





# Hirschsprung's disease

ARM	PSARP	PSARP pull through	ASARP	anoplasty	PSARPVUP
149	128	7	3	3	8



Hirschsprung's disease	Swenson	Soave	Duhamel	
59	41	14	4	









ANORECTAL MALFORMATIONS (ARM) N = 146			
	Ν	%	
Patient Sex			
Male		68	46.57
Female		78	53.42
Colostomy At Presentation			
No colostomy		27	18.49
Colostomy present		119	81.50
Type of ARM		Male(n/%)	Female(n/%)
Recto urethral fistula		31(45.58)	0(0.00)
Perineal fistula		22(32.35)	10(6.84)
Rectovesical fistula		10(14.70)	0(0.00)
Recto vestibular fistula		0(0.00)	58(74.35)
Cloaca		0(0.00)	8(10.26)
Arm with no fistula		5(6.00)	0(0.00)
Rectovaginal fistula		0(0.00)	2(2.56)

N: 146				
	Male	Female	freq.	
Spine	5	14	19	
Renal	5	5	10	
Undescended Testis	5	0	5	
Limb	1	3	4	
GIT	1	1	2	
CHD	1	1	2	
Mayer-Rokitansky- Küster-Hauser	0	2	2	
Currarino Syndrome	1	1	2	
Tethered cord	0	1	1	
Hypospadias	0	1	1	
Tracheomalcia	1	0	1	
VACTERL	0	1	1	
TOTAL	20	30	50	



#### ANORECTAL MALFORMATIONS (ARM) N = 146 Ν % **Surgical Correction Done Primary Procedure** 19 13.01 2 staged PSARP 1 0.68 **3** staged PSARP 126 86.30 Type of procedure **PSARP** 128 87.67 Anoplasty 2 1.36 **PSARP- Pull through** 7 4.79 **PSARVUP** 8 5.47 ASARP 1 0.68 Age At Perineal procedure 1 month to 1year 34 23.28 1 – 5 years 97 66.43 Above 5years 15 10.27







Complications at one month				
Major perineal wound dehiscence	8	Perineal incision wound dehiscence	14	
Surgical injuries	2	Anoplasty dehiscence	5	
Stenosis	1	Stool incontinence	4	
Sepsis	1	Excoriations	2	
Abdominal SSI	1			
Total	13		25	

Complications at six months				
Stoma wound SSI	5	Stoma closure site SSI	11	
Anal stenosis	3	Stool incontinence	10	
Urinary incontinence	3	Excoriations	2	
Anastomotic leak post stoma closure	3	Rectal prolapse	4	
Mislocated anus	2			
Rectal prolapse	2			
Vaginal stenosis	1			
DVT	1			
Total	20		27	



*Colostomy:*Unfortunately, transverse colostomies,loop colostomy and Hartman colostomy are still frequently done for patients with ARM.

The ideal colostomy remains a completely divided proximal sigmoid colostomy (at the junction of the descending and sigmoid segments of colon)

*Anal mapping*: An essential component in the manage- ment of patients with ARM, especially in higher forms where the sphincter location is not obvious on inspection. Lack of availability of such instrumentation is a significant limiting factor in resource-limited settings. Typical replacements for the popular Peña muscle stimulator are the inexpensive anesthesia nerve stimulators that can be attached via alligator clamps to needles or to a bipolar forceps.

*Caring for children with colorectal disease in the context of limited resources Seminars in Pediatric Surgery (2010) 19, 118-127* 



## Hirschsprung's disease

HIRSCHSPRUNG'S DISEASE (HD)

	Ν	(%)
	39	
Surgical Correction Done		
Primary pull through	3	7.69
2 staged Pull through	28	71.79
3 staged Pull through	8	20.51
Type of Pull-through done		
Swenson procedure	22	56.41
Soave procedure	17	43.59
Age At Pull Through		
One to five years	29	74.36
Above 5 years	10	25.64

Complicatations at first month of follow up		
Mortality	1	2.56
SSI/dehiscence	3	7.69
Anastomotic leak	1	2.56
Constipation	2	5.13
Enterocolitis	3	7.69
Anastomotic stenosis	1	2.56
Peri anal excoriation	4	10.26
Others	8	20.51
None	10	25.64
Lost to follow up	6	15.39
Complications at six months of follow up		
Constipation	6	18.18
Soiling	4	12.12
Anal stenosis	2	6.06
Other	5	15.15
HAEC	1	3.03
None	3	9.09
Lost to follow up	12	36.36
Missing data	6	



Abdominal x-ray /Contrast enema:. The combination of extreme gaseous distension, hugely dilated loops of large bowel, and massive fecal loading of the colon is diagnostic for HD. The contrast enema my show a clear transition zone.

*Fecaloma*: The retained feces in the distal colon can be very difficult to remove, especially after a barium enema To do this successfully requires dedication and application from the nursing staff, often supplemented by separate bowel washouts in the operating room and, if necessary, on-table lavage.

*Hystology/Rectal biopsy*: The resources to obtain and interpret suction rectal biopsy specimens are not available, and the only option is a full-thickness posterior rectal biopsy carried out under a general anesthetic. An adequate quantity of the bowel muscle wall must be included for evaluation.





**Caring for children with colorectal disease in the context of limited resources** Seminars in Pediatric Surgery (2010) 19, 118-127



# Hirschsprung's disease (surgery)

*Primary pull-through*: The use of a primary procedure in the malnourished, late-presenting patient with an ex- tremely dilated proximal colon is not advisable. Once the colostomy is performed, the distal colon requires 3 to 12 months to adequately reduce in size for a successful pull-through.

*Transition zone*: Late presentation of long segment HD is rare. The transition zone in most patients who present late is in the rectosigmoid.Usually, the residual fecaloma in dilated bowel is a good clinical indication that the transition zone has been reached. When there is doubt about the level of the transition zone and histologic con- firmation is not available, the bowel at the level of a well-functioning colostomy may be used for the pull- through.

**Caring for children with colorectal disease in the context of limited resources** Seminars in *Pediatric Surgery (2010) 19, 118-127* 









surgeries	151	
	Patiens with 4 surgeries	2
	Paients with 3 surgeries	3
	Patients with 2 surgeries	19
		24

- Treatment consisted in operating on burn sequelae such as contractures, hypertrophic scars and hard cords.
- Impaired mobility was our only indication for the operation.
- The most common methods of surgery were skin grafting and Z-plasty.
- A continued and often long-lasting follow-up by occupational therapists and physiotherapists is highly mandatory in order to guarantee good long-term results

Surgery on burns sequelae in developing countries O.El Ezzi,M.Dolci et Al Annals of burns and fire disasters. 2017 Mar 31;30(1):47-15



Head and Neck	11%
Upper Limb	77%
Trunk	1%
Pelvis	1%
Lower Limb	10%





	procedures	
Z plasty	91	
<ul><li>Skin graft</li><li>Split thikness</li><li>Full thikness</li></ul>	131 27 104	
flap	44	
Scar excision	60	

















# **Cleft lip and palate**

Cleft lip and/or palate (CL/P) is the most common craniofacial congenital anomaly worldwide. Yet, CL/P is undertreated in low- and middle-income countries (LMICs). Untreated children with CL/P experience malnutrition, poor dentition, ear infections, speech deficiencies, and extreme social stigma which has resulted in abandonment or infanticide. These experiences are exacerbated by delays in care.

Cleft lip and palate surgeries	40
Cleft lip repair	13
Cleft palate repair	3
Soft palate repair (two stages)	17
Hard palte repair (two stages)	7











## **Cleft lip and palate**

Surgeons in developing countries modify the timing of their repairs to improve results.

There are indications that early repair of both lip and palate in one surgical session is not inferior in outcome to the traditional staged procedure.

The aim when working in Developing Countries is to work toward the self-sufficiency: teach them a simple basic approach to cleft repair that is reproducible.





Protocols for repair of cleft lip and palate deformities were varied, with Millard's and von Langenbeck's techniques being the preferred approach for the management of cleft lip and palate deformities, respectively. A large proportion of providers have limited access to core cleft care supporting teams, especially speech language pathologists, orthodontists, and audiologists. Several challenging barriers to cleft care were also identified at both the institutional and individual levels and are reported.



## **hypospadias**

types	patients	surgeries
glandular	19	21
coronal	48	56
distal penile	15	17
midshaft	53	61
proximal penile	15	18
penoscrotal	79	114
perineal	24	29

45%	55%	
	45%	45% 55%





# **hypospadias**

techniques	surgeries
TIP	100
Bracka (1°stage)	173
Bracka(2° stage)	62
Onlay Duckett	6
Koyanagi	3
Mathieu	5
others	3

