Clinical Characteristics and Management of Benign Transient Non-Organic Ileus of Neonates: A Single-Center Experience

Hye Kyung Chang,1 Hong Koh,2 Young Ju Hong,3 Eun Young Chang,3 Seok Joo Han,3 and Jung-Tak Oh3

1Department of Surgery, Seoul St. Mary’s Hospital, The Catholic University of Korea College of Medicine, Seoul; 2Departments of Pediatrics and 3Pediatric Surgery, Severance Children’s Hospital, Yonsei University College of Medicine, Seoul, Korea.

Purpose: The term benign transient non-organic ileus of neonates (BTNIN) is applied to neonates who present symptoms and plain radiographic findings of Hirschsprung’s disease, but do not have aganglionic bowel and are managed well by conservative treatment. It can often be difficult to diagnose BTNIN because its initial symptoms are similar to those of Hirschsprung’s disease. The aim of this study is to evaluate the clinical characteristics and proper treatment of BTNIN.

Materials and Methods: A retrospective review was made on the clinical data of 19 neonates who were treated for BTNIN between January 2008 and December 2011 at a single facility.

Results: Abdominal distension occurred in every patient (19/19). Other common symptoms included emesis (5/19), explosive defecation (5/19), and constipation (4/19). The vast majority of patients (15/19) experienced the onset of symptoms between 2 and 4 weeks of age. Radiograph findings from all of the patients were similar to Hirschsprung’s disease. A barium study showed a transition zone in 33.4% (6/18) of the patients. However, rectal biopsy revealed ganglion cells in the distal rectum in 88.2% (15/17) of the patients, and anorectal manometry showed a normal rectoanal inhibitory reflex in 90% (9/10). All patients responded well to conservative treatment. Symptoms disappeared at the mean age of 4.9±1.0 months, and the abdominal radiographs normalized.

Conclusion: BTNIN had an excellent outcome with conservative treatment, and must be differentiated from Hirschsprung’s disease. A rectal biopsy and anorectal manometry were useful diagnostic tools in the differential diagnosis.

Key Words: Ileus, neonate, Hirschsprung’s disease, transient

INTRODUCTION

First described in 1886, Hirschsprung’s disease is generally suspected in neonates and infants with abdominal distension and constipation.1 Pediatricians or pediatric surgeons often encounter patients whose symptoms are similar to Hirschsprung’s disease, however, ganglion cells are present in their rectal biopsies and their clinical course are different form the Hirschsprung’s disease. These patients are variously labeled with pseudo-Hirschsprung’s disease, neonatal intestinal pseudo-obstruction, intestinal neuronal dysplasia, hypoganglionosis, and internal anal sphincter achala-
patients who were treated for BTNIN at Severance Children’s Hospital in Seoul, Korea between January 2008 and December 2011. A diagnosis of BTNIN was established based on previously published criteria: presence of a non-organic ileus of the neonate excluding Hirschsprung’s disease, Hirschsprung’s disease-allied disorders, and functional ileus associated with meconium obstruction. Nineteen patients were treated during the study period, and no patients were excluded from the study. Patients’ demographics, symptoms, diagnostic studies, and clinical courses were analyzed.

This study was approved by the Institutional Review Board of Severance Children’s Hospital.

RESULTS

Clinical characteristics
The clinical characteristics of the patients are shown in Table 1. All of the patients were healthy neonates, and only one patient was born at less than 35 weeks gestational age with a birth weight of 2.5 kg. All but two patients were breast-fed. Abdominal distension occurred in all of the patients. Emesis (5/19), explosive defecation (5/19), and constipation (4/19) were other common symptoms. The onset of majority (79%) of patients’ symptoms was between 2 and 4 weeks of age. Four patients’ symptoms began within the first week of life.

Diagnostic studies (Table 2)
Plain radiograph findings from all of the patients were similar to Hirschsprung’s disease (Fig. 1A, B and C). A barium study was performed on 94.7% (18/19) of the patients, and 33.4% (6/18) of those patients showed the transition zones. Rectal biopsies identified ganglion cells in the distal rectum in 88.2% (15/17) of the patients without findings of any other diseases. The two patients without identified ganglion cells did not undergo repeated rectal biopsies because of normal findings on the barium studies and rapid improvement in symptoms. Anorectal manometry also showed a normal rectoanal inhibitory reflex in 90% (9/10) of the patients.

Clinical outcomes
All patients responded well to conservative treatment of intermittent glycerin enema. The frequency of glycerin enemas...
depended upon the symptoms, and ranged from daily to once weekly. No other medications were administered to relieve the symptoms, and the feeding strategy did not change because the symptoms gradually improved. After 4.4±1.1 months of conservative treatment, the symptoms disappeared by the mean age of 4.9±1.0 months (range, 2.8-6.6 months). The abdominal radiographs also normalized (Fig. 1G, H and I).

**DISCUSSION**

Although the term of ‘benign transient non-organic ileus of neonate’ is not well established yet, it should be considered in the differential diagnosis of Hirschsprung’s disease. First reported in 2002, BTNIN did not receive much attention from pediatric surgeons because of its favorable clinical course, and very few supportive articles have been published.4,5 However, the age at the definitive surgery of Hirschsprung’s disease is far younger than previous decade and one-stage procedure has been popular.6-8 These new treatment strategies require early and reliable diagnosis.9-11 Consequently, it is more important than ever to differentiate Hirschsprung’s disease from other nonsurgical conditions.9-11

The most important characteristics of BTNIN are its similarity to Hirschsprung’s disease. In our present study, the
initial symptoms in order of frequency were abdominal distention, emesis, explosive defecation, and constipation. Unfortunately, however, these symptoms alone are not distinguishable from those of Hirschsprung’s disease. In addition, plain radiographs of the abdomen are also very similar to Hirschsprung’s disease. One-third of our patients showed the typical transition zone in barium studies. However, the presence of ganglion cells in rectal biopsies and the presence of the rectoanal inhibitory reflex in anorectal manometry were excellent discriminators in favor of BTNIN.

BTNIN responds well to conservative treatment and demonstrates an excellent prognosis. The average duration of treatment was only 4.4 months in our study and has been reported to be 5 months in a previous study. Conservative treatment should be periodic or intermittent enema. Prokinetic medication has not shown any remarkable effect.

The etiology of BTNIN has not been fully understood, but it could not be classified as the variants of Hirschsprung’s disease or meconium related disorders. BTNIN occurs in the neonatal period with no association with birth weight or complications during pregnancy. Our results identified only one patient born at less than 35 weeks gestational age with a birth weight of 2.5 kg. Most patients with BTNIN showed normal appearance of ganglion cells without hyper- or hypoganglionosis. In addition, the clinical outcome of BTNIN in this study was more favorable than the other variants of Hirschsprung’s disease.

The one possible cause of BTNIN was allergy-related disorder. Kubota, et al. suggest that cow’s milk allergy is the main cause of BTNIN. In their study, patients who were diagnosed with BTNIN showed a positive drug-induced lymphocyte stimulation test for cow’s milk. The majority of these patients had symptom relief after conversion to formula. Allergic proctitis, a specific form of cow’s milk allergy, could induce symptoms mimicking Hirschsprung’s disease. Eosinophilic infiltration and inflammatory cell infiltration in the lamina propria of rectum have been identified, and rectal suction biopsy is very useful to differentiate this from other disorders.

However, our results did not support the aforementioned etiology. Although we had the limitation that allergy test for cow’s milk was not performed, most patients in our study were breast-fed and improved without a formula change. Although cow’s milk allergy could happen in breast-fed infants, the proportion of breast feeding in our study was too high to consider cow’s milk allergy as the sole etiology of BTNIN. Allergic proctitis could not be considered as the possible cause of BTNIN because no patients showed eosinophilic infiltration in the rectal biopsies in our study. Considering that BTNIN occurs in the neonatal period and disappears with age, immaturity of the ganglion cells is suspected as the likely pathophysiology.

In this retrospective study, we could not perform advanced histopathologic evaluations. Furthermore, the number of patients was not sufficient to generate any statistical results. These are also the limitations of this study. Nevertheless, we think that our results showed important clinical characteristics and management of BTNIN.

In summary, our data suggest that BTNIN has symptoms very similar to but should be differentiated from Hirschsprung’s disease. Rectal biopsy and anorectal manometry were useful diagnostic tools. Patients with BTNIN had excellent outcomes with conservative treatment, and symptoms disappeared by 5 months of age. The pathophysiology of this disease is not fully understood, and further studies are required.

REFERENCES

11. Lewis NA, Levitt MA, Zallen GS, Zafar MS, Iacono KL, Rossman
