Results of COVID-19 Vaccine Effectiveness Studies: An Ongoing Systematic Review

Duration of Protection Weekly Summary Table

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Duration of Protection Studies

These are studies that assess duration of protection criteria as outlined above along with those studies that do not meet aforementioned criteria that are relevant to evaluating duration of protection. Some of these studies are also in the above table but duplicated here for ease. As of April 28, 2022, those studies that provide VE estimates at least 4 months after the primary series or at least 2 months after the booster series are included below.

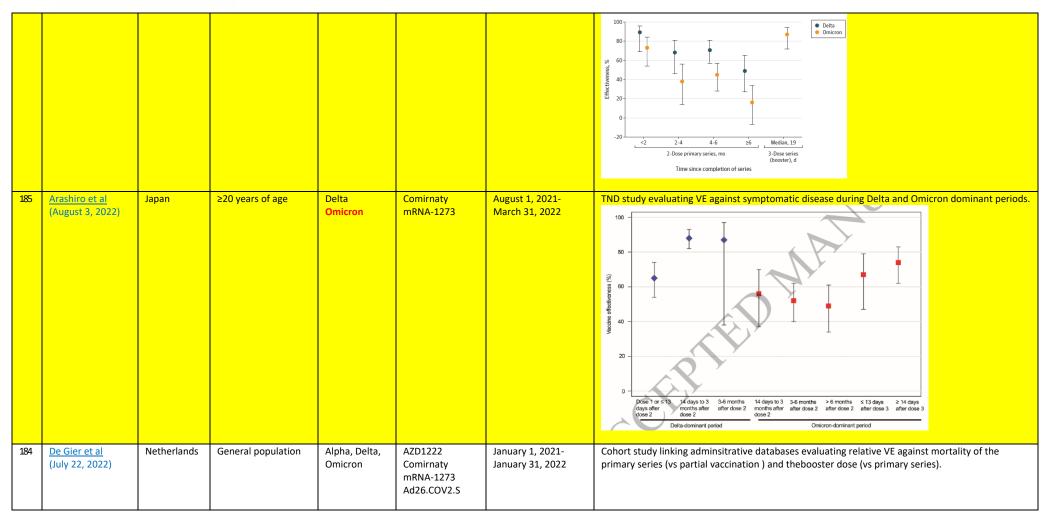
We would like to highlight:

- Countries have implemented different dose intervals and vaccination strategies that can make comparisons across studies challenging.
- Persons who are vaccinated early in a program are different than those who are vaccinated later. For example, many who were vaccinated early were those at highest risk, and this could confound the results. Some of the older individuals also might have some degree of immunosenescence.

#	Reference (date)	Country	Population	Dominant Variants	Vaccine product	Study Period	Descriptive Findings
188	Stirrup et al (August 9, 2022)	UK	LTCF residents and staff	Omicron	AZD1222 Comirnaty mRNA-1273	December 12, 2021- March 31, 2022	VIVALDI cohort study with regular asymptomatic testing in LTCF of staff and residents. Calculated relative VE of 3 rd dose compared to 84+ days post dose 2 against infection, hospitalization and death (providing results stratified by prior infection). In residents without known prior SARS-CoV-2 infection, there was reduced risk of SARS-CoV-2 infection in the periods 14-48 days: VE 72% (62-80) 49-83 days: VE 69% (60-76) after first booster vaccine dose, relative to 2-dose vaccination. The first booster dose reduced risk of hospitalisation 14-48 days: VE 81% (46-93), 49-83 days: VE 85% (68-93), 84+ days: VE 53 (11-76) from receipt of booster dose. The first booster reduced risk of death within 28 days of SARS-CoV-2 infection after 14-48 days VE 88% (66-96), 49-83 days: VE 89% (77-95), 84+ days: VE 63% (38-79).
187	Zambrano et al (August 4, 2022)	USA	5-18 year olds with MISC vs hospitalized negative SARS-CoV-2 controls	Delta, Omicron	Comirnaty	July 1, 2021-April 7, 2022	TND study comparing children with MISC to hospitalized children without SARS-COV-2. 28-120 days post dose 2 VE was 90 (75-96%); 121-200 day post dose 2 VE was 92% (78-97%).
186	Tartof et al (August 3, 2022)	USA	12-17 year old members of Kaiser Permanente Southern California	Delta Omicron	Comirnaty	November 1, 2021, - March 18, 2022	TND study linking administrative databases to evaluate VE against emergency department and urgent care visits (without subsequent hospitalization)

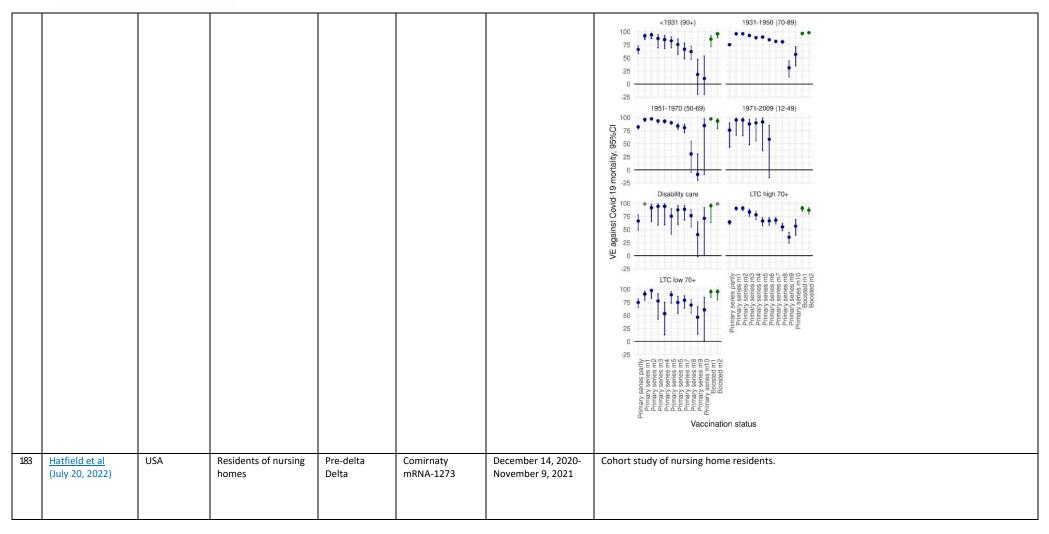
















							Unvaccinated Completed Pfizer-BioNTech, within past 150 days Completed Moderna, within past 150 days	871 1,196 466	57,871 103,668 35,290	Median days contributed per resident (IQR) minance (Dec 14, 2020 51 (21, 122) 95 (87, 404) 86 (73, 89) mance (Jun 21, 2021 - 141 (60, 141) 14 (5, 14) 126 (84, 135) 21 (14, 32) 109 (30, 122)	109 22 6	Vaccine Effectiveness % (95% CI) REF 67% (40%, 82%) 75% (32%, 91%) REF Not Estimated ^b 33% (-2%, 56%) Not Estimated ^b 77% (48%, 91%)
182	Cerqueria-Silva et al (July 18, 2022)	Brazil	≥18 year olds	Omicron	Coronavac followed by Comirnaty booster	January 1-April 17, 2022	First dose ≥ 14 Second dose 14-180 > 180 Booster with BNT162b2 0-13 14-30 31-60 61-90 91-120	Outcomptomatic and Severe adjusted vaccing against symptomatic	-59 years ① ① ① ① ② ② ① ① ③ ③ ② ② ① ② ③ ① ② ③ ② ③ ② ③ ③ ③ ③ ③ ③ ③	60-79 years (D) (O) (O) (O) (O) (O) (O) (O) (O) (O) (O	0 0 1 1-120 >120 0-13 14	≥ 80 years

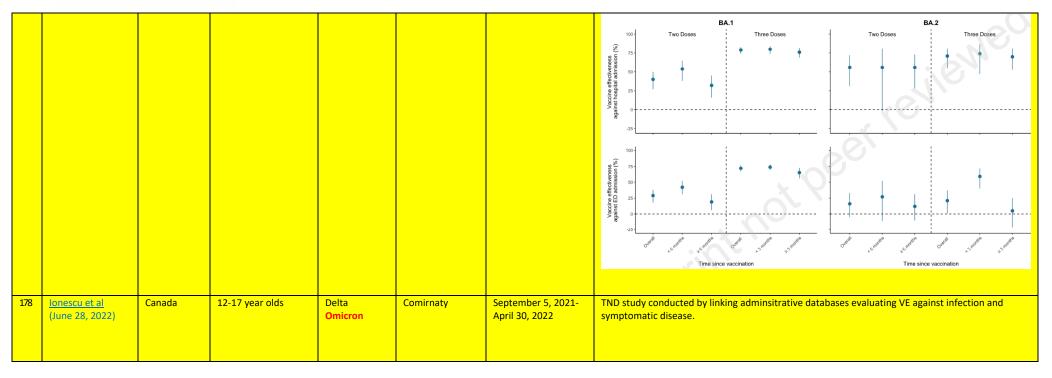




181	Link-Gelles et al	USA	≥18 year olds	Omicron	Comirnaty	December 18, 2021-	TND study in the VISIO	N netw	ork eval	uating	VE agai	inst E	D/urgent	care v	visit and	d hospitali	zaiton.
	(July 15, 2022)			(BA1, BA2 /	mRNA-1273	June 10, 2022	_		Omicron BA.1-	-predomina	ant period¶			Omicron B	A.2/BA.2.12.	1–predominant p	eriod**
	, , , , ,			BA2.12.1)		,	-	No Total	. (%) of positive test results [†]	since l	n interval last dose,	VE %* (95%	CI) Total	No. (%)	of positive results†	Median interval since last dose,	VE %* (95% CI)
							Encounter type ED or UC, age group (days since last dose		test results'	day	/s (IQR)	%* (95%	CI) Iotal	test	results'	days (IQR)	%* (95% CI)
							All ages, yrs	e)									
							Unvaccinated (Ref)	51,359	23,175 (45.1)		-		27,907		01 (12.6)		
								7,286 32,740	2,377 (32.6) 11,365 (34.7)		(76–129) 232–306)	47 (44-5) 39 (37-4			110 (6.2) 84 (12.4)	104 (71-128) 352 (278-398)	51 (38–60) 12 (7–17)
							3 doses (7–119)	29,333	3,667 (12.5)	66	6 (41-89)	84 (83-8	(5) 9,142	2 4	441 (4.8)	94 (72-108)	56 (51-61)
							i i	3,315	217 (6.5)	132 (1	125-142)	73 (68–7	7) 26,654	3,18	86 (11.9)	166 (145-190)	26 (21–30)
							18–49 yrs Unvaccinated (Ref)	33,003	14,236 (43.1)		_		- 18,429	2,26	69 (12.3)	_	_
								4,909	1,621 (33.0)		(76–129)	40 (36-4			75 (6.3)	105 (72–129)	47 (31–60)
								16,313 8,755	5,918 (36.3) 1,259 (14.4)		220–288) 5 (33–79)	24 (21–2 76 (75–7			27 (12.7) 207 (5.0)	332 (254–379) 91 (69–107)	7 (0–14) 55 (47–62)
							3 doses (≥120)	426	39 (9.2)		124–141)	29 (-1-5			96 (14.4)	159 (140-182)	17 (10-25)
							≥50 yrs										
								18,356 2,377	8,939 (48.7) 756 (31.8)	109	(77–129)	59 (54–6	— 9,478 (3) 582		32 (13.0) 35 (6.0)	102 (68–128)	59 (40-71)
							2 doses (≥150)	16,427	5,447 (33.2)	283 (2	248-316)	52 (50-5	9,680	1,15	57 (11.9)	376 (319-414)	18 (10-26)
								20,578 2,889	2,408 (11.7) 178 (6.2)		1 (46–93) 125–143)	87 (86-8 81 (77-8			234 (4.7) 90 (11.0)	96 (73-109) 170 (147-193)	58 (51–64) 32 (26–38)
							4 doses (≥7)††	N/A	178 (0.2)	133 (125-143)	81 (//-8	— 4,094		355 (8.7)	28 (17–42)	66 (60–71)
							Hospitalization, age group (days since la	ist dose)									
							All ages, yrs										
								14,742	6,829 (46.3) 297 (24.0)	105	(73–129)	68 (63-73	- 6,682 3) 343		12 (3.5)	102 (71-128)	- 57 (19-77)
								8,850	2,542 (28.7)		252-322)	61 (58-63			12 (3.5)	371 (308–413)	24 (12–35)
							3 doses (7–119)	9,146	786 (8.6)	72	2 (47-93)	92 (91-93	3) 2,350		72 (3.1)	94 (74-108)	69 (58-76)
								1,425	80 (5.6)	132 (1	125-142)	85 (81-89	9) 7,686	5	19 (6.8)	168 (146-191)	52 (44–59)
							18–49 yrs ^{§§} Unvaccinated (Ref)	4,057	1,515 (37.3)		_	_			-	_	1-1
							2 doses (14–149)	392	83 (21.2)		(67-127)	64 (52-73			55	-	100 to
							2 doses (≥150) 3 doses (7–119)	1,304 812	329 (25.2) 53 (6.5)		226-294) 7 (36-81)	52 (43-59 91 (87-94			_	_	
							3 doses (≥120)	56	1 (1.8)		126-142)	94 (62-99			_	_	1-1
							≥50 yrs ^{§§}	10 505	F 24 4 (40 T)				4.505		102 (0.6)		
							Unvaccinated (Ref) 1 2 doses (14–149)	10,685 844	5,314 (49.7) 214 (25.4)	108	(76–129)	71 (65-75	- 4,595 5) -	. 3	193 (8.6)	_	_
							2 doses (≥150)	7,546	2,213 (29.3)	294 (2	259-325)	63 (60-66	6) 4,139		152 (8.5)	381 (325-418)	22 (8-34)
								8,334 1,369	733 (8.8) 79 (5.8)		3 (49-94) 125-142)	92 (91-93 86 (82-89			57 (2.9) 180 (6.8)	95 (74-108) 169 (147-191)	73 (63–81) 55 (46–62)
							4 doses (≥7) ^{††}	N/A	-	132(-	-	- 1,204		74 (6.2)	27 (17-41)	80 (71–85)
180	Tonnaro et al	San Marino	≥18 year old	Alpha, Delta	Sputnik V	February 21-October	Cohort study of entire	countr	/.		Any vaccir	10			G=	m-COVID-Vac	
	(July 4, 2022)					1, 2021		Perio	1	C	rude		justed ^a	. *	Cru		Adjusteda
									Cases*	VE	95% CI	VE	95% CI	Cases*	VE	95% CI V	95% CI
							SARS-CoV-2 infections										
		İ		Ì		1		<60 da			94.9-97.8 81.0-87.7		82.8-92.6 40.3-60.7	16 117		5.3-98.2 91 7.1-84.4 47	
		ĺ						120+			81.1-88.9		40.3-60.7 36.7-63.8	53		7.1-84.4 47 1.3-89.2 57	
		ĺ						Tota			87.2-91.0		61.8-72.5	186	89.9 8		
							COVID-19 related Hospitalizations	_				21.13	.()				
								<60 da	ys 5	94.5	84.9-98.0	90.6	74.9-96.5	2	97.5 8	8.9-99.4 95	.2 79.1-98.9
		ĺ						60-11			88.4-98.7		73.4-96.6	4		6.5-98.5 87	
								120+			71.5-95.9	76.1	35.1-91.2	2		2.5-99.2 89	
179	Tartof et al	USA	≥18 year old	Omicron (BA1	Comirnaty	December 27, 2021-	TND study evaluating \	/E again	_				77.4-93.5	8	96.4 9		6 81.5-96.2
1/3	(June 30, 2022)	03/	members of Kaiser Permanente	and BA2)	Committaty	June 4, 2022	THE study evaluating t	v L agaii	130 HO3PI	tanzat	.ioir ariu	Cinci	Series de	-par till	ieni (Li	o, autilissi	011.
			southern California														

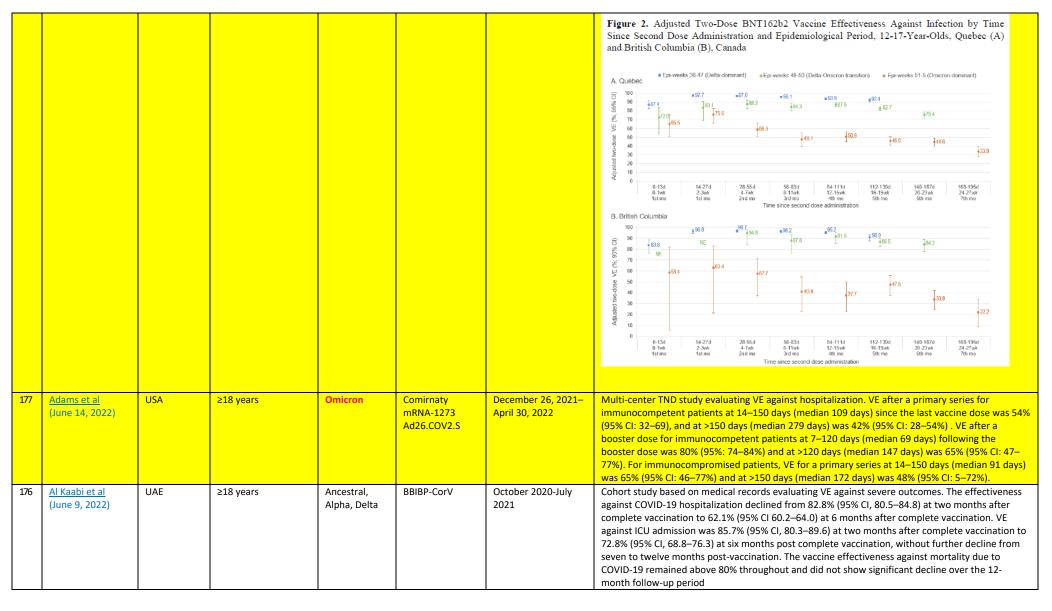






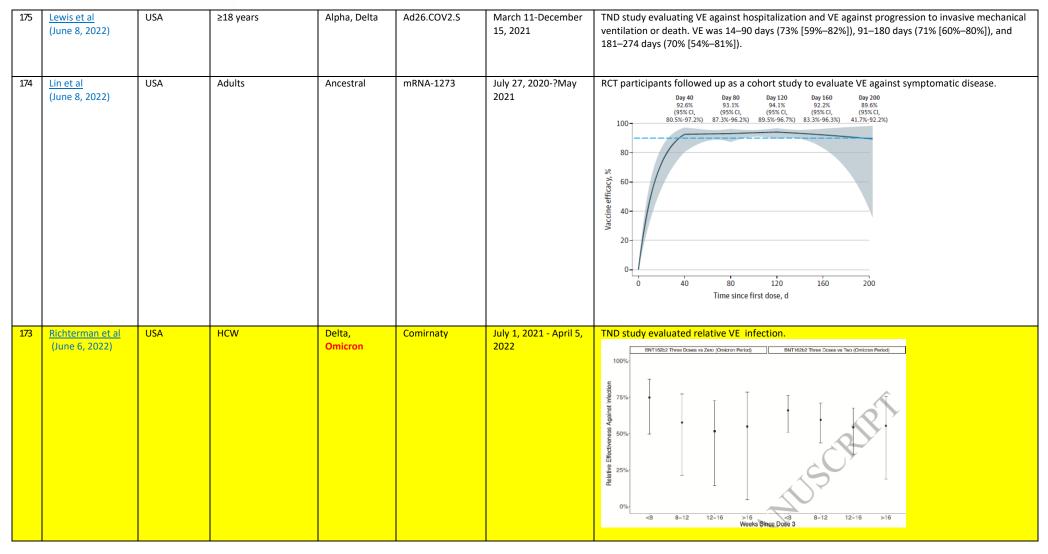
















172	Andrejko et al (June 3, 2022)	USA	12+ year olds	Pre-Omicron	Comirnaty mRNA-1273	February 23- December 5, 2021	reported. The figure vaccination card for	belows sh vaccinatio	ows VE over n data.		that vaccination data wons who were asked to	
171	Accorsi et al (May 25, 2022)	USA	18+ year olds	Omicron	Comirnaty mRNA-1273 Ad26.COV2.S	January 2-March 23, 2022	TND study based on Vaccination Regimen No vaccination Ad26.COV2.S 14 days to 1 mo since last dose 2 to 4 mo since last dose Ad26.COV2.S/Ad26.COV2.S 14 days to 1 mo since last dose 2 to 4 mo since last dose Ad26.COV2.S/mRNA 14 days to 1 mo since last dose mRNA/mRNA 14 days to 1 mo since last dose mRNA/mRNA 14 days to 1 mo since last dose mRNA/mRNA 14 days to 1 mo since last dose	No. of Tests 207,784 se 706 3,100 se 1,017 2,506 se 3,585 9,752	Positive for SARS-CoV-2 % 50.1 47.2 49.8 46.9 41.5 31.5 30.4 27.3 26.6	Vaccine Effect	Reference 17.8 (4.3–29.5) 8.4 (1.5–14.8) 27.9 (18.3–36.5) 29.2 (23.1–34.8) 61.3 (58.4–64.0) 54.3 (52.2–56.3) 68.9 (68.3–69.5) 62.8 (62.2–63.4) 75 100	call.
170	Amir et al (May 25, 2022)	Israel	12-15 year olds	Omicron	Comirnaty	December 26, 2021- January 8, 2022	Cohort study conduct Ages 12-15 3rd dose effect Cohort Confirmed Infections (at-risk days) Univaccinated 2,684 (834,149) 2nd dose (153 (115,371) (14-60 days) 2nd dose (60-120 days) 2nd dose (120-days) (2,003,011) Internal control 494 (180,100) 3rd dose (14-60 days) (166 (171,281) (14-60 days)	Adjusted rate ratio vs. 3rd dose 5.0 [4.3, 5.9] 2.2 [1.8, 2.8] 3.8 [3.3, 4.5] 4.2 [3.6, 4.9] Ref	king admin d	latabases looking a	at risk against infection.	





460	1	1117	D	Alaba Balia	ChAdOx1	D	T TAID II die een dere die differente en latie en 1916 eeu en 1917
169	<u>Lee et al</u> (May 23, 2022)	UK	Persons with cancer	Alpha, Delta		December 8, 2020- October 15, 2021	Two TND studies conducted in different populations with comparison of VE against infection, hospitazliation, and death among the two groups.
	(IVIdy 23, 2022)		and general population		Comirnaty	October 15, 2021	
							100 90 80 70 50 60 50 10- Cancer cohort 10- Control population 0 -8 to -1 0 to 8 9 to 16 17 to 24 25 to 32 33 to 40
							Weeks after second COVID-19 vaccine dose
							Post-second dose (overall) 3-6 months post-second dose Vaccine Vaccine
							Exposed (PCR-positive) Not exposed (PCR-negative) Effectivenes Outcome Post-2 rd Univaccinate Post-2 rd Univaccinate s (%) measure dose (n) d (N) dose (n) d (N) dose (n) d (N) dose (n) d (N) d (N) dose (n) d (N)
							Breakthrough 65.5% 47.0% 465982 65.1-65.9) 12513 31649 347414 465982 46.3-47.6)
							Coronavirus 84.5% Hospitalisation 837 3227 780054 465982 (83.6-85.4) 611 3227 347414 465982 (72.8-76.3)
							Coronavirus 93.5% 90.3% Death 560 5139 780054 465982 (93.0-94.0) 373 5139 347414 465982 (89.3-91.2)
168	Paranthaman et al	England	≥65 years living in	Alpha, Delta	ChAdOx1	December 8, 2020-	Cohort study conducted by linking adminsitrative databases evaluating VE against infection and
	(May 5, 2022)		LTCF		Comirnaty	September 30, 2021	death.
							Table 2. Adjusted HRs for infection by vaccination status for LTCF residents, England
							Vaccination Time since Any ChAdOt-1 BNT162b2
							Person-time in days Events Adjusted HR ^b Person-time in days Events Adjusted HR ^b Person-time in days Events Adjusted HR ^b (unique individuals) ^a (unique individuals) ^a (unique individuals) ^a (unique individuals) ^a
							Unvaccinated 6.958.732 (190.202) 26.765 6.958.732 (190.202) 26.765 6.958.732 (190.202) 26.765
							First dose 1-2 wks 2,070,258 (153,383) 8,190 0.68 (0.62-0.74) 1,427,012 (105,589) 5,256 0.67 (0.6-0.75) 643,246 (47,803) 2,934 0.68 (0.6-0.78) 3 wks 990,274 (143,432) 2,762 0.64 (0.57-0.73) 664,527 (99,949) 1,731 0,73 (0.63-0.86) 395,747 (44,347) 1,051 0,56 (0.48+0.67)
							4 wks 965,091 (139,327) 1,554 0.5 (0.43-0.59) 671,379 (96,744) 921 0.58 (0.48-0.7) 293,712 (42,583) 633 0.48 (0.39-0.59)
							5 wks 948,533 (136,661) 1.057 0.47 (0.4-0.56) 660,612 (95,140) 654 0.59 (0.47-0.73) 287,921 (41,521) 403 0.44 (0.36-0.55) 6-7 wks 185,2109 (134,595) 1.190 0.46 (0.38-0.56) 122,0208 (93,718) 642 0.5 (0.4-0.62) 561,901 (40,877) 548 0.52 (0.41-0.66)
							8-10 wks 2,472,598 (130,173) 815 0.64 (0,5-0.82) 1,715,549 (90,634) 347 0.51 (0,38-0.68) 757,449 (39,539) 468 0.79 (0,59-1.06)
							11+ wks 1,112,436 (86,592) 254 0.83 (0.62-1.11) 768,455 (57.784) 181 0,94 (0.67-1.33) 343,981 (28.718) 73 0.63 (0.44-0.9) Second dose 1-4 wks 3,452,288 (124,173) 259 0.4 (0.29-0.55) 2,401,640 (86,845) 119 0.39 (0.26-0.6) 1,030,648 (37,328) 120 0.38 (0.27-0.54)
							5-10 wis 5,037 822 (122,460) 179 0.47 (0.34-0.64) 3,521.278 (85,615) 134 0.54 (0.37-0.78) 1,516,544 (36,785) 45 0.54 (0.21-0.55) 11-15 wis 4,035,312 (117,469) 384 0.45 (0.34-0.59) 2,810,444 (81,979) 327 0.48 (0.36-0.64) 1,224,868 (35,450) 57 0.31 (0.2-0.48)
							16-20 wls 3,757,167 (111,858) 1384 0.66 (0.54-0.81) 2,599,430 (77.764) 1090 0.72 (0.58-0.9) 1,157,737 (34,094) 294 0.55 (0.39-0.78)
							21+ was 3381,529 (99.696) 2.104 0.6 (0.49-0.74) 2.070,748 (68.221) 1.474 0.71 (0.57-0.9) 1.310,781 (31.475) 630 0.53 (0.42-0.68) Number of unions individually scale for any distriction of time within each time partial. Pulling of the condense of the property of the condense of the con
							*Number of unique individuals at risk for any duration of time within each time period. *Adjusted for gender, age group, case rate in local authority and deprivation, along with a cluster term for care home postcode. See Supplementary Figure S4, Supplementary Tables S1 and S2 in Supplementary data.

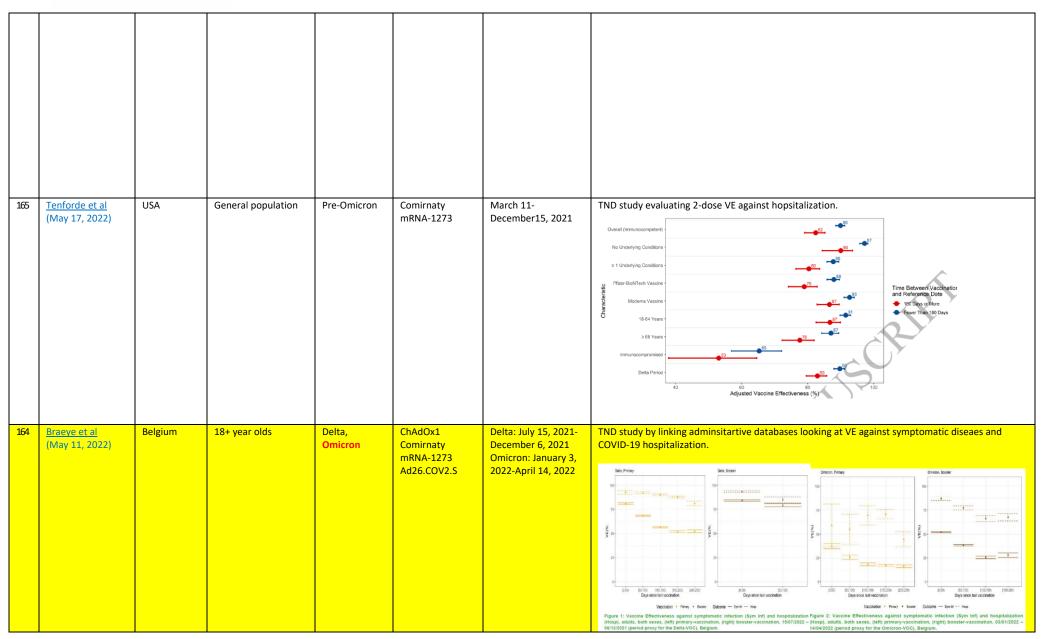




			1			<u> </u>	Table 3. Adi	justed HRs for CO	VID-rela	ited death by	v vaccination s	rarus amo	one LTCF re	sidents. Englan	d	
								since Any			ChAdOx-1		8	BNT162b2		
							11-1: 16-2: 21+1 *Number of uniqu	6,931,978 (190,109) vks 2,070,228 (153,379) vks 1,955,365 (143,880) vks 3,697,628 (137,419) rks 2,668,668 (124,523)	7,425 2,125 812 347 71 18 15 43 193 280 ny duration	0.59 (0.52–0.66) 0.41 (0.35–0.48) 0.35 (0.26–0.41) 0.44 (0.3–0.63) 0.15 (0.67–0.3) 0.19 (0.09–0.41) 0.21 (0.13–0.34) 0.35 (0.24–0.52) 0.37 (0.25–0.53) of time within o	6,931,978 (190,109) 1,426,998 (105,578) 1,355,906 (99,356) 2,575,162 (95,636) 1,844,561 (86,556) 240,1617 (86,843) 3,521,162 (85,610) 2,810,271 (81,971) 2,998,423 (77,717) 1,916,253 (64,662) each time period. ^b ₄	7,425 1,364 485 178 36 9 10 39 155 196 Adjusted for g	0.58 (0.5-0.66) 0.49 (0.4-0.61) 0.37 (0.27-0.5) 0.43 (0.26-0.71) 0.17 (0.06-0.42) 0.18 (0.07-0.47) 0.22 (0.13-0.38) 0.44 (0.3-0.67) gender, age grou	643,230 (47,801) 599,459 (44,556) 1,122,466 (41,783) 824,107 (37,967) 1,030,631 (37,325) 1,516,513 (36,784) 1,224,835 (35,428) 1,137,582 (34,987) 1,230,371 (30,054) up, case rate in local a	7,425 761 327 169 35 9 5 4 38 84 uthority an	0.6 (0.51-0.7) 0.35 (0.29-0.43) 0.34 (0.26-0.45) 0.5 (0.32-0.78) 0.14 (0.06-0.33) 0.19 (0.05-0.7) 0.09 (0.03-0.25) 0.27 (0.16-0.46) 0.31 (0.2-0.49)
167	Martellucci et al (April 22, 2022)	Italy	General population	Alpha, Delta, Omicron	ChAdOx1 Comirnaty mRNA-1273 Ad26.COV2.S	January 2, 2021- December 18, 2021		y conducted by ion, and death.		g adminsit	trative data	bases ev	valuating	VE against i	nfectio	n,
							Var	iables			OVID-19 oitalization	A		COVID-:	19-Rel eath	ated
							Follow-u	p duration ^B			R (95% CI)			OR (9	5% CI)
								s of follow-up								
								ccinated			(Ref. cat.)				f. cat.)	
								doses			(0.02-0.03)			0.01 (0.0		,
								doses of follow-up		0.18	(0.15–0.23)	*		0.15 (0.1	10-0.24	ł) *
								ccinated		1	(Ref. cat.)			1 (Re	f. cat.)	
								doses			(0.26–0.37)	*		0.25 (0.1		
166	Fano et al (May 18, 2022)	Italy	12+ year olds	Alpha, Delta, Omicron	ChAdOx1 Comirnaty mRNA-1273 Ad26.COV2.S	January 1, 2021- January 10, 2022	Figure 2 – Adjusted values after the adminisumvaccinated 100 – 80 – 80 – 80 – 80 – 80 – 80 – 80	y conducted by scaling effectiveness (VE) again stration of the second dose	at SARS-CoV-3	Infection at different ter dose. Reference ter dose. Reference ter dose. Reference ter dose terror t	vent		_			n.











163	Butt et al (May 3, 2022)	USA	Veterans	Omicron	Comirnaty mRNA-1273	January 1-February 20, 2022	their booster v [35-44%] for B negligible for b	BNT-162b2; RVE=30% both vaccines for pati	s of the sta [23-36%] for ents with 4	rt of the period or mRNA-1273 For more mont	d of omicron pr), and protectic ths since receiv	edominance (RVE=40% on against infection was
162	Amir et al (May 5, 2022)	Israel	60+ year olds	Omicron	Comirnaty	January 16, 2022, to March 12, 2022	Cohort study k	oy linking adminsitrat	ive databas	ses evaluating i	relative VE agai	nst severe disease.
	(1717 17 7					, ,			VE	LCI	UCI	
							2nd dose	4+ months		ref		
								0-1 month	57%	38%	71%	
							0	1-2 months	66%	44%	79%	
							3rd dose	2-3 months 3-4 months	68% 67%	55% 58%	78% 73%	
							rd c	4-5 months	64%	60%	73%	
							m	5-6 months	64%	60%	69%	
								6-7 months	68%	58%	76%	
							4th dose	0-2 months	89%	87%	91%	





160 Castillo et al	ed VE of 2 doses of	e that they evaluated VE o	e that they e	e study. Note	art of Sisonk	nducted as p	TND study cor	Novem	Comirnaty	Omicron	HCW	South Africa	Gray et al	161
Bell Prince 18+ year olds Delta, Omirron December 13, 2021 This study linking administrative databases to assess WE against symptom confort study done among cord flooring linking administrative databases to assess WE against symptom confort study done among cord flooring linking administrative databases to assess WE against symptom confort study done among cord flooring linking administrative databases to assess WE against symptom confort study done among cord flooring linking administrative databases to assess WE against symptom confort study done among cord flooring linking administrative databases to assess WE against symptom confort study done among cord flooring linking administrative databases to assess WE against symptom confort study done among cord flooring linking administrative databases to assess WE against symptom confort study done among cord flooring linking administrative databases to assess WE against symptom confort study done among cord flooring linking administrative databases to assess WE against symptom confort study done among cord flooring linking administrative databases to assess WE against symptom confort study done among cord flooring linking administrative databases to assess WE against symptom confort study done among cord flooring linking administrative databases to assess WE against symptom confort study done among cord flooring linking administrative databases to assess WE against symptom confort study done among cord flooring linking administrative databases to assess WE against symptom confort study done among cord flooring linking administrative databases to assess WE against symptom confort study done among cord flooring linking administrative databases to assess WE against symptom confort study done among cord flooring linking								Januar	Ad26.COV2.S					
### December 13, 2021 - The Study linking administrative databases to assess VE against symptom cohort study done among coyol hospitalized cases. ### December 13, 2021 - The Study linking administrative databases to assess VE against symptom cohort study done among coyol hospitalized cases. ### December 13, 2021 - The Study linking administrative databases to assess VE against symptom cohort study done among coyol hospitalized cases. ### December 13, 2021 - The Study linking administrative databases to assess VE against symptom cohort study done among coyol hospitalized cases. ### December 13, 2021 - The Study linking administrative databases to assess VE against symptom cohort study done among coyol hospitalized cases. ### December 13, 2021 - The Study linking administrative databases to assess VE against symptom cohort study done among coyol hospitalized cases. ### December 13, 2021 - The Study linking administrative databases to assess VE against symptom cohort study done among coyol hospitalized cases. ### December 13, 2021 - The Study linking administrative databases to assess VE against symptom cohort study done among coyol hospitalized cases. ### December 13, 2021 - The Study linking administrative databases to assess VE against symptom cohort study done among coyol hospitalized cases. ### December 13, 2021 - The Study linking administrative databases to assess VE against symptom cohort study done among coyol hospitalized cases. ### December 13, 2021 - The Study linking administrative databases to assess VE against symptom cohort study done among coyol hospitalized cases. ### December 13, 2021 - The Study linking administrative databases to assess VE against symptom cohort study done among coyol hospitalized cases. ### December 13, 2021 - The Study linking administrative databases to assess VE against symptom cohort study done among coyol hospitalized cases. ### December 13, 2021 - The Study linking administrative databases to assess VE against symptom cohort study done among coyol hospitali	i–4 Mo ≥5 Mo	-2 Mo 3–4 Mo	-2 Mo				i i						, , , ,	
Both					т	,-	1007							
150 Castillo et al	1 ⁷³ 1 ⁶⁷ 1	70 70 71 71	70 70] ⁷²]	74 88 69	81	Vaccine Effectiveness (%) 80- 00- 00- 00- 00- 00- 00- 00- 00- 00-							
Practical Comment Proposal	anticipi anticipi anticipi	cold and and and and and and and and and an	Pecont State Par	Spy William William	RATISTO BATIS	s.covi.5 Ratisco								
December 13, 2021 - TND study linking administrative databases to assess VE against symptom cohort study done among covid hospitalized cases. December 13, 2021 - TND study linking administrative databases to assess VE against symptom cohort study done among covid hospitalized cases. December 13, 2021 - TND study linking administrative databases to assess VE against symptom cohort study done among covid hospitalized cases. December 14, 2021 - TND study linking administrative databases to assess VE against symptom cohort study done among covid hospitalized cases. December 14, 2021 - TND study linking administrative databases to assess VE against symptom cohort study done among covid hospitalized cases. December 14, 2021 - TND study linking administrative databases to assess VE against symptom cohort study done among covid hospitalized cases. December 14, 2021 - TND study linking administrative databases to assess VE against symptom cohort study done among covid hospitalized cases. December 14, 2021 - TND study linking administrative databases to assess VE against symptom cohort study done among covid hospitalized cases. December 14, 2021 - TND study linking administrative databases to assess VE against symptom cohort study done among covid hospitalized cases. December 14, 2021 - TND study linking administrative databases to assess VE against symptom cohort study done among covid hospitalized cases. December 14, 2021 - TND study linking administrative databases to assess VE against symptom cohort study done among covid hospitalized cases. December 14, 2021 - TND study linking administrative databases to assess VE against symptom cohort study done among covid hospitalized cases. December 14, 2021 - TND study linking administrative databases to assess VE against symptom cohort study done among covid hospitalized cases. December 14, 2021 - TND study linking administrative databases to assess VE against symptom cohort study lincided and study linking administrative databases to assess VE ag		ligh Care Hospital High Ca	ligh Care	Hospital H	ital High Care		Hospital Hig							
April 21, 2022	n or ICU Admission or ICU	or ICU Admission or ICU	or ICU A	Admission	sion or ICU	r ICU Admi:	Admission o							
April 21, 2022														
Ontering	tomatic disease, with a	s VE against symptomatic	ss VE against							1	18+ year olds	France		160
Bits reduced year content of the part of				alized cases.		done among	cohort study o	Januar	mRNA-1273	Omicron			(April 21, 2022)	
	Protection	Delta ^a		Protection										
		Risk reduction ^c against	Risk reduc	1-OR×HR	tion ^c against	Risk reduc								
	ymptomatic	Infaction among symptomatic		Protection(or% CI)	among symptomatic	Symptomatic Infection								
Vaccinated (eff. unwaccinated without prior infection withdexes) Uniform - Jardeys		ORIGER CD	OR ^d (95%CI)	Protection(95%CI)		OR4 (95%CI)								
De. oday - podays De. primorth - promise De. primorth	(5 MCI)	nk (95 leci)		e)		ccinated without pri	Vaccinated (ref.: unva							
D21 membh - 2 membh D32 membh - 2 membh D32 membh - 2 membh														
D2 x months - months D2 x							 							
D2.1 months - months D2.2 months - months D2.2 months - months D2.2 months - months D2.2 m														
D2.5 months														
D216 months														
DB: 6 days - 14 days														
DB:15days-3odiys 0.33 (0.32t00.33) 0.18 (0.14t00.22) 0.94 (0.93t00.95) 0.04 (0.04t00.05) 0.16 (0.11t00.10)							 							
DB: months	1 2 1 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2	71 7 7 7												
DB 3 months	1 11111	11 1 1												
Naturally-acquired and hybrid immunity/tef.:unwaccinated without prior infection evidence) Uwaccinated: NA 0.49 (0.48100.59) 0.45 (0.3910.0.6) 0.78 (0.7010.0.85) 0.11 (0.1110.1.2) 0.43(0.2210.0.0) 0.10 colors (0.2110.0.1.2) 0.43 (0.2100.0.0) 0.20 (0.3210.0.3.6) 0.78 (0.7010.0.85) 0.11 (0.1110.1.2) 0.43 (0.2210.0.0) 0.20 (0.3210.0.3.6) 0.54 (0.9310.0.9.6) 0.56 (0.9310.0.9.6) 0.56 (0.9310.0.0.2) 0.29 (0.3210.0.3.6) 0.94 (0.9310.9.9.6) 0.02 (0.0.210.0.0.2) 0.29 (0.3310.0.0.0.2) 0.29 (0.3310.0.0.2) 0.29 (0.2310.0.2) 0.29 (0.2310.0.2) 0.29 (0.2310.0.2) 0.29 (0.231														
Umaccinated: NA 0.49 (0.48to.50) 0.45 (0.30to.60) 0.78 (0.70to.85) 0.11 (0.11to.12) 0.43(0.22to.0 DB in CD IN A 0.39 (0.31to.34) 0.51 (0.3to.60) 0.89 (0.78to.88) 0.00 (0.08to.00) 0.56 (0.39to.34) 0.51 (0.3to.60) 0.89 (0.78to.88) 0.00 (0.08to.00) 0.56 (0.39to.34) 0.50 DB in CD IN A 0.39 (0.3to.34) 0.51 (0.3to.60) 0.89 (0.78to.68) 0.00 (0.08to.00) 0.56 (0.3to.60) 0.58 (0.3to.60) 0.	06t0 0.15) 0.99 (0.99 to 1.00)	17	17											
DB: NA 0.19 (0.1910-20) 0.29 (0.22100.36) 0.94 (0.93100.90) 0.02 (0.02100.02) 0.29 (0.13100. CI: confidence interval; COVID-19: coronavirus disease; D1: first vaccine dose; D2: second vaccine dose; D8: booster di NA: not applicable; O8: odds ratio; ref.: reference; RT-PCR: reverse-transcription PCR; SARS-CoV-2: severa eutre per septembre productive of Omicron): aboratory-confirmed (RT-PCR) SARS-CoV-2: severa eutre per septembre productive of Omicron): aboratory-confirmed (RT-PCR) SARS-CoV-2: severa eutre per septembre productive of Omicron): aboratory-confirmed (RT-PCR) SARS-CoV-2: severa eutre per septembre productive of Omicron): aboratory-confirmed (RT-PCR) SARS-CoV-2: severa eutre per septembre productive of Omicron): aboratory-confirmed (RT-PCR) SARS-CoV-2: severa eutre per septembre productive of Omicron): aboratory-confirmed (RT-PCR) SARS-CoV-2: severa eutre per severa p	22too.64) 0.95(0.93too.98)													
Cl: confidence Interval; COVID-19: coronavirus disease; D1: first vaccine dose; D2: second vaccine dose; D8: booster d NA: not applicable; OR: odds ratio; ref.: reference; RT-PCR: reverse-transcription PCR; SARS-CoV-2: severe acute res **Delta (respective Omicron): laboratory-confirmed (RT-PCR) SARS-CoV-2 infection with mutation with respective of Omicron) variant [14]. **Duration since receiving the COVID-19 vaccine dose in question, at presentation to the Screening centre. **Risk reductions are relative to symptoms attributable respectively to the Delta or the Omicron variant. **Odds ratios of symptomatic infections, according to the time elapsed since each COVID-19 vaccine dose reception or prior infection. **Hazard ratios of hospitalisations after symptomatic infections, according to the time elapsed since each COVID-19 vaccine dose reception or prior infection.														
**Delta (resportance Omicron): laboratory-confirmed (RT-PCR) SARS-CoV-2 infection with mutation screening indicative of Omicron variant [14]. **Duration since receiving the COVID-19 vaccine dose in question, at presentation to the screening centre. **Risk reductions are relative to symptoms attributable respectively to the Delta or the Omicron variant. **Odds symptomatic infections, according to the time elapsed since each COVID-19 vaccine dose reception or a prior infection. **Hazard radig to eithog after symptomatic infections, according to the time elapsed since each COVID-19 vaccine dose reception or a control of the prior infection.	13t0 0.44) 0.99 (0.99 to 1.00)					'	<u> </u>							
	on or according to evidence of	vith mutation screening indicative of Delta the screening centre. he Omicron variant. OVID-19 vaccine dose reception or accord ne elapsed since each COVID-19 vaccine d	vith mutation screen the screening centriche Omicron variant. (OVID-19 vaccine dos me elapsed since ea	ARS-CoV-2 infection won, at presentation to tively to the Delta or t elapsed since each Cos, according to the tin	onfirmed (RT-PCR) SA ccine dose in questi- attributable respect ccording to the time mptomatic infection	icron): laboratory-co il. ilng the COVID-19 va elative to symptoms omatic infections, a bitalisations after sy ce of prior infection.	Delta (respective Om Omicron) variant [14] Duration since receiv Risk reductions are re Odds ratios of sympt prior infection. Hazard ratios of hosp according to eviden.							





										Omicron ^a			Delta	
								Immune status: time since named	Hospital admission	ICU admission	Death	Hospital admission	ICU admission	Death
								vaccine dose ^b	HR ^c (95%CI)	HR ^c (95%CI)	HR ^c (95%CI)	HR ^c (95%CI)	HR° (95%CI)	HR ^c (95%CI)
								Vaccinated (ref.: unv	accinated without pr	rior infection evidend	:e)			
								D1: 0-28 days	0.99 (0.75 to 1.23)	1.09 (0.49 to 1.69)	1.09 (0.53 to 1.65)	0.66 (0.50 to 0.81)	0.43 (0.21t0 0.65)	0.93 (0.48t01.37)
								D2: 0-30 days	0.72 (0.50t0 0.95)	0.54 (0.06t01.02)	0.71 (0.14 to 1.29)	0.40 (0.23t00.57)	0.32 (0.04 to 0.60)	0.44 (0.01t00.87)
								D2: 1-2 months	0.40 (0.27 to 0.53)	0.32 (0.06 to 0.59)	0.38 (0.10 to 0.67)	0.41 (0.25 to 0.57)	0.52 (0.21t0 0.84)	0.14 (-0.13 to 0.42)
								D2: 2-3 months	0.56 (0.41t0 0.71)	0.22 (0.00 to 0.43)	0.12 (-0.05 to 0.29)	0.36 (0.25 to 0.47)	0.35 (0.16t0 0.54)	0.11 (-0.04 to 0.26)
								D2: 3-4 months	0.58 (0.48 to 0.68)	0.25 (0.09 to 0.42)		0.29 (0.23t00.35)	0.18 (0.10 to 0.26)	0.31 (0.12t0 0.49)
								D2:4-5 months	0.43 (0.36 to 0.49)	0.15 (0.07 to 0.24)	0.30 (0.14 to 0.45)	0.21 (0.17 to 0.24)	0.17 (0.12 to 0.23)	0.37 (0.20t0 0.53)
								D2:5-6 months	0.30 (0.24 to 0.35)		0.32 (0.15 to 0.48)		0.10 (0.07 to 0.13)	
								D2:>6 months	0.50 (0.43t00.56)					0.35 (0.25 to 0.44)
								DB: 1-7 days	0.35 (0.27 to 0.43)		0.29 (0.07 to 0.50)		0.06 (0.03 to 0.10)	
								DB: 8-14 days	0.28 (0.21t0 0.36)		0.14 (0.00t00.28)		0.07 (0.02 to 0.12)	
								DB: 15-30 days DB: 1-2 months	0.18 (0.14 to 0.22)		0.18 (0.08 to 0.28)			0.15 (0.02 to 0.29)
								DB: 1-2 months	0.16 (0.13 to 0.18) 0.18 (0.15 to 0.21)	0.06 (0.03 to 0.08) 0.08 (0.04 to 0.13)		0.14 (0.10 to 0.17) 0.10 (0.06 to 0.14)	0.13 (0.07 t0 0.19) 0.08 (0.00 t0 0.15)	0.16 (0.06t0 0.25)
													0.03	
								DB>3 months	0.14 (0.11t0 0.16)		0.13 (0.08 to 0.17)		(-0.03t00.09)	0.10 (0.01t0 0.19)
								Naturally-acquired o	r hybrid immunity ^a (r	ef.: unvaccinated wit	hout prior infection	evidence)		
								Unvaccinated: NA	0.45 (0.30 to 0.60)	0.14 (-0.05 to 0.33)	0.24 (-0.09t00.58)	0.43 (0.22t00.64)	0.54 (0.10t00.97)	1.06 (0.02 to 2.10)
								D1 or D2: NA	0.51 (0.36 to 0.66)	0.42 (0.12 to 0.72)	0.34 (0.07 to 0.61)	0.56 (0.34t00.77)	0.39 (0.08 to 0.71)	0.90 (0.17 to 1.62)
								DB: NA	0.29 (0.22 to 0.36)	0.16 (0.05 to 0.28)	0.19 (0.06 to 0.32)	0.29 (0.13 to 0.44)	0.13 (-0.05t0 0.30)	0.11 (-0.11 to 0.33)
15	59	Kirsebom et al (April 28, 2022)	England	General population	Omicron Delta	ChAdOx1 Comirnaty mRNA-1273 followed by ChAdOx1	September 13, 2021- February 17, 2022	TND study linki	ng adminsitra	tive databases	to assess VE a	against sympto	matic disease	
						booster								





Spins Spin								Age		Booster	Interval			
Deciding Character Chara								(years		Manufacture	r (days)	Controls Cases OR*	VE (95% CI)	
Dock 2" 19									Unvaccinate	d			Baseline	
### April 1 11.19 7710 (20) 223 133 134 144 145 145 145 145 145 145 145 145 14										n/a	175+	85175 89230 0.94)		
### April 12									Booster	Any***	0-1	11,879 7715 0.83)	23.3)	
### PATH										Anue	2.6	27430 21422 0.78)		
Bit 11022 14.34 Bit 1002 1003 100												0.42 (0.41-	58.2 (57.0 to	
BITTER22 1.6.4 1.6.77 6.640 2.77 6.4.5 BITTER22 1.6.6 1.7.5 2.7.5 2.7.5 2.7.5 BITTER22 1.6.6 1.7.5 2.7.5 2.7.5 2.7.5 BITTER22 1.6.6 1.7.5 2.7.5 2.7.5 2.7.5 BITTER22 1.6.6 2.7.5 2.7.5 2.7.5 2.7.5 BITTER23 1.6.6 2.7.5 2.7.5 2.7.5 2.7.5 BITTER24 1.6.7 2.7.5 2.7.5 2.7.5 2.7.5 BITTER25 1.6.7 2.7.5 2.7.5 2.7.5 BITTER25 1.6.7 2.7.5 2.7.5 2.7.5 BITTER25 1.6.7 2.7.5 BITTER25 1.6.7 2.7.5 2.7.5 BITTER25 2.6.7 2.7.5 2.7.5 BITTER25 2.7.5 2.7.5 2.7.5 BITTER25 2.7.5 2.7.5 2.7.5										BNT162b2	7-13			
BITTERD 35-66 07500 07500 0450 0550 0								40-64		BNT162b2	14-34	86719 66406 0.37)	64.5)	
BRT140202 75-10-10 2200 2307 95-10 247 67 07 0 20 20 20 20 20 20 20 20 20 20 20 20 2								1		BNT162b2	35-69	87592 90787 0.44) 0.43 (0.42-	57.3 (56.4 to 58.2)	
BHT16202 155- 2784 2776 2786 2786 2787 278										DNIT400b0	70.104	0.54 (0.52-	46.4 (45.0 to	
Childry 15 15 15 15 15 15 15 1												0.69 (0.66-	30.6 (26.8 to	
Chadori-15 1-34 710 40 0.580 74.6 73 70 40 0.580 74.6 74 75 75 75 75 75 75 75										BNT162b2	105+	2758 4278 0.73)	34.3)	
CANADYLIS 14-34 100 150 041) 041 0										ChAdOx1-S	7-13	70 40 0.59)	74.6)	
ChadOx1-6 3-5-69 218 218 207 30 415 50 50 615 50 615 615 615 615 615 615 615 615 615 615										ChAdOx1-S	14-34	193 159 0.48 (0.38-	51.7 (38.9 to 61.8)	
ChadOx1-3 70-104 69 07 05 80 043 40 61 68 9 0 ChadOx1-3 105+ 10 14 50 50 10 14 50 80 10 1										100011000000000000000000000000000000000	104010000	0.47 (0.38-	53.0 (42.6 to	
ChadOx1-S 70-104 69 07 03.03 05 50 07 03.03 07 372 (44.1 to ChadOx1-S 105 40 10 03.00) 1 03.00 07 03.0												0.59 (0.43-	40.8 (18.6 to	
Chadoxis 105+ 105+ 10 14 14.4 72e)										ChAdOx1-S	70-104			
Dose 2" n/a 175+ 4460 3053 0.31 (0.73- 19.5 (117-to 20.6)										ChAdOx1-S	105+			
Dose 2" n/a 175 4466 3053 0.88 26.6 Bootster Any*** 0.1 428 110 0.85 5.48 44.8 to Any*** 2.6 110 35 5.5 42.8 16.0 to BNT162b2 14.34 14.911 3010 0.41 30.8 30.8 30.8 30.8 30.8 BNT162b2 14.34 14.911 3010 0.41 30.8 30.8 30.8 30.8 30.8 30.8 30.8 BNT162b2 14.34 14.911 3010 0.41 30.9 30.8 30.8 30.8 30.8 30.8 BNT162b2 14.34 14.911 3010 0.41 30.9 30.8 30.8 30.8 BNT162b2 70.104 14.210 1837 0.65 63.5 44.5 BNT162b2 105* 1970 2780 777 77.7 23.1 (15.1 to ChAdOX1-S 7.13 23 60.83 60.83 60.83 ChAdOX1-S 14.34 55 32.0 (10.8 to ChAdOX1-S 14.34 55 32.0 (10.8 to ChAdOX1-S 14.34 55 32.0 (10.8 to ChAdOX1-S 14.34 56 32.0 (10.8 to ChAdOX1-S 14.34 50.8 to ChAdOX1-S 15.0 to ChADOX1-S 15.									Unvaccinate	d				
Any.*** 0-1 428 110 (a.67) 468) 468) Any.*** 2-6 1140 370 (a.84) 503 (a.61) 60 (a.67)									Dose 2**	n/a	175+	4466 3053 0.88)	26.6)	
Any*** 2-8 1140 370 (0.81) (3.8) (0.81) (3.9) (3.9) (3.8) (3									Booster	Anyses	0.1			
BNT162b2 7-13 1,883 430,48) 63,80 8) 68,75 70 72 72 72 72 72 72 73 73,60 73 70 70 70 70 70 70 70 70 70 70 70 70 70												0.71 (0.61-	28.6 (16.0 to	
BNT162b2 7-13 1,838 433 (A8) 63.8) BNT162b2 14-34 1,4311 3010 0.34 71.2) BNT162b2 35-69 36300 25240 0.49 71.2) BNT162b2 70-104 1420 18317 0.65 55 44.5) BNT162b2 70-104 1420 18317 0.65 55 44.5) BNT162b2 70-104 1420 18317 0.65 55 44.5) BNT162b2 105- 1970 2780 0.85 7.5) ChadOx1-S 7-13 2 0.34 (0.14 0.5 10.5 10.5 10.5 10.5 10.5 10.5 10.5										Any***	2-6			
BNT162b2 14-34 14311 3010 0.34) 71.2) BNT162b2 35-69 36300 25240 0.49) 67.5 to BNT162b2 70-104 14210 18317 0.65 44.5) BNT162b2 105+ 1970 2789 0.85) 44.5) ChAgOx1-S 7-13 23 6 0.34 (0.14 - 66 116 6 to ChAgOx1-S 14-34 53 32 0.79) 70.4 ChAgOx1-S 35-69 88 81 0.78) 68.3 to ChAgOx1-S 35-69 88 81 0.78) 68.3 to ChAgOx1-S 70-104 16 40 2.32) 70.4 to ChAgOx1-S 70-104 16 40 2.32) 70.4 to ChAgOx1-S 70-104 16 40 2.32) 70.7 (-7 27.2 (131.8 to 30.1) 70.4 to ChAgOx1-S 70-104 16 40 2.32) 70.7 (-7 27.2 (131.8 to 30.1) 70.4 to 70								+		BNT162b2	7-13	1,883 433 0.48)	63.8)	
BNT162b2 35-69 36300 25240 049 57.5) BNT162b2 70-104 14210 183170 85) 44.59 BNT162b2 105+ 1970 2789 085) 30.50 ChAdOX1-S 7-13 23 80.33 (0.14-66.1 (16.6 to 80.3) ChAdOX1-S 14-34 55 32.079) 70.4) ChAdOX1-S 35-69 88 810.78) 60.2 ChAdOX1-S 70-104 16 40.232) 30.10 ChAdOX1-S 105+ 3 5 988 0.23 ChAdOX1-S 105+ 3 5 988 0.23 N too low								65		BNT162b2	14-34	14311 3010 0.34)	68.5 (65.7 to 71.2)	
BNT192b2 70-104 14210 18317 0.85 40.1 (35.2 to 4.5) BNT192b2 105+ 1970 2780 0.85 30.5 (0.14-16.6 to 16.6 to 1												0.46 (0.42-	54.1 (50.5 to	
BNT162b2 105+ 1970 2780 0.85										1000175013001301501		0.6 (0.55-	40.1 (35.2 to	
BNT162b2 105+ 1970 2786 0.85 30.5 30.5 ChAdOx1-S 7.13 23 8 0.83 80.83										BNT162b2	70-104	14210 18317 0.65)	44.5)	
ChAdOx1-S 7-13 23 8 (0.83) 86.3) 86.3) 86.3) (1.40 to 1.40 to										BNT162b2	105+	1970 2789 0.85)	30.5)	
ChAdOx1-S 14-34 53 30,48 (0.3-151.6 (20.8 to 7.4) (20.8 to										ChAdOx1-S	7-13	0.34 (0.14- 23 8 0.83)		
ChAdOx1-S 35-69 88 81 0.56 (0.4 44.5 (22.4 to 80.2) ChAdOx1-S 70-104 16 40 2.32) 30.1) ChAdOx1-S 105+ 3 5 4.28) 105+ 3 105+									Ī		1	0.48 (0.3-	51.6 (20.8 to	
Chadox1-S 35-69 88 81 0.78) 60.27 Chadox1-S 70-104 16 40 2.32) 30.1) 18 to Chadox1-S 105+ 3 5 4.28) N too low												0.56 (0.4-	44.5 (22.4 to	
ChadOx1-S 70-104 16 40 2.32 30.1										ChAdOx1-S	35-69	88 81 0.78)	60.2)	
										ChAdOx1-S	70-104	16 40 2.32)	30.1)	
										ChAdOx1-S	105+	0.98 (0.23 5 (4.28)	N too low	
158 Sheikh et al Scotland General population Omicron ChAdOx1 November 1- TND study linking adminsitrative databases to assess VE against symptom											100		1	
158 Sheikh et al Scotland General population Omicron ChAdOx1 November 1- TND study linking adminsitrative databases to assess VE against symptom														
158 Sheikh et al Scotland General population Omicron ChAdOx1 November 1- TND study linking adminsitrative databases to assess VE against symptom														
	158	Sheikh et al	Scotland	General population	Omicron	ChAdOx1	November 1-	TND	study li	inking ad	minsi	trative database	s to assess \	VE against symptomatic
(April 22, 2022) Comirnaty December 19, 2021		(April 22, 2022)				Comirnaty	December 19, 2021							
mRNA-1273		, , , , , , , , , , , , , , , , , , , ,					,							

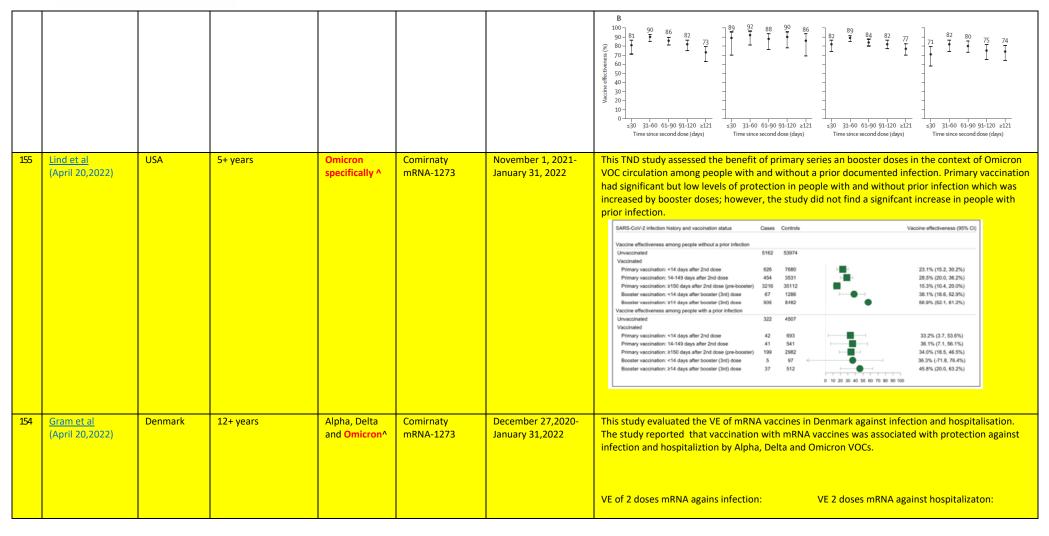




							S-gene-negative infections S-gene-positive infections
							Tested, n Positive, n Relative vaccine Tested, n Positive, n Relative vaccine
							effectiveness, % effectiveness, %
							(95% CI) (95% CI)
							16-49 years Unvaccinated 10 302 1003 22% (14 to 29) 14 583 5284 -98% (-109 to -87)
							Unvaccinated 10 302 1003 22% (14 to 29) 14 583 5284 -98% (-109 to -87) First dose
							0-27 days 550 36 47% (24 to 63) 676 162 -24% (-50 to -3)
							228 days 6570 581 30% (21 to 38) 8339 2350 -39% (-49 to -30)
							Second dose
							0-13 days 732 46 58% (42 to 70) 805 119 31% (16 to 44)
							14-69 days 4248 256 53% (46 to 60) 4258 266 73% (69 to 76) 70-104 days 12 581 814 33% (26 to 40) 13 559 1792 50% (46 to 53)
							70-104 days 12581 814 33% (26 to 40) 13559 1792 50% (46 to 53) 105-139 days 29209 3503 15% (9 to 21) 31963 6257 32% (29 to 36)
							140-174 days 1496 1824 3 (5-1611) 17991 4829 9% (40-13)
							≥175 days 13 183 1435 Reference 15 462 3714 Reference
							Third dose
							0-6 days 3773 515 26% (16 to 34) 4003 745 33% (27 to 39)
							7-13 days 2185 143 62% (54to 68) 2155 113 84% (80 to 87) 214 days 12887 783 56% (51 to 60) 12798 694 83% (81 to 84)
							250years 100/ 763 30% (31000) 12/36 034 63% (61004) 250years
							Unvacinated 716 48 33% (7 to 52) 1158 490 -45% (-65 to -28)
							First dose
							0-27 days 27 4 0 (-230 to 70) 36 13 -16% (-134 to 42)
							28 days 256 13 48% (7 to 72) 343 100 10% (-15 to 30)
							Second dose 0-13 days 23 1 62% (-267 to 95) 23 1 90% (27 to 99)
							0-13 days 23 1 62% (-207 to 95) 23 1 90% (77 to 99) 14-69 days 120 9 5% (-98 to 54) 131 20 62% (38 to 77)
							70-104 days 128 12 88 (-76 to 52) 149 33 40% (10 to 60)
							105-139 days 463 17 35% (-10 to 62) 634 188 20% (4 to 33)
							140-174 days 5513 265 4% (-13 to 19) 8205 2957 4% (-3 to 10)
							=175 days 8007 799 Reference 10 856 3648 Reference
							Third dose 0-6 days 3522 420 0 (-15 to 13) 4352 1250 20% (13 to 26)
							7-13 days 3006 180 545 (46 to 52) 3146 320 77% (47 to 80)
							2.14 days 17.572 1045 57% (52 to 62) 17.504 977 88% (86 to 89)
455	Communication City	D 11	40	0	ChAdOx1	January 1-March 7,	THE state of the s
157	<u>Cerqueria-Silva et</u>	Brazil,	18+ year olds	Omicron	I (nAd()v1	I January 1-March 7	
	<u>al</u>						TND study linking administrative databases.
	al	Scotland			Comirnaty	2022	Symptomatic SARS-CoV-2 Infection Severe COVID-19
		Scotland			Comirnaty		Symptomatic SARS-CoV-2 Infection Severe COVID-19
	(April 14, 2022)	Scotland					
		Scotland			Comirnaty		Symptomatic SARS-CoV-2 Infection Severe COVID-19
		Scotland			Comirnaty		Symptomatic SARS-CoV-2 Infection Severe COVID-19
		Scotland			Comirnaty		Symptomatic SARS-CoV-2 Infection Severe COVID-19
		Scotland			Comirnaty		Symptomatic SARS-CoV-2 Infection Severe COVID-19
		Scotland			Comirnaty		Symptomatic SARS-CoV-2 Infection Severe COVID-19 80 80 60
		Scotland			Comirnaty		Symptomatic SARS-CoV-2 Infection Severe COVID-19 80 80 60
		Scotland			Comirnaty		Symptomatic SARS-CoV-2 Infection Severe COVID-19
		Scotland			Comirnaty		Symptomatic SARS-CoV-2 Infection Severe COVID-19 80 80 60
		Scotland			Comirnaty		Symptomatic SARS-CoV-2 Infection Severe COVID-19 80 80 40
		Scotland			Comirnaty		Symptomatic SARS-CoV-2 Infection Severe COVID-19 100 80 80 40
		Scotland			Comirnaty		Symptomatic SARS-CoV-2 Infection Severe COVID-19 80 80 40
		Scotland			Comirnaty		Symptomatic SARS-CoV-2 Infection Severe COVID-19 80 80 40
		Scotland			Comirnaty		Symptomatic SARS-CoV-2 Infection Severe COVID-19 80 40 40 0
		Scotland			Comirnaty		Symptomatic SARS-CoV-2 Infection Severe COVID-19 80 40 20 0-1 2-4 5-8 9-12 2-13 0-1 2-4 5-8 9-12 2-13
		Scotland			Comirnaty		Symptomatic SARS-CoV-2 Infection Severe COVID-19
		Scotland			Comirnaty		Symptomatic SARS-CoV-2 Infection Severe COVID-19 80 40 20 0-1 2-4 5-8 9-12 2-13 0-1 2-4 5-8 9-12 2-13
		Scotland			Comirnaty		Symptomatic SARS-CoV-2 Infection Severe COVID-19
156	(April 14, 2022)		Patients with		Comirnaty mRNA-1273	2022	Symptomatic SARS-CoV-2 Infection Severe COVID-19 80 40 0-1 2-4 5-8 9-12 ≥ 13 Weeks since booster dose Country Brazil Severe COVID-19 Severe COVID-19 100 80 40 40 40 50 80 80 80 80 80 80 80 80 8
156	(April 14, 2022) Widdifield et al	Canada	Patients with	Alpha, Delta	Comirnaty mRNA-1273	March 1-November	Symptomatic SARS-CoV-2 Infection Severe COVID-19
156	(April 14, 2022)		Patients with rheumatoid arthritis,		Comirnaty mRNA-1273	2022	Symptomatic SARS-CoV-2 Infection Severe COVID-19 80 40 0-1 2-4 5-8 9-12 ≥ 13 Weeks since booster dose Country Brazil Severe COVID-19 Severe COVID-19 100 80 40 40 40 50 80 80 80 80 80 80 80 80 8
156	(April 14, 2022) Widdifield et al		rheumatoid arthritis,		Comirnaty mRNA-1273	March 1-November	Symptomatic SARS-CoV-2 Infection Severe COVID-19 80 40 0-1 2-4 5-8 9-12 ≥ 13 Weeks since booster dose Country Brazil Severe COVID-19 Severe COVID-19 100 80 40 40 40 50 80 80 80 80 80 80 80 80 8
156	(April 14, 2022) Widdifield et al		rheumatoid arthritis, ankylosing		Comirnaty mRNA-1273	March 1-November	Symptomatic SARS-CoV-2 Infection Severe COVID-19 80 40 0-1 2-4 5-8 9-12 ≥ 13 Weeks since booster dose Country Brazil Severe COVID-19 Severe COVID-19 100 80 40 40 40 50 80 80 80 80 80 80 80 80 8
156	(April 14, 2022) Widdifield et al		rheumatoid arthritis, ankylosing spondylitis, psoriasis,		Comirnaty mRNA-1273	March 1-November	Symptomatic SARS-CoV-2 Infection Severe COVID-19 80 40 0-1 2-4 5-8 9-12 ≥ 13 Weeks since booster dose Country Brazil Severe COVID-19 Severe COVID-19 100 80 40 40 40 50 80 80 80 80 80 80 80 80 8
156	(April 14, 2022) Widdifield et al		rheumatoid arthritis, ankylosing		Comirnaty mRNA-1273	March 1-November	Symptomatic SARS-CoV-2 Infection Severe COVID-19 80 40 0-1 2-4 5-8 9-12 ≥ 13 Weeks since booster dose Country Brazil Severe COVID-19 Severe COVID-19 100 80 40 40 40 50 80 80 80 80 80 80 80 80 8
156	(April 14, 2022) Widdifield et al		rheumatoid arthritis, ankylosing spondylitis, psoriasis,		Comirnaty mRNA-1273	March 1-November	Symptomatic SARS-CoV-2 Infection Severe COVID-19 80 40 0-1 2-4 5-8 9-12 ≥ 13 Weeks since booster dose Country Brazil Severe COVID-19 Severe COVID-19 100 80 40 40 40 50 80 80 80 80 80 80 80 80 8

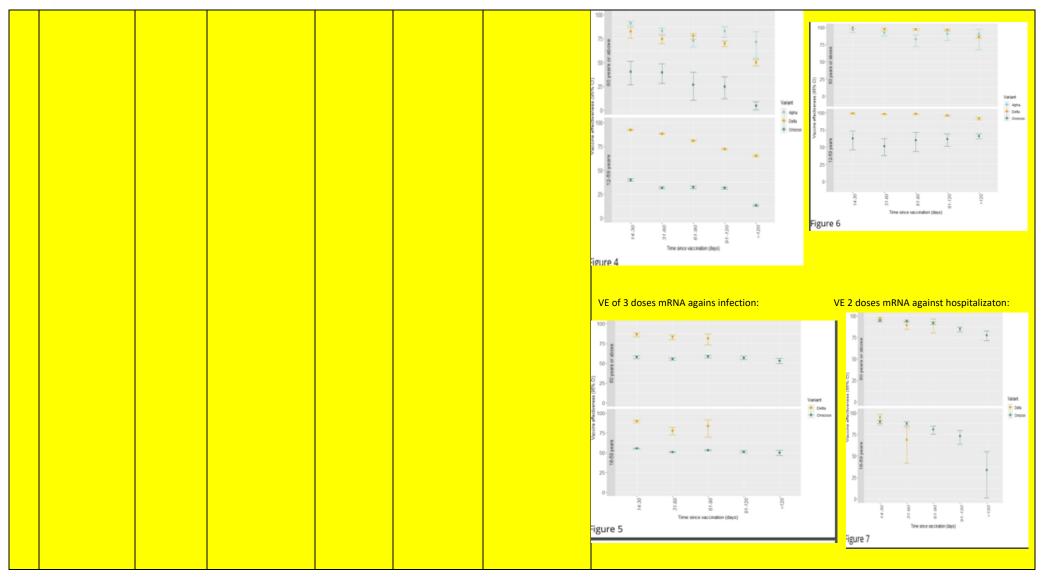












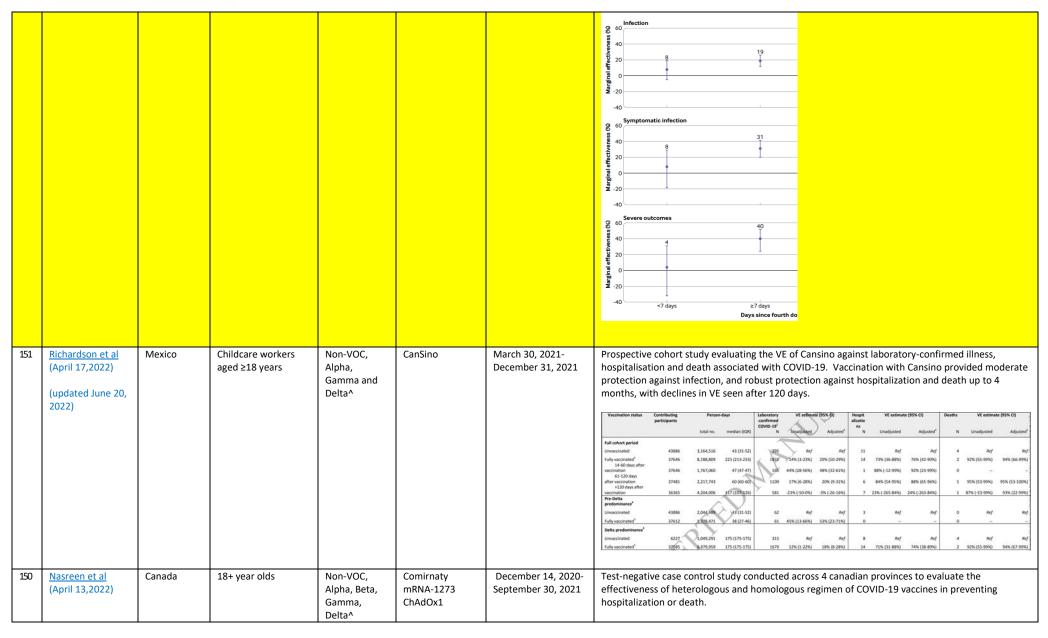




153	Voko et al (April 18,2022) (updated July 22, 2022)	Hungary	18-100 years	Delta^	Comirnaty, mRNA-1273, ChAdOx1, Ad26.COV2.S, Sputnik, Sinopharm	March 4, 2020- December 31, 2021	This study assessed the effectiveness and duration of protection of six different types of vaccines with combinations as primary or booster vaccines against COVID-19 infection, hospitalization and death during a period of Delta variant predominance. (left figure 16-64; right figure 65-100) A 100
152	Grewal et al (April 18,2022) (updated June 1, 2022) (final publication July 6, 2022)	Canada	LTC residents aged ≥60 years	Omicron specifically [^]	Comirnaty, mRNA-1273	December 30, 2021- April 27, 2022	This test-negative case control study estimated the marginal effectiveness of a fourth dose of COVID-19 vaccines relative to individuals with a third dose and or unvaccinated.

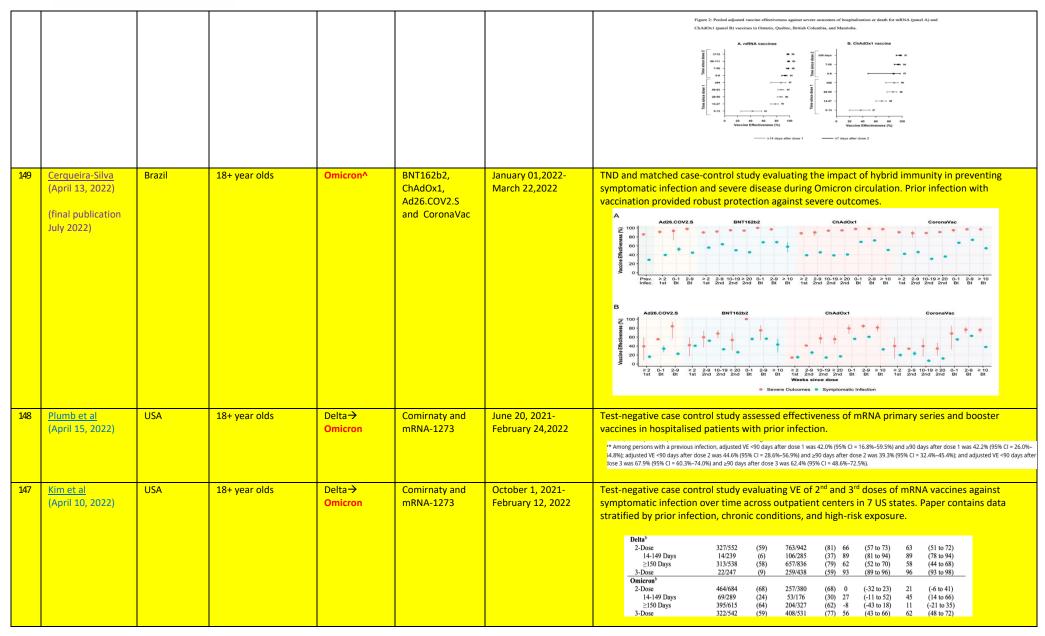






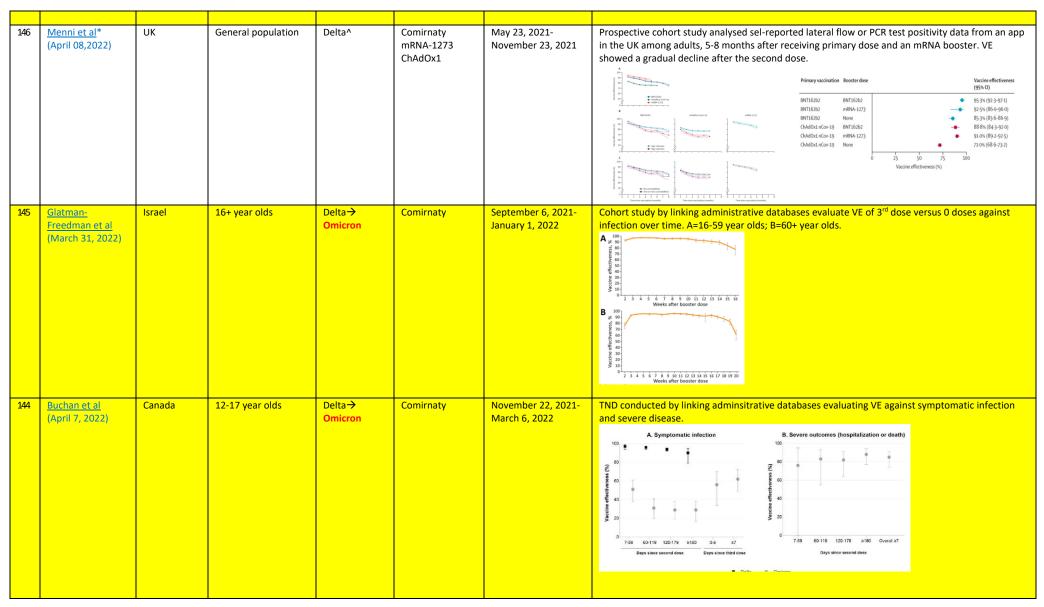
















143	Fabiani et al (April 6, 2022)	Italy	60+ and other priority groups (e.g. hcws)	Delta	Comirnaty mRNA-1273 ChAdOx1 Ad26.COV2.S	July 19, 2021- December 12, 2021	Cohort study among vaccine recipients comparing time intervals to day 4-10 post dose 1. Paper contains data stratified by priority groups. Any SARS-CoV-2 Infection* Severe COVID-19 to
142	Bansal et al (April 6, 2022)	Qatar	General population	Alpha, Beta, Delta, Omicron (but no omicron specific estimate)	Comirnaty mRNA-1273 ChAdOx1 (1.6% of all vaccinated)	January 1, 2021- February 20, 2022	Matched case-control among all cases in Qatar, looking at progression to ICU. VE 89% (95% CI, 85 to 92) between 0-4 months post the second dose. VE 91%; 95% CI 84 to 95) between 4-6 months after the second dose; VE 90%; 95% CI 84 to 94)) at 6 to 9 months after the second dose.
141	Florentino et al (April 5, 2022) (updated to final publication on August 8, 2022)	Brazil, Scotland	12-17 year olds	Delta-> Omicron	Comirnaty	Brazil: September 2, 2021-April 19, 2022 Scotland: August 6, 2021- April 19, 2022	TND study against symptomatic disease: VE against symptomatic disease: Delta-dominant period in Brazil Delta-dominant period in Scotland Delta-dominant





							Vaccine effectiveness (%; 95% CI) Number of tests from unvaccinated individuals Time after first dose O-6 days 20-6 (-152-2 to 75-0) 7-13 days 62-4 (-22-2 to 88-5) 214 days 56-3 (45-9 to 64-6) Time after second dose O-13 days 65-0 (37-2 to 80-5) 14-27 days 75-6 (58-1 to 85-8) 28-41 days 82-8 (72-1 to 89-4) 42-55 days 84-2 (76-3 to 89-5) 56-69 days 83-7 (76-0 to 88-9) 70-83 days 82-0 (72-6 to 88-2) 84-97 days 86-4 (75-2 to 92-6) 298 days 82-7 (68-8 to 90-4)
140	Bar-On et al (April 5, 2022)	Israel	60+ year olds	Omicron	Comirnaty	January 10-March 2, 2022	Relative VE comparing 4 th to 3 rd dose. 7.5 7.0 6.5 6.0 5.5 90
139	Perumal et al (April 1, 2022)	Germany	12+ year olds	Delta, Omicron	Comirnaty mRNA-1273	November 8, 2021- February 13, 2022	Analysis of surveillance data with comparison to aggregate vaccination data to calculate the VE against symptomatic disease, hospitalization, and severe disease. (Note unable to adjust for many confounders).

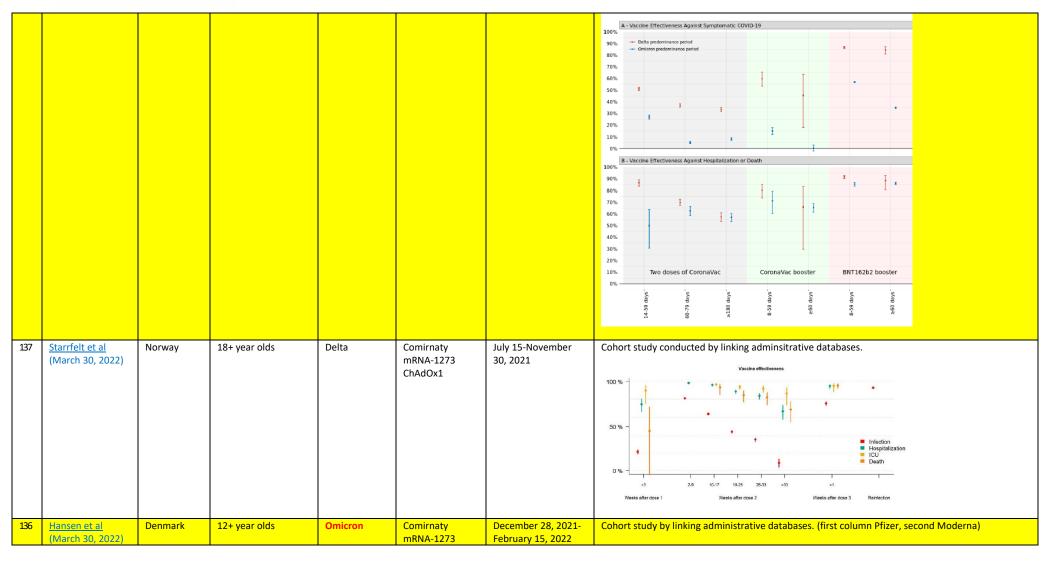




							Table 3: Effectivenes	s of booste	r vaccinatio	n against	symptomatic	SARS-Co	V-2 infection	and CO	OVID-	
							19-associated hospitalizations and severe illness during dominant circulation of the Omicron variant									
							in Germany, CW52/2021-06/2022, by age group and time interval.									
							≥18 vears									
								- -	L2-17 years		All		8-59 years		≥60 years	
								(Cases)	VE (95% C	(Cases	VE (95% CI	(Cases)	VE (95% CI)	(Cases)	VE (95% CI)	
							Symptomatic infection	(cases)		(Cases	1	(Cases)		(cases)		
							Unvaccinated	46,544	Re	f. 166,565	5 Ref	147,877	Ref.	18,688	Ref.	
							Boosted*	2,565	88-3 (86-2-90-2	156,215	5 69-7 (65-2-73-6	131,523	67-4 (62-3-71-8)	26,959	81-6 (77-2-85-2)	
							Boosted, by time interva	_								
							<4 weeks				1 78-7 (75-8-81-3				87-8 (86-0-89-4)	
							4 to <8 weeks	871		_	8 65-9 (62-1-69-4			_	81-3 (79-3-83-2)	
							8 to <12 weeks 12 to <16 weeks		N N		5 56-7 (50-0-62-5	29,713		_	76-4 (73-4-79-0)	
							Hospitalization		N		N.	-	NC.	2,267	75-0 (69-7-79-5)	
							Unvaccinated	222	Re	f. 5,325	5 Ref	2,404	Ref.	2,921	Ref.	
							Boosted*	9	90-5 (86-4-93-6	1,340	94-4 (92-6-95-8	617	89-9 (86-9-92-3)	905	95-9 (94-6-97-0)	
							Boosted by time interval									
							<4 weeks	_	91-4 (85-2-95-6	_	96-4 (94-9-97-6	_	93-7 (92-3-95-0)	_	97-7 (97-0-98-3)	
							4 to <8 weeks	3	83-9 (66-2-93-9		94-8 (93-0-96-1	_	88-6 (86-5-90-5)	_	96-7 (95-9-97-4)	
							8 to <12 weeks 12 to <16 weeks	-	N N	_	91-4 (88-0-94-1	158	77-1 (71-4-82-0) NC	_	94-3 (92-8-95-5)	
							Severe illness		N		N.		NC.	182	85-6 (81-3-89-1)	
							Unvaccinated	5	Re	f. 1,535	5 Ref	. 289	Ref.	1.246	Ref.	
							Boosted*	0		_	97-5 (96-8-98-2	24	96-2 (92-2-98-4)	220	97-7 (97-0-98-2)	
							Boosted by time interval									
							<4 weeks		N	с -	- NO	-	NC	_	98-8 (98-2-99-2)	
							4 to <8 weeks		N	-	- No	-	NC	_	98-1 (97-4-98-6)	
							8 to <12 weeks		N	_	- NO	-	NC		97-3 (96-0-98-2)	
							12 to <16 weeks	A	N N	٠ .	-] N	1	NC.	62	87-9 (83-1-91-6)	
120	Donzoni et el	Drozil	10 Lygar olds	Dolto	Coronovas	Contombor C 2024	TND cturely list	deina -	dunin cit	rotiv	a datala a	505 N	loto bes	cto-	doso \/F	is a valative VE (sampaged to
138	Ranzani et al	Brazil	18+ year olds	Delta,	Coronavac	September 6, 2021-										is a relative VE (compared to
	(April 1, 2022)			Omicron	Comirnaty	March 10, 2022	primary serie	s recip	oients) v	vhile	primary	serie	s VE is co	omp	ared to u	ivaccianted.











					<u> </u>		a) ech)
							Days since
							Not vaccinated (ref) (ref) (ref) Protection against 14-30 37-9 (34-54-22) 37-9 (35-5-98-3) 37-9 (34-5-22) 37-9 (35-5-98-3) 37-9 (34-5-22) 37-9 (35-5-98-3) 37-9 (34-5-22) 37-9 (35-5-98-3) 37-9 (34-5-22) 37-9 (35-5-98-3) 37-9 (34-5-22) 37-9 (35-5-98-3) 37-9 (34-5-22) 37-9 (35-5-98-3) 37-9 (34-5-22) 37-9 (35-5-98-3) 37-9 (34-5-22) 37-9 (35-5-98-3) 37-9 (34-5-22) 37-9 (35-5-98-3) 37-9 (34-5-22) 37-9 (35-5-98-3) 37-9 (34-5-22) 37-9 (35-5-98-3) 37-9 (34-5-22) 37-9 (35-5-98-3) 37-9 (34-5-22) 37-9 (35-5-98-3) 37-9 (34-5-22) 37-9 (35-5-98-3) 37-9 (34-5-22) 37-9 (35-5-98-3) 37-9 (35-5-98
							Protection against infection infection after 2 doses 91-120 25.82(28.82.96) 27.4 (26.3; 28.4) 26.6 (25.3; 27.9) 27.4 (26.3; 28.9) 27.4 (
							121+ 132 (12.3; 14.2) 9.8 (9.2; 10.4)
							Not vaccinated (ref)
							hospitalisation after 2 doses 91-120 42.5 (26.9; 54.9)
							121+ 51.6 (47.2; 55.6)
							Protection against 31-60 47.7 (47.0-88.3) 47.9 (47.4-83.8) 4.50 45.5 (44.9-6.2) 41.0 (40.5-41.5)
							infection after 3 61-90 43.5 (42.2; 44.7) 41.0 (40.3; 41.7) 41.0 (
							Not vaccinated (ref) (ref)
							Protection against 31-60 87.7 (83.3 93.7) 88.5 (87.4 99.6) 85.5 (87.4 99.6) 87.7 (83.3 93.7) 88.5 (87.4 99.6) 87.7 (83.3 93.7
							after 3 doves 91-120 87.6 (0+7.3 70.4) 6-43 (05.1.5 61.3) 6-43 (05.1.5 61.3) 121+ 77.3 (65.1.5 66.1) 78.0 (75.5 81.3)
135	<u>Price et al</u>	USA	5-18 year olds	Delta→	Comirnaty	July 1, 2021-February	TND study at 31 hospitals. Vaccinated Case Vaccinated Control Vaccine Effectiveness
	(March 30, 2022)			Omicron		17, 2022	Subgroup Patients Patients (95% CI) no. of patients, plotal no. (%) %
							Adolescents 12-18 yr of age Age group
							12-15 yr 63/543 (12) 313/828 (38) — 83 (77 to 88) 16-18 yr 59/375 (16) 229/529 (43) — 82 (74 to 88)
							Delta-predominant period 33/684 (5) 442/1161 (8) ■ 92 (89 to 95) 2-22 wk since vaccination 25/676 (4) 372/1091 (34) ■ 93 (89 to 95)
							23-44 wk since vaccination 6/657 (1) 60/779 (8) —■ 92 (80 to 97) Omicron-predominant period 89/234 (38) 100/196 (51) ■ 40 (9 to 60)
							2 - 22 wk since vaccination 35/180 (19) 39/135 (29) 43 (-1 to 68) 23-44 wk since vaccination 52/197 (26) 59/155 (38) 38 (-3 to 62)
							Children 5-11 yr of age Omicron-predominant period 20/267 (7) 50/270 (19)
							-25 0 25 50 75 100
124	Manaki ak al	Namen	12.17	Delta	Carainastu	August 24, 2024	Cohont study of 12.17 year olds and ratios VE assists infection based on limited administrative
134	Veneti et al (March 25, 2022)	Norway	12-17 year olds	Delta→ Omicron	Comirnaty	August 24, 2021- January 16, 2022	Cohort study of 12-17 year olds evaluating VE against infection based on linking adminiistrative databases.
	(IVIdICII 23, 2022)			Officion		January 10, 2022	Age () 12-15 years () 16-17 years
							b) Delta infections,
							25 August 2021 to 16 January 2022 26 November 2021 to 16 January 2022
							100
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							Days after dose 1 Days after dose 2 Days after dose 1 Days after dose 2
							No. of the state o
133	Wang et al	LISΔ	General nonulation	Delta→	Comirnaty	I October 1 2021-	TND study at Cleveland Clinic evaluating risk against intection (ton table, note this can be
133	Wang et al (March 25, 2022)	USA	General population	Delta -> Omicron	Comirnaty mRNA-1273	October 1, 2021-	TND study at Cleveland Clinic evaluating risk against infection (top table, note this can be converted to VE by subtracting the OR from 1) and, death (bottom table, not this is among cases
133	Wang et al (March 25, 2022)	USA	General population	Delta→ Omicron	Comirnaty mRNA-1273	October 1, 2021- January 31, 2022	TND study at Cleveland Clinic evaluating risk against infection (top table, note this can be converted to VE by subtracting the OR from 1) and death (bottom table, not this is among cases only and thus is VE against progression of infection to death).

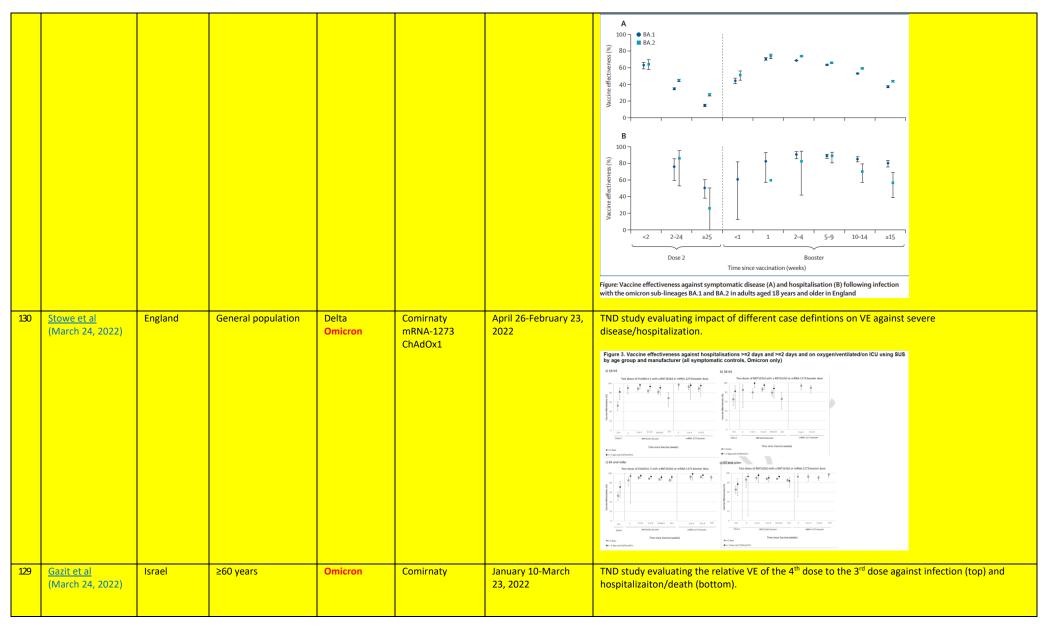




							Patients
132	Ng et al (March 24, 2022)	Singapore	Contacts of cases	Delta	Comirnaty mRNA-1273	March 1-August 31, 2021	Cohort study looking at transmission in households of cases. 1.2
131	Kirsebom et al (March 24, 2022) (updated to final publication May 24, 2022)	England	General population	Omicron (BA.1 vs BA.2)	Comirnaty mRNA-1273 ChAdOx1	January 17-February 17, 2022	TND study comparing VE against symptomatic disease with BA.1 vs BA.2

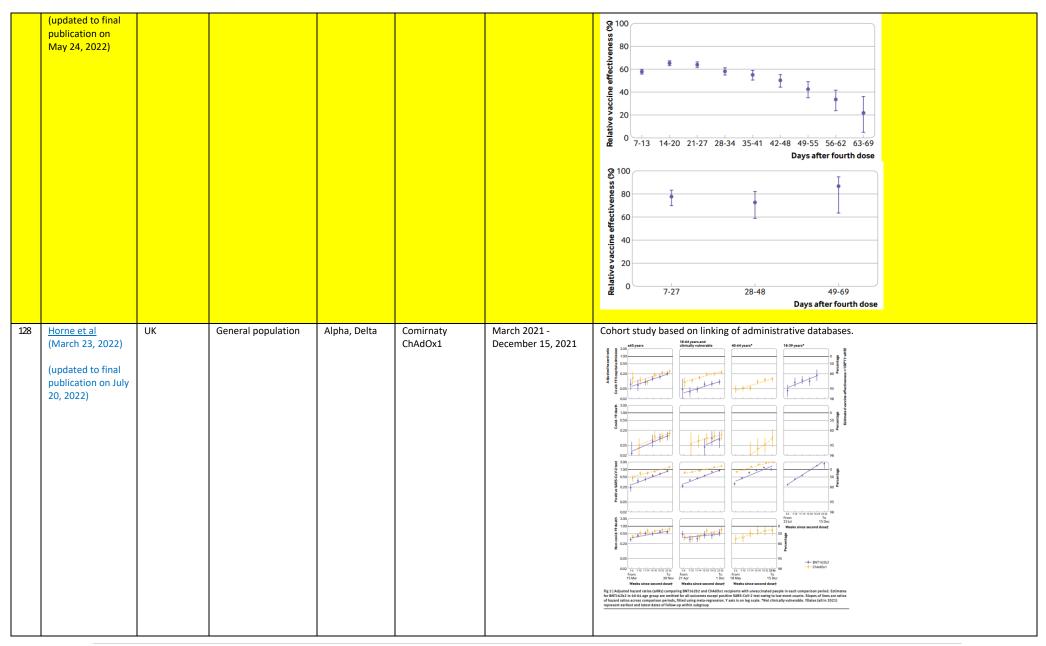






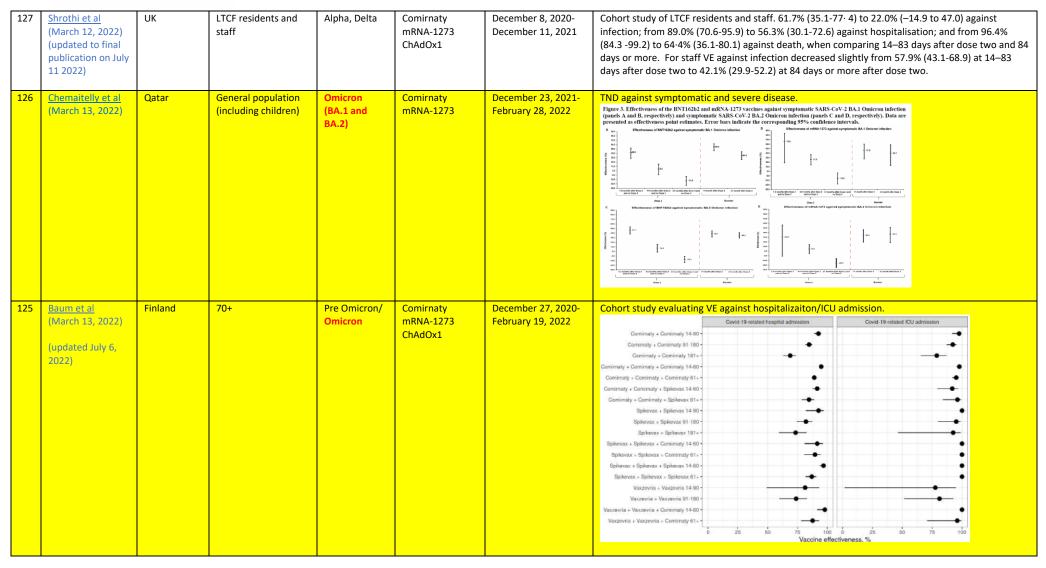












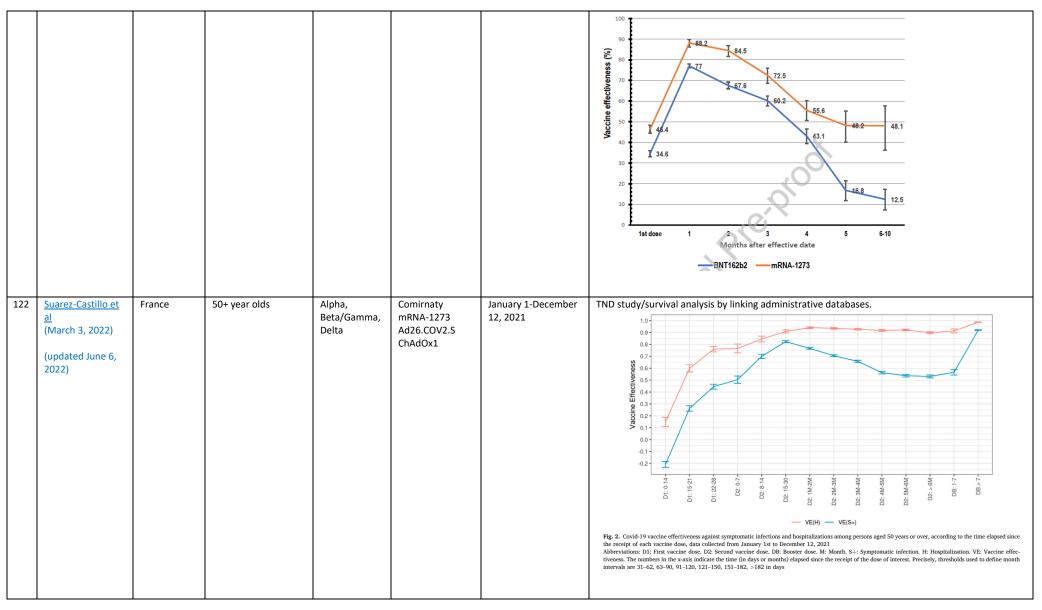




							Supplementary Table 11: VE against Covid-19-related hospital admission in 2022 Q1, i.e., between January 01 and Pebruary 19. Vaccine effectiveness (in %) quantified as 1 minus the hazard ratio adjusted for age, ser, region of residence, residence in a long-turm care facility, infiluenza vaccination in 2019-2020, number of nights hospitalized between 2015 and 2019 and presence of predisposing comorbilities. Committed 145 5121
124	Fowlkes et al (March 11, 2022)	USA	5-15 year olds	Delta, Omicron	Comirnaty	July 25, 2021– February 12, 2022	Cohort study finding the adjusted VE at 14–149 days after receipt of dose 2 was 87% (95% CI = 49%–97%) against Delta infection and 59% (95% CI = 22%–79%) against Omicron infection. Adjusted VE ≥150 days after dose 2 was 60% against Delta infection and 62% against Omicron, with wide CIs that included zero.
123	Syed et al (March 2, 2022)	Qatar	12+	Alpha, Beta/Gamma, Delta	Comirnaty mRNA-1273	December 16, 2020- October 31, 2021	Cohort study linking adminsitrative databases. VEs are unadjusted











121	Klein et al	USA	5-17 year olds	Omicron	Comirnaty	April 2021-January	TND study evaluating VE aga	ainst eme	rgency denai	rtment/urge	nt care visits and hospitalizations.	
121	(March 1, 2022)	03/1	3 17 yeur olus	Delta	Commutey	2022			SARS-CoV-2 test-positive,	VE%*		
							Encounter type/Vaccination status		no. (%)	(95% CI)		
							ED or UC encounters during Delta 5–11 yrs	ED or UC encounters during Delta or Omicron predominance, by age group				
							Unvaccinated (Ref)	8,599	2,652 (30.8)	_		
							2 doses (14–67 days earlier)	582	124 (21.3)	46 (24-61)		
							12–15 yrs Unvaccinated (Ref)	12,064	3,238 (26.8)	_		
							2 doses (14–149 days earlier)	4,547	254 (5.6)	83 (80-85)		
							2 doses (≥150 days earlier) 3 doses (≥7 days earlier)	1,517 10	378 (24.9) 3 (30)	38 (28-48) NC		
							16-17 yrs					
							Unvaccinated (Ref) 2 doses (14–149 days earlier)	7,421 2,692	2,068 (27.9) 193 (7.2)	76 (71–80)		
							2 doses (≥150 days earlier)	1,721	329 (19.1)	46 (36–54)		
							3 doses (≥7 days earlier)	64	13 (20.3)	86 (73-93)		
							ED or UC encounters, by age grou 5–11 yrs**	ıp anα preα	Johnnant Varian			
							Omicron predominant ⁺⁺					
							Univaccinated (Ref) 2 doses (14–67 days earlier)	5,938 486	2,409 (40.6) 118 (24.3)	51 (30-65)		
							12-15 yrs					
							Delta predominant ⁺⁺ Unvaccinated (Ref)	9,633	1,978 (20.5)	_		
							2 doses (14–149 days earlier)	4,060	80 (2.0)	92 (89-94)		
							2 doses (≥150 days earlier)	798	32 (4.0)	79 (68–86)		
							Omicron predominant ⁺⁺ Unvaccinated (Ref)	2,336	1,254 (53.7)	_		
							2 doses (14–149 days earlier)	472	174 (36.9)	45 (30-57)		
							2 doses (≥150 days earlier) 3 doses (≥7 days earlier)	719 10	346 (48.1) 3 (30.0)	−2 (−25−17) NC		
							16-17 yrs					
							Delta predominant ⁺⁺ Unvaccinated (Ref)	5.302	1,191 (22.5)	_		
							2 doses (14–149 days earlier)	2,340	78 (3.3)	85 (81-89)		
							2 doses (≥150 days earlier) 3 doses (≥7 days earlier)	1,156 2	47 (4.1) 0 (—)	77 (67–84) NC		
							Omicron predominant#					
							Unvaccinated (Ref) 2 doses (14–149 days earlier)	1,363 263	771 (56.6) 114 (43.4)	34 (8-53)		
							2 doses (≥150 days earlier)	565	282 (49.9)	-3 (-30-18)		
							3 doses (≥7 days earlier)	62	13 (21.0)	81 (59-91)		
							Hospitalizations during Delta or (5–11 yrs	ormulon pr	econimance, by	age group		
							Unvaccinated (Ref)	262	59 (22.5)			
							2 doses (14–67 days earlier) 12–15 yrs	23	2 (8.7)	74 (-35-95)		
							Unvaccinated (Ref)	496	149 (30)	_		
							2 doses (14–149 days earlier) 2 doses (≥150 days earlier)	182 63	7 (3.8) 13 (20.6)	92 (79-97) 73 (43-88)		
							16–17 yrs	0.5	13 (20.0)	73 (43-00)		
							Unvaccinated (Ref)	437	136 (31.1)	_		
							2 doses (14–149 days earlier) 2 doses (≥150 days earlier)	150 82	7 (4.7) 14 (17.1)	94 (87–97) 88 (72–95)		
							3 doses (≥7 days earlier)	4	1 (25.0)	NC NC		





120	Smid et al	Czech	General population	Omicron	Comirnaty	December 7, 2021-	Cohort study created by linking administrative databases. (<2 months and >=2 months prior to
-20	(February 25,	Republic	of country	Delta	mRNA-1273	February 13, 2022	onset)
	2022)	Периопе	or country	Deita	Ad26.COV2.S	1 Coldary 13, 2022	onset)
	2022)				ChAdOx1		Protection against Delta and Omicron infection
	(updated April 28,				CHAUOXI		1.0
	2022)						0.9-
	2022)						y 0.8-
							89 0.7
							g 0.7-
							<u>1000</u> 0.6-
							\$\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\
							9 0.4·
							g 0.3.
							<u></u> <u> </u>
							0.1
							0.0 Inf6- Inf6+ Full2- Full2+ Booster2- Booster2+
							Fig. 2. Protection provided by vaccination or previous infection against infection by
							the Omicron and Delta variants of the SARS-CoV-2 virus. Inf6-, previous infection <6
							months ago; Inf6+, previous infection > 6 months ago; Full2-, complete vaccination < 2
							months ago; Full2+, complete vaccination >2 months ago; Booster2-, booster dose <2 months ago; Booster2+, booster dose >2 months ago; Booster2+, booster dose >2 months ago; Booster2-,
							of protection with 95% CI.
							of protocolor with 55% of.
							Table 3. Vaccine effectiveness and protection provided by post-
							infection immunity against hospitalization, for the Omicron and Delta
							variants of the SARS-CoV-2 virus, 95% confidence intervals (CI) in
							parentheses.
							Effect ag. Hosp. Omicron Delta
							Full 2- 45% (29-57%) 75% (68-80%)
							Full 2+ 29% (21-37%) 79% (78-81%)
							Booster 2- 87% (84-88%) 98% (97-98%)
							Booster 2+ 79% (75-83%) 97% (95-98%)
							Table 6. Vaccine effectiveness and protection provided by post- infection immunity against hospitalization with a need for oxygen
							therapy, for the Omicron and Delta variants of the SARS-CoV-2 virus,
							95% confidence intervals (CI) in parentheses.
							Effect ag. O_2 Omicron Delta
							Full 2- 57% (32-72%) 82% (76-87%)
							Full 2+ 32% (20-43%) 82% (80-83%)
							Booster 2- 90% (87-92%) 98% (98-98%)
							Booster 2+ 85% (80-88%) 97% (95-98%)





							Table 7. Vaccine effectiveness and protection provided by post- infection immunity against hospitalization with a need for <i>intensive care</i> , for the Omicron and Delta variants of the SARS-CoV-2 virus, 95% confidence intervals (CI) in parentheses. Effect ag. ICU
119	Patalon et al (February 26, 2022) (updated June 9, 2022)	Israel	16+ Maccabi insured patients	Omicron	Comirnaty	January 1-January 21, 2022	Matched TND study to evaluate relative VE against infection and hospitalization/death. All persons had received the primary series by August 1, 2021. Marginal effectiveness against infection of a booster dose given a month before the outcome period was at its peak at 59.4% (95% CI, 54.9%-63.5%). Effectiveness declined gradually with time from inoculation, reaching 16% (95% CI, 12.3%-19.5%) in those vaccinated 5 months prior to the outcome period compared to those not receiving the booster dose. As for the marginal effectiveness against severe disease, it seems that waning exists though to a much lesser degree, as effectiveness declines from 72.2% (95% CI, 37.8%-87.6%) 3 months after inoculation to 54.5% (95% CI, 13.4-76.1) five months after vaccination. However, numbers are small as also reflected by the confidence intervals.
118	Wright et al (February 25, 2022)	USA	18+ hospitalized	Pre Delta; Delta	Comirnaty mRNA-1273 Ad26.COV2.S	April 1-October 26, 2021	Case-control study of patients hospitalized in one large US network of hospitals. 100 90 80 70 90 40 100 90 Waccine type Moderna Pfizer Janssen Vaccine effectiveness against severe COVID-19 by time since vaccination and vaccine type





117	Liu et al (February 18, 2022)	Australia	Persons exposed in two outbreaks (1 at a night club, 1 at a medical school graduation event)	Omicron	Comirnaty mRNA-1273 ChAdOx1	December 8, 2021- December 22, 2021	Unadjusted VE in two or Timing <1 month 1-2 months 2-3 months 3+ months	utbreaks by time since 2' Night club outbreak -33.3 (-141.4-26.3) -18.1 (-85.7-24.8) -5.9 (-67.5-33.1) -36.2 (-114.3-13.4)	dose (combined for al Graduation event outbreak No cases 87.5 (64-95.7) 60 (38-74.2) 32 (22-40.6)	l vaccines)
116	<u>Wu et al</u> (February 2022)	China	18+ year old contacts of cases	Delta	Coronavac BBIBP-CorV	July 31, 2021-? (prior to November 17, 2021)	COVID-19 was 52.32% (25.73-69.39) for ≤3-mon 0-19 pneumonia, VEs wer	th intervals and 49.95%	ation against symptomatic (1.2-74.64) for 4–6-month or ≤3-month and 67.08%
115	Britton et al (February 14, 2022)	USA	12+ year olds	Pre-Delta and Delta	Comirnaty mRNA-1273 Ad26.COV2.S	March 13, April 15, or June 15 (based on age-based vaccine- eligibility October 17, 2021	(note vaccination data by the paper, there is a strain the paper, there is a strain the paper, there is a strain the paper and t	passed on recall and some attification by age group. In m80M-1273 vaccination among the control of the control	e portion of 2 dose recip	ed for each product- and period-specific model, beyond daily OR estimates from days 14 to 60 (initial OR), and
114	Ferdinands et al (February 11, 2022)	USA	18+ years	Delta, Omicron	Comirnaty mRNA-1273	August 26, 2021- January 22, 2022	TND study at 8 VISION r hospitalizations.	network sites evaluating '	VE against emergency ro	oom/urgent care visits nad

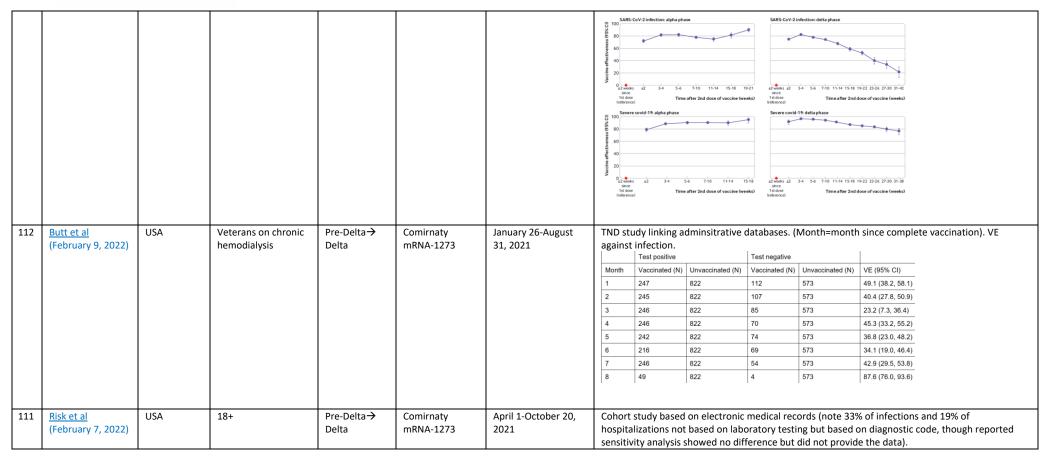




							TABLE 2. mRNA COVID-19 vaccine e care encounters and hospitalization August 2021–January 2022**				
							Characteristic	Total	SARS-CoV-2 positive test result no. (%)	VE fully adjusted % (95% CI)*	Waning trend p value [†]
							ED/UC encounters Overall				
ı							Unvaccinated (Ref)	110,873	43,054 (39)	_	_
1 /							Any mRNA vaccine, 2 doses	105,193 4,808	16,487 (16) 301 (6)	72 (72-73) 88 (87-90)	<0.001
1							2-3 mos	10,644	1,312 (12)	80 (78-81)	
							4 mos	10,175	1,230 (12)	79 (77-80)	
7							≥5 mos Any mRNA vaccine, 3 doses	79,566 25,138	13,644 (17) 2,285 (9)	69 (68-70) 89 (89-90)	<0.001
J							<2 mos 2–3 mos	15,614 8,759	920 (6) 1,120 (13)	92 (91–93) 86 (85–87)	
I							4 mos	736	227 (31)	75 (70-79)	
1.7							≥5 mos Delta-predominant period	29	18 (62)	50 (-7-77)	
							Unvaccinated (Ref)	86,074	29,063 (34)	_	_
							Any mRNA vaccine, 2 doses <2 mos	85,371 4,253	8,136 (10) 144 (3)	80 (79-81) 92 (91-94)	<0.001
							2-3 mos	8,662	527 (6)	88 (86-89)	
							4 mos ≥5 mos	8,941 63,515	721 (8) 6.744 (11)	85 (83-86) 77 (76-78)	
							Any mRNA vaccine, 3 doses	14.207	347 (2)	96 (95-96)	<0.001
I							<2 mos 2–3 mos	10,621 3,542	210 (2) 134 (4)	97 (96–97) 93 (92–94)	
1							≥4 mos	44	3 (7)	89 (64–97)	
ı							Omicron-predominant period Unvaccinated (Ref.)	24,799	13,991 (56)	_	_
I							Any mRNA vaccine, 2 doses	19,822	8,351 (42)	41 (38-43)	<0.001
							<2 mos 2–3 mos	555 1,982	157 (28) 785 (40)	69 (62-75) 50 (45-55)	
							4 mos	1,234	509 (41)	48 (41-54)	
							≥5 mos Any mRNA vaccine, 3 doses	16,051 10,931	6,900 (43) 1,938 (18)	37 (34–40) 83 (82–84)	<0.001
V							<2 mos	4,993	710 (14)	87 (85-88)	(0.001
							2–3 mos 4 mos	5,217 692	986 (19) 224 (32)	81 (79–82) 66 (59–71)	
							≥5 mos	29	18 (62)	31 (-50-68)	
							Hospitalizations Overall				
							Unvaccinated (Ref)	40,125	16,335 (41)	_	_
							Any mRNA vaccine, 2 doses	42,326	4,294 (10) 71 (4)	82 (81-83) 93 (91-94)	<0.001
1							<2 mos 2–3 mos	1,662 3,084	223 (7)	88 (86-90)	
							4 mos ≥5 mos	3,279 34,301	234 (7) 3,766 (11)	89 (87–90) 80 (79–81)	
							Any mRNA vaccine, 3 doses	10,957	471 (4)	93 (92-94)	< 0.001
							<2 mos 2–3 mos	7,332 3,413	221 (3) 211 (6)	95 (94–95) 91 (89–92)	
							2–3 mos ≥4 mos	3,413	39 (18)	81 (72-87)	
							Delta-predominant period				_
							Unvaccinated (Ref) Any mRNA vaccine, 2 doses	36,214 38,707	14,445 (40) 3,315 (9)	85 (84–85)	<0.001
							<2 mos	1,574	49 (3)	94 (92-96)	
							2–3 mos 4 mos	2,790 3,129	154 (6) 192 (6)	91 (89-92) 90 (89-92)	
							≥5 mos	31,214	2,920 (9)	82 (82-83)	
Г							Any mRNA vaccine, 3 doses <2 mos	8,124 6,071	195 (2) 118 (2)	95 (95–96) 96 (95–97)	<0.001
							2–3 mos ≥4 mos	2,030 23	74 (4) 3 (13)	93 (91–95) 76 (14–93)	
Г							Omicron-predominant period			70 (14-93)	
							Unvaccinated (Ref)	3,911	1,890 (48)		
							Any mRNA vaccine, 2 doses <2 mos	3,619 88	979 (27) 22 (25)	55 (50-60) 71 (51-83)	0.01
							2–3 mos 4 mos	294 150	69 (23) 42 (28)	65 (53-74) 58 (38-71)	
J							≥5 mos	3,087	846 (27)	54 (48-59)	
1							Any mRNA vaccine, 3 doses <2 mos	2,833 1,261	276 (10) 103 (8)	88 (86-90) 91 (88-93)	< 0.001
							2–3 mos	1,383	137 (10)	88 (85-90)	
							≥4 mos	189	36 (19)	78 (67–85)	
	- I I I I I		10		0	D 07.055				1 .	
	<u>Fabiani et al</u>	Italy	16+ years	Alpha, Delta	Comirnaty	December 27, 2020-	Cohort study of pe				
1 (1	February 10,				mRNA-1273	November 7, 2021	Used of day 0-<14	days post of	dose 1 as proxy for	unvaccinat	ed group. Pr
					-	,	and risk group in pa				0 - 1
								auei			
2022)							and risk group in p	apci.			











							Vaccine Ef	ffectiveness				HF	(95% CI)	p-value	
								/-2 Infection							
							BNT162b2 pre-delta								
							0-6 mor	nths				0.13	(0.1-0.16) <0.001	
							6+ mon		⊢•				,	3) <0.001	
							post-delta								
							0-6 mor	nths	H∎H			0.36	(0.32-0.42	?) <0.001	
							6+ mon					0.78	(0.67-0.9) 0.002	
							mRNA-127	73							
							pre-delta 0-6 mor	nthe				0.00	(0.06-0.1	3) <0.001	
							6+ mon						,	(a) <0.001	
							post-delta					0.11	(0.00 0.2	, 0.001	
							0-6 mor	nths	+■			0.22	(0.17-0.3	3) <0.001	
							6+ mon	iths	-	→		0.45	(0.33-0.6) <0.001	
									0 0.5	1	1.5	2			
110	Cerqueria-Silva et	Brazil	General population	Gamma, Delta	Coronavac	January 18-	TND stud	dy linking a	administra	ative dat	ahases				
110	al	Diazii	General population	Guillia, Delta	followed by	November 11, 2021		tiveness of Corona\			Table 4 Effe	ctiveness of CoronaV			
	(February 9, 2022)				Comirnaty	11000111501111, 2021	dose vaccination	nfection, by length on on or BNT162b2 bo			dose vaccina	on or death, by length tion or BNT162b2 boo	of time (in da ster dose, str	rs) since two- tified by age	
					booster		group Period after	Overall 18-59	9 60-79	≥80	group Period after		60-79	≥80	
							vaccine (days) Second dose				Vaccine (days) Second dose				
							0-13		% 32.2%		0-13	65.5% 79.6% (64.2-66.6) (77.6-8			
							14-30	(36.9-38.8) (42.4- 55.0% 56.5%	% 55.1%	50.3%	14-30	82.1% 91.4%	81.6%	68.7%	
								(54.3-55.7) (55.6- 51.7% 52.9%	5-57.5) (53.7-56.5) % 51.1%		31-60	(81.4-82.8) (90.3-9 82.6% 89.9%	81.4%	66.5%	
								(51.1-52.4) (52.1- 47.6% 48.99	-53.8) (49.7-52.4) % 45.3%		61-90	(82.1-83.2) (88.9-9 80.5% 87.2%) (64.0-68.9) 63.2%	
								(46.8-48.3) (47.9-	-49.9) (43.6-46.9)	(37.3-44.4)	91-120	(79.8-81.0) (86.0-8			
l							,	(45.3-46.9) (51.3-		(27.3-36.1)		(78.3-79.6) (87.8-9)	.0) (74.3-76.7	(54.7-61.1)	
								41.8% 50.69 (40.8-42.8) (49.3-	% 36.3% 3-51.9) (33.8-38.7)		121-150	77.0% 86.7% (76.1-77.8) (85.2-8)			
								38.0% 44.09 (36.7-39.3) (42.3-	% 35.3% 8-45.6) (32.2-38.2)	15.1% (8.3-21.5)	151-180	75.0% 81.9% (73.9-76.0) (79.8-8			
							>180	34.7 % 34.1%	6 34.5%	10.1%	>180	72.6% 74.8% (71.0-74.2) (72.1-77	72.6%	41.4%	
							Booster (BNT162	(33.1-36.3) (32.2- 2b2)	(29.9-38.7)	(1.1-10.3)	Booster (BNT	62b2)			
								39.6% 40.39 (33.8-44.8) (31.6-	% 35.7% -47.8) (25.2-44.8)		0-6	80.6% 89.1% (76.4-84.0) (76.6-9	79.6% I.9) (73.5-84.2	48.8%) (31.3-61.9)	
							7-13		% 75.9%	59.6%	7-13	91.4% 95.8% (88.5-93.5) (82.9-9	88.3%	78.0%	
							14-30	92.7% 93.5%	% 93.4%	82.0%	14-30	97.3% 97.9%	97.1%	89.5%	
1	1							(91.0-94.0) (90.7- 82.6% 61.8%			>30		92.0%	89.3%	
												(94.1-98.3)			
								(76.9-86.9) (27.2-	(-/9.9) (67.6-89.1)	(49.6-77.5)	*The Classic Co.) (78.6-94.7)	
								(76.9-86.9) (27.2-	-/9.9) (67.6-89.1)	(49.6-77.5)	*The CI could not b	e estimated owing to zero/few ex) (/8.0-94./)	





							Extended Data Table 4 Vaccine effectiveness against death due to COVID-19 using RT-PCR, by length of time (in days) since two-
							dose vaccination or BNT162b2 booster dose
							Period post vaccine (days) Overall 18-59 60-79 ≥80
							Second dose
							0-13 67.3% (65.6-68.9) 86.4% (82.5-89.4) 69.6% (67.6-71.6) 56.0% (51.6-60.0)
							14-30 82.7% (81.7-83.6) 91.4% (88.7-93.5) 84.5% (83.3-85.6) 72.7% (69.8-75.4)
							31-60 83.6% (82.8-84.3) 91.9% (89.7-93.6) 84.8% (83.8-85.7) 70.0% (67.2-72.5)
							61-90 81,4% (80,5-82,2) 92,2% (89,8-94,0) 82,5% (81,3-83,7) 67,2% (64,2-69,9)
							91-120 79.8% (78.7-80.8) 95.0% (93.1-96.4) 81.7% (80.3-83.0) 63.5% (59.9-66.7)
							121-150 78.3% (77.0-79.6) 93.7% (90.9-95.7) 82.0% (80.3-83.5) 58.7% (54.3-62.7)
							151-180 76.8% (75.1-78.4) 92.1% (88.2-94.7) 81.9% (79.7-83.8) 53.9% (48.3-58.9)
							>180 74.8% (72.2-77.2) 90.3% (85.5-93.5) 81.5% (77.6-84.7) 45.5% (37.1-52.8)
							Booster (BNT162b2)
							0-6 80,3% (73,1-85,6) 100% (*) 81,4% (71,3-87,9) 59,9% (39,3-73,5)
							7-13 92.2% (87.4-95.2) 100% (*) 92.3% (83.8-96.3) 80.7% (65.3-89.2)
							14-30 98.3% (96.3-99.2) 81.9% (-31.6-97.5) 99.1% (93.6-99.9) 95.4% (88.7-98.1)
							>30 97.1% (90.5-99.1) 100% (*) 94.3% (58.3-99.2) 93.5% (73.2-98.4)
109	Andeweg et al	Netherlands	General population	Omicron	Comirnaty	November 22, 2021-	TND study linking administrative databases evaluating VE/risk reduction from prior infection
	(February 8, 2022)			(BA.1 and	ChAdOx1	January 19, 2022	and/or vaccination.
	(updated May 12,			BA.2)	mRNA-1273	·	A Dalla Onlinea PA Carbori
	2022)			Delta	Ad26.COV2.S		William Committee Committee
	2022)			Denta	7.020.0072.0		Previous infection, Primary vaccination Booster unascontained Primary vaccination
							90 7 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4
							20 4 4 4 4 4 4 4 4 4 4 4
							8 0
							First start primary vaccination, First inflection, Previous infection,
							then infection then primary vaccination booster
							Agrilos 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2
							30 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1
							10
							Time since last event (days)
							B. Omicron BA.1-BA.2 cohort Variant → Omicron BA.1 → Omicron BA.2
							Previous infection, Primary vaccination Booster unacconstend
							90
							70 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5
							9 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3
							€ 10
							5 ± 5 + + + + + + + + + + + + + + + + +
							First start primary vaccination, First infection, Previous infection, then primary vaccination bossiler
							> 100 100 100 100 100 100 100 100 100 100
							2 75 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1
							40
							<u> </u>
							35 1
							Jacobson Company Compa
							Time since last event (days)





108	Chemaitelly et al	Qatar	General population	Omicron	Comirnaty	December 23, 2021-	Matched TND study based on linking adminsitrative databases.
100	(February 8, 2022)	Qatai	General population	Jilicion	mRNA-1273	February 2, 2022	Figure 1. Effectiveness of the BNT162b2 vaccine against A) symptomatic SARS-CoV-2
	(: 22: 32:) 2/ 2222/						Omicron infection and B) severe, critical, or fatal COVID-19 due to Omicron infection. C)
							Effectiveness of the mRNA-1273 vaccine against symptomatic SARS-CoV-2 Omicron infection. Data are presented as effectiveness point estimates. Error bars indicate the
							corresponding 95% confidence intervals. A Effectiveness of the BNT162b2 vaccine against symptomatic Omicron infection
							86.0 88.0
							75.0 66.0
							\$\frac{a}{a} \qquad \qqqq \qqq
							3 340
							10 10 10 10 10 10 10 10 10 10 10 10 10 1
							450
							14 0 2nd 2nd 4m 5m 5m 6m 7m 6m 5m 10m 10m 10m 10m 10m 10m 10m 10m 10m 10
							Does 2 Booster
							B Effectiveness of the BNT162b2 vaccine against any severe, critical, or fatal 1886 - SARS-CoV-2 Infection 1 1868 - T 1868
							86.0 T 180.7 T 180.7 T 180.7
							2 60.0
							8 40.0 L
							38.0 - 10
							0.0 1-4 months after Dose 2 or success after Dose 2 1-4 meets after Dose 3 or weeks after Dose 3
							One 2 Booster C Effectiveness of the mRNA-1273 vaccine against symptomatic Omicron infection
							M.S.
							76.0 MA -
							50.0 P 50.1 9 50.5 P 50.1
							1 Ms 1 1
							** (4.5
							419
							1-3 roottes where 4 a screttes where 2 is acustas where Closes 2 Dones 2 Dones 3 Dones 3 Dones 3 Dones 3 Dones 2 Dones 2 Dones 2 Dones 3 Dones
							Desa 2 Baseter The time-since-veccination categories were chosen to maximize statistical precision in each analysis as the number of cases was arreal for some of the time-since-veccination status. Therefore, the time-since-veccination categories were different for the INT INT IDEA and
							miRNA-1273 mallyses and for the analysis against any severe, ortical, or faal COVID-19.
							mRNA-1273
							Cases' Controls' (Severe, rifical, or (PCR-negative) Sub-studies' (Severe, rifical, or (PCR-negative) Effectiveness in %
							Sub-studies fatal disease :
							Dose 1
							Dose 2 1-6 months after Dose 2 and no Dose 3 3 105 35 265 76.9 (19.2 to 93.4)
							27 months after Dose 2 and no Dose 3 23 117 139 257 64.0 (39.1 to 78.7) Dose 3 (booster dose)
							16 weeks after Dose 3 1 103 19 270 80 8 (-51.9 to 97.6) ≥7 weeks after Dose 3 0 102 5 278 100.0 (Omitted)¹
107	Lauring et al	USA	≥18 years	Delta (for the	Comirnaty	July 4-December 25,	TND case control study in 21 hospitals in the US (IVY Network). For Delta, VE against
	(February 7, 2022)			duration	mRNA-1273	2021 (for the Delta	hospitalization 88% (95% CI: 86 to 90%) 14-150 days post 2 nd dose; >150 days, VE was 81% (78 to
				analysis	1	analysis)	84%).





	(updated March 9, 2022)						
106		Portugal	≥12 years	Delta→ Omicron	Comirnaty ChAdOx1 mRNA-1273 Ad26.COV2.S	December 6-21, 2021	Compared the odds of vaccination in Delta versus Omicron cases. (higher odds =lower VE of Omicron). Omicron: Delta aOR Complete primary vaccination <113 days 2.3(1.9 to 2.8) Complete primary vaccination 113-168 days 2.0 (1.7 to 2.4) Complete primary vaccination 169+ days 1.9(1.6 to 2.3)
105	Corrao et al (January 27, 2022)	Italy	≥12 years	Alpha → Delta	Comirnaty ChAdOx1 mRNA-1273 Ad26.COV2.S	January 17-October 20, 2021	Cohort study Figure 1: Influence of time since complete vacination on vacation and production from the complete vacination on vacation and production from the complete vacination from production from p





104	Roberts et al (January 31, 2022)	USA	Adults	Multiple	Comirnaty mRNA-1273 (for duration)	January 1-December 31, 2021	TND study evaluating VE against infection (top) and hospitaliation/death (bottom). Note that this is a combination of primary and booster dose VE in quarter 4.
							B VE for Severity Other Control of Control
103	Belayachi et al (January 27, 2022)	Morocco	≥18 year olds	Unknown→ Delta	BBIBP-CorV	February 1-October 1, 20221	TND linking adminsitrative databases to evaluate VE against severe disease. As a function of time after vaccination of second dose vaccination, vaccine effectiveness among persons who had received the second dose 1–30 days earlier was 88% (95% CI, 84-91), 87% (95% CI: 83-90) among those who had received it 31–90 days earlier, 75% (95% CI: 67-80) among those who had received it 91–120 days earlier, 61% (95% CI: 54-67) among those who had received it 121–150 days earlier, 64% (95% CI: 59-69) among those who had received it ≥150 days earlier. Note they attempted to stratify by age (>/< 60 years) showing a trend towards a lower VE gainst severe/critical disease in those over 60 but confidence intervals were overlapping.
102	Lytras et al (January 29, 2022) (updated June 14, 2022)	Greece	≥15 year olds	Alpha→Delta	Comirnaty ChAdOx1 mRNA-1273 Ad26.COV2.S	January-December 2021	Cohort study linking administrative databases evaluating VE against intubation and death. VE provided for 6 months





							Vaccine Effectiveness (comparative)
							Vaccine VE (%) VE (%)
							3-dose BNTHs2b2 (age 15-79) 98.2 (972-98.9) 98.3 (96.8-99.1)
							3-dose BNT162D2 (age 80+) 97.5 (95.5-98.6) 98.4 (97.4-99.0)
							2-dose BNTis/22 (before "delta", age 15-29) 96.7 (95.3-97.6) 96.2 (94.2-97.5)
							2-dose BNT1s202 (Defore "delta", age 60-79) 94.1 (92A-95.4) 93.5 (913-94.8)
							2-dose BNTR2D2 (before "delta", age so+)
							2-dose BNTIG2b2 (age 15-59) 98.1 (97.5-98.6) 96.5 (94.8-97.6)
							2-dose BNTIc2b2 (age 60-79) 96.7 (95.9-97.A) 94.1 (92.7-95.2)
							2-dose BNT16/2b2 (age 80+) 94.2 (92.0-95.7) 91.0 (88.4-93.0)
							2-dose BNTIc2b2 (age 15-59, at 6 months) 95.5 (94.3-96.5) 9 93.8 (91.0-95.7)
							2-dose BNT162b2 (age 60-79, at 6 months) 92.0 (91.0-92.9) 89.4 (87.9-90.8)
							2-dose BNT162b2 (age 80+, at 6 months) 85.9 (83.5-88.0) 86.0 (82.2-85.6)
							2-dose mRNA-1273 (age 15-59) 99.4 (98.2-99.8) 99.3 (94.7-99.9)
							2-dose mRNA-1273 (age 60-79) 98.9 (97.3-99.5) 98.4 (93.5-99.5)
							2-dose mRNA-1273 (age 80+) — 97.9 (90.2-99.5) — 96.7 (87.9-99.1)
							2-dose mRNA-1273 (age 15-59, at 6 months) 90.3 (80.3-99.9) 98.3 (80.3-99.8)
							2-dose mRNA-127 (age 60-79, at 6 months) 95.1 (93.0-96.5) 95.2 (93.6-97.7)
							2-dose mRNA-1273 (age 80+, at 6 months) 92.6 (67.0-97.3) 92.0 (80.0-96.8)
							2-dose ChAdOx1 nCoV-19 (age 15-59) 97.8 (95.3-99.0) - 97.5 (89.7-99.4)
							2-dose Child OX1 nCOV-19 (age 60-79) 97.2 (95.3-98.3) - 95.4 (912-97.6)
							2-dose ChAdOx1 nCoV-19 (age 80+) 97.8 (91.7-99.A) 92.6 (84.2-96.5)
							2-dose ChAdOX1 nCoV-19 (age 15-59, at 6 months) — 92.A (84.0-96.A) — 94.5 (77.2-98.7)
							2-dose ChAdOX1 nCoV-19 (age 60-79, at 6 months) • 90.3 (87.4-92.5) • 89.8 (85.2-93.0)
							2-dose Chadoxi ncoV-19 (age eo-, at s months) = 92.4 (72.7-97.9) = 83.4 (69.6-90.9)
							1-doss Ad26.COV2.5 (age 15-59) — 85.0 (73.9-91.4) = 81.7 (57.5-92.1)
							1-0ose AQ26.COV2.S (age 60-79)
							1-dose Ad2s.COV.2.5 (age 80+)
							1-00se Ad26,COV2.5 (age 60-79, at 6 months)
							1-dose Ad2s.COV2.5 (age 80+7, 46 minitis)
							20 40 80 80 100 20 40 60 80 100
							VE (s.) against VE (s.) against Intubation death
101	Goldhaber-Fiebert	USA	Prison population	Delta	Comirnaty	June 1-November 5,	Matched TND among cases evaluating duration of protection against infection of early vs late fully
	et al		and staff		mRNA-1273	2021	(primary series) vaccinated persons. Among staff, odds of infection increased 25% (Odds Ratio
	(January 23, 2022)						[OR], 1.25; 95% Confidence Interval [CI], 1.13 – 1.40) in each 28-day period post-vaccination;
	(Juliadi y 23, 2022)						
							among residents, the odds increased by 21% (OR, 1.21; 95%Cl 1.08 – 1.36) (Figure 1). Compared
							with individuals within 60 days of being fully vaccinated, odds of infection were over fourfold
							greater ≥181 days since full vaccination for staff (OR, 4.36; 95%CI 1.92 – 9.89) and nearly threefold
							greater for residents (OR, 2.89; 95%CI 1.40 – 5.98)
-			<u> </u>				
100	Bedston et al	Wales	Healthcare Workers	Alpha → Delta	Comirnaty	December 7, 2020-	Cohort study. 2 weeks after dose 2, VE against infection was 67% (aHR 0.33, 95 %CI 0.24–0.44).
	(January 20, 2022)					September 30, 2021	This increased in weeks 2–5 to 86% (aHR 0.14, 95 %CI 0.09–0.21), and decreased to 77% over
	(3.11.11.)						weeks 6–13. After this, vaccine effectiveness decreased from 60% to 53% between weeks 14–25,
							and from week 26 vaccine effective was 45% (aHR 0.55, 95 %CI 0.49–0.61).
99	Accorsi et al	USA	≥18 year olds	Delta →	Comirnaty	December 10-	TND study in ICATT (free testing sites throughout US) against symptomatic disease. Note OR can be
	(January 21, 2022)		,	Omicron	mRNA-1273	January 1, 2022	converted to VE by the formulate VE=1-OR
				O.IIICI OII	111111111111111111111111111111111111111	Juliadi y 1, 2022	Converted to VE by the formulate VE-1 On





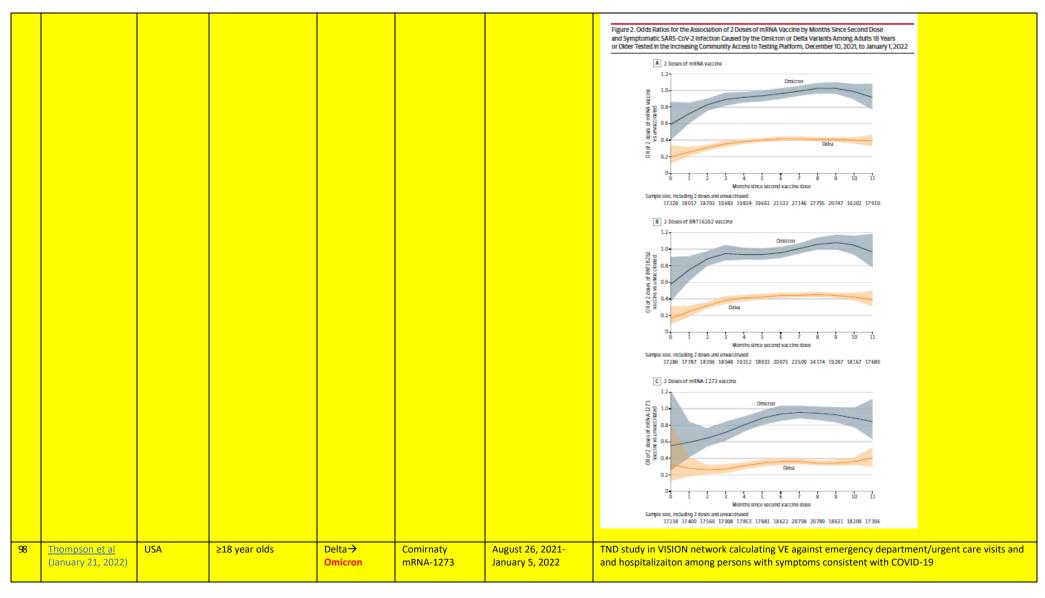






							TABLE 2. mRNA COVID-19 vaccine effectiveness* against labora encounters and hospitalizations among adults aged ≥18 year VISION Network, 10 states, August 2021–January 2022	story-confirmed COVID rs, by number and tin	-19–associated [†] emergency departm ning of vaccine doses ⁹ and vaccine p	nent and urgent care product received —	
							Encounter/Predominant variant period/Vaccination status	Total	SARS-CoV-2 positive test result, no. (%)	VE, %* (95% CI)	
							ED or UC encounters Delta predominant Unvaccinated (Ref) Any mRNA vaccine	98,087	36,542 (37.2)	=	
							2 doses (14–179 days earlier) 2 doses (±180 days earlier) 3 doses Omicron predominant	39,629 52,506 14,523	3,269 (8.2) 6,893 (13.1) 469 (3.2)	86 (85–87) 76 (75–77) 94 (93–94)	
							Unvaccinated (Ref) Any mRNA vaccine 2 doses (14–179 days earlier) 2 doses (2180 days earlier)	6,996 1,746 5,409	3,398 (48.6) 591 (33.9) 2,037 (37.7)	52 (46-58) 38 (32-43)	
							3 doses Hospitalizations Delta predominant Unvaccinated (Ref)	3,876 37,400	520 (13.4) 14,272 (38.2)	82 (79-84)	
							Any mRNA vaccine 2 doses (14–179 days earlier) 2 doses (2180 days earlier) 3 doses	14,645 26,190 8,092	895 (6.1) 2,563 (9.8) 209 (2.6)	90 (89–90) 81 (80–82) 94 (93–95)	
							Omicron predominant Unvaccinated (Ref) Any mRNA vaccine 2 doses (14–179 days earlier)	460 115	174 (37.8) 14 (12.2)	— 81 (65-90)	
							2 doses (±180 days earlier) 3 doses	488 514	86 (17.6) 24 (4.7)	57 (39-70) 90 (80-94)	
9	7 <u>Tartof et al</u> (January 19, 2022)	USA	≥18 year olds enrolled in Kaiser	Delta Omicron	Comirnaty	December 1, 2021- February 6, 2022	TND study of persons admitted COVID-19.	to the eme	ergency room or ho	ospital with	symptoms consistent with
			insurance	Officion		rebluary 6, 2022					
	(updated April 22, 2022)						Prograd administrative to old to (B. 5.6) 27) varient Second dose Third dose 100 75 888889 50 975 975 975 975 975 975 975 975 975 975	Second	+ +		
							E) administra due to delta (E 1.57.27) variant Second due Third dose Third dose 25 - 25 - 4 - 4 - 4 - 4 - 4 - 4 - 4 - 4 - 4 -	ED admission due to omismission due to omismo due to omismission due to omismission due to omismission due t			
							Selfer 35 per special	3. Freedom 3. Spreedom 6. Spreedom	Specified Specif		
g	Amodio et al (January 19, 2022)	Italy	≥18 year olds	Alpha→Delta	Comirnaty mRNA-1273	January 1-September 30, 2021	Cohort study of 3.9 millions adu trends for vaccine effectiveness significant for all the three eval infection; -2·27% per month, pa COVID-19 intubation/death, res	s, measured uated outco =0·029 agai	l as monthly perce omes (-4·76% per i	ntage chan month, p<0	ges, were statistically ·001 against SARS-CoV-2

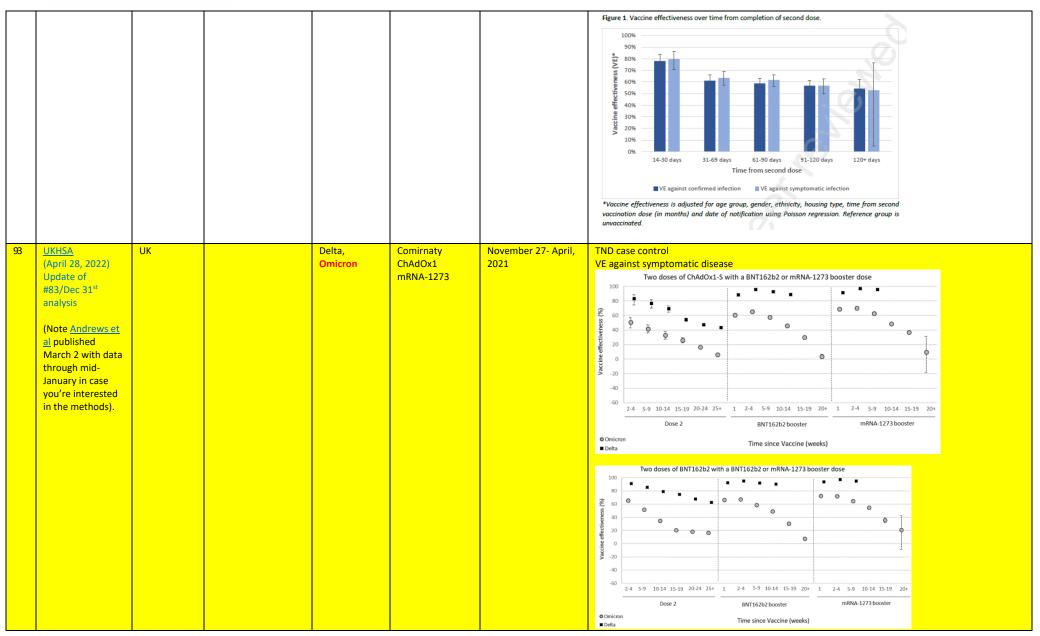




							Figure 4: Vaccine effectiveness estimates after adjustment for age and sex according to the different assessed outcomes and follow-up periods. A Vaccine effectiveness against SARS-CoV-2 infection Fathows period Consumer (2) 132 20881 6 197108 1497374
95	Suah et al (January 16, 2022) (updated June 2022)	Malaysia	General population	Delta	Comirnaty CoronaVac	September 1-30, 2021	Compared early (April-June) vs late (July-August) vaccinated persons (comparing to unvaccinated based on census data). For BNT162b2, crude vaccine effectiveness against COVID-19 infections declined from 90.8% (95% CI 89.4, 92.0) in the late group to 79.1% (95% CI 75.8, 81.9) in the late group. Vaccine effectiveness for BNT162b2 against ICU admission and deaths were comparable between the two different periods. For CoronaVac, crude vaccine effectiveness waned against COVID-19 infections from 74.4% in the late group (95% CI 209 70.4, 77.8) to 30.0% (95% CI 18.4, 39.9) in the early group. It also declined significantly against ICU admission, dropping from 56.1% (95% CI 51.4, 60.2) to 29.9% (95% CI 13.9, 43.0) (adjusted). For deaths, however, CoronaVac's effectiveness did not wane after three to five months of full vaccination. Waning more prominent in 60+.
94	Chiew et al (January 8, 2022)	Singapore	12-18 year olds	Delta	Comirnaty	June 1-November 20, 2021	Cohort study evaluating VE against infection and disease.













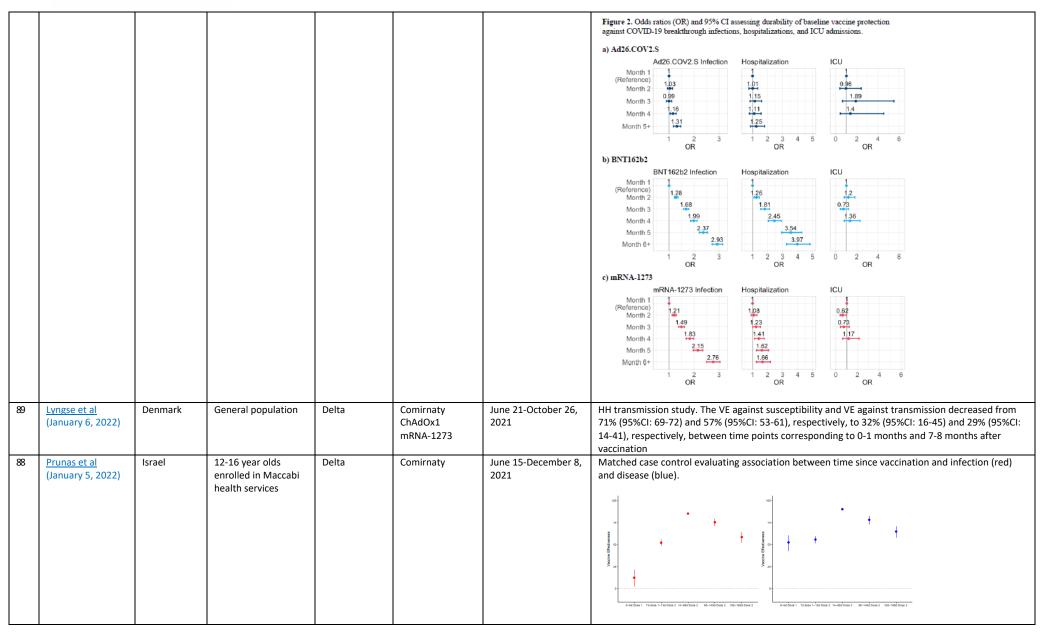




							2 3 3 3	Interval after dose 25+ weeks 2-4 weeks 5-9 weeks 10+ weeks	Odds Ratio 0.52 (0.34-0 0.08 (0.03-0 0.11 (0.07-0 0.12 (0.09-0	.81) 47.9 .12) 93. .17) 88.9	95% CI) (19.3 to 66.4) 6 (88 to 96.6) (83.4 to 92.6) (81.9 to 91.5)			
92	Tseng et al* (February 21, 2022) [update from January 21 preprint]	USA	18+ year olds enrolled in Kaiser insurance	Delta, Omicron	mRNA-1273	December 6-23, 2021	VE agains 2 dose (14-90 91-186 181-27 > 270 d 3 dose 3rd d 3 dose 3 rd cd 3 dose	st infection (14+) days o days 70 days days	60: 82: 63: 61:4 52: 95: 0/21 95: 11 90: setent) 95:	g administra to VE (95% C)) 7 (56.5-64.5) 8 (69.6-90.3) 8 (51.8-72.5) 9 (43.7-60.5) 9 (43.7-60.5) 2 (93.4-96.4) 7 (94.2-96.9) 7 (81.4-95.3) 7 (94.2-96.8) 9 (94.4-97.0) (83.9-97)		7.9) 3.9) 1.4)		
91	Grgič Vitek et al (January 6, 2022)	Slovenia	18+ year olds	Delta	Comirnaty mRNA-1273	October 2021	Note res Age group (Vaccinated a 18-49 50-64 ≥ 65 Vaccinated a 18-49 50-64 ≥ 65 Vaccinated a 18-49 50-64 ≥ 65	ults are unad years) 9 3 months ago 9 9 9 9 9 8 10 10 10 10 10 10 10 10 10	Vaccine effectiveness % 95% CI 97 90-99 14 91-97 13 88-96 14 NA 10 79-95 18 81-88 19 56-97 10 30-54					
90	Zheutlin et al (January 6, 2022)	USA	18+ year olds who had been fully vaccinated	Alpha, Delta, nonVOC	Comirnaty mRNA-1273 Ad26.COV2.S	January 1-September 7, 2021	odds of i	nfection, hos h after full va	I using an admin spitalization, and accination. Note	ICU admissi	on at 28 day	intervals p	ost dose 2 rela	ative to the

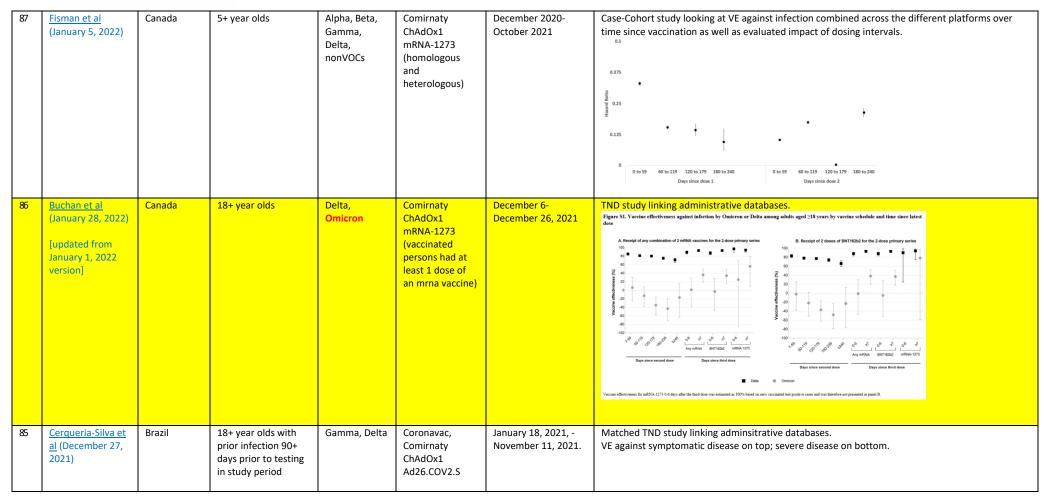












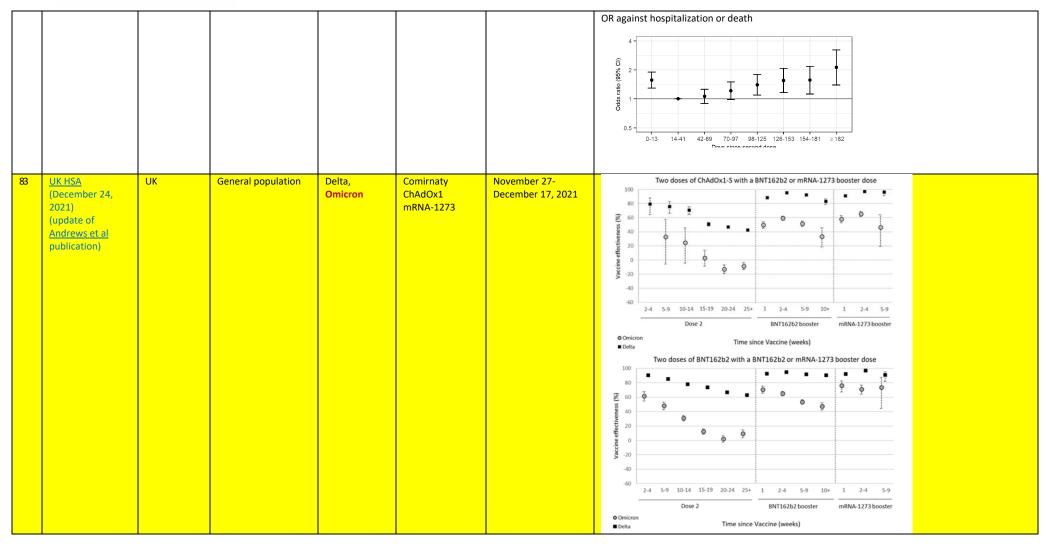




							BNT162b2 $\frac{14.90 \text{ days}}{(54.27\pm 0.0)} \times \frac{90 \text{ days}}{\text{value}}$ BNT162b2 $\frac{64.2\%}{(54.27\pm 0.0)} \times \frac{100\%}{(5)} 0.277$ ChAdOX1 $\frac{55.5\%}{(50.56\pm 0.1)} \times \frac{68.8\%}{(46.6-65.1)} 0.544$ CoronaVac $\frac{40.5\%}{(36.4-44.3)} \times \frac{33.0\%}{(33.1-42.5)} 0.760$ Ad26.COV2.S $\frac{46.1\%}{(32.7-56.7)} \times \frac{30.6\%}{(-12.4-57.1)} 0.420$ Table A4. Vaccine effectiveness ≥14 days after series comp Vaccine waning (time after series completion) 14-90 days >90 days p-value 88.8\% 100\% 0.765 BNT162b2 $\frac{88.8\%}{88.9\%} \times \frac{100\%}{(0.00\%)} \times \frac{95.1\%}{(77.6-92.0)} \times \frac{(84.8-98.4)}{(84.8-98.4)} \times 0.007$ CoronaVac $\frac{86.6\%}{(79.8-90.3)} \times \frac{94.8-94}{(33.3-82.2)} \times 0.012$ Go $\frac{60.2\%}{60.2\%} \times \frac{41.0\%}{41.0\%} \times 0.978$
84	Hitchings et al (December 24, 2021)	Brazil	18+ year olds living in Sao Paulo	Gamma, Delta	Coronavac	January 17- September 30, 2021	TND based on linking administrative databases among persons with 2 doses of coronavac (ref period day 14-41 post dose 2). OR for symptomatic disease. Priority status Non-HCW HCW

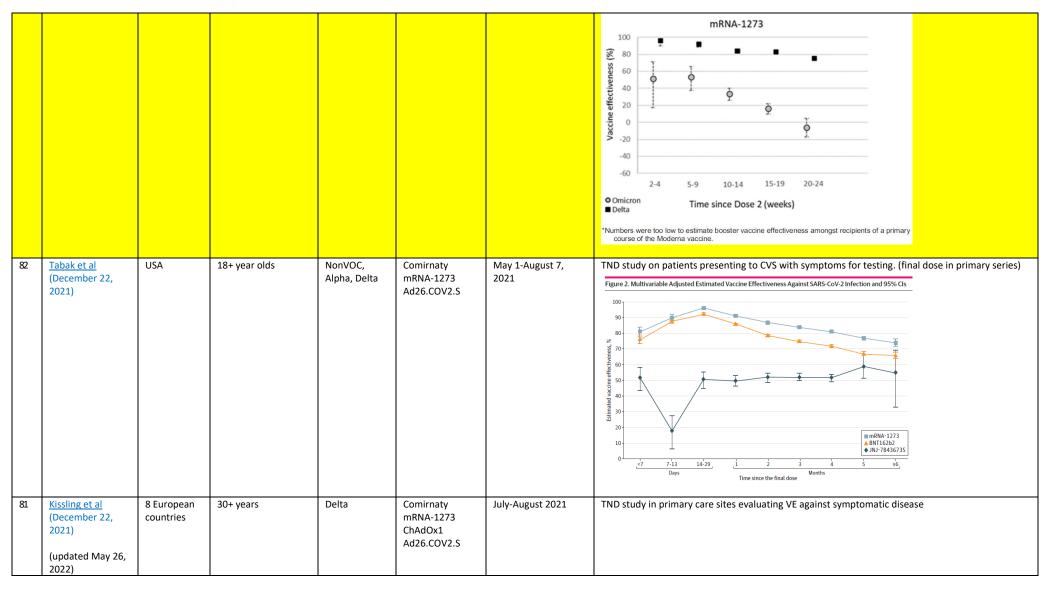






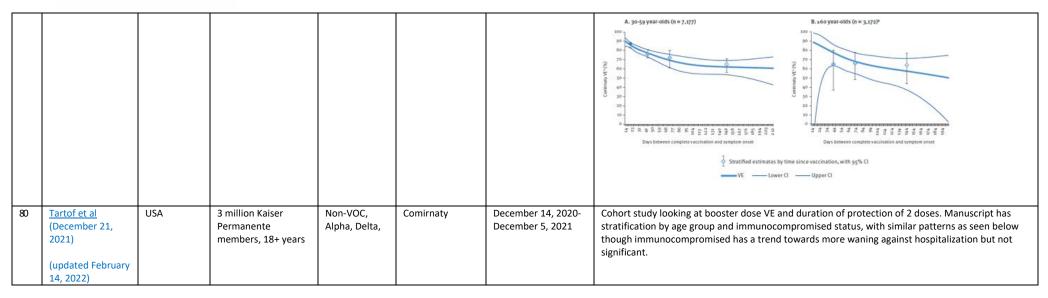






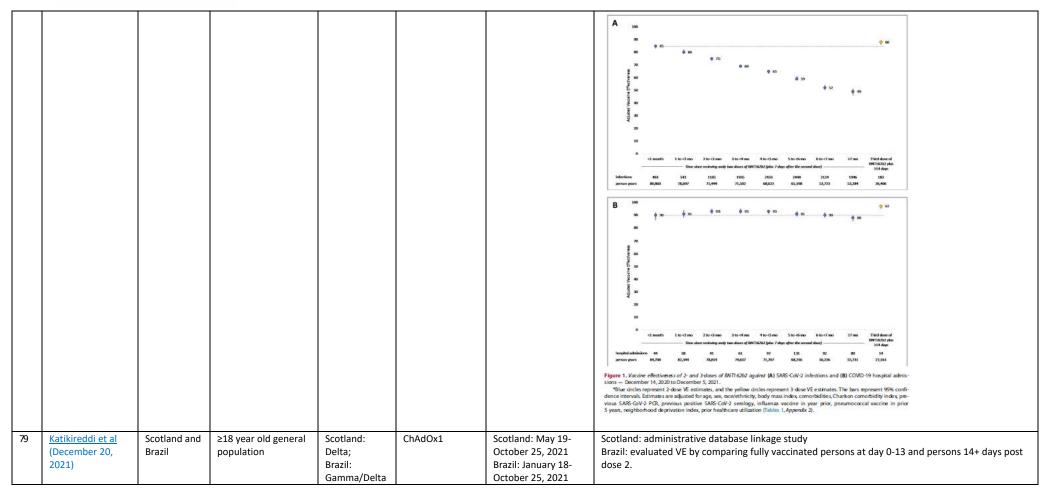












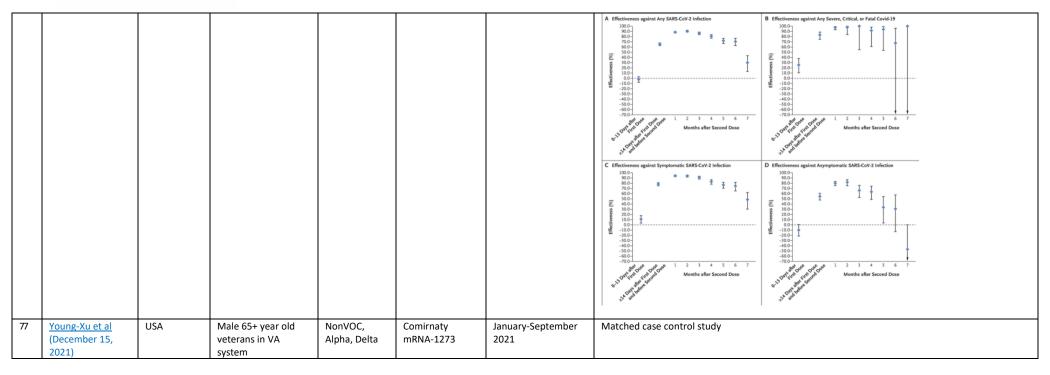




								Scotland			Brazil		
								Person-years	Number of events	Vaccine effectiveness* (95% CI)	Person-years	Number of events	Vaccine effectiveness* (95% CI)
							Unvaccinated	336 942	2245	0% (ref)			-
							0–2 weeks after first dose	6860	39	-15·4% (-60·6 to 17·0)	1849099	21736	0% (ref)
							Partially vaccinated†	94761	420	49·3% (43·3 to 54·6)	11701310	37802	57-9% (56-9 to 58-9)
							0–1 week after second dose	47252	78	77-7% (71-9 to 82-3)	1601585	2688	73-2% (71-9 to 74-5)
							2–3 weeks after second dose	55318	85	83.7% (79.7 to 87.0)	1492259	1095	86-4% (85-4 to 87-3)
							4–5 weeks after second dose	65 698	106	86-6% (83-6 to 89-0)	1338 063	1019	83-5% (82-3 to 84-7)
							6-7 weeks after second dose	71120	134	86-8% (84-2 to 88-9)	1117983	1019	77-9% (76-1 to 79-5)
							8–9 weeks after second dose	73540	245	79-0% (75-9 to 81-7)	862 976	863	75-6% (73-4 to 77-6)
							10-11 weeks after second dose	73212	280	79-6% (76-8 to 82-1)	651213	751	69-3% (66-3 to 72-1)
							12-13 weeks after second dose	71773	337	77-4% (74-6 to 80-0)	445 924	646	60-8% (56-6 to 64-6)
							14-15 weeks after second dose	68114	356	75·9% (72·9 to 78·6)	264128	472	59-7% (54-6 to 64-2)
							16-17 weeks after second dose	63 974	402	70-5% (67-0 to 73-7)	169692	397	50-5% (43-4 to 56-6)
							18-19 weeks after second dose	58608	508	63.7% (59.6 to 67.4)	132 459	275	42-2% (32-4 to 50-6)
							20–21 weeks after second dose	45716	598	53·6% (48·4 to 58·3)			
							Scotland reference group: unvaccinat deprivation, comorbidities, number of from the analysis. In Brazil, vaccine eff and temporal trend. †Partially vaccina Table 2: Vaccine effectiveness esti	f previous tests, in fectiveness was ad ited: ≥2 weeks afte imates for ChAd	iterval between do justed for age, sex, er the first dose and	ses, and temporal trend; individ deprivation, macroregion of re I before the second dose.	luals positive for SA sidence, primary re	ARS-CoV-2 before eason for vaccinat	e Dec 8, 2020, were excluded tion, interval between doses,
							vaccination in Scotland and Brazi	Scotland			Brazil		
								Total samples	Positive sample	es Vaccine effectiveness* (95% CI)	Total samples	Positive samp	eles Vaccine effectiveness* (95% CI)
							Unvaccinated	26130	13 698	0% (ref)	9852053	4920001	0% (ref)
							0–1 week after first dose	911	374	20-9% (8-2 to 31-9)	286 322	151328	-9-6% (-10-5 to -8-8)
							Partially vaccinated†	15714	7176	37-6% (34-6 to 40-5)		398717	37-6% (37-3 to 37-9)
							0–1 week after second dose	5027	2025	50-2% (46-7 to 53-5)	112391	30550	51-3% (50-6 to 52-0)
							2–3 weeks after second dose	7141	2429	67-9% (65-9 to 69-8)	95671	7963	69-8% (69-3 to 70-4)
							4–5 weeks after second dose	8947	3387	67-3% (65-3 to 69-1)	79 298	15 568	68-4% (67-8 to 68-9)
							6–7 weeks after second dose	10622	4346	63-8% (61-7 to 65-7)	60301	12 401	66-8% (66-1 to 67-5)
							8–9 weeks after second dose	11258	4633	63-3% (61-3 to 65-3)	44351	9424	65-4% (64-6 to 66-2)
							10–11 weeks after second dose	14043	6319	59-3% (57-2 to 61-4)	32 832	7103	63-2% (62-2 to 64-2)
							12–13 weeks after second dose	17300	7966	55-3% (53-0 to 57-5)	22 454	5177	58-8% (57-4 to 60-1)
							14-15 weeks after second dose	17421	7670	52-9% (50-4 to 55-2)	15305	3435	59-8% (58-2 to 61-4)
							16-17 weeks after second dose	15442	6554	48-7% (45-9 to 51-4)	10 822	2529	58-7% (56-7 to 60-5)
							18–19 weeks after second dose	14 403	6248	44·6% (41·5 to 47·6)	7458	1852	57-7% (55-4 to 60-0)
							20–21 weeks after second dose		4718	39·1% (35·4 to 42·6)	,430	2032	377.2 (33.4 (0.00-0)
							*In Scotland, vaccine effectiveness w board, interval between doses, and to immunosuppression, cardiac disease appendix 2 (pp 11–15). †Partially vac	as adjusted for age emporal trend. In 8 pregnancy, puerp cinated: ≥2 weeks a	e, sex, deprivation, Brazil, vaccine effect eral period, chronic after the first dose	comorbidities, number of at-risl tiveness was adjusted for age, so it kidney disease, and temporal to and before the second dose.	ex, deprivation, mad rend. Descriptive ch	croregion of resid paracteristics for t	lence, diabetes, obesity, the sample are available in
							Table 3: Vaccine effectiveness est vaccination in Scotland and Braz				! symptomatic in	fection by leng	th of time since two-dose
78	Abu-Raddad et al (December 16, 2021	Qatar	General population	Alpha→Beta →Delta	mRNA-1273	January 1 and December 5, 2021	TND study linkin	g admir	nsitrativ	e databases.			
	Updated January 26,2022)												











ſ								Table. Change	in Estimated Me	ssenger RNA Va	ccine Effectiveness A	Against Labor	ratory-Confirme	ed SARS-CoV-2
									nuary to Septemb	per 2021				
								Month			y month from full vaccin			Contombor
1						1		Month 1	Pre-Delta (Janua 94.5 (90.7-96.7		Rising Delta (May to J 92.1 (87.2-95.1)		gh Delta (July to 2.0 (45.6-73.5)	
						1		2	88.5 (86.1-90.5		90.6 (87.8-92.7)		0.9 (51.5-68.4)	
J								3	87.9 (85.9-89.5		87.3 (80.8-91.7)		7.8 (52.5-62.5)	
								4	NA		86.6 (83.0-89.5)		3.3 (33.5-42.7)	
								5	NA		67.3 (63.2-70.9)	18	3.9 (13.7-23.8)	
								6	NA		NA		3.4 (13.3-23.3)	
								7	NA		NA		3.4 (17.3-29.0)	
								8	NA		NA	24	1.8 (18.8-30.4)	
								SARS-CoV-2 II	ated Messenger R nfection by Delta ptember 2021		ectiveness Against			
								1. tiveness, %		.	Pre-DeltaHigh DeltaRising Del	a –		
								Vaccine effec	T _	Ĭ Ĭ	Į Į Į			
								0	1 2 3 Mon	4 5 iths after full vacci		9		
-	76	Machado et al (December 14,	Portugal	Non-institutionalized 65-<110 year olds	Alpha, Delta	Comirnaty mRNA-1273	February 2 (80+) or March 30 (65-79) -	Cohort st	udy linking	administr	rative databas	ses.		
		The second secon		05-<110 year olds		ChAdOx1		timing post	dise	ease	hospitalizati	ion	dea	iths
		2021)				CHAUOXI	August 2021	dose 2			65-79 years 80-<			
								14-41 days	79 (76-83) 7		95 (90-97) 83 (6		5 (88-98) 8	
								42-69 days	68 (64-71) 6					88 (78-94)
								70+ days			93 (86-96)		3 (87-96)	
ļ								70-97 days 98+ days	59 (53-64) 5 39 (29-48)	3 (43-62)	74 (6	60-84)	8	86 (78-91)
ļ								98+ days 98-123 days		50 (40-59)	74 (5	58-83)	R	30 (71-86)
						1		124+days		34 (29-48)		37-78)		75 (64-82)
									AZ disease					
						1		timing post						
						1		dose 2 14-41 days	year olds 48 (42-54)					
								42-69	33 (23-42)					
						1		70+	34 (10-52)					
						1								
ŀ	<i>7</i> 5	Florea et al	USA	≥18 year olds Kaiser	NonVOC,	mRNA-1273	December 18, 2020-	Cohort st	udv					
	,3	(December 14, 2021)	33/1	Permanente insured patients	Alpha, Delta		September 30, 2021	201101131	····· y					
		(updated April 28, 2022)												





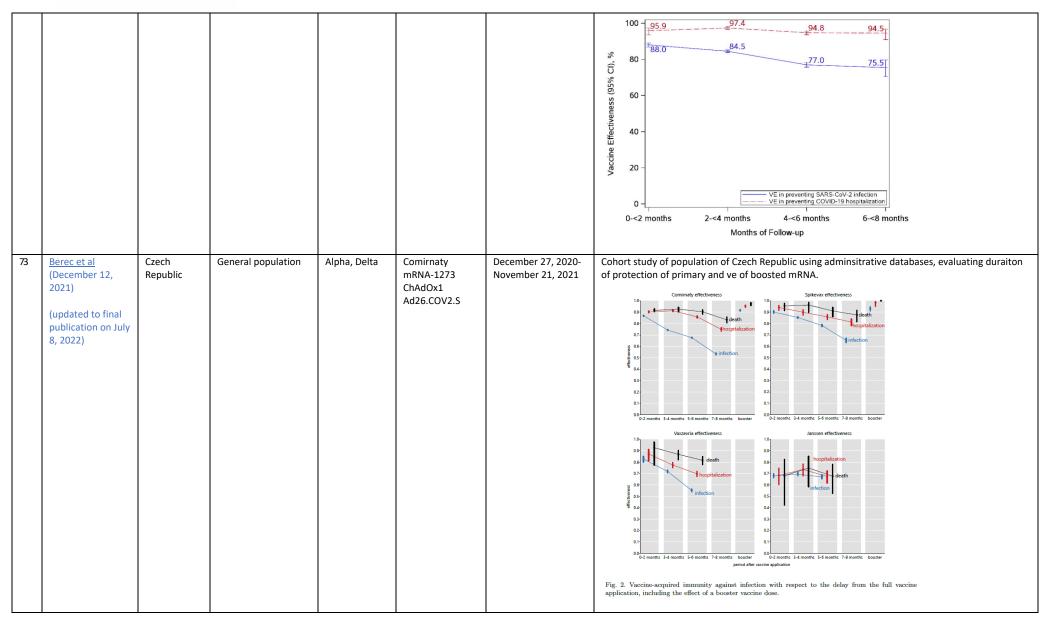


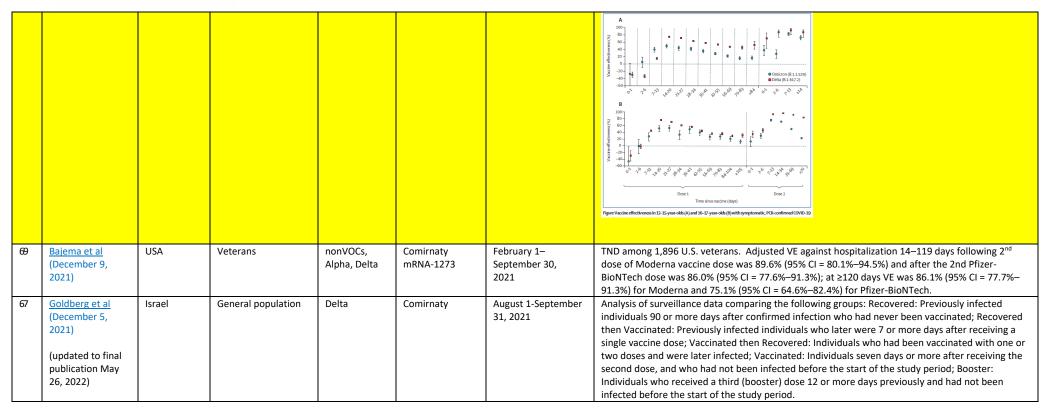




							Table 1. Estimated increase of breakthrough infection hazard ratios (HRs) in times of the SARS-CoV-2 delta variant dominance for age groups having started vaccination in the same month. Vaccine
72	Bjork et al (December 9, 2021) (Updated March 2, 2022)	Sweden	General population	Alpha, Delta	Comirnaty mRNA-1273 ChAdOx1	March 8-November 7, 2021	Case-control study based on surveillance data, matching on age/sex and no adjustment for other confounders. Infection Vaccine type, at least two doses Pfizer BioNTech AstraZeneca
71	Kshirsagar et al (December 9, 2021)	USA	Fully vaccinated persons	NonVOCs, Alpha, Delta	Comirnaty mRNA-1273 Ad26.COV2.S	March 10-October 14, 2021	Cohort study of fully vaccinated persons evaluating risk of reinfection by vaccination. There was an increase in the rate of hospitalization starting ~110-125 days after full vaccination for all three vaccines depending on age group, with a steeper increase for Janssen.
70	Powell et al (February 18, 2022) (updated May 2022)	UK	General population with a focus on adolescents	Delta, Omicron	Comirnaty	Week 32 (~Aug 15) (16-17 yo) and Week 37 (12-15 yo) - January 12, 2022	TND study among adolescents against symptomatic disease

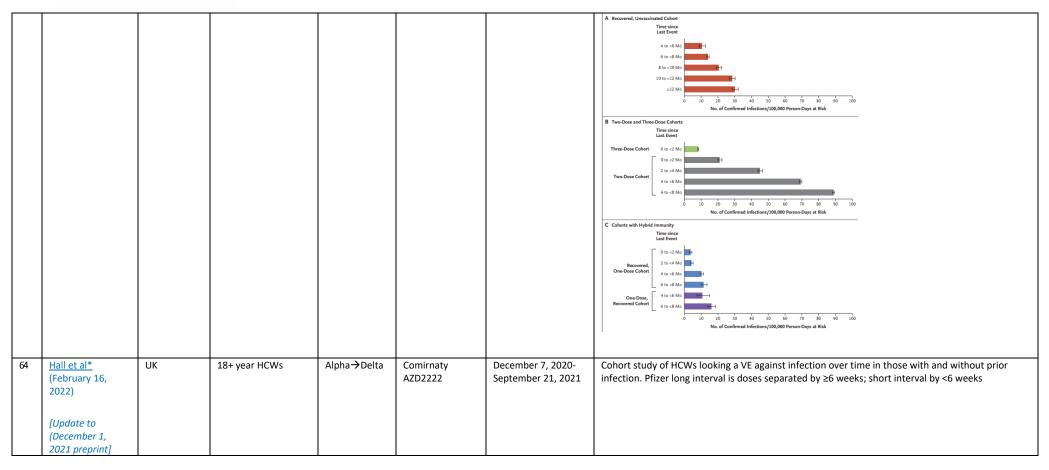
















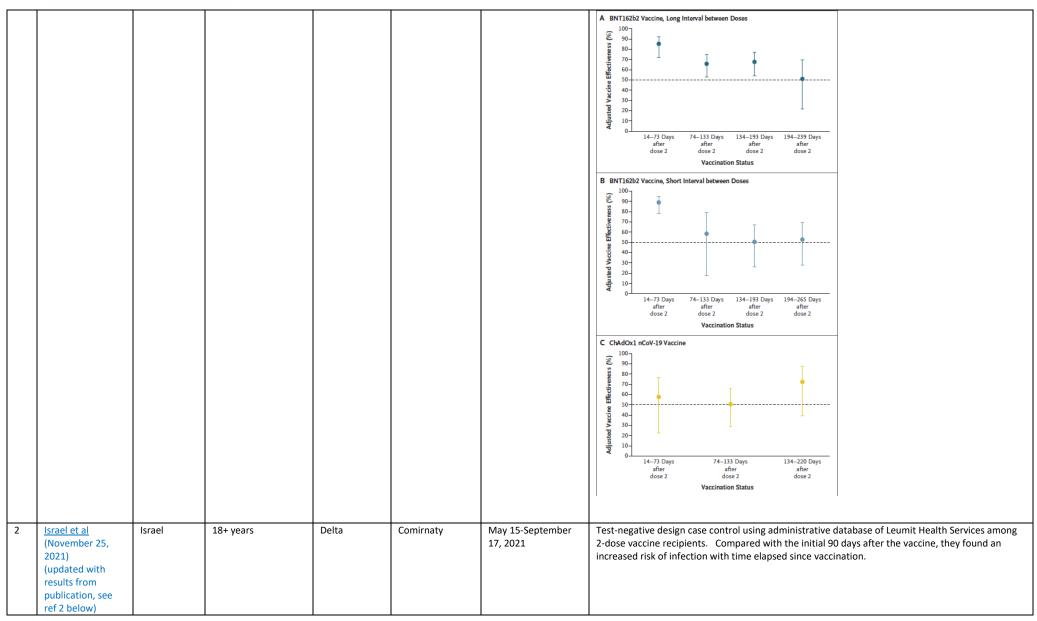


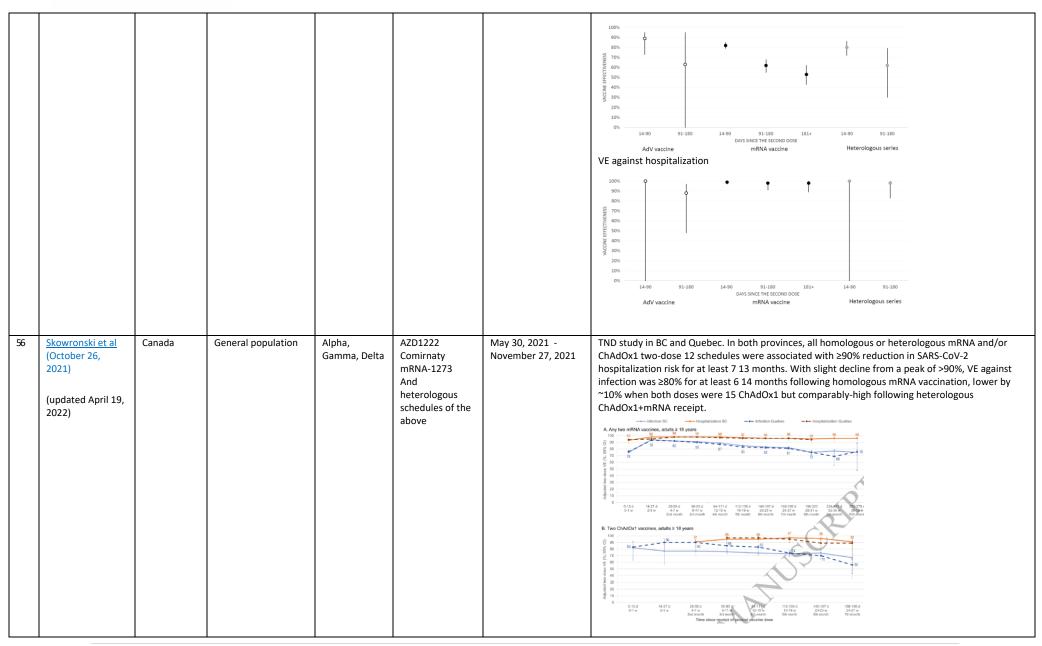




							Table 4 Adjusted odds ratios for risk of SARS-CoV-2 in matched cohort
							Adjusted odds ratio (95% CI) P value
							Time since second vaccine (days):
							21-89 Reference —
							90·119 2.37 (1.67 to 3.36) <0.001 120·149 2.66 (1.94 to 3.66) <0.001
							120-149 2.00 (1.94 to 3.00) 0.001 150-179 2.82 (2.07 to 3.84) (0.001
							≥180 2.82 (2.07 to 3.85) <0.001
							Age (continuous in years) 1.01 (1.00 to 1.01) 0.008
							Male sex 1.05 (0.99 to 1.11) 0.08
							Socioeconomic status (continuous 1-20) 0.97 (0.96 to 0.98) (0.001 Based on a conditional regression model fitted in a cohort matched for week of testing, age category (<18-39,
							40-59, ×60 years), and demographic group.
63	Irizarry et al	USA (Puerto	12+ years	Predelta and	Comirnaty	December 15, 2020-	Analysis of surveillance data linked to immunization registry data. VE against B) Infection c)
	(November 19,	Rico)		delta	mRNA-1273	October 15, 2021	Hospitalizations D) death by time since 2 weeks post complete series completion. Shading
	2021)				Ad26.COV2.S		represents 99% CI.
							B C D
							9 6 75%
							ctiv
							£ 50%
							95 25%
							> 0%
							Days since fully vaccinated
							Vaccine — mRNA-1273 — BNT162b2 — Ad26.COV2.S
61	Andrews et al	UK	50+	Delta	Comirnaty	September 13-	TND booster dose study that also calculated the VE of a 2 nd dose >140 days after receipt of the 2 nd
	(November 15,				AZD2222	November 1, 2021	dose. VE against symptomatic diseaes for two doses of ChAdOx1-S and BNT162b2 ≥20 weeks after
	2021)						being given were 44.1% (41.9 to 46.1) and 62.5% (61.0 to 63.9), respectively.
59	Tenforde et al	USA	Hospitalized patients	Mix, alpha,	Comirnaty	March 11-August 15,	Case-control study among hospitalized patients. When the mRNA-1273 and BNT162b2 vaccines
	(November 4,			and delta	mRNA-1273	2021	were compared, estimated vaccine effectiveness was similar within 120 days of vaccination. In
	2021)						
							contrast, beyond 120 days, the results corresponded to an estimated effectiveness of 85% for the
							contrast, beyond 120 days, the results corresponded to an estimated effectiveness of 85% for the mRNA-1273 and 64% for the BNT162b2 vaccine to prevent COVID-19 hospitalizations.
							mRNA-1273 and 64% for the BNT162b2 vaccine to prevent COVID-19 hospitalizations.
							mRNA-1273 and 64% for the BNT162b2 vaccine to prevent COVID-19 hospitalizations. Vaccine core Vaccine core
							mRNA-1273 and 64% for the BNT162b2 vaccine to prevent COVID-19 hospitalizations. Vaccinate care Vaccina
							mRNA-1273 and 64% for the BNT162b2 vaccine to prevent COVID-19 hospitalizations. Vaccined care prevent covid pr
							mRNA-1273 and 64% for the BNT162b2 vaccine to prevent COVID-19 hospitalizations. Variable Court Variabl
							mRNA-1273 and 64% for the BNT162b2 vaccine to prevent COVID-19 hospitalizations. Variable Court Variabl
							The contract case The
							mRNA-1273 and 64% for the BNT162b2 vaccine to prevent COVID-19 hospitalizations.
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							mRNA-1273 and 64% for the BNT162b2 vaccine to prevent COVID-19 hospitalizations.
58	Poukka et al	Finland	16-69 year old HCWs	Mix and delta	Comirnaty	December 27 2020-	mRNA-1273 and 64% for the BNT162b2 vaccine to prevent COVID-19 hospitalizations.
58	Poukka et al	Finland	16-69 year old HCWs	Mix and delta	Comirnaty	December 27,2020-	mRNA-1273 and 64% for the BNT162b2 vaccine to prevent COVID-19 hospitalizations.
58	(November 4,	Finland	16-69 year old HCWs	Mix and delta	mRNA-1273	August 26 (infection)	mRNA-1273 and 64% for the BNT162b2 vaccine to prevent COVID-19 hospitalizations.
58		Finland	16-69 year old HCWs	Mix and delta	mRNA-1273 AZD2222	August 26 (infection) October 26	mRNA-1273 and 64% for the BNT162b2 vaccine to prevent COVID-19 hospitalizations.
58	(November 4,	Finland	16-69 year old HCWs	Mix and delta	mRNA-1273	August 26 (infection)	mRNA-1273 and 64% for the BNT162b2 vaccine to prevent COVID-19 hospitalizations.







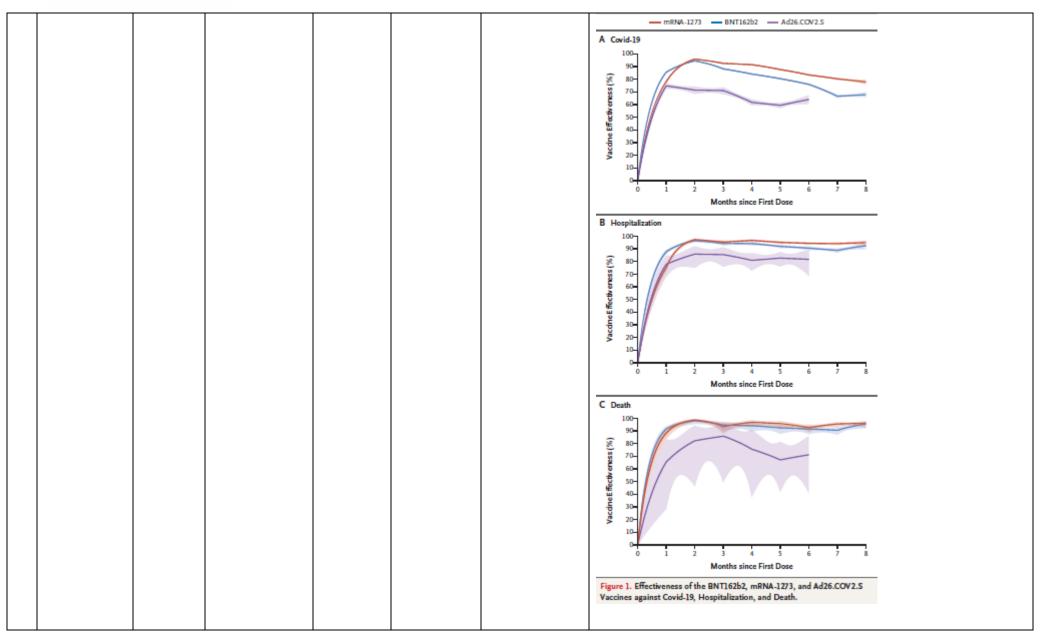




55	<u>Lin et al</u> (October 26,	USA	General population	multiple	Comirnaty mRNA-1273	December 13, 2020- Sept 8, 2021	Administrative database cohort study in North Carolina. For Pfizer two-dose,VE peaks at 94.5% (95% CI, 94.1 to 94.9) at 2 months (post the first dose). VE starts to decline after 2 months and
	2021)				Ad26.COV2.S	33,232	drops to 66.6% (95% CI, 65.2 to 67.8) at 7 months. For Moderna two-dose,VE peaks at 95.9% (95% CI, 95.5 to 96.2) at 2 months. Effectiveness started to decline after 2 months and was
	[updated with						maintained at 80.3% (95% CI, 79.3 to 81.2) at 7 months. For the Janssen one-dose regimen,
	final publication						vaccine effectiveness ramps to a peak level of 74.8% (95% CI, 72.5 to 76.9) at 1 month.
	on January 12,						Effectiveness started to decline after 1 month and decreased to 59.4% (95% CI, 57.2 to 61.5) at 5
	2022}						months.











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54	Nordstrom et al	Sweden	General population	Alpha, Delta,	AZD1222	January 12-October	National cohort study based on database linkage. Vaccine effectiveness of BNT162b2 against
	(October 25,				Comirnaty	4, 2021	infection waned progressively from 92% (95% CI, 92-93, P<0.001) at day 15-30 to 47% (95% CI, 39-
	2021)				mRNA-1273		55, P<0.001) at day 121-180, and from day 211 and onwards no effectiveness could be detected
					And		(23%; 95% CI, -2-41, P=0.07). The effectiveness waned slightly slower for mRNA-1273, being
	[Updated				AZD1222à		estimated to 59% (95% CI, 18-79) from day 181 and onwards. In contrast, effectiveness of
	February 4, 2022}				mRNA-1273		ChAdOx1 nCoV-19 was generally lower and waned faster, with no effectiveness detected from day
							121 and onwards (-19%, 95% CI, -97-28), whereas effectiveness from heterologous ChAdOx1 nCoV-
							19 / mRNA was maintained from 121 days and onwards (66%; 95% CI, 41-80). Overall, vaccine
							effectiveness was lower and waned faster among men and older individuals. For the outcome
							severe Covid-19, effectiveness waned from 89% (95% CI, 82-93, P<0.001) at day 15-30 to 42% (95%
							CI, -35-75, P=0·21) from day 181 and onwards, with sensitivity analyses showing notable waning
							among men, older frail individuals, and individuals with comorbidities.
52	Hulme et al	UK	HCW	Alpha, delta	Comirnaty	January 4-June 13	Comparative VE Cohort study of HCWs based on linking databases who were vaccinated with
1	(October 18,			1 .,	AZD1222	, , , , , , , , , , , , , , , , , , , ,	AZD1222 or Comirnaty between January 4-February 28, 2021 who were followed for 20 weeks.
	2021)						Figure 2: Comparative effectiveness For each outcome based on the fully adjusted model, the marginal
							cumulative incidence for ChAdOx1 and BNT162b2, their difference, and the hazard ratio are shown. Models
							that assumed piecewise-constant hazards gave similar effect estimates (supplementary Figure S2). The models with less extensive confounder adjustment gave very similar estimates (supplementary Figure S1)
							suggesting that recipients of each vaccine were similar after accounting for differences in vaccine allocation over space and time (as did all models).
							Positive SARS-CoV-2 test COVID-19 A&E attendance COVID-19 hospitalisation 25 1.25
							20 0.75
							a pue
							mendan 15 0.75 0.50
							P 2 20 CD 15 0.50 0.50 0.50 0.50
							5 On/dox1 0.25
							0 28 56 84 112 140 00 0 28 56 84 112 140 00 0 28 56 84 112 140
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							10 10 N N N N N N N N N N N N N N N N N
							02 02 02
							0.7 0.28 56 84 112 140 0 28 56 84 112 140 0 28 56 84 112 140
							Days since first dose
51	Robles-Fontan et	USA (Puerto	General population	Multiple, with	Comirnaty	December 15,2020-	Cohort study of Puerto Rican population.
	al	Rico)		delta time	mRNA-1273	October 15, 2021	
	(October 18,	'		frame analysis	Ad26.COV2.S	,	
	2021)						
	/						
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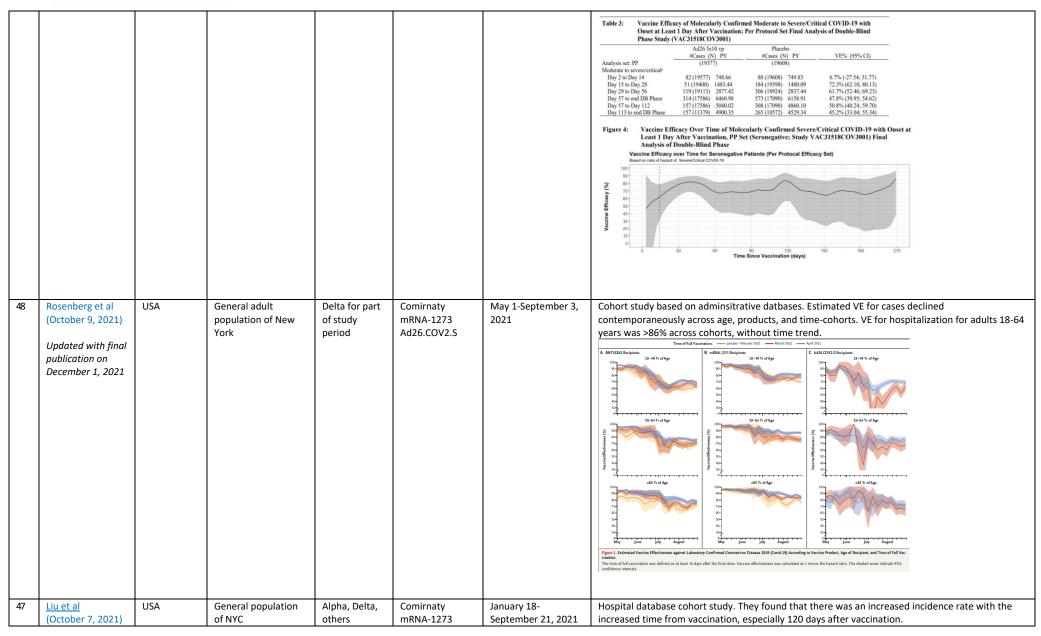




	(updated March 2,						Outcome	Vaccine	Effectiveness o	n first day as fully va	ccinated (CI) E	Effectiveness after 144 days (0	1),
	2022)						Infection	mRNA-1273	90% (88-91%)		7	72% (69—75%)	
	2022)						Infection	BNT162b2	87% (85-88%)			54% (51-57%)	
							Infection	Ad26.COV2.S	64% (58-69%)			36% (31-42%)	
							Hospitalization	mRNA-1273	95% (89-97%)		9	91% (84-95%)	
							Hospitalization	BNT162b2	92% (86-95%)		8	81% (74-86%)	
							Hospitalization	Ad26.COV2.S	82% (61-91%)		6	67% (54-77%)	
							Death	mRNA-1273	99% (89-100%)	1	9	93% (81-97%)	
							Death	BNT162b2	97% (87-99%)			86% (76-92%)	
							Death	Ad26.COV2.S	78% (14-94%)		7	73% (49-86%)	
50	De Cienetal	Netherlands	Constal association	Delta	Comingto	Avenut O Contamban		ffectiveness against in				:	
50	De Gier et al	Netherlands	General population	Deita	Comirnaty	August 9-September							valuate transmission.
	(October 14,				mRNA-1273	24, 2021	They did no	ot have suffic	ent sample	e size but ev	valuated if VE	E against transm	ission differed by time
	2021)				Ad26.COV2.S		since vacci	nation of the	index case				
	,				AZD1222								
					ALDIZZZ							ination of the contact	
											oup of the index cas ull vaccination of the		
							week of notif	ication date of the i	idex case, stratili	ed by time since i	uli vaccination or the	le index case.	
							Analysis	Unvaccinated	Index fully	Index fully	Index fully	Index fully vaccinated	
								index - infected contacts / all	vaccinated < 60 days ago -	vaccinated < 60 days ago -	vaccinated >= 60 days ago - infected	>= 60 days ago - adjusted VET (%)(95%	
								contacts (SAR)	infected	adjusted VET	contacts / all	CI)	
									contacts / all contacts (SAR)	(%) (95% CI)	contacts (SAR)		
							Unvaccinated	547/2517 (22%)	24/209 (11%)	67 (47;79)	14/94 (15%)	55 (19;76)	
							household						
							contacts	164/1505 (11%)	99/1278 (8%)	57 (40;69)	157/792 (20%)	28 (-4;50)	
							vaccinated	104/1505 (1170)	33/12/0 (0/0)	37 (40,03)	1377732 (2070)	20 (-4,50)	
							household						
							contacts						
49	Janssen Briefing	multiple	General population	Multiple	Ad26.COV2.S	September 21, 2020-	Final result	s from BCT					
7.7		manupic	Scheral population	manipic	, 1020.CO V2.3								
	document for US FDA					July 9, 2021	ì	with Onset at Least Final Analysis of Do	1 Day After Vac uble-Blind Phas	cination, PP Set (Seronegative; Stud	ere/Critical COVID-19 ly VAC31518COV3001)	
	(October 14,						Vacci Based of	ine Efficacy over Time on ratio of hazard of Moderate	o for Seronegative to SeverelCritical COVID-1	Patients (Per Proto	ocal Efficacy Set)		
	2021)						100						
	2021/						90-						
							80 70						
							8 60-						
							\$\vec{\vec{\vec{\vec{\vec{\vec{\vec{	1					
							30-						
							20-					~	
							10-						
								30	60	90 120	150	180 210	
									Time	Since Vaccination	95% noin	intwise CI; 95% of events prior to day 189.	
1											Last event: d Base	day 229; Hazard smoothed over 21 days. ed on the methods in Gilbert et al. (2002).	
	ı		IL.	L	1	1	L						

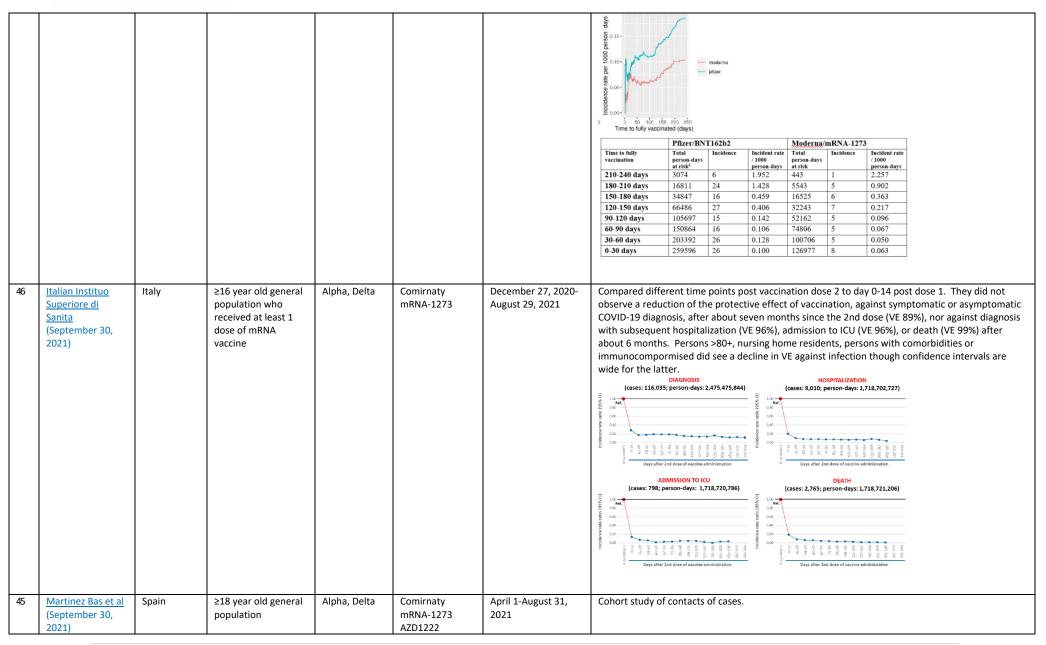












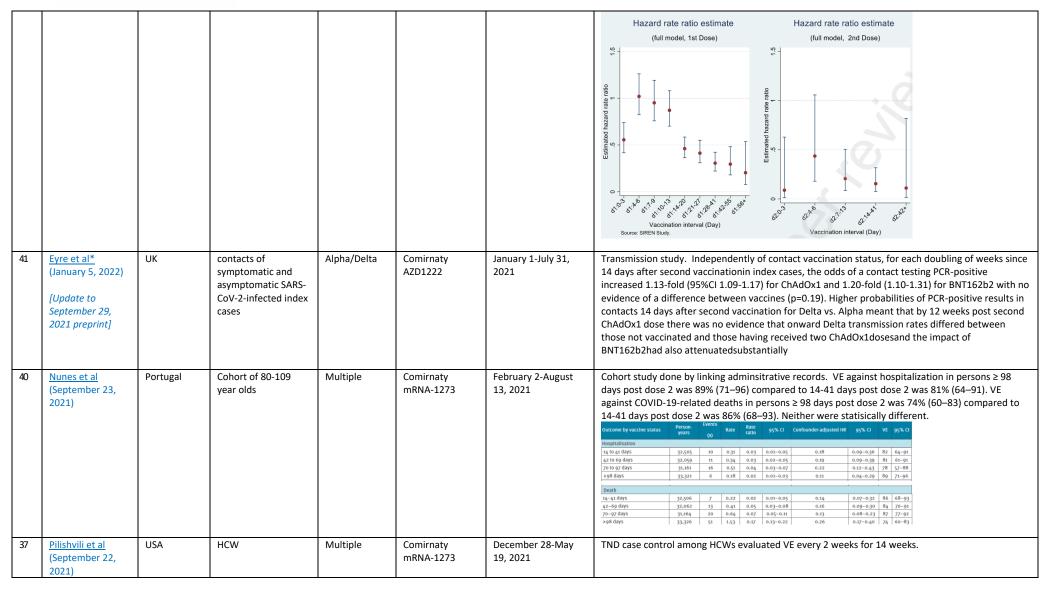




Wind					Ad26.COV2.S		Adjust VE (9	95% CI)	
Lidose of January S2 (44-59) 28 (8-53) 1 1 1 1 1 1 1 1 1						</td <td>90 days since last dose ≥</td> <td>90 days since last dose</td> <td></td>	90 days since last dose ≥	90 days since last dose	
1 1 1 1 1 1 1 1 1 1							REF	REF	
2 2 2 2 2 2 2 2 2 2									
dose of Cominaty 57(51-61) NA Alpha Comirnaty 2 doses of Comiraty 70(67.73) 63 (84-88) 2 doses of Comiraty 70(67.73) 63 (84-88) 2 doses of Vazervia 40 (31-47) 52 (37-64) 2 doses of Vazervia 54 (47-60) NA Alpha									
2 does of Commany 70 (67-73) 63 (58-68) 1 does of Vaxervia 40 (31-47) 52 (37-64) 2 does of Vaxervia 40 (31-47) 52 (37-64) 2 does of Vaxervia 54 (47-60) NA 1 does of Vaxervia 1 does of Vaxervia 54 (47-60) NA 1 does of Vaxervia 1 does of Commany 85 (69-93) NA									
Lidese of Vaxzervia 40 (31-47) 52 (37-64) NA 2 (dose of Vaxzervia 40 (31-47) 54 (47-65) NA 2 (dose of Vaxzervia 40 (47-65) NA 4 (dose of									
Bruxvoort et al (October 1, 2021) USA General population Delta, Alpha+others March 1-July 27, 2021 USA General population Delta, Alpha+others March 1-July 27, 2021 TND study among persons insured by Kaiser Permante Southern California.									
44 Bruxvoort et al (October 1, 2021) USA General population Delta, Alpha+others March 1-July 27, 2021 TND study among persons insured by Kaiser Permante Southern California. 100-100-100-100-100-100-100-100-100-10									
44 Bruxvoort et al (October 1, 2021) USA General population Delta, Alpha+others all (October 1, 2021) USA General population Delta, Alpha+others all (October 1, 2021) Delta, Alpha+others all (October 1, 2021) Warch 1-July 27, 2021 TND study among persons insured by Kaiser Permante Southern California. Variant Homo Delta Ho									
Alpha+others Alpha+others 2021 Alpha+others Alpha Comirnaty December 7, 2020- Cohort study of HCWs						1 dose of Vaxzervia+1 dose of Comirnaty	85 (69-93)	NA	
43 Payne et al UK HCWs Alpha Comirnaty December 7, 2020- Cohort study of HCWs	44	USA	General population		mRNA-1273		by Kaiser Permante	Southern Califorr	ia.
						Variant Delta Non-Delta Unidentified 14-60 days 61-90 days Time		ys 151-160 days	
	43	UK	HCWs	Alpha	Comirnaty	Cohort study of HCWs			











							88 80 70 60 50 10 10 16 24 23 35 24
36	El Sahly et al (September 22, 2021)	USA	RCT participants	Multiple	mRNA-1273	July 27, 2020-March 26, 2021	No. of Controls 541 213 156 137 99 139 88 Findings from the double blinded placebo controlled RCT. VE against disease was similar at 2 weeks-<2 months (91.8%), 2 months-<4 months (94%), and ≥4 months (92.4%) post dose 2
35	Baden et al (September 22, 2021)	USA	≥18-year-old RCT participants	Delta	mRNA-1273	July 1-August 27, 2021	RCT participants were followed after unblinding. Initial vaccine recipients (mRNA-1273e) were vaccinated between 7/27/20-12/16/20 while those vaccinated after unblinding (mRNA-1273p) were vaccianted between 12/29/20-4/30/21. Median follow-up times from the first dose were 13 months in the mRNA-1273e (including double-blind and open-label phases) and 7.9 months in the mRNA-1273p (only open-label phase) groups. While there was a significant difference in disease incidence rates between the groups, there was no difference in severe disease incidence rates
							though numbers are small. mRNA-1273e mRNA-1273p* mRNA-1273p vs
							N=14746 N=11431 mRNA-1273e
							Cases† n yr Person-yr n yr Person-yr incidence rate % (95% CI) All cases 162 2102 77.1 88 1796 49.0 36.4 (17.1-51.5)
							≥18-<65 136 1558 87.3 68 1289 52.8 39.6 (18.6-55.5)
							≥65 yr 26 544 47.8 20 507 39.5 17.4 (-53.9-56.3) Severe 13 2102 6.2 6 1796 3.3 46.0 (-52.4-83.2)
							≥1865 7 1558 4.5 4 1289 3.1 30.9 (471.7-85.2)
							yr 265 yr 6 544 11.0 2 507 3.9 64.2 (-100.2-96.5)
34	Hagan et al	USA	Incarcerated persons	Delta	Comirnaty	July 11-August 14,	Outbreak investigation in a prison found that the attack rate among fully vaccinated persons was
	(September 21, 2021)		sarceratea persons	25114	mRNA-1273 Ad26.COV2.S	2021	significantly higher in those vaccinated 4-6 months ago (89%) compared to those vaccinated 2 weeks-2 months ago (61%). This was combined for 3 vaccines used in the population.
33	Thomas et al	Multiple	≥12-year-old RCT	Multiple	Comirnaty	July 27, 2020-March	Findings from the double blinded placebo controlled RCT. VE against disease was 96.2% (93.3-98.1)
	(September 15,		participants			13, 2021	at 7 days-<2 months, 90.1% (86.6-92.9) at 2 months-<4 months, and 83.7% (74.7-89.9) at ≥4
	2021)						months post dose 2.

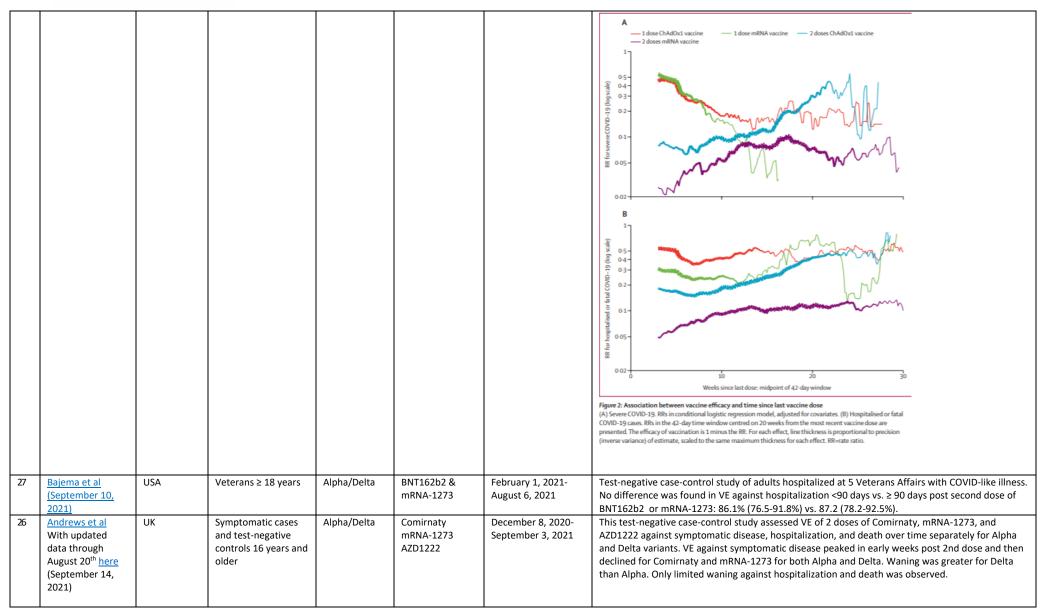




							Efficacy End Point No. of Surveillance No. at No. of Surveillance Operation No. of Surveillance No. at No. of No.
							27 Days after receipt of second dose 82 6.649 22,132 889 6.371 22,001 91.2 (83.9 to 93.0) 27 Days after receipt of second dose to -2 no after 12 2.923 22,112 312 2.884 2.20.01 82.6 31.5 to 98.1) 27 No after receipt of second dose to -4 no after 46 2.696 20,814 449 2.593 20,344 90.1 (86.6 to 92.9) 34 No after receipt of second dose 24 1.030 12,690 128 0.895 11,802 83.7 (74.7 to 89.9 t)
32	Pfizer (September 17, 2021)	Multiple	≥16-year-old RCT participants	Delta	Comirnaty	July 1-August 31, 2021	RCT participants were evaluated for duration of protection against symptomatic disease, with the original placebo recipients receiving the vaccine after unblinding. The mean time from Dose 2 of Comirnaty to 01 July 2021 was approximately 5 months for the crossover group and 10 months for the original group. There was a 26.3% (7.4%- 41.4%) relative vaccine efficacy for the group vaccinated later (crossover group) compared to the group vaccinated earlier (original group), with a difference in incidence rates of -18.6 per 1000 person-years of follow-up.
31	de Gier et al (September 17, 2021)	Netherlands	Hospitalized patients	Delta (just for duration of protection)	Comirnaty mRNA-1273 Ad26.COV2.S AZD1222	July 4-August 29, 2021 (just for duration of protection)	Incidence rate ratios were calculated based on national coverage and vaccination status of hospitalized cases. All 4 vaccines were combined in calculating the VE by time since vacciantion, and VE was only calculated during the delta dominant period when 99% of sequenced isolates were delta. No drop in VE against hospitalization nor in VE against ICU admission was seen between those vaccinated up to 20 weeks since full vacciantion among 15-49, 50-69, ≥70 year olds.
30	Self et al (September 17, 2021)	USA	≥18 years who were hospitalized at 21 U.S. hospitals across 18 states	Alpha, Delta, Non-VOC	Comirnaty mRNA-1273 Ad26.COV2.S	March 11–August 15, 2021	This case-control study found that the for mRNA-1273 vaccine, there was no difference in VE against hospitalization among those were 14-120 days post full vaccination and those who were >120 days post full vaccination. For Comirnaty, VE against hopsitalization was 91% (88-93) for those 14-120 days post full vaccination while it was 77% (67-84) for those >120 das post full vaccination. Ad26.COV2.S did not have enough data to stratify by more than 28 days post full vaccination.
29	Polinski et al (September 12, 2021) (updated March 17, 2022)	USA	≥18 years of age	Alpha/Delta	Ad26.COV2.S	March 1, 2021- August 31, 2021	Retrospective cohort study used insurance claims data linked to health data sources to evaluate VE of Ad26.COV2.S against COVID-19 diagnosis and hospitalization among vaccinated individuals and matched unvaccinated individuals (matched on age, sex, comorbid-risk, calendar date, location, and other risk factors for COVID-19 severity). VE was stable over time up to 152 days after vaccination.
28	McKeigue et al (September 15, 2021) (updated February 25, 2022)	Scotland	Population of Scotland	Alpha/Delta	Comirnaty mRNA-1273 AZD1222	December 1, 2020- September 8, 2021	Matched case-control study (REACT-SCOT) assessed rate ratios over time comparing rate of severe COVID-19 and the rate of hospitalization or death among thoswe full vaccinated with Comirnaty, mRNA-1273, and AZD1222 to unvaccinated persons.

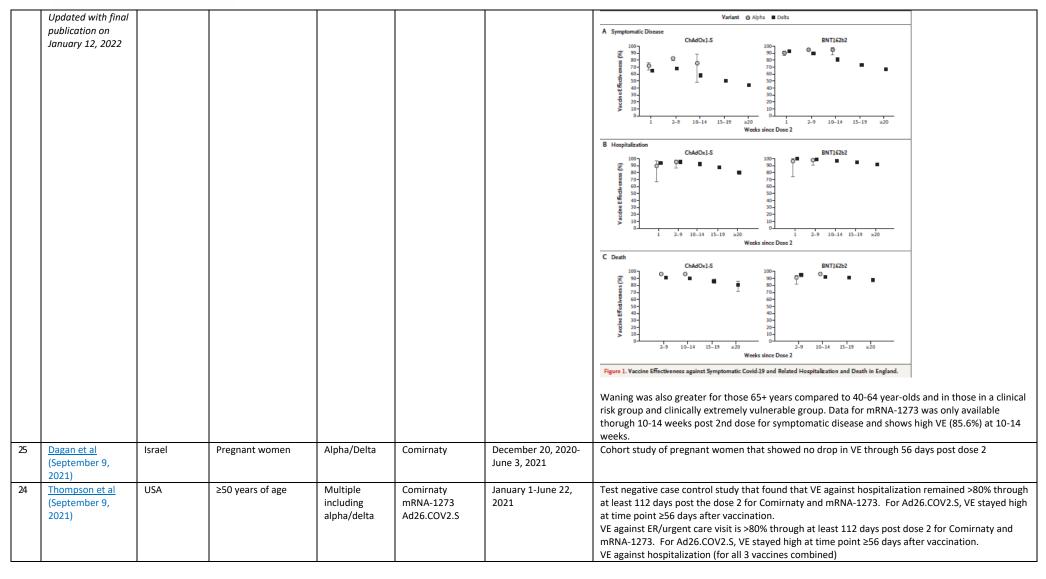












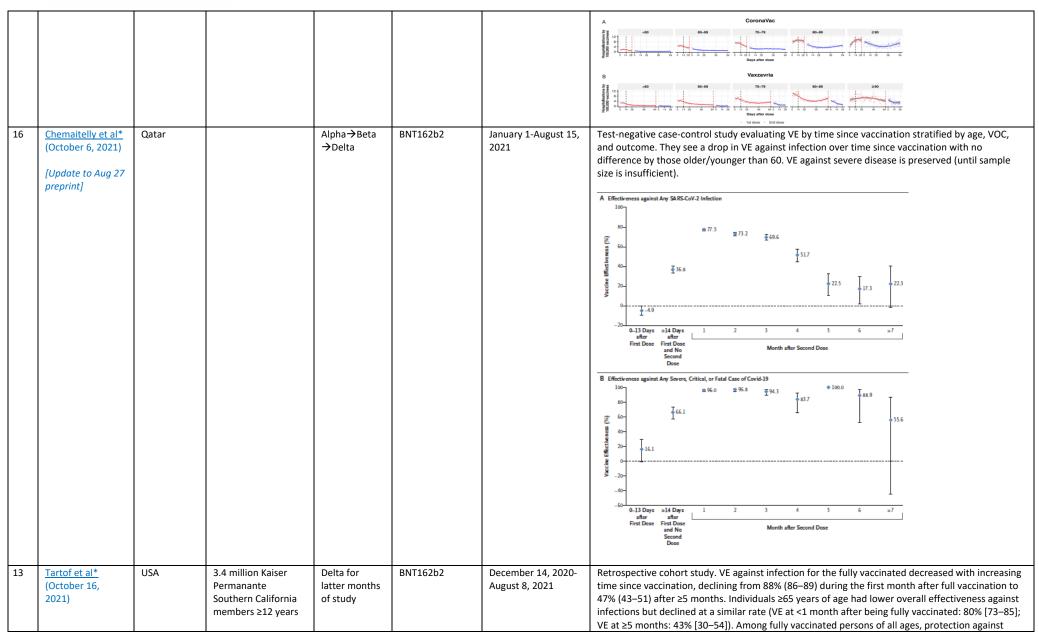




							Fully vaccinated — 14–27 Days after 28–41 Days after 42–55 Days after 42–55 Days after 70–83 Days after 84–97 Days after 84–97 Days after 112 Days after 112 Days after 124–17 Days after 14–27 Days after 42–55 Days after 42–55 Days after 44–97 Days after 84–97 Days after 84–111 Days after 84–111 Days after	r dose 2 2 r dose 2 1 r dose 2 1 greated a 2 r dose 2 1 greated a 2 greated a 3 gre	,198 23 (1.9) ,170 20 (1.7) ,067 18 (1.7) 924 28 (3.0) 667 24 (3.6) 487 13 (2.7) 331 17 (5.1) 221 11 (5.0)	I→I 88 (84 to 92)
23	Puranik et al	USA	Persons ≥14 days	Multiple	Comirnaty	January 1-August 8,	Test negativ	ve case control		duration of protection against symptomatic disease.
	(September 7,		post dose 2 ("full	including	,	2021	_		•	60 after full vaccination.
	2021)		vaccination") who	alpha/delta			Covariate	Level/Category	Symptomatic Int	ection
			received first dose after January 1				Time Relative to Full	Day 0	1 (Reference	e)
							vaccination	Day 30	2.19 (0.89, 5.	36)
								Day 60	3.65 (1.78, 7.	46)
								Day 90	5.58 (2.72, 11	.46)
								Day 120	7.25 (3.47, 15	.18)
								Day 150	10.33 (5.03, 21	.24)
22	Kertes et al	Israel	Fully vaccinated	Delta	Comirnaty	June 9-July 18, 2021	,			days post dose 2 by June 9 and had no history of prior
	(September 7, 2021)		population							anuary-February had odds of infection of 1.61 (1.45- rch-May of testing positive for SARS-CoV-2.
19	Keehner et al	USA	~19,000 employees	Delta	BNT162b2	July -August 26, 2021	· · ·			g symptomatic cases occurring in July, HCW vaccinated in
	(September 1,		of University of		mRNA-1273	, , ,	January or F	, February had ar	n attack rate of	6.7 per 1000 persons (95% CI, 5.9 to 7.8), whereas the
	2021)		California San Diego				attack rate	was 3.7 per 100	00 persons (95%	5 CI, 2.5 to 5.7) among those who completed vaccination
			Health						U	y. Among unvaccinated persons, the July attack rate was
		_							% CI, 11.8 to 22.	,
18	Nunes et al	Portugal	1.5 million ≥65 year olds	Alpha→Delta	BNT162b2 mRNA-1273	?February-August 13, 2021				For those 80+, VE against hospitalization was 82 (64-91)
	(August 29, 2021)		(duration of		IIIKINA-12/3	2021		•		For COVID related mortality, it was 86% (68-93) at day mitations are that data delays could mean that outcomes
			protection on only						•	been recorded for more recent cases. Additionally, only
			those 80+)						•	ted during the study period, making these unvaccinated
									different from t	
17	Cerqueria-Silva et	Brazil	75.9 million	Gamma	CoronaVac	January 18-July 24,		•	•	calculated VE, as well as evaluated the daily
	<u>al</u>		vaccinated in Brazil		AZD1222	2021		•	•	tinees. For CoronaVac, there was low hospitalization
	(August 27, 2021)								•	79 years old. 80-89 and ≥90 age groups lowest creased but were still lower than 1 dose recipients

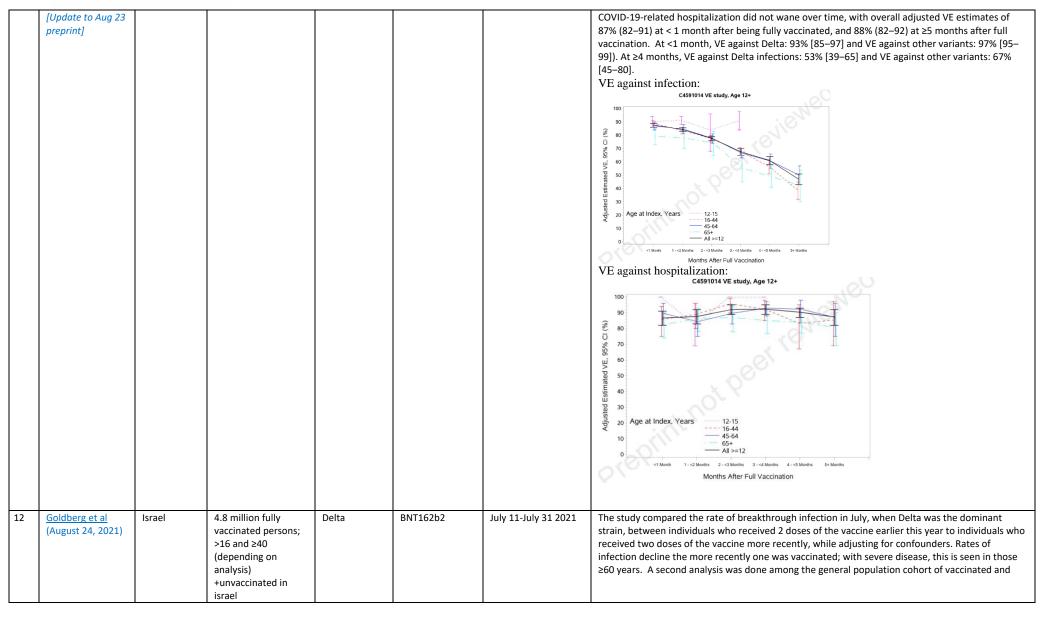












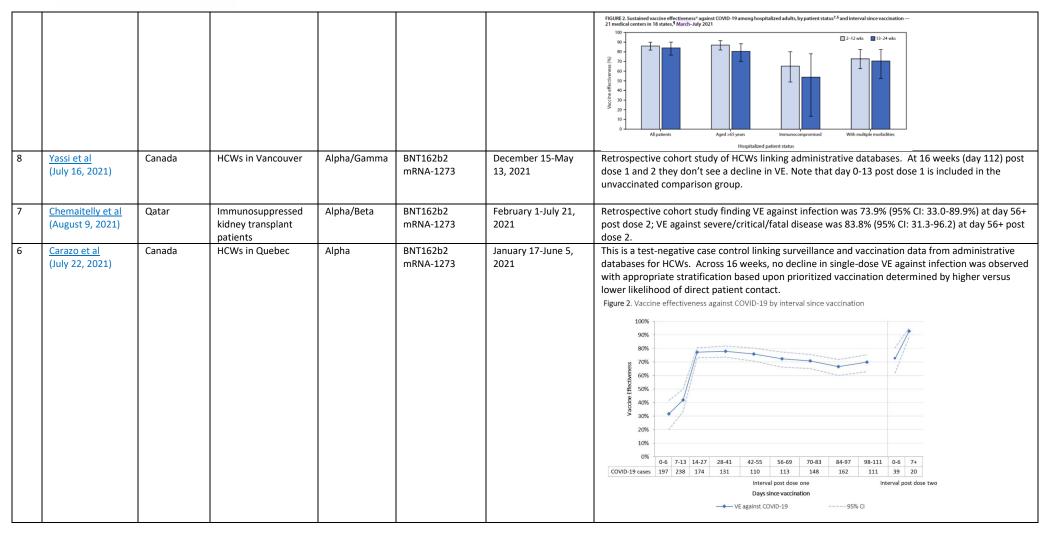




							unvaccinated to calculate VE by age group and month of vaccination. OUTCOME = Positive SARS-Cov.2 PCR test
							Age JanB FebA FebB MarA MarB Apr May
							16-39 50% [45,55] 47% [42,52] 58% [55,62] 62% [59,64] 68% [65,70] 74% [71,77] 73% [67,78]
							40-59 58% [54, 62] 61% [58, 65] 63% [59, 66] 67% [63, 70] 74% [70, 77] 78% [73, 82] 80% [71, 86]
							60+ 57% [52, 62] 63% [57, 67] 65% [57, 71] 73% [66, 78] 72% [64, 77] 73% [63, 81] 75% [58, 85]
							OUTCOME = Severe COVID-19
							Age Jan Feb Mar
							40-59 94% [87, 97] 98% [95, 99] 98% [94, 99]
							60+ 86% [82, 90] 88% [84, 91] 91% [85, 95]
10	Pouwels et al* (October 14, 2021) [Update to Aug 18 preprint]	UK	General adult population	Alpha, Delta	BNT162b2 AZD1222	December 1, 2020- August 1, 2020	COVID-19 infection survey is a household longitudinal survey with testing. During the delta dominant period, in those 18 to 64 years, VE of BNT162b2 against new PCR-positives reduced by 22% (95% CI 6% to 41%) for every 30 days from second vaccination. Reductions were numerically smaller for ChAdOx1 (change -7% per 30 days, 95% CI -18% to +2%) but there was no formal evidence of heterogeneity (p=0.14). Overall BNT162b2 ChAdOx1 Days since 14 days after 2nd dose
9	Tenforde et al (August 18, 2021)	USA	Hospitalized patients	Alpha→Delta	BNT162b2 mRNA-1273	March 11-July 14, 2021	Test-negative design case control study of hospitalized patients. VE against COVID-19– associated hospitalization was 86% (95% CI = 82%–90%) 2–12 weeks and 84% (95% CI = 77%–90%) 13–24 weeks from receipt of the 2^{nd} dose, with no significant change between these periods (p = 0.854). There was no difference in VE by timing since vaccine among those \geq /< 65 years, immunocompromised versus not and among those with \geq /< 3 chronic conditions.

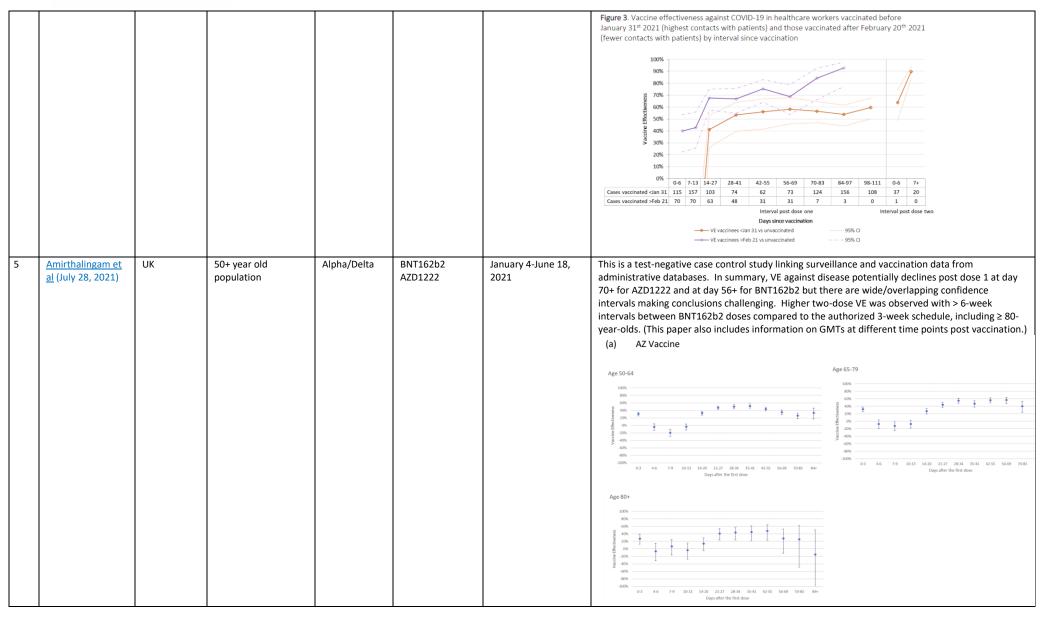






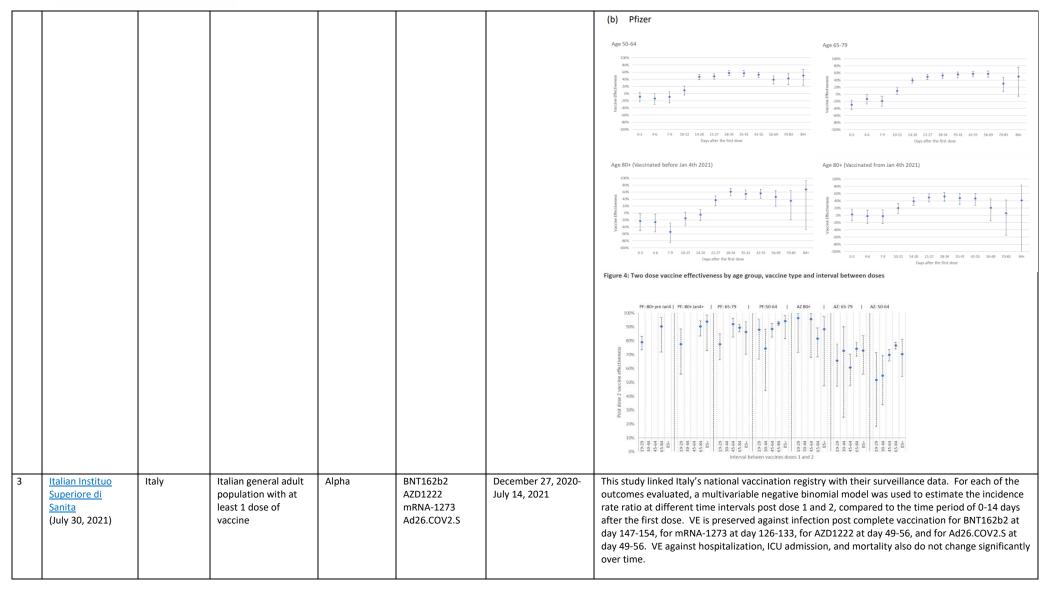






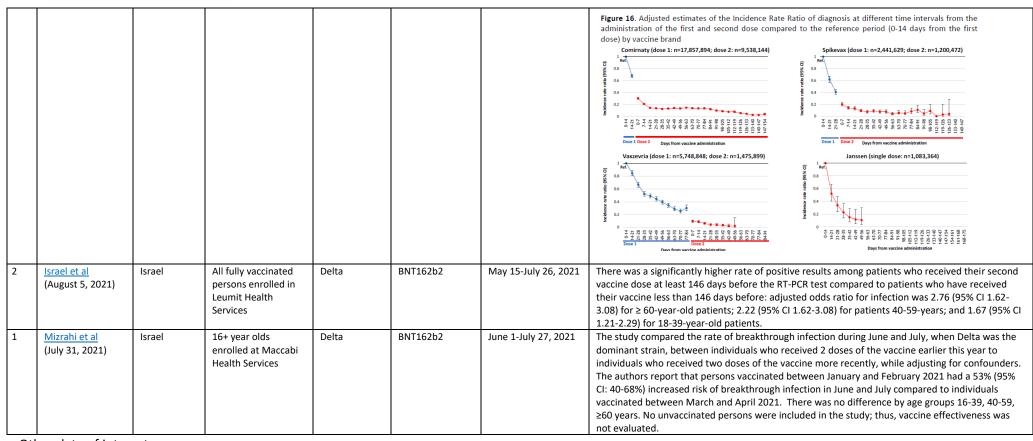












Other data of interest:

- https://www.gov.il/BlobFolder/reports/vpb-12082021/he/files_publications_corona_vpb-12082021-01.pdf
- Salo et al HH transmission study in Finland, showing VE 10 weeks after 1 dose of an mRNA vaccine but is a mix of 1 and 2 dose recipients.
- Pfizer's press announcement of 4 month efficacy in adolescents https://www.pfizer.com/news/press-release-detail/follow-data-phase-3-trial-pfizer-biontech-covid-19-vaccine

Note as of January 7, 2022 version, only true duration of protection analyses are included. Please look at the <u>update</u> from December 30, 2021 if you wish to see full list of previously included studies with other data such as Kaplan-Meier curves. Missing reference numbers in table above indicate studies that have been removed.