

Global Initiative for Chronic Obstructive Lung Disease

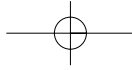


GLOBAL STRATEGY FOR THE DIAGNOSIS, MANAGEMENT, AND PREVENTION OF CHRONIC OBSTRUCTIVE PULMONARY DISEASE

UPDATED 2003

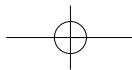
EXECUTIVE SUMMARY

NATIONAL INSTITUTES OF HEALTH
National Heart, Lung, and Blood Institute



EXECUTIVE SUMMARY UPDATED 2003

**GLOBAL STRATEGY FOR THE DIAGNOSIS, MANAGEMENT,
AND PREVENTION OF CHRONIC OBSTRUCTIVE PULMONARY DISEASE**
(Based on an April 1998 NHLBI/WHO Workshop)



**Global Strategy for the Diagnosis, Management, and Prevention of
Chronic Obstructive Pulmonary Disease: NHLBI/WHO Workshop**
National Heart, Lung, and Blood Institute: Claude Lenfant, MD
World Health Organization: Nikolai Khaltaev, MD
April 1998 Workshop Panel

Romain Pauwels, MD, PhD, *Chair*
Ghent University Hospital
Ghent, Belgium

Nicholas Anthonisen, MD
University of Manitoba
Winnipeg, Manitoba, Canada

William C. Bailey, MD
University of Alabama at Birmingham
Birmingham, Alabama, US

Peter J. Barnes, MD
National Heart & Lung Institute
London, UK

A. Sonia Buist, MD
Oregon Health Sciences University
Portland, Oregon, US

Peter Calverley, MD
University Hospital, Aintree
Liverpool, UK

Tim Clark, MD
Imperial College
London, UK

Leonardo Fabbri, MD
University of Modena & Reggio Emilia
Modena, Italy

Yoshinosuke Fukuchi, MD
Juntendo University
Tokyo, Japan

Lawrence Grouse, MD, PhD
University of Washington
Seattle, Washington, US

James C. Hogg, MD
St. Paul's Hospital
Vancouver, British Columbia, Canada

Christine Jenkins, MD
Concord Hospital
Sydney, New South Wales, Australia

Dirkje S. Postma, MD
Academic Hospital Groningen
Groningen, the Netherlands

Klaus F. Rabe, MD
Leiden University Medical Center
Leiden, the Netherlands

Scott D. Ramsey, MD, PhD
University of Washington
Seattle, Washington, US

Stephen I. Rennard, MD
University of Nebraska Medical Center
Omaha, Nebraska, US

Roberto Rodriguez-Roisin, MD
University of Barcelona
Barcelona, Spain

Nikos Siafakas, MD
University of Crete Medical School
Heraklion, Greece

Sean D. Sullivan, PhD
University of Washington
Seattle, Washington, US

Wan-Cheng Tan, MD
National University Hospital
Singapore

GOLD Staff

Sarah DeWeerd
Editor
Seattle, Washington, US

Suzanne S. Hurd, PhD
Scientific Director
Bethesda, Maryland, US

Consultant Reviewers

Individuals

Sherwood Burge (UK)
Maira Chan-Yeung (Hong Kong)
James Donohue (US)
Nicholas J. Gross (US)
Helgo Magnussen (Germany)
Donald Mahler (US)
Jean-Francois Muir (France)
Mrigendra Pandey (India)
Peter Pare (Canada)
Thomas Petty (US)
Michael Plit (South Africa)
Sri Ram (US)
Harold Rea (New Zealand)
Andrea Rossi (Italy)
Maureen Rutten-van Molken (The Netherlands)
Marina Saetta (Italy)
Raj Singh (India)
Frank Speizer (US)
Robert Stockley (UK)
Donald Tashkin (US)
Ian Town (New Zealand)
Paul Vermeire (Belgium)
Gregory Wagner (US)
Scott Weiss (US)
Miel Wouters (The Netherlands)
Jan Zielinski (Poland)

Organizations

American College of Chest Physicians

Suzanne Pingleton

American Thoracic Society

Bart Celli
William Martin

Austrian Respiratory Society

Friedrich Kummer

Arab Respiratory Society

Salem El Sayed

Thoracic Society of Australia and New Zealand

Alastair Stewart
David McKenzie
Peter Frith

Australian Lung Foundation

Robert Edwards

Belgian Society of Pneumology

Marc Decramer
Jean-Claude Yernault

British Thoracic Society

Neil Pride

Canadian Thoracic Society

Louis-Philippe Boulet
Kenneth Chapman

Chinese Respiratory Society

Nan-Shan Zhong
Yuanjue Zhu

Croatian Respiratory Society

Neven Rakusic
Davor Plavec

Czech Thoracic Society

Stanislav Kos
Jaromir Musil
Vladimir Vondra

European Respiratory Society

Marc Decramer (Belgium)

French Speaking Pneumological Society

Michel Fournier
Thomas Similowski

Hungarian Respiratory Society

Pal Magyar

Japanese Respiratory Society

Yoshinosuke Fukuchi

Latin American Thoracic Society

Juan Figueroa (Argentina)
Maria Christina Machado (Brazil)
Ilma Paschoal (Brazil)
Jose Jardim (Brazil)
Gisele Borzone (Chile)
Orlando Diaz (Chile)
Patricio Gonzales (Chile)
Carmen Lisboa (Chile)
Rogelio Perez Padilla (Mexico)
Jorge Rodriguez De Marco (Uruguay)
Maria Victorina Lopez (Uruguay)
Roberto Lopez (Uruguay)

Malaysian Thoracic Society

Zin Zainudin

Norwegian Thoracic Society

Amund Gulsvik
Ernst Omenaas

Polish Phthisiopneumonological Society

Michal Pirozynski

Romanian Society of Pulmonary Diseases

Traian Mihaescu
Sabina Antoniu

Singapore Thoracic Society

Alan Ng
Wei Keong

Slovakian Pneumological and Phthisiological Society

Ladislav Chovan

Slovenian Respiratory Society

Stanislav Suskovic

South African Thoracic Society

James Joubert

Spanish Society of Pneumology

Teodoro Montemayor Rubio
Victor Sobradillo

Swedish Society for Chest Physicians

Kjell Larsson
Sven Larsson
Claes-Goran Lofdahl

Swiss Pulmonary Society

Philippe Leuenberger
Erich Russi

Thoracic Society of Thailand

Ploysongsang Youngyudh

Vietnam Asthma-Allergology and Clinical Immunology Association

Nguyen Nang An

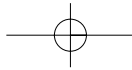


TABLE OF CONTENTS

Preface vi

Methods Used to Update Report; Summary Recommendations vii

Introduction 1

1. Definition and Classification of Severity 2

 Definition 2

 Classification of Severity 2

 Pathogenesis 3

 Pathology 3

 Pathophysiology 3

2. Burden of COPD..... 4

 Epidemiology 4

 Economic and Social Burden of COPD 4

 Risk Factors 5

3. Four Components of COPD Management 6

 Introduction..... 6

Component 1: Assess and Monitor Disease 6

 Diagnosis 6

 Ongoing Monitoring and Assessment 8

Component 2: Reduce Risk Factors 9

 Smoking Prevention and Cessation 9

 Occupational Exposures 10

 Indoor/Outdoor Air Pollution 10

Component 3: Manage Stable COPD 11

 Introduction 11

 Education 11

 Pharmacologic Treatment 11

 Bronchodilators 12

 Glucocorticosteroids 14

 Other Pharmacologic Treatments 14

 Non-Pharmacologic Treatment 15

 Rehabilitation 15

 Oxygen Therapy 15

 Ventilatory Support 16

 Surgical Treatments 16

Component 4: Manage Exacerbations 16

 Diagnosis and Assessment of Severity 16

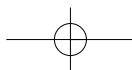
 Home Management..... 17

 Hospital Management..... 18

 Hospital Discharge and Follow-up 20

4. Future Research 21

References 22



PREFACE

Chronic Obstructive Pulmonary Disease (COPD) is a major public health problem. It is the fourth leading cause of chronic morbidity and mortality in the United States¹ and is projected to rank fifth in 2020 as a worldwide burden of disease according to a study published by the World Bank/World Health Organization². Yet, COPD fails to receive adequate attention from the health care community and government officials. With these concerns in mind, a committed group of scientists encouraged the US National Heart, Lung, and Blood Institute and the World Health Organization to form the Global Initiative for Chronic Obstructive Lung Disease (GOLD). Among GOLD's important objectives are to increase awareness of COPD and to help the thousands of people who suffer from this disease and die prematurely from COPD or its complications.

The first step in the GOLD program was to prepare a consensus Workshop Report, *Global Strategy for the Diagnosis, Management, and Prevention of COPD*. The GOLD Expert Panel, a distinguished group of health professionals from the fields of respiratory medicine, epidemiology, socioeconomics, public health, and health education, reviewed existing COPD guidelines, as well as new information on pathogenic mechanisms of COPD as they developed a consensus document. Many recommendations will require additional study and evaluation as the GOLD program is implemented.

A major problem is the incomplete information about the causes and prevalence of COPD, especially in developing countries. While cigarette smoking is a major known risk factor, much remains to be learned about other causes of this disease. The GOLD Initiative will bring COPD to the attention of governments, public health officials, health care workers, and the general public, but a concerted effort by all involved in health care will be necessary to control this major public health problem.

I would like to acknowledge the dedicated individuals who prepared the Workshop Report and the effective leadership of the Workshop Chair, Professor Romain Pauwels. It is a privilege for the National Heart, Lung, and Blood Institute to serve as one of the cosponsors. We look forward to working with the World Health Organization, and all other interested organizations and individuals, to meet the goals of the GOLD Initiative.

Development of the Workshop Report was supported through educational grants to the Department of Respiratory Diseases of the Ghent University Hospital, Belgium (WHO Collaborating Center for the Management of Asthma and COPD) from Altana, Andi-Ventis, AstraZeneca, Aventis, Bayer, Boehringer Ingelheim, Chiesi, GlaxoSmithKline, Merck, Sharp & Dohme, Mitsubishi Pharma, Nikken Chemicals, Novartis, Pfizer, Schering-Plough, and Zambon

Claude Lenfant, MD
Director
National Heart, Lung, and Blood Institute

Methodology and Summary of New Recommendations*

A GOLD Science Committee** was established to review published research that impacts on recommendations for COPD presented in the 2001 GOLD Workshop Report, and to post an updated report yearly on the GOLD website. The 2003 update includes review of publications from June 2000 (approximate time of completion of the 2001 report) through March 2003. Each year, a new update report will be posted; a revision of the entire document will be prepared approximately every 5 years.

Methods: The process included a Pub Med search using search fields established by the Committee: 1) COPD OR chronic bronchitis OR emphysema, All Fields, All Adult, 19+ years, only items with abstracts, Clinical Trial, Human, sorted by Authors; and 2) COPD OR chronic bronchitis OR emphysema AND systematic, All fields, All adult, 19+ years, only items with abstracts, Human, sorted by Author. In addition, publications in peer review journals not captured by Pub Med could be submitted to individual members of the Committee providing an abstract and the full paper were submitted in (or translated into) English.

All members of the Committee received a summary of citations and all abstracts. Each abstract was assigned to 2 Committee members (members were not assigned to a paper where he/she appears as an Author), although any member was offered the opportunity to provide an opinion on any abstract. Members evaluated the abstract or, up to her/his judgment, the full publication, by answering specific written questions from a short questionnaire, and to indicate if the scientific data presented impacted on recommendations in the GOLD Workshop Report. If so, the member was asked to specifically identify modifications that should be made. The entire GOLD Science Committee met on a regular basis to discuss each individual publication that was indicated to have an impact on COPD management by at least 1 member of the Committee, and to reach a consensus on the changes in the report. Disagreements were decided by vote.

Summary of New Recommendations: Between June 2000 and March 2003, 241 articles met the search

*Fabbri LM, Hurd SS. GOLD Scientific Committee. Global Strategy for the Diagnosis, Management, and Prevention of COPD 2003 Update. (Editorial) *Eur Resp J.* 2003;22:1-2

**Members: L. Fabbri, *Chair*; P. Barnes, S. Buist, P. Calverley, Y. Fukuchi, W. MacNee, R. Pauwels, K. Rabe, R. Rodriguez-Roisin, I. Zielink

criteria. Of these, 36 papers were identified to have an impact on the GOLD report, either by: 1) confirming, that is, adding or replacing an existing reference, or 2) modifying, that is, changing the text or introducing a concept that required a new recommendation to the report. The major modifications introduced to the management section include:

- The position of long-acting and short-acting bronchodilators, including the introduction of the new long-acting anticholinergic, tiotropium
- The position of inhaled glucocorticosteroids and combination of inhaled long-acting, β_2 -agonists/glucocorticosteroids
- Evidence related to length of pulmonary rehabilitation programs
- Home vs. hospital care for COPD exacerbations

Because of difficulties encountered using the 2001 GOLD classification by severity in the dissemination process, and in line with the recommendations that are being proposed by the COPD Guidelines Committee nominated jointly by the European Respiratory Society and by the American Thoracic Society, the classification was maintained, but the stages of severity were renamed into *0=At Risk*, *I=Mild*, *II=Moderate*, *III=Severe*, and *IV=Very severe* to replace stages *At Risk* (0), *Mild* (I), *Moderate* (IIA, IIB), and *Severe* (III) respectively.

The 2001 GOLD Workshop Report included the recommendation to use regular treatment with bronchodilators for moderate to severe COPD, mentioning that long-acting bronchodilators were more convenient than short-acting bronchodilators. Based on publications that appeared since June 2000, the updated 2003 GOLD Workshop Report recommends for moderate to very severe COPD use of regular treatment with long-acting bronchodilators, including tiotropium, rather than short-acting bronchodilators (**Evidence A**).

The 2001 Report included the recommendation to use inhaled glucocorticosteroids for patients with moderate COPD or more, providing they had a spirometric response to a short-term course of steroids and/or an $FEV_1 < 50\%$ predicted and frequent exacerbations. This recommendation was assigned (**Evidence B**) reflecting the inconsistency of response to inhaled glucocorticosteroids reported in the literature. Based on publications appearing since June 2000, the 2003 Report recommends use of inhaled glucocorticosteroids only in patients with severe and very severe (Stages III and IV) COPD (called IIB and III in the 2001 Report) and fre-

quent exacerbations, assigning to the recommendation (**Evidence A**), reflecting the consistency of the response to inhaled glucocorticosteroids in more severe patients reported in the literature.

The 2001 Report did not include a specific recommendation for the duration of rehabilitation programs. Based on publications appearing since June 2000, the 2003 Report recommends a duration of at least 2 months for rehabilitation programs, assigning to the recommendation (**Evidence B**), reflecting the limited number of studies available.

The 2001 GOLD Workshop Report did not include a specific recommendation for the nurse administered home care as an alternative to hospitalization of patients with COPD exacerbations. Based on publications appearing since June 2000, the 2003 GOLD Workshop Report suggests that nurse-administered home care represents an effective and practical alternative to hospitalization in selected patients with exacerbations of COPD without acidotic respiratory failure. However, because the exact criteria for home compared to hospital treatment remains uncertain, and may vary by health care setting, no level of evidence was assigned to this recommendation.

Finally, in addition to small changes and correction of mistakes contained in the original document, the Committee identified important issues for which the new scientific evidence reviewed was considered insufficient to change the 2001 Report, but that were judged as priority issues to be addressed in the 2004 update. These include antibiotic treatment of COPD exacerbations, step-up/down of pharmacological treatment, use of walking aids for rehabilitation, and anesthesia in severe COPD patients undergoing surgery.

Prior to its release, the proposed modifications to the 2001 GOLD Workshop Report were submitted to the GOLD Executive Committee for approval. The 2003 GOLD Workshop Report (Executive Summary and Pocket Guide) and the complete list of references examined by the Committee are available on the GOLD website (www.goldcopd.com).

INTRODUCTION

Chronic Obstructive Pulmonary Disease (COPD) is a major cause of chronic morbidity and mortality throughout the world. COPD is currently the fourth leading cause of death in the world³, and further increases in the prevalence and mortality of the disease can be predicted in the coming decades. A unified international effort is required to reverse these trends.

The **Global Initiative for Chronic Obstructive Lung Disease (GOLD)** is conducted in collaboration with the US National Heart, Lung, and Blood Institute (NHLBI) and the World Health Organization (WHO). Its goals are to increase awareness of COPD and decrease morbidity and mortality from this disease. GOLD aims to improve prevention and management of COPD through a concerted worldwide effort of people involved in all facets of health care and health care policy, and to encourage a renewed research interest in this extremely prevalent disease.

The GOLD Workshop Report, *Global Strategy for the Diagnosis, Management, and Prevention of COPD*, presents a COPD management plan with four components: (1) Assess and Monitor Disease; (2) Reduce Risk Factors; (3) Manage Stable COPD; (4) Manage Exacerbations. The Workshop Report is based on the best-validated current concepts of COPD pathogenesis and the available evidence on the most appropriate management and prevention strategies. It has been developed by individuals with expertise in COPD research and patient care and extensively reviewed by many experts and scientific societies. Prior to its release for publication, the Workshop Report was reviewed by the NHLBI and the WHO. This Executive Summary provides key information about COPD; the full Workshop Report provides more details.

In section 3, "Four Components of COPD Management," levels of evidence are assigned to statements, where appropriate, using a system developed by the NHLBI (**Table 1**). Levels of evidence are indicated in boldface enclosed in parentheses after the relevant statement – e.g., (**Evidence A**).

Evidence Category	Sources of Evidence	Definition
A	Randomized controlled trials (RCTs). Rich body of data.	Evidence is from endpoints of well-designed RCTs that provide a consistent pattern of findings in the population for which the recommendation is made. Category A requires substantial numbers of studies involving substantial numbers of participants.
B	Randomized controlled trials (RCTs). Limited body of data.	Evidence is from endpoints of intervention studies that include only a limited number of patients, posthoc or subgroup analysis of RCTs, or meta-analysis of RCTs. In general, Category B pertains when few randomized trials exist, they are small in size, they were undertaken in a population that differs from the target population of the recommendation, or the results are somewhat inconsistent.
C	Nonrandomized trials. Observational studies.	Evidence is from outcomes of uncontrolled or nonrandomized trials or from observational studies.
D	Panel Consensus Judgment.	This category is used only in cases where the provision of some guidance was deemed valuable but the clinical literature addressing the subject was deemed insufficient to justify placement in one of the other categories. The Panel Consensus is based on clinical experience or knowledge that does not meet the above-listed criteria.

1. DEFINITION AND CLASSIFICATION OF SEVERITY

DEFINITION

COPD is a disease state characterized by airflow limitation that is not fully reversible. The airflow limitation is usually both progressive and associated with an abnormal inflammatory response of the lungs to noxious particles or gases.

A diagnosis of COPD should be considered in any patient who has symptoms of cough, sputum production, or dyspnea, and/or a history of exposure to risk factors for the disease. The diagnosis is confirmed by spirometry. The presence of a postbronchodilator FEV₁ < 80% of the predicted value in combination with an FEV₁/FVC < 70% confirms the presence of airflow limitation that is not fully reversible. Where spirometry is unavailable, the diagnosis of COPD should be made using all available tools. Clinical symptoms and signs, such as abnormal shortness of breath and increased forced expiratory time, can be used to help with the diagnosis. A low peak flow is consistent with COPD, but has poor specificity since it can be caused by other lung diseases and by poor performance. In the interest of improving the diagnosis of COPD, every effort should be made to provide access to standardized spirometry. Chronic cough and sputum production often precede the development of airflow limitation by many years, although not all individuals with cough and sputum production go on to develop COPD.

CLASSIFICATION OF SEVERITY

For educational reasons, a simple classification of disease severity into four stages is recommended (Table 2). The management of COPD is largely symptom driven, and there is only an imperfect relationship between the degree of airflow limitation and the presence of symptoms. The staging, therefore, is a pragmatic approach aimed at practical implementation and should only be regarded as an educational tool, and a very general indication of the approach to management. All FEV₁ values refer to post-bronchodilator FEV₁.

Stage 0: At Risk - Characterized by chronic cough and sputum production. Lung function, as measured by spirometry, is still normal.

Stage I: Mild COPD - Characterized by mild airflow limitation (FEV₁/FVC < 70% but FEV₁ ≥ 80% predicted) and usually, but not always, by chronic cough and sputum production. At this stage, the individual may not even be aware that his or her lung function is abnormal.

Stage II: Moderate COPD - Characterized by worsening airflow limitation (50% ≤ FEV₁ < 80% predicted) and usually the progression of symptoms, with shortness of breath typically developing on exertion. This is the stage at which patients typically seek medical attention because of dyspnea or an exacerbation of their disease.

Table 2 - Classification of Severity*

Stage	Characteristics
0: At Risk	<ul style="list-style-type: none"> • normal spirometry • chronic symptoms (cough, sputum production)
I: Mild COPD	<ul style="list-style-type: none"> • FEV₁/FVC < 70% • FEV₁ ≥ 80% predicted • with or without chronic symptoms (cough, sputum production)
II: Moderate COPD	<ul style="list-style-type: none"> • FEV₁/FVC < 70% • 50% ≤ FEV₁ < 80% predicted • with or without chronic symptoms (cough, sputum production)
III: Severe COPD	<ul style="list-style-type: none"> • FEV₁/FVC < 70% • 30% ≤ FEV₁ < 50% predicted with or without chronic symptoms (cough, sputum production)
IV: Very Severe COPD	<ul style="list-style-type: none"> • FEV₁/FVC < 70% • FEV₁ < 30% predicted or FEV₁ < 50% predicted plus chronic respiratory failure

*Classification based on postbronchodilator FEV₁
 FEV₁: forced expiratory volume in one second; FVC: forced vital capacity; respiratory failure: arterial partial pressure of oxygen (PaO₂) less than 8.0 kPa (60 mm Hg) with or without arterial partial pressure of CO₂ (PaCO₂) greater than 6.7 kPa (50 mm Hg) while breathing air at sea level.

Stage III: Severe COPD - characterized by further worsening of airflow limitation ($30\% \leq FEV_1 < 50\%$ predicted), increased shortness of breath, and repeated exacerbations which have an impact on patients' quality of life.

Stage IV: Very Severe COPD - Characterized by severe air-flow limitation ($FEV_1 < 30\%$ predicted) or the presence of chronic respiratory failure. Patients may have very severe (Stage IV) COPD even if the FEV_1 is $> 30\%$ predicted, whenever these complications are present. At this stage, quality of life is appreciably impaired and exacerbations may be life-threatening.

Poorly reversible airflow limitation associated with bronchiectasis, cystic fibrosis, tuberculosis, or asthma is not included except insofar as these conditions overlap with COPD. In many developing countries both pulmonary tuberculosis and COPD are common. Therefore, in all subjects with symptoms of COPD, a possible diagnosis of tuberculosis should be considered especially in areas where this disease is known to be prevalent. In countries in which the prevalence of tuberculosis is greatly diminished, the possible diagnosis of this disease is sometimes overlooked.

PATHOGENESIS

COPD is characterized by chronic inflammation throughout the airways, parenchyma, and pulmonary vasculature. Macrophages, T lymphocytes (predominately CD8⁺), and neutrophils are increased in various parts of the lung. Activated inflammatory cells release a variety of mediators – including leukotriene B₄ (LTB₄)⁴, interleukin 8 (IL-8)⁵⁻⁷, tumor necrosis factor α (TNF- α)^{5,8}, and others – capable of damaging lung structures and/or sustaining neutrophilic inflammation. In addition to inflammation, two other processes thought to be important in the pathogenesis of COPD are an imbalance of proteinases and antiproteinases in the lung, and oxidative stress.

Inflammation of the lungs is caused by exposure to inhaled noxious particles and gases. Cigarette smoke can induce inflammation and directly damage the lungs⁹⁻¹⁴. Although fewer data are available, it is likely that other COPD risk factors initiate a similar inflammatory process¹⁵⁻¹⁹. It is believed that this inflammation can then lead to COPD.

PATHOLOGY

Pathological changes characteristic of COPD are found in the central airways, peripheral airways, lung parenchyma, and pulmonary vasculature.

In the central airways – the trachea, bronchi, and bronchioles greater than 2-4 mm in internal diameter – inflammatory cells infiltrate the surface epithelium^{9,20,21}. Enlarged mucus secreting glands and an increase in the number of goblet cells are associated with mucus hypersecretion. In the peripheral airways – small bronchi and bronchioles that have an internal diameter of less than 2 mm – chronic inflammation leads to repeated cycles of injury and repair of the airway wall²². The repair process results in a structural remodeling of the airway wall, with increasing collagen content and scar tissue formation, that narrows the lumen and produces fixed airways obstruction²³.

Destruction of the lung parenchyma in COPD patients typically occurs as centrilobular emphysema. This involves dilatation and destruction of the respiratory bronchioles²⁴. These lesions occur more frequently in the upper lung regions in milder cases, but in advanced disease they may appear diffusely throughout the entire lung and also involve destruction of the pulmonary capillary bed. An imbalance of endogenous proteinases and antiproteinases in the lung – due to genetic factors or the action of inflammatory cells and mediators – is thought to be a major mechanism behind emphysematous lung destruction. Oxidative stress, another consequence of inflammation, may also contribute²⁵.

Pulmonary vascular changes in COPD are characterized by a thickening of the vessel wall that begins early in the natural history of the disease. Thickening of the intima is the first structural change²⁶, followed by an increase in smooth muscle and the infiltration of the vessel wall by inflammatory cells²⁷. As COPD worsens, greater amounts of smooth muscle, proteoglycans, and collagen²⁸ further thicken the vessel wall.

PATHOPHYSIOLOGY

Pathological changes in the lungs lead to corresponding physiological changes characteristic of the disease, including mucus hypersecretion, ciliary dysfunction, airflow limitation, pulmonary hyperinflation, gas exchange abnormalities, pulmonary hypertension, and cor pulmonale. They usually develop in this order over the course of the disease.

Mucus hypersecretion and ciliary dysfunction lead to chronic cough and sputum production. These symptoms can be present for many years before other symptoms or physiological abnormalities develop.

Expiratory airflow limitation, best measured through spirometry, is the hallmark physiological change of COPD³ and the key to diagnosis of the disease. It is primarily

due to fixed airways obstruction and the consequent increase in airways resistance. Destruction of alveolar attachments, which inhibits the ability of the small airways to maintain patency, plays a smaller role.

In advanced COPD, peripheral airways obstruction, parenchymal destruction, and pulmonary vascular abnormalities reduce the lung's capacity for gas exchange, producing hypoxemia and, later on, hypercapnia. Pulmonary hypertension, which develops late in the course of COPD (*Stage IV: Very Severe COPD*), is the major cardiovascular complication of COPD and is associated with the development of cor pulmonale and a poor prognosis²⁹. The prevalence and natural history of cor pulmonale in COPD are not yet clear.

2. BURDEN OF COPD

EPIDEMIOLOGY

Most of the information available on COPD prevalence, morbidity, and mortality comes from developed countries. Even in these countries, accurate epidemiological data on COPD are difficult and expensive to collect.

Prevalence and morbidity data greatly underestimate the total burden of COPD because the disease is usually not diagnosed until it is clinically apparent and moderately advanced. The imprecise and variable definitions of COPD have made it hard to quantify the morbidity and mortality of this disease in developed³⁰ and developing countries. Mortality data also underestimate COPD as a cause of death because the disease is more likely to be cited as a contributory than as an underlying cause of death, or may not be cited at all³¹.

Prevalence: In the Global Burden of Disease Study conducted under the auspices of the WHO and the World Bank^{2,32}, the worldwide prevalence of COPD in 1990 was estimated to be 9.34/1,000 in men and 7.33/1,000 in women. However, these estimates include all ages and underestimate the true prevalence of COPD in older adults. The prevalence of COPD is highest in countries where cigarette smoking has been, or still is, very common, while the prevalence is lowest in countries where smoking is less common, or total tobacco consumption per individual is low.

Morbidity: The limited data that are available indicate that morbidity due to COPD increases with age and is greater in men than women¹. COPD is responsible for a significant part of physician visits, emergency department visits, and hospitalizations.

Table 3 - Four-Country Comparison of COPD Direct and Indirect Costs

Country (ref)	Year	Direct Cost (US\$ Millions)	Indirect Cost (US\$ Millions)	Total (US\$ Millions)	Per Capita* (US\$)
UK ³³	1996	778	3,312	4,090	65
The Netherlands ³⁴	1993	256	N/A	N/A	N/A [#]
Sweden ³⁵	1991	179	281	460	60
US ¹	1993	14,700	9,200	23,900	87

* Per capita valuation based on 1993 population estimates from the United Nations Population Council and expressed in 1993 US dollars.
The authors did not provide estimates of indirect costs.

Mortality: COPD is currently the fourth leading cause of death in the world², and further increases in the prevalence and mortality of the disease can be predicted in the coming decades^{2,32}. In the US, COPD death rates are very low among people under age 45 but then increase with age, and COPD becomes the fourth or fifth leading cause of death among those over 45¹.

ECONOMIC AND SOCIAL BURDEN OF COPD

Table 3 provides an understanding of the relative economic burden of COPD in four countries with Western styles of medical practice and social or private insurance structures. Similar data from developing countries are not available.

The Global Burden of Disease Study^{2,32} estimated the fraction of mortality and disability attributable to major diseases and injuries using a composite measure of the burden of each health problem, the Disability-Adjusted Life Year (DALY= the sum of years lost because of pre-

Table 4 - Leading Causes of Disability-Adjusted Life Years (DALYs) Lost Worldwide: 1990 and 2020 (Projected)^{2,32}

Disease or Injury	Rank 1990	Percent of Total DALYs	Rank 2020	Percent of Total DALYs
Lower respiratory infections	1	8.2	6	3.1
Diarrheal diseases	2	7.2	9	2.7
Perinatal period conditions	3	6.7	11	2.5
Unipolar major depression	4	3.7	2	5.7
Ischemic heart disease	5	3.4	1	5.9
Cerebrovascular disease	6	2.8	4	4.4
Tuberculosis	7	2.8	7	3.1
Measles	8	2.6	25	1.1
Road traffic accidents	9	2.5	3	5.1
Congenital anomalies	10	2.4	13	2.2
Malaria	11	2.3	19	1.5
COPD	12	2.1	5	4.1
Trachea, bronchus, lung cancer	33	0.6	15	1.8

Excerpted with permission from Murray CJL, Lopez AD. *Science* 1999; 274:740-3. Copyright 1999 American Association for the Advancement of Science

mature mortality and years of life lived with disability, adjusted for the severity of disability). According to projections, COPD will be the fifth leading cause of DALYs lost worldwide in 2020 (in 1990 it ranked twelfth), behind ischemic heart disease, major depression, traffic accidents, and cerebrovascular disease (**Table 4**).

RISK FACTORS

Risk factors for COPD include both host factors and environmental exposures, and the disease usually arises from an interaction between these two types of factors. The host factor that is best documented is a rare hereditary deficiency of alpha-1 antitrypsin. Other genes involved in the pathogenesis of COPD have not yet been identified. The major environmental factors are tobacco smoke; heavy exposure to occupational dusts and chemicals (vapors, irritants, fumes); and indoor/outdoor air pollution.

The role of gender as a risk factor for COPD remains unclear. In the past, most studies showed that COPD prevalence and mortality were greater among men than women³⁶⁻³⁹. More recent studies^{1,40} from developed countries show that the prevalence of the disease is almost equal in men and women, which probably reflects changing patterns of tobacco smoking. Some studies have in fact suggested that women are more susceptible to the effects of tobacco smoke than men^{38,41}. This is an important question given the increasing rate of smoking among women in both developed and developing countries.

Host Factors

Genes: It is believed that many genetic factors increase (or decrease) a person's risk of developing COPD. The genetic risk factor that is best documented is a rare hereditary deficiency of alpha-1 antitrypsin⁴²⁻⁴⁴. Premature and accelerated development of panlobular emphysema and decline in lung function occurs in many smokers and nonsmokers with the severe deficiency, although smoking increases the risk appreciably. Other genes involved in the pathogenesis of COPD have not yet been identified.

Airway Hyperresponsiveness: Asthma and airway hyperresponsiveness, identified as risk factors that contribute to the development of COPD⁴⁵, are complex disorders related to a number of genetic and environmental factors. How they influence the development of COPD is unknown. Airway hyperresponsiveness may also develop after exposure to tobacco smoke or other environmental insults and thus may be a result of smoking-related airways disease.

Lung Growth: Lung growth is related to processes occurring during gestation, birth weight, and exposures during childhood⁴⁶⁻⁵⁰. Reduced maximal attained lung function (as measured by spirometry) may identify individuals who are at increased risk for the development of COPD⁵¹.

Exposures

Tobacco Smoke: Cigarette smokers have a higher prevalence of lung-function abnormalities and respiratory symptoms, a greater annual rate of decline in FEV₁, and higher death rates for COPD than nonsmokers. Pipe and cigar smokers have higher COPD morbidity and mortality rates than nonsmokers, although their rates are lower than those for cigarette smokers⁵². Not all smokers develop clinically significant COPD, which suggests that genetic factors must modify each individual's risk. Passive exposure to cigarette smoke may also contribute to respiratory symptoms and COPD by increasing the lungs' total burden of inhaled particulates and gases^{36,53,54}. Smoking during pregnancy may also pose a risk for the fetus, by affecting lung growth and development *in utero* and possibly the priming of the immune system^{50,55}.

Occupational Dusts and Chemicals: When the exposures are sufficiently intense or prolonged, occupational dusts and chemicals (vapors, irritants, fumes) can cause COPD independently of cigarette smoking and increase the risk of the disease in the presence of concurrent cigarette smoking⁵⁶. Exposure to particulate matter, irritants, organic dusts, and sensitizing agents can cause an increase in airway hyperresponsiveness⁵⁷, especially in airways already damaged by other occupational exposures, cigarette smoke, or asthma.

Outdoor and Indoor Air Pollution: High levels of urban air pollution are harmful to persons with existing heart or lung disease. The role of outdoor air pollution in causing COPD is unclear, but appears to be small when compared with cigarette smoking. Indoor air pollution from biomass fuel, burned for cooking and heating in poorly vented dwellings, has been implicated as a risk factor for the development of COPD⁵⁸⁻⁶⁷.

Infections: A history of severe childhood respiratory infection has been associated with reduced lung function and increased respiratory symptoms in adulthood⁴⁵. However, viral infections may be related to another factor, e.g., low birth weight, that itself is related to COPD.

Socioeconomic Status: There is evidence that the risk of developing COPD is inversely related to socioeconomic

status⁶⁸. It is not clear, however, whether this pattern reflects exposures to indoor and outdoor air pollutants, crowding, poor nutrition, or other factors that are related to socioeconomic status^{67,69}.

3. THE FOUR COMPONENTS OF COPD MANAGEMENT

INTRODUCTION

An effective COPD management plan includes four components: (1) Assess and Monitor Disease; (2) Reduce Risk Factors; (3) Manage Stable COPD; (4) Manage Exacerbations.

The goals of effective COPD management are to:

- Prevent disease progression
- Relieve symptoms
- Improve exercise tolerance
- Improve health status
- Prevent and treat complications
- Prevent and treat exacerbations
- Reduce mortality.

These goals should be reached with a minimum of side effects from treatment, a particular challenge in COPD patients where comorbidities are common. The extent to which these goals can be realized varies with each individual, and some treatments will produce benefits in more than one area. In selecting a treatment plan, the benefits and risks to the individual and the costs, direct and indirect, to the community must be considered.

Patients should be identified before the end stage of the illness, when disability is substantial. However, the benefits of community-based spirometric screening, of either the general population or smokers, are still unclear. Educating patients and physicians to recognize that cough, sputum production, and especially breathlessness are not trivial symptoms is an essential aspect of the public health care of this disease.

Reduction of therapy once symptom control has been achieved is not normally possible in COPD. Further deterioration of lung function usually requires the progressive introduction of more treatments, both pharmacologic and non-pharmacologic, to attempt to limit the impact of these changes. Exacerbations of signs and symptoms, a hallmark of COPD, impair patients' quality of life and decrease their health status. Appropriate treatment and measures to prevent further exacerbations should be implemented as quickly as possible.

COMPONENT 1: ASSESS AND MONITOR DISEASE

KEY POINTS

- Diagnosis of COPD is based on a history of exposure to risk factors and the presence of airflow limitation that is not fully reversible, with or without the presence of symptoms.
- Patients who have chronic cough and sputum production with a history of exposure to risk factors should be tested for airflow limitation, even if they do not have dyspnea.
- For the diagnosis and assessment of COPD, spirometry is the gold standard as it is the most reproducible, standardized, and objective way of measuring airflow limitation. $FEV_1/FVC < 70\%$ and a postbronchodilator $FEV_1 < 80\%$ predicted confirms the presence of airflow limitation that is not fully reversible.
- Health care workers involved in the diagnosis and management of COPD patients should have access to spirometry.
- Measurement of arterial blood gas tensions should be considered in all patients with $FEV_1 < 40\%$ predicted or clinical signs suggestive of respiratory failure or right heart failure.

DIAGNOSIS

A diagnosis of COPD (**Table 5**) should be considered in any patient who has cough, sputum production, or dyspnea, and/or a history of exposure to risk factors for the disease. The diagnosis is confirmed by an objective measure of airflow limitation, preferably spirometry.

Assessment of Symptoms: Chronic cough, usually the first symptom of COPD to develop⁷⁰, may initially be intermittent, but later is present every day, often throughout the day, and is seldom entirely nocturnal. In some cases, significant airflow limitation may develop without the presence of a cough. Small quantities of tenacious sputum are commonly raised by COPD patients after coughing bouts. Dyspnea is the reason most patients seek medical attention and is a major cause of disability

and anxiety associated with the disease. As lung function deteriorates, breathlessness becomes more intrusive. Wheezing and chest tightness are relatively non-specific symptoms and may vary between days and over the course of a single day. An absence of wheezing or chest tightness does not exclude a diagnosis of COPD.

- Social and family support available to the patient
- Possibilities for reducing risk factors, especially smoking cessation.

Table 5 - Key Indicators for Considering a Diagnosis of COPD

Consider COPD and perform spirometry if any of these indicators are present. These indicators are not diagnostic by themselves, but the presence of multiple key indicators increases the probability of a diagnosis of COPD. Spirometry is needed to establish a diagnosis of COPD.

Chronic cough:	Present intermittently or every day. Often present throughout the day; seldom only nocturnal.
Chronic sputum production:	Any pattern of chronic sputum production may indicate COPD.
Dyspnea that is:	Progressive (worsens over time). Persistent (present every day). Described by the patient as: "increased effort to breathe," "heaviness," "air hunger," or "gasping." Worse on exercise. Worse during respiratory infections.
History of exposure to risk factors, especially:	Tobacco smoke. Occupational dusts and chemicals. Smoke from home cooking and heating fuels.

Physical Examination: Though an important part of patient care, a physical examination is rarely diagnostic in COPD. Physical signs of airflow limitation are rarely present until significant impairment of lung function has occurred^{71,72}, and their detection has a relatively low sensitivity and specificity.

Measurement of Airflow Limitation: To help identify patients earlier in the course of the disease, spirometry should be performed for patients who have chronic cough and sputum production and a history of exposure to risk factors, even if they do not have dyspnea. Spirometry should measure the maximal volume of air forcibly exhaled from the point of maximal inhalation (forced vital capacity, FVC) and the volume of air exhaled during the first second of this maneuver (forced expiratory volume in one second, FEV₁), and the ratio of these two measurements (FEV₁/FVC) should be calculated. Patients with COPD typically show a decrease in both FEV₁ and FVC. The presence of a postbronchodilator FEV₁ < 80% of the predicted value in combination with an FEV₁/FVC < 70% confirms the presence of airflow limitation that is not fully reversible. The FEV₁/FVC on its own is a more sensitive measure of airflow limitation, and an FEV₁/FVC < 70% is considered an early sign of airflow limitation in patients whose FEV₁ remains normal (≥ 80% predicted). This approach to defining airflow limitation is a pragmatic one in view of the fact that universally applicable reference values for FEV₁ and FVC are not available.

Medical History: A detailed medical history of a new patient known or thought to have COPD should assess:

- Exposure to risk factors
- Past medical history, including asthma, allergy, sinusitis or nasal polyps, respiratory infections in childhood, and other respiratory diseases
- Family history of COPD or other chronic respiratory disease
- Pattern of symptom development
- History of exacerbations or previous hospitalizations for respiratory disorder
- Presence of comorbidities, such as heart disease and rheumatic disease, that may also contribute to restriction of activity
- Appropriateness of current medical treatments
- Impact of disease on patient's life, including limitation of activity; missed work and economic impact; effect on family routines; and feelings of depression or anxiety

Assessment of Severity: Assessment of severity (Table 2) is based on the level of symptoms, severity of the spirometric abnormality, and the presence of complications such as respiratory failure and right heart failure.

Additional Investigations: For patients in Stage II: Moderate COPD and beyond, the following additional investigations may be useful.

Bronchodilator reversibility testing: Generally performed only once, at the time of diagnosis, this test is useful to help rule out a diagnosis of asthma, to establish a patient's best attainable lung function, to gauge a patient's prognosis, and to guide treatment decisions. However, even patients who do not show a significant FEV₁ response to a short-acting bronchodilator test may benefit symptomatically from long-term bronchodilator treatment.

Glucocorticosteroid reversibility testing: The simplest, and potentially safest, way to identify patients most likely to respond to long-term glucocorticosteroid treatment is with a treatment trial of inhaled glucocorticosteroids for 6 weeks to 3 months, using as criteria for glucocorticosteroid reversibility an FEV₁ increase of 200 ml and 12% above baseline^{73,74}. The response to glucocorticosteroids should be evaluated with respect to the *post*bronchodilator FEV₁ (that is, the effect of treatment with inhaled glucocorticosteroids should be in addition to that of regular treatment with a bronchodilator).

Chest X-ray: A chest X-ray is seldom diagnostic in COPD unless obvious bullous disease is present, but it is valuable in excluding alternative diagnoses. Computed tomography (CT) of the chest is not routinely recommended. However, when there is doubt about the diagnosis of COPD, high resolution CT (HRCT) might help in the differential diagnosis. In addition, if a surgical procedure such as bullectomy or lung volume reduction is contemplated, chest CT is helpful.

Arterial blood gas measurement: In advanced COPD, measurement of arterial blood gases is important. This test should be performed in patients with FEV₁ < 40% predicted or with clinical signs suggestive of respiratory failure or right heart failure. Clinical signs of respiratory failure or right heart failure include central cyanosis, ankle swelling, and an increase in the jugular venous pressure. Clinical signs of hypercapnia are extremely nonspecific outside of exacerbations. Respiratory failure is indicated by a PaO₂ < 8.0 kPa (60 mm Hg) with or without PaCO₂ > 6.7 kPa (50 mm Hg) while breathing air at sea level. Measurement of arterial blood gases should be obtained by arterial puncture; finger or ear oximeters for assessing arterial oxygen saturation (SaO₂) are less reliable.

Alpha-1 antitrypsin deficiency screening: In patients who develop COPD at a young age (< 45 years) or who have a strong family history of the disease, it may be valuable to identify coexisting alpha-1 antitrypsin deficiency. This could lead to family screening and appropriate counseling.

Differential Diagnosis: A major differential diagnosis is asthma. In some patients with chronic asthma, a clear distinction from COPD is not possible using current imaging and physiological testing techniques. In these cases, current management is similar to that of asthma. Other potential diagnoses are usually easier to distinguish from COPD (**Table 6**).

Table 6 - Differential Diagnosis of COPD

Diagnosis	Suggestive Features*
COPD	Onset in mid-life. Symptoms slowly progressive. Long smoking history. Dyspnea during exercise. Largely irreversible airflow limitation.
Asthma	Onset early in life (often childhood). Symptoms vary from day to day. Symptoms at night/early morning. Allergy, rhinitis, and/or eczema also present. Family history of asthma. Largely reversible airflow limitation.
Congestive Heart Failure	Fine basilar crackles on auscultation. Chest X-ray shows dilated heart, pulmonary edema. Pulmonary function tests indicate volume restriction, not airflow limitation.
Bronchiectasis	Large volumes of purulent sputum. Commonly associated with bacterial infection. Coarse crackles/clubbing on auscultation. Chest X-ray/CT shows bronchial dilation, bronchial wall thickening.
Tuberculosis	Onset all ages. Chest X-ray shows lung infiltrate or nodular lesions. Microbiological confirmation. High local prevalence of tuberculosis.
Obliterative Bronchiolitis	Onset in younger age, nonsmokers. May have history of rheumatoid arthritis or fume exposure. CT on expiration shows hypodense areas.
Diffuse Panbronchiolitis	Most patients are male and nonsmokers. Almost all have chronic sinusitis. Chest X-ray and HRCT show diffuse small centrilobular nodular opacities and hyperinflation.

*These features tend to be characteristic of the respective diseases, but do not occur in every case. For example, a person who has never smoked may develop COPD (especially in the developing world, where other risk factors may be more important than cigarette smoking); asthma may develop in adult and even elderly patients.

ONGOING MONITORING AND ASSESSMENT

Monitor Disease Progression and Development of Complications: COPD is usually a progressive disease, and a patient's lung function can be expected to worsen over time, even with the best available care. Symptoms and objective measures of airflow limitation should be

monitored for development of complications and to determine when to adjust therapy.

Follow-up visits should include a discussion of new or worsening symptoms. Spirometry should be performed if there is a substantial increase in symptoms or a complication. Measurement of arterial blood gas tensions should be performed in all patients with an $FEV_1 < 40\%$ predicted or clinical signs of respiratory failure or right heart failure. Elevation of the jugular venous pressure and the presence of pitting ankle edema are often the most useful findings suggestive of right heart failure in clinical practice. Measurement of pulmonary arterial pressure is not recommended in clinical practice as it does not add practical information beyond that obtained from a knowledge of PaO_2 .

Monitor Pharmacotherapy and Other Medical

Treatment: In order to adjust therapy appropriately as the disease progresses, each follow-up visit should include a discussion of the current therapeutic regimen. Dosages of various medications, adherence to the regimen, inhaler technique, effectiveness of the current regime at controlling symptoms, and side effects of treatment should be monitored.

Monitor Exacerbation History: Frequency, severity, and likely causes of exacerbations should be evaluated. Increased sputum volume, acutely worsening dyspnea, and the presence of purulent sputum should be noted. Severity can be estimated by the increased need for bronchodilator medication or glucocorticosteroids and by the need for antibiotic treatment. Hospitalizations should be documented, including the facility, duration of stay, and any use of critical care or intubation.

Monitor Comorbidities: In treating patients with COPD, it is important to consider the presence of concomitant conditions such as bronchial carcinoma, tuberculosis, sleep apnea, and left heart failure. The appropriate diagnostic tools (chest radiograph, ECG, etc.) should be used whenever symptoms (e.g., hemoptysis) suggest one of these conditions.

COMPONENT 2: REDUCE RISK FACTORS

KEY POINTS

- Reduction of total personal exposure to tobacco smoke, occupational dusts and chemicals, and indoor and outdoor air pollutants are important goals to prevent the onset and progression of COPD.
- Smoking cessation is the single most effective – and cost-effective – intervention to reduce the risk of developing COPD and stop its progression (**Evidence A**).
- Brief tobacco dependence treatment is effective (**Evidence A**) and every tobacco user should be offered at least this treatment at every visit to the health care provider.
- Three types of counseling are especially effective: practical counseling, social support as part of treatment, and social support arranged outside of treatment (**Evidence A**).
- Several effective pharmacotherapies for tobacco dependence are available (**Evidence A**), and at least one of these medications should be added to counseling if necessary and in the absence of contraindications.
- Progression of many occupationally induced respiratory disorders can be reduced or controlled through a variety of strategies aimed at reducing the burden of inhaled particles and gases (**Evidence B**).

SMOKING PREVENTION AND CESSATION

Comprehensive tobacco control policies and programs with clear, consistent, and repeated nonsmoking messages should be delivered through every feasible channel. Legislation to establish smoke-free schools, public facilities, and work environments should be encouraged by government officials, public health workers, and the public.

Smoking cessation is the single most effective – and cost-effective – way to reduce the risk of developing COPD and stop its progression. Even a brief, 3-minute period of counseling to urge a smoker to quit can be effective, and at the very least this should be done for every smoker at every visit^{75,76}. Health education, public policy, and information dissemination programs are all vital components in a comprehensive cessation effort.

Guidelines for Smoking Cessation: Guidelines for smoking cessation were published by the US Agency for Health Care Policy and Research (AHCPR) in 1996⁷⁷ and updated in 2000 by the US Public Health Service in *Treating Tobacco Use and Dependence: A Clinical Practice Guideline*⁷⁸.

Smoking Cessation Intervention Process: The Public Health Service Report recommends a five-step program for intervention (**Table 7**), which provides a strategic framework helpful to health care providers interested in helping their patients stop smoking. Three types of counseling are especially effective: practical counseling, social support as part of treatment, and social support arranged outside of treatment⁷⁷⁻⁸¹ (**Evidence A**).

Pharmacotherapy: Numerous effective pharmacotherapies for smoking cessation now exist^{78,82,83} (**Evidence A**). Except in the presence of special circumstances, pharma-

cotherapy is recommended when counseling is not sufficient to help patients quit smoking. Numerous studies indicate that nicotine replacement therapy in any form (nicotine gum, inhaler, nasal spray, transdermal patch, sublingual tablet, or lozenge) reliably increases long-term smoking abstinence rates^{78,83}. The antidepressants bupropion⁸⁴ and nortriptyline have also been shown to increase long-term quit rates, although fewer studies have been conducted with these medications^{78,83}. The effectiveness of the antihypertensive drug clonidine is limited by side effects⁸³. Special consideration should be given before using pharmacotherapy in selected populations: people with medical contraindications, light smokers (fewer than 10 cigarettes/day), and pregnant and adolescent smokers.

OCCUPATIONAL EXPOSURES

Although it is not known how many individuals are at risk of developing respiratory disease from occupational exposures in either developing or developed countries, many occupationally induced respiratory disorders can be reduced or controlled through a variety of strategies aimed at reducing the burden of inhaled particles and gases⁸⁵. Emphasis should be on primary prevention, which is best achieved by the elimination or reduction of exposures to various substances in the workplace. Secondary prevention, achieved through epidemiologic surveillance and early case detection, is also of great importance.

INDOOR/OUTDOOR AIR POLLUTION

Individuals experience diverse indoor and outdoor environments throughout the day, each of which has its own unique set of air contaminants. Although outdoor and indoor air pollution are generally thought of separately, the concept of total personal exposure may be more relevant for COPD. Reducing the risk from indoor and outdoor air pollution requires a combination of public policy and protective steps taken by individual patients.

The health care provider should consider susceptibility (including family history, exposure to indoor/outdoor pollution) for each individual patient⁸⁶. Those who are at high risk should avoid vigorous exercise outdoors during pollution episodes. If various solid fuels are used for cooking and heating, adequate ventilation should be encouraged. Persons with severe COPD should monitor public announcements of air quality and should stay indoors when air quality is poor. Under most circumstances, health care providers should not suggest respiratory protection as a method for reducing the risks of ambient air pollution. Air cleaners have not been shown to have health benefits, whether directed at pollutants generated by indoor sources or at those brought in with outdoor air.

Table 7 - Strategies to Help the Patient Willing to Quit Smoking⁷⁸

1. **ASK:** Systematically identify all tobacco users at every visit.
Implement an office-wide system that ensures that, for EVERY patient at EVERY clinic visit, tobacco-use status is queried and documented.
2. **ADVISE:** Strongly urge all tobacco users to quit.
In a clear, strong, and personalized manner, urge every tobacco user to quit.
3. **ASSESS:** Determine willingness to make a quit attempt.
Ask every tobacco user if he or she is willing to make a quit attempt at this time (e.g., within the next 30 days).
4. **ASSIST:** Aid the patient in quitting.
Help the patient with a quit plan; provide practical counseling; provide intra-treatment social support; help the patient obtain extra-treatment social support; recommend use of approved pharmacotherapy except in special circumstances; provide supplementary materials.
5. **ARRANGE:** Schedule follow-up contact.
Schedule follow-up contact, either in person or via telephone.

COMPONENT 3: MANAGE STABLE COPD

KEY POINTS

- The overall approach to managing stable COPD should be characterized by a stepwise increase in treatment, depending on the severity of the disease.
- For patients with COPD, health education can play a role in improving skills, ability to cope with illness, and health status. It is effective in accomplishing certain goals, including smoking cessation (**Evidence A**).
- None of the existing medications for COPD has been shown to modify the long-term decline in lung function that is the hallmark of this disease (**Evidence A**). Therefore, pharmacotherapy for COPD is used to decrease symptoms and/or complications.
- Bronchodilator medications are central to the symptomatic management of COPD (**Evidence A**). They are given on an as-needed basis or on a regular basis to prevent or reduce symptoms.
- The principal bronchodilator treatments are β_2 -agonists, anticholinergics, theophylline, and a combination of one or more of these drugs (**Evidence A**).
- Regular treatment with long-acting bronchodilators is more effective and convenient than treatment with short-acting bronchodilators, but more expensive (**Evidence A**).
- The addition of regular treatment with inhaled glucocorticosteroids to bronchodilator treatment is appropriate for symptomatic COPD patients with an FEV₁ < 50% predicted (Stage III: *Severe COPD* and Stage IV: *Very Severe COPD*) and repeated exacerbations (**Evidence A**).
- Chronic treatment with systemic glucocorticosteroids should be avoided because of an unfavorable benefit-to-risk ratio (**Evidence A**).
- All COPD patients benefit from exercise training programs, improving with respect to both exercise tolerance and symptoms of dyspnea and fatigue (**Evidence A**).
- The long-term administration of oxygen (> 15 hours per day) to patients with chronic respiratory failure has been shown to increase survival (**Evidence A**).

INTRODUCTION

The overall approach to managing stable COPD should be characterized by a stepwise increase in treatment, depending on the severity of the disease. The management strategy is based on an individualized assessment of disease severity and response to various therapies. Disease severity is determined by the severity of symptoms and airflow limitation, as well as other factors such as the frequency and severity of exacerbations, complications, respiratory failure, comorbidities (cardiovascular disease, sleep-related disorders, etc.), and the general health status of the patient. Treatment depends on the patient's educational level and willingness to apply the recommended management, on cultural and local conditions, and on the availability of medications.

EDUCATION

Although patient education alone does not improve exercise performance or lung function⁸⁷⁻⁹⁰, it can play a role in improving skills, ability to cope with illness, and health status⁹¹. In addition, patient education is effective in accomplishing certain specific goals, including smoking cessation⁴¹ (**Evidence A**), initiating discussions and understanding of advance directives and end-of-life issues⁹² (**Evidence B**), and improving patient responses to exacerbations^{93,94} (**Evidence B**).

Education may take place in many settings: consultations with physicians or other health care workers, home-care or outreach programs, and comprehensive pulmonary rehabilitation programs. It should be tailored to the needs and environment of the patient, interactive, directed at improving quality of life, simple to follow, practical, and appropriate to the intellectual and social skills of the patient and the caregiver. The topics that seem most appropriate for an education program to cover include: smoking cessation; basic information about COPD and pathophysiology of the disease; general approach to therapy and specific aspects of medical treatment; self-management skills; strategies to help minimize dyspnea; advice about when to seek help; self-management and decision-making in exacerbations; and advance directives and end-of-life issues.

PHARMACOLOGIC TREATMENT

Pharmacologic therapy (**Table 8**) is used to prevent and control symptoms, reduce the frequency and severity of exacerbations, improve health status, and improve exercise tolerance. None of the existing medications for COPD (**Table 10**) has been shown to modify the long-

term decline in lung function that is the hallmark of this disease^{41,95-98} (**Evidence A**). However, this should not preclude efforts to use medications to control symptoms.

BRONCHODILATORS

Bronchodilator medications are central to the symptomatic management of COPD⁹⁹⁻¹⁰² (**Evidence A**) (**Table 9**). They are given either on an as-needed basis for relief of persistent or worsening symptoms, or on a regular basis to prevent or reduce symptoms. Dose-response relationships using the FEV₁ as the outcome are relatively flat with all classes of bronchodilators. Side effects are pharmacologically predictable and dose-dependent. Adverse effects are less likely, and resolve more rapidly after treatment withdrawal, with inhaled than with oral treatment. When treatment is given by the inhaled route, attention to effective drug delivery and training in inhaler technique is essential.

Bronchodilator drugs commonly used in treating COPD, β_2 -agonists, anticholinergics, and methylxanthines. The

Table 9 - Bronchodilators in Stable COPD
<ul style="list-style-type: none"> • Bronchodilator medications are central to symptom management in COPD. • Inhaled therapy is preferred. • The choice between β_2-agonist, anticholinergic, theophylline, or combination therapy depends on availability and individual response in terms of symptom relief and side effects. • Bronchodilators are prescribed on an as-needed or on a regular basis to prevent or reduce symptoms. • Long-acting inhaled bronchodilators are more effective and convenient, but more expensive. • Combining bronchodilators may improve efficacy and decrease the risk of side effects compared to increasing the dose of a single bronchodilator.

choice depends on the availability of the medication and the patient's response. All categories of bronchodilators have been shown to increase exercise capacity in COPD, without necessarily producing significant changes in FEV₁^{103,104} (**Evidence A**).

Table 8 - Therapy at Each Stage of COPD					
Old (2001)	0: At Risk	I: Mild	II: Moderate		III: Severe
New (2003)	0: At Risk	I: Mild	IIA	IIB	IV: Very Severe
Characteristics	<ul style="list-style-type: none"> • Chronic symptoms • Exposure to risk factors • Normal spirometry 	<ul style="list-style-type: none"> • FEV₁/FVC < 70% • FEV₁ \geq 80% • With or without symptoms 	<ul style="list-style-type: none"> • FEV₁/FVC < 70% • 50% > FEV₁ < 80% • With or without symptoms 	<ul style="list-style-type: none"> • FEV₁/FVC < 70% • 30% > FEV₁ < 50% • With or without symptoms 	<ul style="list-style-type: none"> • FEV₁/FVC < 70% • FEV₁ < 30% or presence of chronic respiratory failure or right heart failure
	Avoidance of risk factor(s); influenza vaccination				
		Add short-acting bronchodilator when needed			
			Add regular treatment with one or more long-acting bronchodilators		
			Add rehabilitation		
				Add inhaled glucocorticosteroids if repeated exacerbations	
					Add long-term oxygen if chronic respiratory failure Consider surgical treatments

Regular treatment with long-acting bronchodilators is more effective and convenient than treatment with short-acting bronchodilators, but more expensive¹⁰⁵⁻¹⁰⁷ (**Evidence A**). Regular use of a long-acting β_2 -agonist¹⁰⁵ or long-acting anticholinergic improves health status¹⁰⁵⁻¹⁰⁷. Theophylline is effective in COPD, but due to its potential toxicity, inhaled bronchodilators are preferred when avail-

able. All studies that have shown efficacy of theophylline in COPD were done with slow-release preparations.

Combining drugs with different mechanisms and durations of action might increase the degree of bronchodilation for equivalent or lesser side effects. A combination of a short-acting β_2 -agonist and an anticholinergic produces

Table 10 - Commonly Used Formulations of Drugs for COPD

Drug	Inhaler (μ g)	Solution for Nebulizer (mg/ml)	Oral	Vials for injection (mg)	Duration of Action (hours)
β_2-agonists					
Short-acting					
Fenoterol	100-200 (MDI)	1	0.05% (Syrup)		4-6
Salbutamol (albuterol)	100, 200 (MDI & DPI)	5	5mg (Pill) Syrup 0.024%	0.1, 0.5	4-6
Terbutaline	400, 500 (DPI)	-	2.5, 5 (Pill)	0.2, 0.25	4-6
Long-acting					
Formoterol	4.5-12 (MDI & DPI)				12+
Salmeterol	25-50 (MDI & DPI)				12+
Anticholinergics					
Short-acting					
Ipratropium bromide	20, 40 (MDI)	0.25-0.5			6-8
Oxipropium bromide	100 (MDI)	1.5			7-9
Long-acting					
Tiotropium	18 (DPI)				+24
Combination short-acting β_2-agonists plus anticholinergic in one inhaler					
Fenoterol/Ipratropium	200/80 (MDI)	1.25/0.5			6-8
Salbutamol/Ipratropium	75/15 (MDI)	0.75/4.5			6-8
Methylxanthines					
Aminophylline			200-600 mg (Pill)	240 mg	Variable, up to 24
Theophylline (SR)			100-600 mg (Pill)		Variable, up to 24
Inhaled glucocorticosteroids					
Beclomethasone	50-400 (MDI & DPI)	0.2-0.4			
Budesonide	100, 200, 400 (DPI)	0.20, 0.25, 0.5			
Fluticasone	50-500 (MDI & DPI)				
Triamcinolone	100 (MDI)	40		40	
Combination long-acting β_2-agonists plus glucocorticosteroids in one inhaler					
Formoterol/Budesonide	4.5/80, 160 (DPI) (9/320) (DPI)				
Salmeterol/Fluticasone	50/100, 250, 500 (DPI) 25/50, 125, 250 (MDI)				
Systemic glucocorticosteroids					
Prednisone			5-60 mg (Pill)		
Methyl-prednisolone	10-2000 mg		4, 8, 16 mg (Pill)		

MDI=metered dose inhaler; DPI=dry powder inhaler

greater and more sustained improvements in FEV₁ than either alone and does not produce evidence of tachyphylaxis over 90 days of treatment¹¹⁰⁸⁻¹¹¹⁰ (**Evidence A**).

Combination of a β_2 -agonist, an anticholinergic, and/or theophylline may produce additional improvements in lung function^{108,111-115} and health status^{104,108,111,116}. Increasing the number of drugs usually increases costs, and an equivalent benefit may occur by increasing the dose of one bronchodilator when side effects are not a limiting factor. Detailed assessments of this approach have not been carried out.

Increasing the dose of either a β_2 -agonist or an anticholinergic, especially when given by a wet nebulizer, appears to provide subjective benefit in acute episodes¹¹⁷ (**Evidence B**). Some patients may request regular treatment with high-dose, nebulized bronchodilators¹¹⁸, especially if they have experienced subjective benefit from this treatment during an exacerbation. Clear scientific evidence for this approach is lacking, but one option is to examine the improvement in mean daily peak expiratory flow recording during 2 weeks of treatment in the home and continue with nebulizer therapy if a significant improvement occurs¹¹⁸. In general, nebulized therapy for a stable patient is not appropriate unless it has been shown to be better than conventional dose therapy.

GLUCOCORTICOSTEROIDS

Regular treatment with inhaled glucocorticosteroids does not modify the long-term decline of FEV₁ in patients with COPD⁹⁵⁻⁹⁸. However, data from four large studies¹¹⁹⁻¹²² provide evidence that regular treatment with inhaled glucocorticosteroids is appropriate for symptomatic COPD patients with an FEV₁ < 50% predicted (*Stage III: Severe COPD* and *Stage IV: Very Severe COPD*) and repeated exacerbations (for example, 3 in the last three years). This treatment has been shown to reduce the frequency of exacerbations and improve health status (**Evidence A**). In three of these studies^{119,121,122}, glucocorticosteroid combined with a long-acting β_2 -agonist was more effective than the individual components (**Evidence A**). The dose-response relationships and long-term safety of inhaled glucocorticosteroids in COPD are not known.

The present guidelines recommend a trial of 6 weeks to 3 months with inhaled glucocorticosteroids to identify COPD patients who may benefit from long-term inhaled glucocorticosteroid therapy. Many existing COPD guidelines recommend the use of a short course (two weeks) of oral glucocorticosteroids to identify COPD patients

who might benefit from long-term treatment with oral or inhaled glucocorticosteroids. There is mounting evidence, however, that a short course of oral glucocorticosteroids is a poor predictor of the long-term response to inhaled glucocorticosteroids in COPD^{97,123}.

Long-term treatment with oral glucocorticosteroids is not recommended in COPD¹²⁴⁻¹²⁶ (**Evidence A**). There is no evidence of long-term benefit from this treatment. Moreover, a side effect of long-term treatment with systemic glucocorticosteroids is steroid myopathy^{125,126}, which contributes to muscle weakness, decreased functionality, and respiratory failure in patients with advanced COPD.

OTHER PHARMACOLOGIC TREATMENTS

Vaccines: Influenza vaccines can reduce serious illness and death in COPD patients by about 50%¹²⁷. Vaccines containing killed or live, inactivated viruses are recommended¹²⁸, and should be given once (in autumn) or twice (in autumn and winter) each year (**Evidence A**). A pneumococcal vaccine containing 23 virulent serotypes has been used but sufficient data to support its general use in COPD patients are lacking¹²⁹⁻¹³¹ (**Evidence B**).

Alpha-1 Antitrypsin Augmentation Therapy: Young patients with severe hereditary alpha-1 antitrypsin deficiency and established emphysema may be candidates for alpha-1 antitrypsin augmentation therapy. However, this therapy is very expensive, is not available in most countries, and is not recommended for COPD that is unrelated to alpha-1 antitrypsin deficiency (**Evidence C**).

Antibiotics: The use of antibiotics, other than in treating infectious exacerbations of COPD and other bacterial infections, is not recommended^{132,133} (**Evidence A**).

Mucolytic (Mucokinetic, Mucoregulator) Agents (ambroxol, erdosteine, carbocysteine, iodinated glycerol): Although a few patients with viscous sputum may benefit from mucolytics^{134,135}, the overall benefits seem to be very small. Therefore, the widespread use of these agents cannot be recommended on the basis of the present evidence (**Evidence D**).

Antioxidant Agents: Antioxidants, in particular N-acetylcysteine, have been shown to reduce the frequency of exacerbations and could have a role in the treatment of patients with recurrent exacerbations¹³⁶⁻¹³⁹

(**Evidence B**). However, before their routine use can be recommended, the results of ongoing trials will have to be carefully evaluated.

Immunoregulators (Immunostimulators, Immunomodulators): A study using an immunostimulator in COPD showed a decrease in the severity (though not in the frequency) of exacerbations¹⁴⁰, but these results have not been duplicated. Thus, the regular use of this therapy cannot be recommended based on the present evidence¹⁴¹ (**Evidence B**).

Antitussives: Cough, although sometimes a troublesome symptom in COPD, has a significant protective role¹⁴². Thus, the regular use of antitussives is contraindicated in stable COPD (**Evidence D**).

Vasodilators: In patients with stable COPD, inhaled nitric oxide can worsen gas exchange because of altered hypoxic regulation of ventilation-perfusion balance^{143,144} and thus is contraindicated.

Respiratory Stimulants: The use of doxapram, a non-specific respiratory stimulant available as an intravenous formulation, is not recommended in stable COPD (**Evidence D**). Almitrine bismesylate is not recommended for regular use in stable COPD patients¹⁴⁵⁻¹⁴⁷ (**Evidence B**).

Narcotics: The use of oral and parenteral opioids are effective for treating dyspnea in COPD patients with advanced disease. There are insufficient data to conclude whether nebulized opioids are effective¹⁴⁸. However, there are some clinical studies suggesting that morphine used to control dyspnea may have serious adverse effects and its benefits may be limited to a few sensitive subjects¹⁴⁹⁻¹⁵³.

Others: Nedocromil, leukotriene modifiers, and alternative healing methods (e.g., herbal medicine, acupuncture, homeopathy) have not been adequately tested in COPD patients and thus cannot be recommended at this time.

NON-PHARMACOLOGIC TREATMENT

REHABILITATION

The principal goals of pulmonary rehabilitation are to reduce symptoms, improve quality of life, and increase physical and emotional participation in everyday activities. To accomplish these goals, pulmonary rehabilitation covers a range of non-pulmonary problems

including exercise deconditioning, relative social isolation, altered mood states (especially depression), muscle wasting, and weight loss. COPD patients at all stages of disease benefit from exercise training programs, improving with respect to both exercise tolerance and symptoms of dyspnea and fatigue¹⁵⁴ (**Evidence A**). The minimum length of an effective rehabilitation program is two months; the longer the program continues, the more effective the results¹⁵⁵⁻¹⁵⁷ (**Evidence B**). However, as yet, no effective structure has been developed to maintain the effects over time¹⁵⁸. Benefits have been reported from rehabilitation programs conducted in inpatient, outpatient, and home settings¹⁵⁹⁻¹⁶¹.

Ideally, pulmonary rehabilitation should involve several types of health professionals. A comprehensive pulmonary rehabilitation program includes exercise training, nutrition counseling, and education. Baseline and outcome assessments of each participant in a pulmonary rehabilitation program should be made to quantify individual gains and target areas for improvement and should include:

- Detailed medical history and physical examination
- Measurement of spirometry before and after a bronchodilator drug
- Assessment of exercise capacity
- Measurement of health status and the impact of breathlessness
- Assessment of inspiratory and expiratory muscle strength and lower limb strength (e.g., quadriceps) in patients who suffer from muscle wasting (optional).

The first two assessments are important for establishing entry suitability and baseline status but are not used in outcome assessment. The last three assessments are baseline and outcome measures.

OXYGEN THERAPY

The long-term administration of oxygen (>15 hours per day) to patients with chronic respiratory failure has been shown to increase survival¹⁶²⁻¹⁶⁴ (**Evidence A**). It can also have a beneficial impact on hemodynamics, hematologic characteristics, exercise capacity, lung mechanics, and mental state¹⁶⁴.

Long-term oxygen therapy is generally introduced in *Stage IV: Very Severe COPD* for patients who have:

- PaO₂ at or below 7.3 kPa (55 mm Hg) or SaO₂ at or below 88%, with or without hypercapnia; or

- PaO₂ between 7.3 kPa (55 mm Hg) and 8.0 kPa (60 mm Hg) or SaO₂ 89%, if there is evidence of pulmonary hypertension, peripheral edema suggesting congestive heart failure, or polycythemia (hematocrit > 55%).

The goal of long-term oxygen therapy is to increase the baseline PaO₂ to at least 8.0 kPa (60 mm Hg) at sea level and rest, and/or produce SaO₂ at least 90%, which will preserve vital organ function by ensuring an adequate delivery of oxygen.

A decision about the use of long-term oxygen should be based on the waking PaO₂ values. The prescription should always include the source of supplemental oxygen (gas or liquid), the method of delivery, duration of use, and the flow rate at rest, during exercise, and during sleep.

VENTILATORY SUPPORT

To date there is no convincing evidence that mechanical ventilatory support has a role in the routine management of stable COPD.

SURGICAL TREATMENTS

Bullectomy: In carefully selected patients, this procedure is effective in reducing dyspnea and improving lung function¹⁶⁵ (**Evidence C**). A thoracic CT scan, arterial blood gas measurement, and comprehensive respiratory function tests are essential before making a decision regarding a patient's suitability for resection of a bulla.

Lung Volume Reduction Surgery (LVRS): Although there are some encouraging reports (**Evidence C**), LVRS is still an unproven palliative surgical procedure^{166,167}. Several large randomized studies are now underway to investigate the effectiveness and cost of LVRS in comparison to vigorous conventional therapy¹⁶⁹. Patients with an FEV₁ < 20 % predicted and either homogeneous emphysema on HRCT or a DLCO < 20 % predicted were shown to be at high risk for death after LVRS and unlikely to benefit from the intervention¹⁶⁸. Until additional results from these studies are known, LVRS cannot be recommended for widespread use.

Lung Transplantation: In appropriately selected patients with very advanced COPD, lung transplantation has been shown to improve quality of life and functional capacity¹⁶⁰⁻¹⁶³ (**Evidence C**). Criteria for referral for lung transplantation include FEV₁ < 35% predicted, PaO₂ < 7.3-8.0 kPa (55-60 mm Hg), PaCO₂ > 6.7 kPa (50 mm Hg), and secondary pulmonary hypertension¹⁷⁰.

COMPONENT 4: MANAGE EXACERBATIONS

KEY POINTS

- Exacerbations of respiratory symptoms requiring medical intervention are important clinical events in COPD.
- The most common causes of an exacerbation are infection of the tracheobronchial tree and air pollution, but the cause of about one-third of severe exacerbations cannot be identified (**Evidence B**).
- Inhaled bronchodilators (particularly inhaled β₂-agonists and/or anticholinergics), theophylline, and systemic, preferably oral, glucocorticosteroids are effective treatments for exacerbations of COPD (**Evidence A**).
- Patients experiencing COPD exacerbations with clinical signs of airway infection (e.g., increased volume and change of color of sputum, and/or fever) may benefit from antibiotic treatment (**Evidence B**).
- Noninvasive intermittent positive pressure ventilation (NIPPV) in exacerbations improves blood gases and pH, reduces in-hospital mortality, decreases the need for invasive mechanical ventilation and intubation, and decreases the length of hospital stay (**Evidence A**).

COPD is often associated with exacerbations of symptoms¹⁷¹⁻¹⁷⁴. The economic and social burden of COPD exacerbations is extremely high. The most common causes of an exacerbation are infection of the tracheobronchial tree¹⁷⁵⁻¹⁷⁹ and air pollution¹⁸⁰, but the cause of about one-third of severe exacerbations cannot be identified. The role of bacterial infections, once believed to be the main cause of COPD exacerbations, is controversial^{175-179,181-184}. Conditions that may mimic the symptoms of an exacerbation include pneumonia, congestive heart failure, pneumothorax, pleural effusion, pulmonary embolism, and arrhythmia.

DIAGNOSIS AND ASSESSMENT OF SEVERITY

Increased breathlessness, the main symptom of an exacerbation, is often accompanied by wheezing and

chest tightness, increased cough and sputum, change of the color and/or tenacity of sputum, and fever. Exacerbations may also be accompanied by a number of nonspecific complaints, such as malaise, insomnia, sleepiness, fatigue, depression, and confusion. A decrease in exercise tolerance, fever, and/or new radiological anomalies suggestive of pulmonary disease may herald a COPD exacerbation. An increase in sputum volume and purulence points to a bacterial cause, as does a prior history of chronic sputum production¹⁷⁹.

The assessment of the severity of an exacerbation is based on the patient's medical history before the exacerbation, symptoms, physical examination, lung function tests, arterial blood gas measurements, and other laboratory tests. The medical history should cover how long worsening or new symptoms have been present, the frequency and severity of breathlessness and coughing attacks, sputum volume and color, limitation of daily activities, any previous episodes/exacerbations and whether they required hospitalization, and the present treatment regimen. When available, prior measurements of lung function and arterial blood gases are extremely useful for comparison with those made during the acute episode, as an acute change in these tests is more important than their absolute values. In patients with very severe COPD, the most important sign of severe exacerbation is a change in alertness of the patient and this signals a need for immediate evaluation in the hospital.

Lung Function Tests: Even simple lung function tests can be difficult for a sick patient to perform properly. In general, a PEF < 100 L per minute or an FEV₁ < 1.00 L indicates a severe exacerbation¹⁸⁵⁻¹⁸⁷.

Assessment of Arterial Blood Gases: In the hospital, measurement of arterial blood gases is essential to assess the severity of an exacerbation. A PaO₂ < 8.0 kPa (60 mm Hg) and/or SaO₂ < 90% with or without PaCO₂ > 6.7 kPa, 50 mmHg (when breathing room air) indicates respiratory failure. In addition, PaO₂ < 6.7 kPa (50 mm Hg), PaCO₂ > 9.3 kPa (70 mm Hg), and pH < 7.30 point towards a life-threatening episode that needs close monitoring or critical management¹⁸⁸.

Chest X-ray and ECG: Chest radiographs (posterior/anterior plus lateral) are useful in identifying alternative diagnoses that can mimic the symptoms of an exacerbation. An ECG aids in the diagnosis of right ventricular hypertrophy, arrhythmias, and ischemic episodes. Pulmonary embolism can be very difficult to distinguish from an exacerbation, especially in severe COPD, because right ventricular hypertrophy and large

pulmonary arteries lead to confusing ECG and radiographic results. Spiral CT scanning and angiography and perhaps specific D-dimer assays are the best tools presently available for diagnosis of pulmonary embolism in patients with COPD but ventilation-perfusion scanning is of no value. A low systolic blood pressure and an inability to increase the PaO₂ above 8.0 kPa (60 mm Hg) despite high-flow oxygen also suggest pulmonary embolism. If there are strong indications that pulmonary embolism has occurred, it is best to treat for this along with the exacerbation.

Other Laboratory Tests: The whole blood count may identify polycythemia (hematocrit > 55%) or bleeding. White blood cell counts are usually not very informative. The presence of purulent sputum during an exacerbation of symptoms is sufficient indication for starting antibiotic treatment. *Streptococcus pneumoniae*, *Hemophilus influenzae*, and *Moraxella catarrhalis* are the most common bacterial pathogens involved in COPD exacerbations. If an infectious exacerbation does not respond to initial antibiotic treatment, a sputum culture and an antibiogram should be performed. Biochemical tests can reveal whether the cause of the exacerbation is an electrolyte disturbance(s) (hyponatremia, hypokalemia, etc.), a diabetic crisis, or poor nutrition (low proteins), and may suggest a metabolic acid-base disorder.

HOME MANAGEMENT

There is increasing interest in home care for end-stage COPD patients, although economic studies of home care services have yielded mixed results. A major outstanding issue is when to treat an exacerbation at home and when to hospitalize the patient.

Bronchodilator Therapy: Home management of COPD exacerbations involves increasing the dose and/or frequency of existing bronchodilator therapy (**Evidence A**). If not already used, an anticholinergic can be added until the symptoms improve. In more severe cases, high-dose nebulized therapy can be given on an as-needed basis for several days if a suitable nebulizer is available. However, long-term use of nebulizer therapy after an acute episode is not routinely recommended.

Glucocorticosteroids: Systemic glucocorticosteroids are beneficial in the management of exacerbations of COPD. They shorten recovery time and help to restore lung function more quickly¹⁸⁹⁻¹⁹¹ (**Evidence A**). They should be considered in addition to bronchodilators if the patient's baseline FEV₁ is < 50% predicted. A dose of 40 mg prednisolone per day for 10 days is recommended

(Evidence D). One large study indicates that nebulized budesonide may be an alternative to oral glucocorticosteroids in the treatment of nonacidotic exacerbations¹⁹².

Antibiotics: Antibiotics are only effective when patients with worsening dyspnea and cough also have increased sputum volume and purulence¹⁷⁴ (**Evidence B**). The choice of agents should reflect local patterns of antibiotic sensitivity among *S. pneumoniae*, *H. influenzae*, and *M. catarrhalis*.

HOSPITAL MANAGEMENT

The risk of dying from an exacerbation of COPD is closely related to the development of respiratory acidosis, the presence of significant comorbidities, and the need for ventilatory support¹⁹³. Patients lacking these features are not at high risk of dying, but those with severe underlying COPD often require hospitalization in any case. Attempts at managing such patients entirely in the community have met with only limited success¹⁹⁴, but returning them to their homes with increased social

Table 11 - Indications for Hospital Assessment or Admission for Exacerbations of COPD*	
<ul style="list-style-type: none"> • Marked increase in intensity of symptoms, such as sudden development of resting dyspnea. • Severe background COPD. • Onset of new physical signs (e.g., cyanosis, peripheral edema). • Failure of exacerbation to respond to initial medical management. • Significant comorbidities. • Newly occurring arrhythmias. • Diagnostic uncertainty. • Older age. • Insufficient home support. 	

* Local resources need to be considered

Table 12 - Indications for ICU Admission of Patients with Exacerbations of COPD*	
<ul style="list-style-type: none"> • Severe dyspnea that responds inadequately to initial emergency therapy. • Confusion, lethargy, coma. • Persistent or worsening hypoxemia (PaO₂ < 5.3 kPa, 40 mm Hg), and/or severe/worsening hypercapnia (PaCO₂ > 8.0 kPa, 60 mm Hg), and/or severe/worsening respiratory acidosis (pH < 7.25) despite supplemental oxygen and NIPPV. 	

* Local resources need to be considered

support and a supervised medical care package after an initial emergency room assessment has been much more successful¹⁹⁵. However, detailed cost-benefit analyses of these approaches are awaited.

A range of criteria to consider for hospital assessment/admission for exacerbations of COPD are shown in **Table 11**. Some patients need immediate admission to an

Table 13 - Management of Severe but Not Life-threatening Exacerbations of COPD in the Emergency Department or the Hospital*	
<ul style="list-style-type: none"> • Assess severity of symptoms, blood gases, chest X-ray. • Administer controlled oxygen therapy and repeat arterial blood gas measurement after 30 minutes. • Bronchodilators: <ul style="list-style-type: none"> — Increase doses or frequency. — Combine β₂-agonists and anticholinergics. — Use spacers or air-driven nebulizers. — Consider adding intravenous methylxanthine, if needed. • Add glucocorticosteroids <ul style="list-style-type: none"> — Oral or intravenous. • Consider antibiotics <ul style="list-style-type: none"> — When signs of bacterial infection, oral or occasionally intravenous. • Consider noninvasive mechanical ventilation. • At all times: <ul style="list-style-type: none"> — Monitor fluid balance and nutrition. — Consider subcutaneous heparin. — Identify and treat associated conditions (e.g., heart failure, arrhythmias). — Closely monitor condition of the patient. 	

* Local resources need to be considered

intensive care unit (ICU) (**Table 12**). Admission of patients with severe COPD exacerbations to intermediate or special respiratory care units may be appropriate if personnel, skills, and equipment are available to identify and manage acute respiratory failure successfully.

The first actions when a patient reaches the emergency department are to provide controlled oxygen therapy and to determine whether the exacerbation is life-threatening. If so, the patient should be admitted to the ICU immediately.

Otherwise, the patient may be managed in the emergency department or hospital as detailed in **Table 13**.

Controlled Oxygen Therapy: Oxygen therapy is the cornerstone of hospital treatment of COPD exacerbations. Adequate levels of oxygenation ($\text{PaO}_2 > 8.0$ kPa, 60 mm Hg or $\text{SaO}_2 > 90\%$) are easy to achieve in uncomplicated exacerbations, but CO_2 retention can occur insidiously with little change in symptoms. Once oxygen is started, arterial blood gases should be checked 30 minutes later to ensure satisfactory oxygenation without CO_2 retention or acidosis. Venturi masks are more accurate sources of controlled oxygen than are nasal prongs but are more likely to be removed by the patient.

Bronchodilator Therapy: Short-acting, inhaled β_2 -agonists are usually the preferred bronchodilators for the treatment of exacerbations of COPD^{85,133,134} (**Evidence A**). If a prompt response to these drugs does not occur, the addition of an anticholinergic is recommended, even though evidence concerning the effectiveness of this combination is rather controversial^{196,197}. Despite its widespread clinical use, the role of aminophylline in the treatment of COPD exacerbations remains controversial. Most studies of aminophylline have demonstrated minor improvements in lung volumes without showing gas exchange deterioration^{198,199}. In more severe exacerbations, addition of an oral or intravenous methylxanthine to the treatment can be considered. However, close monitoring of serum theophylline is recommended to avoid the side effects of these drugs^{198,200,201}.

Glucocorticosteroids: Oral or intravenous glucocorticosteroids are recommended as an addition to bronchodilator therapy (plus eventually antibiotics and oxygen therapy) in the hospital management of exacerbations of COPD¹⁸⁹⁻¹⁹¹ (**Evidence A**). The exact dose that should be given is not known, but high doses are associated with a significant risk of side effects. Thirty to 40 mg of oral prednisolone daily for 10 to 14 days is a reasonable compromise between efficacy and safety (**Evidence D**). Prolonged treatment does not result in a greater efficacy and increases the risk of side effects.

Antibiotics: Antibiotics are only effective when patients with worsening dyspnea and cough also have increased sputum volume and purulence¹⁴¹. The choice of agents should reflect local patterns of antibiotic sensitivity among *S. pneumoniae*, *H. influenzae*, and *M. catarrhalis*.

Ventilatory Support: The primary objectives of mechanical support in patients with exacerbations in *Stage IV: Very Severe COPD* are to decrease mortality and

morbidity and to relieve symptoms. Ventilatory support includes both noninvasive mechanical ventilation using either negative or positive pressure devices, and invasive (conventional) mechanical ventilation by oro-/naso-tracheal tube or tracheostomy.

Noninvasive Mechanical Ventilation: Noninvasive intermittent positive pressure ventilation (NIPPV) has been studied in many uncontrolled and five randomized controlled trials in acute respiratory failure^{202,203}. The studies show consistently positive results with success rates of 80-85%²⁰⁴. Taken together they provide evidence that NIPPV increases pH, reduces PaCO_2 , reduces the

Table 14 - Indications and Relative Contraindications for NIPPV^{203,204}

Selection criteria

- Moderate to severe dyspnea with use of accessory muscles and paradoxical abdominal motion.
- Moderate to severe acidosis ($\text{pH} < 7.35$) and hypercapnia ($\text{PaCO}_2 > 6.0$ kPa, 45mm Hg)²⁰⁹.
- Respiratory frequency > 25 breaths per minute.

Exclusion criteria

- Respiratory arrest.
- Cardiovascular instability (hypotension, arrhythmias, myocardial infarction).
- Somnolence, impaired mental status, uncooperative patient.
- High aspiration risk; viscous or copious secretions.
- Recent facial or gastroesophageal surgery.
- Craniofacial trauma, fixed nasopharyngeal abnormalities.
- Extreme obesity.

severity of breathlessness in the first 4 hours of treatment, and decreases the length of hospital stay (**Evidence A**). More importantly, mortality – or its surrogate, intubation rate – is reduced by this intervention²⁰⁵⁻²⁰⁸. However, NIPPV is not appropriate for all patients, as summarized in **Table 14**.

Invasive (Conventional) Mechanical Ventilation: Patients who show impending acute respiratory failure and those with life-threatening acid-base status abnormalities and/or altered mental status despite aggressive pharmacologic therapy are likely to be the best candidates for invasive mechanical ventilation. The indications for initiating mechanical ventilation during exacerbations of COPD are shown in **Table 15**, the first being the commonest and most important reason. The

three ventilatory modes most widely used are assisted-control ventilation, and pressure support ventilation alone or in combination with intermittent mandatory ventilation²¹⁰.

The use of invasive ventilation in end-stage COPD patients is influenced by the likely reversibility of the precipitating event, the patient's wishes, and the availability of intensive care facilities. Major hazards include the risk of ventilator-acquired pneumonia (especially

Table 15 - Indications for Invasive Mechanical Ventilation
<ul style="list-style-type: none"> • Severe dyspnea with use of accessory muscles and paradoxical abdominal motion. • Respiratory frequency > 35 breaths per minute. • Life-threatening hypoxemia (PaO₂ < 5.3 kPa, 40 mm Hg or PaO₂/FiO₂* < 200 mm Hg). • Severe acidosis (pH < 7.25) and hypercapnia (PaCO₂ > 8.0 kPa, 60 mm Hg). • Respiratory arrest. • Somnolence, impaired mental status. • Cardiovascular complications (hypotension, shock, heart failure). • Other complications (metabolic abnormalities, sepsis, pneumonia, pulmonary embolism, barotrauma, massive pleural effusion). • NIPPV failure (or exclusion criteria, see Table 14).

when multiresistant organisms are prevalent), barotrauma, and failure to wean to spontaneous ventilation. Contrary to some opinions, mortality among COPD patients with respiratory failure is no greater than mortality among patients ventilated for non-COPD causes. When possible, a clear statement of the patient's own treatment wishes – an advance directive or “living will” – makes these difficult decisions much easier to resolve.

Weaning or discontinuation from mechanical ventilation can be particularly difficult and hazardous in patients with COPD, and the best method to wean patients from the ventilator remains a matter of debate^{211,212}. Whether pressure support or a T-piece trial is used, weaning is shortened when a clinical protocol is adopted (**Evidence A**). Noninvasive ventilation (NIV) has been applied to facilitate the weaning process in COPD patients with acute or chronic respiratory failure²¹³. Compared with invasive pressure support ventilation, noninvasive intermittent positive pressure ventilation (NIPPV) during weaning shortened weaning time, reduced the stay in the intensive care unit, decreased the incidence of nosocomial pneumonia, and improved 60-day survival rates²¹³.

Similar findings have been reported when NIPPV is used after extubation for hypercapnic respiratory failure²¹⁴ (**Evidence C**).

Other Measures: Further treatment measures that can be used in the hospital include: fluid administration (accurate monitoring of fluid balance is essential); nutrition (supplementary when the patient is too dyspneic to eat); low molecular weight heparin in immobilized, polycythemic, or dehydrated patients with or without a history of thromboembolic disease; sputum clearance (by stimulating coughing and low volume forced expirations as in home management). Manual or mechanical chest percussion and postural drainage may be beneficial in patients producing > 25 ml sputum per day or with lobar atelectasis.

HOSPITAL DISCHARGE AND FOLLOW-UP

Insufficient clinical data exist to establish the optimal duration of hospitalization for acute exacerbations of COPD^{171,215,216}. Consensus and limited data support the discharge criteria listed in **Table 16**. **Table 17** provides items to include in a follow-up assessment 4 to 6 weeks

Table 16 - Discharge Criteria for Patients with Exacerbations of COPD
<ul style="list-style-type: none"> • Inhaled β₂-agonist therapy is required no more frequently than every 4 hrs. • Patient, if previously ambulatory, is able to walk across room. • Patient is able to eat and sleep without frequent awakening by dyspnea. • Patient has been clinically stable for 12-24 hrs. • Arterial blood gases have been stable for 12-24 hrs. • Patient (or home caregiver) fully understands correct use of medications. • Follow-up and home care arrangements have been completed (e.g., visiting nurse, oxygen delivery, meal provisions). • Patient, family, and physician are confident patient can manage successfully.

Table 17 - Follow-up Assessment 4-6 Weeks After Discharge from Hospital for Exacerbations of COPD
<ul style="list-style-type: none"> • Ability to cope in usual environment. • Measurement of FEV₁. • Reassessment of inhaler technique. • Understanding of recommended treatment regimen. • Need for long-term oxygen therapy and/or home nebulizer (for patients with very severe COPD).

after discharge from the hospital. Thereafter, follow-up is the same as for stable COPD, including supervising smoking cessation, monitoring the effectiveness of each drug treatment, and monitoring changes in spirometric parameters⁸⁵. Home visits by a community nurse may permit earlier discharge of patients hospitalized with a non-acidotic exacerbation of COPD, without increasing readmission rate²¹⁷⁻²²⁰

If hypoxemia developed during the exacerbation, arterial blood gases should be rechecked at discharge and at the follow-up visit. If the patient remains hypoxemic, long-term oxygen therapy should be instituted. Decisions about continuous domiciliary oxygen based on the severity of the acute hypoxemia during an exacerbation are frequently misleading.

The opportunities for prevention of future exacerbations should be reviewed before discharge with particular attention to future influenza vaccination plans, knowledge of current therapy including inhaler technique^{221,222}, and how to recognize symptoms of exacerbations. Pharmacotherapy known to reduce the number of exacerbations should be considered. Social problems should be discussed and principal caregivers identified if the patient has a significant persisting disability.

4. FUTURE RESEARCH

A better understanding of molecular and cellular pathogenic mechanisms of COPD should lead to many new directions for both basic and clinical investigations. Improved methods for early detection, new approaches for interventions through targeted pharmacotherapy, possible means to identify the "susceptible" smoker, and more effective means of managing exacerbations are needed. Some research recommendations are provided; there are many additional avenues to explore.

■ Until there is a better understanding of the causal mechanisms of COPD, an absolutely rigid definition of COPD, and its relationship to other obstructive airways diseases, will remain controversial. Defining characteristics of COPD should be better identified.

■ The stages of COPD and the disease course will vary from one patient to another. The GOLD Report describes four stages and their clinical utility needs to be evaluated.

■ Surrogate markers of inflammation, possibly derived from sputum (cells, mediators, enzymes) or exhaled condensates (lipid mediators, reactive oxygen species,

cytokines), that may predict the clinical usefulness of new management and prevention strategies for COPD need to be developed.

■ Information is needed about the cellular and molecular mechanisms of inflammation in stable COPD and in exacerbations. Inflammatory responses in nonsmokers, ex-smokers, and smokers with and without COPD should be compared. The mechanisms responsible for the persistence of the inflammatory response in COPD should be investigated. Why inflammation in COPD is poorly responsive to glucocorticosteroids and what treatments other than glucocorticosteroids are effective in suppressing inflammation in COPD are research topics that could lead to new treatment modalities.

■ There is a pressing need to develop drugs that control symptoms and prevent the progression of COPD. Some progress has been made and there are several classes of drugs that are now in preclinical and clinical development for use in COPD patients.

■ Standardized methods for tracking trends in COPD prevalence, morbidity, and mortality over time need to be developed so that countries can plan for future increases in the need for health care services in view of predicted increases in COPD. This need is especially urgent in developing countries with limited health care resources.

■ Longitudinal studies demonstrating the course of COPD are needed in a variety of populations exposed to various risk factors. Such studies would provide insight into the pathogenesis of COPD, identify additional genetic bases for COPD, and identify how genetic risk factors interact with environmental risk factors in specific patient populations. Factors that determine why some, but not all, smokers develop COPD need to be identified.

■ Data are needed on the use, cost, and relative distribution of medical and non-medical resources for COPD, especially in countries where smoking and other risk factors are prevalent. These data are likely to have some impact on health policy and resource allocation decisions. As options for treating COPD grow, more research will be needed to help guide health care providers and health budget managers regarding the most efficient and effective ways of managing this disease. Methods and strategies for implementation of COPD management programs in developing countries will require special attention.

■ While spirometry is recommended to assess and monitor COPD, other measures need to be developed

and evaluated in clinical practice. Reproducible and inexpensive exercise-testing methodologies (e.g., stair-climbing tests) suitable for use in developing countries need to be evaluated and their use encouraged. Spirometers need to be developed that can ensure economical and accurate performance when a relatively untrained operator administers the test.

■ Since COPD is not fully reversible (with current therapies) and slowly progressive, it will become ever more important to identify early cases as more effective therapies emerge. Consensus on standard methods for detection and definition of early disease needs to be developed. Data to show whether or not screening spirometry is effective in directing management decisions in COPD outcomes are required.

■ Primary prevention of COPD is one of the major objectives of GOLD. Investigations into the most cost-effective ways to reduce the prevalence of tobacco smoking in the general population and more specifically in young people are very much needed. Strategies to prevent people from starting to smoke and methods for smoking cessation require constant evaluation and improvement. Research is required to gauge the impact and reduce the risk from growing air pollution, urbanization, recurrent childhood infections, occupational exposures, and use of local cigarette equivalents. Programs designed to reduce exposure to biomass fuel in countries where this is used for cooking and domestic heating should be explored in an effort to reduce exposure and improve ventilation in the homes.

■ The specific components of effective education for COPD patients need to be determined. It is not known, for example, whether COPD patients should be given an individual management plan, or whether these plans are effective in reducing health care costs or improving the outcomes of exacerbations. Developing and evaluating effective tools for physician education concerning prevention, diagnosis, and management of COPD will be important in view of the increasing public health problem presented by COPD.

■ Studies are needed to determine whether education is an essential component of pulmonary rehabilitation. The cost effectiveness of rehabilitation programs has not been assessed and there is a need to assess the feasibility, resource utilization, and health outcomes of rehabilitation programs that can be delivered outside the major teaching hospital setting. Criteria for selecting individuals for rehabilitation should be evaluated, along with methods to modify programs to suit the needs of individual patients.

■ Collecting and evaluating data to set levels of severity for COPD exacerbations would stimulate standardization of this outcome measure that is so frequently used in clinical trials. Better data on outcomes of COPD exacerbations would allow physicians to provide better advice to patients on possible outcomes and appropriateness of various types of treatment. Further exploration of the ethical principles of life support and greater insights into the behavioral influences that inhibit discussion of end-of-life issues are needed, along with studies to define the needs of end-stage COPD patients.

REFERENCES

1. National Heart, Lung, and Blood Institute. *Morbidity & Mortality: Chartbook on Cardiovascular, Lung, and Blood Diseases*. Bethesda, MD: US Department of Health and Human Services, Public Health Service, National Institutes of Health, 1998: Available from: URL: www.nhlbi.nih.gov/nhlbi/seiin/other/cht-book/htm.
2. Murray CJL, Lopez AD. Evidence-based health policy - lessons from the global burden of disease study. *Science* 1996; 274:740-743.
3. *World Health Report*. Geneva: World Health Organization, 2000: Available from URL: <http://www.who.int/whr/2000/en/statistics.htm>.
4. Hill AT, Bayley D, Stockley RA. The interrelationship of sputum inflammatory markers in patients with chronic bronchitis. *Am J Respir Crit Care Med* 1999; 160:893-8.
5. Keatings VM, Collins PD, Scott DM, Barnes PJ. Differences in interleukin-8 and tumor necrosis factor-alpha in induced sputum from patients with chronic obstructive pulmonary disease or asthma. *Am J Respir Crit Care Med* 1996; 153:530-534.
6. Beeh KM, Beier J, Kornmann O, Mander A, Buhl R. Long-term repeatability of induced sputum cells and inflammatory markers in stable, moderately severe COPD. *Chest* 2003; 123:778-83.
7. Yamamoto C, Yoneda T, Yoshikawa M, Fu A, Tokuyama T, Tsukaguchi K, et al. Airway inflammation in COPD assessed by sputum levels of interleukin-8. *Chest* 1997; 112:505-10.
8. Mueller R, Chanez P, Campbell AM, Bousquet J, Heusser C, Bullock GR. Different cytokine patterns in bronchial biopsies in asthma and chronic bronchitis. *Respir Med* 1996; 90:79-85.
9. Mullen JB, Wright JL, Wiggs BR, Pare PD, Hogg JC. Reassessment of inflammation of airways in chronic bronchitis. *Br Med J (Clin Res Ed)* 1985; 291:1235-9.
10. Cosio M, Ghezzi H, Hogg JC, Corbin R, Loveland M, Dosman J, et al. The relations between structural changes in small airways and pulmonary- function tests. *N Engl J Med* 1978; 298:1277-81.
11. Niewoehner DE, Kleinerman J, Rice DB. Pathologic changes in the peripheral airways of young cigarette smokers. *N Engl J Med* 1974; 291:755-8.

12. Wright JL, Lawson LM, Pare PD, Wiggs BJ, Kennedy S, Hogg JC. Morphology of peripheral airways in current smokers and ex-smokers. *Am Rev Respir Dis* 1983; 127:474-7.
13. Ollerenshaw SL, Woolcock AJ. Characteristics of the inflammation in biopsies from large airways of subjects with asthma and subjects with chronic airflow limitation. *Am Rev Respir Dis* 1992; 145:922-7.
14. Hunninghake GW, Crystal RG. Cigarette smoking and lung destruction. Accumulation of neutrophils in the lungs of cigarette smokers. *Am Rev Respir Dis* 1983; 128:833-8.
15. Li XY, Brown D, Smith S, MacNee W, Donaldson K. Short-term inflammatory responses following intratracheal instillation of fine and ultrafine carbon black in rats. *Inhal Toxicol* 1999; 11:709-31.
16. Monn C, Becker S. Cytotoxicity and induction of proinflammatory cytokines from human monocytes exposed to fine (PM_{2.5}) and coarse particles (PM_{10-2.5}) in outdoor and indoor air. *Toxicol Appl Pharmacol* 1999; 155:245-52.
17. Salvi S, Blomberg A, Rudell B, Kelly F, Sandstrom T, Holgate ST, et al. Acute inflammatory responses in the airways and peripheral blood after short-term exposure to diesel exhaust in healthy human volunteers. *Am J Respir Crit Care Med* 1999; 159:702-9.
18. Von Essen SG, O'Neill DP, McGranaghan S, Olenchock SA, Rennard SI. Neutrophilic respiratory tract inflammation and peripheral blood neutrophilia after grain sorghum dust extract challenge. *Chest* 1995; 108:1425-33.
19. Von Essen SG, Robbins RA, Thompson AB, Ertl RF, Linder J, Rennard SI. Mechanisms of neutrophil recruitment to the lung by grain dust exposure. *Am Rev Respir Dis* 1988; 138:921-7.
20. O'Shaughnessy TC, Ansari TW, Barnes NC, Jeffery PK. Inflammation in bronchial biopsies of subjects with chronic bronchitis: inverse relationship of CD8+ T lymphocytes with FEV₁. *Am J Respir Crit Care Med* 1997; 155:852-7.
21. Saetta M, Di Stefano A, Maestrelli P, Ferraresso A, Drigo R, Potena A, et al. Activated T-lymphocytes and macrophages in bronchial mucosa of subjects with chronic bronchitis. *Am Rev Respir Dis* 1993; 147:301-6.
22. Saetta M, Di Stefano A, Turato G, Facchini FM, Corbino L, Mapp CE, et al. CD8+ T-lymphocytes in peripheral airways of smokers with chronic obstructive pulmonary disease. *Am J Respir Crit Care Med* 1998; 157:822-6.
23. Leopold JG, Goeff J. Centrilobular form of hypertrophic emphysema and its relation to chronic bronchitis. *Thorax* 1957; 12:219-235.
24. McLean KA. Pathogenesis of pulmonary emphysema. *Am J Med* 1958; 25:62-74.
25. Repine JE, Bast A, Lankhorst I. Oxidative stress in chronic obstructive pulmonary disease. Oxidative Stress Study Group. *Am J Respir Crit Care Med* 1997; 156:341-57.
26. Wright JL, Lawson L, Pare PD, Hooper RO, Peretz DI, Nelems JM, et al. The structure and function of the pulmonary vasculature in mild chronic obstructive pulmonary disease. The effect of oxygen and exercise. *Am Rev Respir Dis* 1983; 128:702-7.
27. Peinado VI, Barbera JA, Abate P, Ramirez J, Roca J, Santos S, et al. Inflammatory reaction in pulmonary muscular arteries of patients with mild chronic obstructive pulmonary disease. *Am J Respir Crit Care Med* 1999; 159:1605-11.
28. Riley DJ, Thakker-Varia S, Poiani GJ, Tozzi CA. Vascular remodeling. In: Crystal RG, West JB, Barnes PJ, Weibel ER, eds. *The lung: scientific foundations*. Philadelphia: Lippincott-Raven, 1977:1589-1597.
29. MacNee W. Pathophysiology of cor pulmonale in chronic obstructive pulmonary disease. Part two. *Am J Respir Crit Care Med* 1994; 150:1158-68.
30. Pride NB, Vermeire P, Allegra L. Diagnostic labels applied to model case histories of chronic airflow obstruction. Responses to a questionnaire in 11 North American and Western European Countries. *Eur Respir J* 1989; 2:702-9.
31. Mannino DM, Brown C, Giovino GA. Obstructive lung disease deaths in the United States from 1979 through 1993. An analysis using multiple-cause mortality data. *Am J Respir Crit Care Med* 1997; 156:814-818.
32. Murray CJL, Lopez AD, editors. *The global burden of disease: a comprehensive assessment of mortality and disability from diseases, injuries and risk factors in 1990 and projected to 2020*. Cambridge, MA: Harvard University Press, 1996.
33. Office of National Statistics. Mortality Statistics (revised) and 1994. *England and Wales*. London HMSO, 1996.
34. Rutten-van Molken MP, Postma MJ, Joore MA, Van Genugten ML, Leidl R, Jager JC. Current and future medical costs of asthma and chronic obstructive pulmonary disease in The Netherlands. *Respir Med* 1999; 93:779-787.
35. Jacobson L, Hertzman P, Lofdahl C-G, Skoogh B-E, Lindgren B. The economic impact of asthma and COPD in Sweden 1980 and 1991. *Respir Med* 2000; 94:247-255.
36. Buist AS, Vollmer WM. Smoking and other risk factors. In: Murray JF, Nadel JA, eds. *Textbook of respiratory medicine*. Philadelphia: WB Saunders Co, 1994:1259-1287.
37. Thom TJ. International comparisons in COPD mortality. *Am Rev Respir Dis* 1989; 140:S27-34.
38. Xu X, Weiss ST, Rijcken B, Schouten JP. Smoking, changes in smoking habits, and rate of decline in FEV₁: new insight into gender differences. *Eur Respir J* 1994; 7:1056-61.
39. Feinleib M, Rosenberg HM, Collins JG, Delozier JE, Pokras R, Chevarley FM. Trends in COPD morbidity and mortality in the United States. *Am Rev Respir Dis* 1989; 140:S9-18.
40. US Centers for Disease Control and Prevention: *Vital and Health Statistics: Current Estimates from the National Health Interview Survey*: Department of Health and Human Service, Public Health Service, 1995:Publication No 96-1527.
41. Anthonisen NR, Connett JE, Kiley JP, Altose MD, Bailey

- WC, Buist AS, et al. Effects of smoking intervention and the use of an inhaled anticholinergic bronchodilator on the rate of decline of FEV₁. The Lung Health Study. *JAMA* 1994; 272:1497-505.
42. Laurell CB, Eriksson S. The electrophoretic alpha-1 globulin pattern of serum in alpha-1 antitrypsin deficiency. *Scand J Clin Lab Invest* 1963; 15:132-140.
 43. Hubbard RC, Crystal RG. Antiproteases. In: Crystal RB, West JB, Barnes PJ, Cherniack NS, Weibel ER, eds. *The lung: scientific foundations*. New York: Raven Press, Ltd., 1991:1775-1787.
 44. McElvaney NG, Crystal RG. Inherited susceptibility of the lung to proteolytic injury. In: Crystal RG, West JB, Weibel ER, Barnes PJ, eds. *The lung: scientific foundations*. Philadelphia: Lippincott-Raven, 1997:2537-2553.
 45. Orié NGM, Sluiter HJ, De Vreis K, Tammerling K, Wikop J. The host factor in bronchitis. In: Orié NGM, Sluiter HJ, eds. *Bronchitis, an international symposium*. Assen, Netherlands: Royal Vangorcum, 1961:43-59.
 46. Hagstrom B, Nyberg P, Nilsson PM. Asthma in adult life--is there an association with birth weight? *Scand J Prim Health Care* 1998; 16:117-20.
 47. Svanes C, Omenaas E, Heuch JM, Irgens LM, Gulsvik A. Birth characteristics and asthma symptoms in young adults: results from a population-based cohort study in Norway. *Eur Respir J* 1998; 12:1366-70.
 48. Todisco T, de Benedictis FM, Iannacci L, Baglioni S, Esлами A, Todisco E, et al. Mild prematurity and respiratory functions. *Eur J Pediatr* 1993; 152:55-8.
 49. Stein CE, Kumaran K, Fall CH, Shaheen SO, Osmond C, Barker DJ. Relation of fetal growth to adult lung function in South India. *Thorax* 1997; 52:895-9.
 50. Morgan WJ. Maternal smoking and infant lung function. Further evidence for an in utero effect. *Am J Respir Crit Care Med* 1998; 158:689-90.
 51. Tager IB, Segal MR, Speizer FE, Weiss ST. The natural history of forced expiratory volumes. Effect of cigarette smoking and respiratory symptoms. *Am Rev Respir Dis* 1988; 138:837-49.
 52. US Surgeon General. *The health consequences of smoking: chronic obstructive pulmonary disease*. Washington, D.C.: US Department of Health and Human Services, 1984:Publication No. 84-50205.
 53. Leuenberger P, Schwartz J, Ackermann-Liebrich U, Blaser K, Bolognini G, Bongard JP, et al. Passive smoking exposure in adults and chronic respiratory symptoms (SAPALDIA Study). Swiss Study on Air Pollution and Lung Diseases in Adults, SAPALDIA Team. *Am J Respir Crit Care Med* 1994; 150:1222-1228.
 54. Dayal HH, Khuder S, Sharrar R, Trieff N. Passive smoking in obstructive respiratory disease in an industrialized urban population. *Environ Res* 1994; 65:161-171.
 55. Holt PG. Immune and inflammatory function in cigarette smokers. *Thorax* 1987; 42:241-249.
 56. Kauffmann F, Drouet D, Lellouch J, Brille D. Twelve years spirometric changes among Paris area workers. *Int J Epidemiol* 1979; 8:201-212.
 57. Niewoehner DE. Anatomic and pathophysiological correlations in COPD. In: Baum GL, Crapo JD, Celli BR, Karlinsky JB, eds. *Textbook of pulmonary diseases*. Philadelphia: Lippincott-Raven, 1998:823-842.
 58. Chen JC, Mannino MD. Worldwide epidemiology of chronic obstructive pulmonary disease. *Current Opinion in Pulmonary Medicine* 1999; 5:93-99.
 59. Perez-Padilla R, Regalado U, Vedal S, Pare P, Chapela R, Sansores R, et al. Exposure to biomass smoke and chronic airway disease in Mexican women. *Am J Respir Crit Care Med* 1996; 154:701-706.
 60. Dossing M, Khan J, al-Rabiah F. Risk factors for chronic obstructive lung disease in Saudi Arabia. *Respiratory Med* 1994; 88:519-522.
 61. Behera D, Jindal SK. Respiratory symptoms in Indian women using domestic cooking fuels. *Chest* 1991; 100:385-388.
 62. Amoli K. Bronchopulmonary disease in Iranian housewives chronically exposed to indoor smoke. *Eur Respir J* 1998; 11:659-663.
 63. Dennis R, Maldonado D, Norman S, Baena E, Martinez G. Woodsmoke exposure and risk for obstructive airways disease among women. *Chest* 1996; 109:115-119.
 64. Pandey MR. Prevalence of chronic bronchitis in a rural community of the Hill Region of Nepal. *Thorax* 1984; 39:331-336.
 65. Pandey MR. Domestic smoke pollution and chronic bronchitis in a rural community of the Hill Region of Nepal. *Thorax* 1984; 39:337-339.
 66. Samet JM, Marbury M, Spengler J. Health effects and sources of indoor air pollution. *Am Rev Respir Dis* 1987; 136:1486-1508.
 67. Tao X, Hong CJ, Yu S, Chen B, Zhu H, Yang M. Priority among air pollution factors for preventing chronic obstructive pulmonary disease in Shanghai. *Sci Total Environ* 1992; 127:57-67.
 68. Prescott E, Lange P, Vestbo J. Socioeconomic status, lung function and admission to hospital for COPD: results from the Copenhagen City Heart Study. *Eur Respir J* 1999; 13:1109-14.
 69. Strachan DP. Epidemiology: A British perspective. In: Calverley PMA, Pride NB, eds. *Chronic obstructive pulmonary disease*. London: Chapman and Hall, 1995:47-67.
 70. Georgopoulos D, Anthonisen NR. Symptoms and signs of COPD. In: Cherniack NS, ed. *Chronic obstructive pulmonary disease*. Toronto: WB Saunders Co, 1991:357-363.
 71. Loveridge B, West P, Kryger MH, Anthonisen NR. Alteration in breathing pattern with progression of chronic obstructive pulmonary disease. *Am Rev Respir Dis* 1986; 134:930-4.
 72. Kesten S, Chapman KR. Physician perceptions and management of COPD. *Chest* 1993; 104:254-8.
 73. Reis AL. Response to bronchodilators. In: Clausen J, ed.

- Pulmonary function testing: guidelines and controversies.* New York: Academic Press, 1982.
74. American Thoracic Society. Lung function testing: selection of reference values and interpretative strategies. *Am Rev Respir Dis* 1991; 144:1202-18.
 75. Wilson DH, Wakefield MA, Steven ID, Rohrsheim RA, Esterman AJ, Graham NM. "Sick of Smoking": evaluation of a targeted minimal smoking cessation intervention in general practice. *Med J Aust* 1990; 152:518-21.
 76. Britton J, Knox A. Helping people to stop smoking: the new smoking cessation guidelines. *Thorax* 1999; 54:1-2.
 77. Fiore MC, Bailey WC, Cohen SJ. Smoking cessation: information for specialists. Rockville, MD: US Department of Health and Human Services, Public Health Service, Agency for Health Care Policy and Research and Centers for Disease Control and Prevention, 1996:AHCPH Publication No. 96-0694.
 78. The tobacco use and dependence clinical practice guideline panel, staff, and consortium representatives. A clinical practice guideline for treating tobacco use and dependence. *JAMA* 2000; 28:3244-3254.
 79. American Medical Association. Guidelines for the diagnosis and treatment of nicotine dependence: how to help patients stop smoking. Washington DC: American Medical Association, 1994.
 80. Glynn TJ, Manley MW. How to help your patients stop smoking. A National Cancer Institute manual for physicians. Bethesda, MD: US Department of Health and Human Services, Public Health Service, National Institutes of Health, National Cancer Institute, 1990:NIH Publication No. 90-3064.
 81. Glynn TJ, Manley MW, Pechacek TF. Physician-initiated smoking cessation program: the National Cancer Institute trials. *Prog Clin Biol Res* 1990; 339:11-25.
 82. Fiore MC, Smith SS, Jorenby DE, Baker TB. The effectiveness of the nicotine patch for smoking cessation. A meta-analysis. *JAMA* 1994; 271:1940-7.
 83. Lancaster T, Stead L, Silagy C, Sowden A. Effectiveness of interventions to help people stop smoking: findings from the Cochrane Library. *BMJ* 2000; 321:355-8.
 84. Tashkin D, Kanner R, Bailey W, Buist S, Anderson P, Nides M, et al. Smoking cessation in patients with chronic obstructive pulmonary disease: a double-blind, placebo-controlled, randomised trial. *Lancet* 2001; 357:1571-5.
 85. The COPD Guidelines Group of the Standards of Care Committee of the BTS. BTS guidelines for the management of chronic obstructive pulmonary disease. *Thorax* 1997; 52 Suppl 5:S1-28.
 86. Samet J, Utell MJ. Ambient air pollution. In: Rosenstock L, Cullen M, eds. *Textbook of occupational and environmental medicine*. Philadelphia: WB Saunders Co, 1994:53-60.
 87. Ries AL, Kaplan RM, Limberg TM, Prewitt LM. Effects of pulmonary rehabilitation on physiologic and psychosocial outcomes in patients with chronic obstructive pulmonary disease. *Ann Intern Med* 1995; 122:823-32.
 88. Janelli LM, Scherer YK, Schmieder LE. Can a pulmonary health teaching program alter patients' ability to cope with COPD? *Rehabil Nurs* 1991; 16:199-202.
 89. Ashikaga T, Vacek PM, Lewis SO. Evaluation of a community-based education program for individuals with chronic obstructive pulmonary disease. *J Rehabil* 1980; 46:23-7.
 90. Toshima MT, Kaplan RM, Ries AL. Experimental evaluation of rehabilitation in chronic obstructive pulmonary disease: short-term effects on exercise endurance and health status. *Health Psychol* 1990; 9:237-52.
 91. Celli BR. Pulmonary rehabilitation in patients with COPD. *Am J Respir Crit Care Med* 1995; 152:861-4.
 92. Heffner JE, Fahy B, Hilling L, Barbieri C. Outcomes of advance directive education of pulmonary rehabilitation patients. *Am J Respir Crit Care Med* 1997; 155:1055-9.
 93. Stewart MA. Effective physician-patient communication and health outcomes: a review. *CMAJ* 1995; 152:1423-33.
 94. Clark NM, Nothwehr F, Gong M, Evans D, Maiman LA, Hurwitz ME, et al. Physician-patient partnership in managing chronic illness. *Acad Med* 1995; 70:957-9.
 95. Pauwels RA, Lofdahl CG, Laitinen LA, Schouten JP, Postma DS, Pride NB, et al. Long-term treatment with inhaled budesonide in persons with mild chronic obstructive pulmonary disease who continue smoking. European Respiratory Society Study on Chronic Obstructive Pulmonary Disease. *N Engl J Med* 1999; 340:1948-53.
 96. Vestbo J, Sorensen T, Lange P, Brix A, Torre P, Viskum K. Long-term effect of inhaled budesonide in mild and moderate chronic obstructive pulmonary disease: a randomised controlled trial. *Lancet* 1999; 353:1819-23.
 97. Borge PS, Calverley PM, Jones PW, Spencer S, Anderson JA, Maslen TK. Randomised, double blind, placebo controlled study of fluticasone propionate in patients with moderate to severe chronic obstructive pulmonary disease: the ISOLDE trial. *BMJ* 2000; 320:1297-303.
 98. The Lung Health Study Research Group. Effect of inhaled triamcinolone on the decline in pulmonary function in chronic obstructive pulmonary disease: Lung Health Study II. *N Engl J Med* 2000; 343:1902-1909.
 99. Vathenen AS, Britton JR, Ebdon P, Cookson JB, Wharrad HJ, Tattersfield AE. High-dose inhaled albuterol in severe chronic airflow limitation. *Am Rev Respir Dis* 1988; 138:850-5.
 100. Gross NJ, Petty TL, Friedman M, Skorodin MS, Silvers GW, Donohue JF. Dose response to ipratropium as a nebulized solution in patients with chronic obstructive pulmonary disease. A three-center study. *Am Rev Respir Dis* 1989; 139:1188-91.
 101. Chrystyn H, Mulley BA, Peake MD. Dose response relation to oral theophylline in severe chronic obstructive airways disease. *BMJ* 1988; 297:1506-10.
 102. Higgins BG, Powell RM, Cooper S, Tattersfield AE. Effect of salbutamol and ipratropium bromide on airway calibre and bronchial reactivity in asthma and chronic bronchitis.

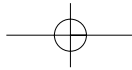
- Eur Respir J* 1991; 4:415-20.
103. Ikeda A, Nishimura K, Koyama H, Tsukino M, Mishima M, Izumi T. Dose response study of ipratropium bromide aerosol on maximum exercise performance in stable patients with chronic obstructive pulmonary disease. *Thorax* 1996; 51:48-53.
 104. Guyatt GH, Townsend M, Pugsley SO, Keller JL, Short HD, Taylor DW, et al. Bronchodilators in chronic air-flow limitation. Effects on airway function, exercise capacity, and quality of life. *Am Rev Respir Dis* 1987; 135:1069-74.
 105. Mahler DA, Donohue JF, Barbee RA, Goldman MD, Gross NJ, Wisniewski ME, et al. Efficacy of salmeterol xinafoate in the treatment of COPD. *Chest* 1999; 115:957-65.
 106. Dahl R, Greefhorst LA, Nowak D, Nonikov V, Byrne AM, Thomson MH, et al. Inhaled formoterol dry powder versus ipratropium bromide in chronic obstructive pulmonary disease. *Am J Respir Crit Care Med* 2001; 164:778-84.
 107. Vincken W, van Noord JA, Greefhorst AP, Bantje TA, Kesten S, Korducki L, et al. Improved health outcomes in patients with COPD during 1 yr's treatment with tiotropium. *Eur Respir J* 2002; 19:209-16.
 108. COMBIVENT Inhalation Aerosol Study Group. In chronic obstructive pulmonary disease, a combination of ipratropium and albuterol is more effective than either agent alone. An 85-day multicenter trial. *Chest* 1994; 105:1411-9.
 109. The COMBIVENT Inhalation Solution Study Group. Routine nebulized ipratropium and albuterol together are better than either alone in COPD. *Chest* 1997; 112:1514-21.
 110. Gross N, Tashkin D, Miller R, Oren J, Coleman W, Linberg S. Inhalation by nebulization of albuterol-ipratropium combination (Dey combination) is superior to either agent alone in the treatment of chronic obstructive pulmonary disease. Dey Combination Solution Study Group. *Respiration* 1998; 65:354-62.
 111. Ulrik CS. Efficacy of inhaled salmeterol in the management of smokers with chronic obstructive pulmonary disease: a single centre randomised, double blind, placebo controlled, crossover study. *Thorax* 1995; 50:750-4.
 112. Taylor DR, Buick B, Kinney C, Lowry RC, McDevitt DG. The efficacy of orally administered theophylline, inhaled salbutamol, and a combination of the two as chronic therapy in the management of chronic bronchitis with reversible air-flow obstruction. *Am Rev Respir Dis* 1985; 131:747-51.
 113. van Noord JA, de Munck DR, Bantje TA, Hop WC, Akveld ML, Bommer AM. Long-term treatment of chronic obstructive pulmonary disease with salmeterol and the additive effect of ipratropium. *Eur Respir J* 2000; 15:878-85.
 114. ZuWallack RL, Mahler DA, Reilly D, Church N, Emmett A, Rickard K, et al. Salmeterol plus theophylline combination therapy in the treatment of COPD. *Chest* 2001; 119:1661-70.
 115. Bellia V, Foresi A, Bianco S, Grassi V, Olivieri D, Bensi G, et al. Efficacy and safety of oxitropium bromide, theophylline and their combination in COPD patients: a double-blind, randomized, multicentre study (BREATH Trial). *Respir Med* 2002; 96:881-9.
 116. Murciano D, Auclair MH, Pariente R, Aubier M. A randomized, controlled trial of theophylline in patients with severe chronic obstructive pulmonary disease. *N Engl J Med* 1989; 320:1521-5.
 117. O'Driscoll BR, Kay EA, Taylor RJ, Weatherby H, Chetty MC, Bernstein A. A long-term prospective assessment of home nebulizer treatment. *Respir Med* 1992; 86:317-25.
 118. Tashkin DP, Bleecker E, Braun S, Campbell S, DeGraff AC, Jr., Hudgel DW, et al. Results of a multicenter study of nebulized inhalant bronchodilator solutions. *Am J Med* 1996; 100:62S-69.
 119. Calverley P, Pauwels R, Vestbo J, Jones P, Pride N, Gulsvik A, et al. Combined salmeterol and fluticasone in the treatment of chronic obstructive pulmonary disease: a randomised controlled trial. *Lancet* 2003; 361:449-56.
 120. Jones PW, Willits LR, Burge PS, Calverley PM. Disease severity and the effect of fluticasone propionate on chronic obstructive pulmonary disease exacerbations. *Eur Respir J* 2003; 21:68-73.
 121. Mahler DA, Wire P, Horstman D, Chang CN, Yates J, Fischer T, et al. Effectiveness of fluticasone propionate and salmeterol combination delivered via the Diskus device in the treatment of chronic obstructive pulmonary disease. *Am J Respir Crit Care Med* 2002; 166(8):1084-91.
 122. Szafranski W, Cukier A, Ramirez A, Menga G, Sansores R, Nahabedian S, et al. Efficacy and safety of budesonide/formoterol in the management of chronic obstructive pulmonary disease. *Eur Respir J* 2003; 21:74-81.
 123. Senderovitz T, Vestbo J, Frandsen J, Maltbaek N, Norgaard M, Nielsen C, et al. Steroid reversibility test followed by inhaled budesonide or placebo in outpatients with stable chronic obstructive pulmonary disease. The Danish Society of Respiratory Medicine. *Respir Med* 1999; 93:715-8.
 124. Rice KL, Rubins JB, Lebahn F, Parenti CM, Duane PG, Kuskowski M, et al. Withdrawal of chronic systemic corticosteroids in patients with COPD: a randomized trial. *Am J Respir Crit Care Med* 2000; 162:174-8.
 125. Decramer M, Lacquet LM, Fagard R, Rogiers P. Corticosteroids contribute to muscle weakness in chronic airflow obstruction. *Am J Respir Crit Care Med* 1994; 150:11-6.
 126. Decramer M, Stas KJ. Corticosteroid-induced myopathy involving respiratory muscles in patients with chronic obstructive pulmonary disease or asthma. *Am Rev Respir Dis* 1992; 146:800-2.
 127. Nichol KL, Margolis KL, Wuorenma J, Von Sternberg T. The efficacy and cost effectiveness of vaccination against influenza among elderly persons living in the community. *N Engl J Med* 1994; 331:778-84.
 128. Edwards KM, Dupont WD, Westrich MK, Plummer WD, Jr., Palmer PS, Wright PF. A randomized controlled trial of cold-adapted and inactivated vaccines for the prevention of influenza A disease. *J Infect Dis* 1994; 169:68-76.

129. Simberkoff MS, Cross AP, Al-Ibrahim M, Baltch AL, Geiseler PJ, Nadler J, et al. Efficacy of pneumococcal vaccine in high-risk patients. Results of a Veterans Administration Cooperative Study. *N Engl J Med* 1986; 315:1318-27.
130. Williams JH, Jr., Moser KM. Pneumococcal vaccine and patients with chronic lung disease. *Ann Intern Med* 1986; 104:106-9.
131. Davis AL, Aranda CP, Schiffman G, Christianson LC. Pneumococcal infection and immunologic response to pneumococcal vaccine in chronic obstructive pulmonary disease. A pilot study. *Chest* 1987; 92:204-12.
132. Isada CM, Stoller JK. Chronic bronchitis: the role of antibiotics. In: Niederman MS, Sarosi GA, Glassroth J, eds. *Respiratory infections: a scientific basis for management*. London: WB Saunders, 1994:621-33.
133. American Thoracic Society. Standards for the diagnosis and care of patients with chronic obstructive pulmonary disease. *Am J Respir Crit Care Med* 1995; 152:S77-121.
134. Siafakas NM, Vermeire P, Pride NB, Paoletti P, Gibson J, Howard P, et al. Optimal assessment and management of chronic obstructive pulmonary disease (COPD). The European Respiratory Society Task Force. *Eur Respir J* 1995; 8:1398-420.
135. Poole PJ, Black PN. Mucolytic agents for chronic bronchitis or chronic obstructive pulmonary disease. *Cochrane Database Syst Rev* 2000; 2:Available from URL www.update-software.com or www.updatasa.com.
136. Hansen NC, Skriver A, Brorsen-Riis L, Balslov S, Evald T, Maltbaek N, et al. Orally administered N-acetylcysteine may improve general well-being in patients with mild chronic bronchitis. *Respir Med* 1994; 88:531-5.
137. British Thoracic Society Research Committee. Oral N-acetylcysteine and exacerbation rates in patients with chronic bronchitis and severe airways obstruction. *Thorax* 1985; 40:832-5.
138. Boman G, Backer U, Larsson S, Melander B, Wahlander L. Oral acetylcysteine reduces exacerbation rate in chronic bronchitis: report of a trial organized by the Swedish Society for Pulmonary Diseases. *Eur J Respir Dis* 1983; 64:405-15.
139. Rasmussen JB, Glennow C. Reduction in days of illness after long-term treatment with N-acetylcysteine controlled-release tablets in patients with chronic bronchitis. *Eur Respir J* 1988; 1:351-5.
140. Collet JP, Shapiro P, Ernst P, Renzi T, Ducruet T, Robinson A. Effects of an immunostimulating agent on acute exacerbations and hospitalizations in patients with chronic obstructive pulmonary disease. The PARI-IS Study Steering Committee and Research Group. Prevention of Acute Respiratory Infection by an Immunostimulant. *Am J Respir Crit Care Med* 1997; 156:1719-24.
141. Anthonisen NR. OM-8BV for COPD. *Am J Respir Crit Care Med* 1997; 156:1713-4.
142. Irwin RS, Boulet LP, Cloutier MM, Fuller R, Gold PM, Hoffstein V, et al. Managing cough as a defense mechanism and as a symptom. A consensus panel report of the American College of Chest Physicians. *Chest* 1998; 114:133S-181S.
143. Barbera JA, Roger N, Roca J, Rovira I, Higenbottam TW, Rodriguez-Roisin R. Worsening of pulmonary gas exchange with nitric oxide inhalation in chronic obstructive pulmonary disease. *Lancet* 1996; 347:436-40.
144. Jones AT, Evans TW. NO: COPD and beyond. *Thorax* 1997; 52 Suppl 3:S16-21.
145. Bardsley PA, Howard P, DeBacker W, Vermeire P, Mairesse M, Ledent C, et al. Two years treatment with almitrine bismesylate in patients with hypoxic chronic obstructive airways disease. *Eur Respir J* 1991; 4:308-10.
146. Watanabe S, Kanner RE, Cutillo AG, Menlove RL, Bachand RT, Jr., Szalkowski MB, et al. Long-term effect of almitrine bismesylate in patients with hypoxemic chronic obstructive pulmonary disease. *Am Rev Respir Dis* 1989; 140:1269-73.
147. Winkelmann BR, Kullmer TH, Kneissl DG, Trenk D, Kronenberger H. Low-dose almitrine bismesylate in the treatment of hypoxemia due to chronic obstructive pulmonary disease. *Chest* 1994; 105:1383-91.
148. Jennings AL, Davies AN, Higgins JP, Gibbs JS, Broadley KE. A systematic review of the use of opioids in the management of dyspnoea. *Thorax* 2002; 57:939-44.
149. Eiser N, Denman WT, West C, Luce P. Oral diamorphine: lack of effect on dyspnoea and exercise tolerance in the "pink puffer" syndrome. *Eur Respir J* 1991; 4:926-31.
150. Young IH, Daviskas E, Keena VA. Effect of low dose nebulised morphine on exercise endurance in patients with chronic lung disease. *Thorax* 1989; 44:387-90.
151. Rice KL, Kronenberg RS, Hedemark LL, Niewoehner DE. Effects of chronic administration of codeine and promethazine on breathlessness and exercise tolerance in patients with chronic airflow obstruction. *Br J Dis Chest* 1987; 81:287-92.
152. Woodcock AA, Gross ER, Gellert A, Shah S, Johnson M, Geddes DM. Effects of dihydrocodeine, alcohol, and caffeine on breathlessness and exercise tolerance in patients with chronic obstructive lung disease and normal blood gases. *N Engl J Med* 1981; 305:1611-6.
153. Poole PJ, Veale AG, Black PN. The effect of sustained-release morphine on breathlessness and quality of life in severe chronic obstructive pulmonary disease. *Am J Respir Crit Care Med* 1998; 157:1877-80.
154. Berry MJ, Rejeski WJ, Adair NE, Zaccaro D. Exercise rehabilitation and chronic obstructive pulmonary disease stage. *Am J Respir Crit Care Med* 1999; 160:1248-53.
155. Behnke M, Taube C, Kirsten D, Lehnigk B, Jorres RA, Magnussen H. Home-based exercise is capable of preserving hospital-based improvements in severe chronic obstructive pulmonary disease. *Respir Med* 2000; 94:1184-91.
156. Finnerty JP, Keeping I, Bullough I, Jones J. The effective-

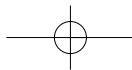
- ness of outpatient pulmonary rehabilitation in chronic lung disease: a randomized controlled trial. *Chest* 2001; 119:1705-10.
157. Green RH, Singh SJ, Williams J, Morgan MD. A randomized controlled trial of four weeks versus seven weeks of pulmonary rehabilitation in chronic obstructive pulmonary disease. *Thorax* 2001; 56:143-5.
 158. Ries AL, Kaplan RM, Myers R, Prewitt LM. Maintenance after pulmonary rehabilitation in chronic lung disease: a randomized trial. *Am J Respir Crit Care Med* 2003; 167:880-8.
 159. Goldstein RS, Gort EH, Stubbing D, Avendano MA, Guyatt GH. Randomised controlled trial of respiratory rehabilitation. *Lancet* 1994; 344:1394-7.
 160. Wijkstra PJ, Van Altena R, Kraan J, Otten V, Postma DS, Koeter GH. Quality of life in patients with chronic obstructive pulmonary disease improves after rehabilitation at home. *Eur Respir J* 1994; 7:269-73.
 161. McGavin CR, Gupta SP, Lloyd EL, McHardy GJ. Physical rehabilitation for the chronic bronchitic: results of a controlled trial of exercises in the home. *Thorax* 1977; 32:307-11.
 162. Nocturnal Oxygen Therapy Trial Group. Continuous or nocturnal oxygen therapy in hypoxemic chronic obstructive lung disease: a clinical trial. *Ann Intern Med* 1980; 93:391-8.
 163. Report of the Medical Research Council Working Party. Long term domiciliary oxygen therapy in chronic hypoxic cor pulmonale complicating chronic bronchitis and emphysema. *Lancet* 1981; 1:681-6.
 164. Tarry SP, Celli BR. Long-term oxygen therapy. *N Engl J Med* 1995; 333:710-4.
 165. Mehran RJ, Deslauriers J. Indications for surgery and patient work-up for bullectomy. *Chest Surg Clin N Am* 1995; 5:717-34.
 166. Benditt JO, Albert RK. Surgical options for patients with advanced emphysema. *Clin Chest Med* 1997; 18:577-93.
 167. Geddes D, Davies M, Koyama H, Hansell D, Pastorino U, Pepper J, et al. Effect of lung-volume-reduction surgery in patients with severe emphysema. *N Engl J Med* 2000; 343:239-45.
 168. National Emphysema Treatment Trial Research Group. Patients at high risk of death after lung-volume-reduction surgery. *N Engl J Med* 2001; 345:1075-83.
 169. The National Emphysema Treatment Trial Research Group. Rationale and design of The National Emphysema Treatment Trial: a prospective randomized trial of lung volume reduction surgery. *Chest* 1999; 116:1750-61.
 170. Maurer JR, Frost AE, Estenne M, Higenbottam T, Glanville AR. International guidelines for the selection of lung transplant candidates. The International Society for Heart and Lung Transplantation, the American Thoracic Society, the American Society of Transplant Physicians, the European Respiratory Society. *Transplantation* 1998; 66:951-6.
 171. Regueiro CR, Hamel MB, Davis RB, Desbiens N, Connors AF, Jr., Phillips RS. A comparison of generalist and pulmonologist care for patients hospitalized with severe chronic obstructive pulmonary disease: resource intensity, hospital costs, and survival. SUPPORT Investigators. Study to Understand Prognoses and Preferences for Outcomes and Risks of Treatment. *Am J Med* 1998; 105:366-72.
 172. Gibson PG, Wlodarczyk JH, Wilson AJ, Sprogis A. Severe exacerbation of chronic obstructive airways disease: health resource use in general practice and hospital. *J Qual Clin Pract* 1998; 18:125-33.
 173. Warren PM, Flenley DC, Millar JS, Avery A. Respiratory failure revisited: acute exacerbations of chronic bronchitis between 1961-68 and 1970-76. *Lancet* 1980; 1:467-70.
 174. Anthonisen NR, Manfreda J, Warren CP, Hershfield ES, Harding GK, Nelson NA. Antibiotic therapy in exacerbations of chronic obstructive pulmonary disease. *Ann Intern Med* 1987; 106:196-204.
 175. Wilson R. The role of infection in COPD. *Chest* 1998; 113:242S-248S.
 176. Soler N, Torres A, Ewig S, Gonzalez J, Celis R, El-Ebiary M, et al. Bronchial microbial patterns in severe exacerbations of chronic obstructive pulmonary disease (COPD) requiring mechanical ventilation. *Am J Respir Crit Care Med* 1998; 157:1498-505.
 177. Smith CB, Kanner RE, Golden CA, Klauber MR, Renzetti AD, Jr. Effect of viral infections on pulmonary function in patients with chronic obstructive pulmonary diseases. *J Infect Dis* 1980; 141:271-80.
 178. MacFarlane JT, Colville A, Guion A, Macfarlane RM, Rose DH. Prospective study of aetiology and outcome of adult lower-respiratory-tract infections in the community. *Lancet* 1993; 341:511-4.
 179. Stockley RA, O'Brien C, Pye A, Hill SL. Relationship of sputum color to nature and outpatient management of acute exacerbations of COPD. *Chest* 2000; 117:1638-45.
 180. Anderson HR, Spix C, Medina S, Schouten JP, Castellsague J, Rossi G, et al. Air pollution and daily admissions for chronic obstructive pulmonary disease in 6 European cities: results from the APHEA project. *Eur Respir J* 1997; 10:1064-71.
 181. Chodosh S, McCarty J, Farkas S, Dreobl M, Tosiello R, Shan M, et al. Randomized, double-blind study of ciprofloxacin and cefuroxime axetil for treatment of acute bacterial exacerbations of chronic bronchitis. The Bronchitis Study Group. *Clin Infect Dis* 1998; 27:722-9.
 182. Walsh EE, Falsey AR, Hennessey PA. Respiratory syncytial and other virus infections in persons with chronic cardiopulmonary disease. *Am J Respir Crit Care Med* 1999; 160:791-5.
 183. Mogulkoc N, Karakurt S, Isalska B, Bayindir U, Celikel T, Korten V, et al. Acute purulent exacerbation of chronic obstructive pulmonary disease and Chlamydia pneumoniae infection. *Am J Respir Crit Care Med* 1999; 160:349-53.
 184. Murphy TF, Sethi S, Klingman KL, Brueggemann AB, Doern GV. Simultaneous respiratory tract colonization by multiple strains of nontypeable Haemophilus influenzae in chronic obstructive pulmonary disease: implications for

- antibiotic therapy. *J Infect Dis* 1999; 180:404-9.
185. Emerman CL, Effron D, Lukens TW. Spirometric criteria for hospital admission of patients with acute exacerbation of COPD. *Chest* 1991; 99:595-9.
 186. Emerman CL, Lukens TW, Effron D. Physician estimation of FEV₁ in acute exacerbation of COPD. *Chest* 1994; 105:1709-12.
 187. Emerman CL, Cydulka RK. Use of peak expiratory flow rate in emergency department evaluation of acute exacerbation of chronic obstructive pulmonary disease. *Ann Emerg Med* 1996; 27:159-63.
 188. Emerman CL, Connors AF, Lukens TW, Effron D, May ME. Relationship between arterial blood gases and spirometry in acute exacerbations of chronic obstructive pulmonary disease. *Ann Emerg Med* 1989; 18:523-7.
 189. Thompson WH, Nielson CP, Carvalho P, Charan NB, Crowley JJ. Controlled trial of oral prednisone in outpatients with acute COPD exacerbation. *Am J Respir Crit Care Med* 1996; 154:407-12.
 190. Davies L, Angus RM, Calverley PM. Oral corticosteroids in patients admitted to hospital with exacerbations of chronic obstructive pulmonary disease: a prospective randomised controlled trial. *Lancet* 1999; 354:456-60.
 191. Niewoehner DE, Erbland ML, Deupree RH, Collins D, Gross NJ, Light RW, et al. Effect of systemic glucocorticoids on exacerbations of chronic obstructive pulmonary disease. Department of Veterans Affairs Cooperative Study Group. *N Engl J Med* 1999; 340:1941-7.
 192. Maltais F, Ostinelli J, Bourbeau J, Tonnel AB, Jacquemet N, Haddon J, et al. Comparison of nebulized budesonide and oral prednisolone with placebo in the treatment of acute exacerbations of chronic obstructive pulmonary disease: a randomized controlled trial. *Am J Respir Crit Care Med* 2002; 165:698-703.
 193. Connors AF, Jr., Dawson NV, Thomas C, Harrell FE, Jr., Desbiens N, Fulkerson WJ, et al. Outcomes following acute exacerbation of severe chronic obstructive lung disease. The SUPPORT investigators (Study to Understand Prognoses and Preferences for Outcomes and Risks of Treatments). *Am J Respir Crit Care Med* 1996; 154:959-67.
 194. Shepperd S, Harwood D, Jenkinson C, Gray A, Vessey M, Morgan P. Randomised controlled trial comparing hospital at home care with inpatient hospital care. I: three month follow up of health outcomes. *BMJ* 1998; 316:1786-91.
 195. Gravil JH, Al-Rawas OA, Cotton MM, Flanigan U, Irwin A, Stevenson RD. Home treatment of exacerbations of chronic obstructive pulmonary disease by an acute respiratory assessment service. *Lancet* 1998; 351:1853-5.
 196. Moayyedi P, Congleton J, Page RL, Pearson SB, Muers MF. Comparison of nebulised salbutamol and ipratropium bromide with salbutamol alone in the treatment of chronic obstructive pulmonary disease. *Thorax* 1995; 50:834-7.
 197. Fernandez A, Munoz J, de la Calle B, Alia I, Ezpeleta A, de la Cal MA, et al. Comparison of one versus two bronchodilators in ventilated COPD patients. *Intensive Care Med* 1994; 20:199-202.
 198. Barbera JA, Reyes A, Roca J, Montserrat JM, Wagner PD, Rodriguez-Roisin R. Effect of intravenously administered aminophylline on ventilation/perfusion inequality during recovery from exacerbations of chronic obstructive pulmonary disease. *Am Rev Respir Dis* 1992; 145:1328-33.
 199. Mahon JL, Laupacis A, Hodder RV, McKim DA, Paterson NA, Wood TE, et al. Theophylline for irreversible chronic airflow limitation: a randomized study comparing n of 1 trials to standard practice. *Chest* 1999; 115:38-48.
 200. Lloberes P, Ramis L, Montserrat JM, Serra J, Campistol J, Picado C, et al. Effect of three different bronchodilators during an exacerbation of chronic obstructive pulmonary disease. *Eur Respir J* 1988; 1:536-9.
 201. Murciano D, Aubier M, Lecocguic Y, Pariente R. Effects of theophylline on diaphragmatic strength and fatigue in patients with chronic obstructive pulmonary disease. *N Engl J Med* 1984; 311:349-53.
 202. Meyer TJ, Hill NS. Noninvasive positive pressure ventilation to treat respiratory failure. *Ann Intern Med* 1994; 120:760-70.
 203. Lightowler JV, Wedzicha JA, Elliott MW, Ram FS. Non-invasive positive pressure ventilation to treat respiratory failure resulting from exacerbations of chronic obstructive pulmonary disease: Cochrane systematic review and meta-analysis. *BMJ* 2003; 326:185.
 204. American Thoracic Society. International Consensus Conferences in Intensive Care Medicine: noninvasive positive pressure ventilation in acute respiratory failure. *Am J Respir Crit Care Med* 2001; 163(1):283-91.
 205. Bott J, Carroll MP, Conway JH, Keilty SE, Ward EM, Brown AM, et al. Randomised controlled trial of nasal ventilation in acute ventilatory failure due to chronic obstructive airways disease. *Lancet* 1993; 341:1555-7.
 206. Brochard L, Mancebo J, Wysocki M, Lofaso F, Conti G, Rauss A, et al. Noninvasive ventilation for acute exacerbations of chronic obstructive pulmonary disease. *N Engl J Med* 1995; 333:817-22.
 207. Kramer N, Meyer TJ, Meharg J, Cece RD, Hill NS. Randomized, prospective trial of noninvasive positive pressure ventilation in acute respiratory failure. *Am J Respir Crit Care Med* 1995; 151:1799-806.
 208. Plant PK, Owen JL, Elliott MW. Early use of non-invasive ventilation for acute exacerbations of chronic obstructive pulmonary disease on general respiratory wards: a multicentre randomised controlled trial. *Lancet* 2000; 355:1931-5.
 209. Plant PK, Owen JL, Elliott MW. Non-invasive ventilation in acute exacerbations of chronic obstructive pulmonary disease: long term survival and predictors of in-hospital outcome. *Thorax* 2001; 56:708-12.
 210. Esteban A, Anzueto A, Alia I, Gordo F, Apezteguia C, Palizas F, et al. How is mechanical ventilation employed in the intensive care unit? An international utilization review. *Am J Respir Crit Care Med* 2000; 161:1450-8.
 211. Ericsson CH, Svartengren K, Svartengren M, Mossberg B,

- Philipson K, Blomquist M, et al. Repeatability of airway deposition and tracheobronchial clearance rate over three days in chronic bronchitis. *Eur Respir J* 1995; 8:1886-93.
212. Brochard L, Rauss A, Benito S, Conti G, Mancebo J, Rekiq N, et al. Comparison of three methods of gradual withdrawal from ventilatory support during weaning from mechanical ventilation. *Am J Respir Crit Care Med* 1994; 150:896-903.
213. Nava S, Ambrosino N, Clini E, Prato M, Orlando G, Vitacca M, et al. Noninvasive mechanical ventilation in the weaning of patients with respiratory failure due to chronic obstructive pulmonary disease. A randomized, controlled trial. *Ann Intern Med* 1998; 128:721-8.
214. Hilbert G, Gruson D, Portel L, Gbikpi-Benissan G, Cardinaud JP. Noninvasive pressure support ventilation in COPD patients with postextubation hypercapnic respiratory insufficiency. *Eur Respir J* 1998; 11:1349-53.
215. Kessler R, Faller M, Fourgaut G, Mennecier B, Weitzenblum E. Predictive factors of hospitalization for acute exacerbation in a series of 64 patients with chronic obstructive pulmonary disease. *Am J Respir Crit Care Med* 1999; 159:158-64.
216. Mushlin AI, Black ER, Connolly CA, Buonaccorso KM, Eberly SW. The necessary length of hospital stay for chronic pulmonary disease. *JAMA* 1991; 266:80-3.
217. Cotton MM, Bucknall CE, Dagg KD, Johnson MK, MacGreagor G, Stewart C, Stevenson RD. Early discharge for patients with exacerbations of chronic obstructive pulmonary disease: a randomized controlled trial. *Thorax* 2000 Nov;55(11):902-6.
218. Hughes SL, Weaver FM, Giobbie-Hurder A, Manheim L, Henderson W, Kubal JD, Ulasevich A, Cummings J. Effectiveness of team-managed home-based primary care: a randomized multicenter trial. *JAMA* 2000 Dec 13;284(22):2877-85.
219. Hermiz O, Comino E, Marks G, Daffum K, Wilson S, Harris M. Randomised controlled trial of home based care of patients with chronic obstructive pulmonary disease. *BMJ* 2002 Oct 26;325(7370):938
220. Hernandez C. et al. Home hospitalizaation of exacerbated COPD patients. *Eur Respir J* 2003;21:58-67.
221. Stoller JK, Lange PA. Inpatient management of chronic obstructive pulmonary disease. *Respir Care Clin N Am* 1998; 4:425-38.
222. Peach H, Pathy MS. Follow-up study of disability among elderly patients discharged from hospital with exacerbations of chronic bronchitis. *Thorax* 1981; 36:585-9.



DISCRIMINATION PROHIBITED: Under provisions of applicable public laws enacted by Congress since 1964, no person in the United States shall, on the grounds of race, color, national origin, handicap, or age, be excluded from participation in, be denied the benefits of, or be subjected to discrimination under any program or activity (or, on the basis of sex, with respect to any education program or activity) receiving Federal financial assistance. In addition, Executive Order 11141 prohibits discrimination on the basis of age by contractors and subcontractors in the performance of Federal contract, Executive Order 11246 states that no federally funded contractor may discriminate against any employee or applicant for employment because of race, color, religion, sex, or national origin and Executive Order 13087 prohibits discrimination based on sexual orientation. Therefore, the National Heart, Lung, and Blood Institute must be operated in compliance with these laws and Executive Orders.



Printing and distribution has been made possible by educational grants from:



Zambon Group

Visit the GOLD website at www.goldcopd.com