**Megacystis Microcolon Intestinal Hypoperistalsis Syndrome (MMIHS): A Case Report in Thailand**

Jirapa Manop MD*,
Sangkae Chamnanvanakij MD*, Charnvit Wattanasarn MD**

* Department of Pediatrics, Phramongkutklao Hospital
** Department of Surgery, Phramongkutklao Hospital

Megacystis microcolon intestinal hypoperistalsis syndrome (MMIHS) is a rare congenital disease with high mortality rate. The authors report a case of a female term infant with massive abdominal distension at birth. Abdominal ultrasonography revealed a huge cystic mass resolved after urinary catheterization. Exploratory laparotomy was performed and intraoperative findings were an enlarged urinary bladder, microcolon, short bowel and malrotation of the small intestine. Ladd’s procedure, ileostomy and vesicostomy were performed. Pathological findings of rectal biopsy revealed normal ganglion cells. Although prokinetic drugs were given for promoting bowel motility, enteral feedings were not tolerated. She died from septicemia at the age of 50 days.

**Keywords**: Megacystis, Microcolon, Hypoperistalsis, MMIHS

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Megacystis microcolon intestinal hypoperistalsis syndrome (MMIHS) is a rare congenital disease characterized by abdominal distension, distended non-obstructive urinary bladder, microcolon, hypoperistalsis and malrotation of the small intestine. The etiology and pathogenesis of this disorder are not clear. Although there is no specific findings for MMIHS, pathological studies of the colon, small intestine and urinary bladder are important for differentiating from other diseases. There is no specific treatment and prognosis is very poor. The authors reported the first case of a Thai infant who had typical manifestation of MMIHS with pathological studies of the intestine.

**Case Report**

A female 2,300-gram infant was born by cesarean section at 31 weeks of gestation. Prenatal history was unremarkable except for polyhydramnios. There was no family history of congenital anomalies.

Physical examination at birth revealed markedly distended abdomen with a large movable, ill-defined mass at the middle part of the lower abdomen. The abdominal wall looked normal. Hypoactive bowel sound was noted.

Blood urea nitrogen (BUN) and serum creatinine on the first day of life were 23.24 mg/dL and 1.8 mg/dL, respectively. She had metabolic acidosis with serum bicarbonate of 16.3 mmol/L. Urinalysis on the same day showed pH of 8.0, and specific gravity of 1.005 without protein, sugar, white blood cell or red blood cell. The infant also had low serum calcium level of 5.96 mg/dL and was treated with 10% calcium gluconate. The levels of BUN and serum creatinine declined gradually to 8 mg/dL and 0.7 mg/dL, respectively within the first week. Although sodium bicarbonate was given, she continued to have metabolic acidosis until 2 weeks of age. Ultrasonography of the abdomen showed a large cystic mass with bilateral hydronephrosis and hydroureret (Fig. 1). Normal liver, gall bladder, spleen and biliary system were noted. There was no ascites.

Computerized tomography of abdomen showed a huge cystic mass sized 10x17x7 cm. The mass had a pressure effect on proximal part of the duodenum causing a markedly dilated stomach.
Voiding cystoureterography showed markedly distended urinary bladder (Fig. 2). The wall of the bladder was smooth without intrinsic or extrinsic lesion. No demonstration of vesicoureteral reflux was noted. The urethra was not evaluated because of no voiding phase.

Upper gastrointestinal study showed a distended stomach with normal gastric contour and duodenal cap (Fig. 3). There was marked delay of gastric emptying time because of narrowing of the second part of duodenum.

Since the first day of life, she was given nothing per oral and parenteral nutrition was started. Abdominal distension decreased dramatically after placing a urine catheter to release 120 milliliters of urine. Urine catheter was retained and urine output was 3-4 ml/kilogram/day. On the 16th day of age, exploratory laparotomy was performed. The operation revealed microcolon, malrotation of small intestine, small terminal ileum, short bowel and a huge urinary bladder. Surgical procedures including Ladd’s procedure, ileostomy and cutaneous vesicostomy were performed. Pathologic findings of bowel biopsy showed no significant pathological changes of ileal, appendix, sigmoid and colon. Rectal biopsy revealed normal ganglion cells.

A week after the operation, she received a very small amount of enteral nutrients by continuous gastric infusion. Prokinetic drugs, erythromycin and cisapride, were given for promoting bowel motility. She couldn’t tolerate enteral feedings and continued to receive total parenteral nutrition.

During the hospital course, she had many episodes of urinary tract infection (UTI). The first episode was at the age of 5 days with Enterobacter cloacae. She was initially treated with cefotaxime and amikacin but changed to ceftazidime and then to imipenem. The infection resolved after a 14-day course of antibiotics. At 27 days of age, she developed the second episode of UTI. Budding yeasts were noted in urine and urine culture was positive for candida albicans. She was treated with Amphotericin B. At 38 days of age, she developed UTI and clinical sepsis. Meropenem was started and cystostomy tube was removed. However, she died 12 days later. Blood and urine cultures before starting antibiotics grew Enterobacter cloacae.

Discussion:

Since Berdon et al reported 5 cases of MMIHS in 1976(1), there were only a hundred cases reported in the medical literature. MMIHS is a rare congenital
disease found more commonly in female infants. Although a few cases have been reported in siblings as autosomal recessive inheritance\(^2,3\), most cases occur sporadically. The etiology and pathogenesis of the disease are not clear. However, many hypothesis have been proposed such as degenerative disease of smooth muscle\(^4,6\), disturbance in bowel innervation\(^5,6\), imbalance of gut peptides\(^7\), in utero intramural inflammatory process of gastrointestinal and urinary tract\(^8\), defect of contractile fiber synthesis\(^9\), and drugs\(^10-12\).

The common manifestations of this disease include megacystis, hydronephrosis or hydroureter, abdominal distension, microcolon, and intestinal hypoperistalsis reported in more than 80% of patients\(^1,2,4,6-9,13-22\). Some patients have other associated anomalies such as megaesophagus\(^22\), hydrometrocolpos\(^21\), and Barrett’s esophagus\(^20\).

Some manifestations of MMIHS can be identified prenatally by routine prenatal ultrasonography such as distended stomach and urinary bladder, increased amniotic fluid, dilated renal calyces and megaureter\(^19\). The most important ultrasonographic finding is distended urinary bladder without oligohydramnios which is different from obstructive uropathy diseases. The earliest time to detect distended urinary bladder and hydronephrosis was at 16-20 weeks\(^19\).

Diagnosis of MMIHS is based on the presence of clinical characteristics including, megacystis, microcolon, intestinal malrotation, short bowel and intestinal hypoperistalsis in the absence of anatomical obstruction. The differential diagnosis depends on the main manifestations. For megacystis, differentiations from other anomalies of genitourinary tract such as prune belly syndrome, urethral agenesis and variant of caudal regression are needed. For microcolon, the differential diagnosis includes intestinal atresia,volvulus and total colon aganglionosis. Other secondary causes of ileus or pseudoobstruction such as hypothyroidism, hypokalemia and narcotics must be excluded.

Manometric studies shows that the frequency and amplitude of spontaneous contraction in the stomach and small intestine are significantly lower or absent compared to healthy controls\(^2,3\).

Pathological findings of gastrointestinal tract are varied. Abnormal findings reported include vacuolar degeneration of smooth muscle cells\(^4,6\) and shrunken neuron\(^9\). However, the most important finding of MMIHS is normal or increased ganglion cells\(^3,4,6-9,13,14,18-22\) which is different from total aganglionosis.

Nutritional support is the mainstay of treatment. Hyperalimentation is required. Palliative surgery such as jejunostomy or cystostomy is generally needed. Prokinetic drugs for promoting bowel motility are ineffective\(^3,4,9,23\). Totally digested enteral nutrients by continuous infusion through jejunal or gastric tube was reported in one case to improve the problem of feeding intolerance\(^17\). A cholinergic drug, bethanechol, was partially effective to improve peristalsis in one patient\(^16\). Only one report showed the success of multivisceral transplantation in MMIHS patients\(^24\). MMIHS is lethal disease. Most patients died within one year after birth from severe infection. Only a few cases have been reported to survive until 4-11 years of age\(^8,17\).

Even though most cases occur sporadically, the recurrence risk in subsequent pregnancies is up to 25% because there are reports of MMIHS in siblings\(^2,3\). Therefore, prenatal ultrasonography is recommended to identify a case in subsequent pregnancies.

Reference

8. Srikanth MS, Ford EG, Isaacs H Jr, Mahour GH. Megacystis microcolon intestinal hypoperistalsis


