

25.1 Definition and Terminology

The term idiopathic constipation refers to the incapacity or difficulty to pass stool regularly and efficiently. In addition, we believe that it also means incapacity to empty the colon.

We use the term “idiopathic” because we do not know the etiology of this condition. We are aware of many proposed explanations to understand the pathophysiology of this condition, but we firmly believe that none of those explanations have solid scientific basis. In agreement with Benjamin Disraeli, we believe that “to be conscious that you are ignorant is a great step to knowledge” [1].

We intentionally avoided other terms frequently used in the literature, basically because many of those names implied an accepted etiology. Some of those names include: “puborectalis spasm syndrome,” “descending perineum syndrome,” “chronic obstipation,” and “spastic pelvic floor syndrome.” In addition and most importantly the treatments that we have to offer to patients with constipation are not different, regardless of the category or type of constipation that the patient suffers from.

We also refer to idiopathic constipation as the central subject of this chapter because, by far, the greatest number of patients coming to our clinic

with a colonic motility disorder, requesting a surgical consultation, belongs to this particular group.

25.2 Incidence, Social Impact, and Relevance

Idiopathic constipation is by far the most common defecation disorder and colonic motility disorder seen in children. It represents a common cause for surgical consultation [2–4]. It affects millions of Americans, as well as patients of other countries, but perhaps most important is the fact that it is an incapacitating condition when it is not treated properly. In fact, it produces a form of fecal incontinence known as encopresis or overflow pseudoincontinence that makes the patient socially rejected and discriminated.

25.3 Etiology

We are aware of multiple publications proposing different possible causes for this condition. However, most of those explanations have no scientific basis, and therefore we do not embrace them. We prefer to take a healthy and potentially more productive attitude by declaring our ignorance about the origin of this condition.

Some authors believe that diet is very important as an etiologic factor of constipation [5, 6].

Electronic supplementary material Supplementary material is available in the online version of this chapter at [10.1007/978-3-319-14989-9_25](https://doi.org/10.1007/978-3-319-14989-9_25).

There is no question that different types of food have either a laxative or a constipating effect on our bodies. In addition, we recognize the existence of personal idiosyncrasies that explain why one type of food may act as a laxative for one individual and have a constipating effect for another one. Although we recognize that diet is important to regulate colonic motility, we believe that the therapeutic value of diet is negligible in the most serious forms of constipation. We must keep in mind that this chapter belongs to a book of surgical treatments of colorectal problems in children. The type of constipation that is manageable by diet belongs to the pediatric clinics. Those patients do not come to our clinic because they are treated either by a pediatrician or gastroenterologist. The patients that come for surgical consultation are patients that have already received all kinds of unsuccessful medical managements previously.

There are also many articles [7–12] that try to explain the problem of idiopathic constipation on psychological basis. The psychodynamic mechanisms proposed are interesting and sometimes picturesque; including, strict demanding parents. Strict demanding parents who impose rigid rules on a child during the toilet training process, children that supposedly retain the stool to manipulate the parents to achieve their own purposes. All these mechanisms may have an element of truth, but we do not believe that they can explain the severe forms of constipation in patients with fecal pseudo-incontinence, giant megacolon, sometimes megabladder, serious nutritional and developmental disturbances, and sometimes death. We think that it is certainly not easy to retain stool voluntarily in an otherwise autonomous normal rectosigmoid with normal peristalsis.

It is true that most patients suffering from idiopathic constipation also have a psychological disorder, but we do not think it is a primary one. Any human being suffering from severe constipation and soiling, ostracized and discriminated understandably, must have a very significant secondary psychological problem.

Surgeons, on the other hand, have proposed different potential mechanisms to explain this problem. For instance, a rather simplistic expla-

nation is that there is a lack of relaxation of the “internal sphincter” also known as achalasia of the internal sphincter [13–16]. This is, obviously, a very attractive and popular idea. In other words, a simplistic logic dictates that incontinence means “lack of sphincter”; therefore, constipation most likely means “too much sphincter.” However, as discussed in the chapter of ultrashort Hirschsprung’s, we do not believe that the “achalasia of the internal sphincter” is an entity that can explain the symptoms of these patients.

Many patients suffering from idiopathic constipation are subjected to rectal manometry. Many of them have no relaxation reflex. The lack of relaxation has been described as a diagnostic of Hirschsprung’s disease, and therefore the next step is usually to take a rectal biopsy. If the rectum has no ganglion cells, the diagnosis of Hirschsprung’s disease is confirmed. On the other hand, if the rectal biopsy shows ganglion cells, the patient then receives the diagnosis of “achalasia of the internal sphincter.”

25.3.1 Ultrashort Segment Hirschsprung’s Disease

Ultrashort Hirschsprung’s disease has been defined as a condition in which a small length of the distal bowel has no ganglion cells. The specific length of this aganglionic zone has not been defined. Under normal circumstances, human beings have an area of aganglionosis above the pectinate line [43–45]. The length of this normal aganglionic zone has not been well established at different ages from premature life to the adult size. Consequently, it is very difficult to know if a biopsy that shows absent ganglion cells was taken from this normal aganglionic area of the rectum. At what point an aganglionic zone is considered normal and at what point is considered typical Hirschsprung’s? Both are unanswered questions.

Many authors believe that this is a common cause of constipation [39–42]. We believe that “ultrashort segment Hirschsprung’s disease” and the so-called internal sphincter achalasia are highly debatable conditions. From the clinical

and radiologic point of view, patients who have been labeled with these conditions cannot be differentiated from those suffering from idiopathic constipation clinically or from the radiologic point of view. In other words, the three groups of patients suffer from severe constipation, they have a tendency to soil the underwear when the constipation is severe, they have a very dilated rectum, and most importantly, the three groups respond to the use of laxatives. Those authors who support the existence of these two conditions (ultrashort segment Hirschsprung's and internal sphincter acatalasia) [13–16, 33–40] claim that they can make the diagnosis based on manometric studies [17–21] and confirmed by sophisticated histochemical techniques not always available to the clinician.

We have serious questions about the existence of these conditions for several reasons:

25.3.2 Rectal Manometry

During a regular rectal manometry study, a balloon is placed inside the lumen of the rectum, and the pressure of the anal canal is recorded. Under normal circumstances, the inflation of the balloon in the rectum elicits a response from the patient consisting in a drop of pressure in the anal canal, which the manometrists interpret as “relaxation of the internal sphincter.” The first question that comes to our mind is how do they know that the internal sphincter is the structure that is relaxing? If we think about the real anatomy of normal individuals (see Chap. 2), we find that the so-called internal sphincter has been defined as a thickening of the circular layer of smooth muscle of the most distal portion of the bowel. However, surrounding the entire rectum, there is an obvious and powerful striated skeletal muscle structure or voluntary sphincter mechanism. During very early manometric studies, some investigators used paralyzing agents in animals and even in human volunteers in order to be able to discriminate the relaxation of the striated muscle, from relaxation of the smooth muscle (internal sphincter), and they concluded that it was the smooth muscle that was relaxing. In addition, they even

reached the conclusion that 85 % of the bowel control depended on the “internal anal sphincter!” Yet, during all the clinical manometric evaluations of patients, muscle relaxants are not used, and therefore, we do not know if the drop of pressure in the anal canal is due to a relaxation of the striated muscle complex or to the smooth muscle.

Another very serious question that comes to our mind when discussing rectal manometry is the fact that most patients evaluated manometrically suffer from constipation, and, by definition, most of them have a very dilated rectum. In order for the patient to have a relaxation reflex, it is necessary to stretch the rectal wall with the inflated balloon. That means that we are supposed to use different volume balloons for different patients depending on the size of the rectum. We are unaware of this kind of methodology. Most manometries use the same size balloon for all patients, and certainly we have never seen 1 or 2 L balloons, yet we have seen many patients with giant rectums. As a consequence, it is conceivable that the inflation of a relatively small balloon that does not stretch the rectum would produce a negative reflex. In other words, the pressure in the anal canal would not drop because actually the rectum was never distended due to the fact that they used a standard size balloon in a megarectum. When we asked these questions to different proponents of manometry, we do not have a reasonable answer.

25.3.3 Doubts and Questions About the Anatomy of the Internal Sphincter

The “internal sphincter” has been defined as a thickening of the circular layer of the smooth muscle of the bowel in its most distal portion, the anorectum. Yet, real photographs documenting the presence of this structure are extremely unusual to see (see Chap. 2). A description of the specific thickness and upper and lower limits of this elusive structure at different ages is inexistent. There are plenty of diagrams but very few photographs, and those photographs do not show

the limits of that structure and the way to separate it from the striated sphincter mechanism. Recently, endoanal sonography illustrates a series of circular structures that are interpreted arbitrarily with an arrow as “internal sphincter.” In addition, most of those endoanal study publications do not specify how much they introduce the device and how the so-called internal sphincter looks at different depths in the anal canal and the rectum.

Our experience in the surgical exploration of the normal rectum via posterior sagittal for the repair of urethral problems or tumors did not allow us to identify a structure that looked similar to the descriptions of the so-called internal sphincter (see Chap. 2). With a posterior sagittal incision, we can identify and clean the entire posterior rectal wall all the way down to the skin and evaluate the thickness of the bowel wall. We have been unable to see a thickening of the bowel wall layer. With the use of an electrical stimulator, we can clearly differentiate what is a striated muscle from a smooth muscle, and, again, we have not seen the so-called internal sphincter. Another problem is that all anatomic structures have different sizes depending on the patient's age, and we are unaware of a study of the characteristics and size of the anatomic structures such as the so-called internal sphincter at different ages.

25.3.4 Questions About Myectomy Technique

The treatment proposed for the treatment of ultrashort Hirschsprung's and internal sphincter achalasia is an operation called myotomy or “myectomy,” myotomy being just dissection of the internal sphincter and myectomy a resection of part of that [22–25]. The description of this operation is rather vague; the surgeon must make an incision at the posterior aspect of the mucocutaneous junction in the anus and create a plane of dissection between the mucosa and the posterior wall of the rectum. All the tissue that remains between the mucosa and the posterior wall of the rectum is smooth muscle, and once the surgeon created those two planes, he is supposed to resect

a strip of that tissue. The length of this strip has not been defined for different ages. That specimen should be oriented and should be sent to pathology. The pathologist is supposed to look for ganglion cells in a well-oriented specimen, and he is going to find absent ganglion cells in the most distal part and present ganglion cells in the proximal portion. However, again, the question is how do we know what is a normal length of aganglionosis in a human being? We are unaware of publications describing the specific technique followed by different pathologists. The study of the specimens must show that only included smooth muscle. In addition, looking at the pictures of the intraoperative operations as well as the description of the surgical technique used by different surgeons, one gets the feeling that every surgeon is doing a different technique, and they do not really know exactly what is included in the specimen that they send to pathology. Interestingly, for those who believe that the internal sphincter is the key for bowel control, it is difficult to conciliate the concept with the idea of doing an operation to cut that “important structure.”

25.3.5 Botulinum Toxin Injection

Recently, the injection of botulinum toxin is becoming popular to paralyze the “internal sphincter” and by doing that to improve the symptoms consecutive to the lack of relaxation of this structure [102–104]. A detailed, meticulous description of the injection has not been published. The injection is performed rather blindly. There are some reports on the endosonographic control of the injection, but again, the fact that we see an endosonographic image does not tell us exactly where we are injecting the toxin. The effect of this toxin, as we know, is to paralyze the muscle, but it also paralyzes the striated muscle. How do we know if we are actually injecting the voluntary sphincter mechanism rather than the internal sphincter? In addition, the effect of the injection of botulinum toxin is transient and must be repeated to continue the effect, and that is another reason why we are so skeptical about it.

As can be concluded from all this discussion, until all these questions can be answered in a satisfactory way, we will continue thinking that we need more scientific and systematic studies to clarify this rather confusing subject.

There are many publications of authors who propose the intestinal neuronal dysplasia (IND) as a potential explanation for some cases of constipation [26–31]. Again, we are rather skeptical about this. A critical, comprehensive evaluation of the literature on neuronal intestinal dysplasia was conducted by us [32]. The most obvious impression that we obtained from this review was that there is no basic agreement between pathologists about this histologic diagnosis [33]. In addition, we are not aware of the existence of topographic studies that describe the extension of this histologic disorder in different patients. For a surgeon to be able to offer a rational treatment for this condition, we need to know the extent of the affected bowel that will be resected. In theory that will cure the patient. This has never been done or has never been reported. In addition, the symptoms described in patients with neuronal intestinal dysplasia vary from patient to patient. The treatments vary from laxatives to enemas and to different types of resections, and finally, the follow-up of the patients has not been consistent. To complicate the problem even more, some patients recover spontaneously. We believe that “neuronal intestinal dysplasia” represents an interesting histologic disorder that deserves further scientific evaluation to try to establish a truthful clinicopathologic correlation, but at the present time, the concept has very little clinical application.

“Hypoganglionosis” has also been invoked as a potential explanation for patients with severe constipation [34–38]. Again, we have raised several serious questions about the existence of such entity. We do not know if the number of ganglion cells remains constant through our lifetime. Assuming that the number of ganglion cells remains the same in patients who develop constipation and megacolon, it is conceivable that in a specimen taken from the colon in a case with a giant megasigmoid the pathologist will see relatively less ganglion cells in every field and that

may give him the impression of dealing with a case of hypoganglionosis. Yet, what the pathologist really sees in that case is only the result of stretching a normal ganglionic colon. We have never heard a coherent answer to this question, and therefore we believe that at the present time we have no basis to make such diagnosis. In addition, the treatment for such condition, assuming that exists, is the same that we offer for idiopathic constipation.

As we discussed in the Chap. 24 and specifically in the paragraph dedicated to “ultrashort Hirschsprung’s,” most of the patients that come to our center for consultation for severe idiopathic constipation already had manometric studies of the colon and the rectum, as well as rectal biopsies, with different inconsistent types of results. The protocol of management that we propose for all these patients is the same. We believe that there is no way clinically to differentiate those patients with idiopathic constipation and so-called ultrashort Hirschsprung’s, and we believe that the treatment should be the same.

We try to follow closely the developments related to newly described histologic disorders that may explain the problem of constipation including a deficiency of the substance P [46], abnormalities found with the use of monoclonal anti-neurofilament antibodies [47], and abnormalities in the cells of Cajal [48]; also we are recently learning more about the increased plasma level of pancreatic polypeptide and a decreased plasma level of motilin in children with encopresis [49]. All these deserve future investigation but at the present time have no clinical application.

We believe that patients with idiopathic constipation are born with a colonic hypomotility, not a well-characterized disorder, that affects mainly the rectosigmoid, but also may extend to the rest of the colon. We also strongly believe that this is a spectrum type of condition that may include patients who have very mild constipation, manageable with diet, as well as very severe cases that overlap with a condition known as “intestinal pseudo-obstruction” and may also kill patients. We also believe that the motility disorder frequently affects also the urinary tract simul-

taneously. We do not believe that the urinary problems seen in some of these patients are consecutive to the mechanical effect of the dilatation of the colon, but rather that these patients suffer from a similar idiopathic malfunction (hypotonic, large) of the bladder associated to the hypomotility of the rectosigmoid.

The concept of spectrum of the disease cannot be overemphasized. Most of the proposed treatments for constipation [50–55] do not take this concept into consideration. The authors rather offer standard therapeutic protocols, like if all patients suffered from the same degree of constipation. In other words, the treatments proposed are not individualized. Interestingly, the different modalities of treatment proposed in the literature [56–64] report percentages of success that varies from 50 to 80 %, but there is always a group of patients who do not respond. We believe that this is another manifestation of a spectrum type of condition. We also believe that we, pediatric surgeons, most likely will be dealing with the group of patients in whom the traditional therapeutic strategies failed.

25.4 Pathogenesis

Although we do not know the cause of idiopathic constipation, we have learned a great deal about its natural history due to the long-term follow-up of our patients. Idiopathic constipation is a self-perpetuating and self-aggravating incurable condition, incurable but manageable. We assume that the babies are born with a primary hypomotility disorder that affects mainly the rectosigmoid but may affect the entire colon. This motility disorder incapacitates the patient to empty the rectum. This produces accumulation of stool, which is responsible for the dilatation of the rectosigmoid or sometimes other portions of the colon.

We have learned that in patients that are born with atresias of different hollow viscus, reconnecting or reanastomosing an extremely dilated, chronically obstructed hollow viscus to a tiny, nonused microintestine produces poor results in terms of function which is attributed to a lack of peristalsis of the most dilated part of the bowel or

hollow viscus. Therefore, the recommendation is to resect the most dilated part of the bowel or to taper it. This has been observed when dealing with the small bowel, colon, esophagus, or ureter. It seems like it is necessary for the hollow viscus to have a specific diameter in order for the peristalsis to be optimal and efficient. It seems to us that the dilatation of the rectosigmoid has the same effect, affecting the peristalsis. In other words, retention of stool produces dilatation, and dilatation produces poor peristalsis. Poor peristalsis produces more retention, and more retention produces more inefficient peristalsis, creating a vicious cycle (Fig. 25.1).

Eventually the passage of a large, hard piece of stool through the anus may produce a laceration (fissure), understandably producing pain during defecation. This explains the voluntary attempt of the patient to retain the stool and avoid bowel movements. In other words, the patient is born with a certain degree of rectosigmoid malfunction, but eventually, another factor is added to the equation which is the voluntary intention to hold the stool to avoid pain. This happens often in these patients, but we do not believe that it is a primary phenomenon (Fig. 25.2). We believe that the patient retains the stool because he learned that passing the stool is painful. Therefore, the treatment of fissures in patients with constipation consists in providing the parents with the necessary information, for them to understand the pathophysiology of this condition. In other

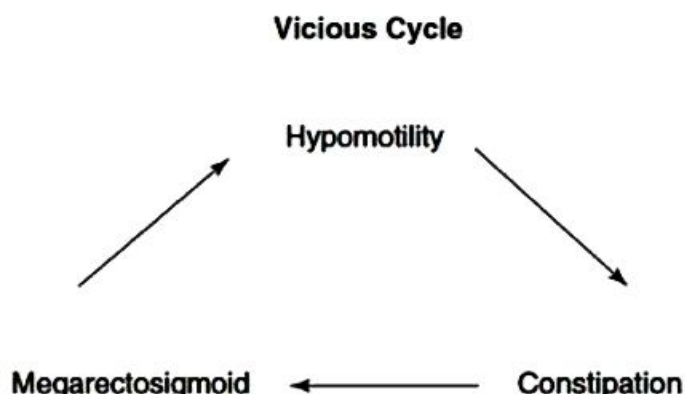


Fig. 25.1 Vicious cycle of idiopathic constipation. Poor colonic motility produces fecal retention; fecal retention produces dilatation; rectal and colonic dilatation produces poor peristalsis which continues producing more severe constipation

words, in order for the fissure to heal, we have to guarantee that the patient does not pass a hard piece of solid stool through the anus, which could reopen the fissures contributing to the aggravation of the symptoms. Stool softeners, laxatives, and time will make the fissure heal, provided the patient does not have another episode of impaction, passing hard stool which will reopen the laceration of the anus (fissure).

The concept of incurability of this condition is also fundamental for a successful management. Not understanding and not accepting the idea that this condition is incurable explains in part the high recurrence rates reported in the literature [56–64]. Treatments are provided frequently on a temporary basis based on the rather naive assumption that the condition is cured. Subsequently, the treatments are tapered or interrupted, assuming that the patient has been cured, only to find that the patient suffers a recurrence. This creates frustration for the patients and parents which may explain why most of these patients go from institution to institution looking for an answer. Sometimes, colostomies or enemas are performed, also on a temporary basis. Opening a proximal colostomy or applying enemas may produce a decrease in the size of the

dilated viscera and give the impression that the patient has been cured. In fact, the patients actually may show symptoms of improvement after the colostomies are closed or when the enemas are discontinued. However, if the patient does not receive further treatment, symptoms most likely will come back.

Figure 25.2 shows a cycle of constipation and megarectum that we believe occurs in these patients. Many publications support the idea that the problem of constipation starts during the toilet training process [50–55]. We believe that the toilet training stage of life is rather the time when the symptoms become more evident; however, we think the patients are born with this condition. Babies who are breastfed may not show symptoms because of the well-known laxative effect of the human breast milk. However, when the breastfeeding is discontinued and the patient receives formulas and other kinds of food, the symptoms become obvious. Babies who have symptoms of constipation while receiving breast milk most likely suffer from a more severe type of constipation. Many times, the parents tell us that the problem started during the preschool years. However, when we inquire specifically about the bowel movement pattern since birth, we frequently find evidence of constipation from very early in life. Actually, the parents remember most vividly the episode of the first fecal impaction, and they may refer to that event as the initiation of symptoms. Yet, we all know that a symptomatic episode of fecal impaction represents the final step of a chain of events that started a long time before.

Many pediatricians believe that normal individuals can go 2 or 3 days without a bowel movement through life without having any significant implications. We believe that that is true for many human beings; however, in dealing with patients with idiopathic constipation, it is extremely important to expect the patient to have bowel movements every day as a manifestation of the response to our treatment. Allowing the patient to go one or several days without bowel movements would generate again the vicious cycle that we have been referring to (Figs. 25.1 and 25.2).

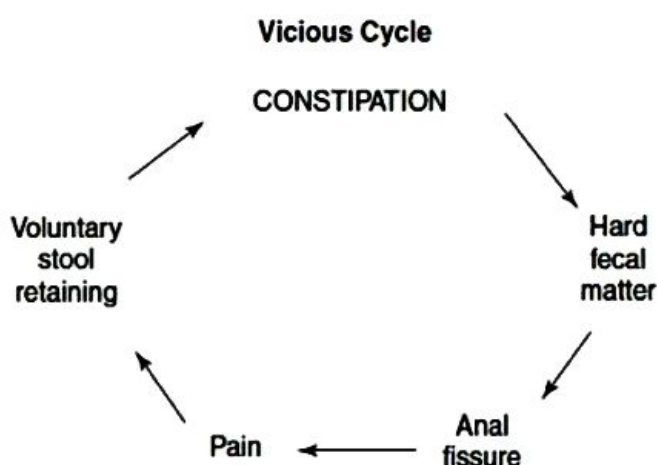


Fig. 25.2 Vicious cycle of patients with idiopathic constipation and painful bowel movements. Poor rectosigmoid motility produces stool retention. This eventually becomes fecal impaction, which is the presence of hard fecal matter in the rectum. When this finally passes through the anus, it produces a fissure. The fissure produces pain, and the pain induces the patient to try to hold the stool and avoid bowel movements, which exacerbates the stool retention

25.5 Natural History and Clinical Manifestations

A meticulous, detailed clinical history may show sometimes that babies did not pass meconium in the first 24 h and started having symptoms of constipation even when they were young babies and were taking breast milk. As the patient grows, the symptoms of constipation become more severe. The parents describe vividly how the patient suffered for the first time a painful bowel movement with blood in the stool. After that, the patient became a "stool retainer"; he goes to a corner of a room and hides while passing stool, trying to avoid the bowel movement. The patient refuses to sit on the toilet because he knows that what follows is a painful experience. Eventually, the patient has the first episode of fecal impaction. This is a very stressful event in the life of the patient and the parents. We refer to fecal impaction as a situation in which the patient has in the rectum a very large, solid piece of stool that has been there for days or weeks. When laxatives are prescribed to a patient who has fecal impaction, the result is exacerbation of severe, crampy abdominal pain and sometimes vomiting. This may resemble the symptomatology of colonic obstruction. For this reason, we consider it contraindicated the use of laxatives in a patient with fecal impaction. Occasionally, the laxatives produce diarrhea, and the patient keeps passing liquid stool around the impacted fecal matter (this is known as "paradoxical diarrhea"). This gives the parents the false impression that they are overusing laxative and may induce them to reduce the dosage, which of course, will exacerbate the problem.

Usually it is during the preschool or school age, when the patient starts showing a very bad prognostic sign which is soiling the underwear also known as "encopresis." This is a phenomenon that we call overflow pseudoincontinence. Sometimes, constipated patients have 1, 2, 3, 5, or even 10 bowel movements every day, giving the parents and the doctors the false impression that the patient is not constipated. Actually, this may be a manifestation of a serious constipation problem. The patient passes a very small amount of stool, but never empties the rectum. Eventually,

he/she will suffer from chronic fecal impaction. The impaction produces dilatation of the entire rectum including the anal canal. The anal canal represents the sensitive part of our bowel that allows us to know when stool is coming down to the anus and allows us to determine when to use our voluntary sphincter. However, we believe that perhaps the presence of a large mass of solid stool stretching the rectum on a chronic basis, eventually, makes the patient accustomed to the presence of that mass and then starts soiling on a chronic basis without the patient's awareness. This is an ominous sign. This means that the condition has already advanced very significantly and will require a very aggressive management. The patient soils the underwear day and night and basically does not have spontaneous bowel movements. At this stage, the patient basically behaves like a fecally incontinent patient, with all the implications that come with this diagnosis. In other words, the patient smells bad, and the family starts fighting with him/her and rejects them. In addition, the patient becomes accustomed to his/her own smell and is not accepted at school, and as a consequence, the patients develop serious psychological sequelae which we believe again are not primary.

Eventually, the parents believe that the patient is intentionally trying to upset them by sitting at home in the living room, obviously smelling very badly and not doing anything to solve the problem. In fact, the patient does not perceive the bad odor. These ideas in the parents are supported sometimes by the explanations given by psychiatrists in the sense that they believe that the patient is trying to manipulate the family by holding the stool intentionally. The emotional interrelations in the family are severely affected, and that is when the psychological problems become worse. We do not believe that the patients do this intentionally. We believe that if an individual wants to manipulate his/her parents, he could select many other ways to do so. Nobody wants to have stool-stained underwear, to smell bad, and to be rejected by society.

The family may put a lot of emphasis on the lack of cooperation from the patient and make the patient feel guilty. By the time these patients come for surgical consultation, they are with-

drawn, shy, negative, and reluctant to be examined by the surgeon. They usually have been subjected to many painful rectal examinations. They have scars from previous fissures in the anus. The family is usually in distress. These patients have also been subjected to unsuccessful therapeutic programs including biofeedback [65–67], behavior modification [68–74], and psychological and sometimes psychiatric treatments without positive results.

As we will see, a successful, efficient, adequate management of these patients will make the problem of encopresis disappear. Very occasionally, we see patients that we treat efficiently; in other words, we are sure that the patient no longer carries large amounts of stool in the rectum and yet they keep behaving like if they are incontinent. Those patients deserve a more meticulous study to rule out neurologic problems such as tethered cord, spina bifida, tumors, or a more severe psychological disorder. Fortunately, this particular situation is rare.

25.6 Diagnosis

The diagnosis of idiopathic constipation is a clinical one, supported by a radiologic evaluation. A patient that presents with the symptoms already described most likely has a problem of idiopathic constipation. Patients with Hirschsprung's disease do not soil. In addition, when left unattended without surgical treatment, patients with Hirschsprung's disease are at risk of dying. They frequently suffer from severe enterocolitis. The patients who survive and go undiagnosed with Hirschsprung's disease are frequently malnourished and have a history of episodes of enterocolitis. Most patients with idiopathic constipation are well nourished. It is extremely unusual to see a clinical picture similar to enterocolitis in patients with idiopathic constipations. We have seen something similar in cases with extremely severe idiopathic constipation.

Obviously, the patient must be examined with special emphasis in the characteristics of the anus, to be sure that there is no stricture and/or anterior mislocation of the anal orifice.

A contrast enema performed with a hydrosoluble material (Fig. 25.3) is the most valuable diagnostic study to confirm the diagnosis of idiopathic constipation. The characteristic image of a contrast enema in a child with a megarectosigmoid is shown in Fig. 25.4. Most of the times the dilatation of the colon affects the rectosigmoid all the way down to the level of the levator muscle which is recognized because it coincides with the pubococcygeal line (Fig. 25.4). The lack of dilatation of the rectum below the levator mechanism (pubococcygeal line) should not be interpreted as a transition zone or non-dilated rectosigmoid. Unfortunately, we have seen many patients that suffer from idiopathic constipation; somebody misinterpreted the radiologic study and erroneously treated the patient like Hirschsprung's disease.

In cases of idiopathic constipation, the rectum, above the anal canal, and the sigmoid are extremely dilated. This provokes an image that has been described many times in the literature as a "posterior



Fig. 25.3 Characteristic image of a contrast enema performed in a patient with severe idiopathic constipation. Typically the most dilated part of the colon is the rectosigmoid, and the descending and transverse colon seem to be normal in caliber



Fig. 25.4 Megarectosigmoid extending all the way down to the pubococcygeal line. Below this line, the rectum is not dilated because it is compressed by the funnel-like sphincter mechanism. Arrow shows the rectum compressed by the sphincter

shelf" (Fig. 25.4). This "posterior shelf" has been interpreted by some authors as evidence of an anteriorly located anus [75–78]. Many surgeons adopt this concept and treat these patients on the basis of that idea. We believe that such diagnosis has no scientific basis. We have never seen a real anteriorly located anus, defined as a normal anus, nonstrictured, with normal anal canal, surrounded by the sphincteric mechanism 360°. The contrast enema in patients with idiopathic constipation shows different degrees of dilatation of the rectosigmoid as expected in this spectrum of disease (Fig. 25.5a, b). Most interestingly, most of the times, there is a dramatic size discrepancy between a normal size transverse and descending colon and the very dilated megarectosigmoid (Fig. 25.5a). These changes are actually the reverse from what we see in Hirschsprung's disease (Fig. 25.6). In Hirschsprung's disease, the aganglionic segment is the most distal part of the rectosigmoid, and the dilatation is located in the proximal colon (normoganglionic).

We learned that patients with a more localized rectosigmoid dilatation have a better prognosis and respond better to the treatment. When the patients have a dilatation of the entire colon, we consider that a bad prognostic sign (Fig. 25.5b). We formally contraindicate the use of barium in these patients. The term "barium enema" is widely used. In addition, adult radiologists like to use barium because the barium lines the mucosa of the colon and allows an accurate diagnosis of mucosal abnormalities such as diverticula, ulcers, and/or polyps. In our patients, on the other hand, we are not looking for mucosal abnormalities; we rather want to see the degree, location, and extension of the dilatation of the colon, and also in addition, we want to know how well or how bad the colon empties after it has been filled up with a contrast material. In other words, we want to see sequential imaging of the filling up of the colon and then a post-evacuation film.

For many years, we took rectal biopsies in these patients as part of our routine evaluation. Now, we have found that study unnecessary when the clinical picture and the radiologic images are characteristics. At the present time, we only perform biopsies when there is a suspicious radiologic image of Hirschsprung's in the contrast enema or when the patient clinically behaves in a way similar to a patient with Hirschsprung's disease. As we previously discussed, we do not perform rectal manometries in these patients because we believe that it has no diagnostic or therapeutic value. Total colonic manometry [79–82] is a promising study. We would like very much to correlate the site, degree, and location of the dilatation of the colon with the colonic manometric abnormalities and subsequently with the histologic abnormalities. In the very few colonic manometric studies performed in our patients, we did not find a correlation between the colonic manometric results, clinical behavior, radiologic findings, and histologic changes. However, we look forward to doing more studies and learning more from this diagnostic modality. We look forward to the improvement of the accuracy of these studies, so we can use them as an aid for therapeutic purposes, but at the present time we believe are not

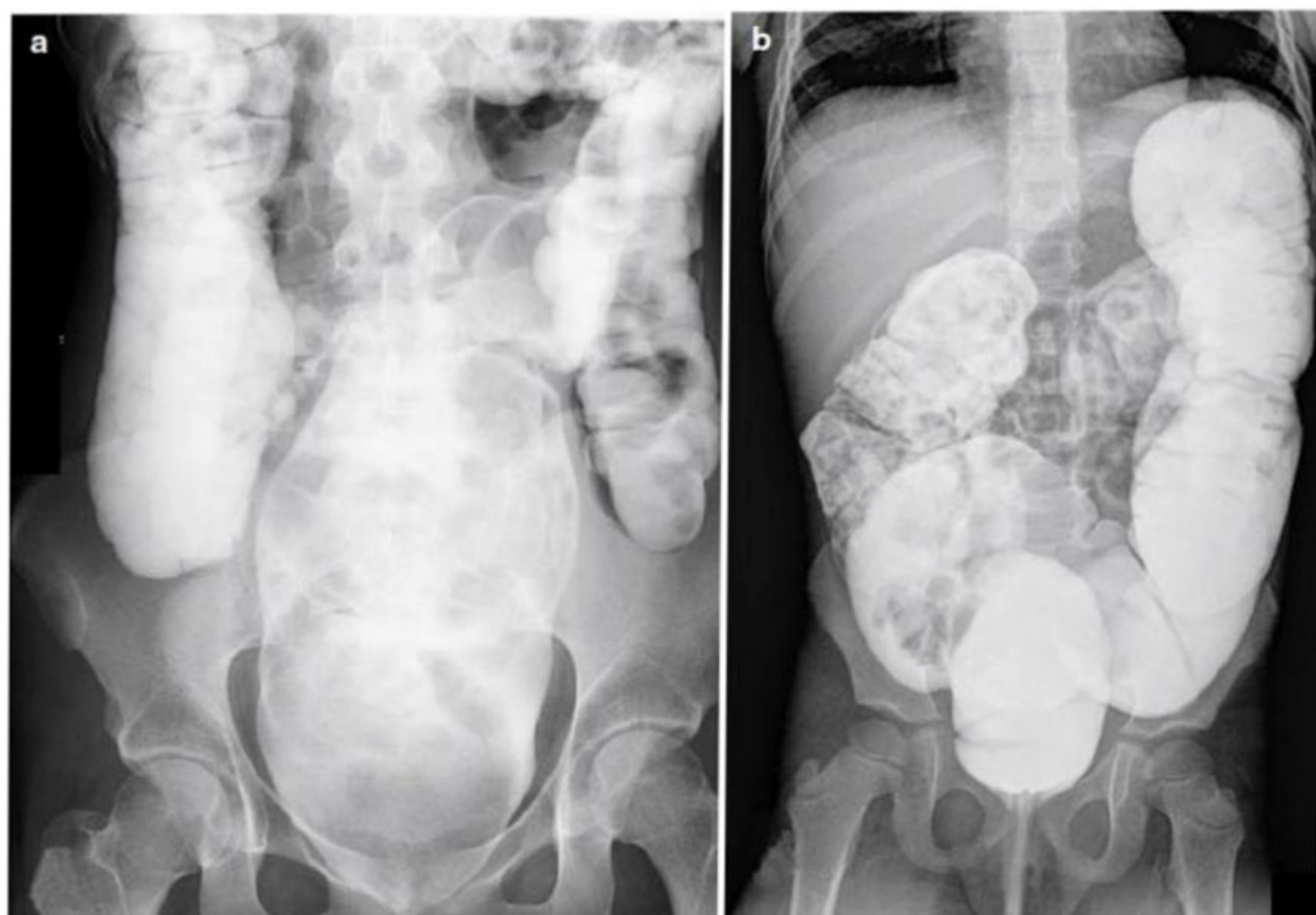


Fig. 25.5 Two different types of colonic dilatation in patients with idiopathic constipation. (a) Dilated rectosigmoid with normal caliber proximal colon. (b) Generalized colonic dilatation

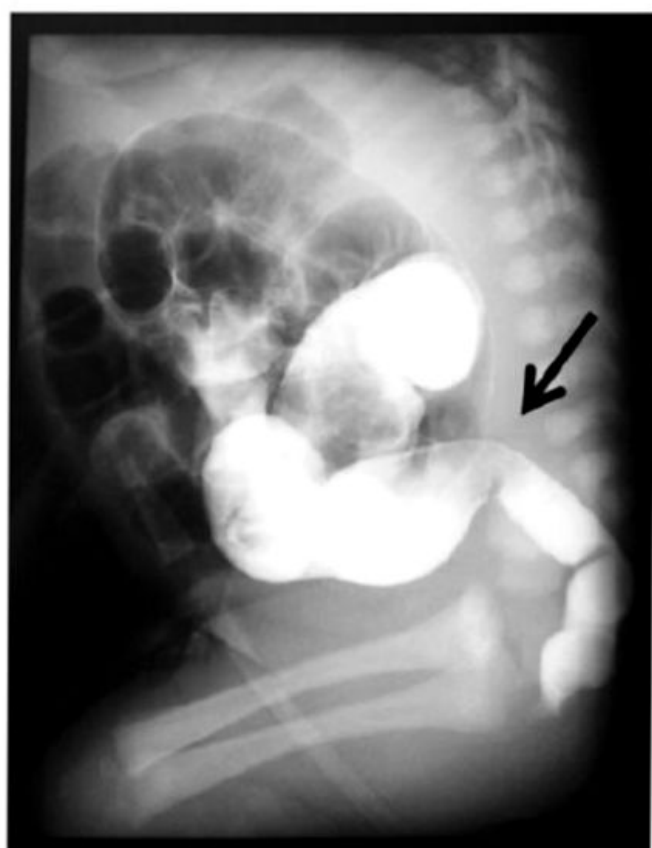


Fig. 25.6 Characteristic changes of the rectosigmoid in a patient with the most common type of Hirschsprung's disease. Arrow shows the transition zone

reliable. We also look into the future to have systematic, thorough, reliable, histologic studies of the colonic specimens from bowel resections, including the rectal portion, the most dilated part, and the non-dilated part of the colon, looking for all those recently described types of histologic abnormalities, including neuronal intestinal dysplasia [26–31], hypoganglionosis [34–38], and “desmosis” [83].

25.6.1 Colonic Transit Time

This is a promising way to evaluate the magnitude and modality of constipation. It has been done in the past using radio markers [84–87]. More recently, this type of measurement has been improved, using nuclear scintigraphy [46, 88–94]. By using this diagnostic modality, Hutson has been able to identify three transit modalities, namely, “normal transit,” “slow transit,” and “functional fecal retention.” He believes that this study helps to determine the type of treatment

that will benefit the patient; "slow transit" patients may benefit more from an operation, whereas "functional fecal retention" responds to medical management. Perhaps even more interesting, Hutson found that about 80 % of patients suffering from "slow transit constipation" had reduced SP immune reactivity in the axons of the colonic muscle and 6 % had heterotopic ganglion cells or hypoplastic ganglia on routine histology [46, 89]. Hutson also found that patients suffering from "slow transit" do not respond to "standard medical therapy." All of this represents a very significant piece of information and deserves to be commended.

We have no experience with the scintigraphic measurement of colonic transit time. However, our evaluations with contrast enemas tend to show two different types of images:

- (a) Those with dilated rectosigmoid and normal caliber proximal colon (Fig. 25.5a)
- (b) Those with a generalized dilatation of the colon (Fig. 25.5b)

In general, we observed that those with generalized dilatation of the colon have less favorable prognosis, and they suffer from a more severe form of constipation. It is conceivable that this group can suffer from "slow transit constipation," whereas those with dilatation of the rectosigmoid and normal caliber proximal colon could suffer from what Hutson calls "anorectal retention." Unfortunately Hutson does not describe what "standard medical therapy" is.

Many of the patients that come to our clinic have been "resistant to a standard medical therapy." The parents say that they tried "all kinds of laxatives" and "nothing worked." Yet, we try our protocol of management in all patients and find that most patients respond. However, the amount of laxative necessary for them to empty the colon is usually much higher than what "the book" says or than "standard" dosages.

We look forward to hear more about the clinical possible implication of the substance P deficiency.

From our point of view, the treatments of patients with idiopathic constipation are the same, regardless what specific type or modality of constipation they suffer from.

25.6.2 The Evaluation of Severity: Search for Objective "Instruments"

One serious concern about the literature related with constipation is the fact that we, the readers, do not know the severity or magnitude of the problem that the authors are referring to. Consequently, we cannot judge their results. Most publications claim a percentage of success and/or failure with different therapeutic modalities. Yet we do not know if the authors were treating patients that belong to the most benign side of the spectrum, patients who respond to a change in diet.

We firmly believe that constipation presents in the form of a spectrum. Benign forms of constipation respond to all kinds of treatments, whereas very severe forms overlap with "intestinal pseudo-obstruction" and represent a therapeutic challenge and a risk of death.

A great deal of enthusiasm has been provoked by the "Rome III criteria" as a validated measure instrument for constipation [95–98]. The efforts of the authors are appreciated. However, we were very disappointed to learn about the signs and symptoms used to evaluate the severity of constipation. They included extreme subjective signs and symptoms such as:

1. "Two or three defecations in the toilet per week" – the presence and frequency of bowel movements does not necessarily reflect the magnitude of the problem. In fact, many patients have several bowel movements, but they are fecally impacted.
2. "At least 1 episode of fecal incontinence per week" – we agree that soiling is a sign of severity; however, there are cases with severe constipation without soiling.
3. "History of retentive posturing or excessive volitional stool retention" – the history of retention attitude does not reflect the magnitude of the problem.
4. "History of painful or hard bowel movements."
5. Presence of large-diameter stools that may obstruct the toilet.

These last three criteria are so subjective that they do not deserve any comment.

We believe that the severity of the problem can be measured by the amount of senna necessary to produce a complete emptying of the colon, radiologically demonstrated.

25.7 Management

The management plan that we offer to parents of our patients is based on the following premises:

- Idiopathic constipation is mostly incurable but manageable.
- It is represented by a spectrum of severity, and therefore the treatment must be individualized.
- Most patients considered non-manageable by “standard methods” respond to an aggressive management with laxatives.
- The most objective way to know the effect of our management is with radiologic monitoring.
- As many as 85 % of our patients have been manageable. Fifteen percent are candidates for surgical treatment.

In this chapter, we concentrate our attention on the treatment of those patients with severe forms of constipation. Drugs like cisapride supposedly increased the motility of the colon but in the type of constipation that we are discussing here, have no good results [99–101]. Some surgeons use botulinum toxin injected in the “anal sphincter” to produce relaxation of the sphincter [102–104]. It is understandable to believe that this type of treatment may facilitate the passing of stool, but usually does not solve the problem of the severe idiopathic constipation, because the relaxation of the sphincter is temporary and the primary problem of the patient has not been solved. In addition, again, that modality of treatment appeals to those who believe in the simplistic idea that constipation is consecutive to the presence of “too much sphincter.” The problem of constipation seems to be less simplistic [105].

During the first visit, we give the parents a specific manuscript designed for them. In that manuscript, we emphasize the unknown nature of this condition, the fact that it is incurable and

therefore the fact that the patients have to accept treatment for life or the possibility of an operation. When the patients come to our clinic, the parents sometimes feel frustrated by the fact that we offer them a medical treatment that they think is not different from previous unsuccessful treatments. We try to convince them that although we will be using the same medications (laxative), we are going to use them with a specific rationale and following a different protocol. The first difference consists in that we adapt the dosage to the patient’s response. The overwhelming majority of patients with severe forms of idiopathic constipation that come to our clinic have been receiving insufficient amount of laxatives. The parents usually tell us that they were treated with “*ALL KINDS OF LAXATIVES* and none of them worked.” That is because they were prescribed following what the book says. We have found that these patients need 2, 3, 4, 5, and 10 times more laxatives than what the books usually says, which is a measure of the magnitude of the severity of the condition. Another very important feature, not described in the literature, is the radiologic monitoring of the patient’s response to our treatment. We become tired of speculating about the efficiency of our treatment. In other words, we used to give laxatives to a patient; the parents came back a few days later to say how great the results were. In retrospect, those patients were passing stool, but they were not emptying their colon. Yet, soon enough, we were very disappointed because the patient came back with new symptoms and are fecally impacted. The only reliable way to know how much stool is in the colon of a patient is by taking abdominal x-ray films.

When the patients come to our clinic fecally impacted (Fig. 25.7), the first step of our routine is to apply our protocol of fecal disimpaction. We explain to the parents that the disimpaction process (Animation 25.1) is going to be cumbersome and very uncomfortable for both the parents and the patient. We also tell them that if they follow our instructions, this will be the last time that the patient will be fecally impacted in his/her life.

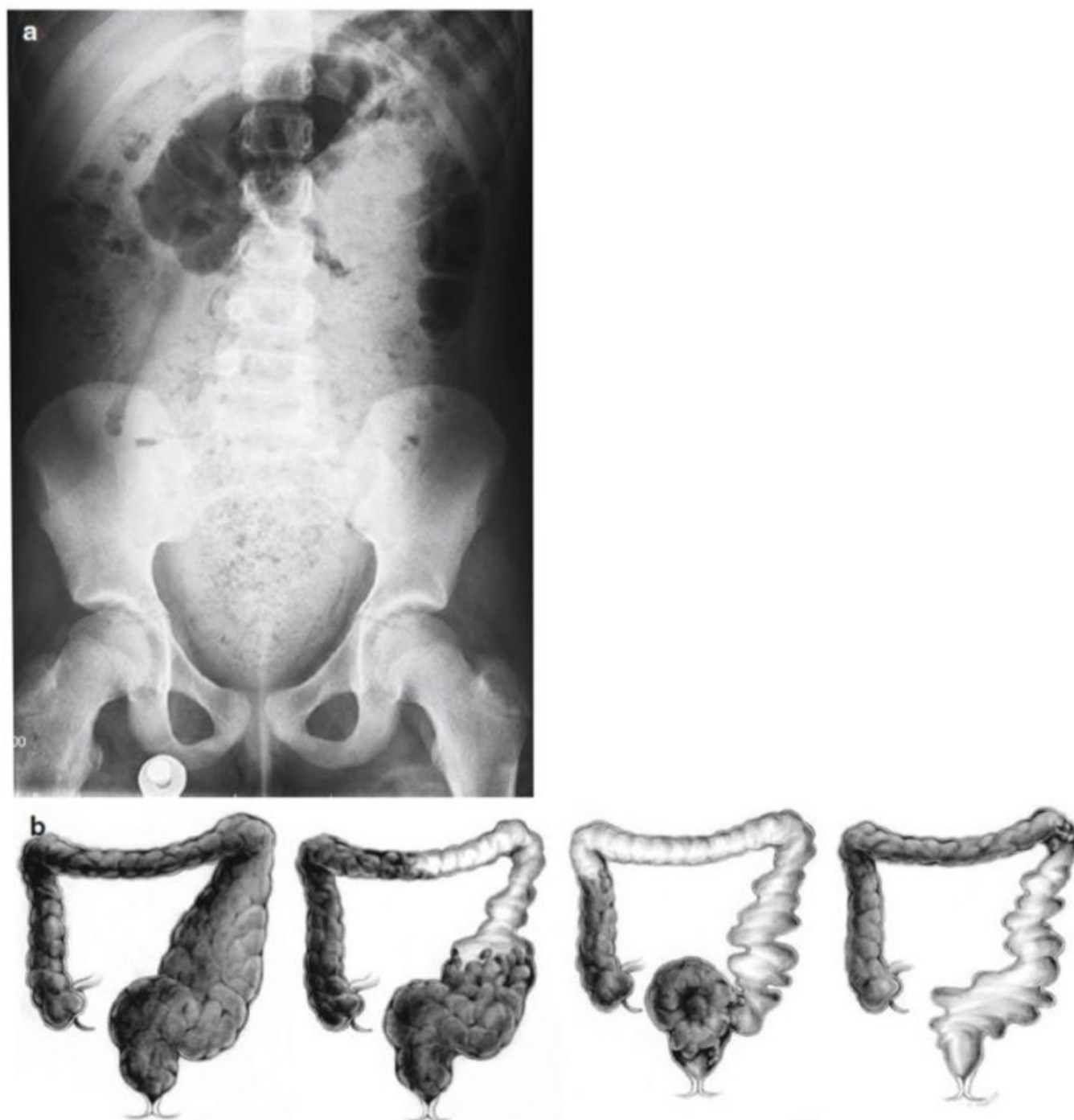


Fig. 25.7 Severe fecal impaction in a patient with idiopathic constipation. (a) Abdominal film. (b) Sequence of events during disimpaction

25.7.1 Fecal Disimpaction Protocol (Animation 20.4)

Our routine includes the administration of three enemas per day. The first enema is administered in the morning and includes the use of saline solution alone; the second enema (in the middle of the day) includes saline solution plus phosphate, and the third enema (before bed) includes saline solution plus glycerin. The volume of the enema depends on the size of the patient and the

degree of colonic dilatation and usually varies from 250 to 1,000 mL of saline solution. The phosphate (Fleet¹ enema) solution amount varies from half pediatric Fleet to 1 adult Fleet depending on the patient's age, and the amount of glycerin also varies from 5 to 30 mL of glycerin, depending on the patient's age. Every day the

¹Fleet – monobasic sodium phosphate 19 g, dibasic sodium phosphate 7 g. C.B. Fleet Company, Inc. Lynchburg, VA, USA.

patient gets an x-ray film of the abdomen, and most of the time, by the second day, the patient is disimpacted. If the patient is not disimpacted after 3 days of this treatment, then we offer to bring the patient to the hospital, continue with the same three enemas per day, and in addition introduce a nasogastric tube and administer GoLYTELY at a rate of 25 mL/kg per hour until disimpaction. In the event in which the patient stays 48 h receiving GoLYTELY and enemas and is still impacted, we offer them disimpaction under anesthesia. It is very unusual for us to do fecal disimpactions under anesthesia because most patients respond to our protocol. Thirty-two percent of our patients required the implementation of this protocol of disimpaction.

Once the patient has been disimpacted as demonstrated by an x-ray film of the abdomen, we start the second part of the management which is the determination of the laxative requirement.

25.7.2 Determination of Laxative Requirements

We prescribe an arbitrary amount of laxative (usually senna) that we think is going to work based on our impression of the degree of dilatation of the colon and patient's age, and watch the patient for the next 24 h. If the patient does not have a bowel movement, it means that the amount of laxative is not sufficient. The amount of laxative is then increased, but we also administer an enema to remove the stool that was produced during the previous 24 h. The basic rule is that the stool, in these extremely constipated patients, should never remain in the rectosigmoid more than 24 h because if it stays there, it will become hard and be more difficult to expel in the following days. We continue the routine, increasing the amount of laxatives and administering an enema every night, until we achieve our goal, which is to produce bowel movements and empty the colon completely as radiologically demonstrated (Fig. 25.8). At that point, the overwhelming majority of the patients stop soiling. The patients and the parents, for the first time, learn what the real individual laxative dosage for the patient is,



Fig. 25.8 Abdominal x-ray film showing a completely clean colon once the right amount of laxatives has been reached

which, as we previously mentioned, represents a measurement of the magnitude of the condition. Because we are dealing with a spectrum type of disease, we find patients with laxative requirements much larger than one might expect. In our series, for school-age children, the average amount of senna that allowed them to empty their colon was 69 mg/day with a range of 15–180 mg. The average dosage of senna that these patients were receiving prior to coming to our center was 33 mg/day. The most popular medication prescribed for our patients, prior to treatment, was polyethylene glycol.

Occasionally, in the process of increasing the amount of laxatives, we find patients who vomit before reaching any positive effect. In these patients we may try a different medication to see whether it is better tolerated, and some patients vomit all kinds of laxative, feel very sick, and have severe cramps, and we never reach the

amount of laxative capable of producing a bowel movement that empties the colon. Those patients are considered medically intractable and therefore candidates for surgical intervention. Approximately 85 % of the time, however, we find the dosage that the patient needs to empty the colon completely, as demonstrated radiologically. Once we have reached that amount, we expect the patient to stop soiling.

Patients that soil the underwear after reaching their laxative requirement, with a radiologically demonstrated empty colon, are very rare. As we previously mentioned, they deserve a more detailed study to rule out unusual neurologic or psychiatric conditions.

Sometimes, the required amount of senna necessary to empty the colon produces diarrhea. At that point, the parents know that they not only reached the necessary amount of laxative to empty the colon, but also perhaps they went a little too far in the amount of laxative, and now they are allowed to reduce a little bit of the amount of laxative that they were giving. In addition, once the patient is responding to the administration of laxatives, it is no longer necessary to give an enema. At this point, the parents, as well as us, have an idea of the magnitude of the problem. We usually use senna derivatives because we found that it is a laxative that has a more controlled effect. Once we have reached the decided amount, the parents are given two choices:

1. To continue with that amount of laxative for an undetermined period of time, perhaps for life, and most probably will still have to increase the laxative as time goes by.
2. The other alternative is to consider the possibility of an operation that may not cure the condition, but at least will help to decrease significantly the amount of laxative that the patient needs.

Confronted with these alternatives, the parents frequently express their concern about the long-term, secondary effects of the administration of laxatives, particularly senna. We share with the parents our concern about the use of this kind of medication, but unfortunately, so far, we do not have a better alternative. They may con-

sider, rather than giving a single type of laxative, to mix different types, trying to achieve the same effect.

In 84 % of our patients, this protocol was successful, which meant we reached the amount of laxative necessary to avoid soiling, keep the colon radiologically clean, and make the patients and families happy. Sixteen percent of our patients did not have a good result; they showed signs of intolerance to the laxative, as previously described; and we were unable to clean their colon.

We also explain to the family that there is no scientific evidence of senna derivatives producing cancer [106–110].

25.7.3 Electric Stimulation

Recently, a new modality of treatment is becoming popular [94, 111]. This has been tried in adults with encouraging results, and some enthusiastic doctors are trying it in children. We have no experience with that methodology. Since the mechanism of action has not been explained, we remain skeptical about its use.

25.8 Surgical Treatment

The most common operations offered to these patients are:

- Continent appendicostomy, cecostomy, or other kinds of procedures for the antegrade administration of enemas [89, 112–114]
- Colonic resections [115–118]
- Total proctocolectomy [119–121]
- Combination of resection plus an operation to administer antegrade enemas [122]

25.8.1 Operations to Administer Antegrade Enemas (ACE Procedures)

This type of procedure is perhaps the most common operation performed nowadays for the treatment of what the authors call unremitting or

intractable constipation. It is done with different modalities. The original operation, called Malone procedure, consisted of using the cecal appendix and connecting it to the skin of the abdominal wall in order for the patient to administer enemas in an antegrade fashion. A plication of the cecum around the appendix is frequently done with the goal to create a one-way valve mechanism to allow the passing of a feeding tube and to prevent fecal soiling through the orifice created in the abdominal wall. This original technique has been modified in an attempt to make it less invasive, and nowadays many doctors preferred to use button cecostomies that are very attractive because they are performed with a minimally invasive technology; however, they have the inconvenience of leaving a foreign body (button) in place. Many surgeons are extremely enthusiastic about the use of this procedure [89, 114]. However, there are also reports of rather poor results with the use of this type of procedures [123, 124].

We believe that ACE procedures or operations designed for the administration of antegrade enemas only represent a different route of administration of enemas. We have two concerns about this type of operations when performed in patients suffering from idiopathic constipation. The first one is that authors say that they perform those operations in cases of “unremitting constipation or intractable constipation,” but they do not define what exactly they mean by that. This means that some of those patients, conceivably, receive an operation, and actually they could have been treated medically and avoided the procedure. In addition, the logical way to proceed would be to try medical management following the guidelines that we expose, and if the patient does not tolerate that, we should try enemas, and once we demonstrate that the enemas are working, meaning that they are capable of keeping the colon completely clean every day as radiologically demonstrated, it would be a choice for the patient or the family to decide whether the enema should be given from below or from one of these orifices in the abdomen (see Chap. 21 on “Operations to Administer Enemas”).

25.8.2 Colonic Resection

This type of procedure has been used through the years and is still used at the present time [115–118]. The results of these operations, as expected in dealing with a spectrum type of condition, are variable, some surgeons being very enthusiastic and others rather skeptical. Our early experience included 237 patients medically treated for constipation following our guidelines; 70 of them underwent a resection of the most dilated part of the sigmoid (Fig. 25.9). Some of these patients received the operation because they were really nonresponsive to the medical treatment. In other words, they became very sick in the process of increasing the amount of laxatives, and they did not pass stool, or in other cases, the parents elected to have the procedure in an attempt to decrease the laxative requirement. Approximately 10 % of the patients did not require any more laxatives after the procedure and had bowel movements every day with no soiling; 30 % of the patients decreased the laxative requirement by 80 %. The remaining 60 % of the patients decreased the laxative requirement by 40 %. The resection included the most dilated part of the sigmoid but respected the entire rectum. In other words, the colon was resected at the level of the peritoneal reflection, and a piece of non-dilated descending colon was anastomosed to the rectum [118].

As previously mentioned, these operations do not cure the patients but rather simply improve them.

In an attempt to find a more radical cure for this condition, total proctocolectomy has been performed [119–121]. Obviously, this kind of operation would have more chances to succeed in eradicating the problem of constipation; however, the price that the patient has to pay for this is very high because the patient will have the problem of liquid stool for life and many bowel movements.

Another more conservative approach is the resection of the rectosigmoid including the most dilated part of the colon (not a total colectomy), performing the operation via transanal like in Hirschsprung's disease (Fig. 25.10). We have

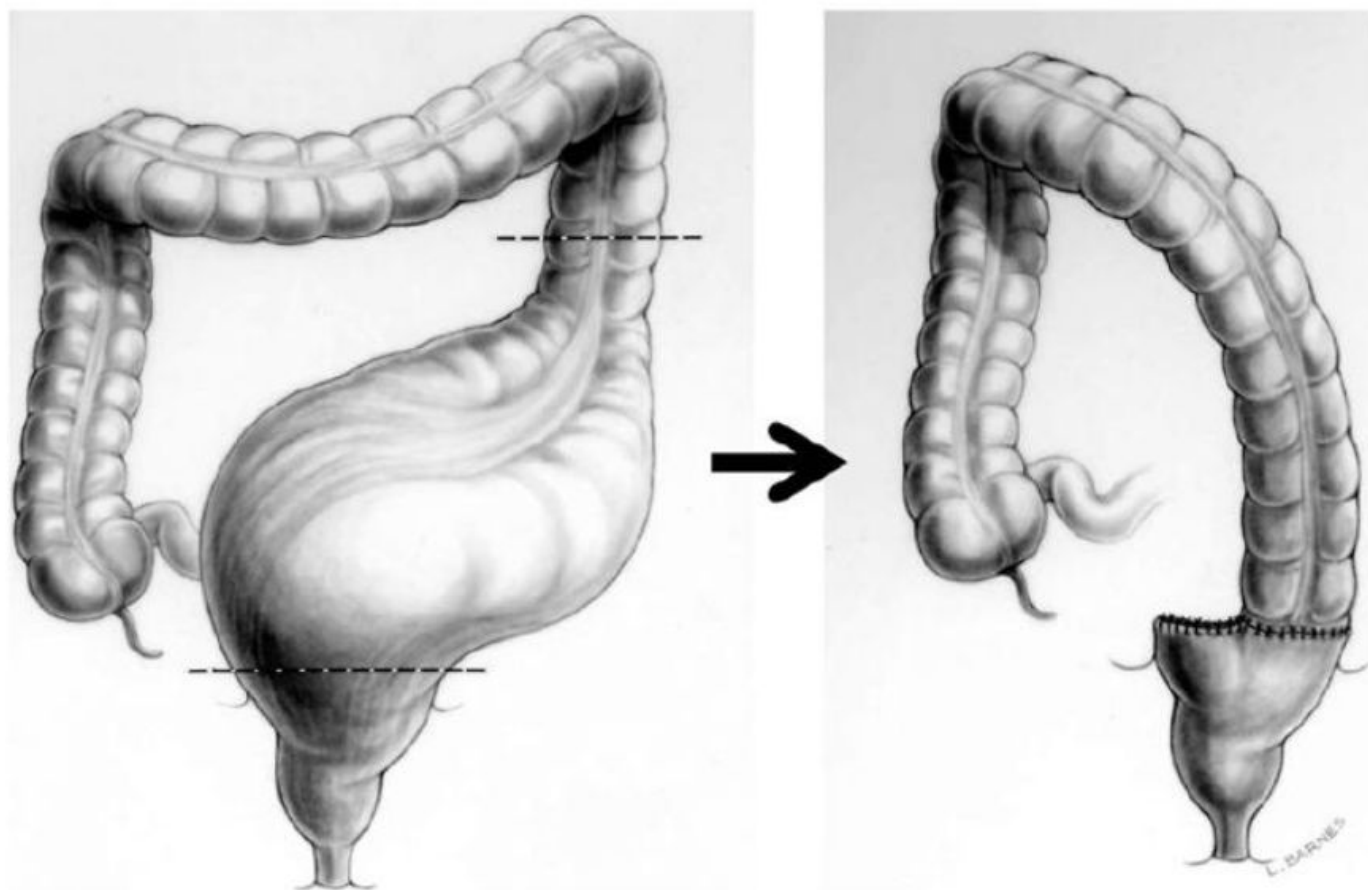


Fig. 25.9 Diagram showing a sigmoid resection. The most dilated part of the sigmoid is resected, and an anastomosis is performed between the normal caliber descending colon

and the rectum at the level of the peritoneal reflection. The (*dashed lines*) in the diagram on the left, show the limits of the resection. The (*arrow*) shows the finished operation



Fig. 25.10 Intraoperative picture of resection of a giant megarectosigmoid

done this in more severe types of constipation. However, the main concern in those cases is the possibility of a patient suffering from a mild degree of fecal incontinence. If this procedure is done, it should be performed in a very meticulous way, being sure to preserve the integrity of the anal canal, performing the anastomosis 2 cm

above the pectinate line and being sure not to stretch too much the anal opening during the procedure.

In summary, patients suffering from idiopathic constipation are clinically diagnosed at our center followed by a contrast enema; if both are characteristic of idiopathic constipation, we do not performed manometric studies, and we do not feel the need to take unnecessary rectal biopsies. The patients then are disimpacted when necessary, and the next step consists of finding the amount of laxative necessary to empty the colon, radiologically demonstrated, which represents a measure of the severity of the problem. For those patients (10–15 %) that demonstrate to be truly resistant to the medical management, we offer them an alternative of enemas given through the rectum or through a Malone or a conservative resection of the most dilated part of the colon. If this is not enough, we may consider the possibility of a more radical transanal resection of the most dilated part of the colon.

References

- Bartlett J (ed) (1919) Familiar quotations, 10th edn. Little, Brown, Boston, p 627
- Bellman M (1966) Studies on encopresis. *Acta Paediatr Scand* 170(Suppl):1-15
- Levine MD (1975) Children with encopresis: a descriptive analysis. *Pediatrics* 56:412-416
- Taitz LS, Water JKH, Urwin OM, Molnar D (1986) Factors associated with outcome in management of defecation disorders. *Arch Dis Child* 61:472-477
- Sandler RS, Jordan MC, Shelton BJ (1990) Demographic and dietary determinants of constipation in the US population. *Am J Public Health* 80:185-189
- Taylor R (1990) Management of constipation. *Br Med J* 300:1063-1064
- Pinkerton P (1957) Pathogenic megacolon in children: the implications of bowel negativism. *Arch Dis Child* 33:371-380
- Berg I, Jones KV (1964) Functional fecal incontinence in children. *Arch Dis Child* 39:465-472
- Gabel S, Hegedus AM, Wald A, Chandra R, Chiponis D (1986) Prevalence of behavior problems and mental health utilization among encopretic children: implications for behavioral pediatrics. *J Dev Behav Pediatr* 7:293-297
- MacNamara M (1965) Notes on a case of lifelong encopresis. *Br J Med Psychol* 38:333-338
- Shane M (1967) Encopresis in a latency boy: an arrest along a developmental line. *Psychoanal Study Child* 22:296-314
- Silber S (1968) Encopresis: rectal rebellion and anal anarchy. *J Am Soc Psychosom Dent Med* 15:97-106
- Holschneider AM, Puri P (2000) Anal sphincter achalasia and ultrashort Hirschsprung's disease. In: Holschneider AM, Puri P (eds) *Hirschsprung's disease and allied disorders*, 2nd edn. Harwood Academic Publishers, The Netherlands, pp 399-424
- Fadda B, Welskop J, Muntefering H, Meier-Ruge W, Engert J (1987) Achalasia of the anal sphincter. *Pediatr Surg Int* 2:81-85
- Hecker WC, Holschneider A, Fendel H, Schauer A, Meister P, Beige H (1973) Die chronische obstipation beim kind durch anal sphinkterachalasie. *Dtsch Med Wochenschr* 98:2334-2340
- Oue T, Puri P (1999) Altered intramuscular innervation and synapse formation in internal sphincter achalasia. *Pediatr Surg Int* 15(3-4):192-194
- Holschneider AM, Puri P (2000) Diagnosis of Hirschsprung's disease and allied disorders. In: Holschneider AM, Puri P (eds) *Hirschsprung's disease and allied disorders*, 2nd edn. Harwood Academic Publishers, The Netherlands, pp 230-245
- Meunier P, Marechal JM, DeBeaujeu MJ (1979) Rectoanal pressures and rectal sensitivity studies in chronic childhood constipation. *Gastroenterology* 77:330-336
- Loening-Baucke VA (1984) Abnormal anorectal function in children recovered from chronic constipation and encopresis. *Gastroenterology* 87:1299-1304
- Keren S, Wagner Y, Heldenberg D, Golan M (1988) Studies of manometric abnormalities of the rectoanal region during defecation in constipated and soiling children: modification through biofeedback therapy. *Am J Gastroenterol* 83:827-831
- Corazziari E, Cucchiara S, Staiano A et al (1985) Gastrointestinal transit time, frequency of defecation, and anorectal manometry in healthy and constipated children. *J Pediatr* 106:379-382
- Lynn HB, van Heerden JA (1975) Rectal myectomy in Hirschsprung's disease: a decade of experience. *Arch Surg* 110:991-994
- Bourdela D, Barbet JP, Gross PH (1994-1995) Constipation of the child: usefulness of anorectal sphincterectomy. *Chirurgie (memoires de l'Academie)* 120:48-52
- Ecaluwe D, Yoneda A, Akl U, Puri P (2001) Internal anal sphincter achalasia: outcome after internal sphincter myectomy. *J Pediatr Surg* 36:736-738
- Pinho M, Yoshioka K, Keighley MRB (1989) Long term results of anorectal myectomy for chronic constipation. *Br J Surg* 7:1163-1164
- Ure BM, Holschneider AM, Schulten D, Meier-Ruge W (1999) Intestinal transit time in children with intestinal neuronal malformations mimicking Hirschsprung's disease. *Eur J Pediatr Surg* 9:91-95
- Stoss F (1990) Neuronal dysplasia: considerations for the pathogenesis and treatment of primary chronic constipation in adults. *Int J Colorectal Dis* 5:106-112
- Fadda B, Maier W, Meier-Ruge W, Scharli A, Daum R (1983) Neuronale Intestinale Displasie: eine kritische 10 Jahre analyse klinischer und biotischer Diagnostik. *Z Kinderchir* 38:305-311
- Dickson JAS, Variend S (1983) Colonic neuronal dysplasia. *Acta Paediatr Scand* 72:635-637
- Scharli AF, Meier-Ruge W (1981) Localized and disseminated forms of neuronal intestinal dysplasia mimicking Hirschsprung's disease. *J Pediatr Surg* 16:164-170
- Pistor G, vonKap-Herr SH, Grussner R, Munakata H, Munterfertigo H (1987) Neuronal intestinal dysplasia: modern diagnosis and therapy: report of 23 patients. *Pediatr Surg Int* 87:352-358
- Csury L, Peña A (1995) Intestinal neuronal dysplasia: myth or reality? *Pediatr Surg Int* 10:441-446
- Schofield DE, Yunis EJ (1991) Intestinal neuronal dysplasia. *J Pediatr Gastroenterol Nutr* 12(2):182-189
- Kubota A, Yamauchi K, Yonekura T, Kosumi T, Oyanagi H, Mushiaki S, Nakayama M, Imura K, Okada A (2001) Clinicopathologic relationship of hypoganglionosis. *J Pediatr Surg* 36(6):898-900
- Scharli AF, Sossai R (1998) Hypoganglionosis. *Semin Pediatr Surg* 7(3):187-191
- Munakata K, Okabe I, Morita K (1992) Hypoganglionosis. *Pediatr Surg Int* 7:8-11
- Meier-Ruge WA, Brunner LA (2001) Morphometric assessment of Hirschsprung's disease: associated