

# Adherence to an eHealth Self-Management Intervention for Patients with Both COPD and Heart Failure: Results of a Pilot Study

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**Background:** Chronic obstructive pulmonary disease (COPD) and chronic heart failure (CHF) often coexist and share periods of symptom deterioration. Electronic health (eHealth) might play an important role in adherence to interventions for the self-management of COPD and CHF symptoms by facilitating and supporting home-based care.

**Methods:** In this pilot study, an eHealth self-management intervention was developed based on paper versions of multi-morbid exacerbation action plans and evaluated in patients with both COPD and CHF. Self-reporting of increased symptoms in diaries was linked to an automated decision support system that generated self-management actions, which was communicated via an eHealth application on a tablet. After participating in self-management training sessions, patients used the intervention for a maximum of four months. Adherence to daily symptom diary completion and follow-up of actions were analyzed. An add-on sensorized (Respiro<sup>®</sup>) inhaler was used to analyze inhaled medication adherence and inhalation technique.

**Results:** In total, 1148 (91%) of the daily diaries were completed on the same day by 11 participating patients (mean age  $66.8 \pm 2.9$  years; moderate (55%) to severe (45%) COPD; 46% midrange left ventricular function (LVF) and 27% reduced LVF). Seven patients received a total of 24 advised actions because of increased symptoms of which 11 (46%) were followed-up. Of the 13 (54%) unperformed advised actions, six were “call the case manager”. Adherence to inhaled medication was 98.4%, but 51.9% of inhalations were performed incorrectly, with “inhaling too shortly” (<1.25 s) being the most frequent error (79.6%).

**Discussion:** Whereas adherence to completing daily diaries was high, advised actions were inadequately followed-up, particularly the action “call the case manager”. Inhaled medication adherence was high, but inhalations were poorly performed. Future research is needed to identify adherence barriers, further tailor the intervention to the individual patient and analyse the intervention effects on health outcomes.

**Keywords:** telemedicine, chronic conditions, self-treatment, disease management, dry powder inhalers

## Introduction

Chronic obstructive pulmonary disease (COPD) and chronic heart failure (CHF) are both progressive diseases that share periods of acute deterioration of symptoms. They are associated with each other (7.5% to 31.3% of patients with COPD also have CHF) which adds to the high burden of both diseases.<sup>1,2</sup> This association may result from shared risks factors (eg smoking) and overlapping symptoms. Dyspnea could,

for example, be related to either a COPD exacerbation or a sudden deterioration of CHF.<sup>3</sup> Overlapping symptoms can easily delay the start of appropriate treatment as it complicates differentiation of both diseases,<sup>4</sup> and may lead to a further increase in patient burden and healthcare costs.<sup>5,6</sup>

Self-management interventions become more significant in the usual care of patients with COPD and CHF.<sup>7</sup> They consist of multiple components (eg optimizing physical activity, self-treatment of exacerbations and improving inhaled medication adherence and technique)<sup>8</sup> and aim to positively change the patient's health behavior and improve patients' self-management skills by a personalized, structured approach.<sup>8</sup> Evidence shows that COPD self-management interventions lead to improved health-related quality of life, respiratory-related hospitalization rate and dyspnea.<sup>9–11</sup> Self-management interventions for patients with CHF have shown beneficial effects on heart failure hospitalization duration for younger patients (<65 years), and time to CHF-related hospitalizations and all-cause death for patients of all ages.<sup>12</sup> Incorporating multi-morbid exacerbation action plans into a self-management intervention tailored for COPD patients with at least one common comorbidity (CHF, ischemic heart disease, anxiety, depression, diabetes mellitus) reduces COPD exacerbation duration and the risk of having at least one respiratory-related hospitalization during follow-up.<sup>13</sup>

Adherence to and uptake of self-management interventions are essential for effective self-management in patients with COPD.<sup>14,15</sup> Studies have shown that adherence to action plans is associated with a reduced exacerbation duration and hospitalization rate in patients with COPD.<sup>14,15</sup> However, previous randomized controlled trials, involving paper-based COPD self-management interventions, have shown poor results with regard to action plan adherence. In the study of Schrijver et al, only 38% of the patients showed (sub)optimal adherence to paper COPD exacerbation action plans<sup>16</sup> and Bischoff et al found similar results on adherence (40%).<sup>14</sup> Adherence to self-management interventions in patients with chronic diseases like COPD and CHF might be influenced by factors, such as age, social support, disease perception and knowledge, the role of the healthcare provider and case manager, (digital) health literacy, and to what level an intervention fits a patient's needs and competences.<sup>17–25</sup>

Electronic health (eHealth) could play an important role in improving adherence, as it makes it easier to

provide tailored information, give reminders, and adapt to patients' needs.<sup>21,26–29</sup> Self-management interventions are increasingly provided to patients with COPD or CHF by using eHealth technology at home to support patients in health communication (eg teleconsultation), self-monitoring (eg symptom diary, wearables), and their medical treatment (eg self-treatment with prednisolone).<sup>30–33</sup> The different intervention components have, however, yet to be robustly combined into a comprehensive eHealth intervention for patients with both COPD and CHF.

In this study, we developed an eHealth self-management intervention for patients with both COPD and CHF that was based on an already evaluated paper-based self-management intervention.<sup>13</sup> The aim of this pilot study was to evaluate patient adherence to this eHealth self-management intervention during a follow-up of a maximum of four months. In addition, adherence to inhaled medication and patients' inhalation technique was assessed.

## Materials and Methods

### Study Design

This is a prospective pilot study in which patients with both COPD and CHF were recruited from two hospitals (Medisch Spectrum Twente (MST) Enschede and Ziekenhuisgroep Twente (ZGT) Almelo and Hengelo) in the Netherlands.

In September and October 2018, patients participated in three self-management training sessions. After the first session, patients started to use an eHealth self-management intervention on a tablet for a period of a maximum of four months. The study was approved by the Medical Ethical Committee Twente and registered with the Netherlands Trial Register (NL6480). The study was conducted in accordance with the Declaration of Helsinki.

### Selection of Patients

In hospital MST, electronic health record data of in- and outpatients from the respiratory department were screened for patient eligibility. In hospital ZGT, outpatients attending the heart failure clinics of the involved nurse practitioner CHF were screened for eligibility. Subsequently, the patient's pulmonary physician and/or cardiologist were asked for permission for inclusion of their patient. Furthermore, patients received information about the study from the involved researchers face-to-face and/or by phone. Written informed consent to participate in the

study was obtained from all patients prior to data collection.

Patients had to fulfill the following inclusion criteria: 1) a clinical diagnosis of COPD defined according to the GOLD criteria;<sup>7</sup> 2) a clinical diagnosis of CHF defined according to the current (2016) ESC guidelines;<sup>3</sup> 3)  $\geq 2$  COPD and/or CHF exacerbations, defined as deterioration of symptoms for which treatment with oral corticosteroids and/or antibiotics (for COPD) or diuretics (for CHF) were necessary, in the two years preceding study entry, and/or  $\geq 1$  hospitalization for COPD and/or CHF in the two years preceding study entry; 4) age  $\geq 40$  years; 5)  $\geq 1$  week after exacerbation of COPD and/or CHF; 6)  $\geq 1$  week after hospitalization; 7)  $\geq 4$  weeks post-rehabilitation; 8) able to understand and read the Dutch language; and 9) able to use a tablet.

Exclusion criteria were as follows: 1) end stage of a disease; 2) other serious lung disease (eg  $\alpha 1$ -antitrypsin deficiency; interstitial lung diseases); 3) expected cardiovascular intervention within three months; 4) currently enrolled in a (randomized controlled) trial; 5) waiting for a heart or lung transplantation; 6) renal dialysis; 7) diabetes mellitus type I; 8) Hospital Anxiety and Depression Scale (HADS)-score<sup>34,35</sup> of  $\geq 11$  for anxiety and/or depression domain symptom scores.

## Self-Management Intervention

### Self-Management Training Sessions

Self-management training sessions consisted of two group sessions (two hours per session) and a one-hour individual session at weekly intervals. These sessions were led by trained case managers (one nurse practitioner COPD, one nurse practitioner CHF). The sessions were based on self-management training sessions that were provided in a previous self-management study in patients with COPD and comorbidities.<sup>13</sup> During these sessions, the significance of early recognition of symptom deterioration and prompt and appropriate treatment was emphasized. First, the patient's individual symptom level in a stable health state was discussed and described on a "what are my 'usual' symptoms" card. Subsequently, patients were trained on how to recognize symptom deterioration by comparing their symptoms over the last 24 hours to their "usual" symptoms. Monitoring of symptoms over a longer period of time (eg recovery from exacerbation to a stable health state) was trained using a specific "monitoring module" on the eHealth application (see details in the paragraph "monitoring module"). During the self-management training sessions, patients received a tablet and were instructed on how to connect it to WiFi internet at home. They also received an add-on sensor for

inhaled medication (if applicable), a Fitbit<sup>®</sup> for measuring step count, and a digital weighing scale. Study technicians assured that all devices were connected to the tablet via Bluetooth. Two weeks after the self-management training sessions, a follow-up phone call with the patients was performed by the case managers to verify usage of and problems with the different components of the self-management intervention. The content of the self-management training sessions is detailed in [Table 1](#).

### eHealth Self-Management Application Modules

The eHealth self-management intervention was offered through a tablet application for patients and a website accessible via PC for the case managers and researchers. Patients required accessibility to the internet to be able to use the self-management application. The different modules of the eHealth self-management application for patients are shown in [Table 2](#), and some are explained in more detail below.

### Self-Management Module

The digital daily symptom diary included symptoms that were related to COPD (eg dyspnea, cough) and CHF (eg weight by digital weighing scale, edema). All diaries included symptoms regarding ischemic heart disease, anxiety and depression, irrespective of formal diagnosis. This was done for safety reasons, as exacerbations of these comorbidities can be triggered by and may show similar symptoms as COPD and CHF exacerbations (eg dyspnea, fatigue).<sup>3,7,36</sup> The latter may confuse patients with proper differentiation and self-treatment of diseases. Every day, participants had to detail whether their symptoms in the last 24 hours were the same, slightly more, or significantly more compared to their "usual" symptoms (defined on their "what are my 'usual' symptoms" card). However, for some symptoms, they only had to differentiate between "normal" or "increased". If necessary, an automated decision support system launched a message with an action for the patient to take (eg initiate self-treatment, perform relaxation exercises, call the case manager). The automated decision support system was based on a paper version of an exacerbation action plan for patients with COPD and comorbidities.<sup>13</sup> It was personalized by establishing per patient the type and dose of diuretics and antibiotics that should be used during symptom deterioration and whether they needed to have nitroglycerine at home (in case of diagnosed ischemic heart disease). The type and dose of diuretics and antibiotics

**Table 1** Content of Self-Management Training Sessions

<b>Week 1</b> Group session	<ul style="list-style-type: none"> <li>• Self-management information regarding COPD and CHF (e.g. medication, risk factors)</li> <li>• Information regarding symptom monitoring, recognition of symptom deterioration and self-treatment of symptoms (e.g. triggers of exacerbations)</li> <li>• Introduction and demonstration on how to use the self-management application on a tablet</li> <li>• Demonstration how to use the digital weighing scale and Fitbit</li> </ul>
<b>Week 2</b> Individual session	<ul style="list-style-type: none"> <li>• Discussion of experiences and addressing perceived problems with the eHealth self-management application, digital weighing scale and Fitbit</li> <li>• Completion of 'what are my "usual" symptoms' card</li> <li>• Training on symptom monitoring, recognition of symptom deterioration, and self-treatment of symptoms</li> <li>• Practicing completing the daily symptom diary and following up advised actions on the self-management application</li> <li>• Practicing the use of the other modules of the self-management application (for more details see paragraph 'eHealth self-management application modules')</li> <li>• Training inhalation technique of inhaled medication</li> </ul>
<b>Week 3</b> Group session	<ul style="list-style-type: none"> <li>• Re-iteration of the use of the self-management application and addressing potential perceived problems</li> <li>• Promoting healthy lifestyle behaviors: diet, physical fitness and exercise, quit smoking</li> <li>• Practicing breathing- and relaxation exercises</li> <li>• Instruction about add-on sensor for inhaled medication (Respiro® by Amiko) for measuring inhaled medication adherence and technique</li> </ul>
<b>Week 5</b> Phone call	<ul style="list-style-type: none"> <li>• Verifying whether the different modules of the self-management application were used (for more details see paragraph 'eHealth self-management application modules')</li> <li>• Verifying problems with the use of the self-management application</li> <li>• Feedback on diary completion and advised actions</li> </ul>

were established in agreement with the patient's healthcare providers from the cardiology and pulmonology department. Patients received prescriptions for the action plan medication from the case manager at the start of the study. The automated decision support system advised patients to call the case manager (or general practitioner outside office hours) when symptoms did not improve after two days of self-treatment. If dyspnea did not improve two days after the start of self-treatment, patients were asked to have NT-proBNP measured at a local laboratory to be able to distinguish between symptoms of COPD or CHF and subsequently call the case manager for their results and further advice. Once a week, patients were asked if they had visited a doctor or started medication. A link to this weekly questionnaire, daily symptom diary, advised actions and a reminder for weighing were listed in the patient self-management application.

For safety reasons, case managers would check online whether patients reported an improvement in symptoms the days after patients had received advice on self-treatment. If symptoms did not improve, case managers would call the patient to provide further advice.

### Monitoring Module

The monitoring module showed individual patients' health status and self-reported symptoms per day, advised and performed actions, and a weight graph.

### Inhaler Module

Inhaled medication adherence and inhalation technique, including duration of flow, peak inspiratory flow, orientation of the inhaler, and closing of the cap, were monitored by an add-on inhaler sensor (Respiro<sup>®37</sup> by Amiko Digital Health Limited, London, UK), which was compatible with the Ellipta<sup>®</sup> (GlaxoSmithKline BV, UK) inhaler. At inclusion, each patient's pulmonary physician was asked whether the patient was eligible for a switch to the Ellipta<sup>®</sup> inhaler. The patient's daily medication schedule was established collaboratively by patient and healthcare provider. The add-on inhaler sensors transferred all data collected via Bluetooth to a paired app installed on the tablet device, which in turn uploaded the data to the cloud. The self-management application showed these data and indicated whether the inhalations were performed correctly. On-sensor audio-visual signs reminded the patients of scheduled inhaled medication doses.

**Table 2** Content of the eHealth Self-Management Intervention Modules

Modules	Contents
Self-management (home view)	<ul style="list-style-type: none"> <li>• Phone number of case manager (during office hours) and General Practitioner (outside office hours)</li> <li>• Daily symptom diary</li> <li>• Automated decision support system that launched a patient's tailored advised action in case of symptom deterioration.</li> <li>• To-do list (e.g. complete diary, initiate self-treatment)</li> <li>• Overview: health status (stable, slight or significant deterioration of symptoms), weight (current and difference with the day before), last action that was launched, inhaled medication adherence</li> </ul>
Monitoring	<ul style="list-style-type: none"> <li>• Detailed overview of health status, self-reported symptoms, advised and performed actions and weight graph</li> </ul>
Inhaler <sup>a</sup>	<ul style="list-style-type: none"> <li>• Graph illustrating performed inhalations per day and whether the inhalation technique was correctly/incorrectly performed, measured by add-on inhaler sensor (Respiro®)</li> </ul>
Information	<ul style="list-style-type: none"> <li>• Individualized overview of 'what are my "usual" symptoms'</li> <li>• General information about COPD, CHF, anxiety, depression, ischemic heart disease, nutrition, physical exercise</li> <li>• Link to instruction videos about inhalation technique for all inhaler devices</li> <li>• Individualized written action plan on which the automated decision support system was based</li> <li>• Copy of patient's informed consent form</li> </ul>
Exercise and physical activity	<ul style="list-style-type: none"> <li>• Instruction videos with exercises specified for patients with CHF and COPD: relaxation exercises, exercises to maintain fit, breathing exercises during deterioration of symptoms</li> <li>• Graph with number of steps taken per day, measured by a Fitbit®</li> <li>• Videos with general information about COPD</li> </ul>
Embodied Conversational Agent	<ul style="list-style-type: none"> <li>• An Embodied Conversational agent (ECA) - defined as a more or less autonomous and intelligent software entity with an embodiment to communicate with the user<sup>38</sup> – gave reminders on completing the to-do list, feedback on inhaled medication adherence and technique, and recommendations to promote healthy behavior</li> </ul>

Notes: <sup>a</sup>Only for patients using an add-on inhaler sensor (Respiro®).

## Outcomes

The primary outcome of this study was adherence to different components of the eHealth self-management intervention: completing digital daily symptom diaries, following the advised actions, and using inhaled medication. Adherence was measured starting from the patient's individual self-management training session (first two weeks of October 2018), till their last completed diary (last week of January/first week of February 2019). Adherence to daily diary completion was reported per month and per patient, calculated by dividing the total number of diary completions by the total number of follow-up days. Patient's adherence to the advised actions was calculated by dividing the number of performed actions by the total number of actions that were advised by the automated decision support system. Inhaled medication adherence levels were calculated by dividing the number of times patients used their inhaler by the number of times they should have used their inhaler according to their physician's prescription.

The patient's inhalation technique, evaluated by the frequency of correct inhalations and by the different types of errors that were made per patient, was a secondary outcome parameter of this study. The correct inhalation technique was defined as duration of flow >1.25 seconds, Peak Inspiratory Flow > 30 L/min, orientation of the inhaler between 45° and 135°, and closing the cap of the Ellipta® inhaler properly. These cut-off values were based on literature and expert opinion.<sup>37</sup> In addition, paper questionnaires at baseline and after a maximum of four months of follow-up were used to assess other secondary outcomes: health-related quality of life (St. George Respiratory Questionnaire,<sup>39</sup> Minnesota Living with Heart Failure Questionnaire<sup>40</sup>), COPD self-management behavior and knowledge (Partners in Health questionnaire<sup>41</sup>), COPD self-efficacy (COPD Self-Efficacy Score<sup>42</sup>) and anxiety/depression score (HADS<sup>35</sup>). Due to the low number of participants and the short follow-up time of this pilot study, we decided a-priori not to analyze differences between baseline and follow-up data using statistical tests.

## Statistical Analyses

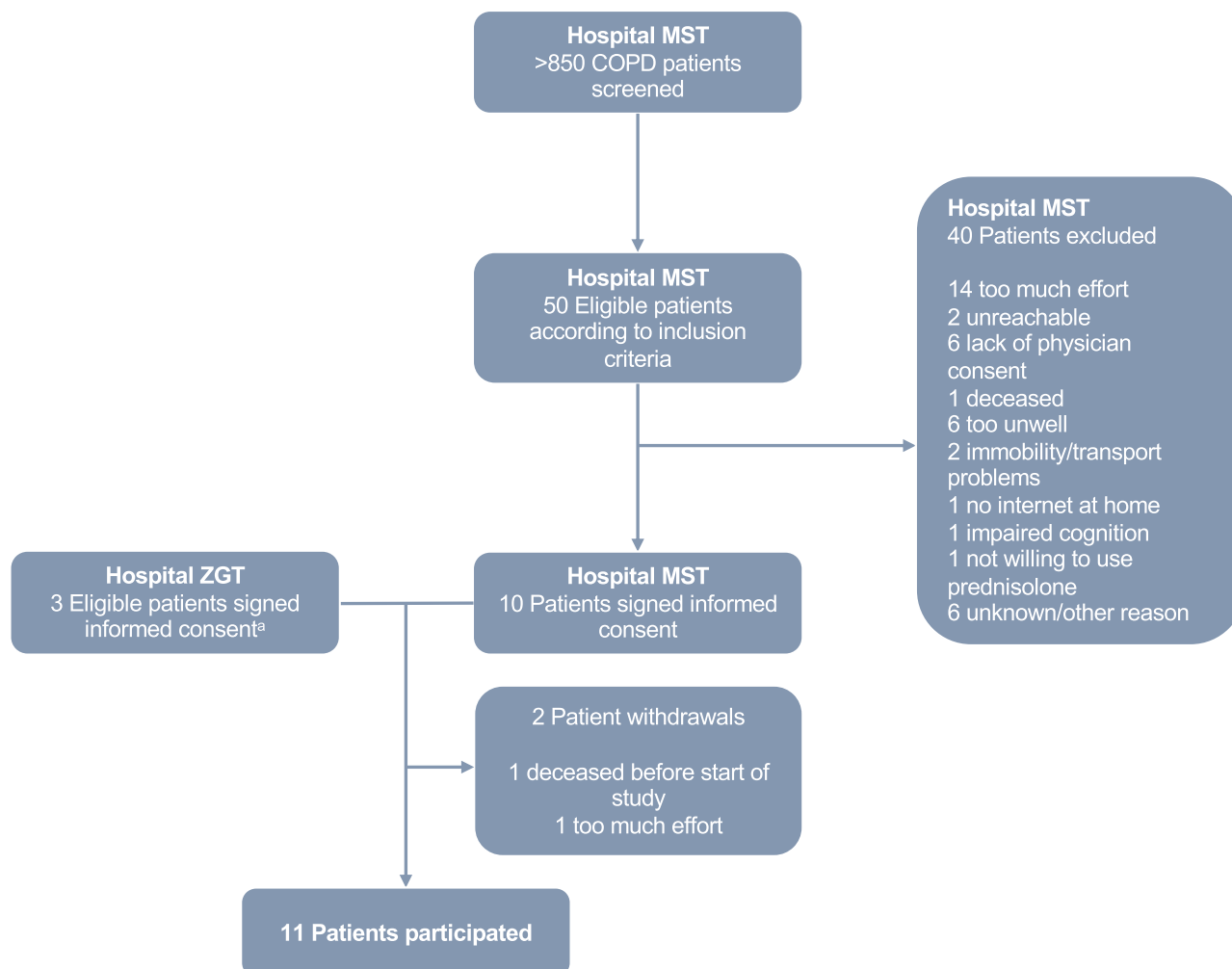
Descriptive statistics were used to report numbers and percentages, means and standard deviations (SD) and medians and interquartile ranges (IQR) of baseline characteristics. Numbers and percentages were calculated for adherence to daily symptom diary completion, following up of advised actions and inhaled medication, number of correct inhalations and types of inhalation errors. These analyses were performed using SPSS version 22. Daily data on inhalation adherence and inhalation technique were delivered by Amiko and subsequently analyzed by SPSS version 22.

## Results

### Patient Characteristics

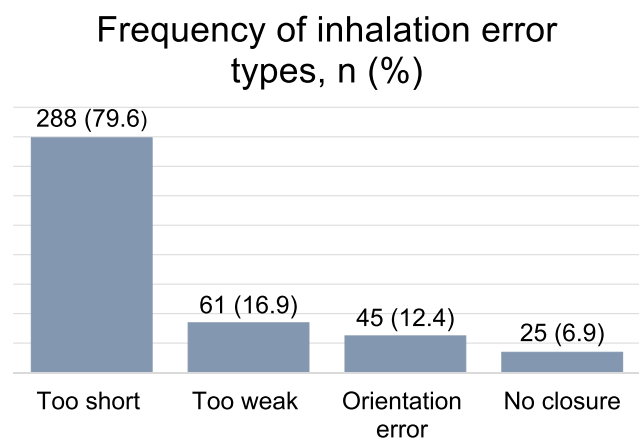
Fifty patients of hospital MST met the study criteria. Of these patients, 40 were excluded for reasons detailed in

Figure 1, Figure 2. Three patients of hospital ZGT were included. Unfortunately, data on screened patients in hospital ZGT were not assessed and are therefore not available. In total, thirteen patients signed informed consent. One patient deceased before the start of the study and another patient decided to withdraw from participation after the first group self-management training session. The remaining 11 patients that completed the training and follow-up had a mean age of  $66.8 \pm 2.9$  years, moderate (55%,  $n=6$ ) to severe (45%,  $n=5$ ) COPD, five (46%) patients had midrange left ventricular function (LVF) and three (27%) patients had reduced LVF (Table 3). Three (27.3%) patients had ischemic heart disease and two (18.2%) had diabetes mellitus as comorbid diseases. Eight patients (72.7%) had at least one COPD exacerbation and six patients (54.5%) had increased the dose of diuretics because of decompensated CHF in the



**Figure 1** Flowchart of patient enrollment.

**Notes:** <sup>a</sup>For hospital ZGT detailed information on screened patients is not available.



**Figure 2** Frequency of inhalation error types. Definition of inhalation errors: Too short: flow <1.25s, Too weak: Peak inspiratory Flow (PIF) <30L/min, Orientation error: Orientation of the inhalator <45° or >135°, No closure: Cap of the inhaler was not closed properly.

**Notes:** Total percentage of errors >100% because during 57 inhalations a combination of errors was made. The “Orientation error” is probably an underestimation because the add-on sensor does not measure the orientation of the inhaler in the following cases: flow <0.25 s, PIF <25 L/min and if the cap of the inhaler was not closed. This was the case during 170 inhalations.

previous year. Three patients (27.3%) had a low digital health literacy score (eHealth Literacy Scale (eHEALS) <26). Six patients (54.5%) were familiar with using a tablet at home.

## Adherence to Daily Symptom Diary

1148 (91.0%) out of 1261 daily diary entries were completed on the actual day (Table 4). Nine patients showed total adherence rates above 95%, and the other two patients completed 59–64% of the diaries on the same day. Because of a system failure, 23 diary days were excluded from adherence analyses as patients were not able to adhere to their intervention for 2 days in a row (for one patient: 3 days). Because of another system failure, diaries were completed twice for 119 diary days. Only one set of these double daily entries was included in adherence analyses.

## Action Plan Adherence

Seven patients received a total of 24 messages with advised actions because of symptom deterioration (see Table 5). Eleven (46%) of these advised actions were actually performed by the patients. Thirteen actions (54%) were not performed, of which six were “call the case manager”. In five cases, patients decided not to perform the action that was advised by the self-management application after having contact with a healthcare provider

and/or because they were hospitalized. An overview of action plan adherence differentiated per advised action and reasons for non-adherence is shown in Table 5.

In six cases (patients n=4) patients increased the dose of diuretics in the absence of an automated advice by the eHealth self-management application. These patients reported significantly increased symptoms during one day instead of two days (n=1), slightly increased instead of significantly increased symptoms (n=4) and/or a gradual instead of a significant weight increase (n=4). In six cases (patients n=5), patients started a course of prednisolone and/or antibiotics, while not being advised by the self-management application to do this. It should be noted that this was always done after consulting the case manager or general practitioner. Symptoms of chest pain occurred twelve times in two patients. Because of a fault in the self-management application, the patients were unable to see the advised action “take nitroglycerine” after reporting their chest pain. No adverse effects (eg hospitalization, death) were reported as a result of this system failure.

## Inhalations

Adherence to taking inhaled medication and using the correct inhalation technique was measured in seven patients. Three of these patients had already used an Ellipta® Inhaler before the start of this study. Prior to study participation, four patients were switched to Ellipta® in agreement with their respiratory physician. All seven patients had to use their inhaled maintenance medication once a day according to their physician’s prescription.

### Inhalation Adherence and Technique

Patients showed an inhaled medication adherence of 98.4% (752 out of 764 days). The range of performed inhalations between patients was 96.3–100%. 390 (51.9%) inhalations were performed correctly. The percentage of correct inhalations varied between and within patients (Table 6). Inhaling too shortly (<1.25 s) was by far the most made inhalation error (79.6%). Inhalation errors are detailed in Figure 1. The add-on sensor of two patients had a low battery for a total of 32 days. Adherence and inhalation technique could therefore not be measured during these days. Double doses (n=10, 1.3% of the total inhalations) were not taken into account in our analyses.

## Secondary Outcomes

The mean and median scores of baseline and follow-up assessments of the secondary outcomes are presented in Table 7.

**Table 3** Baseline Characteristics of Participants Who Completed Follow-Up

A. General baseline characteristics (n=11)	
General characteristics	
Age, mean $\pm$ SD	66.8 $\pm$ 2.9
Gender (male), n (%)	7 (63.6)
Partner (yes), n (%)	8 (72.2)
Tablet at home (yes), n (%)	6 (54.5)
Digital Health Literacy (eHEALS score <sup>43</sup> ), mean $\pm$ SD <sup>a</sup>	27.5 $\pm$ 5.5
Educational level, n (%)	
Secondary Education	2 (18.2)
Trade School	6 (54.5)
Higher Professional education	1 (9.1)
Pre-university education	1 (9.1)
University	1 (9.1)
B. Baseline Characteristics regarding COPD (n=11)	
COPD characteristics	
FEV1%, median (IQR)	51 (43–59)
GOLD classification, n (%) <sup>7b</sup>	
2A	1 (9.1)
2B	3 (27.3)
2C	2 (18.2)
3A	1 (9.1)
3B	1 (9.1)
3D	3 (27.3)
mMRC dyspnea score, n (%) <sup>44</sup>	
1	4 (36.4)
2	4 (36.4)
3	2 (18.2)
4	1 (9.1)
Exacerbations COPD in previous year, n (%) <sup>c</sup>	
0	3 (27.3)
1	3 (27.3)
2	2 (18.2)
3	3 (27.3)
Hospitalizations COPD in previous year, n (%) <sup>d</sup>	
0	10 (90.9)
3	1 (9.1)
C. Baseline characteristics regarding CHF and comorbidities (n=11)	
Chronic Heart Failure (CHF) characteristics	
Left ventricular function, n (%) <sup>e</sup>	
Preserved LVEF	2 (18.2)
Midrange LVEF	5 (45.5)
Reduced LVEF	3 (27.3)

(Continued)

**Table 3** (Continued).

Exacerbations CHF in previous year, n (%) <sup>f</sup>	
0	5 (45.5)
1	4 (36.4)
2	1 (9.1)
3	1 (9.1)
Hospitalizations CHF in previous year, n (%) <sup>g</sup>	
0	8 (72.7)
1	2 (18.2)
3	1 (9.1)
Characteristics comorbidities	
Diabetes Mellitus, n (%)	2 (18.2)
Ischemic Heart Disease, n (%)	3 (27.3)
HADS Anxiety, n (%)	
0–7 (no anxiety disorder)	8 (72.7)
8–10 (possible anxiety disorder)	3 (27.3)
HADS Depression, n (%)	
0–7 (no depressive disorder)	10 (90.9)
8–10 (possible depression disorder)	1 (9.1)

**Notes:** <sup>a</sup>eHEALS, Range of total scores: 8 (low eHealth literacy) to 40 (high eHealth literacy). <sup>b</sup>According to GOLD guidelines 2021.<sup>7</sup> <sup>c</sup>Number of courses of prednisolone/antibiotics for deterioration of COPD symptoms one year prior to participation according to patient questionnaire data. <sup>d</sup>Number of hospitalizations for COPD according to electronic health record data. <sup>e</sup>Preserved LVEF: LVEF  $\geq$ 50%, Midrange LVEF: 40–49%, reduced LVEF <40%.<sup>3</sup> Data is missing for one patient because of inconclusive results of the echocardiogram. <sup>f</sup>Number of increased diuretics doses for deterioration of CHF symptoms one year prior to participation, according to patient questionnaire data. <sup>g</sup>Number of hospitalizations for CHF according to electronic health record data.

**Abbreviations:** COPD, Chronic Obstructive Pulmonary Disease; eHEALS, eHealth Literacy Scale; FEV1%, percentage of predicted Forced Expiratory Volume in 1 second; GOLD, Global Initiative For Chronic Obstructive Lung Disease; mMRC, modified Medical Research Council; IQR, interquartile range; CHF, Chronic Heart Failure; LVEF, Left Ventricular Ejection Fraction; HADS, Hospital Anxiety and Depression Scale; IQR, interquartile range.

## Discussion

In this pilot study, we have tested an eHealth self-management intervention for patients with both COPD and CHF. Whereas our study results showed high patient adherence to completing daily symptom diaries, more than half of the advised actions were not performed by patients, especially the action “call the case manager”. Patients showed high adherence to taking inhaled medication, but their inhaled medication technique was poor.

Patient adherence to completing a symptom diary on a daily basis was high (91%). This was in line with previous findings from studies on eHealth self-management interventions in patients with only COPD.<sup>31,33</sup> Despite including a more complex patient group in our study, the



**Table 4** Adherence to Daily Symptom Diary per Month and in Total

	Number of Diary Days	Diary Completion the Actual Day, n (%)	Diary Completion Next Day or Later, n (%)	Diary Not Completed, n (%)
Month 1 (October)	242	225 (93.0)	1 (0.4)	16 (6.6)
Month 2 (November)	330	312 (94.5)	3 (0.9)	15 (4.5)
Month 3 (December)	339	296 (87.3)	9 (2.7)	34 (10.0)
Month 4 (January)	334	301 (90.1)	15 (4.5)	18 (5.4)
Month 5 (February)	16	14 (87.5)	0	2 (12.5)
<b>Total</b>	<b>1261</b>	<b>1148 (91.0)</b>	<b>28 (2.2)</b>	<b>85 (6.7)</b>

**Notes:** Adherence was measured starting from the patient's individual self-management training session (first two weeks of October 2018), till their last completed diary (last week of January/first week of February 2019). 23 diary days were excluded because the system failed. For 119 diary days, diaries were completed twice because of a system failure. Only one set of these double daily entrees was included.

**Abbreviations:** n, Number of days, (%) percentage of total.

adherence to the diary remained high. Real-time reminders to complete the diary might have positively influenced adherence to symptom diaries. Overall patient adherence to advised actions was relatively low, especially “call the case manager for support”. Non-adherence to actions as self-initiating prednisolone could, however, often be explained for valid reasons (eg, patients already started treatment after consultation with the case manager or patients already received in-hospital treatment). Most

patients ignored the advice to call the case manager because of dizziness. Whereas patients did report their dizziness in the diary, it might not have felt serious enough for them to call the case manager. Emphasizing when patients with dizziness problems should call a case manager (eg, the severity of dizziness that might suggest health deterioration or medication side effects) during the self-management training sessions and providing this information on the eHealth self-management application, might

**Table 5** Action Plan Adherence During Follow-Up Time

Required Action According to Action Plan	Number of Advised Actions	Performed Actions	Unperformed Actions	Reasons Actions Not Performed
Initiate prednisolone course	2	1	1	- Treatment started 2 days ago after consult with case manager (n=1)
Initiate prednisolone and antibiotic courses	3	1	2	- Hospitalized (n=1) - Diary was completed incorrectly and therefore action not performed in agreement with case manager (n=1)
Increase diuretic dose	3	2	1	- Hospitalized (n=1)
Have a NT-proBNP Lab-test	2	0	2	- Hospitalized (n=1) - Unclear (n=1)
Perform relaxation exercises	1	0	1	- Unclear (n=1)
Contact case manager <sup>a</sup> Reason: - Dizziness - Symptoms did not improve - Symptoms and dizziness <sup>b</sup>	10 1 2	5 1 1	5 0 1	- Phone connection failed (n=1) - Unclear (n=5)
Total number of actions	24	11 (46%)	13 (54%)	

**Notes:** <sup>a</sup>Patients were advised to call the case manager in case they reported dizziness and/or symptoms did not improve after two days of self-treatment. <sup>b</sup>Reported dizziness and symptoms did not improve after two days of self-treatment.

**Abbreviation:** %, percentage of total number of actions.

**Table 6** Inhaled Medication Adherence and Technique per Patient During Follow-Up

Patient	Performed inhalations n (%) <sup>a</sup>	Correct inhalations n (%) <sup>b,c</sup>	Range in correct inhalations per week (%)
1*	107 (100)	76 (71.0)	14 – 100
2*	116 (100)	25 (21.6)	0 – 57
3*	111 (98.2)	69 (62.2)	14 – 100
4	105 (96.3)	50 (47.6)	0 – 71
5*	110 (97.3)	76 (69.1)	14 – 100
6	112 (97.4)	19 (17.0)	0 – 57
7	91 (100)	75 (82.4)	71– 100
<b>Total</b>	<b>752 (98.4)</b>	<b>390 (51.9)</b>	

**Notes:** \*Switched to inhaler device Ellipta<sup>®</sup> at the start of the study. <sup>a</sup>Missing data on inhalations because of low battery: patient 4: 5 inhalations, patient 7: 27 inhalations. <sup>b</sup>A correct inhalation is defined as flow >1.25 s, Peak Inspiratory Flow >30 L/min, Orientation between 45° and 135°. <sup>c</sup>Missings because inhalation not performed or low battery: patient 3: 2, patient 4: 9, patient 5: 3, patient 6: 3, patient 7: 27.

improve patients' adherence to contact the case manager for dizziness problems. Moreover, reducing the threshold to call the case manager by improving the availability of case managers' support (eg by using a chat service) might improve adherence. In our study, some patients also initiated actions without an advice from the self-management application, showing their willingness to self-manage their disease. For example, doses of diuretics were increased by patients themselves, while CHF symptoms were only slightly increased or present for only one day. Furthermore, prednisolone and antibiotics were initiated after having contact with a healthcare provider because of increased symptoms. We know that patients use experiential knowledge to recognize exacerbations, we could therefore assume that decisions to start self-treatment without being prompted by the eHealth application may have been driven by previous experiences with

exacerbations.<sup>45</sup> This might suggest that, to improve adherence to action plans and adequate timing of self-treatment, action plans for the self-treatment of symptoms should be further tailored to previous experiences of the individual patient. For example, for specific patients, self-treatment could be started after one day instead of two days of significantly increased symptoms, taking into account the risk of side effects as low blood pressure, with diuretics.

Previous studies showed that more frequent follow-up contact initiated by the case manager increases adherence to action plans by improving disease awareness.<sup>21,22</sup> These contacts with a case manager could be beneficial for the patients in our study as well. However, it might influence the degree to which patients self-manage their disease as patients might become too reliant on their case manager and thereby eliminate the effect of the self-management intervention. However, this was not assessed in our study. The low adherence to advised actions also raises the question whether patients felt completely confident with following up the advice of an eHealth application. This is supported by the findings of a qualitative study on perceptions of patients and healthcare providers on using mobile Health (mHealth) for the self-management of COPD, showing that some patients had limited trust in the advice of mHealth interventions.<sup>46</sup> Moreover, there is a growing body of literature that suggests that the individual patient's needs of COPD self-management (depending on, eg culture, norms and values) misalign with the healthcare providers expectations of behavior change of the patient, which may influence patient adherence.<sup>17,47</sup> Furthermore, the adoption of the self-management intervention by the healthcare provider is highly important and is dependent on the particular healthcare provider's perceived value and

**Table 7** Quality of Life at Baseline and After Follow-Up

Questionnaire	Baseline	Follow-Up (4 Months)
COPD self-efficacy score, mean (SD) <sup>a</sup>	75.2 (26.6)	73.9 (22.8)
Partners in Health, mean (SD) <sup>b</sup>	81.7 (8.1)	86.4 (4.0)
MLHFQ, median (IQR) <sup>c</sup>	37 (3–66)	34 (0–52)
SGRQ, median (IQR) <sup>d</sup>	45.0 (9.2–68.2)	43.1 (6.9–64.0)
HADS anxiety, median (IQR) <sup>e</sup>	5 (2–10)	5 (0–12)
HADS depression, median (IQR) <sup>e</sup>	5 (1–10)	4 (1–9)

**Notes:** Missings: COPD self-efficacy score: 4 patients, MLHFQ: 3 patients. <sup>a</sup>Range of total scores: 34 (low COPD self-efficacy) to 170 (high COPD self-efficacy). <sup>b</sup>Range of total scores: 0 (low self-management behavior and knowledge) to 96 (high self-management behavior and knowledge). <sup>c</sup>Range of total scores: 0 (high heart failure related quality of life) to 105 (low heart failure related quality of life). <sup>d</sup>Range of total score: 0 (high respiratory related quality of life) to 100 (low respiratory related quality of life). <sup>e</sup>HADS: Score 0–7= no anxiety/depression disorder, score 8–10= possible anxiety/depression disorder, score 11–21= probable anxiety/depression disorder.

**Abbreviations:** MLHFQ, Minnesota Living with Heart Failure Questionnaire; SGRQ, St. George Respiratory Questionnaire; HADS, Hospital Anxiety Depression Score.

needs, which should be aligned with the patient's needs and preferences before implementing.<sup>48</sup>

Adherence to inhaled medication was high for all patients (98.4%). This is in contrast to literature,<sup>49–51</sup> possibly because in our study patients received reminders via the Embodied Conversational Agent (ECA) and via audio-visual signs by the sensor to take their inhaled medication. Despite the inhalation instruction provided by the case manager, also on the use of the add-on inhaler sensors, the patient's inhalation technique was poor, similar to other studies.<sup>52,53</sup> More time should therefore be taken for an individual inhalation instruction at the start of the study.<sup>54</sup> This should be repeated during follow-up within the self-management application and/or by the case manager. Also, feedback of the ECA on inhalation medication was not specified on error type and, because of technical problems with the sensor, not always correct (eg a performed inhalation was wrongly registered by the sensor as unperformed, leading to incorrect feedback to the patient by the ECA). In the future, prompt, specified and correct information about individual inhalations that is recorded by the sensor, may be helpful to improve inhalation technique.

In addition to the quantitative results we obtained, this pilot study taught us more about how to further develop the eHealth self-management intervention for patients with multiple chronic conditions, such as COPD and CHF. We experienced that, especially at the start of the study, patients were focused on how to use the tablet and various devices that were connected to this tablet. This might have distracted them from (learning how to) self-report symptoms and follow-up advices.<sup>55</sup> It suggests that some patients need more intensive training and/or an adaptation period before starting the actual self-management intervention. Before the start of a self-management intervention, it is important to assess patient readiness to change behavior, as it is related to adherence to the intervention.<sup>8,56</sup> Motivational interviewing could be used to improve patient readiness.<sup>57</sup> Further, we feel that for patients that suffer from severe COPD and/or CHF, more intensive case manager support (by phone or face-to-face) might be necessary to differentiate between symptoms of COPD and/or CHF. In case of significantly increased symptoms in these patients, the automated decision support system should probably give advice towards case manager contact instead of self-treatment. Further, we aimed to support patients in differentiating between COPD and CHF breathlessness symptoms in case

breathlessness symptoms did not improve after starting self-treatment by adding the laboratory test NT-proBNP to the action plan. During this study, patients were only twice advised to have the lab-test NT-proBNP measured. In both cases, these actions were not performed. A longer follow-up time and a larger group of patients are therefore needed to evaluate the usefulness of NT-proBNP as part of a multi-morbid exacerbation action plan. Moreover, an easy-to-use point-of-care test for measuring NT-proBNP at home in these patients with health deterioration could be advantageous to improve adherence to measuring NT-proBNP.

The limitations of our study also gave us insight into how to further develop the eHealth self-management intervention. First, although including a small patient group fits to the pilot phase of our study, we included less patients than expected. This has limited the generalizability of our results. Many patients in hospital MST declined participation because of logistic issues (eg not being able to visit self-management training because of immobility and transport problems), not being familiar with using a tablet, and because patients presumed it was too much effort. Unfortunately, for ZGT patients this information was not reported. Offering digital and/or face-to-face self-management training sessions at home for a specific group of patients who are immobile or who have no experience in using a tablet might increase the number of patients willing to participate in eHealth self-management interventions. Second, we did not assess (e)health literacy and cognitive impairment during patient enrollment. Future studies should not only measure (e)health literacy and cognitive impairment at enrollment but ideally eHealth self-management interventions should also be adjustable to (e)health literacy and cognitive impairment. This will increase its applicability and uptake in patients with limited (e)health literacy and cognitive impairment.<sup>24,55,56,58</sup> For example, more and tailored self-management training sessions and a simplified version of the eHealth self-management intervention could be offered to these patients. Third, the use of home monitoring devices (add-on sensors for an inhaler device, a smart weighing scale and an activity sensor) led to connection and low-battery problems. In addition, due to system failures, the self-management application could not be used for two days, some diaries were completed double, and the action "take nitroglycerine" in case of chest pain did not appear. Whereas this was recognized as a serious safety issue in this pilot study, fortunately no adverse events were

reported related to this. Although the technical issues correspond to the pilot phase of the technology and technology readiness level,<sup>59,60</sup> it led to some frustrations and distrust amongst patients regarding the eHealth self-management intervention. In follow-up studies, the technology should therefore be intensively tested upfront together with its end users (ie patients, technicians, health-care providers) to ensure its suitability for larger-scale summative evaluation.<sup>61</sup> Finally, a physical activity program could be added to our future eHealth self-management intervention to optimize and preserve physical health.<sup>8,62</sup>

Our results indicate that the eHealth self-management intervention should be further adapted to the needs and competences of the individual patient, which is also suggested by several qualitative studies.<sup>8,46,55,63</sup> This will increase patient adherence, and thereby potentially improve individual health outcomes. Also, the eHealth self-management application itself should be further developed towards an application with a higher technology readiness level, suitable for larger-scale follow-up studies to investigate the clinical added value.<sup>59,60</sup> In addition, further analysis of both clinical and home monitoring data of eHealth self-management interventions can increase our understanding of the development and onset of disease progression and exacerbations of COPD and CHF. It can provide insight into day-to-day fluctuations in COPD and CHF symptoms and behavior (eg adherence), and thereby work towards preventive chronic disease self-management for the patient with COPD and CHF.<sup>64–66</sup>

## Conclusion

Whereas adherence to completing daily diaries was high as part of our eHealth self-management intervention for patients with both COPD and CHF, advised actions were inadequately followed-up, particularly the action “call the case manager”. Inhaled medication adherence was also high, but inhalations were poorly performed. This pilot study gives insight into how to further develop the eHealth self-management intervention. For this, personalizing and tailoring to an individual patient’s needs and competences are essential. Future quantitative and qualitative analyses are necessary to unravel patient adherence and evaluate the effects of a further developed eHealth self-management intervention on health outcomes.

## Abbreviations

eHEALS, eHealth Literacy Scale; FEV1%, FEV1 percentage of predicted; FVC, Forced Vital Capacity; GOLD, Global

Initiative For Chronic Obstructive Lung Disease; mMRC, modified Medical Research Council; COPD, Chronic Obstructive Pulmonary Disease; CHF, Chronic Heart Failure; ECA, Embodied Conversational Agent; eHealth, electronic Health; HADS, Hospital Anxiety and Depression Scale; IQR, interquartile range; LVEF, Left Ventricular Ejection Fraction; n, number; IQR, interquartile range.

## Data Sharing Statement

Deidentified participant data are stored in archived datasets for a total of 15 years and can be requested from the corresponding author.

## Ethics Approval and Informed Consent

This study was approved by the Medical Ethical Committee of Twente (NL17-17). Informed consent for participation was given by all participants.

## Consent for Publication

All authors have seen the content of this article and have given consent for publication.

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## Author Contributions

All authors made a significant contribution to the work reported, whether that is in study conception and design, acquisition of data, analysis and interpretation of data, drafting of the manuscript, critical revision and review of the manuscript or acquisition of funding. All authors gave final approval of the version to be published, have agreed on the journal to which the article has been submitted, and agree to be accountable for all aspects of the work.

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## Disclosure

Martijn Grinovero is an employee (CCO) at Amiko Digital Health Limited. Dr Anke Lenferink reports grants from Pioneers In Healthcare Innovation Fund 2016, during the conduct of the study. The other authors did not report any conflicts of interest related to this work.

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