

Andrew P Dickens¹, Christine Jenkins², Alexander Evans¹, Kerry Hancock³, Anita Sharma⁴, Belinda Cochrane^{5,6}, Paul Leong^{7,8}, Brian Ko⁹, Florian Heraud¹, Porsche Le Cheng¹, Alexander Roussos¹, Sinthia Bosnic-Anticevich¹⁰⁻¹², Fabio Botini¹, Victoria Carter¹, Angelina Catanzariti¹³, Clare Ghisla¹³, Thao Le¹⁴, Chantal Le Lievre¹, Ruth Murray¹, Kanchanamla Ranasinghe^{15,16}, Deb Stewart⁴, Marije van Melle^{17,18}, Rebecca Vella¹, Russell Wiseman¹⁹, and David Price^{1,17,20}

¹Optimum Patient Care Australia, Australia, ²Professor Respiratory Medicine, UNSW, Sydney, Head Respiratory Group, George Institute, Australia, ³Chandlers Hill Surgery, Happy Valley SA 5159, Australia, ⁴Platinum Medical Centre, Chermside QLD 4032, Australia, ⁵Senior Staff Specialist, Department of Respiratory and Sleep Medicine, Campbelltown Hospital (SWSLHD), Australia, ⁶A/Prof, School of Medicine, Western Sydney University, NSW Australia, ⁷Monash Health, Clayton, Victoria, Australia, ⁸School of Clinical Sciences at Monash Health, Monash University, Clayton, Victoria, Australia, ⁹Monash Heart, Monash Cardiovascular Research Centre and Monash University, Monash Health, Clayton, Victoria, Australia, ¹⁰Macquarie Medical School, Faculty of Medicine, Health and Human Sciences, Macquarie University, NSW, Australia, ¹¹Woolcock Institute of Medical Research, Glebe, NSW Australia, ¹²Sydney Local Health District, Camperdown NSW, Australia, ¹³AstraZeneca Pty Ltd, Medical Affairs Biopharmaceuticals Unit, ¹⁴Director, Medical Education, and Events Management Pte Ltd Singapore, ¹⁵School of Medicine, Griffith University, Gold Coast, Australia, ¹⁶Cannon Hill Family Doctors, Cannon Hill, QLD Australia, ¹⁷Observational and Pragmatic Research Institute, Singapore, ¹⁸Connecting Medical Dots BV, Utrecht, the Netherlands, ¹⁹Suncoast Medical Centre, Coolumb Beach, QLD Australia, ²⁰Centre of Academic Primary Care, Division of Applied Health Sciences, University of Aberdeen, Aberdeen United Kingdom

Introduction

- Delayed COPD diagnosis and unrecognised exacerbations before diagnosis are associated with future exacerbations.
- UK and US studies^{1,2} conducted within the COllaboration N on QUality improvement initiative for achieving Excellence in STandards of COPD care (CONQUEST) program³ identified substantial opportunities for earlier recognition of COPD in those with potential high-risk COPD (≥2 exacerbation-like events in the previous 12 months).
- It is unknown if similar opportunities exist within other healthcare systems, such as in Australia.

Methods

- Optimum Patient Care Research Database Australia (OPCRDA) is a primary care database of electronic health record (EHR) data containing 900,000 ever-active patients
- We identified patients with potential COPD (ever smokers aged ≥40) at high risk of future exacerbations (≥2 antibiotic and/or steroid prescriptions in the prior 12 months).
- EHR codes and free text data were analysed to examine COPD inhaled therapy, smoking cessation support, and formal COPD reviews.
- Cross-sectional analyses were conducted on annual patient cohorts between 2015 and 2019 to exclude confounding by COVID-19 in later years.

Aims and Objectives

To compare identification and assessment of patients with potential high-risk COPD in Australia, to the national and international guidelines, and to the CONQUEST quality standards⁴ as follows:

- 1 Identification of patients with high-risk COPD
- 2 Assessment of disease and quantification of future risk
- 3 Pharmacological and non-pharmacological intervention
- 4 Appropriate follow-up

Results

- 6.2% (1045/16816) to 8.3% (834/9998) of patients with no COPD diagnosis were deemed high-risk in 2015-2019 (Figure 1).
- In the annual period when patients first became high-risk, <5% of patients with potential high-risk COPD had recorded lung function, <1% had a respiratory referral, and <50% had an updated smoking status recorded.
- 11-13% of this population were prescribed maintenance inhaled COPD therapy in each study year without a recorded respiratory diagnosis (Figure 2).
- Around 1% of undiagnosed high-risk patients received a COPD diagnosis in each study year (Figure 3).

Figure 1: Number and proportion of undiagnosed patients classified as high-risk in each study year

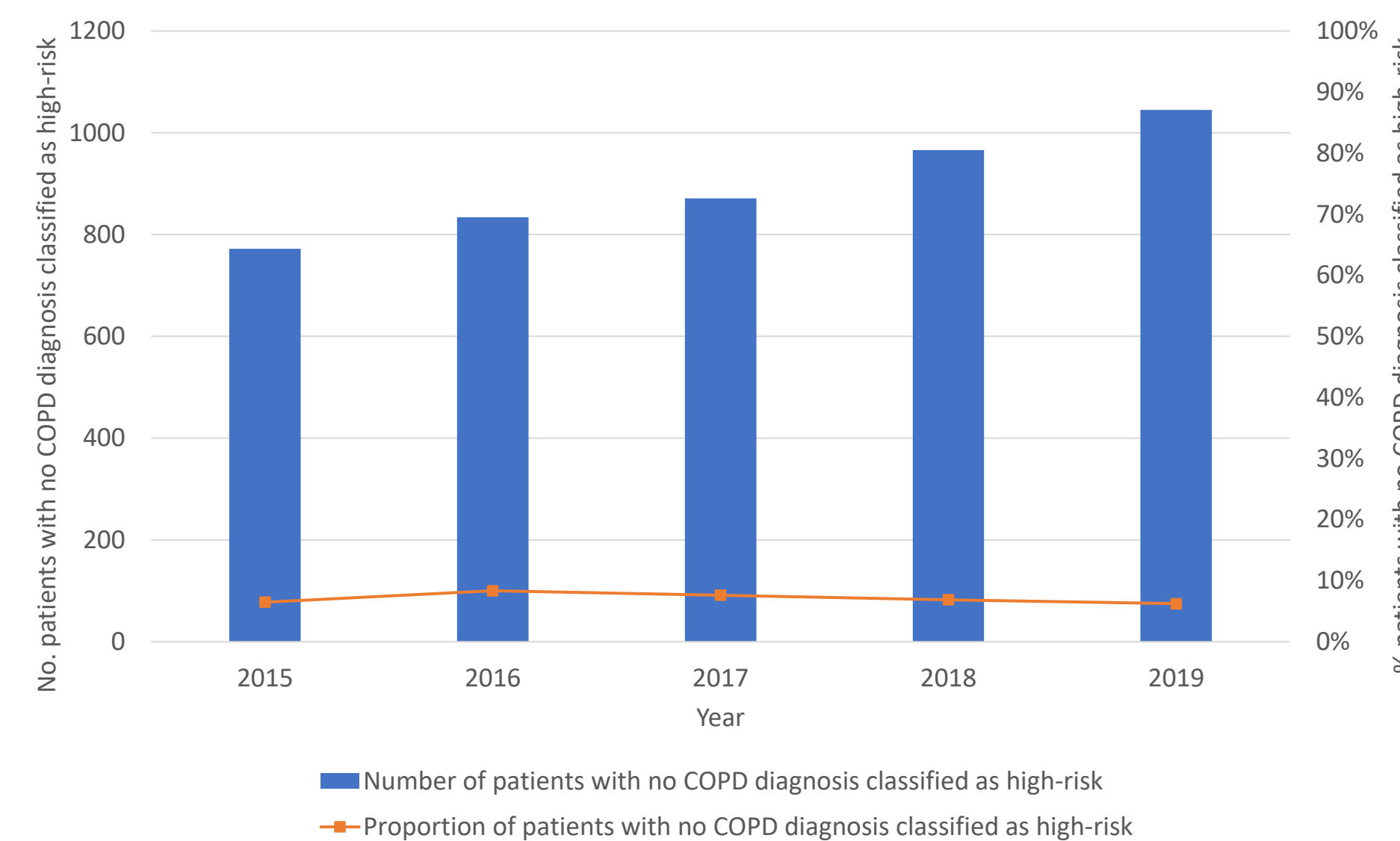


Figure 3: Proportion of undiagnosed high-risk patients receiving a new COPD diagnosis in 12-month follow-up period

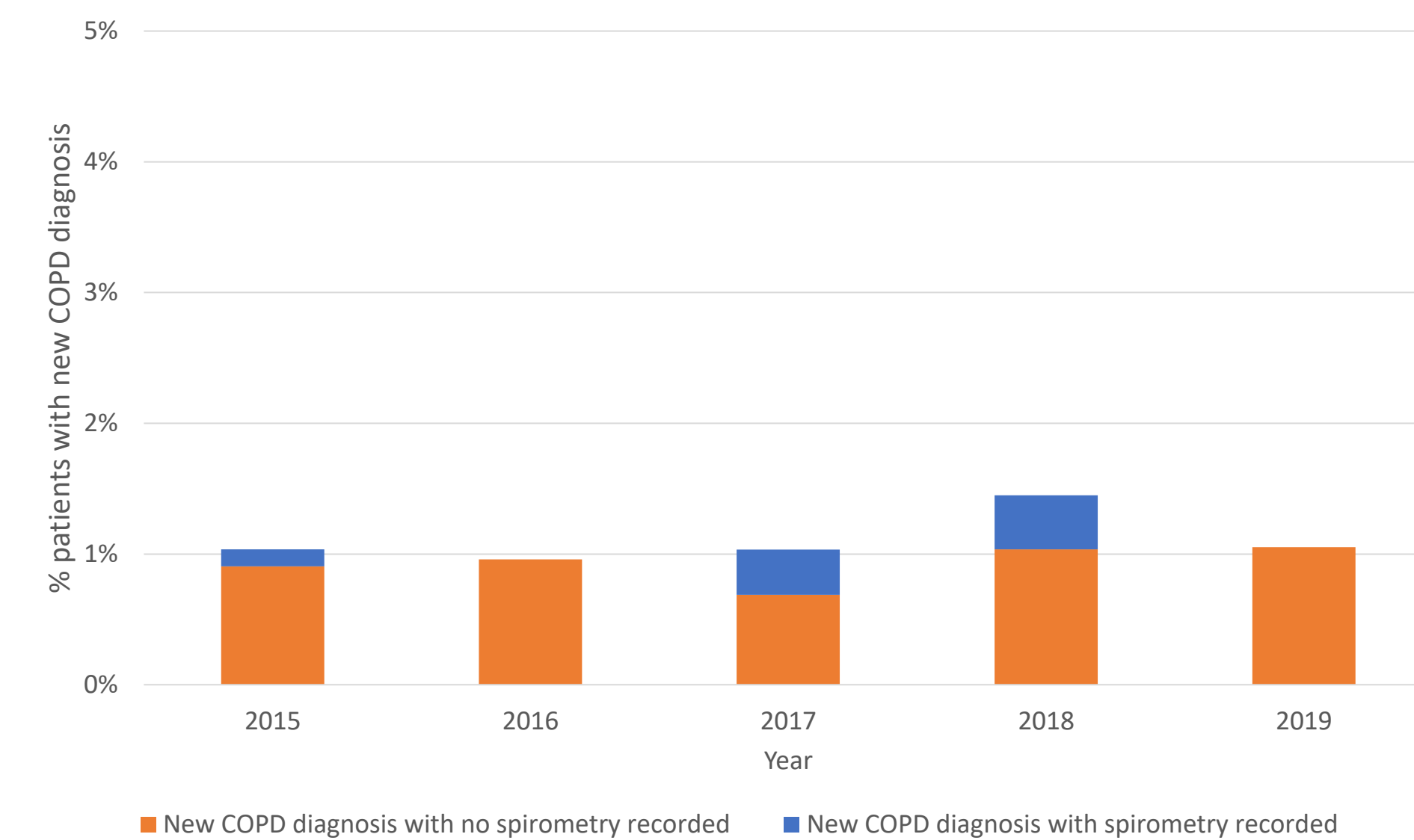
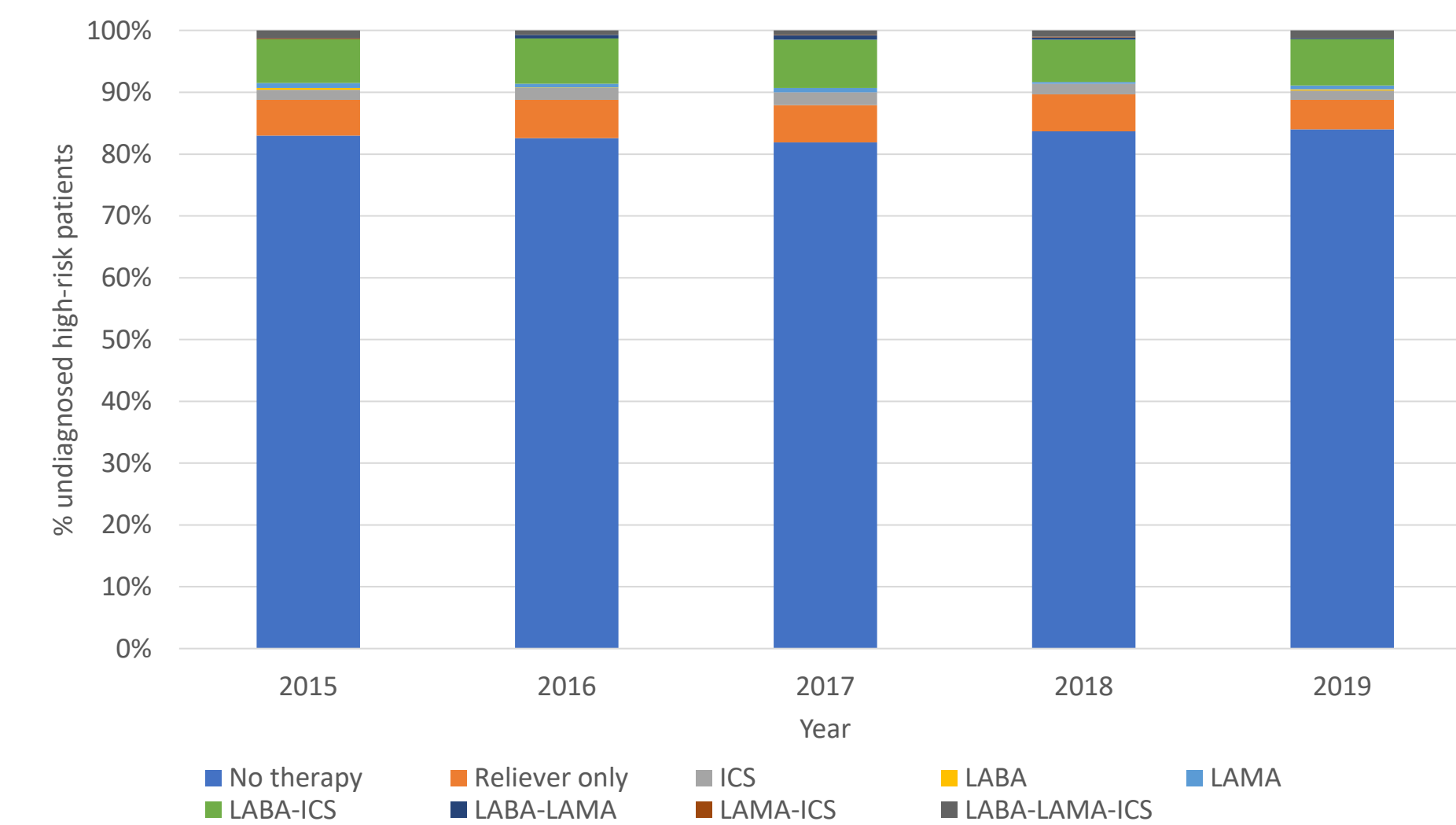


Figure 2: Proportion of undiagnosed high-risk patients prescribed inhaled therapy in the 12 months before 1st January in each study year



Conclusions

- There are opportunities for earlier diagnosis of COPD among patients at high risk of exacerbation in line with Australian and global guidelines and CONQUEST quality standards.
- Pharmacological management of undiagnosed high-risk patients should trigger further diagnostic tests in this population.

Research Approval

ENCePP registration number: ENCePP/DSPP/49365
ADEPT approval reference number: ADEPT1222

Disclosures

Andrew Dickens is an employee of the Observational and Pragmatic Research Institute. This study was conducted by Optimum Patient Care Australia (OPCA) and was partially funded by AstraZeneca and Optimum Patient Care Australia (OPCA).

References

1. Halpin DMG, et al. Lancet Reg Health Eur. 2023;29:100619. <https://doi.org/10.1016/j.lanepe.2023.100619>
2. Kerr M, et al. Lancet Reg Health Am. 2023;24:100546. <https://doi.org/10.1016/j.lana.2023.100546>
3. www.conquest.care (Viewed Mar 2023)
4. Pullen et al. Int J Chron Obstruct Pulmon Dis. 2021;16:2301-2322. <https://doi.org/10.2147/COPD.S313498>

Acknowledgements

Writing, editorial support, and/or formatting assistance in the development of this poster was provided by Ms Shilpa Suresh, MSc, Optimum Patient Care Australia (OPCA), partially funded by AstraZeneca and Optimum Patient Care Australia (OPCA).

Electronic copy of the poster



Audio Narration of the poster



Co-authors conflict of interest

