

Long COVID during Omicron: Patients' characteristics, vaccine uptake and healthcare resource utilisation

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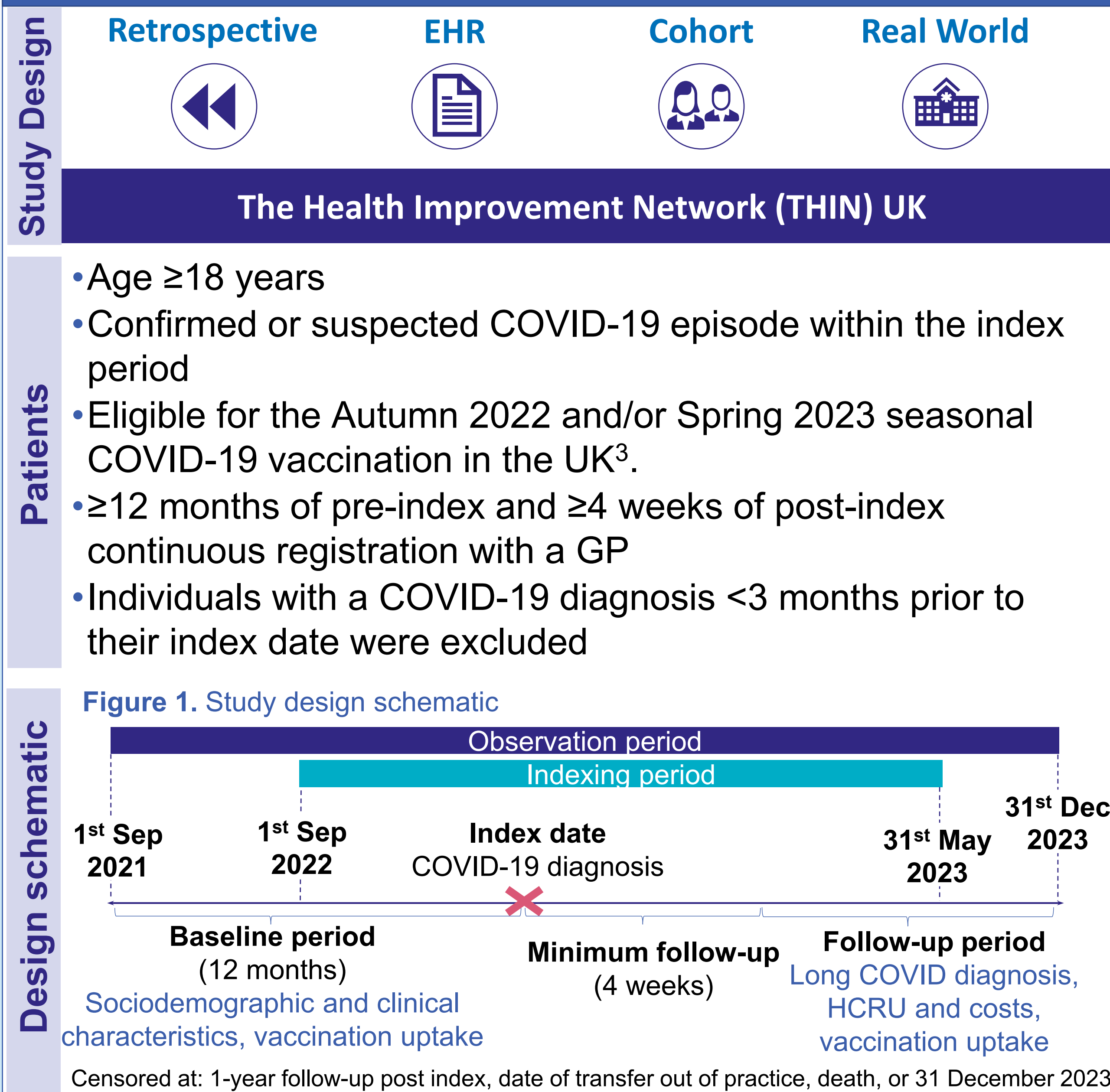
INTRODUCTION

- Recent data from the UK's Office for National Statistics (2024) indicated that long COVID affected 3.3% of the population in England and Scotland (approximately 2 million people)¹.
- Risk factors for long COVID include being female, smoking, immunosuppression, prior hospitalisation, and comorbidities such as obesity, hypertension, cardiovascular, cerebrovascular, respiratory, and kidney disease².
- There is limited research on the impact of vaccine uptake on the development of long COVID, especially during Omicron period.

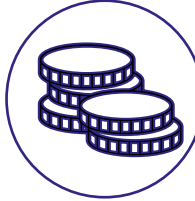
AIM

- To describe characteristics, vaccine uptake and healthcare resource utilisation (HCRU) in patients with and without long COVID in the UK primary care setting.

METHODS



METHODS (continued)

Exposure	Cases (long COVID): ≥1 primary care clinical code for long COVID symptoms ⁴ or a long COVID primary care diagnostic or referral clinical code observed ≥4 weeks ⁵ after index.		
	Controls: Patients that did not develop long COVID after index.		
Outcomes	 Sociodemographics	 HCRU	
	 Clinical characteristics	 Primary care costs	
	 Vaccination uptake		
	Statistical analyses		
Comparisons between patients with long COVID vs those without long COVID were performed using appropriate statistical tests: t-tests for continuous variables, Mann-Whitney U tests for ordinal variables, Fisher's exact tests for categorical variables, and Wald tests for HCRU rates, at a two-sided significance level of 0.05.			

RESULTS

- A total of 5,661 adults with COVID-19 and eligible for seasonal COVID-19 vaccination were included in the study; of whom, 49.0% (n=2,772) developed long COVID during follow-up.
- Statistically significant differences (p<0.05) were observed between long COVID cases and comparators without long COVID (**Table 1**).
- Compared to comparators, a greater proportion of cases were older (≥65 years: 51.5% vs. 46.6%), female (61.3% vs 64.9%), ever-smokers (40.9% vs 46.5%), overweight or obese (41.7% vs. 32.9%) and had a higher comorbidity burden.
- Whilst a similar proportion of cases and comparators had received seasonal COVID-19 vaccination pre-index (44.9% vs 44.3%), the timing of the vaccinations differed. Fewer cases received a COVID-19 vaccination more than 6-months before their index COVID-19 infection (54.8% vs 58.2%). However, a greater proportion were vaccinated after their index infection (58.4% vs 53.7%).
- During the follow-up period, cases had statistically higher primary care consultation rates, mean cost, and hospitalisation rates compared to comparators (p<0.05) (**Table 2**).

RESULTS (continued)

Table 1. Baseline sociodemographic, clinical characteristics and vaccination uptake in long COVID cases and comparators.

	Long COVID cases (n=2,772)	Comparators (n=2,889)	p-value
Age (years) at index			
18 - <50 years	339 (12.2)	428 (14.8)	
50 - <65 years	1,005 (36.3)	1,113 (38.5)	
65 - <75 years	576 (20.8)	525 (18.2)	≤0.001
75 - <85 years	558 (20.1)	507 (17.5)	
85+ years	294 (10.6)	316 (10.9)	
Sex, n (%)			
Female	1,799 (64.9)	1,770 (61.3)	≤0.01
Smoking status, n (%)			
Non-smoker	1333 (50.8)	1536 (56.5)	
Ex-smoker	909 (34.7)	833 (30.6)	≤0.001
Current smoker	380 (14.5)	350 (12.9)	
BMI (kg/m²), n (%)			
Underweight (<18.5)	46 (1.7)	35 (1.2)	
Normal weight (18.5-24.9)	353 (12.7)	345 (11.9)	
Overweight (25.0-29.9)	463 (16.7)	445 (15.4)	≤0.001
Obese (≥30.0)	694 (25.0)	507 (17.5)	
Unknown	1,216 (43.9)	1,557 (53.9)	
Quan-CCI score, n (%)			
0	2,025 (73.1)	2,256 (78.1)	
1-2	646 (23.3)	537 (18.6)	≤0.001
3-4	91 (3.3)	81 (2.8)	
≥5	10 (0.4)	15 (0.5)	
COVID-19 infections prior to index infection			
Mean (SD)	0.2 (0.5)	0.1 (0.4)	≤0.001
Received a 2022-2023 seasonal COVID-19 vaccination			
Yes	1,244 (44.9)	1,279 (44.3)	0.65
Time since last COVID-19 vaccination pre-index, n (%)			
<3 months	546 (19.7)	486 (16.8)	
3-6 months	708 (25.5)	723 (25.0)	≤0.01
>6 months	1,518 (54.8)	1,680 (58.2)	
COVID-19 vaccination uptake post-index, n (%)			
Yes	1,619 (58.4)	1,551 (53.7)	≤0.001

RESULTS (continued)

Table 2. Healthcare resource use (HCRU) and costs during follow-up long COVID cases and comparators.

	Long COVID cases (n=2,772)	Comparators (n=2,889)	p-value
Primary care consultation rate (95% CI), per-person per-month			
	1.61 (1.59, 1.62)	0.91 (0.90, 0.92)	≤0.001
Mean primary care consultation costs (£, SD), per-person per-month			
	82 (63)	54 (69)	≤0.001
Hospitalisation rate (95% CI), per-person per-month			
	0.07 (0.07, 0.07)	0.04 (0.04, 0.05)	≤0.001

LIMITATIONS

- No standardised global definition for long COVID exists, leading to potential misdiagnosis or misclassification.
- Secondary care HCRU information in the THIN UK database is limited, potentially underestimating hospitalisation rates.
- As patients were required to have a diagnosis of COVID-19 (after the end of routine COVID testing/reporting), it is likely that these patients have more health seeking behaviour.

CONCLUSIONS

- Despite Omicron being associated with lower long COVID incidence than earlier variants⁶, HCRU and cost differences were observed between long COVID cases and comparators.
- Adherence to UK COVID-19 vaccination recommendations was generally low across both groups.
- These findings contribute to the evidence base of long COVID in the Omicron period and can contribute to informing policy decisions regarding prevention strategies for long COVID.

References

- Office for National Statistics (ONS). Self-reported coronavirus (COVID-19) infections and associated symptoms, England and Scotland: 2024. Accessed 2025
- Tsampsian, V., et al., Risk Factors Associated With Post-COVID-19 Condition: A Systematic Review and Meta-analysis. JAMA Internal Medicine, 2023. 183(6): p. 566-580
- VACCINE ELIBILITY CRITERIA
- Subramanian, A., et al., Symptoms and risk factors for long COVID in non-hospitalized adults. Nature Medicine, 2022. 28(8):p.1706-1714.
- NICE. COVID-19 rapid guideline: managing the long-term effects of COVID-19 2024. Accessed 2025.
- Antonelli M, Pujol JC, Spector TD, Ourselin S, Steves CJ. Risk of long COVID associated with delta versus omicron variants of SARS-CoV-2. Lancet. 2022;399(10343):2263-4.

Acknowledgments

J Yang, C Harrison, R Butfield, C Reynard, D Gruben, HR. Volkman, and JL Nguyen are employees of Pfizer and may hold stock or stock options. KK. Rai, H Gowman, R Hulme, and I Jimenez are employees of Adelphi Real World, which received funds from Pfizer to conduct the study and develop the manuscript.

Funding: Funding for this study was provided by Pfizer Inc. Pfizer Inc. commissioned Adelphi Real World to independently conduct the study.

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Abbreviations: BMI: Body mass index; CCI: Charlson Comorbidity Index; CI: Confidence interval; COVID: Coronavirus disease; EHR: Electronic health record; GP: General practitioner; HCRU: healthcare resource utilisation; SD: Standard deviation; THIN: The Health Improvement Network