

Article

Not peer-reviewed version

Long COVID and Associated Factors Among Chinese Residents Aged 16 Years and Older in Canada: A Cross-Sectional Online Study

[Matin Shariati](#) , [Kieran Luke Gill](#) , Mark Peddle , [Ying Cao](#) , Fangli Xie , Xiao Han , Nan Lei , [Rachel Prowse](#) , [Desai Shan](#) , [Lisa Fang](#) , [Vita Huang](#) , [Arianna Ding](#) , [Peizhong Peter Wang](#) *

Posted Date: 21 January 2025

doi: [10.20944/preprints202501.1534.v1](https://doi.org/10.20944/preprints202501.1534.v1)

Keywords: COVID-19; long COVID; immigrant health; long COVID risk factors; Chinese immigrants; Canada; LASSO regression



Preprints.org is a free multidisciplinary platform providing preprint service that is dedicated to making early versions of research outputs permanently available and citable. Preprints posted at Preprints.org appear in Web of Science, Crossref, Google Scholar, Scilit, Europe PMC.

Copyright: This open access article is published under a Creative Commons CC BY 4.0 license, which permit the free download, distribution, and reuse, provided that the author and preprint are cited in any reuse.

Disclaimer/Publisher's Note: The statements, opinions, and data contained in all publications are solely those of the individual author(s) and contributor(s) and not of MDPI and/or the editor(s). MDPI and/or the editor(s) disclaim responsibility for any injury to people or property resulting from any ideas, methods, instructions, or products referred to in the content.

Article

Long COVID and Associated Factors Among Chinese Residents Aged 16 Years and Older in Canada: A Cross-Sectional Online Study

Matin Shariati ^{1,†}, Kieran Gill ^{1,†}, Mark Peddle ¹, Ying Cao ², Fangli Xie ², Xiao Han ², Nan Lei ^{1,2}, Rachel Prowse ¹, Desai Shan ¹, Lisa Fang ², Vita Huang ², Arianna Ding ² and Peizhong Peter Wang ^{1,2,3,*}

¹ Division of Population Health and Applied Health Sciences, Faculty of Medicine, Memorial University, 300 Prince Philip Drive, St. John's, Newfoundland and Labrador, Canada, A1B 3V6

² Centre for New Immigrant Well-Being (CNIW), 96 Scarsdale Road, Toronto, Ontario, M3B 2R7

³ Dalla Lana School of Public Health, University of Toronto, 155 College Street, Room 534, Toronto, Ontario, M5T 3M7

* Correspondence: pwang@mun.ca

† Kieran Gill and Matin Shariati made equal contributions.

Abstract: As the COVID-19 pandemic evolved, long COVID emerged as a significant threat to public health, characterized by one or more persistent symptoms impacting organ systems beyond 12 weeks of infection [9–12]. Informative research has been derived from assessments of long COVID among the Chinese populace. However, none of these studies considered the COVID-19 experience of Chinese residents in Canada. **Objectives:** We aimed to fill this literature gap by delineating the long COVID experience, prevalence, and associated factors among a sample of Chinese residing in Canada during the pandemic. **Methods:** The present study employed a cross-sectional online survey questionnaire distributed to a sample of Canadian Chinese using a convenience sampling procedure from December 22, 2022, to February 15, 2023. Respondents were probed for sociodemographic background, health-, COVID-, and vaccine-related characteristics. Logistic LASSO regression was used for model building and multivariate logistic regression was used to identify factors associated with developing long COVID. **Results:** Among 491 eligible participants, 63 (12.83%) reported experiencing long COVID with a mean duration of 5.31 (95% CI: 4.06–6.57) months, and major symptoms including difficulty concentrating (21.67%), pain/discomfort (15.00%), as well as anxiety/depression (8.33%). Our final model identified significant associations between long COVID and two or more COVID-19 infections ($OR = 23.725$, 95% CI: 5.098–110.398, $p < 0.0001$), very severe/severe symptoms ($OR = 3.177$, 95% CI: 1.160–8.702, $p = 0.0246$), over-the-counter medicine ($OR = 2.473$, 95% CI: 1.035–5.909, $p = 0.0416$), and traditional Chinese medicine ($OR = 8.259$, 95% CI: 3.016–22.620, $p < 0.0001$). Further, we identified a significant protective effect of very good/good health status ($OR = 0.247$, 95% CI: 0.112–0.544, $p = 0.0005$). **Conclusions:** Long COVID adversely effected a notable proportion of Canadian Chinese for a prolonged period during the pandemic. Our findings underscore the importance of preexisting health status and reinfection prevention when managing long COVID. Moreover, our work indicates the need for culturally accessible guidance and services pertaining to effective COVID-19 treatment modalities through infection, recovery, and beyond.

Keywords: COVID-19; long COVID; immigrant health; long COVID risk factors; Chinese immigrants; Canada; LASSO regression

1. Introduction

In the months after the first case was detected during December 2019, coronavirus disease 2019 (COVID-19) evolved into an unprecedented pandemic significantly impacting health outcomes,

societies, and economies worldwide [1]. According to the World Health Organization (WHO), as of December 8, 2024, over 777 million people have been infected, and more than 7 million people have died globally [2]. During this period Canada reported around 4.8 million confirmed COVID-19 cases and more than 55,000 deaths. Within Canada, the Greater Toronto Area's (GTA) first coronavirus cases came from mainland China in January 2020, as did most COVID-19 infections during the early pandemic [3]. According to 2021 census records the population of Canadian Chinese residents was around 1.7 million, or approximately 4.7% of the total population [4]. This makes Canadian Chinese one of the largest immigrant populations in Canada. Notably, compared to other Canadians, Chinese immigrants have more frequent and intimate relationships with China, where the first cases of COVID-19 were detected, making them uniquely susceptible to infection as COVID-19 spread around the world [5].

COVID-19 targets respiratory systems, but research indicates other organs are also affected. Symptoms associated with lower respiratory tract infection, including fever, dry cough, and difficulty breathing, were reported in the first case series in Wuhan, China [6]. Further, headache, dizziness, weakness, vomiting, and diarrhea have also been reported [7]. Major complications reported in patients with COVID-19 may also include coagulopathy, laryngeal edema, laryngitis, necrotizing pneumonia, acute respiratory failure, ventilation-associated pneumonia, massive pulmonary embolism, sepsis, and a higher risk of mortality [8]. Additionally, long COVID may occur after experiencing a wide variety of mild to severe COVID-19 symptoms [9]. Long COVID is a symptom that continues or develops beyond 12 weeks from an acute COVID-19 infection, with no explanation from any other diagnosis [10]. Studies have shown respiratory, cardiovascular, neurological, gastrointestinal, and musculoskeletal systems are affected by long COVID [9,11,12]. The most common symptoms include fatigue, cough, sore throat, dyspnea, cardiac abnormalities, sleep turbulences, anxiety, post-traumatic stress disorder, myalgia, arthralgia, cognitive impairment, concentration problems, and headaches [9].

Importantly, COVID-19 infection has been shown to induce production of antibodies able to damage proteins, leading to autoimmune cell damage that could persist following initial illness [13]. Further, research has identified additional potential pathophysiological mechanisms underlying long COVID. These investigations highlight the occurrence of hypercoagulation, characterized by the formation of micro-clots and endothelial dysfunction. Concurrently, disruption of cellular energy metabolism has been identified as a contributing factor, ultimately culminating in a hypoxic state [14].

Considering the significant impact of long COVID on individual health and at-risk groups, it is crucial to develop and implement targeted preventative measures to mitigate the risk of developing long COVID, in addition to efforts to prevent initial infection. Numerous risk factors for developing long COVID have been identified in research conducted on the Chinese populace. More specifically, the risk factors most closely linked to long COVID include being female [15–18], experiencing many initial symptoms of COVID-19, having increased levels of D-dimer and C-reactive protein (CRP), having prior psychiatric disorders, engaging in military and transport jobs, smoking, reporting poor self-perceived health status, having chronic diseases or using medication, having early dyspnea, and experiencing critical severity of COVID-19 [19,20].

In combating the health impacts of COVID-19 various vaccines were created as the pandemic evolved. A comparison of long COVID symptoms between unvaccinated and vaccinated individuals has indicated a robust correlation between vaccination and reducing long COVID symptoms [21–23]. That is, vaccination reduced acute COVID-19 infection severity, which reduced long COVID symptoms. However, the consequence of vaccination in people with long COVID symptoms remains unknown and controversial [24]. For example, one study showed the severity of COVID-19 symptoms improved more in those vaccinated with mRNA-type vaccines than those immunized with an adenovirus vector [25]. Similarly, a systematic review study presented that the effect of vaccination on long COVID symptoms was supported in some studies and not in other studies, which could be due to the variation in the type of vaccine received in different studies [24].

Traditional Chinese beliefs about health and illness are distinct from those of Western societies. For instance, one study exploring the health beliefs of older Chinese Australians found they possessed a holistic view of health and the role of food in preventive care and self-medication in times of illness [26]. Another study found some Chinese residents of the United Kingdom prefer the self-treatment approach of traditional Chinese medicine (TCM) or use TCM when they feel Western medicine offers ineffective treatment [27]. Overall, social images of health and illness are closely linked to cultural identity [28]. Chinese have less desire to integrate into Canadian society, which may make them less willing to seek health services.

To date, although long COVID has been explored in numerous countries, considering diverse factors, most studies assessing the COVID-19 experiences of Chinese individuals have been in the Chinese context [20,29,30]. That is, there is a notable gap in research pertaining to the COVID-19 experience of Chinese residents in Canada. Most importantly, no large-scale study has been conducted on Canadian Chinese regarding long COVID [5]. Accordingly, the current study aims to address the following research question: what are the demographic, medical, and behavioral factors (i.e., sociodemographic, COVID-, health-, and vaccine-related factors) associated with the likelihood of developing long COVID among Chinese residents in Canada during the global COVID-19 pandemic? More specifically, this study addresses two specific objectives: (1) to define and contrast the characteristics of a subset of Chinese residents in Canada in relation to their long COVID experience; (2) to evaluate any associations between the characteristics of Canadian Chinese and development of long COVID. Identifying the factors associated with long COVID among Canadian Chinese could help health professionals and policymakers better understand the unique needs and vulnerabilities of East Asian Canadians, and particularly Chinese residents in Canada. Characterizing the long COVID experience of these vulnerable Canadians is important to plan rehabilitative services which support an efficient return to typical daily activities (e.g., social, academic, vocational) [31,32]. Furthermore, such information may contribute foundational knowledge prerequisite for adequate policies, programs, and services supporting rapidly growing Canadian immigrant populations during future communicable disease outbreaks [33].

2. Materials and Methods

2.1. Study Design and Setting

Our cross-sectional online survey questionnaire was distributed among a sample of Canadian Chinese from December 22, 2022, to February 15, 2023. During this period a multifaceted public health response to COVID-19 was still ongoing; prominently including restricted contact between people (e.g., lockdowns), travel restrictions, and widespread vaccination against COVID-19 [34]. All methodologies employed in the current study strictly adhere to all pertinent guidelines and regulations.

To estimate effective sample size, we used the Cochrane formula, considering the total Chinese population in Canada (1.7 million), confidence coefficients of 95%, precision level of 5%, and 10% dropout [35,36]. We estimated 425 for the effective sample size. Only individuals of self-identified Chinese ethnicity residing in Canada were invited to complete the survey questionnaire. This included Canadian citizens and permanent residents of Chinese ancestry, international students, and individuals holding valid work permits [37,38]. Additional eligibility criteria included individuals 16 years of age or older and living in Canada for at least 6 months at the time of survey questionnaire completion. Our online survey was voluntarily completed by 591 respondents, with nearly 85% of respondents (N=502) meeting all eligibility criteria for inclusion in this study. Among these valid responses, 491 respondents with completed age group and immigration status measures were included in final data analyses (see Supplemental Material—S1).

Utilizing a convenience sampling procedure, our survey questionnaire was distributed leveraging communication channels widely used by Canadian Chinese including social media platforms such as WeChat, emails, and community organizations such as the Consulate General of

China in Toronto, Ontario. Potential respondents were provided with a brief study overview and a consent form detailing principles of anonymity and their rights as participants. Subsequently, informed consent was obtained from all participants involved in the study prior to beginning the survey questionnaire. Further, respondents had the option to withdraw their participation at any time before clicking "Submit" to record their survey questionnaire responses. After submitting survey responses participants could enter their email in an incentive draw with a \$25.00CAD electronic shopping card (10 in total). Utilizing IP addresses, we deterred duplicate entries and fostered genuine responses. To safeguard privacy, all identifiers, such as WeChat IDs, email addresses, and IP addresses were removed and not associated with responses.

For all data collection, we utilized an online survey questionnaire created in Qualtrics TM, and made available in traditional Chinese, simplified Chinese, and English to ensure accessibility to a sufficiently large sample of Canadian Chinese. This questionnaire was also used in previous studies conducted on Canadian Chinese [39–41]. Following informed consent, the questionnaire commenced with questions screening for each eligibility criterion and then proceeded through five sections.

The first section aimed to gather background and general sociodemographic information about Chinese in Canada. Next, the second section focused on experiences of individuals with COVID-19 infection, symptom severity after the most recent infection, COVID-19 infection prevention efforts, and types of treatment. In the third section participants were asked about experiences with COVID-related symptoms more than 12 weeks after being infected with COVID-19. The fourth section contained questions about current health status, diagnosis with underlying diseases, risky health behaviors (e.g., smoking status, regular alcohol consumption), and other health behaviors (e.g., utilizing supplements, traditional Chinese medicine). Additionally, the fifth section explored information on the participants' COVID-19 vaccination history, COVID-19 vaccine type, side effects after receiving the COVID-19 vaccine, and past year Influenza vaccination status.

2.2. Main Outcome Variable – Long COVID

One question assessed experienced long COVID: "Some people still have physical or psychological symptoms more than 12 weeks after being infected with the coronavirus. This phenomenon is called long COVID. Do you think you have or have had long COVID symptoms?" The corresponding binary outcome variable was created with responses "Yes, recovered now" and "Still experiencing long COVID-19" classified as "Yes" and "No" classified as "No".

2.3. Long COVID Experience

The duration of any long COVID symptoms experienced by participants was reported as a continuous interval variable with a unit of months. Symptoms of long COVID were reported as "Fatigue", "Memory problems", "Sleep disorder", "Shortness of breath", "Anxiety and depression", "Pain/discomfort", "Difficulty thinking or concentrating", and "Others". The question assessing symptoms of long COVID allowed for multiple selections. Additionally, participants who selected the response option "Others" (n=24) were prompted to specify additional symptoms via a text box entry. These variables related to long COVID experiences were not included in logistic regression models as they applied only to participants reporting a history of long COVID.

2.4. Covariates

We explored 27 covariates across five broad categories. For improved data analysis and more efficient interpretation, response categories with no or limited data were reclassified into closely related or otherwise appropriate categories. Similarly, questions with too many categories were also reclassified by merging related or similar categories.

2.4.1. Background and General Sociodemographic Information

Age group was assessed as "Under 25", "25 to 34", "35 to 44", "45 to 54", "55 to 64", and "65 or older" and categorized as "Under 45", "45 to 64", and "65 or older". For logistic regression a dummy variable was created for age group using "Under 45" (n=98) as the reference category. Gender responses comprised "Female", "Male", and "Others", and were reclassified as "Women" and "Men". The category "Others" (n=3) was classified as the majority category, "Women". Religiosity responses included "None", "Christianity", "Catholicism", "Islam", "Buddhism", and "Others". Religiosity responses were reclassified to create a new religiosity variable comprised of "Religious" when responses were "Christianity", "Catholicism", "Islam", "Buddhism", and "Others" and "Not religious" when response was "None". Marital status comprised "Single/divorced/widowed", "Married/common law", and "Others". The category "Others" (n=2) was classified as the majority category, "Married/common law". Education responses included "High school or below", "College/university", "Postgraduate", and "Others". The category "Others" (n=2) was classified as the majority category, "College/university". Place of birth was assessed as "Mainland China", "Hong Kong", "Taiwan", "Canada", and "Other regions or countries", and were reclassified as "Mainland China" and "Others".

Whether employment was as a health care worker was assessed as "Yes" and "No". Similarly, whether employment involved contact with the public was assessed as "Yes" and "No". Satisfaction with financial status was assessed on a 5-point Likert scale including "Very dissatisfied", "Dissatisfied", "Neutral", "Satisfied", and "Very satisfied". Financial satisfaction responses were then reclassified as "Satisfied" when responses were "Very satisfied" and "Satisfied" or "Not satisfied" when responses were "Neutral", "Dissatisfied", and "Very dissatisfied".

Immigration status was assessed as "Canadian citizen/permanent resident", "International student", "Family visit or tourist", "Business", and "Others", and were reclassified as "Canadian citizen/permanent resident" and "Others". Period living in Canada was assessed as "Under 5 years", "5 to under 10 years", and "10 years or more". Province of residence was assessed as "Ontario", "Quebec", "British Columbia", "Alberta", "Saskatchewan", "Manitoba", "Atlantic Canada", and "Other provinces of Canada", and were reclassified as "Ontario" and "Others".

Responses to whether there were children aged 16 or under in the household included "No", "Yes – one", "Yes – two", and "Yes – three or more", and were merged into three categories including "No", "Yes – one", and "Yes – two or more". Here, the category "Yes – three or more" (n=4) was merged with "Yes – two". Responses to whether there were people aged 65 or above in the household included "No", "Yes – one", "Yes – two", "Yes – three or more", and were merged into three categories including "No", "Yes – one", and "Yes – two or more". Here, the category "Yes – three or more" (n=1) was merged with "Yes – two". These previous two measures were combined for logistic regression, creating a new variable, household with children (aged \leq 16) or elderly (aged \geq 65), categorized as "Yes" and "No".

2.4.2. COVID-19 Infection Experiences

Number of positive COVID-19 test results responses included "No", "Infected once", "Infected twice", "Infected more than twice", and "Not sure", and were merged into three categories including "None/not sure", "One", and "Two or more". Symptom severity after the most recent COVID-19 infection were rated as "Asymptomatic", "Very mild", "mild", "Serious", and "Very serious", and were merged into three categories including "Asymptomatic/very mild", "Mild", and "Serious/very serious". For logistic regression a dummy variable was created for symptom severity combining "Asymptomatic" (n=13) and "Very mild" (n=41) as the reference category, "Asymptomatic/very mild". Treatment received for most recent COVID-19 infection was assessed as "Did not treat the symptoms", "Treated with traditional Chinese medicine", "Treated with over-the-counter medication", "Treated with prescription medication", and "Hospitalized". The category "Hospitalized" (n=3) was classified as "Treated with prescription medication". For logistic regression

a dummy variable was created for treatment received using “No treatment” (n=123) as the reference category.

2.4.3. Health and Health Behaviors

Health status was assessed as “Very good”, “Good”, “Fair”, “Poor”, “Very poor”, and were reclassified as “Very good/good” and “Fair/poor/very poor”. Responses for underlying health conditions include “Hypertension”, “Diabetes”, “Heart disease”, “Allergy”, “COPD”, “Asthma”, “Arthritis”, “Osteoporosis”, “Cancer”, “Back pain/lumbar pain”, “Obesity”, “Others”, and “None”. The question assessing underlying health conditions allowed for multiple selections. For logistic regression analysis, a binary underlying health conditions variable was created with categories including “Had at least one underlying disease” and “None”.

Regularly taking vitamin supplements or medicine to prevent COVID-19 infection was reported as “Vitamin D”, “Vitamin C”, “Traditional Chinese medicine”, and “Others”. The question assessing vitamin supplement and medication intake allowed for multiple selections. Smoking status was assessed as “Yes”, “No”, “Occasionally”, and “Rather not answer”, and were merged into two categories including “Smoker” and “Nonsmoker”. The category “Rather not answer” (n=2) was classified as the majority category, “Nonsmoker”. Similarly, regular alcohol consumption was assessed as “Yes”, “No”, “Occasionally”, and “Rather not answer”, and were merged into two categories including “Yes” and “No”. Here, the category “Rather not answer” (n=2) was classified as the majority category, “No”.

2.4.4. Vaccination History

COVID-19 vaccination history was assessed as “Never”, “Once”, “Twice”, “Three times”, and “Four or more times”, and were merged into four categories including “Never vaccinated”, “Vaccinated once”, “Vaccinated twice”, and “Three or more”. Type of COVID-19 vaccination included “mRNA-type”, “Vector”, “Inactivated virus”, “Protein”, and “Not sure”. The category “Not sure” (n=4) was classified as the majority category “mRNA-type”. Side effects after receiving COVID-19 vaccine comprised “Yes”, “No”, and “Not sure”, and were merged into two categories including “Yes” and “No/not sure”. Here, the category “Not sure” (n=88) was combined with the majority category, “No”. Finally, influenza vaccination within the year prior to survey completion comprised “Yes”, “No”, and “Not sure”, and were merged into two categories including “Yes” and “No”. The category “Not sure” (n=2) was classified as the majority category, “No”.

2.5. Missing Data

Prior to analyzing the final dataset, we explored missing data frequency and proportion by observation across the final dataset. The pattern and extent of missingness reflected an arbitrary pattern of missingness. Observed data was leveraged to estimate the likelihood of missingness for each covariate and its association with other variables considered in explanatory analyses [42]. Upon inspecting the entire dataset and exploring associations between covariates and missing values in each variable we assumed data in the study were missing at random. Accordingly, we handled all missing data through multiple imputations by fully conditional specification, a valid approach used to handle missing data across categorical variables [43,44].

For the final sample there was a missing data rate of 0.61% to 21.38%, with an overall missing data rate of 8.48% among variables included in explanatory analyses, indicating the potential for multiple imputation to reduce bias and improve validity of parameter estimates for the associations of interest [42,44]. In accordance with the overall proportion of missing values in our dataset, we generated 10 imputed datasets for analysis [44]. Notably, there were no missing values for the main outcome variable, long COVID, or covariate variables including age group and marital status.

2.6. Statistical Analyses

All statistical analyses were performed using SAS (version 9.4) and a significance threshold set at a p -value < 0.05 was applied for all analyses. Descriptive analyses for categorical variables included summarizing the final dataset by frequency and proportion. For the continuous variable, long COVID duration, descriptive analyses included summarizing long COVID duration by mean and 95% confidence interval. Each covariate was stratified by long COVID experience and the difference of each covariate variable between long COVID history groups was assessed using Pearson's Chi-square tests or Fisher's exact test (when more than 20% of cells had expected cell counts lower than 5).

Explanatory analyses were performed to identify factors associated with experiencing long COVID. First, univariate logistic regression models were used to assess the association between each selected covariate and experienced long COVID for the completed dataset. Categorical variables with insufficient numbers in one or more categories (e.g., education, COVID-19 vaccination history, COVID-19 vaccination type, province of residence) were not considered in explanatory analyses [45]. Additionally, based on our knowledge of the present study in the context of relevant literature, variables describing participant characteristics which were either minimally relevant to long COVID or captured in other covariates (e.g., place of birth, immigration status, length of stay in Canada) were not included in logistic regression. Similarly, categorical variables without reference levels facilitating meaningful comparisons (e.g., infection prevention efforts) were not considered in explanatory analyses. Overall, among covariate variables age group, gender, religiosity, marital status, work in health care, contact with the public at work, financial satisfaction, children or elderly in house, positive COVID-19 test results, COVID-19 symptom severity, COVID-19 treatment received, health status, underlying diseases, smoking status, regular alcohol consumption, COVID-19 vaccine side effects, and received Influenza vaccine were considered as predictive variables.

We used logistic LASSO regression to select the predictive variables most significantly associated with experiencing long COVID for inclusion in the final multivariate logistic regression model [46,47]. In logistic LASSO regression the LASSO penalty function can increase prediction accuracy through reducing some coefficients to zero. Subsequently, multivariate logistic regression models were developed to yield odds ratios in the presence of the explanatory variables selected from the LASSO logistic regression model. Finally, sensitivity analysis consisted of creating multivariate logistic regression models for both the completed dataset and the imputed datasets to assess the potential influence of missing data on our parameter estimates. The Hosmer-Lemeshow test was then employed to evaluate the goodness of fit of our final multivariate logistic regression model [48]. Furthermore, the sensitivity and specificity of the final model were assessed utilizing the area under the ROC Curve, with a cut-point of ROC over 0.7 considered acceptable [48].

3. Results

3.1. Participants' Sociodemographic and Health-Related Characteristics

Results of the descriptive analyses characterize participants' sociodemographic, COVID-related, and health-related characteristics stratified by long COVID experience (see Table 1). Amongst the 491 participants, 63 (12.83%) reported a long COVID experience and 428 (87.17%) did not report any long COVID experience. Participants with a history of long COVID were more often women (69.84% vs. 54.35%, $p = 0.0207$), religious (32.65% vs. 18.99%, $p = 0.0275$), with fair, poor, or very poor health status (54.10% vs. 25.27%, $p < 0.0001$), and one or more underlying diseases (70.00% vs. 52.49%, $p = 0.0116$), relative to those who did not report a history of long COVID. Across long COVID groups most participants were married or living common law (84.13% in the long COVID group vs. 80.14% in the no long COVID group), born in Mainland China (98.41% vs. 96.03%), Canadian citizens or permanent residents (95.24% vs. 93.46%), and residing in Ontario (90.48% vs. 82.44%). Further, more than half had stayed in Canada for 10 years or more (71.43% in the long COVID group vs. 77.57% in the no long COVID group), were 45 to 64 years old (63.49% vs. 59.81%), had a college or university education

(52.38% vs. 51.52%), were not satisfied with their financial status (58.73% vs. 56.42%), had no children (aged ≤ 16) in their household (63.49% vs. 68.94%), and had no elderly (aged ≥ 65) in their household (68.25% vs. 67.93%). Moreover, a small proportion of participants reported working in health care (6.35% in the long COVID group vs. 12.22% in the no long COVID group) and contact with the public at work (25.81% vs. 25.76%).

Table 1. Participants' characteristics by long COVID experience.

Variables	N	Long COVID		<i>p</i> -value
		Yes (n = 63)	No	
Age group				0.8320
65 or above	97 (19.76)	12 (19.05)	85 (19.86)	
45 to 64	296 (60.29)	40 (63.49)	256 (59.81)	
Under 45	98 (19.96)	11 (17.46)	87 (20.33)	
Gender				0.0207
Women	275 (56.35)	44 (69.84)	231 (54.35)	
Men	213 (43.65)	19 (30.16)	194 (45.65)	
Religiosity				0.0275
Religious	80 (20.73)	16 (32.65)	64 (18.99)	
Not religious	306 (79.27)	33 (67.35)	273 (81.01)	
Marital status				0.4545
Married/common law	396 (80.65)	53 (84.13)	343 (80.14)	
Single/divorced/widowed	95 (19.35)	10 (15.87)	85 (19.86)	
Education				0.8526
High school or below	13 (2.65)	1 (1.59)	12 (2.81)	
College/university	253 (51.63)	33 (52.38)	220 (51.52)	
Postgraduate	224 (45.71)	29 (46.03)	195 (45.67)	
Place of birth				0.4918 ^a
Mainland China	473 (96.33)	62 (98.41)	411 (96.03)	
Others	18 (3.67)	1 (1.59)	17 (3.97)	
Work in health care				0.1733
Yes	53 (11.42)	4 (6.35)	49 (12.22)	
No	411 (88.58)	59 (93.65)	352 (87.78)	
Contact with the public at work				0.9935
Yes	118 (25.76)	16 (25.81)	102 (25.76)	
No	340 (74.24)	46 (74.19)	294 (74.24)	
Financial satisfaction				0.7313
Satisfied	199 (43.26)	26 (41.27)	173 (43.58)	
Not satisfied	261 (56.74)	37 (58.73)	224 (56.42)	
Immigration status				0.7838 ^a
Citizen/permanent resident	460 (93.69)	60 (95.24)	400 (93.46)	
Others	31 (6.31)	3 (4.76)	28 (6.54)	
Length of stay in Canada				0.3278
Under 5 years	43 (8.76)	5 (7.94)	38 (8.88)	
5 to under 10 years	71 (14.46)	13 (20.63)	58 (13.55)	
10 years or above	377 (76.78)	45 (71.43)	332 (77.57)	
Province of residence				0.1087
Ontario	409 (83.47)	57 (90.48)	352 (82.44)	

Others	81 (16.53)	6 (9.52)	75 (17.56)	
Children (aged ≤ 16) in house				0.1302
Yes – two or more	53 (11.55)	12 (19.05)	41 (10.35)	
Yes – one	93 (20.26)	11 (17.46)	82 (20.71)	
No	313 (68.19)	40 (63.49)	273 (68.94)	
Elderly (aged ≥ 65) in house				0.9986
Yes – two or more	81 (17.65)	11 (17.46)	70 (17.68)	
Yes – one	66 (14.38)	9 (14.29)	57 (14.39)	
No	312 (67.97)	43 (68.25)	296 (67.93)	
Positive COVID-19 test results				<0.0001
Two or more	16 (3.53)	9 (14.29)	7 (1.79)	
One	218 (48.12)	46 (73.02)	172 (44.10)	
None/not sure	219 (48.34)	8 (12.70)	211 (54.10)	
COVID-19 symptom severity				0.0011
Very serious/serious	73 (27.86)	28 (45.16)	45 (22.50)	
Mild	135 (51.53)	21 (33.87)	114 (57.00)	
Asymptomatic/very mild	54 (20.61)	13 (20.97)	41 (20.50)	
COVID-19 treatment received				0.0046
Prescription medicine	22 (8.49)	7 (11.11)	15 (7.65)	
Over-the-counter medicine	82 (31.66)	20 (31.75)	62 (31.63)	
Traditional Chinese medicine	32 (12.36)	15 (23.81)	17 (8.67)	
No treatment	123 (47.49)	21 (33.33)	102 (52.04)	
Health status				<0.0001
Very good/good	309 (70.71)	28 (45.90)	281 (74.73)	
Fair/poor/very poor	128 (29.29)	33 (54.10)	95 (25.27)	
Underlying diseases				0.0116
One or more	232 (54.98)	42 (70.00)	190 (52.49)	
None	190 (45.02)	18 (30.00)	172 (47.51)	
Infection prevention efforts				0.6720
Vitamin D	63 (17.95)	10 (19.23)	53 (17.73)	
Vitamin C	193 (54.99)	29 (55.77)	164 (54.85)	
Traditional Chinese medicine	30 (8.55)	6 (11.54)	24 (8.03)	
Others	65 (18.52)	7 (13.46)	58 (19.40)	
Smoking status				1.0000 ^a
Smoker	15 (3.44)	2 (3.28)	13 (3.47)	
Nonsmoker	421 (96.56)	59 (96.72)	362 (96.53)	
Regular alcohol consumption				1.0000 ^a
Yes	34 (7.80)	4 (6.56)	30 (8.00)	
No	402 (92.20)	57 (93.44)	345 (92.00)	
COVID-19 vaccination history				0.1703 ^a
Three or more	325 (72.38)	39 (63.93)	286 (73.71)	
Vaccinated twice	109 (24.28)	20 (32.79)	89 (22.94)	
Vaccinated once	3 (0.67)	1 (1.64)	2 (0.52)	
Never vaccinated	12 (2.67)	1 (1.64)	11 (2.84)	
COVID-19 vaccination type				0.6751 ^a
mRNA-type	381 (88.19)	54 (90.00)	327 (87.90)	
Vector	17 (3.94)	3 (5.00)	14 (3.76)	
Protein	1 (0.23)	0 (0.00)	1 (0.27)	

Inactivated virus	33 (7.64)	3 (5.00)	30 (8.06)	
COVID-19 vaccine side effects				0.0016
Yes	146 (33.80)	31 (51.67)	115 (30.91)	
No/not sure	286 (66.20)	29 (48.33)	257 (69.09)	
Received Influenza vaccine				0.4664
Yes	192 (43.64)	24 (39.34)	168 (44.33)	
No	248 (56.36)	37 (60.66)	211 (55.67)	

¹ Chi-square tests (or Fisher's exact test^a when greater than 20% of the cells had expected cell counts less than 5) were utilized to evaluate and compare the distribution of categorical covariate variables between long COVID experience groups.

3.2. Participants' COVID-Related and Vaccine-Related Characteristics

As presented in Table 1, participants with long COVID experience more often had one (73.02% vs. 44.10%) or two or more (14.29% vs. 1.79%) positive COVID-19 PCR or rapid antigen detection test results ($p < 0.0001$), compared to those who did not report a history of long COVID. Participants with long COVID experience also more often reported very serious or serious COVID-19 symptoms (45.16% vs. 22.50%, $p = 0.0011$), COVID-19 vaccine side effects (51.67% vs. 30.91%, $p = 0.0016$), and received TCM (23.81% vs. 8.67%) or prescription medicine (11.11% vs. 7.65%) for COVID-19 treatment ($p = 0.0046$), relative to those not reporting long COVID. Among both long COVID groups most participants were nonsmokers (96.72% in the long COVID group vs. 96.53% in the no long COVID group), did not regularly consume alcohol (93.44% vs. 92.00%), and received mRNA-type COVID-19 vaccination (90.00% vs. 87.90%). Additionally, over half of participants reported taking vitamin C supplements (55.77% vs. 54.85%) as part of efforts to prevent COVID-19 infection, and did not receive Influenza vaccination within the previous year (60.66% vs. 55.67%). Finally, although prevalence of three or more doses of COVID-19 vaccination was lower in the long COVID group (63.93%) compared to the no long COVID group (73.71%), this difference was insignificant ($p = 0.1703$).

3.3. Participants' Long COVID Experience, Duration, Symptoms, and Underlying Diseases

Among the 63 participants (12.83%) reporting a history of long COVID, a high proportion (n=51) also reported the duration of their corresponding COVID-19 symptoms. The mean duration of long COVID was 5.31 (95% CI: 4.06–6.57) months. A majority (n=60) of participants reporting long COVID experience also identified the symptoms of COVID-19 they experienced for an extended period. The main symptoms of long COVID were difficulty concentrating (21.67%), pain/discomfort (15.00%), anxiety/depression (8.33%), fatigue (6.67%), shortness of breath (5.00%), and other symptoms (43.34%). Notably, other symptoms (n=26) mainly included cough (30.77%), lethargy (7.69%), chest pain/tightness (7.69%), and hair loss (7.69%). Furthermore, most of the long COVID group (n=42) reported at least one underlying disease primarily including back/lumbar pain (40.48%), allergy (19.05%), diabetes (7.14%), cancer (7.14%), and obesity (7.14%).

3.4. Participant Characteristics Associated with Long COVID Experience

Several potentially significant associations with long COVID were identified from a univariate logistic regression model (see Table 2). Women had significantly higher odds of developing long COVID compared to men (OR = 1.945, 95% CI: 1.099–3.442, $p = 0.0224$). Similarly, individuals who identified as religious had greater odds of experiencing long COVID compared to non-religious individuals (OR = 2.068, 95% CI: 1.073–3.986, $p = 0.0300$). A history of multiple positive COVID-19 test results was strongly associated with long COVID, with odds increasing dramatically for individuals with two or more positive test results (OR = 33.907, 95% CI: 10.070–114.171, $p < 0.0001$). COVID-19 symptom severity also showed a significant association as individuals reporting very serious or serious symptoms had increased odds of long COVID (OR = 6.809, 95% CI: 3.793–12.223, $p < 0.0001$).

Odds of long COVID were also elevated among those who received prescription medicine (OR = 3.442, 95% CI: 1.345–8.808, $p = 0.0099$), over-the-counter medicine (OR = 2.746, 95% CI: 1.515–4.978, $p = 0.0009$), and TCM (OR = 7.555, 95% CI: 3.548–16.090, $p < 0.0001$) compared to those who received no treatment. Notably, good or very good health status appeared to be protective against long COVID experience (OR = 0.287, 95% CI: 0.165–0.500, $p < 0.0001$). Additionally, individuals with one or more underlying diseases had higher odds of long COVID (OR = 2.112, 95% CI: 1.172–3.808, $p = 0.0129$). Finally, experiencing COVID-19 vaccine side effects was positively associated with the odds of experiencing long COVID (OR = 2.389, 95% CI: 1.375–4.149, $p = 0.0020$).

Table 2. Characteristics associated with long COVID experience from the univariate logistic regression model.

Variables	Long COVID		<i>p</i> -value
	OR	95% CI	
Age group			
65 or above	0.949	0.485-1.860	0.8799
45 to 64	1.168	0.675-2.021	0.5777
Under 45	ref		
Gender			
Women	1.945	1.099-3.442	0.0224
Men	ref		
Religiosity			
Religious	2.068	1.073-3.986	0.0300
Not religious	ref		
Marital status			
Married/common law	1.313	0.642-2.688	0.4558
Single/divorced/widowed	ref		
Work in health care			
Yes	0.487	0.169-1.400	0.1817
No	ref		
Contact with the public at work			
Yes	1.003	0.544-1.849	0.9935
No	ref		
Financial satisfaction			
Satisfied	0.910	0.531-1.560	0.7314
Not satisfied	ref		
Children or elderly in house			
Yes	1.074	0.629-1.833	0.7937
No	ref		
Positive COVID-19 test results			
Two or more	33.907	10.070-114.171	<0.0001
One	7.054	3.242-15.346	<0.0001
None/not sure	ref		
COVID-19 symptom severity			
Very serious/serious	6.809	3.793-12.223	<0.0001
Mild	1.377	0.782-2.426	0.2675
Asymptomatic/very mild	ref		
COVID-19 treatment received			
Prescription medicine	3.442	1.345-8.808	0.0099
Over-the-counter medicine	2.746	1.515-4.978	0.0009
Traditional Chinese medicine	7.555	3.548-16.090	<0.0001
No treatment	ref		
Health status			
Very good/good	0.287	0.165-0.500	<0.0001

Fair/poor/very poor	ref		
Underlying diseases			
One or more	2.112	1.172-3.808	0.0129
None	ref		
Smoking status			
Smoker	0.944	0.208-4.290	0.9405
Nonsmoker	ref		
Regular alcohol consumption			
Yes	0.807	0.274-2.377	0.6974
No	ref		
COVID-19 vaccine side effects			
Yes	2.389	1.375-4.149	0.0020
No/not sure	ref		
Received Influenza vaccine			
Yes	0.815	0.469-1.415	0.4669
No	ref		

² A significance threshold set at a *p*-value < 0.05 was applied. Abbreviations: OR = Odds ratio, 95% CI = 95% Confidence interval.

Subsequently, we utilized multiple imputed datasets to estimate parameter coefficients for multivariate logistic regression analysis. Numerous significant associations with long COVID were identified from the imputed multivariate logistic regression model (see Table 3). Multivariate logistic regression models were utilized to produce odds ratios and 95% confidence intervals after a subset of explanatory variables were selected from the LASSO logistic regression model (see Supplemental Material—S2). Notably, in the imputed cases analysis, individuals with two or more positive COVID-19 test results had substantially higher odds of experiencing long COVID compared to those not reporting any positive test results (OR = 23.725, 95% CI: 5.098–110.398, *p* < 0.0001). Unsurprisingly, participants with one positive test result also exhibited increased odds of long COVID experience (OR = 4.286, 95% CI: 1.504–12.216, *p* = 0.0065). Although religiosity approached statistical significance in complete cases analysis, it fell short of the significance threshold for imputed cases analysis (OR = 2.257, 95% CI: 0.993–5.128, *p* = 0.0519). A significant positive association was observed between severe or very severe COVID-19 symptoms and the odds of long COVID experience (OR = 3.177, 95% CI: 1.160–8.702, *p* = 0.0246). Additionally, the use of TCM for COVID-19 treatment was strongly associated with higher odds of long COVID (OR = 8.259, 95% CI: 3.016–22.620, *p* < 0.0001), while the use of over-the-counter medications also showed a significant association (OR = 2.473, 95% CI: 1.035–5.909, *p* = 0.0416). Conversely, participants who reported very good or good health status had significantly lower odds of long COVID compared to those with fair, poor, or very poor health status (OR = 0.247, 95% CI: 0.112–0.544, *p* = 0.0005). However, no significant associations were observed for gender, working in health care, financial satisfaction, underlying diseases, or COVID-19 vaccine side effects in either analysis.

Table 3. Characteristics associated with long COVID experience from the final multivariate logistic regression model.

Variables	Complete Cases Analysis			Imputed Cases Analysis		
	OR	95% CI	<i>p</i> -value	OR	95% CI	<i>p</i> -value
Gender						
Women	1.431	0.590-3.470	0.4272	1.291	0.633-2.634	0.4820
Men	ref			ref		
Religiosity						
Religious	2.611	1.010-6.751	0.0477	2.257	0.993-5.128	0.0519
Not religious	ref			ref		
Work in health care						

Yes	0.300	0.049-1.818	0.1902	0.256	0.063-1.042	0.0570
No	ref			ref		
Financial satisfaction						
Satisfied	1.426	0.614-3.316	0.4091	1.500	0.747-3.013	0.2548
Not satisfied	ref			ref		
Positive COVID-19 test results						
Two or more	53.912	6.901-421.189	0.0001	23.725	5.098-110.398	<0.0001
One	7.328	2.063-26.028	0.0021	4.286	1.504-12.216	0.0065
None/not sure	ref			ref		
COVID-19 symptom severity						
Very serious/serious	2.739	0.840-8.924	0.0946	3.177	1.160-8.702	0.0246
Mild	0.344	0.097-1.222	0.0990	0.860	0.302-2.447	0.7758
Asymptomatic/very mild	ref			ref		
COVID-19 treatment received						
Prescription medicine	3.274	0.725-14.775	0.1229	2.969	0.868-10.156	0.0828
Over-the-counter medicine	1.956	0.682-5.608	0.2118	2.473	1.035-5.909	0.0416
Traditional Chinese medicine	14.781	4.006-54.542	<0.0001	8.259	3.016-22.620	<0.0001
No treatment	ref			ref		
Health status						
Very good/good	0.144	0.055-0.378	<0.0001	0.247	0.112-0.544	0.0005
Fair/poor/very poor	ref			ref		
Underlying diseases						
One or more	1.426	0.560-3.629	0.4570	1.609	0.751-3.445	0.2207
None	ref			ref		
COVID-19 vaccine side effects						
Yes	1.663	0.728-3.801	0.2275	1.738	0.823-3.668	0.1465
No/not sure	ref			ref		

³ A significance threshold set at a *p*-value < 0.05 was applied. Abbreviations: OR = Odds ratio, 95% CI = 95% Confidence interval.

3.5. Sensitivity Analysis

A complete cases analysis was conducted using a dataset with no missing values (N=333) for a multivariate logistic regression model (see Table 3). The results identified religiosity (OR = 2.611, 95% CI: 1.010–6.751, *p* = 0.0477), two or more positive COVID-19 test results (OR = 53.912, 95% CI: 6.901–421.189, *p* = 0.0001), one positive COVID-19 test result (OR = 7.328, 95% CI: 2.063–26.028, *p* = 0.0021), and the use of TCM for COVID-19 treatment (OR = 14.781, 95% CI: 4.006–54.542, *p* < 0.0001) as significant predictors of long COVID. Additionally, participants who reported very good or good health status were significantly less likely to develop long COVID (OR = 0.144, 95% CI: 0.055–0.378, *p* < 0.0001). While these findings largely align with the imputed data analysis, the odds ratios for positive COVID-19 test results and TCM use were notably higher in the complete cases analysis compared to the multiple imputation approach, suggesting potential biases or variability in estimates arising from the exclusion of incomplete cases with missing observations.

Any discrepancies observed between complete cases analysis and imputed cases analysis could stem from reduced statistical power and potential biases associated with excluding participants with incomplete data [49]. For instance, the odds ratio for two or more positive COVID-19 test results was markedly higher in the complete cases analysis (OR = 53.912, 95% CI: 6.901–421.189) compared to the imputed cases analysis (OR = 23.725, 95% CI: 5.098–110.398). Similarly, TCM use had an elevated odds ratio in the complete cases analysis (OR = 14.781, 95% CI: 4.006–54.542) compared to the imputed cases analysis (OR = 8.259, 95% CI: 3.016–22.620). Under the assumption of data missing at random, imputed cases analysis can reduce such biases by incorporating information from incomplete cases into the analysis [49]. Therefore, despite the results of complete cases analysis offering valuable insights, the imputed cases analysis model likely provides more robust and reliable estimates for identifying predictors of long COVID from our data analysis.

4. Discussion

Nearly 13% of Canadian Chinese who participated in our study reported a history of long COVID with an average duration slightly longer than 5 months. Among participants with a history of long COVID, most reported at least one underlying disease, with back/lumbar pain, allergy, diabetes, cancer, and obesity being among the most common. Additionally, the most frequently reported symptoms of long COVID were difficulty concentrating, pain/discomfort, anxiety/depression, fatigue, and shortness of breath. Our results also indicate that the number of positive COVID-19 test results, COVID-19 symptom severity, taking over-the-counter medicine, and receiving TCM were positively associated with long COVID among Canadian Chinese during the pandemic. Further, our results suggest a protective effect of very good/good health status.

4.1. Long COVID Experience, Duration, Symptoms, and Underlying Diseases

Findings from a meta-analysis of 120 studies present a wide range of long COVID prevalence estimates from available literature, ranging from 0 to 93% [50]. Corresponding with this wide range was a pooled prevalence estimate of 42.1%. Such wide variation between studies may reflect commensurate variation in the definition and measurement of long COVID, perhaps diminishing comparability of results between different studies. Focusing on Canada, a retrospective chart review examining the experiences of a COVID-19 cohort in Toronto, Ontario, showed that 27% of those infected with COVID-19 developed long COVID [51]. This is consistent with our findings, as about 26.9% of participants who reported at least 1 positive COVID-19 test result also reported experiencing long COVID. In another meta-analysis of 41 studies, the global prevalence of long COVID was again estimated to be as substantial as 43% [52]. Conversely, in a prospective observational cohort study of prolonged COVID-19 symptoms in the United Kingdom, 13.3% of 4,182 incident cases of COVID-19 reported experiencing symptoms lasting at least 28 days [53]. This estimate is markedly lower than the long COVID prevalence among those with at least 1 positive test result (27%). Such disparities may be attributable to key differences in factors such as participant characteristics and regional differences in long COVID prevalence [50,52,54]. For example, the present study focuses on Canadian Chinese while most available studies focus on general populations, likely differing in a variety of sociodemographic (e.g., age, gender), health-related (e.g., health status), and COVID-related (e.g., number of infections) factors [50,53,54]. These results suggest the prevalence of long COVID among Canadian Chinese was notable during the pandemic.

Another meta-analysis exploring the COVID-19 symptom durations reported in 15 papers described more than 50 COVID-19 symptoms persisting between 14 to 110 days post infection [55]. In the Canadian context, a prospective population-based cohort study assessing acute COVID-19 symptoms and their evolution up to 9 months post infection reported a long COVID duration of approximately 3 months [56]. This estimate is lower than the duration of long COVID of about 5 months reported in our study. Such a differentiation in long COVID duration may be attributable to differences in racial, behavioral, or lifestyle characteristics [57]. These differences could have been further influenced by disparate timing and duration of data collection. That is, our study involved a data collection period between December 22, 2022, and February 15, 2023, whereas Benoit-Piau et al. (2023) recruited participants diagnosed with COVID-19 between November 1, 2020, and May 31, 2021, followed by a data collection period between August and September of 2021 [56]. Accordingly, future studies should further explore racial and ethnic differences in long COVID duration, in the Canadian context, to confirm whether these results represent a true increase in long COVID duration for Canadian Chinese relative to the general population.

Research conducted in the Chinese context has elucidated that long COVID can impact diverse organ systems such as the respiratory system, nervous system, digestive system, as well as mental health [20,29,30]. The most common symptoms of long COVID identified in the Chinese population were fatigue, cough, pharyngitis, lack of concentration, anxiety, myalgia, arthralgia, Sputum, diarrhea, dyspnea, arrhythmias, fever, and hyperhidrosis. Similarly, the most common long COVID

symptoms identified among the Chinese population in the Canadian context included difficulty concentrating, pain/discomfort, anxiety/depression, fatigue, and shortness of breath. These findings offer valuable insights into the impact of long COVID among Canadian Chinese, highlighting the diverse impairments and disruptions to daily functioning associated with the disease.

Concerning underlying diseases, previous studies underscored a higher prevalence of long COVID when the proportion of patients with diabetes, hypertension, obesity, respiratory diseases, liver disease, kidney disease, immune disorders, or allergies was greater [50]. Furthermore, another review study stated that heart disease, diabetes, cancer, COPD, chronic kidney disease, and obesity elevate the risk of both severe COVID-19 symptoms and long COVID [58]. The results of our study further support the comorbidity of long COVID with allergy, diabetes, cancer, and obesity among Canadian Chinese during the pandemic. In addition to these underlying diseases, the most reported comorbidity in our study was back/lumbar pain. This finding is consistent with a growing body of literature suggesting that the incidence of back pain is increasing worldwide and may be accentuated during pandemic lockdowns [59–61].

4.2. Sociodemographic Characteristics, Health-Related Factors, and Long COVID

The present study failed to observe any significant association between long COVID experience and factors such as age group, marital status, work in health care, contact with the public at work, financial satisfaction, children or elderly in the household, smoking status, or regular alcohol consumption. This is inconsistent with a large population-based survey conducted by Wong et al. (2023) aiming to assess the COVID-19 experiences of 2,712 COVID-19 patients across multiple centers in Beijing, Shanghai, Guangzhou, and Hong Kong during the pandemic [20]. Their findings highlighted the multifaceted nature of long COVID, revealing correlations between long COVID susceptibility, such as femininity, engagement in transportation or disciplined labor, living arrangements, smoking habits, overall health perception, presence of chronic diseases, medication use, and COVID-19 severity. Similarly, in their meta-analysis of 120 studies, Woodrow et al. (2023) identified an elevated prevalence of long COVID reported in studies wherein study samples had higher proportions of those older than 50 years of age, men, and people of non-White ethnicity [50]. Conversely, consistent with our findings, Woodrow et al. (2023) reported the prevalence of long COVID did not significantly differ by smoking status. Moreover, the systematic review of 50 studies from Chen et al. (2022) noted positive associations between long COVID and older age, number of COVID-19 symptoms, comorbidity, and pre-existing conditions such as obesity [52]. In addition, Subramanian et al. (2022) conducted a large retrospective matched cohort study using a United Kingdom-based primary care database to select 486,149 adults with a confirmed COVID-19 infection and no COVID-related hospitalization during 2022 [62]. They found the risk of long COVID was notably higher among older adults, females, ethnic minorities, those with lower socioeconomic status, smokers, and individuals with obesity or other comorbidities.

Although there was no significant association found between long COVID and factors including gender, religiosity, and underlying diseases in the final multivariate logistic regression model, these factors had significant effects on long COVID experience in our univariate models. Furthermore, religiosity approached significance in the complete cases analysis for multivariate regression. Employing a survey questionnaire developed by the United Kingdom's Office of National Statistics, a 2022 population-representative survey of 3,042 adults in the United States of America identified associations between long COVID and factors including female gender and underlying diseases [63]. Similarly, results from a study of 7,150 COVID-19 patients in Spain found the probability of long COVID in women was significantly higher compared with men [15]. Intriguingly, a systematic review by David et al. (2023) outlines the potential relevance of religiosity as it pertains to individual responses to the COVID-19 pandemic, identifying numerous articles reporting greater religiosity was associated with poorer adherence to public health behavior guidelines [64]. Relatedly, a multi-national comparison exploring regional variations in religiosity and the spread of COVID-19 during the pandemic, revealed that declared attendance of religious services was associated with more

infections and higher mortality [65]. Of particular importance, the observed association remained when controlling for regional variations in both the number of coronavirus tests per 1 million population and gross domestic product per capita.

The variation observed between the findings of the present study and those described above may be attributed to numerous factors [20,50,52,58]. First, differences in study design, sample size, and population characteristics could significantly influence observed associations. For example, while studies such as Wong et al. (2023) or Chen et al. (2022) utilized large and diverse samples covering multiple centers, our study may have been limited by a smaller or more homogenous sample, focusing on the GTA, likely reducing statistical power and the ability to detect certain associations [50,52]. Second, cultural and healthcare system differences may play a role, as factors such as healthcare access, diagnostic practices, and social determinants of health vary across regions and could influence the reported associations [54]. Third, differences in statistical modeling approaches, including adjustment for missing values, differences in analytical decisions (e.g., variable definition and categorization), the number of covariates, and the selected study sample [66]. Altogether, these considerations and the variations observed between available literature and the current study underscore the importance of contextualizing findings within methodological and regional frameworks of individual studies. Additionally, we recommend future research should continue to explore the complex interactions between individual demographic characteristics, health-related factors, and contextual factors influencing the likelihood of long COVID development.

4.3. Health Status and Long COVID

Among health-related factors assessed in our explanatory analyses, health status was significantly associated with long COVID. More specifically, the odds of long COVID were 75.3% lower for participants with very good or good health status compared to those with fair, poor, or very poor health status. A significant amount of literature has reported similar results regarding a strong negative association between very good or good health status and the development of long COVID [20,67]. Consistent with our findings, using data from 10 longitudinal study samples in the United Kingdom, Thompson et al. (2022) completed a meta-analysis of survey responses from 6,907 individuals with self-reported COVID-19 infection [68]. Notably, they found those with poor or fair pre-pandemic general health and mental health had a greater risk of long COVID. In a prospective single health system observational cohort study Weerahandi et al. (2021) presented that patients (≥ 18 years of age) without good health conditions were more likely to experience long COVID [67]. Further, diverse studies have demonstrated that people with poor health status, such as those who had a history of hospitalization in an intensive care unit (ICU), had underlying diseases, or had multisystem inflammatory syndrome (MIS) during or after the COVID-19 infection were at higher risk of developing long COVID [69–71].

4.4. Number of Positive COVID-19 Test Results and Long COVID

Remarkably, our results indicate a strong association between the number of positive COVID-19 test results and the likelihood of long COVID such that participants with two or more positive test results were about 23.73 times more likely to develop long COVID compared to those not reporting positive test results. This is consistent with emerging literature describing a potentially important association between COVID-19 reinfection and long COVID. A 2023 narrative review article postulated the absolute number of long COVID outcomes substantially increased as a greater proportion of COVID-19 cases caused by the variety of Omicron family subvariants contributed to a considerable increase in COVID-19 reinfections [72]. Interestingly, a 2022 assessment of United States Department of Veterans Affairs databases explored the COVID-19 experience of a population ≥ 50 years of age, illustrating that the likelihood of developing long COVID significantly increased among individuals with reinfections compared to those with primary COVID-19 infections, regardless of COVID-19 vaccination status [73]. Moreover, Su et al. (2022) conducted a longitudinal multi-omic study on 309 COVID-19 patients, from diagnosis to convalescence, and found long COVID patients

may also have lower antibody levels following infection, relative to those without long COVID experience, potentially elevating their risk of reinfection [74]. These results indicate effectively managing the burden of COVID-19 among vulnerable populations should incorporate strategies aimed at limiting reinfection [73].

4.5. COVID-19 Symptom Severity and Long COVID

As it pertains to COVID-19 symptom severity, participants in our study who experienced very serious or serious symptoms had approximately 3.18 times higher odds of developing long COVID compared to those who were asymptomatic or reported very mild symptoms. This result is consistent also with numerous previous studies which have identified symptom severity as a significant predictive risk factor for long COVID [15,20,58,67]. In April 2020, a prospective observational cohort study of 161 hospitalized patients \geq 18 years of age with laboratory-confirmed COVID-19 diagnosis, described how patients with more severe COVID-19 symptoms experienced poorer health status, which in turn increased likelihood of long COVID [67]. Relatedly, Sudre et al. (2020) identified experiencing more than 5 symptoms in the first week of COVID-19 infection among the most important long COVID risk factors across gender and age groups [53]. Similarly, in their systematic review and meta-analysis, Woodrow et al. (2023) identified hospitalization and severity of COVID-19 infection as being the most important factors for long COVID experience [50]. In a multicenter cohort study examining 1,969 inpatient and clinical records of individuals who had recovered from COVID-19 across 5 public hospitals in Spain, noting more than 60% of hospitalized COVID-19 survivors developed long COVID [15]. Collectively, these findings underscore the significant prevalence of long-lasting symptoms among those who have battled severe cases of COVID-19, highlighting the necessity for continued research, medical attention, and additional rehabilitation-oriented services, to address the lingering effects of COVID-19 beyond initial recovery.

4.6. COVID-19 Treatment Received and Long COVID

Additionally, our results underscore the significance of COVID-19 treatment received, with participants who reported treating symptoms with over-the-counter medicine having around 2.47 times higher odds of long COVID compared to those who did not treat symptoms. Remarkably, participants who treated symptoms with TCM were about 8.26 times more likely to develop long COVID compared to those not receiving treatment. To the best of our knowledge, the present study appears to be among the first to consider COVID-19 symptom treatment modalities as predictive risk factors for long COVID development. Around the time of data collection for our study, clinical guidelines primarily emphasized long COVID symptom management with various treatment approaches under active evaluation [75]. Research has highlighted the widespread use of self-prescribed medications for preventing and managing acute COVID-19, including antiretrovirals, antibiotics such as penicillin, vitamin C, traditional remedies, and drugs such as hydroxychloroquine [76–78]. The notable prevalence of self-prescribing behaviors observed during the pandemic were likely driven by the high morbidity and mortality associated with COVID-19, the scarcity of effective treatment guidelines, and limited access to health care during lockdowns. Importantly, self-prescription carries significant risks, including potential adverse drug interactions and the use of ineffective or even harmful therapies [79]. These findings suggest further research is warranted to fully understand the self-management practices that were used to manage COVID-19 symptoms; factors influencing their uptake; and their possible harms, particularly as it pertains to Canadian Chinese, who are less inclined to engage with Western medical services [27,28].

4.7. Vaccine-Related Factors and Long COVID

In contrast, our study failed to observe any significant association between history of long COVID and Influenza vaccination status. COVID-19 and Influenza, while being distinct types of infectious disease, have similarities in epidemiology, clinical manifestations, and pathological

mechanisms [80]. Relatedly, some studies have shown that Influenza vaccination reduces the risk of COVID-19 infection and severity or mortality [81–83]. However, consistent with our results, existing studies indicate there is no significant effect of Influenza vaccination on subsequent long COVID experience.

A retrospective follow-up study of 1,236 adults with long COVID found receiving a COVID-19 vaccination was significantly associated with prolonged COVID-19 symptoms for more than 1 year following initial infection [84]. Nonetheless, a recent systematic review describes numerous observational studies which have reported protective and therapeutic effects of COVID-19 vaccination on long COVID, including reductions in symptom severity, reinfections, and mortality [84,85]. Among our participants, we did not observe sufficient variation in COVID-19 vaccination status to conduct a robust analysis of such effects on the long COVID experience of Canadian Chinese. However, we explored the effect of vaccine side effects on long COVID. Despite COVID-19 vaccine side effects not reaching statistical significance in the final multivariate logistic regression model, vaccine side effects had a significant effect on long COVID history in univariate models. After reviewing other studies, it seems we are among the first to observe a potential association between long COVID and COVID-19 vaccine side effects [85]. Accordingly, we recommend future immunological studies assess vaccine side effects amongst long COVID patients.

4.8. Strengths and Limitations

This study has some important strengths. The study offers valuable insights into the long COVID experience of Chinese residents in Canada during the pandemic, indicating long COVID effected a notable proportion of Canadian Chinese, causing diverse and prolonged impairments and disruptions for effected individuals. Furthermore, our study identifies key characteristics of Canadian Chinese who developed long COVID, and factors associated with increased likelihood of long COVID among Chinese residents in Canada. Available literature identifies associations among factors such as health status, sociodemographic characteristics, number of COVID-19 infections, and symptom severity, which suggests a potential for multicollinearity among covariate variables [30,58,73,86,87]. Accordingly, another strength of the current study is the use of LASSO multivariate logistic regression to address this potential issue in model building.

Despite these strengths, our study also has some notable limitations. Most of our study participants were Chinese residents of Ontario which may not be representative of the total Canadian Chinese population. Relatedly, our use of a convenience sampling procedure could further limit the generalizability of our findings to the total population of Chinese residents in Canada. Additionally, collecting data through an online survey might have introduced selection bias, as individuals who did not have or use electronic devices, as well as those without an internet connection, were either excluded or under-represented in our sample. Nonetheless, an online survey was among the most suitable data collection methodologies during the COVID-19 pandemic, especially considering the ongoing public health response in Canada during our data collection period [34]. Finally, our reliance on self-report measures for data collection introduces the potential for various information biases such as recall or social desirability bias. Overall, we recommend caution when generalizing the results of the current study.

Supplementary Materials: The following supporting information can be downloaded at the website of this paper posted on Preprints.org, Supplemental material—S1: Characteristics of final dataset and study variable distributions; Supplemental material—S2: LASSO variable selection models and sensitivity and specificity analyses.

Author Contributions: Conceptualization, M.P. and P.P.W; Formal analysis, K.G.; Methodology, F.X., X.H. and N.L.; Project administration, Y.C. and P.P.W.; Supervision, P.P.W.; Writing – original draft, M.S. and K.G.; Writing – review & editing, K.G., R.P., D.S., L.F., V.H., and A.D. All authors have read and agreed to the published version of the manuscript.

Funding: This work was supported by the Public Health Agency of Canada (PHAC) through the Immunization Partnership Fund (IPF) with grant number 2223-HQ-000310. Co-first authors were both supported by the Dean's Fellowship Award, Faculty of Medicine, Memorial University.

Institutional Review Board Statement: This study was approved by the Interdisciplinary Committee on Ethics in Human Research (ICEHR), Memorial University of Newfoundland, file number 20201772-ME, 12 March 2020.

Informed Consent Statement: All participants signed consent electronically to the participation statement included in the online survey.

Data Availability Statement: The datasets used and/or analyzed during this study are available from the corresponding author on reasonable request due to restrictions (e.g., privacy reasons).

Acknowledgments: We sincerely thank the members of the study steering board, all collaborators, and especially the study participants, whose time and effort were invaluable and essential to the completion of this study. The basis for this manuscript was derived from Matin Shariati's MSc thesis, and the authors would also like to extend their gratitude to Dr. Peter Daley and Dr. Lin Na for reviewing the early thesis chapters which informed this paper.

Conflicts of Interest: The authors declare no conflicts of interest.

References

1. Alah, M.A.; Abdeen, S.; Kehyayan, V. The first few cases and fatalities of Corona Virus Disease 2019 (COVID-19) in the Eastern Mediterranean Region of the World Health Organization: A rapid review. *J Infect Public Health* **2020**, *13*, 1367–1372. <https://doi.org/10.1016/j.jiph.2020.06.009>.
2. World Health Organization. WHO COVID-19 dashboard | WHO. Available online: <https://data.who.int/dashboards/covid19/> (accessed on 8 December 2024).
3. Wang, L.; Wang, J.; Fatholahi, S.N.; Chapman, M. Assessing the impact of COVID-19 on human activities in the Greater Toronto Area by nighttime light images and active COVID-19 cases. In Proceedings of the Geoscience and Remote Sensing Symposium, Kuala Lumpur, Malaysia, 17 July 2022. <https://doi.org/10.1109/IGARSS46834.2022.9884325>.
4. Statistics Canada. Chinese New Year and quality of life among Chinese in Canada | StatCan. Available online: <https://www.statcan.gc.ca/o1/en/plus/2816-chinese-new-year-and-quality-life-among-chinese-canada> (accessed on 8 December 2024).
5. Lu, C.; McGinn, M.K.; Xu, X.; Sylvestre, J. Living in two cultures: Chinese Canadians' perspectives on health. *J Immigr Minor Health* **2017**, *19*, 423–429. <https://doi.org/10.1007/s10903-016-0386-2>.
6. Huang, C.; Wang, Y.; Li, X.; Ren, L.; Zhao, J.; Hu, Y.; Zhang, L.; Fan, G.; Xu, J.; Gu, X.; et al. Clinical features of patients infected with 2019 novel coronavirus in Wuhan, China. *Lancet* **2020**, *395*, 497–506. [https://doi.org/10.1016/S0140-6736\(20\)30183-5](https://doi.org/10.1016/S0140-6736(20)30183-5).
7. Shi, H.; Han, X.; Jiang, N.; Cao, Y.; Alwalid, O.; Gu, J.; Fan, Y.; Zheng, C. Radiological findings from 81 patients with COVID-19 pneumonia in Wuhan, China: a descriptive study. *Lancet Infect Dis* **2020**, *20*, 425–434. [https://doi.org/10.1016/S1473-3099\(20\)30086-4](https://doi.org/10.1016/S1473-3099(20)30086-4).
8. Azer, S.A. COVID-19: pathophysiology, diagnosis, complications and investigational therapeutics. *New Microbes New Infect* **2020**, *37*, 100738. <https://doi.org/10.1016/j.nmni.2020.100738>.
9. Crook, H.; Raza, S.; Nowell, J.; Young, M.; Edison, P. Long covid—mechanisms, risk factors, and management. *BMJ* **2021**, *374*, n1648. <https://doi.org/10.1136/bmj.n1648>.
10. Mao, L.; Jin, H.; Wang, M.; Hu, Y.; Chen, S.; He, Q.; Chang, J.; Hong, C.; Zhou, Y.; Wang, D.; et al. Neurologic manifestations of hospitalized patients with Coronavirus Disease 2019 in Wuhan, China. *JAMA Neurol* **2020**, *77*, 683–690. <https://doi.org/10.1001/jamaneurol.2020.1127>.
11. Goldstein, D.S. The possible association between COVID-19 and postural tachycardia syndrome. *Heart Rhythm* **2021**, *18*, 508–509. <https://doi.org/10.1016/j.hrthm.2020.12.007>.

12. Sun, B.; Tang, N.; Peluso, M.J.; Iyer, N.S.; Torres, L.; Donatelli, J.L.; Munter, S.E.; Nixon, C.C.; Rutishauser, R.L.; Rodriguez-Barraquer, I.; et al. Characterization and biomarker analyses of post-COVID-19 complications and neurological manifestations. *Cells* **2021**, *10*, 386. <https://doi.org/10.3390/cells10020386>.
13. Ledford, H. Do vaccines protect against long COVID? What the data say. *Nature* **2021**, *599*, 546–548. <https://doi.org/10.1038/d41586-021-03495-2>.
14. Astin, R.; Banerjee, A.; Baker, M.R.; Dani, M.; Ford, E.; Hull, J.H.; Lim, P.B.; McNarry, M.; Morten, K.; O'Sullivan, O.; et al. Long COVID: mechanisms, risk factors and recovery. *Exp Physiol* **2023**, *108*, 12–27. <https://doi.org/10.1113/EP090802>.
15. Fernández-de-Las-Peñas, C.; Martín-Guerrero, J.D.; Pellicer-Valero, Ó.J.; Navarro-Pardo, E.; Gómez-Mayordomo, V.; Cuadrado, M.L.; Arias-Navalón, J.A.; Cigarán-Méndez, M.; Hernández-Barrera, V.; Arendt-Nielsen, L. Female sex is a risk factor associated with long-term post-COVID related-symptoms but not with COVID-19 symptoms: the LONG-COVID-EXP-CM multicenter study. *J Clin Med* **2022**, *11*, 413. <https://doi.org/10.3390/jcm11020413>.
16. Cohen, J.; van der Meulen Rodgers, Y. An intersectional analysis of long COVID prevalence. *Int J Equity Health* **2023**, *22*, 261. <https://doi.org/10.1186/s12939-023-02072-5>.
17. Ortona, E.; Buonsenso, D.; Carfi, A.; Malorni, W.; Long Covid Kids study group. Long COVID: an estrogen-associated autoimmune disease? *Cell Death Discov* **2021**, *7*, 77. <https://doi.org/10.1038/s41420-021-00464-6>.
18. Matschke, J.; Lütgehetmann, M.; Hagel, C.; Sperhake, J.P.; Schröder, A.S.; Edler, C.; Mushumba, H.; Fitzek, A.; Allweiss, L.; Dandri, M.; et al. Neuropathology of patients with COVID-19 in Germany: a post-mortem case series. *Lancet Neurol.* **2020**, *19*, 919–929. [https://doi.org/10.1016/S1474-4422\(20\)30308-2](https://doi.org/10.1016/S1474-4422(20)30308-2).
19. Yong, S.J. Long COVID or post-COVID-19 syndrome: putative pathophysiology, risk factors, and treatments. *Infect Dis (Lond)* **2021**, *53*, 737–754. <https://doi.org/10.1080/23744235.2021.1924397>.
20. Wong, M.C.-S.; Huang, J.; Wong, Y.-Y.; Wong, G.L.-H.; Yip, T.C.-F.; Chan, R.N.-Y.; Chau, S.W.-H.; Ng, S.-C.; Wing, Y.-K.; Chan, F.K.-L. Epidemiology, symptomatology, and risk factors for long COVID symptoms: population-based, multicenter study. *JMIR Public Health Surveill* **2023**, *9*, e42315. <https://doi.org/10.2196/42315>.
21. Aiello, R. Canada locks in thousands more early COVID-19 vaccine doses | CTV. Available online: <https://www.ctvnews.ca/politics/canada-locks-in-thousands-more-early-covid-19-vaccine-doses-1.5231973?cache=1> (accessed on 10 December 2024).
22. Tan, C.Y.; Chiew, C.J.; Lee, V.J.; Ong, B.; Lye, D.C.; Tan, K.B. Comparative effectiveness of 3 or 4 doses of mRNA and inactivated whole-virus vaccines against COVID-19 infection, hospitalization and severe outcomes among elderly in Singapore. *Lancet Reg Health West Pac* **2022**, *29*, 100654. <https://doi.org/10.1016/j.lanwpc.2022.100654>.
23. Kuodi, P.; Gorelik, Y.; Zayyad, H.; Wertheim, O.; Wiegler, K.B.; Jabal, K.A.; Dror, A.A.; Nazzal, S.; Glikman, D.; Edelstein, M. Association between BNT162b2 vaccination and reported incidence of post-COVID-19 symptoms: cross-sectional study 2020-21, Israel. *NPJ Vaccines* **2022**, *7*, 101. <https://doi.org/10.1038/s41541-022-00526-5>.
24. Antonelli, M.; Penfold, R.S.; Merino, J.; Sudre, C.H.; Molteni, E.; Berry, S.; Canas, L.S.; Graham, M.S.; Klaser, K.; Modat, M.; et al. Risk factors and disease profile of post-vaccination SARS-CoV-2 infection in UK users of the COVID Symptom Study app: a prospective, community-based, nested, case-control study. *Lancet Infect Dis* **2022**, *22*, 43–55. [https://doi.org/10.1016/S1473-3099\(21\)00460-6](https://doi.org/10.1016/S1473-3099(21)00460-6).
25. Alah, M.A.; Abdeen, S.; Kehyayan, V. The first few cases and fatalities of Corona Virus Disease 2019 (COVID-19) in the Eastern Mediterranean Region of the World Health Organization: A rapid review. *J Infect Public Health* **2020**, *13*, 1367–1372. <https://doi.org/10.1016/j.jiph.2020.06.009>.
26. Hirawat, R.; Jain, N.; Saifi, M.A.; Rachamalla, M.; Godugu, C. Lung fibrosis: post-COVID-19 complications and evidences. *Int Immunopharmacol* **2023**, *116*, 109418. <https://doi.org/10.1016/j.intimp.2022.109418>.
27. Zhou, P.; Yang, X.-L.; Wang, X.-G.; Hu, B.; Zhang, L.; Zhang, W.; Si, H.-R.; Zhu, Y.; Li, B.; Huang, C.-L.; et al. A pneumonia outbreak associated with a new coronavirus of probable bat origin. *Nature* **2020**, *579*, 270–273. <https://doi.org/10.1038/s41586-020-2012-7>.
28. Wang, C.; Horby, P.W.; Hayden, F.G.; Gao, G.F. A novel coronavirus outbreak of global health concern. *Lancet* **2020**, *395*, 470–473. [https://doi.org/10.1016/S0140-6736\(20\)30185-9](https://doi.org/10.1016/S0140-6736(20)30185-9).

29. Liang, L.; Yang, B.; Jiang, N.; Fu, W.; He, X.; Zhou, Y.; Ma, W.L.; Wang, X. Three-month follow-up study of survivors of coronavirus disease 2019 after discharge. *J Korean Med Sci* **2020**, *35*, e418. <https://doi.org/10.3346/jkms.2020.35.e418>.

30. Zhang, X.; Wang, F.; Shen, Y.; Zhang, X.; Cen, Y.; Wang, B.; Zhao, S.; Zhou, Y.; Baoman Hu, B.; Wang, M. Symptoms and health outcomes among survivors of COVID-19 infection 1 year after discharge from hospitals in Wuhan, China. *JAMA Netw Open* **2021**, *4*, e2127403–e2127403. <https://doi.org/10.1001/jamanetworkopen.2021.27403>.

31. Global Burden of Disease Long COVID Collaborators; Hanson, S.W.; Abbafati, C.; Aerts, J.G.; Al-Aly, Z.; Ashbaugh, C.; Ballouz, T.; Blyuss, O.; Bobkova, P.; Bonsel, G. 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>.

32. Rybkina, J.; Jacob, N.; Colella, B.; Gold, D.; Stewart, D.E.; Ruttan, L.A.; Meusel, L.-A.C.; McAndrews, M.P.; Abbey, S.; Green, R. Self-managing symptoms of Long COVID: an education and strategies research protocol. *Front Public Health* **2024**, *12*, 1106578. <https://doi.org/10.3389/fpubh.2024.1106578>.

33. Statistics Canada. A statistical snapshot of Asians in Canada | StatCan. Available online: <https://www.statcan.gc.ca/o1/en/plus/6178-statistical-snapshot-asians-canada> (accessed on 13 December 2024).

34. Hobgood, D.K., *Personality traits of aggression-submissiveness and perfectionism associate with ABO blood groups through catecholamine activities*. *Med Hypotheses* **2011**, *77*, 294–300. <https://doi.org/10.1016/j.mehy.2011.04.039>.

35. Sapra, R.L. How to Calculate Adequate Sample Size? In *How to Practice Academic Medicine and Publish from Developing Countries?: A Practical Guide*, 1st ed.; Springer: Singapore, 2022; pp. 81–93. https://doi.org/10.1007/978-981-16-5248-6_9.

36. Chan, A.B.; Cooper, C.; Ma, C. Chinese Canadians | TCE. Available online: <https://www.thecanadianencyclopedia.ca/en/article/chinese-canadians> (accessed on 3 December 2024).

37. Statistics Canada. Immigration and ethnocultural diversity: Key results from the 2016 census | StatCan. Available online: <https://www150.statcan.gc.ca/n1/en/daily-quotidien/171025/dq171025b-eng.pdf?st=rzfC1EiM> (accessed on 2 December 2024).

38. Zhang, K. Flows of people and the Canada-China relationship in the 21st century. In *International perspectives on migration: Migration in China and Asia*, 1st ed.; Zhang, J.; Duncan, H., Eds.; Springer: Dordrecht, 2014; Volume 10, pp. 22–50. https://doi.org/10.1007/978-94-017-8759-8_3.

39. Kong, Y.; Shaver, L.G.; Shi, F.; Yang, L.; Zhang, W.; Wei, X.; Zhang, E.; Ozbek, S.; Effiong, A.; Wang, P.P. Knowledge, psychological impacts, and protective behaviours during the first wave of the COVID-19 pandemic among Chinese residents in Canada with dependent school-age children: a cross-sectional online study. *BMC Public Health* **2023**, *23*, 2140. <https://doi.org/10.1186/s12889-023-16923-x>.

40. Na, L.; Yang, L.; Yu, L.; Bolton, K.; Zhang, W.; Wang, P.P. The Appraisal and Endorsement of Individual and Public Preventive Measures to Combat COVID-19 and the Associated Psychological Predictors among Chinese Living in Canada. *Open Public Health J* **2021**, *14*, 592–599. <https://doi.org/10.2174/1874944502114010592>.

41. Kong, Y.; Shaver, L.G.; Shi, F.; Yang, L.; Zhang, W.; Wei, X.; Zhu, Y.; Wang, Y.; Wang, P.P. Attitudes of Chinese immigrants in Canada towards the use of Traditional Chinese Medicine for prevention and management of COVID-19: a cross-sectional survey during the early stages of the pandemic. *BMJ Open* **2021**, *11*, e051499. <https://doi.org/10.1136/bmjopen-2021-051499>.

42. Madley-Dowd, P.; Hughes, R.; Tilling, K.; Heron, J. The proportion of missing data should not be used to guide decisions on multiple imputation. *J Clin Epidemiol* **2019**, *110*, 63–73. <https://doi.org/10.1016/j.jclinepi.2019.02.016>.

43. Van Buuren, S.; Brand JP, L.; Groothuis-Oudshoorn CG, M.; Rubin, D.B. Fully conditional specification in multivariate imputation. *J Stat Comput Simul* **2006**, *76*, 1049–1064. <https://doi.org/10.1080/10629360600810434>.

44. Wright, S.E.; Walmsley, E.; Harvey, S.E.; Robinson, E.; Ferrando-Vivas, P.; Harrison, D.A.; Canter, R.R.; McColl, E.; Richardson, A.; Richardson, M.; et al. Missing Data and Imputation | NIHR. Available online: <https://www.ncbi.nlm.nih.gov/books/NBK333183/> (accessed on 14 December 2024).

45. Ranganathan, P.; Pramesh, C.S.; Aggarwal, R. Common pitfalls in statistical analysis: Logistic regression. *Perspect Clin Res.* **2017**, *8*, 148–151. https://doi.org/10.4103/picr.PICR_87_17.

46. Li, Y.; Lu, F.; Yin, Y. Applying logistic LASSO regression for the diagnosis of atypical Crohn's disease. *Sci Rep* **2022**, *12*, 11340. <https://doi.org/10.1038/s41598-022-15609-5>.

47. International Business Machines. What is Lasso Regression? IBM. Available online: <https://www.ibm.com/topics/lasso-regression> (accessed on 14 December 2024).

48. Hosmer, D.W.; Lemeshow, S.; Sturdivant, R.X. *Applied Logistic Regression*; John Wiley & Sons: Hoboken, NJ, USA, 2013; ISBN 978-0-470-58247-3.

49. Sterne, J.A.C.; White, I.R.; Carlin, J.B.; Spratt, M.; Royston, P.; Kenward, M.G.; Wood, A.M.; Carpenter, J.R. Multiple imputation for missing data in epidemiological and clinical research: Potential and pitfalls. *BMJ* **2009**, *338*, b2393. <https://doi.org/10.1136/bmj.b2393>.

50. Woodrow, M.; Carey, C.; Ziauddeen, N.; Thomas, R.; Akrami, A.; Lutje, V.; Greenwood, D.C.; Alwan, N.A. Systematic review of the prevalence of Long COVID. *Open Forum Infect Dis* **2023**, *10*, ofad233. <https://doi.org/10.1093/ofid/ofad233>.

51. Kozak, R.; Armstrong, S.M.; Salvant, E.; Ritzker, C.; Feld, J.; Biondi, M.J.; Tsui, H. Recognition of long-COVID-19 patients in a Canadian tertiary hospital setting: a retrospective analysis of their clinical and laboratory characteristics. *Pathogens* **2021**, *10*, 1246. <https://doi.org/10.3390/pathogens10101246>.

52. Chen, C.; Haupert, S.R.; Zimmermann, L.; Shi, X.; Fritzsche, L.G.; Mukherjee, B. Global Prevalence of Post-Coronavirus Disease 2019 (COVID-19) Condition or Long COVID: A Meta-Analysis and Systematic Review. *J Infect Dis* **2022**, *226*, 1593–1607. <https://doi.org/10.1093/infdis/jiac136>.

53. Sudre, C.H.; Murray, B.; Varsavsky, T.; Graham, M.S.; Penfold, R.S.; Bowyer, R.C.; Pujol, J.C.; Klaser, K.; Antonelli, M.; Canas, L.S. Attributes and predictors of long COVID. *Nat Med.* **2021**, *27*, 626–631. <https://doi.org/10.1038/s41591-021-01292-y>.

54. Hanson, S.W.; Abbafati, C.; Aerts, J.G.; Al-Aly, Z.; Ashbaugh, C.; Ballouz, T.; Blyuss, O.; Bobkova, P.; Bonsel, G.; Borzakova, S. 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>.

55. Lopez-Leon, S.; Wegman-Ostrosky, T.; Perelman, C.; Sepulveda, R.; Rebollo, P.A.; Cuapio, A.; Villapol, S. More than 50 long-term effects of COVID-19: a systematic review and meta-analysis. *Sci Rep* **2021**, *11*, 16144. <https://doi.org/10.1038/s41598-021-95565-8>.

56. Benoit-Piau, J.; Tremblay, K.; Piché, A.; Dallaire, F.; Bélanger, M.; d'Entremont, M.-A.; Pasquier, J.-C.; Fortin, M.; Bourque, C.; Lapointe, F.; et al. Long-Term Consequences of COVID-19 in Predominantly Immunonaive Patients: A Canadian Prospective Population-Based Study. *J. Clin. Med.* **2023**, *12*, 5939. <https://doi.org/10.3390/jcm12185939>.

57. Brüssow, H.; Timmis, K. COVID-19: long covid and its societal consequences. *Environ. Microbiol.* **2021**, *23*, 4077–4091. <https://doi.org/10.1111/1462-2920.15634>.

58. Hacker, K.A.; Briss, P.A.; Richardson, L.; Wright, J.; Petersen, R. Peer reviewed: COVID-19 and chronic disease: the impact now and in the future. *Prev Chronic Dis* **2021**, *18*, E62. <https://doi.org/10.5888/pcd18.210086>.

59. Dutmer, A.L.; Preuper, H.R.S.; Soer, R.; Brouwer, S.; Bültmann, U.; Dijkstra, P. U.; Copes, M.H.; Stegeman, P.; Buskens, E.; van Asselt, A.D.I.; et al. Personal and societal impact of low back pain: the Groningen spine cohort. *Spine* **2019**, *44*, E1443–E1451. <https://doi.org/10.1097/BRS.0000000000003174>.

60. Battié, M.; Videman, T.; Parent, E. Lumbar Disc Degeneration: Epidemiology and Genetic Influences. *Spine* **2004**, *29*, 2679–2690. <https://doi.org/10.1097/01.brs.0000146457.83240.eb>.

61. Šagát, P.; Bartík, P.; González, P.P.; Tohánean, D.I.; Knjaz, D. Impact of COVID-19 Quarantine on Low Back Pain Intensity, Prevalence, and Associated Risk Factors among Adult Citizens Residing in Riyadh (Saudi Arabia): A Cross-Sectional Study. *Int. J. Environ. Res. Public Health.* **2020**, *17*, 7302. <https://doi.org/10.3390/ijerph17197302>.

62. Subramanian, A.; Nirantharakumar, K.; Hughes, S.; Myles, P.; Williams, T.; Gokhale, K.M.; Taverner, T.; Chandan, J.S.; Brown, K.; Simms-Williams, N.; et al. Symptoms and risk factors for long COVID in non-hospitalized adults. *Nat Med.* **2022**, *28*, 1706–1714. <https://doi.org/10.1038/s41591-022-01909-w>.

63. Robertson, M.M.; Qasmieh, S.A.; Kulkarni, S.G.; Teasdale, C.A.; Jones, H.E.; McNairy, M.; Borrell, L.N.; Nash, D. The epidemiology of long coronavirus disease in US adults. *Clinical Infect Dis.* **2023**, *76*, 1636–1645. <https://doi.org/10.1093/cid/ciac961>.

64. David, A.B.; Park, C.L.; Awao, S.; Vega, S.; Zuckerman, M.S.; White, T.F.; Hanna, D. Religiousness in the first year of COVID-19: A systematic review of empirical research. *CRESP* **2023**, *4*, 100075. <https://doi.org/10.1016/j.cresp.2022.100075>.

65. Linke, M.; Jankowski, K.S. Religiosity and the Spread of COVID-19: A Multinational Comparison. *J Relig Health.* **2022**, *61*, 1641–1656. <https://doi.org/10.1007/s10943-022-01521-9>.

66. Silberzahn, R.; Uhlmann, E.L.; Martin, D.P.; Anselmi, P.; Aust, F.; Awtrey, E.; Bahník, Š.; Bai, F.; Bannard, C.; Bonnier, E.; et al. Many Analysts, One Data Set: Making Transparent How Variations in Analytic Choices Affect Results. *Adv. Methods Pract. Psychol. Sci.* **2018**, *1*, 337–356. <https://doi.org/10.1177/2515245917747646>.

67. Weerahandi, H.; Hochman, K.A.; Simon, E.; Blaum, C.; Chodosh, J.; Duan, E.; Garry, K.; Kahan, T.; Karmen-Tuohy, S.L.; Karpel, H.C.; et al. Post-discharge health status and symptoms in patients with severe COVID-19. *J Gen Intern Med.* **2021**, *36*, 738–745. <https://doi.org/10.1007/s11606-020-06338-4>.

68. Thompson, E.J.; Williams, D.M.; Walker, A.J.; Mitchell, R.E.; Niedzwiedz, C.L.; Yang, T.C.; Huggins, C.F.; Kwong, A.S.F.; Silverwood, R.J.; Di Gessa, G.; et al. Long COVID burden and risk factors in 10 UK longitudinal studies and electronic health records. *Nat Commun* **2022**, *13*, 3528. <https://doi.org/10.1038/s41467-022-30836-0>.

69. Haase, N.; Plovsing, R.; Christensen, S.; Poulsen, L.M.; Brøchner, A.C.; Rasmussen, B.S.; Helleberg, M.; Jensen, J.U.S.; Andersen, L.P.K.; Siegel, H.; et al. Characteristics, interventions, and longer term outcomes of COVID-19 ICU patients in Denmark—a nationwide, observational study. *Acta Anaesthesiol Scand.* **2021**, *65*, 68–75. <https://doi.org/10.1111/aas.13701>.

70. Hassen, H.D.; Welde, M.; Menebo, M.M. Understanding determinants of COVID-19 vaccine hesitancy; an emphasis on the role of religious affiliation and individual's reliance on traditional remedy. *BMC Public Health.* **2022**, *22*, 1142. <https://doi.org/10.1186/s12889-022-13485-2>.

71. Noval Rivas, M.; Porritt, R.A.; Cheng M.H.; Bahar, I.; Arditì, M. Multisystem Inflammatory Syndrome in Children and Long COVID: The SARS-CoV-2 Viral Superantigen Hypothesis. *Front Immunol.* **2022**, *13*, 941009. <https://doi.org/10.3389/fimmu.2022.941009>.

72. Boufidou, F.; Medić, S.; Lampropoulou, V.; Siafakas, N.; Tsakris, A.; Anastassopoulou, C. SARS-CoV-2 Reinfections and Long COVID in the Post-Omicron Phase of the Pandemic. *Int J Mol Sci.* **2023**, *24*, 12962. <https://doi.org/10.3390/ijms241612962>.

73. Bowe, B.; Xie, Y.; Al-Aly, Z. Acute and postacute sequelae associated with SARS-CoV-2 reinfection. *Nat Med.* **2022**, *28*, 2398–2405. <https://doi.org/10.1038/s41591-022-02051-3>.

74. Su, Y.; Yuan, D.; Chen, D.G.; Ng, R.H.; Wang, K.; Choi, J.; Li, S.; Hong, S.; Zhang, R.; Xie, 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>.

75. Crook, H.; Raza, S.; Nowell, J.; Young, M.; Edison, P. Long covid-mechanisms, risk factors, and management. *BMJ.* **2021**, *374*, n1648. <https://doi.org/10.1136/bmj.n1648>.

76. Sadio, A.J.; Gbeasor-Komlanvi, F.A.; Konu, R.Y.; Bakoubayi, A.W.; Tchankoni, M.K.; Bitty-Anderson, A.M.; Gomez, I.M.; Denadou, C.P.; Anani, J.; Kouanfack, H.R.; et al. Assessment of self-medication practices in the context of the COVID-19 outbreak in Togo. *BMC Public Health.* **2021**, *21*, 58. <https://doi.org/10.1186/s12889-020-10145-1>.

77. Quispe-Cañari, J.F.; Fidel-Rosales, E.; Manrique, D.; Mascaró-Zan, J.; Huamán-Castillón, K.M.; Chamorro-Espinoza, S.E.; Garayar-Peceros, H.; Ponce-López, V.L.; Sifuentes-Rosales, J.; Alvarez-Risco, A.; et al. Self-medication practices during the COVID-19 pandemic among the adult population in Peru: A cross-sectional survey. *Saudi Pharm J.* **2021**, *29*, 1–11. <https://doi.org/10.1016/j.jpsps.2020.12.001>.

78. Onchonga, D.; Omwoyo, J.; Nyamamba, D. Assessing the prevalence of self-medication among healthcare workers before and during the 2019 SARS-CoV-2 (COVID-19) pandemic in Kenya. *Saudi Pharm J.* **2020**, *28*, 1149–1154. <https://doi.org/10.1016/j.jpsps.2020.08.003>.

79. Ruiz, M.E. Risks of self-medication practices. *Curr Drug Saf.* **2010**, *5*, 315–323. <https://doi.org/10.2174/157488610792245966>.

80. Bai, Y.; Tao, X. Comparison of COVID-19 and influenza characteristics. *J Zhejiang Univ Sci B.* **2021**, *22*, 87–98. <https://doi.org/10.1631/jzus.B2000479>.

81. Rogers, J.P.; Chesney, E.; Oliver, D.; Pollak, T.A.; McGuire, P.; Fusar-Poli, P.; Zandi, M.S.; Lewis, G.; David, A.S. Psychiatric and neuropsychiatric presentations associated with severe coronavirus infections: a systematic review and meta-analysis with comparison to the COVID-19 pandemic. *Lancet Psychiatry.* **2020**, *7*, 611–627. [https://doi.org/10.1016/S2215-0366\(20\)30203-0](https://doi.org/10.1016/S2215-0366(20)30203-0).

82. Nordberg, P.; Jonsson, M.; Hollenberg, J.; Ringh, M.; Berggren, R.K.; Hofmann, R.; Svensson, P. Immigrant background and socioeconomic status are associated with severe COVID-19 requiring intensive care. *Sci Rep.* **2022**, *12*, 12133. <https://doi.org/10.1038/s41598-022-15884-2>.

83. Đoàn, L.N.; Chong, S.K.; Misra, S.; Kwon, S.C.; Yi, S.S. Immigrant communities and COVID-19: strengthening the public health response. *Am J Public Health.* **2021**, *111*, S224–S231. <https://doi.org/10.2105/AJPH.2021.306433>.

84. Asadi-Pooya, A.A.; Nemati, M.; Shahsavandi, M.; Nemati, H.; Karimi, A.; Jafari, A.; Nasiri, S.; Mohammadi, S.S.; Rahimian, Z.; Bayat, H.; et al. How does COVID-19 vaccination affect long-COVID symptoms? *PLoS One.* **2024**, *19*, e0296680. <https://doi.org/10.1371/journal.pone.0296680>.

85. Byambasuren, O.; Stehlik, P.; Clark, J.; Alcorn, K.; Glasziou, P. Effect of covid-19 vaccination on long covid: systematic review. *BMJ Med.* **2023**, *2*, e000385. <https://doi.org/10.1136/bmjmed-2022-000385>.

86. Karmakar, M.; Lantz, P.M.; Tipirneni, R. Association of Social and Demographic Factors With COVID-19 Incidence and Death Rates in the US. *JAMA Netw Open.* **2021**, *4*, e2036462. <https://doi.org/10.1001/jamanetworkopen.2020.36462>.

87. Bhargava, A.; Fukushima, E.A.; Levine, M.; Zhao, W.; Tanveer, F.; Szpunar, S.M.; Saravolatz, L. Predictors for Severe COVID-19 Infection. *Clin Infect Dis.* **2020**, *71*, 1962–1968. <https://doi.org/10.1093/cid/ciaa674>.

Disclaimer/Publisher's Note: The statements, opinions and data contained in all publications are solely those of the individual author(s) and contributor(s) and not of MDPI and/or the editor(s). MDPI and/or the editor(s) disclaim responsibility for any injury to people or property resulting from any ideas, methods, instructions or products referred to in the content.