



Anorectal Malformation with Rectovestibular Fistula and Vaginal Agenesis: Usage of Rectovestibular Fistula as a Neovagina Followed by PSARP with Preservation of the Anal Sphincter

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Abstract

Keywords

- ▶ anorectal malformation
- ▶ vaginal agenesis
- ▶ posterior sagittal anorectoplasty
- ▶ vaginoplasty

Anorectal malformation with rectovestibular fistula associated with vaginal agenesis is rare. We report on a child in whom this combination was diagnosed at the age of 1 year. After creation of a divided descending colostomy, we chose to leave the rectum-rectovestibular fistula to function as a neovagina, while the sigmoid colon was relocated via modified posterior sagittal anorectoplasty. The colostomy was closed 6 months after the main surgery. After a follow-up of 3 years, the patient is continent for stool and urine. The introitus of the neovagina appears wide and easy to explore. We conclude that our surgical approach may be a good option for these children.

New Insights and the Importance for the Pediatric Surgeon

ARM is one of the most complex congenital anomalies involving several organs. The association of ARM with vaginal genesis, as it is described in our case, is a very rare and complex event to be corrected surgically.

Introduction

Anorectal malformations (ARMs) are rare congenital defects with an estimated worldwide incidence of 2 to 6 per 10,000 live births.¹ They comprise a wide range of pathologies that may involve other parts of the gastrointestinal tract as well as urinary and genital tracts.^{2–9} The most common type of ARM in female patients is a rectovestibular fistula. The incidence of vaginal agenesis associated with ARM is even lower.⁴ The approach to this combination was first described by Cohn and Murphy via a combined laparotomic and perineal operation.¹⁰ The use of rectovestibular fistula as a neovagina has also been reported in combination of a

posterior sagittal anorectoplasty (PSARP),² as described by Levitt et al.⁴ We describe the case of a 1-year-old girl with ARM and rectovestibular fistula in whom we chose to leave the rectum-rectovestibular fistula to function as a neovagina, while the sigmoid colon was relocated via modified PSARP, according to Liem and Hau, and Liem and Quynh technique.^{11,12}

Case Report

The previous term infant (birth weight 3.1 kg) was admitted to the Pediatric Surgery Service at Saint Damien Pediatric Hospital, Port-au-Prince, Haiti, with a diagnosis of vaginal

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agenesis associated to ARM with rectovestibular fistula. Until 10 months of age, the infant defecated through the rectovestibular fistula that required dilations even though it had a considerable caliber. Two months prior to clinical admission, she received a descending colostomy with distal mucous fistula. No other obvious malformations and a normal sacrum were found. Due to the lack of resources at the local hospital, no additional diagnostic imaging, for example, magnetic resonance imaging, was performed.

We chose to leave the rectovestibular fistula in situ to function as a neovagina and to perform a PSARP,^{2,4} modified at preserving the anal sphincter as reported by Liem and Hau, and Liem and Quynh.^{11,12} The patient was placed in a prone jackknife position. After locating the anal sphincter by using a neurostimulator, a posterior sagittal incision of the cutaneous and subcutaneous planes was made 1 to 2 cm over the coccyx to the superior limit area of the anal dimple. The dissection was deepened while paying attention to the midline, to avoid injuries to muscles and nerve fibers, and arrested at the level of the upper limit of the muscle complex. The muscle complex was identified and its lower part was retracted downward to expose the posterior wall of the rectum (**-Fig. 1**). The fibrotic bands between the rectal wall and the coccyx were divided and the coccyx removed aiming at gaining extra space. The rectum appeared considerably dilated to the extent that not all its circumferences could be isolated. As a consequence, it was isolated only from its lateral sides and the anterior surface of the sacrum until the peritoneal reflection which was opened in view of the subsequent laparotomy.

After having temporarily closed the posterior sagittal skin incision to minimize the risk of bacterial contamination, the patient was placed in supine position to open the abdomen through a Pfannenstiel incision. The rectum, which was abnormally dilated until 1 to 2 cm beyond the peritoneal reflection, continued as sigmoid colon whose caliber appeared suddenly normal (**-Fig. 2**). The rectouterine Douglas' pouch was explored; while ovaries were located with fallopian tubes presenting cordlike structures, instead no structures resembling vagina and uterus were found.



Fig. 1 The lower part of the muscle complex was retracted down to expose the posterior wall of the rectum.



Fig. 2 Anatomy of the patient.

The dilated rectum was surgically separated from the sigmoid colon and closed; the sigmoid colon was only temporarily sutured to avoid fecal contamination (**-Fig. 3**).

After binding and dissecting some branches of its vascular arcade, the sigmoid colon was then mobilized, as it needed to be easily passed without traction through the retrorectal space into the previously opened posterior sagittal space (**-Fig. 4**), which was reopened at this point.

Once the suprapubic incision was sutured and the patient was again in the same prone position as before, a second small inverted cutaneous Y-shape incision was made on the anal dimple, after reconfirming the location of the underlying anal muscle complex using the neurostimulator (**-Fig. 5**). Our technique was different from the original technique of PSARP described by Peña and Devries, which envisages to separate the sphincter muscle complex in the midline.² In contrast, we created a tunnel in the center of the sphincter muscle complex through a gentle and progressive dilation of its muscle fibers using Hegar dilators (size 6–12) (**-Fig. 6**)

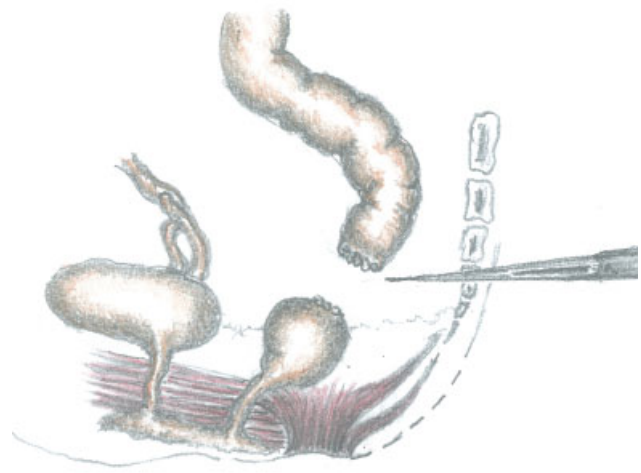


Fig. 3 Sigmoid colon surgically separated from the dilated rectum.

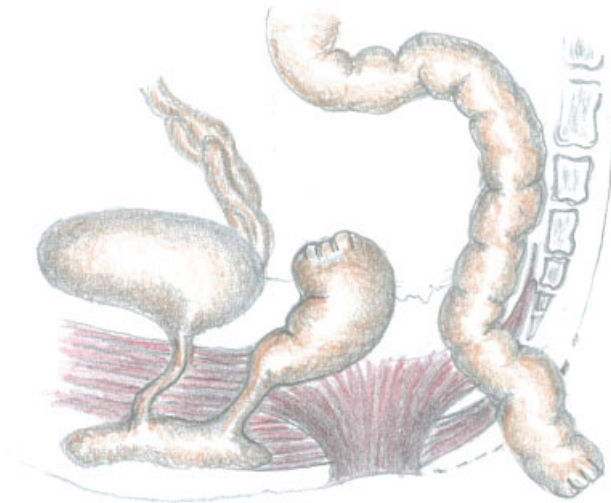


Fig. 4 Mobilized sigmoid colon passed into the posterior sagittal space.

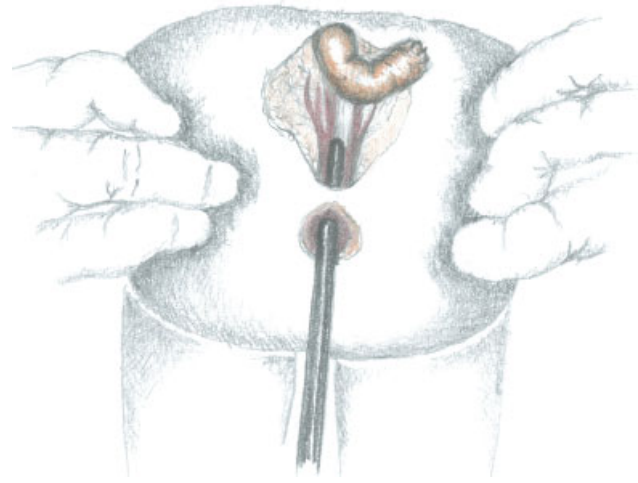


Fig. 6 Progressive dilation of the tunnel created in the center of the sphincter complex.

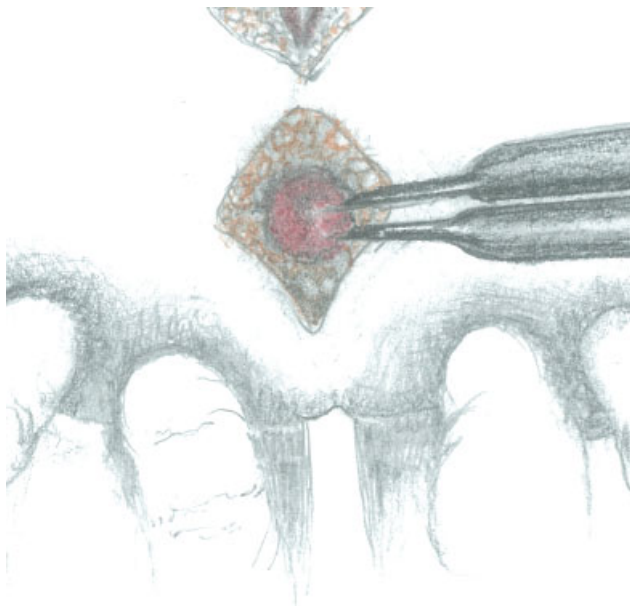


Fig. 5 The center of the sphincter complex is defined using a neurostimulator.

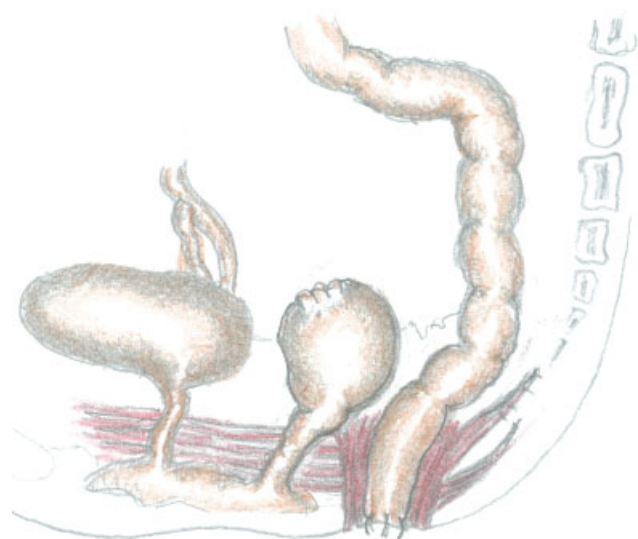


Fig. 7 The sigmoid colon is passed through the dilated tunnel of the sphincter complex and a neoanus is created.

until reaching an acceptable diameter to enable the mobilized sigmoid colon to go through it.

Consequently, the sigmoid colon, that managed to pass through the dilated tunnel of the sphincter muscle complex, was sutured to the external sphincter and to the margins of the small Y-shape cutaneous incision that had already been made in the anal dimple by using interrupted absorbable stitches (**Fig. 7**). The posterior sagittal incision was then closed. Postoperatively, the neoanus was gradually dilated using Hegar dilators from the second week after the surgery for a total period of 1 month. The colostomy was closed after 6 months.

No complications were observed during the early postoperative period. Neither anal stenosis nor other local and general complications were observed.

Two years after the anoplasty and at the age of 3 years, the parents reported that a few months earlier the girl stopped using diapers as she had achieved good bowel control and always evacuated spontaneously. So, no complications occurred such as fecal incontinence (not even partially), no constipation or mucosal prolapse, nor urinary incontinence. The rectovestibular fistula, that had become the vagina introitus, appeared relatively wide and easily to explore (**Fig. 8**).

Discussion

Our case is a subtype of Mayer–Rokitansky–Kuster–Hauser syndrome,^{6,13} caused by embryologic underdevelopment of the Mullerian duct with resultant agenesis of vagina and uterus. Vaginal agenesis associated with ARM is uncommonly seen.^{4–7,9,10,13–17} Banu et al¹⁸ reported 5 patients with a



Fig. 8 Neanus and neovaginal introitus at 2.6 years of age.

rectovestibular fistula and absence of vagina in their series of 563 cases.

For what concerns the reconstruction of vaginal agenesis in cases of vaginal agenesis without ARM, the first-line approach, based on safety reasons, is the primary vaginal elongation by dilations, sometimes with local estrogen, known as Vecchietti procedure^{13,19}; if no satisfactory results are achieved, during adolescence can be indicated an intra-abdominal traction applied to the perineal membrane, causing invagination over the course of a week. This technique often produces an inadequate vaginal length that might interfere with normal sexual life. Due to these reasons, other surgical options include vaginoplasty using a part of the ileus or sigmoid colon.^{5-7,9,13,15,16}

In 1956, Cohn and Murphy¹⁰ proposed for the time to correct vaginal agenesis in ARM rectovestibular fistula type, using the rectum-rectovestibular fistula in situ as a neovagina. In other reports on vaginal agenesis, an abdominoperineal approach^{14,17} and a PSARP according to Levitt et al were used.^{4,20}

This was also our decision: the vaginal agenesis was corrected using the rectum including the rectovestibular fistula to function as the vagina. The sigmoid colon was pulled through and relocated to be the neanus,² as proposed by Levitt et al,⁴ including laparotomy, but preserving the anal sphincter, according to Liem and Hau, and Liem and Quynh.^{11,12}

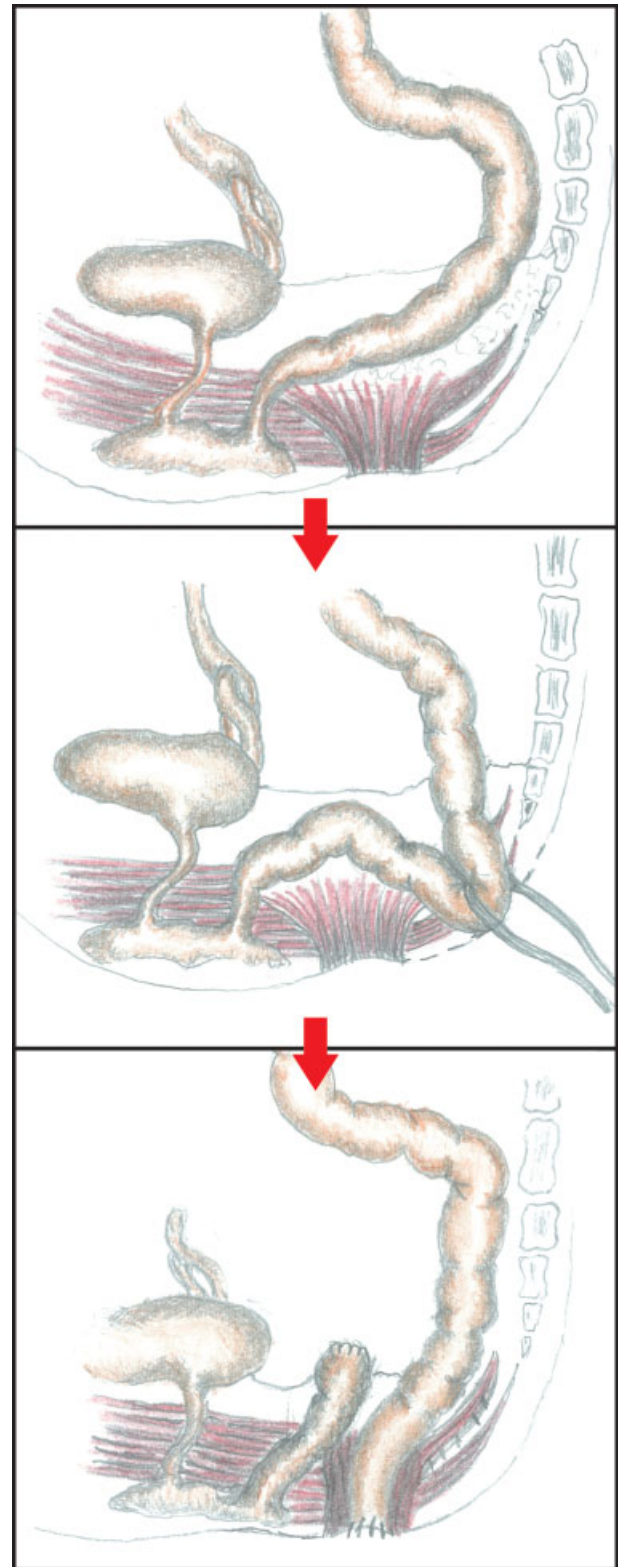


Fig. 9 Sketch of the one-stage posterosagittal approach, preserving the muscular sphincter complex (Liem and Hau, and Liem and Quynh technique), to correct vaginal agenesis and anorectal malformation with rectovestibular fistula.

At 3 years of age follow-up, the patient showed good bowel control, which generally at that age cannot be completely achieved.²¹ No clinical complications such as constipation or fecal incontinence were reported.

Although it represents a unique case, we believe that such satisfactory functional results probably also derived from the surgical technique used besides the fact that no lesions involving the sacrum were observed; unlike what takes place in the Peña and Devries' PSARP technique, based on an incision of the sphincter muscle complex in the midline,² the sphincter muscle complex was not divided and the sigmoid colon was pulled through a tunnel created at its center. Having left the sphincter muscle complex untouched leads us to hypothesize that the function of this important structure was preserved; this technique has already been reported.^{11,12}

Differently from our strategy of using the in situ rectum as a neovagina, another clinical option was to separate the rectum from urethra and relocate it within the limits of the sphincter mechanism. However, the major disadvantage of this technique is that later a more time-consuming surgery for vaginal reconstruction is required, which can also be an intervention of higher complexity. In addition, the separation of the rectum from the urethra might provoke denervation of the lower urinary tract as well as risks of denervation involving the rectum.⁴

As the rectum showed a considerable dilation, we added a suprapubic laparotomy before performing a modified PSARP with preservation of an intact anal sphincter. If the rectum had a normal caliber, we would have pulled down the proximal rectum to the neoanus (**Fig. 9**).

The final result of our procedure was an introitus situated rather anteriorly in the vulva probably due to extra space left by the missing vagina whose space might have been taken.

We hope that the future sexual function and satisfaction will be beneficial as the terminal nerves to the vulva and rectum involved were not subjected to any surgical manipulation, unlike a lot of other surgical techniques currently used for vaginal reconstruction.

Clearly, the diameter of the new vaginal introitus will have to be assessed during puberty; in the case of insufficient diameter, an event highly expected to occur, it might be widened using the wide walls of the rectovestibular fistula.

Also, at long-term follow-up, the sensitivity of the new vaginal introitus will have to be assessed.

Conclusion

Our surgical strategy based on the decision to leave the rectum in situ with its rectovestibular fistula used as a neovagina before creating a neoanus via a PSARP may provide an effective alternative to other vaginal reconstruction strategies using more complex reconstruction techniques.

Conflict of Interest

None declared.

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