

Health-Related Quality of Life and Health State Utility Values in Moderate to Very Severe COPD: A Systematic Literature Review

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Background

- COPD is a leading cause of morbidity and mortality worldwide, and is associated with substantial economic and healthcare burden¹
- COPD exacerbations are associated with reduced health status and long-lasting negative effects, including a significant impact on patients' health-related quality of life^{2,3}
- Burden of disease not only affects patients with COPD but also their caregivers^{4,5}

Aims

This SLR aimed to identify health state utility values and disutilities in patients with moderate to very severe COPD and their caregivers

Study design

GSK ID: 223395

SLR approach

- This SLR was conducted in accordance with PRISMA standards and the Centre for Reviews and Dissemination guidance^{6,7}
- Studies were included based on predefined eligibility criteria in terms of the PICOS framework

Population

- Adult patients with a diagnosis of moderate to very severe COPD
- Caregivers to adults with moderate to very severe COPD

Study designs

- Non-RCTs except if the RCT was non-pharmacological*
- PK/PD studies, economic evaluations, reviews and meta-analyses, comment/letter to the editor were excluded

Intervention/comparators

- Not restricted (non-pharmacological intervention or none)

Sources (last searched July 26, 2024)

- MEDLINE, Embase, and Cochrane databases (searched from 2018–current)
- Conference proceedings from 2021–2024 (ISPOR, CHEST, ATS, ERS)
- Additional sources (including EQ-5D website, reference lists of included studies checked for further studies, HERC database of mapping studies)

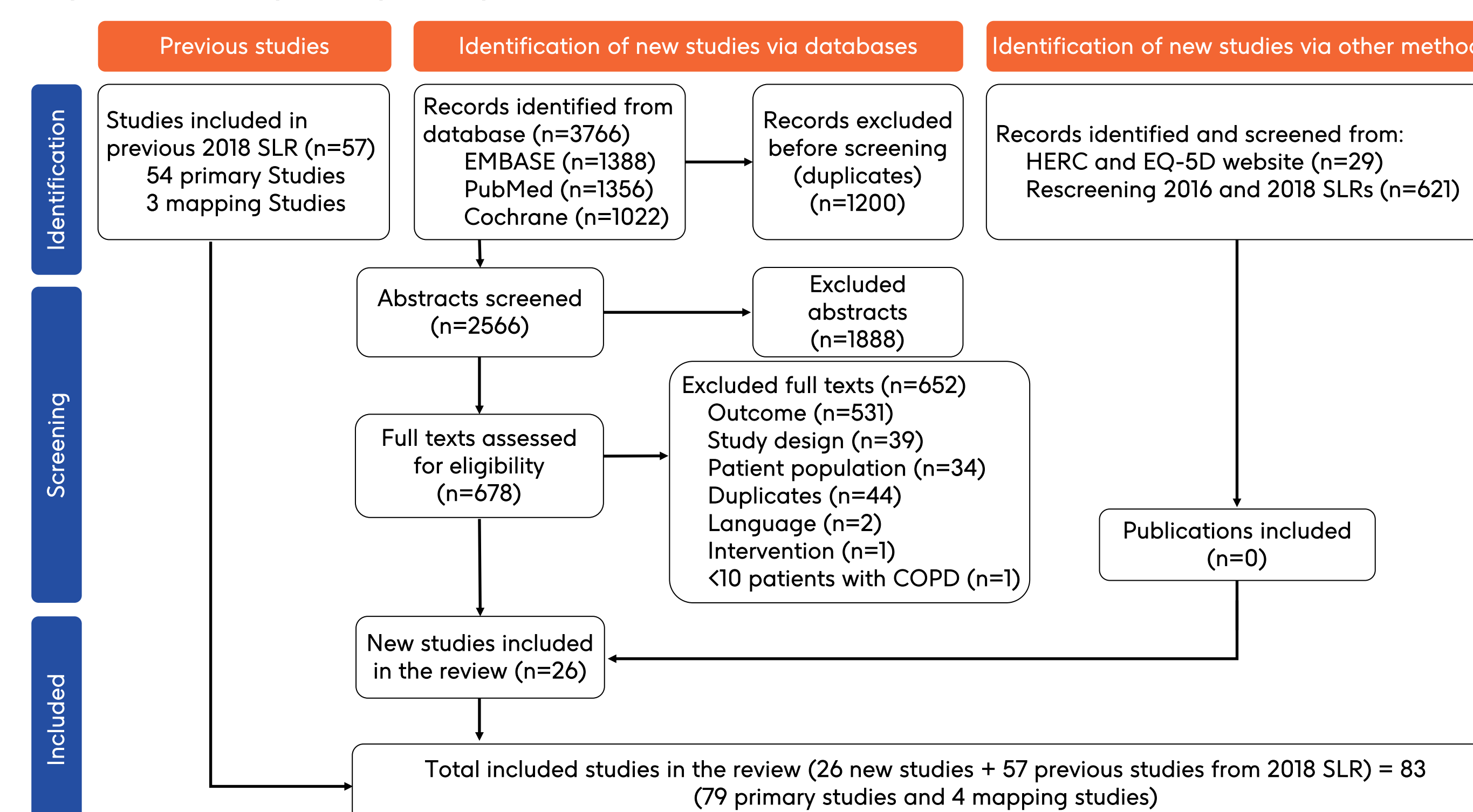
Outcomes

- Utility values were described separately for health states during COPD exacerbations and for stable COPD
- For stable COPD, utility values were further stratified by GOLD severity and by history of exacerbations when the study reported them
- Health-state utility is the value given to a specific health-related quality of life measure, where a value of 1 represents full health and 0 represents dead, with negative values indicating states worse than death⁸

*RCTs assessing non-pharmacological interventions such as home care or pulmonary rehabilitation have been included in this SLR

Results

Figure 1: Seventy-nine primary studies were identified



- Of the 79 primary studies identified, 64 reported utility values for patients with stable moderate to very severe COPD, with 36 of these studies utilizing EQ-5D (utilities ranging from 0.25 to 0.86)

Table 1: Across 6 studies that reported utility values for patients with stable COPD stratified by the number of exacerbations in the previous year, lower utility values were consistently associated with a greater frequency of prior exacerbations

Study/country	Number exacerbations in previous year	Utility, mean (SD) or median (95% CI)	Sample size
Rutten-van Molken, 2009/ Netherlands	1 non-serious exacerbation	0.010 (utility decrement)	NR
	2 non-serious exacerbation	0.021 (utility decrement)	NR
	1 serious exacerbation	0.042 (utility decrement)	NR
Altawalbeh, 2022/ Jordan	1 serious and 1 non-serious exacerbation	0.088 (utility decrement)	NR
	All patients regardless of exacerbation	0.68 (0.36)	203
Esquinas, 2020/ Spain	≥1 hospital admission for COPD	0.48 (0.4)	41
	≥1 ER visit for COPD	0.53 (0.4)	89
	COPD hospital admission: 0	0.73 (0.2)	NR
	COPD hospital admission: 1	0.65 (0.3)	NR
Miravittles, 2014/ Spain	COPD hospital admission: 2	0.54 (0.3)	NR
	COPD hospital admission: >2	0.45 (0.3)	NR
	1–2 exacerbations	0.78	NR
	<2 exacerbations	0.84	NR
Punekar, 2007/ France, Germany, Italy, Spain, UK, US	≥2 exacerbations	0.74	NR
	No exacerbations	0.78 (0.75–0.80)*	NR
	1–2 exacerbations	0.75 (0.72–0.77)*	NR
Solem, 2013/ US	1–2 exacerbations	0.74 (0.72–0.77)*	NR
	≥3 exacerbations	0.73 (0.71–0.76) [†]	NR
	≥3 exacerbations	0.61 (0.59–0.64)*	NR
Solem, 2013/ US	1 moderate/severe exacerbation	0.727 (0.175)	82
	2 moderate/severe exacerbations	0.684 (0.204)	84
	3+ moderate/severe exacerbations	0.638 (0.212)	148

*Patients seeking care from primary care physicians; [†]patients seeking care from respiratory specialists



This SLR highlights the significant impact of disease severity and exacerbations on health-related quality of life in patients with COPD

Digital poster



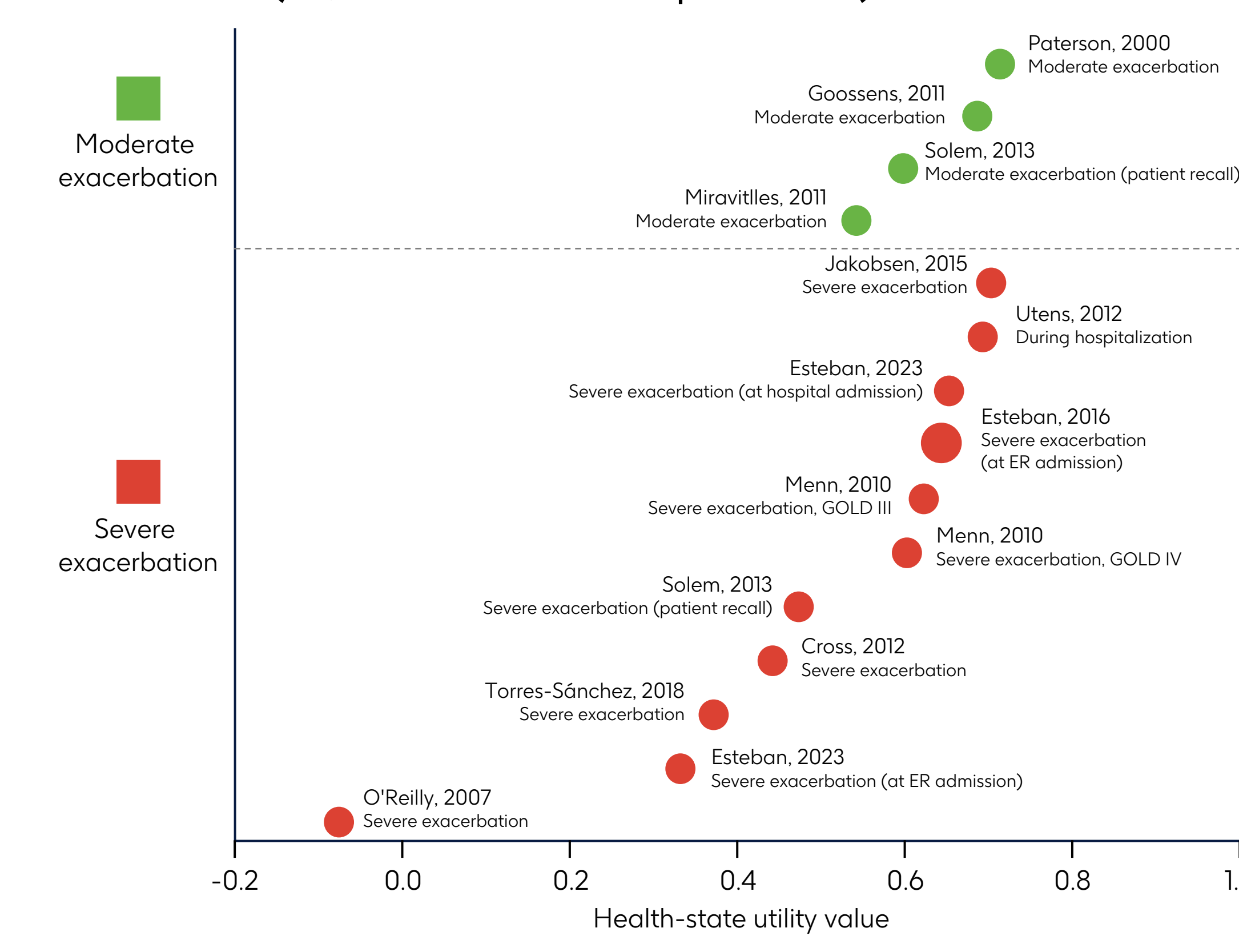
SCAN ME

Figure 2: Seventeen reported utilities stratified by COPD severity defined using GOLD grades. Within all studies reporting severity-specific utilities, patients with more severe COPD consistently reported lower utility values



Each circle represents a study-specific estimate. Circle area is proportional to sample size. NR sample sizes are displayed using minimum marker size

Figure 3: Twelve studies reported utilities during an exacerbation. Mean utilities for exacerbations ranged from –0.120 (assessed at visit to treat exacerbation; patients admitted to hospital during an exacerbation) to 0.71 (assessed at the first visit to treat exacerbation; patients with acute exacerbation at a general practice), with worsening utilities following severe exacerbation (i.e., a COPD-related hospitalization)



Each circle represents a study-specific estimate. Circle area is proportional to sample size. NR sample sizes are displayed using minimum marker size

Table 2: Caregiver utilities were reported in 3 different studies, ranging from 0.76 to 0.88

Study/country	Participant description	No. of participants with response	Utility, mean (SD)
EQ-5D			
Majellano, 2021/ Australia	Caregivers, ≥18 years, providing unpaid care (denoting informal caregiver) to a relative/friend who with severe asthma or COPD and had been in the caring role for ≥6 months	48	0.85 (0.17)
Nakken, 2016/ Netherlands	Closest relatives who spent most time with the patient and/or provided most of the care, assistance and support	193	0.88 (0.15)
Rehman, 2021/ Malaysia	Caregivers of patients with COPD (unpaid voluntary family members and friends) assist the patients to perform daily activities	127	0.76 (0.36)
AQoL			
Nakken 2016/ Netherlands	Closest relatives who spent most time with the patient and/or provided most of the care, assistance and support	193	0.78 (0.14)

Conclusions

This SLR highlights the significant impact of COPD severity and exacerbations on health-related quality of life. The findings provide valuable health state utility values for economic modeling and emphasize the need for tailored interventions to improve health-related quality of life in patients with COPD and their caregivers

Abbreviations

AQoL, The Assessment of Quality of Life; ATS, American Thoracic Society; CHEST, American College of Chest Physicians; CI, confidence interval; COPD, chronic obstructive pulmonary disease; EQ-5D, EuroQoL Five Dimension; ER, emergency room; ERS, European Respiratory Society; GOLD, Global Initiative for Chronic Obstructive Lung Disease; HERC, Health Economics Research Centre; ISPOR, International Society for Pharmacoeconomics and Outcomes Research; NR, not reported; PD, pharmacodynamics; PICOS, Population, Intervention and Comparators, Outcomes and Study design; PK, pharmacokinetics; PRISMA, Preferred Reporting Items for Systematic reviews and Meta-Analyses; RCT, randomized controlled trial; SD, standard deviation; SLR, systematic literature review; US, United States

References

- Global Initiative for Chronic Obstructive Lung Disease. Global Strategy for Prevention, Diagnosis and Management of COPD: 2026 Report; 2026. Available from: <https://goldcopd.org/2026-gold-report-and-pocket-guide/> [Accessed April 23, 2026]
- Müllerová H, et al. *Int J Chron Obstruct Pulmon Dis*. 2019;14:741–55
- Jones PW, et al. *Respir Med*. 2011;105:57–66
- Miravittles M, et al. *Int J Chron Obstruct Pulmon Dis*. 2015;10:347–56
- Gholami S, et al. *COPD*. 2025;22:2481260
- CRD. Systematic reviews: CRD's guidance for undertaking reviews in health care. Available from: https://www.york.ac.uk/media/crd/Systematic_Reviews.pdf [Accessed April 23, 2026]
- Moher D, et al. Preferred reporting items for systematic review and meta-analysis protocols (PRISMA-P) 2015 statement. *Syst Rev*. 2015;4:1
- Wolowacz SE, et al. *Value Health*. 2016;19:704–19

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Disclosures

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