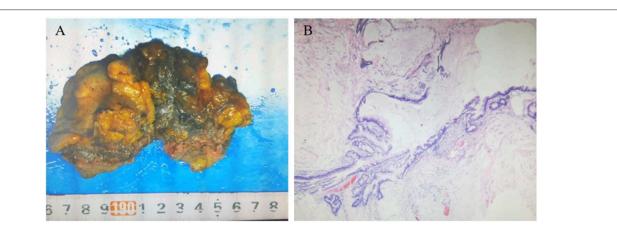
This report provides the characteristic US appearance of urachal cancer. The main features of this patient's ultrasonographic imaging findings were: (1) a solid mass extending between the dome of the bladder and the abdominal wall, with an irregular shape and bladder wall invasion; (2) a hypoechoic, heterogeneous echo pattern with a small amount

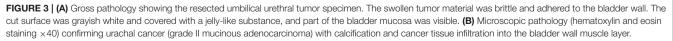












of calcification; and (3) patchy and short-line blood flow signals within the mass. These findings are consistent with previous reports (8, 9, 13) and the key findings of this case to further enrich the imaging features of this disease.

The US and CT examinations showed that the tumor infiltration reached the bladder muscle layer, without obvious lymph node involvement or spread to other organs, consistent with intraoperative findings and surgical pathology. Urachal tumors contain mucin, and calcifications are present in 70% of cases (1, 14). The presence of calcifications in a soft-tissue mass along the course of the abdominal midline is considered pathognomonic for the diagnosis of urachal cancer (8, 9).

CONCLUSION

We have described the typical imaging features of a case of urachal adenocarcinoma. Diagnosing urachal cancer is easier with imaging.

REFERENCES

- Thali-Schwab CM, Woodward PJ, Wagner BJ. Computed tomographic appearance of urachal adenocarcinomas: review of 25 cases. *Eur Radiol.* (2005) 15:79–84. doi: 10.1007/s00330-004-2408-z
- Upadhyay V, Kukkady A. Urachal remnants: an enigma. Eur J Pediatr Surg. (2003) 13:372–6. doi: 10.1055/s-2003-44725
- Molina JR, Quevedo JF, Furth AF, Richardson RL, Zincke H, Burch PA. Predictors of survival from urachal cancer: a Mayo Clinic study of 49 cases. *Cancer*. (2007) 110:2434–40. doi: 10.1002/cncr.23070
- Ashley RA, Inman BA, Sebo TJ, Leibovich BC, Blute ML, Kwon ED, et al. Urachal carcinoma: clinicopathologic features and long-term outcomes of an aggressive malignancy. *Cancer*. (2006) 107:712–20. doi: 10.1002/cncr.22060
- Schubert GE, Pavkovic MB, Bethke-Bedürftig BA. Tubular urachal remnants in adult bladders. J Urol. (1982) 127:40–2. doi: 10.1016/S0022-5347(17)53595-8
- Szarvas T, Módos O, Niedworok C, Reis H, Szendröi A, Szász MA, et al. Clinical, prognostic, 'and therapeutic aspects of urachal carcinoma—a comprehensive review with meta-analysis of 1,010 cases. Urol Oncol. (2016) 34:388–98. doi: 10.1016/j.urolonc.2016.04.012
- Gopalan A, Sharp DS, Fine SW, Tickoo SK, Herr HW, Reuter VE, et al. Urachal carcinoma: a clinicopathologic analysis of 24 cases with outcome correlation. *Am J Surg Pathol.* (2009) 33:659–68. doi: 10.1097/PAS.0b013e31819aa4ae
- 8. Koster IM, Cleyndert P, Giard RW. Best cases from the AFIP: urachal carcinoma. *Radiographics*. (2009) 29:939–42. doi: 10.1148/rg.293085152
- 9. Wong-You-Cheong JJ, Woodward PJ, Manning MA, Davis CJ. From the archives of the AFIP: inflammatory and nonneoplastic bladder

DATA AVAILABILITY STATEMENT

All datasets generated for this study are included in the article/Supplementary Material.

ETHICS STATEMENT

The studies involving human participants were reviewed and approved by appropriate institutional of Shanxi Academy of Medical Sciences and Shanxi DAYI Hospital. The patients/participants provided their written informed consent to participate in this study. Written informed consent was obtained from the individual(s) for the publication of any potentially identifiable images or data included in this article.

AUTHOR CONTRIBUTIONS

XC, CK, and MZ performed image acquisition and completed the manuscript.

masses—radiologic-pathologic correlation. *Radiographics*. (2006) 26:1847–68. doi: 10.1148/rg.266065126

- Dhillon J, Liang Y, Kamat AM, Siefker-Radtke A, Dinney CP, Czerniak B, et al. Urachal carcinoma: a pathologic and clinical study of 46 cases. *Hum Pathol.* (2015) 46:1808–14. doi: 10.1016/j.humpath.2015. 07.021
- Parada Villavicencio C, Adam SZ, Nikolaidis P, Yaghmai V, Miller FH. Imaging of the urachus: anomalies, complications, and mimics. *Radioggraphics*. (2016) 36:2049–63. doi: 10.1148/rg.2016160062
- Siefker-Radtke A. Urachal carcinoma: surgical and chemotherapeutic options. *Expert Rev Anticancer Ther.* (2006) 6:1715–21. doi: 10.1586/14737140.6.12.1715
- Chen D, Li Y, Yu Z, Su Z, Ni L, Gui Y, et al. Investigating urachal carcinoma for more than 15 years. Oncol Lett. (2014) 8:2279–83. doi: 10.3892/ol.2014.2502
- Yu JS, Kim KW, Lee HJ. Urachal remnant diseases: spectrum of CT and US findings. *Radiographics*. (2001) 21:451–61. doi: 10.1148/radiographics.21.2.g01mr02451

Conflict of Interest: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

Copyright © 2019 Chen, Kang and Zhang. This is an open-access article distributed under the terms of the Creative Commons Attribution License (CC BY). The use, distribution or reproduction in other forums is permitted, provided the original author(s) and the copyright owner(s) are credited and that the original publication in this journal is cited, in accordance with accepted academic practice. No use, distribution or reproduction is permitted which does not comply with these terms.