# IMPACT OF ADHERENCE TO COMMUNITY ACQUIRED PNEUMONIA MANAGEMENT GUIDELINES ON PATIENT OUTCOME

## Thesis Submitted for Partial Fulfillment of the Master Degree in Chest Diseases

By
Eman Abd Al Halem AbdAlah Al Kholy
M.B.B.ch
Faculty of Medicine, Ain Shams University

## Supervised by **Prof.Dr.Mohammed Farrag**

Professor of Chest Diseases Faculty of Medicine Ain Shams University

#### Dr. Samar Hasan Sharkawy

Assistant Professor of Chest Diseases Faculty of Medicine Ain Shams University

> Faculty of Medicine Ain Shams University 2012

### بِسْمِ اللَّهِ الرَّحْمَٰنِ الرَّحِيمِ

ا فَرَاْ بِاسْمِ رَبُّكَ الَّذِي فَلَقَ ۞ فَلَقَ الْإِنسَانَ مِنْ عَلَقٍ ۞ ا فَرَا فَرَرُكَ الْأَكْرَهُ ۞ الَّذِي عَلَّمَ بِالْفَلَمِ ۞ عَلَّمَ الْإِنسَانَ مَا لَمْ يَعْلَمْ ۞

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"سورة العلق آية 1 إلى 5 "

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#### **LIST OF ABBREVIATIONS**

| ACIP    | Advisory Committee on Immunization practices   |  |  |
|---------|--|--|--|
| AECB    | Acute- exacerbation of chronic bronchitis      |  |  |
| AIDS    | Acquired immunodeficiency syndrome             |  |  |
| ALI     | Acute lung injury                              |  |  |
| ARDS    | Acute respiratory distress syndrome            |  |  |
| ATS     | American Thoracic Society                      |  |  |
| BOOP    | Bronchiolitis obliterans organizing pneumonia  |  |  |
| CA-MRSA | Community acquired methicillin resistant       |  |  |
|         | Staphylococcus aureus                          |  |  |
| CAP     | Community-acquired pneumonia                   |  |  |
| CDC     | Centers for disease control and prevention     |  |  |
| CI      | Confidence interval                            |  |  |
| COPD    | Chronic obstructive pulmonary disease          |  |  |
| COP     | Cryptogenic organizing pneumonitis             |  |  |
| CPGs    | Clinical practice guidelines                   |  |  |
| CURB-65 | Confusion, uremia, respiratory rate, low blood |  |  |
|         | pressure, age 65or greater                     |  |  |
| DM      | Diabetes mellitus                              |  |  |
| DRSP    | Drug-resistant Strept pneumonia                |  |  |
| EBM     | Evidence-Based Medicine                        |  |  |
| ECG     | Electrocardiography                            |  |  |
| ED      | Emergency department                           |  |  |
| ESR     | Erythrocyte sedimentation rate                 |  |  |
| FDA     | Food and Drug Administration                   |  |  |
| HB      | Hemoglobin                                     |  |  |
| HCAP    | Healthcare-associated pneumonia                |  |  |
| HIV     | Human immunodeficiency virus                   |  |  |
| HPS     | Hantavirus pulmonary syndrome                  |  |  |
| ICU     | Intensive care unit                            |  |  |
| IDSA    | Infectious Diseases Society of America         |  |  |
| LOS     | Length of stay in hospital                     |  |  |
| LRTIs   | Lower respiratory tract infections             |  |  |
| LRT     | Lower respiratory tract                        |  |  |

| MRSA         | Methicillin resistant staphylococcus aureus |  |  |
|--------------|---|--|--|
| NHS          | National health service                     |  |  |
| PCP          | Pneumocystis (carinii) jiroveci pneumonia   |  |  |
| PSI          | Pneumonia Severity Index                    |  |  |
| RICU         | Respiratory intensive care unit             |  |  |
| SARS         | Severe acute respiratory syndrome           |  |  |
| S.createnine | e Serum createnine                          |  |  |
| SD           | Standard deviation                          |  |  |
| SGOT         | Serum glutamic oxaloacetic transaminase     |  |  |
| SGPT         | Serum glutamic pyruvic transaminase         |  |  |
| SLE          | Systemic lupus erythrematosis               |  |  |
| TCS          | Time to clinical stability                  |  |  |
| TLC          | Total leucocytic count                      |  |  |
| VAP          | Ventilator-associated pneumonia             |  |  |

#### LIST OF CONTENTS

| No. | Title   | Pages |
|-----|---|-------|
| 1   | Introduction  | 1     |
| 2   | Aim of the work   | 2     |
| 3   | Review  | 4     |
| •   | Defining terms  | 8     |
| •   | Pneumonia   | 12    |
| •   | Classification of pneumonia                             | 14    |
| •   | Modes of transmission                                   | 21    |
| •   | Pathology and pathogenesis of pneumonia                 | 22    |
| •   | Etiology of CAP   | 26    |
| •   | Epidemiology of CAP                                     | 32    |
| •   | CAP-associated complications                            | 42    |
| •   | Diagnosis of cap  | 47    |
| •   | Management of CAP according to IDSA,ATS guidelines 2007 | 49    |
| •   | Prevention  | 67    |
| 4   | Subject and method                                      | 79    |
| 5   | Results   | 88    |
| 6   | Discussion  | 97    |
| 7   | <b>Summary and Conclusion</b>                           | 109   |
| 8   | Recommendations   | 112   |
| 9   | References  | 113   |
| 10  | Arabic Summary  |       |

#### **LIST OF TABLES**

| No. | Name   | Pages |
|-----|--|-------|
| 1   | The most common pathogens implicated in CAP          | 33    |
|     | and their relative contributions                     |       |
| 2   | Epidemiologic conditions and\or risk factors related | 34    |
|     | to specific pathogens in community acquired          |       |
|     | pneumonia  |       |
| 3   | Most common etiologies of community acquired         | 41    |
|     | pneumonia  |       |
|     |  |       |
| 4   | Criteria for sever community acquired pneumonia      | 43    |
| 5   | Criteria for clinical stability                      | 44    |
| 6   | Clinically relevant outcome parameters in            | 56    |
|     | community acquired Pneumonia                         |       |
| 7   | Recommended empirical Antibiotics for CAP            | 65    |
| 8   | Recommendations for vaccine prevention of            | 77    |
|     | community acquired Pneumonia                         |       |
| 9   | Distribution of age and sex among group A of         | 88    |
|     | patients   |       |
| 10  | Distribution of age and sex among group B of         | 88    |
|     | patients   |       |
| 11  | Distribution of age and sex among patients of        | 88    |
|     | studied groups                                       |       |
| 12  | Distribution of Comorbidities among patients of      | 89    |
|     | studied groups                                       |       |
| 13  | Distribution of comorbidities among patients of      | 90    |
|     | studied groups                                       |       |
| 14  | Description of site of affected lobes and zones      | 91    |
|     | among patients of studied Groups                     |       |
| 15  | Laboratory investigations among group A of           | 92    |
|     | patients   |       |
| 16  | Laboratory investigations among group B of           | 93    |

|    | patients   |    |
|----|--|----|
| 17 | Description of clinical stability and time to clinical | 94 |
|    | stability among Group A patients                       |    |
| 18 | Description of clinical stability and time to clinical | 94 |
|    | stability among Group B patients                       |    |
| 19 | Description of clinical stability and time to clinical | 95 |
|    | stability(T.C.S) among patients of studied groups      |    |
| 20 | Length of stayment in hospital (L.O.S.) among          | 95 |
|    | patients of studied groups                             |    |
| 21 | T.C.S, L.O.S, and clinical stability among patients    | 96 |
|    | of studied groups                                      |    |

#### INTRODUCTION

Community-acquired pneumonia (CAP) is a common illness worldwide, and guidelines for management have been developed in many countries in the past 15 years following the initial development of CAP guidelines in North America in 1993. The most recent guidelines for the United States were published in 2007 as a joint effort of the American Thoracic Society (ATS) and the Infectious Diseases Society of America (IDSA) (Mandell et al., 2007).

The current IDSA/ATS guidelines for CAP divide patients into three groups: outpatients, those admitted to the hospital but not the ICU, and those admitted to the ICU. For each group of patients, there is a list of likely pathogens, and the initial empirical therapy is chosen with these organisms in mind. If a specific pathogen is subsequently identified by diagnostic testing, then therapy can be focused. In this scheme, it is also important to identify patients with healthcare-associated pneumonia (HCAP), and to exclude them from CAP management because these patients require their own management approach, with some needing therapy similar to CAP, whereas others require therapy that is similar to that for nosocomial pneumonia (Niederman and Brito, 2007).

In choosing empirical therapy of CAP, certain principles should guide therapy (Woodhead et al., 2005). The principles that apply to empirical therapy in U.S. guidelines are to give the first dose of therapy rapidly and before the patient leaves the emergency department; all patients should be treated for atypical pathogens and pneumococcus, plus other pathogens based on risk factors; monotherapy with macrolides can be used but should be inpatients outpatients selected limited to and with cardiopulmonary disease or recent antibiotic therapy; antipseudomonal therapy should only be used for patients with pseudomonal risk factors; MRSA therapy (vancomycin/linezolid) should be used cautiously; and no ICU-admitted CAP patient should receive monotherapy. The greatest differences from European guidelines are the recommendation for routine atypical pathogen coverage in North America, and a trend to use penicillins and to avoid quinolones in the United Kingdom (File et al., 2004).

The latest IDSA/ATS guidelines have addressed these performance measures, endorsing some but rejecting others, particularly the standard related to the timing of antibiotic administration (Mandell et al., 2007).

Delayed antibiotic therapy has been associated with increased risk of death. Therefore, a correct and rapid diagnosis is

mandatory. Two major studies have suggested that early diagnosis and early antibiotic administration are associated with improved survival and decreased length of stay in CAP (Houck et al., 2004; Meehan et al., 1997).

The US Centers for Medicare and Medicaid Services use the timing of first administration of antibiotics to patients admitted for pneumonia as a quality core measure for public reporting. In 2004, the Centers for Medicare and Medicaid Services adopted a 4-h antibiotic rule as a quality measure. The 4-h cutoff time for initial antibiotic administration as a quality core measure was questioned in some studies (Welker et al., 2008; Kanwar et al., 2007). Kanwar and colleagues compared hospitalized CAP patients before and after the 4-h rule and found that endorsement of the 4-h rule led to increased use of antibiotics (Kanwar et al., 2007).

#### **AIM OF THE WORK**

Evaluation of adherence to guidelines management of community acquired pneumonia in a random sample of adult patients and its impact on patient outcome in comparison to non-adherent patients expressed as time of clinical stability and length of stay in hospital (LOS).

Evidence-Based Medicine came to the fore in the early 1990s and has become a major driving force for many national healthcare organizations. The term and concept originated at McMaster University. It has been defined as "the integration of best research evidence with clinical expertise and patient values". EBM advocates the use of up-to-date "best" scientific evidence from health care research as the basis for making medical decisions.

#### For supporters, EBM has three main advantages:

1-It offers the surest and most objective way to determine and maintain consistently high quality and safety standards in medical practice.

2-It can help speed up the process of transferring clinical research findings into practice.

3-It has the potential to reduce health-care costs significantly.

The approach, however, is not without its opponents. These consider that EBM risks downplaying the importance of clinical experience and expert opinion, and that the conditions under which clinical trials used to define best practice take place are hard to replicate in routine practice. (*Guyatt G, et al,2004*)

The standard definition of Clinical practice guidelines (CPGs) is: "systematically developed statements to assist practitioners and patient decisions about appropriate health care for specific circumstances" (Field and Lohr,1990). Guidelines are

designed to support the decision-making processes in patient care The content of a guideline is based on a systematic review of clinical evidence - the main source for evidence-based care.

A medical guideline (also called a clinical guideline, clinical protocol or clinical practice guideline) is a document with the aim of guiding decisions and criteria regarding diagnosis, management, and treatment in specific areas of healthcare. Such documents have been in use for thousands of years during the entire history of medicine. However, in contrast to previous approaches, which were often based on tradition or authority, modern medical guidelines are based on an examination of current evidence within the paradigm of evidence-based medicine.

They usually include summarized consensus statements on best practice in healthcare. A healthcare provider is obliged to know the medical guidelines of his or her profession, and has to decide whether or not to follow the recommendations of a guideline for an individual treatment (Council of Europe., 2002).

Modern clinical guidelines identify, summarize and evaluate the highest quality evidence and most current data about prevention, diagnosis, prognosis, therapy including dosage of medications, risk/benefit and cost-effectiveness. Then they define the most important questions related to clinical practice and identify all possible decision options and their outcomes. Some