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شبكة المعلومات الجامعية

# بسم الله الرحمن الرحيم



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# شبكة المعلومات الجامعية التوثيق الالكتروني والميكروفيلم





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شبكة المعلومات الجامعية

# جامعة عين شمس

التوثيق الإلكتروني والميكروفيلم

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# بالرسالة صفحات لم ترد بالأصل



# **Formulation and Stability of Gatifloxacin**

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بسم الله الرحمن الرحيم

\*رب أوزعني أن أشكر نعمتك التي أنعمت علي

و على والدي و أن أعمل صالحا ترضاه

و أدخلني برحمتك في عبادك الصالحين\*

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# **LIST OF CONTENTS**

# CONTENTS

ABSTRACT	I
INTRODUCTION	1
SCOPE OF WORK	16

## CHAPTER I

### Compatibility Study between Gatifloxacin and Commonly Used Excipients in Tablets and Eye Drops

Introduction	18
Experimental	27
Results and Discussion	31
Conclusion	92

## CHAPTER II

### Formulation, Evaluation, Stability and Bioequivalence Studies of Gatifloxacin Tablets

Introduction	94
Experimental	109
Results and Discussion	123
Conclusion	180

## CHAPTER III

### Formulation, Evaluation, and Stability of Gatifloxacin Eye Drops

Introduction	184
Experimental	196
Results and Discussion	203
Conclusion	231
REFERENCES	234
ARABIC SUMMARY	-----

# ABSTRACT



## ABSTRACT

Quinolone antibiotics are widely prescribed drugs because of their safety, good tolerance and broad antibacterial spectrum with less resistance. Fluoroquinolones are bacteriostatic at low concentrations and bactericidal at high concentrations. They are quite stable in both oral and parenteral dosage forms. These compounds are sensitive to strong light and should be protected from light for long-term storage to prevent loss of activity.

The classification of the fluoroquinolones is somewhat informal and nonstandardized. Gatifloxacin is an 8-methoxy fluoroquinolone antibacterial agent. The most distinct difference between gatifloxacin's structure and those of other fluoroquinolones is the 8-methoxy group. It is believed that this group decreases the likelihood of high level resistance. The lack of halogenation at position 8 indicates that gatifloxacin, like ciprofloxacin and levofloxacin, may decrease patients risk of developing phototoxicity.

Gatifloxacin is given by mouth, or by intravenous infusion as a 2 mg per ml solution over 60 minutes, for the treatment of susceptible infections, including respiratory and urinary tract infections. The usual adult dose is 400 mg once daily. Eye drops are available in some countries as 0.3% preserved solution.

The aim of work in this thesis was to overcome the problem of poor flowability of gatifloxacin in order to formulate it as compressed tablets with lower production costs, compatible excipients, no coating