

Post-COVID symptoms prevalence, latent class pattern, and functional impact: A population-based cross-sectional study in two Chilean cities, 2024

Paola Rubilar^a, Mauricio Apablaza^b, Muriel Ramírez-Santana^c, Lina J Cortés^d, Loreto Nuñez-Franz^{e*}

^aCentro de Epidemiología y Políticas de Salud, Facultad de Medicina Clínica Alemana, Universidad del Desarrollo, Santiago, Chile; ^bFacultad de Gobierno, Universidad del Desarrollo, Santiago, Chile; ^cDepartamento de Salud Pública, Facultad de Medicina, Universidad Católica del Norte, Coquimbo, Chile; ^dInstituto de Ciencias e Innovación en Medicina, Facultad de Medicina Clínica Alemana, Universidad del Desarrollo, Santiago, Chile; ^eDepartamento de Salud Pública, Facultad de Ciencias de la Salud, Universidad de Talca, Talca, Chile

ABSTRACT

INTRODUCTION Post-COVID-19 condition affects approximately 6% of individuals globally. Most evidence is based on hospitalized patients and is from countries with relatively low vaccination coverage. This study estimates the prevalence of symptoms compatible with post-COVID-19 and explores symptom patterns in a population-based sample from Chile, a country with high vaccination coverage. Moreover, it examines their associations with daily activities and demographic, social, and clinical factors.

METHODS A cross-sectional study was conducted in May 2024 in two Chilean Cities, involving 654 participants (88.5% response rate) aged $\geq$ seven years, from randomly selected households. Ten post-COVID-19 symptoms that persisted for $\geq$ three months were collected. Latent class analysis identified symptom patterns.

RESULTS Among 277 participants with a confirmed COVID-19 diagnosis, 114 reported at least one symptom lasting $\geq$ 3 months. Population-weighted prevalence of at least one post-COVID-19 symptom was 17.4% in the general population and 40.7% among individuals with prior COVID-19. Among those with post-COVID-19 symptoms, 40.1% reported a reduction in daily activities. Three classes were explored: class 1 (38.4%) characterized by fatigue, ageusia/anosmia, and muscular/joint pain; class 2 (40.7%) with predominant cardiorespiratory symptoms; and class 3 (20.9%) with multisystemic symptoms. Class three was associated with a higher frequency of comorbidities (particularly diabetes), hospitalization, multiple COVID-19 episodes, and Aboriginal affiliation. Classes 2 and 3 were independently associated with younger age and diabetes, class 2 was additionally associated with lower educational attainment and lower BMI, while class 3 was additionally associated with Aboriginal affiliation.

CONCLUSIONS This population-based study reveals a substantial burden of at least one post-COVID-19 symptom, with distinct symptom classes highlighting the need for tailored interventions. These findings underscore the importance of targeted public health policies and personalized clinical approaches, particularly for severe symptom classes that impact quality of life and contribute critical data from Latin America to the global understanding of post-COVID-19 condition.

KEYWORDS Long COVID, Chile, Cluster Analysis, prevalence



* Corresponding author lnunezf@utalca.cl

Citation Rubilar P, Apablaza M, Ramírez-Santana M, Cortés LJ, Nuñez-Franz L. Post-COVID symptoms prevalence, latent class pattern, and functional impact: A population-based cross-sectional study in two Chilean cities, 2024. Medwave 2026;26(8):e3224
DOI 10.5867/medwave.2026.08.3234

Submitted May 1, 2026, **Accepted** Sep 3, 2026,

Published Sep 23, 2026

Postal address Avenida Lircay s/n Universidad de Talca Depto. Salud Pública, Talca, Chile

INTRODUCTION

The post-COVID-19 condition [1,2], also known as long COVID, is characterized by a set of symptoms that persist after the acute phase of SARS-CoV-2 infection, typically for 4 to 12 weeks or longer. Despite its widespread recognition, the condition lacks a standardized definition [1,3], likely because its symptoms are heterogeneous and multisystemic, affecting the respiratory, cardiovascular, neurological, gastrointestinal, and musculoskeletal systems. Among these varied and heterogeneous manifestations, general symptoms such as

MAIN MESSAGES

- Understanding the population-level burden of post-COVID-19 symptoms is critical for public health response
- Prevalence of post-COVID-19 symptoms reached 40.7% in those with a diagnosis of COVID-19 and 17.4% in the whole population.
- The three explored phenotypes require distinct clinical protocols, enabling prioritization of high-risk subgroups for intensive monitoring and comprehensive care.
- The main limitation is the cross-sectional design, which precludes causal inference and assessment of symptom progression over time.
- The most severe class (the multisystemic phenotype) was independently associated with younger age, diabetes, and Aboriginal affiliation.

fatigue are common, with fatigue being the most frequently reported [4].

The World Health Organization (WHO) estimates that approximately 6.2% of individuals who survive acute COVID-19 infection develop post-COVID-19 condition [5,6]. Several factors have been described as influencing the type and severity of post-COVID-19 symptoms, including gender, age, severity of acute infection, pre-existing comorbidities, socioeconomic status, ethnicity, vaccination status, and the SARS-CoV-2 variant responsible for the infection [7–10]. These factors contribute to the variability in clinical presentation and underscore the need for tailored approaches to diagnosis and treatment.

Due to the heterogeneity of symptoms and affected systems, researchers have attempted to identify symptom patterns by using various clustering methodologies. Among the methods used are exploratory factor analysis [11], agglomerative hierarchical clustering (Ward's method) [12–16], non-negative factorization matrices [17], latent class analysis (LCA) [18,19], and partitioning around medoids [20] (k-medoids), including multi-methods [15,16].

Common classes include respiratory symptoms, neurological/mental/psychiatric symptoms, cardiovascular/cardiac/circulatory symptoms, musculoskeletal/nervous symptoms, digestive/diarrhea symptoms, and other isolated symptoms such as fatigue, pain, and minor symptoms. At the same time, studies have reported that respiratory symptoms are more frequent in hospitalized patients and those with severe acute infections [14]. Likewise, neuropsychiatric symptoms were more frequent in young people, in women, and in outpatients [20].

The available literature on post-COVID-19 conditions consists primarily of studies conducted in countries outside Latin America, with most research focusing on hospitalized patients, self-reported individuals with COVID-19, or utilizing data from hospitals or healthcare centers. Nevertheless, there remains limited evidence from population-based studies that include individuals who have not engaged with the health system, a gap that is particularly relevant in countries with high vaccine coverage and persistent health inequalities, like Chile [21,22]. This study aims to report a) the prevalence of symptoms compatible with the post-COVID-19 condition in a population-based sample from two Chilean cities, b) explore symptom

classes, and c) examine their relationship with the reduction of daily activities and demographic, social, and clinical factors.

METHODS

Study design and setting

This cross-sectional study presents a secondary analysis of a population-based COVID-19 seroprevalence survey conducted in La Serena/Coquimbo and Talca, Chile, in May 2024 [22].

Participants

All participants from a prior data collection wave [23] were re-invited. Non-responders were replaced using two-stage random sampling (census block selection, then systematic household selection within blocks). Non-participating households were systematically replaced either in the same block or, if necessary, through random block replacement. All household members aged $\geq$ seven years were eligible to participate. The same recruitment methodology was employed in previous study round [21,24].

Study size

The current study is a secondary analysis of a sample originally powered for seroprevalence. The sample size was calculated based on the 97% seroprevalence [23]. Assuming a margin of error of 3%, 95% confidence level, and a design effect of 2.0 to account for the two-stage sampling. The minimum required sample size was 249 participants per city (498 total). Sample size calculation was performed using the OpenEpi software.

Variables

Dependent Variables: Participants self-reported COVID diagnosis (polymerase chain reaction, antigen, or clinical diagnosis). Ten post-COVID-19 symptoms persisting for > three months after infection were assessed using Household Pulse Survey questions developed by the National Center for Health Statistics and the Census Bureau in the United States of America [25]. Items were translated directly into Spanish and culturally adapted to local language use where necessary. Each symptom was rated as present or absent (yes/no): cognitive impairment, memory impairment, vertigo or dizziness, anosmia

or ageusia/altered taste, dyspnea, chest pain, palpitations, fatigue, exercise intolerance, and musculoskeletal pain. These items were selected for their brevity and prior use in large-scale population surveillance, facilitating comparability with international post-COVID-19 condition estimates; however, they were not subjected to formal cross-cultural adaptation or psychometric validation in the Chilean population. Reduction in daily activities was an additional outcome variable.

Independent variables: sex (male/female/other/prefer not to answer), age (< 30, 30 to 59, $\geq$ 60), aboriginal affiliation (yes/no), education (participants $\geq$ 18: high school education or less/technical/professional-postgraduate), health insurance (public/private/armed forces), Body Mass Index (BMI) (self-reported: underweight/normal/overweight/obese), smoking (yes/no), comorbidities (neoplastic disease/heart disease/chronic kidney disease/hypertension/obesity/autoimmune disease/diabetes), vaccination (none/first dose and booster/bivalent or omicron), COVID-19 episodes (one vs $\geq$ two), and due to COVID-19 hospitalization (yes/no).

Data sources and measurements

Trained undergraduate health science students conducted standardized 15-to-20-minute home interviews using questionnaires. Interviewers were trained in consistent data collection and entered data in real time into REDCap using tablets. Data collected included: demographics, COVID-19 history, COVID-19 symptoms, vaccine history, post-COVID-19 symptoms, and risk factors. Vaccination status was verified using two approaches: (1) vaccine card or participant accessing their records through the national immunization registry portal or presenting vaccination cards; (2) when this was not possible, participants authorized the research team to obtain records directly from the national immunization program. The data manager and principal investigator performed daily quality checks; when necessary, participants were recontacted to resolve inconsistencies or data-entry errors.

Statistics

Prevalence rates were calculated based on the population expansion of the sample. Expansion factors were calculated considering a two-stage, structured probabilistic sampling design: 1) conditional household selection probability within census district, 2) individual selection probability ($\geq$ 7 years) assuming equivalent probabilities within the household, 3) differential nonresponse adjustment based on the number of actual participants in each household, 4) calibration based on 2024 population projections estimated by the National Institute of Statistics.

The expansion factors were derived from the sampling design used for the seroprevalence and were not recalculated for nonresponse specific to the symptom questionnaire. Since all participants answered the symptom questionnaire, no additional non-response adjustment was necessary. Descriptive statistics were computed for each symptom, including

proportions and standard errors, to summarize the prevalence of post-COVID-19 symptoms. All symptom variables were dichotomous (yes/no). Latent class analysis (LCA) was utilized to identify unobserved subgroups based on the co-occurrence of post-COVID-19 symptoms. Latent class analysis models the conditional independence structure within each latent class, enabling it to identify symptom types where specific symptom classes co-occur. Furthermore, latent class analysis classes directly correspond to interpretable patient phenotypes with clear probabilistic symptom profiles that will be further explored in the paper. The latent class analysis model was specified without covariates, assuming conditional independence between symptoms within each latent class. Several strategies were explored to determine the number of classes, including the elbow method and the Bayesian Information Criterion (BIC). While the Bayesian Information Criterion continued to improve beyond three classes, the final 3-class solution was selected based on interpretability and clinical relevance [26] rather than statistical fit alone. Finally, we combined the elbow method with a normative decision, acknowledging that identified groups are meaningful, interpretable, and useful for policy or clinical applications. Conditional response probabilities for each symptom across latent classes were estimated, and individuals were assigned to the most likely latent class based on posterior probabilities. Sociodemographic, daily activities reductions, and clinical characteristics were then summarized for each class to provide comparative profile. To examine the association between sociodemographic and clinical characteristics and membership in the latent post-COVID-19 symptom classes, survey-weighted multinomial logistic regression models were estimated using the complex sampling design of the study population. Class membership was specified as the dependent variable, with Class 1 serving as the reference category. Exponentiated coefficients are presented as relative rate ratios (RRRs), representing the relative likelihood of belonging to Class 2 or Class 3 compared with Class 1. The fully adjusted model included all variables associated with class membership at $p < 0.10$ in the bivariate comparison: aboriginal affiliation, educational attainment, number of previous COVID-19 episodes, smoking status, history of hospitalization due to COVID-19, diabetes, and body mass index (BMI). Sex and age were retained in the model despite not reaching statistical significance in the bivariate analysis, given their role as potential confounders. Survey weights were incorporated in all analyses to account for the sampling design and produce population-representative estimates. To ensure representativeness, analyses applied survey weights and were conducted in RStudio (version 2024.12.1+563) using R (version 4.5.0) or in STATA/IC 15 (StataCorp LLC).

Ethics approval

All procedures performed in studies involving human participants were in accordance with the ethical standards of the institutional and/or national research committee (1) Scientific ethical committee from the Faculty of Medicine,

Universidad Católica del Norte, Resolution number 63/2023, dated October 16th, 2023. (2) Scientific ethical committee from the Faculty of Medicine, Universidad del Desarrollo, dated December 13th, 2023, and 3) Scientific ethical committee from Universidad de Talca, Folio 30-2023-E, dated April 17th, 2024. Additionally, it was approved by the Institutional Committee of Biosecurity of Universidad Católica del Norte 07/2023 dated October 2023, Universidad del Desarrollo, CIB-FORM-01B, dated 14 November 2023, and Universidad de Talca 20-CBS-2023 dated November 9th, 2023) and with the 1964 Helsinki declaration and its later amendments or comparable ethical standards. Informed consent was obtained from all subjects involved in the study, including assent from minors (7 to 17 years).

RESULTS

A total of 654 participants were recruited in May 2024 (response rate 88.5%): 318 from La Serena-Coquimbo and 336 from Talca (Supplementary data. online supplementary figure 4). Across both cities, 277 reported prior COVID-19 infections confirmed by Polymerase Chain Reaction, antigen test, or medical diagnosis. Among them, 117 reported at least one post-COVID-19 symptom, with a mean age of 49.9 years (range: 7 – 95). Three women reported menstrual changes as their sole symptoms (these cases were excluded to avoid gender-specific bias). The final analytic sample comprises 114 individuals (Figure 1).

In the expanded sample ($n = 722\ 430$), 52.0% were female. The population-based prevalence of at least one post-COVID-19 symptom was 17.4%, and 40.7% among those with prior COVID-19. Figure 2 presents the prevalence of post-COVID-19 condition symptoms as percentages among individuals with prior COVID-19, and those who have experienced at least one symptom lasting for at least three months. Fatigue was the most common (66.7%), followed by dyspnea (59.4%), musculoskeletal pain (35.7%), and exercise intolerance (28.2%). Chest pain (26.4%) and palpitations (23.6%) were also frequent. Neuro-sensorial symptoms included cognitive impairment (24.7%), memory issues (22.2%), olfactory or taste alterations (18.5%), and dizziness (10.3%).

Regarding symptoms interfering with daily activities, 40.1% reported limitations. Individuals without daily activity reduction reported the fewest symptoms (median two), concentrated between one and three (mean: 3.04). Those with a slight reduction in daily activity had a median of three, interquartile range (IQR) two to three (mean: 3.2). In contrast, individuals with a major reduction also had a median of three symptoms but a higher mean (4.8) and wider IQR (three to eight), indicating greater heterogeneity and symptom burden (Supplementary information. online supplementary figure 5).

Three main classes emerge: Class 1 grouped symptoms related to fatigue and physical performance, including chest pain, fatigue, dyspnea, and exercise intolerance. Class 2 includes symptoms such as palpitations, memory and cognitive

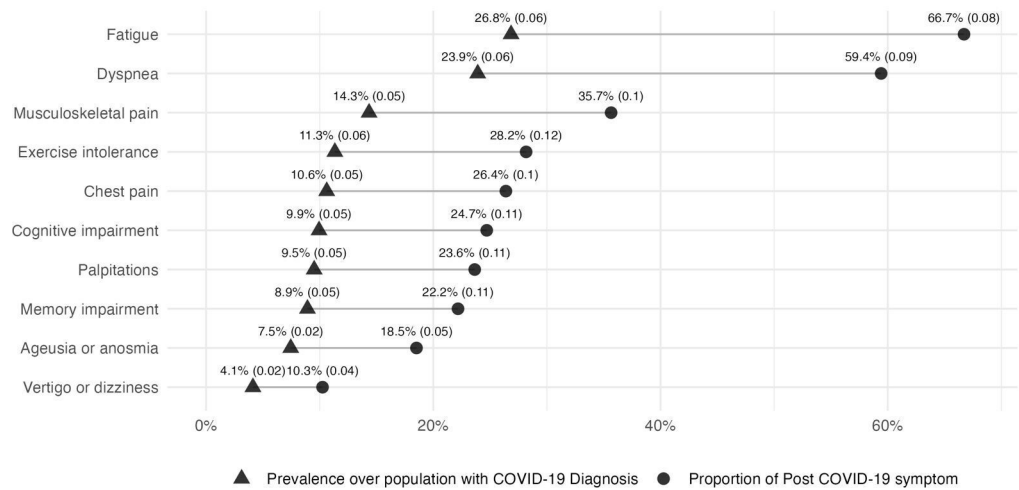
impairment, musculoskeletal pain, and vertigo or dizziness, reflecting a combination of cardiorespiratory and neurological manifestations. Class 3 comprised taste alteration and ageusia, consistent with neuro-sensorial alterations.

Following a normative decision, the latent class analysis seems to have converged on a three-class solution after multiple iterations with different starting values were run to ensure model stability (Supplementary Material). The latent class analysis identified three latent classes: Class 1 (38.4%, $n = 48,230$); Class 2 (40.7%, $n = 51,115$); and Class 3 (20.8%, $n = 26,177$) (Table 1 and Figure 2). Conditional item response probabilities reveal three distinct post-COVID-19 condition symptom structures. Class 1 exhibited high probabilities of fatigue (42.7%), ageusia or anosmia (39.5%), and musculoskeletal pain (29.5%), but relatively low probabilities of cognitive symptoms, suggesting a predominantly physical symptoms profile. Class 2 represents a phenotype with high probabilities across various domains, such as dyspnea (95.8%), fatigue (73%), and exercise intolerance (32.4%), but with almost no neurological manifestations. Finally, Class 3 includes a group with very high probabilities across all symptoms (mostly over 70%).

Table 1 presents the distribution of the study variables across latent class analysis classes. Class 1 and 2 had the highest proportion with High school or less education. Class 3 had the highest proportion of participants with Aboriginal affiliation, technician education, comorbidity, diabetes, multiple COVID-19 episodes, and hospitalization. It also included a higher proportion of men and tobacco users, although these differences were not statistically significant. All symptoms varied significantly across classes, with Class 3 showing the highest prevalence of symptoms.

Multinomial logistic regression analysis used class 1 as the reference category. Age was retained in the model as a potential confounder despite not reaching significance in the bivariate analysis (Table 2). In the adjusted model, younger participants were more likely to belong to class 2 (RRR = 0.95; 95% CI: 0.90 to 0.99; $p = 0.04$) and 3 (RRR = 0.92; 95% CI: 0.86 to 0.99; $p = 0.03$), relative to class 1. Diabetes was independently associated with both class 2 (RRR = 35.25; 95% CI: 3.57 to 348.0; $p = 0.00$) and class 3 (RRR = 11.75; 95% CI: 1.44 to 95.84; $p = 0.02$). Class 2 was additionally associated with lower body mass index (RRR = 0.83; 95% CI: 0.75 to 0.93; $p = 0.00$) and with educational attainment: compared to participants with high school education or less (reference), having higher technical education was associated with substantially lower relative risk of class 2 membership (RRR = 0.08; 95% CI: 0.01 to 0.42; $p = 0.00$), while professional/post-graduate education showed no significant association (RRR = 1.34; 95% CI: 0.37 to 4.80; $p = 0.65$). Class 3 was additionally associated with Aboriginal affiliation (RRR = 10.98; 95% CI: 1.74 to 69.44; $p = 0.01$). No significant associations were found for sex, number of previous COVID-19 episodes, smoking status, or history of hospitalization due to COVID-19 in either class.

Figure 1. Population prevalence of post-COVID-19 symptoms among individuals with prior COVID-19 diagnosis and symptoms proportion among those reporting persistent symptoms lasting ≥3 months, Chile, May 2024.



Graph of post-COVID-19 symptoms prevalence in total COVID-19 cases (left) and in those with persistent symptoms (right), ordered by descending frequency.
Source: Prepared by the authors of this study.

Figure 2. Latent class analysis of post-COVID-19 symptoms: Probability of symptoms by class, Chile, 2024.

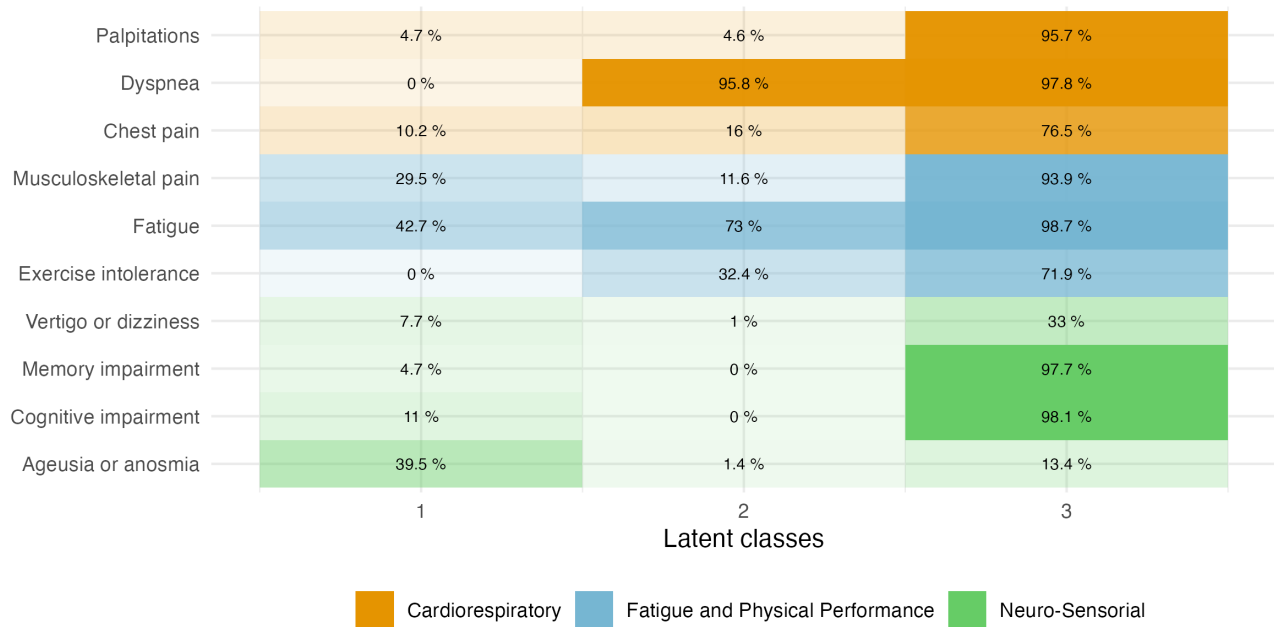


Chart displaying the probability of conditional symptoms across three latent classes. The horizontal axis depicts the three classes (1 to 3) in columns from left to right. The vertical axis depicts symptom probability. Each column represents the probability of symptom occurrence, with dark colors indicating a higher probability. Symptoms are colored by domain: cardiorespiratory (brown), fatigue and physical performance (blue), and neuro-sensorial (green).
Source: Prepared by the authors of this study.

DISCUSSION

Among individuals with prior COVID-19 infection, 40% experienced at least one symptom persisting for three months, underscoring post-COVID-19 as a significant public health concern. Latent class analysis identifies three

phenotypes. The most severe class, the multisystemic phenotype, had high prevalence of comorbidities, hospitalization due to COVID-19, multiple episodes of COVID-19, and socio-demographic variables. After adjustment, multinomial logistic regression confirmed younger age and diabetes as independent correlates of both symptom classes (2 and 3), with class 2

Table 1. Population sociodemographic and clinical characteristics stratified by post-COVID-19 symptom classes resulting from LCA and bivariate comparisons across class, Chile, May 2024.

Variable	Categories	Total		Class 1		Class 2		Class 3		p value ^b
		N	%	%	%	%	%	%	%	
Total		125522	100	38.4 (24.4 to 54.7)	40.7 (24.3 to 59.5)	20.9 (6.7 to 49.0)				
Sex	Male	56779	45.2	40.3 (24.8 to 57.9)	40.7 (15.8 to 71.5)	63.2 (17.2 to 93.4)				0.61
	Female	68743	54.8	59.7 (42.1 to 75.2)	59.3 (28.5 to 84.2)	36.8 (6.6 to 82.8)				
Age group	< 30 years	28473	22.7	15.0 (6.1 to 32.2)	32.8 (9.2 to 70.2)	17.1 (1.7 to 70.5)				0.40
	30 to 59 years	85505	68.1	70.9 (54.6 to 83.1)	58.0 (28.2 to 82.9)	82.8 (29.6 to 98.2)				
	60 years and over	11544	9.2	14.1 (6.7 to 27.3)	9.2 (3.7 to 21.3)	0.1 (0.006 to 0.9)				
Aboriginal affiliation	Yes	34652	27.6	14.0 (8.3 to 22.5)	13.9 (4.9 to 33.5)	79.4 (41.2 to 95.5)				0.00
	No	90870	72.4	86.0 (77.5 to 91)	86.1 66.5 to 95.1)	20.6 (4.5 to 58.8)				
Education (age 18 years or more)	High school or less	63467	51.6	56.4 (37.9 to 73.3)	65.2 (42.0 to 82.9)	17.0 (3.8 to 51.3)				0.00
	Higher Technician	39287	31.9	28.6 (13.4 to 50.8)	11.3 (6.0 to 20.4)	76.9 (36.9 to 95.0)				
	Professional/postgraduate	20324	16.5	15.0 (8.7 to 24.5)	23.5 (10.8 to 43.7)	6.1 (0.6 to 41.1)				
Insurance	Public	113037	91.0	92.2 (83.7 to 96.4)	88.6 (74.2 to 95.5)	93.4 (58.7 to 99.3)				0.89
	Private	10274	8.3	7.2 (3.1 to 15.6)	10.2 (3.9 to 24.1)	6.6 (0.7 to 41.2)				
	Armed forces	915	0.7	0.6 (0.08 to 4.6)	1.2 (0.1 to 8.8)	0.0				
Body mass index	Emaciated	737	0.6	1.5 (0.3 to 8.3)	0.0	0.0				0.08
	Normal	37954	30.4	18.3 (9.5 to 32.4)	54.1 (29.2 to 77.1)	6.9 (0.8 to 40.2)				
	Overweight	40128	32.1	46.9 (29.5 to 65.0)	23.2 (10.8 to 42.9)	22.2 (3.1 to 71.4)				
	Obese	46070	36.9	33.3 (21.2 to 48.0)	22.7 (10.6 to 41.9)	70.9 (23.3 to 95.1)				
Tobacco	Yes	44058	35.1	29.2 (14.1 to 50.8)	21.4 (10.6 to 38.5)	72.7 (25.1 to 95.5)				0.08
	No	81464	64.9	70.8 (49.1 to 85.9)	78.6 (61.5 to 89.3)	27.3 (4.5 to 74.9)				
Comorbidities	Yes	86148	68.6	53.8 (36.8 to 70.0)	70.3 (47.9 to 85.9)	92.6 (66.8 to 98.7)				0.03
	No	39374	31.4	46.2 (30.0 to 63.2)	29.7 (14.0 to 52.1)	7.4 (1.3 to 33.1)				
Neoplastic disease	Yes	702	0.6	0.0	0.9 (0.5 to 1.5)	0.9 (0.1 to 8.1)				0.34
	No	124820	99.4	100.0	99.1 (98.5 to 99.5)	99.1 (91.8 to 99.9)				
Heart disease	Yes	5977	4.8	5.7 (0.8 to 31.3)	4.6 (1.1 to 17.1)	3.3 (0.3 to 26.8)				0.93
	No	119545	95.2	94.3 (68.6 to 99.2)	95.4 (82.8 to 98.9)	96.7 (73.2 to 99.7)				
Chronic respiratory disease	Yes	30056	23.9	9.9 (3.2 to 26.4)	37.6 (13.0 to 70.8)	23.3 (3.3 to 72.9)				0.28
	No	95466	76.1	90.1 (73.6 to 96.8)	62.4 (29.2 to 87.0)	76.7 (27.1 to 96.7)				
Chronic kidney disease	Yes	2240	1.8	1.9 (0.2 to 12.9)	0.9 (0.2 to 4.3)	3.3 (0.3 to 26.8)				0.64
	No	123282	98.2	98.1 (87.1 to 99.7)	99.1 (95.6 to 99.8)	96.7 (73.2 to 99.7)				
Diabetes	Yes	26059	20.8	4.6 (1.3 to 14.8)	16.4 (8.1 to 30.2)	59.0 (13.9 to 92.8)				0.01
	No	99463	79.2	95.4 (85.2 to 98.7)	83.6 (69.8 to 91.8)	41.0 (72.0 to 86.1)				
Hypertension	Yes	22014	17.5	25.6 (10.7 to 49.6)	14.0 (5.7 to 30.5)	9.6 (2.1 to 34.1)				0.34
	No	103508	82.5	74.4 (50.4 to 89.3)	86.0 (69.5 to 94.3)	90.4 (65.9 to 97.9)				
Self-reported obesity	Yes	11618	9.3	14.6 (6.8 to 28.3)	4.3 (1.6 to 10.6)	9.2 (1.6 to 38.0)				0.25
	No	113904	90.7	85.4 (71.6 to 93.2)	95.7 (89.3 to 98.3)	90.8 62.0 to 98.3)				
Autoimmune disease	Yes	1908	1.5	1.7 (0.2 to 12.1)	2.1 (0.3 to 14.5)	0.0				0.79
	No	123614	98.5	98.3 (87.9 to 99.8)	97.9 (85.5 to 99.7)	100.0				
Vaccination	No vaccine	2356	1.9	0.0	2.6 (0.3 to 17.6)	3.9 (0.3 to 30.1)				0.32
	First dose+ booster	70900	56.5	51.7 (33.9 to 69.1)	72.0 (52.1 to 85.9)	34.9 (6.1 to 81.3)				
	Bivalent or Omicron	52266	41.6	48.3 (30.9 to 66.0)	25.3 (12.8 to 44.0)	61.2 (15.5 to 93.1)				
N° COVID episodes	1	81357	64.8	70.4 (48.5 to 85.7)	80.0 (61.2 to 91.0)	24.8 (3.9 to 73.0)				0.04
	2 or more	44165	35.2	29.6 (14.3 to 51.5)	20.0 (9.0 to 38.8)	75.2 (27.0 to 96.1)				
	Yes	19517	15.5	6.0 (1.4 to 22.0)	2.4 (0.4 to 13.5)	58.9 (13.8 to 92.8)				0.00
Hospitalization	No	106005	84.5	94.0 (78.0 to 98.6)	97.6 (86.4 to 99.6)	41.1 (7.2 to 86.2)				

(Continued)

(Continued)

Variable	Categories	Total		Class 1		Class 2		Class 3		p value ^b
		N	%	%	%	%	%	%		
Presence of symptoms	Fatigue or tiredness	83739	66.7	42.7 (25.3 to 62.0)	73.1 (54.1 to 86.2)	98.7 (87.4 to 99.9)		0.00		
	Concentration difficulties	31009	24.7	11.0 (4.0 to 26.8)	0	98.1 (83.1 to 99.8)		0.00		
	Memory loss	27834	22.2	4.7 (1.4 to 14.1)	0	97.7 (83.8 to 99.7)		0.00		
	Ageusia/anosmia	23247	18.5	39.5 (24.7 to 56.4)	1.4 (0.3 to 6.3)	13.4 (2.5 to 48.2)		0.00		
	Dyspnea	74583	59.4	0.0 –	95.8 (89.3 to 98.4)	97.8 (87.3 to 99.6)		0.00		
	Chest pain	33124	26.4	10.2 (7.0 to 14.7)	16.0 (6.2 to 35.1)	76.5 (27.5 to 96.5)		0.00		
	Palpitations	29679	23.6	4.7 (1.3 to 16.1)	4.6 (1.4 to 14.4)	95.7 (70.7 to 99.5)		0.00		
	Dizziness	12867	10.2	7.7 (2.9 to 19.0)	1.0 (0.2 to 4.7)	33.0 (5.8 to 79.6)		0.01		
	Menstrual change ^a	7344	10.7	4.2 (0.9 to 18.0)	1.0 (0.1 to 7.1)	60.5 (19.0 to 90.9)		0.00		
	Inability to exercise	35359	28.2	0.0	32.4 (8.9 to 70.0)	71.9 (24.5 to 95.3)		0.01		
Reduction in daily activities	Joint and muscle pain	44749	35.6	29.5 (13.6 to 52.5)	11.6 (4.6 to 26.4)	93.9 (58.9 to 99.4)		0.00		
	Yes, a little	39092	32.4	14.2 (5.7 to 31.2)	54.6 (27.3 to 79.2)	26.5 (4.3 to 74.3)		0.14		
	Yes, a lot	9290	7.7	3.8 (0.8 to 15.9)	7.9 (19.5 to 27.1)	14.5 (3.0 to 48.6)				

^a Only women.

^b Chi2, two sided.

Source: Prepared by the authors of this study.

Table 2. Multinomial logistic regression analysis of classes of symptoms compatible with post-COVID-19 condition.

Variable	Class 2 (vs. Class 1)			Class 3 (vs. Class 1)		
	RRR	p-value	95% CI	RRR	p-value	95% CI
Age	0.95	0.04	0.90 to 0.99	0.92	0.03	0.86 to 0.99
Female	2.97	0.09	0.83 to 10.61	6.88	0.24	0.27 to 174.70
Indigenous affiliation	3.18	0.18	0.56 to 17.95	10.98	0.01	1.74 to 69.44
Higher technical education	0.08	0.00	0.01 to 0.42	0.45	0.49	0.04 to 4.75
Professional/postgraduate	1.34	0.65	0.37 to 4.80	1.51	0.68	0.21 to 11.10
≥2 COVID-19 episodes	1.32	0.72	0.29 to 6.03	3.83	0.18	0.52 to 28.08
Tobacco use	0.93	0.92	0.25 to 3.43	1.13	0.91	0.13 to 9.79
Hospitalization	0.32	0.38	0.02 to 4.13	12.90	0.14	0.44 to 378.20
Diabetes	35.25	0.00	3.57 to 348.00	11.75	0.02	1.44 to 95.84
BMI	0.83	0.00	0.75 to 0.93	0.85	0.08	0.70 to 1.02

RRR: relative rate ratio; CI: confidence interval; BMI: body mass index.

Source: Prepared by the authors of this study.

additionally associated with lower educational attainment and lower BMI, and class 3 with Aboriginal affiliation.

The prevalence of symptoms compatible with post-COVID-19 condition observed in our study was consistent with that reported in an independent systematic review and meta-analysis, which showed a prevalence of 45% regardless of hospitalization status [27].

The identification of distinct latent classes reflects the heterogeneous clinical presentation of post-COVID-19 condition, in which symptoms are grouped according to severity or pathological mechanisms rather than affected organs or systems involved [15,18]. Our findings are consistent with existing literature. The identified phenotypes align with cardiorespiratory, systemic/inflammatory, and neurological classes described in a recent meta-analysis [28], and with a two-year follow-up study of severe COVID-19 cases in Latin America [29]. Class 1 is characterized by low symptom burden, suggesting a milder clinical phenotype. However, its high prevalence, combined with the presence of sensory and musculoskeletal manifestations, may significantly impair quality of life and limit the resumption of routine activities.

These symptoms are frequently associated with low-grade inflammation, endothelial dysfunction, and mitochondrial impairment, leading to fatigue and exercise intolerance [30]. Phenotypes dominated by somatic or sensory symptoms are more common among younger individuals, women, and those without significant comorbidities [31,32]. This aligns with findings by Ito et al. [33] [33] who identified a class characterized by taste and smell disorders, primarily comprising young women with mild acute symptoms.

Cardiorespiratory symptoms, predominantly respiratory in Class 2, are commonly reported [32]. This arises from persistent inflammation, cardiac thrombi, endothelial dysfunction, autoimmunity, or viral persistence [30]. Such symptoms are also associated with pre-existing chronic respiratory disease [31–34]. Although this study found no statistical inter-class differences in chronic respiratory comorbidities, a higher proportion of affected individuals was observed in Class 2 -characterized by frequency of dyspnea- compared to the other two classes.

Class 3, characterized by the highest symptom burden with predominantly neurological and cardiorespiratory manifestations, exhibits multisystemic involvement associated with significant functional impairment. This is probably attributable to neuroinflammation and autonomic dysfunction [30], consistent with findings reported by Arango-Ibañez et al. [29]. This class showed statistical differences with a higher proportion of individuals with Aboriginal affiliation and diabetes. Ethnic minority status has been identified as a risk factor for post-COVID-19 condition, with increased likelihood of cardiopulmonary symptoms as observed in our Class 3 [35].

Hospitalized COVID-19 survivors demonstrated higher post-COVID-19 risk than non-hospitalized patients [36]. Hospitalization severity further increases post-COVID-19 risk [37], although this association may be confounded with Post-Intensive Care Unit Syndrome [38]. In contrast, hospitalization was significantly associated with class membership in the bivariate analysis, but lost significance in the multivariate model once other severity-related variables (e.g., diabetes, number of COVID-19 episodes) were included, suggesting that its apparent effect was largely confounded by these factors rather than reflecting an independent association.

Class 3 had a higher proportion of individuals with two or more COVID-19 infections compared to other classes. This aligns with disease pathophysiology, as damage increases with repeated acute infection. Babalola et al., in a retrospective study of 2,511 essential workers, demonstrated that repeated SARS-CoV-2 infections, combined with a reduced viral clearance capacity, may promote viral persistence and contribute to post-COVID-19 syndrome development [39]. A similar pattern emerged in bivariate analysis, but the association did not persist after multivariable adjustment, indicating that repeated infections are not an independent predictor of symptom phenotype.

While Classes 1 and 2 showed a higher proportion of individuals with high school or less education, Class 3 had the lowest proportion with professional education. However, no existing studies explicitly explain this educational disparity within post-COVID-19 symptoms classes. Some evidence

suggests reduced family income may contribute to persistent COVID-19 symptoms [40].

Comorbidities are associated with increased risk of post-COVID-19 symptoms. A meta-analysis of 34 studies showed that pre-existing conditions, including anxiety and/or depression, asthma, chronic obstructive pulmonary disease, diabetes, ischemic heart disease, and immunosuppression, significantly increased post-COVID-19 risk [41]. Similarly, Falsetti et al. showed that a higher Charlson Comorbidity Index score is associated with greater likelihood of developing post-COVID-19 [42]. Diabetes has been particularly associated with post-COVID-19 symptoms [43]. The underlying mechanisms may involve weakened immune response or proinflammatory state that, combined with virus-triggered inflammation, promote prolonged responses to SARS-CoV-2 [44,45]. In our study, diabetes was independently associated with classes 2 and 3, relative to class 1, on multivariable analysis. While one study suggests this association reflects higher diabetes prevalence in older population [33] we found no significant differences in diabetes prevalence across age groups.

Although no significant differences were found between classes, Class 3 (high cardiorespiratory and neurological symptom burden) showed higher male proportions, contrary to previous reports [18]. Women were more prevalent in the two most common classes with less severe symptoms, while the most severe class had the highest proportion of people with comorbidities. Psychological and psychosomatic factors critically influence long COVID symptoms persistence, potentially acting as triggers and highlighting the need for a multidisciplinary approach. These factors are predominantly associated with female sex, though this study did not specifically assess these pathologies; however, previous studies have established this link [46,47]. No age differences were observed; specifically, Class 3 did not contain higher elderly proportion. Other studies similarly found no association between symptom prevalence and older ages [29,48].

Vaccination showed no classes differences, consistent with similar study [49]. This may reflect our categorization approach (first dose/booster versus bivalent/Omicron) rather than vaccination status (yes/no). Categorizing vaccinated versus unvaccinated was impossible given the high national vaccination coverage (98,3%). Two systematic reviews indicate inconclusive evidence; however, studies suggest pre-infection vaccination may reduce post-COVID-19 risk [50,51].

Although smoking increases the post-COVID-19 symptoms [52], tobacco use showed no significant differences among classes; however, the prevalence of smokers in Class 3 was 2.5 and 3.5 times higher than in Class 2 and Class 1, respectively. A possible explanation is that smoking affects symptom severity through pathophysiological pathway rather than creating symptom patterns.

Excess weight is generally associated with post-COVID-19 symptoms. A recent systematic review and meta-analysis found obesity most strongly associated with smell disorder (OR = 1.16;

95% CI: 1.11 to 1.22) and taste disorder (OR = 1.22; 95% CI: 1.08 to 1.38) [53]. In our study, anosmia/ageusia was a defining feature of class 1, which may partly explain why lower, rather than higher, BMI was associated with class 2 membership. However, since BMI was self-reported, potential misclassification cannot be ruled out.

Post-COVID-19 symptoms occur in severity-based classes rather than by symptom number or system, requiring differentiated interventions. This research guides personalized therapeutic strategies. The healthcare system should develop targeted interventions for each class, with priority given to individuals reporting substantial quality of life impairment, independent of hospitalization status during acute COVID-19 infection. In Chile, *Garantías Explicitas de Salud* (Explicit Health Guarantees) cover post-COVID-19 diagnosis and treatment only for patients who were hospitalized during the acute infection [54]. Although only 15% of COVID-19-positive individuals overall were hospitalized, 60% of Class 3 patients were, yet the remaining 40%, despite similar severity, went without guaranteed care.

This is the first population-based investigation to assess post-COVID-19 condition symptoms in Chile and provide evidence from Latin America, a region frequently underrepresented in scientific literature. Symptoms class analysis enables the identification of patterns rather than simply counting. While deterministic methods are commonly used for their simplicity and transparency, LCA offers greater flexibility, adopting probabilistic framework to uncover latent patterns. This flexibility comes at a cost: class number is not fully data-determined, and larger future studies should assess phenotype reproducibility.

This study has limitations. Its cross-sectional design precludes causal inference and assessment of symptom progression. Recall bias is an inherent limitation, as symptoms, prior infections, and their timing were self-reported—often months to years after the acute episode—which may have led to underreporting or symptom misattribution and affected the accuracy of prevalence estimates. Symptoms lacked clinical validation and adjustment for previous history. Most participants neither required hospitalization nor experienced severe COVID-19. Focusing on survivors introduces survival bias. Older adults are likely underrepresented, given the high mortality in this group. Symptom severity was not directly measured, though daily activity reduction was used as a proxy. BMI was self-reported, which may have introduced misclassification bias. The exclusion of rural areas limits generalizability to Chile's urban context. Longitudinal studies are needed to address these limitations. We acknowledge potential barriers to participation in this study among the most marginalized groups (homeless, undocumented migrants, institutionalized populations). Future studies should employ targeted outreach strategies to ensure full inclusion of historically excluded populations. As the sample was not originally designed for post-COVID-19 symptoms prevalence estimation or latent class analysis, precision and class stability may be limited; however, the achieved sample

size supports exploratory analysis, and the 100% questionnaire response rate—using expansion factors from the seroprevalence study's design—likely limits this impact. Data on time since infection, viral variant, reinfection history, mental health, and post-ICU syndrome were not systematically collected, potentially contributing to residual confounding in symptom heterogeneity; future studies should incorporate these factors for a more comprehensive understanding.

CONCLUSIONS

There is a substantial burden of post-COVID-19 conditions among individuals who experienced COVID-19. A three-class structure offers the clearest clinical interpretation among the models tested, though this classification should be treated as exploratory rather than definitive. These three symptom classes identified may guide clinical diagnosis and treatment. Class 1 (38.4%) includes individuals with fatigue, ageusia, and musculoskeletal pain (physical/inflammatory phenotype). Class 2 (40.7%) was characterized by predominantly dyspnea, cardiorespiratory symptoms, and fatigue. Class 3 (20%), though less prevalent, involves multisystemic phenotype symptoms and a high frequency of reduced activities of daily living. This more severe class was associated with diabetes and Aboriginal affiliation in the adjusted multivariable analysis. Care strategies should prioritize individuals whose daily activities are impaired, with attention to factors linked to the most severe symptom class. Regardless of other factors, diabetes emerged as a variable of interest across class 2 and class 3.

Author contributions Conceptualization, LNF and MRS; Methodology, LNF, MRS, PR and MA; Software, PR and MA; Validation, PR and MA; Formal Analysis, PR and MA; Investigation, LNF, MRS and LC; Resources, LNF and MRS; Data Curation, PR and MA; Writing – Original Draft Preparation, PR, LNF and MRS; Writing – Review & Editing, LNF, MRS, PR and MA; Visualization, MA, PR, LNF and MRS; Supervision, LNF and MRS; Project Administration, LNF and MRS; Funding Acquisition, LNF. All authors reviewed and approved the final version of the manuscript for publication.

Acknowledgements The authors acknowledge the institutional support for this research provided by Universidad de Talca, Universidad Católica del Norte, and Universidad del Desarrollo. We thank the field work team for their assistance in the data collection and participants recruitment. We also extend our gratitude to the laboratory personnel in both cities for their technical support throughout the study. Finally, we are grateful to all study participants who generously shared their time and experience, making this research possible.

Competing interests The authors have no competing interests to declare that are relevant to the content of this article.

Funding The Chilean National Research Agency supported this work, Grant code FONIS SA23I0063 and Anillo-ATE 220061. The Chilean Ministry of Health provided nasal swab tests to detect SARS-CoV-2 antigens.

Declaration of artificial intelligence model use Artificial intelligence tools (ChatGPT-5, OpenAI) were used to revise the manuscript's English grammar. The content was fully reviewed and verified by the authors.

Ethics The study was conducted in accordance with the Declaration of Helsinki, and the study protocol was reviewed and approved by

three independent Scientific Ethical Committees: 1) Scientific ethical committee from the Faculty of Medicine, Universidad Católica del Norte, Resolution number 63/2023, dated October 16th, 2023. 2) Scientific ethical committee from the Faculty of Medicine, Universidad del Desarrollo, dated December 13th, 2023, and 3) Scientific ethical committee from Universidad de Talca, Folio 30-2023-E, dated April 17th, 2024. Additionally, it was approved by the Institutional Committee of Biosecurity of Universidad Católica del Norte 07/2023 dated October 2023, Universidad del Desarrollo, CIB-FORM-01B, dated 14 November 2023, and Universidad de Talca 20-CBS-2023 dated November 9th, 2023.

Data availability statement The datasets used and/or analyzed during the current study are available from the corresponding author on reasonable request. Code availability: <https://doi.org/10.5281/zenodo.1776466>

Language of submission English.

Peer review and provenance Not commissioned. Externally peer-reviewed by two reviewers. Reviewed by the journal's statistical editor. All reviews were conducted in a double-anonymous modality.

REFERENCES

- Burton C, Dawes H, Goodwill S, Thelwell M, Dalton C. Within and between-day variation and associations of symptoms in Long Covid: Intensive longitudinal study. *PLoS ONE*. 2023;18. <https://doi.org/10.1371/journal.pone.0280343>
- Haunhorst S, Bloch W, Wagner H, Ellert C, Krüger K, Vilser DC, et al. Long COVID: a narrative review of the clinical aftermaths of COVID-19 with a focus on the putative pathophysiology and aspects of physical activity. *Oxf Open Immunol*. 2022;3. <https://doi.org/10.1093/oxfimm/iqac006>
- Katsarou M-S, Iasonidou E, Osarogue A, Kalafatis E, Stefanatou M, Pappa S, et al. The Greek Collaborative Long COVID Study: Non-Hospitalized and Hospitalized Patients Share Similar Symptom Patterns. *J Pers Med*. 2022;12. <https://doi.org/10.3390/jpm12060987>
- Saricaoglu EM, Cinar G, Azap A, Bayar MK, Tokay-Isıkay C, Kutlayacaksin S, et al. Dark Side of the COVID-19 Pandemic; 'Long COVID'. *Infect Dis Clin Microbiol*. 2023;5: 205–211. <https://www.idcmjournal.org/issue/september-2023> <https://doi.org/10.36519/idcm.2023.213>
- Wulf Hanson S, Abbafati C, Aerts JG, Al-Aly Z, Ashbaugh C, Ballouz T, et al. Estimated Global Proportions of Individuals With Persistent Fatigue, Cognitive, and Respiratory Symptom Clusters Following Symptomatic COVID-19 in 2020 and 2021. *JAMA*. 2022;328: 1604–1615. <https://doi.org/10.1001/jama.2022.18931>
- World Health Organization. Post COVID-19 condition (long COVID) Newsroom Fact sheet. In: [https://www.who.int/newsroom/fact-sheets/detail/post-covid-19-condition-\(long-covid\)](https://www.who.int/newsroom/fact-sheets/detail/post-covid-19-condition-(long-covid)). 2025.
- France K, Glick M. Long COVID and oral health care considerations. *J Am Dent Assoc*. 2022;153: 167–174. <https://doi.org/10.1016/j.adaj.2021.08.007>
- Kogevinas M, Karachaliou M, Espinosa A, Iraola-Guzmán S, Castaño-Vinyals G, Delgado-Ortiz L, et al. Risk, determinants, and persistence of long-COVID in a population-based cohort

- study in Catalonia. *BMC Med.* 2025;23. <https://doi.org/10.1186/s12916-025-03974-7>
9. Baratta JM, Tomparay A, Siano S, Floris-Moore M, Weber DJ. Postacute Sequelae of COVID-19 Infection and Development of a Physiatry-Led Recovery Clinic. *Am J Phys Med Rehabil.* 2021;100: 633–634. <https://doi.org/10.1097/PHM.0000000000001778>
10. Rigoni M, Torri E, Nollo G, Donne LD, Rizzardo S, Lenzi L, et al. In: Long COVID" follow-up pathway: results of longitudinal cohort study to investigate 6- and 12-month outcomes after hospitalization for SARS-CoV-2 infection [Internet]. 2021. <http://europepmc.org/abstract/PPR/PPR397467%0Ahttps://doi.org/10.21203/rs.3.rs-908078/v1%0Ahttps://europepmc.org/article/PPR/PPR397467>
11. Busatto GF, de Araujo AL, Castaldelli-Maia JM, Damiano RF, Imamura M, Guedes BF, et al. Post-acute sequelae of SARS-CoV-2 infection: relationship of central nervous system manifestations with physical disability and systemic inflammation. *Psychol Med.* 2022;52: 2387–2398. <https://doi.org/10.1017/S0033291722001374>
12. Zhang H, Zang C, Xu Z, Zhang Y, Xu J, Bian J, et al. Data-driven identification of post-acute SARS-CoV-2 infection subphenotypes. *Nat Med.* 2023;29: 226–235. <https://doi.org/10.1038/s41591-022-02116-3>
13. Zhang H, Zang C, Xu Z, Zhang Y, Xu J, Bian J, et al. Machine Learning for Identifying Data-Driven Subphenotypes of Incident Post-Acute SARS-CoV-2 Infection Conditions with Large Scale Electronic Health Records: Findings from the RECOVER Initiative. *medRxiv.* 2022; 05. <https://doi.org/10.1101/2022.05.21.22275412>
14. Frontera JA, Thorpe LE, Simon NM, de Havenon A, Yaghi S, Sabadia SB, et al. Post-acute sequelae of COVID-19 symptom phenotypes and therapeutic strategies: A prospective, observational study. *PLoS ONE.* 2022;17. <https://doi.org/10.1371/journal.pone.0275274>
15. Ozonoff A, Jayavelu ND, Liu S, Melamed E, Milliren CE, Qi J, et al. Features of acute COVID-19 associated with post-acute sequelae of SARS-CoV-2 phenotypes: results from the IMPACC study. *Nat Commun.* 2024;15. <https://doi.org/10.1038/s41467-023-44090-5>
16. Kenny G, McCann K, O'Brien C, Savinelli S, Tinago W, Yousif O, et al. Identification of Distinct Long COVID Clinical Phenotypes Through Cluster Analysis of Self-Reported Symptoms. *Open Forum Infect Dis.* 2022;9. <https://doi.org/10.1093/ofid/ofac060>
17. Huang Y, Pinto MD, Borelli JL, Asgari Mehrabadi M, Abraham HL, Dutt N, et al. COVID Symptoms, Symptom Clusters, and Predictors for Becoming a Long-Hauler Looking for Clarity in the Haze of the Pandemic. *Clin Nurs Res.* 2022;31: 1390–1398. <https://doi.org/10.1177/10547738221125632>
18. Sun X, DeShazo JP, Anatole-Tardiff L, Di Fusco M, Allen KE, Porter TM, et al. Latent class analysis of post-acute sequelae of SARS-CoV-2 infection. *J Biopharm Stat.* 2025;35: 902–917. <https://doi.org/10.1080/10543406.2024.2424844>
19. O'Neil ST, Madlock-Brown C, Wilkins KJ, McGrath BM, Davis HE, Assaf GS, et al. Finding Long-COVID: temporal topic modeling of electronic health records from the N3C and RECOVER programs. *NPJ Digit Med.* 2024;7: 296. <https://doi.org/10.1038/s41746-024-01286-3>
20. Danesh V, Arroliga AC, Bourgeois JA, Boehm LM, McNeal MJ, Widmer AJ, et al. Symptom Clusters Seen in Adult COVID-19 Recovery Clinic Care Seekers. *J Gen Intern Med.* 2023;38: 442–449. <https://doi.org/10.1007/s11606-022-07908-4>
21. Vial P, González C, Icaza G, Ramírez-Santana M, Quezada-Gaete R, Núñez-Franz L, et al. Seroprevalence, spatial distribution, and social determinants of SARS-CoV-2 in three urban centers of Chile. *BMC Infect Dis.* 2022;22: 99. <https://doi.org/10.1186/s12879-022-07045-7>
22. Núñez-Franz L, Rubilar P, Apablaza M, Canales L, Cortés LJ, Molina X, et al. Population-based seroprevalence survey: post-pandemic COVID-19 vaccination, related factors, and geographic distribution of vaccine acceptability in Chile. *BMC Public Health.* 2025;25. <https://doi.org/10.1186/s12889-025-22314-1>
23. Núñez-Franz L, Ramírez-Santana M, Rubilar P, Vial C, Apablaza M, González C, et al. Seroprevalence of Natural and Acquired Immunity against the SARS-CoV-2 Virus in a Population Cohort from Two Chilean Cities, 2020–2022. *Viruses.* 2023;15: 2020–2022. <https://doi.org/10.3390/v15010201>
24. Aguilera X, González C, Apablaza M, Rubilar P, Icaza G, Ramírez-Santana M, et al. Immunization and SARS-CoV-2 Antibody Seroprevalence in a Country with High Vaccination Coverage: Lessons from Chile. *Vaccines (Basel).* 2022;10: 1002. <https://doi.org/10.3390/vaccines10071002>
25. National Center for Health Statistic. In: Long COVID Household Pulse Survey [Internet]. 2024. https://www.cdc.gov/nchs/covid19/pulse/long-covid.htm#technical_notes
26. Bailly S, Dey T. Latent Class Analysis to Identify Subgroups in Clinical Research: A Data-Driven Approach. *Chest.* 2026;169: 37–40. <https://doi.org/10.1016/j.chest.2025.07.4085>
27. O'Mahoney LL, Routen A, Gillies C, Ekezie W, Welford A, Zhang A, et al. The prevalence and long-term health effects of Long Covid among hospitalised and non-hospitalised populations: A systematic review and meta-analysis. *EClinicalMedicine.* 2023;55: 101762. <https://doi.org/10.1016/j.eclinm.2022.101762>
28. Kuodi P, Gorelik Y, Gausi B, Bernstine T, Edelstein M. Characterization of post-COVID syndromes by symptom cluster and time period up to 12 months post-infection: A systematic review and meta-analysis. *Int J Infect Dis.* 2023;134: 1–7. <https://doi.org/10.1016/j.ijid.2023.05.003>
29. Arango-Ibanez JP, Córdoba-Melo BD, Gutiérrez Posso JM, Barbosa-Rengifo MM, Herrera CJ, Quintana Da Silva MA, et al. Long COVID Clusters of Symptoms Persist beyond Two Years after Infection: Insights from the CARDIO COVID 20-21 Registry. *Viruses.* 2024;16. <https://doi.org/10.3390/v16071028>
30. Li J, Zhou Y, Ma J, Zhang Q, Shao J, Liang S, et al. The long-term health outcomes, pathophysiological mechanisms and

- multidisciplinary management of long COVID. *Sig Transduct Target Ther.* 2023;8. <https://doi.org/10.1038/s41392-023-01640-z>
31. Escoda T, Chiche L, Faralli H, Cohen F, Halfon P, Pegliasco H, et al. Cluster analysis of post-COVID-19 physical and mental health outcomes 3-6 months after SARS-CoV-2 infection: results of the French Prospective ALCOVID Cohort Study. *BMJ Open.* 2025;15. <https://doi.org/10.1136/bmjopen-2024-089136>
 32. Yelin D, Margalit I, Nehme M, Bordas-Martínez J, Pistelli F, Yahav D, et al. Patterns of Long COVID Symptoms: A Multi-Center Cross Sectional Study. *JCM.* 2022;11: 898. <https://doi.org/10.3390/jcm11040898>
 33. Ito F, Terai H, Kondo M, Takemura R, Namkoong H, Asakura T, et al. Cluster analysis of long COVID in Japan and association of its trajectory of symptoms and quality of life. *BMJ Open Resp Res.* 2024;11: e002111. <https://doi.org/10.1136/bmjresp-2023-002111>
 34. Gentilotti E, Górka A, Tami A, Gusinow R, Mirandola M, Rodríguez Baño J, et al. Clinical phenotypes and quality of life to define post-COVID-19 syndrome: a cluster analysis of the multinational, prospective ORCHESTRA cohort. *EClinicalMedicine.* 2023;62: 102107. <https://doi.org/10.1016/j.eclinm.2023.102107>
 35. Mkoma GF, Agyemang C, Benfield T, Rostila M, Cederström A, Petersen JH, et al. Risk of long COVID and associated symptoms after acute SARS-COV-2 infection in ethnic minorities: A nationwide register-linked cohort study in Denmark. *PLoS Med.* 2024;21. <https://doi.org/10.1371/journal.pmed.1004280>
 36. Yuan N, Lv Z-H, Sun C-R, Wen Y-Y, Tao T-Y, Qian D, et al. Post-acute COVID-19 symptom risk in hospitalized and non-hospitalized COVID-19 survivors: A systematic review and meta-analysis. *Front Public Health.* 2023;11. <https://doi.org/10.3389/fpubh.2023.1112383>
 37. Huang C, Huang L, Wang Y, Li X, Ren L, Gu X, et al. 6-month consequences of COVID-19 in patients discharged from hospital: a cohort study. *The Lancet.* 2023;401: e21–e33. [https://doi.org/10.1016/S0140-6736\(23\)00810-3](https://doi.org/10.1016/S0140-6736(23)00810-3)
 38. Vrettou CS, Jolley SE, Mantziou V, Dimopoulou I. Clinical Comparison of Post-intensive Care Syndrome and Long Coronavirus Disease. *Crit Care Clin.* 2025;41: 89–102. <https://doi.org/10.1016/j.ccc.2024.08.009>
 39. Babalola TK, Clouston SAP, Sekendiz Z, Chowdhury D, Soriolo N, Kawuki J, et al. SARS-COV-2 re-infection and incidence of post-acute sequelae of COVID-19 (PASC) among essential workers in New York: a retrospective cohort study. *The Lancet Regional Health - Americas.* 2025;42: 100984. <https://doi.org/10.1016/j.lana.2024.100984>
 40. Ziauddeen N, Gurdasani D, O'Hara ME, Hastie C, Roderick P, Yao G, et al. Characteristics and impact of Long Covid: Findings from an online survey. *PLoS ONE.* 2022;17: e0264331. <https://doi.org/10.1371/journal.pone.0264331>
 41. Tsampasian V, Elghazaly H, Chattopadhyay R, Debski M, Naing TKP, Garg P, et al. Risk Factors Associated With Post-COVID-19 Condition: A Systematic Review and Meta-analysis. *JAMA Intern Med.* 2023;183: 566–580. <https://doi.org/10.1001/jamainternmed.2023.0750>
 42. Falsetti L, Zacccone V, Santoro L, Santini S, Guerrieri E, Giuliani L, et al. The Relationship between Post-COVID Syndrome and the Burden of Comorbidities Assessed Using the Charlson Comorbidity Index. *Medicina (B Aires).* 2023;59: 1583. <https://doi.org/10.3390/medicina59091583>
 43. Su Y, Yuan D, Chen DG, Ng RH, Wang K, Choi J, et al. Multiple early factors anticipate post-acute COVID-19 sequelae. *Cell.* 2022;185: 881–895. <https://doi.org/10.1016/j.cell.2022.01.014>
 44. Buetti N, Trimboli P, Mazzuchelli T, Lo Priore E, Balmelli C, Trkola A, et al. Diabetes mellitus is a risk factor for prolonged SARS-CoV-2 viral shedding in lower respiratory tract samples of critically ill patients. *Endocrine.* 2020;70: 454–460. <https://doi.org/10.1007/s12020-020-02465-4>
 45. Rizvi A, Ziv Y, Crawford JM, Trindade AJ. Gastrointestinal and Hepatobiliary Symptoms and Disorders with Long (Chronic) COVID Infection. *Gastroenterol Clin North Am.* 2023;52: 139–156. <https://doi.org/10.1016/j.gtc.2022.09.002>
 46. Fleischer NL, Roux AVD. Using directed acyclic graphs to guide analyses of neighbourhood health effects: an introduction. *J Epidemiol Community Health.* 2008;62: 842–846. <https://doi.org/10.1136/jech.2007.067371>
 47. Engelmann P, Reinke M, Stein C, Salzmann S, Löwe B, Toussaint A, et al. Psychological factors associated with Long COVID: a systematic review and meta-analysis. *EClinicalMedicine.* 2024;74. <https://doi.org/10.1016/j.eclinm.2024.102756>
 48. Hirschtick JL, Xie Y, Slocum E, Hirschtick RE, Power LE, Elliott MR, et al. A statewide population-based approach to examining Long COVID symptom prevalence and predictors in Michigan. *Prev Med.* 2023;177: 107752. <https://doi.org/10.1016/j.ypmed.2023.107752>
 49. Tawfiq E, Chen R, Honeyman DA, Dawson R, Kunasekaran M, Notaras A, et al. Long Covid Symptom Clusters, Correlates and Predictors in a Highly Vaccinated Australian Population in 2023. *Health Expect.* 2025;28. <https://doi.org/10.1111/hex.70273>
 50. Notarte KI, Catahay JA, Velasco JV, Pastrana A, Ver AT, Pangilinan FC, et al. Impact of COVID-19 vaccination on the risk of developing long-COVID and on existing long-COVID symptoms: A systematic review. *eClinicalMedicine.* 2022;53: 101624. <https://doi.org/10.1016/j.eclinm.2022.101624>
 51. Byambasuren O, Stehlik P, Clark J, Alcorn K, Glasziou P. Effect of covid-19 vaccination on long covid: systematic review. *BMJ Med.* 2023;2. <https://doi.org/10.1136/bmjmed-2022-000385>
 52. Trofor AC, Robu Popa D, Melinte OE, Trofor L, Vicol C, Grosu-Creangă IA, et al. Looking at the Data on Smoking and Post-COVID-19 Syndrome-A Literature Review. *J Pers Med.* 2024;14. <https://doi.org/10.3390/jpm14010097>

53. Ronca DB, Mesquita LO, Oliveira D, Figueiredo ACMG, Wen J, Song M, et al. Excess weight is associated with neurological and neuropsychiatric symptoms in post-COVID-19 condition: A systematic review and meta-analysis. PLoS One. 2025;20: e0314892. <https://doi.org/10.1371/journal.pone.0314892>
54. In: Problemas de Salud - AUGE - Ministerio de Salud [Internet]. Sep 2025. <https://auge.minsal.cl/problemasdesalud/index/87>

Prevalencia de síntomas post-COVID, agrupamiento sintomático e impacto funcional: Estudio transversal de base poblacional en dos ciudades chilenas, 2024

RESUMEN

INTRODUCCIÓN La condición post-COVID-19 afecta aproximadamente al 6% de las personas a nivel mundial. La mayor parte de la evidencia previa se basa en pacientes hospitalizados y proviene de países con una cobertura de vacunación relativamente menor. Este estudio estima la prevalencia de síntomas compatibles con la condición post-COVID-19 y explora patrones sintomáticos en una muestra poblacional de Chile, un país con alta cobertura de vacunación, y examina la asociación con las actividades diarias y factores demográficos, sociales y clínicos.

MÉTODOS Se realizó un estudio transversal en mayo de 2024 en dos ciudades chilenas, con la participación de 654 individuos (tasa de respuesta: 88,5%) de ≥ 7 años, provenientes de hogares seleccionados aleatoriamente. Se recopilaban diez síntomas post-COVID-19 persistentes durante ≥ 3 meses. Se utilizó análisis de clases latentes para identificar patrones sintomáticos.

RESULTADOS Entre 277 participantes con diagnóstico confirmado de COVID-19, 114 reportaron al menos un síntoma con duración ≥ 3 meses. La prevalencia ponderada de al menos un síntoma post-COVID-19 fue de 17,4% en la población general y de 40,7% entre quienes habían tenido COVID-19. Entre aquellos con síntomas post-COVID-19, el 40,1% reportó una reducción en sus actividades diarias. Se exploraron tres clases: la clase 1 (38,4%), caracterizado por fatiga, ageusia/anosmia y dolor muscular/articular; la clase 2 (40,7%), con predominio de síntomas cardiorrespiratorios; y la clase 3 (20,9%), con síntomas multisistémicos. La clase 3 se asoció con mayor frecuencia de comorbilidades (particularmente diabetes), hospitalización, múltiples episodios de COVID-19 y pertenencia a pueblos originarios. Las clases 2 y 3 se asociaron de forma independiente con menor edad y diabetes; el clúster 2 se asoció además con menor nivel educativo y menor IMC, y el clúster 3 con afiliación aborigen.

CONCLUSIONES Este estudio poblacional evidencia una carga sustancial de al menos un síntoma post-COVID-19, con patrones sintomáticos diferenciados que enfatizan la necesidad de intervenciones específicas. Estos hallazgos subrayan la importancia de políticas públicas dirigidas y enfoques clínicos personalizados, especialmente para los conglomerados de mayor gravedad que afectan la calidad de vida, y aportan evidencia relevante desde América Latina a la comprensión global de la condición post-COVID-19.



This work is licensed under a Creative Commons Attribution 4.0 International License.