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Congenital primary obstructed megaureter presenting as inguinal hernia in an infant

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Abstract

A 4-month-old boy presented with left inguinal swelling, and the examination was notable for an inguinal hernia; ultrasonography and CT revealed that the hernia content was hugely dilated ureter of ureteral-inguinal hernia caused by primary obstructed megaureter. Imaging of pediatric inguinal hernia elucidates contents, etiology, and guide for proper surgery.

K E Y W O R D S

congenital, pediatrics, primary obstructed megaureter, ureteral-inguinal hernia

1 | INTRODUCTION

A 4-month-old male child presented to the hospital with his caregiver complaining of left-sided inguinal

swelling. Physical examination was notable for left inguinal hernia. Abdominal ultrasonography (US) and computed tomography (CT) showed that the left ureter was hugely dilated and was viewed inside left

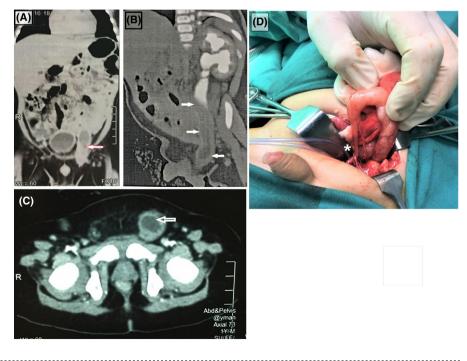


FIGURE 1 Abdominal computed tomography findings showing the hugely dilated ureter of congenital primary obstructed megaureter forming a left inguinal hernia in an infant (A, B, C, arrows). Intraoperative finding of the hugely dilated ureter with obstructed lower ureteric segment of congenital primary obstructed megaureter (D, asterisk)

This is an open access article under the terms of the Creative Commons Attribution-NonCommercial-NoDerivs License, which permits use and distribution in any medium, provided the original work is properly cited, the use is non-commercial and no modifications or adaptations are made. © 2022 The Authors. *Clinical Case Reports* published by John Wiley & Sons Ltd. inguinal canal with advanced hydroureteronephrosis (Figure 1A,B,C, arrows). At laparotomy through pubic incision, the ureter was found dilated and entrapped inside the internal inguinal ring. The ureter was delivered intra-abdominal, and the diagnosis of primary obstructed megaureter (POM) was confirmed (Figure 1D, asterisk). Surgical correction of POM was done, and the internal inguinal ring was closed. At a follow-up visit after 6-months, the child was doing well; US showed resolution of hydroureteronephrosis.

Infantile ureteral-inguinal hernia is extremely rare with 6 case reports in literature.¹ It is predominantly diagnosed during the repair of inguinal hernia; it is crucial to be recognized preoperatively in order to avoid accidental ureteral injury during hernia repair.^{1,2}

In children with inguinal hernia, the possibility of the ureter being the content may be considered and US followed by computerised tomography should be considered to elucidate primary cause, to plan proper surgery, and to avoid ureteral injury during hernia repair.

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CONFLICT OF INTEREST Nothing to declare.

AUTHOR CONTRIBUTIONS

MW managed the patient, contributed significantly to draft preparation, manuscript editing, and reviewed the final version.

ETHICAL APPROVAL

Written informed consent was obtained from the patient's caregiver for publication of this clinical image, this report was conducted in accordance with the declaration of Helsinki.

CONSENT

Written consent for publication was obtained from caregiver of the patient.

DATA AVAILABILITY STATEMENT

Data are available on request from the corresponding author.

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