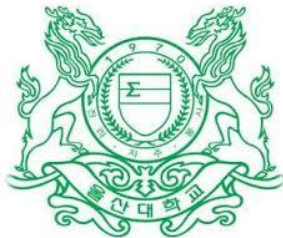


Airway Clearance

- update 및 실제 적용 -

이세원



울산대학교 의과대학
서울아산병원



Bronchiectasis

- Morphologic term used to describe abnormal, irreversibly dilated and thick-walled bronchi
- Represent the end stage of a variety of pathologic process that cause destruction of the bronchial wall and its surrounding supporting tissues
- Anatomic definition that evolved from **Lannec**'s original description in 1819 of ectatic bronchi in pathologic specimens



Fishman's pulmonary diseases

Bronchiectasis: Definition in guidelines

This guideline refers to the investigation and management of patients with **symptoms of persistent or recurrent bronchial sepsis** related to **irreversibly damaged and dilated bronchi** - namely, clinical bronchiectasis

Pasteur et al., Thorax.2010;65 Suppl 1:i1-58

Bronchiectasis is a **suppurative lung disease** with heterogeneous phenotypic features. Bronchiectasis is diagnosed on axial images of HRCT scans

McShane et al., Am J Respir Crit Care Med. 2013;188(6):647-56

Bronchiectasis is a chronic respiratory disease characterized by a clinical syndrome of **cough, sputum production, and bronchial infection**, and radiologically by **abnormal and permanent dilatation of the bronchi**

Polverino et al., Eur Respir J. 2017;50(3).

Definition of BE: Observational Cohorts in Europe


- **Inclusion criteria**

- Aged 18 years and older
- Radiological diagnosis made on **HRCT**
- **Clinical history** consistent with **bronchiectasis**

- **Exclusion criteria**

- Patients with cystic fibrosis or traction BE due to pulmonary fibrosis were all excluded in all four cohort

Europe Bronchiectasis Registry (EMBARC)



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**STUDY PROTOCOL
BRONCHIECTASIS**

The EMBARC European Bronchiectasis Registry: protocol for an international observational study

James D. Chalmers^{1,33}, Stefano Aliberti^{2,33}, Eva Polverino^{3,33}, Montserrat Vendrell⁴, Megan Crichton¹, Michael Loebinger⁵, Katerina Dimakou⁶, Ian Clifton⁷, Menno van der Eerden⁸, Gernot Rohde⁹, Marlene Murriss-Espin¹⁰, Sarah Masefield¹¹, Eleanor Gerada¹², Michal Shteinberg¹³, Felix Ringshausen¹⁴, Charles Haworth¹⁵, Wim Boersma¹⁶, Jessica Rademacher¹⁴, Adam T. Hill¹⁷, Timothy Aksamit¹⁸, Anne O'Donnell¹⁹, Lucy Morgan²⁰, Branislava Milenkovic^{21,22}, Leandro Tramma¹, Joao Neves²³, Rosario Menendez²⁴, Perluigi Paggiaro²⁵, Victor Botnaru²⁶, Sabina Skrgat²⁷, Robert Wilson⁹, Pieter Goeminne²⁸, Anthony De Soyza^{29,30}, Tobias Welte¹⁴, Antoni Torres³, J. Stuart Elborn³¹ and Francesco Blasi³², on behalf of EMBARC

Affiliations: ¹College of Medicine, University of Dundee, Ninewells Hospital and Medical School, Dundee, UK. ²Dept of Health Science, University of Milan Bicocca, Clinica Pneumologica, AD San Gerardo, Monza, Italy. ³Fundació Clinic, IDIBAPS, CIBERES, Hospital Clinic de Barcelona, Barcelona, Spain. ⁴Bronchiectasis Girona Biomedical Research Institute (IDIBGI), Dr Trueta University Hospital, Girona, Spain. ⁵Host Defect Unit, Royal Brompton Hospital, London, UK. ⁶5th Pulmonary Department, 'Sotiria' Chest Hospital, Athens, Greece. ⁷Leeds Centre for Respiratory Medicine, St James's Hospital, Leeds, UK. ⁸Dept of Pulmonary Medicine, Erasmus MC Rotterdam, Rotterdam, The Netherlands. ⁹Dept of Respiratory Medicine, Maastricht University Medical Center, Maastricht, The Netherlands. ¹⁰Pôle des Voies Respiratoires, Hôpital de Larr CHU de Toulouse, Toulouse, France. ¹¹European Lung Foundation, Sheffield, UK. ¹²Dept of Respiratory Medicine, Mater Dei Hospital, Msida, Malta. ¹³Pulmonary Institute, Carmel Medical Center, Haifa, Israel. ¹⁴Dept of Respiratory Medicine, Hannover Medical School, Member of the German Centre for Lung Research, Hannover, Germany. ¹⁵Cambridge Centre for Lung Infection, Papworth Hospital, Cambridge, UK. ¹⁶Dept of Pulmonary Diseases, Medical Centre Alkmaar, Alkmaar, The Netherlands. ¹⁷Royal Infirmary of Edinburgh, University of Edinburgh, Edinburgh, UK. ¹⁸Mayo Clinic College of Medicine, Rochester, MN, USA. ¹⁹Division of Pulmonary, Critical Care and Sleep Medicine, Georgetown University Medical Center, Washington, DC, USA. ²⁰Dept of Respiratory Medicine, Concord Hospital, Concord Clinical School, University of Sydney, Sydney, Australia. ²¹Faculty of Medicine, University of Belgrade, Belgrade, Serbia. ²²Clinic for Pulmonary Diseases, Clinical Center of Serbia, Belgrade, Serbia. ²³Medicina Interna, Centro Hospitalar do Porto, Porto, Portugal. ²⁴Pneumology Service, University and Polytechnic Hospital La Fe, Valencia, Spain. ²⁵Dept of Surgery, Medicine, Molecular Biology and Critical Care, University of Pisa, Pisa, Italy. ²⁶Division of Pneumology, Dept of Internal Medicine, State Medical University of Medicine and Pharmacy 'Nicolae Testemitanu' Chisinau, Republic of Moldova. ²⁷University Clinic of Respiratory and Allergic Diseases Golnik, Golnik, Slovenia. ²⁸Dept of Respiratory Medicine, AZ Nikolaas, Sint-Niklaas, Belgium. ²⁹Institute of Cellular Medicine, Newcastle University, Newcastle upon Tyne, UK. ³⁰Bronchiectasis Service, Freeman Hospital, Newcastle upon Tyne, UK. ³¹School of Medicine, Dentistry and Biomedical Sciences, Queen's University of Belfast, Belfast, UK. ³²Respiratory Unit, IRCCS Fondazione Cà Granda Milano, Dept of Pathophysiology and Transplantation, University of Milan, Milan, Italy. ³³These authors contributed equally.

Correspondence: James D. Chalmers, College of Medicine, University of Dundee, Ninewells Hospital and Medical School, Dundee, DD1 9SY, UK. E-mail: jchalmers@dundee.ac.uk

The European Bronchiectasis Registry will recruit 10 000 patients over 5 years.

The inclusion criteria of this study are:

- a **clinical history** consistent with bronchiectasis (**cough, chronic sputum production** and/or **recurrent respiratory infections**)

- **computed tomography** of chest demonstrating bronchiectasis (bronchial dilatation) affecting one or more lobes.

The exclusion criteria are:

- bronchiectasis due to known cystic fibrosis;
- age <18 years

US Bronchiectasis Registry

[Original Research **Bronchiectasis**]



Adult Patients With Bronchiectasis A First Look at the US Bronchiectasis Research Registry



Timothy R. Aksamit, MD; Anne E. O'Donnell, MD; Alan Barker, MD; Kenneth N. Olivier, MD; Kevin L. Winthrop, MD; M. Leigh Anne Daniels, MD, MPH; Margaret Johnson, MD; Edward Eden, MD; David Griffith, MD; Michael Knowles, MD; Mark Metersky, MD; Matthias Salathe, MD; Byron Thomashow, MD; Gregory Tino, MD; Gerard Turino, MD; Betsy Carretta, MPH; and Charles L. Daley, MD; for the Bronchiectasis Research Registry Consortium

OBJECTIVES: We sought to describe the characteristics of adult patients with bronchiectasis enrolled in the US Bronchiectasis Research Registry (BRR).

METHODS: The BRR is a database of patients with non-cystic-fibrosis bronchiectasis (NCFB) enrolled at 13 sites in the United States. Baseline demographic, spirometric, imaging, microbiological, and therapeutic data were entered into a central Internet-based database. Patients were subsequently analyzed by the presence of NTM.

RESULTS: We enrolled 1,826 patients between 2008 and 2014. Patients were predominantly women (79%), white (89%), and never smokers (60%), with a mean age of 64 ± 14 years. Sixty-three percent of the patients had a history of NTM disease or NTM isolated at baseline evaluation for entry into the BRR. Patients with NTM were older, predominantly women, and had bronchiectasis diagnosed at a later age than those without NTM. Gastroesophageal reflux disease (GERD) was more common in those with NTM, whereas asthma, primary immunodeficiency, and primary ciliary dyskinesia were more common in those without NTM. Fifty-one percent of patients had spirometric evidence of airflow obstruction. Patients with NTM were more likely to have diffusely dilated airways and tree-in-bud abnormalities. *Pseudomonas* and *Staphylococcus aureus* isolates were cultured less commonly in patients with NTM. Bronchial hygiene measures were used more often in those with NTM, whereas antibiotics used for exacerbations, rotating oral antibiotics, steroid use, and inhaled bronchodilators were more commonly used in those without NTM.

CONCLUSIONS: Adult patients with bronchiectasis enrolled in the US BRR are described, with differences noted in demographic, radiographic, microbiological, and treatment variables based on stratification of the presence of NTM. CHEST 2017; 151(5):982-992

KEY WORDS: airways; bronchiectasis; nontuberculous mycobacteria; *Pseudomonas*; registry

Adult patients with a **physician-established diagnosis of bronchiectasis**

The exclusion of patients with primary CF was established based on clinical history, previous sweat chloride test results, genetic testing results, or a combination

Patients registered with an NTM patient advocacy group and participating in online surveys are not necessarily representative of the general bronchiectasis population and may be skewed toward patients with NTM disease.

KMBARC (Korean Multicenter Bronchiectasis Audit and Research Collaboration)

Open access

Protocol

BMJ Open KMBARC registry: protocol for a multicentre observational cohort study on non-cystic fibrosis bronchiectasis in Korea

Hyun Lee,¹ Hayoung Choi,² Yun Su Sim,² Shinhee Park,³ Woo Jin Kim,⁴ Kwang Ha Yoo,⁵ Seung Jun Lee,⁶ Tae-Hyung Kim,⁷ Bumhee Yang,⁸ Ina Jeong,⁹ Soo-Jung Um,¹⁰ Deog Kyeom Kim,¹¹ Ji-Hyun Lee,¹² Byoung Soo Kwon,¹³ Young-Jae Cho,¹³ Hye Yun Park,¹⁴ Chang-Hoon Lee,¹⁵ Chin Kook Rhee,¹⁶ Sang Haak Lee,¹⁷ Ju Ock Na,¹⁸ An-Soo Jang,¹⁹ Ji Ye Jung,²⁰ Seung Won Ra,²¹ Ji-Ho Lee,²² Sang-Ha Kim,²² Changhwan Kim,²³ Youlim Kim,²⁴ Chang Youl Lee,²⁴ Hyun Kuk Kim,²⁵ Jae Seung Lee,²⁶ Sei Won Lee,²⁶ Yeon-Mok Oh,²⁶ on behalf of the KMBARC

To cite: Lee H, Choi H, Sim YS, et al. KMBARC registry: protocol for a multicentre observational cohort study on non-cystic fibrosis bronchiectasis in Korea. *BMJ Open* 2020;10:e034090. doi:10.1136/bmjopen-2019-034090

► Prepublication history and additional material for this paper are available online. To view these files, please visit the journal online (<http://dx.doi.org/10.1136/bmjopen-2019-034090>).

HL and HC contributed equally.

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For numbered affiliations see end of article.

Correspondence to
Dr Yeon-Mok Oh;
yeonmok.oh@gmail.com

ABSTRACT

Introduction Despite the significant disease burden of bronchiectasis in Korea, no large-scale, representative prospective cohort studies have been conducted to evaluate the clinical characteristics of Korean patients with bronchiectasis, indicating an urgent need for cohort studies on bronchiectasis.

Methods and analysis The Korean Multicenter Bronchiectasis Audit and Research Collaboration (KMBARC) is a prospective, non-interventional observational cohort study on bronchiectasis in Korea. The inclusion criteria of this registry are as follows: (1) adult patients (aged ≥ 18 years) with or without respiratory symptoms (cough, chronic sputum and/or recurrent respiratory infection) and chest computed tomography revealing bronchiectasis affecting one or more lobes and (2) stable status at the time of registration; patients with bronchiectasis who were admitted for a respiratory aetiology can be enrolled at least 4 weeks after hospital discharge. The exclusion criteria are as follows: (1) bronchiectasis due to cystic fibrosis; (2) traction bronchiectasis associated with interstitial lung disease; (3) patients actively being treated for pneumonia, pulmonary tuberculosis or non-tuberculous mycobacterial infection; (4) patients who are unable or unwilling to provide informed consent; and (5) pregnant patients. Although the KMBARC questionnaires for baseline and annual follow-up data are similar to the European Multicenter Bronchiectasis Audit and Research Collaboration questionnaires, KMBARC has distinctive features such as use of Bronchiectasis Health Questionnaires, measurement with fatigue and depression scales, blood tests, use of consensus definition of exacerbations and information on emergency room or hospitalisation.

We aim to recruit at least 1200 patients over the study period from more than 26 hospitals in South Korea. Patients will undergo a detailed baseline and yearly assessment for up to 5 years. The study objectives of the KMBARC registry are as follows: (1) uncovering the

Strengths and limitations of this study

- This is the first prospective cohort study on patients with bronchiectasis in Korea.
- We will recruit and follow-up patients with bronchiectasis annually using a standardised protocol to improve the quality of data collection.
- Sharing similar case-reporting forms with European Multicenter Bronchiectasis Audit and Research Collaboration (EMBARC) will allow collaboration studies with EMBARC.
- Distinctive features of the Korean Multicenter Bronchiectasis Audit and Research Collaboration registry (KMBARC) will provide several novel findings of bronchiectasis, which might be difficult to be elucidated using other registries.
- This study is limited by a lack of collecting patient samples.

natural course of bronchiectasis; (2) aiding in establishing evidence-based bronchiectasis guidelines in Korea; and (3) encouraging and facilitating studies on bronchiectasis in Korea.

Ethics and dissemination This study received necessary approval from the Institutional Review Boards of all participating institutions. The Asan Medical Center Institutional Review Board gave overall approval for the study. Results will be disseminated via peer-reviewed publications and conference presentations.
Trial registration number KCT0003088.

INTRODUCTION

Bronchiectasis is a chronic respiratory disease characterised by irreversible bronchial dilatation.¹ Patients with bronchiectasis often develop chronic respiratory

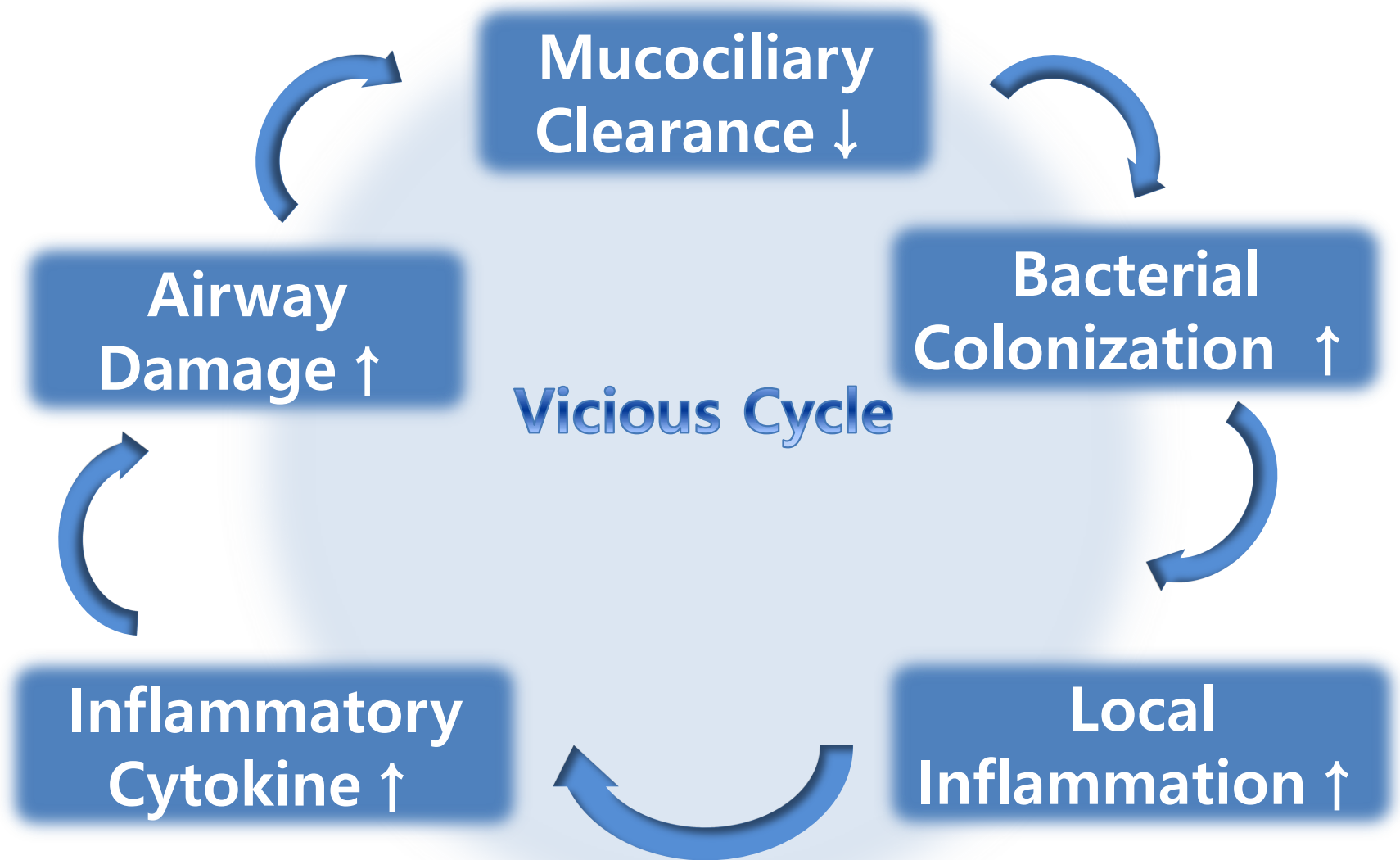
Inclusion

- Adult patients (aged ≥ 18 years) with or without respiratory symptoms (**cough, chronic sputum and/or recurrent respiratory infection**)
- Chest CT: bronchiectasis affecting one or more lobes
- Stable status at the time of registration

Exclusion

- Bronchiectasis due to cystic fibrosis
- Traction bronchiectasis associated with ILD
- Actively being treated for pneumonia, TB or NTM

Pathogenesis: Vicious cycle hypothesis



Guidelines

BTS guidelines

British Thoracic Society guideline for non-CF bronchiectasis

M C Pasteur¹, D Hilton², A T Hill³ on behalf of the British Thoracic Society

BTS Guideline

British Thoracic Society Guideline for bronchiectasis in adults

Adam T Hill,¹ Anita L Sullivan,² James D Chalmers,³ Anthony De Soya,⁴ J Stuart Elborn,⁵ R Andres Floto,^{6,7} Lizzie Grillo,⁸ Kevin Gruffydd-Jones,⁹ Alex Harvey,¹⁰ Charles S Haworth,⁷ Edwin Hiscocks,¹¹ John R Hurst,¹² Christopher Johnson,⁷ W Peter Kelleher,^{13,14,15} Pallavi Bedi,¹⁶ Karen Payne,¹⁷ Hashem Saleh,⁹ Nicholas J Screaton,¹⁸ Maeve Smith,¹⁹ Michael Tunney,²⁰ Deborah Whitters,²¹ Robert Wilson,¹⁴ Michael R Loebinger¹⁴

SUMMARY OF RECOMMENDATIONS AND GOOD PRACTICE POINTS

How should the diagnosis of bronchiectasis be determined?

- Recommendations – Imaging
- > Perform baseline chest X-ray in patients with suspected bronchiectasis. (D)
 - > Perform a thin section computed tomography scan (CT) to confirm a diagnosis of bronchiectasis when clinically suspected. (C)
 - > Perform baseline imaging during clinically stable disease as this is optimal for diagnostic and serial comparison purposes. (D)

Good practice points

CT imaging protocol

- ✓ The most appropriate imaging protocol will vary according to scanner technology and patient factors.
- ✓ When using volumetric CT, dose reduction techniques including adaptive mA and kV and iterative reconstruction should be utilised where available.
- ✓ Typical CT imaging parameters for the diagnosis of bronchiectasis are:
 - Slice thickness: ≤1mm
 - Reconstruction algorithm: – high spatial frequency
 - kVp: 100–140
 - mAs (or effective mAs): 100 – 200
 - Gantry rotation time: <0.5s

CT features of bronchiectasis

- ✓ Bronchiectasis is defined by bronchial dilatation as suggested by one or more of the following:
 - Bronchoarterial ratio >1 (internal airway lumen vs adjacent pulmonary artery)
 - Lack of tapering
 - Airway visibility within 1cm of costal pleural surface or touching mediastinal pleura.
- ✓ The following indirect signs are commonly associated with bronchiectasis:
 - Bronchial wall thickening
 - Mucus impaction
 - Mosaic perfusion / air trapping on expir-

General

- ✓ CT scanning can also aid in identifying an aetiology of bronchiectasis eg Allergic bronchopulmonary aspergillosis (ABPA), Non-tuberculous mycobacteria (NTM), primary ciliary dyskinesia, alpha one antitrypsin deficiency, Williams Campbell syndrome and a foreign body.

In whom should the diagnosis of bronchiectasis be suspected?

Recommendations

- > Consider investigation for bronchiectasis in patients with persistent production of mucopurulent or purulent sputum particularly with relevant associated risk factors. (D)
- > Consider investigation for bronchiectasis in patients with rheumatoid arthritis if they have symptoms of chronic productive cough or recurrent chest infections. (C)
- > Consider investigation for bronchiectasis in patients with Chronic Obstructive Pulmonary Disease (COPD) with frequent exacerbations (two or more annually) and a previous positive sputum culture for *P. aeruginosa* whilst stable. (B)
- > Consider investigation for bronchiectasis in patients with inflammatory bowel disease and chronic productive cough. (C)

Good practice points

- ✓ In at risk groups, if bronchiectasis is suspected, bronchiectasis needs confirmation.
- ✓ In patients with COPD, investigation for bronchiectasis may be appropriate especially in the presence of chronic productive cough with positive sputum cultures for potentially pathogenic microorganisms (PPM) whilst stable or 2 or more exacerbations in the preceding 12 months.
- ✓ In patients with asthma, investigation for bronchiectasis may be appropriate with severe or poorly-controlled disease.
- ✓ In patients with a history of HIV-1 infection, solid organ and bone marrow transplant, and history of immunosuppressive therapy for lymphoma and vasculitis, investigation for bronchiectasis may be appropriate with symptoms of chronic productive cough or recurrent

European Respiratory Society guidelines for the management of adult bronchiectasis

Eva Polverino¹, Pieter C. Goeminne^{2,3}, Melissa J. McDonnell^{4,5,6}, Stefano Aliberti⁷, Sara E. Marshall⁸, Michael R. Loebinger⁹, Marlene Murriss¹⁰, Rafael Cantón¹¹, Antoni Torres¹², Katerina Dimakou¹³, Anthony De Soya^{14,15}, Adam T. Hill¹⁶, Charles S. Haworth¹⁷, Montserrat Vendrell¹⁸, Felix C. Ringshausen¹⁹, Dragan Subotic²⁰, Robert Wilson⁹, Jordi Vilaró²¹, Bjorn Stallberg²², Tobias Welte¹⁹, Gernot Rohde²³, Francesco Blaszi⁷, Stuart Elborn^{9,24}, Marta Almagro²⁵, Alan Timothy²⁵, Thomas Ruddy²⁵, Thomy Tonia²⁶, David Rigau²⁷ and James D. Chalmers²⁸

@ERSpublications

The publication of the first ERS guidelines for bronchiectasis <http://ow.ly/wQSO30dU0nE>

Cite this article as: Polverino E, Goeminne PC, McDonnell MJ, et al. European Respiratory Society guidelines for the management of adult bronchiectasis. *Eur Respir J* 2017; 50: 1700629 [https://doi.org/10.1183/13993003.00629-2017].

ABSTRACT Bronchiectasis in adults is a chronic disorder associated with poor quality of life and frequent exacerbations in many patients. There have been no previous international guidelines.

The European Respiratory Society guidelines for the management of adult bronchiectasis describe the appropriate investigation and treatment strategies determined by a systematic review of the literature.

A multidisciplinary group representing respiratory medicine, microbiology, physiotherapy, thoracic surgery, primary care, methodology and patients considered the most relevant clinical questions (for both clinicians and patients) related to management of bronchiectasis. Nine key clinical questions were generated and a systematic review was conducted to identify published systematic reviews, randomised clinical trials and observational studies that answered these questions. We used the GRADE approach to define the quality of the evidence and the level of recommendations. The resulting guideline addresses the investigation of underlying causes of bronchiectasis, treatment of exacerbations, pathogen eradication, long term antibiotic treatment, anti-inflammatories, mucocactive drugs, bronchodilators, surgical treatment and respiratory physiotherapy.

These recommendations can be used to benchmark quality of care for people with bronchiectasis across Europe and to improve outcomes.



▶ <http://dx.doi.org/10.1183/2018-212468>

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To cite: Hill AT, Sullivan AL, Chalmers JD, et al. *Thorax* 2018;69:1-22.

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¹Norfolk and Norwich Unk Hospital, Norfolk, UK
²Royal Brompton Hospital, London, UK
³Royal Infirmary of Edinburgh, Edinburgh, UK

Correspondence to M C Pasteur, Norfolk and Norwich University Hospital, Cahey Lane, Norwich NR4 7UK; mark.pasteur@nruh.nhs.uk

Received 29 January 2010
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For numbered affiliations see end of article.

Correspondence to Professor Adam T Hill, Respiratory Medicine, Royal Infirmary of Edinburgh, Edinburgh and University of Edinburgh, EH16 4SA, UK; adam.hill318@rifs.net

Non-pharmacologic Treatment of Bronchiectasis

- Active cyclic breathing
- Oscillating positive expiratory devices
- Postural drainage
- Forced expiration technique
- Pulmonary rehabilitation
- Inspiratory muscle training

Non-pharmacologic Treatment of Bronchiectasis

- Active cyclic breathing
- Oscillating positive expiratory devices
- Postural drainage
- Forced expiration technique
- Pulmonary rehabilitation
- Inspiratory muscle training

: ACBT with OPAP is emphasized

- The active cycle of breathing techniques (plus postural drainage) and oscillating positive expiratory devices (plus postural drainage and the forced expiration technique) should be considered when offering individuals with non-CF bronchiectasis effective airway clearance techniques.
- Recombinant human DNase should not be used in adults with bronchiectasis.
- The evidence base remains weak for the majority of interventions used in daily clinical practice.

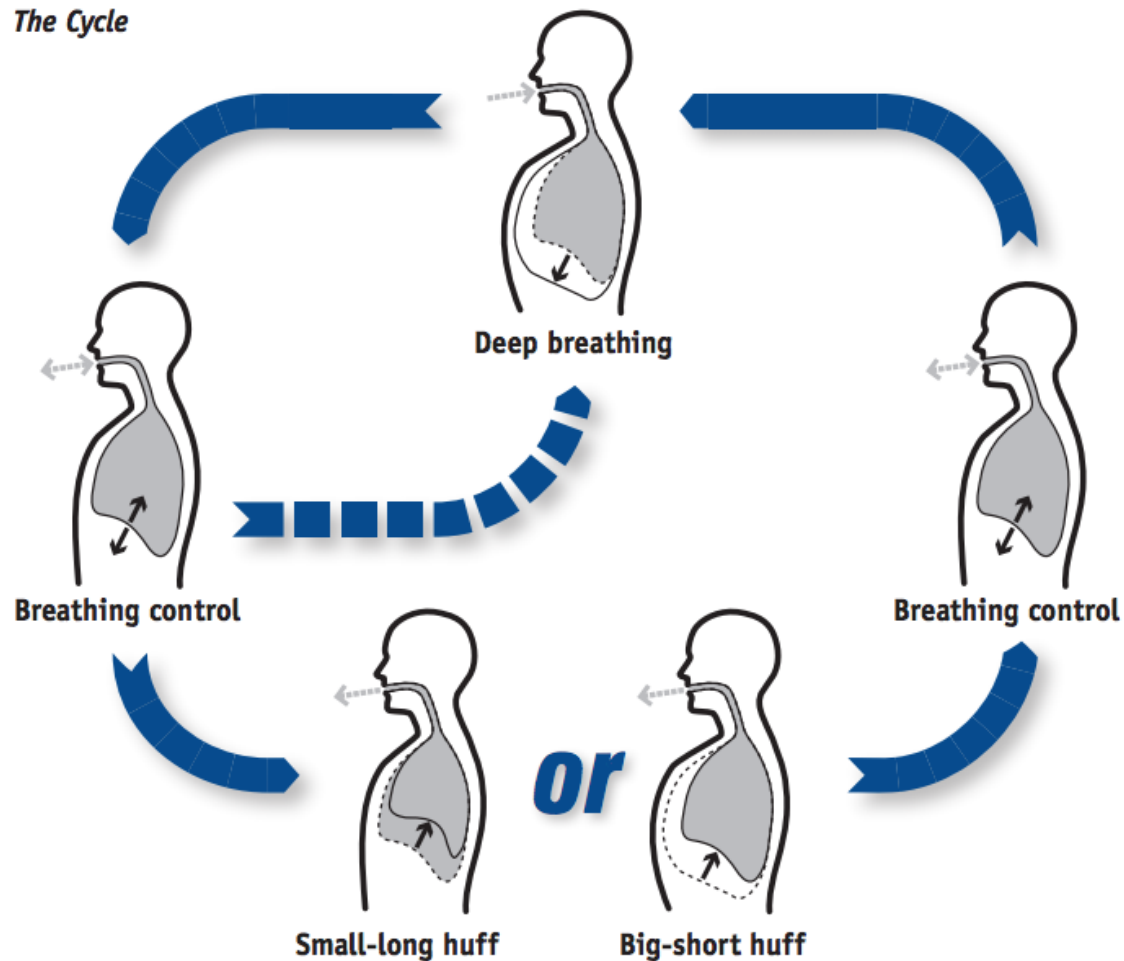
: Physiotherapy is also emphasized

- The inclusion of postural drainage should be considered for all airway clearance techniques.
- The inclusion of the forced expiration technique should be considered for all airway clearance techniques.

Physiotherapy

가래배출법	능동적	<ul style="list-style-type: none">▫ 기침법▫ 허핑▫ 가슴팽창운동▫ 능동주기 호흡법 (ACBT)▫ 자가 배출법▫ 보조기침법과 덧대기 기침법
	수동적	<ul style="list-style-type: none">▫ 자세변경 배출법▫ 두드리기·진동법 및 흔들기▫ 가슴팽창운동과 자세변경 배출법의 활용
	도구	<ul style="list-style-type: none">▫ 에어로비카 · 플루터 · 아카펠라▫ BPEP▫ HFCO · IPPB

Active Cyclic Breathing

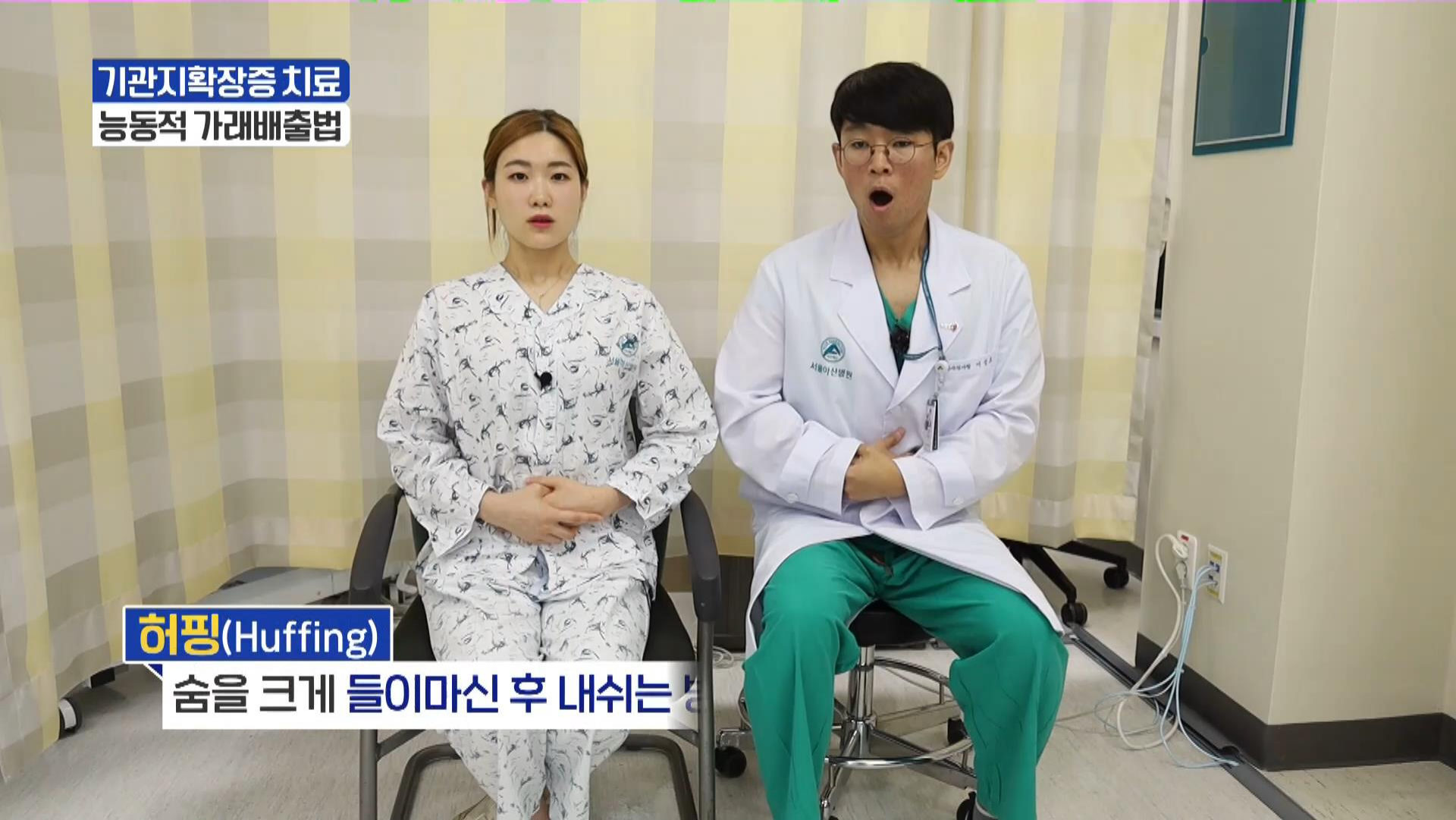


Small-long puff and Big-Short puff

기관지확장증 치료
능동적 가래배출법

허핑(Huffing)

숨을 크게 들이마신 후 내쉬는 방법



Active Cyclic Breathing

기관지확장증 치료
능동적 가래배출법

능동주기 호흡기술(ACBT : active cycle of breathing technique)

의식 장애가 없고 기도분비물이 많은 환자에게
호흡의 크기를 바꾸어 가래배출을 촉진하는 방법

Airway clearance in bronchiectasis: a randomized crossover trial of active cycle of breathing techniques (incorporating postural drainage and vibration) versus test of incremental respiratory endurance

JE Patterson¹, JM Bradley^{1,2} and JS Elborn^{2,3*}

¹School of Rehabilitation Sciences, University of Ulster at Jordanstown, Northern Ireland BT37 0QB, UK;

²Regional Adult CF Centre, Belfast City Hospital, Belfast, Northern Ireland BT9 7AB, UK; and

³Department of Respiratory Medicine, Queen's University Belfast, University Street, Belfast, Northern Ireland BT7 1SU, UK

Table 1 Weights of sputum expectorated with ACBT and TIRE

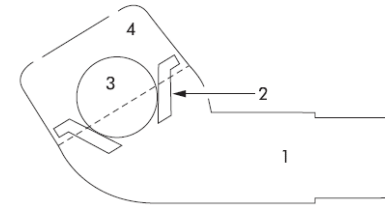
	ACBT	TIRE	Mean difference (95% CI)	P-value
Sputum weight (during treatment) (g)	6.34 (6.63)	4.04 (4.26)	2.30 (0.52 to 4.08)	0.01
Sputum weight (30 min post-treatment) (g)	2.66 (2.61)	2.50 (3.26)	0.16 (-1.02 to 1.34)	0.78
Total (g)	8.98 (7.82)	6.53 (6.83)	2.44 (0.43 to 4.45)	0.02

Table 2 Pre- and post-treatment lung function and SpO₂

	ACBT Mean difference (95% CI)	P-value	TIRE Mean difference (95% CI)	P-value
FEV₁ (%pred)	2.25 (-2.48 to 6.98)	0.33	-0.03 (-2.56 to 1.96)	0.78
FVC (%pred)	0.65 (-2.80 to 4.10)	0.66	1.45 (-1.07 to 3.97)	0.22
FVC (L)	0.006 (-0.12 to 0.14)	0.70	0.03 (-0.04 to 0.10)	0.24
PEF (%pred)	2.95 (-0.44 to 6.34)	0.93	3.65 (-0.95 to 8.25)	0.34
PEF (L/min)	10.30 (-3.69 to 24.29)	0.08	15.70 (-3.60 to 35.00)	0.11
SpO ₂ (%)	1.20 (-1.07 to 3.47)	0.14	1.90 (-0.32 to 4.12)	0.11
		0.28		0.89

Randomised crossover study of the Flutter device and the active cycle of breathing technique in non-cystic fibrosis bronchiectasis

C S Thompson, S Harrison, J Ashley, K Day, D L Smith



Sputum weights: 26.6 g (15.0–45.2) for ACBT
23.4 g (16.8–36.2) for the Flutter ($p>0.05$).

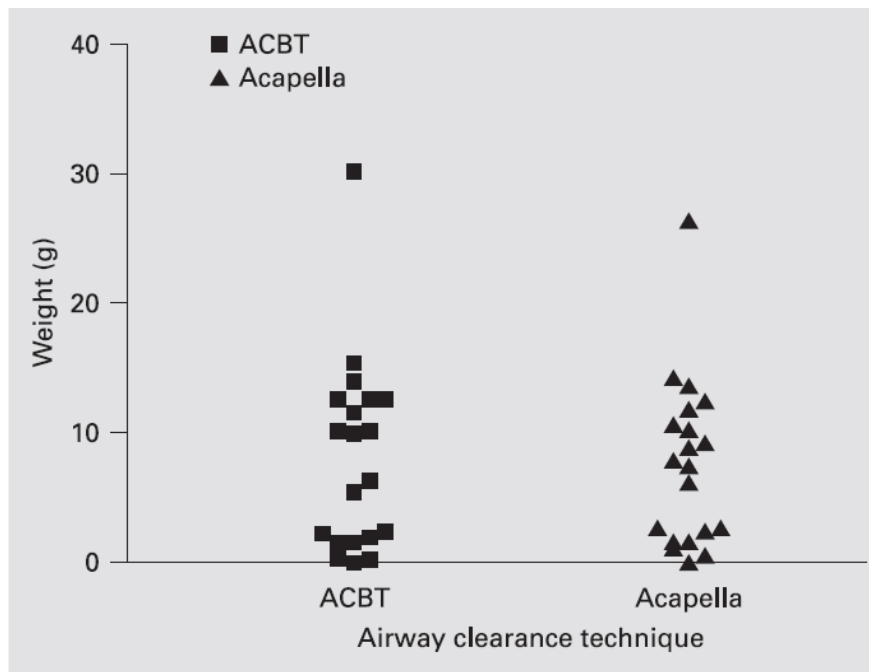
There was a statistically significant improvement in FEV₁ with the Flutter, but this did not achieve a clinically meaningful change.

11/17: patients preferred the Flutter for routine daily use,
3/17: preferred ACBT,
3/17: three had no preference.

Airway Clearance in Bronchiectasis: A Randomized Crossover Trial of Active Cycle of Breathing Techniques versus Acapella®

Janet E. Patterson Judy M. Bradley Oonagh Hewitt Ian Bradbury
J. Stuart Elborn

Health and Rehabilitation Sciences Research Institute, University of Ulster, Newtownabbey, Northern Ireland, UK



	ACBT before treatment mean ± SD	ACBT after treatment mean ± SD	p value (95% CI)	Acapella before treatment mean ± SD	Acapella after treatment mean ± SD	p value (95% CI)
FEV ₁ , % predicted	65±20	65±22	0.56 (-2.02 to 1.12)	65±22	65±22	0.88 (-2.42 to 2.82)
FVC, % predicted	81±20	81±20	0.40 (-2.23 to 0.93)	80±19	81±20	0.48 (-3.42 to 1.82)
PEF, % predicted	67±19	68±19	0.89 (-3.50 to 2.20)	69±20	70±18	0.34 (-5.48 to 1.98)
SpO ₂ , %	97±1	96±2	0.23 (-0.18 to 0.68)	97±1	97±1	0.06 (-0.81 to 1.31)

A randomized evaluation of the acute efficacy, acceptability and tolerability of Flutter and active cycle of breathing with and without postural drainage in non-cystic fibrosis bronchiectasis

Table 2 (a) Comparison of acute sputum clearance measures with Flutter and ACBT with and without PD

<i>Results expressed as mean (SD)</i>	<i>Flutter</i>	<i>ACBT</i>	<i>ACBT-PD</i>
Sputum wet wt (g) clearance	4.4 (6.1)	4.1 (5.7)	10.0 (12.4)
Sputum wet wt (g) post-30 min	1.2 (1.8)	1.5 (2.1)	11.7 (15.1)
Total wet wt (g)	5.6 (7.5)	5.6 (6.7)	11.2 (13.3)
Sputum vol (mL) clearance	6.0 (8.5)	5.4 (8.0)	11.1 (15.1)
Sputum vol (mL) post-30 min	1.9 (3.1)	1.9 (2.4)	12.6 (15.9)
Total vol (mL)	7.9 (11.4)	7.3 (9.6)	12.6 (15.9)
Duration (min)	15.0 (7.8)	15.8 (7.6)	

(b) Comparison of mean differences in acute sputum clearance between Flutter and ACBT with and without PD

<i>Results expressed as mean difference (SD)</i>	<i>Flutter versus ACBT</i>	<i>Flutter versus ACBT-PD</i>	<i>ACBT versus ACBT-PD</i>
Sputum wet wt (g)	0.3 (3.4)	-5.6 (8.2)*	-5.9 (9.6)*
Sputum wet wt (g) post-30-min	-0.3 (2.0)	0.01 (1.6)	0.3 (1.6)
Total wet wt (g)	0.0 (3.7)	-5.6 (8.5)**	-5.6 (9.2)**
Sputum wet vol (mL)	0.6 (3.9)	-5.1 (8.8)*	-5.7 (10.5)*
Sputum wet vol (mL) post-30-min	0.1 (2.2)	0.5 (2.5)	0.4 (1.8)
Total vol (mL)	0.9 (4.7)	-4.9 (8.2)**	-5.3 (9.9)**
Duration	-0.7 (7.0)	-1.7 (7.3)	-1.4 (9.5)

P-value: * <0.01 ; ** <0.001 .

A randomized evaluation of the acute efficacy, acceptability and tolerability of Flutter and active cycle of breathing with and without postural drainage in non-cystic fibrosis bronchiectasis

Results expressed as mean (SD) Likert scale
(1 = extremely to 7 = not at all)

	Flutter	ACBT	ACBT-PD
Useful at clearing sputum	Flutter 2.4 (1.2)	ACBT 2.3 (1.1)	ACBT-PD 2.8 (1.6)
Easy to perform	4.7 (1.8)	5.4 (1.3)	5.0 (1.6)
Discomfort	Flutter 6.3 (0.9)	ACBT 6.0 (1.3)	ACBT-PD 5.3 (1.5)
Time-consuming	5.9 (1.6)	6.0 (1.6)	5.4 (2.1)
Interferes with everyday life			
Embarrassing			

(b) Comparison of mean differences in acceptability and tolerability of Flutter and ACBT with and without PD

Results expressed as mean difference (SD)	Flutter versus ACBT	Flutter versus ACBT-PD	ACBT versus ACBT-PD
Useful at clearing sputum	-0.3 (2.3)	0.6 (2.0)	1.0 (1.9)*
Easy to understand	0.2 (0.8)	0.2 (0.9)	-0.1 (0.7)
Easy to perform	0.1 (1.4)	-0.4 (1.8)	-0.5 (1.9)
Tiring	-0.7 (1.7)	-0.3 (2.0)	0.4 (1.8)
Discomfort (why* text)	-0.4 (1.6)	0.3 (1.6)	0.7 (1.4)*
Time-consuming	-0.2 (1.6)	1.1 (1.8)**	1.3 (1.4)**
Interferes with everyday life	0.4 (1.1)	1 (1.6)***	0.6 (1.3)
Embarrassing	-0.1 (1.3)	0.5 (2.0)	0.6 (2.2)



A randomised crossover trial of chest physiotherapy in non-cystic fibrosis bronchiectasis

M.P. Murray*, J.L. Pentland[#] and A.T. Hill*

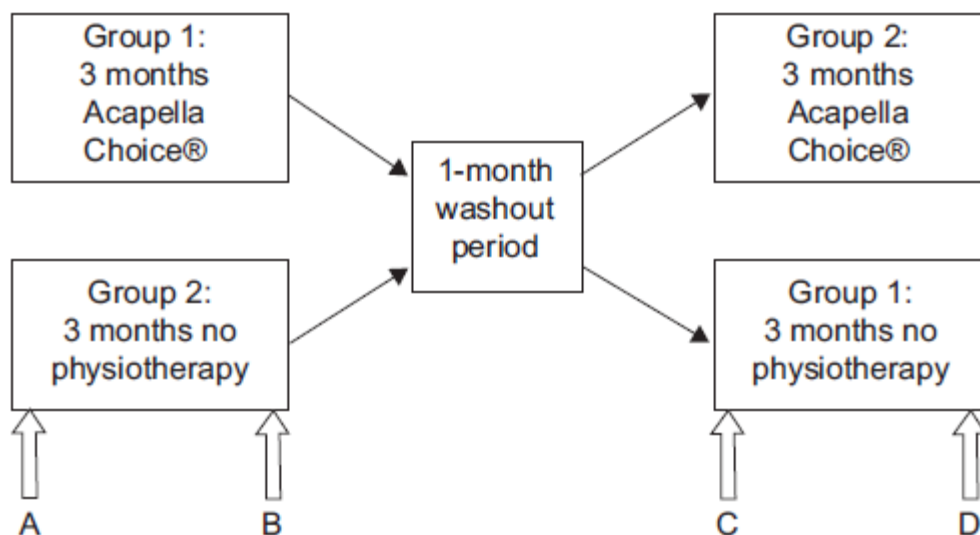
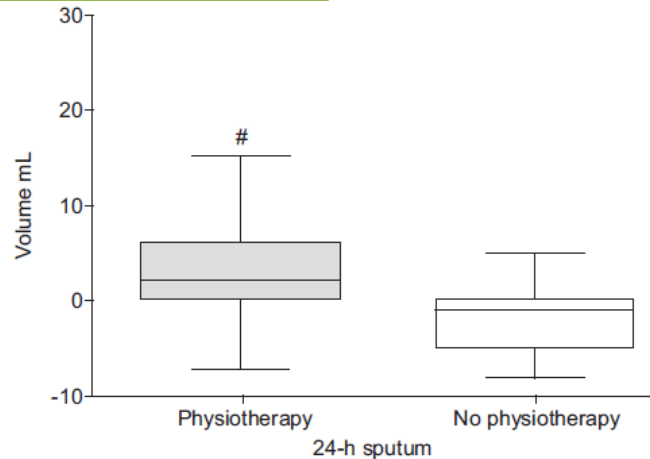


TABLE 1 Patient characteristics

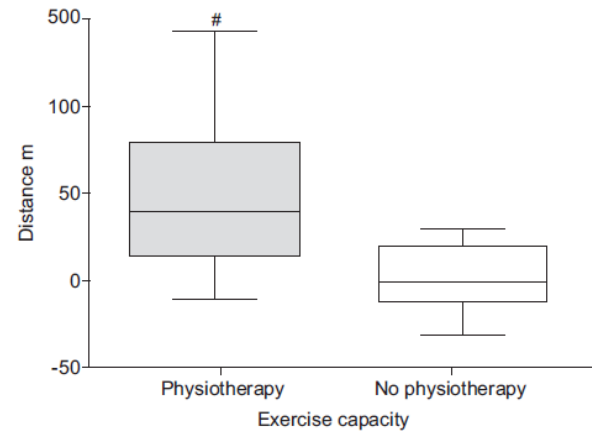
Study participants[#]	20
Male	12 (60)
Age yrs	73 (72–77)
Ex-smokers	8 (40)
Chronic cardiac disease	3 (15)
Neurological disease	1 (5)
Chronic renal impairment	1 (5)
Diabetes mellitus	0
Inhaled corticosteroid therapy	12 (60)
Systemic corticosteroid therapy	0
Long-term antibiotic therapy	2 (10)
Infective exacerbations requiring antibiotic treatment in preceding 12 months	2 (1.5–3)
Lobes affected with bronchiectasis on HRCT	4 (3–4.75)
Varicose or cystic dilatation affecting ≥ 1 lobe	15 (75)
Chronically colonised with pathogenic organisms in sputum when stable	14 (70)
<i>Pseudomonas aeruginosa</i>	6 (42.9)
<i>Haemophilus influenzae</i>	5 (35.7)
<i>Staphylococcus aureus</i>	2 (14.3)
<i>Moraxella catarrhalis</i>	1 (7.1)
Aetiology of bronchiectasis	
Post-infective	10 (50)
Idiopathic	8 (40)
Inactive allergic bronchopulmonary aspergillosis	1 (5)
Inflammatory bowel disease	1 (5)

OPAP is effective in sputum expectoration and improving QOL and exercise capacity

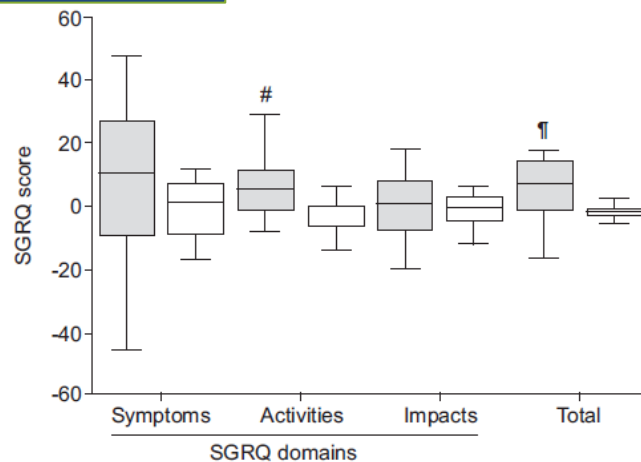
Sputum volume



Walking distance



SGRQ



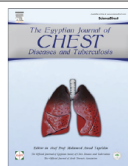
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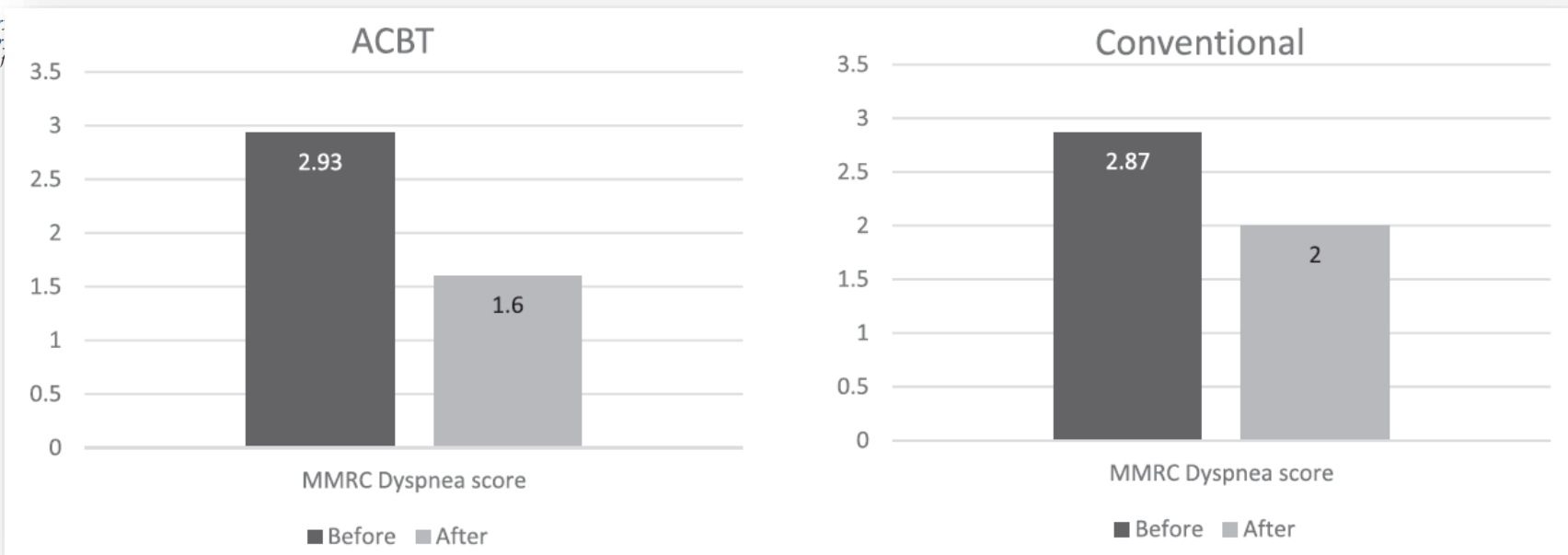
ORIGINAL ARTICLE

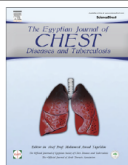
Comparison between active cycles of breathing with postural drainage versus conventional chest physiotherapy in subjects with bronchiectasis



Hesham A. AbdelHalim ^{a,*}, Heba H. AboElNaga ^b, Karim A. Fathy ^c

^a Pulmonar,
^b Pulmonar,
^c Faculty of





ORIGINAL ARTICLE

Comparison between active cycles of breathing with postural drainage versus conventional chest physiotherapy in subjects with bronchiectasis

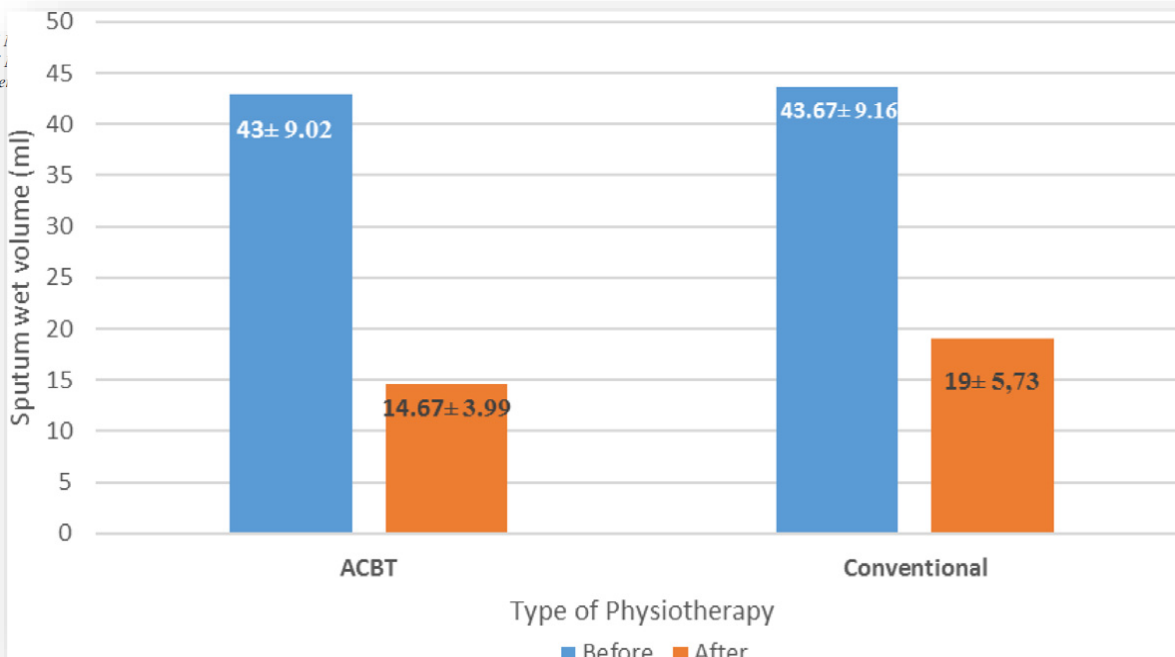


Hesham A. AbdelHalim ^{a,*}, Heba H. AboElNaga ^b, Karim A. Fathy ^c

^a Pulmonary Medicine Department, Faculty of Medicine, Assiut University, Assiut, Egypt

^b Pulmonary Medicine Department, Faculty of Medicine, Assiut University, Assiut, Egypt

^c Faculty of Physical Therapy, October 6 University, Giza, Egypt

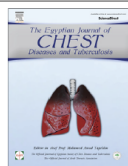


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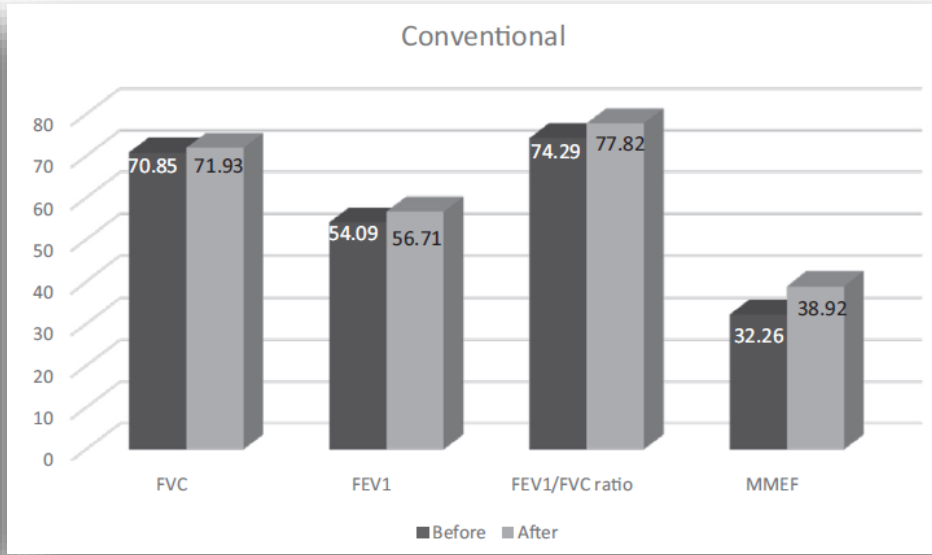
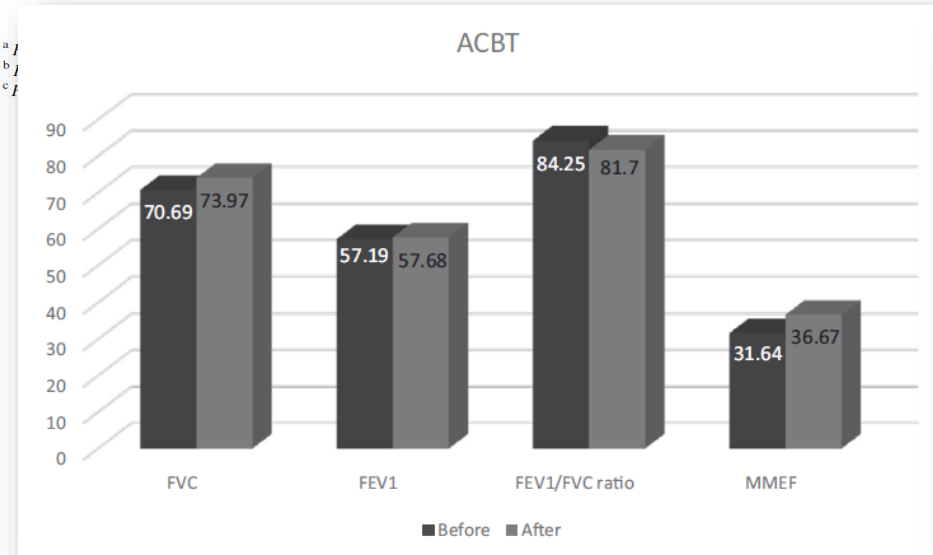


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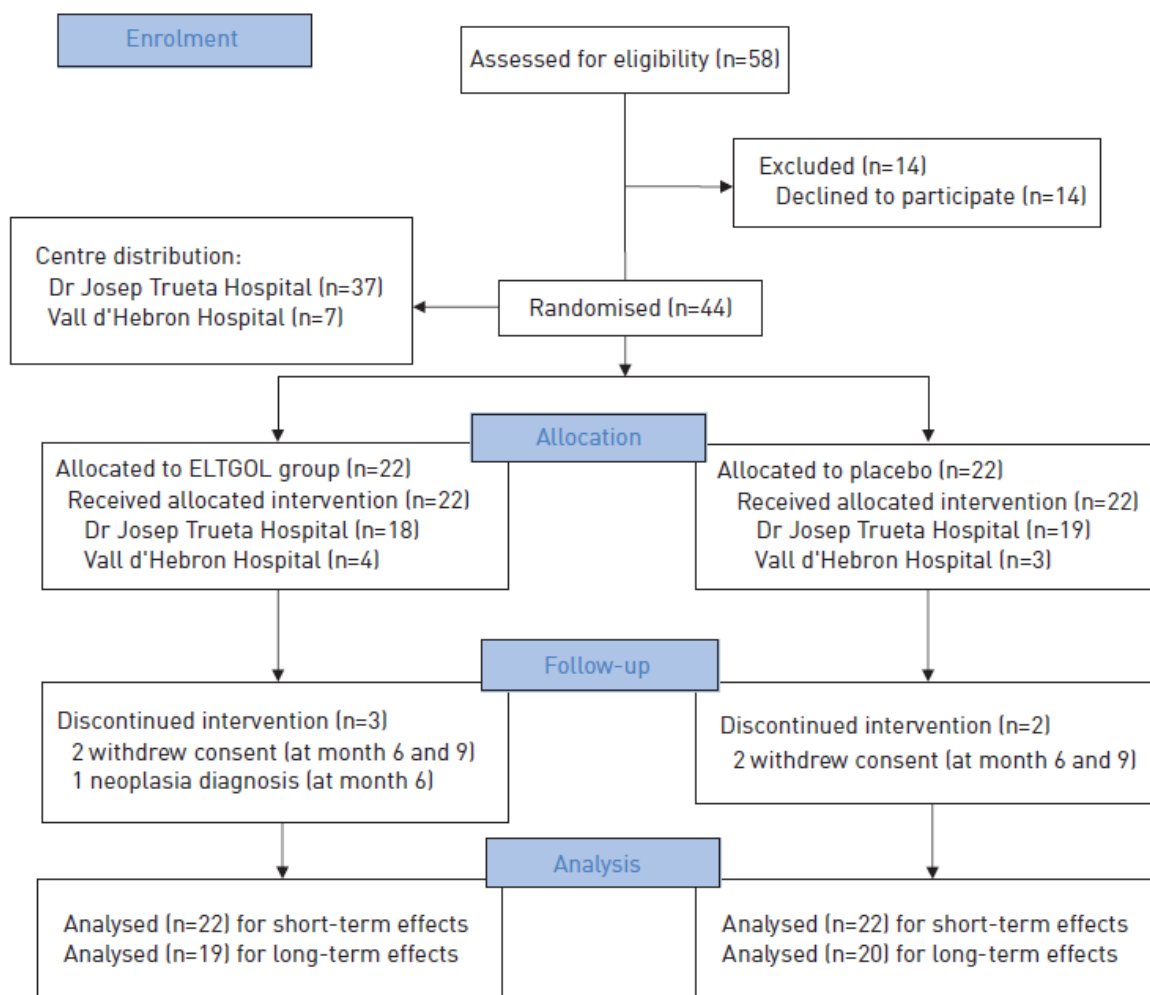


Hesham A. AbdelHalim ^{a,*}, Heba H. AboElNaga ^b, Karim A. Fathy ^c



Long-term benefits of airway clearance in bronchiectasis: a randomised placebo-controlled trial

Gerard Muñoz^{1,2}, Javier de Gracia^{3,4,5}, Maria Buxó⁶, Antonio Alvarez^{3,4} and Montserrat Vendrell^{1,3}



Intervention (ELTGOL)

Slow and prolonged expirations with the glottis opened, from the functional residual capacity to the residual volume, in the lateral decubitus position with the affected lung in the inferolateral position.

자세 변경 가래 배출법

기관지확장증 치료
수동적 가래배출법

수동적 가래배출법



자세변경 가래배출법



Long-term benefits of airway clearance in bronchiectasis: a randomised placebo-controlled trial

Gerard Muñoz^{1,2}, Javier de Gracia^{3,4,5}, Maria Buxó⁶, Antonio Alvarez^{3,4} and Montserrat Vendrell^{1,3}

	Sputum volume mL		p-value
	ELTGOL group	Placebo group	
Baseline 24-h	20 (15–40)	15 (15–20)	0.061
Visit 2 overall 24-h	40 (23.75–60)	12.5 (0–20)	<0.001
During intervention	12.27±11.93	0	
24 h later	30 (20–45)	12.5 (0–20)	<0.001
Difference between visit 2 and baseline⁺	17.5 (10–26.25)	–5 (–11.25–0)	<0.001
Month 12 overall 24-h	35 (30–50)	15 (10–20)	<0.001
During intervention	10.83±5.21	0	
24 h later	25 (20–40)	15 (10–20)	0.001
Difference between month 12 and baseline[#]	10 (–5–25)	0 (–10–3.75)	0.015
Difference between month 12 and visit 2[¶]	–5 (–30–5)	5 (5–10)	0.019

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	ELTGOL			Placebo			p-value [#]
	Baseline	Month 12	Between-group difference	Baseline	Month 12	Between-group	
SGRQ total score			-6.8 (-15.1-1.5)⁺			11.4 (6.9-15.9)⁺	
LCQ total score	40.2±13.7	33.7±15.7	1.96 (0.2-3.8)⁺	35.0±9.9	47.6±12.8	-2 (-2.8- -1.2)⁺	<0.001
Exacerbations	2 (1-3.25)	1 (0-2)	-0.8 (-1.5- -0.1) [¶]	1 (0.75-2.25)	2 (1-3)	0.35 (-0.5-0.35) [¶]	0.042
FEV ₁ % predicted	58.1±22.9	57.9±25	-0.4 (-3.5-2.8) ⁺	64.6±21.1	61.3±21	-2.5 (-4.7- -0.2) ⁺	0.262
FEV ₁ L	1.6±0.8	1.6±0.8	-0.004 (-0.1-0.03) ⁺	1.5±0.4	1.5±0.4	-0.1 (-0.2-0.004) ⁺	0.407
mMRC	1 (0-1.25)	1 (0-1)	0 (-0.5-0) [¶]	1 (1-1.25)	1 (1-2)	0.5 (0-0.5) [¶]	0.127
6MWT m	417.8±67	423.5±84.9	2.3 (-16.7-21.2) ⁺	382.9±76.9	377.8±57.3	-2.6 (-29.3-24.1) ⁺	0.746
ESR mm	22.3±26.5	17.1±17.5	9 (7-23) ⁺	25.5±22.3	23.9±17.6	24 (7.3-34.5) ⁺	0.863
Leukocytes ×10 ³ μL ⁻¹	6.9±2	7.5±2	0.03 (-0.8-0.9) ⁺	7.5±2.2	7.7±2.7	0.6 (-0.2-1.3) ⁺	0.641
Neutrophils %	59.7±8.7	60±8.9	-1.6 (-6.6-3.3) ⁺	58.5±8.4	57.9±12.1	-1.4 (-6-3.2) ⁺	0.945
CRP mg·dL ⁻¹	0.7±0.9	1.7±2.7	0.7 (-0.7-2.2) ⁺	0.6±0.5	0.7±0.6	0.06 (-0.3-0.4) ⁺	0.619
Fibrinogen mg·dL ⁻¹	425.5±69	468.6±1000.5	43.9 (-31.3-119) ⁺	449.6±930.5	492.6±125.2	59.3 (-13.8-132.3) ⁺	0.756

Long-term benefits of airway clearance in bronchiectasis: a randomised placebo-controlled trial

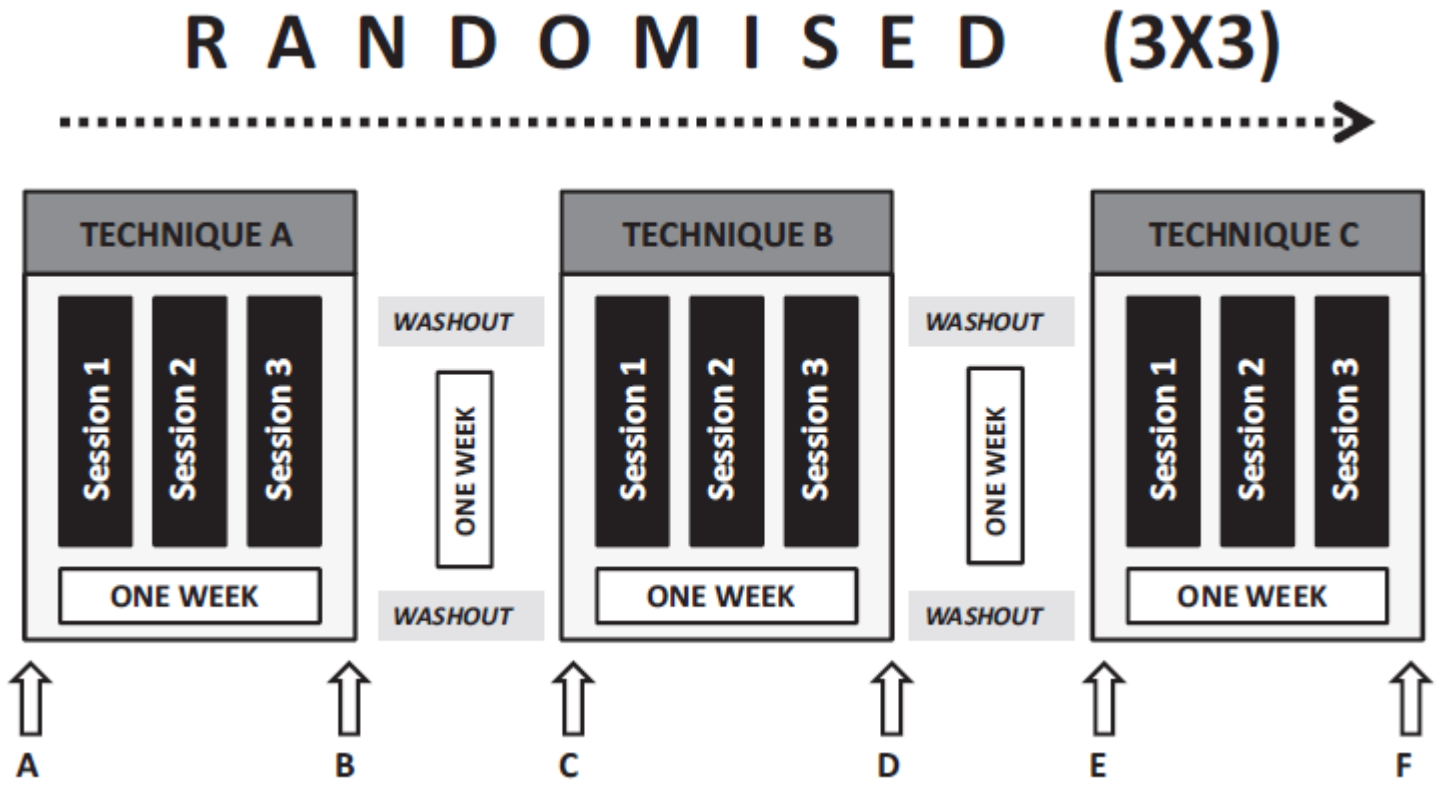
Gerard Muñoz^{1,2}, Javier de Gracia^{3,4,5}, Maria Buxó⁶, Antonio Alvarez^{3,4} and Montserrat Vendrell^{1,3}

	Month 6 – visit 2				Month 12 – month 6			
	ELTGOL [#]	Placebo [#]	Mean difference (95% CI) [¶]	p-value ⁺	ELTGOL [#]	Placebo [#]	Mean difference (95% CI) [¶]	p-value ⁺
SGRQ total	-7.7±8.6	7.8±10.5	-15.51 (-22.10; -8.92)	<0.001	0.8±14	1.6±9.6	-0.78 (-9.07; 7.50)	0.848
SGRQ symptoms	-8.3±21.4	6.2±10.3	-14.50 (-26.11; -2.89)	0.017	-1±17.9	0.7±15.8	-1.65 (-9.04; 9.54)	0.775
SGRQ activity	-4.5±12	8.6±14.5	-13.06 (-22.17; -3.95)	0.006	0.4±14.4	2.5±15.8	-2.10 (-12.46; 8.26)	0.683
SGRQ impact	-9.3±10.5	7.9±13.2	-17.23 (-25.40; -9.06)	<0.001	1.6±16.4	1.4±9.5	0.25 (-9.04; 9.54)	0.957
LCQ total	1.9±4	-1.6±2.6	3.53 (1.20; 5.86)	0.004	0.002±3.5	-0.5±2.2	0.48 (-1.54; 2.50)	0.633

Short-term effects of three slow expiratory airway clearance techniques in patients with bronchiectasis: a randomised crossover trial

B. Herrero-Cortina^{a,b}, J. Vilaró^c, D. Martí^{a,*}, A. Torres^{a,d},
M. San Miguel-Pagola^b, V. Alcaraz^a, E. Polverino^a

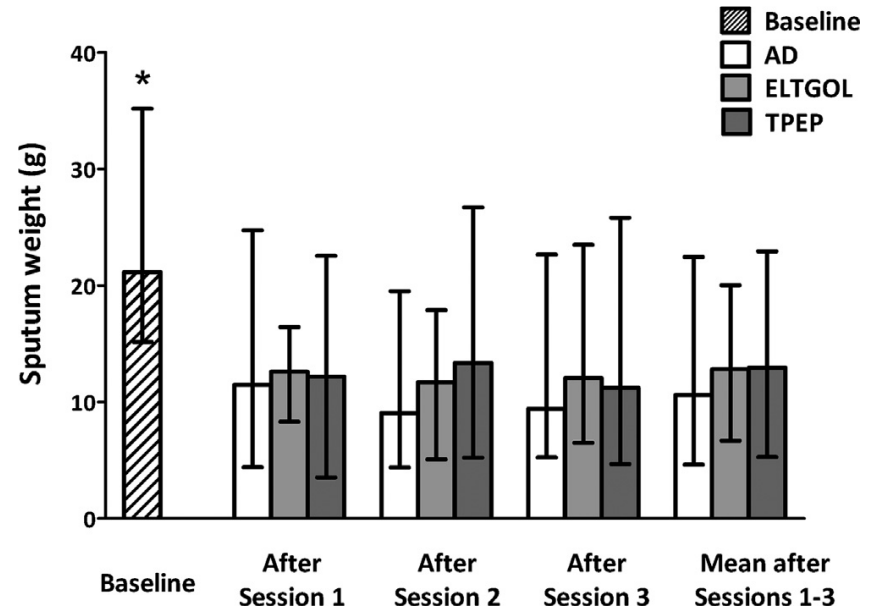
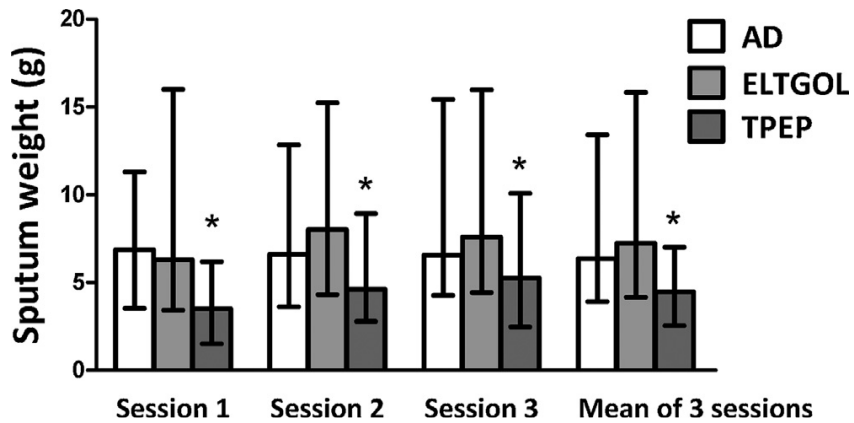
^a Pneumology Department, Clinic Institute of Thorax, Hospital Clinic of Barcelona, Institut d'Investigacions Biomèdiques August Pi I Sunyer – Centro de Investigación Biomédica En Red-Enfermedades Respiratorias, Barcelona, Spain
^b Health Sciences Faculty, San Jorge University, Zaragoza, Spain
^c Health Sciences Faculty Blanquerna, Physiotherapy Research Group, Ramon Llull University, Barcelona, Spain
^d University of Barcelona, Barcelona, Spain



Short-term effects of three slow expiratory airway clearance techniques in patients with bronchiectasis: a randomised crossover trial

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 M. San Miguel-Pagola^b, V. Alcaraz^a, E. Polverino^a

^a Pneumology Department, Clinic Institute of Thorax, Hospital Clinic of Barcelona, Institut d'Investigacions Biomèdiques August Pi I Sunyer – Centro de Investigación Biomédica En Red-Enfermedades Respiratorias, Barcelona, Spain
^b Health Sciences Faculty, San Jorge University, Zaragoza, Spain
^c Health Sciences Faculty Blanquerna, Physiotherapy Research Group, Ramon Llull University, Barcelona, Spain
^d University of Barcelona, Barcelona, Spain



AD, auto-genic drainage

ELTGOL, slow expiration with glottis opened in lateralposture;

TPEP, temporary positive expiratory pressure.

Short-term effects of three slow expiratory airway clearance techniques in patients with bronchiectasis: a randomised crossover trial

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^a *Pneumology Department, Clinic Institute of Thorax, Hospital Clinic of Barcelona, Institut d'Investigacions Biomèdiques August Pi I Sunyer – Centro de Investigación Biomédica En Red-Enfermedades Respiratorias, Barcelona, Spain*
^b *Health Sciences Faculty, San Jorge University, Zaragoza, Spain*
^c *Health Sciences Faculty Blanquerna, Physiotherapy Research Group, Ramon Llull University, Barcelona, Spain*
^d *University of Barcelona, Barcelona, Spain*

Table 2
Change in Leicester Cough Questionnaire score after 1 week of treatment for each slow expiratory airway clearance technique.

	Total score			Physical score			Psychological score			Social score		
	Median difference	95% CI	P-value	Median difference	95% CI	P-value	Median difference	95%	P-value	Median difference	95% CI	P-value
AD	0.5	0.1 to 0.5	0.01	0.1	0.0 to 0.3	0.1	0.1	-0.1 to 0.4	0.1	0.0	-0.1 to 0.5	0.2
ELTGOL	0.9	0.5 to 2.1	0.001	0.4	-0.1 to 0.6	0.006	0.3	0.1 to 0.6	0.001	0.2	0.1 to 0.9	0.001
TPEP	0.4	0.1 to 1.2	0.04	0.1	-0.1 to 0.3	0.3	0.1	0.0 to 0.5	0.06	0.1	0.0 to 0.5	0.02

AD, autogenic drainage; ELTGOL, slow expiration with glottis opened in lateral posture; TPEP, temporary positive expiratory pressure; CI, confidence interval. Wilcoxon's tests were performed. $P < 0.05$ was considered to indicate statistical significance.

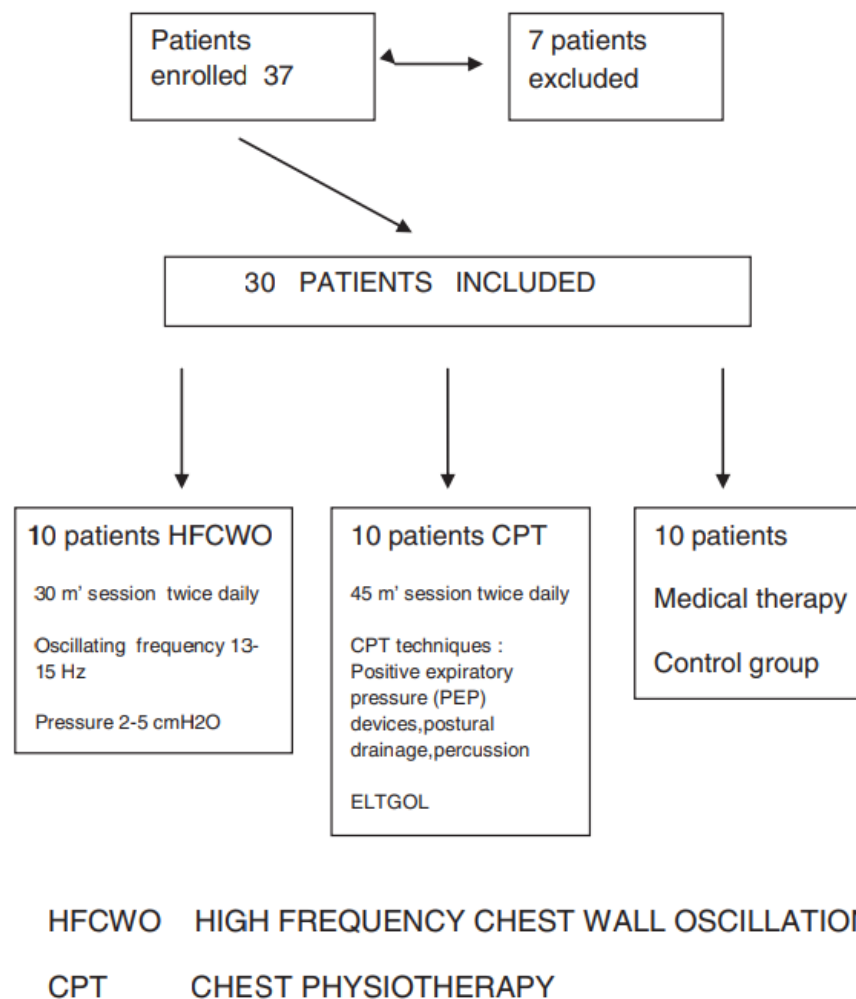
AD, auto-genic drainage

ELTGOL, slow expiration with glottis opened in lateral posture;

TPEP, temporary positive expiratory pressure.

Effectiveness of treatment with high-frequency chest wall oscillation in patients with bronchiectasis

Antonello Nicolini^{1*}, Federica Cardini¹, Norma Landucci¹, Sergio Lanata², Maura Ferrari-Bravo³ and Cornelius Barlascini⁴



Physiotherapy

가래배출법	능동적	<ul style="list-style-type: none">▫ 기침법▫ 허핑▫ 가슴팽창운동▫ 능동주기 호흡법 (ACBT)▫ 자가 배출법▫ 보조기침법과 덧대기 기침법
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	도구	<ul style="list-style-type: none">▫ 에어로비카 · 플루터 · 아카펠라▫ BPEP▫ HFCO · IPPB

보조 기침법

기관지확장증 치료
능동적 가래배출법



Evidence A in BTS guideline 2019

- Teach individuals with bronchiectasis to perform airway clearance. (D)
- Offer active cycle of breathing techniques or oscillating positive expiratory pressure to individuals with bronchiectasis. (D)
- Consider gravity assisted positioning (where not contraindicated) to enhance the effectiveness of an airway clearance technique. (D)
- **Do not** routinely **use recombinant human DNase** in adults with bronchiectasis.

Non-pharmacologic Treatment of Bronchiectasis

- Active cyclic breathing
- Oscillating positive expiratory devices
- Postural drainage
- Forced expiration technique
- Pulmonary rehabilitation
- Inspiratory muscle training

Evidence B in BTS guideline 2010

- **Pulmonary rehabilitation** should be offered to individuals who have breathlessness affecting their activities of daily living.
- **Inspiratory muscle training** can be used in conjunction with conventional pulmonary rehabilitation to enhance the maintenance of the training effect.

Evidence B in BTS guideline 2019

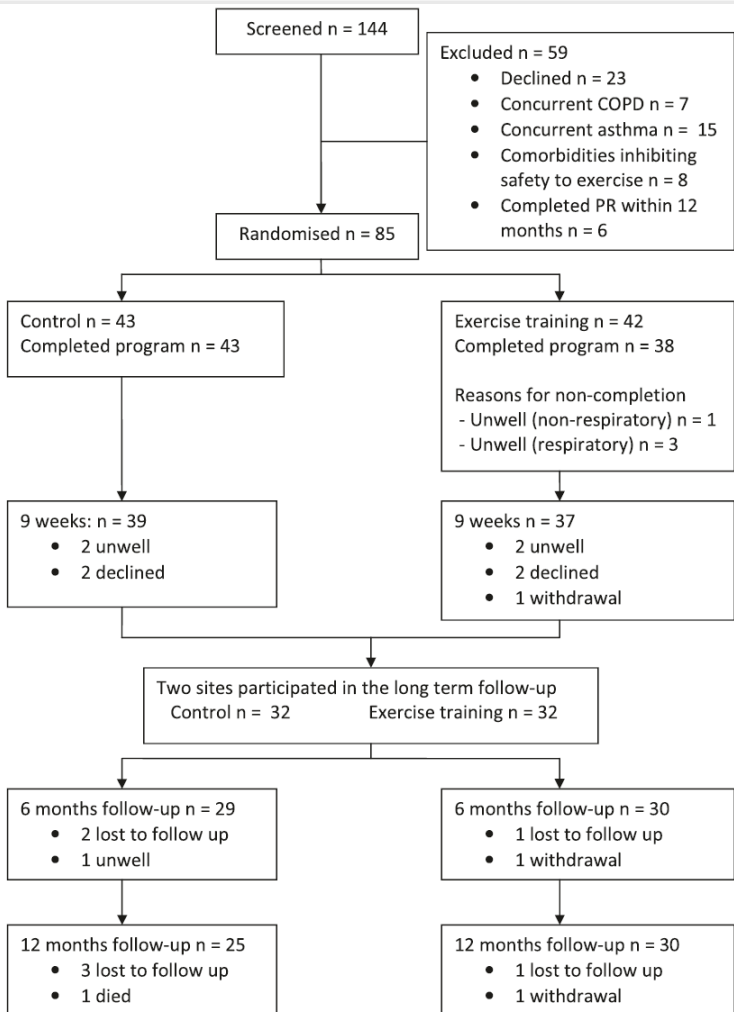
- Offer **pulmonary rehabilitation** to individuals who are functionally limited by shortness of breath (MMRC ≥ 1).
- Consider the use of **inspiratory muscle training** in conjunction with conventional pulmonary rehabilitation to enhance the maintenance of the training effect.

Research Recommendations in BTS 2019

- The role of education, self management plans and who delivers the pulmonary rehabilitation needs to be explored.
- The role of pulmonary rehabilitation after exacerbations requiring hospital admission needs to be explored.
- The incidence of cross-infection of respiratory pathogens in the group exercise setting should be investigated in the bronchiectasis population.

The short and long term effects of exercise training in non-cystic fibrosis bronchiectasis – a randomised controlled trial

Annemarie L. Lee^{1,2,3*}, Catherine J Hill^{2,4}, Nola Cecins^{5,6}, Sue Jenkins^{5,6,7}, Christine F McDonald^{2,4}, Angela T Burge¹, Linda Rautela^{2,4}, Robert G Stirling^{1,8}, Philip J Thompson^{5,6,7} and Anne E Holland^{1,2,9}



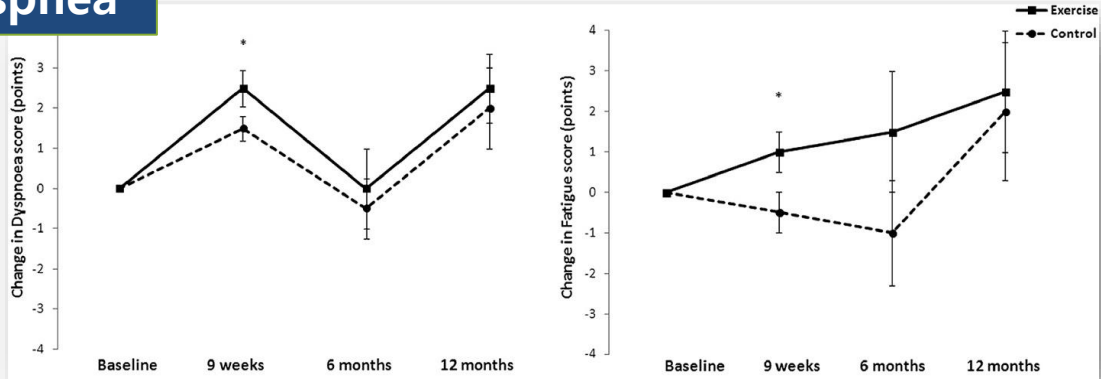
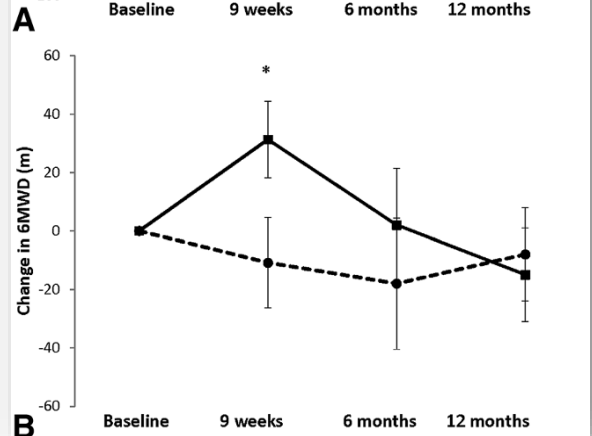
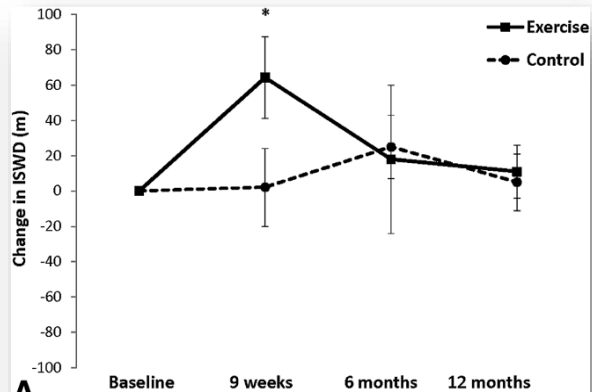
Flow of participants.

- 8 weeks of supervised exercise training
- Twice-weekly
- Treadmill or land-based walking
- Initial intensity set to 75% of the maximal speed

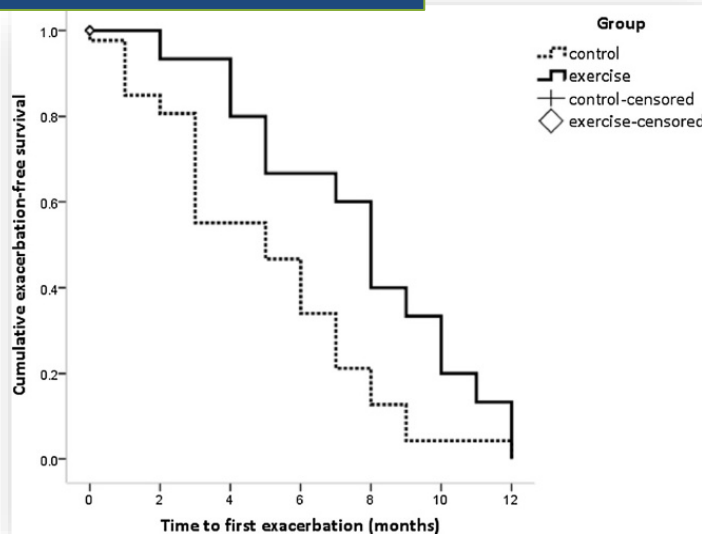
Pulmonary rehab improved exercise capacity and dyspnea in BE

Dyspnea

ISWD



Exacerbation free



Exercise training and inspiratory muscle training in patients with bronchiectasis

C Newall, R A Stockley, S L Hill

Thorax 2005;60:943–948. doi: 10.1136/thx.2004.028928

Table 1 Mean (SD) baseline physiological characteristics of patients who completed the study (n = 32)

Parameter	PR-SHAM	PR-IMT	Control
M/F	4/7	2/10	0/9
Smoking status	4 EX/7 NS	2 EX/10 NS	2 EX/7 NS
Age (years)	63.1 (3.5)	57.3 (2.4)	62.9 (3.9)
FEV ₁ (l)	1.44 (0.77)	1.23 (0.74)	1.49 (0.61)
FEV ₁ (% predicted)	64	54	69
FEV ₁ /VC%	57.0 (0.20)	54.1 (0.20)	66.2 (0.09)
RV (l)	2.14 (0.97)	2.51 (0.95)	1.83 (0.87)
RV (% predicted)	111	133	99
TLC (l)	5.04 (1.68)	4.92 (1.10)	4.26 (1.37)
TLC (% predicted)	97	98	88
TlCO (mmol/min/kPa)	6.32 (2.07)	5.88 (1.61)	5.92 (2.13)
TlCO (% predicted)	79	76	76
Kco (mmol/min/kPa/l)	1.65 (0.44)	1.63 (0.29)	1.66 (0.35)
Kco (% predicted)	86	82	82
P _{imax} (cm H ₂ O)	66.4 (22.0)	78.0 (17.7)	77.2 (24.7)
P _{imax} (% predicted)	87	98	103
P _{Emax} (cm H ₂ O)	97.3 (37.5)	75.8 (28.5)	87.4 (22.9)
P _{Emax} (% predicted)	66	53	63
Peak V _{O₂} (ml/min/kg)	15.6 (6.2)	19.8 (8.3)	19.0 (5.0)
Peak V _{O₂} (% predicted)	77	88	92

EX, ex-smoker; NS, never smoked; FEV₁, forced expiratory volume in 1 second; VC, vital capacity; TLC, total lung capacity; RV, residual volume; TlCO, lung carbon monoxide transfer factor; Kco, carbon monoxide transfer coefficient; P_{imax}, P_{Emax}, maximum inspiratory and expiratory pressures; V_{O₂}, oxygen consumption.

- 8 weeks of supervised exercise training
- Three times weekly
- IMT was performed at home for 15 minutes
- 80% of the peak heart rate

Exercise capacity maintained with pulmonary rehab

PR-IMT

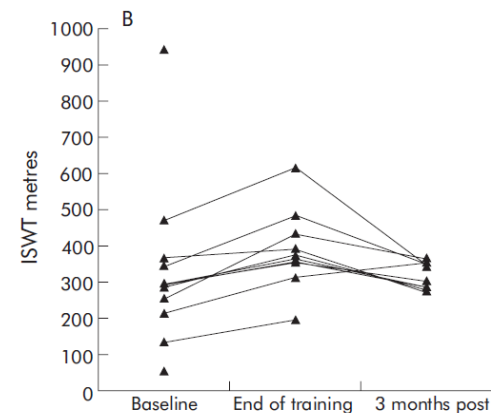
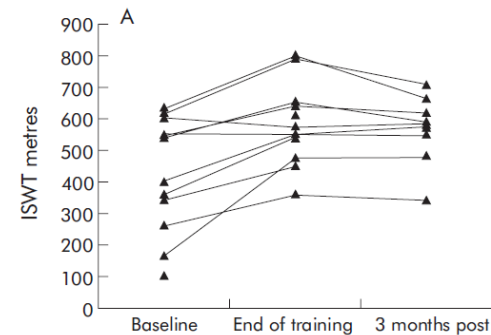
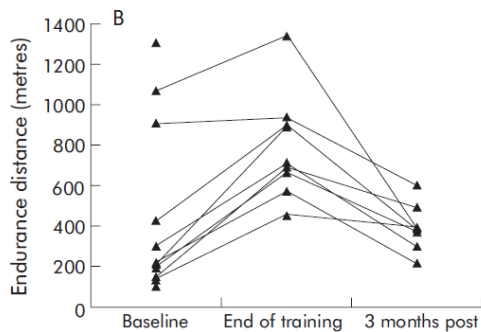
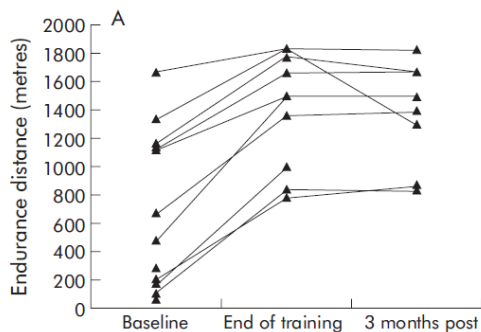
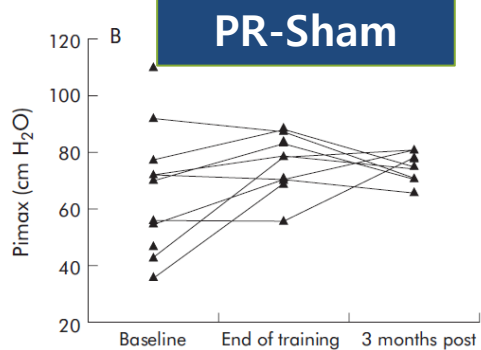
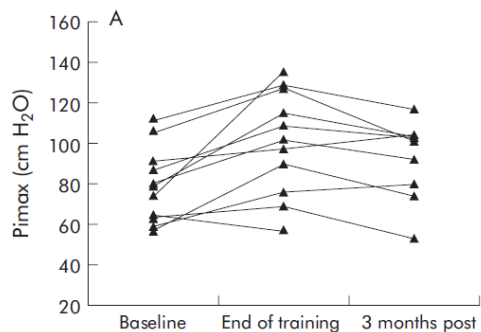


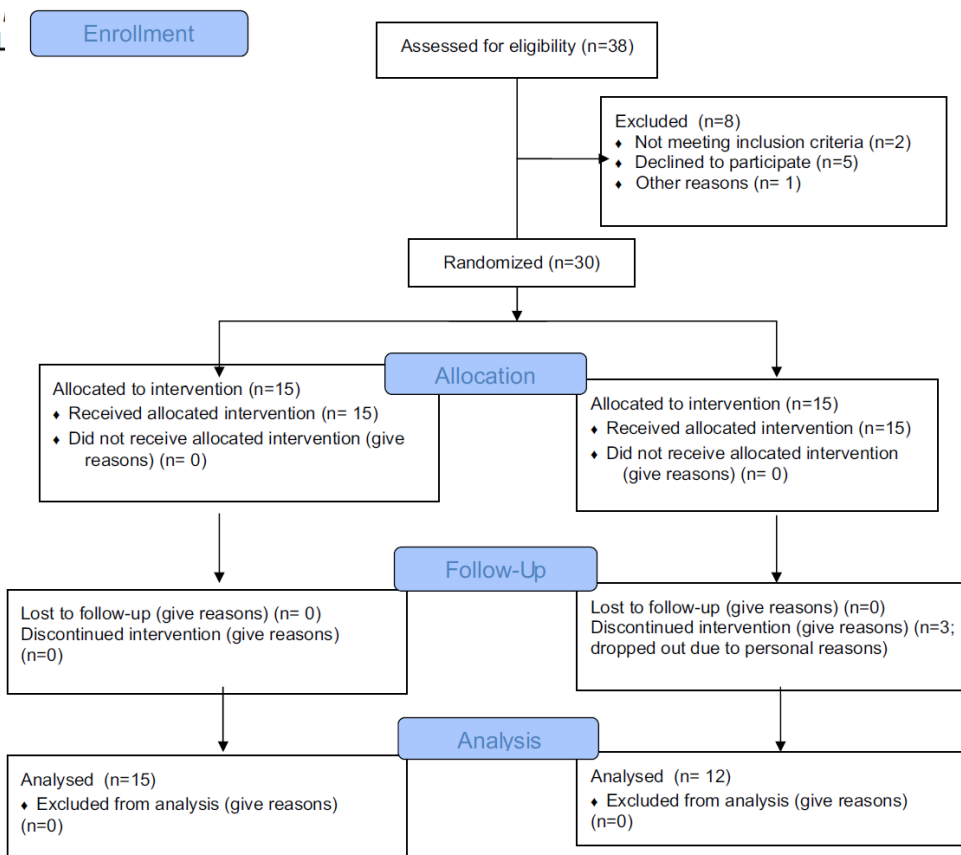
Figure 1 Individual changes in Pimax during training in (A) the PR-IMT group and (B) the PR-SHAM group. Individual values for Pimax during training are shown. The lines join data for individual patients.

Figure 2 Individual changes in endurance capacity during training in (A) the PR-IMT group and (B) the PR-SHAM group. Individual data and changes in endurance capacity are shown before and after 8 weeks training.



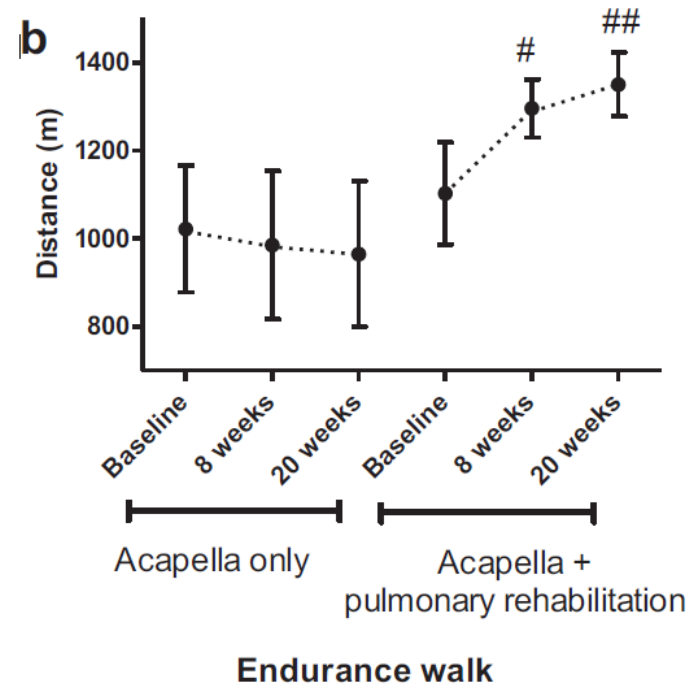
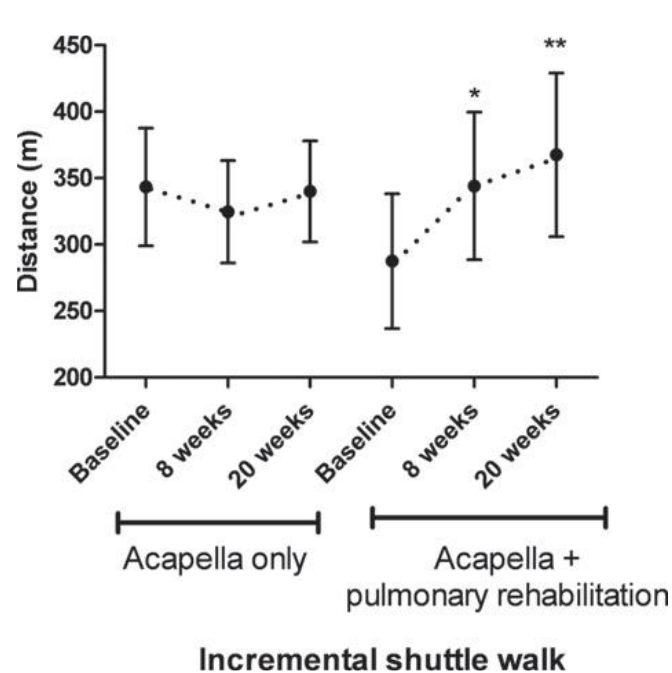
A pilot study of pulmonary rehabilitation and chest physiotherapy versus chest physiotherapy alone in bronchiectasis

P. J.L.



- 8 weeks
- 1:1 randomization
- Physiotherapy \pm PR

Exercise capacity improved with pulmonary rehab



Exercise capacity and QOL improved with pulmonary rehab

Table 2 *p* values represent difference between groups at time points indicated.

	Group receiving acapella (<i>n</i> = 15)			Group receiving acapella + pulmonary rehabilitation (<i>n</i> = 12)		
	Baseline	8 weeks	20 weeks	Baseline	8 weeks	20 weeks
ISWT (m)	343.3 (44.4)	338.7 (42.2)	343.3 (39.8)	287.5 (50.6)	344.2 (115.5)*	367.5 (61.5)**
EWT (m)	1021.4(144.4)	985 (168.6)	964.3 (165.7)	1102.5 (116.2)	1295.8 (65.8) #	1350 (72.6) ##

p* = 0.03; *p* = 0.04; #*p* = 0.01; ##*p* = 0.003.

Table 3 *p* values represent difference between groups at time points indicated.

	Group receiving acapella (<i>n</i> = 15)			Group receiving acapella + pulmonary rehabilitation (<i>n</i> = 12)		
	Baseline	8weeks	20weeks	Baseline	8weeks	20 weeks
SGRQ	40.6 (3.9)	39.2 (4.5)	45.2 (4.5)	38.6 (6.4)	30.6 (6.6)*	34.6 (7.7)*
LCQ	14.4 (1.5)	14.6 (1.4)	13.62 (1.4)	12.3 (2.3)	14.9 (2.3)*	16.7 (1.8)*

**p* < 0.001.



REVIEW ARTICLE (META-ANALYSIS)

Pulmonary Rehabilitation in Individuals With Non-Cystic Fibrosis Bronchiectasis: A Systematic Review

Annemarie L. Lee, PhD,^{a,b,c} Catherine J. Hill, PhD,^{c,d} Christine F. McDonald, PhD,^{c,e}
Anne E. Holland, PhD^{c,f,g}

From the ^aWest Park Healthcare Centre, Toronto, Ontario, Can-
Canada; ^bInstitute for Breathing and Sleep, and Departments
Heidelberg, Victoria, Australia; ^cDepartment of Physiotherapy,
of Rehabilitation, Nutrition and Sport, La Trobe University, M

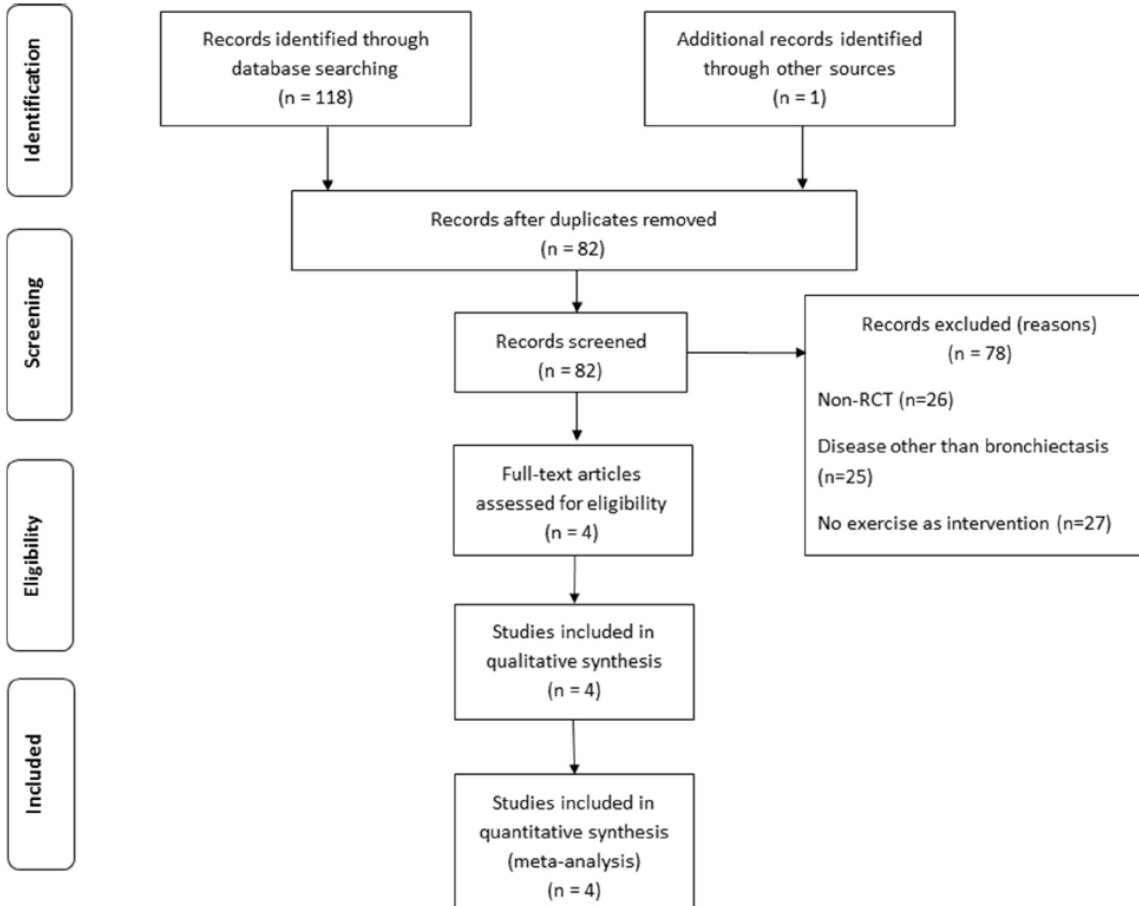


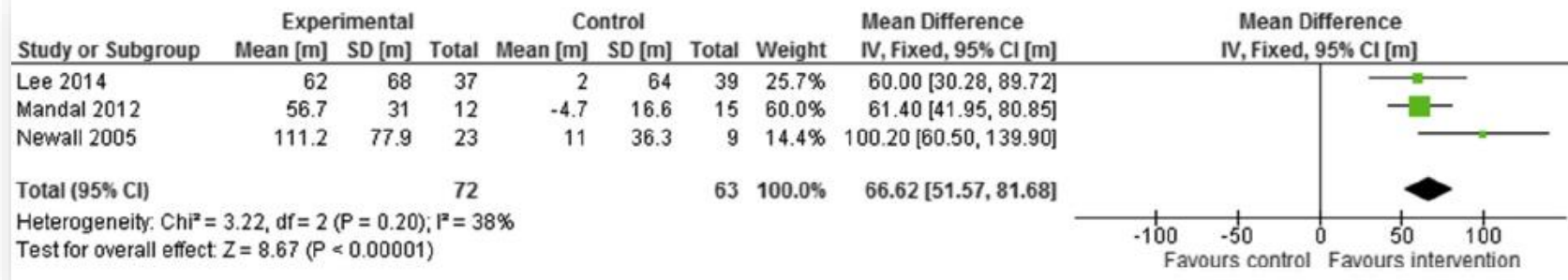
Fig 1 Study flow from identification to inclusion. Abbreviation: RCT, randomized controlled trial.

Exercise capacity and QOL improved with pulmonary rehab in meta-analysis

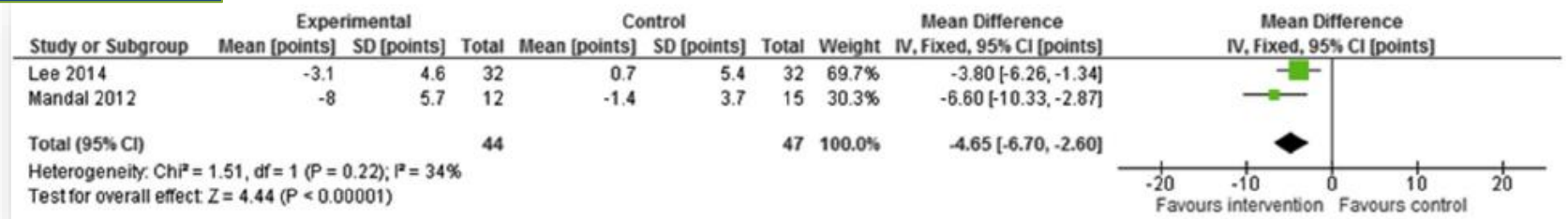
Table 3 Quality assessment (Cochrane risk-of-bias tool)

Study	Randomization Sequence	Allocation Concealment	Blinding			Incomplete Data	Selective Reporting	Other Biases
			Participants	Therapists	Outcome Assessors			
Newall, ¹⁶ 2005	Low	Unclear	Unclear	Unclear	Unclear	Low	Low	None
Mandal, ¹⁷ 2012	Low	Unclear	Unclear	Unclear	Unclear	Low	Low	None
Lee, ¹⁸ 2014	Low	Low	High	Unclear	Low	Low	Low	None
Greening, ²⁶ 2014	Unclear	Unclear	High	Unclear	Low	Low	Low	None

ISWD

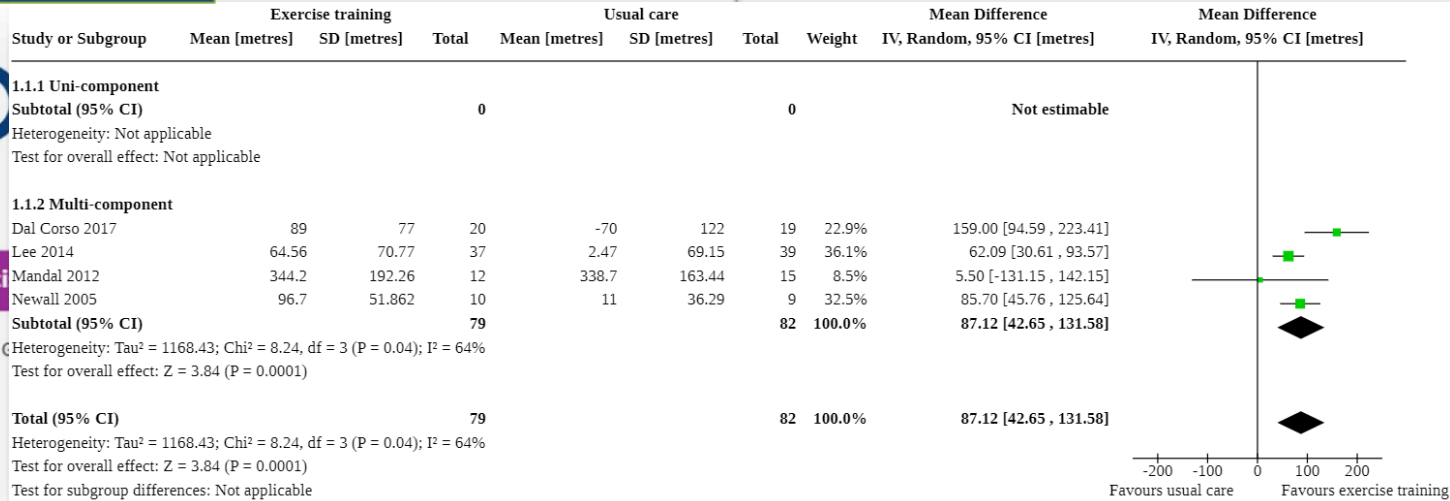


SGRQ



Exercise capacity and QOL improved with pulmonary rehab in meta-analysis

ISWD

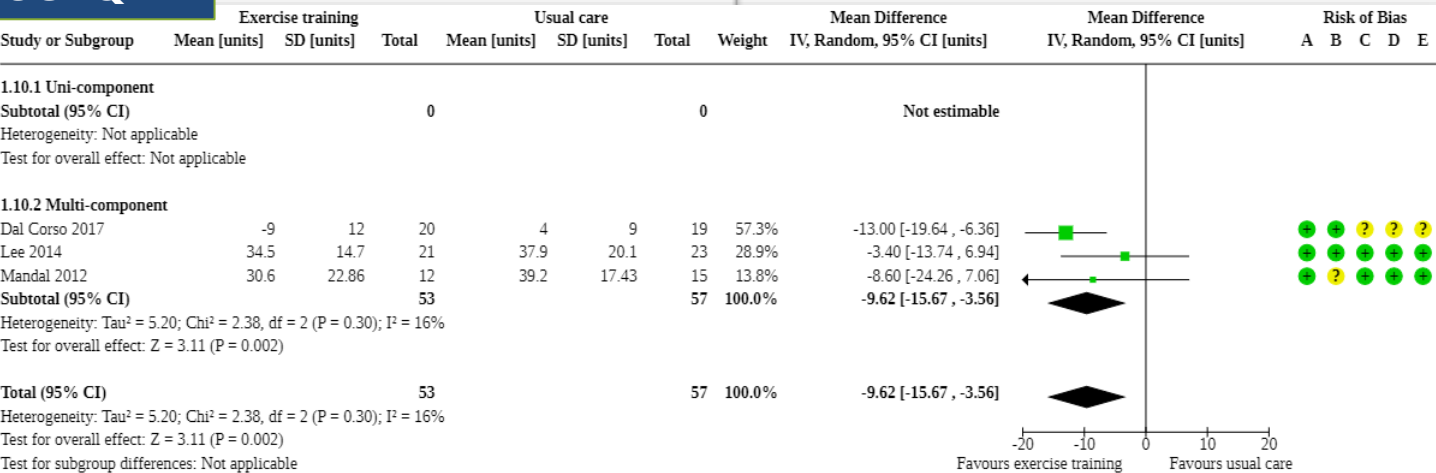


Exerci

Lee AL, C

SGRQ

Lee AL, C
Exercise
Cochran
DOI: 10.



호흡재활 보험고시 (2016)

대상

만성폐쇄성폐질환

기관지확장증

천식

폐동맥고혈압

간질성폐질환

결핵성폐질환

폐암 및 폐암의 수술 전후

폐이식 폐용적감소수술

제한성폐질환

(전후척추측만증)

*호흡곤란, 일상생활 어려움

소요인력

의사 간호사 물리치료사

내용

호흡재활 전문의사

환자의 운동능력과 호흡곤란 정도,

기저질환 및 동반질환을 고려

개인별 맞춤 운동프로그램

유산소 운동 :

- 최대운동능력의 60% 이상의 강도

- 20~60분의 운동

- 주 3~5회시행

- 고강도 운동이 불가능한 환자에서는 저강도 운동
근력 운동 :

- 최대 근력의 60~70%의 강도

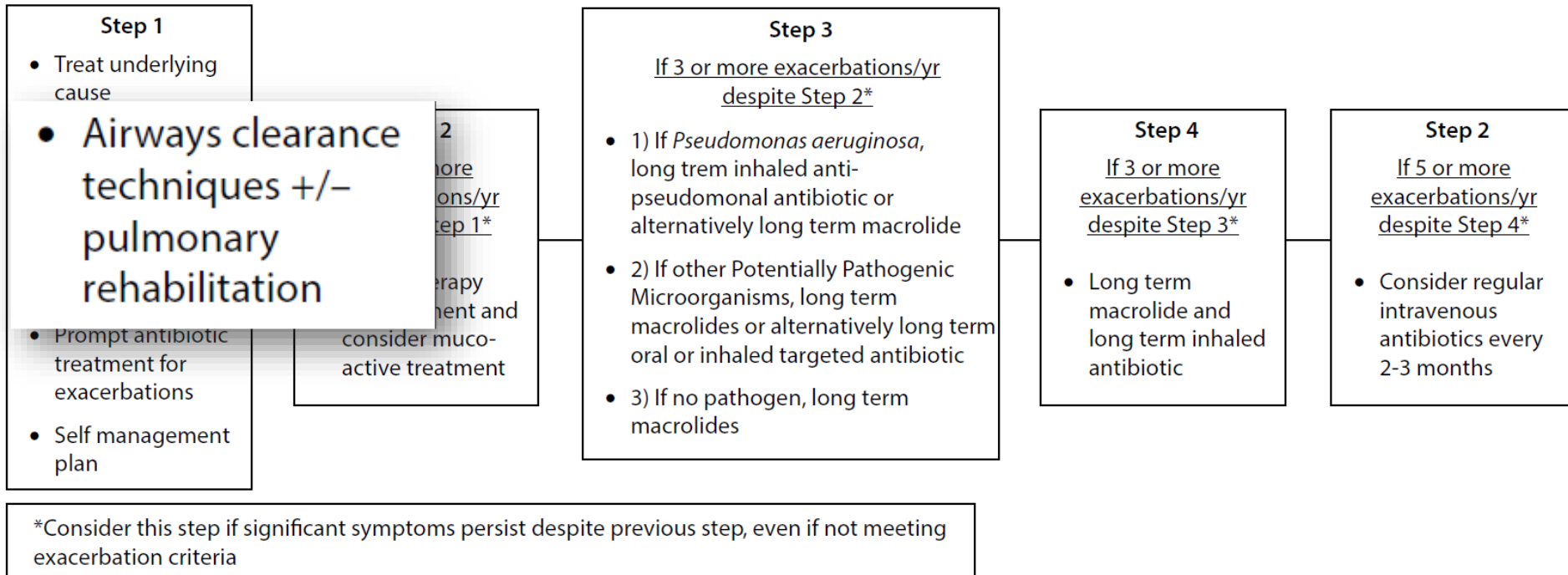
- 1세트 10회 이상

- 세트 2~3회 시행

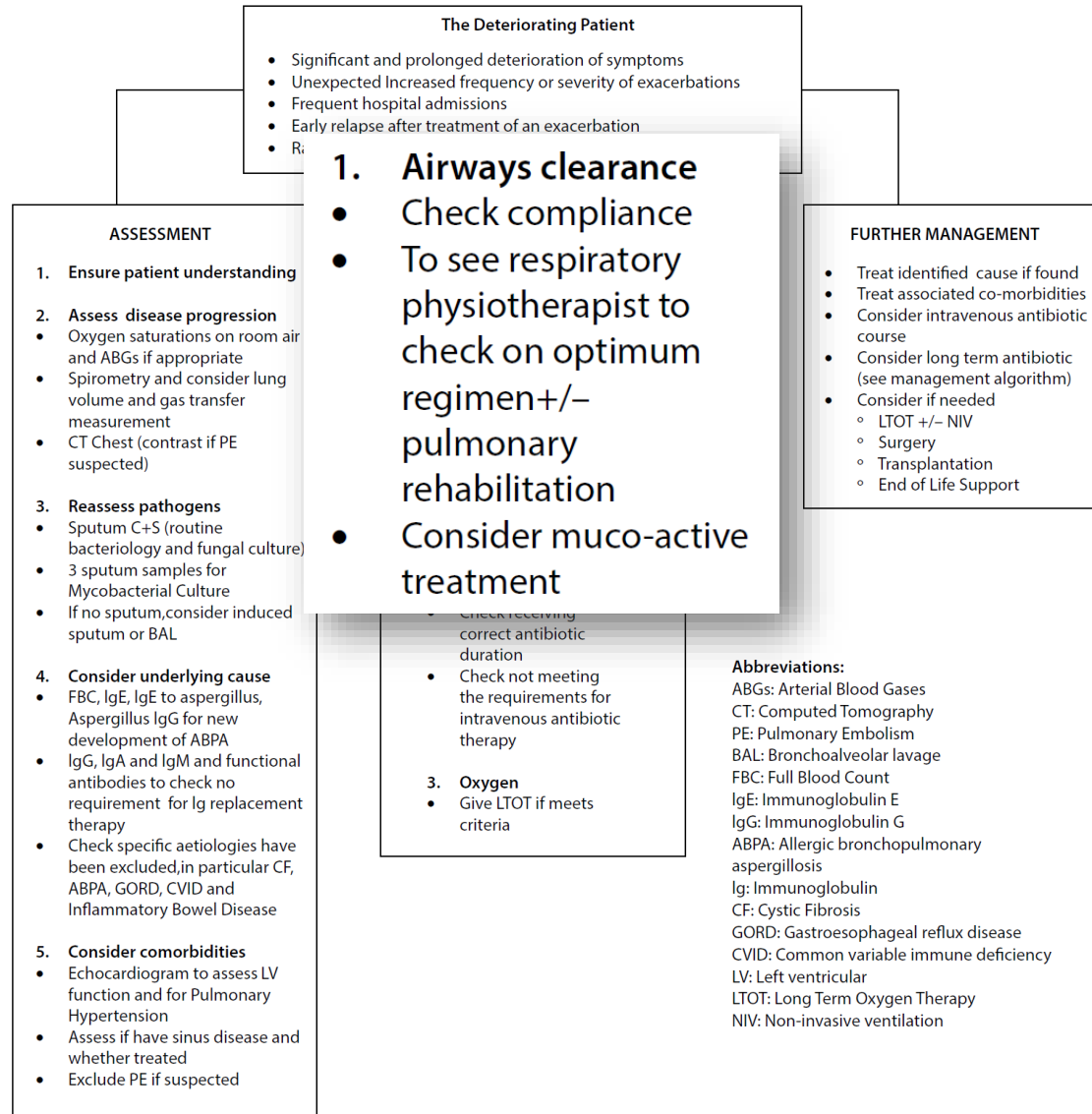
Non-pharmacologic Treatment of Bronchiectasis

- Active cyclic breathing
- Oscillating positive expiratory devices
- Postural drainage
- Forced expiration technique
- Pulmonary rehabilitation
- Inspiratory muscle training

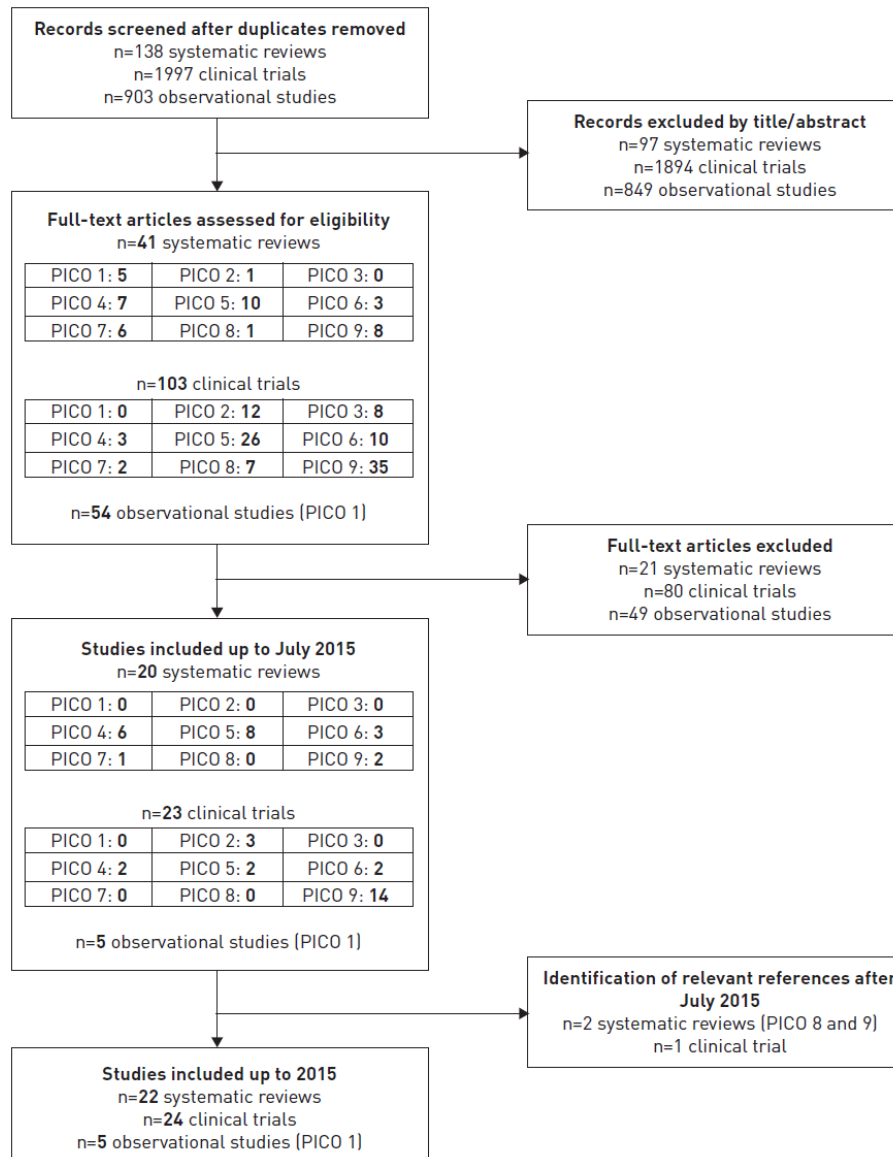
Stepwise management in BTS 2019



Management of deteriorating patients



ERS guidelines: Key Questions



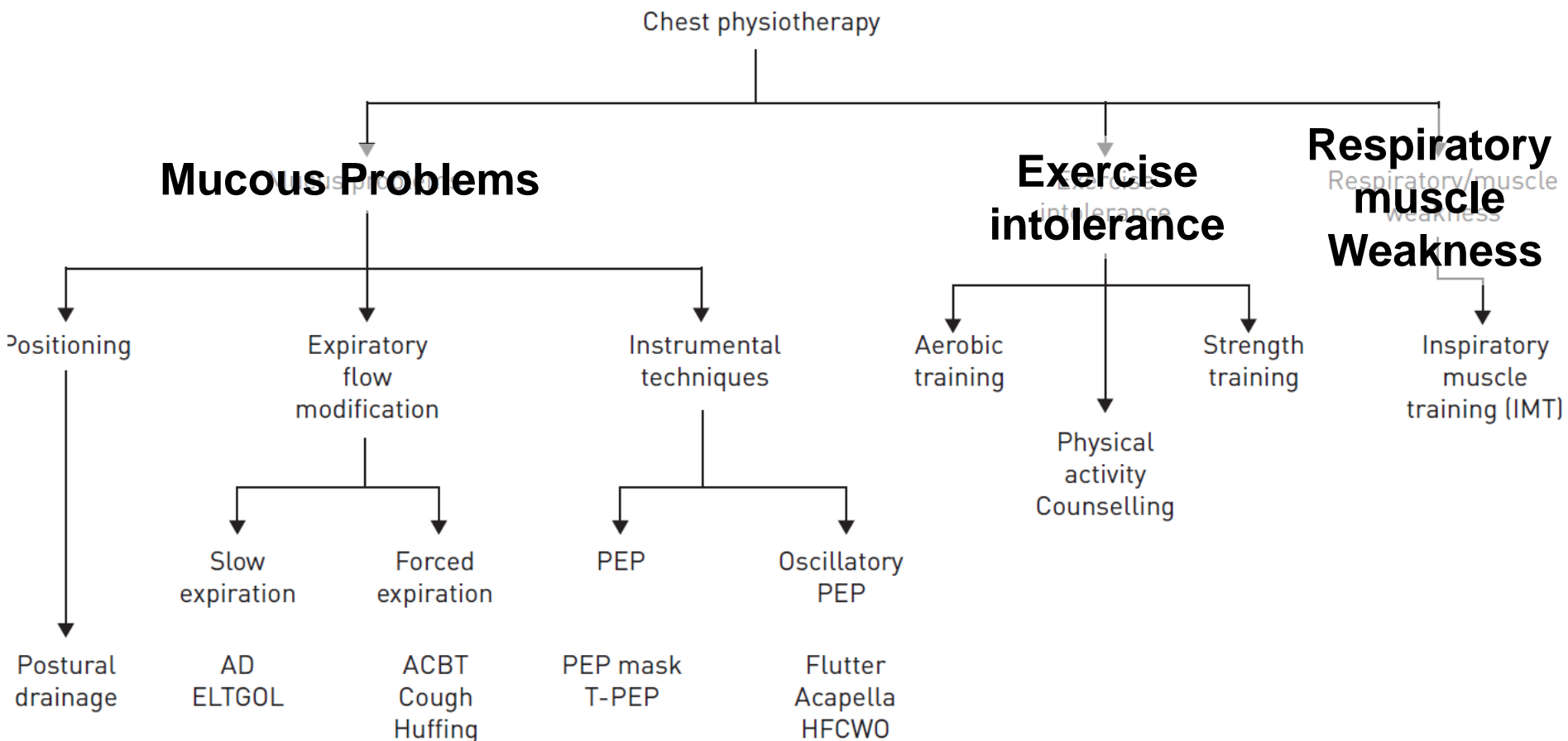
Key Questions 9:

Question 9: Is regular physiotherapy (airway clearance and/or pulmonary rehabilitation) more beneficial than control (no physiotherapy treatment) in adult bronchiectasis patients?

Answers to Key Questions 9

Patients with chronic productive cough or difficulty to expectorate sputum should be taught an airway clearance technique (ACT) by a trained respiratory physiotherapist to perform once or twice daily.

Chest physiotherapy interventions flow chart



Summary

- Decreased mucociliary clearance and associated vicious cycle are the main pathogenesis of BE
- Physiotherapy to reduce sputum burden should be one of major treatment in BE.
- Pulmonary rehab can improve not only exercise capacity but also quality of life.