

Effect of tele–health care on quality of life in patients with severe COPD: a randomized clinical trial

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Background and objective: Telemonitoring (TM) of patients with COPD has gained much interest, but studies have produced conflicting results. We aimed to investigate the effect of TM with the option of video consultations on quality of life (QoL) in patients with severe COPD.

Patients and methods: COPD patients at high risk of exacerbations were eligible for the 6-month study and a total of 281 patients were equally randomized to either TM (n=141) or usual care (n=140). TM comprised recording of symptoms, oxygen saturation, spirometry, and video consultations. Algorithms generated alerts if readings breached thresholds. Both groups filled in a health-related QoL questionnaire (15D[®]) and the COPD Assessment Test (CAT) at baseline and at 6 months. Within-group differences were analyzed by paired *t*-test.

Results: Most of the enrolled patients had severe COPD (86% with Global Initiative for Chronic Obstructive Lung Disease stage 3 or 4 and 45% with admission for COPD within the last year, respectively). No difference in drop-out rate and mortality was found between the groups, and likewise there was no difference in 15D or CAT at baseline. At 6 months, a significant improvement of 0.016 in 15D score ($p=0.03$; minimal clinically important difference 0.015) was observed in the TM group (compared to baseline), while there was no improvement in the control group -0.003 ($p=0.68$). After stratifying 15D score at baseline to <0.75 or ≥ 0.75 , respectively, there was a significant difference in the <0.75 TM group of 0.037 ($p=0.001$), which is a substantial improvement. No statistically significant changes were found in CAT score.

Conclusion: Compared to the nonintervention group, TM as an add-on to usual care over a 6-month period improved QoL, as assessed by the 15D questionnaire, in patients with severe COPD, whereas no difference between groups was observed in CAT score.

Keywords: COPD, tele–health care, video consultations, quality of life

Introduction

Trailing in the wake of decline of lung function in COPD patients, diminishing physical, emotional, and social functioning leads to a deterioration in quality of life (QoL).¹ Furthermore, a substantial prevalence of psychological distress further adds to the considerable adverse effects on QoL.^{2,3}

Telemonitoring (TM) may have the potential to alleviate some of the burdens that COPD places on patients QoL, not only by providing rapid medical intervention in the early phase of deterioration in the patient's condition, and by that prevent hospital admissions, but also a way of reducing costs and time spent in transportation.⁴

To date, TM has been assessed in several different designs and setups, but results are conflicting, though mostly negative. Review articles published to date have struggled

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in extracting significant conclusions and, as it goes, have recommended additional and more extensive studies.⁵⁻⁷

Three large studies have examined TM as an alternative to either home or outpatient clinic visits, with differing results. Pinnock et al⁸ and Trappenburg et al⁹ published their findings from 2 large randomized studies with COPD patients and reported that TM had no significant effects on QoL. McDowell's et al¹⁰ reported from a large 2-center randomized trial that TM has a positive impact on health-related QoL and anxiety. As TM involves substantial healthcare costs and commitment, there is a need for well-designed longer-term studies with appropriate follow-up to determine whether it is suitable for widespread application.^{7,11,12} Further studies are, therefore, clearly needed in order to establish whether or not TM has clinically relevant impact on QoL in patients with COPD.

The present randomized study aimed to investigate the impact of TM with the option of video consultations on QoL in patients with severe and very severe COPD recruited from respiratory outpatient clinics.

Patients and methods

Eligible patients

The NetKOL study was a randomized controlled trial recruited patients from 4 hospitals with specialized pulmonary wards in Copenhagen (Hvidovre, Bispebjerg, Herlev, and Amager Hospitals).

The target population were patients with severe and very severe, stable COPD at high risk of exacerbations and hospital admissions. The inclusion criteria were as follows:

1. A diagnosis of COPD defined according to the Global Initiative for Chronic Obstructive Lung Disease criteria¹³ (post-bronchodilator forced expiratory volume in 1 second [FEV₁]/forced vital capacity <0.7).
2. Post-bronchodilator FEV₁ <60% of predicted (pred) value.
3. Hospital admission due to COPD exacerbation within the previous 36 months and/or prescribed long-term oxygen therapy (LTOT) due to chronic respiratory failure for at least 3 months.
4. Regularly scheduled visits to the respiratory outpatient clinics.
5. COPD judged by the study staff as the leading cause of disability.
6. Residents in 1 of 6 municipalities in the Copenhagen area and the catchment area of 1 of the 4 recruiting hospitals.

The exclusion criteria were as follows:

1. An exacerbation of COPD within the 3 weeks before enrollment.

2. Unwilling to provide informed consent for participating in the study.
3. Unable to use a tablet computer.
4. Planned stay outside the catchment area for 2 weeks or more during the study period.
5. Inability to participate due to language barrier or cognitive disorders.
6. Impossible to setup a functioning telephone line.

In case of hospital admission, patients continued in the study immediately after discharge from the hospital.

Patient recruitment and randomization

Over a 6-month inclusion period, we identified 560 eligible patients, of whom 281 (51.2%) patients gave written informed consent to study participation and were randomized to either TM or usual care (control group) with a 1:1 allocation for 6 months. Randomized patients were divided into 3 categories:

Group 1 – Hospital admission for COPD in the previous year.

Group 2 – Hospital admission for COPD in the past 3 years, but not within the year before enrollment.

Group 3 – LTOT regardless of the number of hospitalisations for COPD in previous years.

Block randomization was used to ensure equal numbers of patients from the 3 groups were allocated to TM and usual care.

Standard management of COPD

All trial patients were managed and treated according to national and international guidelines. Usual care included outpatient pulmonary rehabilitation for patients with FEV₁ <50% predicted and Medical Research Council dyspnea score ≥3¹⁴ and supported discharge to selected in-hospital patients to reduce the risk of early readmission.¹⁵

Usual care group

Patients were seen for scheduled visits at the outpatient clinics once or twice a year and for unscheduled visits as required. In case of acute respiratory problems, patients could contact the staff at the outpatient clinic Monday to Friday, during the daytime. Depending on their mobility, patients on LTOT were seen either at the outpatient clinics or at home by a respiratory care nurse.

TM group (intervention group)

Patients randomized to the TM group were not seen at regularly scheduled visits at the outpatient clinics during the 6-month study period, but unscheduled visits were arranged

if the TM consultation was considered inadequate by the health care professional.

The TM equipment comprised a tablet computer with a web camera, a microphone, and measurement equipment (spirometer, pulse oximeter, and bathroom scale). Patients reported changes in dyspnea, sputum color, volume, and purulence on the tablet computer. Measurements, including lung function, oxygen saturation, and body weight, and video consultations, together with the other measurements, were conducted at regular intervals, ie, once a week for the first 4 weeks of the study period and then every 4 weeks.¹⁶ Consultations without video, but with the other measurements, were performed every week throughout the study period. Patients were free to carry out additional measurements at any time or phone the call center during open hours. The observations were transferred to the call center at each participant's local hospital and automatically categorized and prioritized (ie, coded green, yellow, or red), and aberrant measurements were responded to by the specialist nurse either by phone or video consultation and if relevant conferred with a respiratory specialist. All measurements were stored in a secured database; access was granted to the patient's general practitioner and the municipal's social and health workers.

A more detailed description of the design of the present study is given in Ringbaek et al.¹⁶

Measurements of outcome variables

To measure the QoL, we selected 2 questionnaires, the 15D[®] and the COPD assessment test (CAT). The 15D questionnaire is a standardized 15-dimensional self-administered generic questionnaire. It assesses mobility, vision, hearing, breathing, sleeping, eating, speech, excretion, activities (usual), mental function, discomfort and symptoms, depression, distress, vitality, and sexual activity. There are 5 levels of each area of assessment (scored 1–5), and the resulting score is between 0 and 1. One is equal to perfect health and well-being; a score of 0 is equal to a state of death.¹⁷ The 15D questionnaire has previously been found to be comparable to the AQ20 score as a generic measure of health-related QoL.¹⁸ The CAT, a disease-specific health status questionnaire, provides a measure of disease severity from the patient's perspective. It contains 8 questions (cough, phlegm, chest tightness, breathlessness, activities, confidence, sleep, and energy) that pertain to a patient's COPD symptom burden. Each question consist of a 6-point differential scale, and the final score is between 0 and 40.¹⁹

Outcomes and ethics

The primary outcome was the change in health-related QoL as measured by the 15D questionnaire, with change in CAT

score as the secondary efficacy parameter. Patients completed both questionnaires at baseline and again at 6-months (end of study period).

The study was approved by the Research Ethics Committee of the Capital Region of Denmark (H-4-2013-052).

Statistics

Data were analyzed with the statistical package SPSS version 24.0 (IBM Corporation, Armonk, NY, USA). The Fisher's exact test, unpaired *t*-test, and Mann-Whitney *U* test were used, as appropriate, for comparisons between groups at baseline. For analyses of changes in 15D and CAT scores (including standard deviation), within-group differences were calculated using paired *t*-test. For between-group differences at follow-up, we calculated the values using analysis of covariance to account for baseline 15D and CAT scores. The 0.75 cut-point for 15D was based on the rounded median of the groups. A 2-sided *p*-value of <0.05 was considered significant.

Results

Baseline characteristics

A total of 207 COPD patients (85.7%) enrolled in the study had an FEV₁<50% pred, 148 patients (52.7%) had a Medical Resrach Council dyspnea score ≥4, and 126 patients (44.9%) had at least 1 hospital admission for COPD in the year before enrollment (Table 1). More females were allocated to TM; further comparisons between the 2 groups are given in Table 1.

Study completion and withdrawals

Four patients (3 in the TM group) withdrew their consent within the first 2 weeks of the study period, and 2 patients dropped out due to technical issues. Seventeen patients died during the study period, and 10 patients dropped out for other reasons (technical issues after the first 2 weeks or did not have the energy to carry on anymore). No significant differences in drop-out rates (*p*=0.79) or mortality were found between the groups (*p*=0.87).

15D QoL questionnaire

In total, 140 patients in the TM group and 139 patients in the usual care group completed baseline CAT and the 15D questionnaire with no significant differences in scores between the 2 groups (Table 2). Of the enrolled patients, 123 from the TM group and 123 from the usual care group completed the follow-up assessment questionnaires at 6 months. We found a statistically significant (*p*=0.03) within-group difference of 0.0162 by paired *t*-test in the TM group's change from

Table 1 Baseline characteristics of the 281 patients enrolled in the study and randomized to either TM (n=141) or usual care (n=140)

Characteristics	TM group (N=141)	Usual care group (N=140)	P-value
Females, n (%)	86 (61%)	64 (45%)	0.01 ^a
Age in years, mean (SD)	69.8 (9.0)	69.4 (10.1)	0.75 ^b
BMI, kg/m ² , mean (SD)	24.9 (6.3)	26.9 (7.0)	0.01 ^b
FEV ₁ , % predicted, mean (SD)	34.9 (13.3)	33.8 (12.0)	0.48 ^b
MRC dyspnea score, mean (SD)	3.5 (0.8)	3.7 (0.9)	0.09 ^c
Pack-years, mean (SD)	42.9 (26.4)	41.0 (22.7)	0.72 ^b
LTOT, n (%)	37 (26%)	38 (27%)	0.86 ^a
Charlson comorbidity index, mean (SD)	1.70 (1.49)	1.96 (1.51)	0.13
Living alone	58%	52%	0.27 ^c
Number of hospital admissions for COPD exacerbation last year, mean (range)	0.91 (0–7)	1.22 (0–23)	0.61 ^a

Note: ^aMann–Whitney *U* test. ^bUnpaired *t*-test. ^cFisher's exact test.

Abbreviations: BMI, body mass index; FEV₁, forced expiratory volume in 1 second; LTOT, long-term oxygen therapy; MRC, Medical Research Council; TM, telemonitoring.

baseline (Table 2). We found a statistically significant difference between groups at follow-up; the TM group had a higher mean 15D score of 0.0222 (95% CI: 0.0023–0.0421, $p=0.03$) compared with the usual care group. There was a numerical, but statistically nonsignificant difference in CAT score at follow-up, and the TM group had a mean score of -0.59 (95% CI: -1.86 to 0.67 , $p=0.36$) lower than the usual care group.

After stratifying subjects by 15D score at baseline using a cut-off of 0.75, we found a statistically significant within-group difference in 15D in the TM group having a baseline score below 0.75 of 0.037 ($p=0.001$). The 0.75 cut-point was chosen based on it being the rounded median for the groups. There was no significant change in 15D scores for the above 0.75 TM group or both control groups.

CAT

There was no significant difference in CAT score between groups at baseline, though the TM group had a 1.6 point lower mean CAT score than the control group. There was no significant within-group change between baseline and follow-up in paired *t*-test. Both groups had a similar change in CAT score between baseline and follow-up (Table 3).

Discussion

The present randomized clinical trial revealed that TM with close contact between the patient and the health care providers at the hospital had a positive impact on QoL in patients with more severe COPD. To date, there have been multiple studies looking at the usefulness and potential application of TM. Results from these studies show that TM is not

Table 2 QoL, as assessed by the 15D[®] score, at baseline, follow-up, as change within-group from baseline for the TM and usual care groups and difference at follow-up between TM and usual care

	TM group (N=141)		Usual care group (N=140)	
15D baseline, mean (SD)	0.735 ^a (0.101)		0.726 ^a (0.113)	
15D 6-month follow-up, mean (SD)	0.761 ^b (0.107)		0.728 ^c (0.121)	
Within-group difference, mean	0.016 ^b (0.002–0.031) ^d $p=0.03^e$		-0.003^c (-0.018 to 0.012) ^d $p=0.68^e$	
Between-group difference at follow-up, mean	0.0222 (0.0049–0.0622) ^d $p=0.02^f$			
TM + usual care groups stratified by baseline 15D score	<0.75 (N=62)	≥0.75 (N=60)	<0.75 (N=71)	≥0.75 (N=52)
15D baseline, mean (SD)	0.666 (0.065)	0.824 (0.057)	0.656 (0.076)	0.834 (0.054)
15D 6-months follow-up, mean (SD)	0.703 (SD =0.095)	0.819 (SD =0.083)	0.661 (SD =0.100)	0.819 (SD =0.082)
15D change to 6-month follow-up, mean	0.037 (0.016–0.059) ^d $p=0.001^e$	-0.005 (-0.024 to 0.013) ^d $p=0.569^e$	0.006 (-0.015 to 0.026) ^d $p=0.588^e$	-0.015 (-0.037 to 0.007) ^d $p=0.186^e$

Notes: ^aNonsignificant difference. ^bOnly 122 patients completed follow-up 15D. ^cOnly 123 patients completed follow-up 15D. ^d95% confidence interval of the difference. ^ePaired *t*-test. ^fAnalysis of covariance.

Abbreviations: QoL, quality of life; TM, telemonitoring.

Table 3 QoL, as assessed by the CAT score, at baseline, follow-up and as change within-group from baseline for the TM and usual care groups

	TM group (N=141)	Usual care group (N=140)
CAT baseline, mean (SD)	26.9 ^a (6.7)	28.5 ^a (7.9)
CAT 6-month follow-up, mean (SD)	25.6 ^b (6.9)	27.5 ^c (7.5)
Within-group difference, mean	-0.76 (-1.68 to 0.15) ^d <i>p</i> =0.101 ^e	-0.73 (-1.74 to 0.28) ^d <i>p</i> =0.154 ^e
Between-group difference, mean	-0.59 (-1.86 to 0.67) ^d <i>p</i> =0.36 ^f	

Notes: ^aNonsignificant difference. ^bOnly 122 patients completed follow-up CAT. ^cOnly 123 patients completed follow-up CAT. ^d95% confidence interval of the difference. ^ePaired t-test. ^fAnalysis of covariance.

Abbreviations: CAT, COPD Assessment Test; TM, telemonitoring; QoL, quality of life.

a clear-cut and straightforward solution that will improve all COPD patients' QoL, prevent exacerbations, and reduce hospitals costs. However, there are evident inclinations that TM will have a valuable place in specific, selected circumstances. The question is who can TM be beneficial to and in which areas can we expect TM to be successful? Based on our findings we will discuss the potential place for TM to improve QoL in selected severe and very severe COPD patients.

Changes in QoL

COPD patients in the TM group had a significant, but a small improvement in the QoL.

The minimum important change (MIC) in the 15D questionnaire has previously been evaluated by Alanne et al.²⁰ They found the MIC of 15D to be ± 0.015 and anchored the results to a 5-level subjective global assessment scale (1: much better, 2: slightly better, 3: much the same, 4: slightly worse, and 5: much worse) (Figure 1). The change seen in 15D score in the TM group equates, only just, to a

Minimally important differences in 15D score

1. Much better =>0.035
2. Slightly better =0.015 to -0.035
3. No change =>-0.015 to ≤ 0.015
4. Slightly worse =-0.035 to -0.015
5. Much worse =<-0.035

Figure 1 Clinically significant changes in the 15D[®] QoL score anchored to a global assessment scale of 1–5, by standardized algorithm transformed into an index from 0 to 1.

Abbreviation: QoL, quality of life.

global assessment score of 2 or slightly better. However, we find a group with potential for substantial improvement when selecting patients with a lower than median (<0.75) QoL score as measured by the 15D questionnaire. The TM patients with a lower than median 15D score at baseline had a change equivalent to a global assessment score of 1, or much better, at follow-up. The between-group difference equates to a clinically significant difference. However, there is at present no available study that has examined whether the MIC found by Alanne et al²⁰ applies to between-group differences, but the assumption is that for now the same MIC applies.

Both McDowell et al¹⁰ with Saint George Respiratory Questionnaire (SGRQ) and Hospital Anxiety and Depression Scale (HADS) and Koff et al²¹ with SGRQ found similar improvements in QoL, with a similar TM setup, where monitoring and follow-up are maintained by the same respiratory nurses. Pinnock et al⁸ found an improvement in QoL, but with no difference between treatment arms, ie, TM and control group. In this setup, information was only monitored by the study group, and in case of aberrant findings, they were relayed to the patient's general practitioner and action was at the GP's discretion. In our study and that of McDowell et al,¹⁰ monitoring was performed and followed up by the designated respiratory nurses in each center, in cooperation with a respiratory physician. Trappenburg et al⁹ used the clinical COPD questionnaire to determine the change in health-related QoL and found no significant effect of TM. The clinical COPD questionnaire score is based on the degree of symptoms and physical functioning, but not on aspects such as mood that is registered by the 15D, HADS, and SGRQ questionnaires. As we have found based on our patients' CAT score, TM has no significant effect on symptoms and functioning but improves when accounting for mood and outlook. We speculate that the reason we find a significant improvement, lies in the setup of the TM (that is very different between studies) with regular face-to-face or telephone interaction and follow-up action taken by the same team.

Changes in CAT score

We found no significant difference in CAT score between the TM and the usual care group at follow-up, and showed a similar nonsignificant decrease between baseline and 6-month follow-up. However, this is likely due to dropouts from the trial were patients with the highest baseline CAT score (median 30 for dropouts' vs 27 for completion to follow-up). Seeing as our intervention was merely monitoring and not for a new medicine or a new type of rehabilitation, it is not expected to improve symptom burden.

Conclusion

Our findings suggest that TM has a potential role in improving the QoL of COPD patients overall, despite not improving symptoms and exacerbations, and not least in patients with a lower than median score of QoL. We found that patients who were randomized to TM in addition to usual care showed a clinically and statistically significant improvement in the QoL. Future studies should aim at determining the optimal cut-off point for 15D or other health-related QoL assessments for selecting TM candidates.

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Disclosure

The authors report no conflicts of interest in this work.

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