Supplementary Online Content

Nasserie T, Hittle M, Goodman SN. Assessment of the frequency and variety of persistent symptoms among patients with COVID-19: a systematic review. *JAMA Netw Open*. 2021;4(5):e2111417. doi:10.1001/jamanetworkopen.2021.11417

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This supplementary material has been provided by the authors to give readers additional information about their work.

eTable 1. Literature Search Strategy

Database	Terms/Code
PubMed	(((COVID-19) OR (SARS-CoV-2) OR (coronavirus) OR (2019-nCoV)) AND ((long-term) OR ("long term") OR ("long haul*") OR ("after recovery") OR (prolong*) OR (persist*)) AND ((outcome*) OR (symptom*) OR (disease*) OR (illness*)) AND ((cohort) OR (follow up) OR (longitudinal)))
Web of Science	Query #1: TS=((COVID-19) OR (SARS-CoV-2) OR (coronavirus) OR (2019-nCoV)) Query #2: TS=((long-term) OR ("long term") OR ("long haul*") OR ("after recovery") OR (prolong*) OR (persist*)) Query #3: TS=((outcome*) OR (symptom*) OR (disease*) OR (illness*)) Query #4: TS=((cohort) OR (follow up) OR (longitudinal)) Final query: #1 AND #2 AND #3 AND #4

eTable 2. Studies Excluded from Review

Reference	Summary Comment for Exclusion
Shang Y, Xu C, Jiang F, et al. Clinical characteristics and changes of chest CT features in 307 patients with common COVID-19 pneumonia infected SARS-CoV-2: A multicenter study in Jiangsu, China. <i>Int J Infect Dis IJID Off Publ Int Soc Infect Dis</i> . 2020;96:157-162. doi:10.1016/j.ijid.2020.05.006	Insufficient follow-up length (maximum of to 3 weeks after symptom onset)
Boscolo-Rizzo P, Borsetto D, Fabbris C, et al. Evolution of altered sense of smell or taste in patients with mildly symptomatic COVID-19. JAMA Otolaryngol Head Neck Surg. 2020;146(8):729-732. doi:10.1001/jamaoto.2020.1379	Insufficient follow-up length (4 weeks after symptom onset)
Guo T, Fan Y, Chen M, et al. Cardiac complications in patients hospitalised with COVID-19. <i>JAMA Cardiol.</i> 2020;5(7):811-818. doi: 10.1001/jamacardio.2020.1017	Does not assess prevalence of persistent symptoms
Cellai M, O'Keefe JB. Characterization of Prolonged COVID-19 Symptoms in an Outpatient Telemedicine Clinic. Open Forum Infect Dis. 2020;7(10):ofaa420. doi:10.1093/ofid/ofaa420	Insufficient follow-up length (6 weeks after symptom onset)
Chung TW-H, Sridhar S, Zhang AJ, et al. Olfactory Dysfunction in Coronavirus Disease 2019 Patients: Observational Cohort Study and Systematic Review. <i>Open</i> <i>Forum Infect Dis.</i> 2020;7(6):ofaa199. doi:10.1093/ofid/ofaa199	Insufficient follow-up length (7-9 days after symptoms subsided)
Cicco S, Vacca A, Cittadini A, Marra AM. Long-Term Follow-Up May be Useful in Coronavirus Disease 2019 Survivors to Prevent Chronic Complications. Infect Chemother. 2020;52(3):407-409. doi:10.3947/ic.2020.52.3.407	Review of current knowledge about chronic complications among survivors
Frija-Masson J, Debrary MP, Gilbert M, et al. Functional characteristics of patients with SARS-CoV-2 pneumonia at 30 days post-infection. <i>Eur Respir J</i> . 2020;56(2). doi: 10.1183/13993003.01754-2020	Insufficient follow-up length (30 days after infection)

George PM, Barratt SL, Condliffe R, et al. Respiratory follow-up of patients with COVID-19 pneumonia. <i>Thorax</i> . 2020, doi: 10.1136/thoraxjnl-2020-215314	Review of current knowledge about long-term respiratory outcomes
Karimi-Galougahi M, Safavi Naini A, Ghorbani J, Raad N, Raygani N. Emergence and Evolution of Olfactory and Gustatory Symptoms in Patients with COVID-19 in the Outpatient Setting. <i>Indian J Otolaryngol Head Neck Surg</i> . Published online September 28, 2020:1-7. doi:10.1007/s12070-020-02166-4	Insufficient follow-up length (two weeks after initial symptoms)
Khoo B, Tan T, Clarke SA, et al. Thyroid function before, during and after COVID- 19. <i>J Clin Endocrinol Metab</i> . Published online November 12, 2020. doi:10.1210/clinem/dgaa830	Insufficient length of follow-up for some patients (IQR is 52 to 108 days after hospital admission)
Konstantinidis I, Delides A, Tsakiropoulou E, Maragoudakis P, Sapounas S, Tsiodras S. Short-term follow-up of self-isolated COVID-19 patients with smell and taste dysfunction in Greece: Two phenotypes of recovery. ORL. Published online October 13, 2020:1-9. doi:10.1159/000511436	Insufficient follow-up length (4 weeks after diagnosis)
Mizrahi B, Shilo S, Rossman H, et al. Longitudinal symptom dynamics of COVID-19 infection. <i>Nature Comm</i> . 2020:11, 6208. doi: 10.1038/s41467-020-20053-y	Insufficient follow-up length (mean 31.4 \pm 20.1 days after discharge)
Małek ŁA, Marczak M, Miłosz-Wieczorek B, et al. Cardiac involvement in consecutive elite athletes recovered from Covid-19: A magnetic resonance study. <i>J Magn Reson Imaging JMRI</i> . Published online January 20, 2021. doi:10.1002/jmri.27513	Insufficient follow-up length (median: 32 days after diagnosis, IQR: 22 to 62 days)
Manson JJ, Crooks C, Naja M, et al. COVID-19-associated hyperinflammation and escalation of patient care: a retrospective longitudinal cohort study. <i>Lancet Rheumatol</i> . 2020;2(10):e594-e602. doi:10.1016/S2665-9913(20)30275-7	Length of follow-up is not clearly reported. Minimum follow time is provided, but not range

Mazza MG, De Lorenzo R, Conte C, et al. Anxiety and depression in COVID-19 survivors: Role of inflammatory and clinical predictors. <i>Brain Behav Immun</i> . 2020;89:594-600. doi:10.1016/j.bbi.2020.07.037	Insufficient follow-up length (mean 31.29 +/ 15.7 days after discharge)
Negrini F, Ferrario I, Mazziotti D, et al. Neuropsychological features of severe hospitalized COVID-19 patients at clinical stability and clues for post-acute rehabilitation. Arch Phys Med Rehabil. Published online September 26, 2020. doi:10.1016/j.apmr.2020.09.376	Insufficient follow-up length (minimum of 30 days after hospital admission)
Paderno A, Mattavelli D, Rampinelli V, et al. Olfactory and gustatory outcomes in COVID-19: A prospective evaluation in nonhospitalized subjects. Otolaryngol Head Neck Surg. Published online June 30, 2020. doi:10.1177/0194599820939538	Insufficient follow-up length (maximum 45 days after symptom onset)
Panda S, Mohamed A, Sikka K, et al. Otolaryngologic Manifestation and Long-Term Outcome in Mild COVID-19: Experience from a Tertiary Care Centre in India. <i>Indian</i> <i>J Otolaryngol Head Neck Surg Off Publ Assoc Otolaryngol India</i> . Published online October 14, 2020:1-6. doi:10.1007/s12070-020-02217-w	Insufficient follow-up length (1 month after hospital admission)
Pellaud C, Grandmaison G, Pham Huu Thien HP, et al. Characteristics, comorbidities, 30-day outcome and in-hospital mortality of patients hospitalised with COVID-19 in a Swiss area - a retrospective cohort study. <i>Swiss Med Wkly</i> . 2020;150:w20314. doi:10.4414/smw.2020.20314	Insufficient follow-up length (7-9 days after symptom onset)
Prescott C, Sussman JB, Joost Wiersinga W. Post-critical illness vulnerability. Curr Opin Crit Care. 2020;26(5):500-507. doi: 10.1097/MCC.000000000000076	Review of current knowledge about long-term outcomes among critical illness survivors
Rajpal S, Tong MS, Borchers J, et al. Cardiovascular Magnetic Resonance Findings in Competitive Athletes Recovering From COVID-19 Infection. JAMA Cardiol. Published online September 11, 2020. doi:10.1001/jamacardio.2020.4916	Insufficient follow-up length (some patients only followed for 11 days after diagnosis)

Shaw B, Daskareh M, Gholamrezanezhad A. The lingering manifestations of COVID-19 during and after convalescence: update on long-term pulmonary consequences of coronavirus disease 2019 (COVID-19). Radiol Med (Torino). Published online October 1, 2020. doi:10.1007/s11547-020-01295-8	Review of current knowledge about long-term pulmonary outcomes
Tenforde MW. Symptom Duration and Risk Factors for Delayed Return to Usual Health Among Outpatients with COVID-19 in a Multistate Health Care Systems Network — United States, March–June 2020. <i>MMWR Morb Mortal Wkly Rep</i> . 2020;69. doi:10.15585/mmwr.mm6930e1	Insufficient follow-up length (14-21 days after test date)
Wang X, Xu H, Jiang H, et al. Clinical features and outcomes of discharged coronavirus disease 2019 patients: a prospective cohort study. <i>QJM Mon J Assoc Physicians</i> . 2020;113(9):657-665. doi:10.1093/qjmed/hcaa178	Insufficient follow-up length (some patients only followed for 3 weeks)
Xia L, Chen J, Friedemann T, et al. The Course of Mild and Moderate COVID-19 Infections-The Unexpected Long-Lasting Challenge. <i>Open Forum Infect Dis</i> . 2020;7(9):ofaa286. doi:10.1093/ofid/ofaa286	Insufficient follow-up length (minimum of one month after hospital admission)
Yan CH, Prajapati DP, Ritter ML, DeConde AS. Persistent Smell Loss Following Undetectable SARS-CoV-2. <i>Otolaryngol Head Neck Surg</i> . Published online June 9, 2020:194599820934769. doi:10.1177/0194599820934769	Insufficient follow-up length (average of 16.2 days after diagnosis, IQR (9-22.3 days)
Li Y, Li M, Wang M, et al. Acute cerebrovascular disease following COVID-19: a single center, retrospective, observational study. Stroke Vasc Neurol. 2020;5(3):279-284. doi:10.1136/svn-2020-000431	Length of follow-up is not clearly reported
Yang SS, Lipes J, Dial S, et al. Outcomes and clinical practice in patients with COVID-19 admitted to the intensive care unit in Montréal, Canada: a descriptive analysis. <i>CMAJ Open</i> . 2020;8(4):E788-E795. doi:10.9778/cmajo.20200159	Does not assess prevalence of persistent symptoms

Zou R, Chen F, Chen D, Xu C-L, Xiong F. Clinical characteristics and outcome of hemodialysis patients with COVID-19: a large cohort study in a single Chinese center. Ren Fail. 2020;42(1):950-957. doi:10.1080/0886022X.2020.1816179	Insufficient follow-up length (3 weeks for some patients)
Hopkins C, Surda P, Vaira LA, et al. Six month follow-up of self-reported loss of smell during the COVID-19 pandemic. Rhinology. Published online December 15, 2020. doi:10.4193/Rhin20.544	Study sample was not limited to individuals who tested positive for COVID-19
Goërtz YMJ, Van Herck M, Delbressine JM, et al. Persistent symptoms 3 months after a SARS-CoV-2 infection: the post-COVID-19 syndrome? ERJ Open Res. 2020;6(4). doi:10.1183/23120541.00542-2020	Selected for patients with existing persistent symptoms
Lovato A, Galletti C, Galletti B, de Filippis C. Clinical characteristics associated with persistent olfactory and taste alterations in COVID-19: A preliminary report on 121 patients. Am J Otolaryngol. 2020;41(5):102548. doi:10.1016/j.amjoto.2020.102548	Insufficient follow-up length (mean of 38 days after diagnosis)
Sheng W-H, Liu W-D, Wang J-T, Chang S-Y, Chang S-C. Dysosmia and dysgeusia in patients with COVID-19 in northern Taiwan. J Formos Med Assoc Taiwan Yi Zhi. 2021;120(1 Pt 2):311-317. doi:10.1016/j.jfma.2020.10.003	Assesses median duration of symptoms, not prevalence of persistent symptoms
Reiter ER, Coelho DH, Kons ZA, Costanzo RM. Subjective smell and taste changes during the COVID-19 pandemic: Short term recovery. Am J Otolaryngol. 2020;41(6):102639. doi:10.1016/j.amjoto.2020.102639	Insufficient follow-up length (1 month after symptom onset)
Han X, Fan Y, Alwalid O, et al. Six-Month Follow-up Chest CT findings after Severe COVID-19. Pneumonia. Radiology. Published online January 26, 2021:203153. doi:10.1148/radiol.2021203153	Assessing outcomes among individuals with pneumonia

Brandão Neto D, Fornazieri MA, Dib C, et al. Chemosensory Dysfunction in COVID- 19: Prevalences, Recovery Rates, and Clinical Associations on a Large Brazilian Sample. OtolaryngolHead Neck Surg Off J Am Acad Otolaryngol-Head Neck Surg. Published online September 1, 2020:194599820954825. doi:10.1177/0194599820954825	Insufficient follow-up length for some patients (36 to 119 days after symptom onset)
Liu D, Baumeister RF, Veilleux JC, et al. Risk factors associated with mental illness in hospital discharged patients infected with COVID-19 in Wuhan, China. Psychiatry Res. 2020;292:113297. doi:10.1016/j.psychres.2020.113297	Length of follow-up unclear (minimum not reported)
Udwadia ZF, Koul PA, Richeldi L. Post-COVID lung fibrosis: The tsunami that will follow the earthquake. <i>Lung India</i> . 2021;38(Supplement):S41-S47. doi:10.4103/lungindia.lungindia_818_20	Review of current knowledge about long term pulmonary outcomes
Trinkmann F, Müller M, Reif A, et al. Residual symptoms and lower lung function in patients recovering from SARS-CoV-2 infection. <i>Eur Respir J</i> . 2021;57(2). doi:10.1183/13993003.03002-2020	Insufficient follow up length (68 ± 16 days after symptom onset)
Makaronidis J, Firman C, Magee CG, et al. Distorted chemosensory perception and female sex associate with persistent smell and/or taste loss in people with SARS-CoV-2 antibodies: a community based cohort study investigating clinical course and resolution of acute smell and/or taste loss in people with and without SARS-CoV-2 antibodies in London, UK. <i>BMC Infect Dis.</i> 2021;21(1):221. doi:10.1186/s12879-021-05927-w	Insufficient follow up length (4-6 weeks after positive test)

Lampl BMJ, Buczovsky M, Martin G, Schmied H, Leitzmann M, Salzberger B. Clinical and epidemiological data of COVID-19 from Regensburg, Germany: a retrospective analysis of 1084 consecutive cases. <i>Infection</i> . Published online March 5, 2021. doi:10.1007/s15010-021-01580-2	Insufficient follow up length (6 weeks after symptom onset)
Jaywant A, Vanderlind WM, Alexopoulos GS, Fridman CB, Perlis RH, Gunning FM. Frequency and profile of objective cognitive deficits in hospitalized patients recovering from COVID-19. <i>Neuropsychopharmacology</i> . Published online February 15, 2021. doi:10.1038/s41386-021-00978-8	Insufficient follow up length (43 days after hospital admission)
Zhou M, Wong C-K, Un K-C, et al. Cardiovascular sequelae in uncomplicated COVID-19 survivors. <i>PLoS One</i> . 2021;16(2):e0246732. doi:10.1371/journal.pone.0246732	Insufficient follow up length (1-4 weeks after hospital discharge)
Logue JK, Franko NM, McCulloch DJ, et al. Sequelae in Adults at 6 Months After COVID-19 Infection. <i>JAMA Netw Open</i> . 2021;4(2):e210830. doi:10.1001/jamanetworkopen.2021.0830	Insufficient follow up length (less than 30 days or 30-60 days after symptom onset for some patients)
Hall J, Myall K, Lam JL, et al. Identifying patients at risk of post-discharge complications related to COVID-19 infection. <i>Thorax</i> . Published online February 4, 2021. doi:10.1136/thoraxjnl-2020-215861	Selected for patients with persistent symptoms
Stavem K, Ghanima W, Olsen MK, Gilboe HM, Einvik G. Persistent symptoms 1.5-6 months after COVID-19 in non-hospitalised subjects: a population-based cohort study. <i>Thorax</i> . Published online December 3, 2020. doi:10.1136/thoraxjnl-2020-216377	Insufficient follow up length (1.5-6 months after symptom onset)

Leite VF, Rampim DB, Jorge VC, et al. Persistent symptoms and disability after	Insufficient follow up length (21.8±11.7 days after hospital discharge)
COVID-19 hospitalization: data from a comprehensive telerehabilitation program.	
Arch Phys Med Rehabil. Published online March 9, 2021.	
doi:10.1016/j.apmr.2021.03.001	

eTable 3. Study and Patient Characteristics

Study	Country	Cohort Diagnostic Type, Criteria	Cohort Type, Study	Severity of COVID-19 illness			Patient Demograph	ics & Unde	erlying Comorbidities	
		Arms		Asymp %	Mild or Moderate - %	Severe - %	Critical - %	Age in Years - Mean (SD)	Male - %	BMI & Comorbidities
Akter ¹	Bangladesh	Non- concurrent, single-arm	Positive SARS-CoV- 2 real-time reverse transcriptas e PCR	Not reported				NR	76.0	Diabetes: 19.89% Cancer: 1.4% Cardiovascular diseases: 9.1% Respiratory disease: 6.1% Liver diseases: 2.2% Heart attack history: 2.5% Other chronic diseases: 7.9%
Arnold ²	United Kingdom	Concurrent, single-arm	Positive SARS-CoV- 2 real-time reverse transcriptas e PCR or radiological diagnosis	0	84	16	0	47 (range: 32 to 61)	61.8	Hypertension: 15% Mean BMI (kg/m ²): 31.2 Heart disease: 22% Chronic lung disease: 15% Severe liver disease: 0% Severe kidney disease: 3.7% HIV/AIDS: 0%
Carfi ³	Italy	Concurrent, single-arm	Positive SARS-CoV- 2 real-time reverse transcriptas e PCR	Not reported	·			56.5 (14.6)	62.9	Diabetes: 7% Hypertension: 35% BMI (kg/m ²), mean SD: 26.3 (4.4) Chronic heart disease: 4.9% Atrial fibrillation: 2.8% Heart failure: 2.8% Stroke: 1.4% Kidney failure: 2.1% Thyroid disease: 18.2% COPD: 9.1% Active cancer: 3.5% Immune disorders: 11.2%

Carvalho- Schneider ⁴	France	Concurrent, single-arm	Positive SARS-CoV- 2 real-time reverse transcriptas e PCR	0	78	22	0	49 (15)	47.7	Comorbid conditions: None: 49.1% 1: 31.5% 2 or more: 19.2%
Chen⁵	China	Concurrent, single-arm	Lab- confirmed	0	91	9	0	47.2 (13.0)	51.5	BMI, kg/m², mean (SD): 23.64 (3.31) History of chronic disease: 31.9%
Chiesa- Estomba ⁶	Spain	Concurrent, single-arm	Positive SARS-CoV- 2 real-time reverse transcriptas e PCR or positive immunoglo bulin G/M test	0	100	0	0	41 (13)	36.5	Diabetes: 2.5% Heart problems: 2.3% Chronic obstructive pulmonary disorder: 0.8% Hypertension: 6.3% Asthma: 5.6% Hypothyroidism: 6.1% Autoimmune disease: 3.9%
Chopra ⁷	United States	Concurrent, single-arm	Not stated	Not reported				Median: 62 (IQR: 50-72)	51.8	Diabetes: 34.9% Hypertension: 64% Cardiovascular disease: 24.1% Moderate/severe kidney disease: 23% Asthma: 13.4% Congestive heart failure/cardiomyopathy: 11.6% Chronic obstructive pulmonary disease: 10.4% Cerebrovascular disease/paraplegia: 10.4% Dementia: 7.7% Cancer: 7.1% None: 14.3%

D'Cruz ⁸	United Kingdom	Concurrent, single-arm	Positive SARS-CoV- 2 real-time reverse- transcriptas e PCR	0	0	66	34	58.7 (14.4)	62.2	Median BMI ((kg/m ²): 30.0 (IQR: 25.9 to 35.2) Current or former smoker: 25.4% Cardiovascular disease: 45.4% Diabetes: 34.5% Immunosuppressed: 13.4% Obstructive lung disease: 10.9% Malignancy: 10.1% End-stage renal failure: 6.7% Thyroid disease: 5.9% Mental health conditions: 5.0% Cerebrovascular disease: 4.2%
Daher ⁹	Germany	Concurrent, single-arm	Positive SARS-CoV- 2 real-time reverse transcriptas e PCR	Not reported				64 (3)	66.7	Hypertension: 59% Diabetes mellitus: 25% COPD: 9% Bronchial asthma: 13% Heart failure: 9% Atrial fibrillation: 9% Chronic kidney disease: 22% Coronary artery disease: 19%
de Graaf ¹⁰	Netherlands	Concurrent, single-arm	Positive SARS-CoV- 2 real-time reverse- transcriptas e PCR	Not reported				60.8 (13)	63.0	BMI: 27.8 ± 4.5 Hypertension: 34% Hypercholesterolemia: 27% Diabetes: 23% Smoking: 11% Alcohol: 23% Cardiovascular disease: 28% Heart failure: 1% Atrial fibrillation: 5% Valvular heart disease: 6% Myocardial infarction: 4% Stroke or transient ischemic attack: 10% Peripheral vascular disease:

										2% Chronic kidney injury: 11%
Garrigues ¹¹	France	Concurrent, single-arm	Positive SARS-CoV- 2 real-time reverse transcriptas e PCR and/or abnormaliti es on chest CT scan	Not reported				63.2 (15.7)	62.5	Hypertension: 46.7% Diabetes: 21.7% BMI (kg/m ²): <25: 29.2% >/ 25: 47.5% Missing: 23.3%
Gherlone ¹²	Italy	Concurrent, single-arm	Positive SARS-CoV- 2 real-time reverse- transcriptas e PCR	0	0	75.4	24.6	Median: 62.5 (IQR: 53.9 to 74.1)	75.4	Hypertension: 41% Coronary artery disease: 9.8% Diabetes mellitus: 13.9% Chronic kidney disease: 7.4% Neoplasia: 5.7% Chronic obstructive pulmonary disease: 6.6% Smoking: 39.3%
Gonzalez ¹³	Spain	Concurrent, single-arm	Not reported	0	0	0	100	Median: 60 (IQR: 48 to 65)	74.2	Median BMI ((kg/m ²): 28.2 (IQR: 25.4 to 32.6) Current or former smoker: 56.7% Hypertension: 37.1% Diabetes mellitus: 14.5% Chronic heart disease: 9.7% Asthma: 4.8% COPD: 4.8%
Halpin ¹⁴	United Kingdom	Concurrent, single-arm	Positive SARS-CoV- 2 real-time reverse transcriptas e PCR	Not reported		- -	·	Ward - Median: 70.5 (range: 20 to 93) ICU - Median: 58.5 (range: 34 to 84)	54.0	BMI: Underweight: 3% Healthy weight: 25% Overweight: 35% Obese: 24% Unknown: 13%

										Active cancer: 7% Active or previous cancer: 21% Heart failure: 5% Hyperlipidemia: 4% Ischemic heart disease: 10% Hypertension: 41% Tachyarrhythmias: 11% Valvular heart disease: 3% Venous thromboembolism:5%
Huang ¹⁵	China	Concurrent, single-arm	Labconfirm ed	0	25.3	70.7	4	Median: 57 (IQR: 47 to 65)	51.8	Hypertension: 29% Diabetes: 12% Cardiovascular diseases: 7% Cerebrovascular diseases: 3% Malignant tumour: 3% COPD: 2% Chronic kidney disease: 2% Past smoker: 3% Current smoker: 6%
Jacobs ¹⁶	United States	Concurrent, single-arm	Positive SARS-CoV- 2 real-time transcriptas e PCR	0	87.4	12.6	0	Median: 57 (IQR: 48 to 68)	60.9	Overweight: 36.1% Obese: 49.2% Hypertension: 47.5% Diabetes: 28.4% CAD or history of MI: 11.5% Arrhythmia: 4.9% Heart failure: 2.7% Hyperlipidemia: 10.9% Asthma: 10.4% Cancer: 9.8% Immunodeficiency: 4.4% Hypothyroidism: 4.4% Psychiatric disorders: 4.4% Obstructive sleep apnea: 3.3% COPD: 3.8% Thromboembolic disease: 1.6%

Lechien ¹⁷	Belgium	Concurrent, single-arm	Positive SARS-CoV- 2 real-time transcriptas e PCR	0	100	0	0	46.2 (11.2)	33.0	Diabetes: 2.7% Hypertension: 6.8% Asthma: 6.8% Reflux: 10.8% Allergic rhinitis: 6.8% Hypothyroidism: 2.7%
Lerum ¹⁸	Norway	Concurrent, single-arm	Diagnostic codes U07.1, U07.2, or J12.x	Not reported				Median: 59 (IQR: 49 to 72)	52.4	Median BMI (IQR): 25.8 (23.9 to 29.6) Current smoker: 3.4% Past smoker: 39% Hypertension: 35% Diabetes: 8%
Liang ¹⁹	China	Concurrent, single-arm	Positive SARS-CoV- 2 real-time reverse transcriptas e PCR	0	91	9		41.3 (13.8)	27.6	Hypertension: 6.6% Cardiovascular disease: 1.2% Diabetes: 3.9% Thyroid disease: 3.6% Pulmonary disease: 14.5% Digestive system disease: 18.4% Smoking: 0%
Lu ²⁰	China	Concurrent, two-arm	Positive SARS-CoV- 2 real-time reverse transcriptas e PCR	0	78	20	2	44.1 (16)	56.7	Hypertension: 21.67% Diabetes: 10.00%
Mandal ²¹	United Kingdom	Concurrent, single-arm	Positive SARS-CoV- 2 real-time reverse transcriptas e PCR	0	67	30	0	59.9 (16.1)	62.0	Hypertension: 41% Diabetes mellitus: 27.2% Asthma and/or COPD: 17.5% Chronic kidney disease: 11% Ischaemic heart disease: 9.7% Obese: 8% None: 34%
Mazza ²²	Italy	Concurrent, single-arm	Positive SARS-CoV-	Not reported	58.5 (12.8)	65.9	Not reported			

			2 real-time reverse- transcriptas e PCR							
Mendez ²³	Spain	Non- concurrent, single-arm	Positive SARS-CoV- 2 real-time reverse- transcriptas e PCR	Not reported				Median: 57 (IQR: 49 to 67)	58.7	Hypertension: 32.4% Diabetes: 16.2% Dyslipidemia: 29/1% Chronic heart disease: 5.6% Chronic renal disease: 1.7% Chronic liver disease: 1.7% Cancer: 1.7% Chronic respiratory disease: 11.7%
Moreno- Perez ²⁴	Spain	Concurrent, single-arm	Positive SARS-CoV- 2 real-time reverse transcriptas e PCR	0	34.3	65.7	0	Median: 62 (range: 53 to 72)	52.7	Hypertension: 36.5% Diabetes: 11.6% Obesity: 30.6% Cardiovascular disease: 6.9% Chronic respiratory disease: 18.1% Immunosuppression: 4.1%
Munro ²⁵	United Kingdom	Concurrent, single-arm	Positive SARS-CoV- 2 real-time reverse transcriptas e PCR or chest radiology consistent with COVID-19	Not reported				Not reported	NR	Not reported
Nguyen ²⁶	France	Concurrent, single-arm	Positive SARS-CoV- 2 real-time reverse transcriptas	Not reported				Median: 36 (IQR: 27 to 48)	44.8	Hypertension: 12.8% Diabetes: 9.6% Chronic respiratory disease: 12% Chronic cardiac disease: 2.4%

			e PCR or serological test							Cancer: 0.8% Obesity: 9.6%
Poncet- Megemont ² 7	France	Non- concurrent, single-arm	Lab confirmed or based on chest CT scan	0	59	37	4	48.5 (15.3)	37.4	Not reported
Puntmann ²⁸	Germany	Concurrent, two-arm	Positive SARS-CoV- 2 real-time reverse transcriptas e PCR	18	49	33	0	49 (14)	53.0	Hypertension: 22% Diabetes: 18% BMI, (kg/m ²), median (IQR): 25 (23 to 28) Hypercholesterolemia: 22% Known coronary artery disease: 135 Smoking: 22% COPD or asthma: 21%
Qu ²⁹	China	Concurrent, single-arm	Positive SARS-CoV- 2 real-time reverse- transcriptas e PCR	0	90.6	9.4	0	Median: 47.5 (IQR: 37 to 57)	50.0	Not reported
Raman ³⁰	United Kingdom	Concurrent, two-arm	Positive SARS-CoV- 2 real-time reverse transcriptas e PCR	Not reported				55.4 (13.2)	58.6	Hypertension: 37.9% Diabetes: 15.5% Current/ex smoker: 34.5% Coronary artery disease: 3.4% Cerebrovascular disease: 1.7% Asthma: 34.5% COPD: 5.2% Previous cancer: 3.4% Depression: 5.2%

Rosales- Castillo ³¹	Spain	Non- concurrent, single-arm	Positive SARS-CoV- 2 real-time reverse transcriptas e PCR or serological test	Not reported				60.16 (15.08)	55.9	Hypertension: 50% Hypercholesterolemia (28%) Type 2 diabetes: 22% Asthma: 14.4% Sleep apnea-hypopnea syndrome: 8.5% COPD: 6% Ischemic heart disease: 6% Chronic kidney disease: 6%
Shah ³²	Canada	Concurrent, single-arm	Positive SARS-CoV- 2 real-time reverse transcriptas e PCR	Not reported				Median: 67 (IQR: 54 to 74)	68.3	Median BMI: 25 (IQR: 23-29) Hypertension: 35% Diabetes: 22% Chronic pulmonary disease: 13% Coronary heart disease: 10% Malignancy: 10% Chronic kidney disease: 7% Ever smoker: 38%
Sonnwebbe r, Eur Resp J ³³	Austria	Concurrent, single-arm	Positive SARS-CoV- 2 real-time reverse transcriptas e PCR and typical clinical presentatio n	0	50.3	27.6	22.1	57 (14)	54.9	None: 23% Mean BMI (SD): 26 (5) Current smoker: 3% Past smoker: 39% Hypertension: 30% Diabetes mellitus: 17% Cardiovascular disease: 40% Chronic obstructive pulmonary disease: 6% Asthma: 7% Interstitial lung disease: 1% Hypercholesterolemia: 19% Chronic kidney disease: 7% Chronic liver disease: 6% Malignancy: 12% Immunodeficiency: 6%
Sonnweber, Resp Res ³⁴	Austria	Concurrent, single-arm	Positive SARS-CoV- 2 real-time	0	51	32	17	58 (14)	59.6	Hypertension: 29% Diabetes mellitus: 18% Cardiovascular disease: 40%

			reverse transcriptas e PCR Symptom presentatio n							Pulmonary disease: 19% Endocrine disease: 45% Chronic kidney disease: 6% Chronic liver disease: 6% None: 19%
Sykes ³⁵	England	Concurrent, single-arm	Positive SARS-CoV- 2 real-time reverse transcriptas e PCR	Not reported				59.6 (14.0)	65.7	Type 1 diabetes: 0.7% Type 2 diabetes: 22.0% Ischaemic heart disease: 16.4% COPD: 8.2% Asthma: 14.2% Hypertension: 41.0% CKD: 4.5% History of venous thromboembolism: 2.2% Cancer: 5.2% Cardiovascular disease: 4.5% Smoking history: 44.0% Alcohol use: 42.5%
Taboada ³⁶	Spain	Concurrent, single-arm	Positive SARS-CoV- 2 real-time reverse- transcriptas e PCR	0	0	0	100	65.5 (10.4)	64.8	Hypertension: 55.5% Hyperlipidemia: 44% Diabetes: 23.1% Asthma: 6.6% COPD: 8.8% Heart disease: 20.9% Obesity: 38.5%
Tomasoni ³⁷	Italy	Concurrent, single-arm	Positive SARS-CoV- 2 real-time reverse transcriptas e PCR	Not reported				55 (range: 43 to 65)	73.3	Charlson Comorbidity Score, median (IQR): 1 (0 to 2.5)
Townsend ³⁸	Ireland	Concurrent, single-arm	Positive SARS-CoV-	Not reported				49.15 (15)	46.1	BMI (kg/m²), mean (+/- SD): 28.7 +/- 5.3

			2 real-time reverse transcriptas e PCR							Total number of comorbidities, median (IQR): 1 (0-2)
Ugurlu ³⁹	Turkey	Concurrent, single-arm	Positive SARS-CoV- 2 real-time transcriptas e PCR	Not reported				41.2 (14.6)	45.2	Not reported
Vaira ⁴⁰	Italy	Concurrent, single-arm	Positive SARS-CoV- 2 real-time reverse transcriptas e PCR	Not reported				51.2 (8.8)	49.3	Diabetes: 10.9% BMI > 30 kg/m ² : 29% Cardiovascular disorder: 26.8% Pulmonary disorder: 15.2%
van den Borst ⁴¹	Netherlands	Concurrent, single-arm	Positive SARS-CoV- 2 real-time reverse transcriptas e PCR or clinical diagnosis	0	62.9	21	16.1	59 (14)	59.7	Diabetes mellitus: 14% Hypertension: 28% Cardiovascular disease: 24% Asthma: 10% Chronic obstructive pulmonary disease: 6% Other lung disease: 3% Immunocompromised: 15% Chronic kidney failure: 8% Cancer: 20%
Weerahand j ⁴²	United States	Concurrent, single-arm	Lab- confirmed	0	0	100	0	62 (IQR: 50 to 67)	62.5	Hypertension: 60.25% Diabetes: 36.65% Chronic kidney disease: 8.07% Cancer: 7.45% Coronary artery disease: 9.32% Heart failure: 4.97% Hyperlipidemia: 46.58% Asthma or COPD: 24.22% BMI (kg/m ²):

										<25: 14.29% 25 to <30: 30.43% 30 to <40: 40.99% >/ 40: 13.66% Missing: 0.62%
Wong ⁴³	Canada	Concurrent, single-arm	Lab- confirmed	Not reported				62 (16)	64.1	Diabetes: 26% Pulmonary disease: 8% History of heart attack: 8% None: 59%
Xiong ⁴⁴	China	Concurrent, two-arm	Not stated	0	61.5	33.5	5	Median: 52 (IQR: 95 - 102)	45.5	Hypertension: 15.2% Diabetes: 7.4% Chronic obstructive lung disease: 4.1% Coronary heart disease: 3.3% Chronic kidney disease: 2.2% Carcinoma: 0.9%
Zhao ⁴⁵	China	Non- concurrent, single-arm	Positive SARS-CoV- 2 real-time reverse transcriptas e PCR	0	93	7	0	47.7 (15.5)	58.2	Diabetes mellitus: 3.64% Hypertension: 10.91% Cardiovascular disease: 3.64% Smoking: 4% Pulmonary diseases: 0%

eTable 4. Selection Criteria

Study	Source of Participants	Inclusion & Exclusion Criteria	No. of Reported Eligible Participants	Number Excluded & Reasons for Non- Participation	Final Sample Size (Proportion of Total Eligible)
Akter ¹	Hospitalized: 100% (% in non-ICU and ICU not indicated)	Inclusion: Patients with confirmed infection	Not reported	Not reported	734 (NR)
Arnold ²	Hospitalized: 100% (% in non-ICU and ICU not indicated)	Inclusion: Hospitalized patients Exclusion: <18 years old; inability to consent to study participation; nursing home residents; unwilling or unable to undergo chest CT or provide blood sample	131	N=21 Declined follow-up (N=18) Unreachable (N=3)	110 (83.9%)
Carfi ³	Hospitalized: 87.4% in non-ICU, 12.6% in ICU	Inclusion: Patients discharged from hospital after recovery (based on WHO guidelines)	157	N=14 Declined participation (N=14)	143 (91.1%)
Carvalho- Schneider ⁴	Outpatient: 64.6% Hospitalized: 35.4% (ICU patients excluded)	Inclusion: Patients with confirmed infection Exclusion: <18 years old; ICU patients; residents of nursing homes or long-term care facilities;	174	N=44 Lost to follow-up (N=44)	130 (74.7%)

		patients transferred to other healthcare facilities			
Chen⁵	Hospitalized: 100% (% in non-ICU and ICU not indicated)	Not clearly stated	503	N=142 Lost to follow-up (N=131) Provided incomplete data (N=11)	361 (71.8%)
Chiesa- Estomba ⁶	Hospitalized and outpatient (numbers not provided)	Inclusion: Hospitalized patients or outpatients Exclusion: Admitted to ICU; pre-existing olfactory or gustatory dysfunction	1, 222	N = 471 Incomplete follow-up data (N=362) Unreachable (N=61) Declined participation (N=48)	751 (61.5%)
Chopra ⁷	Hospitalized: 100% (% in non-ICU and ICU not indicated)	Inclusion: Hospitalized patients alive 60 days after discharge Exclusion: Residing in a health care or correctional facility	1167	N = 679 Reasons not reported	488 (41.8%)
D'Cruz ⁸	Hospitalized: 65.5% in non-ICU, 35.5% in ICU	Inclusion: Patients with severe COVID- 19 (hospital length of stay ≥48 h and maximum fraction of inspired oxygen ≥40% or ICU admission) Exclusion: <18 years	143	N = 24 Unreachable or declined participation (N=24)	119 (83.2%)

		old; mild or moderate COVID-19; unwilling or unable to undergo CT scan			
Daher ⁹	Hospitalized: 100% (% in non-ICU and ICU not indicated)	Inclusion: Patients with confirmed infection Exclusion: patients with Acute Respiratory Distress Syndrome who needed mechanical ventilation in the ICU; unwilling or unable to	Not reported	Not reported	33 (NR)
		undergo CT scan, echocardiography, or whole-body plethysmography or provide blood sample			
de Graaf ¹⁰	Hospitalized: 100% (59% in non-ICU, 41% in ICU)	Inclusion: Patients discharged from hospital Excluded: Patients transferred to hospice; re-admitted patients; <18 years old; unable or unwilling to undergo chest CT scan	98	N = 17 Declined follow-up due to lack of symptoms (N = 7) Administrative errors (N=10)	81 (82.7%)
Garrigues ¹¹	Hospitalized: 80% in non-ICU, 20% in ICU	Inclusion: Hospitalized patients Exclusion: Deceased, unreachable, demented,	192	N=72 Unreachable (N=69) Declined participation (N=2) Other (N=1)	120 (62.5%)

		bedridden, and non- French speaking patients; patients admitted directly to ICU			
Gherlone ¹²	Hospitalized: 100% (75.4% in non-ICU, 24.6% in ICU)	Inclusions: patients admitted to emergency department Exclusion: <17 years	Not reported	Not reported	122 (NR)
Gonzalez ¹³	Hospitalized: 100% in ICU	Inclusion: Patients discharged from ICU; acute respiratory distress syndrome due to COVID-19 Exclusion: <18 years; transferred to another hospital during hospitalization; palliative care patients; severe mental illnesses; unable or unwilling to undergo CT scan	75	N = 13 Unreachable or refused participation (N=13)	62 (82.7%)
Halpin ¹⁴	Hospitalized: 68% in non-ICU, 32% in ICU	Inclusion: Patients discharged from hospital; resides in Leeds Metropolitan District Exclusion criteria: <18 years old; dementia, learning disability, cognitive or	158	N = 58 Unreachable (N=56) Declined participation (N=2)	100 (63.3%)

		communication impairment			
Huang ¹⁵	Hospitalized: 95.6% in non-ICU, 4.4% in ICU	Inclusion: Patients discharged from hospital Exclusion: individuals with psychotic disorders, dementia; re-admitted to hospital due to underlying diseases; immobile or unable to move freely due to osteoarthropathy, stroke, or embolism; living outside of Wuhan or in nursing or welfare homes; unwilling or unable to undergo CT scan	2, 142	N = 409 Declined participation (N=347) Unreachable (N=62)	1733 (80.9%)
Jacobs ¹⁶	Hospitalized: 100% (% in ICU and non-ICU not stated)	Inclusion: Patients discharged from hospital Exclusion: Hospital stay less than 3 days; non-English speakers; individuals with dementia or delirium	351	N = 168 Reasons not reported	183 (52.1%)
Lechien ¹⁷	Outpatient: 100%	Inclusion: Patients with mild or moderate COVID-19 presenting with sudden loss of smell	95	N = 7 Did not complete study (N=7)	88 (92.6%)

Lerum ¹⁸	Hospitalized: 85.4 in non-ICU, 14.6% in ICU	Inclusion: Patients discharged from hospital Exclusion: <18 years old; admitted for less than 8 hours; live outside the hospitals' catchment areas; inability to provide informed consent; participation in WHO Solidarity trial; unable or unwilling to undergo CT scan	Not reported	Not reported	103 (NR)
Liang ¹⁹	Hospitalized: 90.8% in non-ICU, 9.2% in ICU	Inclusion: Patients discharged from hospital after recovery Exclusion: <18 years old; history of pulmonary resection; neurological or psychiatric disease; unwilling or unable to undergo CT scan or provide blood sample	134	N = 58 Unreachable (N=11) Not nearby during study period (N=6) Declined participation (N=33) Did not complete study (N=8)	76 (56.7%)
Lu ²⁰	Hospitalized: 100% (% in non-ICU and ICU not indicated)	Inclusion: Patients who recovered from COVD-19Recovered COVID-19 patients Exclusion: Unwilling or unable to undergo MRI scan	155	N=95 Reasons not reported	60 (38.7%)
Mandal ²¹	Hospitalized: 85.5% in	Inclusion: Patients	878	N=494	384 (43.7%)

	non-ICU, 14.5% in ICU	with confirmed infection Exclusion: Unwilling to undergo chest radiograph or provide blood sample		Follow-up not completed for logistic reasons (N=430) Unreachable by phone (N=53) Declined follow-up (N=11)	
Mazza ²²	Outpatient: 21.7%* Hospitalized: 78.3% (% in non-ICU and ICU not indicated) *Diagnosed in hospital ER	Inclusion: Patients with confirmed COVID diagnosed in the emergency department Exclusion: <18 years old	402	N=176 Reasons not reported	226 (56.2%)
Mendez ²³	Hospitalized: 81% in non-ICU, 19% in ICU	Inclusion: Hospitalized patients Exclusion: ≥ 85 years old or <18 years old; non-Spanish speaking; nursing home residents; pre- existing dementia or cognitive decline or brain injury; current alcohol or substance use disorder (except nicotine); history of major psychiatric disorders	216	N = 37 Declined participation (N=5) Unreachable (N=18) Withdrew consent (N=9) Did not complete study (N=5)	179 (82.9%)
Moreno- Perez ²⁴	Hospitalized: 91.3% in non-ICU, 8.7% in ICU	Inclusion: Hospitalized patients Exclusion: Severe comorbidities	326	N = 49 Declined participation (N=4) Did not attend face- to-face assessment (N=15)	277 (85.0%)

				Lost to follow-up (N=30)	
Munro ²⁵	Hospitalized: 98.3% in non-ICU, 1.7% in ICU	Inclusion: Patients discharged from hospital	Not reported	Not reported	121 (NR)
Nguyen ²⁶	Outpatient: 100%	Inclusion: Patients who reported anosmia and/or ageusia in the acute phase of infection	200	N=75 Lost to follow-up (N=75)	125 (62.5%)
Poncet- Megemont ²⁷	Outpatient: 54.7% Hospitalized: 41% non- ICU, 4.3% ICU	Inclusion: Patients with confirmed infection	161	N=22 Declined participation (N=5) Unreachable (N=17)	139 (86.3%)
Puntmann ²⁸	Outpatient: 67% Hospitalized: 33% (% in non-ICU and ICU not indicated)	Inclusion: Minimum of 2 weeks from original diagnosis; resolution of respiratory symptoms; negative result on COVID-19 swab test Exclusion: Unwilling or unable to take MRI scan	Not reported	Not reported	100 (NR)
Qu ²⁹	Hospitalized: 100% (% in non-ICU and ICU not indicated)	Inclusion: Patients discharged from hospital Exclusion: Patients transferred to another medical facility for treatment of concurrent issues;	573	N = 33 Declined participation (N=14) Did not complete study (N=19)	540 (94.2%)

		confirmed hepatitis B, C, AIDS or other viral infections; pregnant women			
Raman ³⁰	Hospitalized: 64% in non-ICU, 36% in ICU	Inclusion: patients with moderate to severe confirmed infection	Not reported	Not reported	58 (NR)
		Exclusion: unable or unwilling to undergo MRI; severe comorbidities			
Rosales- Castillo ³¹	Hospitalized: 92.4% in non-ICU, 7.6% in ICU	Inclusion: Patients discharged from hospital	Not reported	Not reported	118 (NR)
Shah ³²	Hospitalized: 100% (% in non-ICU and ICU not reported)	Inclusion: Hospitalized patients Exclusion: Unwilling or unable to undergo CT scan	82	N = 22 Declined participation (N=10) Unreachable (N=7) Did not complete study (N=5)	60 (73.2%)
Sonnwebber, Eur Resp J ³³	Outpatient: 25.6% Hospitalized: 52.6% in non-ICU, 21.8% in ICU	Inclusion: Hospitalized patients or outpatients with persisting symptoms	190	N = 57 Unable to participate (N=27) Unavailable for follow up (N=12) Declined participation (N=18)	133 (70 %)
Sonnweber, Resp Res ³⁴	Outpatient: 20.2% Hospitalized: 63.3% in non-ICU, 16.5% in ICU	Inclusion: Patients with confirmed infection Exclusion: Unwilling	186	N=77 Lost to follow-up (N=59) Declined participation (N=18)	109 (58.6%)

		to have blood sample taken or to undergo CT scan			
Sykes ³⁵	Hospitalized: 100% (79.9% in non-ICU, 20.1% in ICU)	Inclusion: Patients with confirmed infection treated for COVID-19 pneumonia Exclusion: care home residents; Clinical Frailty Score ≥ 6;	190	N=56 Lost to follow-up (N=56)	134 (70.5%)
Taboada ³⁶	Hospitalized: 100% (100% in ICU)	Inclusion: Patients with COVID-19 induced acute respiratory distress syndrome requiring treatment in an ICU	92	N = 1 Declined participation (N=1)	91 (98.9%)
Tomasoni ³⁷	Hospitalized: 100% (% in non-ICU and ICU not indicated)	Inclusion: Patients who recovered from COVID-19 Recovered patients COVID-19 patients	Not reported	Not reported	105 (NR)
Townsend ³⁸	Outpatient: 44.5% Hospitalized: 41.4% in non-ICU, 14.1% in ICU	Inclusion: Minimum of 6 weeks after date of last acute COVID- 19 symptoms for outpatients or date of discharge for patients admitted to hospital during illness	223	N=95 Reasons not reported	128 (57.4%)
Ugurlu ³⁹	Hospitalized: 100% (ICU patients	Inclusion: Hospitalized patients	42	N=0	42 (100%)

	excluded)	with olfactory dysfunction Exclusion: <18 years of age or >60 years of age; history of nasal surgery or olfactory dysfunction; chronic sinusitis; neurological or psychiatric diseases			
Vaira ⁴⁰	Outpatient: 77% Hospitalized: 23% non- ICU (*ICU patients excluded)	Inclusion: severe COVID-19 diagnosis; symptomatic patients presenting within 4 days of symptom onset Exclusion: <18 years old; admitted to ICU; patients with history of previous trauma, surgery, or radiotherapy in oral and nasal cavities; patients with allergic rhinitis or rhinosinusitis; patients with mental health illnesses	146	N=8 Lost to follow-up (N=8)	138 (94.5%)
van den Borst ⁴¹	Outpatient: 21.8% Hospitalized: 62.1% in non-ICU, 16.1% in ICU	Inclusion: Hospitalized patients, or patients referred by general practitioners Exclusion: unable or	197	N = 73 Declined participation (N=69) Unreachable (N=2) Unknown (N=2)	124 (62.9%)

		unwilling to undergo chest CT (hospitalized patients) or to provide blood sample			
Weerahandi ⁴²	Hospitalized: 53.9% in non-ICU, 46.1% in ICU	Inclusion: Hospitalized patients needing at least 6L of oxygen at any point during hospitalization Exclusion: <18 years old; deceased, demented, or communication impaired patients; patients discharged to hospice; residents of long-term care facilities prior; patients fully dependent in daily living activities prior to hospitalization; re- hospitalized	390	N=238 Unreachable (N=135) Declined participation (N=94) Lost to follow-up (N=9)	152 (39.0%)
Wong ⁴³	Hospitalized: 100% (% in non-ICU and ICU not indicated)	Inclusion: Hospitalized patients Exclusion: <18 years old; inability to complete surveys in English	96	N=18 Declined participation (N=10) Could not complete questionnaires because seen virtually (N=8)	78 (81.3%)
Xiong ⁴⁴	Hospitalized: 100% (% in non-ICU and ICU not stated)	Inclusion: Patients discharged from hospital	706	N = 168 Unreachable (N=87) Declined participation (N=75)	538 (76.2%)

		Exclusion: <20 years old or >80 years old; pregnant or lactating patients; transferred to another hospital for treatment; recent surgery or chemotherapy; presence of other serious disease; incomplete medical record		Inability to describe symptoms clearly (N=6)	
Zhao ⁴⁵	Hospitalized: 100% (*ICU patients excluded)	Inclusion: Patients with confirmed infection Exclusion: <18 years old; ICU patients; unable or unwilling to undergo chest CT	73	N=18 Unreachable (N=4) Declined participation (N=14)	55 (75.3%)

eTable 5. Follow-Up and Outcome Measurement

Study	Time Zero	Length of Final Follow-Up	Determination of End of Follow-Up	Outcome Measurement
Akter ¹	Recovery	4 weeks after recovery	All participants followed for same amount of time	Self-report
Arnold ²	Hospital admission	8 to 12 weeks after admission Minimum: 56 days	Varied for each participant; based on date of examination	X-ray (chest abnormalities); blood sample (lab assessments); SF-36 Survey (QOL); self-report (other outcomes)
Carfi ³	Symptom onset	Mean: 60.3 days after symptom onset (SD: 13.6) or 36.1 days after hospital discharge Minimum: not stated	Varied for each participant; based on date of examination	EQ-VAS (QOL); patient reporting survey (other outcomes)
Carvalho- Schneider ⁴	Symptom onset	Mean: 59.7 days after symptom onset (range: 57 to 67 days) Minimum: 57 days	Varied for each participant, although intended to be at day 60; based on date of examination	mMRC Dyspnea Scale (dyspnea); 10-point analog scale (chest pain, anosmia, ageusia)
Chen⁵	Hospital discharge	1 month after hospital discharge	All participants followed for same amount of time	SF-36 (QOL)
Chiesa- Estomba ⁶	Recovery	Mean 47 (IQR:30,71) after first consultation Minimum of 30 days after negative test	Varied for each participant; based on date of examination	sQOD-NS (olfactory function)
Chopra ⁷	Hospital discharge	Sixty days after hospital discharge	All participants followed for same amount of time	Self-report
D'Cruz ⁸	Hospital discharge	61 days after hospital discharge (IQR: 51- 67)	Varied for each participant; based on date of examination	mMRC Dyspnea Scale (dyspnea); PHQ-9 (depression); TSQ (trauma); GAD-7 (anxiety); 6-CIT (cognitive impairment); CT scans (organ functioning); 4MGS (gait speed); 1-minute Sit to Stand Test (mobility)
Daher ⁹	Hospital discharge	6 weeks after discharge	All participants followed for same amount of time	PHQ-9 (depression); GAD-7 (anxiety); EQ-5D-5L (QOL); 6 minute walk test (mobility); blood sample (lab assessments); electrocardiography, CT scans (organ functioning)
de Graaf ¹⁰	Hospital discharge	6 weeks after hospital discharge	All participants followed for the same amount of time	CT scans (organ functioning); pulmonary function tests; GAD-7 (anxiety); PHQ-9 (depression); PCL_5 (PTSD); CFQ-25 (cognitive impairment); IQ- CODE-N (cognitive impairment among elderly patients); NYHA (dyspnea)
Garrigues ¹¹	Hospital admission	Mean: 110.9 days after admission Minimum: 100 days	Varied for each participant; based on date of examination	mMRC Dyspnea Scale (dyspnea); EQ-5D-5L (QOL); self-report (other outcomes)
Gherlone ¹²	Hospital discharge	Median: 104 days after hospital discharge (IQR: 95 to 132)	Varied for each participant; based on date of examination	Extraoral and intraoral physical examination (facial abnormalities)
Gonzalez ¹³	Hospital discharge	3 months after hospital discharge	All participants followed for the same amount of time	SF-12 (QOL); HADS (depression); CT scan (organ functioning); mMRC Dyspnea scale (dyspnea); pulmonary function test
Halpin ¹⁴	Hospital discharge	Mean: 48 days(Range: 29,71) (SD: 17 days)	Varied for each participant; based on date of examination	EQ-5D-5L (QOL); Telephone screening tool (all other symptoms)
Huang ¹⁵	Symptom onset	Median: 186 days after symptom onset (IQR: 175 to 199)	Varied for each participant; based on date of examination	mMRC (dyspnea); EQ-5D-5L (QOL, anxiety, depression); EQ- VAS (QOL); blood sample (lab assessments); CT scans (organ function); 6-min walk test (mobility)
Jacobs ¹⁶	Hospital discharge	Mean: 35 day after	Varied for each participant;	PROMIS (all outcomes)

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		discharge (range: 30 to 40 days)	based on date of examination	
Lechien ¹⁷	Diagnosis	Two months	All participants followed for the same amount of time	SNOT-22 (sinonasal outcomes); sQOD-ns (olfactory function); NHANES (olfactory and gustatory function); 16-item Sniffiin-Sticks identification test (psychosocial olfactory evaluation)
Lerum ¹⁸	Hospital admission	Median: 83 days after hospital admission (IQR: 73 to 90 days)	Varied for each participant; based on date of examination	mMRC Dyspnea Scale (dyspnea); EQ-5D-5L (QOL); chest CT scan (organ function);
Liang ¹⁹	Hospital discharge	3 months	All participants followed for same amount of time	Spirometry (pulmonary function); CT scans (organ functioning); blood sample (lab assessments)
Lu ²⁰	Diagnosis	Three months after diagnosis	All participants followed for same amount of time	MRI scan (cerebral activity); self- report (other outcomes)
Mandal ²¹	Hospital discharge	Median: 54 days after discharge ((IQR 47– 59))	Varied for each participant; based on date of examination	X-ray (chest abnormalities); blood sample (lab assessments); PHQ-2 (depression); self-report (other outcomes)
Mazza ²²	Hospital discharge	90.1 days after hospital discharge (SD: 13.4)	Varied for each participant; based on date of examination	IES-R (distress); PCL-5 (PTSD); ZSDS (depression); BDI-13 (depression); STAI-Y (anxiety); WHIIRS (insomnia); OCI (obsessive compulsive disorder); BACS (cognitive function); clinical charts (inflammatory markers)
Mendez ²³	Hospital discharge	2 months (±1 month) after hospital discharge	Varied for each participant; based on date of examination	SF-12 (QOL); SCIP (verbal memory); ANT (verbal fluency); WAIS-III (working memory); GAD-7 (anxiety); PHQ-2 (depression); DTS (PTSD)
Moreno-Perez ²⁴	Symptom onset	Median: 76 days (IQR: 72 to 83 days)	Varied for each participant; based on date of final examination	EQ-VAS (QOL); x-ray (chest abnormalities); blood sample (lab assessments); pulmonary function test
Munro ²⁵	Hospital discharge	8 weeks	All participants followed for the same amount of time	General questionnaire
Nguyen ²⁶	Symptom onset	Mean: 221.7 days after symptom onset (SD: 10.9, range: 201-234)	Varied for each participant, based on date of final examination	Self-report
Poncet- Megemont ²⁷	Recovery	Mean: 1 month after recovery (range: 30- 35 days) Minimum: 30 days	Varied for each participant; based on date of examination	Self-report
Puntmann ²⁸	Diagnosis	Median: 71 days after diagnosis (IQR: 64 to 92 days) Minimum: not stated	Varied for each participant; based on date of examination	MRI scan (cardiac activity); self- report (other outcomes)
Qu ²⁹	Hospital discharge	3 months after hospital discharge	All participants followed for the same amount of time	SF-36 (QOL); self-report (all other symptoms)
Raman ³⁰	Symptom onset	Median: 2 to 3 months after symptom onset (IQR: 2.05 to 2.53 months)	Varied for each participant; based on date of examination	MRI scan (organ activity); spirometry (lung functioning); 6 minute walk test (mobility); PHQ- 9 (depression); GAD-7 (anxiety); MoCA (cognitive functioning); mMRC Dyspnea Scale (dyspnea); FSS (fatigue); SF-36 (QOL)
Rosales- Castillo ³¹	Hospital discharge	50.8 days after hospital discharge (SD: 6.02)	Varied for each participant, based on date of final examination	Self-report
Shah ³²	Symptom onset	Mean: 11.7 weeks after symptom onset (range: 8 to 12 weeks)	Varied for each participant; based on date of examination	Detailed pulmonary function testing (pulmonary function); 6 minute walk test (mobility); CT scans (organ functioning)
Sonnwebber, Eur Resp J ³³	Diagnosis	Mean: 103 days after diagnosis (SD: 21)	Varied for each participant; based on date of examination	mMRC Dyspnea Scale (dyspnea); spirometry, blood plethysmography (pulmonary

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				function); chest CT scan (organ function); blood sample (lab assessments); trans-thoracic echocardiography (cardiac function)
Sonnweber, Resp Res ³⁴	Symptom onset	60 days (SD ± 12) after symptom onset	Varied for each participant, although intended to be at day 60; based on date of examination	6-minute walk test (mobility); CT scan (lung functioning); blood sample (lab assessments)
Sykes ³⁵	Hospital discharge	Median: 113 days post-discharge, range: 46-167 days	Varied for each participant; based on date of examination	X-ray (chest abnormalities; mMRC Dyspnea Scale (dyspnea); EQ-5D-5L (QOL); direct questioning (all other symptoms)
Taboada ³⁶	Hospital discharge	6 months after hospital discharge	All participants followed for the same amount of time	EQ-5D-5L (QOL); PCFS (functional status)
Tomasoni ³⁷	Recovery	Median: 46 days after recovery (IQR: 43 to 48 days) Minimum: 30 days	Varied for each participant; based on date of examination	HADS (anxiety and depression); MMSE (cognitive disorders)
Townsend ³⁸	Hospital discharge and diagnosis	Median: 72 days after hospital discharge for hospitalized patients or timepoint of 14 days following diagnosis for outpatients (IQR: 62 to 87 days) Minimum: 42 days	Varied for each participant; based on date of examination	CFQ-11 (fatigue)
Ugurlu ³⁹	Hospital admission	Three months	All participants followed for the same amount of time	BSIT (olfactory function)
Vaira ⁴⁰	Symptom onset	60 days after initial symptom onset	All participants followed for same amount of time	Self-administered olfactory and gustatory psychosocial tests (anosmia and ageusia/dysgeusia, outpatients); CCCRC Olfactory Test (anosmia and ageusia/dysgeusia, hospitalized patients)
van den Borst ⁴¹	Hospital discharge and symptom onset	Mean of 10 weeks (SD: 1.7 weeks) since hospital discharge (hospitalized patients - 78.2% of participants) Mean of 14.7 weeks (SD: 2.2 weeks) since symptom onset for referred patients (21.8% of participants)	Varied for each participant; based on date of examination	Resting pulse-oximetry, spirometry (pulmonary functioning); mMRC Dyspnea Scale (dyspnea); CT scan and x- ray (chest functioning); CFS (frailty); HADS (anxiety and depression); TICS, CFQ (cognitive functioning); PCL-5 and IES-R (PTSS); SF-36 (QOL); blood sample (lab assessments)
Weerahandi ⁴²	Hospital discharge	Median: 37 days (range: 30 to 43 days) after hospital discharge Minimum: 30 days	Varied for each participant; based on date of examination	PROMIS (all outcomes)
Wong ⁴³	Symptom onset	Median: 13 weeks after symptom onset (IQR: 11-14 weeks)	Varied for each participant; based on date of examination	EQ-5D-5L (QOL); Frailty Index, UCSD (frailty); SOB Questionnaire, UCSD (shortness of breath); Pittsburgh Sleep Quality Index (sleep quality); PHQ-9 (depression)
Xiong ⁴⁴	Hospital discharge	Median: 97 days after hospital discharge (range: 95 to 102 days)	Varied for each participant; based on date of examination	Self-report
Zhao ⁴⁵	Hospital discharge	Range: 64 to 93 days after hospital discharge Minimum: 64 days	Varied for each participant; based on date of examination	Medical records; CT scan (chest functioning); spirometry (pulmonary functioning); self- report (other outcomes)

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eTable 6. Reported Outcomes and Frequencies at Follow-Up

Study	A kt er 1	Ar n ol d ²	C ar fi ³	C ar va lh o- S c h n ei d er ₄	C h e n⁵	C hi es a- Es to m ba 6	C ho pr a ⁷	D' Cr uz 8	Dah er ⁹	de Gr aa f ¹⁰	G ar ri g u es 11	G h er lo n e ¹ 2	G o nz al ez ¹³	Hal pin¹ ₄	Hua ng¹⁵	J a c o b s ¹ 6	L ec hi e n ¹ ⁷	Lerum ¹⁸	Li a n g ¹ ⁹	L u ² 0	M a n d al 21	Maz za ²²	M e d ez ²³	M or e n o- P er ez 24	M u nr o ² 5	N g u ye f 6	P once t- M e g e m on ² 7	P unt m a n ² ⁸	Q u ² 9	R a m a ³ ₀	R o sa le s- C as til lo 31	Sh ah ³²	Sonnweber, RespRes	Sonnweber, EurRespJ ³	Syk es ³⁵	T b o a d a ³ 6	T o m as o ni ³⁷	T o w n se n d ³ 8	U gu la 3	V ra ⁴⁰	va n e n B or st 41	Weerahandi 42	W o n g ⁴ 3	Xi o n g ⁴	Z h a o⁴ ₅
Sample Size	73 4	11 0	14 3	13 0	36 1	75 1	48 8	11 9	33	81	12 0	12 2	62	100	1733	1 8 3	88	10 3	76	60	38 4	226	17 9	27 7	12 1	12 5	13 9	10 0	54 0	58	11 8	60	10 9	14 5	134	91	10 5	12 8	42	13 8	12 4	15 2	78	53 8	55
Interpretation: % of study s	sampl	e cor	ntinu	ing to	o exp	erier	nce o	utco	ome at	the e	end o	f the	indiv	vidual	or s	tudy	follov	w-up	perio	od un	less	indic	ated	othe	rwise																				
Symptom Persistence (≥ 1 symptom at follow-up)		74 %	87 %	66 %				89 %					73 %		76%						72 %			50 .9 %					57 .6 %		62 .5 %			41 %	86%	84 %	52 .4 %						76 %	49 .6 %	
Cardiac																	·	·																											
Atypical chest pain		12 .7 % †	21 .7 %	13 .1 %					18%	18 %	10 .8 %				2.9%													17 %							17.9 %	8. 8 %								12 .3 %	
Palpitations				10 .9 %						14 %					8.9%				62 %									20 %	20 .4 %																
Chest tightness																			62 %																										
Chest distress																																												14 .1 %	
Resting heart rate increase																																												11 .2 %	

Discontinuous flushing																				4. 8
																				%
Cardiac Magnetic Resonan	ce Im	agin	g			•							<u>.</u>		<u>.</u>					· ·
Myocardial inflammation (abnormal native T1 and T2)														60 %						
Myocardial LGE														32 %						
Non-ischemic LGE														20 %						
Ischemic LGE														12 %						
Pericardial LGE														22 %						
Pericardial effusion														20 %						
Neurological				<u> </u>	<u> </u>							<u> </u>				<u> </u>			<u>_</u>	
Memory loss/impairment	19 .2 %						34 .2 %			28 .3 %							37.3 %			
Immediate verbal memory impairment										10%	49 .2 %									
Delayed memory impairment											14 .6 %									
Semantic verbal fluency impairment										32%	43 %									
Working memory impairment										24%	7. 2 %									
Attention and information processing impairment										33%										
Psychomotor coordination impairment										57%										
Executive functions impairment										50%										

Concentration issues	25 .6 %							2	28 %		22%																	25.4 %								
Confusion													8. 7 4 %																							
Cognitive deficit						21 .6 %	18%	27 %																				9.7 %		17 .1 %			15 %			
Mood change/low mood															16 .7 %	5												39.6 %								
Anxiety						22 .1 %		5 %		22 .1 %							30%	29 .6 %						14 %				47.8 %		29 %			10 %		6. 5 %	
Depression						18 %		17 %		15 .2 %						14 .6 %	9%	26 .8 %						19 %						11 %			12 %		4. 3 %	
Anxiety or depression	21 .6 %										23%	21.1 %																							14 .3 %	
Panic attack	13 .4 %																																			
Fatigue/exhaustion/asthe nia		39 .1 % †	53 .1 %	40 %		68 %	45%	Ę	55 %	29 .5 %	64%	59.9 %	4 4. 8 %	59 %	9 26 .7 %	6 69 %		34 .8 %	4			36 %	29 .4 %	55 %	30 .5 %			39.6 %	37 .4 %	31 .4 %	52 .3 %		69 %		28 .3 %	16 .4 %
PTSD						24 .8 %					31%						13%	25 .1 %																		
Dysphoria																																			1. 7 %	
Respiratory								·	·	·					·	·			·	·	·	·		·			·		·					·		
Shortness of breath/dyspnea		39 .1 % †	43 .4 %	30 %			33%	62 4 % .	41 7 %		50%		3 1. 7 %		-	53 %		3. .4 %	4			36 %	26 .1 %		31 .4 %	20 %			57 .1 %	27 .6 %				74 .3 %	50 %	14 .6 %

mMRC ≥ 0										69.0 %		54 %																
mMRC ≥ 1****								46 .7 %		24.2 %		19 %																
mMRC ≥ 2**		7. 7 %					29 %												64 %									
mMRC unspecified					44 %																	36 %	6 9.7 %					
Cough	11 .8 % †			15 .4 %	44 2. 6 %	33%	16 .7 %	16 .4 %	16.2 %		2 5. 1 %		60 %	34 %	2 .3 %	1		11 .7 %		5 2	20 %	17 %	7 35.1 %	1 14 .4 %			23 %	7. 1 %
Rhinorrhea						12%																						
Hemoptysis						0%																						
Sore throat						9%				4%													12.7 %	7				3. 2 %
Post-activity polypnea																												21 .4 %
Nonmotor polypnea																												4. 7 %
Lung Imaging																												
Abnormal chest radiograph appearance														38 %														
Deteriorating chest radiograph appearance														9 %														
Consolidation	1 %																											
Reticulation	7 %																											
Atelectasis	5 %																											

-																	
Pleural effusion	1 %																
Neuromuscular																	
Myalgia	22 .7 % †			15%	2.3	3%		25 %			13 %		37 .4 %	7		4 5 9	k. 5 6
Arthralgia	4. 1 5 % % †	6 3 6											28 .6 %	3		7 6 9	7. 5 %
Myalgia or arthralgia									19 .6 %			5´%	1.5				
Impaired mobility	16 .9 %				37% 6.5	5%		6. 7 %					56 %	5			
Numbness in extremities								6. 7 %									
Dysbasia											.4. 1 %						
Tremor								1. 7 %									
Joint pain	27 .3 %				8.9	9% 1 5. 8 %					26 .3 %						
Muscle pain						2 1. 3 %											
Headache	1. 82 % †			15%	1.9	9% 1 2. 6 %		10 %		3. 6 %							18 .1 8 %
Chills																4 6 9	k. 3 16

Limb oedema																									2. 6 %
Dizziness								5.8%																	2. 6 %
Trigeminal neuralgia							2 %																		
Gastrointestinal					-		 ·		•	 		 		 			 , ,	,				 t	t		
Diarrhea		0. 91 % †	3. 85 %			9%			3. 8 %		26 %		10 .5 %		679	5. 7 %									
Diarrhea or vomiting								4.6%											9 %						
Abdominal or stomach pain		1. 82 % †				3%																			
Nausea		0 % †				6%																			
Appetite issues								8% 8%																	
Quality of Life										_															
Returned to work				60 %			67 .9 %	20%													69 %				10 0 %
Worsened self-care	9. 6 %							16%												13 %					
Resumed sports							71 .8 %																		
Physical health (composite)		40 .2 *†	55 .9 (7 .2 4) *					45 .9 (3 6. 1 to 54																43 .8 (9 .3)** *	

						.4)																
Mental health (composite)	44 .8 *†	48 .9 2 (1 0. 81) *				55 .8 (4 0. 6 to 58)														2 (()) *	47 3 9 3 ***	
Poor or fair physical health								2 7. 1 %			44 .1 %		1: .4 %	5								
Poor or fair mental health								1 6. 9 %			39 .1 %		3: .6 %	2								
Pain/discomfort				49 .6 %			19% 24.9 %									24 %	48 %	3				
Circulatory and Immune System	I																					
Iron deficiency																30 %						
Anemia																9. 2 %						
Hyperferritinemia																38 %						
Elevated IL6 (inflammation marker)					0%											12 %						
Elevated CRP (inflammation marker)	1. 8 %									9. 5 %						16 %						
Elevated d-dimer (blood clot marker)										30 .1 %												
Lymphopenia	1. 8 %									7. 9 %												

Other																														
Visual loss																5. 4 %														
Eye irritation											8. 2 0 %																			
Phlegm											1 4. 8 %																			
Pharyngalgia					0	%																								
Ageusia/Dysgeusia					3 %	3.3 6	1 .: 9	0 1 [*] 3 %	7 0 5 %	6.9%	2 2. 8 %			25 %			1 - c	11 ´ 5 . % 9	19 .8 %		2 %	9. %	0		9. 5 %		10 .7 *			
Anosmia	11 .8 % †			37 %	5	0%	1 .: 9	3 14 3 %	4 1. 5 6 %	10.2 %	2 6. 2 %	20 .5 %		10 0 %				23 2 3 . % 9	23 .8 %		4 %	44 9. .4 % %	7	11 .0 %	10 .5 %	14 .3 %	9. 5 %			
Hair loss 9. 7 %							2	0		20.7 %																			28 .6 %	
Sleep disorders/insomnia	23 .6 % †	;			56 .5 %		3	0 3 6		25.2 %					3.1 %							28 3: % %	5.1	30 .8 %						
Night sweats																						24 %								
Somnipathy																													17 .7 %	
Fever	0. 91 % †		0 %		3	%			1. 6 %		1. 0 9 %		20 %			0 %						0 10 % %).4		0 %					
Ulcer											1. 0 9 %																			

Salivary gland ectasia						38 %											
Dry mouth						30 %											
TMJ abnormalities						9 %											
Masticatory muscle weakness						19 %											
Facial tingling						3 %											
Facial asymmetry						0. 8 %											
Burning Pain																	
Vision change										1. 7 %							
Hearing loss										1. 67 %			6. 6 %				
Tinnitus													6. 6 %				
Difficulty swallowing							8%										
Laryngeal sensitivity							17%										
Voice change							20%										
Difficulty communicating							6%										
Bowel control issues							3%										
Bladder control issues							10%							İ			

* 36 item short-fom survey (SF-36): lower scores indicate higher disability, score of 100 represents the best possible health status. Values presented represent mean (SD). ** mMRC ≥ 2: Walks slower than people of the same age because of dyspnoea or has to stop for breath when walking at own pace *** Patient-Reported Outcomes Measurement Information System 10-Question Short-Form (PROMIS-10 Global Health): Converted normed t-scores, where mean score of 50 and SD of 10 represents the mean for the general US population

[†]weighted average of mild, moderate, and severe baseline COVID-19 patient groups





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