

Supplementary Materials

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Figure s1. PRISMA Flow Diagram

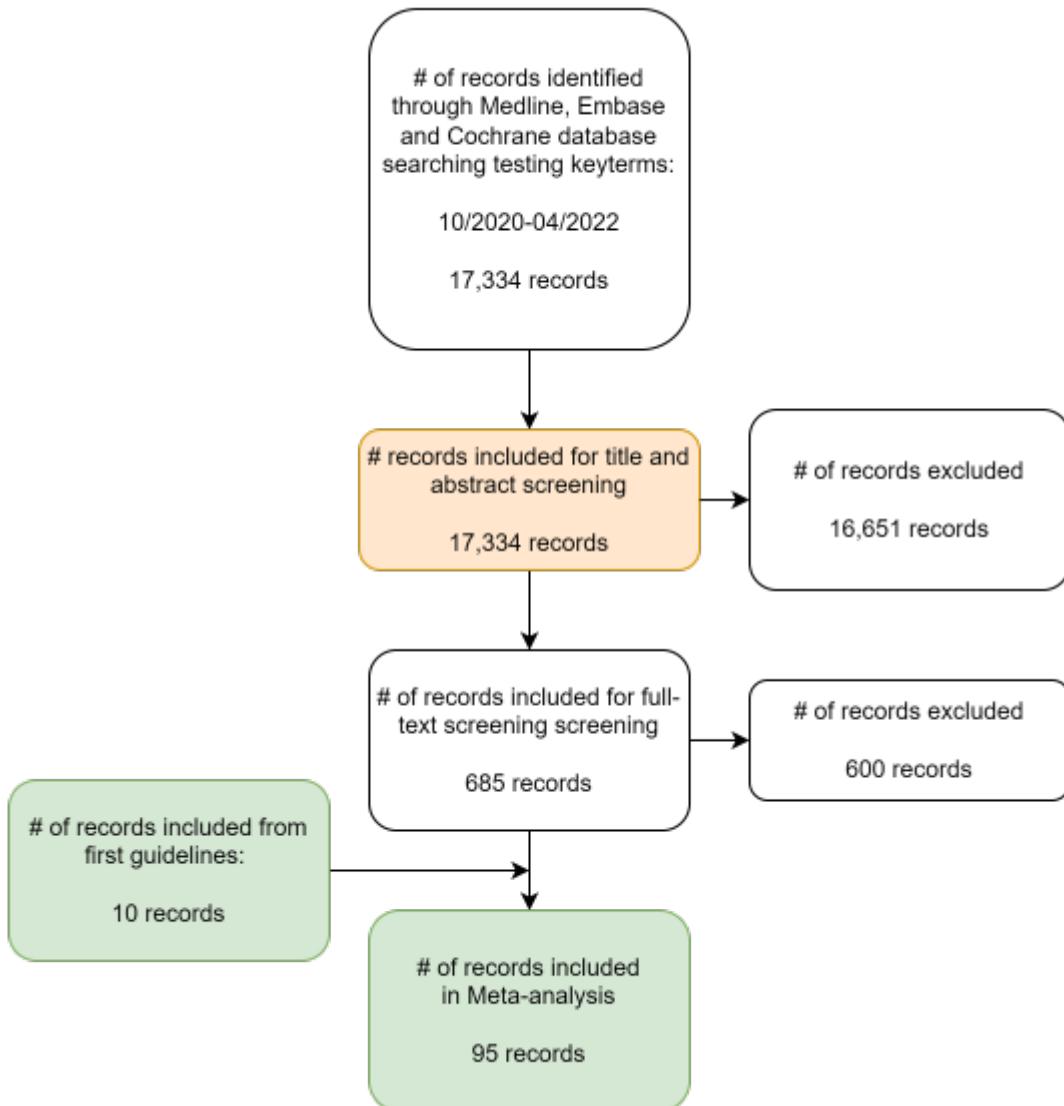


Table s1. PICO Questions Identified by the Panel

PICO	Population	Test/strategy 1	Test/strategy 2	Outcome	Recommendation #
1.	Symptomatic	Ag	No testing	Diagnostic test accuracy	Recommendation #1
2.	Symptomatic	Ag	NAAT	Diagnostic test accuracy	Recommendation #2
3.	Symptomatic	Ag strategy, repeat testing	NAAT	Diagnostic test accuracy	Recommendation #3
4.	Asymptomatic	Ag	No testing	Diagnostic test accuracy	Recommendation #4
5.	Asymptomatic	Ag	NAAT	Diagnostic test accuracy	Recommendation #5
6.	Asymptomatic	Ag strategy, repeat testing	NAAT	Diagnostic test accuracy	Recommendation #6
7.	Students in educational settings and employees in workplaces	Ag strategy, repeat testing	No testing	Patient important outcomes, transmission events	Recommendation #7 (Evidence gap)
8.	Asymptomatic individuals planning to attend large gatherings	Ag	No testing	Patient important outcomes, transmission events	Recommendation #8 (Evidence gap)
9.	Symptomatic	Point-of-care Ag	laboratory-based Ag	Patient important outcomes, transmission events	Both absorbed into Recommendation #9
10.	Asymptomatic	Point-of-care Ag	laboratory-based Ag	Patient important outcomes, transmission events	
11.	Symptomatic	Observed self-collection	Unobserved self-collection	Diagnostic test accuracy	Both absorbed into

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12.	Asymptomatic	Observed self-collection	Unobserved self-collection	Diagnostic test accuracy	Recommendation #10
13.	Symptomatic	Home testing #1	Home testing # 2	Diagnostic test accuracy	Deprioritized question

Ag: antigen [testing]; **NAAT:** nucleic acid amplification test

Table s2. Search Strategy

EMBASE	
Set #	Search Strategy
1	('coronavirus disease 2019'/exp OR 'Severe acute respiratory syndrome coronavirus 2'/exp)
2	((corona* OR corono*) AND (viral* OR viridae* OR virinae* OR virus*)):ti,ab
3	('2019 novel*' OR '2019-ncov' OR 2019ncov OR betacoronavirus* OR Coronaviridae* OR coronavirinae* OR coronavirus* OR coronovirus* OR corvid19 OR 'corvid-19' OR cov OR covid19 OR covid2019 OR 'covid 2019' OR 'covid-19' OR 'hcov-19' OR hcov19 OR ncorona* OR ncorono* OR 'n-cov' OR 'ncov-2019' OR ncov OR ncov2019 OR ncovchina* OR ncovchinese* OR ncovhubei* OR ncovor OR ncovwuhan* OR 'novel betacoronavirus' OR 'Novel Coronavirus' OR 'novel CoV' OR 'sarscov-19' OR 'sarscov19' OR 'sars-cov-19' OR sarscov19 OR 'sarscov2' OR 'sarscov-2' OR 'sars-cov2' OR 'sars-cov-2' OR 'wn-cov' OR wncov):ti,ab
4	((epidem* OR outbreak* OR pandem* OR wildlife*) NEAR/1 (china* OR chinese* OR huanan* OR pneumonia OR respirator*)):ti,ab
5	(('food market*' OR 'seafood market*') NEAR/10 (china* OR chinese* OR huanan* OR hubei* OR wuhan*)):ti,ab
6	OR/1-5
7	('virus antigen'/exp)
8	antigen:ti,ab
9	OR/7-8
10	('delayed diagnosis'/exp OR 'diagnosis'/de OR 'diagnostic procedure'/de OR 'differential diagnosis'/exp OR 'early diagnosis'/exp OR 'examination'/exp OR 'laboratory diagnosis'/exp OR 'physical examination'/exp OR 'qualitative diagnosis'/exp OR 'quantitative diagnosis'/exp OR 'screening'/exp OR 'symptom assessment'/exp OR 'virus diagnosis'/exp)
11	(case* OR 'case finding' OR casefinding OR detect* OR diagnos* OR screen* OR test*):ti,ab
12	OR/10-11
13	6 AND 9 AND 12
14	[english]/lim
15	13 AND 14
16	[animals]/lim NOT [humans]/lim
17	15 NOT 16

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	('abstract report'/exp OR 'animal experiment'/exp OR 'book'/exp OR 'case finding'/exp OR 'case report'/exp OR 'case study'/exp OR 'conference paper'/exp OR 'editorial'/exp OR 'feasibility study'/exp OR 'in vitro study'/exp 'letter'/exp OR 'meta analysis'/exp OR 'meta analysis topic'/exp OR 'meta analysis (topic)'/exp OR 'note'/exp OR 'practice guideline'/exp OR 'review'/exp OR 'systematic review'/exp OR 'systematic review topic'/exp OR 'systematic review (topic)'/exp OR 'veterinary clinical trial'/exp OR 'veterinary study'/exp OR [conference abstract]/lim OR [conference paper]/lim OR [conference review]/lim OR [editorial]/lim OR [letter]/lim OR [note]/lim OR [short survey]/lim OR meta*analys*:ti,ab OR (integrative NEAR/5 research NEAR/5 review*):ti,ab OR (methodologic* NEAR/5 overview*):ti,ab OR (methodologic* NEAR/5 review*):ti,ab OR (quantitativ* NEAR/5 overview*):ti,ab OR (quantitativ* NEAR/5 review*):ti,ab OR (quantitativ* NEAR/5 synthesis*):ti,ab OR (research NEAR/5 integration):ti,ab OR (systematic* NEAR/5 overview*):ti,ab OR (systematic* NEAR/5 review*):ti,ab)
18	17 NOT 18
20	[22-2-2021]/sd NOT [29-9-2021]/sd
21	19 AND 20

PUBMED	
Set #	Search Strategy
1	"COVID-19"[Mesh] OR "SARS-CoV-2"[Mesh] OR "Coronavirus Infections"[Mesh] OR "Betacoronavirus"[Mesh]
2	((corona*[tiab] OR corono*[tiab]) AND (viral*[tiab] OR viridae*[tiab] OR virinae*[tiab] OR virus*[tiab]))
3	"2019 novel*"[tiab] OR "2019-ncov"[tiab] OR 2019ncov[tiab] OR betacoronavirus*[tiab] OR Coronaviridae*[tiab] OR coronavirinae*[tiab] OR coronavirus*[tiab] OR coronavirus*[tiab] OR cov[tiab] OR covid19[tiab] OR covid2019[tiab] OR "covid 2019" [tiab] OR "covid-19"[tiab] OR "hcov-19"[tiab] OR hcov19[tiab] OR "n-cov"[tiab] OR "ncov-2019"[tiab] OR ncov[tiab] OR ncov2019[tiab] OR "novel betacoronavirus"[tiab] OR "Novel Coronavirus"[tiab] OR "novel CoV"[tiab] OR "sars-cov19"[tiab] OR "sars-cov-19"[tiab] OR sarscov19[tiab] OR "sarscov2"[tiab] OR "sarscov-2"[tiab] OR "sars-cov2"[tiab] OR "sars-cov-2"[tiab]
4	(epidem* OR outbreak* OR pandem* OR wildlife*) AND (china* OR chinese* OR huanan* OR pneumonia OR respirator*)
5	(("food market*"[tiab] OR "seafood market*"[tiab]) AND (china*[tiab] OR chinese*[tiab] OR huanan*[tiab] OR hubei*[tiab] OR wuhan*[tiab]))
6	OR/1-5
7	"Antigens, Viral"[Mesh]
8	antigen[tiab]
9	OR/7-8

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10	"COVID-19 Testing"[Mesh] OR "Delayed Diagnosis"[Mesh] OR "Diagnosis"[Mesh:NoExp] OR "Diagnosis, Differential"[Mesh] OR "Diagnostic Techniques and Procedures"[Mesh] OR "Early Diagnosis"[Mesh]
11	"diagnosis"[Subheading]
12	case*[tiab] OR "case finding"[tiab] OR casefinding[tiab] OR detect*[tiab] OR diagnos*[tiab] OR screen*[tiab] OR test*[tiab]
13	OR/10-12
14	6 AND 9 AND 13
15	English[Language]
16	14 AND 15
17	animals[Mesh] NOT humans[Mesh]
18	16 NOT 17
19	"2021/02/22"[PDAT] : "3000/12/31"[PDAT]
20	18 AND 19
21	("Academic Dissertation"[Publication Type] OR "address"[Publication Type] OR "Anecdotes"[Publication Type] OR "Animation"[Publication Type] OR "autobiography"[Publication Type] OR "bibliography"[Publication Type] OR "biography"[Publication Type] OR "Book Illustrations"[Publication Type] OR "Book Review"[Publication Type] OR "Bookplate"[Publication Type] OR "Cartoon"[Publication Type] OR "Case Reports"[Publication Type] OR "Catalog"[Publication Type] OR "Chart"[Publication Type] OR "Comment"[Publication Type] OR "congress"[Publication Type] OR "consensus development conference"[Publication Type] OR "consensus development conference, nih"[Publication Type] OR "dictionary"[Publication Type] OR "directory"[Publication Type] OR "editorial"[Publication Type] OR "Expression of Concern"[Publication Type] OR "Guideline"[Publication Type] OR "Handbook"[Publication Type] OR "interactive tutorial"[Publication Type] OR "interview"[Publication Type] OR "Juvenile Literature"[Publication Type] OR "lecture"[Publication Type] OR "legal case"[Publication Type] OR "legislation"[Publication Type] OR "letter"[Publication Type] OR "Meeting Abstract"[Publication Type] OR "Meta-Analysis"[Publication Type] OR "news"[Publication Type] OR "newspaper article"[Publication Type] OR "overall"[Publication Type] OR "patient education handout"[Publication Type] OR "periodical index"[Publication Type] OR "personal narrative"[Publication Type] OR "portrait"[Publication Type] OR "Review"[Publication Type] OR "Scientific Integrity Review"[Publication Type] OR "Systematic Review"[Publication Type] OR "Unpublished Work"[Publication Type] OR "hascommenton"[All Fields] OR "Cartoons as Topic"[Mesh] OR "Meta-Analysis as Topic"[Mesh] OR "Review Literature as Topic"[Mesh] OR "Systematic Reviews as Topic"[Mesh] OR "case report*"[tiab] OR "integrative research review*"[tiab] OR "integrative review*"[tiab] OR "literature review"[tiab] OR meta-analys*[tiab] OR "meta analys*"[tiab] OR metaanalys*[tiab] OR "narrative review"[tiab] OR "research integration"[tiab] OR "scoping review"[tiab] OR ((methodologic*[tiab] OR quantitative*[tiab] OR systematic*[tiab])) AND (overview*[tiab] OR review*[tiab] OR synthesis*[tiab])))

22 | 20 NOT 21**Cochrane**

Set #	Search Strategy
1	MeSH descriptor: [Betacoronavirus] explode all trees
2	MeSH descriptor: [Coronavirus Infections] explode all trees
3	((corona* OR corono*) NEXT (viral* OR viridae* OR virinae* OR virus*)):ti,ab
4	("2019 novel*" OR "2019-ncov" OR 2019ncov OR betacoronavirus* OR Coronaviridae* OR coronavirinae* OR coronavirus* OR coronavirus* OR corvid19 OR "corvid-19" OR cov OR covid19 OR covid2019 OR "covid 2019" OR "covid-19" OR "hcov-19" OR hcov19 OR ncorona* OR ncorono* OR "n-cov" OR "ncov-2019" OR ncov OR ncov2019 OR ncovchina* OR ncovchinese* OR ncovhubei* OR ncovor OR ncovwuhan* OR "novel betacoronavirus" OR "Novel Coronavirus" OR "novel CoV" OR "sarscov-19" OR "sars-cov19" OR "sars-cov-19" OR sarscov19 OR "sarscov2" OR "sarscov-2" OR "sars-cov2" OR "sars-cov-2" OR "wn-cov" OR wncov):ti,ab
5	(epidem* OR outbreak* OR pandem* OR wildlife*) NEAR/1 (china* OR chinese* OR huanan* OR pneumonia OR respirator*):ti,ab
6	(("food market*" OR "seafood market*") NEAR/10 (china* OR chinese* OR huanan* OR hubei* OR wuhan*)):ti,ab
7	OR/1-6
8	"Antigens, Viral"[Mesh]
9	antigen:ti,ab
10	OR/8-9
11	MeSH descriptor: [Delayed Diagnosis] explode all trees
12	MeSH descriptor: [Diagnosis] this term only
13	MeSH descriptor: [Diagnosis, Differential] explode all trees
14	MeSH descriptor: [Diagnostic Techniques and Procedures] explode all trees
15	MeSH descriptor: [Early Diagnosis] explode all trees
16	Any MeSH descriptor in all MeSH products and with qualifier(s): [diagnosis - DI]
17	case* OR "case finding" OR casefinding OR detect* OR diagnos* OR screen* OR test*
18	OR/11-17
19	7 AND 10 AND 18
20	MeSH descriptor: [Animals] explode all trees
21	MeSH descriptor: [Humans] explode all trees
22	20 NOT 21
23	19 NOT 22
24	February 22, 2021 to Current

25

23 AND 24

Table s3. QUADAS-2 risk of bias assessment for included studies (95 studies)

	Author, Year	Patient Selection Risk of Bias	Index Test Risk of Bias	Reference Standard Risk of Bias	Flow & Timing Risk of Bias
1.	Albert 2021 [1]	Low	Low	Unclear	Low
2.	Alghounaim 2021 [2]	High	Low	Unclear	Low
3.	Allan-Blitz 2021 [3]	Low	Low	Unclear	Low
4.	Almendares 2022 [4]	Low	Low	Unclear	Low
5.	Alqahtani 2021 [5]	Low	Low	Unclear	Low
6.	Amer 2021 [6]	Low	Low	Unclear	Low
7.	Aoki 2021 [7]	High	Unclear	Low	Unclear
8.	Aoki 2021 [8]	High	Unclear	Unclear	Low
9.	Aranaz-Andrés 2022 [9]	Low	Unclear	Unclear	Low
10.	Baro 2021 [10]	Low	Low	Low	Low
11.	Beck 2021 [11]	Low	Low	Low	Low
12.	Bianco 2021 [12]	Low	Low	Unclear	Low
13.	Brihn 2021 [13]	Low	Low	Unclear	Low
14.	Bulilete 2021 [14]	Low	Low	Unclear	Low
15.	Carbonell-Sahuquillo 2021 [15]	Low	Low	Unclear	Low
16.	Caruana 2021 [16]	Low	Low	Unclear	Low
17.	Chiu 2021 [17]	Low	Unclear	Unclear	Unclear
18.	Courtellemont 2021 [18]	Low	Low	Unclear	Low
19.	Di Domenico 2021 [19]	Low	Low	Low	Low
20.	Dierks 2021 [20]	High	Low	Low	Low
21.	Drain 2021 [21]	Low	Low	Low	Low
22.	Drain 2021 [22]	Low	Low	Low	Low
23.	Escribano 2022 [23]	Low	Low	Low	Low
24.	Fernandez-Montero 2021 [24]	Low	Low	Low	Low

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25.	Fernández-Rivas 2021 [25]	Low	Low	Low	Low
26.	Ferté 2021 [26]	Low	Low	Low	Low
27.	Fitoussi 2021 [27]	Low	Low	Low	Low
28.	Ford 2021 [28]	Low	Low	Low	Low
29.	Fourati 2021 [29]	High	Low	Low	Unclear
30.	Fourati 2022 [30]	High	High	Low	Low
31.	García-Fiñana 2021 [31]	Low	Low	Low	Low
32.	Gili 2021 [32]	Low	Low	Low	Low
33.	González-Donapetry 2021 [33]	Low	Low	Low	Low
34.	Hagbom 2022 [34]	Low	Low	Low	Low
35.	Harris 2021 [35]	Low	Low	Low	Low
36.	Hirotsu 2021 [36]	Low	Low	High	Unclear
37.	Holzner 2021 [37]	Low	Low	High	Low
38.	Homza 2021 [38]	Low	Low	Low	Low
39.	Ifko 2021 [39]	Low	Low	Low	Low
40.	Ishii 2021 [40]	High	Low	High	High
41.	Jakobsen 2021 [41]	Low	Low	Low	Low
42.	James 2022 [42]	Low	Low	Low	Low
43.	Kahn 2021 [43]	Low	High	Low	Low
44.	Kernéis 2021 [44]	Low	Low	Low	Low
45.	Kim 2021 [45]	Low	Unclear	Low	Low
46.	Kim 2022 [46]	High	High	Low	Low
47.	Klajmon 2022 [47]	Low	Low	Low	low
48.	Klein 2021 [48]	Low	Low	Low	low
49.	Kolwijck 2021 [49]	Low	Low	Low	low
50.	Koskinen 2021 [50]	Low	High	Low	Low
51.	Krüger 2021 [51]	Low	Low	Low	Low
52.	Kumar 2021 [52]	Low	Low	Low	Low

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53.	Landaas 2021 [53]	Low	Low	Low	Low
54.	Leber 2021 [54]	Low	Low	Low	Low
55.	Leixner 2021 [55]	Low	Low	Low	Low
56.	Leli 2021 [56]	Low	Low	Low	Low
57.	Masiá 2021 [57]	Low	Low	Low	Low
58.	Mboumba 2021 [58]	High	Low	Low	Low
59.	Merino 2021 [59]	Low	Low	Low	Low
60.	Merino-Amador 2021 [60]	Low	Low	Low	Low
61.	Mitchell 2021 [61]	Low	Low	Low	Low
62.	Møller 2022 [62]	Low	Low	Low	Low
63.	Montalvo 2021 [63]	Low	Unclear	Unclear	Low
64.	Mungomklang 2021 [64]	Low	Unclear	Low	Low
65.	Murillo-Zamora 2021 [65]	Low	Unclear	Low	Low
66.	Nikolai 2021 [66]	Low	Unclear	Low	Low
67.	Nörz 2021 [67]	Low	Unclear	Low	Low
68.	Osterman 2021 [68]	High	Unclear	Unclear	Low
69.	Paul 2021 [69]	Low	High	Low	Low
70.	Peacock 2022 [70]	Low	Low	Low	Low
71.	Peña 2021 [71]	Low	Low	Unclear	Low
72.	Pérez-García 2021 [72]	High	Unclear	Unclear	Low
73.	Pérez-García 2021 [73]	Low	High	Low	Low
74.	Petronnet 2022 [74]	High	High	Low	Low
75.	Pilarowski 2021 [75]	Low	Low	Low	Low
76.	Pollock 2021 [76]	Low	Low	Low	Low
77.	Pray 2021 [77]	Low	Low	Unclear	Low
78.	Prince-Guerra 2021 [78]	Low	Low	Low	Low
79.	Quentin 2022 [79]	Low	Low	Low	Low
80.	Rahman 2021 [80]	Low	Low	Low	Low

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81.	Regev-Yochay 2022 [81]	Low	Low	Low	Low
82.	Schuit 2021 [82]	Low	Low	Low	Low
83.	Shaikh 2021 [83]	Low	Unclear	High	Low
84.	Siddiqui 2021 [84]	Low	Low	Low	Low
85.	Smith 2021 [85]	Low	Unclear	Unclear	Low
86.	Tinker 2021 [86]	Low	Unclear	Unclear	Low
87.	Tonen-Wolyec 2021 [87]	Low	Unclear	Unclear	Low
88.	Turcato 2022 [88]	Low	Unclear	Unclear	Low
89.	Van Der Moeren 2021 [89]	Low	Unclear	Unclear	Low
90.	Venekamp 2022 [90]	Low	Low	Low	Low
91.	Villaverde 2021 [91]	Low	Low	Low	Low
92.	Von Ahnen 2021 [92]	Low	Low	Low	Low
93.	Von Ahnen 2022 [93]	Low	Low	Low	Low
94.	Wachinger 2021 [94]	Low	Low	Low	High
95.	Winkel 2021 [95]	Low	Low	High	High

*Reference list is located after Table s4. Baseline Characteristics of the Included Studies

Table s4. Baseline Characteristics of the Included Studies

Author (Country)	Patient Selection	Index test	Reference Standard
Author, year: Albert 2021 [1] Country: Spain Study Design: Non-randomized/ observational study with comparator (e.g. case control)	Number of patients: 412 Age: median 36, range 17-91 Gender (%male): 42% male Symptomatic or Asymptomatic or Mix: Symptomatic Days since symptom onset (if applicable): tested if symptoms within past week	Test name(s): Panbio EUA certified: No CE certified: Yes Platform: Lateral flow immunochromatographic assay Target antigen: nucleocapsid antigen COI: NR Turnaround time: 15min Sample site: NP	Test name(s): TaqPath COVID-19 Kit Target gene: ORF1ab, N and S genes Ct threshold: NR Sample site: NP Type of swab: flocked swab Transport media: NA Self vs HCW: HCW

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		Type of swab: flocked swabs Transport media: NA HCW collection: Yes Sampling sequence: Unclear which one obtained first	
Author, year: Alghounaim 2021 [2] Country: Kuwait Study Design: Non-randomized/ observational study with comparator (e.g. case control)	Number of patients: 897 (Same patients as below row - all pts were swabbed twice for the two tests, not a separate cohort of patients) Age: median 40.2 (IQR 32.3-47.8) Gender (%male): 94% male Symptomatic or Asymptomatic or Mix: Mix	Test name(s): Diasorin LIAISON EUA certified: No CE certified: Yes Platform: chemiluminescence sandwich-immunoassay (CLIA) Target antigen: NR	Test name(s): TaqPath COVID-19 Kit Target gene: ORF1ab, N and S genes Ct threshold: NR Sample site: NP Type of swab: polyester-tipped 3-dimensionally printed swabs Transport media: NR - Collected samples intended for laboratory-based tests were

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	<p>Days since symptom onset (if applicable): 3% of patients symptomatic</p>	<p>COI: NR</p> <p>Turnaround time: 15-30min</p> <p>Sample site: NP</p> <p>Type of swab: polyester-tipped 3-dimensionally printed swabs</p> <p>Transport media: NR - Collected samples intended for laboratory-based tests were stored at a site and transported at 2–8°C immediately to the Jaber Innovation Laboratory at Jaber Alahmad Hospital and were processed within 12 h</p> <p>HCW collection: Yes</p> <p>Sampling sequence: Unclear which one obtained first</p>	<p>stored at a site and transported at 2–8°C immediately to the Jaber Innovation Laboratory at Jaber Alahmad Hospital and were processed within 12 h</p> <p>Self vs HCW: HCW</p>
	<p>Number of patients: 18457</p>	<p>Test name(s): BinaxNow</p>	<p>Test name(s): modified CDC protocol</p>

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Author, year: Allan-Blitz 2021 [3]	Age: NR	EUA certified: Yes	Target gene: NR
Country: FL, USA	Gender (%male): NR	CE certified: No	Ct threshold: 40
Study Design: Non-randomized/observational study with comparator (e.g. case control)	Symptomatic or Asymptomatic or Mix: Mix Days since symptom onset (if applicable): NR	Platform: lateral flow immunoassay Target antigen: NR COI: NR Turnaround time: NR Sample site: AN Type of swab: flocked swabs Transport media: NR	Sample site: AN Type of swab: flocked swabs Transport media: NR Self vs HCW: Mix - health care worker-observed self-collected oral fluid swabs, self-collected anterior nares swabs, and clinician-collected nasopharyngeal swabs.

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		HCW collection: Yes Sampling sequence: Ag first	
Author, year: Almendares 2022 [4] Country: AZ, USA Study Design: Non-randomized/observational study with comparator (e.g. case control)	Number of patients: 3419 Age: median 41 (range 10-95) Gender (%male): 37.7% male, 49.2% female, 13.1% undisclosed Symptomatic or Asymptomatic or Mix: Mix Days since symptom onset (if applicable): NR	Test name(s): BinaxNow EUA certified: Yes CE certified: No Platform: lateral flow immunoassay Target antigen: NR COI: NR Turnaround time: 15-30min Sample site: AN	Test name(s): CDC 2019-nCoV rRT-PCR diagnostic panel (CDC rRT-PCR assay) for detection of SARS-CoV-2 (n = 2,582) or the Fosun COVID19 rRT-PCR detection kit (Fosun rRT-PCR assay) (n = 837) Target gene: Statistical analysis was limited to data collected from specimens tested by the CDC rRT-PCR assay when using the nucleocapsid 1 (N1) cycle threshold value as a variable, because the two rRT-PCR assays detect different SARS-CoV-2 gene targets and CT value Ct threshold: NR Sample site: NP Type of swab: NR

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		<p>Type of swab: Swab provided in collection kit was used</p> <p>Transport media: NA</p> <p>HCW collection: Yes</p> <p>Sampling sequence: Ag first</p>	<p>Transport media: NR</p> <p>Self vs HCW: HCW</p>
<p>Author, year: Alqahtani 2021 [5]</p> <p>Country: Bahrain</p> <p>Study Design: Non-randomized/ observational study with comparator (e.g. case control)</p>	<p>Number of patients: 4183</p> <p>Age: median 30.9 +/- 14.5</p> <p>Gender (%male): 56.5% male</p> <p>Symptomatic or Asymptomatic or Mix: Symptomatic</p> <p>Days since symptom onset (if applicable): median 2 days (range 1-3)</p>	<p>Test name(s): Panbio</p> <p>EUA certified: No</p> <p>CE certified: Yes</p> <p>Platform: Lateral flow immunochromatographic assay</p> <p>Target antigen: Nucleocapsid protein</p>	<p>Test name(s): TaqPath COVID-19 Kit</p> <p>Target gene: E gene - If the E gene was detected, the sample was then confirmed by RdRP and N genes</p> <p>Ct threshold: Ct values >40 were considered negative</p> <p>Sample site: NP</p> <p>Type of swab: NR</p>

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		COI: NR Turnaround time: 15min Sample site: AN Type of swab: Swab provided with kit Transport media: NA HCW collection: Yes Sampling sequence: Ag first	Transport media: NA Self vs HCW: HCW
Author, year: Amer 2021 [6] Country: Egypt	Number of patients: 83 Age: median 55.5 +/- 18.4	Test name(s): Standard Q EUA certified: No	Test name(s): real-time PCR kit (Primerdesign Ltd, Ref: Z-Path-COMD-19CE, UK) in Stratagene Mx3000P qPCR System (Agilent). Target gene: RdRp gene

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Study Design: Non-randomized/observational study with comparator (e.g. case control)	Gender (%male): 59% male Symptomatic or Asymptomatic or Mix: Mix Days since symptom onset (if applicable): 50.6% of patients with symptoms, 6.2% asymptomatic, 43.3% not reported. Of the people symptomatic, 45.7% had symptoms within 0-7 days, 32.5% had symptoms within 8-16, 4.8% had symptoms >16 days, 1.2% with symptoms after sampling	CE certified: Yes Platform: Lateral flow immunochromatographic assay Target antigen: Nucleocapsid protein COI: NR Turnaround time: 15-30min Sample site: NP and OP swab mixed into single VTM for testing Type of swab: NR Transport media: VTM Self vs HCW: HCW	Ct threshold: The analyzed samples were considered negative if they have a Ct value ≥40 or when no Ct values were reported. For positive samples, SARS-CoV-2 RNR content was categorized according to the Ct values into strong (Ct < 29), moderate (Ct = 29–36) and weak (Ct ≥ 37)
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		Sampling sequence: Same sample for both	
Author, year: Aoki 2021a [7]	Number of patients: 548 Age: NR Gender (%male): NR Symptomatic or Asymptomatic or Mix: Symptomatic Days since symptom onset (if applicable): collected from COVID-19 patients or suspected patients with COVID-19	Test name(s): Lumipulse G EUA certified: No CE certified: Yes Platform: chemiluminescent enzyme immunoassay (CLEIA) Target antigen: NR COI: Using a cut-off value of 1.34 pg/ml Turnaround time: NR Sample site: NP Type of swab: NR	Test name(s): One-step RT-qPCR was performed using QuantStudio® 5 (Applied Biosystems™, U.S.A.) or BD MAX™ open system, respectively. TaqMan Fast Virus 1-Step Master Mix (Thermo Fisher Scientific, U.S.A.) and BD MAX™ TNR MMK (SPC) were used as one-step RT-qPCR master m Target gene: N gene Ct threshold: NR Sample site: NP Type of swab: NR Transport media: NR Self vs HCW: NR

		Transport media: NR HCW collection: NR Sampling sequence: Same sample used for both tests	
Author, year: Aoki 2021b [8] Country: Japan Study Design: Non-randomized/observational study with comparator (e.g. case control)	Number of patients: 129 Age: NR Gender (%male): NR Symptomatic or Asymptomatic or Mix: Symptomatic Days since symptom onset (if applicable): All	Test name(s): Espline EUA certified: No CE certified: Yes Platform: Immunochromatographic assay Target antigen: N antigen COI: NR	Test name(s): One-step RT-qPCR was performed using QuantStudio® 5 (Applied Biosystems™, U.S.A.) or BD MAX™ open system, respectively. TaqMan Fast Virus 1-Step Master Mix (Thermo Fisher Scientific, U.S.A.) and BD MAX™ TNR MMK (SPC) were used as one-step RT-qPCR master m Target gene: N gene Ct threshold: NR Sample site: NP Type of swab: NR

		<p>Turnaround time: 30min</p> <p>Sample site: NP</p> <p>Type of swab: NR</p> <p>Transport media: Some frozen samples taken from UVT</p> <p>HCW collection: NR</p> <p>Sampling sequence: NR</p>	<p>Transport media: Some frozen samples taken from UVT</p> <p>Self vs HCW: NR</p>
Author, year: Aranaz-Andrés 2022 [9]	Number of patients: 541 Age: median 76.7 Gender (%male): 56% male	Test name(s): Panbio EUA certified: No CE certified: Yes	Test name(s): TaqPath COVID-19 Kit Target gene: Orf1ab, N, S gene Ct threshold: Only samples with Ct value ≤ 35 were considered positives. Sample site: NP

Supplementary Materials

observational study with comparator (e.g. case control)	Symptomatic or Asymptomatic or Mix: Asymptomatic Days since symptom onset (if applicable): NA	Platform: Lateral flow immunochromatographic assay Target antigen: NR COI: NR Turnaround time: 15min Sample site: NP Type of swab: NR Transport media: NA HCW collection: Yes Sampling sequence: Random - switches which one is first	Type of swab: NR Transport media: NA Self vs HCW: HCW

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Author, year: Baro 2021 [10]	Number of patients: 286 Age: NR Gender (%male): NR Symptomatic or Asymptomatic or Mix: Asymptomatic Days since symptom onset (if applicable): NA	Test name(s): Multiple EUA certified: Yes CE certified: Yes Platform: Multiple Target antigen: NR COI: NR Turnaround time: 15min Sample site: NP Type of swab: NR Transport media: VTM	Test name(s): Allplex 2019-nCoV assay Target gene: NR Ct threshold: NR Sample site: NP Type of swab: NR Transport media: VTM Self vs HCW: HCW
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		HCW collection: Yes Sampling sequence: NR - RT PCR test completed before Ag but unclear which sample was obtained first or if same sample used	
Author, year: Beck 2020 [11] Country: US Study Design: Non-randomized/ observational study with comparator (e.g. case control)	Number of patients: 347 Age: range 1-90 years Gender (%male): NR Symptomatic or Asymptomatic or Mix: Symptomatic Days since symptom onset (if applicable): 0-5, >5days	Test name(s): Sofia EUA certified: Yes CE certified: Yes Platform: Lateral Flow Assay Target antigen: Nucleocapsid COI: NR Turnaround time: NR	Test name(s): Hologic Aptima, Cepheid Xpert Xpress used for discrepant results (data from CXN not used) Target gene: ORF1ab Ct threshold: NR Sample site: NP Type of swab: mini-tip nylon flocked swab Transport media: Amies bacterial transport medium

		<p>Sample site: AN</p> <p>Type of swab: Provided by manufacturer</p> <p>Transport media: dry swab</p> <p>HCW collection: Unclear</p> <p>Sampling sequence: PCR first</p>	<p>Self vs HCW: Unclear</p>
<p>Author, year: Bianco 2021 [12]</p> <p>Country: Italy</p> <p>Study Design: Non-randomized/ observational study with</p>	<p>Number of patients: 907</p> <p>Age: mean 47.9</p> <p>Gender (%male): 55.7% male</p> <p>Symptomatic or Asymptomatic or Mix: Mix</p>	<p>Test name(s): LumiraDx</p> <p>EUA certified: No</p> <p>CE certified: Yes</p> <p>Platform: fluorescence immunoassay (FIA)</p>	<p>Test name(s): Xpert Xpress SARS-CoV-2 assay (Cepheid, Sunnyvale, CA)</p> <p>Target gene: NR</p> <p>Ct threshold: NR</p> <p>Sample site: NP</p> <p>Type of swab: NR</p>

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comparator (e.g. case control)	<p>Days since symptom onset (if applicable): At specimen collection, 676 (74.5%) participants were asymptomatic and 231 (25.5%) reported experiencing one or more COVID-19 symptoms. The median interval from symptom onset to specimen collection was 4 days (interquartile range 2-7).</p>	<p>Target antigen: Nucleocapsid protein</p> <p>COI: NR</p> <p>Turnaround time: NR</p> <p>Sample site: AN</p> <p>Type of swab: NR</p> <p>Transport media: NR</p> <p>HCW collection: Yes</p> <p>Sampling sequence: NR</p>	<p>Transport media: NR</p> <p>Self vs HCW: HCW</p>
Author, year: Brihn 2021 [13]	<p>Number of patients: 2039</p> <p>Age: median 56 (range 16-107)</p>	<p>Test name(s): Multiple</p> <p>EUA certified: No</p>	<p>Test name(s): Fulgent COVID-19 RT-PCR (Fulgent Genetics) (RT-PCR test)</p> <p>Target gene: N1 and N2 gene</p>

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Country: California, USA	Gender (%male): 45% male Symptomatic or Asymptomatic or Mix: Mix Days since symptom onset (if applicable): 15% had covid symptoms	CE certified: Yes Platform: Multiple Target antigen: NR COI: NR Turnaround time: NR Sample site: AN Type of swab: NR Transport media: NA HCW collection: Yes	Ct threshold: Ct values <40 Sample site: NP Type of swab: NR Transport media: NA Self vs HCW: HCW
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		Sampling sequence: Ag first, NP second	
Author, year: Bulilete 2021 [14]	Number of patients: 1369 Age: mean 42.5 +/- 14.9 Gender (%male): 45.7% male Symptomatic or Asymptomatic or Mix: Mix Days since symptom onset (if applicable): 36.7% with symptoms, 54.8% close contact, 8.5% unknown symptoms. Of patients with symptoms, 70.6% with testing 5 days or less since symptom onset or close contact, 15.7% with >5 days.	Test name(s): Panbio EUA certified: No CE certified: Yes Platform: Lateral flow immunochromatographic assay Target antigen: NR COI: NR Turnaround time: NR Sample site: NP Type of swab: NR	Test name(s): TaqPath COVID-19 Kit Target gene: ORF, N, S gene Ct threshold: NR Sample site: NP Type of swab: NR Transport media: NR Self vs HCW: HCW

		Transport media: NR HCW collection: Yes Sampling sequence: NR	
Author, year: Carbonell-Sahuquillo 2021 [15] Country: Spain Study Design: Non-randomized/observational study with comparator (e.g. case control)	Number of patients: 357 Age: median 2 (IQR 1-6) Gender (%male): 56.5% Symptomatic or Asymptomatic or Mix: Symptomatic Days since symptom onset (if applicable): median 1 days since symptom onset (IQR 1-3)	Test name(s): Panbio EUA certified: No CE certified: Yes Platform: Lateral flow immunochromatographic assay Target antigen: NR COI: NR	Test name(s): TaqPath COVID-19 Kit Target gene: ORF, N, S gene Ct threshold: NR Sample site: NP Type of swab: NR Transport media: NR

		Turnaround time: NR Sample site: NP Type of swab: NR Transport media: NR HCW collection: Yes Sampling sequence: NR	Self vs HCW: HCW
Author, year: Caruana 2021 [16] Country: Switzerland	Number of patients: 532 Age: median 67 (IQR 48.5, 81.0) for patients without symptoms, median 75 (IQR 61.0, 85.0) for patients with symptoms	Test name(s): Multiple EUA certified: No CE certified: Yes	Test name(s): VIASURE SARS-CoV-2 (N1 + N2) Real-Time PCR Detection Kit for BD MAX (Becton Dickinson, Franklin Lake, NJ, USA) or GeneXpert SARS-CoV-2 test (Cepheid, Sunnyvale, CA, USA) Target gene: NR Ct threshold: NR

Supplementary Materials

Study Design: Non-randomized/observational study with comparator (e.g. case control)	Gender (%male): 56.1% male for patients without symptoms, 55.3% male for patients with symptoms Symptomatic or Asymptomatic or Mix: Mix Days since symptom onset (if applicable): NR	Platform: Lateral Flow Assay Target antigen: NR COI: NR Turnaround time: 15-30min for Standard Q and Panbio, 20min for Exdia and BD Veritor Sample site: NP Type of swab: wet swab Transport media: VTM HCW collection: NR Sampling sequence: Same swab for both tests	Sample site: NP Type of swab: wet swab Transport media: VTM Self vs HCW: NR

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Author, year: Chiu 2021 [17]	Number of patients: 329 symptomatic in Cali and 22,994 asymptomatic in Hong Kong	Test name(s): INDICAID EUA certified: No CE certified: Yes Platform: lateral flow immunoassay Target antigen: NR COI: Positive Ct median 18.13 [IQR 16.29 – 22.96] for San Fran/Oakland group. Positive Ct median 19.55 [IQR 17.07 – 24.35] for San Fernando.	Test name(s): RT-PCR Target gene: NR Ct threshold: NR Sample site: AN Type of swab: NR Transport media: NR Self vs HCW: HCW and self collected
Country: California and Hong Kong	Age: San Fran and Oakland population - 83 patients total, 2 removed due to samples lost in transport, median age 32 (IQR 25,44). San Fernando - 270 patients total, 2 removed due to samples lost in transport, median 35yo (IQR 24,50)		
Study Design: Non-randomized/observational study with comparator (e.g. case control)	Gender (%male): San Fran and Oakland population - 83 patients total, 2 removed due to samples lost in transport, 55.6% male. San Fernando - 270 patients total, 2 removed due to samples lost in transport, 47.4% male		
	Symptomatic or Asymptomatic or Mix: Mix	Turnaround time: 20min	
	Days since symptom onset (if applicable): within 5 days for symptomatic patients	Sample site: AN Type of swab: NR	

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		Transport media: NR HCW collection: Yes Sampling sequence: NR	
Author, year: Courtellemont 2021 [18]	Number of patients: 248 Age: NR Gender (%male): 47.1% male Symptomatic or Asymptomatic or Mix: Mix Days since symptom onset (if applicable): Ninety seven patients were symptomatic, and 24 were totally asymptomatic. The median time of symptom duration before	Test name(s): COVID VIRO EUA certified: No CE certified: Yes Platform: Lateral flow immunochromatographic assay Target antigen: NR COI: NR	Test name(s): TaqPath COVID-19 Kit Target gene: ORF1ab, S, N gene Ct threshold: Samples showing an exponential growth curve and a cycle threshold (Ct) value < 37 were considered positive. A unique Ct value > 37 was considered negative. Sample site: NP (but subset had OP/saliva for comparison) Type of swab: A polyester-tipped flexible (viral transport medium tube with swab VTM, Sun-Trine®) was inserted

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	<p>sampling was 5 days (mean, 6 days; range, 1–20).</p>	<p>Turnaround time: 15min</p> <p>Sample site: NP (but subset has OP/saliva too for comparison)</p> <p>Type of swab: A polyester-tipped flexible (viral transport medium tube with swab VTM, Sun-Trine®) was inserted</p> <p>Transport media: NA</p> <p>HCW collection: Yes</p> <p>Sampling sequence: RT-PCR sample first, Ag second</p>	<p>Transport media: NA</p> <p>Self vs HCW: HCW</p>
<p>Author, year: Di Domenico 2021 [19]</p> <p>Country: Italy</p>	<p>Number of patients: 433</p> <p>Age: median age—37 years, range—15–78</p>	<p>Test name(s): Panbio</p> <p>EUA certified: No</p> <p>CE certified: Yes</p>	<p>Test name(s): RT-PCR assay for SARS-CoV-2 (QuantStudioTM, Thermo Fisher Scientific, Waltham, MA, USA)</p> <p>Target gene: Orf-1ab, N Protein and S Protein</p>

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Study Design: Non-randomized/observational study with comparator (e.g. case control)	Gender (%male): NR Symptomatic or Asymptomatic or Mix: Asymptomatic Days since symptom onset (if applicable): NA	Platform: Lateral flow immunochromatographic assay Target antigen: Nucleocapsid protein COI: NR Turnaround time: 15-20 min Sample site: NP Type of swab: NR Transport media: NR HCW collection: Yes Sampling sequence: antigen first	Ct threshold: 35 Sample site: NP Type of swab: NR Transport media: NR Self vs HCW: HCW

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Author, year: Dierks 2021 [20]	Number of patients: 444 Age: NR Gender (%male): NR Symptomatic or Asymptomatic or Mix: Asymptomatic Days since symptom onset (if applicable): NA	Test name(s): Multiple EUA certified: No CE certified: Yes Platform: Lateral Flow Assay Target antigen: Nucleocapsid protein COI: NR Turnaround time: NR Sample site: NP Type of swab: NR Transport media: NR	Test name(s): Roche Cobas SARS-CoV-2 PCR, the Genesig Real-Time PCR Coronavirus (COVID-19) assay, or the Cepheid Xpert Xpress SARS-CoV-2 PCR Target gene: NR Ct threshold: 30 Sample site: NP Type of swab: NR Transport media: NR Self vs HCW: NR
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		HCW collection: NR Sampling sequence: NR	
Author, year: Drain 2021a [21] Country: USA Study Design: Non-randomized/observational study with comparator (e.g. case control)	Number of patients: 222 Age: mean (SD) 38.7 (17.3) Gender (%male): 82 (26.9%) Symptomatic or Asymptomatic or Mix: Asymptomatic Days since symptom onset (if applicable): NA	Test name(s): LumiraDx EUA certified: No CE certified: Yes Platform: fluorescence immunoassay (FIA) Target antigen: Nucleocapsid protein COI: NR Turnaround time: 12min	Test name(s): TaqPath COVID-19 Kit or Cobas 6800 Roche Target gene: NR Ct threshold: 33 Sample site: AN Type of swab: NR Transport media: NR Self vs HCW: Yes

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		Sample site: AN Type of swab: NR Transport media: NR HCW collection: Yes Sampling sequence: same time	
Author, year: Drain 2021b [22] Country: Multiple sites, UK and USA Study Design: Non-randomized/observational study with	Number of patients: 512 Age: 0-90yo, mean 34 (\pm 15.7) for ANS, mean 33.2 (\pm 19.4) for NPS Gender (%male): (44%) Symptomatic or Asymptomatic or Mix: Mix	Test name(s): LumiraDx EUA certified: Yes CE certified: Yes Platform: Microfluidic Immunofluorescence Assay Target antigen: Nucleocapsid	Test name(s): Roche cobas SARS-CoV-2 Target gene: ORF1ab, E Ct threshold: NR Sample site: AN, NP Type of swab: ANS: Copan FLOQ swab, NP: NR

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comparator (e.g. case control)	Days since symptom onset (if applicable): Average duration of four days at testing (for symptomatic)	COI: NR Turnaround time: NR Sample site: AN, NP Type of swab: NR Transport media: 0.7 mL of a proprietary extraction buffer HCW collection: Yes Sampling sequence: same time	Transport media: 3 ml of VTM Self vs HCW: HCW
Author, year: Escribano 2022 [23] Country: Spain	Number of patients: 900 Age: NR	Test name(s): Multiple EUA certified: No	Test name(s): TaqPath COVID-19 Kit Target gene: NR Ct threshold: 20

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Study Design: Non-randomized/observational study with comparator (e.g. case control)	Gender (%male): 361 (40%) Symptomatic or Asymptomatic or Mix: Mix Days since symptom onset (if applicable): NA	CE certified: Yes Platform: Lateral flow Assay Target antigen: Nucleocapsid protein COI: NR Turnaround time: NR Sample site: NP Type of swab: NR Transport media: NR HCW collection: Yes Sampling sequence: NR	Sample site: NP Type of swab: NR Transport media: NR Self vs HCW: HCW
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Author, year: Fernandez- Montero 2021 [24]	Number of patients: 2288 Age: Mean (SD) 28 (12.3)	Test name(s): Roche SD EUA certified: No	Test name(s): RT-PCR Target gene: NR
Country: Spain	Gender (%male): 896 (38%)	CE certified: Yes	Ct threshold: 35 Sample site: OP
Study Design: Non- randomized/ observational study with comparator (e.g. case control)	Symptomatic or Asymptomatic or Mix: Asymptomatic Days since symptom onset (if applicable): NA	Platform: Lateral flow immunochromatographic assay Target antigen: NR COI: NR Turnaround time: NR Sample site: NP Type of swab: NR	Type of swab: NR Transport media: NR Self vs HCW: HCW

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		Transport media: NR HCW collection: Yes Sampling sequence: NR	
Author, year: Fernández-Rivas 2021 [25]	Number of patients: 861 Age: median: 44 years old, range 1-94 Country: Spain Study Design: Non-randomized/ observational study with comparator (e.g. case control)	Test name(s): Diasorin LIAISON EUA certified: No Gender (%male): 245 (28%) Symptomatic or Asymptomatic or Mix: Asymptomatic Days since symptom onset (if applicable): NA	Test name(s): Allplex 2019-nCoV assay Target gene: NR Ct threshold: 40 Sample site: AN Type of swab: NR Platform: chemiluminescence sandwich-immunoassay (CLIA) Target antigen: Nucleocapsid protein COI: 100 TCID50/ml Turnaround time: NR

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		Sample site: AN Type of swab: NR Transport media: NR HCW collection: NR Sampling sequence: NR	
Author, year: Ferté 2021 [26] Country: France Study Design: Non-randomized/ observational study with	Number of patients: 688 Age: Mean (SD) 22.95 (5.52) Gender (%male): 236 (34.0%) Symptomatic or Asymptomatic or Mix: Mix	Test name(s): Panbio EUA certified: No CE certified: Yes Platform: Lateral flow immunochromatographic assay	Test name(s): ARGENE SARS-CoV-2 R-GENE, BioM'erieux, France Target gene: RT-PCR was considered positive when both nucleocapsid (N) and RNA-dependent RNR polymerase (RdRp) genes were detected Ct threshold: 33 Sample site: NP

Supplementary Materials

comparator (e.g. case control)	Days since symptom onset (if applicable): NR	Target antigen: NR COI: NR Turnaround time: NR Sample site: NP Type of swab: NR Transport media: NR HCW collection: Yes Sampling sequence: NR	Type of swab: NR Transport media: NR Self vs HCW: HCW
Author, year: Fitoussi 2021 [27] Country: France	Number of patients: 967 Age: Median 34 Range (18–83)	Test name(s): BIOSYNEX BSS EUA certified: Yes	Test name(s): Real-Time Multiplex RT-PCR Kit (detection for three genes) (Liferiver and Shanghai ZJ Bio-Tech Co., Ltd), Target gene: (E, N and RdRP)

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Study Design: Non-randomized/observational study with comparator (e.g. case control)	Gender (%male): 469 (48.5%) Symptomatic or Asymptomatic or Mix: Mix Days since symptom onset (if applicable): 0-20days	CE certified: Yes Platform: immunochromatographic assay Target antigen: Nucleocapsid protein COI: NR Turnaround time: most seen within 5min Sample site: NP Type of swab: swab provided in the BIOSYNEX Ag-RDT Transport media: NR HCW collection: Yes	Ct threshold: 41 Sample site: NP Type of swab: NR Transport media: NR Self vs HCW: HCW
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		Sampling sequence: Antigen first	
Author, year: Ford 2021 [28]	Number of patients: 2110 Age: NR Gender (%male): NR Symptomatic or Asymptomatic or Mix: Mix Days since symptom onset (if applicable): NA	Test name(s): BinaxNow EUA certified: Yes CE certified: No Platform: lateral flow immunoassay Target antigen: NR COI: NR Turnaround time: NR Sample site: NP Type of swab: NR	Test name(s): TaqPath COVID-19 Kit Target gene: ORF1ab, S-gene, and N-gene Ct threshold: 37 Sample site: AN Type of swab: NR Transport media: NR Self vs HCW: Supervised Self

		Transport media: NR HCW collection: No Sampling sequence: same time	
Author, year: Fourati 2021 [29]	Number of patients: 634 Age: NR Gender (%male): NR Symptomatic or Asymptomatic or Mix: Symptomatic Days since symptom onset (if applicable): NR	Test name(s): Multiple EUA certified: No CE certified: Yes Platform: Lateral Flow Assay Target antigen: NR COI: NR	Test name(s): RealStar SARS-CoV2 RT PCR Kit 1.0, AltoNR Diagnostics GmbH, Hamburg, Germany Target gene: NR Ct threshold: 31 Sample site: NP Type of swab: NR Transport media: NR

		Turnaround time: 30min Standard Q, Abbott Panbio 15-20min, all others 15min Sample site: NP Type of swab: NR Transport media: NR HCW collection: Yes Sampling sequence: NR	Self vs HCW:
Author, year: Fourati 2022 [30]	Number of patients: 1763 Age: NR Gender (%male): NR	Test name(s): VITROS EUA certified: No CE certified: Yes	Test name(s): commercially available NAAT (TMA-based Aptima™ SARS CoV-2 Assay, Hologic, San Diego, California; or PCRbased Alinity m® SARS CoV-2 Assay Abbott, Germany). Target gene: NR Ct threshold: 35

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study with comparator (e.g. case control)	Symptomatic or Asymptomatic or Mix: Symptomatic Days since symptom onset (if applicable): NR	Platform: chemiluminescent immunoassay (CLIA) Target antigen: NR COI: NR Turnaround time: 48min Sample site: NP Type of swab: NR Transport media: NR HCW collection: NR Sampling sequence: NR	Sample site: NP Type of swab: NR Transport media: NR Self vs HCW: NR
	Number of patients: 5869	Test name(s): Innova	Test name(s): TaqPath COVID-19 Kit

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Author, year: García-Fiñana 2021 [31]	Age: mean 50 SD 18	EUA certified: Yes	Target gene: NR
Country: UK	Gender (%male): 2700 (46%)	CE certified: Yes	Ct threshold:
Study Design: Non-randomized/ observational study with comparator (e.g. case control)	Symptomatic or Asymptomatic or Mix: Asymptomatic Days since symptom onset (if applicable): NA	Platform: Lateral flow immunochromatographic assay Target antigen: NR COI: NR Turnaround time: 30min Sample site: Throat + nose Type of swab: NR Transport media: NR	Sample site: Throat and nose combo Type of swab: NR Transport media: NR Self vs HCW: Supervised Self

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		HCW collection: No Sampling sequence: NR	
Author, year: Gili 2021 [32] Country: Italy Study Design: Non-randomized/observational study with comparator (e.g. case control)	Number of patients: 226 Age: NR Gender (%male): NR Symptomatic or Asymptomatic or Mix: Mix Days since symptom onset (if applicable): NR	Test name(s): Lumipulse G EUA certified: No CE certified: Yes Platform: chemiluminescent enzyme immunoassay (CLEIA) Target antigen: NR COI: 1.340 ng/ml Turnaround time: NR Sample site: NP	Test name(s): Allplex 2019-nCoV assay Target gene: NR Ct threshold: 40 Sample site: NP Type of swab: NR Transport media: NR Self vs HCW: NR

		Type of swab: NR Transport media: NR HCW collection: NR Sampling sequence: NR	
Author, year: González-Donapetry 2021 [33]	Number of patients: 440 Age: Median: 3 IQR: (1–7)	Test name(s): Panbio EUA certified: No	Test name(s): Allplex 2019-nCoV assay Target gene: NR Ct threshold: 40
Country: Spain	Gender (%male): 59.1%	CE certified: Yes	Sample site: NP
Study Design: Non-randomized/ observational study with	Symptomatic or Asymptomatic or Mix: Symptomatic	Platform: Lateral flow immunochromatographic assay Target antigen: NR	Type of swab: NR

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comparator (e.g. case control)	Days since symptom onset (if applicable): Median of 1 (IQR:1-3)	COI: 1.340 ng/ml Turnaround time: NR Sample site: NP Type of swab: NR Transport media: NR HCW collection: NR Sampling sequence: NR	Transport media: NR Self vs HCW: NR
Author, year: Hagbom 2022 [34]	Number of patients: 39 Age: Median 57.0 range (32–84)	Test name(s): BTNX EUA certified: No	Test name(s): QIAmp Viral RNR kit Target gene: NR

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Country: Sweden Study Design: Non-randomized/observational study with comparator (e.g. case control)	Gender (%male): 28 (71.8%) Symptomatic or Asymptomatic or Mix: Symptomatic Days since symptom onset (if applicable): Median 11.0 Range(5–30)	CE certified: Yes Platform: immunochromatographic assay Target antigen: NR COI: NR Turnaround time: 15min Sample site: saliva Type of swab: NR Transport media: NR HCW collection: Yes	Ct threshold: NR Sample site: saliva Type of swab: NR Transport media: NR Self vs HCW: NR
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Supplementary Materials

		Sampling sequence: NR	
Author, year: Harris 2021 [35]	Number of patients: 2436 Age: NR Gender (%male): NR Symptomatic or Asymptomatic or Mix: Mix Days since symptom onset (if applicable): NA	Test name(s): Sofia EUA certified: No CE certified: Yes Platform: Fluorescent Immunoassay (FIA) Target antigen: NR COI: NR Turnaround time: NR Sample site: AN Type of swab: Dry swabs	Test name(s): CDC 2019-Novel Coronavirus (2019-nCoV) Real-Time RT-PCR Diagnostic Panel Target gene: NR Ct threshold: 20-23 Sample site: NP Type of swab: NR Transport media: NR Self vs HCW: HCW

Supplementary Materials

		Transport media: NR HCW collection: No Sampling sequence: PCR first	
Author, year: Hirotsu 2021 [36] Country: Japan Study Design: Non-randomized/ observational study with comparator (e.g. case control)	Number of patients: 637 Age: Gender (%male): Symptomatic or Asymptomatic or Mix: Mix Days since symptom onset (if applicable):	Test name(s): Lumipulse G EUA certified: No CE certified: Yes Platform: chemiluminescent enzyme immunoassay (CLEIA) Target antigen: NR COI: NR	Test name(s): StepOnePlus Real-Time PCR System Target gene: nucleocapsid gene of SARS-CoV-2 (NC_045512.2) Ct threshold: NR Sample site: NP Type of swab: cotton swabs Transport media: VTM

		Turnaround time: NR Sample site: NP Type of swab: cotton swabs Transport media: VTM HCW collection: NR Sampling sequence: NR	Self vs HCW: HCW
Author, year: Holzner 2021 [37]	Number of patients: 2375 Age: NR Gender (%male): NR	Test name(s): Standard Q EUA certified: No CE certified: Yes	Test name(s): Alinity in the SARS-CoV-2 assay (Abbott), Cepheid (GeneXpert), and RealStar SARS-CoV-2RT-PCR (AltoNR Diagnostics). Target gene: NR Ct threshold: 30 Sample site: NP

Supplementary Materials

study with comparator (e.g. case control)	Days since symptom onset (if applicable): NA	Target antigen: NR COI: NR Turnaround time: 15-30min Sample site: NP Type of swab: NR Transport media: NR HCW collection: Yes Sampling sequence: 1st	Type of swab: NR Transport media: NR Self vs HCW: HCW
Author, year: Homza 2021 [38]	Number of patients: 494	Test name(s): Ecotest	Test name(s): Multiplex RT-PCR Kit (DiaNR Biotechnologies, Czech Republic)

Supplementary Materials

Country: Czech Republic	Age: mean age: 42.2±15.1years (min. 7; max. 81)	EUA certified: No CE certified: Yes Platform: Lateral flow immunoassay Target antigen: NR COI: NR Turnaround time: NR Sample site: NP Type of swab: NR Transport media: Medium supplied with the antigen test HCW collection: Yes	Target gene: genes coding the Spike protein and EndoRNase. Ct threshold: NR Sample site: NP Type of swab: HCW
Study Design: Non-randomized/observational study with comparator (e.g. case control)	Gender (%male): Symptomatic or Asymptomatic or Mix: Mix Days since symptom onset (if applicable): NR		

		Sampling sequence: 1st	
Author, year: Ifko 2021 [39]	Number of patients: 455 Age: NR Gender (%male): NR	Test name(s): ALLTEST EUA certified: No CE certified: Yes - NADAL and Alltest Platform: Lateral flow assay Target antigen: NR COI: NR Turnaround time: NR Sample site: NP	Test name(s): Seegene Allplex 2019-nCoV test, Cobas 6800 SARS-CoV-2 Test, SARS-CoV-2 RT-PCR assay (PCR Biosystems Ltd, London, UK), Target gene: NR Ct threshold: NR Sample site: NP Type of swab: NR Transport media: NR Self vs HCW: HCW

Supplementary Materials

		Type of swab: NR Transport media: NR HCW collection: NR Sampling sequence: 1st	
Author, year: Ishii 2021 [40] Country: Japan Study Design: Non-randomized/ observational study with comparator (e.g. case control)	Number of patients: 486 Age: NR Gender (%male): NR Symptomatic or Asymptomatic or Mix: Mix Days since symptom onset (if applicable): NR	Test name(s): Lumipulse G EUA certified: No CE certified: Yes Platform: chemiluminescent enzyme immunoassay (CLEIA) Target antigen: NR COI: NR	Test name(s): TaqPath COVID-19 Kit Target gene: NR Ct threshold: 35 Sample site: NP and saliva Type of swab: NR Transport media: NR

		Turnaround time: NR Sample site: NP Type of swab: NR Transport media: NR HCW collection: NR Sampling sequence: NR	Self vs HCW: HCW
Author, year: Jakobsen 2021 [41] Country: Denmark	Number of patients: 4811 Age: median age: 45 years (interquartile range: 30-56) Gender (%male): 47	Test name(s): Standard Q EUA certified: No CE certified: Yes	Test name(s): LuNR Universal Probe One-step RT-qPCR kit Target gene: NR Ct threshold: 10-38 Sample site: OP

Supplementary Materials

Study Design: Non-randomized/observational study with comparator (e.g. case control)	Symptomatic or Asymptomatic or Mix: Mix Days since symptom onset (if applicable): NR	Platform: Lateral flow immunochromatographic assay Target antigen: NR COI: NR Turnaround time: NR Sample site: NP Type of swab: NR Transport media: NR HCW collection: NR Sampling sequence: 2nd	Type of swab: NR Transport media: NR Self vs HCW: HCW
	Number of patients: 2339	Test name(s): BinaxNow	

Supplementary Materials

Author, year: James 2022 [42]	Age: (median, 37 years) age range 16-98 Gender (%male): NR Symptomatic or Asymptomatic or Mix: Mix Days since symptom onset (if applicable): NR	EUA certified: Yes CE certified: No Platform: lateral flow immunoassay Target antigen: nucleocapsid protein antigen COI: NR Turnaround time: NR Sample site: AN Type of swab: NR Transport media: NR	Test name(s): PerkinElmer SARS-CoV-2 real-time RT-PCR assay. Target gene: N and Orf1 gene Ct threshold: 42 Sample site: AN Type of swab: flocked swab Transport media: NR Self vs HCW: HCW
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Supplementary Materials

		HCW collection: NR Sampling sequence: random	
Author, year: Kahn 2021 [43]	Number of patients: 3630 Age: NR Gender (%male): NR Symptomatic or Asymptomatic or Mix: Mix Days since symptom onset (if applicable):	Test name(s): Standard F EUA certified: No CE certified: Yes Platform: Fluorescent Immunoassay (FIA) Target antigen: SARS-CoV-2 nucleoprotein COI: 1 Turnaround time: NR Sample site: OP	Test name(s): cobas SARS-CoV2 (Roche), Aptima SARS-CoV2 assay (Hologic), GeneXpert SARS-CoV2 (Cepheid), RealStar SARS-CoV2-RT-PCR kit (Altona). Target gene: NR Ct threshold: NR Sample site: OP Type of swab: NR Transport media: NR Self vs HCW: HCW

		Type of swab: NR Transport media: NR HCW collection: NR Sampling sequence: 1st	
Author, year: Kernéis 2021 [44] Country: Germany Study Design: Non-randomized/ observational study with comparator (e.g. case control)	Number of patients: 1041 Age: median 36 [IQR 26–50] Gender (%male): 696 (48%) Symptomatic or Asymptomatic or Mix: Mix Days since symptom onset (if applicable): NR	Test name(s): Standard Q EUA certified: No CE certified: Yes Platform: Lateral flow immunochromatographic assay Target antigen: Nucleocapsid	Test name(s): TaqPath™ COVID 19 CE IVD RT PCR Kit Target gene: NR Ct threshold: NR Sample site: NP Type of swab: NR Transport media: NR

Supplementary Materials

		COI: NR Turnaround time: 15-30min Sample site: NP Type of swab: NR Transport media: NR HCW collection: Yes Sampling sequence: NR	Self vs HCW: HCW
Author, year: Kim 2021 [45] Country: India, Korea	Number of patients: 330 Age: NR Gender (%male): NR	Test name(s): GenBody EUA certified: Yes CE certified: Yes	Test name(s): Allplex 2019-nCOV in Korea, EURORealTime SARS-CoV-2 in India Target gene: NR Ct threshold: 30

Supplementary Materials

Study Design: Non-randomized/observational study with comparator (e.g. case control)	Symptomatic or Asymptomatic or Mix: Mix Days since symptom onset (if applicable): NR	Platform: Immunochromatographic assay Target antigen: Nucleocapsid COI: NR Turnaround time: NR Sample site: NP Type of swab: NP Transport media: VTM HCW collection: NR Sampling sequence: PCR first	Sample site: NP Type of swab: Same as index Transport media: VTM Self vs HCW: HCW

Supplementary Materials

Author, year: Kim 2022 [46]	Number of patients: 165 Age: NR Gender (%male): NR Symptomatic or Asymptomatic or Mix: Mix Days since symptom onset (if applicable): NR	Test name(s): Standard Q EUA certified: No CE certified: Yes Platform: Lateral flow immunochromatographic assay Target antigen: NR COI: NR Turnaround time: NR Sample site: NP Type of swab: NR Transport media: VTM	Test name(s): Allplex 2019-nCOV Target gene: RdRp, N genes Ct threshold: 30 Sample site: NP Type of swab: NR Transport media: VTM Self vs HCW: NR
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		HCW collection: NR Sampling sequence: PCR first	
Author, year: Klajmon 2022 [47]	Number of patients: 192 Age: mean age was 64.7 years (± 13.9). Country: Poland Study Design: Non-randomized/ observational study with comparator (e.g. case control)	Test name(s): Humasis EUA certified: No Gender (%male): NR Symptomatic or Asymptomatic or Mix: Symptomatic Days since symptom onset (if applicable): NR	Test name(s): Vitassay qPCR SARS-CoV-2 kit Target gene: ORF1ab, N Ct threshold: 35 Sample site: NP Type of swab: NR Transport media: Nucliswab standard transport media Self vs HCW: NR

Supplementary Materials

		<p>Sample site: NP</p> <p>Type of swab: Flocked swab</p> <p>Transport media: Nucliswab standard transport media</p> <p>HCW collection: NR</p> <p>Sampling sequence: NR</p>	
Author, year: Klein 2021 [48] Country: Germany Study Design: Non-randomized/ observational study with	Number of patients: 290 Age: average age of 42.7 years (standard deviation (SD) 14.6) Gender (%male): NR Symptomatic or Asymptomatic or Mix: Mix	Test name(s): Panbio EUA certified: No CE certified: Yes Platform: Lateral flow immunochromatographic assay	Test name(s): Abbott Panbio HCW collected NP swab Target gene: E and N Ct threshold: NR Sample site: NP Type of swab: IMPROSWAB

Supplementary Materials

comparator (e.g. case control)	Days since symptom onset (if applicable): NA	Target antigen: NR COI: NR Turnaround time: NR Sample site: Both AN and NP (Compared) Type of swab: Multiple Transport media: NR HCW collection: Yes - NP Sampling sequence: NR	Transport media: NR Self vs HCW: HCW
Author, year: Kolwijk 2021 [49]	Number of patients: 825 Age: >16	Test name(s): Panbio EUA certified: No	Test name(s): Abbott Panbio HCW collected NP swab Target gene: E and N

Supplementary Materials

Country: Netherlands	Gender (%male): NR Symptomatic or Asymptomatic or Mix: Symptomatic Days since symptom onset (if applicable): NA	CE certified: Yes Platform: Lateral flow immunochromatographic assay Target antigen: NR COI: NR Turnaround time: NR Sample site: Both AN and NP (Compared) Type of swab: Multiple Transport media: NR HCW collection: Yes - NP	Ct threshold: NR Sample site: NP Type of swab: IMPROSWAB Transport media: NR Self vs HCW: HCW
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Supplementary Materials

		Sampling sequence: NR	
Author, year: Koskinen 2021 [50]	Number of patients: 259 Age: NR Gender (%male): NR Symptomatic or Asymptomatic or Mix: Symptomatic Days since symptom onset (if applicable): NR	Test name(s): MariPoc EUA certified: Yes CE certified: Yes Platform: Enzyme immunoassay (EIA) Target antigen: Nucleoprotein COI: NR Turnaround time: 20-55 min Sample site: NP Type of swab: NR	Test name(s): Allplex™ 2019-nCoV RT-PCR assay Target gene: E, N and RdRP gene Ct threshold: 25 Sample site: NP Type of swab: NR Transport media: Saline, and dry Self vs HCW: NR

		Transport media: Saline, and dry HCW collection: NR Sampling sequence: PCR first	
Author, year: Krüger 2021 [51] Country: Germany Study Design: Non-randomized/ observational study with comparator (e.g. case control)	Number of patients: 1108 Age: Mean 39.4 (14.1) Gender (%male): 551 (49.3%) Symptomatic or Asymptomatic or Mix: Symptomatic Days since symptom onset (if applicable): average 4.01 (SD 3)	Test name(s): Panbio EUA certified: No CE certified: Yes Platform: Lateral flow immunochromatographic assay Target antigen: NR COI: NR	Test name(s): Alplex SARS-CoV-2 Assay, the cobas1 6800 or 8800 system Target gene: NR Ct threshold: 30 Sample site: NP, OP Type of swab: NR Transport media: NR Self vs HCW: HCW

		<p>Turnaround time: 15 minutes</p> <p>Sample site: NP</p> <p>Type of swab: NR</p> <p>Transport media: NR</p> <p>HCW collection: Yes</p> <p>Sampling sequence: PCR first</p>	
Author, year: Kumar 2021 [52]	Number of patients: 204 Country: India Study Design: Non-randomized/ observational study with	Test name(s): Standard Q EUA certified: No CE certified: Yes Platform: Lateral flow immunochromatographic assay	Test name(s): CFX96 real time PCR (Bio Rad). Target gene: NR Ct threshold: 40 Sample site: NP and throat

Supplementary Materials

comparator (e.g. case control)	Symptomatic or Asymptomatic or Mix: Asymptomatic Days since symptom onset (if applicable): NA	Target antigen: Nucleocapsid proteins COI: NR Turnaround time: 30 min Sample site: NP and throat Type of swab: NR Transport media: VTM HCW collection: Unclear Sampling sequence: NR	Type of swab: NR Transport media: VTM Self vs HCW: HCW
Author, year: Landaas 2021 [53]	Number of patients: 3991	Test name(s): Panbio	Test name(s): Aria Dx Real-Time PCR System; Cobas® SARS-CoV-2 kit on the Cobas® 6800 system

Supplementary Materials

Country: Norway Study Design: Non-randomized/ observational study with comparator (e.g. case control)	Age: NR	EUA certified: No	Target gene: NR
	Gender (%male): NR	CE certified: Yes	Ct threshold: 30
	Symptomatic or Asymptomatic or Mix: Mix	Platform: Lateral flow immunochromatographic assay	Sample site: NP/throat
	Days since symptom onset (if applicable):	Target antigen: Nucleocapsid proteins	Type of swab: NR
		COI: NR	Transport media: NR
		Turnaround time: NR	Self vs HCW: HCW
		Sample site: NP/throat	
		Type of swab: NR	
		Transport media: NR	
		HCW collection: Yes	

		Sampling sequence: NR	
Author, year: Leber 2021 [54] Country: Austria Study Design: Non-randomized/ observational study with comparator (e.g. case control)	Number of patients: 1037 Age: NR Gender (%male): NR Symptomatic or Asymptomatic or Mix: Symptomatic Days since symptom onset (if applicable): >1 day	Test name(s): Multiple EUA certified: No CE certified: Yes Platform: Lateral flow Assay Target antigen: NR COI: NR Turnaround time: NR Sample site: NP	Test name(s): Roche LightCycler http://www.roche.com ; Switzerland), BD MAXTM System using original SARS-CoV2 reagents of BD Target gene: NR Ct threshold: 40 Sample site: NR Type of swab: NP Transport media: NR Self vs HCW: NR

		Type of swab: NR Transport media: NR HCW collection: Yes Sampling sequence: NR	
Author, year: Leixner 2021 [55] Country: Austria Study Design: Non-randomized/ observational study with comparator (e.g. case control)	Number of patients: 392 Age: 70 (55-80) median (25th–75th percentiles) Gender (%male): 200 (51%) Symptomatic or Asymptomatic or Mix: Symptomatic Days since symptom onset (if applicable): 1 (0-3) median (25th–75th percentiles)	Test name(s): AMP EUA certified: No CE certified: Yes Platform: chromatographic immunoassay Target antigen: NR COI: NR	Test name(s): cobas® Liat®, GeneXpert®, Liason® MDX, BD MaxTM, cobas® z480 MagNR Pure 24 Target gene: E, ORF1a/b, E, N2, S, ORF1a/b, N1, N2, E Ct threshold: 30 Sample site: NP/OP Type of swab: NR Transport media: NR

		<p>Turnaround time: NR</p> <p>Sample site: NP</p> <p>Type of swab: NR</p> <p>Transport media: NR</p> <p>HCW collection: Yes</p> <p>Sampling sequence: same time</p>	<p>Self vs HCW: HCW</p>
Author, year: Leli 2021 [56] Country: Italy Study Design: Non-randomized/	Number of patients: 792 Age: median age was 71 years (IQR: 53–82.7) Gender (%male): 403 (50.9%)	Test name(s): LumiraDx EUA certified: No CE certified: Yes	Test name(s): NR Target gene: NR Ct threshold: 35 Sample site: NP

Supplementary Materials

observational study with comparator (e.g. case control)	Symptomatic or Asymptomatic or Mix: Mix Days since symptom onset (if applicable): NR	Platform: fluorescence immunoassay (FIA) Target antigen: Nucleocapsid protein COI: NR Turnaround time: NR Sample site: NP Type of swab: standard dry swab Transport media: Universal Transport Medium for Viruses, Chlamydia, Mycoplasma, and Ure- aplasma (Copan UTM®system; Copan, Italy).	Type of swab: NR Transport media: Universal Transport Medium for Viruses, Chlamydia, Mycoplasma, and Ure- aplasma (Copan UTM®system; Copan, Italy). Self vs HCW: NR

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Author, year: Masiá 2021 [57]	Number of patients: 913 Age: 40.6 (23–55.6) median (Q1–Q3)	Test name(s): Panbio EUA certified: No	Test name(s): LightMix Modular SARS-CoV (COVID19) E gene; TIB MOLBIOL, Berlin, Germany, distributed by Roche Target gene: E gene
Country:			Target gene: E gene
Study Design:	Gender (%male): 80 (22.2%) Symptomatic or Asymptomatic or Mix: Mix Days since symptom onset (if applicable): NR	CE certified: Yes Platform: Lateral flow immunochromatographic assay Target antigen: COI: NR Turnaround time: NR Sample site: NP Type of swab: NR Transport media: NR Self vs HCW: HCW	Ct threshold: 35 Sample site: NP Type of swab: NR Transport media: NR Self vs HCW: HCW
		 Sample site: NP Type of swab: NR Transport media: NR	

		HCW collection: Yes Sampling sequence: same time	
Author, year: Mboumba 2021 [58]	Number of patients: 100 Age: NR Gender (%male): NR Symptomatic or Asymptomatic or Mix: Symptomatic Days since symptom onset (if applicable): NR	Test name(s): Sienna-Clarity EUA certified: No CE certified: Yes Platform: Lateral Flow Assay Target antigen: NR COI: NR Turnaround time: NR	Test name(s): CE IVD-marked AllplexTM 2019-nCoV Assay (Seegene, Séoul, Korée) Target gene: (E, RdRP and N genes) Ct threshold: 33 Sample site: NP Type of swab: NR Transport media: NR Self vs HCW: NR

Supplementary Materials

		Sample site: NP Type of swab: NR Transport media: NR HCW collection: Unclear Sampling sequence: NR	
Author, year: Merino 2021 [59] Country: Spain Study Design: Non-randomized/ observational study with comparator (e.g. case control)	Number of patients: 958 Age: 40 (32) median (interquartile range) Gender (%male): 371 (38.7%) Symptomatic or Asymptomatic or Mix: Mix	Test name(s): Panbio EUA certified: No CE certified: Yes Platform: Lateral flow immunochromatographic assay Target antigen:	Test name(s): TaqMan™ 2019-nCoV assay (Applied Biosystems, Pleasanton, CA, USA), Allplex™ 2019-nCoV Assay (Seegene, Seoul, South Korea), GENOMICA S.A.U. (Madrid, Spain), SARSCOV-2 Real Time PCR KIT (Vircell, Granada, Spain), TaqPath COVID-19 Combo Kit (Thermo Fisher), Target gene: NR Ct threshold: 25

Supplementary Materials

	Days since symptom onset (if applicable): NR	COI: NR Turnaround time: NR Sample site: NP Type of swab: swab provided by PanbioRT Transport media: NR HCW collection: Yes Sampling sequence: same time	Sample site: NP Type of swab: NR Transport media: NR Self vs HCW: HCW
Author, year: Merino-Amador 2021 [60] Country: Spain	Number of patients: 450 Age: 65 (44) median (interquartile range)	Test name(s): CLINITEST EUA certified: No	Test name(s): Allplex™ 2019-nCoV Assay (Seegene, Seoul, South Korea), GENOMICA S.A.U. (Madrid, Spain), TaqPath COVID-19 Combo Kit (Thermo Fisher, Waltham, MA, USA), GeneXpert (Cepheid, Sunnyvale, CA, USA), and Cobas 6800

Supplementary Materials

Study Design: Non-randomized/observational study with comparator (e.g. case control)	Gender (%male): 187 (42%) Symptomatic or Asymptomatic or Mix: Mix Days since symptom onset (if applicable): NA	CE certified: Yes Platform: Lateral flow immunochromatographic assay Target antigen: nucleocapsid protein COI: NR Turnaround time: NR Sample site: NP Type of swab: swab provided by ClinitestRT Transport media: NR HCW collection: Yes Sampling sequence: same time	Target gene: NR Ct threshold: 25 Sample site: NP Type of swab: NR Transport media: NR Self vs HCW: HCW
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Author, year: Mitchell 2021 [61] Country: US Study Design: Non-randomized/ observational study with comparator (e.g. case control)	Number of patients: 148 symptomatic and 144 asymptomatic adults Age: mean, range 44.1 (18–83) Gender (%male): Symptomatic or Asymptomatic or Mix: Mix Days since symptom onset (if applicable): NR	Test name(s): Sofia EUA certified: CE certified: Yes Platform: Fluorescent Immunoassay (FIA) Target antigen: nucleocapsid antigen COI: NR Turnaround time: NR Sample site: Nasal Type of swab: NR	Test name(s): Cepheid Xpert Xpress SARS-CoV-2/Flu/RSV RT-PCR (Cepheid). Target gene: NR Ct threshold: NR Sample site: NP Type of swab: NR Transport media: NR Self vs HCW: HCW
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Supplementary Materials

		Transport media: NR HCW collection: Y Sampling sequence: 2nd	
Author, year: Møller 2022 [62]	Number of patients: 827 Age: NR Gender (%male): 409 (49%)	Test name(s): Multiple EUA certified: No CE certified: Yes Platform: Lateral Flow Assay Target antigen: NR COI: NR Turnaround time: NR	Test name(s): NR Target gene: NR Ct threshold: NR Sample site: OP Type of swab: NR Transport media: NR Self vs HCW: HCW

		<p>Sample site: Nasal</p> <p>Type of swab: NR</p> <p>Transport media: NR</p> <p>HCW collection: No</p> <p>Sampling sequence: PCR first</p>	
Author, year: Montalvo 2021 [63]	Number of patients: 523 Age: mean = 36.71, range 5 months to 96 years Gender (%male):	Test name(s): Elecsys EUA certified: No CE certified: Yes Platform: electrochemiluminescence immunoassay (ECLIA)	Test name(s): STAT-NAT® COVID-19 MULTI (SENTINEL Diagnostic) multiplex assay Target gene: RdRP and ORF1b genes Ct threshold: 40 Sample site: NP

Supplementary Materials

comparator (e.g. case control)	Days since symptom onset (if applicable): NR	Target antigen: NR COI: 1 Turnaround time: NR Sample site: NP Type of swab: NR Transport media: NR HCW collection: Unclear Sampling sequence: NR	Type of swab: NR Transport media: NR Self vs HCW: NR
Author, year: Mungomklang 2021 [64]	Number of patients: 1100 Age: Median 33.39 (range 13–60)	Test name(s): Standard Q EUA certified: No	Test name(s): CFX96 Touch instrument with T1000 Thermocycler (Bio-Rad, Hercules, CA) Target gene: ORF1ab

Supplementary Materials

Country: Thailand	Gender (%male): 618 (56.18%)	CE certified: Yes	Ct threshold: 40
Study Design: Non-randomized/observational study with comparator (e.g. case control)	Symptomatic or Asymptomatic or Mix: Asymptomatic Days since symptom onset (if applicable): NA	Platform: Lateral flow immunochromatographic assay Target antigen: NR COI: NR Turnaround time: NR Sample site: NP Type of swab: NR Transport media: 2 mL of viral transport media (VTM) HCW collection: Unclear	Sample site: NP Type of swab: Same as index Transport media: 2 mL of viral transport media (VTM) Self vs HCW: Same as index

		Sampling sequence: NR	
Author, year: Murillo-Zamora 2021 [65]	Number of patients: 15408 Age: mean age (\pm standard deviation) was 47.2 ± 18.5 years. Country: Study Design:	Test name(s): Multiple EUA certified: No CE certified: Yes Platform: Multiple Target antigen: Multiple COI: Multiple Turnaround time: NR Sample site: NP	Test name(s): 7500 Fast RealTime PCR System Target gene: NR Ct threshold: NR Sample site: NP Type of swab: Same as index Transport media: NR Self vs HCW: Same as index

Supplementary Materials

		Type of swab: NR Transport media: NR HCW collection: Unclear Sampling sequence: NR	
Author, year: Nikolai 2021 [66]	Number of patients: 228 Age: average 34.6 Gender (%male): NR	Test name(s): Standard Q EUA certified: No CE certified: Yes	Test name(s): NR Target gene: NR Ct threshold: NR Sample site: NR
Country: Germany	Symptomatic or Asymptomatic or Mix: Symptomatic Days since symptom onset (if applicable): Average 3.4 days (SD 3.0).	Platform: Lateral flow immunochromatographic assay Target antigen: NR COI: NR	Type of swab: NR Transport media: NR

		<p>Turnaround time: NR</p> <p>Sample site: Multiple</p> <p>Type of swab: NR</p> <p>Transport media: NR</p> <p>HCW collection: Yes</p> <p>Sampling sequence:</p>	Self vs HCW: HCW
Author, year: Nörz 2021 [67]	Number of patients: 3139 Age: NR Gender (%male): NR	Test name(s): Elecsys EUA certified: No CE certified: Yes	Test name(s): the cobasÒ 6800 system (Roche Molecular Systems, Inc., Branchburg, NJ, USA) Target gene: E gene Ct threshold: 33

Supplementary Materials

randomized/ observational study with comparator (e.g. case control)	Symptomatic or Asymptomatic or Mix: Mix Days since symptom onset (if applicable): Multiple	Platform: electrochemiluminescence immunoassay (ECLIA) Target antigen: NR COI: NR Turnaround time: as low as 18 min Sample site: NP+OP Type of swab: Flocked swab Transport media: UTM HCW collection: Unclear Sampling sequence: NR	Sample site: NP Type of swab: NR Transport media: UTM Self vs HCW: NR
	Number of patients: 1186	Test name(s): Multiple	Test name(s): Taqman 7500 (Thermo Fisher Scientific, Waltham, USA), and the

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Author, year: Osterman 2021 [68]	Age: adults and pediatric	EUA certified: No	Xpert Xpress SARS-CoV-2 run on the GeneXpert System.
Country: Germany	Gender (%male): NR	CE certified: Yes	Target gene: N, RdRp
Study Design: Non-randomized/observational study with comparator (e.g. case control)	Symptomatic or Asymptomatic or Mix: Symptomatic Days since symptom onset (if applicable): NR	Platform: Multiple Target antigen: NR COI: NR Turnaround time: Multiple Sample site: NP+OP Type of swab: eSwab™ (Copan Diagnostics, Murrieta, California, USA), ImproViral™ (Improve Medical, Guangzhou, Republic of China), dry swabs inserted into sterile 0.9% NaCl,	Ct threshold: NR Sample site: NP+OP Type of swab: same as index Transport media: VTM Self vs HCW: same as index

Supplementary Materials

		Transport media: VTM HCW collection: Yes Sampling sequence: NR	
Author, year: Paul 2021 [69] Country: India Study Design: Non-randomized/ observational study with comparator (e.g. case control)	Number of patients: 148 Age: Median 35 (IQR 12-87) Gender (%male): 99/148 Symptomatic or Asymptomatic or Mix: Mix Days since symptom onset (if applicable): NR	Test name(s): VITROS EUA certified: No CE certified: Yes Platform: chemiluminescent immunoassay (CLIA) Target antigen: NR COI: NR Turnaround time: NR	Test name(s): RT-PCR Target gene: E and S gene Ct threshold: 35 Sample site: NP+OP Type of swab: same as index Transport media: VTM Self vs HCW: same as index

		<p>Sample site: NP+OP</p> <p>Type of swab: NR</p> <p>Transport media: VTM</p> <p>HCW collection: Yes</p> <p>Sampling sequence: NR</p>	
Author, year: Peacock 2022 [70]	Number of patients: 753 Age: 47 (16.6)	Test name(s): BinaxNow EUA certified: Yes CE certified: No Platform: lateral flow immunoassay	Test name(s): Abbott RealTimeSARS-CoV-2 test Ct \geq 23 Target gene: NR Ct threshold: 23 Sample site: NP Type of swab: NR

Supplementary Materials

	Days since symptom onset (if applicable): NA	Target antigen: NR COI: NR Turnaround time: 15 minutes Sample site: MT Type of swab: NR Transport media: NR HCW collection: Yes Sampling sequence: NR	Transport media: NR Self vs HCW: NR
Author, year: Peña 2021 [71] Country: Chile	Number of patients: 842 Age: mean age: 36.67 years; SD: 16.48 years	Test name(s): SD Biosensor EUA certified: Yes	Test name(s): GenomeCov19 Detection Kit ABM Target gene: N and S gene

Supplementary Materials

Study Design: Non-randomized/observational study with comparator (e.g. case control)	Gender (%male): 429 (51.0%) Symptomatic or Asymptomatic or Mix: Asymptomatic Days since symptom onset (if applicable): NA	CE certified: Yes Platform: Lateral flow immunochromatographic assay Target antigen: COI: NR Turnaround time: NR	Ct threshold: 40 Sample site: NP Type of swab: NR Transport media: None Self vs HCW: HCW
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Supplementary Materials

		Sampling sequence: NR	
Author, year: Pérez-García 2021a [72]	Number of patients: 356 Age: NR Gender (%male): NR Symptomatic or Asymptomatic or Mix: Mix Days since symptom onset (if applicable): Multiple	Test name(s): Multiple EUA certified: No CE certified: Yes Platform: Multiple Target antigen: NR COI: NR Turnaround time: NR Sample site: NP Type of swab: NR	Test name(s): Allplex SARS-CoV-2 assay (Seegene, which detected SARS-CoV-2 E, N and RdRP genes), Viasure SARS-CoV-2RealTime PCR Detection Kit (Certest Biotech S.L.; detected genes: ORF1ab and N)and GeneFinder COVID-19 Plus RealAmp Kit (Osang Healthcare Co.; detected ge Target gene: Multiple Ct threshold: Sample site: NP Type of swab: NR Transport media: UTM Self vs HCW: NR

		Transport media: UTM HCW collection: Unclear Sampling sequence: NR	
Author, year: Pérez-García 2021b [73]	Number of patients: 320 Age: media 51 (IQR 38-68) Gender (%male): 89 (52.4%) Symptomatic or Asymptomatic or Mix: Mix Days since symptom onset (if applicable): NA	Test name(s): Multiple EUA certified: No CE certified: Yes Platform: NR Target antigen: NR COI: NR	Test name(s): three RealTime PCR platforms: Viasure SARS-CoV-2 Real Time PCR Detection Kit (Certest Biotech S.L., Zaragoza, Spain; which detected SARS-CoV-2 ORF1ab and N genes), Allplex SARS-CoV-2 assay (Seegene, Seoul, South Korea; detected genes: E, RdRP, S and N) Target gene: Assay dependant Ct threshold: NR Sample site: same as index

		Turnaround time: NR Sample site: NP Type of swab: NR Transport media: UTM HCW collection: Unclear Sampling sequence: NR	Type of swab: same as index Transport media: UTM Self vs HCW: same as index
Author, year: Petonnet 2022 [74]	Number of patients: 242 Age: median 57 (IQR 40-68)	Test name(s): Multiple EUA certified: No	Test name(s): Cobas® SARS-CoV-2 (Roche Diagnostics, Branchburg, NJ, USA), Simplexa™ COVID-19 Direct (DiaSorin, Saluggia, Italy), BioFire® SARS-CoV-2 (BioMerieux, Salt Lake City, UT, USA), and NeumoDX® SARS-CoV-2 (QIAgen, Hilden, Germany). Target gene: S, RdRp
Country: France Study Design: Non-randomized/ observational	Gender (%male): 129/242 Symptomatic or Asymptomatic or Mix: Mix	CE certified: Yes Platform: Multiple	Ct threshold: NR

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study with comparator (e.g. case control)	Days since symptom onset (if applicable): NA	Target antigen: NR COI: NR Turnaround time: NR Sample site: NP Type of swab: NR Transport media: VTM HCW collection: Unclear Sampling sequence: NR	Sample site: NP Type of swab: NR Transport media: VTM Self vs HCW: same as index
Author, year: Pilarowski 2021 [75]	Number of patients: 3302	Test name(s): BinaxNow	Test name(s): RenegadeBio using RenegadeXPTM technology.

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Country: USA	Age: 99 were aged <13 years, 110 aged 13 to 18 years, and 3093 aged >18 years.	EUA certified: Yes CE certified: No	Target gene: NR Ct threshold: NR
Study Design: Non-randomized/observational study with comparator (e.g. case control)	Gender (%male): Symptomatic or Asymptomatic or Mix: Mix Days since symptom onset (if applicable): NA	Platform: lateral flow immunoassay Target antigen: NR COI: NR Turnaround time: NR Sample site: AN Type of swab: NR Transport media: NR Self vs HCW: HCW	Sample site: AN Type of swab: NR Transport media: NR HCW collection: Yes

		Sampling sequence: Ag first	
Author, year: Pollock 2021 [76] Country: US Study Design: Non-randomized/ observational study with comparator (e.g. case control)	Number of patients: 2482 Age: Any Gender (%male): NR Symptomatic or Asymptomatic or Mix: Mix Days since symptom onset (if applicable): NA	Test name(s): BinaxNow EUA certified: Yes CE certified: No Platform: lateral flow immunoassay Target antigen: NR COI: NR Turnaround time: 15 min Sample site: AN	Test name(s): CRSP SARS-CoV-2 Real-time Reverse Transcriptase (RT)-PCR Diagnostic Assay Target gene: N2 gene Ct threshold: 40 Sample site: an Type of swab: Same as index Transport media: NR Self vs HCW: HCW

		Type of swab: Nylon swab Transport media: NR HCW collection: Yes Sampling sequence: Antigen first	
Author, year: Pray, 2021 [77] Country: US Study Design: Non-randomized/ observational study with comparator (e.g. case control)	Number of patients: 1105 Age: Range 17-64 Gender (%male): 41% Symptomatic or Asymptomatic or Mix: Symptomatic Days since symptom onset (if applicable): median 3 days (IQR 1-6), and in 72.4% ≤ 5 days	Test name(s): Sofia EUA certified: Yes CE certified: Yes Platform: Lateral Flow Assay Target antigen: Nucleocapsid COI: NR	Test name(s): CDC 2019-nCoV (University A) and Thermo Fisher TaqPath COVID-19 (University B) Target gene: N1 and N2 of NP (University A) and Thermo ORF1ab, NP and S (University B) Ct threshold: NR Sample site: MT Type of swab: NR

		<p>Turnaround time: 15 minutes</p> <p>Sample site: MT</p> <p>Type of swab: Regular tipped flocked swab</p> <p>Transport media: dry swab</p> <p>HCW collection: Yes</p> <p>Sampling sequence: PCR first</p>	<p>Transport media: VTM</p> <p>Self vs HCW: HCW</p>
<p>Author, year: Prince-Guerra 2021 [78]</p> <p>Country: US</p> <p>Study Design: Non-</p>	<p>Number of patients: 3419</p> <p>Age:</p> <p>Gender (%male):</p>	<p>Test name(s): BinaxNow</p> <p>EUA certified: Yes</p> <p>CE certified: No</p>	<p>Test name(s): CDC 2019-nCoV Real-Time RT-PCR Diagnostic Panel for detection of SARS-CoV-2 (5) (2,582 swabs) or the Fosun COVID-19 RT-PCR Detection Kit</p> <p>Target gene: NR</p> <p>Ct threshold: NR</p>

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randomized/ observational study with comparator (e.g. case control)	Symptomatic or Asymptomatic or Mix: Mix Days since symptom onset (if applicable): NA	Platform: lateral flow immunoassay Target antigen: NR COI: NR Turnaround time: <30 minutes Sample site: AN Type of swab: NR Transport media: NR HCW collection: Yes Sampling sequence: Antigen first	Sample site: NP Type of swab: NR Transport media: NR Self vs HCW: HCW
	Number of patients: 1009	Test name(s): BioSpeedia	Test name(s): ABI7500 fast thermocycler (Thermofisher)

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Author, year: Quentin 2022 [79]	Age: mean 3.7 (4.4)	EUA certified: No	Target gene: NR
Country: France	Gender (%male): 547 (54.3%)	CE certified: Yes	Ct threshold: NR
Study Design: Non-randomized/ observational study with comparator (e.g. case control)	Symptomatic or Asymptomatic or Mix: Mix Days since symptom onset (if applicable): NA	Platform: Lateral Flow Assay Target antigen: NR COI: NR Turnaround time: NR Sample site: NP Type of swab: Transport media:	Sample site: same as index Type of swab: same as index Transport media: NR Self vs HCW: HCW

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		HCW collection: Yes Sampling sequence: NR	
Author, year: Rahman 2021 [80]	Number of patients: 900 Age: median 35 SD 13.94	Test name(s): Standard Q EUA certified: No	Test name(s): CFX96 Touch™ Real-time PCR Detection System (Bio-Rad Laboratories, Inc., Hercules, CA, USA) w Target gene: RdRp, N

Country:
Bangladesh**Gender (%male):****CE certified:** Yes**Ct threshold:** 37**Study Design:****Symptomatic or Asymptomatic or Mix:** Symptomatic**Platform:** Lateral flow immunochromatographic assay**Sample site:** NP**Days since symptom onset (if applicable):** Any**Target antigen:** NR**Type of swab:** same as index**COI:** NR**Transport media:** VTM**Turnaround time:** 15-30 min**Self vs HCW:** same as index**Sample site:** NP

		Type of swab: NR Transport media: VTM HCW collection: Unclear Sampling sequence: NR	
Author, year: Regev-Yochay 2022 [81] Country: NR Study Design: Non-randomized/ observational study with comparator (e.g. case control)	Number of patients: 5142 Age: Mean 56.7, median 50.1 Gender (%male): 2026 (41.97%) Symptomatic or Asymptomatic or Mix: Mix Days since symptom onset (if applicable): NR	Test name(s): Multiple EUA certified: CE certified: Yes Platform: Target antigen: NR	Test name(s): Allplex™ 2019-nCoV, NeuMoDx™ SARS-CoV-2 assay, Xpert® Xpress SARS-CoV-2 Target gene: NR Ct threshold: NR Sample site: NR Type of swab: NR

		COI: NR Turnaround time: NR Sample site: NR Type of swab: NR Transport media: NR HCW collection: Yes Sampling sequence: NR	Transport media: NR Self vs HCW: HCW
Author, year: Schuit 2021 [82] Country: Netherlands	Number of patients: 4645 Age: >16 years Gender (%male): 2153 (50%)	Test name(s): Multiple EUA certified: No CE certified: Yes	Test name(s): the cobas® SARS-CoV-2 test on the cobas® 8800 platform, cobas 6800® platform (Roche Diagnostics International, Rotkreuz, Switzerland). Target gene: NR Ct threshold: NR

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Study Design: Non-randomized/observational study with comparator (e.g. case control)	Symptomatic or Asymptomatic or Mix: Mix Days since symptom onset (if applicable): NR	Platform: NR Target antigen: NR COI: NR Turnaround time: NR Sample site: NP+OP Type of swab: NR Transport media: VTM HCW collection: Yes Sampling sequence: PCR first	Sample site: same as index Type of swab: same as index Transport media: VTM Self vs HCW: HCW

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Author, year: Shaikh 2021 [83]	Number of patients: 199 (subgroup: pediatrics)	Test name(s): BinaxNow EUA certified: Yes CE certified: No Platform: lateral flow immunoassay Target antigen: NR COI: NR Turnaround time: NR Sample site: MT Type of swab: NR Transport media: NR	Test name(s): Roche Cobas or Hologic Panther platforms Target gene: NR Ct threshold: NR Sample site: NP Type of swab: NR Transport media: NR Self vs HCW: Yes
Country: USA	Age: 2 months - 20 years		Target gene: NR
Study Design: randomized/ observational study with comparator (e.g. case control)	Gender (%male): Symptomatic or Asymptomatic or Mix: Symptomatic Days since symptom onset (if applicable): < 7 days		

		HCW collection: Yes Sampling sequence: NR	
Author, year: Siddiqui 2021 [84] Country: US Study Design: Non-randomized/ observational study with comparator (e.g. case control)	Number of patients: 6099 Age: ASx: 31 (26–41), range / Sx: 31 (26–40) Gender (%male): 48 Symptomatic or Asymptomatic or Mix: Mix Days since symptom onset (if applicable): 3 (IQR, 4–5)	Test name(s): BinaxNow EUA certified: Yes CE certified: No Platform: lateral flow immunoassay Target antigen: nucleocapsid (N) protein COI: NR Turnaround time: 15 minutes	Test name(s): (CDC) 2019 Novel Coronavirus Real-Time RT- PCR Diagnostic Panel Target gene: viral N gene Ct threshold: 25 (24.5 for symptomatic and 27 for asymptomatic participants) Sample site: anterior nares Type of swab: NR Transport media: NR Self vs HCW: HCW

		Sample site: AN Type of swab: NR Transport media: NR HCW collection: yes Sampling sequence: 1st	
Author, year: Smith 2021 [85] Country: US Study Design: Non-randomized/ observational study with comparator (e.g. case control)	Number of patients: 43 Age: NR Gender (%male): NR Symptomatic or Asymptomatic or Mix: Mix	Test name(s): Sofia EUA certified: Yes CE certified: Yes Platform: Fluorescent Immunoassay (FIA) Target antigen: NR	Test name(s): NR Target gene: NR Ct threshold: NR Sample site: NP Type of swab: NR

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	Days since symptom onset (if applicable): NR	COI: NR Turnaround time: NA Sample site: AN Type of swab: NR Transport media: HCW collection: yes Sampling sequence: NR	Transport media: NR Self vs HCW: HCW
Author, year: Tinker 2021 [86] Country: Georgia	Number of patients: 1540 Age: NR	Test name(s): BinaxNow EUA certified: Yes	Test name(s): CDC Influenza SARS-CoV-2 (Flu SC2) Multiplex Assay Target gene: NR Ct threshold: 40

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Study Design: Non-randomized/observational study with comparator (e.g. case control)	Gender (%male): Symptomatic or Asymptomatic or Mix: Asymptomatic Days since symptom onset (if applicable): NA	CE certified: No Platform: lateral flow immunoassay Target antigen: NR COI: NR Turnaround time: Students received BinaxNOW results after 15–30 minutes Sample site: AN Type of swab: NR Transport media: NR HCW collection: no Sampling sequence: random	Sample site: Anterior nasal Type of swab: NR Transport media: NR Self vs HCW: Supervised self
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Author, year: Tonen-Wolyec 2021 [87]	Number of patients: 106 Age: median age : 40 years, range 21 - 59 years Country: France Study Design: Non-randomized/ observational study with comparator (e.g. case control)	Test name(s): BIOSYNEX BSS EUA certified: No CE certified: Yes Platform: Immunochromatographic assay Target antigen: nucleocapsid protein COI: NR Turnaround time: Overall, the mean time of antigenic self-test performance (since the opening of the box until the migration step) was 8.1 (SD: 1.3) minutes Sample site: nasal mid-turbinale Type of swab: NR	Test name(s): BIOSYNEX AmpliQuick® SARS-CoV-2 (Biosynex Swiss SA) Target gene: gene (E), and RNA-dependent RNR polymerase gene (ORF1ab of RdRP gene) Ct threshold: NR Sample site: NP Type of swab: flocked swab Transport media: NR Self vs HCW: HCW
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		Transport media: NR HCW collection: no Sampling sequence: 1st	
Author, year: Turcato 2022 [88]	Number of patients: 3899 Age: median (IQR) 69 (49–82) Gender (%male):	Test name(s): Standard Q EUA certified: No CE certified: Yes Platform: Lateral flow immunochromatographic assay Target antigen: NR COI: NR	Test name(s): XPRSARS-COV2-10 (Cepheid, CA, US). Target gene: NR Ct threshold: NR Sample site: NP Type of swab: Laboratory processed swab Transport media: Saline Self vs HCW: HCW

		Turnaround time: NA Sample site: NP Type of swab: NR Transport media: Saline HCW collection: yes Sampling sequence: 1st	
Author, year: Van der Moeren 2021(60)	Number of patients: 352 Age: NR	Test name(s): BD Veritor EUA certified: Yes	Test name(s): Cobas 6800 (Roche) platform using Cobas1 SARS-CoV-2–192 PCR assay (Roche diagnostics), Target gene: RdRp and E-genes. Ct threshold: NR
Country: Netherlands	Gender (%male): NR	CE certified: Yes	
Study Design: Non-randomized/	Symptomatic or Asymptomatic or Mix: Symptomatic	Platform: chromatographic digital immunoassay	Sample site: NP

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observational study with comparator (e.g. case control)	Days since symptom onset (if applicable): NR	Target antigen: nucleocapsid protein COI: NR Turnaround time: The manual prescribes interpretation of the results after 15 minutes with a reading device provided by the manufacturer Sample site: nasal, throat Type of swab: NR Transport media: NR HCW collection: yes Sampling sequence: 2nd	Type of swab: NR Transport media: NR Self vs HCW: HCW
	Number of patients: 7005	Test name(s): BD Veritor	

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Author, year: Venekamp 2022 [90]	Age: mean 41.1 (16.3)	EUA certified: No	Test name(s): Roche cobas 6800/8800 (Rotterdam and Breda, respectively) and ABI-7500 (Zwolle) for RT-PCR and the Hologic Panther system (Aptima SARS-CoV2 assay)
Country: Netherlands	Gender (%male):	CE certified: Yes	Target gene: NR
Study Design: Non-randomized/observational study with comparator (e.g. case control)	Symptomatic or Asymptomatic or Mix: Mix	Platform: chromatographic digital immunoassay	Ct threshold: NR
	Days since symptom onset (if applicable): NR	Target antigen: NR	Sample site: NP, OP-N
		COI: NR	Type of swab: NR
		Turnaround time: NA	Transport media: NR
		Sample site: OP-N	Self vs HCW: HCW
		Type of swab: NR	
		Transport media: NR	

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		HCW collection: yes Sampling sequence: 1st	
Author, year: Villaverde 2021 [91]	Number of patients: 1620 Age: NR Gender (%male): NR Symptomatic or Asymptomatic or Mix: Symptomatic Days since symptom onset (if applicable): <5 days	Test name(s): Panbio EUA certified: No CE certified: Yes Platform: Lateral flow immunochromatographic assay Target antigen: nucleocapsid protein COI: NR Turnaround time: NA Sample site: NP	Test name(s): NR Target gene: E and RdRp genes Ct threshold: NR Sample site: NP Type of swab: NR Transport media: NR Self vs HCW: HCW

		Type of swab: NR Transport media: NR HCW collection: yes Sampling sequence: 1st	
Author, year: von Ahnen 2021 [92] Country: Germany Study Design: Non-randomized/ observational study with comparator (e.g. case control)	Number of patients: 919 Age: mean 41.5 years (range 18–66) Gender (%male): 24	Test name(s): Roche SD EUA certified: No CE certified: Yes	Test name(s): Roche qPCR Target gene: N-Gen and E-Gen Ct threshold: 35 Sample site: NP
	Symptomatic or Asymptomatic or Mix: Asymptomatic	Platform: Lateral flow immunochromatographic assay	Type of swab: NR
	Days since symptom onset (if applicable): NA	Target antigen: nucleocapsid (N) antigen	Transport media: NR

		COI: NR Turnaround time: NA Sample site: NP Type of swab: NR Transport media: NR HCW collection: yes Sampling sequence: NR	Self vs HCW: HCW
Author, year: von Ahnen 2022 [93] Country: Germany	Number of patients: 919 Age: mean age: 41.5 years (range 18–66). Gender (%male): 24	Test name(s): Roche SD EUA certified: Yes CE certified:	Test name(s): Eruofins ViroBOAR kit Target gene: N-Gen Ct threshold: 35

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Study Design: Non-randomized/observational study with comparator (e.g. case control)	Symptomatic or Asymptomatic or Mix: Asymptomatic Days since symptom onset (if applicable): NA	Platform: Lateral flow immunochromatographic assay Target antigen: Nucleocapsid antigen COI: NR Turnaround time: NR Sample site: NP Type of swab: NR Transport media: NR HCW collection: Y Sampling sequence: NR	Sample site: NP Type of swab: NR Transport media: NR Self vs HCW: HCW

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Author, year: Wachinger 2021 [94]	Number of patients: 27421 Age: NR	Test name(s): Standard Q EUA certified: No	Test name(s): TibMolbiol (Berlin, Germany), the Allplex SARS-CoV-2 Assay from Seegene (Seoul, South Korea) or the Abbott (Illinois, USA) RealTime 2019-nCoV assay Target gene: NR
Country: Germany	Gender (%male): NR	CE certified: Yes	 Ct threshold: 33
Study Design: Non-randomized/ observational study with comparator (e.g. case control)	Symptomatic or Asymptomatic or Mix: Asymptomatic Days since symptom onset (if applicable): NA	Platform: Lateral flow immunochromatographic assay Target antigen: NR COI: NR Turnaround time: NA Sample site: NP Type of swab: NR Transport media: NR	 Sample site: NP Type of swab: NR Transport media: NR Self vs HCW: HCW

		HCW collection: yes Sampling sequence: Antigen 1st	
Author, year: Winkel 2021 [95]	Number of patients: 824 Age: median age : 27 years (range 16–80 years, IQR 21–40) Gender (%male): 94 Symptomatic or Asymptomatic or Mix: Asymptomatic Days since symptom onset (if applicable): NA	Test name(s): Panbio EUA certified: No CE certified: Yes Platform: Lateral flow immunochromatographic assay Target antigen: nucleocapsid (N) antigen COI: NR Turnaround time: Test results were recorded after 15min of assay initiation and documented by photograph.	Test name(s): Euro- fins (Brugge, Belgium; commercially available platform Viasure, CerTest Biotech, Spain, according to manufacturer's instructions, as well as laboratory-developed platform), Synlab Laboratories (Luik, Belgium; commercially available platform TaqP) Target gene: NR Ct threshold: NR Sample site: NP and OP Type of swab: NR Transport media: NR

	<p>Sample site: NP</p> <p>Type of swab: NR</p> <p>Transport media: NR</p> <p>HCW collection: yes</p> <p>Sampling sequence: 1st</p>	Self vs HCW: HCW
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References

- Albert ET, I.: Bueno, F.: Huntley, D.: Molla, E.: Fernández-Fuentes, MÁ: Martínez, M.: Poujois, S.: Forqué, L.: Valdivia, A.: Solano de la Asunción, C.: Ferrer, J.: Colomina, J.: Navarro, D. Field evaluation of a rapid antigen test (Panbio™ COVID-19 Ag Rapid Test Device) for COVID-19 diagnosis in primary healthcare centres. *Clin Microbiol Infect* **2021**; 27(3): 472.e7-e10.
- Alghounaim M, Bastaki H, Bin Essa F, Motlagh H, Al-Sabah S. The Performance of Two Rapid Antigen Tests During Population-Level Screening for SARS-CoV-2 Infection. *Front Med (Lausanne)* **2021**; 8: 797109.
- Allan-Blitz LTK, J. D. A Real-World Comparison of SARS-CoV-2 Rapid Antigen Testing versus PCR Testing in Florida. *Journal of Clinical Microbiology* **2021**; 59(10).
- Almendares O, Prince-Guerra JL, Nolen LD, et al. Performance Characteristics of the Abbott BinaxNOW SARS-CoV-2 Antigen Test in Comparison to Real-Time Reverse Transcriptase PCR and Viral Culture in Community Testing Sites during November 2020. *J Clin Microbiol* **2022**; 60(1): e0174221.
- Alqahtani M, Abdulrahman A, Mustafa F, Alawadhi AI, Alalawi B, Mallah SI. Evaluation of Rapid Antigen Tests Using Nasal Samples to Diagnose SARS-CoV-2 in Symptomatic Patients. *Front Public Health* **2021**; 9: 728969.

Supplementary Materials

6. Amer RM, Samir M, Gaber OA, et al. Diagnostic performance of rapid antigen test for COVID-19 and the effect of viral load, sampling time, subject's clinical and laboratory parameters on test accuracy. *J Infect Public Health* **2021**; 14(10): 1446-53.
7. Aoki KN, T.: Ishii, Y.: Yagi, S.: Okuma, S.: Kashiwagi, K.: Maeda, T.: Miyazaki, T.: Yoshizawa, S.: Tateda, K. Clinical validation of quantitative SARS-CoV-2 antigen assays to estimate SARS-CoV-2 viral loads in nasopharyngeal swabs. *J Infect Chemother* **2021**; 27(4): 613-6.
8. Aoki KN, T.: Ishii, Y.: Yagi, S.: Kashiwagi, K.: Miyazaki, T.: Tateda, K. Evaluation of clinical utility of novel coronavirus antigen detection reagent, Espline® SARS-CoV-2. *J Infect Chemother* **2021**; 27(2): 319-22.
9. Aranaz-Andrés JM, Chávez ACF, Laso AM, et al. Analysis of the diagnostic accuracy of rapid antigenic tests for detection of SARS-CoV-2 in hospital outbreak situation. *Eur J Clin Microbiol Infect Dis* **2022**; 41(2): 305-12.
10. Baro B, Rodo P, Ouchi D, et al. Performance characteristics of five antigen-detecting rapid diagnostic test (Ag-RDT) for SARS-CoV-2 asymptomatic infection: a head-to-head benchmark comparison. *Journal of Infection* **2021**; 82(6): 269-75.
11. Beck ET, Paar W, Fojut L, Serwe J, Jahnke RR, Miller MB. Comparison of the Quidel Sofia SARS FIA Test to the Hologic Aptima SARS-CoV-2 TMA Test for Diagnosis of COVID-19 in Symptomatic Outpatients. *Journal of Clinical Microbiology* **2021**; 59(2): e02727-20.
12. Bianco GB, M.: Barbui, A. M.: Scozzari, G.: Riccardini, F.: Coggiola, M.: Lupia, E.: Cavallo, R.: Costa, C. Evaluation of an antigen-based test for hospital point-of-care diagnosis of SARS-CoV-2 infection. *Journal of Clinical Virology* **2021**; 139.
13. Brihn AC, J.: K, O. Yong: Balter, S.: Terashita, D.: Rubin, Z.: Yeganeh, N. Diagnostic Performance of an Antigen Test with RT-PCR for the Detection of SARS-CoV-2 in a Hospital Setting - Los Angeles County, California, June-August 2020. *MMWR Morb Mortal Wkly Rep* **2021**; 70(19): 702-6.
14. Bulilete OL, P.: Leiva, A.: Carandell, E.: Oliver, A.: Rojo, E.: Pericas, P.: Llobera, J. Panbio™ rapid antigen test for SARS-CoV-2 has acceptable accuracy in symptomatic patients in primary health care. *Journal of Infection* **2021**; 82(3): 391-8.
15. Carbonell-Sahuquillo S, Lázaro-Carreño MI, Camacho J, et al. Evaluation of a rapid antigen detection test (Panbio™ COVID-19 Ag Rapid Test Device) as a point-of-care diagnostic tool for COVID-19 in a pediatric emergency department. *J Med Virol* **2021**; 93(12): 6803-7.
16. Caruana GC, A.: Kampouri, E.: Kritikos, A.: Opota, O.: Foerster, M.: Brouillet, R.: Senn, L.: Lienhard, R.: Egli, A.: Pantaleo, G.: Carron, P. N.: Greub, G. Implementing SARS-CoV-2 Rapid Antigen Testing in the Emergency Ward of a Swiss University Hospital: The INCREASE Study. *Microorganisms* **2021**; 9(4).
17. Chiu RYT, Kojima N, Mosley GL, et al. Evaluation of the INDICAID COVID-19 Rapid Antigen Test in Symptomatic Populations and Asymptomatic Community Testing. *Microbiology Spectrum* **2021**; 9(1): 1-10.
18. Courtellemont LG, J.: Guillaume, C.: Giaché, S.: Rzepecki, V.: Seve, A.: Gubavu, C.: Baud, K.: Le Helloco, C.: Cassuto, G. N.: Pialoux, G.: Hocqueloux, L.: Prazuck, T. High performance of a novel antigen detection test on nasopharyngeal specimens for diagnosing SARS-CoV-2 infection. *Journal of Medical Virology* **2021**; 93(5): 3152-7.
19. Di Domenico MDR, A.: Di Gaudio, F.: Internicola, P.: Bettini, C.: Salzano, N.: Castrianni, D.: Marotta, A.: Boccellino, M. Diagnostic accuracy of a new antigen test for sars-cov-2 detection. *International Journal of Environmental Research and Public Health* **2021**; 18(12).

Supplementary Materials

20. Dierks SB, O.: Schwanbeck, J.: Groß, U.: Weig, M. S.: Mese, K.: Lugert, R.: Bohne, W.: Hahn, A.: Feltgen, N.: Torkieh, S.: Denker, F. R.: Lauermann, P.: Storch, M. W.: Frickmann, H.: Zautner, A. E. Diagnosing sars-cov-2 with antigen testing, transcription-mediated amplification and real-time pcr. *Journal of Clinical Medicine* **2021**; 10(11).
21. Drain P, Sulaiman R, Hoppers M, Lindner NM, Lawson V, Ellis JE. Performance of the LumiraDx Microfluidic Immunofluorescence Point-of-Care SARS-CoV-2 Antigen Test in Asymptomatic Adults and Children. *Am J Clin Pathol* **2021**.
22. Drain PK, Ampajwala M, Chappel C, et al. A Rapid, High-Sensitivity SARS-CoV-2 Nucleocapsid Immunoassay to Aid Diagnosis of Acute COVID-19 at the Point of Care: A Clinical Performance Study. *Infect Dis Ther* **2021**; 10(2): 753-61.
23. Escribano P, Sánchez-Pulido AE, González-Leiva J, et al. Different performance of three point-of-care SARS-CoV-2 antigen detection devices in symptomatic patients and close asymptomatic contacts: a real-life study. *Clin Microbiol Infect* **2022**.
24. Fernandez-Montero AA, J.: Rodríguez, J. A.: Ariño, A. H.: Moreno-Galarraga, L. Validation of a rapid antigen test as a screening tool for SARS-CoV-2 infection in asymptomatic populations. Sensitivity, specificity and predictive values. *EClinicalMedicine* **2021**; 37.
25. Fernández-Rivas G, Barallat J, Gonzalez V, et al. Analytical Performance of Quantitative DiaSorin Liaison SARS-COV-2 Antigen Test for the Asymptomatic Population. *Front Public Health* **2021**; 9: 788581.
26. Ferté TR, V.: Cazanave, C.: Lafon, M. E.: Bébéar, C.: Malvy, D.: Georges-Walryck, A.: Dehail, P. Accuracy of COVID-19 rapid antigenic tests compared to RT-PCR in a student population: The StudyCov study. *J Clin Virol* **2021**; 141: 104878.
27. Fitoussi F, Tonen-Wolyec S, Awaida N, Dupont R, Bélec L. Analytical performance of the point-of-care BIOSYNEX COVID-19 Ag BSS for the detection of SARS-CoV-2 nucleocapsid protein in nasopharyngeal swabs: a prospective field evaluation during the COVID-19 third wave in France. *Infection* **2021**: 1-9.
28. Ford LW, M. J.: Shah, M. M.: Salvatore, P. P.: Segaloff, H. E.: Delaney, A.: Currie, D. W.: Boyle-Estheimer, L.: O'Hegarty, M.: Morgan, C. N.: Meece, J.: Ivacic, L.: Thornburg, N. J.: Tamin, A.: Harcourt, J. L.: Folster, J. M.: Medrzycki, M.: Jain, S.: Wong, P.: Goffard, K.: Gieryn, D.: Kahrs, J.: Langolf, K.: Zochert, T.: Tate, J. E.: Hsu, C. H.: Kirking, H. L. Antigen Test Performance Among Children and Adults at a SARS-CoV-2 Community Testing Site. *Journal of the Pediatric Infectious Diseases Society* **2021**.
29. Fourati SL, C.: Audureau, E.: Challine, D.: Michel, J.: Soulier, A.: Ahnou, N.: Désveaux, I.: Picard, O.: Ortonne, V.: Gourgeon, A.: Mills, C.: Hémery, F.: Rieux, C.: Pawlotsky, J. M.: Malou, N.: Chevaliez, S. Performance of six rapid diagnostic tests for SARS-CoV-2 antigen detection and implications for practical use. *Journal of Clinical Virology* **2021**; 142.
30. Fourati S, Soulier A, Gourgeon A, et al. Performance of a high-throughput, automated enzyme immunoassay for the detection of SARS-CoV-2 antigen, including in viral “variants of concern”: Implications for clinical use. *Journal of Clinical Virology* **2022**; 146.
31. Garcíá-Fiñana MH, D. M.: Cheyne, C. P.: Burnside, G.: Stockbridge, M.: Fowler, T. A.: Fowler, V. L.: Wilcox, M. H.: Semple, M. G.: Buchan, I. Performance of the Innova SARS-CoV-2 antigen rapid lateral flow test in the Liverpool asymptomatic testing pilot: Population based cohort study. *The BMJ* **2021**; 374.
32. Gili AP, R.: Russo, C.: Cenci, E.: Pietrella, D.: Graziani, A.: Stracci, F.: Mencacci, A. Evaluation of Lumipulse® G SARS-CoV-2 antigen assay automated test for detecting SARS-CoV-2 nucleocapsid protein (NP) in nasopharyngeal swabs for community and population screening. *International Journal of Infectious Diseases* **2021**; 105: 391-6.

Supplementary Materials

33. González-Donapetry PG-C, P.: Bloise, I.: García-Sánchez, C.: Sánchez Castellano, M. Á: Romero, M. P.: Gutiérrez Arroyo, A.: Mingorance, J.: De Ceano-Vivas La Calle, M.: García-Rodriguez, J. Think of the Children: Evaluation of SARS-CoV-2 Rapid Antigen Test in Pediatric Population. *Pediatric Infectious Disease Journal* **2021**; 385-8.
34. Hagbom M, Carmona-Vicente N, Sharma S, et al. Evaluation of SARS-CoV-2 rapid antigen diagnostic tests for saliva samples. *Heliyon* **2022**; 8(2): e08998.
35. Harris DTB, M.: Jernigan, B.: Sprissler, R.: Edwards, T.: Cohen, R.: Paul, S.: Merchant, N.: Weinkauf, C. C.: Bime, C.: Erickson, H. E.: Bixby, B.: Parthasarathy, S.: Chaudhary, S.: Natt, B.: Cristan, E.: El Aini, T.: Rischard, F.: Campion, J.: Chopra, M.: Insel, M.: Sam, A.: Knepler, J. L.: Knox, K.: Mosier, J.: Spier, C.: Dake, M. D. SARS-CoV-2 Rapid Antigen Testing of Symptomatic and Asymptomatic Individuals on the University of Arizona Campus. *Biomedicines* **2021**; 9(5).
36. Hirotsu YM, M.: Shibusawa, M.: Amemiya, K.: Nagakubo, Y.: Hosaka, K.: Sueki, H.: Hayakawa, M.: Mochizuki, H.: Tsutsui, T.: Kakizaki, Y.: Miyashita, Y.: Omata, M. Prospective study of 1308 nasopharyngeal swabs from 1033 patients using the LUMIPULSE SARS-CoV-2 antigen test: Comparison with RT-qPCR. *International Journal of Infectious Diseases* **2021**; 105: 7-14.
37. Holzner CP, D.: Anastasiou, O. E.: Dittmer, U.: Manegold, R. K.: Risse, J.: Fistera, D.: Kill, C.: Falk, M. SARS-CoV-2 rapid antigen test: Fast-safe or dangerous? An analysis in the emergency department of an university hospital. *Journal of Medical Virology* **2021**.
38. Homza MZ, H.: Janosek, J.: Tomaskova, H.: Jezo, E.: Kloudova, A.: Mrazek, J.: Svagera, Z.: Prymula, R. Five antigen tests for sars-cov-2: Virus viability matters. *Viruses* **2021**; 13(4).
39. Ifko M, Tkalčić Švabek Ž, Friščić I, et al. Diagnostic validation of two SARS-CoV-2 immunochromatographic tests in Slovenian and Croatian hospitals. *Croat Med J* **2021**; 62(5): 513-7.
40. Ishii TS, M.: Yamada, K.: Kato, D.: Osuka, H.: Aoki, K.: Morita, T.: Ishii, Y.: Tateda, K. Immunochromatography and chemiluminescent enzyme immunoassay for COVID-19 diagnosis. *Journal of Infection and Chemotherapy* **2021**; 27(6): 915-8.
41. Jakobsen KKJ, J. S.: Todsen, T.: Tolsgaard, M. G.: Kirkby, N.: Lippert, F.: Vangsted, A. M.: Martel, C. J.: Klokke, M.: von Buchwald, C. Accuracy and cost description of rapid antigen test compared with reverse transcriptase-polymerase chain reaction for SARS-CoV-2 detection. *Dan Med J* **2021**; 68(7).
42. James AE, Gulley T, Kothari A, Holder K, Garner K, Patil N. Performance of the BinaxNOW coronavirus disease 2019 (COVID-19) Antigen Card test relative to the severe acute respiratory coronavirus virus 2 (SARS-CoV-2) real-time reverse transcriptase polymerase chain reaction (rRT-PCR) assay among symptomatic and asymptomatic healthcare employees. *Infect Control Hosp Epidemiol* **2022**; 43(1): 99-101.
43. Kahn MS, L.: Bartenschlager, C.: Zellmer, S.: Frey, R.: Freitag, M.: Dhillon, C.: Heier, M.: Ebigbo, A.: Denzel, C.: Temizel, S.: Messmann, H.: Wehler, M.: Hoffmann, R.: Kling, E.: Römmele, C. Performance of antigen testing for diagnosis of COVID-19: a direct comparison of a lateral flow device to nucleic acid amplification based tests. *BMC Infectious Diseases* **2021**; 21(1).
44. Kernéis S, Elie C, Fourgeaud J, et al. Accuracy of saliva and nasopharyngeal sampling for detection of SARS-CoV-2 in community screening: a multicentric cohort study. *Eur J Clin Microbiol Infect Dis* **2021**; 40(11): 2379-88 %7 20210803 %8 Nov %! Accuracy of saliva

Supplementary Materials

- and nasopharyngeal sampling for detection of SARS-CoV-2 in community screening: a multicentric cohort study %@ 0934-9723 (Print) 0934-9723.
45. Kim DL, J.: Bal, J.: Seo, S. K.: Chong, C. K.: Lee, J. H.: Park, H. Development and Clinical Evaluation of an Immunochromatography-Based Rapid Antigen Test (GenBody™ COVAG025) for COVID-19 Diagnosis. *Viruses* **2021**; 13(5).
46. Kim HW, Park M, Lee JH. Clinical Evaluation of the Rapid STANDARD Q COVID-19 Ag Test for the Screening of Severe Acute Respiratory Syndrome Coronavirus 2. *Annals of laboratory medicine* **2022**; 42(1): 100-4.
47. Klajmon A, Olechowska-Jarząb A, Salamon D, Sroka-Oleksiak A, Brzychczy-Włoch M, Gosiewski T. Comparison of antigen tests and qpcr in rapid diagnostics of infections caused by sars-cov-2 virus. *Viruses* **2022**; 14(1).
48. Klein JAFK, L. J.: Tobian, F.: Gaeddert, M.: Lainati, F.: Schnitzler, P.: Lindner, A. K.: Nikolai, O.: Knorr, B.: Welker, A.: de Vos, M.: Sacks, J. A.: Escadafal, C.: Denkinger, C. M. Head-to-head performance comparison of self-collected nasal versus professional-collected nasopharyngeal swab for a WHO-listed SARS-CoV-2 antigen-detecting rapid diagnostic test. *Medical Microbiology and Immunology* **2021**; 210(4): 181-6.
49. Kolwijck EB-B, M.: Broertjes, J.: van Heeswijk, K.: Runderkamp, N.: Meijer, A.: Hermans, M. H. A.: Leenders, Acap. Validation and implementation of the Panbio COVID-19 Ag rapid test for the diagnosis of SARS-CoV-2 infection in symptomatic hospital healthcare workers. *Infect Prev Pract* **2021**; 3(2): 100142.
50. Koskinen JM, Antikainen P, Hotakainen K, et al. Clinical validation of automated and rapid mariPOC SARS-CoV-2 antigen test. *Sci Rep* **2021**; 11(1): 20363.
51. Krüger LJG, M.: Tobian, F.: Lainati, F.: Gottschalk, C.: Klein, J. A. F.: Schnitzler, P.: Kräusslich, H. G.: Nikolai, O.: Lindner, A. K.: Mockenhaupt, F. P.: Seybold, J.: Corman, V. M.: Drosten, C.: Pollock, N. R.: Knorr, B.: Welker, A.: de Vos, M.: Sacks, J. A.: Denkinger, C. M. The Abbott PanBio WHO emergency use listed, rapid, antigen-detecting point-of-care diagnostic test for SARS-CoV-2—Evaluation of the accuracy and ease-of-use. *PLoS ONE* **2021**; 16(5 May).
52. Kumar KKS, U. C.: Maganty, V.: Prakash, A. A.: Basumatary, J.: Adappa, K.: Chandraprabha, S.: Neeraja, T. G.: Guru Prasad, N. S.: Preethi, B.: Gangasagara, S. B.: Sujatha Rathod, B. L. Pre-Operative SARS CoV-2 Rapid Antigen Test and Reverse Transcription Polymerase Chain Reaction: A conundrum in surgical decision making. *Indian journal of ophthalmology* **2021**; 69(6): 1560-2.
53. Landaas ETS, M. L.: Tollånes, M. C.: Barlinn, R.: Kran, A. M. B.: Bragstad, K.: Christensen, A.: Andreassen, T. Diagnostic performance of a SARS-CoV-2 rapid antigen test in a large, Norwegian cohort. *Journal of Clinical Virology* **2021**; 137.
54. Leber WL, O.: Siebenhofer, A.: Redlberger-Fritz, M.: Panovska-Griffiths, J.: Czypionka, T. Comparing the diagnostic accuracy of point-of-care lateral flow antigen testing for SARS-CoV-2 with RT-PCR in primary care (REAP-2). *EClinicalMedicine* **2021**; 38: 101011.
55. Leixner GV-G, A.: Bonner, E.: Kreil, A.: Zadnikar, R.: Viveiros, A. Evaluation of the AMP SARS-CoV-2 rapid antigen test in a hospital setting. *International Journal of Infectious Diseases* **2021**; 108: 353-6.
56. Leli CDM, L.: Gotta, F.: Cornaglia, E.: Vay, D.: Megna, I.: Pensato, R. E.: Boverio, R.: Rocchetti, A. Performance of a SARS-CoV-2 antigen rapid immunoassay in patients admitted to the emergency department. *International Journal of Infectious Diseases* **2021**; 110: 135-40.

Supplementary Materials

57. Masiá MF-G, M.: Sánchez, M.: Carvajal, M.: García, J. A.: Gonzalo-Jiménez, N.: Ortiz De La Tabla, V.: Agulló, V.: Candela, I.: Guijarro, J.: Gutiérrez, J. A.: De Gregorio, C.: Gutiérrez, F. Nasopharyngeal Panbio COVID-19 Antigen Performed at Point-of-Care Has a High Sensitivity in Symptomatic and Asymptomatic Patients with Higher Risk for Transmission and Older Age. *Open Forum Infectious Diseases* **2021**; 8(3).
58. Mboumba Bouassa RSV, D.: Péré, H.: Bélec, L. Analytical performances of the point-of-care SIENNA™ COVID-19 Antigen Rapid Test for the detection of SARS-CoV-2 nucleocapsid protein in nasopharyngeal swabs: A prospective evaluation during the COVID-19 second wave in France. *International Journal of Infectious Diseases* **2021**; 106: 8-12.
59. Merino PG, J.: Muñoz-Gallego, I.: González-Donapetry, P.: Galán, J. C.: Antona, N.: Cilla, G.: Hernández-Crespo, S.: Díaz-de Tuesta, J. L.: Gual-de Torrella, A.: González-Romo, F.: Escribano, P.: Sánchez-Castellano, MÁ: Sota-Busselo, M.: Delgado-Iribarren, A.: García, J.: Cantón, R.: Muñoz, P.: Folgueira, M. D.: Cuenca-Estrella, M.: Oteo-Iglesias, J. Multicenter evaluation of the Panbio™ COVID-19 rapid antigen-detection test for the diagnosis of SARS-CoV-2 infection. *Clin Microbiol Infect* **2021**; 27(5): 758-61.
60. Merino-Amador PG-D, P.: Domínguez-Fernández, M.: González-Romo, F.: Sánchez-Castellano, MÁ: Seoane-Estevez, A.: Delgado-Iribarren, A.: García, J.: Bou, G.: Cuenca-Estrella, M.: Oteo-Iglesias, J. Clinitest rapid COVID-19 antigen test for the diagnosis of SARS-CoV-2 infection: A multicenter evaluation study. *J Clin Virol* **2021**; 143: 104961.
61. Mitchell SL, Orris S, Freeman T, et al. Performance of SARS-CoV-2 antigen testing in symptomatic and asymptomatic adults: a single-center evaluation. *BMC Infectious Diseases* **2021**; 21(1).
62. Møller IJB, Utke AR, Rysgaard UK, Østergaard LJ, Jespersen S. Diagnostic performance, user acceptability, and safety of unsupervised SARS-CoV-2 rapid antigen-detecting tests performed at home. *International Journal of Infectious Diseases* **2022**; 116: 358-64.
63. Montalvo Villalba MC, Sosa Glaria E, Rodriguez Lay LLA, et al. Performance evaluation of Elecsys SARS-CoV-2 Antigen immunoassay for diagnostic of COVID-19. *J Med Virol* **2021**.
64. Mungomklang A, Trichaisri N, Jirachewee J, Sukprasert J, Tulalamba W, Viprakasit V. Limited Sensitivity of a Rapid SARS-CoV-2 Antigen Detection Assay for Surveillance of Asymptomatic Individuals in Thailand. *Am J Trop Med Hyg* **2021**.
65. Murillo-Zamora ET, X.: Huerta, M.: Ríos-Silva, M.: Mendoza-Cano, O. Performance of Antigen-Based Testing as Frontline Diagnosis of Symptomatic COVID-19. *Medicina (Kaunas, Lithuania)* **2021**; 57(8).
66. Nikolai O, Rohhardt C, Tobian F, et al. Anterior nasal versus nasal mid-turbinate sampling for a SARS-CoV-2 antigen-detecting rapid test: does localisation or professional collection matter? *Infect Dis (Lond)* **2021**; 53(12): 947-52 %7 20210827 %8 Nov-Dec %! Anterior nasal versus nasal mid-turbinate sampling for a SARS-CoV-2 antigen-detecting rapid test: does localisation or professional collection matter? %@ 2374-4235 (Print) 2374-4243.
67. Nörz D, Olearo F, Perisic S, et al. Multicenter Evaluation of a Fully Automated High-Throughput SARS-CoV-2 Antigen Immunoassay. *Infectious Diseases and Therapy* **2021**; 10(4): 2371-9 %9 Article %! Multicenter Evaluation of a Fully Automated High-Throughput SARS-CoV-2 Antigen Immunoassay %@ 193-6382 2193-8229.
68. Osterman A, Igihaut M, Lehner A, et al. Comparison of four commercial, automated antigen tests to detect SARS-CoV-2 variants of concern. *Med Microbiol Immunol* **2021**; 210(5-6): 263-75.

Supplementary Materials

69. Paul D, Gupta A, Rooge S, Gupta E. Performance evaluation of automated chemiluminescence immunoassay based antigen detection - Moving towards more reliable ways to predict SARS-CoV-2 infection. *J Virol Methods* **2021**; 298: 114299.
70. Peacock WF, Soto-Ruiz KM, House SL, et al. Utility of COVID-19 antigen testing in the emergency department. *J Am Coll Emerg Physicians Open* **2022**; 3(1): e12605.
71. Peña MA, M.: Garcés, C.: Gaggero, A.: García, P.: Velasquez, M. S.: Luza, R.: Alvarez, P.: Paredes, F.: Acevedo, J.: Farfán, M. J.: Solari, S.: Soto-Rifo, R.: Valiente-Echeverría, F. Performance of SARS-CoV-2 rapid antigen test compared with real-time RT-PCR in asymptomatic individuals. *International Journal of Infectious Diseases* **2021**; 107: 201-4.
72. Pérez-García FR, J.: Moya Gutiérrez, H.: Labrador Ballesteros, A.: Pérez Ranz, I.: González Arroyo, J.: González Ventosa, V.: Pérez-Tanoira, R.: Domingo Cruz, C.: Cuadros-González, J. Comparative evaluation of Panbio and SD Biosensor antigen rapid diagnostic tests for COVID-19 diagnosis. *Journal of Medical Virology* **2021**; 93(9): 5650-4.
73. Pérez-García FR, J.: Gómez-Herruz, P.: Arroyo, T.: Pérez-Tanoira, R.: Linares, M.: Pérez Ranz, I.: Labrador Ballesteros, A.: Moya Gutiérrez, H.: Ruiz-Álvarez, M. J.: Cuadros-González, J. Diagnostic performance of CerTest and Panbio antigen rapid diagnostic tests to diagnose SARS-CoV-2 infection. *Journal of Clinical Virology* **2021**; 137.
74. Petonnet D, Marot S, Leroy I, et al. Comparison of Rapid and Automated Antigen Detection Tests for the Diagnosis of SARS-CoV-2 Infection. *Diagnostics (Basel)* **2022**; 12(1).
75. Pilarowski G, Marquez C, Rubio L, et al. Field Performance and Public Health Response Using the BinaxNOW™ Rapid Severe Acute Respiratory Syndrome Coronavirus 2 (SARS-CoV-2) Antigen Detection Assay During Community-Based Testing. *Clin Infect Dis* **2021**; 73(9): e3098-e101 %8 Nov 2 %! Field Performance and Public Health Response Using the BinaxNOW™ Rapid Severe Acute Respiratory Syndrome Coronavirus 2 (SARS-CoV-2) Antigen Detection Assay During Community-Based Testing %@ 1058-4838 (Print) 1058-4838.
76. Pollock NRJ, J. R.: Tran, K.: Cranston, A. E.: Smith, S.: O’Kane, C. Y.: Roady, T. J.: Moran, A.: Scarry, A.: Carroll, M.: Volinsky, L.: Perez, G.: Patel, P.: Gabriel, S.: Lennon, N. J.: Madoff, L. C.: Brown, C.: Smole, S. C. Performance and implementation evaluation of the Abbott BinaxNOW rapid antigen test in a high-throughput drive-through community testing site in Massachusetts. *Journal of Clinical Microbiology* **2021**; 59(5).
77. Pray IW, Ford L, Cole D, et al. Performance of an Antigen-Based Test for Asymptomatic and Symptomatic SARS-CoV-2 Testing at Two University Campuses - Wisconsin, September–October 2020. *MMWR Morb Mortal Wkly Rep* **2021**; 69(5152): 1642-7.
78. Prince-Guerra JL AO, Nolen LD, et al. Evaluation of Abbott BinaxNOW Rapid Antigen Test for SARS-CoV-2 Infection at Two Community-Based Testing Sites — Pima County, Arizona, November 3–17, 2020. *MMWR Morb Mortal Wkly Rep* **2021**; 70:100–105.
79. Quentin O, Sylvie P, Olivier M, et al. Prospective evaluation of the point-of-care use of a rapid antigenic SARS-CoV-2 immunochromatographic test in a pediatric emergency department. *Clin Microbiol Infect* **2022**.
80. Rahman MM, Hoque AF, Karim Y, et al. Clinical evaluation of SARS-CoV-2 antigen-based rapid diagnostic test kit for detection of COVID-19 cases in Bangladesh. *Heliyon* **2021**; 7(11): e08455 %7 20211122 %8 Nov %! Clinical evaluation of SARS-CoV-2 antigen-based rapid diagnostic test kit for detection of COVID-19 cases in Bangladesh %@ 2405-8440 (Print) 2405-8440.

Supplementary Materials

81. Regev-Yochay G, Kriger O, Mina MJ, et al. Real World Performance of SARS-CoV-2 Antigen Rapid Diagnostic Tests in Various Clinical Settings. *Infect Control Hosp Epidemiol* **2022**; 1-20.
82. Schuit E, Venekamp RP, Veldhuijzen IK, et al. Accuracy and usability of saliva and nasal rapid antigen self-testing for detection of SARS-CoV-2 infection in the general population: a head-to-head comparison. **2021**.
83. Shaikh NF, E. J.: Tate, P. J.: Liu, H.: Chang, C. H.: Wells, A.: Hoberman, A. Performance of a Rapid SARS-CoV-2 Antigen Detection Assay in Symptomatic Children. *Pediatrics* **2021**; 148(3).
84. Siddiqui ZK, Chaudhary M, Robinson ML, et al. Implementation and Accuracy of BinaxNOW Rapid Antigen COVID-19 Test in Asymptomatic and Symptomatic Populations in a High-Volume Self-Referred Testing Site. *Microbiol Spectr* **2021**; 9(3): e0100821.
85. Smith RDJ, J. K.: Clay, C.: Girio-Herrera, L.: Stevens, D.: Abraham, M.: Zimand, P.: Ahlman, M.: Gimigliano, S.: Zhao, R.: Hildenbrand, C.: Barrueto, F.: Leekha, S. Clinical evaluation of Sofia Rapid Antigen Assay for detection of severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) among emergency department to hospital admissions. *Infect Control Hosp Epidemiol* **2021**; 1-6.
86. Tinker SCS, C. M.: Litvintseva, A. P.: Drenzek, C.: Voccio, G. E.: Hunter, M. A.: Briggs, S.: Heida, D. E.: Folster, J.: Shewmaker, P. L.: Medrzycki, M.: Bowen, M. D.: Bohannon, C.: Bagarozzi, D.: Petway, M.: Rota, P. A.: Kuhnert-Tallman, W.: Thornburg, N.: Prince-Guerra, J. L.: Barrios, L. C.: Tamin, A.: Harcourt, J. L.: Honein, M. A. Point-of-care antigen test for sars-cov-2 in asymptomatic college students. *Emerging Infectious Diseases* **2021**; 27(10): 2662-5.
87. Tonen-Wolyec S, Dupont R, Awaida N, Batina-Agasa S, Hayette MP, Bélec L. Evaluation of the practicability of biosynex antigen self-test covid-19 ag+ for the detection of sars-cov-2 nucleocapsid protein from self-collected nasal mid-turbinate secretions in the general public in france. *Diagnostics* **2021**; 11(12).
88. Turcato G, Zaboli A, Pfeifer N, et al. Rapid antigen test to identify COVID-19 infected patients with and without symptoms admitted to the Emergency Department. *Am J Emerg Med* **2022**; 51: 92-7.
89. Van der Moeren NZ, V. F.: Lodder, E. B.: Van den Billiaardt, W.: Van Esch, Hrjm: Stohr, Jjjm: Pot, J.: Welschen, I.: Van Mechelen, P. M. F.: Pas, S. D.: Kluytmans, Jajw. Evaluation of the test accuracy of a SARS-CoV-2 rapid antigen test in symptomatic community dwelling individuals in the Netherlands. *PLoS One* **2021**; 16(5): e0250886.
90. Venekamp RP, Veldhuijzen IK, Moons KGM, et al. Detection of SARS-CoV-2 infection in the general population by three prevailing rapid antigen tests: cross-sectional diagnostic accuracy study. *BMC Med* **2022**; 20(1): 97.
91. Villaverde SD-R, S.: Sabrido, G.: Pérez-Jorge, C.: Plata, M.: Romero, M. P.: Grasa, C. D.: Jiménez, A. B.: Heras, E.: Broncano, A.: Núñez, M. D. M.: Illán, M.: Merino, P.: Soto, B.: Molina-Arana, D.: Bermejo, A.: Mendoza, P.: Gijón, M.: Pérez-Moneo, B.: Moraleda, C.: Tagarro, A.: Calvo, C.: Mellado, M. J.: Rodríguez-Molino, P.: del Rosal, T.: Santos, M.: Navarro, M.: Rincón, E.: Santiago, B.: Saavedra-Lozano, J.: Aguilera-Alonso, D.: Epalza, C.: Blázquez-Gamero, D.: Villanueva, S.: Rojo, P.: Calleja, G.: Alonso, J. A.: de la Torre, M.: Sanz-Santaeufemia, F. J.: Iglesias, M. I.: Herrero, B.: Alonso, M.: Soriano-Arandes, T.: Pujol, J. M.: Melendo, S.: Soler-Palacin, P.: Simó, S.: Fumadó, V.: Lanaspa, M.: Urretavizcaya, M.: Herranz, M.: Pareja, M.: Ara, F.: Cabañas, S.: del Valle, R.: Barrios, A.: Otheo, E.: Vázquez, J. L.: Falcón, L.: Neth, O.: Olbrich, P.: Goicoechea, W.: Martín, L.: Figueroa, L.: Llorente, M.: Penin, M.: García, C.: García, M.: Alvaredo, T.: Olmedo, M. I.: López, A.: Cobo, E.: Tovizi, M.: Galán, P.: Guillén, S.: Navas, A.: García, M. L.: Pérez, S.: Hernández, M. J.: Berzosa, A.: Gallego, N.: López, A.: Ruiz, B.:

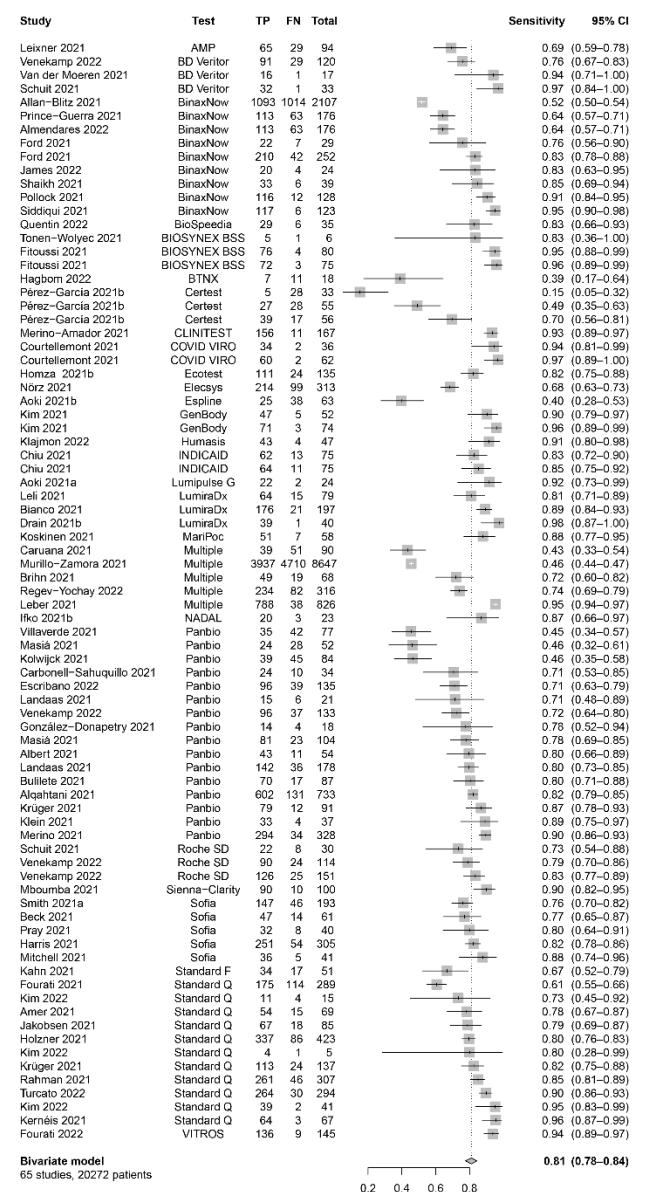
Supplementary Materials

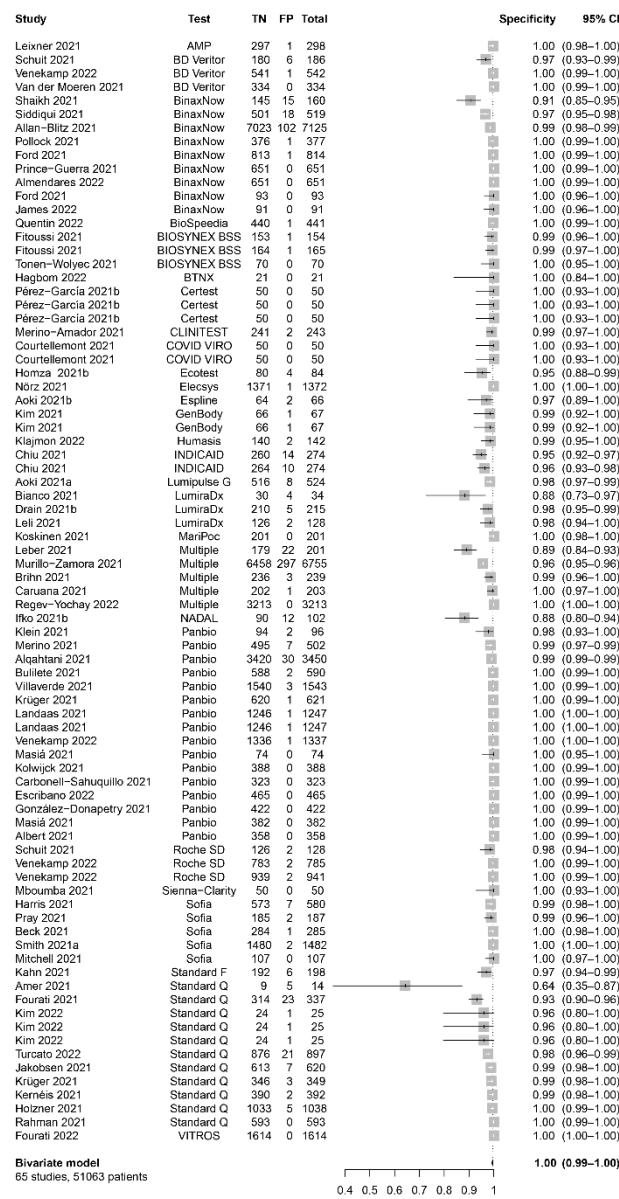
- Alfayate, S.: Menasalvas, A.: Cervantes, E.: Méndez, M.: Hurtado, Á: Ruiz, Y.: García, C.: Amich, I.: Oltra, M.: Villaroya, Á: Ocaña, A.: Romero, I.: Guzmán, M. F.: Pascual, M. J.: Sánchez-Códez, M.: Montesinos, E.: Jensen, J.: Rodríguez, M.: Caro, G.: Rius, N.: Gómez, A.: Bretón, R.: Rodríguez, M.: Romero, J.: Campos, A.: García, M.: Velasco, R. M.: Lobato, Z.: Centeno, F.: Pérez, E.: Vidal, P.: Rey, C.: Vivanco, A.: Alonso, M.: Alcalá, P.: de Dios, J. G.: Solé, E.: Minguell, L.: Astigarraga, I.: Vázquez, M. Á: Sánchez, M.: Díaz, E.: Consuegra, E.: Cabanillas, M.: Peña, L.: Garrote, E.: Goicoechea, M.: Centelles, I.: Lapeña, S.: Gutiérrez, S.: Cavalle, A.: Olmos, J. M.: Cobo, A.: Díaz, S.: Jiménez, B.: González, R.: Lafuente, M.: Bustillo, M.: Pons, N.: Morata, J.: Segura, E. Diagnostic Accuracy of the Panbio Severe Acute Respiratory Syndrome Coronavirus 2 Antigen Rapid Test Compared with Reverse-Transcriptase Polymerase Chain Reaction Testing of Nasopharyngeal Samples in the Pediatric Population. *Journal of Pediatrics* **2021**; 232: 287-9.e4.
92. von Ahnen TvA, M.: Wirth, U.: Schardey, H. M.: Herdtle, S. Evaluation of a rapid-antigen test for COVID-19 in an asymptomatic collective : A prospective study. *Wien Med Wochenschr* **2021**: 1-4.
93. von Ahnen T, von Ahnen M, Wirth U, Schardey HM, Herdtle S. Evaluation of a rapid-antigen test for COVID-19 in an asymptomatic collective : A prospective study. *Wien Med Wochenschr* **2022**; 172(3-4): 70-3.
94. Wachinger JO, I. D.: Horner, S.: Schnitzler, P.: Heeg, K.: Denkinger, C. M. The potential of SARS-CoV-2 antigen-detection tests in the screening of asymptomatic persons. *Clinical Microbiology and Infection* **2021**.
95. Winkel B, Schram E, Gremmels H, et al. Screening for SARS-CoV-2 infection in asymptomatic individuals using the Panbio COVID-19 antigen rapid test (Abbott) compared with RT-PCR: a prospective cohort study. *BMJ Open* **2021**; 11(10): e048206.

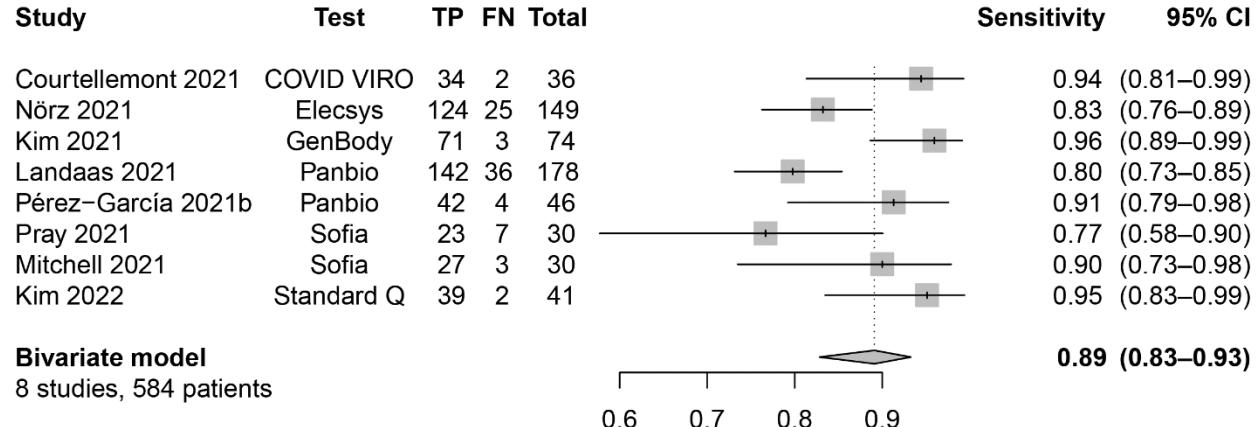
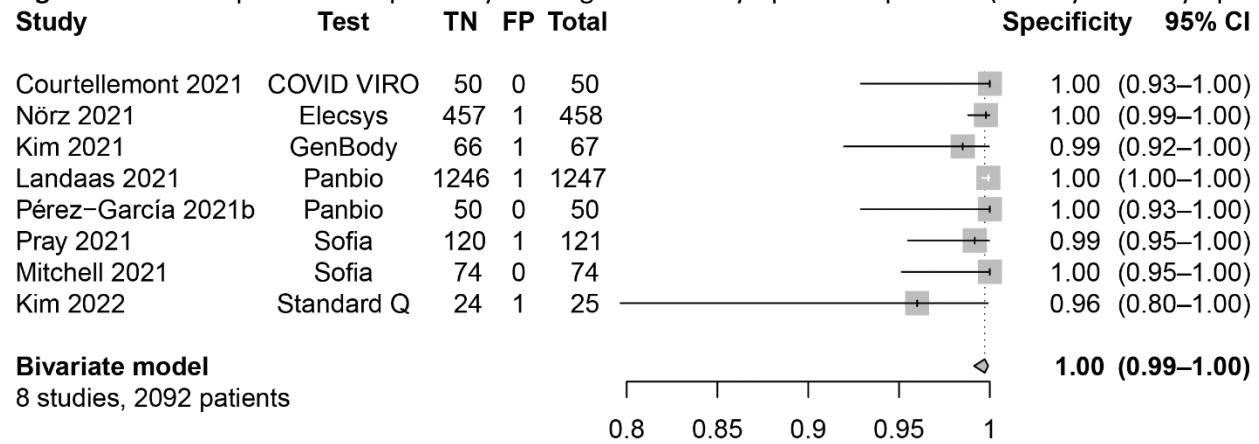
Supplement B

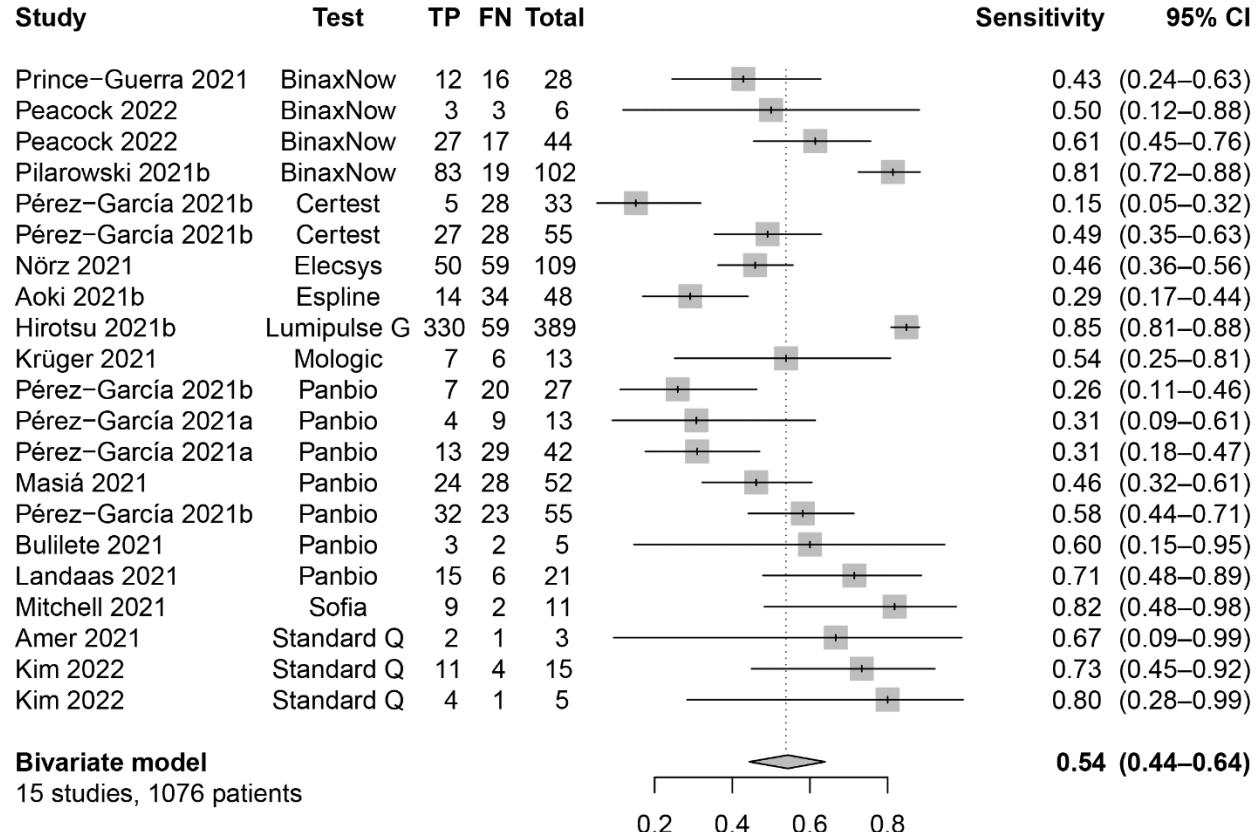
Recommendation 1: For symptomatic individuals suspected of having COVID-19, the IDSA panel recommends a single Ag test over no test.
(strong recommendation, moderate certainty evidence)

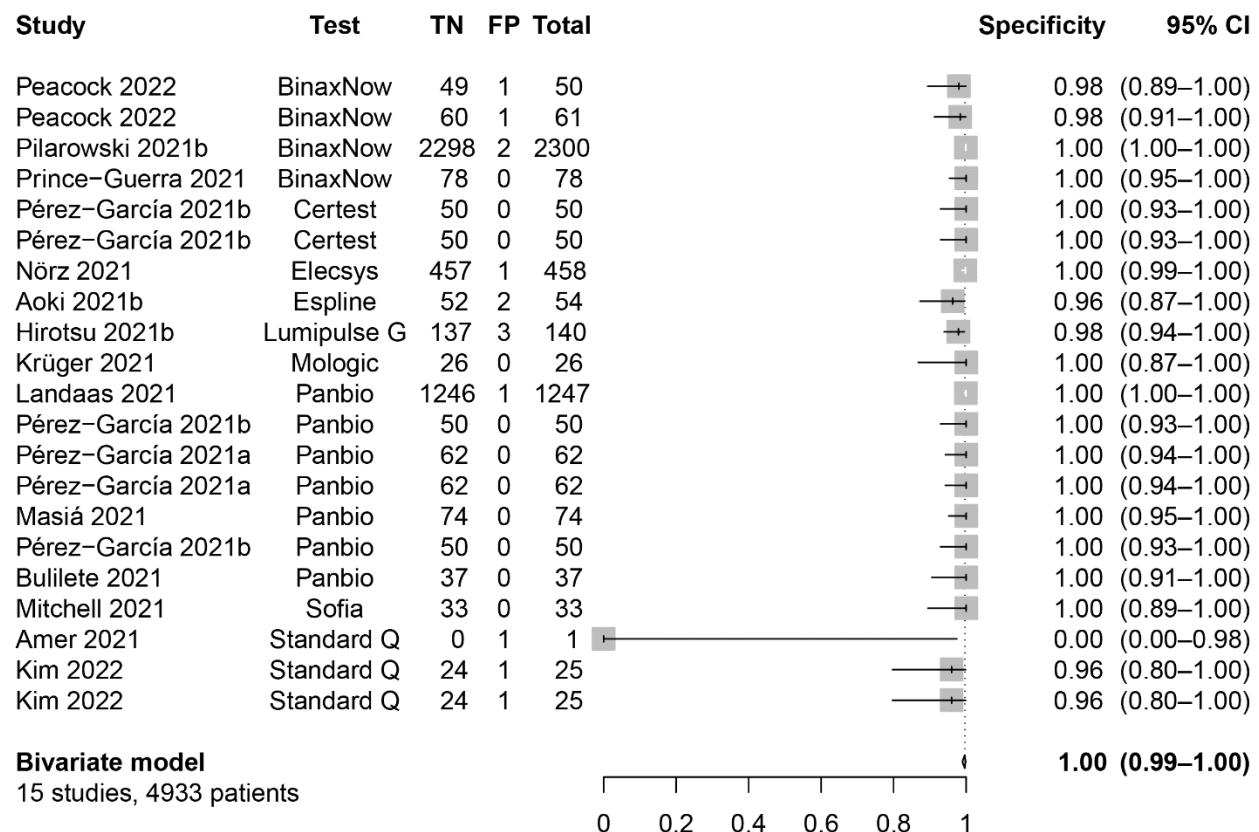
Supplementary Materials

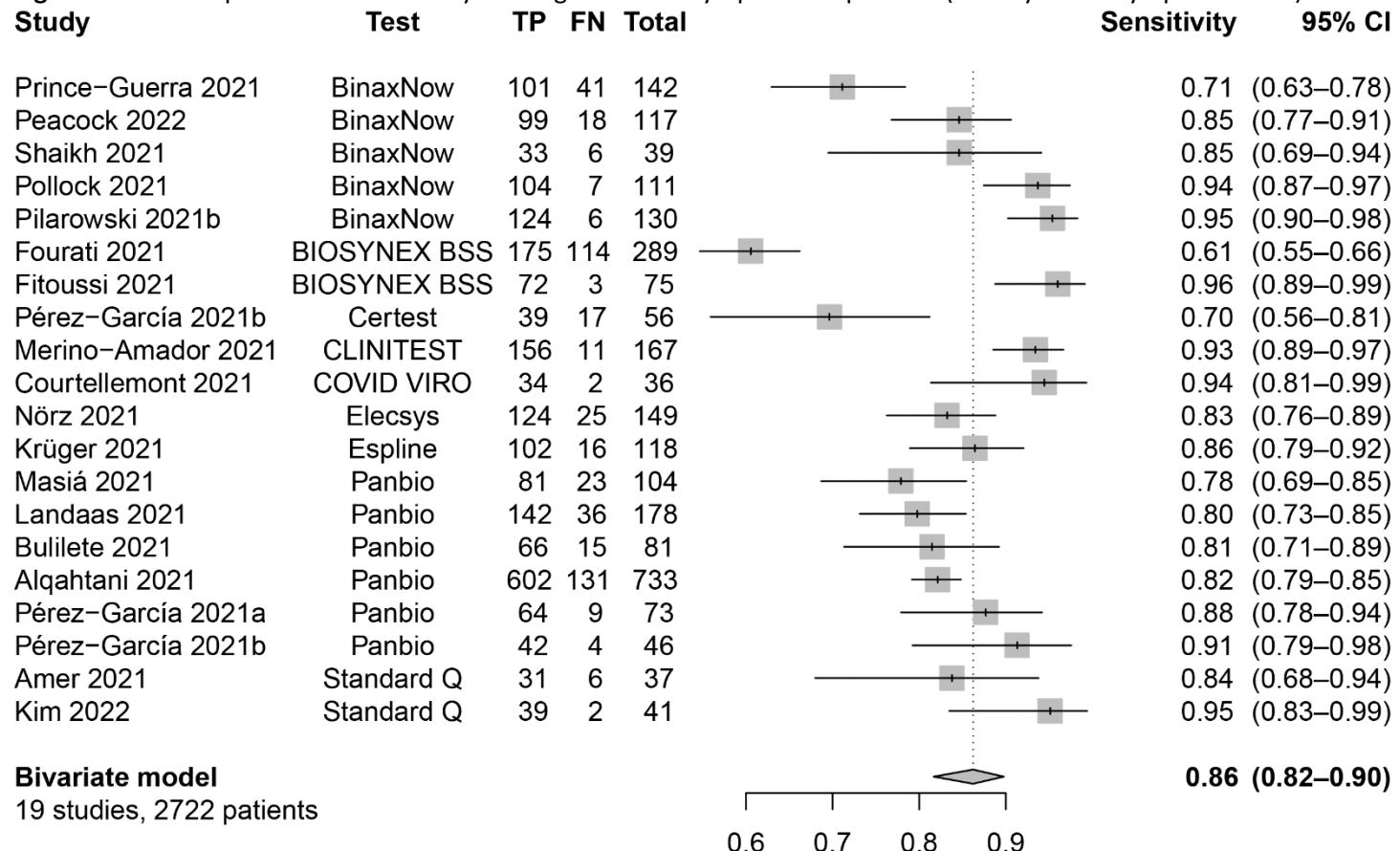
Figure s2a. Forest plot for the overall sensitivity of antigen tests in symptomatic patients

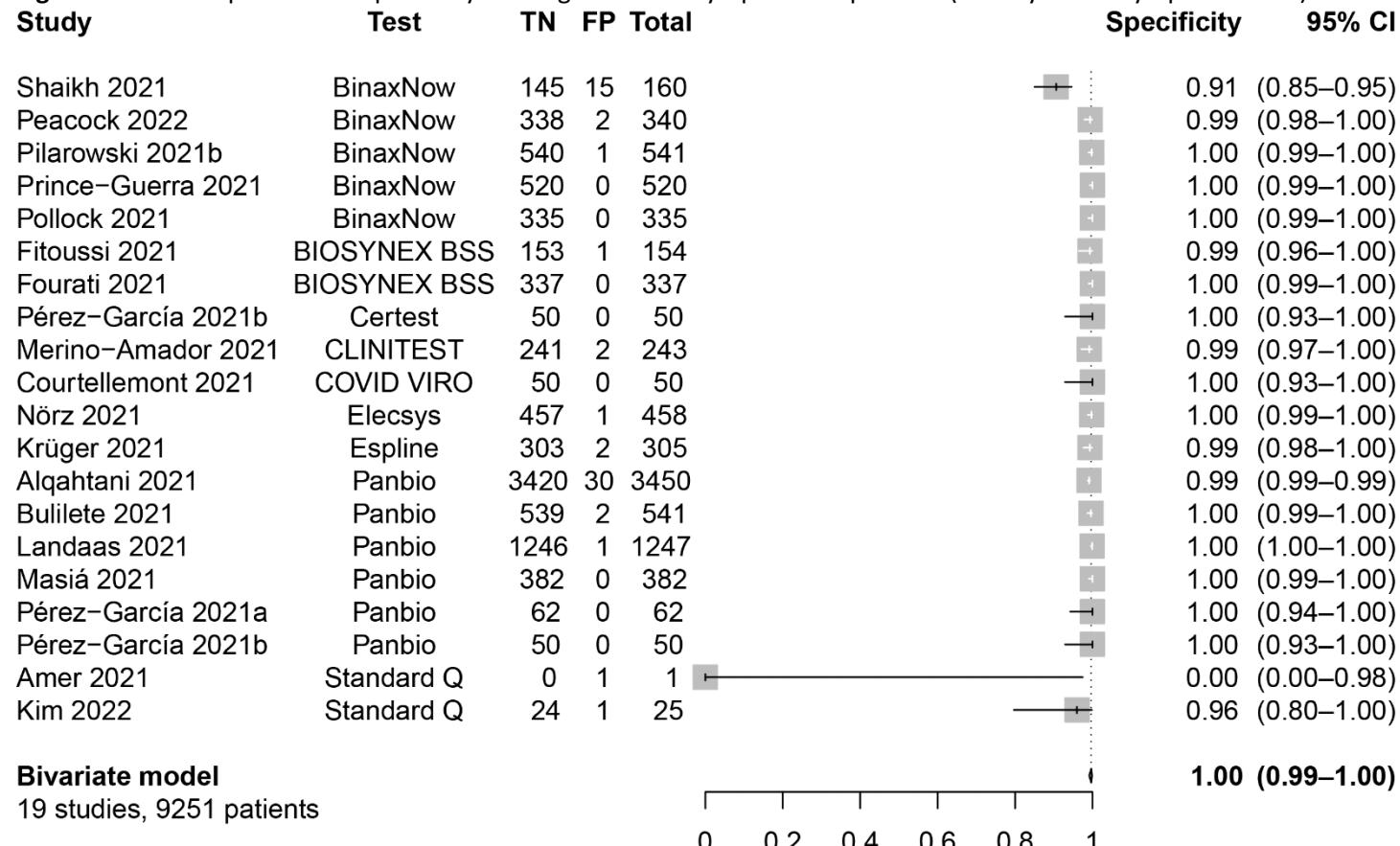
*Supplementary Materials***Figure s2b.** Forest plot for the overall specificity of antigen tests in symptomatic patients

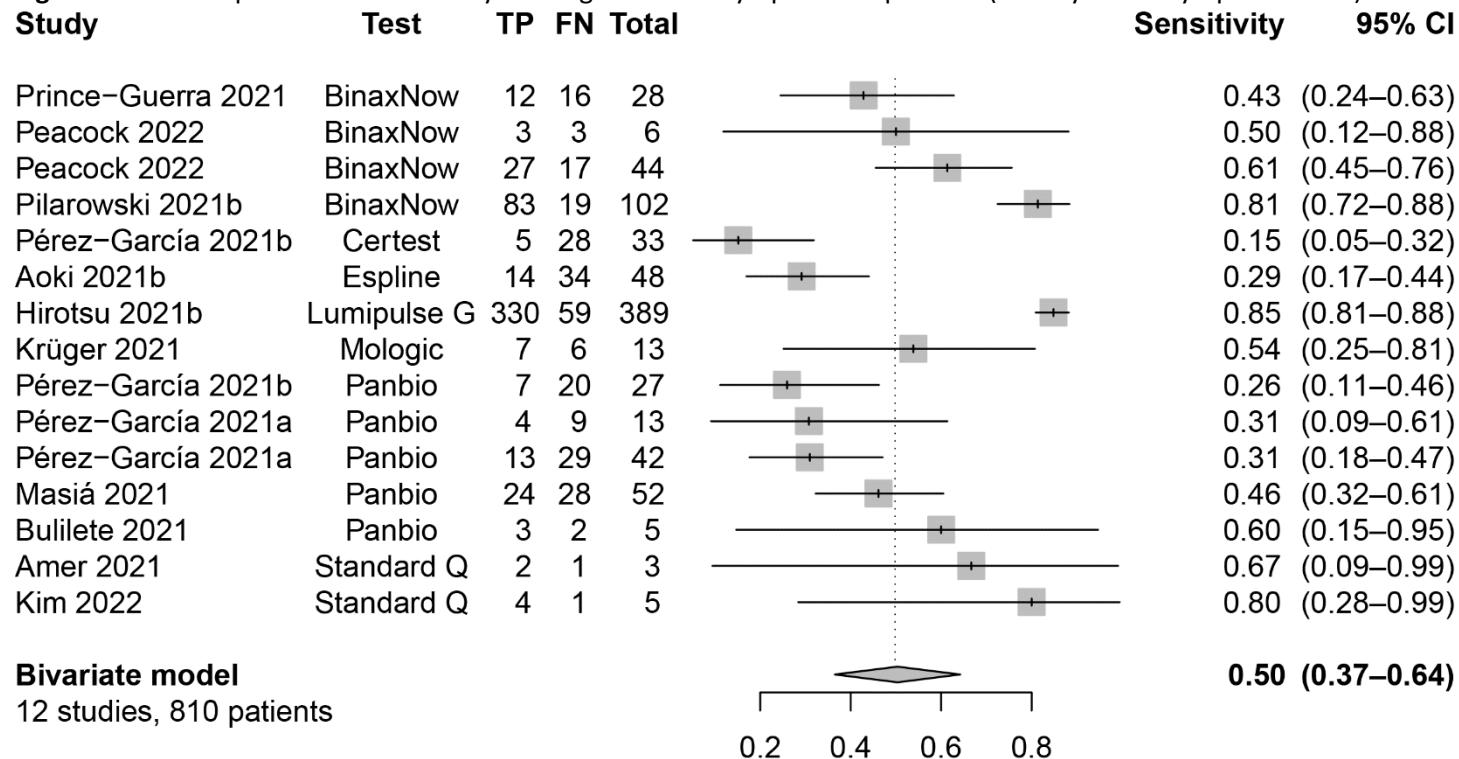
*Supplementary Materials***Figure s3a.** Forest plot for the sensitivity of antigen tests in symptomatic patients (≤ 5 days since symptom onset)**Figure s3b.** Forest plot for the specificity of antigen tests in symptomatic patients (≤ 5 days since symptom onset)

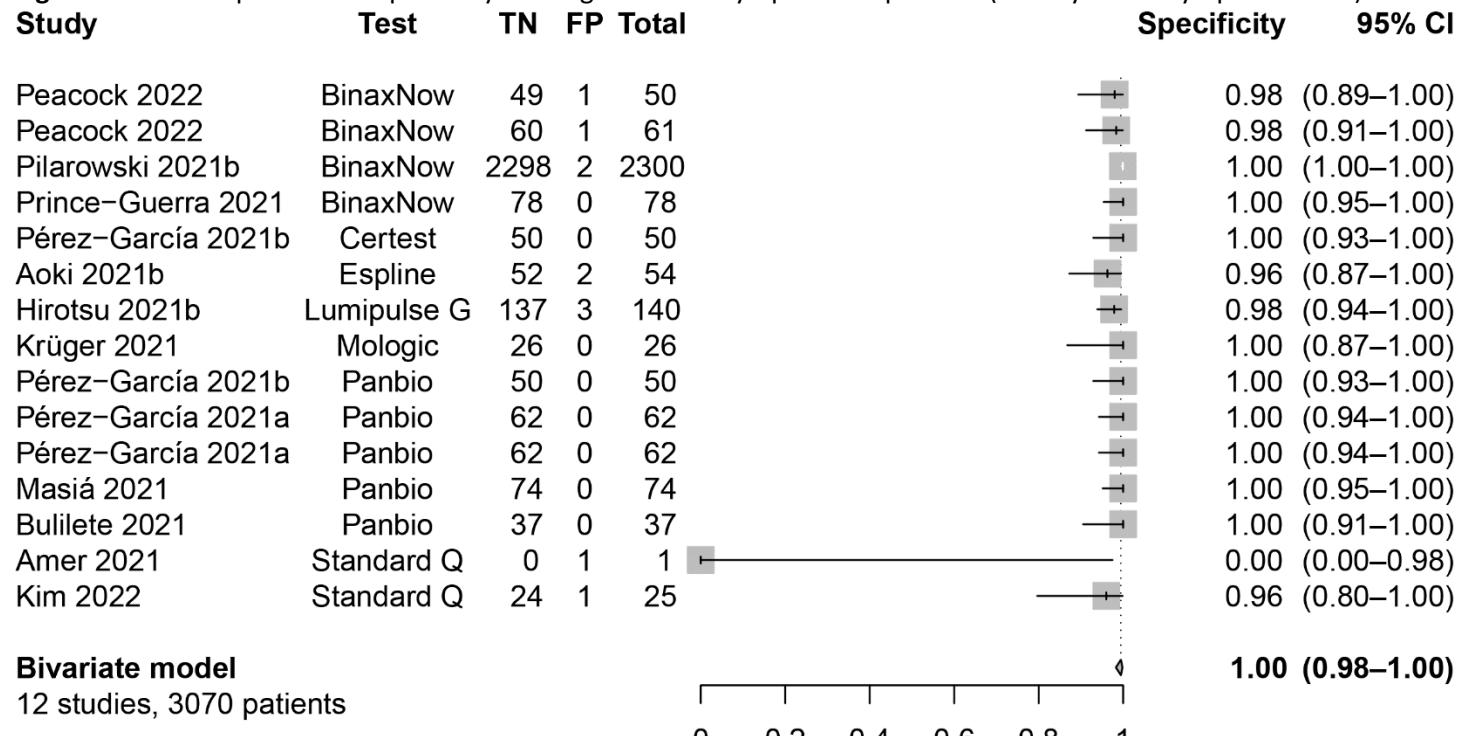
*Supplementary Materials***Figure s4a.** Forest plot for the sensitivity of antigen tests in symptomatic patients (>5 days since symptom onset)

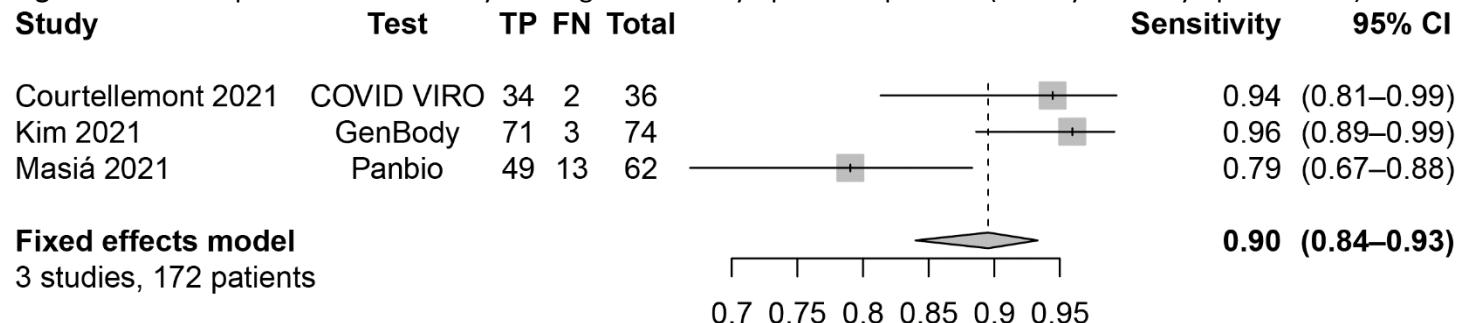
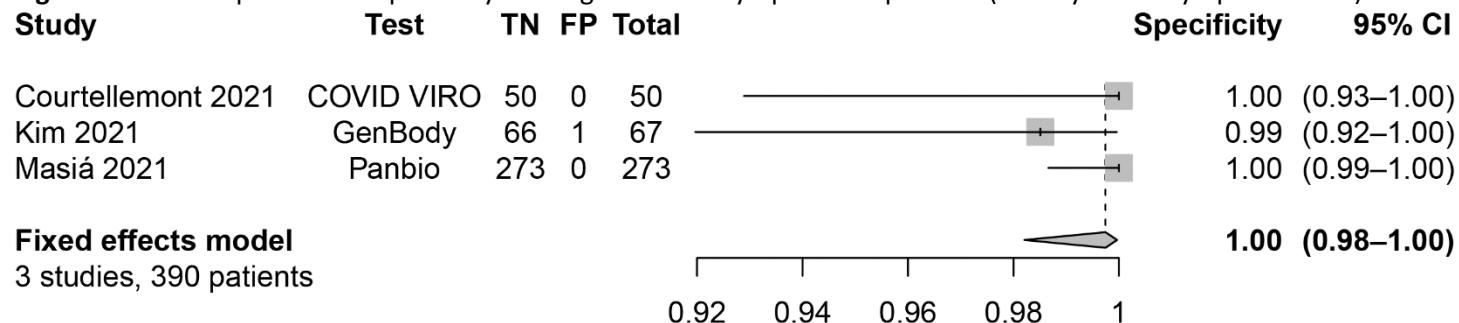
*Supplementary Materials***Figure s4b.** Forest plot for the specificity of antigen tests in symptomatic patients (>5 days since symptom onset)

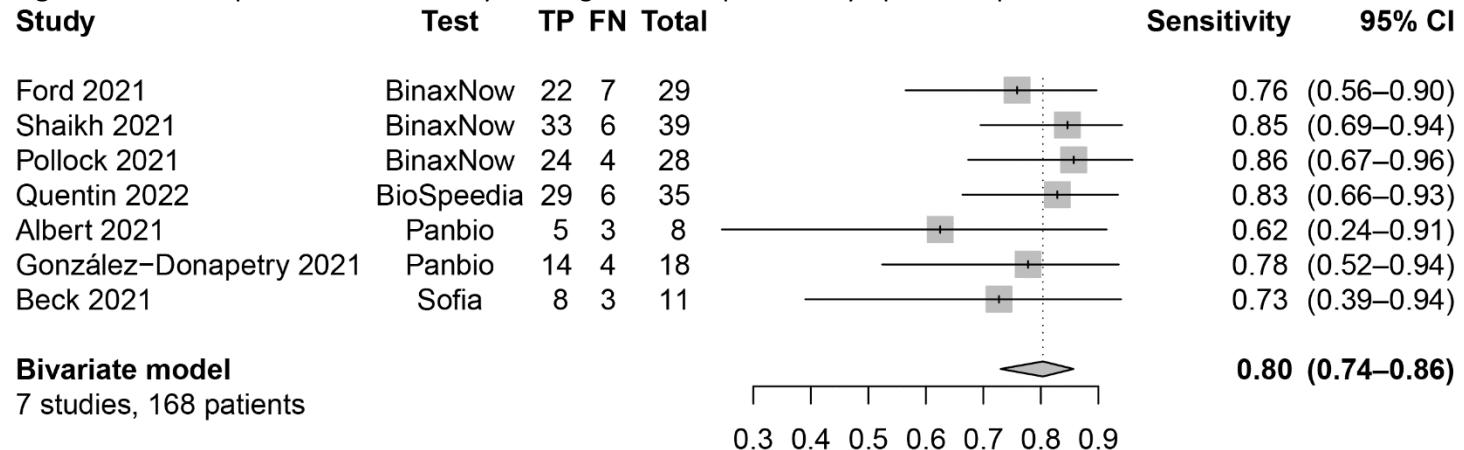
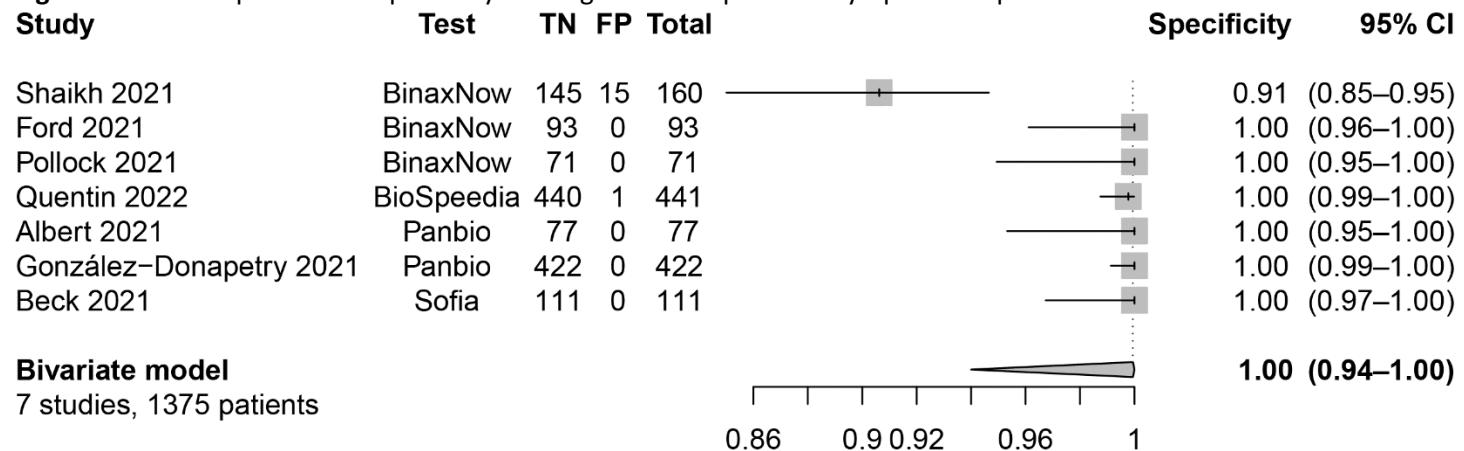
*Supplementary Materials***Figure s5a.** Forest plot for the sensitivity of antigen tests in symptomatic patients (≤ 7 days since symptom onset)

*Supplementary Materials***Figure s5b.** Forest plot for the specificity of antigen tests in symptomatic patients (≤ 7 days since symptom onset)

*Supplementary Materials***Figure s6a.** Forest plot for the sensitivity of antigen tests in symptomatic patients (> 7 days since symptom onset)

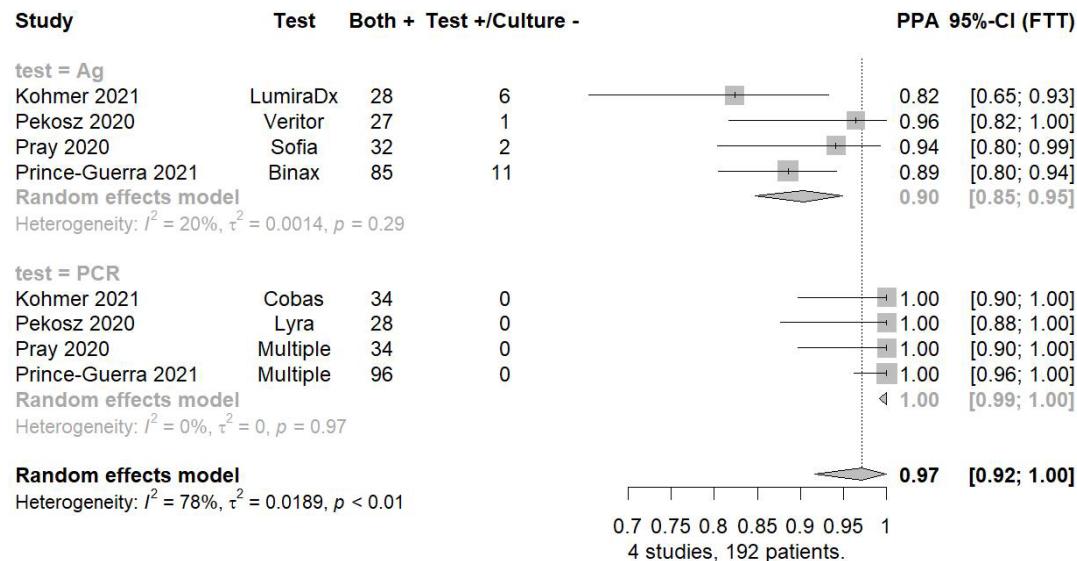
*Supplementary Materials***Figure s6b.** Forest plot for the specificity of antigen tests in symptomatic patients (> 7 days since symptom onset)

*Supplementary Materials***Figure s7a.** Forest plot for the sensitivity of antigen tests in symptomatic patients (≤ 3 days since symptom onset)**Figure s7b.** Forest plot for the specificity of antigen tests in symptomatic patients (≤ 3 days since symptom onset)

*Supplementary Materials***Figure s8a.** Forest plot for the sensitivity of antigen tests in pediatric symptomatic patients**Figure s8b.** Forest plot for the specificity of antigen tests in pediatric symptomatic patients

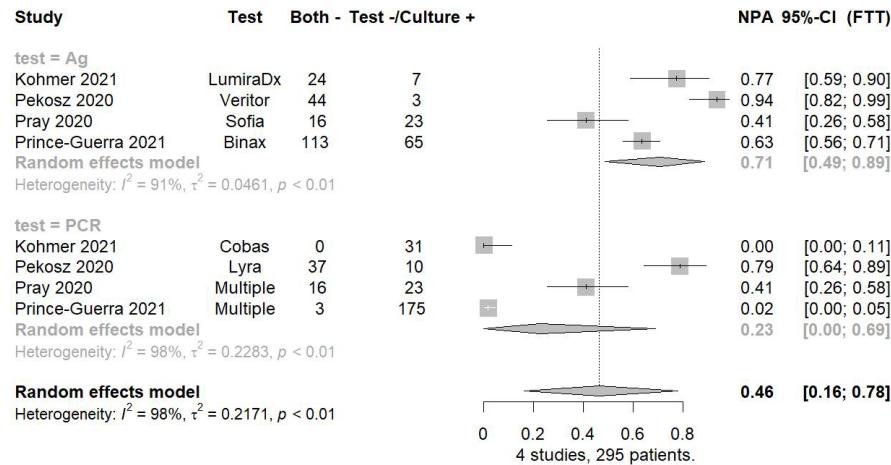
Supplementary Materials

Figure s9a. Forest plot for the agreement on positive results of antigen tests and standard NAAT (either rapid RT-PCR or laboratory-based NAAT) vs. viral culture (all patients)



Supplementary Materials

Figure s9b. Forest plot for the agreement on the negative results of rapid antigen tests and standard NAAT (either rapid RT-PCR or laboratory-based NAAT) vs. viral culture (all patients)



Supplement C

Recommendation 2: For symptomatic individuals suspected of having COVID-19, the IDSA panel suggests using standard NAAT (i.e., rapid RT-PCR and laboratory-based NAAT) over a rapid Ag test (conditional recommendation, low certainty evidence).

For the antigen test accuracy forest plots (symptomatic), refer to Figures s2a and s2b.

Figure s10a. Forest plot for the sensitivity of standard laboratory based NAAT in all patients

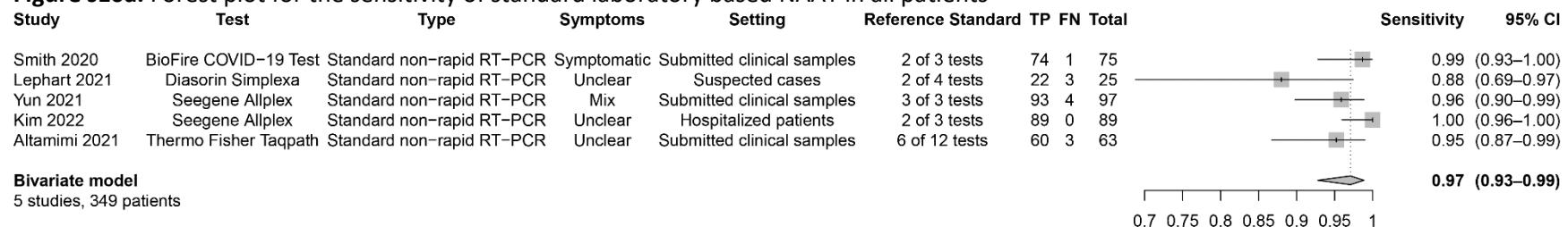
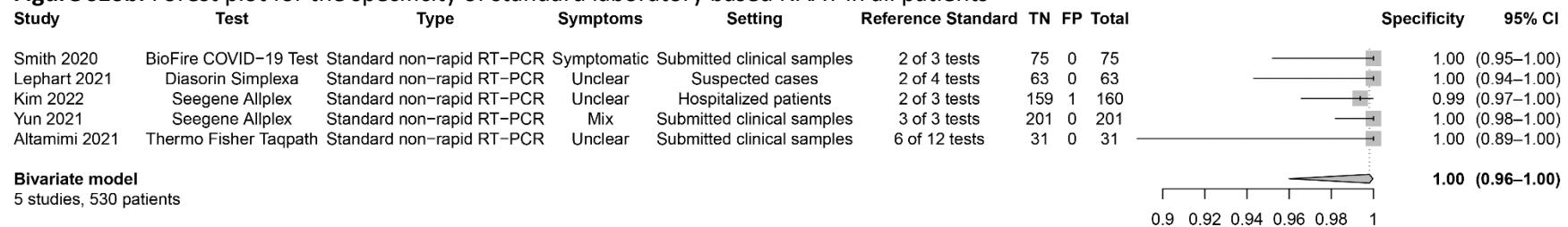


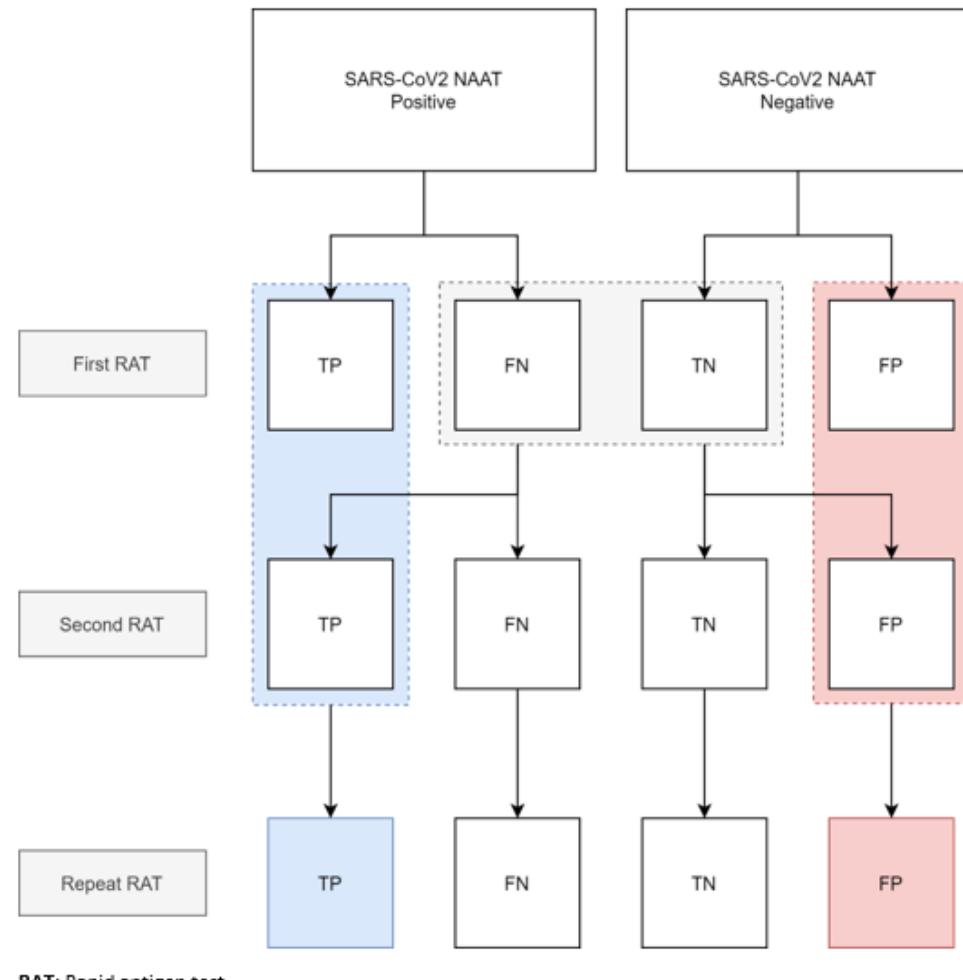
Figure s10b. Forest plot for the specificity of standard laboratory based NAAT in all patients



Supplement D

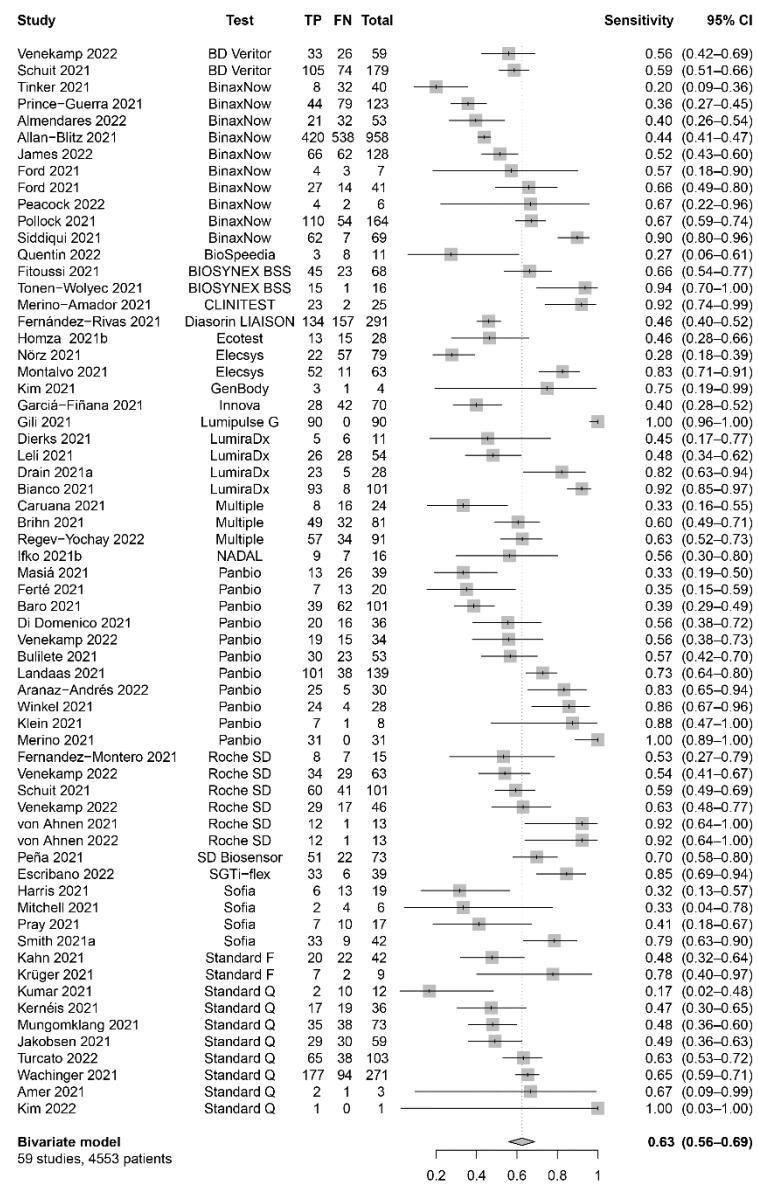
Recommendation 3: For symptomatic individuals suspected of having COVID-19, the IDSA panel suggests using a single standard NAAT (i.e., rapid RT-PCR and laboratory-based NAAT) rather than a strategy of two consecutive rapid Ag tests (conditional recommendation, very low certainty evidence).

Figure s11a. Repeat testing algorithm



Supplement E

Recommendation 4: For asymptomatic individuals with known exposure to SARS-CoV-2 infection, the IDSA panel suggests using a single (i.e., one-time) Ag test over no testing in specific situations (conditional recommendation, moderate certainty evidence).

*Supplementary Materials***Figure s12a.** Forest plot for the overall sensitivity of antigen tests in asymptomatic patients

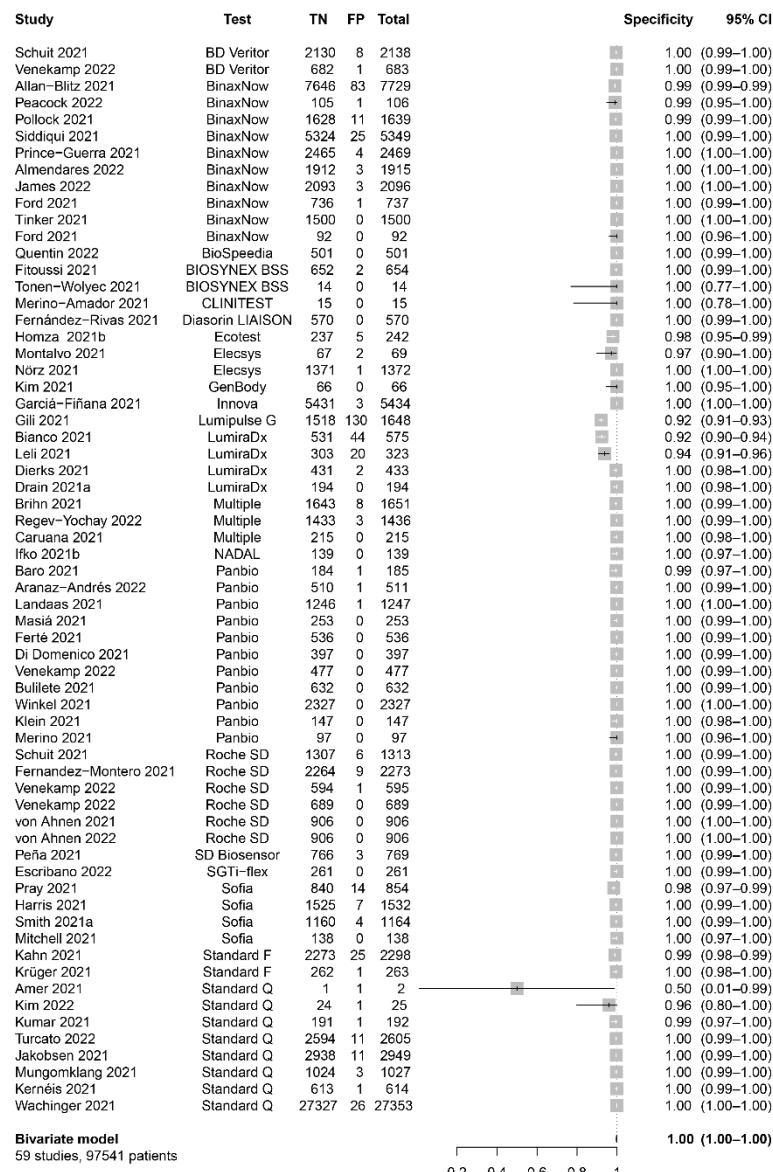
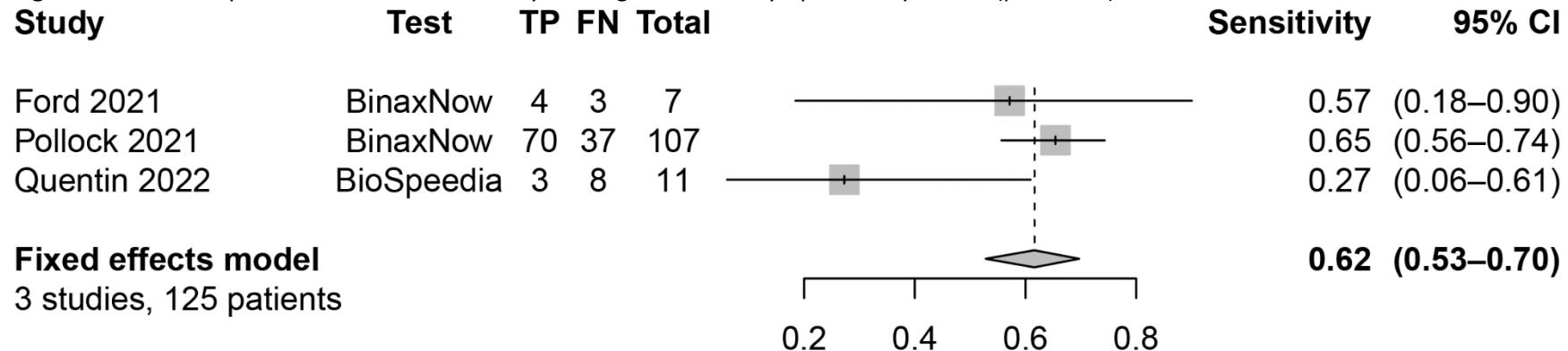
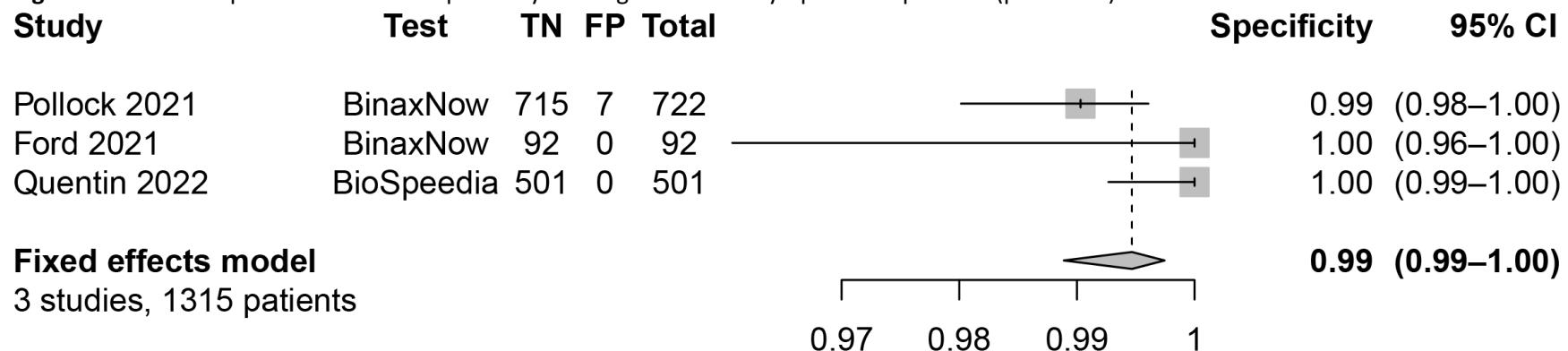
*Supplementary Materials***Figure s12b.** Forest plot for the overall specificity of antigen tests in asymptomatic patients

Figure s13a. Forest plot for the overall sensitivity of antigen tests in asymptomatic patients (pediatrics)**Figure s13b.** Forest plot for the overall specificity of antigen tests in asymptomatic patients (pediatrics)

Supplement F

Recommendation 5: For asymptomatic individuals with known exposure to SARS-CoV-2 infection, the IDSA panel suggests using a single standard NAAT (i.e., rapid RT-PCR and laboratory-based NAAT) over a single rapid Ag test (conditional recommendation, low certainty evidence).

For the antigen test accuracy forest plots (asymptomatic), refer to [Figure s12a](#) and [Figure s12b](#).

For the NAAT accuracy forest plots, refer to [Figure s10a](#) and [Figure s10b](#).

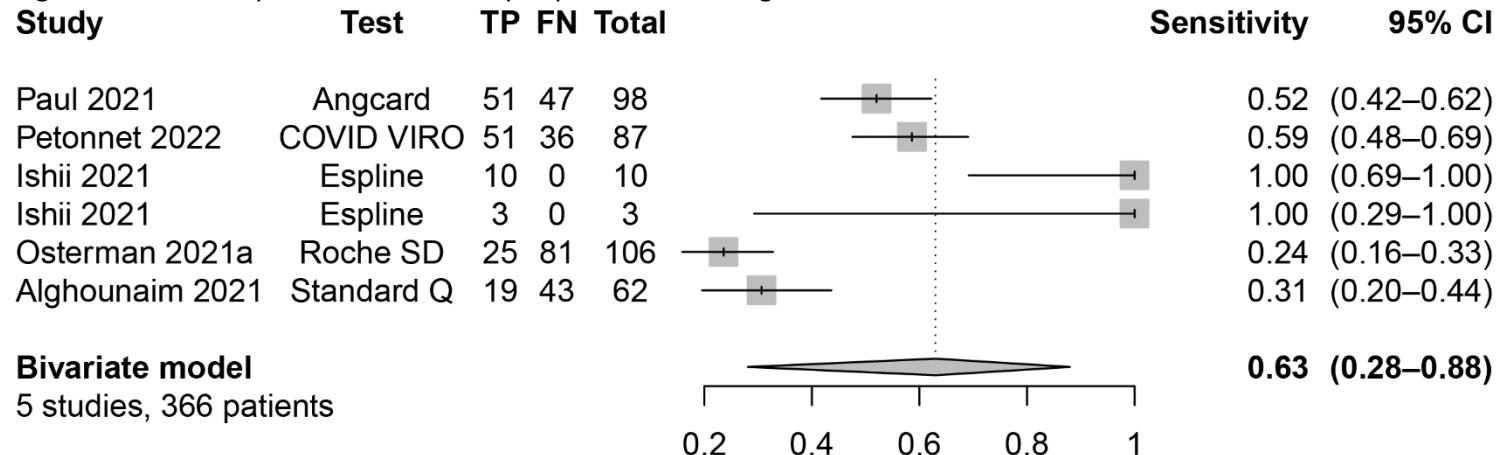
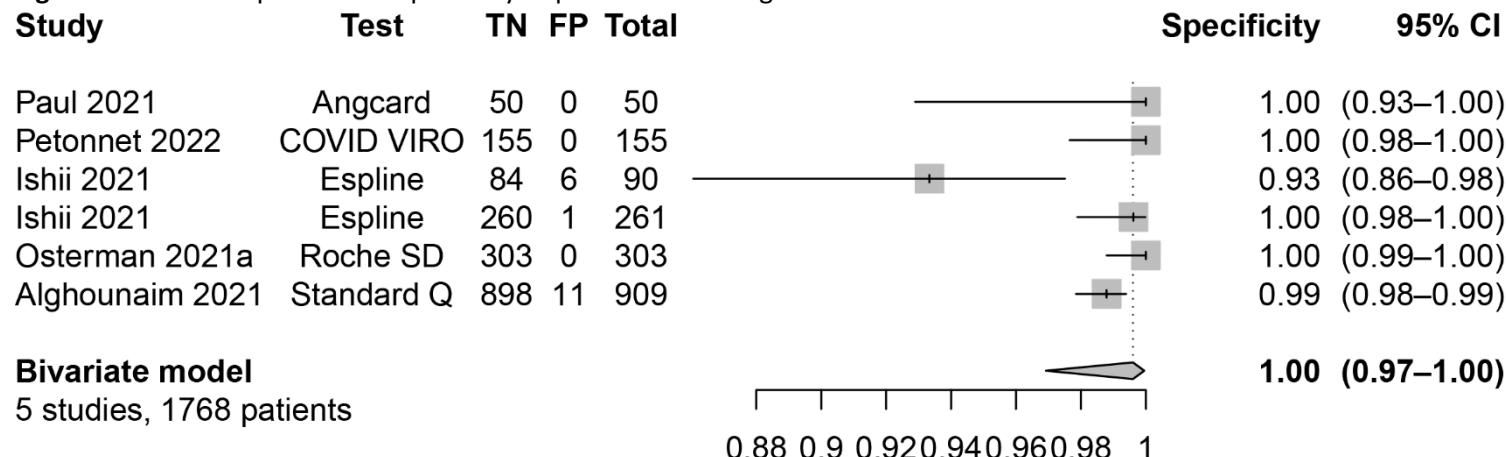
Supplement G

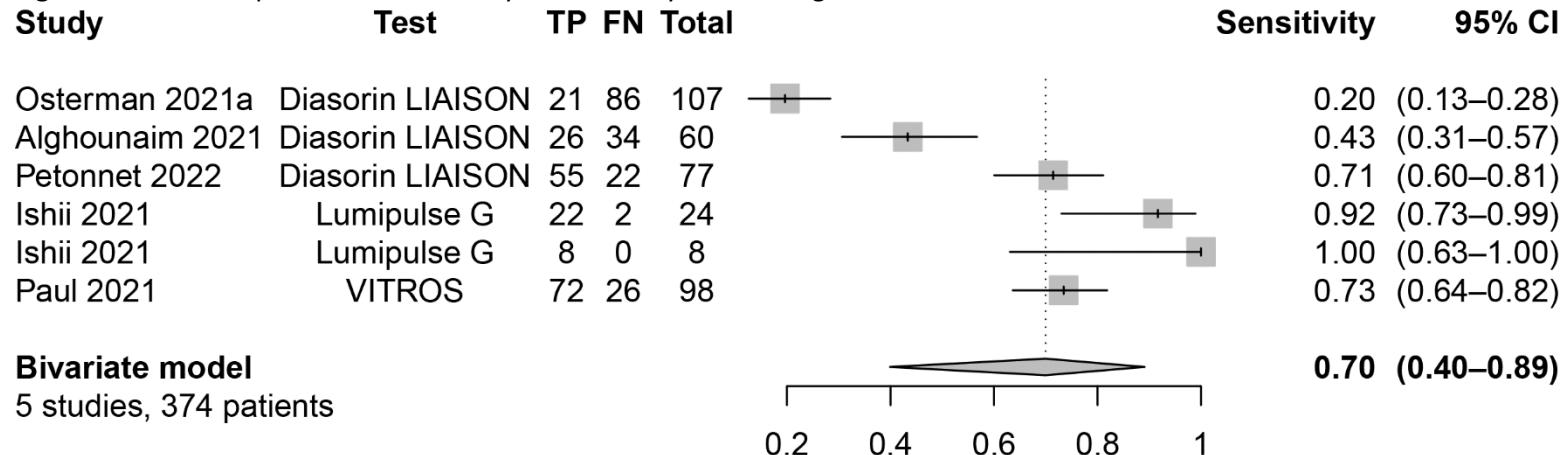
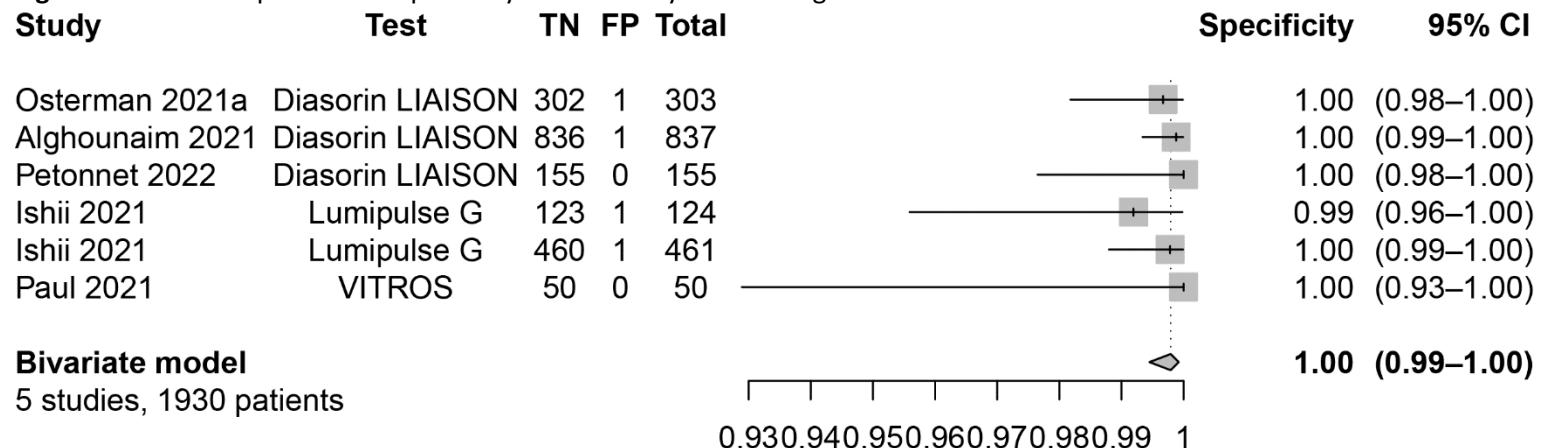
Recommendation 6: In asymptomatic individuals with a known exposure to SARS-CoV-2, if standard NAAT testing or results are not available in a timely manner and a first Ag test is negative, the IDSA panel suggests for repeating Ag testing (conditional recommendation, very low certainty evidence).

Refer to [Figure s11a](#) for repeat testing algorithm

Supplement H

Recommendation 9: For individuals for whom Ag testing is desired, the IDSA panel suggests for either point-of-care or laboratory-based Ag testing (conditional recommendation, low certainty evidence).

*Supplementary Materials***Figure s14a.** Forest plot for the sensitivity of point of care antigen tests**Figure s14b.** Forest plot for the specificity of point of care antigen tests

*Supplementary Materials***Figure s15a.** Forest plot for the sensitivity of laboratory-based antigen tests**Figure s15b.** Forest plot for the specificity of laboratory-based antigen tests

Supplement I

Recommendation 10: The IDSA panel suggests either observed or unobserved self-collection of swab specimens for Ag testing (conditional recommendation, low certainty evidence).

Figure s16a. Forest plot for the sensitivity of observed self-collection

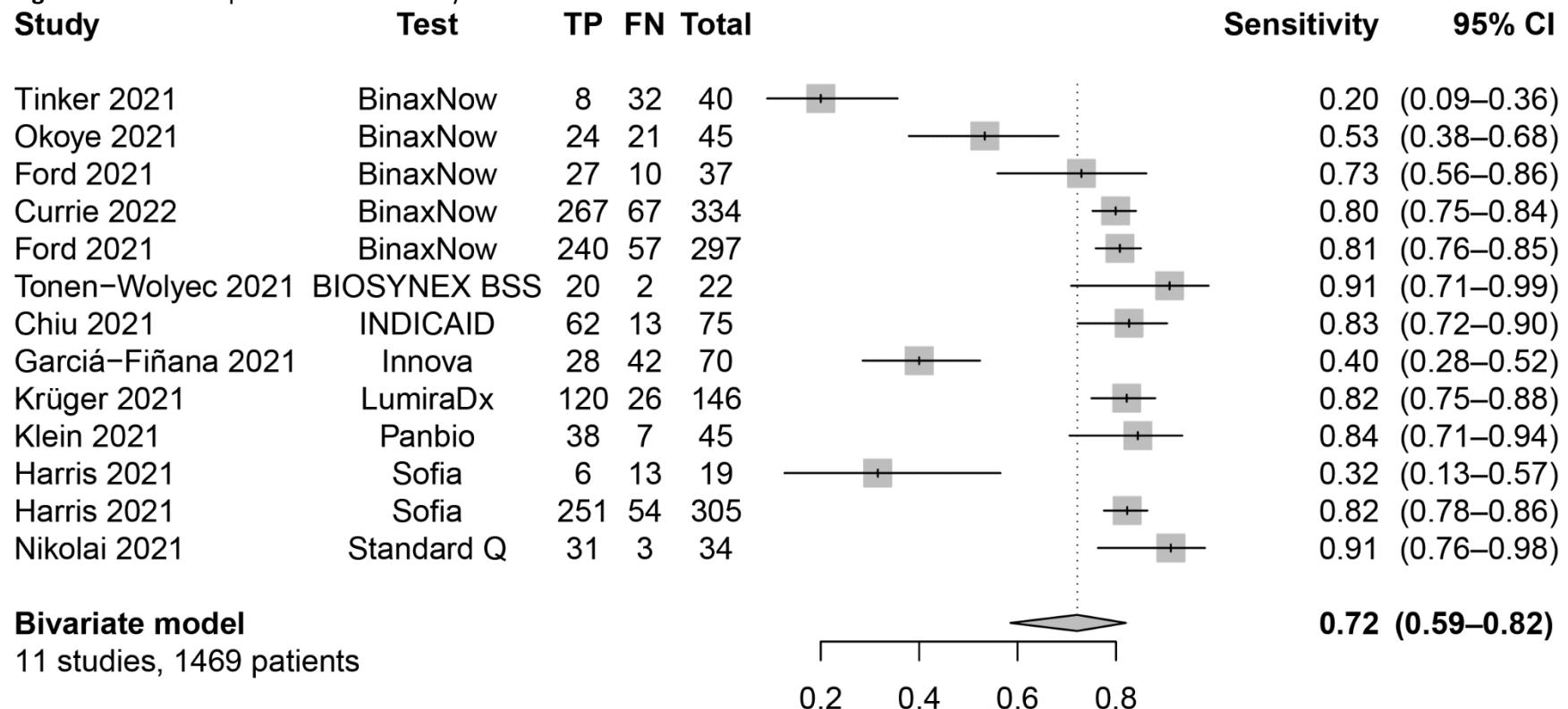


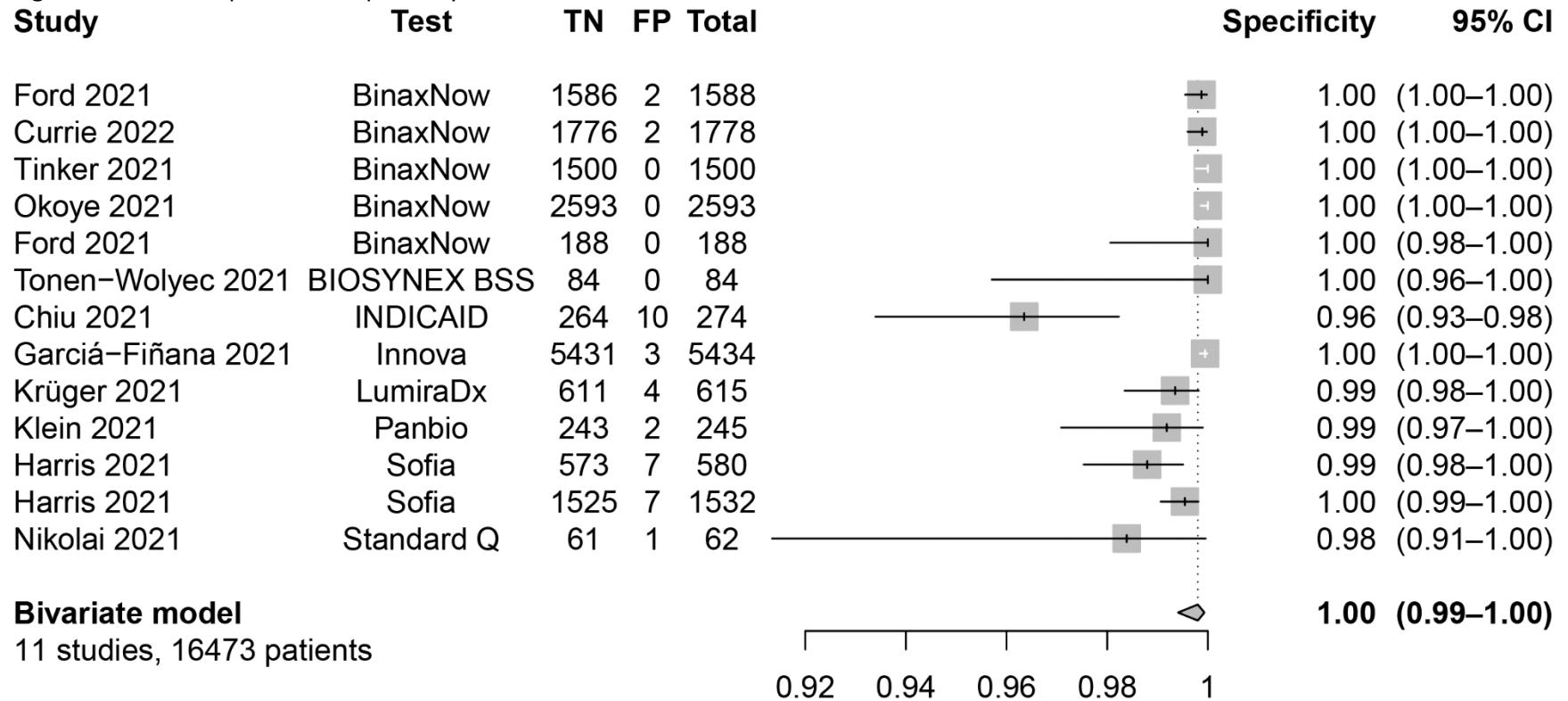
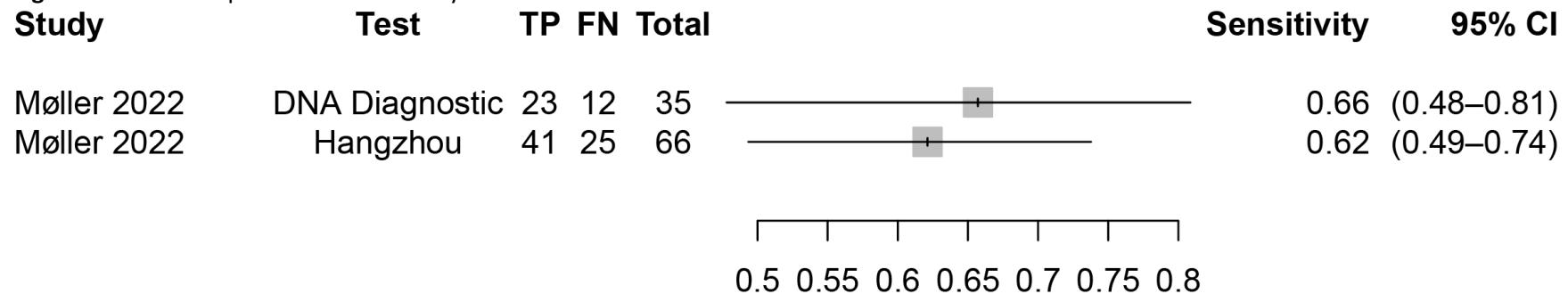
Figure s16b. Forest plot for the specificity of observed self-collection

Figure s17a. Forest plot for the sensitivity of unobserved self-collection**Figure s17b.** Forest plot for the specificity of unobserved self-collection