

The burden of persistent cough in idiopathic pulmonary fibrosis (IPF) and other interstitial lung diseases (ILDs): a systematic evidence synthesis

Nick Pooley,¹ Michael Baldwin,² Rhiannon Green,¹ Maureen PMH Rutten-van Mölken,³ Nina Patel,⁴ Marlies S Wijzenbeek⁵

¹Market Access, Maverex Limited, Manchester, United Kingdom; ²Value and Patient Access, Boehringer Ingelheim International GmbH, Ingelheim am Rhein, Germany; ³Erasmus School of Health Policy and Management, Erasmus University Rotterdam, Rotterdam, The Netherlands; ⁴Inflammation Medicine, Boehringer Ingelheim Pharmaceuticals Inc, Ridgefield, CT, USA; ⁵Respiratory Medicine, Erasmus Medical Center, Rotterdam, The Netherlands

INTRODUCTION

- Cough remains a prevalent and persistent symptom in patients with IPF and other ILDs¹⁻³
- To inform future research, treatment and care models, it is vital to understand the burden of persistent cough in these patient groups

AIM

- To provide the first systematic synthesis of evidence on the burden of cough in IPF and other ILDs

METHODS



Protocol pre-registered on PROSPERO as CRD42022369379

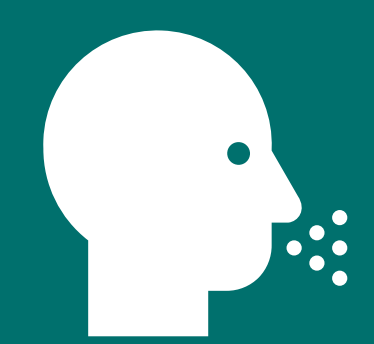


Literature search performed for English-language articles published between Jan 2010 and Aug 2022, using databases including:

- Embase
- MEDLINE
- Cochrane



The literature search included observational and interventional studies reporting cough-related measures in IPF and other ILDs



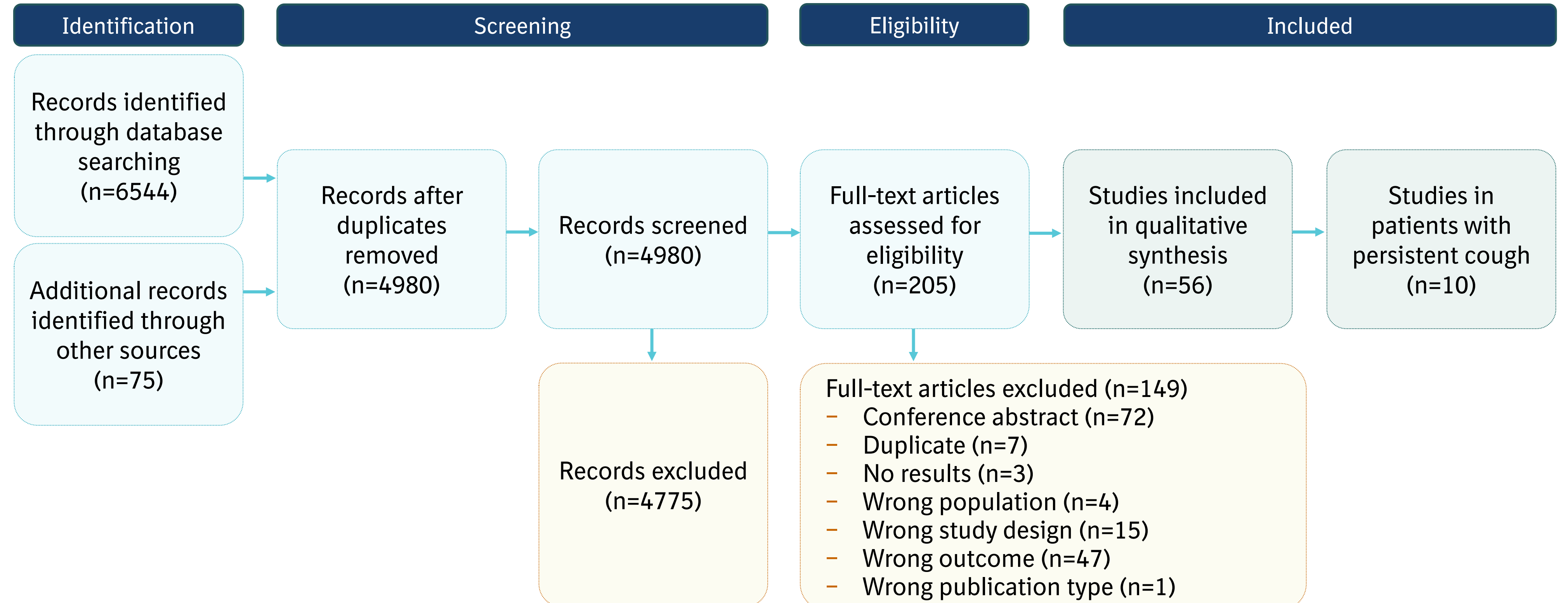
This narrative synthesis focuses on studies in patients with persistent cough investigating associations between cough and impact/health-related QoL measures

CONCLUSIONS

- Our study highlights the heterogeneity in assessing cough and its impact in IPF and other ILDs
- The findings confirm the association of cough with health-related QoL in IPF, with indications of a similar association in other ILDs
- Our synthesis underscores the need for standardised assessment along with dedicated studies, particularly in non-IPF ILDs and on the economic burden of cough

RESULTS

Literature search



Study descriptions

- Ten studies were identified in patients with IPF/ILD and persistent cough

Definitions of persistent cough

Self-reported chronic cough

Stable cough for >4 weeks

Cough for >8 weeks

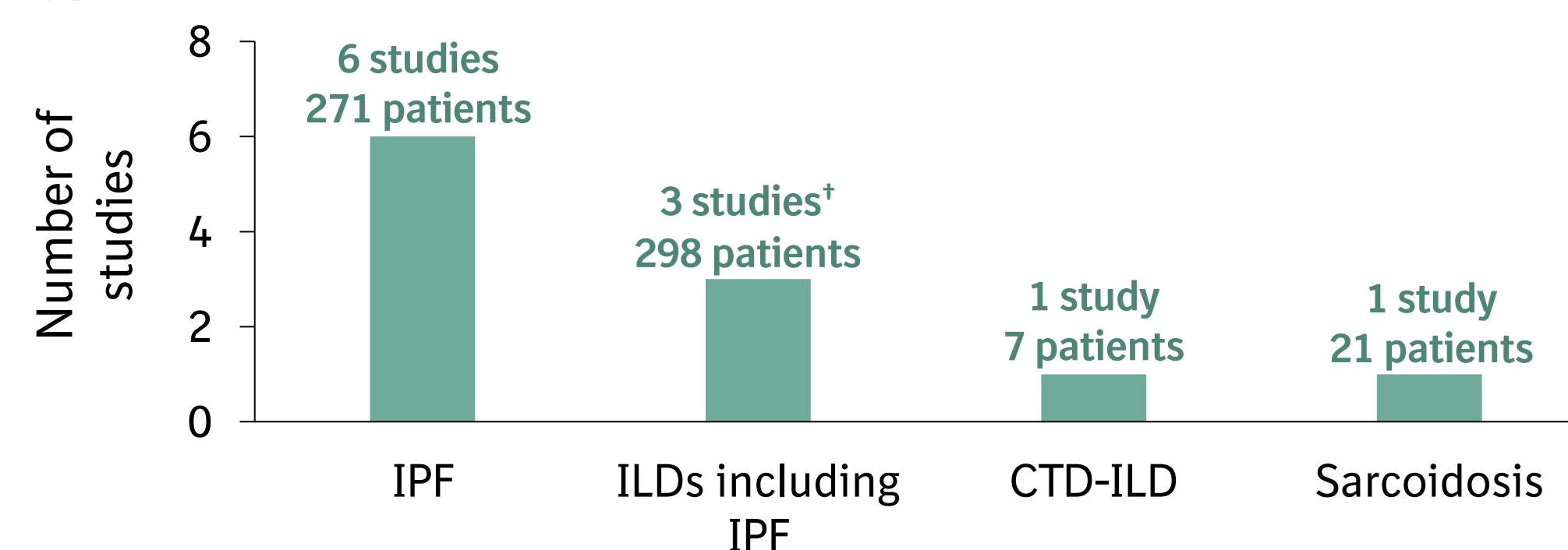
Some studies required additional criteria, such as:

Refractory cough

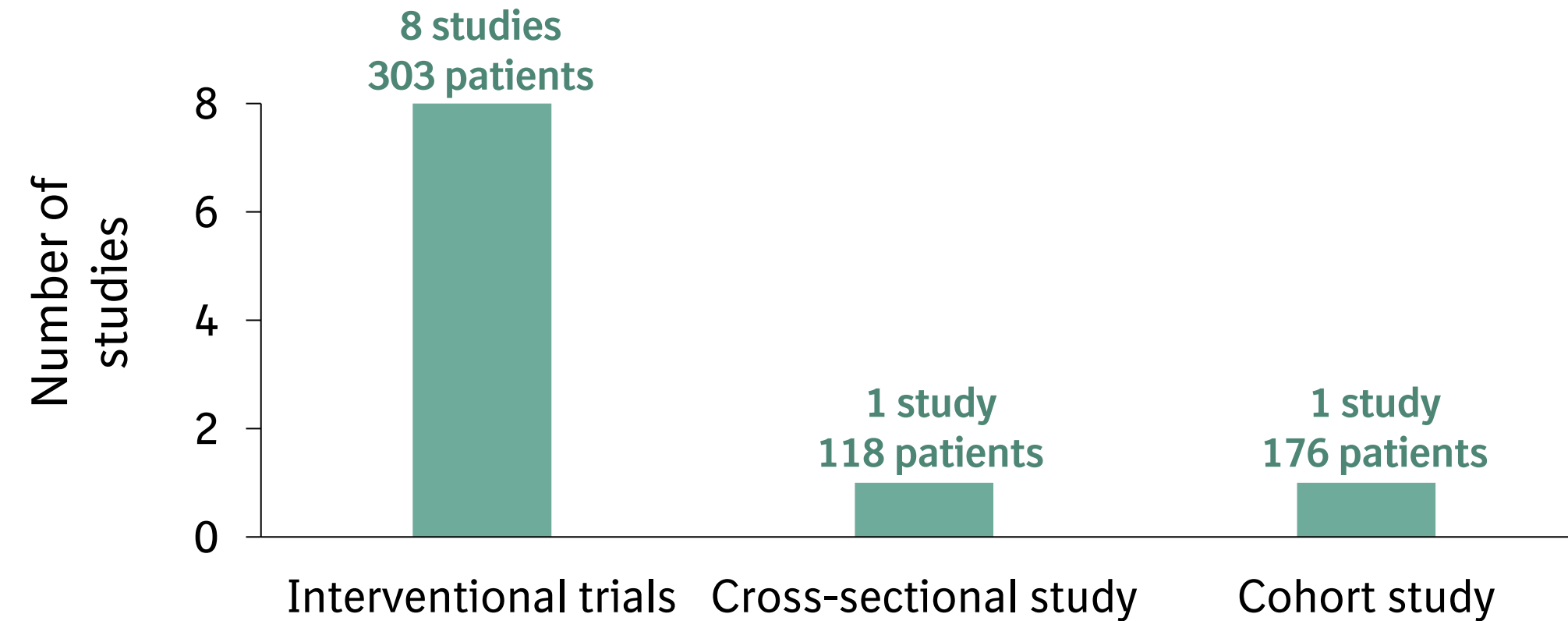
24-hour cough count of >10/15 coughs/hour

Cough severity VAS >40 mm

Types of ILD*



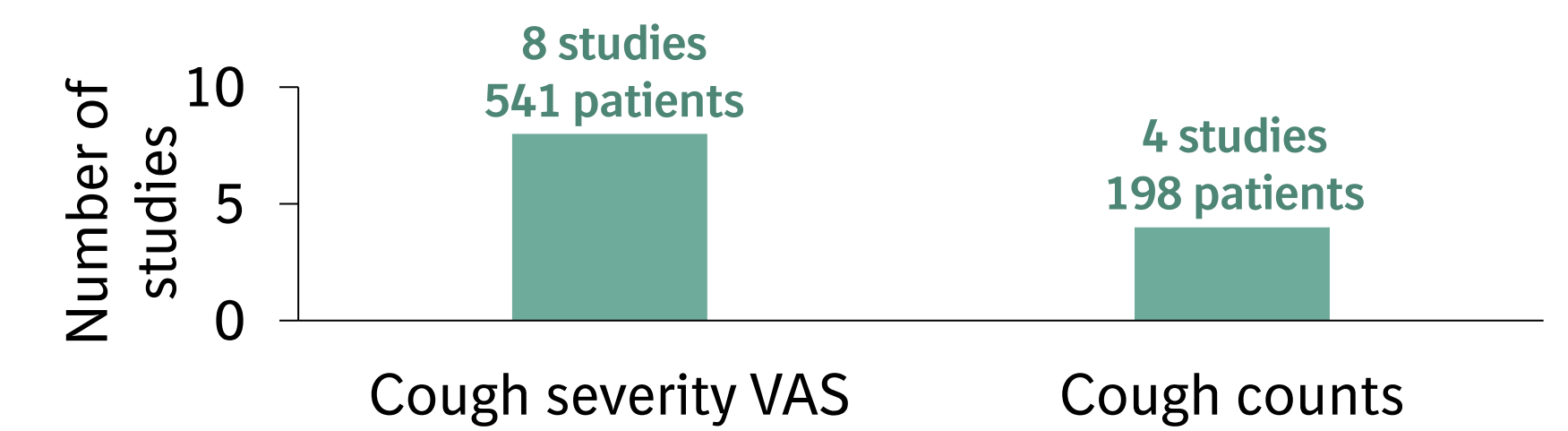
Types of studies*



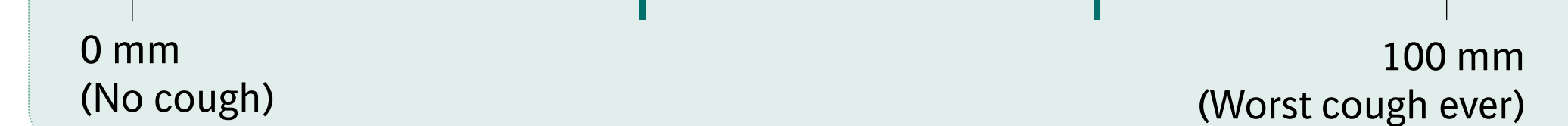
*Patient numbers shown in bar charts only include patients with IPF or other ILDs. Other indications were excluded. *Two patients with IIPs including IPF and two patients with IIPs with unclassifiable idiopathic interstitial pneumonia were included. *The SGRQ has been validated in IPF and SSC-ILD. The CQLQ has been validated in IPF, but not other ILDs. Neither cough severity VAS nor the LCQ have been validated in IPF or other ILDs.

Cough-related measures and PROs*

Most frequently used cough measures*

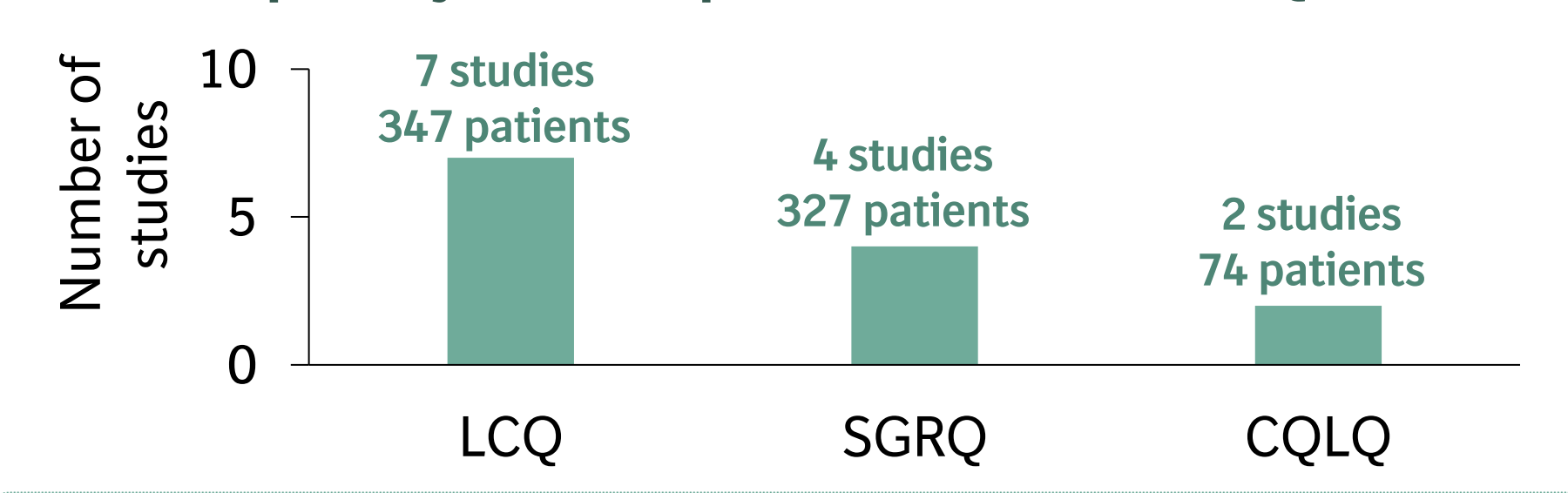


Cough severity VAS mean range: 38.8–73.4



24-hour cough count mean range: 10–40 coughs/hour

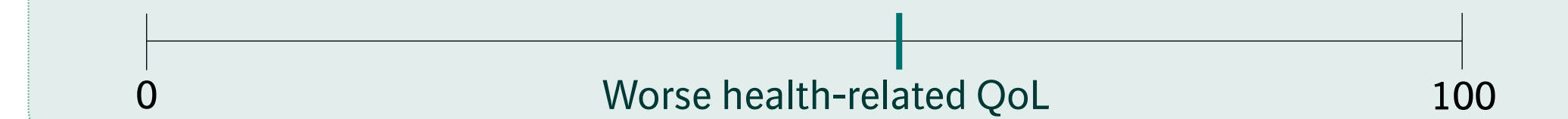
Most frequently used impact/health-related QoL measures*



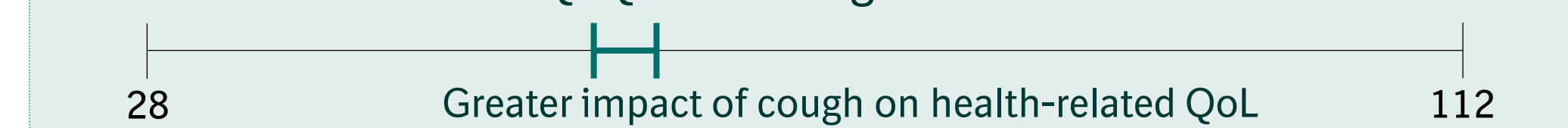
LCQ mean range: 11.0–15.3



SGRQ mean range: 57.2–57.4



CQLQ mean range: 56.5–60.5



Association between cough and impact/health-related QoL measures

- Four studies assessed concurrent/baseline associations between cough and impact/HRQoL measures in patients with persistent cough, with all of these reporting at least one significant association
- Additionally, in a cross-sectional study in ILD, 31% of patients ranked cough as the worst symptom³
- None of the studies examined the economic burden of cough

Author, year	N	Study design	Burden of cough
Birring 2017 ⁵	IPF: 24 CIC: 27	Crossover trial	Significant correlation between daytime cough frequency and VAS cough severity (r=0.683; p=0.0003) and LCQ (r=-0.682; p=0.00002)
Lechtzin 2013 ⁶	IPF: 23	Crossover trial	CQLQ significantly correlated with cough VAS (r=0.63; p<0.01), SGRQ total score (r=0.79; p<0.01) and all subscales of the SGRQ (r range=0.72–0.81; p<0.05)
Guler 2021 ⁷	IPF: 20	Crossover trial	LCQ correlated with VAS cough severity (r=-0.42; p<0.001) and SGRQ (r=-0.70; p<0.001); SGRQ correlated with VAS (r=0.42; p<0.001)
Cheng 2017 ⁸	IPF: 77 cHP: 32 SSc-ILD: 67	Observational study	Cough severity VAS correlated with SGRQ in IPF (r=0.33; p=0.19) and SSc-ILD (r=0.51; p=0.02) but not in cHP (r=-0.1; p=0.62). Cough severity VAS remained an independent predictor of SGRQ after adjustment for age, sex, ILD severity and dyspnoea

INTERACTIVE

Scan QR code or visit URL for a device-friendly version of this poster



<https://bit.ly/48o5gHt>

Scan QR code or visit URL for a webpage featuring all BI-supported publications at ISPOR Europe 2023



<https://bit.ly/3Lztsdn>

REFERENCES

- Birring SS, et al. Chest. 2018; 154:904–917.
- Mann J, et al. Front Rehabil Sci. 2021; 2:751–798.
- Lan NSH, et al. Intern Med J. 2021; 51:923–929.
- Sato R, et al. Intern Med J. 2021; 60:3701–3707.
- Birring SS, et al. Lancet Respir Med. 2017; 5:806–815.
- Lechtzin N, et al. Chest. 2013; 143:1745–1749.
- Guler SA, et al. Ann Am Thorac Soc. 2021; 18:2018–2026.
- Cheng JZ, et al. Respiriology. 2017; 22:1592–1597.

ABBREVIATIONS

cHP, chronic hypersensitivity pneumonitis; CIC, chronic idiopathic cough; CQLQ, Cough Quality-of-Life Questionnaire; CTD-ILD, connective tissue disease-associated ILD; IIP, idiopathic interstitial pneumonia; ILD, interstitial lung disease; IPF, idiopathic pulmonary fibrosis; LCQ, Leicester Cough Questionnaire; PRO, patient-reported outcome; QoL, quality of life; SGRQ, St. George's Respiratory Questionnaire; SSc-ILD, systemic sclerosis-associated ILD; VAS, visual analogue scale.

ACKNOWLEDGEMENTS & DISCLOSURES

This study was supported and funded by Boehringer Ingelheim International GmbH (BI). The authors meet criteria for authorship as recommended by the International Committee of Medical Journal Editors (ICMJE). The authors did not receive payment related to the development of the poster. Eleni Tzouramani, MSc, of Nucleus Global, provided writing, editorial support and formatting assistance, which was contracted and funded by BI. BI was given the opportunity to review the poster for medical and scientific accuracy as well as intellectual property considerations. RG and NP report personal fees from BI outside the submitted work. MB and NP are employees of BI. MRM reports consulting fees from BI outside the submitted work. MSW reports grants or contracts from The Netherlands Organisation for Health Research and Development, The Dutch Lung Foundation, The Dutch Pulmonary Fibrosis organisation, Sarcoidosis.nl, BI, Hoffman-La Roche and AstraZeneca-Dalich; consulting fees from Bristol Myers Squibb, BI, Galapagos, Galecto, Hoffman-La Roche, Horizon Therapeutics, Kinevant Sciences, Molecule, NeReTherapeutics, Novartis, PureTech Health, Thyron, Trevi and Vicore; payments or honoraria from BI, CSL Behring, Hoffman-La Roche and Novartis; support for attending meetings/travel from BI, Hoffman-La Roche and Galapagos; and participation on a Data Safety Monitoring or advisory board for Savara and Galapagos, outside the submitted work. MSW also reports that she is Chair of the Idiopathic Interstitial Pneumonia group of the European Respiratory Society, a member of the board of the Netherlands Respiratory Society, a member of the scientific advisory board of the European Idiopathic Pulmonary Fibrosis and Related Disorders Federation, Chair of the educational committee of the European Reference Network for Rare Lung Diseases, and part of an advisory board for the Dutch Lung Fibrosis and Sarcoidosis patient associations.