## Risk factors for wound dehiscence after limited posterior sagittal anorectoplasty in female patients with low and intermediate anorectal anomalies

#### **A Thesis**

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# **List of Contents**

Pa	ge
List of Abbreviations	i
List of Figures	iii
List of Tables	v
Introduction	1
Aim of the Work	5
Review of Literature	6
Chapter 1: Anatomy and embryology	6
Chapter 2:Incidence and classifications	14
Chapter 3: Associated anomalies	21
Chapter 4:Diagnosis	26
Chapter 5: Management39	
Chapter 6:Post-operative complications	57
Patients and Methods	69
Results	74
Discussion	94
Summary and Conclusion	104
Recommendations	106
References	107
Arabic Summary	

### **List of Abbreviations**

ADA : Anterior displacement of the anus

AM : Anterior meningocele

ARMs : Anorectal malformations

ASARP : Anterior sagittal anorectoplasty approach

ASD : Atrial septal defect

CM : Cloacal membrane

CS : Currarino syndrome

EAS : External anal sphincter

EMG : Electromyography

HSD : Hirschsprung's disease

IAS : Internal anal sphincter

IV : Intravenous fluids

MCU : Micturating cystourethrogram

NPO : Nil per os

PC : Pubococcygeal distance

PDA : Patent ductus arteriosus

PFO : Patent foramen ovale

### List of Abbreviations (Cont.)

PRS : Posterior rectal shelf

PS : Pulmonary stenosis

PSARP : Posterior sagittal anorectoplasty

SSIs : Surgical site infections

TOF : Tetralogy of Fallot

VACTERL: Vertebral, anal, cardiac, tracheoesophageal,

renal and limb anomalies

VSD : Ventricular septal defect

VUR : Vesicoureteric reflux

# **List of Figures**

Fig.	Title	Page
1	Normal embryology of the hind gut.	7
2	Muscles of the anal canal.	12
3	Scheme of measurement of the API for	28
	females.	
4	Perineal fistulain females.	28
5	Perineal fistulain a female patient.	28
6	Vestibular fistula in females.	32
7	Vestibular fistula in a female patient.	32
8	Mid-sagittal slice of the pelvis	35
	demonstrates the hypothesis for the	
	pathogenesis of constipation in children	
	with anteriorly displaced anus.	
9	A, Diagram of an ideal descending	43
	colostomy. <b>B</b> , Picture of patient with ideal	
	descending colostomy.	
10	Repair of vestibular fistula.	47
11	Repair of vestibular fistula.	48
12	Repair of vestibular fistula.	48
13	Repair of vestibular fistula.	49
14A	Exposure of the posterior wall of rectum.	53
14B	Exposed posterior wall of	53
	rectum.Complete mobilization of the	
	rectum is vital till the rectal and vaginal	
	wall are separated.	
14C	Perineal body reconstruction.	53
14D	After complete reconstruction.	53
15	Showing superficial dehiscence of the	57
	wound without retraction of the neoanus	
	after ASARP for vestibular fistulae.	
16	Showing complete dehiscence of the	58
	wound with retraction of the neoanus after	
	ASARP for vestibular fistulae.	

# List of Figures (Cont.)

Fig.	Title	Page
17	<b>A</b> , Megasigmoid. <b>B</b> , Colorectal anastomosis after sigmoid resection.	65
18	Grade of recto-sigmoid dilatation in children with or without major dehiscence.	80
19	Incidence of surgery-related injuries in patients with or without major dehiscence.	82
20	Relation between performing protective colostomy and the incidence of wound dehiscence	84
21	Incidence of major dehiscence and dehiscence of any grade in children undergoing standard or extended mobilization.	89
22	Kaplan-Meier curves for the time to wound dehiscence of any grade in children undergoing standard or extended mobilization.	91
23	Kaplan-Meier curves for the time to major wound dehiscence in children undergoing standard or extended mobilization.	93

# **List of Tables**

Table	Title	Page
1	Wingspread Conference classification.	18
2	Peña's classification.	19
3	Size of anal dilators according to age.	55
4	Characteristics of patients with or without	79
	major dehiscence.	
5	Incidence of surgery-related injury in	81
	patients with or without major dehiscence.	
6	Relation between performing protective	83
	colostomy and the incidence of wound	
	dehiscence.	
7	Characteristics of children undergoing	86
	standard or extended mobilization.	
8	Incidence and timing of wound	88
	dehiscence in children undergoing	
	standard or extended mobilization.	
9	Kaplan-Meier analysis for the time to	90
	wound dehiscence of any grade.	
10	Kaplan-Meier analysis for the time to	92
	majorwound dehiscence.	

### Introduction

Anorectal malformations (ARMs) are congenital anomalies that occur in approximately 1 in 5000 live births. ARMs comprise a wide spectrum of diseases, which can affect boys and girls, and involve the distal anus and rectum as well as the urinary and genital tracts. Defects range from the very minor and easily treated with an excellent functional prognosis, to those that are complex, difficult to manage, are often associated with other anomalies, and have a poor functional prognosis (**Levitt and Peña, 2007**).

The Krickenbeck conference, participants came to the following conclusions. The international Wingspread classification is still useful in the choice of the surgical approach (This classification distinguished between high, intermediate, and low anomalies in the male and female, with special groups established for cloacal and rare anomalies). However, to develop a system for comparable follow-up studies, a modification of the classification of Peña according to the type of the fistula and including rare/regional variants was proposed. The major clinical groups were classified as perineal (cutaneous) fistulas, rectourethral fistulas (prostatic and bulbar), rectovesical

fistulas, vestibular fistulas, cloacal anomalies, patients with no fistula, and anal stenosis. Rare/regional variants were subclassified as pouch colon, rectal atresia/stenosis, rectovaginal fistulas, H-type fistulas and others. This new international classification enables the different operative procedures to be more comparable to each other than with the Wingspread classification (**Peña**, **1995**).

Most of the literature concerning management of ARMs is centered on the treatment and outcome of high anomalies. The management of low anomalies has been significantly less challenging than high considered anomalies. Also, the outcome of low anomalies has traditionally been considered good. However, recent more critical long-term follow up reports show a different picture. Many patients with low anomalies suffer from problems, long-term anorectal functional especially constipation but also soiling that occurs in a significant percentage of patients (Pakarinen and Rintala, 2010).

Vestibular fistula is the most common form of ARMs in female children and is associated with the best prognosis. It is estimated that 93% of patients with vestibular fistula will develop voluntary bowel movements by the age of 3 years (**Peña and Hong, 2000**).

Also one of the most common ARMs seen in female children is the perineal fistula, which is an anterior opening of the rectum on the perineum. We use the term fistula because the opening is not completely surrounded by sphincter muscle. It is usually narrower than the normal anus, and there is no true anal canal with a pectinate line (Peña, 2004).

The posterior sagittal anorectoplasty (PSARP), devised by **Peña and deVries** (1982), has revolutionized the management of ARMs by providing complete exposure of the anatomy of the anorectal region during surgery. This is the most widely used method and provides exact visualization of the fistula between the rectum and the female genital tract and place the rectum within the confines of the striated muscle complex giving best chance to achieve continence (**DeVriesand Peña**, 1982).

LowARM, despite its simplicity, is also likely to have the highest chances of failure leading to sever sequelae if proper reconstruction is not done at the first attempt. The perceived disadvantages of primary repair include wound infection and wound dehiscence. This makes secondary repair difficult because of fibrosis, which further decreases the chances of normal continence. A colostomy is therefore



traditionally advised so as to get the best results (**Peña and Hong, 2000**).

It must be taken into account, however, that complications. colostomies have Besides the extra operation that is needed for colostomy closure (as well as creating one), there are complications such as prolapse, skin excoriation, and the burden for parents in dealing with the colostomy. The question is whether these disadvantages outweigh the protective effect of a colostomy on wound healing after anal reconstruction (Chandramouli, et al., 2004).

# Aim of the Work

The aim of this study was to identify the risk factors for wound dehiscence after limited posterior sagittal anorectoplasty in female patients with low and intermediate anorectal anomalies in Ain Shams Universityhospitals.

### **Anatomy and Embryology**

### **Embryology of the hindgut:**

The normal embryology of the hindgut always has been a matter of debate. Two major theories exist to explain the differentiation of the hindgut into the urogenital (ventral) and anorectal (dorsal) part: The theory of the septation of the cloaca; and the theory of the migration of the rectum (**Kluth**, **2010**).

### The "Anorectal Septum" of the Hindgut:

In very young embryos, the hindgut is a simple structure. Cranially, it is in continuity with the midgut; caudally, it is in direct contact with the ectoderm, thus forming the "cloacal membrane." When development progresses, the caudal part of the hindgut, the "cloaca," differentiates into two separate organ systems— the urogenital tract and the anorectal tract (**Kluth**, **2010**).

Since the work of Tourneux (**Tourneux**, **1888**) and Retterer (**Retterer**, **1890**) at the end of the nineteenth century, it has been generally accepted that the normal development of these tracts depends upon the proper subdivision of the cloaca by a septum, the so-called urorectal septum. According to this theory, abnormal septal

development should always result in abnormal cloacal development. However, there is no agreement among investigators about the nature of this septum and the way it develops (**Kluth**, **2010**).

However, it was noted that neither lateral ridges nor signs of fusing lateral wall components could be discerned. Therefore, clear proof of this process of septation is still missing. It is more likely that a normal-looking septum is the result of normal cloacal development rather than its cause (**Kluth**, **2010**).

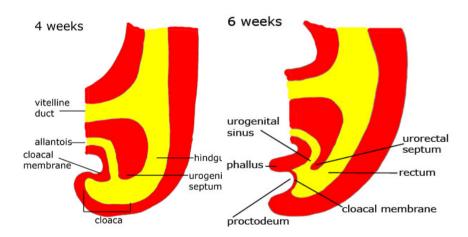


Fig. (1): Normal embryology of the hind gut (Sadler, 2012)

### The "migration" of the rectum:

Studying the morphology of an orectal malformations (ARMs) in human newborns, **Bill and Johnson in 1958** and later **Gans and Friedman in 1961** stated that in most forms of ARM the fistula may present an "ectopic" anal