Neonatal Gallstones

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Abstract
A term male infant was born by spontaneous vaginal delivery to a 23 year old gravida 2 para 2 mother. His weight, length, and head circumference were 2720g (8%), 47cm (6%) and 31.5cm (1%) respectively; the remainder of his physical examination was unremarkable. The pregnancy was complicated by intra-uterine growth retardation, gestational diabetes controlled with glyburide, and gestational thrombocytopenia. The mother had a past medical history of obesity, BMI of 32 (>95%), HPV infection, and atypical squamous cells of the cervix. A fetal ultrasound at 38 weeks gestational age showed echogenic foci consistent with cholelithiasis (Figure 1); a previous US at 37 weeks gestational age was normal. On routine laboratory testing the mother’s blood type was O positive and the infant’s was A+ (positive). His transcutaneous bilirubin was 4.3 mg/dL at 36 hours of life. Both mother and son were discharged after two days without problem.

Key words: spontaneous vaginal delivery; gestational diabetes; cholelithiasis; transcutaneous bilirubin; down syndrome

Introduction
A term male infant was born by spontaneous vaginal delivery to a 23 year old gravida 2 para 2 mother. His weight, length, and head circumference were 2720g (8%), 47cm (6%) and 31.5cm (1%) respectively; the remainder of his physical examination was unremarkable. The pregnancy was complicated by intra-uterine growth retardation, gestational diabetes controlled with glyburide, and gestational thrombocytopenia. The mother had a past medical history of obesity, BMI of 32 (>95%), HPV infection, and atypical squamous cells of the cervix. A fetal ultrasound at 38 weeks gestational age showed echogenic foci consistent with cholelithiasis (Figure 1); a previous US at 37 weeks gestational age was normal. On routine laboratory testing the mother’s blood type was O positive and the infant’s was A+ (positive). His transcutaneous bilirubin was 4.3 mg/dL at 36 hours of life. Both mother and son were discharged after two days without problem. Figure 1 shows, prenatal ultrasound of fetus at 38 weeks. Arrow indicating gallstones.
At seven days of age the infant was brought to the clinic for jaundice. A transcutaneous bilirubin measured 14.9 mg/dL; the total serum bilirubin was 15.6 mg/dL with a direct measurement of 0.2 mg/dL. He was diagnosed with breast milk jaundice and no intervention was indicated. At a one month follow-up his total serum bilirubin was 12.7 mg/dL with a direct measurement of 0.3 mg/dL. An abdominal US showed no evidence of cholelithiasis or sludge.

At five months of age he presented to the emergency department with vomiting. An abdominal US showed ‘mobile echogenic material consistent with biliary sludge’. (Figure 2) No stones were seen. He was diagnosed with ileus and discharged home. Figure 2 shows, postnatal abdominal ultrasound at 5 months old. Arrow indicates biliary sludge.
Figure 2. Abdominal ultrasound at 5 months of age. Arrow indicates mobile echogenic material consistent with biliary sludge.

Discussion

With the routine use of ultrasound (US) examinations in obstetric practice, fetal gallstones are occasionally found. In a study of almost 4,000 consecutive third-trimester US studies, 19 (0.42%) fetuses had echogenic material in the gallbladder. Eleven had echogenic foci and eight had biliary sludge [1]. Although the etiology of fetal gallstones and sludge is unknown, increased bilirubin levels may be a factor. Unconjugated bilirubin is insoluble and at higher concentrations forms calcium salt precipitates. Estrogen also plays a role by stimulating cholesterol production and reducing the synthesis of biliary acids [3]. Stone formation can occur once the concentration of cholesterol overtakes the ability of bile to maintain it in solution [4,5]. As bile becomes saturated, cholesterol vesicles form and interact with surrounding proteins such as mucin and crystallize [6,7]. Other risk factors include certain medications and maternal conditions (Table 1) [8,9]. Additionally, children with Down Syndrome are at increased risk [10].

Gallbladder sludge, thought to be the precursor of stones, is composed of calcium, pigment and cholesterol elements (calcium bilirubinate and cholesterol crystals) [11]. Sonographically it presents as diffuse homogeneous echogenic material filling the gallbladder.

| Maternal risk factors associated with neonatal gallstones |
|----------------------------------------------------------|
| Placental abruption                                       |
| Increased estrogen and progestin levels                   |
| Narcotic use                                              |
| Hemolytic diseases                                        |
| History of cholelithiasis                                |
| Enteral nutrition                                         |
| Intrahepatic cholestasis of pregnancy                    |
| Pharmacological treatment: prostaglandin E2, ceftriaxone, furosemide |
| Intoxication with denatured oil treated with steroid      |
| Sepsis                                                   |
| Prolonged fast                                           |
| Diabetes of any kind                                     |
| Twin pregnancy                                           |
| Twin pregnancy with fetal demise of one twin             |

However, unlike stones that usually cause ‘acoustic shadowing’ or ‘comet tail artifacts’, sludge lacks this sonographic finding. This may be due to small, undetectable stones that are less than 3mm or stones that escape detection because they do not lie in the center of, or in the focal range of,
the US beam. Therefore, it is difficult to distinguish sludge from small non-shadowing stones [12]. Additionally, the development of crystals that ultimately become stones is not a uniform process [6,7]. After birth, when the infant begins to feed, increasing plasma levels of cholecystokinin cause the gallbladder to contract and release the viscid material into the duodenum. This resolution may be associated with a colic pain [1].

The short-term prognosis of fetal gallstones is good. More than 70% of patients have spontaneous resolution within two months after birth and more than 90% within six months [1,8]. The prognosis for the resolution of sludge is less clear as not all infants with sludge have a follow-up US. Pharmacologic treatment with ursodeoxycholic acid and laparoscopic surgery have been reported [2, 13]. However, the risks of adverse events with medications and surgery and anesthesia in the infant population need to be considered. The long-term prognosis in older children is unclear but there are reports of older children requiring a cholecystectomy for persistent gallstones [14].

Conclusion

Although spontaneous resolution of neonatal gallstones is common in the first year of life, our patient’s US at five months of age showed that the sludge had reappeared. Knowing that sludge may progress to stone formation or that it may represent small ‘non-shadowing’ stones, periodic US will be done until complete resolution is demonstrated. Future studies are needed to determine if this neonatal finding is a risk factor for the development of gallstones in childhood.

References

1. Sepulveda W, Wong AE. (2018). Echogenic material in the fetal gallbladder: prevalence, sonographic spectrum, and perinatal outcome in an unselected third-trimester population. J Matern Fetal Neonatal Med. 2018;1-201.
2. Munjuluri N, Elgharaby N, Acolet D, Kadir RA. (2005). Fetal gallstones. Fetal Diagn Ther 2005; 20:241-243.
3. Brown DL, Teele RL, Doublet PM, DiSalvo DN, Benson CB, Vam Alstyne GA. (1992). Echogenic material in the fetal gallbladder: sonographic and clinical observations. Radiology 1992; 182:73-76.
4. St-Vil D, Yazbeck S, Luks F, Hancock B, Filiatrault D, Youssef S. (1992). Cholelithiasis in newborns and infants. J Pediatr Surg 1992; 27(10):1305-1307.
5. Halpern Z, Vinograd Z, Laufer, H, Gilat T, Moskowitz M, Bujanover Y. (1996). Characteristics of Gallbladder Bile of Infants and Children. J Pediat Gastroenter Nutr, 1996; 23(2):147-150.
6. Johnston D, Kaplan M. (1993). Pathogenesis and Treatment of Gallstones. New England Journal of Medicine. 1993; 328(6):412-421.
7. Lee Sum P, Maher K, Nicholls JF. (1988). Origin and fate of biliary sludge. Gastroenterology. 1988; 94(1):170-176.
8. Hurni Y, Vigo F, Wattenwyl BV, Ochsenbein N, Canonica C. (2017). Fetal Cholelithiasis: Antenatal Diagnosis and Neonatal Follow-Up in a Case of Twin Pregnancy – A Case Report and Review of the Literature. Ultrasound International Open. 2017; 03(01).
9. Triunfo S, Rosati P, Ferrara P, Gatto A, Scambia G. (2013). Fetal Cholelithiasis: A Diagnostic Update and a Literature Review. Clinical Medicine Insights: Case Reports. 2013; 6:153-158.
10. Boéchát Márcia Cristina Bastos, Silva Kátia Silveira da, Llerena Jr, Juan Clinton, & Boéchát, Paulo Roberto Mafra. (2007). Cholelithiasis and biliary sludge in Down’s syndrome patients. Sao Paulo Medical Journal. 2007; 125(6):329-332.
11. Jeanty, Cerine. Derderian, S. Christopher, Courtier, Jessie, Hirose, Shinjiro. (2015). Clinical management of infantile cholelithiasis. Journal of Pediatric Surgery. Volume 50 (8) pp 1289-1292.
12. Suma V, Marini A, Bucco N, Toffolutti T, Talenti E. (1998). Fetal gallstones: sonographic and clinical observations. Ultrasound Obstet Gynecol 1998; 12:439-441.
13. Gertner M, Farmer D. Laparoscopic cholecystectomy in a 16 day-old infant with chronic cholelithiasis. J Pediat Surg. 2004; 39:17-19.
14. Troyano-Luque J, Padilla-Perez A, Martinez-Wallin I, et al. (2014). Short and long term outcomes associated with fetal cholelithiasis: a report of two cases with antenatal diagnosis and post-natal follow-up. Case Rep Obst Gynecol. 2014; 2014:714271.