CHAPTER 61 NEONATAL INTESTINAL OBSTRUCTION

Daniel Sidler Miliard Debrew Kokila Lakhoo

Introduction

Neonatal intestinal obstruction (NIO) is one of the most common emergency conditions a paediatric surgeon is called upon to assess during the nermatal period. Successful management of NIO depends on timely diagnosis and referred for therapy. The diagnosis is based on history (symptoms) and physical examination (signs) confirmed by some investigations such as radiographic and historpathological studies. Catastrophic events such as volvulus, ischaemic loop of bowel, presumoperitonesum, and/or presumonia from aspiration and malnutrium could be overcome through efficient and timely resuscitation and argent transport to a specialised unit.

The desired goal of healthy survival of neonatal intestinal obstruction requires a coordinated interaction of medical, norsing, and orbabilitative specialities in an organised team. Early surgical intervention is paramount and may mean all the difference between intestinal salvage and original short gut syndrome. The typical case of neonatal bowel obstruction is generally straightforward, and the outcome is potentially excellent. Only very preterm babies and those of extremely low birth weight may succumb. However, in Africa, late presentations and poor resources lend to a mortality of up to 50%. Few dedicated productic hospitals exist in developing countries.

Intestinal obstruction can be complete (atresia, anorectal malformation (ARMI) or incomplete (stemosis, web). Obstruction may be intraluminal (meconium ileus or meconium play syndrome) or functional (Blaschoprung's disease (BID)). Proximal obstruction presents with earlier vomiting and less abdominal distersion, whereas distal bowel obstruction lends itself to late-emesis and greater abdominal distention. This chapter provides an overview on neonatal bowel obstruction. Each specific condition is covered in chapters elsewhere in this book, specifically, Chapter 58 (inguinal and fernoral hemias and hydroceles), Chapter 62 (duodenal atresia and stenosis), Chapter 63 (intestinal intersia and stenosis), Chapter 65 (intestinal malrotation and midgat volvulas), Chapter 67 (meconium disease), Chapter 76 (Hinschoprung's disease), and Chapter 77 (anorectal anomalies).

Demographics

The incidence of NIO is approximately 1 in 5,000 live births. The true incidence in Africa is unknown, but a recent report from Tanzania has above that it is still the most common neonatal surgical emergency.' Many cases still die undiagnosed and untreated.

Aetiology/Pathophysiology

Neonatal intestinal obstruction has varied actiology, so the pathophysiology is diverse,

The gastrointestinal tract (GIT) arises from the yolk sac. At 3 to 4 weeks' gestation, it becomes a distinct entity. A connection, the vitelline (omphalomesenteric) duct, may persist as a Mockel's diverticulum. The alimentary tube is divided according to its blood supply into the foregut, midgut, and hindget.

The foregut comprises the oesophagus, stomach, and dandesum. These are vascularised by multiple sources—the thyrocervical, intercostal, celiac usis, and superior mesenteric vessels.

The midgat comprises the jejumum and ileum as well as the ascending and proximal transverse coton. These are supplied by the superior measureric vessels.

The hindgest comprises the distal colon, which is supplied by the interior meansteric vessels, and the rectam, which is supplied by the internal ilias vessels.

Aberrations of foregat formation include dandenal stenssis, dacdenal atresia, and annular pancreas. Maldevelopment of the midgat includes malretation as well as jejanal and iteal atresia. Meconium iteas involves the distal iteam. Hirschsprung's disease (aganglionic megacolon), meconium plug syndrome, and imperforate arms involve the hindgat. Emeric duelications occur in all three locations.

Jejunoilleal atresia is a condition acquired during focial development due to disraption of the mesenteric blood supply. In their classic work on foctal dogs in 1955, Louw and Barnard from Cape Town, South Africa, clarified the pathophysiology of jejunoileal atresia. Other abdominal conditions occurring in stero, such as gestroschois, volvulus, or intususception, may be associated with intestinal atresia due to kinking, stockhing, or otherwise disrapting the blood flow to the foctal bowel. Chromosomal attenualies are rate (<1%) in bubies with jejunoileal atresia.

Meconium fleus is the earliest manifestation of cystic fibrosis (CF), an autonomal recessive condition characterised by absormalities in cellular recentrase physiology and obloride ion transport that contribute to progressive respiratory failure, derangements in cellular secretory patterns, and diminished mucosal motility. In developed countries, 10–20% of newborns with CF present with meconium items, an association first described by Landsteiner in 1905. A cystic fibrosis gene that is different from that in the caracterium population has been identified in the black. African population: however, the incidence in Africa is much reduced, and in many sub-Sahutan countries, the disease is not reported.

Abnormalities in the cystic fibrosis trustmembrane regulator (CFTR) disrupt transmembrane flux of the chloride ion, which subsequently affects sedium transport as well. The reconsum of affected babies is thick and sticky; this, coupled with the poor motifity of an immature intestine, leads to intraluminal observacion of the terminal fleum. A contrast enema might reveal the characteristic finding of a microcolon. It is exsential to push the contrast up to the level of obstruction, namely, the distal illeum to be therapeutic by flushing out the meconium plugs (Figures 61.) to 61.4).

Meconium plug syndrome refers to inspissated recentum obstrucing the colon; it may denote HD but not CF. Conditions that predispose to dysmotility of the neonatal bowel te.g., maternal pre-ectampsia, diabetes mellitus, administration of magnesium sulfate, prematurity, sepsis, and hypothyroidism) may be responsible for the formation of the meconium plug. A water-soluble continut enema can be both diagnostic and therapeutic for this condition.

Hirschsprung's disease is a disorder of the neuroenteric pathways in the distal color that results in a bowel that is tonically contracted. Bowel peristals is controlled by neuroenteric ganglion cells, which

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Chapter 61 Neonatal Intestinal Obstruction

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