

شبكة المعلومات الجامعية التوثيق الإلكتروني والميكروفيلو

بسم الله الرحمن الرحيم





MONA MAGHRABY



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MONA MAGHRABY



Prophylactic Use of Haloperidol versus Atypical Antipsychotics (Quetiapine) in Prophylaxis against ICU Delirium in High Risk Patients

Thesis

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List of Abbreviations

Abb. Full term
5HT1A Serotonin 5HT1A receptor antagonist.
APA American Psychiatric Association.
APACHE II The Acute physiology and chronic health evaluation II
APOE apolipoprotein E.
ASE Attention screening examination.
AUC Area under the curve
CAM-ICU Confusion Assessment Method for the ICU.
Cmax Maximum or peak serum concentration that adrug achieves after has been administered and before administeration of asecond dose.
D1,2,3,4 receptor Dopamine 1,2,3,4 receptor
DHF decompansated heart failure.
DKA Diabetic keto acidosis.
DRS Delirium Rating Scale
DSM Diagnostic and Statistical Manual of Mental Disorders.
ECG Electrocardiography.
EEG Electroencephalography.
EPS Extrapyramidal symptoms
ER Extended release.
GABAγ-aminobutyric acid.
GCS Glascow coma score.
GnrHA Gonadotropin releasing hormone
H1receptor Histamine 1 receptor antagonist
ICDSC Intensive Care Delirium Screening Checklist
ICU Intensive care unit
RCTS Randamised Controlled Trials.

List of Abbreviations Cont...

Abb.	Full term
IQR	. Interquartile range
LAT1	. Large neutral amino acid transporter type 1.
LOS	. Length of stay.
MACh	. Muscarinic acetylcholine receptor antagonist.
NMS	. Neuroleptis malignant syndrome
PRE-DELIRIC	Prediction of Delirium in ICU patients.
RASS	. Richmond Agitation-Sedation Scale.
RF	. Respiratory failure.
SAS	. Sedation-Agitation Scale
SCCM	. Society of Critical Care Medicine.
T max	. Time taken to reach maximum concentration



Introduction

elirium is defined as acute brain dysfunction featured by disturbances of attention, awareness and cognition with a fluctuating course caused by an underlying medical condition (APA, 2013), occurs frequently in the intensive care unit, is associated with impaired patient outcome, and substantially increases healthcare costs. Given these consequences, delirium prevention is crucial (Salluh et al., *2015*).

The prevalence of delirium reported in medical and surgical ICU has varied from 20% to 80%, depending upon severity of illness observed and diagnostic methods used.

Mechanisms involved in the development of delirium, are thought related to neurotransmitters imbalances (an excess of dopamine or depletion of acetyl choline), inflammation and Impaired oxidative metabolism (Ely et al., 2001).

Multiple factors contribute to the development of delirium, including preexisting cognitive dysfunction, alcohol, drug withdrawal, sedative use, inadequate sleep, painful procedures, infection and shock state (Ouimet et al., 2007).

Delirium preventive measures are important for all ICU patients. However a delirium prediction model may facilitate early recognition of the patients who may benefit the most from delirium prevention. Non pharmacologic reduction strategies

and medication-based strategies may be most relevant for patients who have an increased risk of developing delirium (Mistraletti et al., 2012).

Various pharmacologic agents (such as Antipsychovics, acetylcholinestrase inhibitors, sleep-wake cycle regulators, and others) have been assessed for potential roles in delirium prevention (Inouve, 2006).

In this study Halopredol and Qutapiene used as preventive measures for prophylaxis of ICU delirium in high risk patients.

Halopridol is a typical antipsychotic medication. Halopridol is used in the treatment of schizophrenia, mania in bibolar disorder, delirium, agitation, acute psychosis and hallucinations in alcohol withdrawal. Despite limitations due to adverse events, including QT prolongation with the potential to trigger ventricular tachycardia, hypotension and extrapyramidal side effects (Milbrandt et al., 2005).

Quetiapine is an atypical antipsychotic drug used in treating schizophrenia, mania, depression and acute delirium outside the ICU (Maze et al., 2001).

Quetiapine anti-psychotics like other inhibits communication between nerves of the brain. It does this by blocking receptors on the nerves for neurotransmitters and chemicals that nerves use to communicate with each other. It is



thought that its beneficial effect is due to blocking of the dopamine type 2 (D2) and serotonin type 2 (5-HT2) receptors so it is associated with lower ratings of agitation, shorter duration of delirium and Increased likelihood of discharge to home or rehabilitation (Kim et al., 2003).

Quetiapine has few extrapyramidal side effects or anticholinergic symptoms, a short half-life and is mildly sedating (Kasliwal et al., 2010).

AIM OF THE WORK

The aim of study is to compare the efficacy and safety of Quetiapine in prophylaxis against ICU delirium in high risk patient with that of haloperidol.