



HAL
open science

Lower airway clinical outcome measures for use in primary ciliary dyskinesia research: a scoping review

Florian Gahleitner, James Thompson, Claire L Jackson, Jana F Hueppe, Laura Behan, Eleonora Dehlink, Myrofora Goutaki, Florian Halbeisen, Ana Paula L. Queiroz, Guillaume Thouvenin, et al.

► To cite this version:

Florian Gahleitner, James Thompson, Claire L Jackson, Jana F Hueppe, Laura Behan, et al.. Lower airway clinical outcome measures for use in primary ciliary dyskinesia research: a scoping review. ERJ Open Research, 2021, 7 (4), pp.00320-2021. 10.1183/23120541.00320-2021] . hal-03465132

HAL Id: hal-03465132

<https://hal.sorbonne-universite.fr/hal-03465132>

Submitted on 3 Dec 2021

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.



Lower airway clinical outcome measures for use in primary ciliary dyskinesia research: a scoping review

Florian Gahleitner^{1,2}, James Thompson^{1,3}, Claire L. Jackson^{1,3}, Jana F. Hueppe ^{1,3}, Laura Behan^{1,3}, Eleonora Dehlink⁴, Myrofora Goutaki ^{5,6}, Florian Halbeisen ^{5,6,7}, Ana Paula L. Queiroz³, Guillaume Thouvenin ⁸, Claudia E. Kuehni ^{5,6}, Philipp Latzin ^{5,6}, Jane S. Lucas ^{1,3} and Bruna Rubbo ^{1,3,9}

¹Primary Ciliary Dyskinesia Centre, NIHR Biomedical Research Centre, University Hospital Southampton NHS Foundation Trust, Southampton, UK. ²Paediatric Respiratory Medicine, Royal Hospital for Children and Young People, Edinburgh, UK. ³School of Clinical and Experimental Science, University of Southampton, Faculty of Medicine, Southampton, UK. ⁴Division of Pediatric Pulmonology, Allergy and Endocrinology, Dept of Pediatrics and Adolescent Medicine, Medical University of Vienna, Vienna, Austria. ⁵Institute of Social and Preventive Medicine, University of Bern, Bern, Switzerland. ⁶Paediatric Respiratory Medicine, Children's University Hospital of Bern, University of Bern, Bern, Switzerland. ⁷Basel Institute for Clinical Epidemiology and Biostatistics, Dept of Clinical Research, University Hospital Basel, University of Basel, Basel, Switzerland. ⁸AP-HP, Pneumologic Unit, Trousseau Hospital, Sorbonne Universités, INSERM, Centre de Recherche Saint-Antoine, Paris, France. ⁹School of Health Sciences, University of Southampton, Faculty of Environmental and Life Sciences, Southampton, UK.

Corresponding author: Jane S. Lucas (jlucas1@soton.ac.uk)



Shareable abstract (@ERSpublications)

Measurement and reporting of lower airway outcome measures in primary ciliary dyskinesia research are not standardised. Validated, disease-specific clinical outcomes are needed to monitor disease progression in future trials and prospective cohorts. <https://bit.ly/3yHERQm>

Cite this article as: Gahleitner F, Thompson J, Jackson CL, *et al.* Lower airway clinical outcome measures for use in primary ciliary dyskinesia research: a scoping review. *ERJ Open Res* 2021; 7: 00320-2021 [DOI: [10.1183/23120541.00320-2021](https://doi.org/10.1183/23120541.00320-2021)].

Copyright ©The authors 2021

This version is distributed under the terms of the Creative Commons Attribution Non-Commercial Licence 4.0. For commercial reproduction rights and permissions contact permissions@ersnet.org

This article has supplementary material available from openres.ersjournals.com

Received: 7 May 2021
Accepted: 31 July 2021

Abstract

Objectives Disease-specific, well-defined and validated clinical outcome measures are essential in designing research studies. Poorly defined outcome measures hamper pooling of data and comparisons between studies. We aimed to identify and describe pulmonary outcome measures that could be used for follow-up of patients with primary ciliary dyskinesia (PCD).

Methods We conducted a scoping review by systematically searching MEDLINE, Embase and the Cochrane Database of Systematic Reviews online databases for studies published from 1996 to 2020 that included ≥10 PCD adult and/or paediatric patients.

Results We included 102 studies (7289 patients). 83 studies reported on spirometry, 11 on body plethysmography, 15 on multiple-breath washout, 36 on high-resolution computed tomography (HRCT), 57 on microbiology and 17 on health-related quality of life. Measurement and reporting of outcomes varied considerably between studies (*e.g.* different scoring systems for chest HRCT scans). Additionally, definitions of outcome measures varied (*e.g.* definition of chronic colonisation by respiratory pathogen), impeding direct comparisons of results.

Conclusions This review highlights the need for standardisation of measurements and reporting of outcome measures to enable comparisons between studies. Defining a core set of clinical outcome measures is necessary to ensure reproducibility of results and for use in future trials and prospective cohorts.

Introduction

Primary ciliary dyskinesia (PCD) is a rare, genetic, multisystem disease that affects motile cilia lining the upper and lower airways, and the eustachian and fallopian tubes [1, 2]. Symptoms start in early infancy, with progressive suppurative lung disease invariably leading to bronchiectasis [3, 4]. Current management of patients with PCD broadly follows that used for patients with cystic fibrosis (CF) and bronchiectasis (formally non-CF bronchiectasis) [5–8]. Therefore, studies have adopted similar outcome measures to



monitor the natural history and disease progression in PCD, even though PCD has a unique pathophysiology and disease pattern [9].

There is no minimum core set of disease-specific outcome measures in PCD research. This is particularly problematic because the choice of outcome measures informs the selection of data sources from which study information can be collected; the appropriateness, frequency and length of follow-up measurements; and the required number of patients. Appropriate sample size relies on the expected frequency and natural variability of outcomes, and on the effect of interest (or the minimal clinically important difference) [10]. The quality of the knowledge generated by research strongly relies on the selection of appropriate outcomes.

Well-defined and validated disease-specific outcome measures are the most efficient and accurate way to assess new therapies and management options. While true for all diseases, this is particularly poignant for rare diseases, where the number of patients available is limited [11]. An outcome measure that is valid for another disease might not be appropriate to measure the effect of interest, or sensitive enough to detect a subtle effect. For example, spirometry is routinely used to monitor disease progression, but is thought to be an insensitive surrogate marker for early lung disease in CF and, more recently, in PCD [12–15].

The aim of this scoping review was to systematically identify and describe the evidence in this area. In addition, we aimed to highlight the most commonly used pulmonary and related outcomes and to examine the consistency of definitions across studies and the variations on the use and reporting of clinical outcome measures in the PCD literature.

Methodology

Search strategy

We followed the *a priori* scoping review protocol, which is available from the authors upon request. A pilot search included only terms related to the disease (items 1–4 of search terms, supplementary box 1) and one reviewer (BR) scanned the first 1000 abstracts to identify key terms that could be used to build the full search strategy, designed for use in Embase and adapted to MEDLINE. We used Embase subject headings (Emtree) and Medical Subject Headings (MeSH) along with individual terms to develop the search strategy, with limitations applied (supplementary box 1). We used EndNote (version 9.2; Thomas Reuters, Philadelphia, PA, USA) as citation manager.

We performed the search on 2 June 2020. We used a standardised data extraction form developed *a priori* in Excel, which was piloted on five randomly selected studies and then refined. Data were recorded for the following: publication details (authors, title, year of publication, country and journal), study characteristics (data collection period, study design, countries that contributed with data, inclusion criteria, clinic type, sample size, population characteristics and diagnostic data), and outcome details (outcomes reported, definitions used, correlation between different outcome measures, equipment used and measurement details).

Two reviewers independently assessed titles and abstracts for eligibility. Full text was obtained for all studies deemed relevant by either reviewer or if there was uncertainty on eligibility. Where disagreements remained after full text review, the manuscripts were discussed with a third person. One reviewer manually searched the reference lists of all eligible studies for additional manuscripts. Three reviewers extracted data for a third of the eligible studies each. A fourth reviewer extracted data from 10% of the total manuscripts included in the study, and their extractions were compared with those extracted by the other three reviewers to ensure consistency.

Inclusion and exclusion criteria

We included studies describing clinical outcome measures in PCD if they 1) had a study population of ≥ 10 PCD patients, 2) were published in English, 3) were published after 1996 and 4) were conducted on humans. We did not include studies prior to 1996 because the diagnosis of PCD has changed in the past 20 years, therefore older manuscripts may contain a high proportion of patients that would no longer fulfil the current diagnostic criteria [16]. Details of diagnostic data for each of the included studies were recorded (supplementary table 1).

We excluded conference abstracts, studies that were not original research and those for which full texts were irretrievable. Studies reporting on multiple disease groups were excluded if the PCD data could not be clearly identified. Manuscripts that reported exclusively on ear, nose and throat symptoms were also excluded.

TABLE 1 Clinical outcome measures used in studies included in this review, grouped by main outcome measure

First author (year) [ref.]	Study outcomes	Population descriptors
Spirometry and/or body plethysmography		
DAVIS (2015) [20]	Anthropometry, spirometry, CT	Microbiology
DAVIS (2019) [21]	Spirometry, anthropometry, microbiology	None
ELLERMAN (1997) [22]	Spirometry	Chest radiography, microbiology
FUGER (2018) [23]	Capillary blood test, spirometry	CT, anthropometry, microbiology
GOKDEMIR (2014) [24]	Spirometry, comfort and efficacy, S_{pO_2}	Anthropometry
HALBEISEN (2018) [25]	Spirometry	Anthropometry
HELLINCX (1998) [26]	Spirometry, body plethysmography	None
KOH (2000) [27]	Spirometry	None
LOPES (2015) [28]	Spirometry and body plethysmography, HRCT, dyspnoea	Anthropometry, treatment
MAGLIONE (2014) [29]	Anthropometry, spirometry, microbiology	None
MARTHIN (2010) [30]	Spirometry	HRCT, chest radiography
OLVEIRA (2017) [31]	Spirometry, microbiology, treatment, CT	Anthropometry
PHILLIPS (1998) [32]	Spirometry	None
PIFFERI (2012) [33]	Body plethysmography, HRCT	Microbiology
SHAH (2016) [34]	Body plethysmography, HRCT, microbiology	None
SUNTHER (2016) [35]	Spirometry	Anthropometry, microbiology, treatment
TAMALET (2001) [36]	Spirometry, blood gas	CT, treatment
VALLET (2013) [37]	Spirometry, blood gas, CT	None
VIDEBAEK (2019) [38]	Spirometry, microbiology	None
MBW		
AHMAD (2015) [39]	MBW	None
ANAGNOSTOPOULOU (2018) [40]	MBW	Anthropometry
GREEN (2012) [41]	SF ₆ MBW, spirometry	Anthropometry, microbiology
GREEN (2016) [42]	MBW, spirometry	Anthropometry
IRVING (2013) [43]	Spirometry, MBW, HRCT	Microbiology
IRVING (2017) [44]	Spirometry, MBW	None
IRVING (2018) [45]	MBW	Spirometry, microbiology
KOBBERNAGEL (2019) [46]	Spirometry, MBW	Microbiology, anthropometry
KOUČKÝ (2020) [47]	MBW, spirometry, endobronchial thickness, bronchoalveolar lavage	Anthropometry, microbiology
NYILAS (2017) [48]	MBW/SBW, body plethysmography	Microbiology, treatment
NYILAS (2018) [49]	Structural and functional MRI, MBW, spirometry	Anthropometry
SMITH (2018) [50]	MRI, MBW, spirometry	Anthropometry
HRCT		
BOON (2015) [12]	Spirometry, N ₂ MBW, HRCT	Anthropometry
COHEN-CYMBERKNOH (2014) [51]	HRCT, spirometry, microbiology	Anthropometry
DETTMER (2018) [52]	CT	Microbiology, spirometry, anthropometry, number of exacerbations
HOANG-THI (2018) [53]	Spirometry, CT	Anthropometry
JAIN (2007) [54]	Chest radiography, HRCT	Microbiology
KENNEDY (2007) [55]	HRCT	Spirometry, microbiology, lobectomy
LI (2005) [56]	HRCT	Spirometry, microbiology
MAGLIONE (2012) [57]	Spirometry, HRCT	None
MAGLIONE (2017) [58]	MRI, CT	Spirometry, anthropometry, treatment, microbiology
MAGNIN (2012) [59]	Spirometry, arterialised capillary blood gases, CT	None
MONTELLA (2009) [60]	HRCT, MRI, body plethysmography	Microbiology
MONTELLA (2009) [61]	HRCT, MRI	None
SANTAMARIA (2008) [62]	HRCT	Spirometry, microbiology
TADD (2019) [63]	CT	None
Microbiology		
ALANIN (2015) [64]	Microbiology	Spirometry
COHEN-CYMBERKNOH (2017) [65]	Microbiology, spirometry, CT	Anthropometry
RODEN (2019) [66]	Microbiology, spirometry	None

Continued

TABLE 1 Continued

First author (year) [ref.]	Study outcomes	Population descriptors
ROGERS (2013) [67]	Microbiology	Spirometry
Anthropometry		
GOUTAKI (2017) [68]	Anthropometry, spirometry	None
SVOBODOVÁ (2013) [69]	Anthropometry	None
HRQoL		
ALANIN (2017) [70]	HRQoL, microbiology, spirometry, anthropometry	None
BEHAN (2017) [71]	HRQoL	Microbiology, spirometry
CAROTENUTO (2013) [72]	HRQoL	Anthropometry
IOANNOU (2020) [73]	HRQoL	Spirometry
KENİŞ COŞKUN (2019) [74]	HRQoL	Spirometry, anthropometry, microbiology
MAGLIONE (2014) [75]	HRQoL, spirometry, exercise testing	Exacerbations, microbiology
McMANUS (2003) [76]	HRQoL	Treatment
McMANUS (2006) [77]	HRQoL	None
PIFFERI (2010) [78]	HRQoL	Treatment
VALERO-MORENO (2020) [79]	HRQoL	Spirometry
WHALLEY (2006) [80]	HRQoL	None
ZENGİN AKKUS (2019) [81]	HRQoL, sleep	None
Sleep disorder		
COHEN-CYMBERKNOH (2019) [82]	Sleep questionnaires, HRQoL	Spirometry
OKTEM (2013) [83]	Body plethysmography, sleep questionnaire, PSQI, polysomnography, HRCT	Anthropometry
SANTAMARIA (2014) [84]	Respiratory polysomnography, sleep questionnaire, HRCT	Spirometry, anthropometry, treatment, microbiology
ŞIŞMANLAR (2018) [85]	Sleep, attention deficit	Spirometry, radiography
Inflammatory markers		
BUSH (2006) [86]	Inflammatory markers, sputum biophysical and transport properties	Spirometry, microbiology
COCKX (2017) [87]	Inflammatory markers	Spirometry, microbiology
COCKX (2017) [88]	Inflammatory markers	Spirometry, microbiology
PAFF (2017) [89]	HRQoL, LRTI-VAS, exacerbations, inflammatory markers in blood, inflammatory markers in sputum, spirometry, adverse events, adherence	Anthropometry, MRC dyspnoea scale score, HRCT or chest radiography
RATJEN (2016) [90]	Inflammatory markers from sputum, spirometry, microbiology	None
ZIHLIF (2006) [91]	Inflammatory markers from exhaled breath condensate and sputum	Spirometry
Exacerbations		
KOBBERNAGEL (2020) [92]	Number of exacerbations, spirometry, body plethysmography, MBW, HRQoL, inflammatory markers, microbiology	Pulse oximetry saturation, respiratory rates, anthropometry
PIATTI (2020) [93]	Number of exacerbations, CT, spirometry, microbiology	Anthropometry
Exercise testing		
LOOMBA (2017) [94]	Spirometry, exercise testing	None
MADSEN (2013) [95]	N ₂ MBW, spirometry, body plethysmography, exercise testing, HRQoL	Anthropometry, microbiology
RING (2018) [96]	Exercise testing, spirometry	Anthropometry, microbiology
SİMSEK (2018) [97]	Spirometry, exercise testing, physical activity level	Anthropometry
VALERIO (2012) [98]	Spirometry, exercise testing, physical activity assessment	Anthropometry
WELLS (2011) [99]	Exercise testing	Spirometry, anthropometry, Habitual Activity Estimation Scale questionnaire
Others		
JOENSEN (2014) [100]	Breath profiles, microbiology, number of exacerbations	Spirometry
KAWAKAMI (1996) [101]	Chronic sputum production, sputum and nasal scores	Fertility
KENNEDY (2007) [102]	Lithoptysis, radiographic findings	Spirometry, microbiology, lobectomy
MARINO (2019) [103]	Nutrition, spirometry, anthropometry, inflammatory markers	None
MIRRA (2015) [104]	Vitamin D, body plethysmography, HRQoL, physical activity assessment, microbiology	Anthropometry, HRCT, treatment
	Breath profiles	

Continued

TABLE 1 Continued

First author (year) [ref.]	Study outcomes	Population descriptors
MONTUSCHI (2014) [105]		Spirometry, microbiology, anthropometry, treatment
NOONE (1999) [106]	Clearance during cough, sputum production rate	Spirometry, cough questionnaire
PAFF (2013) [107]	Exhaled breath profile	Spirometry, microbiology, pulmonary exacerbations
PIFFERI (2017) [108]	Spirometry, HRCT, body plethysmography, microbiology, extracellular matrix	None
SHOEMARK (2009) [109]	F_{eNO}	Anthropometry, spirometry, treatment, microbiology, nasal NO
SMIT (1996) [110]	Lung resection, symptoms questionnaire	Spirometry, bronchiectasis, dyspnoea index
ZIHLIF (2005) [111]	Cough frequency, cough symptom score	Spirometry, eNO, inflammatory markers, microbiology
No main study outcome		
ABITBUL (2016) [112]	None	CT, fertility, microbiology, spirometry
BOON (2014) [113]	None	Anthropometry, spirometry, microbiology, chest radiographs and CT
EDEN (2019) [114]	None	Spirometry, microbiology, number of exacerbations
EMIRALIOĞLU (2020) [115]	None	Spirometry, anthropometry, microbiology, lobectomy, CT
FRIJA-MASSON (2017) [116]	None	Spirometry, microbiology, HRCT, dyspnoea score, treatment, fertility, lobectomy, mortality
KNOWLES (2014) [117]	None	Spirometry, fertility
NOONE (2004) [118]	None	Spirometry, microbiology, radiographs, cough
PIFFERI (2015) [119]	None	Spirometry, HRCT, microbiology
YIALLOUROS (2015) [120]	None	CT, microbiology, spirometry, anthropometry, lobectomy

MBW: multiple-breath washout; HRCT: high-resolution computed tomography, HRQoL: health-related quality of life; CT: computed tomography; S_{pO_2} : peripheral oxygen saturation; SF₆: sulfur hexafluoride; SBW: single-breath washout; MRI: magnetic resonance imaging; N₂: nitrogen; PSQI: Pittsburgh Sleep Quality Index; LRTI: lower respiratory tract infection; VAS: visual analogue scale; MRC: Medical Research Council; F_{eNO} : exhaled nitric oxide fraction; NO: nitric oxide; eNO: exhaled nitric oxide.

Definition of outcome measures and classification into subgroups

Outcome measures were defined as any clinical measure used to monitor patients over time or as a marker of disease severity. Outcome measures were classified as 1) study outcomes, defined *a priori* as study outcome measures; or 2) study population descriptors. The latter indicates measures that were used to characterise the study population (*e.g.* baseline measures of forced expiratory volume in 1 s (FEV₁)) and those that could potentially be used in future studies (*e.g.* cough frequency). The supplementary tables contain detailed information on study characteristics and definitions of outcome measures for all studies included in this review.

Statistical analysis

Critical appraisal of individual studies was not conducted because our focus was on the variety of outcomes used and how these were measured and reported, and not on disease progression, prognosis or treatment effects [17, 18]. Information about scoping reviews is outlined in the supplementary files.

Descriptive and summary data were analysed in R statistical package (version 3.2.3; www.r-project.org). Continuous variables were reported as median and interquartile range (IQR). Categorical variables were reported as proportions. Results were reported according to the Preferred Reporting Items for Systematic Reviews and Meta-Analysis Extension for Scoping Reviews checklist [19]. Figures were plotted in R and Tableau 2019 v4.0.

Results

3150 abstracts were identified, of which 2706 were reviewed after exclusion of 444 duplicates. 198 manuscripts were reviewed in full, of which 102 met the inclusion criteria and were therefore included (supplementary figure 1) [12, 20–120].

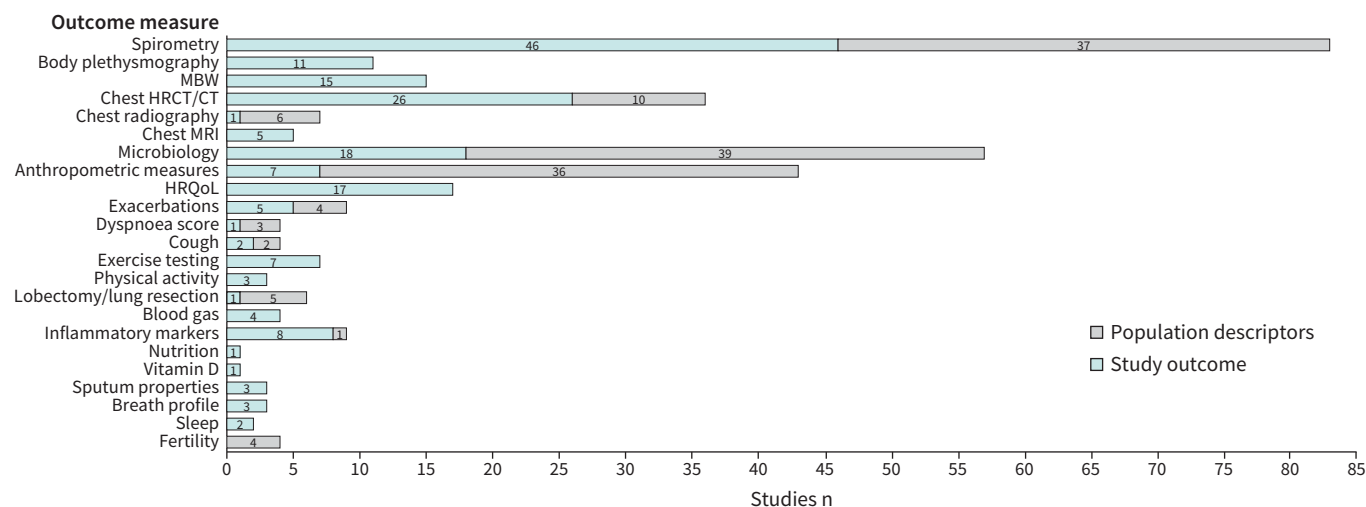


FIGURE 1 Number of studies that reported outcome measures in primary ciliary dyskinesia as either study outcome or population descriptor. Studies often reported on more than one outcome measure and might therefore be featured in more than one instance. MBW: multiple-breath washout; HRCT: high-resolution computed tomography; CT: computed tomography; MRI: magnetic resonance imaging; HRQoL: health-related quality of life.

Study characteristics

The manuscripts included information on 7289 patients with PCD, with a median of 32 PCD patients per manuscript (IQR 20–62, range 10–1609; supplementary table 1). However, some patients were described in several studies and were included more than once if the studies described different outcome measures.

Manuscripts contained data collected from 23 different countries, but most (89%) presented single-centre data. Publications with a higher number of PCD patients were from multicentre studies, with the two largest studies containing data derived from a large international PCD cohort study [121].

Outcome measures reported

93 studies reported on a total of 23 main study outcomes (table 1, figure 1), with 19 presenting data exclusively on population descriptors. 67 presented both study outcomes and population descriptors (table 1, supplementary table 2).

Spirometry-derived parameters were the most frequently reported clinical outcome measures, followed by chest high-resolution computed tomography (HRCT). Microbiology and anthropometric measures were more often reported as descriptors than as study outcomes (figure 1).

Standardised definitions

Definitions of outcome measures varied considerably between studies (supplementary table 2). Of 18 studies reporting on microbiology as study outcomes, 14 provided data on chronic colonisation by potentially pathogenic bacteria [29, 31, 34, 38, 59, 64, 65, 70, 75, 82, 96, 100]. The terms chronic colonisation and chronic infection were sometimes used interchangeably, and the classification used to define them varied. Four studies used the Leeds CF criteria (or a modified version) [38, 64, 65, 70]; two used the European consensus for antibiotic therapy against *Pseudomonas aeruginosa* in CF [75, 104]; and one used the Copenhagen criteria [100]. The remaining studies developed a study-specific criterion, such as pathogen cultured in $\geq 50\%$ of samples from the past year [46, 47] or isolation of the same pathogen on at least two occasions, ≥ 3 months apart in a 1-year period [93].

Sampling frequency of sputum and other microbiological specimens varied between studies. Some studies did not record whether patients had a pulmonary exacerbation at the time of sampling. These differences are likely to have biased results, particularly when reporting prevalence of different respiratory pathogens.

Details on the definitions of all clinical outcome measures reported as study outcomes or population descriptors are provided in supplementary table 2. In the next sections, we focus on the most frequently reported outcome measures.

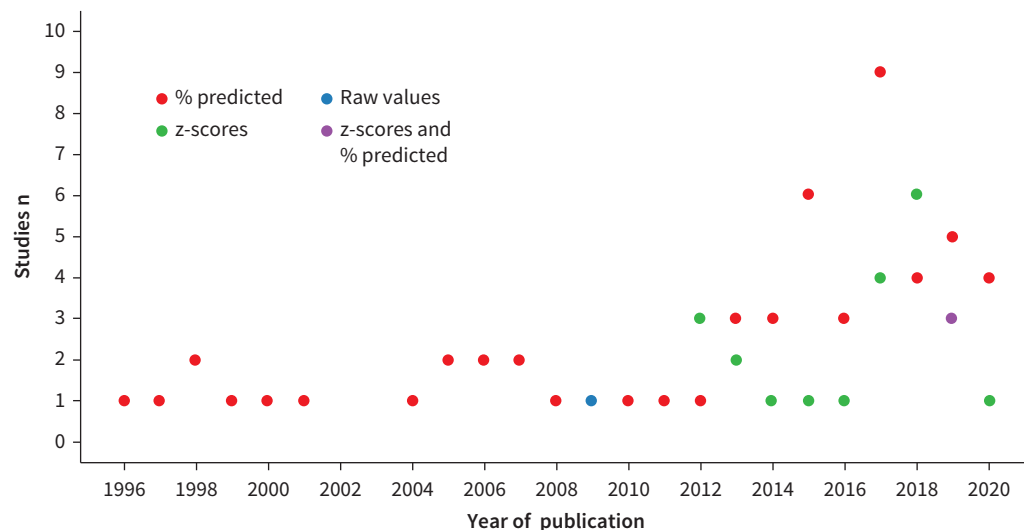


FIGURE 2 Number of studies that reported forced expiratory volume in 1 s as study outcomes or population descriptors (n=83). Circles are coloured according to the spirometric indices presented in the studies.

Lung function

Spirometry

Of the 83 studies reporting spirometry data, 42 reported adherence to European Respiratory Society/American Thoracic Society guidelines [122] (supplementary table 2). 25 reported on FEV₁ z-scores; 18 as a study outcome. FEV₁ % predicted was used as an outcome as often as a descriptor (32 versus 31). Studies reporting on FEV₁ z-scores were published more recently (figure 2). 14 studies reported calculating z-scores based on the Global Lung Function Initiative (GLI) reference equations [123], two used national references and eight used other equations, while the remainder did not detail the equation used (supplementary table 2).

Five studies compared FEV₁ before and after the use of bronchodilators. Studies used inhaled salbutamol [32, 119] or albuterol [28] to assess reversibility and one study performed methacholine challenge before and after the use of salbutamol and placebo [27]. One study did not report which bronchodilator was used [114].

Body plethysmography

11 studies reported on body plethysmography parameters as study outcomes (table 1, supplementary table 2). Lung residual volume % predicted was reported by all studies; total lung capacity % predicted was reported in four studies; and the remaining 19 parameters were rarely reported. Three of these studies additionally measured forced vital capacity (FVC) and FEV₁ using plethysmography devices.

Multiple-breath washout

15 studies reported on parameters derived from the multiple-breath washout (MBW) test as study outcomes (table 1, supplementary table 2), of which 47% were published in the past 2 years. Lung clearance index (LCI) was most frequently reported. Eight studies presented z-scores for LCI, and five presented values for S_{cond} and S_{acin} z-scores (representatives of ventilation inhomogeneity in small conducting and acinar airways, respectively). Studies used different inert tracer gases and equipment; nine of them used nitrogen (N₂), five used 0.2% sulfur hexafluoride and the other study did not report which tracer gas was used.

Chest imaging

40 manuscripts reported on radiological findings, with 26 presenting them as study outcomes and 14 as population descriptors (table 1, supplementary table 2). Of the latter, five studies had spirometry measures as outcomes and provided information on presence or absence of bronchiectasis, diagnosed through chest HRCT/CT or radiography. One study mentioned chest radiography to determine the presence of bronchiectasis; however, the main outcomes were sleep activity and attention deficit scales [85]. The remaining studies did not report on specific outcomes, with data on descriptors only including spirometry, microbiology, anthropometric measurements and fertility.

Four studies reported on both chest radiography and HRCT and two studies on both chest HRCT and magnetic resonance imaging (MRI).

Radiography

Bronchiectasis, seen on chest radiography, was used in one study as study outcome and in six as population descriptor (table 1, supplementary table 2). Similar to other imaging modalities, there are no PCD-specific radiography scoring systems, so studies used different scales to report findings. For example, JAIN *et al.* [54] used a modified version of the Chrispin–Norman score, which was developed for CF [124], while KENNEDY *et al.* [55] developed a study-specific score for bronchiectasis severity.

HRCT

Chest HRCT and/or CT was used as study outcome in 25 studies, with an additional 10 studies reporting it as population descriptor (figure 1, table 1, supplementary table 2). Studies adopted modified versions of different scoring scales as there are no PCD-specific scoring systems available. Seven studies used modifications of the Brody score [125], six used the Bhalla score [126], another four applied the Helbich score [127] and a further six used other systems. Of the latter, four used a study-specific score, one by combining the Brody and Bhalla scores [59]. Two studies did not provide any detail on the scoring system used [37, 56] and therefore were not included in figure 3.

The use of different measurement scales resulted in inconsistent reporting of subscores (figure 3). For example, extent of bronchiectasis was measured by 1) number of bronchopulmonary segments affected; 2) percentage of each lobe involved; 3) scores from 0 to 3; 4) percentage of central lung and peripheral lung involvement; or 5) size of largest and average bronchopulmonary segment involved. Mucus plugging was measured as size of plug (*i.e.* small, large), location of plug (Brody score: largest airways, small airways, peripheral lung, central lung; or Helbich score: number of segments) or a mucus classification score (Bhalla score).

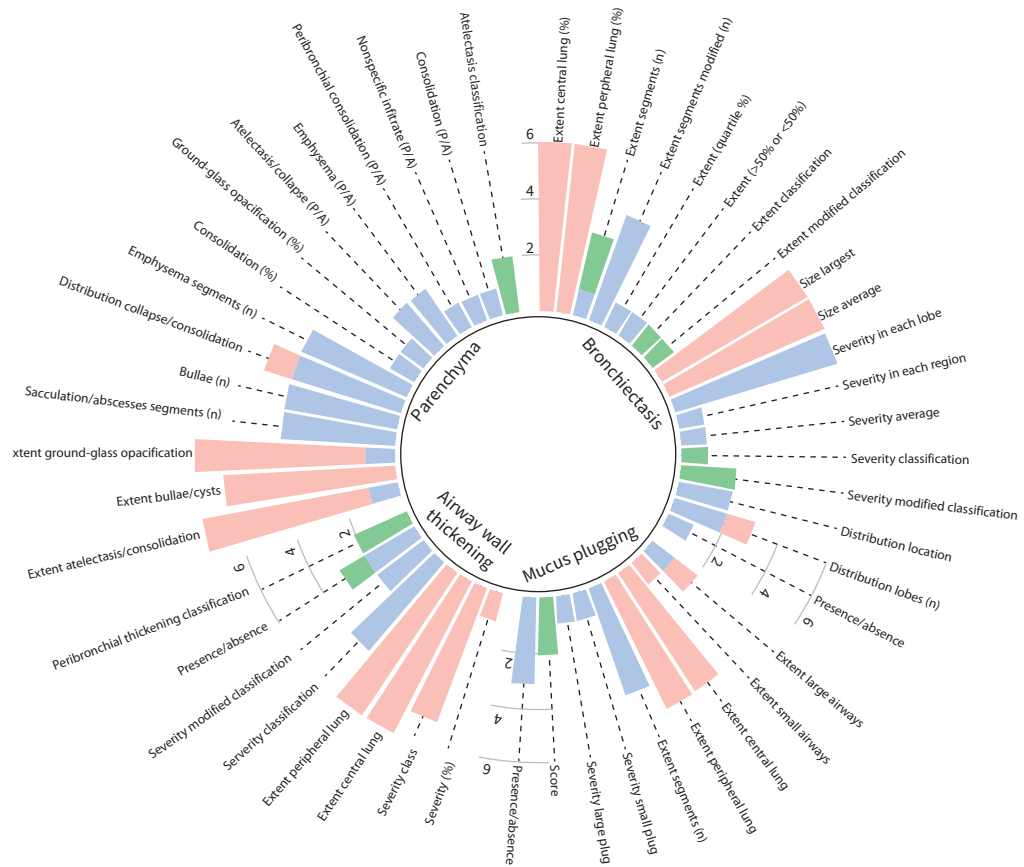


FIGURE 3 High-resolution computed tomography (HRCT) and computed tomography (CT) outcome measures from 23 studies that reported on HRCT/CT scans as study outcomes and had sufficient data on scoring system adopted. Stacked bars represent the number of studies that reported on each subscore component. P: present; A: absent.

As illustrated in figure 3, not all studies using the Brody score reported on the same subscore components, probably due to study-specific modifications. For example, one study [12] classified the location of the mucus plug as small or large airways, while five other studies used number of central and peripheral lobes involved. In another study that used a modified combination of the Brody and Bhalla scores, the partition of lungs into different segments followed a regional approach as opposed to the commonly used pulmonary segmentation approach in order to expedite the time needed for scoring each scan in routine clinical practice [59].

Unsurprisingly, all 23 studies that presented information on their scoring system reported on bronchiectasis. Studies classified bronchiectasis as mild for those with airway diameter slightly greater than the diameter of adjacent blood vessel, moderate for airway lumen two or three times the diameter of the vessel and severe for those at least three times the diameter of the vessel. However, the extent of bronchiectasis varied, with some reporting the number of bronchopulmonary segments, while others reported percentage of compromised area (figure 3). The second most common features described were airway wall thickening and mucus plugging (n=19).

Two studies comparing CT scores in PCD and CF patients found no differences in the global Brody score. A third study used a study-specific system to analyse CT scans from patients with PCD and CF and then assess results against the Brody and Bhalla scores [63]. They found that bronchial wall thickening, bronchiectasis, mucus plugging, atelectasis and air trapping, features commonly seen in CF patients, were even more common in patients with PCD.

MAGLIONE *et al.* [57] reported a significantly higher subscore for severity of collapse or consolidation in PCD compared to CF, and COHEN-CYMBERKNOH *et al.* [82] found that the lower and middle lobes were more frequently affected in PCD compared to the typical upper lobes compromise seen in CF [34, 55, 62, 82]. TADD *et al.* [63] reported a higher frequency of extensive tree-in-bud pattern of mucus plugging, bronchocoeles or nodules, thickening of interlobar and intralobular septa, and atelectasis or collapse of the whole lobes in PCD; these are uncommon in CF patients.

Magnetic resonance imaging

Only five studies reported on chest MRI as study outcome (table 1, supplementary table 2). Four studies applied a modified Helbich scoring system, while SMITH *et al.* [50] developed a study-specific scoring system for three-dimensional volumetric hyperpolarised MRI. When looking at subscores, MAGLIONE *et al.* [75] found no significant difference between total MRI scores and subscores in 20 PCD and 20 mild CF patients, aside from a higher score for severity of collapse/consolidation in PCD patients. In a smaller study of 11 PCD children, all presented with mostly small and heterogeneous abnormalities on ventilation MRI [50].

Microbiology

57 studies reported on microbiology; 18 as study outcomes and 39 as population descriptors (table 1, supplementary table 2). Studies reported most commonly on *Haemophilus influenzae* and *P. aeruginosa*, followed by *Staphylococcus aureus*. Some studies distinguished between mucoid and nonmucoid strains of *P. aeruginosa*, while others simply reported on *P. aeruginosa* infection. Similarly, *S. aureus* subtypes were inconsistently stratified across studies, with some reporting methicillin-sensitive and -resistant strains separately. Not all studies stratified pathogen prevalence by age group (supplementary figure 2).

Other outcome measures

Health-related quality of life scores

17 studies reported on health-related quality of life (HRQoL) as study outcomes (table 1). However, only two studies [74, 92] used QOL-PCD, as most were published before the disease-specific tool was validated [71, 128, 129]. The most common instruments adopted were the St George's Respiratory Questionnaire (SGRQ) (eight studies) and the 36-item short form survey (SF-36) (seven studies) (supplementary table 2).

Pulmonary exacerbations

Nine studies reported on pulmonary exacerbations; five as study outcomes. However, none used the PCD-specific consensus, as the studies included in this review pre-date it [130]. Two randomised controlled trials (RCTs) used pulmonary exacerbation as a primary outcome. PAFF *et al.* [89] defined an exacerbation as respiratory symptoms that led to initiation of systemic antibiotic treatment irrespective of culture results, or a decline of $\geq 10\%$ in FEV₁ % predicted compared to baseline at screening and randomisation [89], while KOBBERNAGEL *et al.* [92] defined it as worsening of respiratory symptoms leading to initiation of antibiotic treatment in the week prior to the clinical appointment up to the day of the

appointment. RATJEN *et al.* [90] studied a subset of patients who experienced an episode of exacerbation, defined as an increase in lower airway symptoms treated with oral antibiotics. JOENSEN *et al.* [100] applied a definition developed for CF studies [61] (supplementary table 2). SUNTHER *et al.* [35] only included patients with pulmonary exacerbation, defined as change in respiratory status for which intravenous antibiotics were needed.

Comparison between outcome measures

Most studies comparing outcome measures used spirometry as the reference to which the other outcomes were compared (figure 1). 12 studies describing imaging modalities reported on agreements or correlations with other outcome measures [12, 33, 43, 49, 53, 55, 57, 60, 93, 108, 119]. The most common comparison was between spirometry-derived FEV₁ and HRCT, with studies presenting contradictory findings. Four studies [33, 53, 55, 93] found agreement between the two outcomes, one of which used an automated CT scoring for adults with PCD [53]. Three studies used a modified Bhalla system and the other a study-specific scoring system. The other four studies [43, 57, 62, 119] reported no association.

FEV₁ was compared to MBW-derived LCI in eight studies, also with contradictory results. Two studies reported no association [41, 44], while the other six [12, 35, 42, 43, 46, 49] found correlations between some parameters. Both BOON *et al.* [12] and KOBBERNAGEL *et al.* [46] found a significant negative correlation between LCI, FEV₁ and FEV₁/FVC ratio z-scores, while IRVING *et al.* [43] only found a correlation between LCI and forced expiratory flow at 25–75% of FVC (FEF_{25–75}) z-scores. GREEN *et al.* [42] did not find any correlation between LCI and FEV₁ z-scores in PCD patients, but reported a significant correlation between LCI_{2.5} (LCI at 2.5%, standard LCI) and FEV₁/FVC ratio and FEF_{25–75} z-scores.

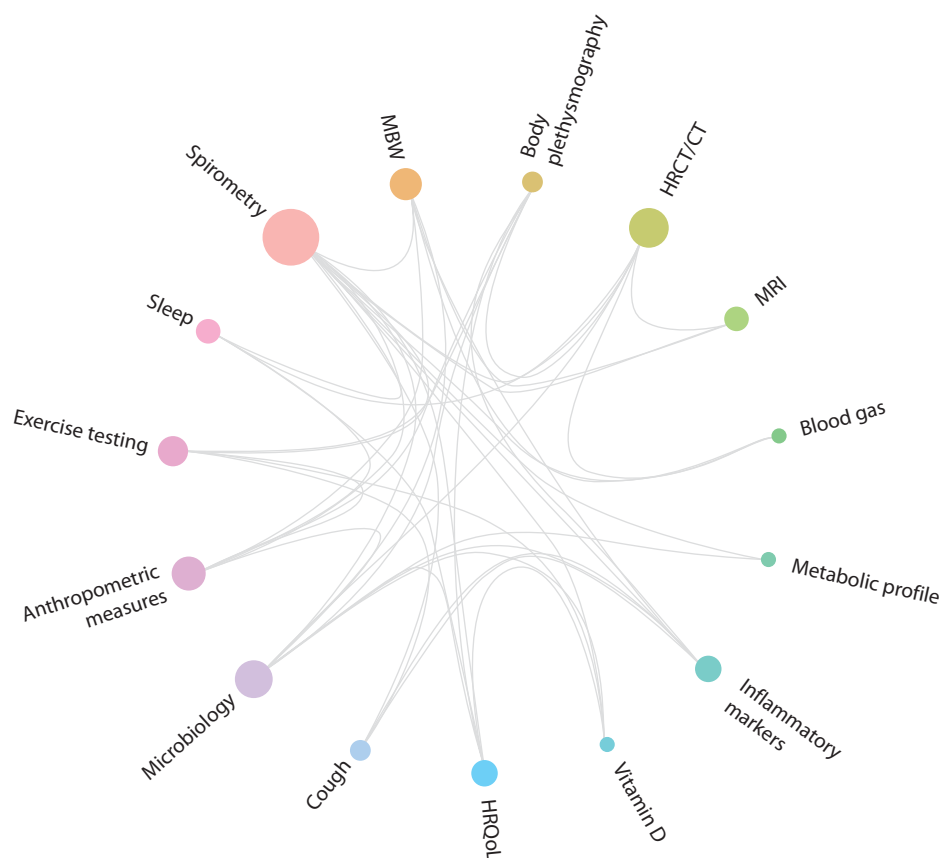


FIGURE 4 Correlations or associations between outcomes measures. Connections between circles depicts the correlated outcomes, with the size of each circle representing the number of studies that reported correlations or associations of that particular outcome. MBW: multiple-breath washout; HRQoL: health-related quality of life; MRI: magnetic resonance imaging; HRCT: high-resolution computed tomography; CT: computed tomography.

MBW-derived LCI might be more sensitive to detect early or milder disease. NYILAS *et al.* [49] found that more than half of the patients with abnormal LCI values and MRI scores had normal FEV₁ z-scores. In another study, five (15%) patients had abnormal LCI, but normal FEV₁ z-scores [43]. In addition, LCI was shown to be more sensitive than FEV₁ to detect lung structure abnormalities [12]. BOON *et al.* [12] reported that LCI z-scores were concordant with total CFCT scores (a variant of the Brody score) in 83% of the patients, while KOBBERNAGEL *et al.* [46] and IRVING *et al.* [43] found no correlation.

Studies comparing HRCT to indices derived from body plethysmography, chest MRI and microbiology found significant correlations. However, these were generally limited to subscores (*e.g.* bronchiectasis on HRCT and body plethysmography and collapse/consolidation on HRCT and MRI) as opposed to the global score.

Other associations between outcome measures are shown in figure 4.

Randomised controlled trials

Only five of the included studies were RCTs, of which four adopted a crossover design (supplementary table 3).

The efficacy of 6 months azithromycin maintenance therapy in reducing the number of respiratory exacerbations in patients with PCD was assessed in a double-blind, parallel-group placebo-controlled RCT at six European PCD centres [92]. Secondary outcomes included changes in spirometry, body plethysmography, N₂ MBW, HRQoL, audiometry, sputum microbiology and inflammatory markers.

The effect of hypertonic saline on HRQoL in PCD adults was investigated in a 28-week double-blind crossover RCT with a washout period of 4 weeks. HRQoL was measured by the SGRQ and Quality of Life Questionnaire – Bronchiectasis [89].

GOKDEMIR *et al.* [24] assessed spirometry measurements (FEV₁, FVC, peak expiratory flow and FEF_{25–75}) in children with PCD using two different airway clearance methods. Half performed conventional pulmonary rehabilitation for 5 days in hospital followed by a 2-day washout period and then high-frequency chest-wall oscillation for another 5 days at home. However, techniques differed between the settings. Another crossover RCT investigated differences in FEV₁ % predicted and in bronchial hyperresponsiveness after the use of salbutamol compared to placebo in children with PCD at both 3 and 6 weeks compared to pre-treatment measurements [27].

NOONE *et al.* [106] reported on mean whole-lung clearance rates of a radionuclide marker after inhalation of uridine-5'-triphosphate compared to placebo during a series of controlled coughs to induce mucociliary clearance in PCD adolescents and adults.

Discussion

This scoping review identified 23 clinical outcome measures used in PCD research. We found a high degree of heterogeneity in the definitions of outcome measures.

Spirometry and chest HRCT were most frequently reported as study outcomes. Spirometry is widely available, relatively easy to perform and does not require expensive equipment [122, 123]; however, researchers have questioned its appropriateness as a measure to monitor disease progression in PCD [35, 49, 57]. A meta-analysis found that mean FEV₁ ranged from 51% predicted to 96% predicted, with high heterogeneity between studies that could not be explained by age or other factors [131]. Studies that did not report on which reference values they used or those that did not provide information on quality control reported lower mean FEV₁ values. Clinical status at the time of measurement was rarely reported and therefore could not be included in the meta-regression. The largest study to date investigating lung disease in PCD patients found consistently low FEV₁ z-scores in patients with PCD compared to reference data, similar to those seen in CF patients [25].

To our knowledge, no study has investigated the timing of physiotherapy in relation to spirometry, which is a significant limitation as, anecdotally, airway clearance techniques can improve spirometric indices. An ongoing multicentre prospective cohort is investigating variability of lung function in stable PCD patients, adjusting for factors such as timing of inhaled medication and respiratory physiotherapy [132, 133] (<https://clinicaltrials.gov/ct2/show/NCT03704896>). Another potential source of variability when using spirometry-derived measurements are the adopted reference equations, as variations between the GLI and national reference equations can occur. Evidence from a longitudinal CF cohort highlighted the disparity

between reference equations, demonstrating the need for a standardised approach to interpreting spirometric measurements to facilitate appropriate comparisons both within and between centres and countries [134].

Chest HRCT has been proposed as a surrogate outcome measure in the assessment of lung disease. However, there are no validated scoring systems for PCD. All studies included in this review used CF-derived scoring systems [125–127], despite significant pathophysiological differences between the two conditions [9, 135]. Additionally, studies do not report the lung volumes at which the CT scans are obtained, with no details on the standard operating procedures used to record the images.

Location, distribution and frequency of features seen in HRCT scans of patients with PCD differ from those with CF [135]. The weights applied to each feature might not be suitable for PCD as CF-derived scoring systems do not reflect the range and severity of structural changes in PCD. Studies found that extensive tree-in-bud pattern of mucus plugging, bronchocoeles or nodules, thickening of interlobar and interlobular septa and atelectasis mostly seen as collapse of whole lobes were frequently described in PCD, but uncommon in CF [52, 63, 135]. Reporting only the global CT scores might be misleading, as some components of the score might be more relevant to clinical outcome, particularly when using a non-disease-specific score. These findings underscore the need for disease-specific CT scoring systems. HOANG-THI *et al.* [53] highlighted the fact that visual scores such as the ones routinely used in the assessment of PCD and CF patients can be highly subjective. In response, they developed an automated CT scoring for adults with PCD, which had moderate to good correlation with FEV₁ and FVC.

MRI scans of the chest have historically been considered of limited value due to intrinsic characteristics of the pulmonary tissue, and the presence of physiological motion resulting in poor resolution and motion artefacts. Research has focused on improving techniques to obtain better-quality images [136, 137].

Lack of agreement between spirometry, HRCT, MBW and MRI parameters reported by some studies might reflect variations on measurement and reporting of outcomes. Discrepancies could be explained by different scoring systems for HRCT, differences in tracer gas for MBW, variations in measurements, inability of some of the outcome measures to accurately monitor lung disease progression in PCD or true variability between populations (*e.g.* underlying genetics, differences in disease severity or treatment). Interpretation of findings was limited by the retrospective nature of most studies. In some cases, there was a significant time lag between measurements performed with the methods that were compared [12], or tests were applied to different subpopulations (*e.g.* HRCT scans conducted only in the older population with more severe lung disease [43] or conducted at different time points of clinical stability [57]). Contradictory results could also be attributed to variations in study design, inclusion criteria or small sample sizes, resulting in variability due to chance.

Recent studies have focused on MBW, with almost half of them published in the past few years. NYILAS *et al.* [49] found that LCI was not able to distinguish between reversible and irreversible lung damage, despite being more sensitive than spirometry to detect changes. A limitation of LCI is the long washout time, and therefore test duration, which is particularly problematic for patients with compromised lung capacity and young children. Studies looking at shorter washout periods have shown promising results, with LCI₅ providing a good alternative to the more conventional LCI_{2.5} [39, 41, 109]. However, as NYILAS *et al.* [49] demonstrated, combining different modalities (*e.g.* MRI and MBW) can be necessary to accurately capture changes in the lungs of PCD patients.

In terms of microbiological outcomes, studies should present a breakdown of pathogens by age group, since the prevalence of bacterial species changes with age [66, 85]. Studies included in this review were also limited by the lack of a universal panel that could be applied consistently across different centres, particularly when reporting the prevalence of each pathogen isolated. ROGERS *et al.* [67] highlighted that some of dominant genera of bacteria found in the sputum of PCD patients were from those unlikely to be detected without specific growth conditions being present. Variations in the frequency of specimen collection and type of specimen (*e.g.* expectorated sputum, cough swab, bronchoalveolar lavage) will also probably affect pathogen prevalence.

Small sample sizes were a common limitation in most studies, highlighting the importance of national and international disease registries, large collaborative multicentre studies and standardised definitions that enable pooling of data [121, 138–140]. Few studies included sample-size calculations, hampering the interpretation of statistically insignificant results due to underpowered samples.

The number of larger multicentre studies has increased in recent years, highlighting the important role of PCD networks such as Better Experimental Screening and Treatment for Primary Ciliary Dyskinesia (BESTCILIA), Better Experimental Approaches to Treat Primary Ciliary Dyskinesia (BEAT-PCD) and the Genetic Disorders of Mucociliary Clearance Consortium in advancing collaborative research in the field [132, 133, 141, 142]. Such collaborations were featured in two studies that used data derived from the international PCD cohort [121].

RCTs and prospective cohort studies with long follow-up periods are uncommon in rare diseases due to the small sample sizes available, high costs and limited commercial interest from pharmaceutical industries [5, 143, 144]. As a result, the majority of PCD studies are cross-sectional case-control studies or small cohort studies with limited follow-up. Interventional studies are currently being designed, but will require close international collaboration and data sharing. The success of these and of future trials will depend on the selection of appropriate outcome measures.

Our review was limited by the quality of the information provided in the studies. As our aim was to identify the evidence available and describe definitions for clinical outcome measures used in PCD research, we opted to conduct a scoping review and therefore we did not critically appraise the studies included in this review. We did not perform quantitative analysis, as studies were heterogeneous, impeding a formal meta-analysis. In fact, the aim of this review was to highlight this heterogeneity. Another limitation was that the broad nature of this review impeded us from focusing on any one clinical outcome measure, and therefore systematic reviews with or without meta-analysis are still needed for the more commonly used and promising outcome measures. A separate review to evaluate upper airways clinical outcome measures is underway, and therefore we deliberately excluded these studies from our scoping review. Despite our attempts not to restrict the search to specific outcomes, our review is limited to the clinical outcomes that were included as search terms.

Recommendations

We advocate that outcome measures for use in future prospective trials must fulfil the following criteria: 1) measured across different studies in a standardised manner (*e.g.* using the standardised PCD data collection tool FOLLOW-PCD [145]); 2) used and reported regularly by a sufficient number of studies; 3) use currently recommended definitions (*e.g.* z-scores based on GLI recommendations); and 4) be embedded within the current knowledge of PCD pathophysiology and natural history (table 2). This will require consensus statements, which are currently being developed by a BEAT-PCD working group.

Spirometry was the outcome most frequently used for disease monitoring, but there were major problems with standardisation on measuring and reporting FEV₁. Large studies are needed to investigate the suitability of spirometry-derived parameters as accurate and sensitive surrogate markers.

Chest HRCT might be a good candidate for longitudinal follow-up of lung disease progression in PCD, particularly modalities using low radiation [146, 147]. However, a disease-specific scoring system must be developed. Agreement between HRCT and other outcomes were limited to subscores as opposed to global score, emphasising the need for PCD-specific scores that consider the distribution, frequency and patterns of lung compromise in this population, and that can be easily applied by clinicians without being unnecessarily time-consuming.

The United States Food and Drug Administration and the European Medicines Agency encourage the inclusion of patient-centred outcome measures. A systematic review on the patient's experience of PCD reported worsening of respiratory symptoms with age, which was also associated with decline in the physical and mental domains [148]. QOL-PCD, a HRQoL instrument, is the only validated disease- and age-specific cross-cultural clinical outcome measure in PCD [71, 128, 129, 149, 150]. QOL-PCD correlated well with the Sino-Nasal Outcome Test (SNOT-20) for upper airway symptoms, the COPD-specific SGRQ for lower airways symptoms and SF-36 for physical functioning, role functioning and mental health [71].

An expert consensus on the definition of pulmonary exacerbations in PCD for children and adults was developed recently [130]. The importance of disease-specific definitions was highlighted by the fact that studies that used exacerbations as an outcome adopted different definitions for pulmonary exacerbations [35, 90, 92]. There is now a need to validate the proposed definition and develop a separate definition for upper airway exacerbations.

TABLE 2 Summary of clinical outcome measures for use in primary ciliary dyskinesia (PCD) research

	Strengths	Limitations	Future directions
Spirometry	Routinely measured Reported in 78% of studies included	Unknown accuracy and sensitivity as surrogate marker for lung disease Unstandardised reporting of indices	Investigate appropriateness of spirometry to monitor disease progression Standardised reporting of spirometric indices (<i>i.e.</i> use of z-scores) Report and investigate appropriateness of other spirometric indices (<i>e.g.</i> FVC, FEV ₁ /FVC ratio) Routine performance and reporting of quality control
HRCT	Can assess structural lung damage Reported in 37% of studies included	Reliance on established CF scoring systems Frequent high doses of radiation if used routinely for disease monitoring; however, low-dose radiation modules are being developed	Identify the most relevant features and subscores for PCD Develop and validate PCD-specific scoring system
HRQoL	Patient-centred outcome measure QOL-PCD was developed and validated for use in PCD	Lack of a minimal clinically relevant difference	Adopt QOL-PCD as outcome measure in prospective longitudinal studies Use translated and cultural validated versions, where available Calculate minimal clinically relevant difference
Pulmonary exacerbations	Developed for use in PCD	Has not been validated	Validate definition Use in future prospective studies

HRCT: high-resolution computed tomography; HRQoL: health-related quality of life; FVC: forced vital capacity; FEV₁: forced expiratory volume in 1 s; CF: cystic fibrosis; QOL-PCD: Quality of Life – Primary Ciliary Dyskinesia.

Conclusions

This scoping review highlights the variety of outcomes and definitions used in PCD research. It also underscores significant differences in measurement and reporting of outcomes. Validated disease-specific clinical outcome measures are needed to monitor disease progression in PCD in future prospective cohort studies and clinical trials. Appropriate outcomes need to be chosen based on the specific patient groups and the study intervention. New studies should aim to measure and report outcomes using standardised methods to build up the body of evidence needed to meaningfully compare results. New promising outcome measures should also be used, such as MBW-derived LCI and microbiology, to assess and better understand the appropriateness of these for long-term monitoring in PCD.

Acknowledgement: This work was presented in poster discussion session at the ERS International Congress on 29 September 2019.

Provenance: Submitted article, peer reviewed.

Author contributions: B. Rubbo, P. Latzin and J.S. Lucas conceived the review; B. Rubbo, J. Thompson and C.L. Jackson performed the literature search; B. Rubbo, C.L. Jackson, E. Dehlink, P. Latzin and J.S. Lucas discussed and agreed on the inclusion and exclusion criteria; B. Rubbo, F. Gahleitner, J. Thompson and C.L. Jackson screened publications for eligibility; B. Rubbo, F. Gahleitner, J. Thompson, C.L. Jackson and A.P.L. Queiroz extracted the data; B. Rubbo, F. Gahleitner and J.F. Hueppe analysed the data; B. Rubbo and F. Halbeisen plotted the figures; B. Rubbo drafted the manuscript; F. Gahleitner, J. Thompson, C.L. Jackson, L. Behan, E. Dehlink, M. Goutaki, F. Halbeisen, G. Thouvenin, C. Kuehni, P. Latzin and J.S. Lucas critically reviewed the manuscript. All authors reviewed and approved the final manuscript.

Conflict of interest: M. Goutaki and C.E. Kuehni report that the BEAT-PCD COST Action supported their travel to and attendance of meetings during the period 2015–2019. P. Latzin reports personal fees from Vertex, Novartis, Roche, Polyphor, Vifor, Gilead, Schwabe, Zambon and Santhera, and grants from Vertex, outside the submitted work. J.S. Lucas reports that on 2 December 2019 she attended the Rare Lung and Airway Diseases on Childhood meeting at the Primary Ciliary Dyskinesia and Neonatal Interstitial Lung Diseases Conference in Jerusalem, Israel, to deliver an invited lecture on monitoring disease progression in PCD; her local travel and accommodation were arranged by the conference organisers, and her international travel was supported by BEAT-PCD. All other authors have nothing to disclose.

Support statement: B. Rubbo, J. Thompson, C.L. Jackson, J.F. Hueppe, L. Behan, E. Dehlink, M. Goutaki, F. Halbeisen, G. Thouvenin, C. Kuehni, P. Latzin and J.S. Lucas participate in the Better Evidence to Advance Therapeutic Options for PCD (BEAT-PCD) network (COST action BM 1407 and ERS Clinical Research Collaboration). B. Rubbo received a short-term scientific mission, provided through BEAT-PCD funding, to travel to Switzerland to discuss the review with M. Goutaki, F. Halbeisen and C. Kuehni.

References

- 1 Shoemark A, Lucas JS. Diagnosis of primary ciliary dyskinesia: current recommendations and future perspectives. *In: Chalmers JD, Polverino E, Aliberti S, eds. Bronchiectasis (ERS Monograph). Sheffield, European Respiratory Society, 2018; pp. 267–281.*
- 2 Lucas JS, Walker WT, Kuehni CE, *et al.* Primary ciliary dyskinesia. *In: Cordier JF, ed. Orphan Lung Diseases (ERS Monograph). Sheffield, European Respiratory Society, 2011; pp. 201–217.*
- 3 Lucas JS, Davis SD, Omran H, *et al.* Primary ciliary dyskinesia in the genomics age. *Lancet Respir Med* 2020; 8: 202–216.
- 4 Goutaki M, Meier AB, Halbeisen FS, *et al.* Clinical manifestations in primary ciliary dyskinesia: systematic review and meta-analysis. *Eur Respir J* 2016; 48: 1081–1095.
- 5 Kuehni CE, Goutaki M, Rubbo B, *et al.* Management of primary ciliary dyskinesia: current practice and future perspectives. *In: Chalmers JD, Polverino E, Aliberti S, eds. Bronchiectasis (ERS Monograph). Sheffield, European Respiratory Society, 2018; pp. 282–299.*
- 6 Lucas JS, Alanin MC, Collins S, *et al.* Clinical care of children with primary ciliary dyskinesia. *Expert Rev Respir Med* 2017; 11: 779–790.
- 7 Barbato A, Frischer T, Kuehni CE, *et al.* Primary ciliary dyskinesia: a consensus statement on diagnostic and treatment approaches in children. *Eur Respir J* 2009; 34: 1264–1276.
- 8 Shapiro AJ, Zariwala MA, Ferkol T, *et al.* Diagnosis, monitoring, and treatment of primary ciliary dyskinesia: PCD foundation consensus recommendations based on state of the art review. *Pediatr Pulmonol* 2016; 51: 115–132.
- 9 Lucas JS, Carroll M. Primary ciliary dyskinesia and cystic fibrosis: different diseases require different treatment. *Chest* 2014; 145: 674–676.
- 10 Valentgas P, Dreyer N, Wu AW. Outcome definition and measurement. *In: Valentgas P, Dreyer N, Nourjah P et al., eds. Developing a Protocol for Observational Comparative Effectiveness Research: a User's Guide. Rockville, Agency for Healthcare Research and Quality (US), 2013. Available from: www.ncbi.nlm.nih.gov/books/NBK126186/*
- 11 Kuehni CE, Goutaki M, Kobbernagel HE. Hypertonic saline in patients with primary ciliary dyskinesia: on the road to evidence-based treatment for a rare lung disease. *Eur Respir J* 2017; 49: 1602514.
- 12 Boon M, Vermeulen FL, Gysemans W, *et al.* Lung structure-function correlation in patients with primary ciliary dyskinesia. *Thorax* 2015; 70: 339–345.
- 13 Maglione M, Bush A, Montella S, *et al.* Progression of lung disease in primary ciliary dyskinesia: is spirometry less accurate than CT? *Pediatr Pulmonol* 2012; 47: 498–504.
- 14 Nyilas S, Schlegtehdal A, Yammine S, *et al.* Further evidence for an association between LCI and FEV₁ in patients with PCD. *Thorax* 2015; 70: 896.
- 15 Mayer-Hamblett N, Ramsey BW, Kronmal RA. Advancing outcome measures for the new era of drug development in cystic fibrosis. *Proc Am Thorac Soc* 2007; 4: 370–377.
- 16 Lucas JS, Barbato A, Collins SA, *et al.* European Respiratory Society guidelines for the diagnosis of primary ciliary dyskinesia. *Eur Respir J* 2017; 49: 1601090.
- 17 Munn Z, Peters MDJ, Stern C, *et al.* Systematic review or scoping review? Guidance for authors when choosing between a systematic or scoping review approach. *BMC Med Res Methodol* 2018; 18: 143.
- 18 Arksey H, O'Malley L. Scoping studies: towards a methodological framework. *Int J Soc Res Methodol* 2005; 8: 19–32.
- 19 Tricco AC, Lillie E, Zarin W, *et al.* PRISMA extension for scoping reviews (PRISMA-ScR): checklist and explanation. *Ann Intern Med* 2018; 169: 467–473.
- 20 Davis SD, Ferkol TW, Rosenfeld M, *et al.* Clinical features of childhood primary ciliary dyskinesia by genotype and ultrastructural phenotype. *Am J Respir Crit Care Med* 2015; 191: 316–324.
- 21 Davis SD, Rosenfeld M, Lee HS, *et al.* Primary ciliary dyskinesia: longitudinal study of lung disease by ultrastructure defect and genotype. *Am J Respir Crit Care Med* 2019; 199: 190–198.
- 22 Ellerman A, Bisgaard H. Longitudinal study of lung function in a cohort of primary ciliary dyskinesia. *Eur Respir J* 1997; 10: 2376–2379.
- 23 Fuger M, Aupiais C, Thouvenin G, *et al.* Gas exchanges in children with cystic fibrosis or primary ciliary dyskinesia: a retrospective study. *Respir Physiol Neurobiol* 2018; 251: 1–7.
- 24 Gokdemir Y, Karadag-Saygi E, Erdem E, *et al.* Comparison of conventional pulmonary rehabilitation and high-frequency chest wall oscillation in primary ciliary dyskinesia. *Pediatr Pulmonol* 2014; 49: 611–616.

- 25 Halbeisen FS, Goutaki M, Spycher BD, *et al.* Lung function in patients with primary ciliary dyskinesia: an iPCD Cohort study. *Eur Respir J* 2018; 52: 1801040.
- 26 Hellinckx J, Demedts M, De Boeck K. Primary ciliary dyskinesia: evolution of pulmonary function. *Eur J Pediatr* 1998; 157: 422–426.
- 27 Koh YY, Park Y, Jeong JH, *et al.* The effect of regular salbutamol on lung function and bronchial responsiveness in patients with primary ciliary dyskinesia. *Chest* 2000; 117: 427–433.
- 28 Lopes AJ, Camilo GB, de Menezes SL, *et al.* Impact of different etiologies of bronchiectasis on the pulmonary function tests. *Clin Med Res* 2015; 13: 12–19.
- 29 Maglione M, Montella S, Mirra V, *et al.* Long-term assessment of quality of life in primary ciliary dyskinesia: time for new tools? *Chest* 2014; 146: e232–e233.
- 30 Marthin JK, Petersen N, Skovgaard LT, *et al.* Lung function in patients with primary ciliary dyskinesia: a cross-sectional and 3-decade longitudinal study. *Am J Respir Crit Care Med* 2010; 181: 1262–1268.
- 31 Oliveira C, Padilla A, Martínez-García MA, *et al.* Etiology of bronchiectasis in a cohort of 2047 patients. An analysis of the Spanish historical bronchiectasis registry. *Arch Bronconeumol* 2017; 53: 366–374.
- 32 Phillips GE, Thomas S, Heather S, *et al.* Airway response of children with primary ciliary dyskinesia to exercise and β_2 -agonist challenge. *Eur Respir J* 1998; 11: 1389–1391.
- 33 Pifferi M, Bush A, Pioggia G, *et al.* Evaluation of pulmonary disease using static lung volumes in primary ciliary dyskinesia. *Thorax* 2012; 67: 993–999.
- 34 Shah A, Shoemark A, MacNeill SJ, *et al.* A longitudinal study characterising a large adult primary ciliary dyskinesia population. *Eur Respir J* 2016; 48: 441–450.
- 35 Sunther M, Bush A, Hogg C, *et al.* Recovery of baseline lung function after pulmonary exacerbation in children with primary ciliary dyskinesia. *Pediatr Pulmonol* 2016; 51: 1362–1366.
- 36 Tamalet A, Clement A, Roudot-Thoraval F, *et al.* Abnormal central complex is a marker of severity in the presence of partial ciliary defect. *Pediatrics* 2001; 108: E86.
- 37 Vallet C, Escudier E, Roudot-Thoraval F, *et al.* Primary ciliary dyskinesia presentation in 60 children according to ciliary ultrastructure. *Eur J Pediatr* 2013; 172: 1053–1060.
- 38 Videbaek K, Buchvald F, Holgersen MG, *et al.* The impact of mannose-binding lectin polymorphisms on lung function in primary ciliary dyskinesia. *Pediatr Pulmonol* 2019; 54: 1182–1189.
- 39 Ahmad F, Irving S, Alton E, *et al.* Multiple breath washouts in children can be shortened without compromising quality. *Eur Respir J* 2015; 46: 1814–1816.
- 40 Anagnostopoulou P, Kranz N, Wolfensberger J, *et al.* Comparison of different analysis algorithms to calculate multiple-breath washout outcomes. *ERJ Open Res* 2018; 4: 00021-02017.
- 41 Green K, Buchvald FF, Marthin JK, *et al.* Ventilation inhomogeneity in children with primary ciliary dyskinesia. *Thorax* 2012; 67: 49–53.
- 42 Green K, Ejlertsen JS, Madsen A, *et al.* Abbreviation modalities of nitrogen multiple-breath washout tests in school children with obstructed lung disease. *Pediatr Pulmonol* 2016; 51: 624–632.
- 43 Irving SJ, Ives A, Davies G, *et al.* Lung clearance index and high-resolution computed tomography scores in primary ciliary dyskinesia. *Am J Respir Crit Care Med* 2013; 188: 545–549.
- 44 Irving S, Carr S, Hogg C, *et al.* Lung clearance index (LCI) is stable in most primary ciliary dyskinesia (PCD) patients managed in a specialist centre: a pilot study. *Lung* 2017; 195: 441–443.
- 45 Irving S, Dixon M, Fassad MR, *et al.* Primary ciliary dyskinesia due to microtubular defects is associated with worse lung clearance index. *Lung* 2018; 196: 231–238.
- 46 Kobbernagel HE, Green K, Ring AM, *et al.* One-year evolution and variability in multiple-breath washout indices in children and young adults with primary ciliary dyskinesia. *Eur Clin Respir J* 2019; 6: 1591841.
- 47 Koucký V, Uhlík J, Hoňková L, *et al.* Ventilation inhomogeneity and bronchial basement membrane changes in chronic neutrophilic airway inflammation. *Chest* 2020; 157: 779–789.
- 48 Nyilas S, Schlegteal A, Singer F, *et al.* Alternative inert gas washout outcomes in patients with primary ciliary dyskinesia. *Eur Respir J* 2017; 49: 1600466.
- 49 Nyilas S, Bauman G, Pusterla O, *et al.* Structural and functional lung impairment in primary ciliary dyskinesia. Assessment with magnetic resonance imaging and multiple breath washout in comparison to spirometry. *Ann Am Thorac Soc* 2018; 15: 1434–1442.
- 50 Smith LJ, West N, Hughes D, *et al.* Imaging lung function abnormalities in primary ciliary dyskinesia using hyperpolarized gas ventilation MRI. *Ann Am Thorac Soc* 2018; 15: 1487–1490.
- 51 Cohen-Cyberknob M, Simanovsky N, Hiller N, *et al.* Differences in disease expression between primary ciliary dyskinesia and cystic fibrosis with and without pancreatic insufficiency. *Chest* 2014; 145: 738–744.
- 52 Dettmer S, Ringshausen F, Vogel-Claussen J, *et al.* Computed tomography in adult patients with primary ciliary dyskinesia: typical imaging findings. *PLoS One* 2018; 13: e0191457.
- 53 Hoang-Thi TN, Revel MP, Burgel PR, *et al.* Automated computed tomographic scoring of lung disease in adults with primary ciliary dyskinesia. *BMC Pulm Med* 2018; 18: 194.
- 54 Jain K, Padley SP, Goldstraw EJ, *et al.* Primary ciliary dyskinesia in the paediatric population: range and severity of radiological findings in a cohort of patients receiving tertiary care. *Clin Radiol* 2007; 62: 986–993.

- 55 Kennedy MP, Noone PG, Leigh MW, *et al.* High-resolution CT of patients with primary ciliary dyskinesia. *AJR Am J Roentgenol* 2007; 188: 1232–1238.
- 56 Li AM, Sonnappa S, Lex C, *et al.* Non-CF bronchiectasis: does knowing the aetiology lead to changes in management? *Eur Respir J* 2005; 26: 8–14.
- 57 Maglione M, Bush A, Montella S, *et al.* Progression of lung disease in primary ciliary dyskinesia: is spirometry less accurate than CT? *Pediatr Pulmonol* 2012; 47: 498–504.
- 58 Maglione M, Montella S, Mollica C, *et al.* Lung structure and function similarities between primary ciliary dyskinesia and mild cystic fibrosis: a pilot study. *Ital J Pediatr* 2017; 43: 34.
- 59 Magnin ML, Cros P, Beydon N, *et al.* Longitudinal lung function and structural changes in children with primary ciliary dyskinesia. *Pediatr Pulmonol* 2012; 47: 816–825.
- 60 Montella S, Santamaria F, Salvatore M, *et al.* Lung disease assessment in primary ciliary dyskinesia: a comparison between chest high-field magnetic resonance imaging and high-resolution computed tomography findings. *Ital J Pediatr* 2009; 35: 24.
- 61 Montella S, Santamaria F, Salvatore M, *et al.* Assessment of chest high-field magnetic resonance imaging in children and young adults with noncystic fibrosis chronic lung disease: comparison to high-resolution computed tomography and correlation with pulmonary function. *Invest Radiol* 2009; 44: 532–538.
- 62 Santamaria F, Montella S, Tiddens H, *et al.* Structural and functional lung disease in primary ciliary dyskinesia. *Chest* 2008; 134: 351–357.
- 63 Tadd K, Morgan L, Rosenow T, *et al.* CF derived scoring systems do not fully describe the range of structural changes seen on CT scans in PCD. *Pediatr Pulmonol* 2019; 54: 471–477.
- 64 Alanin MC, Nielsen KG, von Buchwald C, *et al.* A longitudinal study of lung bacterial pathogens in patients with primary ciliary dyskinesia. *Clin Microbiol Infect* 2015; 21: 1093.
- 65 Cohen-Cymerknoh M, Weigert N, Gileles-Hillel A, *et al.* Clinical impact of *Pseudomonas aeruginosa* colonization in patients with primary ciliary dyskinesia. *Respir Med* 2017; 131: 241–246.
- 66 Roden L, Görlich D, Omran H, *et al.* A retrospective analysis of the pathogens in the airways of patients with primary ciliary dyskinesia. *Respir Med* 2019; 156: 69–77.
- 67 Rogers GB, Carroll MP, Zain NM, *et al.* Complexity, temporal stability, and clinical correlates of airway bacterial community composition in primary ciliary dyskinesia. *J Clin Microbiol* 2013; 51: 4029–4035.
- 68 Goutaki M, Halbeisen FS, Spycher BD, *et al.* Growth and nutritional status, and their association with lung function: a study from the international Primary Ciliary Dyskinesia Cohort. *Eur Respir J* 2017; 50: 4170659.
- 69 Svobodová T, Djakow J, Zemková D, *et al.* Impaired growth during childhood in patients with primary ciliary dyskinesia. *Int J Endocrinol* 2013; 2013: 731423.
- 70 Alanin MC, Aanaes K, Høiby N, *et al.* Sinus surgery can improve quality of life, lung infections, and lung function in patients with primary ciliary dyskinesia. *Int Forum Allergy Rhinol* 2017; 7: 240–247.
- 71 Behan L, Leigh MW, Dell SD, *et al.* Validation of a health-related quality of life instrument for primary ciliary dyskinesia (QOL-PCD). *Thorax* 2017; 72: 832–839.
- 72 Carotenuto M, Esposito M, Di Pasquale F, *et al.* Psychological, cognitive and maternal stress assessment in children with primary ciliary dyskinesia. *World J Pediatr* 2013; 9: 312–317.
- 73 Ioannou P, Kouis P, Kakkoura MG, *et al.* Health related quality of life in adult primary ciliary dyskinesia patients in Cyprus: development and validation of the Greek version of the QOL-PCD questionnaire. *Health Qual Life Outcomes* 2020; 18: 105.
- 74 Keniş Coşkun Ö, Gençer Atalay K, Erdem E, *et al.* Caregiver burden in children with cystic fibrosis and primary ciliary dyskinesia. *Pediatr Pulmonol* 2019; 54: 1936–1940.
- 75 Maglione M, Bush A, Nielsen KG, *et al.* Multicenter analysis of body mass index, lung function, and sputum microbiology in primary ciliary dyskinesia. *Pediatr Pulmonol* 2014; 49: 1243–1250.
- 76 McManus IC, Mitchison HM, Chung EM, *et al.* Primary ciliary dyskinesia (Siewert's/Kartagener's syndrome): respiratory symptoms and psycho-social impact. *BMC Pulm Med* 2003; 3: 4.
- 77 McManus IC, Stubbings GF, Martin N. Stigmatization, physical illness and mental health in primary ciliary dyskinesia. *J Health Psychol* 2006; 11: 467–482.
- 78 Pifferi M, Bush A, Di Cicco M, *et al.* Health-related quality of life and unmet needs in patients with primary ciliary dyskinesia. *Eur Respir J* 2010; 35: 787–794.
- 79 Valero-Moreno S, Castillo-Corullón S, Montoya-Castilla I, *et al.* Primary ciliary dyskinesia and psychological well-being in adolescence. *PLoS One* 2020; 15: e0227888.
- 80 Whalley S, McManus IC. Living with primary ciliary dyskinesia: a prospective qualitative study of knowledge sharing, symptom concealment, embarrassment, mistrust, and stigma. *BMC Pulm Med* 2006; 6: 25.
- 81 Zengin Akkus P, Gharibzadeh Hizal M, Ilter Bahadır E, *et al.* Developmental and behavioral problems in preschool-aged primary ciliary dyskinesia patients. *Eur J Pediatr* 2019; 178: 995–1003.
- 82 Cohen-Cymerknoh M, Atia O, Gileles-Hillel A, *et al.* Sleep disorders in patients with primary ciliary dyskinesia, cystic fibrosis with and without pancreatic insufficiency. *Respir Med* 2019; 151: 96–101.
- 83 Oktem S, Karadag B, Erdem E, *et al.* Sleep disordered breathing in patients with primary ciliary dyskinesia. *Pediatr Pulmonol* 2013; 48: 897–903.

- 84 Santamaria F, Esposito M, Montella S, *et al.* Sleep disordered breathing and airway disease in primary ciliary dyskinesia. *Respirology* 2014; 19: 570–575.
- 85 Şişmanlar Eyüboğlu T, Aslan AT, Ceylan A, *et al.* Neurocognitive disorders and sleep in children with primary ciliary dyskinesia. *Pediatr Pulmonol* 2018; 53: 1436–1441.
- 86 Bush A, Payne D, Pike S, *et al.* Mucus properties in children with primary ciliary dyskinesia: comparison with cystic fibrosis. *Chest* 2006; 129: 118–123.
- 87 Cockx M, Gouwy M, Godding V, *et al.* Neutrophils from patients with primary ciliary dyskinesia display reduced chemotaxis to CXCR2 ligands. *Front Immunol* 2017; 8: 1126.
- 88 Cockx M, Gouwy M, Ruytinx P, *et al.* Monocytes from patients with primary ciliary dyskinesia show enhanced inflammatory properties and produce higher levels of pro-inflammatory cytokines. *Sci Rep* 2017; 7: 14657.
- 89 Paff T, Daniels JM, Weersink EJ, *et al.* A randomised controlled trial on the effect of inhaled hypertonic saline on quality of life in primary ciliary dyskinesia. *Eur Respir J* 2017; 49: 1601770.
- 90 Ratjen F, Waters V, Klingel M, *et al.* Changes in airway inflammation during pulmonary exacerbations in patients with cystic fibrosis and primary ciliary dyskinesia. *Eur Respir J* 2016; 47: 829–836.
- 91 Zihlif N, Paraskakis E, Tripoli C, *et al.* Markers of airway inflammation in primary ciliary dyskinesia studied using exhaled breath condensate. *Pediatr Pulmonol* 2006; 41: 509–514.
- 92 Kobbernagel HE, Buchvald FF, Haarman EG, *et al.* Efficacy and safety of azithromycin maintenance therapy in primary ciliary dyskinesia (BESTCILIA): a multicentre, double-blind, randomised, placebo-controlled phase 3 trial. *Lancet Respir Med* 2020; 8: 493–505.
- 93 Piatti G, De Santi MM, Farolfi A, *et al.* Exacerbations and *Pseudomonas aeruginosa* colonization are associated with altered lung structure and function in primary ciliary dyskinesia. *BMC Pediatr* 2020; 20: 158.
- 94 Loomba RS, Danduran M, Nielsen KG, *et al.* Cardiopulmonary exercise testing in fontan patients with and without isomerism (heterotaxy) as compared to patients with primary ciliary dyskinesia and subjects with structurally normal hearts. *Pediatr Cardiol* 2017; 38: 410–417.
- 95 Madsen A, Green K, Buchvald F, *et al.* Aerobic fitness in children and young adults with primary ciliary dyskinesia. *PLoS One* 2013; 8: e71409.
- 96 Ring AM, Buchvald FF, Holgersen MG, *et al.* Fitness and lung function in children with primary ciliary dyskinesia and cystic fibrosis. *Respir Med* 2018; 139: 79–85.
- 97 Simsek S, Inal-Ince D, Cakmak A, *et al.* Reduced anaerobic and aerobic performance in children with primary ciliary dyskinesia. *Eur J Pediatr* 2018; 177: 765–773.
- 98 Valerio G, Giallauria F, Montella S, *et al.* Cardiopulmonary assessment in primary ciliary dyskinesia. *Eur J Clin Invest* 2012; 42: 617–622.
- 99 Wells GD, Wilkes DL, Schneiderman JE, *et al.* Skeletal muscle metabolism in cystic fibrosis and primary ciliary dyskinesia. *Pediatr Res* 2011; 69: 40–45.
- 100 Joensen O, Paff T, Haarman EG, *et al.* Exhaled breath analysis using electronic nose in cystic fibrosis and primary ciliary dyskinesia patients with chronic pulmonary infections. *PLoS One* 2014; 9: e115584.
- 101 Kawakami M, Hattori Y, Nakamura S. Reflection of structural abnormality in the axoneme of respiratory cilia in the clinical features of immotile cilia syndrome. *Intern Med* 1996; 35: 617–623.
- 102 Kennedy MP, Noone PG, Carson J, *et al.* Calcium stone lithoptysis in primary ciliary dyskinesia. *Respir Med* 2007; 101: 76–83.
- 103 Marino LV, Harris A, Johnstone C, *et al.* Characterising the nutritional status of children with primary ciliary dyskinesia. *Clin Nutr* 2019; 38: 2127–2135.
- 104 Mirra V, Caffarelli C, Maglione M, *et al.* Hypovitaminosis D: a novel finding in primary ciliary dyskinesia. *Ital J Pediatr* 2015; 41: 14.
- 105 Montuschi P, Paris D, Montella S, *et al.* Nuclear magnetic resonance-based metabolomics discriminates primary ciliary dyskinesia from cystic fibrosis. *Am J Respir Crit Care Med* 2014; 190: 229–233.
- 106 Noone PG, Bennett WD, Regnis JA, *et al.* Effect of aerosolized uridine-5'-triphosphate on airway clearance with cough in patients with primary ciliary dyskinesia. *Am J Respir Crit Care Med* 1999; 160: 144–149.
- 107 Paff T, van der Schee MP, Daniels JM, *et al.* Exhaled molecular profiles in the assessment of cystic fibrosis and primary ciliary dyskinesia. *J Cyst Fibros* 2013; 12: 454–460.
- 108 Pifferi M, Bush A, Caramella D, *et al.* Matrix metalloproteinases and airway remodeling and function in primary ciliary dyskinesia. *Respir Med* 2017; 124: 49–56.
- 109 Shoemark A, Wilson R. Bronchial and peripheral airway nitric oxide in primary ciliary dyskinesia and bronchiectasis. *Respir Med* 2009; 103: 700–706.
- 110 Smit HJ, Schreurs AJ, Van den Bosch JM, *et al.* Is resection of bronchiectasis beneficial in patients with primary ciliary dyskinesia? *Chest* 1996; 109: 1541–1544.
- 111 Zihlif N, Paraskakis E, Lex C, *et al.* Correlation between cough frequency and airway inflammation in children with primary ciliary dyskinesia. *Pediatr Pulmonol* 2005; 39: 551–557.
- 112 Abitbul R, Amirav I, Blau H, *et al.* Primary ciliary dyskinesia in Israel: prevalence, clinical features, current diagnosis and management practices. *Respir Med* 2016; 119: 41–47.

- 113 Boon M, Smits A, Cuppens H, *et al.* Primary ciliary dyskinesia: critical evaluation of clinical symptoms and diagnosis in patients with normal and abnormal ultrastructure. *Orphanet J Rare Dis* 2014; 9: 11.
- 114 Eden E, Choate R, Barker A, *et al.* The clinical features of bronchiectasis associated with alpha-1 antitrypsin deficiency, common variable immunodeficiency and primary ciliary dyskinesia – results from the U.S. *Chronic Obstr Pulm Dis* 2019; 6: 145–153.
- 115 Emiralioglu N, Taşkıran EZ, Koşukcu C, *et al.* Genotype and phenotype evaluation of patients with primary ciliary dyskinesia: first results from Turkey. *Pediatr Pulmonol* 2020; 55: 383–393.
- 116 Frija-Masson J, Bassinet L, Honoré I, *et al.* Clinical characteristics, functional respiratory decline and follow-up in adult patients with primary ciliary dyskinesia. *Thorax* 2017; 72: 154–160.
- 117 Knowles MR, Ostrowski LE, Leigh MW, *et al.* Mutations in RSPH1 cause primary ciliary dyskinesia with a unique clinical and ciliary phenotype. *Am J Respir Crit Care Med* 2014; 189: 707–717.
- 118 Noone PG, Leigh MW, Sannuti A, *et al.* Primary ciliary dyskinesia: diagnostic and phenotypic features. *Am J Respir Crit Care Med* 2004; 169: 459–467.
- 119 Pifferi M, Bush A, Michelucci A, *et al.* Mannose-binding lectin 2 gene polymorphism and lung damage in primary ciliary dyskinesia. *Pediatr Pulmonol* 2015; 50: 179–186.
- 120 Yiallourous PK, Kouis P, Middleton N, *et al.* Clinical features of primary ciliary dyskinesia in Cyprus with emphasis on lobectomized patients. *Respir Med* 2015; 109: 347–356.
- 121 Goutaki M, Maurer E, Halbeisen FS, *et al.* The international primary ciliary dyskinesia cohort (iPCD Cohort): methods and first results. *Eur Respir J* 2017; 49: 1601181.
- 122 Miller MR, Hankinson J, Brusasco V, *et al.* Standardisation of spirometry. *Eur Respir J* 2005; 26: 319–338.
- 123 Quanjer PH, Stanojevic S, Cole TJ, *et al.* Multi-ethnic reference values for spirometry for the 3–95-year age range: the Global Lung Function 2012 equations. *Eur Respir J* 2012; 40: 1324–1343.
- 124 Chrispin AR, Norman AP. The systematic evaluation of the chest radiograph in cystic fibrosis. *Pediatr Radiol* 1974; 2: 101–105.
- 125 Brody AS, Kosorok MR, Li Z, *et al.* Reproducibility of a scoring system for computed tomography scanning in cystic fibrosis. *J Thorac Imaging* 2006; 21: 14–21.
- 126 Bhalla M, Turcios N, Aponte V, *et al.* Cystic fibrosis: scoring system with thin-section CT. *Radiology* 1991; 179: 783–788.
- 127 Helbich TH, Heinz-Peer G, Eichler I, *et al.* Cystic fibrosis: CT assessment of lung involvement in children and adults. *Radiology* 1999; 213: 537–544.
- 128 Dell SD, Leigh MW, Lucas JS, *et al.* Primary ciliary dyskinesia: first health-related quality-of-life measures for pediatric patients. *Ann Am Thorac Soc* 2016; 13: 1726–1735.
- 129 Lucas JS, Behan L, Dunn Galvin A, *et al.* A quality-of-life measure for adults with primary ciliary dyskinesia: QOL-PCD. *Eur Respir J* 2015; 46: 375–383.
- 130 Lucas JS, Gahleitner F, Amorim A, *et al.* Pulmonary exacerbations in patients with primary ciliary dyskinesia: an expert consensus definition for use in clinical trials. *ERJ Open Res* 2019; 5: 00147-2018.
- 131 Halbeisen FS, Jose A, de Jong C, *et al.* Spirometric indices in primary ciliary dyskinesia: systematic review and meta-analysis. *ERJ Open Res* 2019; 5: 00231-2018.
- 132 Halbeisen F, Hogg C, Alanin MC, *et al.* Proceedings of the 2nd BEAT-PCD conference and 3rd PCD training school: part 1. *BMC Proc* 2018; 12: Suppl 2, 1.
- 133 Farley H, Rubbo B, Bukowy-Bieryllo Z, *et al.* Proceedings of the 3rd BEAT-PCD conference and 4th PCD training school. *BMC Proc* 2018; 12: Suppl 16, 64.
- 134 Stanojevic S, Stocks J, Bountziouka V, *et al.* The impact of switching to the new global lung function initiative equations on spirometry results in the UK CF Registry. *J Cyst Fibros* 2014; 13: 319–327.
- 135 Robinson P, Morgan L. Bronchiectasis in PCD looks different to CF on CT scan. *Multidiscip Respir Med* 2018; 13: Suppl. 1, 24.
- 136 Campbell-Washburn AE, Ramasawmy R, Restivo MC, *et al.* Opportunities in interventional and diagnostic imaging by using high-performance low-field-strength MRI. *Radiology* 2019; 293: 384–393.
- 137 Edelman RR, Hatabu H, Tadamura E, *et al.* Noninvasive assessment of regional ventilation in the human lung using oxygen-enhanced magnetic resonance imaging. *Nat Med* 1996; 2: 1236–1239.
- 138 Werner C, Wallmeier J, Raidt J, *et al.* An international patient-registry for primary ciliary dyskinesia. *Pediatr Pulmonol* 2014; 49: S71–S72.
- 139 Goutaki M, Eich MO, Halbeisen FS, *et al.* The Swiss Primary Ciliary Dyskinesia registry: objectives, methods and first results. *Swiss Med Wkly* 2019; 149: w20004.
- 140 Goutaki M, Halbeisen FS, Kuehni CE. The time is right for an international PCD disease registry: insight and ongoing research activities. *Eur Respir J* 2017; 49: 1700357.
- 141 Rubbo B, Behan L, Dehlink E, *et al.* Proceedings of the COST action BM1407 inaugural conference BEAT-PCD: translational research in primary ciliary dyskinesia – bench, bedside, and population perspectives. *BMC Proc* 2016; 10: Suppl. 9, 66.
- 142 Gardner LE, Horton KL, Shoemark A, *et al.* Proceedings of the 4th BEAT-PCD conference and 5th PCD training school. *BMC Proc* 2020; 14: Suppl. 8, 7.

- 143 Rubbo B, Lucas JS. Clinical care for primary ciliary dyskinesia: current challenges and future directions. *Eur Respir Rev* 2017; 26: 170023.
- 144 Amirav I, Roduta Roberts M, Mussaffi H, et al. Collecting clinical data in primary ciliary dyskinesia – challenges and opportunities. *F1000Res* 2016; 5: 2031.
- 145 Goutaki M, Papon J-F, Boon M, et al. Standardised clinical data from patients with primary ciliary dyskinesia: FOLLOW-PCD. *ERJ Open Res* 2020; 6: 00237-2019.
- 146 Kuo W, Ciet P, Tiddens HA, et al. Monitoring cystic fibrosis lung disease by computed tomography. Radiation risk in perspective. *Am J Respir Crit Care Med* 2014; 189: 1328–1336.
- 147 Kubo T, Ohno Y, Kauczor HU, et al. Radiation dose reduction in chest CT – review of available options. *Eur J Radiol* 2014; 83: 1953–1961.
- 148 Behan L, Rubbo B, Lucas JS, et al. The patient's experience of primary ciliary dyskinesia: a systematic review. *Qual Life Res* 2017; 26: 2265–2285.
- 149 Emiralioğlu N, Karadağ B, Özçelik HU. Quality of life questionnaire for Turkish patients with primary ciliary dyskinesia. *Turk Thorac J* 2017; 18: 19–22.
- 150 Queiroz APL, Athanazio RA, Olm MAK, et al. Translation of the quality-of-life measure for adults with primary ciliary dyskinesia and its application in patients in Brazil. *J Bras Pneumol* 2019; 45: e20170358.