

شبكة المعلومات الجامعية التوثيق الإلكتروني والميكروفيلو

بسم الله الرحمن الرحيم





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جامعة عين شمس التوثيق الإلكتروني والميكروفيلم قسم

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Comparative Study between Intranasal Dexmedetomidine and Intranasal Ketamine as a Premedication for Anxiolysis and Sedation before Pediatric General Anesthesia

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List of Contents

Title	Page No.
List of Abbreviations	i
List of Figures	iii
List of Tables	v
Introduction	1
Aim of the Study	4
Review of Literature	5
Patients and Methods	27
Results	37
Discussion	64
Summary	78
Conclusion	82
References	83
Arabic Summary	

List of Abbreviations

Abb.	Full term
ASA	. American society of anesthesia
CNS	. Central nervous system
CYP2A6	. Cytochrome P450
D	. Dexmedetomidine
ECG	. Electrocardiogram
GABA	. Gamma aminobutyric acid
GDRS	. Groningen distress rating scale
hr	. hour
HS	. Highly significant
K	. Ketamine
kg	. kilogram
MAP	. Mean arterial blood pressure
mcg	. Microgram
mg	. Milligram
Mg2+	. Magnesium
ml	. Milliliter
MRSS	. Modified Ramsay Sedation Scores
N2O	. Nitrous oxide
NMDA-R	. N-methyl-D-aspartate receptor
No	. Number
non REM	. non rapid eye movement
NS	. Non significant
OR	. Operating Room
PACU	. Post anesthesia care unit
RR	. Respiratory rate
S	. Significant
SD	. Standard deviation

List of Abbreviations Cont...

Abb.	Full term
SPO2	Arterial oxygen saturation
t1/2	Half life
UGT2B10), UGT1A4 Uridine 5'-diphospho-glucuronosyltransferase
c2A-AR	Alpha 2 A adrenergic receptor
œB-AR	Alpha 2 B adrenergic receptor

List of Figures

Fig. No.	Title	Page No.
Figure (1):	Child fear and anxiety from surge anesthesia	
Figure (2):	Pediatric sedation and hypnosis	9
Figure (3):	Nasal cavity	11
Figure (4):	Histology of the nasal mucosa	12
Figure (5):	Dexmedetomidine	14
Figure (6):	Chemical structure of dexmedeton	nidine15
Figure (7):	Pharmacological action of dexmedeto through alpha adrenoceptors	
Figure (8):	Physiological actions of alp adrenoreceptors	
Figure (9):	Ketamine vial	21
Figure (10):	Chemical structure of ketamine	22
Figure (11):	NMDA receptor	23
Figure (12):	Comparison between group D and K as regards gender	
Figure (13):	Comparison between group D and K as regards age and weight	_
Figure (14):	Comparison between group D and K as regards ASA status	-
Figure (15):	Comparison between group D and K as regards heart rate (HR)	-
Figure (16):	Comparison between group D and K as regards mean arterial properties (MAP).	ressure
Figure (17):	Comparison between group D and K as regards respiratory rate (RR)	0 1
Figure (18):	Comparison between group D and K as regards arterial oxygen satur	l group

List of Figures Cont...

Fig. No.	Title	Page No.
Figure (19):	Comparison between group D and K as regards modified Ramsay s score.	edation
Figure (20):	Comparison between sedation baseline before giving the drug minutes after intranasal applicadrug as regards dexmedete group	and 30 ation of omidine
Figure (21):	Comparison between sedation baseline before giving the drug minutes after intranasal applicadrug as regards ketamine group.	scores and 30 ation of
Figure (22):	Comparison between group D and K as regards cannulation score	· -
Figure (23):	Comparison between group D and K as regards parental separation	
Figure (24):	Comparison between group D and K as regards parental satisfaction	d group
Figure (25):	Comparison between group D and K as regards percentage of vomition	· -

List of Tables

Table No.	Title	Page No.
Table (1): Table (2): Table (3):	Modified Ramsay Sedation Scores (I Groningen distress rating scale (GD Parental separation score	RS)33
Table (4):	Parent satisfaction score	34
Table (5):	Comparison between group D and g as regards demographic data:	•
Table (6):	Comparison between group D and g as regards heart rate (HR)	•
Table (7):	Comparison between group D and g as regards mean arterial pressure (I	_
Table (8):	Comparison between group D and g as regards respiratory rate (RR):	roup K
Table (9):	Comparison between group D and g as regards arterial oxygen saturation	•
Table (10):	Comparison between group D and g as regards modified Ramsay so score:	edation
Table (11):	Comparison between sedation preoperatively before giving the dr 30 minutes after intranasal applicathe drug as regards dexmedeto group.	ug and ation of omidine
Table (12):	Comparison between sedation preoperatively before giving the dr 30 minutes after intranasal applicathe drug as regards ketamine group	scores ug and ation of
Table (13):	Comparison between group D and g as regards cannulation score, p separation score, parental satis score and vomiting:	arental sfaction

Introduction

Premedication in children is helpful for both separating the child from their parents and reducing the child's stress and anxiety, thus facilitating smooth induction of anesthesia. Even though intended procedures are explained to children in appropriate details, they are anxious about needle sticks and are often technically challenging to sedate. Furthermore, the drugs given for this purpose should have little effect on hemodynamics and respiration so as to allow the child to recover quickly and to facilitate early discharge without side effects (*Jun et al.*, 2017).

Anxiety of the young pediatric patient can add to the challenging nature of procedures performed before induction of general anesthesia. Pharmacologic and non-pharmacologic means of distraction and anxiolysis are commonly used to optimize the patient and family experience as well as to allow for the successful procedure completion. Intranasal medication delivery has been described as safe and effective and provides high patient and provider satisfaction (*Neville et al.*, 2016).

A lot of drugs can be taken by the intranasal route such as glucocorticoids, nasal decongestants, naloxone, midazolam, ketamine and dexmedetomidine. Intranasal route

is a very effective route for administration of drugs. The nasal mucosa can be used for non-invasive systemic administration of drugs. The surface of the nasal mucosa is a tissue well supplied by blood vessels. This ensures a rapid absorption of most drugs which generate high systemic blood levels and avoids the first pass metabolism (*Marx et al.*, 2015).

Dexmedetomidine is a selective alpha 2 agonist similar to clonidine, but with greater affinity to the alpha 2 receptor. As a sedative agent dexmedetomidine has a favorable working profile. It provides sedative properties similar to natural sleep and attenuates the stress response to the anxiolysis with minimal procedure and respiratory depression. Through the action of dexmedetomidine on the central and peripheral alpha 2 receptors it leads to reduction of heart rate and a decrease of systemic vascular resistance. The most common side effects of dexmedetomidine are hypotension, bradycardia, nausea and vomiting (Bos et al., 2017).

The ability of dexmedetomidine to maintain spontaneous ventilation and upper airway tone makes dexmedetomidine an attractive choice for procedural sedation, sleep endoscopy and imaging studies especially in pediatrics (*Absalom & Mason*, 2017).

Ketamine has been endorsed for its potential ability to offer multimodal analgesia rather than the targeted therapy, focused solely on the opioid receptors as offered by opioid medications. Ketamine is known to interact with multiple receptors, including the N-methyl-D-aspartate receptor (NMDA-R) causing a dissociative anesthesia. Ketamine is postulated to reduce central sensitization to pain, prevent opioid induced hyperalgesia, and possibly decrease overall opioid utilization (*Reynolds et al.*, 2017).