



**Rehabilitation in lung diseases: In-hospital and post
exacerbation pulmonary rehabilitation**

Journal:	<i>Respirology</i>
Manuscript ID	RES-18-1008.R1
Manuscript Type:	Invited Review
Date Submitted by the Author:	22-Jan-2019
Complete List of Authors:	Ibrahim, Wadah; University of Leicester, Department of Respiratory Science Harvey-Dunstan, Theresa; University of Nottingham, Physiotherapy and Rehabilitation Science, School of Health Sciences Greening, Neil; University of Leicester, Department of Respiratory Science; University Hospitals of Leicester NHS Trust, Centre for Exercise and Rehabilitation Science, Institute for Lung Health
Subject Category – Select <i>up to 3 subject categories</i> that best match your manuscript and list them <i>in order of preference</i>.:	COPD
Keywords - Select up to 5 keywords:	COPD, Exercise and Pulmonary Rehabilitation, Clinical Respiratory Medicine

SCHOLARONE™
Manuscripts

Editorial Office Notes:

RES -18-1008.R1

Article type: Invited review

INVITED REVIEW SERIES: Rehabilitation in chronic respiratory diseases.

SERIES EDITORS: Frits M.E. Franssen and Jennifer A. Alison

Received: 20 December 2018

Invited to revise: 16 January 2019

Revised: 22 January 2019

Accepted: 13 February 2019

Publication Fee Waiver: YES

Volume number: 24

1
2
3
4 **Rehabilitation in chronic respiratory diseases: In-hospital and post exacerbation pulmonary**

5
6
7 **rehabilitation**

8
9
10
11 Wadah Ibrahim.^{1*}, Theresa C. Harvey-Dunstan^{2*}, Neil J. Greening^{1,3}

12
13
14
15
16 ¹Department of Respiratory Science, University of Leicester, Leicester, United Kingdom

17
18 ²Division of Physiotherapy and Rehabilitation Sciences, School of Health Sciences, University of
19
20 Nottingham, UK

21
22 ³Centre for Exercise and Rehabilitation Science, Institute for Lung Health, Leicester, UK

23
24
25 *Contributed jointly to manuscript

26
27
28
29 Corresponding Author

30
31 Dr Neil Greening

32
33 NIHR Leicester Biomedical Research Centre- Respiratory

34
35
36 Glenfield Hospital, Groby Road

37
38 Leicester, LE3 9AP

39
40 United Kingdom

41
42 neil.greening@leicester.ac.uk

Abstract

Exacerbations of chronic obstructive pulmonary disease (COPD) that require hospitalisation are important events for patients. Functional impairment and skeletal muscle dysfunction can increase the risk of hospitalisation and readmission, independent of lung function. In addition, once a patient is admitted multiple factors can lead to worsening outcome including immobility, systemic inflammation and nutritional depletion. These non-pulmonary factors are potentially amenable to exercise therapy, as part of pulmonary rehabilitation. Peri-exacerbation pulmonary rehabilitation has an important role in the management of exacerbations of COPD.

In this review we explore how functional limitation and skeletal muscle dysfunction affects patients having a severe exacerbation of COPD, the systemic impact of hospitalisation on patients including potential aetiologies and the role of pulmonary rehabilitation around the time of an exacerbation. This includes rehabilitation during the inpatient phase, post-exacerbation rehabilitation and rehabilitation bridging hospital discharge. We also describe potential future developments in peri-exacerbation pulmonary rehabilitation.

Introduction

Worldwide, COPD constitutes one of the largest health burdens with 3.17 million deaths in 2015¹. In the UK, an estimated 1.2 million people (2% of the population) have diagnosed COPD making it the second commonest lung disease, after asthma².

The natural history of COPD is punctuated by acute episodes of worsening symptoms and increased airways inflammation known as exacerbations and are a major priority in the treatment of COPD. For an individual with COPD it results in marked worsening of symptoms and disease burden, far beyond the burden when in the stable state. It is also now clear that certain individuals are prone to recurrent exacerbations- known as the frequent exacerbator phenotype³. The strongest predictor of future exacerbations is a history of previous exacerbation. Patients with severe exacerbations requiring admission to hospital demonstrate this high risk, with 43% of patients in the UK readmitted within three months⁴. For healthcare organisations exacerbations account for more than half of all COPD costs. As well as considerable expense, the challenge on healthcare systems provide a logistic challenge, with exacerbations accounting for over a million bed days each year across the UK, presenting in an unpredictable fashion².

The primary pathophysiology of COPD is within the lungs and the diagnosis of COPD is confirmed by detecting airflow limitation using spirometry. However, over the past decades it has been increasingly recognised that COPD has a number of extra-pulmonary manifestations that have considerable impact on patients and the disease burden. Exercise limitation is a key symptom of COPD. It is clear that this is in part driven through leg fatigue and skeletal muscle dysfunction, with quadriceps function being an independent marker of mortality⁵.

Muscle dysfunction is more common in patients with advanced COPD, as are exacerbations of COPD. The combination of these two aspects of COPD may have important consequences, as well as specific therapeutic interventions. In this review we describe the functional limitation and skeletal

1
2
3 muscle dysfunction in patients hospitalised for an exacerbation of COPD, the acute systemic effects
4
5 on hospitalisation and the role of peri-exacerbation pulmonary rehabilitation.
6
7

8 9 10 The Impact of Skeletal Muscle Dysfunction and Functional Impairment on Exacerbations of COPD

11 Independent of disease severity, systemic markers of disease may increase the risk for patients to be
12 admitted to hospital with an exacerbation. Physical inactivity and increased sedentary time are well
13 recognised and described in COPD. Patients with reduced physical activity are at an increased risk of
14 hospital admission, readmission to hospital following an index admission and death^{6, 7}. These studies
15 have corrected for factors, such as disease severity and symptom burden.
16
17

18 Functional measures, such as the six minute walk test, have been shown to be predictive of
19 hospitalisation and prognosis. Indeed the most commonly used prognostic tool, the BODE score, is
20 part comprised of walking distance⁸. The Four meter Gait Speed (4MGS) is a well validated measure,
21 and a key part of the frailty assessment^{9, 10}. In patients hospitalised with an exacerbation of COPD,
22 4MGS taken at the time of discharge is a marker of readmission, independent of disease severity¹¹.
23 Low skeletal muscle mass is also known to be an independent risk factor for hospital admission and
24 mortality. Body mass index (BMI) is a crude measure of body composition, and like 6MWT, is one of
25 the components of the BODE score, which predicts hospitalisation and mortality¹². Direct measures
26 of the skeletal muscle, such as the fat free mass and quadriceps function, have been shown to be a
27 better prognostic indicator than BMI^{13, 14}. In those admitted to hospital, patients with the smallest
28 muscles measured at the bedside using ultrasound are more likely to be readmitted or die,
29 independent of other markers of disease severity or risk of readmission¹⁵.
30
31
32
33
34
35
36
37
38
39
40
41
42
43
44
45
46
47
48
49
50
51

52 Systemic Consequences of Exacerbations of COPD

53
54
55
56
57 Exacerbations of COPD present as increasing dyspnoea as a result of increased airways inflammation
58 and require additional therapy to treat the lung. However, it is clear that exacerbations impact
59
60

1
2
3 various body systems with a number of acute non-pulmonary effects. Similar to stable disease, the
4 skeletal muscle system and function to walk are important factors affected during the exacerbating
5 state. In particular severe exacerbations, which require admission to hospital, have a profound
6 effect with multiple insults all contributing to adverse impact on the whole body. This leads to a
7 number of important consequences, such as reduced ability to protect against future insults, i.e.
8 reduced resilience, and increased risk of hospital readmission and death (figure 1).
9
10
11
12
13
14
15
16
17
18

19 **figure 1 here**

20 21 22 23 Factors that impact on Systemic Factors of COPD in Exacerbations

24 25 26 27 Immobility

28 Enforced bed rest has been used as a treatment since the 19th century for a range of medical
29 conditions since Victorian times including hysteria¹⁶ and tuberculosis¹⁷. Bed rest has continued as a
30 mainstay of therapy until the late 20th century¹⁸. As such public perception of bed rest as therapy
31 for acute illness continues to today.

32 Bed rest studies in healthy populations have shown significant reductions in both young¹⁹ and
33 older²⁰ healthy volunteers, with larger decreases in the muscles of ambulation compared to other
34 muscle groups¹⁹. These findings are echoed in the critical care population with acute skeletal muscle
35 loss²¹ and persistent long-term functional deficit^{22, 23}.

36 Whilst the effects of acute immobility are well recognised in those previously fit and well, the effects
37 may be more marked in those with long-term conditions. In patients with COPD in the stable state
38 there is preferential loss of oxidative type 1 muscle fibres and loss of mitochondria, which may
39 impede the ability to protect against the detrimental effects of bedrest^{24, 25}. In the acute care setting
40 patients with COPD have increased symptoms of fatigue²⁶ and breathlessness, resulting in them
41 having the perception of being too ill or frail or disabled to take part in physical activity²⁶.
42
43
44
45
46
47
48
49
50
51
52
53
54
55
56
57
58
59
60

Systemic inflammation

The mechanisms for the systemic effects of COPD are varied and multi-factorial. It has been shown that a sub-set of patients with COPD have an excess of systemic inflammation and those with persistent systemic inflammation have a poorer prognosis and are more prone to exacerbation²⁷.

Systemic inflammation has also been linked with increased risk of co-morbidities, such as cardiovascular disease²⁸.

At the time of exacerbations, systemic inflammation is particularly marked even in those without persistent inflammation. In the most severe exacerbations, such as those undergoing hospitalisation, inflammatory markers measures such as C - reactive protein are increased to a mean of 81 mg/l (SD 99)²⁹.

An increased systemic inflammatory response is associated with worse skeletal muscle function, with higher circulating Interleukin-8 associated with lower quadriceps strength³⁰. Whether this inflammatory effect directly impacts on skeletal muscle or is an associated inflammatory response based on severity of exacerbation is unknown. Indeed, no increase in inflammatory cells are seen in quadriceps muscle at the time of exacerbation, questioning the direct impact of inflammation on acute muscle dysfunction³¹.

Nutritional depletion

Involuntary weight loss, prevalent in patients with advanced COPD, is a well-recognised poor prognostic factor³²⁻³⁴ and is associated with higher mortality³⁵.

Weight loss in COPD is multifactorial, with reduced oral intake, increased work of breathing and associated systemic inflammation. Acutely, exacerbations increase basal metabolic expenditure, when unmatched with adequate caloric intake can lead to nutritional depletion, muscle wasting and dysfunction³⁶. Although nutritional supplementation in stable cachectic COPD patients is recommended, studies on nutritional repletion in the acute state have not shown efficacy.

1
2
3 Vermeeren et al demonstrated no additional improvements in lung function or muscle strength with
4 caloric supplements during exacerbations³⁷. Rutten et al described loss of intestinal integrity in COPD
5 patients, so profound it was suggested that this is added as a new component in the multisystem
6 disorder, and that simply increasing calorie intake may be inadequate³⁸.
7
8
9

10 11 12 13 14 Hospital-Associated Disability

15
16 The detrimental effects described above lead to a loss of physical function (figure 2). In addition to
17 this, patients in hospital have altered sleep patterns, acute cognitive impairment and loss of
18 autonomy. The resulting syndrome has been described as a number of phenomena, including
19 hospital-associated disability, hospital acquired disability and post hospitalisation syndrome.
20
21 Hospital-associated disability (HAD) is described in the geriatric population as a loss of activity of
22 daily living (ADL) due to admission to hospital³⁹. HAD may take months to recover or lead to
23 irreversible changes. Despite the clinical recognition of the problem the process and natural history
24 of HAD is poorly understood. Whilst HAD has been most widely described in the geriatric population
25 it is likely to be a key component in chronic diseases, including COPD.
26
27
28
29
30
31
32
33
34
35
36
37
38

39 **Figure 2 here**

40 41 42 43 Rationale for exercise training

44
45 Currently the treatment for exacerbations of COPD is mainly focused on the primary insult to the
46 lungs. Exercise therapy potentially lends itself to the treatment of HAD and patients with advanced
47 COPD and pre-existing functional limitation.
48
49

50
51 Pulmonary rehabilitation in the stable setting is one of the highest value therapies in COPD and is
52 recommended worldwide in symptomatic patients^{40, 41}. Pulmonary rehabilitation consists of
53 individualised exercises, educational and behavioural elements targeted at minimising symptom
54 burden, maximising exercise performance, promoting autonomy and enhancing quality of life ⁴⁰.
55
56
57
58
59
60

1
2
3 Therefore applying pulmonary rehabilitation around the time of an exacerbation, also known as
4
5 peri-exacerbation pulmonary rehabilitation, has been trialled and is now internationally
6
7 recommended. The remainder of this review will examine the evidence for pulmonary rehabilitation
8
9 initiated following hospitalisation for an exacerbation of COPD-post exacerbation pulmonary
10
11 rehabilitation (PEPR), pulmonary rehabilitation initiated during hospitalisation, and possible future
12
13 developments.
14
15
16
17
18

19 Post Exacerbation Pulmonary Rehabilitation

20
21 PEPR has had a major impact on the treatment of severe exacerbations of COPD and is
22
23 recommended in both National and International guidelines^{40, 41}. Trials of PEPR have typically
24
25 performed PR programmes similar in structure and duration to PR programmes in the stable state,
26
27 but within the recovery period from severe exacerbations (defined as requiring hospitalisation).
28
29 The first randomised controlled trial to investigate PEPR was Man et al in 2004⁴². 42 patients were
30
31 recruited within 10 days of discharge from hospital and underwent an eight week PR programme. In
32
33 this landmark study, patients in the usual care group had not recovered at eight weeks, with a
34
35 median incremental shuttle walk (ISWT) of only 90 meters. In comparison patients who underwent
36
37 PEPR ISWT improved by 90m, double the minimal clinically important difference. Large clinically and
38
39 statistically significant differences were also noted in health related quality of life.
40
41
42

43
44 The results of Man et al have been replicated in a number of other similar sized trials (Murphy 2005,
45
46 Ghanem 2010, Deepak 2014), which use a similar model of 2-3 supervised exercise sessions per
47
48 week starting following discharge⁴³⁻⁴⁵. A number of the trials have used supervised home-based
49
50 exercise, rather than in a formal setting, potentially increasing uptake of patients who do not need
51
52 to travel^{43, 45, 46}. Only one trial (Ko et al 2011⁴⁷) did not show benefit of PEPR in terms of exercise
53
54 capacity or health related quality of life.
55

56
57 One of the secondary outcome measures in the trial by Man et al was readmission to hospital. The
58
59 PEPR group had a significant reduction in rate of hospital admission at 3 months. The rationale, as
60

1
2
3 described above, was likely to be training the non-pulmonary deficits occurring following
4
5 exacerbation and increasing resilience. Seymour et al⁴⁸, part of the same research group as Man et al
6
7 performed a larger RCT, powered to detect a significant reduction in readmission to hospital. They
8
9 recruited 60 patients in a similar model across four hospitals in London. Exercise capacity,
10
11 quadriceps strength, health related quality of life improved similarly to the previous trial. An 85%
12
13 reduction in hospital readmission in the PEPR group was seen during the intervention period.
14
15 The largest trial of PEPR was reported by Ko et al in 2017⁴⁶. PEPR was a component of a larger
16
17 comprehensive care programme, which also included specialist medical review, telephone support
18
19 line, and regular specialist follow-up. 180 patients were included in the study. 33% (n=30) of the
20
21 intervention group received supervised outpatient physiotherapy and 67% (n=60) unsupervised
22
23 home exercise. Similar to the Seymour study a significant reduction in the rate of hospital
24
25 readmissions was seen in the intervention group (incident rate ratio 0.668), with the greatest
26
27 difference seen in the first three months. Improvements were also seen in health related quality of
28
29 life questionnaires, but not in exercise capacity.

30
31
32 The strength of these trials has unequivocally shown strong efficacy of PEPR for severe
33
34 exacerbations of COPD. However, real-life clinical implementation has proven more challenging.
35
36 Recruitment and patient uptake of PEPR has been low in the UK. Jones et al demonstrated that only
37
38 9.6% of participants received and completed PEPR despite an active Pulmonary Rehabilitation
39
40 service⁴⁹. This low uptake had been indicated in Seymour et al who took three years to recruit 60
41
42 patients across four London hospitals⁴⁸. Therefore, strategies to improve uptake to PEPR or
43
44 alternative models of delivering treatment are needed.
45
46
47
48
49
50
51

52 Pulmonary Rehabilitation and Exercise Training During the Hospitalisation

53
54 As described above trials of PEPR have yielded considerable benefits for the patients who attend a
55
56 programme. Whilst it helps the recovery from the impact of hospital associated disability and the
57
58 impact of the exacerbation, it does not prevent or minimise the primary loss of physical function. In
59
60

1
2
3 order to target this, intervention may be required to start during the period of hospitalisation,
4
5 ideally as early as possible. Due to nature of a severe exacerbation and acute illness of inpatients
6
7 conventional models of pulmonary rehabilitation programmes are difficult to implement and
8
9 interventions have been highly varied. For the purposes of this review we have considered exercise
10
11 interventions performed during the hospitalisation only (i.e. ended at time of discharge), including
12
13 non-volitional techniques such as neuromuscular electrical stimulation (NMES) and training that
14
15 includes both during the inpatient and post-discharge periods.
16
17
18
19
20

21 Training during the inpatient phase only

22
23 Two trials of inpatient rehabilitation reported outcomes 20 years ago^{50, 51}. Nava et al recruited 80
24
25 patients in a respiratory intensive care unit with COPD. Length of hospital stay in these patients was
26
27 more than one month. Those who received exercise intervention (comprising progressive aerobic
28
29 training and respiratory muscle training) improved their walking capacity, whereas those in the usual
30
31 care group (not including aerobic training) had no improvement in walking. Baseline measures were
32
33 conducted as soon as patients were able, as 76% had required mechanical ventilation initially⁵⁰.
34
35 Kirsten et al also demonstrated improvements in walking capacity following aerobic training over
36
37 usual care in 29 patients admitted with an exacerbation of COPD. Both trials delivered their
38
39 intervention later in the admission, following initial recovery; Kirsten et al starting at days 6-8 of
40
41 admission and Nava et al once able to perform cycling⁵¹.
42
43
44
45 More recently Troosters et al⁵² used resistance training to target the skeletal muscle to prevent loss
46
47 of quadriceps strength. Patients were recruited in their first day on a respiratory ward and
48
49 underwent a mean of 6 days of training, initially at 70% of 1 repetition max (1RM). Efficacy was
50
51 shown with a 10% increase in strength in the intervention group. This was sustained at one month
52
53 following discharge. Improvement in 6MWT was also seen in the intervention group, though this
54
55 was not statistically different to the control group. No difference in hospitalisation rate was seen at
56
57 six months.
58
59
60

1
2
3 He et al⁵³ also recruited patients on their second day of admission in 101 patients admitted with
4 COPD. Patients underwent twice daily aerobic and resistance training. Those who received inpatient
5 training improved their 6MWT, whereas those who had usual care showed no difference, though
6 between groups were not reported. 6MWT on at time of admission was relatively well preserved
7 (approximately 250m) and duration of intervention was not reported. In a similar study Liao et al⁵⁴
8 performed twice daily aerobic and resistance training for four days in 61 patients admitted with an
9 exacerbation. Large increases (120m) in 6MWT were seen in the intervention group but not the
10 control group. Similar improvements were also seen in dyspnoea.

11
12 Overall short-term the trials have shown that supervised training in patients who are inpatients
13 benefit in terms of function, though the longer term effects are unknown.
14
15
16
17
18
19
20
21
22
23
24
25
26
27

28 Non-Volitional training

29
30 With the increased symptom burden at the time of an exacerbation training techniques that reduce
31 the ventilatory burden of training are attractive. Neuromuscular electrical stimulation (NMES)
32 directly stimulates skeletal muscle using electrical current resulting in muscle contraction with
33 minimal ventilatory demand⁵⁵. NMES has been used in the stable state in COPD and showed short
34 term improvements in muscle function. While the effects are short-lived, it may be well placed to
35 bridge the acute hospital admission before PEPR. Three trials have explored the use of NMES in the
36 peri-exacerbation period; one in the intensive care unit⁵⁶, one during hospitalisation on a medical
37 ward⁵⁷ and one in a rehabilitation unit following hospitalisation⁵⁸. All three trials showed short-term
38 benefit of NMES clinical efficacy in terms of statistically significant differences in muscle quadriceps
39 strength and walking distance. However, use of NMES in the clinical setting at this time are unlikely
40 as conclusions using this modality are limited with small numbers of patients in all trials.
41
42
43
44
45
46
47
48
49
50
51
52
53
54
55
56
57
58
59
60

57 Training crossing both inpatient and outpatient phase

1
2
3 Several trials have attempted to cover the wider peri-exacerbation period, including both during
4 hospitalisation and the post-hospitalisation recovery period⁵⁹⁻⁶¹. In the earliest trial by Behnke et al,
5
6 30 participants underwent 10 days of inpatient training, followed by six months of unsupervised
7
8 home-based training⁵⁹. They were recruited during the recovery phase of the admission (days 4-7).
9
10 In a per-protocol analysis (65% of recruited patients) significant improvements in walking
11
12 performance and symptoms were seen in the intervention group, when compared with the control
13
14 group. The differences in exercise capacity were seen in the first 10 days of intervention, with an
15
16 improvement in the intervention group of 225m on the 6MWT, but no change in the control group.
17
18 Two other, more recent trials, have trialled the wider peri-exacerbation period. Both are two of the
19
20 larger RCTs in the field of peri-exacerbation pulmonary rehabilitation and also attempted to recruit
21
22 earlier into the hospitalisation than the Behnke trial. Eaton et al⁶⁰ recruited 97 patients which
23
24 included inpatient rehabilitation followed by a standard PEPR programme. No difference was seen
25
26 between groups, for hospital readmission, exercise capacity or quality of life. However, in the per-
27
28 protocol sub-analysis of the 19 participants who attended the PEPR aspect a reduction in hospital
29
30 readmission was seen, providing encouraging results.
31
32 In the largest trial in the peri-exacerbation pulmonary rehabilitation literature Greening et al⁶¹
33
34 incorporated several inpatient aspects of therapy including NMES, resistance training and aerobic
35
36 training. Following hospital discharge, in an attempt to increase participation and stop the high
37
38 drop-out associated with PEPR a home-based programme with telephone support, but unsupervised
39
40 training was provided. 389 participants were recruited. However, length of hospital stay was lower
41
42 than in other trials (median 5 days), meaning the inpatient (supervised) intervention was limited.
43
44 The home-based programme was poorly adhered to. Similar to Eaton et al, no between group
45
46 differences in exercise capacity, health status or hospital readmission were seen. Surprisingly,
47
48 separation in mortality was seen starting more than 6 months after the intervention finished, in
49
50 favour of the usual care group. This is unlikely to be due to the intervention, as was separated
51
52 temporally from the intervention. No signal was seen in the per-protocol analysis and is more likely
53
54
55
56
57
58
59
60

1
2
3 explained by case-mix of the groups, as those in the intervention group had higher baseline MRC
4 score. Patients in the intervention group were also less likely to uptake standard supervised
5 pulmonary rehabilitation after three months which may have impacted longer term outcomes.
6
7 Another key difference in both the Eaton and Greening trials was that large increases in exercise
8 capacity and quality of life in usual care group were observed. This is different from other reported
9 trials, where the control groups have remained unimproved months later. In the trial by Greening et
10 al the usual care groups received daily ward-based physiotherapy, as per usual UK practice, which
11 may also explain the recovery seen in the usual care groups and lack of benefit with additional
12 rehabilitation strategies.
13
14 Trials of peri-exacerbation pulmonary rehabilitation that include both inpatient and outpatient
15 training have so far proven less successful than trials focussing on either the inpatient or
16 outpatient components. Difficulties in adapting training across changing environments, without
17 resulting in large drop-out need to be addressed before this is likely to be as successful as other
18 areas of peri-exacerbation pulmonary rehabilitation.
19
20
21
22
23
24
25
26
27
28
29
30
31
32
33
34
35
36

37 **Table 1 here**

38 Peri-Exacerbation Pulmonary Rehabilitation in Non-COPD Patients

39
40
41 Until the last few years research in pulmonary rehabilitation in the stable state has almost
42 exclusively focussed on COPD. Echoing this, peri-exacerbation pulmonary rehabilitation has not been
43 trialled in non-COPD respiratory conditions. However, the rationale for it is no less strong in other
44 respiratory conditions as it targets the non-pulmonary elements of the disease, other respiratory
45 conditions experience exacerbations, and is effective in other conditions in the stable state. 15% of
46 patients in Greening et al⁶¹ had other chronic respiratory diseases (bronchiectasis, interstitial lung
47 disease and chronic asthma) with no difference in outcome between those with COPD and those
48 who didn't. Trials in other respiratory conditions such as bronchiectasis are underway and awaited⁶².
49
50
51
52
53
54
55
56
57
58
59
60

1
2
3 Similarly, rehabilitation around the time of hospitalisation has been successfully implemented in
4
5 other diseases and elderly populations^{63, 64}.
6
7

8 9 10 The Future Direction of Peri-Exacerbation Pulmonary Rehabilitation

11
12 More so than pulmonary rehabilitation in the stable state, peri-exacerbation pulmonary
13
14 rehabilitation has a number of unanswered questions and is likely to evolve over the next few years.
15
16 PEPR has the best efficacy, but with poor uptake has limited application to the wider population.
17
18 Strategies to increase uptake to post-exacerbation pulmonary rehabilitation are therefore
19
20 important. One pragmatic approach to increasing uptake post-exacerbation rehabilitation would be
21
22 to delay the start of rehabilitation until recovery from the hospitalisation is complete, when patients
23
24 may be more willing to attend classes. This would ensure increased participation in pulmonary
25
26 rehabilitation in a high-risk, symptomatic population who would gain benefit from a more standard
27
28 programme of pulmonary rehabilitation. Trials comparing early and late post exacerbation
29
30 pulmonary rehabilitation have proven challenging^{65, 66}. One trial showed no long-term difference
31
32 between patients recruited to either immediate post exacerbation pulmonary rehabilitation or six
33
34 months later, but recruited only 15% of its total planned population⁶⁶. Results of other ongoing trials
35
36 of early versus late pulmonary rehabilitation are awaited⁶⁷.
37
38

39
40
41 Reasons for lack of uptake to PEPR are numerous, linked to both patient perceptions of their current
42
43 illness and high risk of early hospital readmission²⁶. Health coaching following hospitalisation has
44
45 recently been shown as a possible intervention to address these aspects. In a study by Benzo et al⁶⁸
46
47 patients received a motivational interviewing based health coaching intervention. During the
48
49 intervention period there was a reduction in hospital readmission rate, as well improvements in
50
51 health related quality of life. No difference in physical function was found. Techniques such as health
52
53 coaching, may therefore work as a bridge to pulmonary rehabilitation, which may then offer
54
55 additional benefits, such as increased exercise capacity and longer term benefits.
56
57
58
59
60

1
2
3 Growing internet use and web literacy, including among the older population make web-based
4 interventions a potential avenue. Successful trials in the stable state have been published and may
5
6 interventions a potential avenue. Successful trials in the stable state have been published and may
7 allow patients who struggle to attend rehabilitation programmes during the recovery phase a more
8 structured and supervised home-based programme⁶⁹. However, increased transient cognitive
9
10 impairment is common in the recently hospitalised population⁷⁰ and may impact on the ability to
11 deliver more self-management directed interventions. Web literacy may also be lower than reported
12
13 in this population⁷¹.

14
15 Improving the inpatient aspect of rehabilitation also requires further research. The acuity of the
16
17 illness may prevent training similar to an outpatient PR programme, with some trials struggling to
18
19 recruit⁷², but other strategies may continue to develop. Anabolic agents have been trialled in the
20
21 stable state^{73, 74}, including those with advanced disease⁷⁵. Whilst effective at increasing muscle mass,
22
23 a lack of functional improvement above exercise therapy has limited their use in the stable state.
24
25 However, in the early acute care setting they may be able to protect against HAD when ability to
26
27 exercise may be limited.

28 29 30 31 32 33 34 35 36 37 Summary

38
39 Acute exacerbations of COPD are a major event for patients with COPD. They have considerable
40
41 adverse effects on the whole body, which is historically not treated with medical therapy. Post-
42
43 exacerbation pulmonary rehabilitation offers excellent efficacy and improvements in exercise
44
45 capacity, health related quality of life and reduced readmission to hospital⁷⁶. Strategies to improve
46
47 uptake to PEPR are needed however, as this has been limited. Interventions during the
48
49 hospitalisation seem to offer short term functional benefit, but trials attempting to cross both
50
51 inpatient and outpatient settings have been limited, probably due to lack of supervised training. In
52
53 summary it is important that all potentially eligible patients are given every opportunity and support
54
55 to enrol in pulmonary rehabilitation programme following an exacerbation of COPD.
56
57
58
59
60

Funding

Dr Greening is funded by a NIHR post-doctoral fellowship (PDF-2017-10-052). Dr Greening and Dr Ibrahim are supported by the NIHR Leicester Biomedical Research Centre- Respiratory Theme. The views expressed in this publication are those of the author(s) and not necessarily those of the NHS, the National Institute for Health Research, Health Education England or the Department of Health.

The Authors:

Dr Wadah Ibrahim is a clinical research fellow at the University of Leicester. He is currently taking time out of his specialist training in respiratory medicine to study for a PhD. His work mainly focuses on patients with acute exacerbations of airways disease and identifying volatile organic compounds in breath. Dr Theresa Harvey-Dunstan is an Assistant Professor and Physiotherapist at the University of Nottingham. She has spent the last ten years in research within pulmonary rehabilitation, including peri-exacerbation pulmonary rehabilitation. More recently she has completed her PhD, which compared the sensitivity and response of different exercise tests to pulmonary rehabilitation in COPD. Dr Neil Greening is an Associate Professor and Consultant Respiratory Physician at the University of Leicester. His main research interests are understanding systemic changes to patients with exacerbations of COPD. In 2017 he was awarded a national fellowship by the National Institute of Health Research (NIHR) to continue his work in this field.

References

- 1 World Health Organisation. Chronic Obstructive Pulmonary Disease: Key facts. 2016.
- 2 Snell N, Strachan D, Hubbard R, Gibson J, Gruffydd-Jones K, Jarrold I. S32
Epidemiology of chronic obstructive pulmonary disease (COPD) in the uk: findings from the
british lung foundation's 'respiratory health of the nation' project. *Thorax*. 2016; **71**: A20-A.
- 3 Hurst JR, Vestbo J, Anzueto A, et al. Susceptibility to exacerbation in chronic
obstructive pulmonary disease. *The New England journal of medicine* 2010; **363**(12): 1128-
38.
- 4 2015–18 NCAPscw. COPD: Who cares when it matters most? – outcomes report.
Royal College of Physicians, 2017.
- 5 Swallow EB, Reyes D, Hopkinson NS, Man WD, Porcher R, Cetti EJ, Moore AJ,
Moxham J, Polkey MI. Quadriceps strength predicts mortality in patients with moderate to
severe chronic obstructive pulmonary disease. *Thorax*. 2007; **62**: 115-20.
- 6 Pitta F, Troosters T, Probst VS, Spruit MA, Decramer M, Gosselink R. Physical activity
and hospitalization for exacerbation of COPD. *Chest*. 2006; **129**: 536-44.
- 7 Chawla H, Bulathsinghala C, Tejada JP, Wakefield D, ZuWallack R. Physical activity as
a predictor of thirty-day hospital readmission after a discharge for a clinical exacerbation of
chronic obstructive pulmonary disease. *Ann Am Thorac Soc*. 2014; **11**: 1203-9.
- 8 Celli BR, Cote CG, Marin JM, Casanova C, Montes de Oca M, Mendez RA, Pinto Plata
V, Cabral HJ. The body-mass index, airflow obstruction, dyspnea, and exercise capacity index
in chronic obstructive pulmonary disease. *The New England journal of medicine*. 2004; **350**:
1005-12.
- 9 Abellan van Kan G, Rolland Y, Andrieu S, Bauer J, Beauchet O, Bonnefoy M, Cesari M,
Donini LM, Gillette Guyonnet S, Inzitari M, Nourhashemi F, Onder G, Ritz P, Salva A, Visser
M, Vellas B. Gait speed at usual pace as a predictor of adverse outcomes in community-
dwelling older people an International Academy on Nutrition and Aging (IANA) Task Force. *J
Nutr Health Aging*. 2009; **13**: 881-9.
- 10 Kon SSC, Patel MS, Canavan JL, Clark AL, Jones SE, Nolan CM, Cullinan P, Polkey MI,
Man WD-C. Reliability and validity of 4-metre gait speed in COPD. *European Respiratory
Journal*. 2013; **42**: 333-40.
- 11 Kon SS, Jones SE, Schofield SJ, Banya W, Dickson MJ, Canavan JL, Nolan CM,
Haselden BM, Polkey MI, Cullinan P, Man WD. Gait speed and readmission following
hospitalisation for acute exacerbations of COPD: a prospective study. *Thorax*. 2015; **70**:
1131-7.
- 12 Schols AM, Wouters EF, Soeters PB, Westerterp KR. Body composition by
bioelectrical-impedance analysis compared with deuterium dilution and skinfold
anthropometry in patients with chronic obstructive pulmonary disease. *The American
journal of clinical nutrition*. 1991; **53**: 421-4.
- 13 Schols AM, Broekhuizen R, Weling-Scheepers CA, Wouters EF. Body composition and
mortality in chronic obstructive pulmonary disease. *The American journal of clinical
nutrition*. 2005; **82**: 53-9.
- 14 Marquis K, Debigare R, Lacasse Y, LeBlanc P, Jobin J, Carrier G, Maltais F. Mid thigh
muscle cross-sectional area is a better predictor of mortality than body mass index in

- 1
2
3 patients with chronic obstructive pulmonary disease. *American journal of respiratory and*
4 *critical care medicine*. 2002; **166**: 809-13.
- 5
6 15 Greening NJ, Harvey-Dunstan TC, Chaplin EJ, Vincent EE, Morgan MD, Singh SJ,
7 Steiner MC. Bedside Assessment of Quadriceps Muscle Using Ultrasound Following
8 Admission for Acute Exacerbations of Chronic Respiratory Disease. *American journal of*
9 *respiratory and critical care medicine*. 2015.
- 10
11 16 Mitchell SW. *Rest in the Treatment of Nervous Disease*. G.P. Putnam's Sons, 1875.
- 12
13 17 Hurt R. Tuberculosis sanatorium regimen in the 1940s: a patient's personal diary.
14 *Journal of the Royal Society of Medicine*. 2004; **97**: 350-3.
- 15
16 18 Deyo RA, Diehl AK, Rosenthal M. How many days of bed rest for acute low back
17 pain? A randomized clinical trial. *N Engl J Med*. 1986; **315**: 1064-70.
- 18
19 19 LeBlanc AD, Schneider VS, Evans HJ, Pientok C, Rowe R, Spector E. Regional changes
20 in muscle mass following 17 weeks of bed rest. *Journal of applied physiology*. 1992; **73**:
21 2172-8.
- 22
23 20 Kortebein P, Ferrando A, Lombeida J, Wolfe R, Evans WJ. Effect of 10 days of bed rest
24 on skeletal muscle in healthy older adults. *JAMA : the journal of the American Medical*
25 *Association*. 2007; **297**: 1772-4.
- 26
27 21 Puthuchery ZA, Rawal J, McPhail M, Connolly B, Ratnayake G, Chan P, Hopkinson
28 NS, Phadke R, Dew T, Sidhu PS, Velloso C, Seymour J, Agle CC, Selby A, Limb M, Edwards
29 LM, Smith K, Rowlerson A, Rennie MJ, Moxham J, Harridge SD, Hart N, Montgomery HE.
30 Acute skeletal muscle wasting in critical illness. *JAMA : the journal of the American Medical*
31 *Association*. 2013; **310**: 1591-600.
- 32
33 22 Herridge MS, Cheung AM, Tansey CM, Matte-Martyn A, Diaz-Granados N, Al-Saidi F,
34 Cooper AB, Guest CB, Mazer CD, Mehta S, Stewart TE, Barr A, Cook D, Slutsky AS. One-Year
35 Outcomes in Survivors of the Acute Respiratory Distress Syndrome. *New England Journal of*
36 *Medicine*. 2003; **348**: 683-93.
- 37
38 23 Herridge MS, Tansey CM, Matté A, Tomlinson G, Diaz-Granados N, Cooper A, Guest
39 CB, Mazer CD, Mehta S, Stewart TE, Kudlow P, Cook D, Slutsky AS, Cheung AM. Functional
40 Disability 5 Years after Acute Respiratory Distress Syndrome. *New England Journal of*
41 *Medicine*. 2011; **364**: 1293-304.
- 42
43 24 Gosker HR, Zeegers MP, Wouters EF, Schols AM. Muscle fibre type shifting in the
44 vastus lateralis of patients with COPD is associated with disease severity: a systematic
45 review and meta-analysis. *Thorax*. 2007; **62**: 944-9.
- 46
47 25 Whittom F, Jobin J, Simard PM, Leblanc P, Simard C, Bernard S, Belleau R, Maltais F.
48 Histochemical and morphological characteristics of the vastus lateralis muscle in patients
49 with chronic obstructive pulmonary disease. *Medicine and science in sports and exercise*.
50 1998; **30**: 1467-74.
- 51
52 26 Benzo R, Wetzstein M, Neuenfeldt P, McEvoy C. Implementation of physical activity
53 programs after COPD hospitalizations: Lessons from a randomized study. *Chronic*
54 *respiratory disease*. 2015; **12**: 5-10.
- 55
56 27 Agustí A, Edwards LD, Rennard SI, MacNee W, Tal-Singer R, Miller BE, Vestbo J,
57 Lomas DA, Calverley PM, Wouters E, Crim C, Yates JC, Silverman EK, Coxson HO, Bakke P,
58 Mayer RJ, Celli B, Investigators EoCLtIPSEE. Persistent systemic inflammation is associated
59 with poor clinical outcomes in COPD: a novel phenotype. *PloS one*. 2012; **7**: e37483.
- 60
61 28 Sin DD, Man SF. Why are patients with chronic obstructive pulmonary disease at
62 increased risk of cardiovascular diseases? The potential role of systemic inflammation in
63 chronic obstructive pulmonary disease. *Circulation*. 2003; **107**: 1514-9.

- 1
2
3 29 Bafadhel M, Greening NJ, Harvey-Dunstan TC, Williams JE, Morgan MD, Brightling CE,
4 Hussain SF, Pavord ID, Singh SJ, Steiner MC. Blood Eosinophils and Outcomes in Severe
5 Hospitalized Exacerbations of COPD. *Chest*. 2016; **150**: 320-8.
6
7 30 Spruit MA, Gosselink R, Troosters T, Kasran A, Gayan-Ramirez G, Bogaerts P, Bouillon
8 R, Decramer M. Muscle force during an acute exacerbation in hospitalised patients with
9 COPD and its relationship with CXCL8 and IGF-I. *Thorax*. 2003; **58**: 752-6.
10
11 31 Crul T, Spruit MA, Gayan-Ramirez G, Quarck R, Gosselink R, Troosters T, Pitta F,
12 Decramer M. Markers of inflammation and disuse in vastus lateralis of chronic obstructive
13 pulmonary disease patients. *European journal of clinical investigation*. 2007; **37**: 897-904.
14
15 32 Vermeeren MA, Creutzberg EC, Schols AM, Postma DS, Pieters WR, Roldaan AC,
16 Wouters EF, Group CS. Prevalence of nutritional depletion in a large out-patient population
17 of patients with COPD. *Respir Med*. 2006; **100**: 1349-55.
18
19 33 Wagner PD. Possible mechanisms underlying the development of cachexia in COPD.
20 *Eur Respir J*. 2008; **31**: 492-501.
21
22 34 Schols AM. Nutrition in chronic obstructive pulmonary disease. *Curr Opin Pulm Med*.
23 2000; **6**: 110-5.
24
25 35 Vandenberg E, Van de Woestijne KP, Gyselen A. Weight changes in the terminal
26 stages of chronic obstructive pulmonary disease. Relation to respiratory function and
27 prognosis. *The American review of respiratory disease*. 1967; **95**: 556-66.
28
29 36 Gea J, Agustí A, Roca J. Pathophysiology of muscle dysfunction in COPD. *J Appl*
30 *Physiol* (1985). 2013; **114**: 1222-34.
31
32 37 Vermeeren MA, Wouters EF, Geraerts-Keeris AJ, Schols AM. Nutritional support in
33 patients with chronic obstructive pulmonary disease during hospitalization for an acute
34 exacerbation; a randomized controlled feasibility trial. *Clinical nutrition*. 2004; **23**: 1184-92.
35
36 38 Rutten EPA, Lenaerts K, Buurman WA, Wouters EFM. Disturbed intestinal integrity in
37 patients with COPD: effects of activities of daily living. *Chest*. 2014; **145**: 245-52.
38
39 39 Chodos AH, Kushel MB, Greysen SR, Guzman D, Kessell ER, Sarkar U, Goldman LE,
40 Critchfield JM, Pierluissi E. Hospitalization-Associated Disability in Adults Admitted to a
41 Safety-Net Hospital. *J Gen Intern Med*. 2015; **30**: 1765-72.
42
43 40 Spruit MA, Singh SJ, Garvey C, ZuWallack R, Nici L, Rochester C, Hill K, Holland AE,
44 Lareau SC, Man WD, Pitta F, Sewell L, Raskin J, Bourbeau J, Crouch R, Franssen FM, Casaburi
45 R, Vercoulen JH, Vogiatzis I, Gosselink R, Clini EM, Effing TW, Maltais F, van der Palen J,
46 Troosters T, Janssen DJ, Collins E, Garcia-Aymerich J, Brooks D, Fahy BF, Puhan MA,
47 Hoogendoorn M, Garrod R, Schols AM, Carlin B, Benzo R, Meek P, Morgan M, Rutten-van
48 Molken MP, Ries AL, Make B, Goldstein RS, Dowson CA, Brozek JL, Donner CF, Wouters EF,
49 Rehabilitation AETFoP. An official American Thoracic Society/European Respiratory Society
50 statement: key concepts and advances in pulmonary rehabilitation. *American journal of*
51 *respiratory and critical care medicine*. 2013; **188**: e13-64.
52
53 41 Bolton CE, Bevan-Smith EF, Blakey JD, Crowe P, Elkin SL, Garrod R, Greening NJ,
54 Heslop K, Hull JH, Man WD, Morgan MD, Proud D, Roberts CM, Sewell L, Singh SJ, Walker PP,
55 Walmsley S, British Thoracic Society Pulmonary Rehabilitation Guideline Development G,
56 British Thoracic Society Standards of Care C. British Thoracic Society guideline on pulmonary
57 rehabilitation in adults. *Thorax*. 2013; **68 Suppl 2**: ii1-30.
58
59 42 Man WD, Polkey MI, Donaldson N, Gray BJ, Moxham J. Community pulmonary
60 rehabilitation after hospitalisation for acute exacerbations of chronic obstructive pulmonary
disease: randomised controlled study. *Bmj*. 2004; **329**: 1209.

- 1
2
3
4 43 Murphy N, Bell C, Costello RW. Extending a home from hospital care programme for
5 COPD exacerbations to include pulmonary rehabilitation. *Respiratory medicine*. 2005; **99**:
6 1297-302.
- 7 44 Deepak TH, Mohapatra PR, Janmeja AK, Sood P, Gupta M. Outcome of pulmonary
8 rehabilitation in patients after acute exacerbation of chronic obstructive pulmonary disease.
9 *Indian J Chest Dis Allied Sci*. 2014; **56**: 7-12.
- 10 45 Ghanem M, Elaal EA, Mehany M, Tolba K. Home-based pulmonary rehabilitation
11 program: Effect on exercise tolerance and quality of life in chronic obstructive pulmonary
12 disease patients. *Ann Thorac Med*. 2010; **5**: 18-25.
- 13 46 Ko FWS, Cheung NK, Rainer TH, Lum C, Wong I, Hui DSC. Comprehensive care
14 programme for patients with chronic obstructive pulmonary disease: a randomised
15 controlled trial. *Thorax*. 2017; **72**: 122-8.
- 16 47 Ko FW, Dai DL, Ngai J, Tung A, Ng S, Lai K, Fong R, Lau H, Tam W, Hui DS. Effect of
17 early pulmonary rehabilitation on health care utilization and health status in patients
18 hospitalized with acute exacerbations of COPD. *Respirology*. 2011; **16**: 617-24.
- 19 48 Seymour JM, Moore L, Jolley CJ, Ward K, Creasey J, Steier JS, Yung B, Man WD, Hart
20 N, Polkey MI, Moxham J. Outpatient pulmonary rehabilitation following acute exacerbations
21 of COPD. *Thorax*. 2010; **65**: 423-8.
- 22 49 Jones SE, Green SA, Clark AL, Dickson MJ, Nolan AM, Moloney C, Kon SS, Kamal F,
23 Godden J, Howe C, Bell D, Fleming S, Haselden BM, Man WD. Pulmonary rehabilitation
24 following hospitalisation for acute exacerbation of COPD: referrals, uptake and adherence.
25 *Thorax*. 2014; **69**: 181-2.
- 26 50 Nava S. Rehabilitation of patients admitted to a respiratory intensive care unit.
27 *Archives of physical medicine and rehabilitation*. 1998; **79**: 849-54.
- 28 51 Kirsten DK, Taube C, Lehnigk B, Jörres RA, Magnussen H. Exercise training improves
29 recovery in patients with COPD after an acute exacerbation. *Respiratory medicine*. 1998; **92**:
30 1191-8.
- 31 52 Troosters T, Probst VS, Crul T, Pitta F, Gayan-Ramirez G, Decramer M, Gosselink R.
32 Resistance training prevents deterioration in quadriceps muscle function during acute
33 exacerbations of chronic obstructive pulmonary disease. *American journal of respiratory
34 and critical care medicine*. 2010; **181**: 1072-7.
- 35 53 He M, Yu S, Wang L, Lv H, Qiu Z. Efficiency and safety of pulmonary rehabilitation in
36 acute exacerbation of chronic obstructive pulmonary disease. *Medical science monitor :
37 international medical journal of experimental and clinical research*. 2015; **21**: 806-12.
- 38 54 Liao L-Y, Chen K-M, Chung W-S, Chien J-Y. Efficacy of a respiratory rehabilitation
39 exercise training package in hospitalized elderly patients with acute exacerbation of COPD: a
40 randomized control trial. *International journal of chronic obstructive pulmonary disease*.
41 2015; **10**: 1703-9.
- 42 55 Maddocks M, Gao W, Higginson IJ, Wilcock A. Neuromuscular electrical stimulation
43 for muscle weakness in adults with advanced disease. *Cochrane database of systematic
44 reviews*. 2013; **1**: CD009419.
- 45 56 Abdellaoui A, Prefaut C, Gouzi F, Couillard A, Coisy-Quivy M, Hugon G, Molinari N,
46 Lafontaine T, Jonquet O, Laoudj-Chenivresse D, Hayot M. Skeletal muscle effects of
47 electrostimulation after COPD exacerbation: a pilot study. *Eur Respir J*. 2011; **38**: 781-8.
- 48 57 Giavedoni S, Deans A, McCaughey P, Drost E, MacNee W, Rabinovich RA.
49 Neuromuscular electrical stimulation prevents muscle function deterioration in exacerbated
50 COPD: a pilot study. *Respiratory medicine*. 2012; **106**: 1429-34.
- 51
52
53
54
55
56
57
58
59
60

- 1
2
3
4 58 Vivodtzev I, Pepin JL, Vottero G, Mayer V, Porsin B, Levy P, Wuyam B. Improvement
5 in quadriceps strength and dyspnea in daily tasks after 1 month of electrical stimulation in
6 severely deconditioned and malnourished COPD. *Chest*. 2006; **129**: 1540-8.
- 7 59 Behnke M, Taube C, Kirsten D, Lehnigk B, Jorres RA, Magnussen H. Home-based
8 exercise is capable of preserving hospital-based improvements in severe chronic obstructive
9 pulmonary disease. *Respiratory medicine*. 2000; **94**: 1184-91.
- 10 60 Eaton T, Young P, Fergusson W, Moodie L, Zeng I, O'Kane F, Good N, Rhodes L, Poole
11 P, Kolbe J. Does early pulmonary rehabilitation reduce acute health-care utilization in COPD
12 patients admitted with an exacerbation? A randomized controlled study. *Respirology*. 2009;
13 **14**: 230-8.
- 14 61 Greening NJ, Williams JE, Hussain SF, Harvey-Dunstan TC, Bankart MJ, Chaplin EJ,
15 Vincent EE, Chimera R, Morgan MD, Singh SJ, Steiner MC. An early rehabilitation
16 intervention to enhance recovery during hospital admission for an exacerbation of chronic
17 respiratory disease: randomised controlled trial. *Bmj*. 2014; **349**: g4315.
- 18 62 Chalmers J. 2018. ayside Rehabilitation in Bronchiectasis Exacerbations (TRIBE) : a
19 Randomized Controlled Trial
20 (TRIBE). <https://clinicaltrials.gov/ct2/show/study/NCT02179983>. 2019.
- 21 63 Fearon KCH, Ljungqvist O, Von Meyenfeldt M, Revhaug A, Dejong CHC, Lassen K,
22 Nygren J, Hausel J, Soop M, Andersen J, Kehlet H. Enhanced recovery after surgery: A
23 consensus review of clinical care for patients undergoing colonic resection. *Clinical nutrition*.
24 2005; **24**: 466-77.
- 25 64 Martínez-Velilla N, Casas-Herrero A, Zambom-Ferraresi F, et al. Effect of exercise
26 intervention on functional decline in very elderly patients during acute hospitalization: A
27 randomized clinical trial. *JAMA Internal Medicine*. 2019; **179**: 28-36.
- 28 65 Revitt O, Sewell L, Singh S. Early versus delayed pulmonary rehabilitation: A
29 randomized controlled trial - Can we do it? *Chronic respiratory disease*. 2018; **15**: 323-6.
- 30 66 Puhan MA, Spaar A, Frey M, Turk A, Brandli O, Ritscher D, Achermann E, Kaelin R,
31 Karrer W. Early versus late pulmonary rehabilitation in chronic obstructive pulmonary
32 disease patients with acute exacerbations: a randomized trial. *Respiration; international
33 review of thoracic diseases*. 2012; **83**: 499-506.
- 34 67 Kjaergaard JL, Juhl CB, Rosenberg M, Rohde H, Riber H, Poulsen L, Wilcke T. Early
35 pulmonary rehabilitation after acute exacerbation of COPD. *European Respiratory Journal*.
36 2018; **52**: OA1618.
- 37 68 Benzo R, Vickers K, Novotny PJ, Tucker S, Hout J, Neuenfeldt P, Connett J, Lorig K,
38 McEvoy C. Health Coaching and Chronic Obstructive Pulmonary Disease Rehospitalization. A
39 Randomized Study. *American journal of respiratory and critical care medicine*. 2016; **194**:
40 672-80.
- 41 69 Chaplin E, Hewitt S, Apps L, Bankart J, Pulikottil-Jacob R, Boyce S, Morgan M,
42 Williams J, Singh S. Interactive web-based pulmonary rehabilitation programme: a
43 randomised controlled feasibility trial. *BMJ open*. 2017; **7**: e013682-e.
- 44 70 Dodd JW, Charlton RA, van den Broek MD, Jones PW. Cognitive dysfunction in
45 patients hospitalized with acute exacerbation of COPD. *Chest*. 2013; **144**: 119-27.
- 46 71 Houchen-Wolloff L, Orme M, Clinch L, Gardiner N, Singh S. S11 Feasibility of a web-
47 based self-management programme, as a 'bridge' to starting pulmonary rehabilitation, for
48 individuals hospitalised with an acute exacerbation of chronic obstructive pulmonary
49 disease (AECOPD). *Thorax*. 2018; **73**: A8-A9.
- 50
51
52
53
54
55
56
57
58
59
60

- 1
2
3 72 Cox M, O'Connor C, Biggs K, Hind D, Bortolami O, Franklin M, Collins B, Walters S,
4 Wailoo A, Channell J, Albert P, Freeman U, Bourke S, Steiner M, Miles J, O'Brien T,
5 McWilliams D, Schofield T, O'Reilly J, Hughes R. The feasibility of early pulmonary
6 rehabilitation and activity after COPD exacerbations: external pilot randomised controlled
7 trial, qualitative case study and exploratory economic evaluation. *Health Technol Assess.*
8 2018; **22**: 1-204.
9
10 73 Polkey MI, Praestgaard J, Berwick A, Franssen FME, Singh D, Steiner MC, Casaburi R,
11 Tillmann H-C, Lach-Trifilieff E, Roubenoff R, Rooks DS. Activin Type II Receptor Blockade for
12 Treatment of Muscle Depletion in COPD: A Randomized Trial. *American journal of*
13 *respiratory and critical care medicine.* **0**: null.
14
15 74 Casaburi R, Bhasin S, Cosentino L, Porszasz J, Somfay A, Lewis MI, Fournier M, Storer
16 TW. Effects of testosterone and resistance training in men with chronic obstructive
17 pulmonary disease. *American journal of respiratory and critical care medicine.* 2004; **170**:
18 870-8.
19
20 75 Pison CM, Cano NJ, Cherion C, Caron F, Court-Fortune I, Antonini MT, Gonzalez-
21 Bermejo J, Meziane L, Molano LC, Janssens JP, Costes F, Wuyam B, Similowski T, Melloni B,
22 Hayot M, Augustin J, Tardif C, Lejeune H, Roth H, Pichard C, Investigators I. Multimodal
23 nutritional rehabilitation improves clinical outcomes of malnourished patients with chronic
24 respiratory failure: a randomised controlled trial. *Thorax.* 2011; **66**: 953-60.
25
26 76 Puhan M, Gimeno-Santos E, Cates CJ, Troosters T. Pulmonary rehabilitation following
27 exacerbations of chronic obstructive pulmonary disease. 2016; **12**: CD005305.
28
29 77 Borges RC, Carvalho CR. Impact of resistance training in chronic obstructive
30 pulmonary disease patients during periods of acute exacerbation. *Archives of physical*
31 *medicine and rehabilitation.* 2014; **95**: 1638-45.
32
33 78 Domingo-Salvany A, Lamarca R, Ferrer M, Garcia-Aymerich J, Alonso J, Felez M,
34 Khalaf A, Marrades RM, Monso E, Serra-Batlles J, Anto JM. Health-related quality of life and
35 mortality in male patients with chronic obstructive pulmonary disease. *American journal of*
36 *respiratory and critical care medicine.* 2002; **166**: 680-5.
37
38
39
40
41
42
43
44
45
46
47
48
49
50
51
52
53
54
55
56
57
58
59
60

1
2
3 **List of Figures**
4
5
6
7

8 **Figure 1:** Multi-factorial insults and consequences on individuals following admission to hospital with
9 an exacerbation of COPD. Blue markings demonstrate a selection of non-pulmonary organs affected
10 (heart, skeletal muscles, bone/spine, pancreas, brain)
11
12
13
14

15
16 **Figure 2:** Schematic of time-course of hospital associated disability, including symptoms, insults and
17 function
18
19
20
21
22
23
24
25
26
27
28
29
30
31
32
33
34
35
36
37
38
39
40
41
42
43
44
45
46
47
48
49
50
51
52
53
54
55
56
57
58
59
60

1
2
3
4
5
6
7
8
9
10
11
12
13
14
15
16
17
18
19
20
21
22
23
24
25
26
27
28
29
30
31
32
33
34
35
36
37
38
39
40
41
42
43
44
45
46

Table 1: Randomised controlled trials of peri-exacerbation pulmonary rehabilitation

Study	n=	Follow up	Primary Outcome	Outcome of Results		
				Healthcare Utilisation	Exercise capacity/ Strength	Symptoms
In-patient AECOPD						
Nava⁵⁰	80	Hospital discharge	Exercise capacity	Length of hospital stay: No difference	6MWT: Improved in intervention but not control group. Groups not compared.	Dyspnoea: Improved in both groups. Groups not compared
Kirsten⁵¹	29	10 days	Exercise capacity	Not reported	6MWT & minute ventilation: Greater improvement in the intervention group	Dyspnoea: Greater improvement in the intervention group
Troosters⁵²	36	1 month	Quadriceps force	Hospital readmission: No difference at 6 months	Quadriceps strength: Greater improvement in intervention group. 6MWT: No difference between groups	Not reported
Abdellaoui⁵⁶	15	6 weeks	Quadriceps strength	Not reported	Quadriceps force & 6MWT: Greater improvement in the intervention group	Not reported
Giavedoni⁵⁷	11	14 days	Quadriceps strength	Length of hospital stay: No difference	Quadriceps force: Greater improvement in the intervention group	Not reported
Borges⁷⁷	29	30 days	Quadriceps strength	Not reported	6MWT & strength: Greater improvement in the intervention group	SGRQ: No difference between groups
He⁵³	97	Hospital discharge	Exercise capacity	Not reported	6MWT: Improved in intervention but not control group. Groups not compared.	CRQ & ADL: Improved in intervention but not control group. Groups not compared.
Liao⁵⁴	61	4 days	Dyspnoea	Not reported	6MWT: Greater improvement in the intervention group	Dyspnoea: Greater improvement in the intervention group
In-patient progressing to Out-Patient AECOPD						
Behnke⁵⁹	46	6 months	Exercise capacity	Not reported	6MWT: Greater improvement in the intervention group	CRQ: Greater improvement in the intervention group
Eaton⁶⁰	97	3 months	Re-admission	Hospital readmission: No difference	6MWT: No difference between groups	CRQ: No difference between groups
Greening⁶¹	389	12 months	Re-admission	Hospital readmission: No difference	ISWT & Quadriceps strength: No difference between groups	SGRQ: No difference between groups

Cox⁷⁸	58	90 days	Exercise capacity	Hospital readmission: No difference	6MWT: No difference between groups	CAT: No difference between groups
Out-patient AECOPD						
Man⁴²	42	8 weeks	Exercise capacity	Hospital readmission: No difference between groups	ISWT: Greater improvement in the intervention group	CRQ & SGRQ: Greater improvement in the intervention group
Murphy⁴³	26	6 weeks	Exercise capacity	Exacerbation: No difference at 3 or 6 months between groups	ISWT: Improved in intervention but not control group. Groups not compared.	SGRQ: Improved in both groups. Groups not compared
Vivodtzev⁵⁸	17	4 weeks	Quadriceps strength	Not reported	Quadriceps force & 6MWT: Greater improvement in the intervention group	Dyspnoea: Greater improvement in the intervention group
Seymour⁴⁸	60	3 months	Re-admission	Hospital readmission: Fewer readmissions in intervention group	ISWT & quadriceps force: Greater improvement in the intervention group	SGRQ: Greater improvement in the intervention group
Ghanem⁴⁵	39	2 months	Exercise capacity	Not reported	6MWT: Greater improvement in the intervention group	CRQ: Greater improvement in the intervention group
Puhan⁶⁶	36	18 months	Exacerbations	Exacerbation: No difference	Not reported	CRQ: No difference between groups
Ko (2011)⁴⁷	60	12 months	Re-admission	Hospital Readmission: No difference between groups	6MWT & VO _{2max} : No difference between groups	SGRQ: Greater improvement in the intervention group
Deepak⁴⁴	60	3 months	Exercise capacity	Not reported	6MWT: Greater improvement in the intervention group	SGRQ: Greater improvement in the intervention group
Ko (2017)⁴⁶	180	12 months	Re-admission	Hospital readmission: Fewer readmissions in intervention group	6MWT: No difference between groups	SGRQ: Greater improvement in the intervention group

6MWT=six-minute walk test; VO_{2max}=oxygen consumption; ISWT=incremental shuttle walk test; CRQ=chronic respiratory questionnaire; SGRQ= St. Georges respiratory Questionnaire; ADL scale=Activity of Daily Living

1
2
3
4
5
6
7
8
9
10
11
12
13
14
15
16
17
18
19
20
21
22
23
24
25
26
27
28
29
30
31
32
33
34
35
36
37
38
39
40
41
42
43
44
45
46
47
48
49
50
51
52
53
54
55
56
57
58
59
60

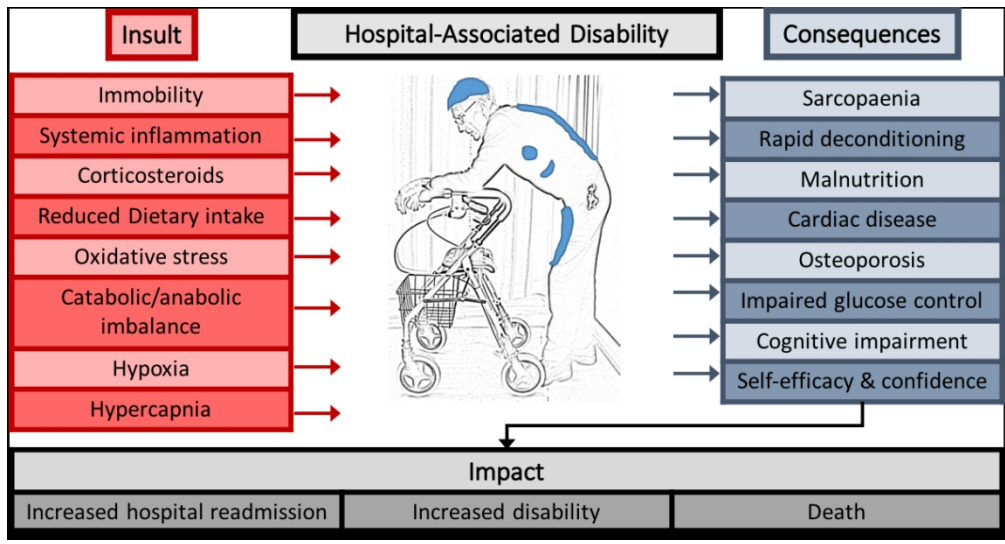
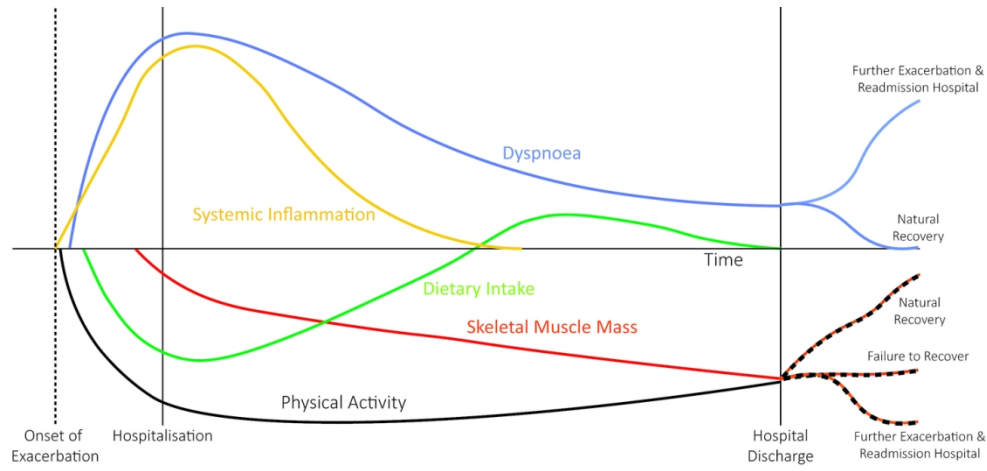


Figure 1: Multi-factorial insults and consequences on individuals following admission to hospital with an exacerbation of COPD. Blue markings demonstrate a selection of non-pulmonary organs affected (heart, skeletal muscles, bone/spine, pancreas, brain)

204x108mm (150 x 150 DPI)



Schematic of time-course of hospital associated disability, including symptoms, insults and function

248x114mm (150 x 150 DPI)