



#### > No Disclosures



#### Neonatal urinary ascitis

Rare condition

Etiology: Obstructive uropathy (bladder outlet obstruction, neurogenic bladder, ureteric obstruction), latrogenic, spontaneous bladder perforation

 Clinical emergency, immediate resuscitation and management (potential life threatening, reversible & good prognosis)

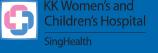


Aim – To share 2 cases of perinatal urinary ascites with unusual etiology and discuss management challenges



# Case 1 History

- 17 days old, male, from neighboring country presented with failure to pass urine after birth
- Past history:
  - Born at 35 weeks via LSCS, 1.9 kg
  - -Antenatal scan @ 32 weeks- distended bladder
  - -Post natal USG- full bladder, b/l severe hydronephrosis
  - Failed catheterisation x2, Suprapubic cystostomy drained urine x 2 weeks



#### History

- Exploratory laparotomy @ 2 weeks for suspected catheter migration- catheter in place, urine in retroperitoneal space
- Bladder closed with retroperitoneal drain, IDC in situ, post op abdominal distension, abdominal and scrotal wall edema, urine from drain and wound, worsening renal function, acidosis, and sepsis
- Transferred to Singapore



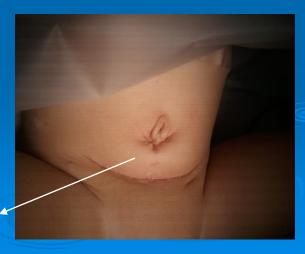
Imaging revealed ascitis, MCU showed bladder rupture in peritoneal cavity

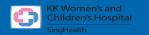
Exploration, repair of posterior bladder rent & mini vesicostomy done

Post operative recovery good, renal function stabilized



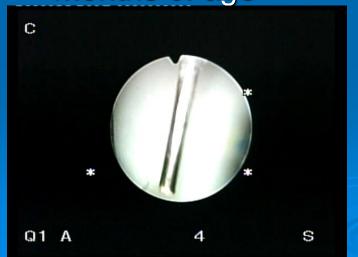
Urinary leak

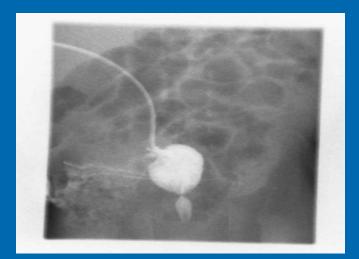




 4 weeks later cystogram done showed dilated posterior urethra with PUV

Fulguration done at 3 months of age

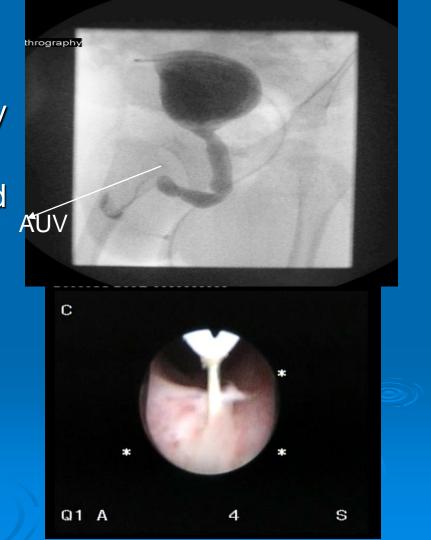


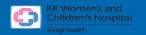






- Post fulguration, poor urinary stream on clamping vesicostomy
- Repeat cystogram through vesicostomy at 9 month, showed anterior urethral obstruction
- Cystoscopy & AUV fulgurated, vesicostomy was closed
- child is voiding well. Had high resting pressures, now on oxybutynin, renal functions normal.





#### Discussion

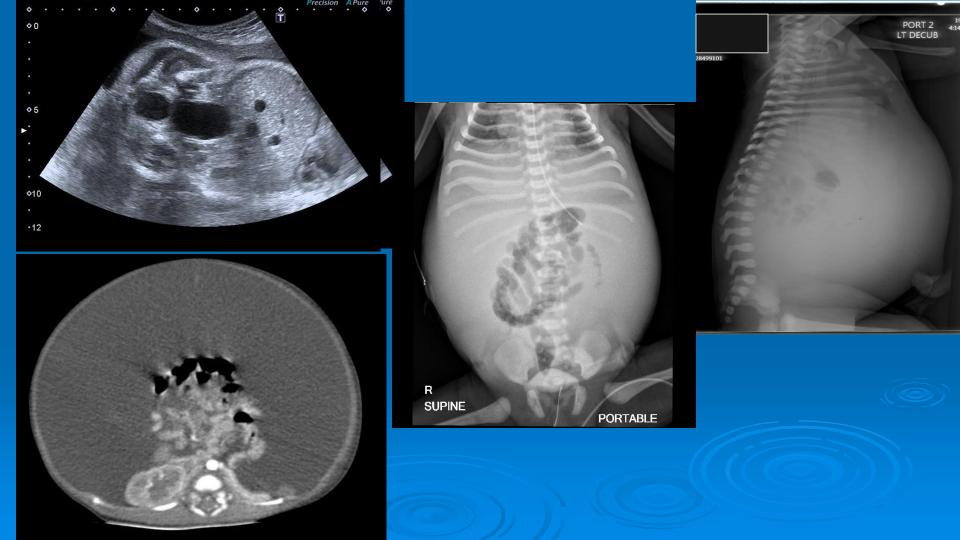
AUV & PUV association with urinary ascitis is rare ( European J of Ped Surg 2005,15:449-451, Kumar et al)

We missed the AUV during first time cystoscopy (radiologically not demonstrated),as we need to carefully look for the flap valve with a hook electrode.

Patency of distal urethra should be ensured before closing vesicostomy

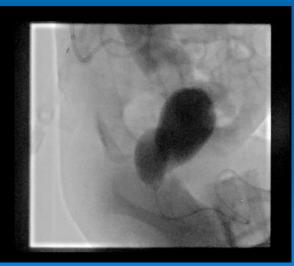
## Case 2 History

- A full term, male neonate with Klinfelter Syndrome presented with a large urinary ascites during perinatal period along with a large cyst next to the suspected bladder detected on antenatal scans.
- > He had renal agenesis on left side.
- Severe acidosis, Respiratory distress, poor renal function
- He was managed with Intubation, urgent peritoneal drainage to improve the ventilation and urethral catheterization at birth.

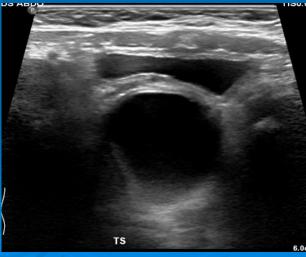


#### Management

- He needed two weeks of ventilatory support due to gross abdominal distension and three weeks of peritoneal fluid drainage.
- > MRI and MCU- bladder outlet obstructed by a large utricular cyst.







- The utricle cyst was not visualized in earlier post natal scans in the presence of ascites and urethral catheter but it remanifested later
- > The renal function also normalized gradually thereafter.
- He underwent a vesicostomy and is planned to be managed with excision of the cyst at a later date.

### Conclusions

- Neonatal urinary ascites is uncommon and usually presents as a clinical emergency.
- The diagnosis and treatment can be challenging if the cause is rare.
- The management of these cases should be planned carefully based on the anatomical and functional status of the urinary system.

#### References

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