Frequency and Etiology of Nosebleeds Among Children in Pediatric Out-Patient Clinic of Ain Shams University Hospitals

Thesis

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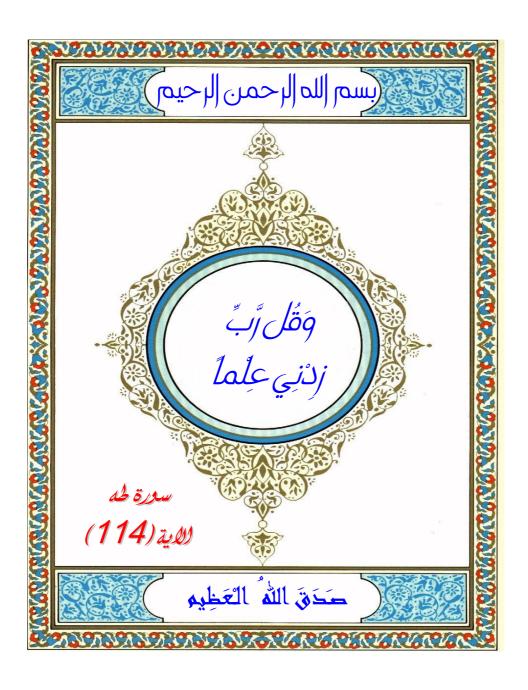
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List of Abbreviations

Abb.	Full term
ADP	Adenosine diphosphate
AG	
	Activated partial thromboplastin time
	Area under curve
	Bone marrow
<i>BS</i>	Bleeding score
	Complete blood count
	Central nervous system
<i>DBP</i>	Diastolic blood pressure
	1-Desamino-8d-azginine vasopresis
	Emergency department
<i>ENT</i>	Ear, nose and throat
FWA	Federal wide assurance
GT	Glanzmann's thrombasthenia
<i>HHT</i>	Heridteray hemorrhagic talangectasia
	Human leukocyte antigen
HS	Highly significant
IBM SPSS	International statistical package for social science business machine
<i>IgE</i>	Immunoglobulin E
<i>INR</i>	International normalization ratio
<i>ISTH</i>	International society of htormbosis and hemostasis
<i>ITP</i>	Immune thrombocytopenic purpura
<i>K2EDT</i>	Potassium ethylene diamine tetracetic acid
<i>LTA</i>	Light transmission aggregometry
<i>MPV</i>	Mean platelet volume
<i>NS</i>	Non significance
<i>PCP</i>	Primary care physician

List of Abbreviations (Cont...)

Abb.	Full term
<i>PFD</i>	Platelet function defect
PI	Prothrombin time
<i>PPP</i>	Platelet poor plasma
<i>PRP</i>	Platelet rich plasma
P-value	Probability volume
<i>RCO</i>	Ristocetin cofactor activity
<i>RFD</i>	Rare factor defeciences
<i>ROC</i>	Receiver operating characteristic curve
S	Significant
<i>SP</i>	Systolic blood pressure
<i>TT</i>	Thrombin time
<i>URRI</i>	upper respiratory tract infection
<i>USA</i>	United states of America
<i>VwD</i>	Vonwillebrand disease
<i>VwF</i>	Vonwillebrand factor
<i>WBC</i>	$White\ blood\ count$
X2	Chi square



INTRODUCTION

Epistaxis, or bleeding from the nose, is a common complaint. Most nosebleeds are benign, self-limiting, and spontaneous, but some can be recurrent. It is rarely life threatening but may cause significant concern, especially among parents of small children (Moreau et al., 1998).

As it is a common problem among children and represents a common emergency in otolaryngology. It is crucial to diagnose the cause of bleeding being local or secondary to inherited systemic coagulopathy, as it strongly reflects the management plan being totally followed by otolaryngologist or pediatric hematologist (Hussain et al., 2006).

The true prevalence of epistaxis is not known, because most episodes are self-limited and thus are not reported. When medical attention is needed, it is usually because of either the recurrent or severe nature of the problem (*Pope et al.*, 2005).

There is limited evidence regarding the prevalence of nosebleeds in children. The incidence of epistaxis declines in adulthood, but approximately one-half of all adults with epistaxis had nosebleeds during childhood (Beran and **Pertruson**, 1986). Epistaxis is rare in children younger than two years (approximately 1 per 10,000) and should prompt consideration of trauma (intentional or unintentional) or serious illness (e.g. thrombocytopenia) (Paranjothy et al., 2009). Some



pediatric healthcare providers believe that when a young baby bleeds from the nose, or is reported to have had a nosebleed, child abuse must be considered (Weindling, 2010).

As regard the data available in Egypt, a study was conducted at Hematology clinic of Ain Shams University Pediatrics Hospital revealed that epistaxis represented 15% of the presenting symptoms of inherited coagulation disorders (Mokhtar et al., 2012) and reported in 10% of pediatric ITP managed at the same hospital (Elalfy, 2013).

Another study was conducted at Cairo University to study recurrent epistaxis in children, results showed 47.6% of patients had bleeding disorders, 39.7% had ENT causes, and 12.7% had idiopathic causes. (Abdel Wahab et al., 2014).

A study was performed at the National Research Centre, Sausan Mubarak children's hospital in Cairo, Egypt and the King Abdul-Aziz University Hospital, Jeddah, Kingdom of Saudi Arabia to study the spectrum of inherited bleeding disorders reported that bruising and epistaxis were the main symptoms in all children with VWD (El-Bostany et al., 2008).

Inherited bleeding disorders include coagulation factors deficiency, platelet deficiency, platelet dysfunction and von Willebrand disease (VWD). Most of severe cases will be diagnosed in an early stage of life, providing the diagnosis is clinically suspected and laboratory diagnostic facilities are



available. However, moderate and mild cases of bleeding disorders may not be detected (Katsanis et al., 1988; Bowman et al., 2009).

We believe that in our region the prevalence of bleeding disorders is higher than the international prevalence, due to higher rate of marriages with family relatives; however, the exact rate is unknown. This study may increase the diagnosis rate of inherited bleeding disorders in children presented with epistaxis and eventually may improve the care given to those children.

AIM OF THE WORK

- 1. To measure the frequency of epistaxis as the presenting symptom among children who attended pediatric outpatient clinic.
- 2. To estimate the severity of epistaxis using severity score as an assessment tool.
- 3. To determine possible underlying etiology of epistaxis and estimate frequency of secondary epistaxis due to bleeding disorder.
- 4. Using a pediatric bleeding questionnaire as a screening tool for detecting bleeding tendency in children with epistaxis.