

ORIGINAL ARTICLE

Nocturnal non-invasive ventilation in COPD patients with prolonged hypercapnia after ventilatory support for acute respiratory failure: a randomised, controlled, parallel-group study

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ABSTRACT

Introduction The effectiveness of non-invasive positive pressure ventilation (NIV) in COPD patients with prolonged hypercapnia after ventilatory support for acute respiratory failure (ARF) remains unclear. We investigated if nocturnal NIV in these patients prolongs the time to readmission for respiratory causes or death (primary endpoint) in the following 12 months.

Methods 201 COPD patients admitted to hospital with ARF and prolonged hypercapnia >48 h after termination of ventilatory support were randomised to NIV or standard treatment. Secondary outcomes were daytime arterial blood gasses, transcutaneous PCO₂ during the night, lung function, health-related quality-of-life (HRQL), mood state, daily activities and dyspnoea.

Results 1 year after discharge, 65% versus 64% of patients (NIV vs standard treatment) were readmitted to hospital for respiratory causes or had died; time to event was not different (p=0.85). Daytime PaCO₂ was significantly improved in NIV versus standard treatment (PaCO₂ 0.5 kPa (95% CI 0.04 to 0.90, p=0.03)) as was transcutaneous PCO₂ during the night. HRQL showed a trend (p=0.054, Severe Respiratory Insufficiency questionnaire) in favour of NIV. Number of exacerbations, lung function, mood state, daily activity levels or dyspnoea was not significantly different.

Discussions We could not demonstrate an improvement in time to readmission or death by adding NIV for 1 year in patients with prolonged hypercapnia after an episode of NIV for ARF. There is no reason to believe the NIV was not effective since daytime PaCO₂ and night-time PCO₂ improved. The trend for improvement in HRQL favouring NIV we believe nevertheless should be explored further.

Trial registration number NTR1100.

INTRODUCTION

Non-invasive positive pressure ventilation (NIV) has become an established treatment in patients with COPD who are admitted to hospital with acute respiratory failure (ARF) due to an exacerbation.¹⁻⁴ As readmission and mortality rates in these patients are high,^{5,6} the application of long-term nocturnal NIV in stable hypercapnic COPD has been subject of several studies. However, in a meta-analysis no clear benefits were documented.⁷⁻¹¹ Reasons for these negative results were: (1) insufficient ventilatory support; (2) lack of proper monitoring during

Key messages

What is the key question?

- What is the effect of nocturnal non-invasive positive pressure ventilation (NIV) alongside standard medical treatment in COPD patients with prolonged hypercapnia after ventilatory support for acute respiratory failure?

What is the bottom line?

- We could not demonstrate an improvement in time to readmission or death by adding NIV for 1 year in COPD patients with prolonged hypercapnia after an episode of NIV for acute respiratory failure.

Why read on?

- NIV did improve daytime PaCO₂ and night-time transcutaneous PCO₂. This and the trend for improvement in health-related quality-of-life favouring NIV should be explored further.

the night; (3) baseline PaCO₂ too low; and (4) low adherence to NIV. The two long-term randomised controlled trials (RCTs) comparing NIV in addition to long-term oxygen therapy (LTOT) showed conflicting results: one demonstrated small improvements in daytime arterial carbon dioxide pressure (PaCO₂), dyspnoea and health-related quality-of-life (HRQL)¹² in favour of NIV whereas the other found an improvement in survival but at the cost of worsening HRQL.¹³

Two other studies investigated the effects of long-term NIV in the most severe subgroup of COPD patients; that is, patients with acute-on-chronic respiratory failure. These indicated a benefit from long-term NIV in patients who previously required mechanical ventilation for treatment of ARF and who remained hypercapnic thereafter.^{6,14} Unfortunately, these were both small studies and given the huge impact of COPD on mortality and morbidity¹⁵ further studies are urgently needed.

We hypothesised that providing nocturnal NIV alongside standard medical treatment to patients who remain hypercapnic after ventilatory support during ARF would prolong the time to readmission

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for respiratory causes or death. Therefore, we conducted an RCT comparing nocturnal NIV with standard treatment for 1 year.

METHODS

The RESCUE trial (REspiratory Support in COPD after acUte Exacerbation) was a multicentre, prospective, randomised controlled study recruiting COPD patients who were admitted to hospital with ARF and treated with ventilatory support throughout the Netherlands between 1 December 2007 and 1 July 2012. A total of 201 patients with severe COPD from 47 hospitals were enrolled by three centres for home mechanical ventilation (University Medical Center Groningen, Maastricht University Medical Center and the Erasmus Medical Center, Rotterdam) with a follow-up of 12 months. Inclusion criteria were: (1) COPD, GOLD stage 3 and 4;¹⁵ (2) >48 h independence from ventilatory support (invasive or non-invasive) for ARF; and (3) prolonged hypercapnia ($\text{PaCO}_2 > 6.0$ kPa) during daytime at rest without oxygen or ventilatory support. The study was approved by the local Medical Ethics Committee of the University Medical Centre Groningen, University of Groningen, patients gave informed consent to participate in the study and the trial was registered at the Nederland's Trial Register (NTR1100). For a more detailed description of the methods used, see online supplementary file.

Randomisation and intervention

Patients were randomised to nocturnal NIV or standard treatment by a computer with minimisation for FEV_1 (≤ 0.8 L or > 0.8 L), PaCO_2 (≤ 7.0 kPa or > 7.0 kPa), body mass index (BMI) (≤ 30 kg/m² or > 30 kg/m²), ventilatory support (invasive or non-invasive ventilation) and rehabilitation (yes/no). Standard treatment consisted of optimal medical treatment as usual, with LTOT in case of clinically stable patients with a $\text{PaO}_2 < 8.0$ kPa as indicated according to the Dutch Guideline.¹⁶

Nocturnal NIV was started by a highly experienced nurse practitioner from the nearest home mechanical ventilation centre who visited the hospital where the patient was admitted. NIV was initiated by bi-level positive airway pressure in the spontaneous/timed mode (Synchrony, Respironics, INC, Murrysville, Pennsylvania, USA) starting with a low back up frequency of 12/min with the aim to: (1) maximally support respiration and (2) and achieve normocapnia. NIV was initiated with an inspiratory positive airway pressure (IPAP) of 14 cm H₂O and gradually increased during trials to a maximal tolerated level. Expiratory positive airway pressure (EPAP) was started at 4 cm H₂O and increased if auto-PEEP was present or when patients used respiratory muscles to trigger the ventilator. Respiratory rate (RR) was set as close as possible to the RR of the patient, with adjustments if the patient triggered too much. Inspiration to expiration time was 1:3, with a short rise time and then titrated on comfort and effectiveness. All patients were initially given ventilation by means of a full-face mask and humidification to maximise comfort. Effectiveness with NIV during the night was observed by monitoring the transcutaneous O₂ saturation and PCO₂ (PtCO₂) with the TOSCA device (TOSCA 500, Linde Medical Sensors AG, Basel, Switzerland).¹⁷ Adherence to NIV was derived from time counters in the NIV device.

Measurements

At baseline, arterial and lung function measurements were performed just before randomisation. Baseline HRQL was assessed after randomisation while transcutaneous measurements were

performed during the night after patients were adjusted to NIV but before discharge of the initial admission for ARF. Lung function and arterial blood gas measurements were repeated after 3, 6 and 12 months during visits to their regular pulmonologist. Arterial blood gases were taken while breathing room air; however, when patients were unable to stop oxygen therapy even for short periods, measurements were taken with their usual oxygen flow rate. The nurse practitioner also visited all patients at home after 3, 6 and 12 months to collect survival, admission and exacerbation rates, HRQL and transcutaneous measurements, and if necessary to adjust NIV.

The long recruitment period enabled us to extend survival follow-up beyond 12 months to maximally 68 months for the first patients who entered the study.

HRQL was assessed by the Clinical COPD Questionnaire (CCQ),¹⁸ the Mageri Respiratory Failure Questionnaire-28 (MRF-28),¹⁹ the Chronic Respiratory Questionnaire (CRQ) self-reported²⁰ and the Severe Respiratory Insufficiency (SRI) questionnaire.²¹ We also measured mood state (Hospital Anxiety and Depression scale),²² daily activity level (Groningen Activity and Restriction Scale)²³ and dyspnoea (Medical Research Council).²⁴

Sample size calculations

Time to event (readmission for respiratory cause or death) as primary outcome was used to determine the sample size based on the assumption of an event-free survival of 20% in the standard treatment group⁵ after 12 months and 40% in the NIV group. With an α of 0.05 and a power of 80%, we needed 174 patients to detect the 20% difference in event-free survival between groups. Estimating a maximum dropout rate of 15%, 200 patients were needed.

Statistical analysis

The primary outcome and survival analysis were based on intention-to-treat (ITT) using Kaplan–Meier estimates and the log-rank test. A complete case analysis was performed for the remaining secondary outcomes comparing data of the patients who completed the 12 months (completers) as the number of dropouts was considerable due to progression of the disease and reluctance to adhere to study requirements. Arterial blood gas measurements could not always be performed under similar conditions for every patient. Therefore, data were analysed in two ways; first regardless of oxygen use or the amount of oxygen and second only comparing baseline and 12 month data when either the measurements were both on room air or both on oxygen at the same flow rate (also see online supplementary file).

As not all patients were compliant to NIV, either not using it every night or refusing treatment altogether, a per protocol analysis comparing patients in the NIV group who used ventilation >5 h/night with standard treatment was conducted based on results of a recent meta-analysis showing positive effects on change in arterial blood gasses in this subgroup.²⁵ All statistical calculations were performed using IBM SPSS Statistics V20 (IBM, New York, USA).

RESULTS

Patients

In all, 201 patients met eligibility criteria, 101 were randomised to NIV and 100 to standard treatment (figure 1). At baseline, patient characteristics and demographics did not differ significantly between treatment groups for ITT as well as the completers (table 1) except for the total number of admission days in

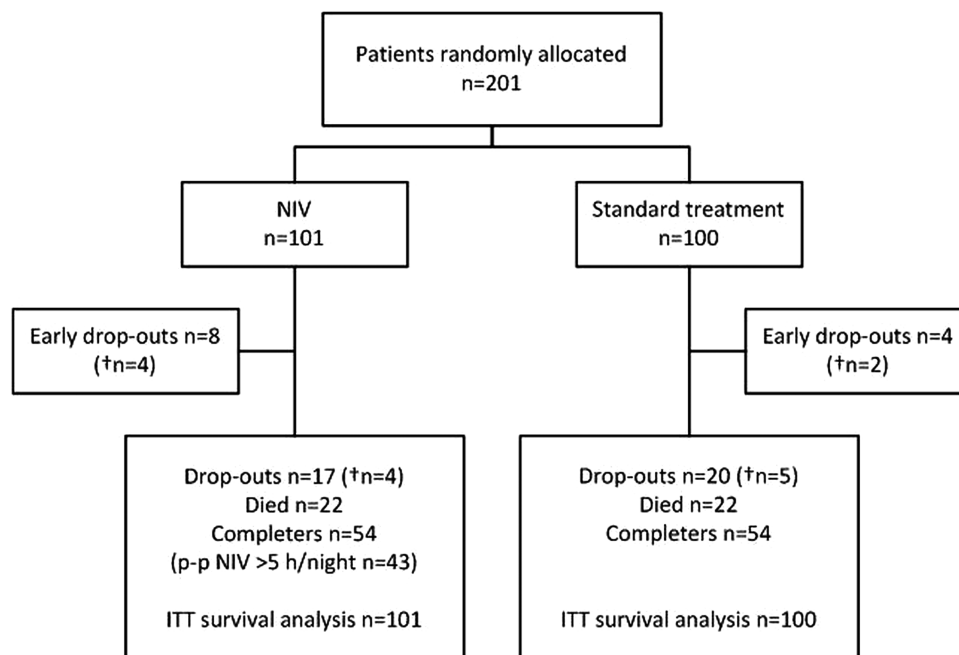


Figure 1 CONSORT flow diagram. Early dropouts: When patients refused or were not able to continue the protocol (either NIV or standard treatment) within 10 days after inclusion. Completers: number of patients included in the complete case analysis. Four patients classified as dropouts crossed from standard therapy over to the NIV treatment during the study. ITT: number of patients included in the intention-to-treat analysis; n, numbers for analysis; †n, number of patients who died within the dropouts; NIV, non-invasive positive pressure ventilation; p-p, included in the per protocol analysis of patients who used ventilation >5 h/night.

Table 1 Baseline characteristics

	ITT NIV n=101	ITT controls n=100	Completers NIV n=54	Completers controls n=54
Age (years)	63.92 (8.6)	63.5 (7.9)	61.9 (7.9)	62.3 (7.9)
Gender, n (% female)	60 (59%)	58 (58%)	36 (67%)	34 (63%)
Pack years*	38 (0–140)	44 (0–125)	38 (0–125)	38 (0–125)
BMI (kg/m ²)	24.6 (5.4)	24.8 (6.3)	24.7 (5.5)	24.2 (5.3)
LTOT, n (%)	76 (75%)	78 (78%)	38 (70%)	37 (69%)
Invasive, n (%)†	13 (13%)	12 (12%)	7 (13%)	10 (19%)
Duration ventilation*‡ (days)	5.0 (1–19)	5.0 (1–24)	5.5 (2–19)	5.0 (1–23)
Admissions* (year ⁻¹)	2.0 (1–9)	2.0 (1–10)	2.0 (1–9)	1.0 (1–6)
Total admission days/patient* (year ⁻¹)	28.5 (8–108)	22.0 (6–115)§	29.0 (8–100)	20.0 (7–115)§
Rehabilitation, n (%)	50 (50%)	51 (51%)	25 (46%)	30 (56%)
FEV ₁ (L)	0.67 (0.23)	0.65 (0.23)	0.70 (0.24)	0.70 (0.25)
FEV ₁ (% pred)	25.6 (7.8)	25.7 (8.6)	26.9 (7.7)	27.6 (8.6)
FVC (% pred)	64.3 (19.8)	63.6 (17.0)	65.7 (18.4)	67.8 (17.1)
FEV ₁ to FVC (%)	32.5 (9.0)	33.0 (9.5)	33.9 (9.6)	33.8 (9.7)
PaCO ₂ ¶ (kPa)	7.9 (1.2)	7.7 (1.3)	8.0 (1.2)	7.5 (1.0)
PaO ₂ ¶ (kPa)	7.9 (2.1)	7.5 (1.7)	8.3 (1.6)	7.9 (1.8)
Medication, n (%)				
Inhaled corticosteroid	86 (85%)	84 (84%)	43 (81%)	48 (89%)
Oral corticosteroids	53 (53%)	38 (38%)	27 (50%)	17 (32%)
Beta agonists	98 (97%)	97 (97%)	52 (98%)	53 (98%)
Anticholinergics	94 (93%)	94 (94%)	49 (93%)	52 (96%)
Theophylline	37 (37%)	27 (27%)	20 (38%)	16 (30%)

Data presented as mean (SD) unless otherwise stated.

*Pack years, duration ventilation and admissions for respiratory causes including index admission in median (range).

†Invasive ventilation during acute respiratory failure. Other patients received NIV.

‡Ventilation during acute respiratory failure of index admission.

§p<0.05 significant difference compared with NIV.

¶While breathing room air (in both groups blood samples from 21 patients were taken with oxygen as they were not able to come of oxygen).

% pred, % predicted; BMI, body mass index; Controls, standard treatment; ITT, intention-to-treat; LTOT, long-term oxygen therapy; n, numbers for analysis; NIV, non-invasive positive pressure ventilation; PaCO₂, arterial carbon dioxide pressure; PaO₂, arterial oxygen pressure.

the previous year. Reasons for dropping-out of the study are listed in the online supplementary table S1. Four patients classified as dropouts crossed from standard treatment over to the NIV treatment during the study. They remained in the standard treatment group for all ITT analyses.

NIV settings+treatment adherence

Patients randomised for NIV were discharged with a mean±SD IPAP of 19.2±3.4 and EPAP of 4.8±1.0 cm H₂O, respectively, mean RR on NIV of 15±3 breaths/minute, an inspiration time of 1.1±0.3 s and a rise time of 0.2±0.11 s. Settings were not different between the completers and the dropouts. Overall, 68% received oxygen combined with NIV during the night. Nearly all patients were ventilated through a full-face mask, one using a total face mask. Minor adjustments were made during the study to improve adherence, blood gasses and/or comfort. After 1 year, the remaining 54 patients were ventilated with a mean IPAP of 21.0±3.4 and an EPAP of 5.2±1.2 cm H₂O. Two additional patients switched to a total face mask.

Mean duration of NIV use per night until death or last follow-up was 6.3±2.4 h/night in the total group and 6.9±2.1 in the completers. Patients included in the per protocol analysis (NIV >5 h) reached an average of 7.7±1.5 h/night (n=43).

Time to event

One year after discharge, 65% versus 64% of patients (NIV vs standard treatment) were either readmitted to hospital for COPD or had died. Figure 2 shows the ITT event-free survival of both groups in the first year, which was not significantly different (mean number of days to event was 192 and 198 for the NIV and standard treatment groups, respectively, p=0.85).

Hospital admissions

After 1 year, readmission to hospital for respiratory causes occurred in 56% versus 57% in the NIV and the standard treatment groups, respectively. The median (range) number of readmissions during the 12 months of follow-up did not differ

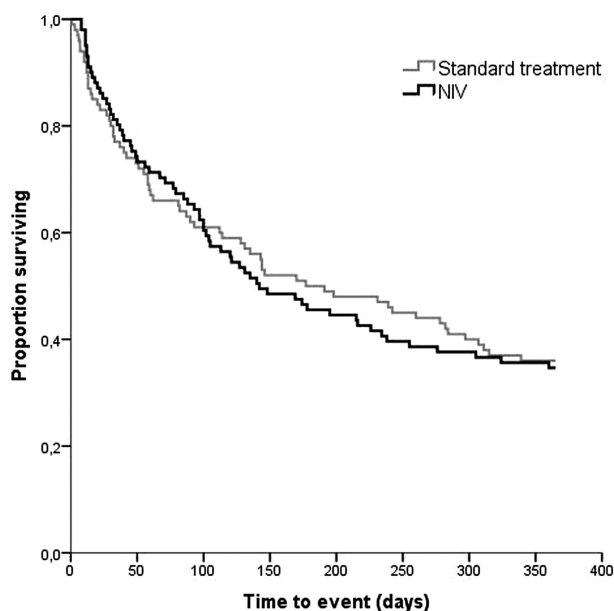


Figure 2 Kaplan–Meier plot of time to event (readmission for respiratory cause or death) in patients randomised to non-invasive positive pressure ventilation (NIV) and standard treatment.

significantly between groups (1.0 (0–9) in NIV vs 1.0 (0–6) in the standard treatment group p=0.23) nor did the median number of total days spent in hospital (7.0 (0–107) in NIV vs 3.5 (0–77) for standard treatment) (p=0.087).

Survival

The number of deaths after 1 year was 30 and 29 in the NIV and standard treatment groups, respectively, with a mean number of survival days of 299 for NIV and 291 days for the standard treatment group (p=0.99) (see online supplementary table S2). Long-term survival is shown in figure 3, with 63 deaths in the NIV group versus 58 in the standard treatment group.

Exacerbations

No difference was observed between the annual median (range) number of exacerbations at home: 1.0 (0–9) for the NIV and 2.0 (0–14) for the standard treatment group (p=0.26).

Daytime arterial blood gases

After 12 months, daytime PaCO₂ improved significantly more in the NIV group compared with standard treatment (table 2, figure 4).

Nocturnal gas exchange measurements

The mean nocturnal PtcCO₂ measured during the night before discharge out of hospital in the ITT group was significantly better in patients on NIV compared with standard treatment (mean difference –0.8 kPa (95% CI –1.3 to –0.4; p<0.01)). In the completers, this effect was not found at baseline but was present after 12 months; mean nocturnal PtcCO₂ was significantly better in the NIV group compared with standard treatment (mean difference –0.6 kPa (95% CI –1.1 to –0.1; p=0.03)).

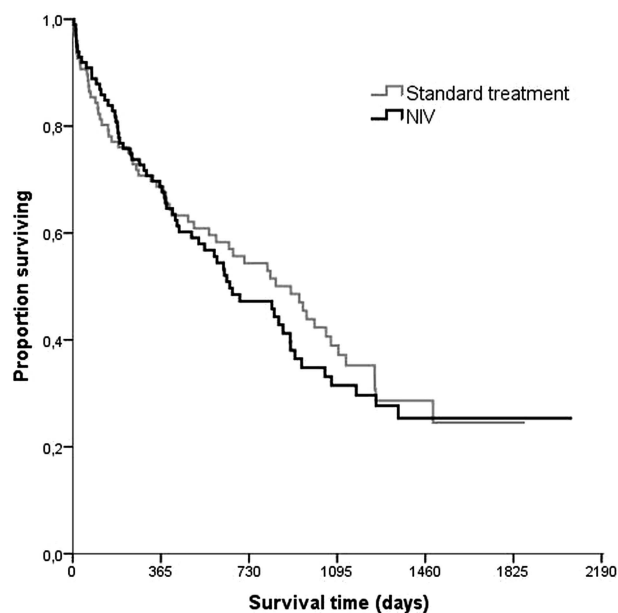


Figure 3 Kaplan–Meier plot of long-term survival curves of patients randomised to non-invasive positive pressure ventilation (NIV) and standard treatment. Because of small numbers of patients followed up after 3 years, the right-hand end of the survival plots remains uncertain.

Table 2 Changes in daytime arterial blood gases

	ITT baseline	n	Completers baseline	n	Completers 12 months	n	Change over 1 year*	n	Change over 1 year similar circumstances†	n
pH										
NIV	7.38±0.04	93	7.38±0.04	46	7.40±0.04	46	0.026 (0.010 to 0.042)‡	46	0.022 (0.005 to 0.039)‡	37
Controls	7.39±0.04	99	7.39±0.04	48	7.39±0.03	48	−0.005 (−0.016 to 0.006)	48	−0.005 (−0.020 to 0.010)	37
<i>Treatment effect</i>							0.03 (0.012 to 0.050)§		0.030 (0.005 to 0.050)§	
PaCO ₂ (kPa)										
NIV	7.9±1.2	100	7.7±1.1	50	6.4±1.0	50	−1.3 (−1.6 to −0.9)‡	50	−1.0 (−1.4 to −0.7)‡	38
Controls	7.7±1.3	99	7.4±0.9	48	6.6±1.0	48	−0.8 (−1.1 to −0.5)‡	48	−0.9 (−1.2 to −0.5)‡	37
<i>Treatment effect</i>							−0.5 (−0.9 to −0.04)§		−0.2 (−0.6 to 0.3)	
PaO ₂ (kPa)										
NIV	7.9±2.1	93	8.3±1.6	45	8.3±1.6	45	0 (−0.6 to 0.6)	45	0.3 (−0.3 to 0.9)	35
Controls	7.5±1.6	99	7.9±1.9	48	8.4±1.4	48	0.6 (−0.1 to 1.2)	48	0.6 (−0.1 to 1.3)	37
<i>Treatment effect</i>							0.6 (−1.4 to 0.2)		−0.3 (−1.2 to 0.6)	
HCO ₃ (mmol/L)										
NIV	33.7±4.6	94	33.0±4.5	47	29.3±3.5	47	−3.7 (−5.2 to −2.1)‡	47	−3.0 (−4.8 to −1.2)‡	36
Controls	33.6±4.8	98	32.6±4.1	48	28.8±4.2	48	−3.8 (−5.0 to −2.6)‡	48	−4.1 (−5.5 to −2.7)‡	37
<i>Treatment effect</i>							0.1 (−1.8 to 2.1)		1.0 (−1.2 to 3.3)	
BE										
NIV	7.8±4.8	90	7.0±3.2	38	4.5±2.5	38	−2.5 (−3.6 to −1.3)‡	38	−2.2 (−3.4 to −0.9)‡	31
Controls	7.7±3.8	93	6.9±3.2	44	3.9±3.2	44	−2.9 (−4.0 to −1.8)‡	44	−3.2 (−4.5 to −1.9)‡	35
<i>Treatment effect</i>							0.5 (−1.1 to 2.0)		1.0 (−0.8 to 2.8)	
Sat (%)										
NIV	87.4±6.4	95	90.0±4.4	43	91.0±5.1	43	1.1 (−0.9 to 3.0)	43	2.2 (0.2 to 4.2)‡	34
Controls	88.4±6.6	88	88.4±6.1	43	91.1±4.8	43	2.7 (0.5 to 5.0)‡	43	2.7 (−0.03 to 5.5)	33
<i>Treatment effect</i>							−1.6 (−4.6 to 1.3)		−0.5 (−3.9 to 2.8)	

Data presented as means (±SD) and mean change (95% CIs); controls, standard treatment.

*Measurement performed regardless of oxygen use.

†Only comparing baseline and 12-month data when the measurements were both on room air or both on oxygen at the same flow rate. N=37 (out of 38 NIV) and n=35 (out of 37 controls) for which both measurements were performed on room air.

‡p<0.05 significant change after 12 months within the group.

§p<0.05 significant difference in change after 12 months between the groups (treatment effect).

BE, base excess; HCO₃, bicarbonate (mmol/L); ITT, intention-to-treat; n, numbers for analysis; NIV, non-invasive positive pressure ventilation; PaCO₂, arterial carbon dioxide pressure (kPa); PaO₂, arterial oxygen pressure (kPa); Sat, saturation (%).

Lung function

Mean FEV₁ and vital capacity did not change significantly after 12 months within the NIV group or standard treatment group nor was there a significant change between groups (table 3).

Health-related quality-of-life

HRQL, activities of daily living, anxiety/depression scores and dyspnoea improved in both groups but there was no significant difference between the groups (table 4 and see online supplementary tables S3–S7). HRQL measured by the SRI total score showed a clear trend (p=0.054) with additionally significant differences between groups on the subdomains ‘attendant symptoms and sleep’ and ‘social relationships’ in favour of the NIV group (see online supplementary file).

Per protocol analysis

All results were similar in the per protocol analysis comparing patients with NIV>5 h/night with standard treatment.

DISCUSSION

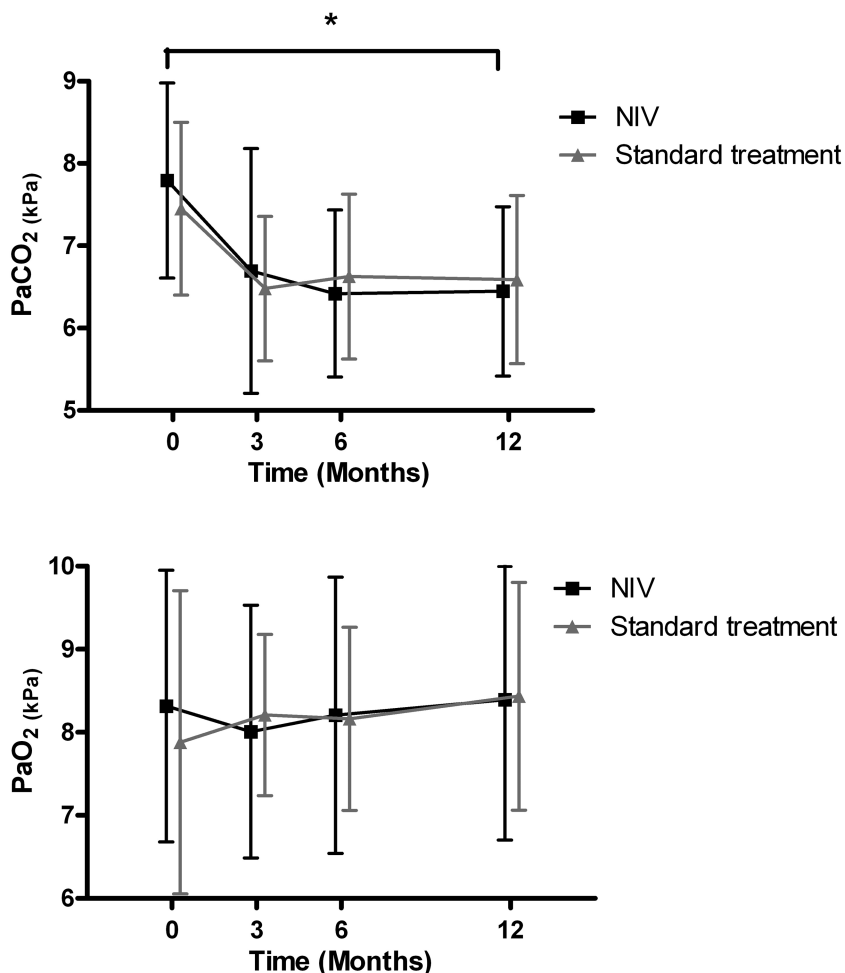
Thus far, this is the largest randomised, controlled multicentre study reporting on the effects of NIV in COPD patients with prolonged hypercapnia after an episode of NIV for ARF. It showed that nocturnal NIV does not prolong the time to readmission for respiratory causes or death in COPD patients who remain hypercapnic after ventilatory support during ARF. Compared with standard treatment alone, NIV improves

daytime PaCO₂ and nocturnal transcutaneous PCO₂ but does not improve survival, number of respiratory readmissions, exacerbations, lung function, HRQL, mood state, daily activity levels or dyspnoea.

In this large study, the design was practical and executed as in ‘real-life’, which make outcomes more broadly applicable (patients remained in their own hospital and blood gas and lung function measurements were performed during their regular visits to their own pulmonologist), but does make it difficult to compare our outcomes with previous studies in this field and therefore necessitates further discussion and comparison.

First, Funk *et al*¹⁴ reported positive results in an RCT including similar patients (COPD patients remaining hypercapnic after ARF requiring mechanical ventilation), showing a higher probability of clinical worsening in the control (withdrawal) group compared with the NIV group. A crucial difference in trial designs is that all patients first received NIV for 6 months after which one group was randomised to stop NIV (withdrawal group). Clinical worsening was defined as (1) an intensive care unit admission for treatment of acute-on-chronic respiratory failure requiring mechanical ventilation, but for the withdrawal group escalation additionally comprised of (2) voluntary resumption of NIV due to severe dyspnoea and (3) re-institution of NIV due to progressive hypercapnia. A pitfall of this approach was that, as the authors also note themselves, by definition the achievable endpoints depended on the group allocation of the patient as the NIV group could only qualify for the first

Figure 4 Daytime PaCO₂ and PaO₂ at baseline, 3, 6 and 12 months (means±SD). Measurements performed in complete case group (n=54 per group) regardless of oxygen use. NIV, non-invasive positive pressure ventilation; PaCO₂, arterial carbon dioxide pressure (kPa); PaO₂, arterial oxygen pressure (kPa). *Change in PaCO₂ was significantly better in the NIV group after 12 months (p<0.05).



endpoint. Results after 12 months follow-up showed that 10 patients in the withdrawal group versus two in the NIV group experienced clinical worsening (p=0.0018). But upon looking more closely and, comparing only the endpoint which was achievable for both groups, only three out of the initial 10 patients in the withdrawal group (vs two) were admitted to the ICU for acute-on-chronic respiratory failure. Other positive outcomes in favour of NIV were a significant difference in 6-min walking distance after 3 months and a slightly lower resting daytime pH in the withdrawal group after 12 months. Our data show a higher daytime pH in the NIV group, but we did not consider this improvement clinically relevant (as patients were

not acidic at baseline). We could not perform the 6-min walking test since most patients were too weak to perform this test adequately and safely, raising ethical concerns.

Second, in a randomised pilot trial by Cheung and colleagues⁶ with similar inclusion criteria as in our study, chronic home NIV was compared with placebo NIV (CPAP of 5 cm H₂O). In this study, primary outcome was recurrent severe COPD exacerbation with acute hypercapnic respiratory failure (AHRF), resulting in NIV, intubation or death within the following year. The proportion of patients developing this condition in 1 year was 38.5% in the NIV versus 60.2% in the placebo group (p=0.039). Compared with our study, these patients

Table 3 Changes in lung function

	ITT baseline	n	Completers baseline	n	Completers 12 months	n	Change over 1 year	n
FEV₁ (litres)								
NIV	0.67 (0.23)	99	0.70 (0.24)	48	0.71 (0.30)	48	0.01 (-0.05 to 0.08)	48
Controls	0.65 (0.23)	95	0.72 (0.26)	46	0.76 (0.32)	46	0.04 (-0.03 to 0.11)	46
Treatment effect							-0.024 (-0.12 to 0.07)	
VC (litres)								
NIV	2.1 (0.7)	98	2.1 (0.70)	49	2.2 (0.78)	49	0.09 (-0.09 to 0.27)	49
Controls	2.3 (0.8)	95	2.3 (0.75)	46	2.4 (0.74)	46	0.13 (-0.08 to 0.34)	46
Treatment effect							-0.036 (-0.31 to 0.23)	

Data presented as means (±SD) and mean change (95% CIs); controls, standard treatment. There were no significant changes after 12 months in lung function within or between the groups. FEV₁, forced expiratory volume in one second, postbronchodilator; ITT, intention-to-treat; n, numbers for analysis; NIV, non-invasive positive pressure ventilation; VC, vital capacity.

Table 4 Changes in health-related quality-of-life (HRQL) measurements

	ITT baseline	n	Completers baseline	n	Completers 12 months	n	Change over 1 year	n
CCQ-Total								
NIV	3.37±1.2	96	3.4±1.2	48	2.9±1.1	48	-0.5 (-0.2 to -0.8) *	48
Controls	3.31±1.06	90	3.2±1.0	51	2.8±1.1	51	-0.5 (-0.8 to -0.1) *	51
<i>Treatment effect</i>							-0.04 (-0.5 to 0.4)	
MRF-28-Total								
NIV	60.88±23.60	99	58.3±24.3	50	51.0±24.8	50	-7.3 (-12.5 to 2.1) *	50
Controls	60.28±23.89	90	55.3±24.2	51	49.4±24.2	51	-5.8 (-10.8 to -0.8)*	51
<i>Treatment effect</i>							-1.5 (-8.6 to 5.7)	
CRQ-Total								
NIV	3.47±1.10	100	3.5±1.1	50	4.2±1.2	50	0.7 (0.4 to 1.1)*	50
Controls	3.59±1.08	89	3.6±1.1	50	4.4±1.1	50	0.7 (0.5 to 1.0)*	50
<i>Treatment effect</i>							0.01 (-0.4 to 0.4)	
SRI-Total								
NIV	48.14±14.97	100	47.9±15.1	50	55.0±15.4	50	7.0 (3.4 to 10.7)*	50
Controls	51.33±15.87	90	53.6±16.9	51	55.8±16.3	51	2.2 (-1.2 to 5.6)	51
<i>Treatment effect</i>							4.8 (-0.1 to 9.7)	
GARS-Total								
NIV	36.66±7.50	100	36.3±9.4	50	34.6± 9.4	50	-1.6 (-3.3 to 0.1)	50
Controls	36.76±8.54	90	34.7±8.8	51	32.7±8.2	51	-2.0 (-4.1 to 0.1)	51
<i>Treatment effect</i>							0.4 (-2.3 to 3.0)	
HADS-Total								
NIV	15.88±9.24	98	15.6±8.8	48	13.9±8.8	48	-1.7 (-4.0 to 0.6)	48
Controls	14.48±8.83	87	13.1±9.1	50	12.7±9.3	50	-0.4 (-2.2 to 1.3)	50
<i>Treatment effect</i>							-1.3 (-4.1 to 1.6)	
MRC								
NIV	3.9±1.0	98	3.8±1.0	49	3.4±1.4	49	-0.4 (-0.7 to -0.01)*	49
Controls	3.8±1.0	90	3.6±1.1	51	3.3±1.2	51	-0.3 (-0.7 to 0.1)	51
<i>Treatment effect</i>							-0.05 (-0.6 to 0.5)	

Data presented as means (±SD) and mean change (95% CIs).

Scores represent: CCQ and MRF-28↓=HRQL↑; CRQ and SRI↑=HRQL↑; GARS↓=ADL↑; HADS↓=mood↑; MRC↓=dyspnoea↓.

*p<0.05 significant change after 12 months within the group. There was no significant difference in change after 12 months between the groups (treatment effect).

CCQ, Clinical COPD Questionnaire; Controls, standard treatment; CRQ, Chronic Respiratory Questionnaire; GARS, Groningen Activity and Restriction Scale; HADS, Hospital Anxiety and Depression Scale (separate scores for anxiety and depression domain); ITT, intention-to-treat; MRC, Medical Research Council dyspnoea scale; MRF-28, Mageri Respiratory Failure Questionnaire-28; n, numbers for analysis; NIV, non-invasive positive pressure ventilation; SRI, Severe Respiratory Insufficiency questionnaire.

were older (mean age 70) and had a lower mean BMI (19.2) suggesting a group of patients in worse condition than in the present study. The authors point out that ethical concerns led them to be prudent and to act upon severe enough exacerbations by instituting NIV, which could indicate a bias towards earlier conversion to NIV in the CPAP group and a possible explanation for the difference in proportions between groups. Furthermore, this trial did not find a difference in survival, arterial blood gasses and adverse events between groups.

Third, our power analysis was based on the study by Chu *et al*⁵ who found that 80% of COPD patients with AHRF who were successfully treated by NIV were readmitted to hospital for respiratory problems within the following year. Our primary outcome consisted of readmission for respiratory causes but also death, probably leading to higher percentages of patients in which 'the event' would occur. Ideally, it would be more appropriate to power on pilot data from our own country or a similar European Healthcare system but as this was the only study at that time we had to work with these numbers. We hypothesised that in our study, the NIV group would have a 20% lower prevalence of readmission for respiratory causes than the standard treatment group (ie, 60% prevalence of readmission). However, our study shows a much lower prevalence of readmission in both the NIV and the standard treatment groups (56%

and 57%, respectively). A possible explanation could be that patients recruited in the study by Chu *et al*⁵ were older (mean age 73.2) and had a lower BMI (20.2), indicating a worse physical condition. Our results do show similarities with a Spanish study²⁶ which was published during our inclusion phase, reporting readmission rates of 66% due to respiratory complications in COPD patients who survived an acute exacerbation with ARF that required NIV.

As patients were included during an admission for ARF, we anticipated blood gasses to improve in the first months after discharge from hospital in both groups. Although PaCO₂ was better in the NIV group after 12 months, we did not expect to see the same level of improvement in PaCO₂ in the standard treatment group in the first 3 months (figure 4). If we assume that the improvement in the control group is the natural recovery of the disease, one might conclude that the included patients might not be the right target group for chronic NIV. The patients in this trial are possibly a mixed cohort of acute-on-chronic respiratory failure and ARF, and therefore a subgroup analysis might be interesting. However, the hypothesis tested in this study is still legitimate as hypercapnia persisting postexacerbation is an increasingly common reason for initiation of NIV in patients with COPD.

Another possible subgroup analyses for the future concerns the need for supplementary oxygen. This subgroup is likely to

indicate a more severe disease state and this would be a group expected to have greater therapeutic benefit with chronic NIV.

In a subgroup analysis from a recent meta-analysis on the effects of NIV in stable COPD,²⁵ a significant change in PaCO₂ after 3 months within the NIV group was found for patients with a baseline PaCO₂ of at least 7.3 kPa (55 mm Hg) for patients who used NIV for at least 5 h per night as well as for patients ventilated with IPAP levels of at least 18 cm H₂O, when compared with patients with lower IPAP levels, lower adherence or lower levels of hypercapnia. In the current study, baseline PaCO₂ and change in PaCO₂ after 12 months of ventilation were significantly correlated ($r=-0.6$, $p<0.001$). Levels of IPAP and the number of hours of NIV/night were not correlated with change in PaCO₂ (data not shown). Our data show that 26% of all patients in the standard treatment group had become eucapnic after 3 months time. If we extrapolate this number to the NIV group, this proportion of patients is less likely to benefit from chronic NIV. This failure to demonstrate a dose-response in number of hours and change in PaCO₂ is therefore likely due to the ‘dilution’ of the prolonged hypercapnic patient cohort by patients who recover and improve and become eucapnic over a longer timeframe (>48 h in the current trial) after an acute decompensated exacerbation, that is, patients who did not have acute-on-chronic respiratory failure but AHRE. Possibly, in future studies, selection of patients should be based on the level of hypercapnia after 3 months of recovering at home, before initiating home NIV. However, data from within the EU have demonstrated the high mortality of this patient group in the months following an AE requiring ventilatory support and although this trial may have evaluated patients too earlier the optimum timing remains to be proven.²⁷

A limitation in our study was our inability to obtain blood gasses on patient’s standard oxygen level as patients were reluctant to undergo testing twice. Although our aim was to only obtain samples on room air, a portion of patients refused to stop their oxygen, making comparison of data difficult. Nevertheless, patients on NIV were adequately ventilated during the night as their mean nocturnal PtcCO₂ was significantly better compared with standard treatment.

In an RCT on NIV in stable COPD, Clini *et al*¹² did not find a difference in PaCO₂ between NIV and the control group (LTOT) during room air breathing whereas they did show a significant difference when measured on patients’ usual oxygen regimen.

Also, long-term oral corticosteroid treatment is suggested to negatively impact on readmission rates and outcome. Since NIV patients more frequently received oral corticosteroids (ITT 53% vs 38%) this could also have affected the results.

The fact that we were able to obtain extensive data on HRQL is unique. During our study, we published an article²⁸ in which reliability and validity of the CCQ, CRQ, MRF-28 and SRI questionnaires were assessed and compared from our patients at baseline. Although, all four questionnaires were feasible in these types of patients, the SRI scored tested most appropriate. This is reflected in our data after 12 months which show a clear trend towards improved HRQL as measured by the SRI total score. Two out of the seven subdomains were significantly improved in the NIV group compared with standard treatment. The very opposite was reported in the RCT by McEvoy *et al*¹³ who demonstrated a survival advantage of chronic NIV in hypercapnic stable COPD but accompanied by a deterioration in HRQL (measured by the more general St. Georges Respiratory Questionnaire). We believe this emphasises the need to

consistently use severity-specific HRQL questionnaires like the SRI and MRF-28 to further explore the possible benefits of NIV in future research, for instance, the supportive role of NIV in advanced care planning also.

CONCLUSIONS

We could not demonstrate an improvement in time to readmission or death by adding NIV for 1 year in COPD patients who remained hypercapnic after an episode of NIV for ARF. There is no reason to believe the NIV was not effective since daytime PaCO₂ and night-time PCO₂ improved. The trend for improvement in HRQL favouring NIV we believe nevertheless should be explored further.

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Contributors FS was the coordinator of the study, directly involved in the project design, patient recruitment, data collection and analysis, and is the main author of the manuscript. GB contributed to data collection and reviewed the final version. The following authors were the main investigators in the collaborating centres and directly involved in patient recruitment and data collection and reviewed and approved the final version of the manuscript: RS, MZ, JA and NC. HK was involved in the design and execution of the study. JV and HK supervised the data interpretation and reviewed and approved the final version of the study. PW was the main investigator, led the study group, contributed to the design of the project, and contributed to and approved the final study; he is the guarantor of the entire study.

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Patient consent Obtained.

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Online data supplementary file

TITLE: Nocturnal noninvasive ventilation in COPD patients with prolonged hypercapnia after ventilatory support for acute respiratory failure: a randomised, controlled, parallel-group study

Acronym: RESCUE REspiratory Support in COPD after acUte Exacerbation.

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METHODS

Patients

Exclusion criteria were: 1) age < 18 or => 80 years; 2) significant bronchiectasis with recurrent infections; 3) significant heart failure; 4) kyphoscoliosis; 5) neuromuscular disease; 6) obstructive sleep apnea (Apnea Hypopnea Index: AHI >15 /hr); 7) current use of Continuous Positive Airway Pressure (CPAP) or Bi-level Positive Airway Pressure (BiPAP); 8) insufficient motivation for chronic ventilatory support; 9) social circumstances making chronic ventilatory support at home impossible; 10) other diagnoses limiting life expectations.

Home Mechanical Ventilation Center

In The Netherlands we have 4 legitimated and highly experienced Home Mechanical Ventilation Centers. Patients requiring chronic ventilation at home are sent in from all hospitals in the country to be established on mechanical ventilation by these centers. The nurse practitioners are allowed to establish long-term NIV also in external hospitals and deliver care at home. For this trial as a form of quality assurance, we set up a standardized protocol for establishment and NIV set up and organized a start up meeting to make sure everybody worked according to the same standard and regulations. Also, all centers worked with the same machines.

Intervention

Patients were instructed to use NIV during the night whilst asleep and, were advised to use NIV during the day/nap times if desired.

Measurements

An overnight polygraphy or Polysomnography was performed in patients with a body mass index $\geq 30 \text{ km/m}^2$, or in patients with complaints of excessive snoring, disrupted sleep or morning headache to exclude Obstructive Sleep Apnea Syndrome.

Survival, respiratory admission rates and days in hospital were checked in hospital registers and with patients general practitioner. Exacerbations at home were registered in a diary by the patient themselves.

An exacerbation was defined using a modified version of the definition of Rodriguez-Roisin, as an event in the natural course of the disease characterized by a change in the patient's baseline dyspnoea, cough, and/or sputum that is beyond day-to-day variations, is acute in onset, and treated with antibiotics and/or prednisolone in patients with COPD.[1]

Lung function

Lung function measurements included routine spirometry by means of a pneumotachograph and according to ERS criteria.[2] Absolute values and % pred. according to normal values.

Transcutaneous measurement

Transcutaneous monitoring during the night was performed using the TOSCA[3] to measure skin-surface PO_2 and PCO_2 to provide estimates of arterial partial pressure of oxygen and carbon dioxide (PaO_2 and PaCO_2) and saturation. The device induces hyperperfusion by local heating of the skin of the earlobe and measures the partial pressure of oxygen and carbon dioxide electrochemically. The mean nocturnal PtcCO_2 was measured during the night before discharge out of hospital to see if patients were adequately adjusted to NIV. Patients in the standard treatment did not receive NIV during this measurement.

Health related quality of life questionnaires

Patients completed the following questionnaires concerning health related quality of life; the Clinical COPD Questionnaire (CCQ),[4] the Chronic Respiratory Questionnaire self reported (CRQ),[5] the Mageri Respiratory Failure Questionnaire -28 (MRF-28)[6] and the Severe Respiratory Insufficiency (SRI) questionnaire.[7] The CCQ is a self-administered, 10-item questionnaire which can be divided into three domains: symptom, functional state and mental state. Scores range from 0 to 6 with high scores indicating extremely poor health status. The CRQ (self reported) contains 20 items and measures physical function and emotional function, divided into four domains: dyspnoea, fatigue, emotion and mastery. Scores range from 1 (worse) to 7 (best). The MRF-28 contains 28 items which are divided into 3 domains; daily activity, cognitive function and invalidity. The scores range from 0 (best) to 100 (worse). The SRI contains seven domains covering 49 items: respiratory complaints, physical functioning, attendant symptoms and sleep, social relationships, anxiety, psychological well-being and social functioning. Scoring ranges between 0-100, with high scores representing better HRQL.

Other measurements

The Groningen Activity and Restriction Scale (GARS) assesses activity and disability of daily living and consists of 18 items.[8] The Hospital Anxiety and Depression Scale (HADS) was used to determine levels of depression and anxiety.[9] It consists of 14 questions from which seven are on detection of anxiety and seven on depression (two subscales). Dyspnoea was measured using the Medical Research Council scale (MRC).[10]

We could not perform the 6-minute walking test since most patients were too weak to perform this test adequately and safely, raising ethical concerns. Our alternative of measuring walking distance with a pedometer also fell short, as we noticed that in the patients who

barely walk, shuffling occurs leading to an underestimation as the pedometer does not always detect their effort and thus real number of steps per day.

Statistical analysis

Continuous variables were summarized as means and standard deviations, or medians and ranges depending on their distribution. Changes within groups were compared using the paired sample T-test. Between group comparisons of continuous variables were performed using the independent samples T-test for variables with a normal distribution or the Mann-Whitney U test for variables with a non-normal distribution. Difference in change within groups and treatment effect between groups are shown as means with associated 95% confidence intervals (C.I.) and p-value ($p < 0.05$ was considered statistically significant).

Supplementary file Table S1: Reasons for dropping out of study

	NIV n=25	Standard treatment n=24
Lack of motivation	15	14
Unable to come for testing	0	6
Discomfort associated with treatment	8	-
Critical intercurrent illness:		
- Dementia	1	0
- CVA	1	0
Switch to NIV	-	4

CVA cerebrovasculair accident; n, numbers for analysis; NIV, noninvasive positive pressure ventilation.

Supplementary file Table S2: Causes of death

	ITT NIV n=30	ITT Standard treatment n=29	Non-drop outs NIV n=22	Non-drop outs Standard treatment n=22
Respiratory causes	21	25	15	20
Natural cause	3	3	3	1
Pneumothorax	1	1	1	1
Cardiac disease	3		1	
Lung carcinoma	1		1	
Missing	1		1	

ITT, intention-to-treat; n, numbers for analysis; NIV, noninvasive positive pressure ventilation.

Supplementary file Table S3: Changes in the Clinical COPD Questionnaire (CCQ) sub domains

	ITT	n	Completers	n	Completers		Change over	n
	Baseline		Baseline		12 months		1 year	
CCQ-Total								
NIV	3.4 ± 1.2	96	3.4 ± 1.2	48	2.9 ± 1.1	48	-0.5 (-0.2; -0.8) [¶]	48
Controls	3.3 ± 1.2	90	3.2 ± 1.0	51	2.8 ± 1.1	51	-0.5 (-0.8; -0.1) [¶]	51
Treatment effect							-0.04 (-0.5; 0.4)	
Symptom								
NIV	3.1 ± 1.2	99	3.0 ± 1.2	51	2.7 ± 1.2	51	-0.3 (-0.6; 0.1)	51
Controls	3.1 ± 1.1	90	3.2 ± 1.1	51	2.8 ± 1.1	51	-0.4 (-0.7; -0.04) [¶]	51
Treatment effect							-0.09 (-0.4; 0.6)	
Functional								
NIV	4.1 ± 1.4	99	4.1 ± 1.5	49	3.5 ± 1.5	49	-0.5 (-0.9; -0.2) [¶]	49
Controls	4.0 ± 1.4	90	3.9 ± 1.4	51	3.4 ± 1.4	51	-0.5 (-0.9; -0.1) [¶]	51
Treatment effect							-0.07 (-0.6; 0.5)	
Mental								
NIV	2.6 ± 1.8	96	2.7 ± 1.8	50	1.7 ± 1.3	50	-1.0 (-1.4; -0.6) [¶]	50
Controls	2.3 ± 1.7	90	2.2 ± 1.6	51	1.6 ± 1.5	51	-0.6 (-1.0; -0.2) [¶]	51
Treatment effect							-0.40 (-1.0; 0.2)	

Data presented as means (± standard deviation) and mean change (95% confidence intervals).
 CCQ, Clinical COPD Questionnaire; ITT, intention-to-treat; n, numbers for analysis; NIV, noninvasive positive pressure ventilation.

[¶] p<0.05 significant change after 12 months within the group.

*p<0.05 significant difference in change after 12 months between the groups (treatment effect).

Low CCQ scores indicate high quality of life. A negative treatment effect signifies a bigger improvement after 12 months for the NIV group compared to standard treatment.

Supplementary file Table S4: Changes in the Mageri Respiratory Questionnaire-28 (MRF-28) sub domains

	ITT	n	Completers	n	Completers	n	Change over	n
	Baseline		Baseline		12 months		1 year	
MRF-28-Total								
NIV	60.9 ± 23.6	99	58.3 ± 24.3	50	51.0 ± 24.8	50	-7.3 (-12.5; -2.1) [¶]	50
Controls	60.3 ± 23.9	90	55.3 ± 24.2	51	49.4 ± 25.0	51	-5.8 (-10.8; -0.8) [¶]	51
Treatment effect							-1.5 (-8.6; 5.7)	
Daily activities								
NIV	62.3 ± 28.4	99	62.0 ± 29.9	50	54.5 ± 35.4	50	-7.5 (-15.09 ; 0.03)	50
Controls	61.7 ± 30.5	90	55.6 ± 30.7	51	46.3 ± 29.2	51	-9.3 (-16.1; -2.4) [¶]	51
Treatment effect							1.7 (-8.4; 11.8)	
Cognition								
NIV	48.5 ± 41.0	99	44.5 ± 40.8	50	33.7 ± 36.3	50	-10.8 (-22.5; 0.8)	50
Controls	43.3 ± 35.9	90	39.7 ± 35.4	51	36.3 ± 35.5	51	-3.4 (-12.7; 5.8)	51
Treatment effect							-7.4 (-22.0; 7.2)	
Invalidity								
NIV	74.2 ± 31.2	99	70.6 ± 34.0	50	69.5 ± 34.8	50	-1.1 (-9.4; 7.2)	50
Controls	68.7 ± 33.5	90	59.2 ± 37.7	51	57.3 ± 39.4	51	-2.0 (-10.0; 6.1)	51
Treatment effect							0.9 (-10.6; 12.3)	

Data presented as means (± standard deviation) and mean change (95% confidence intervals).

ITT, intention-to-treat; MRF-28, Mageri Respiratory Questionnaire-28; n, numbers for analysis;

NIV, noninvasive positive pressure ventilation.

[¶] p<0.05 significant change after 12 months within the group.

*p<0.05 significant difference in change after 12 months between the groups (treatment effect).

Low MRF-28 scores indicate high quality of life. A negative treatment effect signifies a bigger improvement after 12 months for the NIV group compared to standard treatment.

Supplementary file Table S5: Changes in the Clinical Respiratory Questionnaire (CRQ) sub domains

		ITT Baseline	n	Completers Baseline	N	Completers 12 months	n	Change score 12 months	n
CRQ-Total									
	NIV	3.5 ± 1.1	100	3.5 ± 1.1	50	4.2 ± 1.2	50	0.7 (0.4; 1.1) [¶]	50
	Controls	3.6 ± 1.1	89	3.6 ± 1.1	50	4.4 ± 1.2	50	0.7 (0.4; 1.0) [¶]	50
	Treatment effect							0.01 (-0.4; 0.4)	
Dyspnoea									
	NIV	3.0 ± 2.7	100	3.1 ± 1.6	49	3.5 ± 1.7	49	0.4 (0.03; 0.8) [¶]	49
	Controls	3.25 ± 1.4	85	3.4 ± 1.5	46	3.9 ± 1.5	46	0.5 (-0.002; 0.9)	46
	Treatment effect							-0.02 (-0.6; 0.6)	
Fatigue									
	NIV	2.70 ± 1.3	100	2.8 ± 1.3	50	3.8 ± 1.5	50	1.0 (0.5; 1.4) [¶]	50
	Controls	2.62 ± 1.2	89	2.5 ± 1.0	50	3.7 ± 1.3	50	1.1 (0.8; 1.5) [¶]	50
	Treatment effect							-0.2 (-0.7; 0.4)	
Emotion									
	NIV	3.93 ± 1.3	100	3.9 ± 1.3		4.6 ± 1.3	50	0.7 (0.3; 1.1) [¶]	50
	Controls	4.09 ± 1.2	89	4.1 ± 1.3	50	4.7 ± 1.4	50	0.6 (0.3; 0.9) [¶]	50
	Treatment effect							0.1 (-0.4; 0.6)	
Mastery									
	NIV	3.86 ± 1.4	100	3.8 ± 1.4	50	4.6 ± 1.4	50	0.8 (0.4; 1.2) [¶]	50
	Controls	4.01 ± 1.4	89	4.0 ± 1.4	50	5.0 ± 1.4	50	0.9 (0.5; 1.3) [¶]	50
	Treatment effect							-0.1 (-0.7; 0.4)	

Data presented as means (± standard deviation) and mean change (95% confidence intervals).

CRQ, Clinical Respiratory Questionnaire; ITT, intention-to-treat; n, numbers for analysis; NIV, noninvasive positive pressure ventilation.

[¶] p<0.05 significant change after 12 months within the group.

*p<0.05 significant difference in change after 12 months between the groups (treatment effect).

High CRQ scores indicate high quality of life. A positive treatment effect signifies a bigger improvement after 12 months for the NIV group compared to standard treatment.

Supplementary file Table S6: Changes in the Severe Respiratory Insufficiency questionnaire (SRI) sub domains

	ITT	n	Completers	N	Completers	n	Change score	n
	Baseline		Baseline		12 months		12 months	
SRI-Total								
NIV	48.1 ± 15.0	100	47.9 ± 15.1	50	55.0 ± 15.4	50	7.0 (3.4; 10.7) [¶]	50
Controls	51.3 ± 15.9	90	53.6 ± 16.9	51	55.8 ± 16.3	51	2.2 (-1.2; 5.6)	51
Treatment effect							4.8 (-0.1; 9.7)	
Respiratory complaints								
NIV	47.9 ± 19.6	100	48.9 ± 21.2	50	55.6 ± 19.4	50	6.7 (2.1; 11.2) [¶]	50
Controls	44.0 ± 18.6	90	45.7 ± 18.5	51	54.3 ± 16.7	51	8.6 (3.8; 13.5) [¶]	51
Treatment effect							-2.0 (-8.6; 4.6)	
Physical functioning								
NIV	32.0 ± 18.5	100	32.7 ± 21.6	50	36.5 ± 21.6	50	3.8 (-1.4; 9.0)	50
Controls	32.5 ± 21.9	90	37.3 ± 21.4	51	41.3 ± 21.6	51	4.1 (-1.7; 9.9)	51
Treatment effect							0.3 (-8.0; 7.4)	
Attendant symp+sleep								
NIV	57.1 ± 18.4	100	54.4 ± 17.9	50	64.8 ± 18.0	50	10.4 (5.3; 15.5) [¶]	50
Controls	58.6 ± 20.0	90	59.9 ± 22.3	51	61.7 ± 18.4	51	1.7 (-2.9; 6.4)	51
Treatment effect							8.7 (1.9; 15.4)[*]	
Social relationships								
NIV	58.3 ± 17.1	100	56.3 ± 14.0	50	60.9 ± 18.9	50	4.6 (-0.2; 9.4)	50
Controls	66.5 ± 17.1	90	68.2 ± 17.3	51	64.3 ± 17.4	51	-3.9 (-7.7; 0.03)	51
Treatment effect							8.4 (2.4; 14.5)[*]	
Anxiety								
NIV	46.2 ± 22.5	100	47.0 ± 24.6	50	59.8 ± 22.1	52	12.8 (7.1; 18.5) [¶]	50
Controls	50.4 ± 25.4	90	52.6 ± 26.3	51	58.2 ± 23.0	51	5.7 (0.6; 10.7) [¶]	51
Treatment effect							7.1 (-0.4; 14.6)	
Well-being								
NIV	52.6 ± 20.1	100	52.2 ± 19.1	50	58.1 ± 22.9	50	5.9 (0.7; 11.1) [¶]	50
Controls	59.3 ± 20.4	90	60.4 ± 22.1	51	60.3 ± 21.3	51	-0.1 (-4.7; 4.4)	51
Treatment effect							6.0 (-0.8; 12.8)	
Social functioning								
NIV	43.1 ± 40.6	100	44.0 ± 19.4	50	49.1 ± 20.1	50	5.1 (-0.2; 10.4)	50
Controls	48.0 ± 20.4	90	51.4 ± 21.6	51	50.7 ± 23.1	51	-0.8 (-5.9; 4.4)	51
Treatment effect							5.9 (-1.4; 13.2)	

Data presented as means (± standard deviation) and mean change (95% confidence intervals). Attendant symp+sleep, attendant symptoms and sleep sub domain; ITT, intention-to-treat; n, numbers for analysis; NIV, noninvasive positive pressure ventilation; SRI, Severe Respiratory Insufficiency questionnaire.

[¶] p<0.05 significant change after 12 months within the group.

^{*} p<0.05 significant difference in change after 12 months between the groups (treatment effect).

High SRI scores indicate high quality of life. A positive treatment effect signifies a bigger improvement after 12 months for the NIV group compared to standard treatment.

Supplementary file Table S7: Changes in the Groningen Activity and Restriction Scale (GARS), Hospital Anxiety and Depression Scale (HADS) and Medical Research Council (MRC) sub domains

	ITT Baseline	n	Completers Baseline	n	Completers 12 months	n	Change score 12 months	n
GARS-Total								
NIV	36.7 ± 7.5	100	36.3 ± 8.3	50	34.6 ± 9.4	50	-1.6 (-3.3; 0.1)	50
Controls	36.8 ± 8.5	90	34.7 ± 1.2	51	32.7 ± 8.2	51	-2.0 (-4.1; 0.1)	51
Treatment effect							0.4 (-2.3; 3.0)	
HADS-Total								
NIV	15.9 ± 9.2	98	15.6 ± 8.5	48	13.9 ± 8.8	48	-1.7 (-4.0; 0.6)	48
Controls	14.5 ± 8.8	87	13.1 ± 9.1	50	12.7 ± 9.3	50	-0.4 (-2.2; 1.3)	50
Treatment effect							-1.3 (-4.1; 1.6)	
HADS anxiety								
NIV	7.8 ± 5.0	99	7.9 ± 4.7	49	6.6 ± 4.7	49	-1.3 (-2.3; -0.2) [¶]	49
Controls	7.3 ± 4.6	88	6.8 ± 4.8	51	5.9 ± 4.8	51	-0.9 (-1.8; 0.02)	51
Treatment effect							-0.4 (-1.8; 1.0)	
HADS depression								
NIV	8.1 ± 4.9	99	7.5 ± 4.8	50	7.1 ± 4.6	50	-0.4 (-1.7; 0.9)	50
Controls	7.1 ± 4.7	87	6.3 ± 4.7	50	6.7 ± 5.1	50	0.4 (-0.6; 1.5)	50
Treatment effect							-0.8 (0.8; -2.5)	
MRC								
NIV	3.9 ± 1.0	98	3.8 ± 1.4	49	3.5 ± 1.4	49	-0.4 (-0.7; -0.02) [¶]	49
Controls	3.8 ± 1.0	90	3.6 ± 1.1	51	3.3 ± 1.2	51	-0.3 (-0.7; 0.08)	51
Treatment effect							-0.05 (-0.6; 0.5)	

Data presented as means (± standard deviation) and mean change (95% confidence intervals).

GARS, Groninger Activity Restriction Scale; HADS, Hospital Anxiety and Depression Scale; ITT, intention-to-treat; n, numbers for analysis; NIV, noninvasive positive pressure ventilation; MRC, Medical Research Council dyspnoea scale.

[¶] p<0.05 significant change after 12 months within the group.

*p<0.05 significant difference in change after 12 months between the groups (treatment effect).

Low GARS scores indicate better daily activity levels. Low HADS scores indicate better mood state. Lower MRC scores represent less dyspnoea. A negative treatment effect signifies a bigger improvement after 12 months for the NIV group compared to standard treatment.

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