

Research Article

External versus Transnasal Endoscopic Management of Nasolacrimal Obstruction

Adel Abd-Elbaki Abd-Allah, Balegh Hamdy Ali, Mohammad Farag Khalil, Ahmed Mustafa Eid Dessouki and Moustafa Talaat Abd El- Hakeem

Department of E.N.T. El-Minia Faculty of Medicine

Abstract

Background: Dacryocystorhinostomy (DCR) has been the standard procedure for acquired nasolacrimal duct obstruction. Lacrimal sac can be approached (1) Conventional External (Ex-DCR) or (2) Endoscopic (EN-DCR). **Objective:** To evaluate the differences between external and transnasal endoscopic approach in management of nasolacrimal obstruction.

Materials and Methods: This study included Sixty patients with distal lacrimal passage obstruction underwent DCR, forty patients with primary transnasal EnDCR (30 females and 10 males) and twenty patients (12 females and 8 males) with ExDCR from the period of January 2011 till January 2013 they were followed up at 3 and 6 months for surgical outcome. **Results:** This study included 60 patients. The patients were divided randomly into two groups (20 underwent EnDCR which subdivided in to two sub groups 10 underwent endoscopic silicone tube, 10 underwent endoscopic otologic t-tube) and the other 20 underwent external DCR. It was observed that the age of the patients ranged from 18 to 70 years with a mean age of 36.61 years, 56.7% of them were in the 3rd and 4th decades of life. There were 11 males and 49 females in the study. The success rate defined as absence of epiphora in external DCR was 80% at 3 months and the same at 6 months and in transnasal endoscopic DCR was 87.5% at 3 months and 85% at 6 months. **Conclusion:** both external and endoscopic DCR are effective surgical approaches for nasolacrimal duct obstruction with comparable success rate but endoscopic DCR with silicone tube in our study more superior than external and endoscopic otologic t-tube

KeyWords: DCR, Ex-DCR, EN-DCR

Introduction

The lacrimal drainage system includes the upper and lower punctum, superior and inferior canaliculus, common canaliculus, lacrimal sac, and nasolacrimal duct (NLD). Any obstruction in this system results in complaints of epiphora and mucoid or purulent discharge, as well as recurrent swelling in the medial canthal region. If the patient does not benefit from topical or systemic antibiotic treatment, surgery is usually the treatment of choice. Dacryocystorhinostomy (DCR) is performed either externally or transnasally (Sprekelsen and Barberan, 1996; Rice, 1990).

DCR is the standard surgical procedure for the nasolacrimal outflow tract obstruction in

which the lacrimal sac is connected directly to the nose by removing the layers of bone and mucosa that separate these two structures. The earliest surgical intervention for this purpose performed by Toti in 1904, was an external DCR (ExDCR) procedure. The external approach involves a skin incision, drilling or rongeur the bone of the anterior lacrimal crest and lacrimal sac fossa, and suturing anterior and/or posterior flaps to create a mucosal fistula into the nose (Dolman, 2003).

ExDCR is considered the mainstay of the surgical treatment in chronic dacryocystitis and in more than 90% of cases; the surgical outcome is successful (Watkins et al., 2003). However, this procedure is not without drawbacks such as external scar

formation, injury to the medial canthal ligament and periorbital echymosis that are overcome by the alternative endoscopic method (Cokkeser et al., 2003).

The endonasal approach was introduced by Caldwell at 1893 and later modified by West (Wielgosz and Mroczkowski, 2006). Today, EnDCR is typically performed by otorhino-laryngologists with the use of a nasal endoscope (Wormald, 2006; Tripathi et al., 2007; Muellner et al., 2000).

EnDCR has evolved as an alternative treatment option with significant advantages, including wide surgical field, minimal intraoperative bleeding, avoidance of scarring, and preservation of the pumping action of the orbicularis oculi muscle (Bakri et al., 1999; Hartikainen et al., 1998).

Patient and methods

Study design, prospective, interventional, clinical study was conducted in department of otorhinolaryngology in conjunction with ophthalmology, in minia university hospital. Duration of study: two years, from January 2011 till January 2013, No. of cases: 70 cases (40 EnDCR, 30 ExDCR),

Preoperative assessment

Inclusion criteria:

- 1- Adults aged over 16 years
- 2- Failed conservative treatment in the form of systemic antibiotics, steroid/antibiotic eye drops, decongestant nasal drops and local nasal steroid spray.
- 3- Distal nasolacrimal passage obstruction
- 4- History of nasal obstruction in the same side of epiphora
- 5- Fit for surgery under general anesthesia
- 6- Patients consenting for operation

Exclusion criteria:

- 1- Previous nasolacrimal surgery
- 2- Tumors of the lacrimal passage, nose or paranasal sinus
- 3- Nasal polyposis
- 4- Presacal obstruction
- 5- Eye disease causing increased lacrimation and eyelid malpositions
- 6- History of nasal trauma
- 7- CNLDO.
- 8- Mucocele or pyocele of the lacrimal sac

Subjective assessment:

The chief complaint was epiphora. Detailed medical history taking included:

- Onset, course and duration of epiphora.
- Side of lacrimal obstructive whether left, right or bilateral.
- History of allergy, history of eye disease and medication, known systemic disease.
- History of previous lacrimal operations (type, side and time of operation).
- History of radiation, nasal trauma and nasal surgeries.

Objective assessment

- Distal lacrimal passage obstruction was confirmed by the presence of positive regurgitation of pus, mucous or clear fluid through one or both puncta.
- The patency of the nasolacrimal system is assessed by syringing.
- A blunt lacrimal needle (20-gauge) is introduced into the inferior punctum and saline is injected. If the lacrimal system is obstructed, reflux of saline will occur through the upper punctum. If saline passes into the nose without reflux, the lacrimal system is patent but not necessarily functional.
- Routine preoperative nasal endoscopy to exclude intranasal pathology and detect cases that may need additional procedures e.g hypertrophied middle turbinate or significant degree of deviated septum.
- Ocular causes of epiphora were excluded with the help of an ophthalmologist.
- Only cases with confirmed distal obstruction were included in the study.

Results

Laterality of symptoms:

70% of the patients (36) presented with left sided symptomatology as compared to 30% (24) with right sided symptomatology.

40 patients underwent EnDCR 20 of them by silicone tube (8 males and 12 females) and other 20 by T-tube (3 males and 17 females) while 30 patients underwent ExDCR (8 males and 22 females) as shown

in fig. (34)

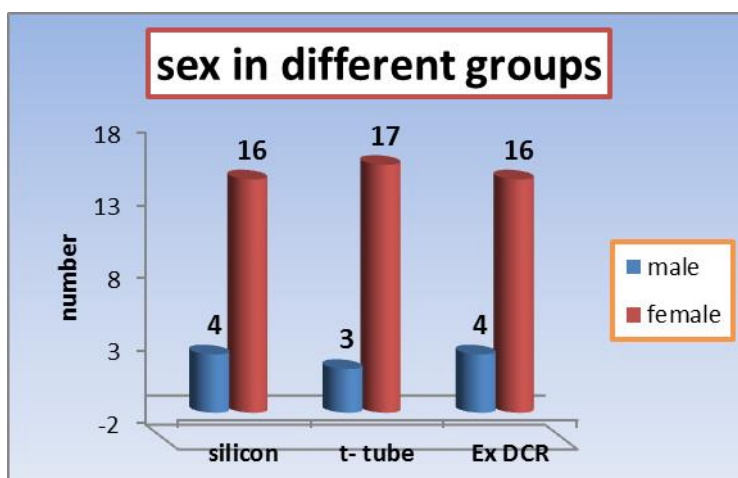


Fig (34): Showing sex in different groups

Table (2): Time taken for surgery

Duration in minutes	ExDCR		EnDCR	
	NO	%	NO	%
< 30 Minutes	-	-	8	20%
30 - 40 Minutes	7	34%	16	40%
40-60 Minutes	10	50%	12	30%
> 60	3	15%	4	10%
P-Value	0.002 **			

It was observed that EnDCR took less time (mean duration = 39.70m ± 13.5m) as compared to ExDCR (mean duration = 52.0m ± 10.1m), which was statistically significant (P = 0.002), as the duration of surgery was shorter in EnDCR.

The mean duration of follow up was 7 months were documented.
months ranging from 0 to 8 months for all patients in this study. Findings at 3 and 7 months were documented.

**Results at 3 months:
Subjective Assessment**

Table (3): Showing subjective outcome between endoscopic and ExDCR

	EnDCR	ExDCR	P value
Subjective assessment:			
Symptom free.	23 (57.5%)	9 (45%)	0.794
Significantly improved.	12 (30%)	7 (35%)	
Slightly improved.	4 (10%)	3 (15%)	
No improved.	1 (2.5%)	1 (5%)	
Worse.	0 (0%)	0 (0%)	
Fate: n (%)			
Success:	35 (87.5%)	16 (80%)	0.443
Failed.	5 (12.5%)	4 (20%)	

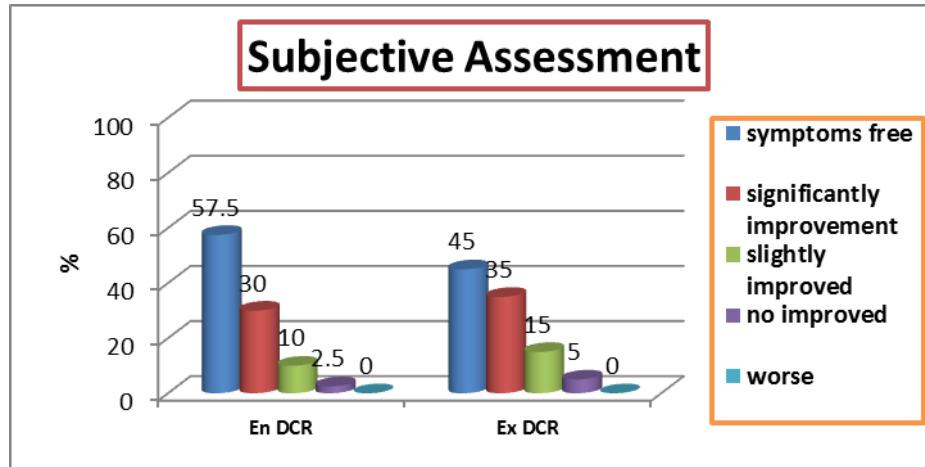


Fig (35): Chart showing subjective outcome between endoscopic and ExDCR shows no statistically significant difference between surgical outcomes of the 2 surgical groups at 3 months on the basis of subjective evaluation ($p > 0.05$).

Table (4): Showing comparison between subjective outcomes of ExDCR with endoscopic silicone DCR

	Endoscopic silicone DCR	ExDCR	P value
Subjective assessment:			
Symptom free.	16 (80%)	9 (45%)	0.110
Significantly improved.	2 (10%)	7 (35%)	
Slightly improved.	2 (10%)	3 (15%)	
No improved.	0 (0%)	1 (5%)	
Worse.	0 (0%)	0 (0%)	
Fate: n (%)			
Success:	18 (90%)	16 (80%)	0.376
Failed.	2 (10%)	4 (20%)	

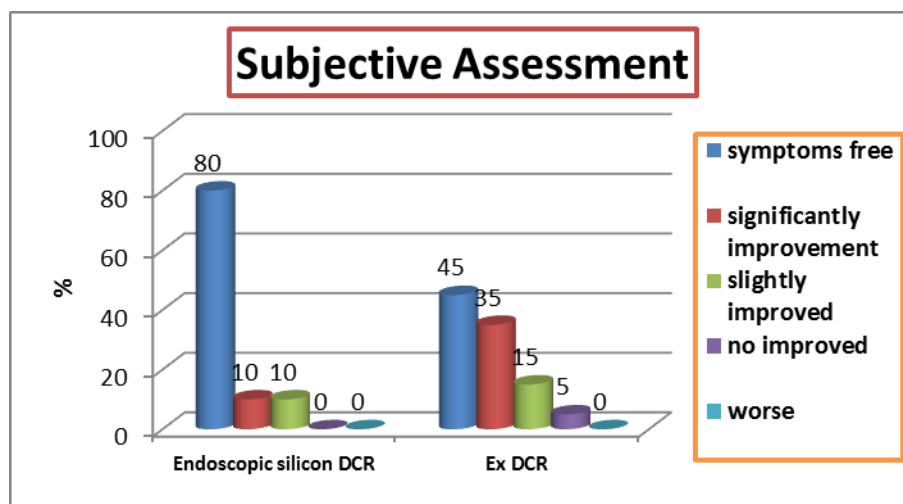


Fig (36): Showing subjective assessment of En DCR with silicone and Ex DCR at 3 months

Table (5): Showing comparison between subjective outcomes of ExDCR with endoscopic otologic T-tube DCR

Subjective assessment:	ExDCR	endoscopic otologic T-tube DCR	P value
Symptom free.	9 (40%)	7 (30%)	0.806
Significantly improved.	7 (30%)	10 (50%)	
Slightly improved.	3 (10%)	2 (10%)	
No improved.	1 (0%)	1 (0%)	
Worse.	0 (0%)	0 (0%)	
Fate: n (%)			
Success:	16 (80%)	17 (85%)	0.677
Failed.	4 (20%)	3 (15%)	

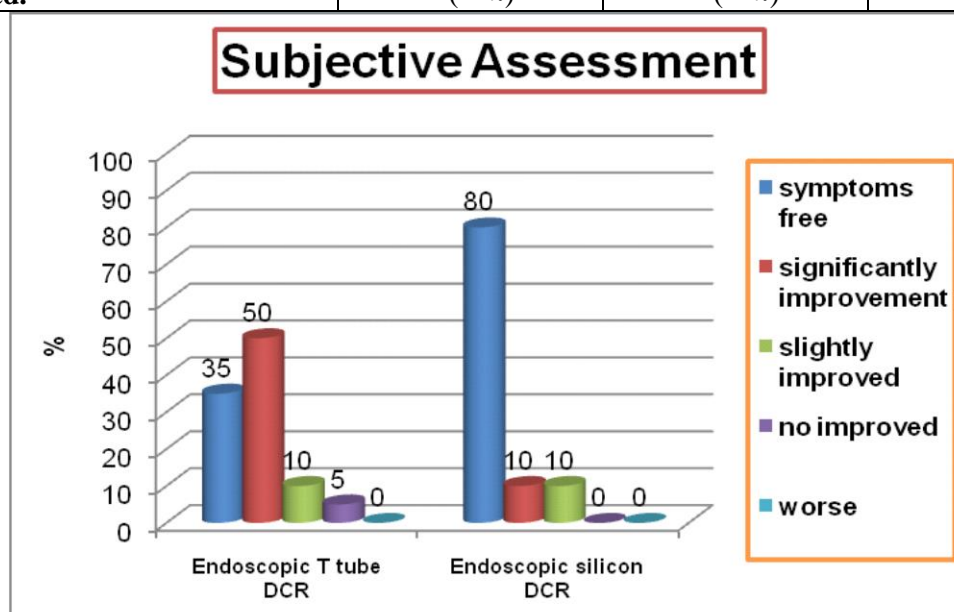


Fig (3A): There was no statistically significant difference between surgical outcomes of the 3 surgical endoscopic groups at 3 months on the basis of subjective evaluation ($p > 0.05$).

Table (6): Showing comparison between subjective outcomes of endoscopic silicone DCR with endoscopic otologic T-tube DCR

Subjective assessment:	3 month		P value
	endoscopic silicone DCR	Endoscopic otologic T-tube DCR	
Symptom free.	16 (80%)	7 (30%)	0.020*
Significantly improved.	2 (10%)	10 (50%)	
Slightly improved.	2 (10%)	2 (10%)	
No improved.	0 (0%)	1 (0%)	
Worse.	0 (0%)	0 (0%)	
Fate: n (%)			
Success:	18 (90%)	17 (85%)	0.633
Failed.	2 (10%)	3 (15%)	

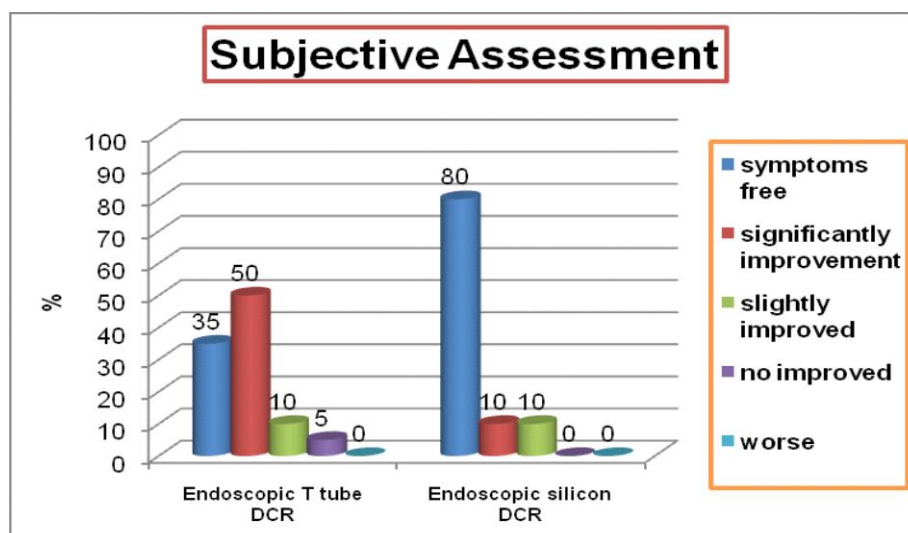


Fig (38): There was no statistically significant difference between surgical outcomes of the 2 surgical endoscopic groups at 3 months on the basis of subjective evaluation ($p > .05$).

Table (5): showing objective assessment between External and EnDCR at 3 months duration.

	EnDCR	ExDCR	P value
Florescence flow: n (%)			
+ve flow:	20 (87.5%)	16 (80%)	.443
No flow:	0 (12.5%)	4 (20%)	
Granulations: n (%)			
Present:	12 (30%)	7 (35%)	.790
Absent:	28 (70%)	13 (65%)	
Rhinostomy: n (%)			
Visible:	20 (62.5%)	11 (55%)	.076
Invisible:	10 (37.5%)	9 (45%)	
Synechia: n (%)			
Present:	11 (27.5%)	9 (45%)	.170
Absent:	29 (72.5%)	11 (55%)	

There was no statistically significant difference between surgical outcomes of the 2 surgical groups at 3 months regarding the objective assessment

Table (6): Showing objective assessment between External and endoscopic silicone DCR at 3 months duration

Objective Assessment	Endoscopic silicone group	ExDCR	P value
Florescence flow: n (%)			
+ve flow:	18 (90%)	16 (80%)	.376
No flow:	2 (10%)	4 (20%)	
Granulations: n (%)			
Present:	9 (45%)	7 (35%)	.019
Absent:	11 (55%)	13 (65%)	
Rhinostomy: n (%)			
Visible:	16 (80%)	11 (55%)	.091
Invisible:	4 (20%)	9 (45%)	
Synechia: n (%)			
Present:	7 (35%)	9 (45%)	.019
Absent:	13 (65%)	11 (55%)	

There was no statistically significant difference between surgical outcomes of the 2 surgical groups at 3 months regarding the objective assessment

Table (9): Showing objective assessment between External and endoscopic otologic T-tube DCR at 3 months duration

Objective Assessment	ExDCR	Endoscopic otologic T-tube DCR	P value
Florescence flow: n (%)			
+ve flow:	16 (80%)	17 (80%)	0.777
No flow.	4 (20%)	3 (15%)	
Granulations: n (%)			
Present.	7 (35%)	3 (15%)	0.144
Absent.	13 (65%)	17 (80%)	
Rhinostomy: n (%)			
Visible:	11 (55%)	9 (45%)	0.527
Invisible.	9 (45%)	11 (55%)	
Synechia: n (%)			
Present.	9 (45%)	4 (20%)	0.091
Absent.	11 (55%)	17 (80%)	

There was no statistically significant difference between surgical outcomes of the 3 surgical groups at 3 months regarding the objective assessment

Table (10): Showing objective assessment between endoscopic silicone and endoscopic otologic T-tube DCR at 3 months duration

objective assessment	Endoscopic silicone DCR	Endoscopic otologic T-tube	P value
Florescence flow: n (%)			
+ve flow:	18 (90%)	17 (80%)	0.633
No flow.	2 (10%)	3 (15%)	
Granulations: n (%)			
Present.	9 (45%)	3 (15%)	0.38*
Absent.	11 (55%)	17 (80%)	
Rhinostomy: n (%)			
Visible:	16 (80%)	9 (45%)	0.022*
Invisible.	4 (20%)	11 (55%)	
Synechia: n (%)			
Present.	7 (35%)	4 (20%)	0.288
Absent.	13 (65%)	17 (80%)	

There was no statistically significant difference between surgical outcomes of the 3 surgical groups at 3 months regarding the objective assessment except visibility of rhinostomy that shows statistically significant difference .

Table (11): Showing subjective outcome between endoscopic and ExDCR

subjective assessment	EnDCR	ExDCR	P value
Symptom free.	24 (60%)	9 (45%)	0.392
Significantly improved.	10 (25%)	7 (35%)	
Slightly improved.	0 (0%)	3 (15%)	
No improved.	1 (2.5%)	1 (5%)	
Worse.	0 (0%)	0 (0%)	
Fate: n (%)			
Success:	34 (80%)	16 (80%)	0.724
Failed.	6 (15%)	4 (20%)	

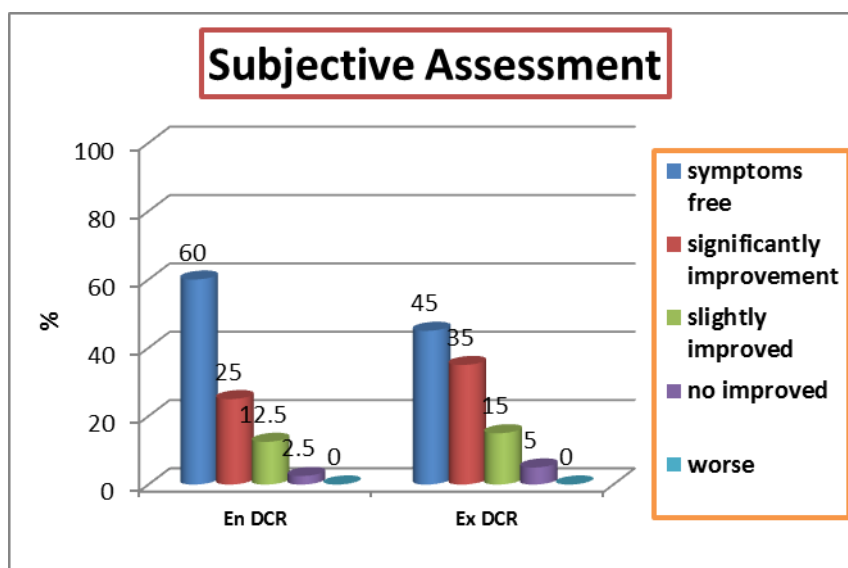


Fig (39): Chart showing subjective outcome between endoscopic and ExDCR shows no statistically significant difference between surgical outcomes of the 2 surgical groups at 7 months on the basis of subjective evaluation ($p > 0.05$).

Table (12): showing comparison between subjective outcomes of ExDCR with endoscopic silicone DCR

Subjective assessment:	Endoscopic silicone DCR	ExDCR	P value
Symptom free.	16 (80%)	9 (45%)	0.110
Significantly improved.	2 (10%)	7 (35%)	
Slightly improved.	2 (10%)	3 (15%)	
No improved.	0 (0%)	1 (5%)	
Worse.	0 (0%)	0 (0%)	
Fate: n (%)			0.376
Success:	18 (90%)	16 (80%)	
Failed.	2 (10%)	4 (20%)	

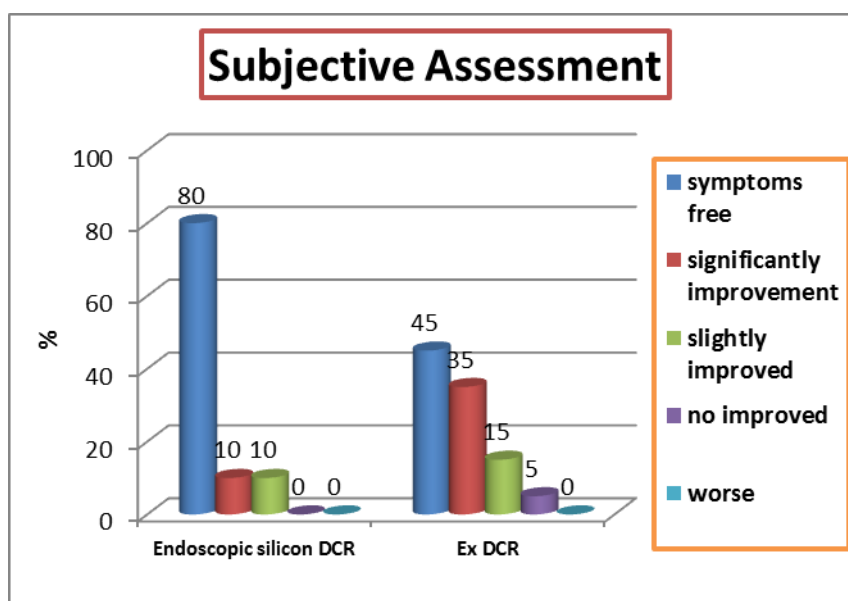


Fig (40): Showing subjective assessment of En DCR with silicone and Ex DCR at 7 months

Table (13): Showing comparison between subjective outcomes of ExDCR with endoscopic otologic T-tube DCR

Subjective assessment:	ExDCR	endoscopic otologic T-tube DCR	P value
Symptom free.	9 (40%)	7 (30%)	.919
Significantly improved.	7 (30%)	9 (40%)	
Slightly improved.	3 (10%)	3 (10%)	
No improved.	1 (5%)	1 (5%)	
Worse.	0 (0%)	0 (0%)	
Fate: n (%)			1.000
Success:	16 (80%)	16 (80%)	
Failed.	4 (20%)	4 (20%)	

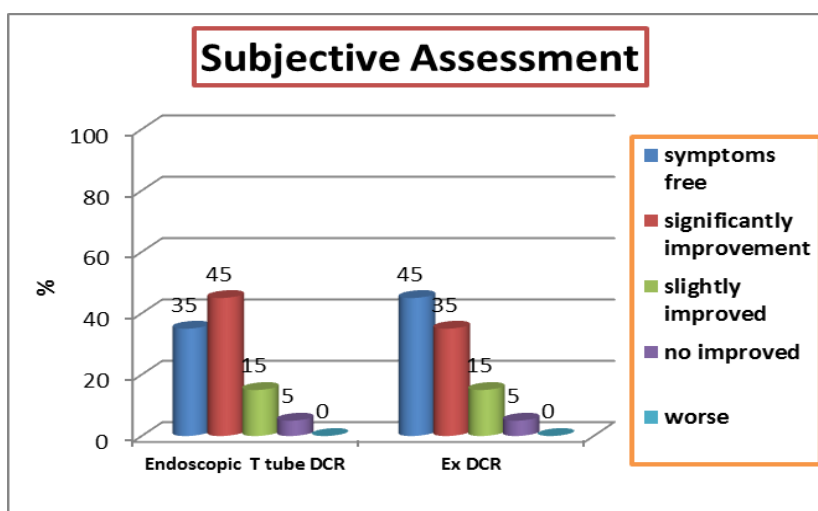


Fig (41): There was no statistically significant difference between surgical outcomes of the 2 surgical groups at 7 months on the basis of subjective evaluation (p > 0.05).

Table (14): showing comparison between subjective outcomes of endoscopic silicone DCR with endoscopic otologic T-tube DCR

Subjective assessment:	endoscopic silicone DCR	Endoscopic otologic T-tube DCR	P value
Symptom free.	16 (80%)	7 (30%)	.027*
Significantly improved.	2 (10%)	9 (40%)	
Slightly improved.	2 (10%)	3 (10%)	
No improved.	0 (0%)	1 (5%)	
Worse.	0 (0%)	0 (0%)	
Fate: n (%)			.376
Success:	18 (90%)	16 (80%)	
Failed.	2 (10%)	4 (20%)	

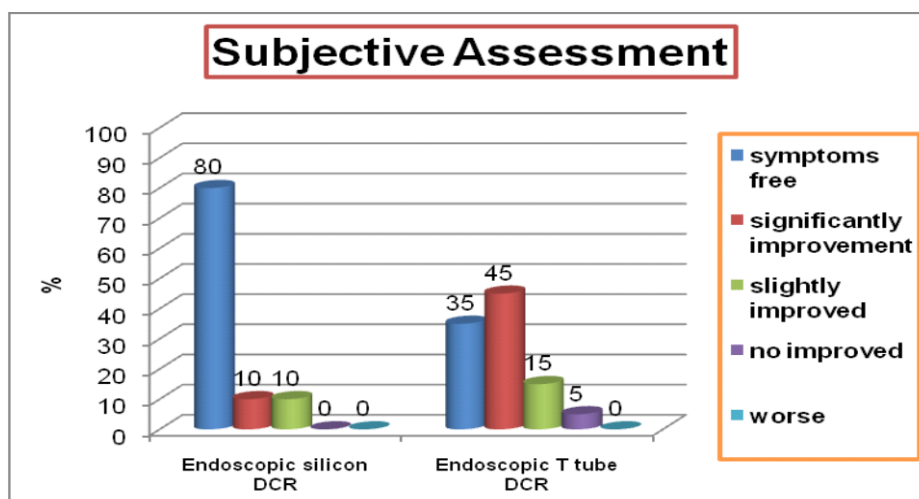


Fig (42): There was no statistically significant difference between surgical outcomes of the 2 surgical endoscopic groups at 6 months on the basis of subjective evaluation ($p > 0.05$).

Objective Assessment (at 6 months)

Table (10): Showing objective assessment between External and EnDCR at 6 months duration.

Objective Assessment	EnDCR	ExDCR	P value
Florescence flow: n (%)			
+ve flow:	34 (80%)	16 (80%)	0.624
No flow.	6 (10%)	4 (20%)	
Granulations: n (%)			
Present.	18 (45%)	10 (50%)	0.714
Absent.	22 (55%)	10 (50%)	
Rhinostomy: n (%)			
Visible:	19 (47.5%)	10 (50%)	0.800
Invisible.	21 (52.5%)	10 (50%)	
Synechia: n (%)			
Present.	19 (47.5%)	9 (45%)	0.800
Absent.	21 (52.5%)	11 (55%)	

There was no statistically significant difference between surgical outcomes of the 2 surgical groups at 6 months regarding the objective assessment

Table (11): Showing objective assessment between External and endoscopic silicone DCR at 6 months duration

Objective Assessment	Endoscopic silicone group	ExDCR	P value
Florescence flow: n (%)			
+ve flow:	18 (90%)	16 (80%)	0.376
No flow.	2 (10%)	4 (20%)	
Granulations: n (%)			
Present.	11 (55%)	10 (50%)	0.720
Absent.	9 (45%)	10 (50%)	
Rhinostomy: n (%)			
Visible:	13 (65%)	10 (50%)	0.337
Invisible.	7 (35%)	10 (50%)	
Synechia: n (%)			
Present.	11 (55%)	9 (45%)	0.527
Absent.	9 (45%)	11 (55%)	

There was no statistically significant difference between surgical outcomes of the 2 surgical

groups at 3 months regarding the objective assessment

Table (17): Showing objective assessment between External and endoscopic otologic T-tube DCR at 3 months duration

objective assessment	ExDCR	endoscopic otologic T-tube DCR	P value
Florescence flow: n (%)			
+ve flow:	16 (80%)	16 (80%)	1.000
No flow.	4 (20%)	4 (20%)	
Granulations: n (%)			
Present.	10 (50%)	7 (35%)	0.337
Absent.	10 (50%)	13 (65%)	
Rhinostomy: n (%)			
Visible:	10 (50%)	6 (30%)	0.197
Invisible.	10 (50%)	14 (70%)	
Synechia: n (%)			
Present.	9 (45%)	8 (40%)	0.749
Absent.	11 (55%)	12 (60%)	

There was no statistically significant difference between surgical outcomes of the 3 surgical groups at 3 months regarding the objective assessment

Table (18): Showing objective assessment between endoscopic silione and endoscopic otologic T-tube DCR at 3 months duration

objective assessment	Endoscopic silione DCR	Endoscopic otologic T-tube	P value
Florescence flow: n (%)			
+ve flow:	18 (90%)	16 (80%)	0.376
No flow.	2 (10%)	4 (20%)	
Granulations: n (%)			
Present.	11 (55%)	7 (35%)	0.204
Absent.	9 (45%)	13 (65%)	
Rhinostomy: n (%)			
Visible:	13 (65%)	6 (30%)	0.027*
Invisible.	7 (35%)	14 (70%)	
Synechia: n (%)			
Present.	11 (55%)	8 (40%)	0.342
Absent.	9 (45%)	12 (60%)	

There was no statistically significant difference between surgical outcomes of the 3 surgical groups at 3 months regarding the objective assessment except visibility of rhinostomy that shows statistically significant difference.

Table (19): Regarding septoplasty there is a significant difference between successes rate in patient underwent septoplasty and patients do not.

Fate of the operation	Septoplasty		P value
	Yes (24)	No (6)	
Success	22 (91.7%)	2 (33.3)	0.001*
failed	2 (8.3%)	4 (66.7)	

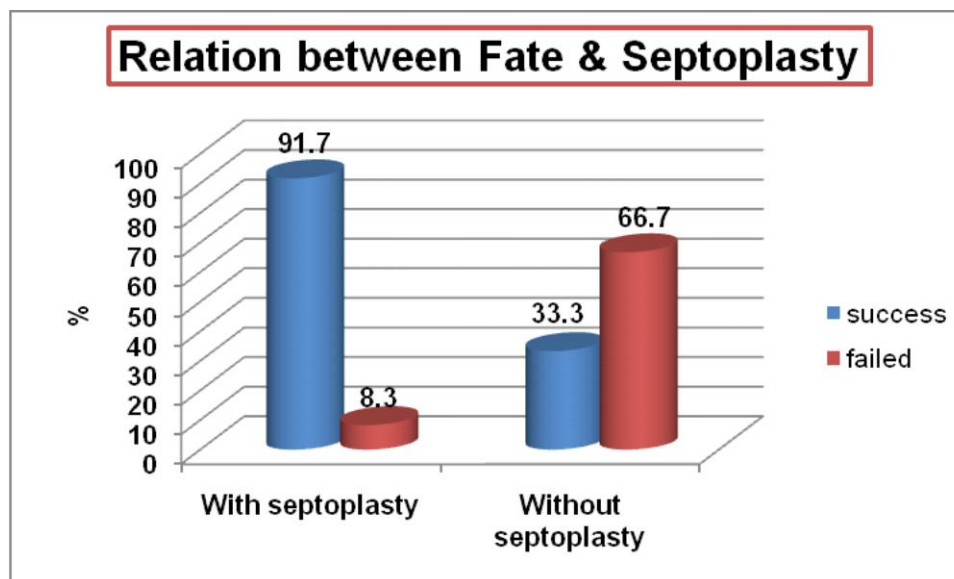


Fig (43): Showing relation of septoplasty and fate of the operation

Intra operative complication of external and EnDCR:

Table (20): Showing intraoperative complication of external and EnDCR

Intra operative complication	ExDCR		EnDCR		P-value
	NO	%	NO	%	
Moderate bleeding	7	30%	12	30%	0.690
Severe bleeding	3	10%	1	0%	0.67
Trauma of the middle turbinate	2	10%	4	10%	1.000
Accidental entry into anterior ethmoidal air cells	2	10%	1	2.0%	0.209
Laceration of punctum	-	-	-	-	-

Post operative complication of external and EnDCR:

Table (21): Post operative complication of external and EnDCR

Post operative complication	ExDCR		Endoscopic DCR		P-value
	NO	%	NO	%	
Epistaxis	1	0%	-	-	0.104
Wound infection	-	-	-	-	-
Obstruction at rhinostomy site	4	20%	6	10%	0.624
Synechiae	9	40%	19	47.0%	0.800
Granulations	10	50%	18	40%	0.714

Post operative patency rates:

The lacrimal drainage system was patent in 34 patients (80%) in Endoscopic DCR and patent in 16 patients (80%) in ExDCR at the end of 6 months, hence the success rate was 80% E

Discussion

Epiphora is an annoying symptom, embracing the patient both socially and functionally (Cokkeser et al, 2003)

The definitive treatment for most of lacrimal system disorders is surgical (Watkins et al., 2003)

DCR, which has been performed for the past hundred years, is a surgical procedure by which lacrimal flow is diverted into the nasal cavity through an artificial Extern ling made at the level of the lacrimal sac. The operation can be carried out using either an external or endoscopic trans nasal surgical approach (Watkins et al., 2003).

ExDCR, originally described by Toti in 1904 has a successful rate, when performed by properly trained ophthalmologists, of about 90% (Feretis, et al., 2006; Yigit, et al., 2007). EnDCR was described prior to this, in 1893 by Caldwell, but poor equipment and subsequent good results from the external approach led it to being abandoned. McDonogh and Meiring were the first modern surgeons credited with introducing EnDCR in the late 1940s (McDonogh and Meiring, 1949). EnDCR has a success rate of 83-94% (Smirnov, et al., 2008), and has been demonstrated to offer similar outcomes when compared to ExDCR, with low complication rates (Feretis, et al 2006; Yigit, et al., 2007). EnDCR is a good option for the treatment of primary nasolacrimal duct obstruction but it is also considered an acceptable procedure for the treatment of failure of ExDCR (Demarco et al., 2007). EnDCR has been gaining popularity, largely due to technological advances in endoscopes and other modern instruments of rhinologic surgery (Watkins et al., 2003).

In order to evaluate the endoscopic procedure versus the external surgical approach sixty patients with primary nasolacrimal duct obstruction were enrolled from Minia University ENT department and ophthalmology department.

In this study the age of the patients ranges from 16-64y and this correlate with Kuldeep et al, 2011 where the age of the patients ranged for 16-64y

The female sex was more predominant than males accounting 82% Vs 18% respectively. This came in correlation with results of Kuldeep et al, 2011 who found that 80% of the patients were females and 20% were males. The striking predilection for females can be explained by the narrower lumen of the bony nasolacrimal canal. It is also possible that endocrine factors may be playing a role in the etiology of chronic dacryocystitis.

Successful EnDCR appears to be dependent on several important factors: (1) a thorough understanding of the endoscopic anatomy and location of the lacrimal sac, (2) complete removal of the frontal process of the maxilla to expose the medial wall of the lacrimal sac, and (3) precise Externaling of the lacrimal sac to achieve adequate exposure of the common internal punctum (Mansour et al., 2006).

Another important key factor to success in EnDCR is the indication. Transnasal endoscopic approach to the lacrimal system obstruction is indicated mainly in patients with postsaccal obstruction (Mannor and Millman, 1992).

The success rate in the present study is 87.0% in EnDCR Vs 80 % in the ExDCR at 3 months and 80% in in ExDCR Vs 80 % in the ExDCR at 6 months and this correlate with Cokkeser et al, 2003 who found success rate of about 87% with EnDCR. There was no statistically significant difference between these success rates. Dolman (2003) found no statistically significant differences in the outcome between EnDCR and ExDCR.

The success rate of the EnDCR with silione is 90% vs 80% success rate of the ExDCR and this correlate with Vijay et al, 2007 that reported 93% success rate for the EnDCR with silione intubation. The use of silione

stents is considered routine in many institutions (Yigit et al., 2007). Although controversial, silicone stents are used to keep the neo-ostium patent after the procedure and are thought to maintain the patency of the ostium by preventing circular stenosis of the neo-ostium in the post-operative healing period. Prolonged silicone intubation adds to the risk of granulation tissue formation at the neo-ostium, and has been described as a cause of failure (Onceri et al., 2000). There is no general agreement regarding the duration of the stenting. Recommendations range from 4 weeks to 3 months. Kim et al, 2007 also Linberg et al, 1982 recommended the policy to remove the tubes at 12 weeks. They believed that rhinostomy patent at 12 weeks stay patent, Whittet et al 1993 recommend removal of the tubes at 3 months and lastly Metin et al, 2000 considered that long intubation period was one of the causes of DCR failure and said that the tubes should not be left for more than 3 months and this correlates with our study as the duration of stenting either for ExDCR or EnDCR was 3 months.

The success rate of the EnDCR using otologic T-tube was 80% at 3 months and 80% at 6 months follow up while ExDCR showed 80% success rate and this show no statistically significant difference between the 2 groups and this correlate with Tamura et al, 2003 that show success rate 86% for EnDCR with the otologic T-tube and the success rate for EnDCR using otologic T-tube is higher than the success rate of Kishore and McGarry (2001) reported success rate 73% and this decline in the success rate was due to spontaneous loss of the otologic T-tube and this was avoided in the current study by warning the patient not to blow the nose forcibly and prescription of medication that inhibit sneezing.

It was observed that EnDCR took less time (mean duration = 39.70m) as compared to ExDCR (mean duration = 54.0m), which was statistically significant ($P = 0.002$), as the duration of surgery was shorter in EnDCR and this correlates with Hurwitz (1996) who reported 38 min. for EnDCR and with Kuldeep et al., (2011) and Vivek et al., (2013) as the mean duration for

EnDCR is 40 min. and for ExDCR is 50 min.

Postoperative care is considered a crucial factor for the success of DCR. However, general guide lines for adequate post-operative treatment do not exist and therefore practices vary widely. The postoperative administration of systemic antibiotics and intranasal steroid spray have been recommended. Cleaning the rhinostomy site 1 week after surgery, local irrigation of the nasal cavity with saline spray and antibiotic-steroid eye drops for 2 weeks postoperatively. It is well founded that the success of surgery may be established at 6 months after the operation (Smirnov et al., 2008). In this study similar guidelines were undertaken postoperative and follow up periods ranged from 6 months to 1 year.

In the current study, it was found that bleeding was the most common Intra operative complication (0%) in the ExDCR and 30% in the EnDCR and this correlate with Kuldeep et al., (2011) that was found bleeding was the most common Intra operative complication (40%) in the ExDCR and 40% in the EnDCR.

In the current study it was observed that no major postoperative complication. epistaxis after surgery as reported as 0% of cases in ExDCR and this correlate with Fatih Qghan and Fatih Ozcura, 2008.

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