Cardio-Pulmonary Sequelae in Recovered COVID-19 Patients: Considerations for Primary Care

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Abstract

Background: Current literature lacks characterization of the post-recovery sequelae among COVID-19 patients. This review characterizes the course of clinical, laboratory, radiological findings during the primary infection period, and the complications post-recovery. Primary care findings are presented for long-COVID care. **Methods:** Adhering to PRISMA guidelines, 4 databases were searched (PubMed, Embase, CINAHL Plus, Scopus) through December 5, 2020, using the keywords "COVID-19 and/or recovered and/or cardiovascular and/or long-term and/or sequelae and/or sub-acute and/or complication." We included published peer-reviewed case reports, case series, and cross-sectional studies providing the clinical course of COVID-19 infection, and cardiopulmonary complications of patients who recovered from COVID-19, while making healthcare considerations for primary care workers. Results: We identified 29 studies across 9 countries including 37.9% Chinese and 24.1% U.S. studies, comprising 655 patients (Mean Age = 45) with various ethnical backgrounds including Asian and European. Based on the WHO COVID-19 severity classification scale, initial disease severity was mild for 377 patients and severe for 52 patients. Treatments during primary infection included corticosteroids, oxygen support, and antivirals. The mean value (in days) for complication onset after acute recovery was 28 days. Complete blood counts and RT-PCR tests were the most common laboratory results described. In 22 of the studies, patients showed signs of clinical improvement and were prescribed medications such as anticoagulants or corticosteroids. Conclusion: Postrecovery infectious complications are common in long-COVID-19 patients ranging from mild infections to life-threatening conditions. International thoracic and cardiovascular societies need to develop guidelines for patients recovering from COVID-19 pneumonia, while focused patient care by the primary care physician is crucial to curb preventable adverse events. Recommendations for real-time and lab-quality diagnostic tests are warranted to establish point-of-care testing, detect early complications, and provide timely treatment.

Keywords

primary care, long COVID, cardiac, pulmonary, COVID-19

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Introduction

Toward the end of December 2019, the COVID-19 outbreak resulted in massive fear worldwide. Although this outbreak initially occurred in Wuhan, a city in Hubei province, China, it rapidly spread throughout the world. The World Health Organization (WHO) announced that the COVID-19 pandemic as a health emergency of public concern on January 30, 2020. The pandemic has drastically impacted all industries and sectors worldwide, placing an

immense burden on the health care system.² COVID-19 has a wide array of clinical presentations ranging from non-specific symptoms such as fever, fatigue, and diarrhea to severe respiratory and cardiovascular complications, including acute respiratory distress syndrome (ARDS) and multiple organ failure leading to mortality.³ As a result of the data gathered from numerous clinical trials conducted worldwide, there has been ongoing concern regarding the persistence of the SARS-CoV-2 virus in the post-acute period.⁴ Due to the limited understanding of the clinical

course of COVID-19, the possible long-term health impacts have not been elucidated. Our systematic review recognizes the sequelae of COVID-19 patients in the post-acute period to identify their persistent symptoms and clinical outcomes and to provide recommendations as the COVID-19 pandemic continues to spread.

Methods

For this systematic review, a literature search was performed from December 1, 2019, through December 5, 2020, to identify published articles that reported the outcomes of COVID-19 patients after recovery. A systematic literature search was conducted on PubMed, Embase, CINAHL Plus, and Scopus to retrieve articles under the Preferred Reporting Items for Systematic Reviews (PRISMA) checklist criteria. The following search terms were used: "COVID-19 and/or recovered and/or cardiovascular and/or long-term and/or sequelae and/or sub-acute and/or complication." The search terms were kept broad to encompass all possibilities for applicable studies. There were no restrictions on the language of articles published.

Duplicates were removed using Endnote v9. After eliminating duplicates, 3 investigators (AS, ZS, and MKC) independently reviewed all titles and abstracts. The full texts of articles regarded as potentially eligible for consideration were extracted and screened for further analysis. Some records were also retrieved via cross-references from published papers (Umbrella review). Thereafter, eligible articles were selected for final analysis according to predefined inclusion and exclusion criteria. Disagreements between the authors were resolved through consensus and active discussion.

Only human studies with clearly defined recovery, survival, other outcomes from acute COVID-19 were included. The patient population had mild, moderate, severe, or critical COVID-19 disease and a delineated of persistent symptoms and clinical outcomes in the post-acute period was essential for inclusion. The exclusion criteria consisted of review articles; animal studies; and the absence or unclear reporting of recovery status of COVID-19. As proposed by the National Institutes of Health, we employed a uniform severity classification, ranging from mild, moderate, severe, and critical.⁵

- 1. Mild Illness: Individuals who have any of the various signs and symptoms of COVID-19 (eg, fever, cough, sore throat, malaise, headache, muscle pain, nausea, vomiting, diarrhea, loss of taste, and smell) but who do not have shortness of breath, dyspnea, or abnormal chest imaging.
- 2. Moderate Illness: Individuals who show evidence of lower respiratory disease during clinical assessment or imaging and who have an oxygen saturation (SpO2) ≥94% on room air at sea level.
- 3. Severe Illness: Individuals who have SpO2 <94% on room air at sea level, a ratio of arterial partial pressure of oxygen to fraction of inspired oxygen (PaO2/FiO2) <300 mm Hg, respiratory frequency >30 breaths/minutes, or lung infiltrates >50%.
- Critical Illness: Individuals who have respiratory failure, septic shock, and/or multiple organ dysfunction.

The only exception to the severity scale employed is the paper by Wei⁶ which relied on the "Diagnosis and Treatment Protocol of Novel Coronavirus issued by the National Health Commission of the People's Republic of China." Data were tabulated into a shared spreadsheet customized for this review. All tables were optimized before commencing data entry using 6 sample articles included in the review to ensure optimal presentation of findings. Articles selected for the final analysis were independently graded by 2 authors (AS and ZS) for quality using the Newcastle–Ottawa Scale (NOS).

No funding was obtained for this study.

Results

The search yielded 1091 studies. After removing 431 duplicates, 660 articles were reviewed for title and abstract. After the initial screening, only 167 articles met the inclusion criteria and underwent full-text evaluation. Records that consisted of commentaries, editorials, or reviews that did not meet our criteria were removed, generating a final list of 29 articles (Figure 1).

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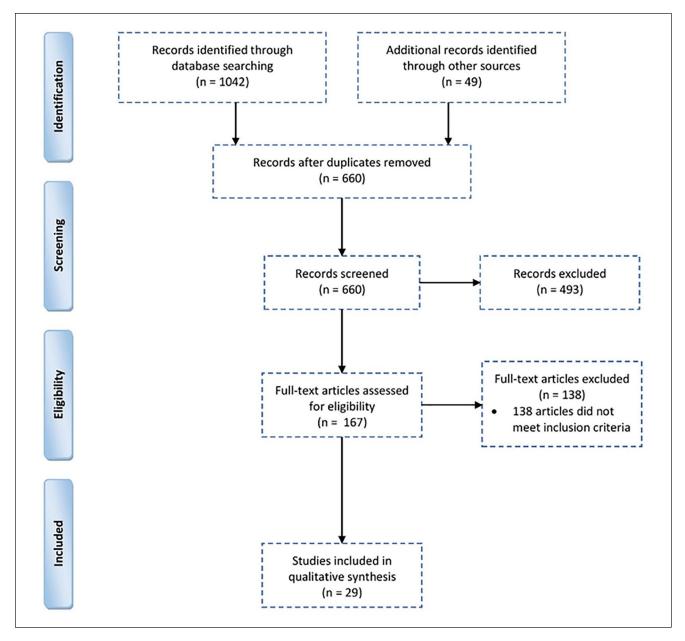


Figure 1. PRISMA flow diagram.

Characteristics of Included Studies

All studies included in this review were published through December 5, 2020. Twenty studies were case reports, 5 were retrospective cohorts, 2 were prospective observational studies, 1 was a case-control, and 1 was a case series (Table 1). Nine countries were represented with the largest data derived from studies in China (37.9%) and the United States (24.1%). The total number of included participants was 655, with a mean age of 45, comprising of various ethnical backgrounds including Asian and European. A total of 419 patients with mild symptoms and 59 patients with

severe symptoms were included. Patient outcomes were notably well due to recovery from infection in 22 of the studies where they recovered or were prescribed medications for underlying thromboembolic conditions including anticoagulants.

Clinical Course and Pulmonary Complications of COVID-19 Recovered Patients

Table 2 summarizes the clinical course of SARS-CoV-2 infection in pulmonary cases, 8 out of these 14 studies had

Table 1. Characteristics of included studies.

o Z	Author	Type of study	Sample (n) Region	Region	Ethnicity	Age (years)	Gender or male (%)	Comorbidities
_	Kosiigi et al7	Case-control	145	Brazil	Not specified	Median: 36 (IOR 30-44)	Male 46 3%	5 1% Aerhma 5 1% HTN
. 2	Brancatella et al ⁸	Case report	<u> </u>	Italy	European	81	Female	
~	Alfano et al ⁹	Case report	-	Italy	European	72	Male	CVD, HTN, DM, CKD, ICH, AF
4	Garg et al ¹⁰	Case report	-	NS	Not specified	78	Male	HTN, DM, CKD, AF CHA2DS2-VASc of 4, ILD
2	Hollingshead and Hanrahan ¹¹	Case report	-	SN	Not specified	50	Male	I
9	He et al ¹²	Case report	-	China	Chinese	39	Female	ICH
7	Sardari et al ¹³	Case report	-	Iran	Iranian	31	Male	1
∞	Cavalagli et al ¹⁴	Case report	-	Italy	European	69	Male	CVD, CeVD
6	Beckman et al ¹⁵	Case report	-	Sweden	European	51	Male	I
0	Liu et al ¹⁶	Case report	-	China	Asian	35	Male	1
=	Li et al ¹⁷	Case report	-	China	Asian	14	Male	1
12	Fujikura et al ¹⁸	Case report	-	SN	Not specified	77	Female	CeVD, HTN, DM, Breast Cancer
<u>~</u>	Pohlan et al ¹⁹	Case report	-	SN	Not specified	64	Female	1
4	Zhou et al ²⁰	Case report	-	China	Not specified	40	Male	I
15	Chen et al ²¹	Case report	-	China	Not specified	38	Male	I
91	May ²²	Case report	-	š	European	12	Male	I
	Xia et al ²³	Case report	2	China	Not specified	70, 42	Male 50%	I
<u>8</u>	Dou et al ²⁴	Case report	-	China	Chinese	34	Male	DΜ
6	Abushahin et al ²⁵	Case report	-	NSA	American	49	Male	DM
70	Insausti-García et al ²⁶	Case report	-	Spain	European	40	Male	I
21	Gervasio et al ²⁷	Case report	2	Italy	Not specified	54, 43	Male 100%	HTN 50%, DM 50%
22	Takeda ²⁸	Case series	9	Brazil	Not specified	Mean (SD): 46.2 (16)	Male 83.3%	HTN 33.3%, BA 16.7%, ICH 16.7%
23	Puntmann et al ²⁹	Prospective Observational cohort study	001	Germany	European	Median: 49 (IQR 45-53)	Male 53%	CVD 13%, HTN 22%, DM 18%, BA 21%, COPD 21%
24	Lu³0	Prospective study	66	China	Chinese	Cases mean (SD): 44.1 (16), controls mean (SD): 45.88 (13.9)	Male cases 56.7%, controls 56.4%	HTN 29.3%, DM 7.1%
25	Mo et al ^{3 l}	Retrospective cohort	01	China	Asian	Mean (SD): 49.1 (14)	Male 50%	CVD 2.7%, CeVD 2.7%, HTN 23.6%, DM 8.2%, Ca 0.9%, BA 0.9%, COPD 2.7%, CLD 5.5%, CKD 1.8%
76	Zhao et al ³²	Retrospective cohort	22	China	Asian	Mean (SD): 47.74 (15.49)	Male 58.1%	CVD 3.64%, HTN 10.91%, DM 3.64%
27	Clark et al ³³	Retrospective cohort	99	SN	N = 16, Non-Hispani	N=16, Non-Hispanic Median: 20 (IQR 19-21)	Male 63.6%	I
78	Rajpal et al ³⁴	Retrospective cohort	76	SN	Not specified	Mean (SD): 19.5 (1.5)	Male 57.7%	I
29	Huang et al ³⁵	Retrospective, observational	26	China	Asian	Median: 38 (IQR 32-45)	Male 38%	MIN 8%

Abbreviations: AF, atrial fibrillation; BA, bronchial asthma; Ca, Cancer; CeVD, cerebrovascular disease; CKD, chronic kidney disease; CLD, chronic liver disease; COPD, chronic obstructive pulmonary disease: CVD, cardiovascular disease; DM, diabetes mellitus; HTN: hypertension; ICH, immuno-compromised host; ILD, interstitial lung disease.

 Table 2. Clinical Course of Pulmonary COVID-19 Among Recovered Patients.

ġ	Author	Disease severity	Status	Clinically recovered (%)	ICU Admission (%) and length of ICU stay	Length of hospital stay (days)	Complications during stay	Treatments during hospital stay	Months of primary infection	Complication onset after primary infection (days)
_	Kosugi et al ⁷	Mild	Course of COVID 19 (+) patients with acute olfactory alterations	52.6%	I	Median: 31 (IQR 10.5-39)*	I	I	I	
2	Garg et al ¹⁰	I	Presents to ER after I month of previous confirmed COVID 19 by nasopharyngeal RT-PCR $\left(+\right)$	%00I	100%, 5 days	I	AF, deterioration, and respiratory distress	Vancomycin, cefepime, MPD, enoxaparin started with till INR of 3	I	30 days
m	Hollingshead and Hanrahan''	Mild	Presents to ER after I month of previous COVID 19 infection	%00I	I	I		Oxygen therapy	Late April 2020	30 days
4	Beckman et al ¹⁵	Asymptomatic	Suspected COVID 19, not confirmed	%00I		49	I	Tinzaparin, LMWH	Beginning of April	7 weeks
2	Liu et al ¹⁶	Mild	Reoccurrence, change of status after RNA SARS-CoV-2 (-) test	%00I	I	91	ı	LPV, INF α 2b, AHG, MPD	Late January 2020	30 days
9	Li et al ¹⁷	Severe	Reoccurrence of infection	%00I	1	7	I	INF-A, TCM, oxygen therapy	Late January 2020	25 days
7	Fujikura et al ¹⁸	Severe	Development of thrombotic state caused by lack of adherence to recommend anticoagulant	%00I	I	17	I	Hydroxychloroquine, azithromycin, vancomycin, oxygen, heparin	I	28 days
∞	Pohlan et al ¹⁹	Mild	Reoccurrence of infection	%001	%00I	I	I	LWMH, sultamicillin	Beginning of April	13 days after initial COVID onset
6	Zhou et al ²⁰	Severe	Reoccurrence of infection by RT-PCR assay due to insufficient antibody production	%001	I	8	1	BiPAP ventilator, MPD, immunoglobulin	I	27 days
<u>o</u>	Dou et al ²⁴	Severe	Reoccurrence of infection by RT-PCR assay in a diabetic patient	I	I	39	I	Arbidol, ribavirin, cefuroxime, chloroquine, AMP, INF-A	Beginning of February	30 days after discharge
=	Abushahin et al ²⁵	Severe	Presents to ER after 21 days of previous COVID 19 infection	%001	100%, 8 days	21	Respiratory failure	Oxygen, piperacillin/tazobactam, vancomycin, azithromycin, hydroxychloroquine	I	21 days
2	Takeda ²⁸	PΙΙ	Recurrent clinical COVID 19 symptoms in health care workers after RT PCR confirmed cases	% 0001	l6.70%	I	Sudden Hypoxemia	Oseltamivir, azithromycin, hydroxychloroquine, prednisone, levofloxacin, piperacillin-tazobactam, MPD, prophylactic anticoagulation, ivermectin	March, Beginning of April	Median (IQR): 56.5 (53-70)
<u>=</u>	Mo et al ³¹	Mild 21.8%, Pneumonia 60.9%, Severe pneumonia 17.3%	Pulmonary testing at time of discharge, impairment of diffusion capacity, restrictive ventilatory defects	%00I	I	Mean (SD): 27 (9)	I	I	I	1
4	Zhao et al ³²	Mild 7.3%, cases of pneumonia 85.4%, severe pneumonia 7.3%	Pulmonary testing after 3 months of discharge	%00I	I	1	I	Oxygen therapy, MPD, TCM	1	3 months

Abbreviations: (+), positive; (-), negative; AHG, arbidol hydrochloride granules; AMP, adenosine monophosphate; BiPAP, bilevel positive airway pressure; INF-A, interferon alpha; LMWH, low molecular weight heparin; LPV, lopinavir; MPD, methylprednisolone; TCM, traditional Chinese medicine.
*Follow up of survey data collection.

a male predominance. About 12 studies reported complete clinical remission, 4 of them reported intensive care unit admission (ICU), 9 mentioned hospital admission, and an average of 34 days from the primary infection to the presence of complication was reported across 11 studies. The major complication during hospital stay was a respiratory failure even though it was only reported in only 2 studies. Treatment was provided in detail in 12 studies, encompassing oxygen support, antivirals, antibiotics, corticosteroids, and anticoagulants. Most patients described shortness of breath, upper respiratory tract like symptoms, and dyspnea as main presenting features. RT-PCR tests and Computed Tomography (CT) Scans were the most commonly recommended tests (Table 3).

Clinical Course and Complications of Cardiovascular COVID-19 Patients

Table 4 summarizes the clinical course and complications of cardiovascular COVID 19 patients. During our search, only 5 studies were classified in the cardiovascular section, with 1 study reported to have a severe COVID-19 population. Chest pain, shortness of breath, and chest pain were noted, with patients in these groups aged 50 years or younger. Complete blood count (CBC) and inflammatory cardiac markers were recurrently elevated and 60% of the cases presented lesions visible through Cardiac Magnetic Resonance (CMR), diagnosed with myocarditis in 3 studies. An approximate period of a month was described from the period of primary infection to the onset of a complication. Treatments during hospital stay were not reported, except for Huang et al³⁵ which reported antiviral, antibiotic, and oxygen support management during the hospital stay (not shown in the table).

Complications and Clinical Course of COVID-19 Patients with Specific Characteristics

The complications and clinical course of all included COVID-19 patients with specific characteristics such as endocrine, hematological, ophthalmological, and neurological details are described in Supplemental Table 1. Of all the aforementioned patients, 70% of them were hospitalized, and 30% of them were admitted to the intensive care unit (ICU). The patients were treated mainly with oxygen supportive therapy, steroids, antibacterial, and antivirals. All patients were confirmed to have COVID-19 with a positive PCR test, initial complete blood work was followed, in addition to inflammatory markers and special panels according to each underlying comorbidity. Lung CT scans were followed in 60% of cases; 70% of the patients develop resolution of the acute clinical syndrome, even though a longer follow up study is suggested to observe the full resolution of symptoms or complete basal return. In 4 (28.6%) of the pulmonary studies, pulmonary embolism (PE) was present. In 2 (14.3%) of studies, pneumothorax was noted, and recurrence of infection was observed in 5 (35.7%) of the studies. Of the cardiovascular studies, 4 (80%) required an additional test to detect abnormal functionality (Tables 3 and 4). Relevant important points for primary care were extracted and described for cardiovascular and pulmonary cases (Tables 3 and 4). Despite the heterogeneity of the included articles, the need for guidelines, and the demand for follow up studies to describe duration or clinical course evolution is significant.

Discussion

This systematic review analyzes the cardio-pulmonary sequelae in recovered COVID-19 patients while making considerations for primary care. With the data gathered from numerous COVID-19 clinical studies conducted across the world, researchers have been able to demonstrate that despite apparent complete recovery from COVID-19, patients may suffer from post-recovery sequelae. However, there has been no consensus on the defining characteristics of post-acute COVID-19 syndrome. Commonly reported symptoms in the post-acute COVID-19 phase are fatigue, dyspnea, joint pain, and chest pain. 36 Cardiopulmonary consequences have been reported in the recovery phases, and have considerable implications for long-term health consequences.^{29,37} A survey conducted by the Centers for Disease Control and Prevention (CDC) identified the contribution of 3 or more underlying medical conditions and age greater than 50 years as important predictors of not achieving baseline health until 3 weeks after testing COVID-19 RT-PCR positive. Nevertheless, 20% of young individuals have not returned to baseline health after a median time of 16 days since testing positive for COVID-19.³⁸ A critical discussion of post-acute COVID-19 syndrome is imperative as sequelae are observed among patients along the infection severity scale (ie, mild to critical).

Pulmonary

Wu et al³⁹ report ARDS in 40% of patients infected with SARS-CoV-2 infection, with over 20% in the older age groups. Burnham et al⁴⁰ find that persistent inflammation of alveolar epithelium and hypercytokinemia in SARS-CoV-2 infection leads to fibroproliferative changes which include fibroblast accumulation and deposition of extracellular matrix collagen in the lung parenchyma. The resulting disruption of the balance between profibrotic and antifibrotic factors leads to pulmonary fibrosis which is observed in a subset of patients as protracted sequelae to ARDS.⁴⁰ Pulmonary fibrosis is characterized by the presence of an unusual interstitial pneumonia-like fibrotic abnormality on high-resolution computed tomography (HRCT) and this is

 Table 3. Pulmonary Complications in COVID-19 Patients.

Nasopharyageal PCR negative, PT 38 CR Bilarenal opacifies, ECK AF, CT Fr with mitigle segmental PE on right lung PE on right lung PE on right lung PE on warfarin 1200 mg/L, BNP 1200. CT Diffuse GGO, 10 cm localized Presumothorax Pe on warfarin Pe	Author		Signs	Symptoms	Laboratory findings	Radiological findings	Diagnosis		Outcomes, discharge indications	Primary care considerations
Nanophalyrigeal PCR registre, FT 38	_	I		86.9% acute anosmia, 13.1% acute hyposmia	I	I	Sudden olfactory dysfunction	I5 (mean)	Recovered, follow up time of 31 days	Patients presented with hyposmia recovered in lesser time than those that presented with acute anosmia
Toponin T 1200 mg L BNP 1200. To District SCO. 10 cm boulsed not of the control o	Garg et al ¹⁰ O2 90 nas: irre witl	O2 90 nas: irre with	% on 4L al cannula, gular pulse h tachycardia	SOB, nonproductive cough		CXR Bilateral opacities, ECK AF, CT multiple segmental PE on right lung	RF with multiple segmental PE on warfarin	I	Recovered, apixaban on discharge	Thromboprophylaxis duration and scheme changes should be given in a case-by-case basis till there is more information in literature
Positive SARS CoV.2 (+) on day 1. CT GGO and patchy lyperdense Positive SARS CoV.2 (+) on day 1. CT GGO and patchy lyperdense Positive SARS CoV.2 (+) on day 1. CT GGO and patchy lyperdense Positive SARS CoV.2 (+) on day 1. CT GGO and patchy lyperdense Positive SARS CoV.2 (+) on day 1. CT GGO and patchy lyperdense Positive SARS CoV.2 (+) on day 1. CT GGO and patchy lyperdense Positive SARS CoV.2 (+) on day 1. CT GGO and patchy lyperdense Positive SARS CoV.2 (+) on day 1. CT GGO and patchy lyperdense Positive SARS CoV.2 (+) on day 2. CT Satistered patches and ground. Positive SARS CoV.2 (+) on day 1. CT GGO and patchy lyperdense Positive SARS CoV.2 (+) on day 2. CT Satistered patches and ground. Positive SARS CoV.2 (+) on day 2. CT Satistered patches and ground. Positive SARS CoV.2 (+) on day 2. CT Satistered patches and ground. Positive SARS CoV.2 (+) on day 2. CT Satistered patches and ground. Positive SARS CoV.2 (+) on day 2. CT Satistered patches and ground. Positive SARS CoV.2 (+) on day 2. CT Satistered patches and ground. Positive SARS CoV.2 (+) on day 2. CT Satistered patches and ground. Positive SARS CoV.2 (+) on day 2. CT Satistered patches and ground. Positive SARS CoV.2 (+) on day 2. CT Satistered patches and ground. Positive SARS CoV.2 (+) on day 2. CT Satistered patches and ground. Positive SARS CoV.2 (+) on day 2. CT Satistered patches and ground. Positive SARS CoV.2 (+) on day 2. CT Satistered patches and ground. Positive SARS CoV.2 (+) on day 2. CT Satistered patches and ground. Positive SARS CoV.2 (+) on day 2. CT Satistered patches and ground. Positive SARS CoV.2 (+) on day 2. CT Satistered patches and ground. Positive SARS CoV.2 (R) on day 2. CT Satistered patches and ground. Positive SARS CoV.2 (R) on day 2. CT Satistered patches and ground. Positive SARS CoV.2 (R) on day 2. CT Satistered patches and ground. Positive SARS CoV.2 (R) on day 2. CT Satistered patches and ground. P	Hollingshead — and Hanrahan''	I		SOB	I	CT Diffuse GGO, 10cm loculated posterior right pneumothorax	Pneumothorax	I	Improvement post chest tube placement	High grade of suspicion based on presentation
RAPA SRS C-OV 2 (+) on day 1. RAPE Recovered from the second of the second o		02 93	%8	Exercise induced dyspnea (~4 weeks) and dyspnea at rest (~48 hours)		CT Bilateral segmental PE and GGOs. Two negative PCR	Widespread PE	I	Recovered, Apixaban for 6 months	Pulmonary sequelae can persist after acute infection; the presence of dyspnea could be key for the timely detection of patients at an early stage
R-PCR (nast) souths, spoulm, stool) Af first viel. LDH (804 LUI, D-dimer 934 ingnite that seed and ground- all (+); RT-PCR (nast) souths, spoulm, stool) Af first viel. LDH (804 LUI, D-dimer 934 ingnite that seed inflammatory markers, reduced (17 Phasse and ground- and problem of antibodies against SARS-Cov-2. gain after 5 days of GGO and consolidation in the bilateral inflammatory markers, reduced CT: Bilateral multiple irregular areas (antibodies against SARS-Cov-2. gain after 5 days of GGO in the bilateral inflammatory markers, after and readial inflammatory markers, after CYOVID-19 was confirmed by RT-PCR (as than 50% extension) Increased inflammatory markers, after CYOVID-19 as confirmed by RT-PCR (as than 50% extension) Increased inflammatory markers and readial inflammatory markers, after CYOVID-19 was confirmed by RT-PCR (as than 50% extension) Anomalies: DLCO abnormalities 47.2%. Anomalies: DLCO abnormalities 47.2%. Anomalies: DLCO impairment (i=3), interestial particle inflammatory markers (i=3), interestial particle inflammatory markers and eveals. DLCO impairment (i=3). Anomalies: DLCO impairment (i=3) All patients with abnormal CT (i=3); pure GGO in the bilateral and levels: DLCO inflammatory markers and eveals inflammatory markers and eveals inflammatory markers. After (i=3) interestial particle inflammatory markers and readial inflammatory markers. After (i=3) interestial particle inflammatory markers and readial inflammatory markers. After (i=3) interestial particle inflammatory markers and readial inflammatory markers. After (i=3) interestial particle inflammatory markers and readial inflammatory markers. After (i=3) interestial particle inflammatory markers and readial inflammatory markers with abnormalities in Covid-19 particle inflammatory markers and particle inflammatory markers with abnormalities in Covid-19 patients where discharged patients and high BUN (i=15), crazy paving (i=3) interestial patients where discharged patients and high BUN (i=15), crazy paving (i=3) interestial p	Liu et al ¹⁶ —			URTI like symptoms	RNA SARS-CoV-2 (+) on day 1, negative on days 14, 15; Positive again on day 30	CT GGO and patchy hyperdense areas	Positive SARS-CoV-2 RNA recurs repeatedly in a case recovered from COVID-19: dynamic results from 108 days of follow-up	95	Patient recovered	The consequences or involvement of dyslipidemia and liver injury must be discerned in greater depth
A first visit. LDH (804 UJ), D-diner (234 µgm). D-diner (220 soudle PE ECHO: MConnell's Late presentation of saddle — Recovered tachycarda (150 months). D-diner (234 µgm). To design the patent (150 months). The control of saddle (150 months). The control of saddle (150 months). The control of saddle (150 months). The control of supplier X-cute (150 months). The control of the co	Li et al ¹⁷ Fevel	Feve	r, O2 90%	Chest pain, cough	RT-PCR (nasal swabs, sputum, stool) all (+); RT-PCR throat swabs were (-), B cells increased, and NIK cells decreased	CT: Scattered patches and ground- glass opacity on both lungs, Septal Line*	Reoccurrence of COVID-19	>35	Recovered, symptoms improved after 8 days	I
NT-pro BNP 3936 ng/L, D dimer 18.3 mg/L 18.4 mg/L 18	Fujikura Tacl et al ¹⁸ 8	Tach 8	ıycardiac, O2 5%	Severe SOB	At first visit LDH (804 U/L), D-dimer (>20.00 (rg/mL), and CPR (11.7 mg/dL). Readmission: D-dimer 9.34 trg/mL	Large saddle PE. ECHO: McConnell's sign. Sausage-like mass in RA, moderater TR, PASP 62 mm Hg. Venous duplex Doppler: Acute occlusive thrombosis	Late presentation of saddle PE and thrombus-in- transit straddle the patent foramen	I	Recovered, tachycardia resolved, discharged home with enoxaparin	Hypercoagulable state even after acute infection, may be of concern in follow-up patients
Dry cough, dyspnea, and Elevated inflammatory markers, reduced CT: Bilateral multiple irragular areas diarrhea by RARS-CoV-2 again after 5 days discharge from hospital, lower levels of GGO and consolidation and probable recurrence of positive from hospital, lower levels of an example of an every of an example of an example of an example of an every of an example of an every of an example of an every of an e	hlan et al ¹⁹ Tacl	Tacl	hycardia, RR 30	Severe SOB and right leg pain	NT-pro BNP 3936 ng/L, D dimer 18.3 mg/L	CT Massive PE, bilateral infiltrates. US DVT of right leg. ECHO: RV strain	PE and DVT	I	Angiographic aspiration thrombectomy, normalization of functional ECHO parameters	I
No symptoms at Increased inflammatory markers, after CT Multiple GGO in the bilateral recurrence 2 consecutive negative results with 2 cough, \$208 and residual infiltrates and patents discomfort, dry COVID-19 was confirmed by RT-PCR CXR large left-sided pneumothorax COVID-19, pneumothorax — Recovered after pigtall chest individual presented (RT)-PCR tests, but reverted to being slight scattered and bilateral GGO. Anomalies: DLCO abnormalities 47.2%, — Anomalies: DLCO abnormalities 47.2%, and small airway function 7.3% G(30.9%), headache Patients with abnormal CT had low (n=15), crazy paving (n=3) abnormalities in Covid-19 recovered patients were discharged patients with abnormal CT had low (n=15), crazy paving (n=3) recovered patients and light BUN (n=15), crazy paving (n=3) recovered patients and levels. DLCO impairment (n=9) recovered patients and levels. DLCO impairment (n=9) recovered patients are of positive and patients were discharged patients and levels. DLCO impairment (n=9) recovered patients and patients were discharged patients and patients were discharged patients and levels. DLCO impairment (n=9) recovered patients and patients were discharged patients and patients were discharged patients and patient	Zhou et al ²⁰ O2	0	%08 >	Dry cough, dyspnea, and diarrhea	Elevated inflammatory markers, reduced Lymphocytes count. (+) RNA test for SARS-CoV-2 again after 5 days discharge from hospital, lower levels of antibodies against SARS-CoV-2.	CT: Bilateral multiple irregular areas of GGO and consolidation	Recurrent pneumonia and probable immunodeficiency	I	Recovered, SARS-CoV-2 remained negative after 14days of further isolation at home	Low tiers anti SARS COV 2 doesn't exclude COVID 19 relapse in all patients
Chest discomfort, dry COVID-19 was confirmed by RT-PCR CXR large left-sided pneumothorax COVID-19, pneumothorax — Recovered after pigtal chest — insertion cough. SOB testing from nasopharyngeal swab and residual infiltrates and residual infiltrat	Dou et al ²⁴ —			No symptoms at recurrence	Increased inflammatory markers, after 2 consecutive negative results with SARS-CoV-2, RT-PCR assay sample was positive	CT Multiple GGO in the bilateral lungs, partially absorbed	Recurrence of positive SARS-CoV-2 (RNA)	73	Recovered, following second discharge, tested negative for SARS-CoV-2 on qRT-PCR for 5 weeks	High probability of severe COVID 19 stages when diabetes mellitus is present
Two discharged patients had negative Patient 3 presented a (HRCT) with Peurrent clinical symptoms — All improved without Ph hospitalization except one positive more evident in peripheral regions and bilateral GGO, of COVID-19 who improved after NIV (less than 50% extension) Anomalies: DLCO abnormalities 47.2%, — (less than 50% extension) Anomalies: DLCO abnormalities 47.2%, — (less than 50% extension) Anomalies: DLCO abnormalities 47.2%, — (less than 50% extension) In C pred 2.1%, FLVI pred 13.6%, PVC 14.9%, and since and in the serum sodium, high serum sodium, (n = 7), interstitial thickening elevated D-dimer and high BUN (n = 15), crazy paving (n = 3) In coovered patients All improved without Ph hospitalization except one who improved after NIV hospitalization except one hospitalization except one who improved after NIV hospitalization except one hospitaliza	Abushahin Hyp et al ²⁵ to	[#]	oxemia, achypnea	Chest discomfort, dry cough, SOB	COVID-19 was confirmed by RT-PCR testing from nasopharyngeal swab	CXR large left-sided pneumothorax and residual infiltrates	COVID-19, pneumothorax	I	Recovered after pigtail chest insertion	I
Anomalies: DLCO abnormalities 47.2%, — Impaired diffusing capacity: — All patients were discharged 13.6%, FVI pred 13.6%, and small airway function 7.3%	Takeda ²⁸ —			URTI like symptoms, I individual presented hypoxemia	Two discharged patients had negative (RT)-PCR tests, but reverted to being positive	Patient 3 presented a (HRCT) with slight scattered and bilateral GGO, more evident in peripheral regions (less than 50% extension	Recurrent clinical symptoms of COVID-19	I	All improved without hospitalization except one who improved after NIV	Phenoms of reoccurrence, viral reactivation or reinfection must be investigated when recurrent clinical symptoms are present
Patients with abnormal CT had low Abnormal CT (n=39): pure GGO Radiographic and — All patients were discharged .4%), serum albumin, high serum sodium, (n=7), interstitial thickening physiological elevated D-dimer and high BUN (n=15), crazy paving (n=3) abnormalities in Covid-19 recovered patients	Mo et al ³¹ —	1		I	Anomalies: DLCO abnormalities 47.2%, TLC pred 25%, FEVI pred 13.6%, FVC pred 9.1%, FEVI/FVC 4.5%, and small airway function 7.3%	Ī	Impaired diffusing capacity: 30.4% (mild illness), 42.4% (pneumonia), and 84.2% (severe pneumonia)	I	All patients were discharged	Prevailing of impaired diffusing lung capacity at time of discharge
	Zhao et al ³² —			Gl (30.9%), headache (18.2%), fatigue (16.4%), exertional dyspnea (16.4%), and cough and sputum (1.8%)	Patients with abnormal CT had low serum abunin, high serum sodium, elevated D-dimer and high BUN levels. DLCO impairment (n=9)	Abnormal CT (n = 39): pure GGO (n = 7), interstital thickening (n = 15), crazy paving (n = 3)	Radiographic and physiological abnormalities in Covid-19 recovered patients	I	All patients were discharged	Higher D-dimer levels on admission could be predictors for impaired DLCO 3 months post discharge

^{*}Pulmonary complications in COVID-19 patients.

 Table 4. Clinical Course and Complications of Cardiovascular COVID-19 Patients.

o Ž	Author	Disease severity	Symptoms	Laboratory findings	Radiological findings	Diagnosis	Complication onset after primary infection (days)	Outcomes, discharge indications	Primary care considerations
_	Sardari et al ¹³	I	Dyspnea on exertion, low grade fever	CRP 3.3 mg/L, Troponin T < 0.03 ng/ mL; RT – PCR negative for SARS-CoV-2	TTE: Mild left ventricular (LV) dysfunction. CMR: T ₂ -weighted showed edema/inflammation in the mid inferoseptal and inferior wall, subepicardial fibrosis in the mid inferior wall	Active myocarditis	21 days from previous hospital admission	Started with bisoprolol and lisinopril	Cardiovascular follow up could be important if sufficient data is gathered regarding cardiovascular complications such as myocarditis, even in mild disease
2	Puntmann et al ²⁹	Mild	I	I	I	COVID-19	14 days	Lower left ventricular ejection fraction, higher left ventricle volumes, higher left ventricle mass, and raised native 7, and 7; CMR revealed cardiar involvement (73%) and ongoing myocardial inflammation (60%)	Cardiovascular involvement after infection could be present independently to severity of disease
м	Clark et al ²⁷	Mild 77%, asymptomatic 23%	I	Troponin I < 99% for age (82%); No alterations in BNP or CRP	CMR. Late gadolinium enhancement found in % of cases; ECC: Median LVEF of 59% LVEDVI 94 mL/m², RVEF 52%	9% patients with myocardial inflammation or fibrosis; 5% met formal criteria for myocarditis	I	All were improved before CMR was done (a median of \$2 days after \$ARS-CoV-2 infection)	Cardiovascular compromise may be missed if there is no opportunity to perform specific cardiac studies (ie, ECG, T1, and strain echocardiography)
4	Rajpal et al ³⁴	Mild 26.9%, Asymptomatic 53.8%	About 2 of 4 athletes with myocardial inflammation had SOB	Normal troponin I	CMR. 15% had myocarditis based on Lake Louise Criteria. About 7.5% had percardial effusion. Mean (SD) Tio those with suspected myocarditis was 59(3) ms; Negative findings on electrocardiogram, normal range TTE	15% with myocarditis	I	Competitive athletes: all patients recovered	CMR could use as a stratification tool for patients at high risk, such as competitive athletes
'n	Huang et al ¹⁵	Moderate 85%, Severe 15%	Chest pain, palpitation, and chest distress	Normal blood work and inflammatory median markers		COMD-19	Median (IQR): 47 (36-58)	58% with abnormal CMR findings: myocardial edema 54%, LGE 31%; Decreased right ventricle functional parameters in patients with positive conventional CMR findings. Global native T ₁ , T ₂ , and ECV were significantly elevated in patients with positive conventional CMR findings; CMR manifestation included myocardial edema, fibrosis, and impaired right ventricle function	The presence of cardiac tissue involvement at this stage could be early in course, could be early in course, discern the correct follow up duration

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linked to a more increased incidence of rapid progression of the disease. 41-44

A prospective study was conducted by Liu et al including 149 patients who were discharged from various hospitals in Wuhan, China following recovery from COVID-19 pneumonia. Patients were followed up during the first, second, and third-weeks post-discharge. The results showed that patients younger than 44 years had a significantly higher percentage of complete radiological resolution than patients who were older than 44 years. Out of the various lesions observed, the most predominant were ground-glass opacity in 83.9% of patients followed by a fibrous stripe in 54.4% and only 22.1% showed thickening of the adjacent pleura. The number of patients testing positive for each lesion decreased gradually over 3 weeks and the radiological resolution was more evident in patients with ground-glass opacity and fibrous stripe in the first and third weeks, respectively. Most residual lung abnormalities were absorbed throughout recovery, suggesting the importance of radiological assessment 2 weeks post-discharge.³⁰ The importance of determining the timing and standpoints of radiological resolution in COVID-19 patients can provide further insight into deciding which patient groups need long term monitoring.

Mo et al³¹ observed that COVID-19 survivors who had been discharged from the hospital demonstrated pulmonary function abnormalities, the most common among those being diffusion capacity impairment, followed by restrictive ventilatory defects. Zhao et al³² demonstrated both physiologic and radiographic abnormalities in a significant number of COVID-19 patients in discharged COVID 19 patients after a 3-month follow-up period. In the INBUILD trial, Kolb and Vašáková⁴⁵ note increased rates of forced vital capacity (FVC) decline and mortality over the 52 weeks in patients with usual interstitial pneumonia (UIP) like lesions when compared to those patients with other patterns of fibrosis. Several factors are involved in the extent to which pulmonary complications present in clinically recovered patients; Salehi et al⁴⁶ document the most significant being patients' age, presence of comorbidities, and type of medications administered during COVID-19 management. Another point of contention is those patients with long-term sequelae who did not have either ARDS or PE during acute infection. While there is a paucity of pulmonary sequelae among patients who did not have either ARDS or PE during the acute infection, Han et al⁴⁷ conducted a 6-month followup of chest CT findings after severe COVID-19 pneumonia. Patients without ARDS were less likely to be associated with post-COVID radiological and functional changes.⁴⁷ There is a lack of detailed knowledge on the pathophysiology of the disease, taking into consideration the available studies, following up on discharged COVID-19 patients, assessing them using radiological and pulmonary tests, and managing potential issues are all necessary precautionary measures.

Cardiovascular

Myocardial injury is defined by laboratory testing, namely by an increased troponin level, which has been documented as a severe COVID-19 sequela. 48 A median of 71 days post COVID-19 diagnosis, cardiac involvement was reported by Puntmann et al²⁹ in 78% patients, and 60% revealed myocardial inflammation on cardiac magnetic resonance imaging in a Germany study. Long et al⁴⁹ reported common cardiovascular sequelae in COVID-19 as myocarditis, thromboembolic disease, and arrhythmias. The alterations in normal cardiovascular physiology, coagulation state, and basal metabolism observed, including those with RNA-sequencing analysis, reveal increased levels of angiotensin-converting enzyme-2 (ACE-2) in cardio-myocytes thereby increasing the possibility of direct myocardial tissue infection in high-risk patients. 47,48 As per Siripanthong et al,50 the ACE-2 receptor on the myocardium may be used by SARS-CoV-2 to gain entry into cells, triggering the release of massive amounts of pro-inflammatory cytokines, immune hyper-activation, and myocardial tissue damage. Cardiac tropism, an impaired cardio-protective function of ACE2 downregulated by SARS-CoV-2, and elevated proinflammatory cytokine release are potential pathological mechanisms. 51-53

Sardari et al¹³ write that myocarditis may occur due to the residual myocardial inflammation in COVID-19. Evidence of SARS-CoV-2 mRNA in the myocardium has been observed by Tavazzi et al54 and Wenzel et al55 on autopsy and endomyocardial biopsies, suggestive of myocarditis. Underlying cardiovascular disease (CVD) may predispose patients to SARS-CoV-2 myocarditis.⁵⁵ Further, cardiovascular complications caused due to COVID-19 or prevalent before infection must be addressed. A meta-analysis finds that the pooled incidence of acute cardiovascular complications during the COVID-19 pandemic is 1.4%. However individuals with COVID-19 who presented with concomitant stroke were more likely to have pre-existing cardiovascular comorbidities not limited to arrhythmias, myocarditis, and thromboembolism, linked to severe COVID-19 infection. The patterns of large vessel occlusions and multi-territory infracts suggests that thromboembolism and/or cerebral thrombosis may potentially be causative pathways for COVID-19 disease. 56 The mortality and morbidity due to cardiovascular complications in COVID-19 patients are higher when there is evidence of myocardial involvement, as implicated by Mitrani et al⁵⁷ In the setting of myocarditis, myocyte lysis releases proteins with abnormal epitopes and with a similar structure to those present in the virus. Błyszczuk⁵⁸ and Gangaplara et al⁵⁹ state that these can be recognized by major histocompatibility complex and seemingly add autoimmune components to cardiac injury. Chapman et al⁶⁰ presume that these mechanisms of tissue damage cause long-term complications in

COVID-19 patients. Varga et al⁶¹ form an association between endothelitis and multi-organ failure with evidencing direct viral infection of the endothelium. Additionally, possible cytopathic effects and increased myocardial cell death after inducible SARS-CoV-2 on induced pluripotent stem cells (iPSC) have also been identified by Bose and McCarthy.⁶²

Multiple studies have shown an increased rate of arrhythmias in patients recovering from viral myocarditis. A study by Blagova et al⁶³ revealed that among 19 cases of "idiopathic" arrhythmias, 78.9% were identified as immunemediated inflammatory after the endomyocardial biopsy was performed, finding cardiotropic viruses as etiologic agents. Since the beginning of the COVID-19 pandemic, thrombotic complications have been reported in critically ill patients. The term MicroCLOTS (microvascular COVID-19 Lung Vessels Obstructive Thromboinflammatory syndrome) due to microvascular pulmonary thrombosis is currently being used in literature to describe manifestations of the disease. The high incidence of micro thrombosis in the lungs was noted in 80% autopsies of COVID-19 cases by Dolhnikoff et al⁶⁴ Klok et al⁶⁵ report the incidence of venous thromboembolism as 31%, leading to pulmonary embolism (80%) and arterial thrombosis (3.7%). Further studies are needed to specifically describe cardiovascular complications post COVID-19, especially screening for the incidence of heart failure. Close monitoring of patients is pertinent, even in cases with no signs of cardiovascular impairment.

Considerations for Primary Care

Although the impact of long-term COVID-19 sequelae is still unclear, a large number of patients will likely experience persistent symptoms, especially in those geographical regions where large outbreaks occurred. Various strategies are being evaluated not only to treat COVID-19 patients but also to prevent as many detrimental complications as possible. Greenhalgh et al describe post-acute COVID-19 as "Long COVID," which is a multisystem disease. 66 Notably, cardio-pulmonary complications may occur after mild infections as well, requiring attention by primary care providers.66 For a general practitioner, it is essential to note improvements over 3 weeks and subsequent weeks/months. The symptoms may manifest in any bodily organ, however, the most common ones include cough (dry versus productive), fever, fatigue during long COVID. Uncommon symptomatology worth considering by the primary care practitioner includes mood swings, palpitations, anxiety, hypotension, and/or skin rashes (ie, maculopapular). The nonspecific (ie, lung burn, brain fog) versus the specific complications (ie, stroke, pulmonary embolism, myocarditis, heart failure) must be distinguished clinically upon a visit to a primary care practitioner and specific presentations recommended to specialist centers. A clinical insight for *long COVID* is the fact that the presentation during *acute COVID* does not directly relate to the severity of findings 6 months later (ie, if a patient had severe pneumonia on the acute onset, the sequelae may be less severe as compared to those who had less severe acute complications). Based on the limited evidence in published literature so far, patients with *long COVID* may have no access to specialist care. To improve patient outcomes, it is essential to adopt a community-based, interprofessional rehabilitation pattern of care by promoting remote technologies, telehealth appointments with a primary care physician, and adopting otherwise locally relevant solutions.

Recommendations

The implementation of a multi-disciplinary approach to identify, and prevent the impact of long-term health outcomes of COVID-19 across different organs, including cardiopulmonary, as well as the well-being of recovered patient groups is necessary to elucidate the potential classification of complications in post-acute COVID-19.67 An example of how rehabilitation can successfully be applied to post-COVID-19 patient management is offered by Shan et al,⁶⁸ who describes the case of a woman at about 80 years of age gradually returning to her premorbid independent condition after severe COVID-19 complicated by acute respiratory distress syndrome (ARDS) following 10 days of post-acute inpatient pulmonary rehabilitation. Interestingly, current knowledge on the long-term impact of the SARS-CoV-2 infection to an important degree comes from previous epidemics, caused by severe acute respiratory syndrome (SARS) and the Middle East respiratory syndrome (MERS). However, deficits associated with the current pandemic are expected to be more critical, given the magnitude of the pandemic.

A practical approach is to follow up on all hospitalized patients with chest radiography and oxygen saturation measurements, as recommended by the British Thoracic Society.⁶⁹ However, the American Thoracic Society/ European Respiratory Society does not make any suggestion in favor of or against routine pulmonary function tests (PFTs), chest CT scans, transthoracic echocardiography (TTE), and cardiopulmonary exercise testing (CPET) within 30 to 60 days. 70 Preliminary evidence suggests the development of respiratory sequelae as early as 2 weeks after discharge, especially among patients with elevated inflammatory markers and D-dimer levels.31,32 Data suggests the frequent occurrence of pulmonary fibrosis, arrhythmias, and cardiomyopathies among survivors over 2 months after recovery.²⁹ All patients requiring further investigation may be referred to as pulmonologists and cardiologists for further evaluation of pulmonary (PFTs and

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chest CT scans) and cardiac (TTE and CPET) assessment within 2 to 3 months of recovery. It is also necessary to establish a baseline time to follow recovery. Admittedly, resource-limited settings may not be able to conduct such tests. Consequently, evidence suggests routine testing of pulmonary and cardiac sequelae in COVID-19 is beneficial within 30 to 90 days of discharge in all previously hospitalized patients.

Limitations

In the United States, 3 different phases of the pandemic have been documented. At the onset, in April, 2020 there was limited testing. During the second phase in July, 2020, social distancing among other factors contributed to infectivity. However, testing was widely available to most of the public. The third phase at the onset of 2021 was potentially connected with seasonal respiratory viruses. These are uncontrollable limitations in post-infection pulmonary-cardiovascular tests that are presented with external factors contributing to varied test outcomes. The other limitation of our study relates to its retrospective nature. Many of the included studies were case reports and, therefore, findings may not be generalizable and the causality within those may not be fully proven. As the majority of the studies were from China, some of the presentations could be influenced by the ethnicity-related characteristics of the region.

Conclusion

With limited insight on the sequelae of COVID-19 infection, a holistic approach is required that incorporates primary care. Complications are frequently observed in patients recovering from COVID-19, ranging from mild to life-threatening conditions. Patients who presented with ARDS, PE, arrhythmia, myocarditis, or other cardiopulmonary complications during the acute phase necessitate multi-speciality care and follow-up for long-COVID. It is currently unclear whether ARDS or other complications due to COVID-19 or other causative factors will have different long-term outcomes. Follow-up of clinically recovered patients with cardiopulmonary imaging studies could be crucial for the evaluation of adverse events caused by SARS-CoV-2 infection. Primary health care forms a critical foundation for gatekeeping timely and effective support during the COVID-19 pandemic. Through primary care services, it is possible to provide rehabilitation and support to recovering patients in light of a greater volume of patients requiring rehabilitation in the post-acute COVID-19 infectious phase. Physicians across all healthcare levels should be alert on and monitor patients for possible adverse events after recovery from SARS-CoV-2 infection. International societies ought to develop

guidelines for patients recovering from COVID-19 and recommend the routine monitoring of these patients to establish cardiopulmonary testing, detect early complications, and provide timely treatment. Importantly, observational studies and clinical trials reporting data on acute and post-acute infectious phases will help elucidate the health outcomes, attributable to COVID-19 and underlying medical conditions.

Appendix

Hs-CRP

Hs-cTnI

IL6

INR

INF

IV Ig

LDH

LPV

LVEF

LMWH

LVEDVi

LYMPH

List of Abbrev	viations
1C	first case
2C	second case
ABGq	arterial blood gases
AD	axial diffusivity
ALT	aminotransferase
AHG	arbidol hydrochloride granules
BPM	breaths per minute
CMR	cardiac magnetic resonance
CP	convalescent plasma
CRP	C-reactive protein
CT	computed tomography
DLCO	diffusing capacity of the lung for carbon
	monoxide
DTI	diffusion tensor imaging
DVT	deep vein thrombosis
ECHO	echocardiography
ECMO	extracorporeal membrane oxygenation
ELISA	enzyme-linked immunoassay
ESR	erythrocyte sedimentation rate
FA	fractional anisotropy
FEV1	FORCED expiratory volume in 1 s
FO	foramen ovale
FT3	free triiodothyronine
FT4	qfree thyroxine
FVC	forced vital capacity
GGOs	ground glass opacity
GMV	bilateral gray matter volumes
Hb	hemoglobin
HRCT	high-resolution chest tomography
TT ODD	

high-sensitivity C-reactive protein

High-sensitivity cardiac troponin I

international normalized ratio

intravenous immunoglobulin

low molecular weight heparin

LV end diastolic volume index

left ventricular ejection fraction

lactate dehydrogenase

inerfertin-6

interferon

lopinavir

lymphocytes

MCH mean corpuscular hemoglobin MCV mean corpuscular volume

MD mean diffusivity
ms milliseconds
MPD methylprednisolone
MV mechanical ventilation
NIV non-invasive ventilation

NT-pro BNP amino-terminal pro-brain natriuretic

peptide

SAT subacute thyroiditis

OCT optical coherence tomography
PASP pulmonary artery systolic pressure
PCO2 carbon dioxide partial pressure

PE pulmonary embolism

PEEP positive end-expiratory pressure PO2 partial pressure oxygen (Po2)

PT prothrombin time

PTE pulmonary thromboembolism PTTa activated plasma thrombin time

RA right atrium RD radial diffusivity RR respiratory rate

RT-PCR reverse transcriptase-polymerase chain

reaction

RV right ventricular

RVEF right ventricular ejection fraction

SOB shortness of breath Tg thyroglobulin

TgAb thyroglobulin antibodies TLC total lung capacity

TTE transthoracic echocardiogram
TTE transthoracic echocardiography

US ultrasonography
WBC white cell blood count

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Author Contributions

The authors contributed equally to all aspects of the article. AS and ZS are co-guarantors of the paper.

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Supplemental Material

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