

Epidemiology, Survival, and Healthcare Resource Use of Patients With Pulmonary Hypertension Associated With Lung Disease and/or Hypoxia in the UK

EPH38



D.G. Kiely¹, S.J. Wort², D.F. Funes³, N. Petrica⁴, C. Proenca⁵, M. Pini⁴, M. Fernández Delgado³, G. Bacchini Jeanneret³

¹NiHR Biomedical Research Centre, Sheffield Teaching Hospitals NHS Foundation Trust, and Department of Clinical Medicine, University of Sheffield, United Kingdom, ²Royal Brompton Hospital and Imperial College, London, United Kingdom; ³Ferrer Internacional, Barcelona, Spain, ⁴Alira Health, Paris, France; ⁵Alira Health, Basel, Switzerland.

Background

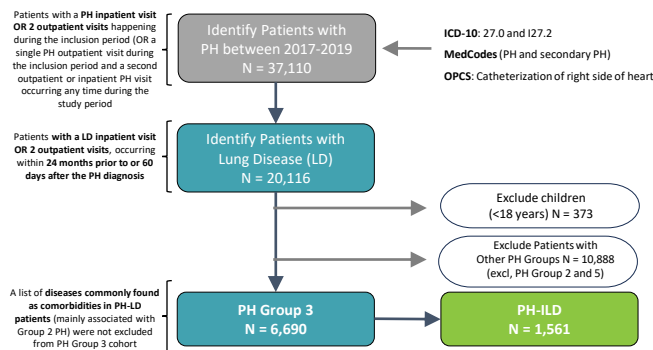
Pulmonary Hypertension associated with Lung Disease and/or Hypoxia (PH-LD), including PH with interstitial lung disease (PH-ILD), are severe conditions with a high mortality. However, data on epidemiology and healthcare management in the real-world setting are limited.

Objective

To estimate the epidemiology and the healthcare resource use (HCRU) of patients with PH Group 3 and PH-ILD in the UK.

Results

Decision Tree to Identify PH Group 3 and PH-ILD Patients

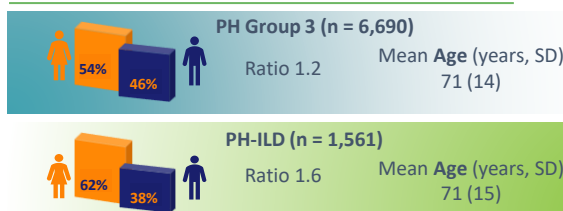


Epidemiology – Yearly Prevalence and Incidence Rate

PH Group 3	2017	2018	2019
Incidence rate per 10,000 person years (95% CI)	0.68 (0.65-0.71)	0.69 (0.66-0.72)	0.81 (0.78-0.84)
Prevalent rate per 10,000 patients (95% CI)	0.96 (0.93-0.99)	1.46 (1.42-1.5)	1.41 (1.37-1.45)
PH-ILD			
Incidence rate per 10,000 person years (95% CI)	0.16 (0.15-0.18)	0.15 (0.14-0.17)	0.19 (0.18-0.21)
Prevalent rate per 10,000 patients (95% CI)	0.26 (0.24-0.27)	0.37 (0.35-0.39)	0.36 (0.33-0.38)

Yearly incidence rates included only new cases without a prior diagnosis in the preceding 24 months

Age and Sex Distribution



Abbreviations: *phosphodiesterase-5 inhibitors (PDE5i), soluble guanylate cyclase stimulators (sGCs), calcium channel blockers (CCBs)

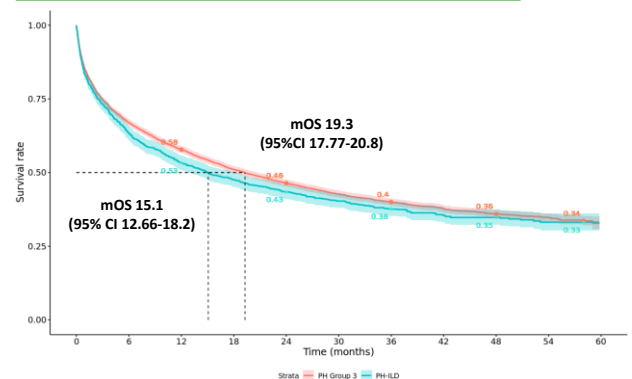
Methods

Data source: Retrospective observational study using the UK Clinical Practice Research Datalink (CPRD) linked to Hospital Episode Statistics (HES), including >18 million patients.

Study design: An algorithm, developed with six PH medical experts, identified PH Group 3 and PH-ILD patients from January 2017 to December 2019 based on a combination of PH and LD ICD-10 codes. The index date was the first PH diagnosis, with a 24-month minimum lookback period and up to 60 months follow-up until end of study, loss to follow-up or death. The study period spanned from 2015 to 2022.

Yearly prevalence rates were estimated for 2017-2019. Patients with active diagnoses codes were identified each year. Prevalence rates were calculated as the number of annual cases per 10,000 people in the CPRD-contributing practices population, with 95% confidence intervals.

Overall Survival



Mean Frequency of visits during the first-year of follow-up

Hospitalization rates decreased over time.

Mean Per patient-year	PH Group 3, Rate (95% CI)	PH-ILD, Rate (95% CI)
All-cause Inpatient hospitalizations	3.46 (3.4-3.51)	2.93 (2.82-3.03)
Length of stay, Mean(SD)	12 (16) days	11 (14) days
PH-related Inpatient Hospitalizations	1.87 (1.83-1.91)	1.78 (1.7-1.87)
Length of stay, Mean(SD)	13 (16) days	12 (14) days
LD-related Inpatient Hospitalizations	3 (2.89-2.99)	2.51 (2.41-2.6)
Length of stay, Mean(SD)	12 (16) days	11 (14) days

Note: PH and LD related hospitalization are not mutually exclusive

Drugs Prescriptions (PDE5i, sGCs, CCBs, and Anticoagulants*)

In the realm of PAH treatments, predominantly handled by specialists, analysis of GP prescriptions showed that fewer than 20% of patients had documented relevant drug prescriptions, regardless of cohort. Among these, PDE5i prescriptions were noted in 4% to 8% of patients with PH Group 3 and PH-ILD.

Conclusion

This study demonstrates that although rare, pulmonary hypertension associated with lung disease and the subgroup of patients with interstitial lung disease, have poor survival outcomes and suffer a significant clinical burden.