

شبكة المعلومات الجامعية





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### جامعة عين شمس

التوثيق الالكتروني والميكروفيلم

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شبكة المعلومات الجامعية التوثيق الالكتروني والميكروفيلم

### وثائق لم ترد بلاصل

## يصعب قراءة بعض الوثائق

# The neuromuscular blocking effects of desflurane and isoflurane alone and combined with vecuronium and atracurium.

Thesis submitted for partial fulfillment of M.D. degree in Anaesthesia

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# بسم الله الرحمن الرحيم

وقل مربى أدخلني مدخل صدق وأخرجني مخرج

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صدق الله العظيم

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### **Table of Contents**

1. Abstract3				
2. Introduction and aim of the work 4				
3. Physiological anatomy of the skeletal muscle and the neuromuscular				
junction 6				
4. Physiology of neuromuscular transmission 16				
5. Pharmacology of atracurium besylate 33				
6. Pharmacology of vecuronium bromide 39				
7. Neuromuscular monitoring 45				
8. Physiochemical and pharmacodynamic properties of desflurane 62				
9. Physiochemical and pharmacodynamic properties of isoflurane 69				
0. Potentiation of neuromuscular blockade by volatile anaesthetic agents - 77				
1. Methods 96				
2. Results 103				
3. Discussion 112				
4. Summary and conclusion124				
5. References 127				
6. Arabic summary 149				

#### ABSTRACT

This study was designed to study the effect of equipotent concentrations (I MAC) of desflurane and isoflurane on the neuromuscular blockade of vecuronium and atracurium. Desflurane caused more potentiation of vecuronium neuromuscular blockade than isoflurane. Mean (SD) onset time was significantly shorter [2.3 (0.6) vs. 3.7 (0.9) min] and clinical duration [40 (8) vs. 28 (4) min] and recovery index [22 (4) vs. 15 (5) min] were significantly longer in the desflurane with vecuronium vs. the vecuronium group (p<0.05). Equipotent concentrations of desflurane and isoflurane had no statistically significant effect on atracurium neuromuscular blockade. In the absence of neuromuscular blockers, there was a significant decrease (p<0.05) in T1 amplitude in the desflurane group at all times compared to the isoflurane group. In conclusion, in equipotent concentrations, desflurane caused more potentiation of vecuronium-induced neuromuscular blockade than isoflurane, while the effects of desflurane and isoflurane on atracuriuminduced neuromuscular blockade were similar. In the absence of neuromuscular blocking agents desflurane causes significant neuromuscular blockade compared to isoflurane.

Keywords:

Anaesthetics; volatile, desflurane, isoflurane.

Neuromuscular blocking agents; nondepolarizing: vecuronium, atracurium.

INTRODUCTION AND AIM OF THE WORK

#### INTRODUCTION AND AIM OF THE WORK

Various volatile anaesthetics have been known for a long time to potentiate the neuromuscular (NM) blockade produced by non-depolarizing blocking agents to different degrees in a concentration-dependent manner. Volatile anaesthetics usually have little or no effect on the pharmacokinetics of a muscle relaxant, and this potentiation is supposed to be a pharmacodynamic one. This potentiation results from central depression as well as from peripheral inhibition of motor endplate depolarization.

When the volatile anaesthetics are eliminated, there is a rapid reversal of the potentiated NM blockade. The potentiation of NM blockade during administration of volatile anaesthetics and the reversal of this potentiation on their withdrawal are clinically relevant, as they may reduce the intraoperative NM blocking agent dose requirements for a given effect and decrease the risk of postoperative residual curarization.

Desflurane is a relatively newly introduced potent volatile anaesthetic derived from isoflurane by substitution of a single fluorine for a chlorine on the  $\alpha$ -ethyl moiety, resulting in its lower solubility in blood and tissues and less potency. Desflurane produces a dose-related impairment of myoneural transmission and can be used to supply relaxation sufficient for endoscopy or intracavity procedures at deeper levels of anaesthesia. The

lower solubility of desflurane suggests that when it is rapidly eliminated from the NM junction the enhancing effect also is rapidly eliminated.

There are only a few previous studies comparing the potentiating effects of desflurane and isoflurane on the NM blocking action of vecuronium and atracurium and their results are inconclusive. Some previous studies suggested that desflurane and isoflurane potentiate the action of vecuronium and atracurium equally, while another report concluded that desflurane has more potentiating effect on vecuronium-induced NM blockade than isoflurane.

This study was designed to study the potentiating effect of equipotent concentrations of desflurane and isoflurane on the NM blockade produced by a single dose of vecuronium and atracurium. Also, the NM blocking effects of desflurane and isoflurane in the absence of NM blocking drugs were studied.

### Physiological anatomy of the skeletal muscle and the neuromuscular junction