
THE EFFECT OF DEXAMETHASONE ON POST TONSILLECTOMY MORBIDITIES IN CHILDREN

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Abstract

This study aimed to evaluate the effect of preoperative single dose of Dexamethasone on postoperative morbidities in children undergoing tonsillectomy.

This is a prospective study considering 90 children with chronic tonsillitis who underwent tonsillectomy between June 2010 and December 2010 at the Department of Otolaryngology in Al-Diwaniah Teaching Hospital in Al-Diwaniah city, Iraq.

Compared with placebo, Dexamethasone group have lower percentage of post tonsillectomy morbidities (pain, postoperative nausea & vomiting and uvular edema) but without statistical significant.

In conclusion, there is no statistical beneficial effect of preoperative dexamethasone on the postoperative morbidities in children undergoing tonsillectomy.

Introduction

Tonsillectomy is one of the most common operations done in ENT specialty. Pain, nausea, vomiting, edema of uvula and poor oral intake regarded as the most common morbidities following this operation¹. Postoperative nausea and vomiting (PONV) can result in dehydration, delayed discharging, tension in suture line, venous hypertension, bleeding and pulmonary aspiration. Several studies had been done to prevent or decrease these morbidities. The use of dexamethasone is one of the methods to eliminate these morbidities depending on the anti inflammatory effect and anti emetic effect for chemotherapy induced vomiting². The aim of this study is to assess the effect of a single preoperative dose of Dexamethasone on postoperative morbidities (pain, vomiting, bleeding and edema of uvula) in children underwent tonsillectomy using a standardized anesthetic and surgical techniques³.

Materials and methods

This study is prospective in nature, it involves 90 children with chronic

tonsillitis, they had been admitted to the ENT Department at Al-Diwania Teaching Hospital for tonsillectomy by dissection and/or adenoidectomy and/or myringotomy between June 2010 and December 2010. The age ranged from 3 years to 17 years. They were 45 males and 45 females. Children who received anti emetic steroid, antihistamine and antibiotic were excluded from this study. Study design is randomized double blind and placebo control. Patients are prospectively randomized to receive either 0.5mg/kg (max. dose 8mg of dexamethasone) or an equivalent volume of normal saline. In our study the Dexamethasone or saline is given immediately after IV access was established by the anaesthetist. Patient was transferred to the ENT ward after full recovery from GA. Pain assessment either by objective pain scale (ops 0-10) in patients below 8 years (appendix 1)⁴ or by visual analogue scale (vas 0-100) in patients above 8 years (figure 1)⁵. For analysis, patients in each group (dexamethasone & control) are divided

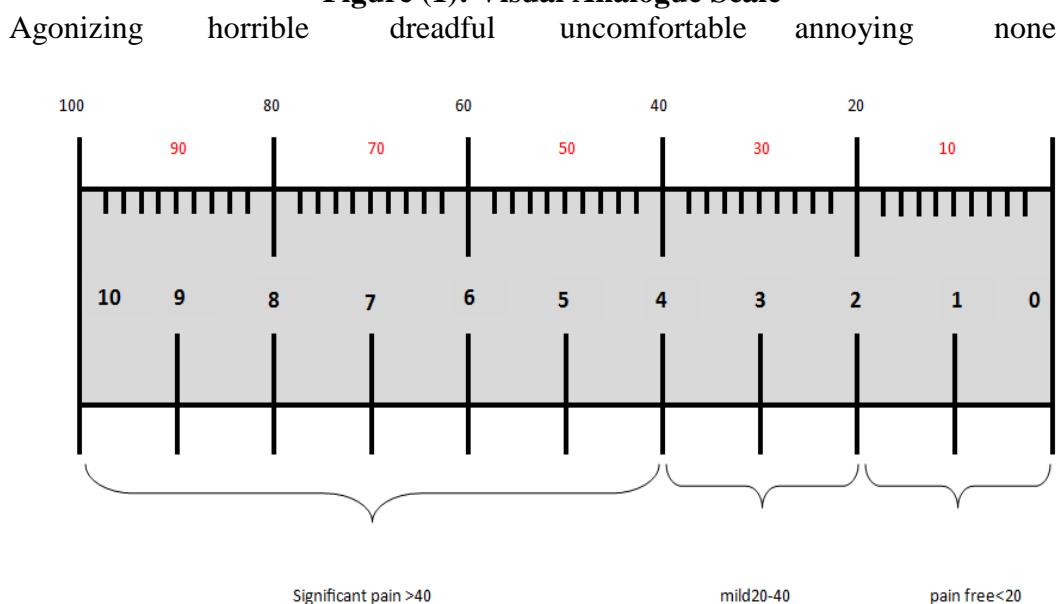
into three pain groups for early period (0-6 hrs) and late period (6-24 hrs). Significant pain = ops > 6 or vas >40, mild pain = ops 4-5 or vas >20<40, pain free = ops 0-3 or vas <20. The nurse record the incidence of vomiting (vomiting repeated within 1-2 minutes recorded as a single episode) nausea is not recorded because it is difficult to assess in children. Uvula edema was

assessed postoperatively by direct examination of oropharynx. Hemorrhage if occur was recorded. All morbidities (pain, vomiting, uvular edema and hemorrhage) are recorded in early period and late period after operation. The comparison of parameter data in this study was performed with chi-square test, a p value less than 0.05 was considered significant.

Appendix (1): Objective pain scale

Appendix 1	Objective pain scale	
observation	Criteria	points
Blood pressure	+10% of preoperative	0
	>20% of preoperative	1
	>30% of preoperative	2
Crying	Not crying	0
	Crying but respond to tender loving care	1
	Crying & not respond to tender loving care	2
Movement	Not	0
	Restless	1
	Thrashing	2
agitation	Patient asleep or calm	0
	Mild	1
	hysterical	2
Verbalizes pain	Asleep or no verbalizes	0
	Cannot localize pain	1
	Localized pain	2

Figure (1): Visual Analogue Scale



Visual Analogue Scale (VAS) is usually a horizontal line, 100mm in length, anchored by word descriptors at each end the patient mark on the line the point that they feel represent their perception of their current state. The VAS score is determined by measuring in millimetres from the left hand end of the line to the

point that the patient marks.

Results

Except for age, all demographic characteristic are comparable for both groups (table I).

Age is slightly higher in the Dexamethasone as shown in table I.

Table I: Patients demographic and clinical data

Parameter	control	Dexamethasone	P value
Age(years)	5.2	9.5	>0.001
Weight(kg)	17.2	22	0.5>p>0.1
M/f No. of patient	21/24	24/21	0.5>p>0.1
Duration of surgery (minutes)	19.6	16.8	P<0.5
Surgery type T	30	39	0.05>p>0.5
T+A	12	3	
T+A+ M	3	3	
1 st oral intake(hrs)	3	3.7	0.1>p>0.005
Method of analysis of pain Ops/vas	36/9	9/36	0.9>p>0.5(ops) p>0.01(vas)

T=tonsillectomy A=Adenoidectomy M= myringotomy Ops=objective pain scale vas=visual analogue scale. Age, weight, duration of surgery & time of first oral intake presented as mean.

Assessment of pain was done using ops in patients below eight years of age (control n=36, Dexamethasone n=9) and using vas in patients above eight years of

age (control n=9, Dexamethasone n=36). Dexamethasone group have lower % of late period without statistical significance as shown in table II.

Table II: Severity of post tonsillectomy pain

pain	Early (0-6)hr		Late(6-24)		P value
	Control No/%	Dexa. No/%	Control No/%	Dexa. No/%	
significant	27 60%	24 53.3%	6 13.3%	Zero	0.5>p>0.1
mild	15 33.3%	21 46.6%	39 86.6%	18 40%	0.5>p>0.1
Pain free	3 6.6%	Zero	Zero	27 60%	p>0.9

The % of incidence of early & late pov is slightly higher in control group

without statistical significance as shown in table III.

Table III: Incidence and severity of pov

pov	control		Dexa.		P value
	No.	%	No.	%	
Early	24	53%	30	66%	0.5>p>0.1
None					
Once	15	33%	6	13.3%	
multiple	6	13.3%	9	20%	
Late	39	86.6%	42	93.3%	0.95>p>0.5
None					
Once	6	13.3%	3	6.6%	
multiple	0	0%	0	0%	
Total no. of vomiting episodes	36		30		

The percentage of incidence of uvular edema is slightly higher in control group in early period (control=60%, Dexamethasone=20%) and in late period (control=13%, dexamethasone=0%) but without statistical significance ($p>0.05$). There is one case of early postoperative bleeding in Dexamethasone group without statistical significance.

Discussion

Tissue injury induces acute inflammation. Nerve irritation and spasm of exposed pharyngeal muscle is known to play a role in genesis of post tonsillectomy pain. Oropharyngeal pain & irritation of gastric mucosa by swallowed blood are two main contributors toward high incidence of pov following tonsillectomy. It was postulated that; inhibiting phospholipase enzyme, corticosteroid block both cyclo-oxygenase & lipo-oxygenase pathways of prostaglandin production, this decreases tissue damage by inflammation & decreases pain. Corticosteroids are used in post chemotherapy & radiotherapy to decrease vomiting & nausea, but by which mechanism it is not fully understood⁶. We select Dexamethasone as it is highly potent & has long half life (36-72) hrs. So that, the effect

would remain even after the discharge of the patient. Single iv dose was used as it is devoid of side effects like gastritis and adrenal suppression. It has been found in the present study that, there is no significant decrease in post tonsillectomy pain & uvular edema by using preoperative Dexamethasone (0.5mg/kg up to 8mg). This result agreed with Galin FI et al. (1991)⁷, Volk MS, et al. (1993)⁸. Ohlms LA, et al. (1995)⁹. These three studies shows that, there's no beneficial effect of Dexamethasone on the incidence of postoperative pain and uvular edema after tonsillectomy in children. our results in contrast with Goldman AC et al. (2000)¹⁰, Stewart DI et al (2001)¹¹, Mokhtar MD. et al. (2003)¹² & Anila D et al (2005)¹³. These four studies showed that, Dexamethasone provide significant analgesia for patients who underwent tonsillectomy. Also in our results, we found that no significant difference was noted between pov in Dexamethasone and control groups. This result agreed with Galin FI et al (1991)⁷, Volk MS, et al (1993)⁸, Ohlms LA, et al (1995)⁹, Anila D (2005)¹³. But this result disagreed with Splinter WM. et al (1991)¹⁴, Pappas, ALS, et al (1998)¹⁵, Henzi, et al (2000)¹⁶, Stewart DL, et al (2001)¹³, Aouad MT, et al (2001)¹⁷ & Mokhtar, MD et al (2003)¹⁴.

These six studies showed that preoperative Dexamethasone significantly reduced ponv in children who underwent tonsillectomy, and for the relative safety and low cost most authors recommend routine use of cs during pediatric tonsillectomy¹⁸. Also in our study, we found that there's no significant effect of Dexamethasone on postoperative bleeding. These results

agreed with most of the above studies^{7-11,13-15}.

Conclusion

We concluded that, a single preoperative iv dose (0.5mg/kg up to 8mg) of Dexamethasone have no effect in decreasing post tonsillectomy morbidities in children undergoing tonsillectomy.

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