

## Early View

Original research article

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# **A practice test and selection of a core set of outcome-based quality indicators in Dutch primary care physical therapy for patients with COPD: a cohort study**

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## Take home message

The major finding is that all participants in the focus groups accepted the quality indicators as a quality improvement tool based on their perceived added value and selected a core set of seven outcome-based quality indicators for patients with COPD.

**Keywords:** Outcome-based quality indicators, COPD, Primary care, Physical therapy

## **Abstract**

**Aim** to estimate the comparability and discriminability of outcome-based quality indicators by performing a practice test in Dutch physical therapy primary care, and to select a core set of outcome-based quality indicators that are well-accepted by physical therapists based on their perceived added value as a quality improvement tool.

**Methods** First, a list of potential quality indicators was defined, followed by the determination of the comparability (case-mix adjusted multi-level analysis) and discriminability (intraclass correlation coefficient (ICC)). Second, focus group meetings were conducted with stakeholders (physical therapists and senior researchers) to select a core set of quality indicators.

**Results** Overall, 229 physical therapists from 137 practices provided 2651 treatment episodes.

Comparability: in 10 of the 11 case-mix adjusted models, the ICC increased compared with the intercept-only model. Discriminability: the ICC ranged between 0.01 and 0.34, with five of the 11 ICCs being > 0.10. The majority of physical therapists in each focus group preferred the inclusion of seven quality indicators in the core set, including three process and four outcome indicators based upon the Six-Minute-Walk-Test (6MWT), the Clinical COPD Questionnaire (CCQ), and the determination of quadriceps strength using a hand-held dynamometer (HHD).

**Conclusion** This is the first study that describes the comparability and discriminability of the outcome-based quality indicators selected for patients with COPD treated in primary care physical therapy practices. Future research should focus on increasing data collection in daily practice and on the development of tangible methods to use as the core set of a quality improvement tool.

## Background

The routine use of outcome measures can play an important role in improving healthcare quality;(1) for example, they can enable the comparison of providers' performances to stimulate improvement initiatives.(2) A fundamental prerequisite of the use of outcome measures is the collection, aggregation, and comprehensive understandable presentation of data.(1) Using quality indicators may stimulate the routine data collection of patient-reported outcome measures (PROMs) by healthcare providers. Quality indicators can be used on an aggregated level to show changes in clinical practice over time.(1, 3, 4)

In a previous study, we developed a standard set of outcome domains and associated measures, including PROMs and physical performance measures, for patients with chronic obstructive pulmonary diseases (COPD) in primary care physical therapy practice.(4) It is still unclear which quality indicators can be selected from the standard set and which quality indicators have perceived added value as quality improvement tools for such patients, however.

In this study we focused on outcome-based quality indicators chosen from the standard set of PROMs and physical performance measures for patients with COPD.(5) PROMs are often combined with other clinician-assessed, impairment-based or physical performance-based measures, such as the Six-Minute Walk Test (6MWT), to provide a more complete interpretation of patient outcomes.(6)

Currently, most strategies for the development of quality indicators are based on an evidence-based consensus between stakeholders in procedures, such as the RAND/UCLA Delphi procedure.(7-9) This is true for recommendations for clinical practice guidelines too, such as the recently published Dutch clinical practice guideline (CPG) for primary care physical therapists treating patients with COPD,(10) and can provide an important basis for the development of quality indicators.(11) In addition, a practice test, including the collection of real-world data prior to selection, is an essential step for the evaluation of the comparability, discriminability, and feasibility of potential quality indicators in daily practice.(9, 12) A practice test can support the usefulness and feasibility of quality indicators in daily practice and gain insight into the psychometric properties of outcome-based quality indicators. (3, 11, 13) Although to our knowledge, no specific definition of a practice test is reported in previous research, there are several examples of using a practice test in development of quality indicators. (11, 13, 14) Such an example is the study of Meerhoff et al. 2021 in which a practice test was conducted to explore the reliability, validity and

discriminability of patient reported outcomes for the development of quality indicators in patients with non-specific low back pain.(13) We defined comparability as the extent to which the quality indicator is comparable between practices, and discriminability as the extent to which the quality indicator is able to discriminate between practices.

Here, we develop outcome-based quality indicators for patients with COPD in physical therapy primary care. The aims of this study are therefore:

- a) To estimate the comparability and discriminability of outcome-based quality indicators included in a previously selected standard set of measures;
- b) To select a core set of outcome-based quality indicators that is well-accepted by physical therapists based on the perceived added value of this core set as a quality improvement tool.

## **Method**

### **Design**

In this mixed methods study, we used a sequential explanatory design taking a previously selected standard set of outcome domains and measures as the basis for defining and selecting a core set of quality indicators. The standard set was developed in two consecutive steps between February 2018 and April 2020,(4) and was registered on the Core Outcome Measures in Effectiveness Trials (COMET) website.(15) In phase 1 of the present study, potential quality indicators were defined, and we estimated their comparability and discriminability with prospectively collected cohort data between February 2018 and December 2019. To enhance the comparability, we adjusted for differences in patient characteristics using a case-mix correction. Furthermore, we calculated whether the quality indicator was able to discriminate the outcomes of patients between practices and could therefore be used as an instrument for quality improvement. In phase 2, we explored the perceived added value of the indicators in focus group meetings with physical therapists. We then actively involved the participants in the selection of a core set of quality indicators.

### **Setting**

A total of 229 Dutch physical therapists working in 137 primary care practices collected the treatment

outcomes of patients with COPD. All participants in the project were recruited via stakeholder organizations in Dutch primary physical therapy care. Participating physical therapists were instructed to treat their patients according to Dutch clinical guideline recommendations for patients with COPD. (16) We only measured outcomes of the treatment; the physical therapists individually decided which treatment was needed for their patients. All procedures were conducted according to the Declaration of Helsinki and approved by the Medical Ethical Committee of Radboud university medical center (Registration # 2019-5455). The STROBE-checklist was used to report the current study. (17) Furthermore, a framework with tools to support the selection and implementation of patient-reported outcome measures was used as guidance for conduction of this study.(3)

### **Data collection**

Data on the treatment outcomes were anonymously collected through electronic health records (EHRs) via three databases, the national data registry (LDK) of the Association for Quality in Physical Therapy (SKF), the national data registry (LDF) of the Royal Dutch Society for Physical Therapy (KNGF), and the database of Spot On Medics (SOM), which is one of the EHR vendors. The EHRs uploaded to the national registries only contain anonymized data. Furthermore, to ensure the uniformity of the provided data, all data in the registries were collected based on predefined technical specifications.(18) Informed consent was obtained and registered in the EHR from all participating patients included in the current study.

### Outcome domains and measures

The outcome domains in the standard set were based on the consensus between stakeholders (patients, physical therapists, policy makers, researchers, and health insurers),(4) and on the recommendations in the Dutch CPG for the physical therapy treatment of patients with COPD.(16) After the development of the standard set, the KNGF published an update for this CPG,(10) in which the suggested outcome domains and associated measures to evaluate physical therapy treatment are in line with the outcome domains from the developed standard set.

The standard set of outcome measures consisted of three mandatory measures for the total population, two conditional measures that depended on the treatment goal, and two exploratory measures that were used as pilot in a small subgroup. In the current study, only the mandatory and conditional

measures were used for the development of quality indicators, since the exploratory measure was only used in a small subgroup of practices. The three mandatory measures for all patients with COPD were the 6MWT for measuring physical capacity, the Clinical COPD Questionnaire (CCQ) for measuring health-related quality of life, and the Global Perceived Effect - Dutch Version (GPE-DV) for measuring the perceived effect. The two conditional measures were the hand-held dynamometer (with Microfet™) for measuring quadriceps strength, and the Medical Research Council dyspnea scale (MRC) for measuring dyspnea. All measures were completed pre- and post-treatment to monitor the changes in outcomes over time, except the GPE-DV, which was only measured after the treatment. For a description of each measure and the measurement protocol, see Appendix A. All physical therapists followed a specific protocol to standardize the testing procedure.(19)

#### Inclusion and exclusion criteria

All patients with COPD (GOLD I–IV), as diagnosed by a medical doctor, who received physical therapy in one of the participating primary care practices between February 2018 and December 2019 were included. Participating physical therapy practices were instructed to collect at least all mandatory and conditional measures from the standard set as presented in the data collection, according to the measurement protocol described in Appendix A. Based on a rule of thumb, a minimum of 30 patients should preferably be included for each practice to allow a valid comparison. (20, 21) However, it was expected that this inclusion requirement could not be reached due to the short inclusion period, and the fact that routine data collection in primary physical therapy care for patients with COPD is relatively new. Therefore, we used a lower threshold and physical therapy practices were excluded from the analysis for a specific quality indicator if they included fewer than 10 patients with COPD

## **Phase 1: Defining quality indicators and estimating their comparability and discriminability.**

#### Defining potential quality indicators

We used national and international standards for defining potential quality indicators.(2, 5, 22, 23) Quality

indicators can be described using mean values and between-relative differences, or quantified and expressed as a proportion in which the numerator describes the number of 'correct' scores and the denominator is the number of persons for which the quality indicator is applicable.(3) See Table 1 for an example of a quality indicator for physical capacity measured with the 6MWT.

**Table 1 Example of a quality indicator monitoring the repeated measurement of the 6MWT**

<b>Definition</b>	<b>The proportion of patients with COPD who underwent physical therapy treatment and who completed the 6MWT pre- and post-treatment to evaluate physical capacity</b>
<b>Rationale</b>	Improvement of physical capacity is an important goal in physical therapy treatment for patients with COPD. Physical capacity is measured with the 6MWT
<b>Numerator</b>	The number of patients who underwent physical therapy treatment and who completed the 6MWT pre- and post-treatment
<b>Denominator</b>	All patients who underwent physical therapy treatment
<b>Specification</b>	Physical capacity is measured in all patients using the 6MWT, a physical performance test where the patients walk for six minutes in a comfortable way
<b>Type of indicator</b>	Process

For each of the five measures in the standard set, we defined four types of quality indicators: 1) by monitoring the process, i.e., whether the outcome was actually measured pre- and post-treatment; 2) by using mean end scores of the outcome, reflecting patient functioning at the end of treatment; 3) by using the mean pre- to post-treatment change in the outcome score, reflecting improvement or decline in the outcome; and 4) by using the minimally clinical important difference (MCID) of the outcome, i.e., the proportion of patients who experienced clinically relevant improvements, stabilizations, or deteriorations (see Box 1 for an example). The change score and the MCID were not defined for the GPE-DV, as this measure was only completed after the treatment. For the MRC, no MCID was defined, since a MCID has not yet been established for the MRC.(24) Hence, in total, we defined 17 potential quality indicators. See Appendix B for an extensive description of each potential quality indicator.

**Box 1: Potential process and outcome quality indicators at the physical therapist or practice level**

**a) Process indicator: proportion of repeated measures**

The proportion of patients with COPD who underwent physical therapy treatment in which a pre- and post-measurement was used.

Example 1: In 60% of the patients, physical capacity was measured pre- and post-treatment with the 6MWT



**b) Outcome indicator: mean end scores**

The mean end score (with 95% confidence intervals (CI)) of patients with COPD after a physical therapy treatment.

Example 2 The mean end score of the health-related quality of life of patients with COPD measured with the CCQ is 2.2 points ( $\pm 0.9$  points)

**c) Outcome indicator: mean change scores**

The mean change score (with 95% CI) of patients with COPD between the pre- and post-physical therapy stages.

Example 3 The mean change score in the symptoms of dyspnea in patients with COPD measured with the MRC is 2.5 points of improvement ( $\pm 1.0$  points)

**d) Outcome indicator: MCID**

The proportion (with 95% CI) of patients with COPD who experienced a MCID improvement between the pre- and post-treatment stages.

Example 4: In 70% ( $\pm 7\%$ ) of the patients, a clinically relevant change in quadriceps strength was reported after treatment, as measured with the HHD.

Abbreviations: 6MWT: Six-Minute Walk Test; CCQ: Clinical COPD Questionnaire; GPE-DV: Global Perceived Effect - Dutch Version; HHD: hand-held dynamometer; MRC: Medical Research Council dyspnea scale; MCID: minimal clinical important difference; CI: confidence intervals.

**Estimating the comparability and discriminability of the quality indicators**

For each measure, at least 30 physical therapy practices needed to be included, this was based on a rule of thumb in multilevel analysis for general calculation. (20, 21) Descriptive statistics were used to determine whether the thresholds for the completeness of the measures were met for estimating the indicator scores. For the analysis, we used measures collected at the beginning and/or end of the treatment. When the treatment episode had not ended, we used the last provided data. The treatment episode is a unique episode of a patient being treated by a physical therapist

### Comparability

In a linear and logistic multi-level analysis, patients were clustered within physical therapy practices to compare the outcomes of the quality indicators between practices.(25) The quality indicators were adjusted for patient characteristics that influence the outcome but were not under the control of the physical therapist or physical therapy practice (so called explanatory variables).(23) Explanatory variables, such as age, gender, and baseline scores for each measure, were used for the adjustment of the multi-level analyses. In the analysis, we started with an intercept-only model that estimates only the intercept and the random variation around the intercept. The inclusion of age,(26, 27) gender,(26) and the baseline score (28, 29) of each measure for the adjustment of a multi-level analysis is common in the field of quality indicator development and the comparison of provider performance.(2, 25-30) Next, all explanatory variables were added to the adjusted model, and the influence of the explanatory variables was evaluated by the amount of the random intercept variance that was explained.(25)

For each physical therapy practice in the case-mix adjusted multi-level analysis, the mean scores were estimated with a 95% confidence interval (CI).

### Discriminability

For the estimation of the variation in the outcomes between physical therapy practices, the intraclass correlation coefficient (ICC) was calculated. The ICC for physical therapy practices was defined by dividing the variance between practices by the summation of the variance between and the variance within physical therapy practices.(25) In multi-level analyses, most ICCs are between .05 and .20, and ICCs  $>.10$  can be interpreted as adequate, indicating that the quality indicator is able to discriminate outcomes between physical therapists or practices.(25, 31) The ICC was also used to compare the intercept-only model with the adjusted model containing the explanatory variables (case-mix).

### Visual representation of indicator scores

To present the mean outcomes for each defined quality indicator of each physical therapy practice in one graph, we used caterpillars plots, as they are found to be user friendly and easy to interpret.(2, 29, 31, 32) We used relative norms by presenting the plots in three colors: blue (95% CI significantly lower than average), purple (no significant 95% CI difference from average), and green (95% CI significantly higher

than average). The plots were used to present the outcomes of the cohort data to the participants in phase 2 of the study.

## **Phase 2: Selecting a core set of quality indicators**

Semi-structured focus group interviews were conducted with purposefully selected participating physical therapists who collected data in phase 1. We also organized one focus group meeting with Dutch senior physical therapists and senior researchers who were members of the development group of the revised Dutch physical therapy guideline for COPD. The senior researchers were asked to comment on the set of quality indicators from a scientific perspective. The senior physical therapists and researchers had at least 10 years of experience in the treatment of and/or research into patients with COPD.

We aimed to conduct four focus group meetings with 6–10 members in every meeting. The primary goal of the focus groups was to reflect on the added value of using the presented indicators in daily practice for quality improvement, and most importantly, to select a core set of quality indicators from the described 17 potential indicators. The focus group meetings were audio recorded and summarized by researcher AV, the summaries of the different focus groups were discussed and interpreted in several meetings with researchers AV (physical therapist and PhD student), SvD (physical therapist and senior researcher), and PvdW (physical therapist and professor of allied health sciences). The identities of the physical therapists were considered confidential; therefore, the answers given by the physical therapists during the interviews and in the survey were processed anonymously. The focus groups were part of the process of reaching consensus on the selection of the core set.

The research members AV, SvD, PvdW, HK (physical therapist and senior researcher), RN (physical therapist and em. professor of allied health sciences) were trained and had experience in conducting qualitative research.

In each focus group meeting, we presented each potential quality indicator in a caterpillar plot, with scores at the levels of the physical therapist and the physical therapy practice, and compared them with the scores for the other practices. The participants interpreted the comparability and discriminability of the potential quality indicators. Finally, we asked participants to select their preferred quality indicators for the core set from the potential quality indicators as described in box 1. During each meeting, the chairman (AV) summed up all the preferred quality indicators and asked the group whether they accepted or

declined the proposed core set. A consensus was reached if >80% of the participants accepted the selection of each quality indicator in the core set.

#### Patient and public involvement

For the development of this standard set we interviewed patients about their perspectives on the selection of patient outcomes. (4, 15) Furthermore, during the conduction of this study a steering committee with representatives from important stakeholders, including the association for patients with COPD Netherlands Patients Federation (Longfonds), advised during the selection process. During the meetings, we discussed the views and perspectives of stakeholders regarding the value and implementation of outcome-based quality indicators for Dutch physical therapy

## Results

### **Phase 1: Estimating the comparability and discriminability of the quality indicators**

#### Descriptive statistics

Table 2 shows descriptive statistics of the included treatment episodes and the number of physical therapists and physical therapy practices who provided the data. The treatment episode is a unique episode of a patient being treated by a physical therapist. The current national data registries cannot detect recurrences of patients over time due to privacy regulations; therefore, the number of unique patients may be lower. Overall, 229 physical therapists from 137 practices provided 4651 treatment episodes of patients with COPD.

**Table 2 Descriptive statistics of the included patients and the number of participating physical therapists and physical therapy practices**

<b>Number of treatment episodes in the dataset</b>	4651
<b>Female patients</b>	2440 (52.5%)
<b>Age, years</b>	67.9 (9.4)

<b>Treatment sessions</b>	49.2 (58.2)
<b>Episode duration, weeks</b>	46.6 (50.3)
<b>Physical therapists who provided the data</b>	229
<b>Physical therapy practices that provided the data</b>	137

Data are presented as means (standard deviation (SD)) or numbers and percentages of the total population.

Table 3 presents the characteristics and unadjusted outcomes of patients with COPD on each measure of the standard set. The number of patients with end scores differed between the measures. Each measure reached the threshold of at least 30 included physical therapy practices that provided  $\geq 10$  cases, except for the HHD, for which only 10 physical therapy practices provided  $\geq 10$  cases and therefore no ICC was calculated. See Table 3 for the number of practices and provided cases that were included in the multi-level analysis.

**Table 3 Descriptive characteristics and unadjusted outcomes of patients with COPD for each measure of the standard set**

	<b>6MWT</b>	<b>CCQ</b>	<b>GPE-DV</b>	<b>HHD</b>	<b>MRC</b>
<b>Female patients</b>	1344 (51.1%)	1786 (52.1%)	636 (50.6%)	223 (51.3%)	1237 (52.7%)
<b>Age, years</b>	67.8 (9.2)	68.1 (9.4)	68.2 (9.4)	68.0 (9.2)	68.2 (9.3)
<b>Treatment episodes with baseline scores</b>	2628	3427	N.A.	435	2348
<b>Treatment episodes with end scores</b>	1822	2408	1256	218	1385
<b>Range of scores on each measure</b>	6–780 m	0–6 points	1–7 points	131–542 Nm	1–5 points
<b>Baseline scores<sup>a</sup></b>	370.8 m (126.3 m)	2.4 points (0.9)	N.A.	284.2 Nm (96.4 Nm)	3.0 points (1.1)
<b>End scores</b>	373.5 m	2.2 points	3.5 points	298.4 Nm	3.0 points

	(130.3 m)	(0.9)	(1.1)	(95.5)	(1.1)
<b>Change (<math>T_{\text{end}}-T_0</math>)</b>	2.7 (86.8)	-0.1 (0.8)	N.A.*	8.4 (50.9)	0.2 (1.3)
<b>MCID improvement</b>	533 (28.7%) <sup>b</sup>	818 (34%) <sup>c</sup>	N.A.*	99 (45.4%) <sup>d</sup>	N.A.**
<b>MCID stabilization</b>	807 (44.3%) <sup>b</sup>	1052 (43.7%) <sup>c</sup>	N.A.*	43 (19.9%) <sup>d</sup>	N.A.**
<b>MCID deterioration</b>	490 (26.9%) <sup>b</sup>	537 (22.3%) <sup>c</sup>	N.A.*	76 (34.7%) <sup>d</sup>	N.A.**
<b>Physical therapists who provided data</b>	145 (63.3%)	202 (88.2%)	117 (51.0%)	46 (20.0%)	168 (46.7%)
<b>Practices that provided data</b>	86 (62.8%)	126 (92%)	72 (52.6%)	28 (20.4%)	107 (78.1%)
<b>Practices that provided <math>\geq 10</math> cases</b>	44 (19.2%)	61 (26.6%)	35 (15.3%)	10 (4.4%)	43 (18.8%)
<b>Patients included in the multi-level analysis</b>	1679 (36.0%)	2201 (47.3%)	1110 (23.8%)	160 (3.4%)	1226 (26.4%)

Data are presented as means (standard deviation (SD)) or numbers and percentages of patients with baseline measures

6MWT: Six-Minute Walk Test; CCQ: Clinical COPD Questionnaire; GPE-DV: Global Perceived Effect - Dutch Version; HHD: hand-held dynamometer; MRC: Medical Research Council dyspnea scale; MCID: minimal clinical important difference; CI: confidence intervals; N.A: not applicable.

\*GPE-DV was only analyzed at the end of the treatment

\*\* The MCID for the MRC is yet to be established (24)

<sup>a</sup> For treatment episodes with end scores

<sup>b</sup> For the multi-level analysis of the 6MWT, we used an MCID of  $\pm \geq 30$  m (33)

<sup>c</sup> For the multi-level analysis of the CCQ, we used an MCID of  $\pm \geq 0.4$  points (34)

<sup>d</sup> For the multi-level analysis of the HHD, we used an MCID of  $\pm \geq 7.5$  Nm (35)

### Comparability

Table 4 presents the ICC calculations of the intercept-only models and the models adjusted with the explanatory variables. For the process measures, similar to the HHD, no ICC was calculated due to insufficient data. In total, 11 models could be estimated for the outcome indicators based on scores of the 6MWT (four indicators), CCQ (four indicators), GPE (one indicator for the end score), and MRC (two indicators; MCID could not be calculated). In 10 of the 11 case-mix adjusted models, the ICC increased compared with the intercept-only model, thus improving the comparability between practices, i.e., the random intercept variance of physical therapy practices increased in the adjusted models.

**Table 4 ICCs for the intercept-only model and adjusted model for the change, end, and MCID scores for each measure of the total population in practices that provided  $\geq 10$  patients**

	Intercept-only model	Adjusted model
6MWT end score	0.08	0.17 <sup>a</sup>
6MWT change score	0.00	0.01 <sup>a</sup>
6MWT MCID improvement <sup>b</sup>	0.03	0.04 <sup>a</sup>
6MWT MCID deterioration <sup>b</sup>	0.06	0.06
CCQ end score	0.11	0.20 <sup>a</sup>
CCQ change score	0.06	0.09 <sup>a</sup>
CCQ MCID improvement <sup>c</sup>	0.05	0.07 <sup>a</sup>
CCQ MCID deterioration <sup>c</sup>	0.03	0.05 <sup>a</sup>
GPE-DV end score	0.14	0.15 <sup>a</sup>
MRC end score	0.08	0.12 <sup>a</sup>
MRC change score	0.23	0.34 <sup>a</sup>

ICC: intraclass correlation coefficient; 6MWT: Six-Minute Walk Test; CCQ: Clinical COPD Questionnaire; GPE-DV: Global Perceived Effect - Dutch Version; HHD: hand-held dynamometer; MRC: Medical Research Council dyspnea scale; MCID: minimal clinical important difference; CI: confidence intervals; N.A: not applicable.

<sup>a</sup> Increase in the ICC compared with the intercept-only model following the adjustment for the case-mix variables age, gender, and baseline score of the measure.

<sup>b</sup> For the multi-level analysis of the 6MWT, we used an MCID of  $\pm \geq 30$  m (33)

<sup>c</sup> For the multi-level analysis of the CCQ, we used an MCID of  $\pm \geq 0.4$  points (34)

### Discriminability

Five of the 11 case-mix adjusted ICCs were  $>0.10$ , ranging between 0.12 and 0.32, which can be interpreted as an adequate discriminability. All adjusted models were used for the visual representation of the quality indicators in the focus group interviews. Also, the quality indicators presenting the outcomes of the HHD, for which no multi-level analysis was conducted, were presented in the focus group interviews. All defined potential quality indicators are presented as caterpillar plots (figures 2a-c). Each graph shows that a wide range of differences in outcomes exist between physical therapy practices.

## **Phase 2: Selecting a core set of quality indicators**

In total, four focus group interviews were conducted with 20 (out of 22 invited) physical therapists and three (out of five invited) senior researchers. The non-acceptance of invited participants was due to the

date and time of the focus groups that did not fit with the agenda of the potential participants. The mean duration of the focus groups was 90 minutes (range 80 – 95) Nine were female, the mean age of the participants was 39 years (range 23–60 years), and they had an average work experience of 14 years (range 1–35 years). In total, 16 of the 20 participating physical therapists also provided data for the practice test. See appendix C for an overview of the characteristics of the participants. Almost all the participants expressed that the presented quality indicators were user friendly and had value for quality improvement in daily practice. Still, several issues surrounding the presented quality indicators were discussed.

#### Using patient profiles for the comparison of patient outcomes

The participants mentioned that, in future research, it would be helpful to stratify patients based on the Dutch model, a profiling system to enhance the comparability between physical therapy practices. In 2020, an ad hoc task force of experts in the field of physical therapy, exercise therapy, rehabilitation science, respiratory medicine, general medicine, and elderly care medicine, as well as patient representatives, developed a profiling system (the “Dutch model”) for patients with COPD to allocate patients into subgroups for exercise-based care.(36) The participants of the current study suggested that baseline measures and patient characteristics needed for allocating patients to subgroups according to the profiling system should be included as process indicators in the core set. They stated that the stratification of patients into subgroups based on these profiles would enhance the comparability between practices.

#### Conditions for interpreting outcomes

Another reported problem was the limited amount of provided outcome data in the study, especially for the HHD. A possible reason could be that for the data collection we used real-world data via national data registries. These registries used predefined technical specifications. (18) During the conduction of the study, the HHD was a new measure implemented in the data registries. Potentially, this may have resulted in the low amount of provided data, which was also mentioned in the focus groups. Participant therefore suggested that the implementation of process measures is needed to stimulate routine data collection as a first step in quality improvement. When comparing outcomes, the participants were interested in the



background information of patients with COPD, such as smoking status, exacerbations, and body weight, for better interpretation of the differences in, for example, the change or end scores. When using these outcomes as a learning tool, the education of physical therapists is needed to gain knowledge about the interpretation of the outcomes. Furthermore, to enhance the comparability between practices, participants suggested to include only outcomes of patients that were treated for  $\geq 3$  months.

#### Including the percentage of a predicted value

Absolute outcomes were used to calculate the end and change scores for the 6MWT and HHD. The participants suggested that outcomes should be presented as percentages of predicted values based on reference data from the healthy population.(37, 38) These normative values are based on previous research and can be calculated according to gender, age, and body weight.

#### Selection of the core set

After discussing the outcomes, all (100%) of the physical therapists in each focus group preferred the inclusion of seven quality indicators in the core set: three process indicators for the routine measurement of the 6MWT, CCQ, and HHD; three outcome indicators using the pre- to post-treatment change in the 6MWT, CCQ, HHD scores; and a combined process indicator to monitor the baseline measurement of three measures (6MWT, CCQ, and an accelerometer (steps per day)) and patient characteristics (age, gender, body weight, and number of exacerbations in the past year) to allocate patients into subgroups based on the profiling system of the Dutch model.(36) The final core set of seven quality indicators is shown in Table 5. Figures 1a (6MWT) 1b(CCQ) 1c(HHD) presents the proportion of patients with pre- and post-treatment and 2a (6MWT), 2b (CCQ) and 2c (HHD) presents the caterpillar plots of the quality indicators in the final core set.

**Table 5 Selected core set of quality indicators accepted by stakeholders based on the perceived added value as quality improvement tools**

Type of indicator	Quality indicator description	Overall mean/ Percentage*	Range*
	Physical capacity measured with the 6MWT		

Process	The proportion of patients with COPD who underwent physical therapy treatment and who completed the 6MWT pre- and post-treatment to evaluate physical capacity	60.7%	26.1–88.8%
Outcome	The mean change score $\pm$ 95% CI of patients with COPD who underwent physical therapy treatment and pre- and post-treatment measurement with the 6MWT to evaluate physical capacity	2.8 meters	–5.4 – 13.4
Health-related quality of life measured with the CCQ			
Process	The proportion of patients with COPD who underwent physical therapy treatment and who completed the CCQ pre- and post-treatment to evaluate aspects of health-related quality of life	62.6%	14.8–88.7%
Outcome	The mean change score $\pm$ 95% CI of patients with COPD who underwent physical therapy treatment and pre- and post-treatment measurement with the CCQ to evaluate health-related quality of life	–0.1	0.3 – –0.6
Quadriceps strength measured with the HDD			
Process	The proportion of patients with COPD who underwent physical therapy treatment and who completed the HDD pre- and post-treatment to evaluate quadriceps strength	31.4%	5.9–87.5%
Outcome	The mean change score $\pm$ 95% CI of patients with COPD who underwent physical therapy treatment and pre- and post-treatment measurement with the HDD to evaluate quadriceps strength	7.5 Nm	2.7–13.1
Baseline measures for the 6MWT, CCQ, accelerometer, and patient characteristics that can be used in a profiling system to stratify patients into subgroups for care**			
Process	The proportion of patients with COPD who underwent physical therapy treatment and who completed the baseline measurements for the 6MWT, CCQ, accelerometer, gender, age, body weight, and number of exacerbations in the past year	2.4%	

6MWT: Six-Minute Walk Test; CCQ: Clinical COPD Questionnaire; GPE-DV: Global Perceived Effect - Dutch Version; HDD: hand-held dynamometer; MRC: Medical Research Council dyspnea scale; MCID: minimal clinical important difference; CI: confidence intervals. Only outcomes of patients that were included that were treated for  $\geq 3$  months

\* The overall mean/percentage and range are the outcomes of the physical therapy practices that provided  $\geq 10$  cases, used for describing and selecting the quality indicators for the core set

\*\* Baseline measures and patient characteristics selected to allocate patients into subgroups based on the Dutch model for exercise-based care in primary care (36)

## Discussion

The major finding in this study is that all participants in the focus groups accepted the quality indicators as a quality improvement tool based on their perceived added value, and selected a core set of seven outcome-based quality indicators for patients with COPD. The final core set includes a process and outcome indicator for three outcomes: physical capacity measured with the 6MWT, health-related quality of life measured with the CCQ, and quadriceps strength measured with the HDD. A combined process

indicator was included to monitor the baseline measurement of three measures used to allocate patients into subgroups based on the Dutch model profiling system.(36) To our knowledge, this is the first study to develop a core set of outcome-based quality indicators including a practice test for patients with COPD in physical therapy primary care practice. With the use of the core set, it is possible to compare standardized outcomes for patients between practices.

Several studies have developed quality indicators for COPD care,(39-44) but most sets were developed for the evaluation of processes or structures of care, e.g., monitoring the proportion of patients for whom smoking status was recorded or the availability of exercise equipment.(39, 41-44) These studies differed in their care focus areas, which were hospitalized care, end-of life care, transitional care after hospitalization, pulmonary rehabilitation, vulnerable elders, or primary care in general.(39-44) None of these publications performed a practice test. In one indicator set, developed for pulmonary rehabilitation, some similar domains (physical capacity, strength, and health-related quality of life) and measures (6MWT) were described.(43) The selection of change scores in the core set and the use of caterpillar plots is in line with other research describing the development of quality indicators based on PROMs. (2, 5, 29) (2, 5, 29) A difference is that in the current study we focused in specific on development quality indicators based on outcomes of care, while other studies are more focused on evaluating processes of care. (7, 45-49) None of these studies aimed to develop a core set of outcome-based quality indicators to be used as quality improvement tools for healthcare providers, however. Quality indicators can also be developed for pay-for-performance initiatives, policy reports, insight into practice variation/delivered care, or the identification of differences in delivered care.

Despite the fact that our core set was developed in a Dutch environment, physical therapists in other countries could potentially use the indicator set in their daily practice. Nonetheless, the context of each country needs to be taken into account, specifically cultural or clinical practice differences between countries, such as differing guidelines or educational levels of physical therapists.(50)

A strength of the current study is that we used a standard set of outcome domains and associated measures. The standard set was explicitly developed for patients with COPD being treated in primary care physical therapy practices, which was designed to be used as a basis for the further development of quality indicators.(4) The standard set is based on recommendations in guidelines and the supporting

literature, and was selected in a RAND/UCLA Delphi procedure, which is one of the most common methods for the development of quality indicators.(11, 51)

Another strength of our study is that we collected real-world data to perform a practice test prior to the selection of the core set, which was judged as an essential step in evaluating the validity, reliability, and feasibility of the indicators.(9) The interpretation of the practice test was discussed with end-users and guideline developers in focus groups. Including stakeholders in the development process is an important step for the successful implementation of quality indicators.(51) In the current study, we explicitly focused on the development of an indicator set for learning and quality improvement purposes for physical therapists. When quality indicators are designed for other purposes, such as a support tool for patients to choose providers, future research should also include other stakeholders (i.e., patients, policy makers, and health insurers) for the evaluation of their usefulness in daily practice.

A limitation of our study is that in multi-level analyses, a general rule of the thumb for the calculation of outcomes is the 30/30 rule (i.e., 30 physical therapy practices including a minimum of 30 patients each), allowing a valid comparison of indicator scores between practices.(20, 21) We did not use this rule of thumb as the threshold for estimating the case-mix adjusted scores for each quality indicator. The routine collection of clinical data by Dutch physical therapists treating patients with COPD is still in its infancy; therefore, we concluded that the 30/30 rule would not have been achievable in our study. Here, the collected data was only used as supporting tool for the selection of the core set, so we decided to include physical therapy practices that had  $\geq 10$  patients with COPD.

It is important to note that many practices did not reach the threshold of providing measurements for  $\geq 10$  patients with COPD. When the process indicators, as presented in Table 5, were based on all participating practices, the proportion of repeated measures was 39% for the 6MWT, 52% for the CCQ, and 5% for the HHD. In our view, future implementation strategies must be conducted to improve the amount of data provided; for example, by giving feedback to practices with process indicators as presented in Table 5.

Furthermore, due to the amount of data provided, we chose to compare the outcomes between physical therapy practices and not between physical therapists. When the amount of available data increases, the opportunity to compare outcomes between physical therapists, both between and within

practices, will arise. When sufficient data within practices is provided, physical therapists are able to learn from their own outcomes in comparison with peers who are employed in the same practice. We expect that, when comparing outcomes between physical therapists, the variability will be larger than between practices.

Another limitation is that we were not able to collect data that allowed us to allocate patients into subgroups based on their burden of disease, physical activity, and physical capacity.(4) Hypothetically, the comparability and discriminability of the quality indicators would increase when allocating patients into subgroups. The participants of the focus groups underlined this hypothesis and suggested the inclusion of the Dutch profiling system for patients with COPD in the core set;(36) however, the Dutch profiling system had not yet been developed at the start of the data collection for this study. Future research could evaluate the core set for each subgroup to compare more homogeneous patient groups on their baseline characteristics. Another aspect to increase the comparability is to include more patient characteristics for case-mix adjustment. As suggested by patients with COPD and physical therapists, potential relevant case-mix variables are, for example, smoking history, comorbidities and number of exacerbations.(4)

### **Implications for practice**

Outcome-based quality indicators based on real-world data, as provided in this study, can be used as a learning tool by comparing the collected patient outcomes between physical therapists or practices. This can, for example, be accomplished by discussing outcomes in peer assessment meetings of physical therapists to improve the quality of care. In such meetings, physical therapists critically appraise their peers' performance and give them constructive feedback. (52-54) In our opinion, Dutch physical therapists treating patients with COPD should first focus on expanding the amount of data collected. Giving feedback information can help to stimulate physical therapy practices to increase data collection. When sufficient data is provided and the comparison of outcomes in patient subgroups is established, the usability of the core set will increase. Future research should focus on the development of methods to improve the use of outcomes between peers and to set up specific actions to improve the quality of care.

### **Conclusion**

This is the first study to describe and select a core set of seven outcome-based quality indicators for patients with COPD treated in primary care physical therapy practice. This core set includes process and outcome indicators related to measuring physical capacity, health-related quality of life, and quadriceps strength, and a process measure for profiling patients within subgroups. To further evaluate the core outcome set, future research should explore different strategies to promote data collection, including providing feedback of the outcomes to physical therapists.

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### **Authors' contributions**

Concept/research design: A.C. Verburg, S.A. van Dulmen, H. Kiers, P.J. van der Wees

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Data analysis: A.C. Verburg

Projectmanagement: A.C. Verburg, P.J. van der Wees

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Providing participants: A.C. Verburg, S.A. van Dulmen, H. Kiers

Providing facilities/equipment: A.C. Verburg, S.A. van Dulmen, H. Kiers

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Clerical/secretarial support: A.C. Verburg

Consultation (including review of manuscript before submitting): S.A. van Dulmen, H. Kiers, P.J. van der Wees

### **Availability of data and materials**

The datasets used and/or analyzed during the current study are available from the corresponding author upon reasonable request.

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### **Ethics approval and consent to participate**

All procedures were conducted according to the Declaration of Helsinki. The study protocol was approved by the Medical Ethical Committee of Radboud university medical center (registration no. 2019-5455). Informed consent was obtained from all participants included in the current study.

### **Consent for publication**

Not applicable

**Competing interest**

The authors declare that they have no competing interests.



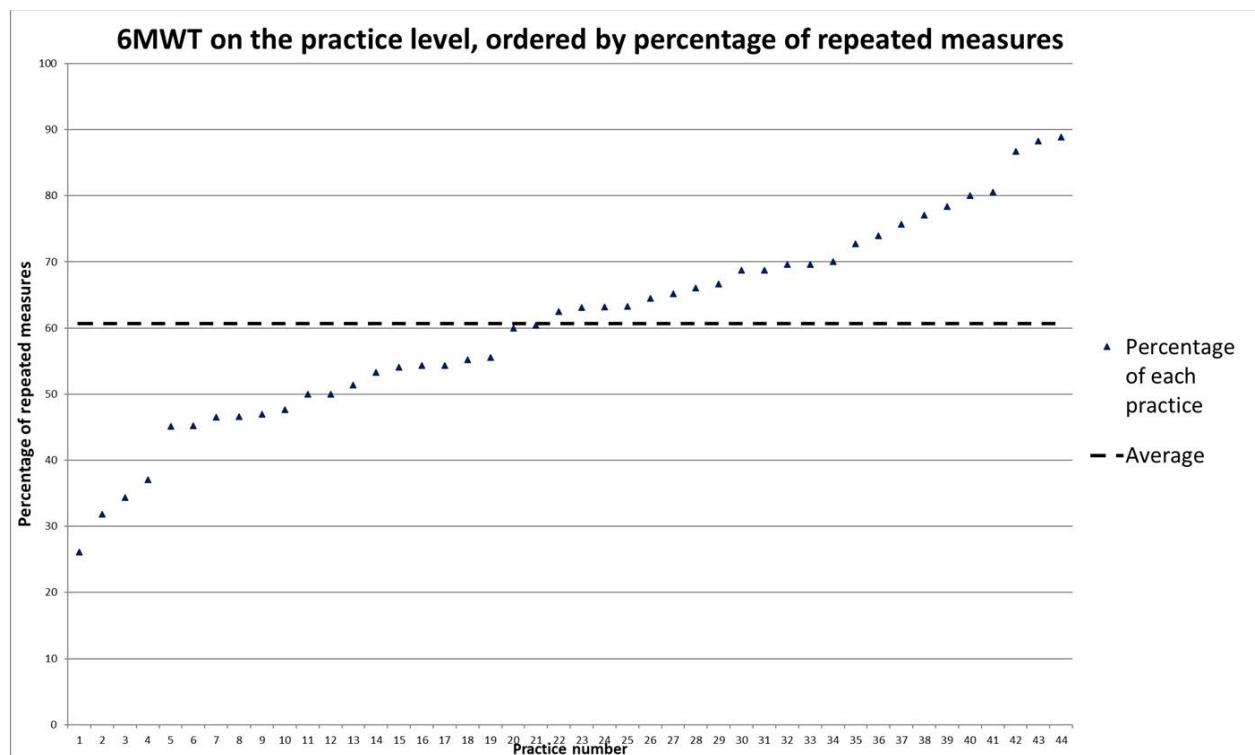
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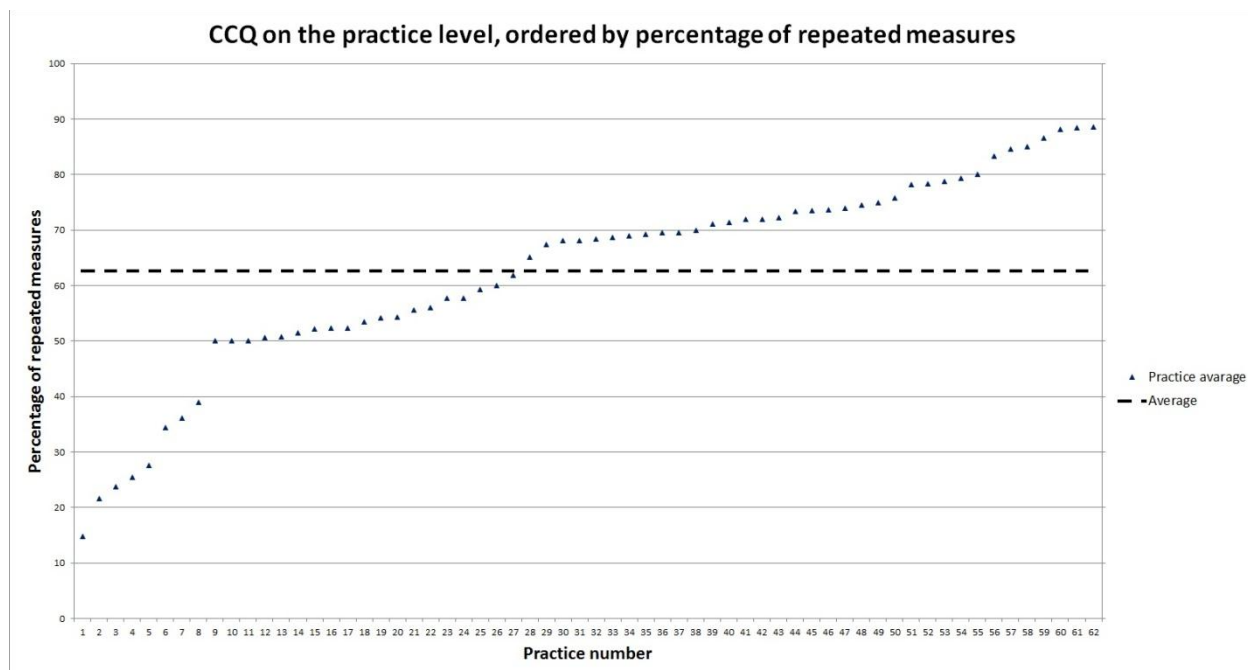
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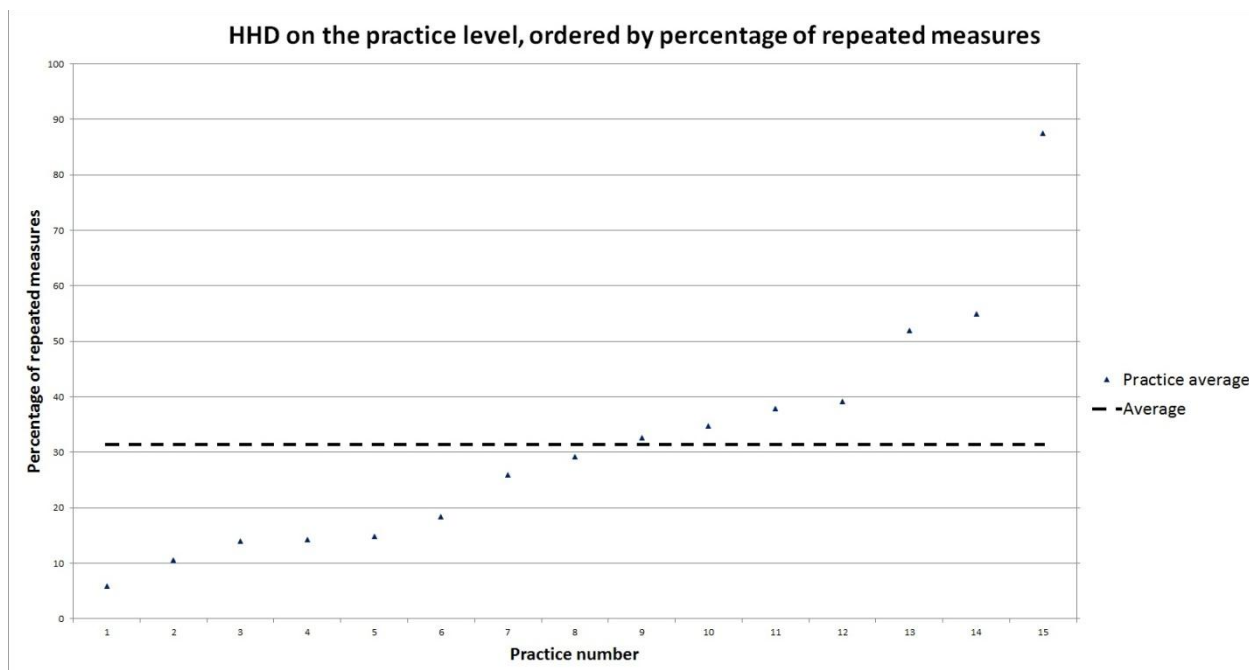
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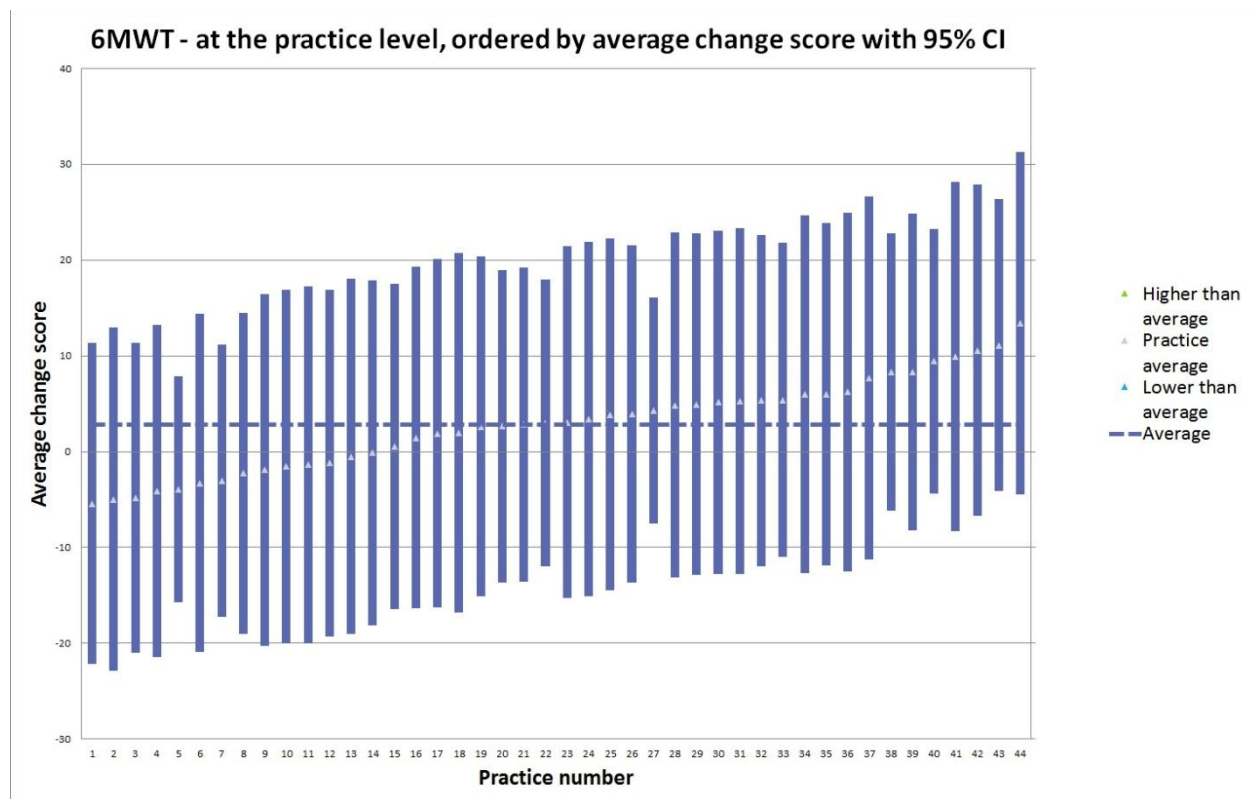
The proportion of patients with COPD who underwent physical therapy treatment in which a pre and/or post the measure was provided



The proportion of patients with COPD who underwent physical therapy treatment in which a pre and/or post the measure was provided for the CCQ

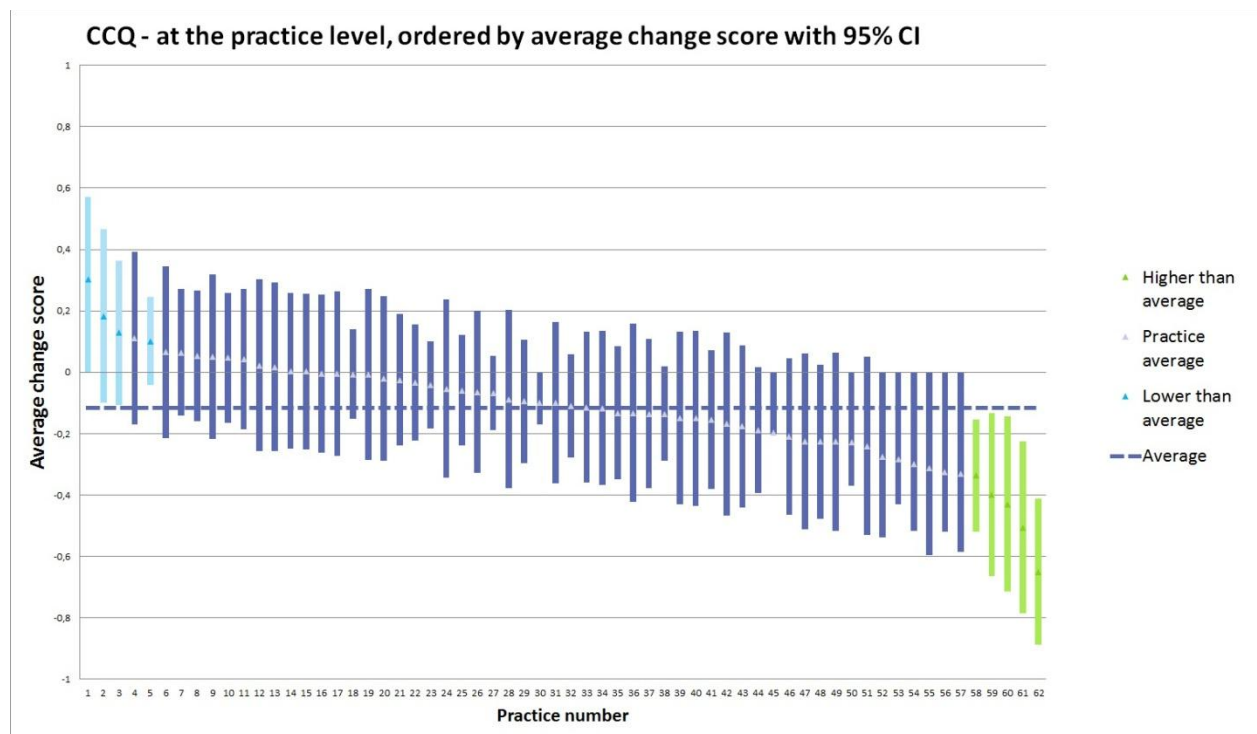


The proportion of patients with COPD who underwent physical therapy treatment in which a pre and/or post the measure was provided for the HHD

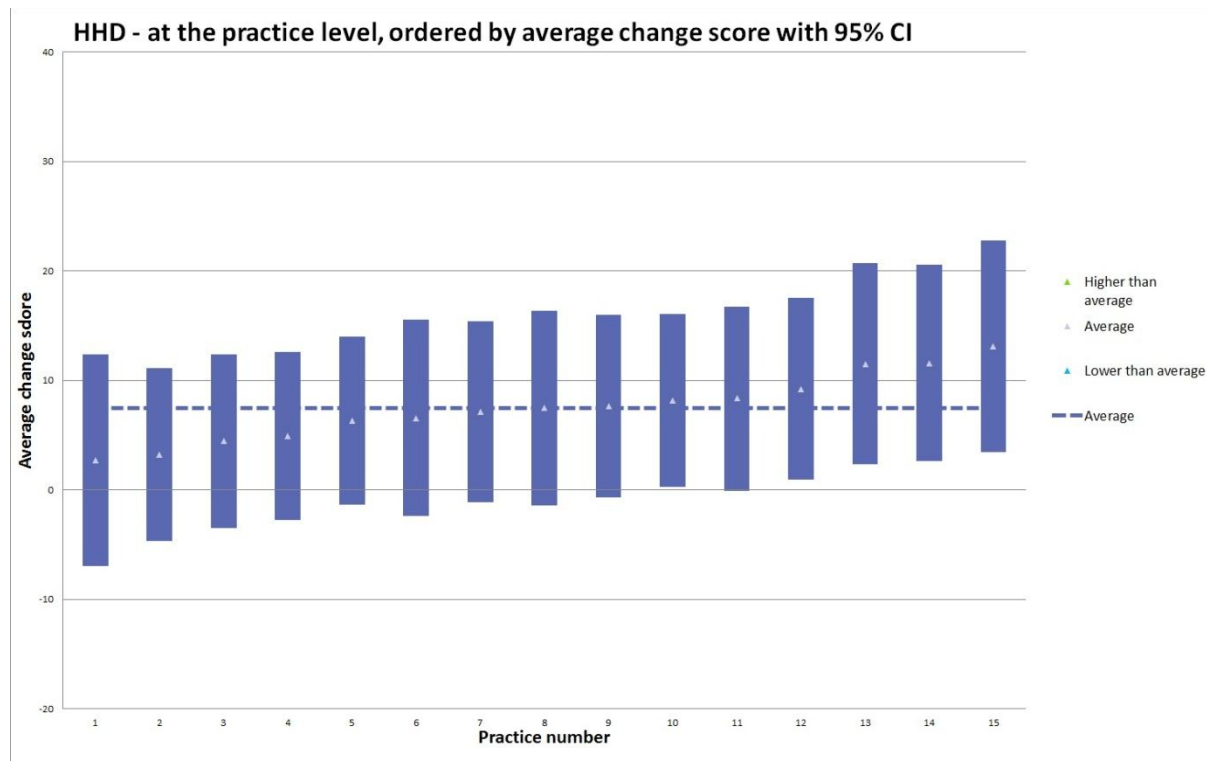


The mean change score on the 6MWT with 95% CI of patients with COPD between pre- and post-physical therapy treatment





The mean change score on the CCQ with 95% CI of patients with COPD between pre- and post-physical therapy treatment



The mean change score on the HHD with 95% CI of patients with COPD between pre- and post-physical therapy treatment

## Appendix A

Table 1 Standard set of outcome domains and associated measures for patients with COPD accepted by all stakeholders as described in recent research[1]

Domain	Measure	Guided measurement protocol		
	A: mandatory for all patients with COPD	Intake	Every 3 months	End
Practice/physical therapist level	Characteristics of practices and physical therapists	Once a year		
Physical capacity	6-Minute Walk Test (6 MWT)	X	X	X
Health-related quality of life	Clinical COPD Questionnaire (CCQ)	X	X	X <sup>1</sup>
Experience	Global Perceived Effect (GPE)		X	X
	B: Conditional measures			
Quadriceps strength	Hand-Held Dynamometer (HHD) (with a Microfet™)	X	X	X
Dyspnea	Medical Research Council Dyspnea (MRC)	X	X	X
	C: Exploratory measures <sup>2</sup>			
Physical activity	Accelerometer (for physical activity in daily life)			
ABC-Tool	The Assessment of Burden of COPD tool			

<sup>1</sup> After treatment ≥12 months the CCQ needs only to be measured once a year.

<sup>2</sup> The exploratory measures were not included in this study

Table 2 Description of PROMs and physical performance measures

Patient Reported Outcome Measures (PROMs) and physical performance measures	Description
6-Minute Walk Test (6 MWT)[2]	Physical performance test where the patients walks for 6 minutes, according to the structured guidelines of Butland et al.[2]
Clinical COPD Questionnaire (CCQ)[3]	The CCQ is an 10 item questionnaire scores the health-related quality of life for patients with COPD, scored on a 7-point Likert scale, score 0 is “very good health status” and 6 “extremely poor health status”
Global Perceived Effect Dutch Version (GPE-DV) [4]	This 2-item questionnaire is about patient satisfaction and the patient experienced effect of the treatment on a 7-point likert scale
Hand Held Dynamometrie (HHD)[5]	Physical performance test where the patients scores his maximal isometric quadriceps strength
Medical Research Council Dyspnea (MRC)[6]	Single item scores the patient experienced symptoms of dyspnoea during physical activity on a 5-point Likert scale, score 1 is “symptoms of dyspnoea during heavy activities” and 5 “to symptoms of dyspnoea to leave the house”
The Assessment of Burden of COPD index [7]	A 15 item questionnaire scores the experienced burden of the COPD disease
Accelerometer	Tool to measure physical activity in daily practice in average steps per day
Characteristics of practices and physical therapist	This structure measure scores the education level of the physical therapists and identifies the characteristics of the practice (e.g. equipment’s, training facilities)

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## Appendix B Description of defined quality indicators for patients with COPD

### 1. Monitoring the process of repeated measurement of the 6MWT

<i>Definition</i>	The proportion of patients with COPD who underwent physical therapy treatment and who completed the 6-Minute Walk Test (6 MWT) pre- and post-treatment to evaluate physical capacity
<i>Rationale</i>	Improvement of physical capacity is an important goal in physical therapy treatment for patients with COPD. Physical capacity is measured with the 6MWT
<i>Numerator</i>	The number of patients with COPD who underwent physical therapy treatment for ≥3 months, and who completed the 6MWT pre- and post-treatment measured
<i>Denominator</i>	All patients who underwent physical therapy treatment for COPD
<i>Specification</i>	Physical capacity is measured in all patients with COPD with the 6MWT, a physical performance test where the patients walks for 6 minutes in a comfortable way
<i>Type of indicator</i>	Process

### 2. Monitoring the mean end score of the 6MWT

<i>Definition</i>	Describing the mean end score and 95% CI of patients with COPD who underwent physical therapy treatment for ≥3 months, and are pre- and post-measured with the 6-Minute Walk Test (6 MWT) to evaluate physical capacity
<i>Rationale</i>	Improvement of physical capacity is an important goal in physical therapy treatment for patients with COPD. Physical capacity is measured with the 6MWT
<i>Specification</i>	Physical capacity is measured with the 6MWT in all patients with COPD, a physical performance test where the patients walks for 6 minutes in a comfortable way
<i>Type of indicator</i>	Outcome

### 3. Monitoring the mean change score of the 6MWT

<i>Definition</i>	The mean change score and 95% CI of patients with COPD who underwent physical therapy treatment for ≥3 months, and are pre- and post-measured with the 6-Minute Walk Test (6 MWT) to evaluate physical capacity
<i>Rationale</i>	Improvement of physical capacity is an important goal in physical therapy treatment for ≥3 months, for patients with COPD. Physical capacity is measured with the 6MWT
<i>Specification</i>	Physical capacity is measured with the 6MWT in all patients with COPD, a physical performance test where the patients walks for 6 minutes in a comfortable way
<i>Type of indicator</i>	Outcome

### 4. Monitoring the minimal clinical important difference of the 6MWT

<i>Definition</i>	The proportion of patients who underwent physical therapy treatment and experienced an minimal clinical important difference (MCID) on physical capacity measured with the 6-Minute Walk Test (6 MWT)
<i>Rationale</i>	Improvement of physical capacity is an important goal in physical therapy treatment for patients with COPD. Physical capacity is measured with the 6MWT
<i>Numerator</i>	The number of patients with COPD who underwent physical therapy treatment for ≥3 months, and had a physical capacity difference of ≥30 meter between pre- and post-treatment measured with the 6MWT
<i>Denominator</i>	Total number of patients with COPD who underwent physical therapy treatment ≥3 months, and was measured on pre- and post-treatment with the 6MWT
<i>Specification</i>	Physical capacity is measured with the 6MWT in all patients with COPD, a physical performance test where the patients walks for 6 minutes in a comfortable way
<i>Type of indicator</i>	Outcome

## 5. Monitoring the process of repeated measurement of the CCQ

<i>Definition</i>	The proportion of patients with COPD who underwent physical therapy treatment and who completed the Clinical COPD Questionnaire (CCQ) pre- and post-treatment to evaluate health-related quality of life
<i>Rationale</i>	Monitoring health-related quality of life is an important goal in physical therapy treatment for patients with COPD. Health-related quality of life is measured with the CCQ
<i>Numerator</i>	The number of patients with COPD who underwent physical therapy treatment for $\geq 3$ months, and who completed the CCQ pre- and post-treatment measured
<i>Denominator</i>	All patients who underwent physical therapy treatment for patients with COPD
<i>Specification</i>	The CCQ is an 10 item questionnaire scores aspects of health-related quality of life for patients with COPD, scored on a 7-point Likert scale, score 0 is "very good health status" and 6 "extremely poor health status"
<i>Type of indicator</i>	Process

## 6. Monitoring the mean end score of the CCQ

<i>Definition</i>	Describing the mean end score and 95% CI of patients with COPD who underwent physical therapy treatment for $\geq 3$ months, and are pre- and post-measured with the Clinical COPD Questionnaire (CCQ) to evaluate health-related quality of life
<i>Rationale</i>	Monitoring health-related quality of life is an important goal in physical therapy treatment for patients with COPD. Health-related quality of life is measured with the CCQ
<i>Specification</i>	The CCQ is an 10 item questionnaire scores aspects of health-related quality of life for patients with COPD, scored on a 7-point Likert scale, score 0 is "very good health status" and 6 "extremely poor health status"
<i>Type of indicator</i>	Outcome

## 7. Monitoring the mean change score of the CCQ

<i>Definition</i>	The mean change score and 95% CI of patients who underwent physical therapy treatment for $\geq 3$ months, and are pre- and post-measured with the Clinical COPD Questionnaire (CCQ) to evaluate health-related quality of life
<i>Rationale</i>	Monitoring health-related quality of life is an important goal in physical therapy treatment for patients with COPD. Health-related quality of life is measured with the CCQ
<i>Specification</i>	The CCQ is an 10 item questionnaire scores aspects of health-related quality of life for patients with COPD, scored on a 7-point Likert scale, score 0 is "very good health status" and 6 "extremely poor health status"
<i>Type of indicator</i>	Outcome

## 8. Minimal clinical important difference of the CCQ

<i>Definition</i>	The proportion of patients with COPD who underwent physical therapy treatment and experienced an minimal clinical important difference (MCID) on health-related quality of life and measured with the Clinical COPD Questionnaire (CCQ)
<i>Rationale</i>	Improving physical functioning is an important goal after physical therapy treatment. Health-related quality of life is measured with the CCQ
<i>Numerator</i>	The number of patients with COPD who underwent physical therapy treatment for $\geq 3$ months, and had a health-related quality of life difference of $\geq 0.4$ points between pre- and post-treatment measured with the CCQ
<i>Denominator</i>	Total number of patients with COPD who underwent physical therapy treatment $\geq 3$ months, and health-related quality of life was measured between pre- and post-treatment with the CCQ
<i>Specification</i>	The CCQ is an 10 item questionnaire scores aspects of health-related quality of life for patients with COPD, scored on a 7-point Likert scale, score 0 is "very good health status" and 6 "extremely poor health status"
<i>Type of indicator</i>	Outcome

## 9. Monitoring the process of end measurement of the GPE-DV

<i>Definition</i>	The proportion of patients with COPD who underwent physical therapy treatment and who completed the Global Perceived Effect Dutch Version (GPE-DV) post-treatment to evaluate the perceived treatment effect
<i>Rationale</i>	Patient with COPD with perceived treatment effect is a generic way to monitor the result of the treatment
<i>Numerator</i>	The number of patients with COPD who underwent physical therapy treatment and who completed the GPE-DV post-treatment measured
<i>Denominator</i>	All patients who underwent physical therapy treatment for patients with COPD
<i>Specification</i>	This questionnaire about patients experienced treatment effect is scored on a 7-point Likert scale, score 1 is "much improved" and 7 "worse than ever"
<i>Type of indicator</i>	Process

## 10. Monitoring the mean end score of the GPE-DV

<i>Definition</i>	Describing the mean end score and 95% CI of patients with COPD who underwent physical therapy treatment for $\geq 3$ months, and are post measured with the Global Perceived Effect Dutch Version (GPE-DV) to evaluate the perceived treatment effect
<i>Rationale</i>	Patient with COPD with perceived treatment effect is a generic way to monitor the result of the treatment
<i>Specification</i>	This questionnaire about patient experienced treatment effect is scored on a 7-point Likert scale, score 1 is "much improved" and 7 "worse than ever"
<i>Type of indicator</i>	Outcome

## 11. Monitoring the process of repeated measurement of the HHD

<i>Definition</i>	The proportion of patients with COPD who underwent physical therapy treatment and who completed the hand-held dynamometer (HHD) pre- and post-treatment to evaluate quadriceps strength
<i>Rationale</i>	Monitoring muscle strength is an important goal in physical therapy treatment for patients with COPD. Muscle strength is measured with the HHD (with a Microfet™)
<i>Numerator</i>	The number of patients with COPD who underwent physical therapy treatment for ≥3 months, and who completed the HHD pre- and post-treatment measured
<i>Denominator</i>	All patients with COPD who underwent physical therapy treatment
<i>Specification</i>	Quadriceps strength is measured in all patients using the hand-held dynamometer (HHD), with a Microfet™
<i>Type of indicator</i>	Process

## 12. Monitoring the mean end score of the HHD

<i>Definition</i>	Describing the mean end score and 95% CI of patients with COPD who underwent physical therapy treatment for ≥3 months, and are pre- and post-measured with the hand-held dynamometer (HHD) to evaluate quadriceps strength
<i>Rationale</i>	Monitoring muscle strength is an important goal in physical therapy treatment for patients with COPD. Muscle strength is measured with the HHD (with a Microfet™)
<i>Specification</i>	Quadriceps strength is measured in all patients using the hand-held dynamometer (HHD), with a Microfet™
<i>Type of indicator</i>	Outcome

## 13. Monitoring the mean change score of the HHD

<i>Definition</i>	The mean change score and 95% CI of patients with COPD who underwent physical therapy treatment for ≥3 months, and are pre- and post-measured with the hand-held dynamometer (HHD) to evaluate quadriceps strength
<i>Rationale</i>	Monitoring muscle strength is an important goal in physical therapy treatment for patients with COPD. Muscle strength is measured with the HHD (with a Microfet™)
<i>Specification</i>	Quadriceps strength is measured in all patients using the hand-held dynamometer (HHD), with a Microfet™
<i>Type of indicator</i>	Outcome

## 14. Monitoring the minimal clinical important difference of the HHD

<i>Definition</i>	The proportion of patients with COPD who underwent physical therapy treatment and experienced an minimal clinical important difference (MCID) on physical capacity measured with the hand-held dynamometer (HHD) to evaluate quadriceps strength
<i>Rationale</i>	Monitoring muscle strength is an important goal in physical therapy treatment for patients with COPD. Muscle strength is measured with the HHD (with a Microfet™)
<i>Numerator</i>	The number of patients with COPD who underwent physical therapy treatment for ≥3 months, and had a quadriceps strength difference of ≥7.5 Nm between pre- and post-treatment measured with the HHD
<i>Denominator</i>	Total number of patients with COPD who underwent physical therapy treatment ≥3 months, and quadriceps strength was measured on pre- and post-treatment with the HHD (with a Microfet™)
<i>Specification</i>	Quadriceps strength is measured in all patients using the hand-held dynamometer (HHD), with a Microfet™
<i>Type of indicator</i>	Outcome



## 15. Monitoring the process of repeated measurement of the MRC

<i>Definition</i>	The proportion of patients with COPD who underwent physical therapy treatment and who completed the Medical Research Council Dyspnea (MRC) pre- and post-treatment to evaluate symptoms of dyspnoea
<i>Rationale</i>	Monitoring symptoms of dyspnoea during physical activity is an important goal in physical therapy treatment, for patients with COPD. Symptoms of dyspnoea was measured with the MRC.
<i>Numerator</i>	The number of patients with COPD who underwent physical therapy treatment for $\geq 3$ months, and who completed the MRC pre- and post-treatment measured
<i>Denominator</i>	All patients who underwent physical therapy treatment for patients with COPD
<i>Specification</i>	Single item scores the patients with COPD experienced symptoms of dyspnoea during physical activity on a 5-point Likert scale, score 1 is "breathless during heavy activities" and 5 "to breathless to leave the house"
<i>Type of indicator</i>	Process

## 16. Monitoring the mean end score of the MRC

<i>Definition</i>	Describing the mean end score and 95% CI of patients with COPD who underwent physiotherapy treatment for $\geq 3$ months, and are pre- and post-measured with the Medical Research Council Dyspnea (MRC) to evaluate symptoms of dyspnoea
<i>Rationale</i>	Monitoring symptoms of dyspnoea during physical activity is an important goal in physical therapy treatment, for patients with COPD. Symptoms of dyspnoea was measured with the MRC
<i>Specification</i>	Single item scores the patients with COPD experienced symptoms of dyspnoea during physical activity on a 5-point Likert scale, score 1 is "breathless during heavy activities" and 5 "to breathless to leave the house"
<i>Type of indicator</i>	Outcome

## 17. Monitoring the mean change score of the MRC

<i>Definition</i>	The mean change score and 95% CI of patients with COPD who underwent physical therapy treatment for $\geq 3$ months, and are pre- and post-measured with the Medical Research Council Dyspnea (MRC) to evaluate symptoms of dyspnoea
<i>Rationale</i>	Monitoring symptoms of dyspnoea during physical activity is an important goal in physical therapy treatment, for patients with COPD. Symptoms of dyspnoea was measured with the MRC
<i>Specification</i>	Single item scores the patient with COPD experienced symptoms of dyspnoea during physical activity on a 5-point Likert scale, score 1 is "breathless during heavy activities" and 5 "to breathless to leave the house"
<i>Type of indicator</i>	Outcome

## **Appendix C** characteristics of participants in the focus groups

Table 1 characteristics of participants in the focus groups

<b>ID</b>	<b>Gender</b>	<b>Age, years</b>	<b>Work experience, years</b>
P.01	Female	33	6
P.02	Female	33	3
P.03	Male	30	5
P.04	Male	33	5
P.05	Male	34	10
P.06	Female	25	0
P.07	Man	54	26
P.08	Female	37	15
P.09	Female	58	35
P.10	Female	32	9
P.11	Male	30	7
P.12	Male	23	1
P.13	Male	39	15
P.14	Male	35	4
P.15	Female	40	13
P.16	Female	48	23
P.17	Female	29	2
P.18	Male	58	34
P.19	Male	57	31
P.20	Male	60	34