Radiological diagnosis, etiology and pathophysiology of encopresis in children. Review.

Abstract The term "encopresis," in analogy with the term "enuresis," was introduced by Weisenberg in 1926 to designate those forms of fecal continence disorder in which there is no organic damage to the elements involved in fecal retention. Most often, encopresis occurs in children with chronic constipation. However, the pathophysiology and pathogenesis of this phenomenon are not well understood. Less commonly observed is non-retentive fecal incontinence, the etiology and pathogenesis of which are unknown. It is not accompanied by fecal retention in the rectum, and therefore there is no constipation. This article is based on a review of our published studies using radiometric analysis, from which we came to the following conclusions. In children with chronic constipation large amounts of feces accumulate in the rectum due to delayed bowel movements. In the dilated rectum (megacolon - megarectum), large fecal stones form, which stretch the pelvic floor muscles. A stretch of the puborectalis muscle (PRM) weakens fecal continence, leading to stool leaking into panties. Various degrees of damage to the PRM function are shown up to the development of descending perineal syndrome. In children with encopresis without constipation, a decrease in the width of the rectum and left half of the colon was found. In most of them, proctoscopy revealed macroscopic signs of inflammation. We hypothesize that the inflammatory process in them leads to increased intestinal tone and rapid movement of the bolus into the rectum, and high rectal tone stimulates the defecation reflex with small volumes of feces. The data obtained allows the use of pathogenetic treatment.

Keywords: Encopresis; functional constipation; barium enema, nonretentive fecal incontinence; pathophysiology.

Introduction The term "encopresis," by analogy with the term "enuresis," was first introduced by Weisenberg in 1926 to designate those forms of continence disorder in which there is no organic damage to the elements involved in fecal retention [quote from 1]. In childhood, the term fecal incontinence is most often used, which includes both organic damage and functional disorder [2,3]. Among modern pediatric doctors, the idea is firmly established that: - "The pathophysiology of functional fecal incontinence related to constipation in children is not clear" [4,5]. In 10% of patients present encopresis as a single symptom without any organic cause or sign of constipation and is currently classified as functional nonretentive fecal incontinence (FNRFI) and neither etiology nor pathogenesis of it is known [6, 7]. This means that there is no pathogenetic treatment, and a trial-and-error search is used.

The purpose of this article is to show the effectiveness and reliability of radiometric analysis of a contrast enema in the diagnosis of various forms of encopresis and its role in determining the cause of encopresis in both groups.

Material This review is devoted to the radiometric analysis of children with encopresis who were treated at the children's surgical center in Minsk (Belarus). Research materials were published in 1983-2020. From this study were excluded patients after surgery on the anorectal area, because of which the muscles involved in fecal continens and defecation are damaged. Their fecal incontinence was of an organic nature. All 166 patients with encopresis were divided into two groups according to the clinical picture and the results of barium enema. In 127 (77%) patients of group 1, encopresis was combined with constipation, and a barium enema revealed megacolon of varying degrees. The 39 (23%) patients of group 2 had no history of constipation and no dilatation of the rectum according to barium enema [1,8,9, 10, 11].

Method. Radiometric analysis is based on a standardized barium enema technique and standards for the width of different parts of the colon and rectum, as well as the length of the anal canal in children of different ages (control), which were determined using this technique [12].

The study was conducted by the hydrostatic barium enema. Barium was introduced from the graduated bag. The bottom of this bag at the beginning of the study was located 40 cm above the deck of the table. Barium was introduced into the colon up to the reflux into the terminal ileum. The difference of barium volume in the bag before and after the colon filling

corresponds to the colon volume. The radiopaque marker is strung on the tip of the enema. It is in contact with the anus during the study. At least two radiographs (frontal and lateral) have been made after the filling of the colon. On a frontal radiograph the widths of the different parts of the colon are measured (Figure 1 a). Since the rectum forms two bends in two projections, it cannot be differentiated on the frontal radiograph, which makes it impossible to measure its width. On a lateral radiograph were measured the maximum width of the vertical portion of the rectum, as well as interval not containing contrast medium between the rectum and a contrast marker on the posterior contour of the tip of the enema (Figure 1b). This distance is due to the contraction of the anal canal and is equal to the length of the anal canal measured during a manometric study.



Figure 1. (a). The frontal radiograph shows the measurement sites: c - caecum, a - ascending colon, t - transverse colon, d - descending colon, s - sigmoid colon. **(b).** On a lateral radiograph, the measurement location for the descending branch of the rectum is shown as (R). The length of the anal canal is measured between the rectum and a radiopaque marker located near the anus.

Filling of the colon occurs under low hydrostatic pressure, resulting in slow colonic filling that is easily tolerated by children of all ages. Therefore, in children with encopresis there was no case of contrast agent leakage, and thus there was no need for an inflated of the rectal balloon. This made it possible to measure the length of the anal canal.

Method of the analysis of the radiographs

In children without anorectal pathology, the distance between the marker (anus) and the rectum, which does not contain a contrast agent, is equal to the anal canal length measured by

the manometric method [12]. This is a zone of the anal canal contraction. The true values were obtained by multiplying of the size measured on radiographs, on a factor of projection magnification (k), which is equal to the ratio of the true diameter of the marker to its value on the radiograph. To compare the different studies as well as studies of the same patient at different ages, we calculate the constant (C), which is the integral characteristic of the colon value. It is calculated using the following formula:

 $\mathbf{V} \times \mathbf{R} \times \mathbf{\kappa}$ $\mathbf{C} = -----$

Where: \mathbf{C} – constant, \mathbf{V} – colon volume (ml).

 \mathbf{R} – rectal width (cm).

 κ – projection distortion factor, which is the ratio of the true width of the marker to its image on the radiograph;

h – patient height (cm).

In healthy children, "Constant" was in the range of 17-31, regardless of age. Megacolon determined if «Constant», exceeds 31. It has been possible to differentiate megacolon varying degrees depending on the constant (C):1st degree - (C = 32 - 45); 2nd degree - (C = 45-60) and 3rd degree - C > 60.

Since the sigmoid colon over the age of 5 years is normally located in the pelvis, its location in the abdominal cavity always indicates its elongation. The lengthening of the sigmoid colon is due to the presence of the mesentery. Since it is not fixed to the wall of the abdominal cavity, it lengthens simultaneously with expansion. Thus, elongation of the sigmoid colon is evidence of overfilling of the colon with feces during megacolon, or in the past, after which the volume of the intestine has decreased because of the inflammatory process. The transverse colon also has a mesentery and can elongate and sag into the pelvis in very severe forms of megacolon in adults. Filling the colon prior to barium reflux into the ileum has proven to be very useful in determining the etiology and pathogenesis of functional megacolon (FM). Since the width of the rectum does not change after filling the splenic angle, to determine the megarectum it is enough to fill the colon to the splenic angle. **Table 1** shows the normal true dimensions of the rectum and anal canal, which in practice are sufficient to judge the megarectum and the condition of the pelvic floor muscles, including puborectalis muscle (PRM).

Age	n	Rectal width (cm)	n	Anal canal length (cm)
5 days-11 months	12	1,3-3.0 (2.24±0.09)	7	1.7-2.5 (2.21±0.15)
1-3 years	9	3.0-3.7 (3.21±0.11)	7	2.3-2.8 (2.55±0.10)
4-7 years	9	3.0-3.9 (3.43±0.14)	8	2.3-3.6 (3.11±0.10)
8 – 10 years	9	3.2-4.1(3.70±0.06)	8	2.6-3.7 (3.07±0.11)
11 – 15 years	19	3.6-4.6 (39.5±0.07)	18	3.1-3.9 (3.43±0.10)

Table 2. The true rectal width and anal canal length at different ages.

The statistical study was conducted using Student's t-test. Statistical significance was determined at p < 0.05.

Encopresis in children with functional megacolon (1st group).

It appears that the cause of chronic constipation (CC) in early childhood has different origins. Very rarely CC begins in the first year of life, most often after cessation of breastfeeding, when the stool becomes formed. As shown in the studies of Duhamel [13] and Clayden with Lawson [14], in some children with severe megacolon, the cause of constipation is minor forms of anorectal malformations, such as anterior anus, anal or rectal stenosis with a relatively wide, but the not normal of the hole width. Clayden and Lawson found secondary megacolon in 5% of children who were initially diagnosed with functional megacolon [15]. As shown in Figure 2 B, FM begins after one year of age during potty training, with a peak between 3 and 8 years of age during social adaptation to the group. Encopresis appears 1–5 years after the onset of constipation. Figure 2 C shows that the more severe the degree of megacolon, the more likely it is to be accompanied by encopresis [8].



Figure 2. A). Graphic representation of the frequency of megacolon depending on the time of occurrence of constipation - (a), and the time of going to the surgical hospital (b). B). Graphical representation of megacolon frequency depending on the time of onset of constipation- (a), and the time of onset of encopresis - (b). (C). The relationship between

numbers of patients with encopresis - (a), and without encopresis - (b) depending on the degree of megacolon.

Analysis of the above graphs shows that encopresis mainly occurs over the age of 3 years, sometime after the onset of constipation, because of the progression of megacolon. The greater the degree of megacolon, the more likely it is that the disease will be complicated by encopresis. Because of the dilation of the rectum, its anterior wall is pushed forward, because of which the lower horizontal branch of the rectum is absent (Figure 3 b-e). This figure also shows the different degrees of damage to the pelvic floor muscles in sequential order.



Figure 3. Lateral radiographs of the anorectal area. (a) In normal two branches of the rectum are identified: vertical (v) and horizontal (h). The length of the anal canal is measured from the anorectal angle (arrow) to the radiopaque marker located near the anus. In the upper part of the anal canal in front of the enema tip, penetration of barium is determined, which is due to relaxation of the internal anal sphincter (IAS). The posterior wall of the anal canal at this level is pressed against the tip of the enema by the contracted PRM. The lower part of the anal canal is closed by contraction of the external anal sphincter (EAS). This is the radiological manifestation of the rectoanal inhibitory reflex [15]. (b-c). Dilatation of the rectum is accompanied by the disappearance of the horizontal branch and the appearance of the PRM. (d-e). The expansion of the rectum is accompanied by a sharp shortening of the anal canal because of weakness (d) or incompetence (e) PRM.

Eighty-three patients (65%) had grade 1–2 megacolon, with the length of the anal canal falling within the age-appropriate range. Among them, encopresis was not observed. Various degrees of PRM weakness were identified by the appearance of barium in the anorectal angle (Figure 3c). In 18 patients, barium leakage into the upper part of the anal canal, behind the enema tip, was evident, with the length of the anal canal near the lower limit of normal

(Figure 3d). In 26 cases, a nearly two-fold reduction in the length of the anal canal compared to the norm was detected (Figure 3e). Encopresis in patients with FM was always associated with varying degrees of PRM weakness. A significant shortening of the anal canal is due to PRM and levator plates insufficiency, a condition referred to in the literature as descending perineal syndrome (DPS) [16, 17]. **Figure 4** illustrates the mechanism of DPS development and our proposed diagnostic method.



Figure 4. A 15-year-old patient with FM and encopresis. The true diameter of the X-ray contrast marker located near the anus is 1.6 cm. The width of the fecal stone is 9 cm is 2 times greater than the maximum normal limit for the width of the rectum (4.6 cm). A peristaltic wave starting from the rectosigmoid sphincter (red arrows) pushes the stone under great pressure. However, the width of the anal canal does not allow a stone of this width to pass through. The distance between the pubococcygeal line and marker is 4 cm and it is equal to the length of the normal anal canal. However, the bougie effect of the stone led to stretching of the pelvic floor muscles (PRM and levator plates). As a result of the weakness of the PRM, only the lower part of the anal canal functions to retain feces. Therefore, liquid barium under pressure passes along the enema tip outward (shown by white lines), which manifests itself as functional fecal incontinence (encopresis).

In adults, descending perineal syndrome (DPS) is defined as the descent of the contrastenhanced rectum several centimeters below the pubococcygeal line during defecation. This condition is typically assessed using defecography [17] or MRI defecography [18], performed while the patient defecates on a radiolucent potty. However, this method has two major drawbacks: first, it exposes patients to long-term ionizing radiation, and second, determining bone landmarks for drawing the pubococcygeal line can be challenging, complicating the evaluation. Due to these factors, defecography is not used to diagnose DPS in children. Our method allows us to diagnose DPS as the shortening of the anal canal from the marker near the anus to the barium in the intestine during a conventional barium enema.

An analysis of the literature and our own research indicates that in patients with FM, encopresis occurs because of rectal overflow with feces, a condition invariably associated with megacolon. Large fecal stones stretch the pelvic floor muscles, leading to the weakening of their function. Without pathogenetic treatment, this condition progresses from minimal damage to DPS, which in turn causes encopresis. DPS represents irreversible damage, and its treatment typically involves either puborectalis muscle (PRM) plication or transposition of the gracilis muscle. Symptomatic treatments, such as bowel management or antegrade enemas for DPS, often complicate the lives of patients without addressing the underlying problem. It should also be noted that stretching of the levator plates impairs the function of opening the anal canal during defecation attempts [15, 19, 20, 21], significantly increasing resistance to fecal passage through the anal canal. This resistance is one of the key factors contributing to the chronicity of FM.

Encopresis in children without constipation (2nd group).

In group 2, consisting of 39 patients with a mean age of 9.2 ± 2.2 years, 37 patients (95%) experienced periodic non-localized abdominal pain in addition to fecal incontinence. The group included eight times more boys than girls. In 20 patients (51%), encopresis was also accompanied by enuresis. Using the previously described X-ray method, only 5 patients (13%) had rectal widths within the age-appropriate range. In the remaining cases, the width of the rectum and sigmoid colon was below the minimum age norm. In 26 patients (67%), the descending colon was narrower than normal, with asymmetric haustration. These findings indicated a high tone in the left side of the colon and led to the recommendation for proctoscopy. Macroscopic signs of inflammation were observed during proctoscopy in 29 patients (74%).

An analysis of the literature shows that the differentiation between encopresis in fecal incontinence (FM) and nonretentive encopresis is based solely on the absence of a history of constipation. There is no single diagnostic method with proven accuracy for this distinction. Since the underlying cause of this pathology remains unknown, current treatment approaches

cannot be considered pathogenetic. In fact, none of the available methods have demonstrated significant breakthroughs. For example, the guidelines approved with "high consensus" include only one clear recommendation: "Laxatives can worsen outcomes and should be avoided" [22]. Even with biofeedback or laxative treatment, only 29% of patients achieved successful outcomes after two years of intensive therapy. Despite improvement in most patients after puberty, by age 18, 15% still experienced fecal incontinence [23].

Moreover, the temporary use of additional rectal enemas did not significantly enhance treatment success compared with conventional therapy alone [24], which consists of patient education, a non-accusatory approach, and a toileting program that includes a daily bowel diary and a reward system. Special attention is recommended for addressing psychosocial and behavioral problems, as these issues are common in affected children [25]. Jørgensen et al. found that transanal irrigation showed some effectiveness in treating encopresis in a small number of patients [26]. Positive outcomes were also noted with transcutaneous functional electrical stimulation [27] and transcutaneous posterior tibial nerve stimulation [7].Our data show that encopresis without constipation is clinically characterized by periodic nonlocalized abdominal pain, an acute need to defecate and the inability to prevent defecation, as well as a frequent combination with enuresis, and a significant predominance of boys. Trial-and-error treatment results in "significant (p <0.05)" symptom relief in some patients according to the results of their questionnaire. However, most often the symptoms do not disappear completely, and in some cases, they reappear after some time. It is known that laxatives can worsen symptoms.

Our X-ray analysis revealed that 13% of patients had rectal widths within the normal range, while in 87%, the rectal width and the left half of the colon were below the minimum age norm, indicating increased tone in these sections. Macroscopic signs of inflammation were observed in the rectum and/or sigmoid colon in 74% of cases. The combination of clinical and radiological findings helps clarify the pathophysiological process. Inflammation increases the tone of the left colon and rectum, causing the stool to move rapidly into the rectum. Since defecation pressure is triggered at a lower bolus volume in a narrowed rectum [15], a rapidly administered bolus triggers a defecation reflex, like what occurs in acute diarrhea [28].

This understanding of the pathophysiology of encopresis without constipation explains why laxatives, which further increase already high intestinal tone, exacerbate encopresis symptoms. In contrast, loperamide has a positive effect by slowing intestinal motility and increasing anal canal tone [29]. Based on these findings, there is reason to believe that anti-inflammatory treatments, delivered orally or via transanal irrigation, could be effective for children with encopresis without constipation.

Conclusion

Encopresis in children, defined as fecal incontinence of a functional nature, is divided into two distinct conditions with different origins. In most cases, it occurs in children with chronic constipation, where delayed defecation leads to the accumulation of large fecal masses in the rectum. This results in a dilated rectum (megacolon or megarectum), where fecal stones form which stretch the pelvic floor muscles. The stretching of the puborectalis muscle (PRM) weakens its ability to maintain fecal continence, causing fecal leakage into the underwear. Varying degrees of PRM dysfunction, preceding the development of descending perineal syndrome (DPS), have been observed. In children without constipation, a narrowing of the rectum and the left side of the colon has been noted. Proctoscopy in most of these cases revealed macroscopic signs of inflammation. The high tone of the intestines leads to the rapid advancement of the stool into the rectum, and increased rectal tone triggers the defecation reflex even with small volumes of feces. This process resembles acute diarrhea, where bowel movements occur quickly and uncontrollably. These findings provide the foundation for pathogenetic treatment approaches.

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