

Table S1. Summary of results for essential and toxic elements: COVID-19 patients

Element	Patients (disease status)	Region	Number Sex	Age (mean/median/range)	MeCon (mean/median)*	Samples	Normal values**	Unit	Ref.
Fe									
	Control	Moscow, Russia	n = 43, 27 (♂), 16 (♀)	55.67	1.87	Serum	n/a	µg/mL	[171]
	Mild		n = 50, 25 (♂), 25 (♀)	50.47	1.32	-/-		-/-	
	Moderate		n = 50, 31 (♂), 19 (♀)	54.22	1.34	-/-		-/-	
	Severe		n = 50, 25 (♂), 25 (♀)	64.5	1.33	-/-		-/-	
	Mild	n/a	n = 22	58	18	Serum	n/a	µmol/L	[35]
	Moderate		n = 34		16	-/-		-/-	
	Severe		n = 53		15	-/-		-/-	
	Non-severe	Wuhan, Hubei	n = 202, 88 (♂), 114 (♀)	69	436.8	WB	406.5- 576.7	mg/L	[163]
	Severe	Province, China	n = 104, 60 (♂), 44 (♀)	58	370.24	-/-		-/-	
	Mild		n = 19	49 (36-65)	6.6	Serum	7.8-32.3	µmol/L	[159]
	Severe		n = 18	55 (46-63)	4.9	-/-		-/-	
	Critical		n = 13	66 (56-74)	5.2	-/-		-/-	
Fe as FER									
	Mild	n/a	n = 22	58	139	Serum	n/a	µg/L	[35]
	Moderate		n = 34		260	-/-		-/-	
	Severe		n = 53		317	-/-		-/-	
	Non-severe	Riyadh, Saudi Arabia	n = 45	47.3	331.24	Serum	12-300	ng/mL	[157]
	Severe		n = 35	58.2	1174.95	-/-		-/-	
	Mild / Moderate	Sakai, Osaka, Japan	n = 22, 12 (♂), 10 (♀)	≥ 65	458	Serum	n/a	ng/mL	[162]
	Severe		n = 7, 4 (♂), 3 (♀)	≥ 65	956	-/-		-/-	
Zn									
	Control	Jigawa, Northwestern	n = 21	35.8	64.9	Plasma	n/a	µg/dL	[172]
	Positive	Nigeria	n = 50	43.8	58.1	-/-		-/-	
	Control	Chennai, India	n = 45	32	105.8	Serum	80-120	µg/dL	[38]
	Positive		n = 47	34	74.5	-/-		-/-	
	Control	Shoushtar city, Iran	n = 186	51	86.66	Serum	70-127	µg/dL	[173]
	Positive		n = 93		67.61	-/-		-/-	
	Control	Chennai, India	n = 45	34 (18-77)	105.8	Serum	n/a	µg/dL	[38]
	Positive		n = 47	32 (18-60)	74.5	-/-		-/-	
	Control	Moscow, Russia	n = 43, 27 (♂), 16 (♀)	55.67	0.96	Serum	n/a	µg/mL	[171]
	Mild		n = 50, 25 (♂), 25 (♀)	50.47	0.92	-/-		-/-	
	Moderate		n = 50, 31 (♂), 19 (♀)	54.22	0.90	-/-		-/-	
	Severe		n = 50, 25 (♂), 25 (♀)	64.5	0.87	-/-		-/-	
	Control	Germany	n = 35	70	975.7	Serum	n/a	µg/L	[80]
	Positive			70	717.4	-/-		-/-	
	Non-survivors			89	< 638.7	-/-		-/-	
	Non-severe	Riyadh, Saudi Arabia	n = 45	47.3	124.57	Serum	66-110	µg/dL	[157]

Severe		n = 35	58.2	116.37	-/-	-/-	
Non-severe	Wuhan, Hubei	n = 202, 88 (♂), 114 (♀)	69	6.61	WB	4.3-7.8	mg/L [163]
Severe	Province, China	n = 104, 60 (♂), 44 (♀)	58	6.18	-/-	-/-	
Mild / Moderate	Sakai, Osaka, Japan	n = 22, 12 (♂)	≥ 65	87.7	Serum	n/a	μg/dL [162]
Severe		n = 7, 4 (♂)	≥ 65	62.4	-/-	-/-	
Mild	Qena, Upper Egypt	n = 45, 24 (♂), 21 (♀)	31.8	0.67	Serum	n/a	μg/mL [167]
Common		n = 57, 33 (♂), 24 (♀)	47.8	0.62	-/-	-/-	
Severe		n = 21, 15 (♂), 6 (♀)	59.1	0.73	-/-	-/-	
Extremely severe		n = 11, 6 (♂), 5 (♀)	69.5	0.72	-/-	-/-	
GCO group	n/a	n = 200	49.4	970	Plasma	n/a	μg/L [158]
PCO group		n = 75	64.1	840	-/-	-/-	
Non-severe	Tabriz, Iran	n = 114, 58 (♂), 56 (♀)	56.72	68.42	Serum	72.6-127 (♂)	μg/dL [174]
Severe		n = 112, 56 (♂), 56 (♀)	56	67.30	-/-	77.0-114 (♀)	-/-
Recovered				69.66	-/-	-/-	-/-
Deceased				62.43	-/-	-/-	-/-
Non-severe	Wuhan, Hubei	n = 202, 88 (♂), 114 (♀)	69	6.61	WB	4.3-7.8	mg/L [163]
Severe	Province, China	n = 104, 60 (♂), 44 (♀)	58	6.18	-/-	-/-	
Cu							
Control	Jigawa, Northwestern	n = 21	35.8	136.5	Plasma	n/a	μg/dL [172]
Positive	Nigeria	n = 50	43.8	128.3	-/-	-/-	
Control	Moscow, Russia	n = 43, 27 (♂), 16 (♀)	55.67	1.13	Serum	n/a	μg/mL [171]
Mild		n = 50, 25 (♂), 25 (♀)	50.47	1.27	-/-	-/-	
Moderate		n = 50, 31 (♂), 19 (♀)	54.22	1.46	-/-	-/-	
Severe		n = 50, 25 (♂), 25 (♀)	64.5	1.33	-/-	-/-	
Non-severe	Riyadh, Saudi Arabia	n = 45	47.3	18.35	Serum	10–22	μmol/L [157]
Severe		n = 35	58.2	18.2	-/-	-/-	
Non-severe	Wuhan, Hubei	n = 202, 88 (♂), 114 (♀)	69	838.55	WB	634.1-999.4	μg /L [163]
Severe	Province, China	n = 104, 60 (♂), 44 (♀)	58	929.73	-/-	-/-	
Survivors	Aschaffenburg-	n = 28, 13 (♂), 15 (♀)	69	1475.9	Serum	897.8-1906.0	μg /L [175]
Non-survivors	Alzenau, Germany	n = 7, 2 (♂), 5 (♀)	89	1317.9	-/-	-/-	
Intermediate care group	n/a	n = 35	65	~ 140 #	Serum	n/a	μg/dL [168]
ICU group				< 100	-/-	-/-	
Non-severe	Wuhan, Hubei	n = 70, 38 (♂), 32 (♀)	60	15.84	Urine	4-21.42	μg/L [164]
	Province, China			15.55 ^κ	-/-	4.39-13.37 ^κ	μg/g
Severe		n = 68, 41 (♂), 27 (♀)	65	32.14	-/-	-/-	μg/L
				77.71 ^κ	-/-	-/-	μg/g
Se							
Control	India	n = 30, 14 (♂), 16 (♀)	33.5 (37.5-43)	79.1	Serum	70-150	ng/mL [176]
Positive		n = 30, 24 (♂), 6 (♀)	40.5 (26-37)	69.3	-/-	-/-	
Control	Jigawa, Northwestern	n = 21	35.8	29.1	Plasma	n/a	ng/dL [172]
Positive	Nigeria	n = 50	43.8	25.3	-/-	-/-	
Control	Moscow, Russia	n = 43, 27 (♂), 16 (♀)	55.67	0.102	Serum	n/a	μg/mL [171]
Mild		n = 50, 25 (♂), 25 (♀)	50.47	0.093	-/-	-/-	
Moderate		n = 50, 31 (♂), 19 (♀)	54.22	0.090	-/-	-/-	

Se-P	Severe	Riyadh, Saudi Arabia	n = 50, 25 (♂), 25 (♀)	64.5	0.087	-/-	-/-	
	Non-severe		n = 45	47.3	134	Serum	70-150	µg/L [157]
	Severe		n = 35	58.2	162	-/-	-/-	
	Non-severe	Wuhan, Hubei Province, China	n = 70, 38 (♂), 32 (♀)	60	25.55 27.65 ^{&}	Urine -/-	10.46-82.71 15.86-38.13 ^{&}	µg/L µg/g [164]
	Severe		n = 68, 41 (♂), 27 (♀)	65	20.27 45.63 ^{&}	-/- -/-	µg/L µg/g	
	Mild		Tehran, Iran	n = 23 (♂), 15 (♀)	51	47.07	Serum	70-150
	Moderate	n = 12 (♂), 15 (♀)		59	47.36	-/-	-/-	
	Severe	n = 12 (♂), 7 (♀)		81	29.86	-/-	-/-	
	Survivors	Alzenau, Germany	n = 27, 12 (♂), 15 (♀)	69	53.3	Serum	45.7–131.6	µg/L [51]
	Non-survivors		n = 6, 2 (♂), 4 (♀)	89	40.8	-/-	-/-	
Survivors	Alzenau, Germany	n = 27, 12 (♂), 15 (♀)	69	3.3	Serum	2.56–6.63	mg/L [51]	
Non-survivors		n = 6, 2 (♂), 4 (♀)	89	2.1	-/-	-/-		
Mn								
	Control, Positive	Jigawa, Northwestern Nigeria	n = 21, n = 50	35.8, 43.8	2.10, 1.64	Plasma	n/a	mg/dL [172]
	Non-severe	Wuhan, Hubei Province, China	n = 202, 88 (♂), 114 (♀)	69	12.96	WB	8.1-18.5	µg /L [163]
	Severe		n = 104, 60 (♂), 44 (♀)	58	10.56	-/-	-/-	
	Non-severe	Wuhan, Hubei Province, China	n = 70, 38 (♂), 32 (♀)	60	<0.53 0.32 ^{&}	Urine -/-	<0.53-1.92 0.09-1.99 ^{&}	µg/L µg/g [164]
	Severe		n = 68, 41(♂), 27 (♀)	65	0.83 1.82 ^{&}	-/- -/-	µg/L µg/g	
Cr								
	Non-severe	Wuhan, Hubei Province, China	n = 202, 88 (♂), 114 (♀)	69	0.81	WB	<0.5-0.76	µg /L [163]
	Severe		n = 104, 60 (♂), 44 (♀)	58	1.03	-/-	-/-	
	Non-severe	Wuhan, Hubei Province, China	n = 70, 38 (♂), 32 (♀)	60	0.88 0.87 ^{&}	Urine -/-	<0.24-0.5 0.05-0.43 ^{&}	µg/L µg/g [164]
	Severe		n = 68, 41(♂), 27 (♀)	65	1.96 4.01 ^{&}	-/- -/-	µg/L µg/g	
Cd								
	Non-severe	Wuhan, Hubei Province, China	n = 202, 88 (♂), 114 (♀)	69	0.71	WB	0.22 -6.44	µg /L [163]
	Severe		n = 104, 60 (♂), 44 (♀)	58	0.7	-/-	-/-	
	Non-severe	Wuhan, Hubei Province, China	n = 70, 38 (♂), 32 (♀)	60	0.95 1.01 ^{&}	Urine -/-	0.34-3.39 0.27-2.23 ^{&}	µg/L µg/g [164]
	Severe		n = 68, 41 (♂), 27 (♀)	65	1.49 2.96 ^{&}	-/- -/-	µg/L µg/g	
Hg								
	Non-severe	Wuhan, Hubei Province, China	n = 202, 88 (♂), 114 (♀)	69	1.68	WB	<1.15-5.97	µg /L [163]
	Severe		n = 104, 60 (♂), 44 (♀)	58	1.45	-/-	-/-	
	Non-severe	Wuhan, Hubei Province, China	n = 70, 38 (♂), 32 (♀)	60	0.75 0.81 ^{&}	Urine -/-	<0.72-2.16 0.15-1.62 ^{&}	µg/L µg/g [164]
	Severe		n = 68, 41 (♂), 27 (♀)	65	<0.72 1.54 ^{&}	-/- -/-	µg/L µg/g	
Pb								

Non-severe	Wuhan, Hubei	n = 202, 88 (♂), 114 (♀)	69	12.37	WB	8.7-48.1	µg /L	[163]
Severe	Province, China	n = 104, 60 (♂), 44 (♀)	58	10.57	-/-		-/-	
Non-severe	Wuhan, Hubei	n = 70, 38 (♂), 32 (♀)	60	0.82	Urine	0.24-2.29	µg/L	[164]
	Province, China			0.86 ^κ	-/-	0.26-1.91 ^κ	µg/g	
Severe		n = 68, 41 (♂), 27 (♀)	65	0.83	-/-		µg/L	
				1.85 ^κ	-/-		µg/g	

*, ** All values are given as reported in the cited publications; [#] approximate value. n: number of patients; n/a: not available; Fe: iron; Zn: zinc; Cu: copper; Se: selenium; Mn: manganese; Cr: chromium; Cd: cadmium; Hg: mercury; Pb: lead; FER: ferritin; MeCon: metal concentration; WB: whole blood; Se-P: selenoprotein P; ICU: Intensive Care Unit; GCO group: good clinical outcome group; PCO group: poor clinical outcome group; ^κ creatinine adjusted analyte level.

Table S2. Summary of results for macroelements: COVID-19 patients

Element	Patients (disease status)	Region	Number Sex	Age (mean/median/range)	MeCon (mean/median)*	Samples	Normal values**	Unit	Ref.
Mg									
	Non-severe	Wuhan, Hubei	n = 202, 88 (♂), 114 (♀)	69	39.46	WB	31.9- 48.0	mg/L	[163]
	Severe	province, China	n = 104, 60 (♂), 44 (♀)	58	38.33	-//-	-//-	-//-	
	Discharge	Tehran, Iran	n = 396, 271 (♂), 125 (♀)	60.4	1.83	Blood	1.5- 2.5	mmol/L	[187]
	Expire		n = 63, 49 (♂), 14 (♀)	70.6	1.61	-//-	-//-	-//-	
	Moderate	Nancy, France	n = 43, 16 (♂), 27 (♀)	67.4	0.73	Serum	0.85- 0.95	mmol/L	[186]
	Severe		n = 108, 63 (♂), 45 (♀)	71.6	0.77	-//-	-//-	-//-	
	Critical		n = 149, 104 (♂), 45 (♀)	63	0.79	-//-	-//-	-//-	
Na									
	Non-severe	Riyadh, Saudi	n = 45	47.3	138.0	Serum	n/a	mmol/L	[157]
	Severe	Arabia	n = 35	58.2	136.58	-//-	-//-	-//-	
	Mild	Zhejiang	n = 82	49	139.6	Serum	137- 147	mmol/L	[188]
	Severe	province, China	n = 9	66	137.85	-//-	-//-	-//-	
	Mild	Chongqing	n = 45	44	139	Plasma	n/a	mmol/L	[192]
	Severe	China	n = 40	56	136.5	-//-	-//-	-//-	
K									
	Control	Modena, Italy	n = 171	64.1	4.0	Serum	3.5-5.3	mmol/L	[198]
	Hypokalemic		n = 119	65.8	3.1	-//-	-//-	-//-	
	Non-severe	Riyadh, Saudi	n = 45	47.3	4.4	Serum	n/a	mmol/L	[157]
	Severe	Arabia	n = 35	58.2	4.2	-//-	-//-	-//-	
	Discharge	Tehran, Iran	n = 396, 271 (♂), 125 (♀)	60.4	3.95	Blood	3.6- 5	mmol/L	[187]
	Expire		n = 63, 49 (♂), 14 (♀)	70.6	3.65	-//-	-//-	-//-	
	Moderate / Mild	Wenzhou China	n = 135	46	3.5	Plasma	> 3.5	mmol/L	[155]
	Severe / Critical		n = 40		3.2	-//-	-//-	-//-	
	Mild	Chongqing	n = 45	44	4	Plasma	n/a	mmol/L	[192]
	Severe	China	n = 40	56	3.8	-//-	-//-	-//-	
Ca									
	Control	Hialeah, Florida	n = 72	36.9	< 10	Serum	8.8–10.2	mg/dL	[199]
	Non-severe		n = 72	37.5	< 9	-//-	-//-	-//-	
	Negative	Monza, Italy	n = 165	66 (18-101)	2.28	Serum	n/a	mmol/L	[154]
	Positive		n = 420		2.15	-//-	-//-	-//-	
	Negative	Shoushtar city,	n = 186	51	9.50	Serum	8.6- 10.3	mg/dL	[173]
	Positive	Iran	n = 93		9.14	-//-	-//-	-//-	

	Mild	Zhejiang	n = 82	49	2.17	Serum	2.11- 2.52	mmol/L	[188]
	Severe	province, China	n = 9	66	2.01	-/-		-/-	
	Non-severe	Wuhan, Hubei	n = 202, 88 (♂), 114 (♀)	69	63.55	WB	46.9- 66.8	mg/L	[163]
	Severe	province, China	n = 104, 60 (♂), 44 (♀)	58	68.20	-/-		-/-	
	Survivors	Blida, Algeria	n = 83	59.5	2.2	Serum	2.21-.2.48	mmol/L	[189]
	Non-survivors		n = 37	68.6	2.02	-/-		-/-	
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Ca²⁺									
	Negative	Monza, Italy	n = 165	66 (18-101)	1.17	WB	n/a	mmol/L	[154]
	Positive		n = 420		1.12	-/-		-/-	
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Cl									
	Non-severe	Riyadh, Saudi	n = 45	47.3	103.3	Serum	n/a	mmol/L	[157]
	Severe	Arabia	n = 35	58.2	97.7	-/-		-/-	

*, ** All values are given as reported in the cited publications. n: number of patients; n/a: not available; Mg: magnesium; Na: sodium; K: potassium; Ca: calcium; Cl: chloride; MeCon: metal concentration; WB: whole blood.

Table S3. Summary of clinical trials with zinc (Zn) for COVID-19

ClinTI	Official Title	Country	Agents	Zn compound	Zn dose	Frequency/Duration of Zn administration	Patient details
Participants with COVID-19 / at high risk of COVID-19							
NCT04370782*	A Randomized Study Evaluating the Safety and Efficacy of Hydroxychloroquine and Zinc in Combination With Either Azithromycin or Doxycycline for the Treatment of COVID-19 in the Outpatient Setting	USA	Zn/HCQ/AZM Zn/HCQ/DOXY	ZnS	220 mg	OD for 5 d	n = 750 ≥ 30 yr ♂/♀
NCT04342728*	Coronavirus Disease 2019- Using Ascorbic Acid and Zinc Supplementation (COVIDatoZ) Research Study A Randomized, Open Label Single Center Study	USA	Zn and/or AscA ST	ZnGLU	50 mg	OD at <i>h.s.</i> for 10 d	n = 520 ≥ 18 yr ♂/♀
NCT04621461*	A Randomized, Placebo-Controlled Study Evaluating the Efficacy of Zinc for the Treatment of COVID-19 in the Outpatient Setting	USA	Zn Placebo	ZnS	220 mg	OD for 5 d	n = 3 ≥ 30 yr ♂/♀
CTRI/2020/05/025215*	A prospective, single centre, randomized open labelled comparative clinical study to evaluate the effectiveness of Siddha medicine, Kabasura kudineer and vitamin c-zinc supplementation in the management of asymptomatic COVID 19 patients.	India	KSK VitC Zn	n/a	100 mg	OD for 14 d	n = 60 18-55 yr ♂/♀
IRCT20180425039414N2*	The effect of zinc on the treatment and clinical course of patients with SARS-CoV-2 (COVID-19)	Iran	Zn HCQ	n/a	220 mg	BID	n = 80 ≥ 18 yr ♂/♀
NCT04446104*	A Randomized Open-label Prophylaxis Trial Among Migrant Workers at High-risk of COVID-19 (DORM Trial)	Singapore	Zn/VitC HCQ IVM PI VitC	n/a	80 mg	OD for 42d	n = 5000 21-60 yr ♂
NCT04584567*	COVID-19 Infection Prophylaxis With Low Dose of Doxycycline and Zinc in Health Care Workers	Tunisia	Zn + DOXY DOXY (placebo) Placebo	n/a	15 mg	daily	n = 1100 18-65 yr ♂/♀
NCT04468139†	The Study of Quadruple Therapy Zinc, Quercetin, Bromelain and Vitamin C on the Clinical Outcomes of Patients Infected With COVID-19	Saudi Arabia	Zn QUE BM VitC	elemental	50 mg	daily	n = 60 ≥ 18 yr ♂/♀
NCT04447534†	Does Zinc Supplementation Enhance the Clinical Efficacy of Chloroquine / Hydroxychloroquine in Treatment of COVID-19?	Egypt	HCQ + Zn HCQ	n/a	n/a	n/a	n = 200 ≥ 18 yr ♂/♀

NCT04384458 [†]	Comparative Study of Hydroxychloroquine and Ivermectin in COVID-19 Prophylaxis	Brazil	HCQ+Zn IVM+Zn	active Zn active Zn	20 mg 20 mg	BID for 50 d BID for 45 d	n = 400 18-70 yr ♂/♀
NCT04472585 [†]	Sub-cutaneous Ivermectin in Combination With and Without Oral Zinc: a Placebo Randomized Control Trial on Mild to Moderate COVID-19 Patients	Pakistan	Zn/ST IVM/ST Zn + IVM/ST Placebo/ST	ZnS	20 mg	TID	n = 180 18-60 yr ♂/♀
NCT04395768 [†]	Therapies to Prevent Progression of COVID-19, Including Hydroxychloroquine, Azithromycin, Zinc, Vitamin D, Vitamin B12 With or Without Vitamin C, a Multi-centre, International, Randomized Trial: The International ALLIANCE Study	Australia	Zn HCQ AZM VitD3 VitB12 (+)(-) VitC	ZnCIT	30 mg	PO daily	n = 200 ≥ 18 yr ♂/♀
NCT04334512 [†]	A Randomized, Double-Blind, Placebo-Controlled Phase IIa Study of Quintuple Therapy to Treat COVID-19 Infection	USA	Zn HCQ AZM VitC VitD Zn, VitC, D (placebo)	n/a	n/a	10 d	n = 600 ≥ 18 yr ♂/♀
NCT04621149 [†]	A Phase 2 Screening Study of Candidate Non-prescription Treatments for COVID-19: A Patient-driven, Randomized, Factorial Study Evaluating Patient-reported Outcomes (PROFACT-01)	USA	Zn ClO ₂ FMT LF GTE Placebo	ZnACE	n/a	7 d	n = 120 20-70 yr ♂/♀
NCT04641195 [†]	A Randomized Trial to Determine the Effect of Vitamin D and Zinc Supplementation for Improving Treatment Outcomes Among COVID-19 Patients in India	India	Zn + VitD3 Zn (placebo) VitD3 (placebo) Placebo	ZnGLU	40 mg	OD up to 8 wk	n = 700 ≥ 18 yr ♂/♀
NCT04323228 [†]	Anti-inflammatory/Antioxidant Oral Nutrition Supplementation on the Cytokine Storm and Progression of COVID-19: A Randomized Controlled Trial	Saudi Arabia	ONS with Zn Placebo	n/a	5.7 mg	14 d	n = 40 18-65yr ♂/♀
NCT04335084 [†]	A Randomized, Double-Blind, Placebo-Controlled Phase IIa Study of Hydroxychloroquine, Vitamin C, Vitamin D, and Zinc for the Prevention of COVID-19 Infection	USA	Zn, HCQ, VitC, VitD Zn, VitC, D (placebo)	n/a	n/a	12 wk	n = 600 ≥ 18 yr ♂/♀
ACTRN126200000454976 [†]	High-dose intravenous zinc (HDIVZn) as adjunctive therapy in COVID-19 positive critically ill patients: A pilot randomized controlled trial	Australia	Zn Placebo	ZnC	0.5 mg/kg	daily for 7 d	n = 160 ≥ 18 yr ♂/♀
NCT04558424 [#]	Randomized, Double -Blind, Placebo Controlled, Trial to Evaluate the Effect of	Bangladesh	Zn + AscA/ST Placebo/ST	ZnGLU	220 mg	OD for 10 d	n = 50 18-70 yr

	Zinc and Ascorbic Acid Supplementation in COVID-19 Positive Hospitalized Patients in BSMMU						♂/♀
PACTR202005622389003 [#]	Efficacy and safety of Hydroxychloroquine, Azithromycin and Zinc for the treatment of patients with SARS-CoV2 infection in Senegal: a dose ranging randomised trial. COVID-19	Senegal	Zn/ST HCQ+AZM	n/a	20 mg	OD for 6 d	n = 128 18-65 yr ♂/♀
CTRI/2020/07/026340 [#]	Prospective study to assess therapeutic role of Zinc in COVID-19 patients	India	Zn/ST	ZnS	100 mg	OD	n = 100 18-80 yr ♂/♀
NCT04898023 [#]	Evaluation of Combination Zinc and Green Tea Extract Supplementation on Reduction in Symptom Duration and Severity Associated With Community Respiratory Viral Infections: a Randomized Control Trial (ZiPhenol Study)	USA	Zn+GTE+AscA Placebo	ZnCIT	50 mg	2 cap BID for 5 d	n = 100 ≥ 18 yr ♂/♀
NCT04377646 [#]	A Study of Hydroxychloroquine and Zinc in the Prevention of COVID-19 Infection in Military Healthcare Workers (COVID-Milit)	Tunisia	Zn Zn (placebo) HCQ HCQ (placebo)	elemental	15 mg	OD up to 2 mo	n = 660 18-65 yr ♂/♀
NCT04542993 ^{††}	Can SARS-CoV-2 Viral Shedding in COVID-19 Disease be Reduced by Resveratrol-assisted Zinc Ingestion, a Direct Inhibitor of SARS-CoV-2-RNA Polymerase? A Single Blinded Phase II Protocol (Reszinate Trial)	USA	Zn + RSV Zn + RSV (placebo)	ZnPIC	50 mg	TID for 5 d	n = 60 18-75 yr ♂/♀
NCT04482686 ^{††}	A Phase I Double-Blind Randomized Placebo-Controlled Trial of Combination Therapy to Treat COVID-19 Infection	USA	Zn IVM DOXY-Hcl VitC VitD3 Zn, VitC, D3 (placebo)	ZnS	n/a	10 d	n = 31 ^{&} ≥ 18 yr ♂/♀

ClinTI: Clinical Trial Identifier; n: number of patients; yr: year; mo: month; d: day; cap: capsules; *h.s.*: *hora somni*; OD: once daily; BID: twice a day; TID: three times a day; PO: *per os*; BSMMU: Bangabandhu Sheikh Mujib Medical University; QUE: quercetin; BM: bromelain; HCQ: hydroxychloroquine; IVM: ivermectin; DOXY: doxycycline; DOXY-Hcl: doxycycline hyclate; FMT: famotidine; LF: lactoferrin; RSV: resveratrol; AZM: azithromycin; AscA: ascorbic acid; ClO₂: chlorine dioxide; PI: povidone iodine; KSK: Kabasura Kudineer; VitB12: vitamin B12; VitC: vitamin C; VitD: vitamin D; VitD3: vitamin D3; Zn: zinc; GTE: green tea extract; ZnGLU: zinc gluconate; ZnCIT: zinc citrate; ZnPIC: zinc picolinate; ZnS: zinc sulfate; ZnCl: zinc chloride; ZnACE: zinc acetate; ONS: oral nutrition supplement; (+): with, (-): without ST: standard treatment; n/a: not available. Recruitment status: * complete, † recruiting, †† active, not recruiting. & Actual enrollment.

Table S4. Summary of clinical trials with copper (Cu) for COVID-19

ClinTI	Official Title	Country	Agents	Cu dose	Frequency/Duration of Cu administration	Patient details
Participants with COVID-19						
CTRI/2020/07/026514 [#]	A Phase-II, Open Label, Randomized Controlled Trial of Resveretrol-Copper Plus Standard Treatment Or Sodium-Copper-Chlorophyllin Plus Standard Treatment Versus Standard Treatment in Cancer patients with SARS-CoV-2 Infection who are Symptomatic Or have Pneumonia for COVID 19	India	RSV-Cu/ST CHLN/ST ST without RSV-Cu	560 ng	1 tablet once every 6 hr from the date of randomization	n = 200 18-99 yr ♂/♀
CTRI/2020/05/025337 [#]	A Phase-II, Open Label, Randomized Controlled Trial Of Resveretrol-Copper Plus Standard Treatment Or Sodium-Copper-Chlorophyllin Plus Standard Treatment Versus Standard Treatment In Hospitalized Patients With Pneumonia Due To SARS-CoV-2 (COVID-19)	India	RSV-Cu/ST CHLN/ST ST without RSV-Cu	560 ng	1 tablet QID from the date of randomization	n = 200 18-99 yr ♂/♀
CTRI/2020/05/025336 [#]	A Phase-III, Open Label, Randomized Controlled Trial Of Resveretrol-Copper Plus Standard Treatment Or Sodium-Copper-Chlorophyllin Plus Standard Treatment Versus Standard Treatment In Asymptomatic Or Mildly Symptomatic Patients With SARS-CoV-2 Infection (COVID-19)	India	RSV-Cu/ST CHLN/ST ST without RSV-Cu	560 ng	1 tablet QID from the date of randomization	n = 300 18-99 yr ♂/♀

ClinTI: Clinical Trial Identifier; n: number of patients; hr: hours; yr: year; ST: standard treatment; RSV-Cu: resveratrol-copper tablets; CHLN: chlorophyllin; QID: four times a day.
 Recruitment status: [#] not yet recruiting.

Table S5. Summary of clinical trials with selenium (Se) for COVID-19

ClinTI	Official Title	Country	Agents	Se dose	Frequency/Duration of Se administration	Patient details
Participants with COVID-19						
IRCT20210427051100N1*	The effect of selenium supplementation on inflammatory markers and blood cells in patients with COVID-19 double-blind randomized clinical trial	Iran	Se/ST CG/Placebo/ST	200 µg	daily for 14 d	n = 40 20-60 yr ♂/♀
IRCT20190418043307N1*	Investigating the effectiveness of selenium on recovery of hospitalized patients with COVID-19	Iran	Se/ST CG/ST	200 µg	daily for 14 d	n = 100 18-65 yr ♂/♀
IRCT20160706028815N5*	The effect of addition of selenium to treatment regimen of hospitalized patients with COVID-19 on the outcome of disease compared with control group	Iran	Se/ST CG/ST	1000 µg / 500 µg	every 12 hr on the 1st d / every 12 d from the 2nd d	n = 40 ≥ 18 yr ♂/♀
IRCT20190312043030N2*	The effect of combination of selenium, vitamin C and methylprednisolone in acute respiratory distress syndrome mortality and morbidity from COVID-19	Iran	Se/VitC/MPS/ST CG/ST	1 mg	daily for 7 d	n = 40 18-90 yr ♂/♀
IRCT20160919029870N3*	Evaluation of the effectiveness of selenium added to intravenous nutrition therapy on mortality and duration of ICU hospitalization in patients with COVID-19 disease	Iran	Se CG/ST	1 mg / 500 µg	at the beginning / IVI daily during 15 min at 2 PM every d for 5 d	n = 80 no age limit ♂/♀
NCT04869579#	Selenium as a Potential Treatment for Moderately-ill, Severely-ill, and Critically-ill COVID-19 Patients	USA	SeA/ST Placebo/ST	2000 µg 1000 µg	on d 1 d 2-14	n = 100 ≥ 18 yr ♂/♀

ClinTI: Clinical Trial Identifier; n: number of patients; d: day; hr: hours; min: minutes; Se: selenium; ST: standard treatment; CG: control group; VitC: vitamin C; MPS: methylprednisolone; IVI: intravenous infusion. Recruitment status: * complete, # not yet recruiting.

Table S6. Summary of clinical trials with magnesium (Mg) for COVID-19

ClinTI	Official Title	Country	Agents	Mg compound	Mg dose	Frequency/Duration of Mg administration	Patient details
Participants with COVID-19							
IRCT20191211045691N1*	Evaluation of the efficacy and safety of inhaled magnesium sulfate in combination with standard treatment in COVID-19 patients: a clinical trial	Iran	Mg/ST CG/ST	MgS	5 cc of a 20% INJ-V or 2 cc of a 50% INJ-V	every 8 h for 5 d	n = 100 18-80 yr ♂/♀
IRCT20210702051763N1*	The effect of vitamin D and magnesium supplementation on clinical symptoms, inflammatory markers and oxidative stress in patients with COVID-19: double-blind randomized control clinical trial	Iran	Mg+VitD VitD+Mg (placebo) Mg+VitD (placebo) Placebo	n/a	300 mg	for 3 wk	n = 104 18-65 yr ♂/♀
NCT04941703†	Investigation of Choice Alteration of the Gut Metagenome on COVID-19 Severity	USA	Mg/PB Placebo	MgCIT	296 ml	1 bottle PO once within a 4-h period	n = 30 18-99 yr ♂/♀
CTRI/2020/06/026189#	Randomized, Double Blind, Parallel Group Study of Vitamin D3 & Magnesium in COVID-19 Infection	India	VitD3+Mg	MgGly	250 mg	BID for 14 d	n = 210 20-60 yr ♂/♀

ClinTI: Clinical Trial Identifier; n: number of patients; wk: weeks; d: day; h: hour; BID: twice a day; PO: *per os*; PB: probiotic; INJ-V: injectable vial; cc: cubic centimeter; MgS: magnesium sulfate; MgCIT: magnesium citrate; MgC: magnesium carbonate; MgGly: magnesium glycinate VitD: vitamin D; ST: standard treatment; CG: control group. Recruitment status: * complete, † recruiting, # not yet recruiting.