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



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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 realworld 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.

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.

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	0,68	0,69	0,81
years (95% CI)	(0,65-0,71)	(0,66-0,72)	(0,78-0,84)
Prevalent rate per 10,000	0,96	1,46	1,41
patients (95% CI)	(0.93 - 0.99)	(1.42 - 1.5)	(1.37 - 1,45)
PH-ILD			
Incidence rate per 10,000 person	0,16	0,15	0,19
years (95% CI)	(0,15-0,18)	(0,14 - 0,17)	(0,18-0,21)
Prevalent rate per 10,000	0,26	0,37	0,36
patients (95% CI)	(0.24 - 0,27)	(0,35-0,39)	(0,33 - 0,38)

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

Age and Sex Distribution



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.



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% Cl)	PH-ILD, Rate (95% Cl)	
All-cause Inpatient hospitalizations	3.46 (3.4-3.51)	2.93 (2.82-3.03)	
All-cause 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.