## Vaccine efficacy reported in trials

Vaccine efficacies (VE) reported in phase III clinical trials depend on the outcome, time interval, number of received doses and subgroups. The overview in table 1, lists the overall VE estimates after full vaccination focused on the prespecified time interval after full vaccination as described in the study protocol.

Table 1. Vaccine efficacy (VE) after complete vaccination against symptomatic SARS-CoV-2 infections, hospitalization, death and severe COVID19 for the COVID19vaccines developed by Pfizer, Moderna, AstraZenca and Janssen.

	Vaccine											
	Pfizer <sup>£</sup>			Moderna <sup>£</sup>			AstraZeneca <sup>£,α</sup>			Janssen <sup>£</sup> (one dose scheme)		
Outcome	Time interval	VE [ref]	95% CI	Time interval	VE [ref]	95% CI	Time interval	VE [ref]	95% CI	Time interval	VE [ref]	95% CI
Symptomatic SARS-CoV-2 infection	≥7 days after d2	95,0 [ <u>Polack</u> ]	(90,3- 97,6)	≥14 days after d2	94,1 [ <u>Baden</u> ]	(89,3- 96,8)	≥15 days after d2	62.1 [ <u>Voysey]</u>	(41.0- 75.7)	≥14 days after vaccinati on	66.9 [ <u>FDA</u> J]	(59.0- 73.4)
	≥14 days after d2	94,2 [ <u>EPAR P</u> ]	(88,7- 97,2)							≥28 days after vaccinati on	66.1 [ <u>EMA J]</u>	(55.0- 74.8)
		-		1			1					
	-	Pfizer		Moderna			AstraZeneca			Janssen (one dose scheme)		
Outcome	Time interval	VE [ref]	95% Cl	Time interval	VE [ref]	95% CI	Time interval	VE [ref]	95% CI	Time interval	VE [ref]	95% CI
Hospitalization							≥15 days after d2	100 [ <u>EPAR AZ</u> ]	(42.65, NE)	≥14 days after vaccinati on	81.8 [ <u>FDA</u> <u>J</u> ]	(16.7- 98.0)
										≥28 days after vaccinati on	100 [ <u>FDA</u> <u>J</u> ]	(74.3- 100.0)
				1			1					
	Pfizer			Moderna		AstraZeneca			Janssen (one dose scheme)			
Outcome	Time interval	VE [ref]	95% Cl	Time interval	VE [ref]	95% CI	Time interval	VE [ref]	95% CI	Time interval	VE [ref]	95% CI

Death										≥14 days	80.0 [FDA	(29.4,
										after	J]	96.3)
										vaccinati	-	
										on		
										≥28 days	75 [FDA J]	(-25.2,
										after		97.4)
										vaccinati		
										on		
		Pfizer			Moderna			AstraZeneca		Jansser	n (one dose s	cheme)
Outcome	Time	Pfizer VE [ref]	95% Cl	Time	Moderna VE [ref]	95% CI	Time	AstraZeneca VE [ref]	95% CI	Jansser Time	n (one dose s VE [ref]	cheme) 95% Cl
Outcome	Time interval		95% CI	Time interval		95% CI						
Outcome Severe COVID19			95% CI (-124,8-			95% CI (NE,	Time			Time		
	interval	VE [ref]		interval	VE [ref]		Time interval	VE [ref]	95% CI	Time interval	VE [ref]	95% CI
	interval ≥7 days	VE [ref] 66,4 [FDA	(-124,8-	interval ≥14 days	VE [ref] 100	(NE,	Time interval ≥15 days	VE [ref] 100.0	95% CI (-	Time interval ≥14 days	VE [ref] 76.7 [FDA	95% CI (54.6,
	interval ≥7 days	VE [ref] 66,4 [FDA	(-124,8-	interval ≥14 days	VE [ref]	(NE,	Time interval ≥15 days	VE [ref] 100.0	95% CI (- 3742.53,	Time interval ≥14 days after	VE [ref] 76.7 [FDA	95% CI (54.6,
	interval ≥7 days	VE [ref] 66,4 [FDA	(-124,8-	interval ≥14 days	VE [ref]	(NE,	Time interval ≥15 days	VE [ref] 100.0	95% CI (- 3742.53,	Time interval ≥14 days after vaccinati	VE [ref] 76.7 [FDA	95% CI (54.6,
	interval ≥7 days	VE [ref] 66,4 [FDA	(-124,8-	interval ≥14 days	VE [ref]	(NE,	Time interval ≥15 days	VE [ref] 100.0	95% CI (- 3742.53,	Time interval ≥14 days after vaccinati on	VE [ref] 76.7 [ <u>FDA</u> <u>J</u> ]	95% CI (54.6, 89.1)
	interval ≥7 days	VE [ref] 66,4 [FDA	(-124,8-	interval ≥14 days	VE [ref]	(NE,	Time interval ≥15 days	VE [ref] 100.0	95% CI (- 3742.53,	Time interval ≥14 days after vaccinati on ≥28 days	VE [ref] 76.7 [FDA J] 85.4 [FDA	95% CI (54.6, 89.1) (54.2,

\*Complete vaccination reflects two doses for the vaccines developed by Pfizer, Moderna and AstraZeneca and one dosis for the vaccine developed by Janssen. <sup>e</sup>Median age in years of the study populations are 52 (16-91) [Polack], NR [18-70+] [Voysey] and 53 (18-100) [FDA J] for the vaccines developed by Pfizer, AstraZeneca and Janssen respectively. Participants in the trial of the Moderna vaccine had a mean age of 51.4 (range 18-95) year [Baden]. "Standard dose – standard dose scheme. NE, not estimated since there were 0 cases in the intervention or placebo arm. NR, not reported.

## Vaccine effectiveness reported in post-marketing studies

Note: no post-marketing studies have been found for the Moderna and Janssen vaccines. VEs estimates after full vaccination in post-marketing studies are limited for the vaccines developed by Pfizer and AstraZeneca. In table 2, VEs are reported in time intervals that are as similar as possible to those used in phase III trials, i.e.  $\geq$ 7 days or  $\geq$ 14 days after dose 2. Additionally, VE estimates after one dose are provided if no relevant information after two doses is available. VEs after one dose are listed for time intervals around one or two weeks after dose 1 and the maximum time interval after dose 1 provided in the study.

Table 2. Vaccine effectiveness (VE) obtained from preprints (unless stated otherwise) against symptomatic SARS-CoV-2 infections, hospitalization, death and severe
COVID19 for the COVID19- vaccines developed by Pfizer and AstraZeneca.

		Pfize	r		AstraZeneca				
Outcome	Population [ref]	Time interval	VE	95% CI	Population	Time interval	VE	95% Cl	
Symptomati c SARS-CoV- 2 infection	Israel, ≥16 yr [ <u>Chodick</u> *]	13-24 days after dose 1	51.4	(-7.2-78)	England, ≥70 yr [ <u>Lopez</u> ]	7-9 days after d1	-3	(-16, 1)	
	Israel, ≥16 yr [ <u>Hunter</u> *]	21 days after dose 1	91	(83-98) <sup>°</sup>	England, ≥70 yr [ <u>Lopez]</u>	14-20 days after d1	22	(11-32)	
	Israel, ≥16 yr [ <u>Dagan<sup>β</sup>]</u>	≥7 days after d2	94	<mark>(</mark> 87-98)	England, ≥70 yr [ <u>Lopez]</u>	≥35 days after d1	73	(27-90)	
	England, >80 yr [ <u>PHE</u> ]	≥7 days after d2	88	(84-90)					
	England, ≥80 yr [Lopez]	≥14 days after d2	85	(79-89)					
	Israel, HCW [ <u>Amit</u> ¥]	1-14 days after d1	47	(17-66)					
	Israel, HCW [ <u>Amit</u> <sup>*</sup> ]	15-28 d after d1	85	(71-92)					
	Denmark, HCW, 47 (36-57) <sup>Σ</sup> [ <u>Moustsen]</u>	>14 days after d1 – d2	17	(4-28)					
	Denmark, HCW,	≥7 days after d2	90	(82-95)					

	47 (36-57) <sup>Σ</sup>									
	Moustsen									
	England, HCW,	≥7 days after d2	85	(74-96)						
	±45° [ <u>Hall</u> *]									
	Denmark, LTCF,	>14 days after	21	(-11-44)						
	<b>84 (77-90)</b> <sup>Σ</sup>	d1 – d2								
	Moustsen									
	Denmark, LTCF,	≥7 days after d2	64	(14-84)						
	84 (77-90) <sup>Σ</sup>									
	[Moustsen]									
			1.1							
		Pfiz			-	AstraZ				
	Population [ref]	Time interval	VE	95% CI	Population	Time interval	VE	95% CI		
Hospitalizati	Israel, ≥16 yr	≥7 days after d2	87	(55–100)	England, ≥80 yr	≥14 days after	37	(3-59)		
on	[Dagan <sup>B</sup> ]			And the second second	[Lopez]	d1		18.01.000.000.001.21		
	England, ≥80 yr	≥14 days after	71.4	(43.1-86.2)	England, ≥80 yr	≥14 days after	80.4	(36.4-94.5)		
	[Hyams]	d1			[Hyams]	d1				
	Scotland, adults	7-13 days after	38	(28-47)	Scotland, adults	7-13 days after	70	(63-76)		
	[Vasileiou]	d1			[Vasileiou]	d1				
	Scotland, adults	14-20 days after	60	(50-68)	Scotland, adults	14-20 days after	74	(66-81)		
	[Vasileiou]	d1			[Vasileiou]	d1		(=,)		
	Scotland, adults	42+ days after	64	(49-75)	Scotland, adults	28-34 days after	94	(71-99)		
	[Vasileiou]	d1			[Vasileiou]	d1				
	1	Pfiz			AstraZeneca					
	Population [ref]	Time interval	VE	95% CI	Population	Time interval	VE	95% CI		
Death	England, ≥80 yr	≥14 days after	51	(37-62)	Population	Time Interval	VE	93% CI		
Death	[Lopez]	d1	51	(57-02)						
	[ <u>copez</u> ]	ui								
		Pfiz	ər		AstraZeneca					
	Population [ref]	Time interval	VE	95% CI	Population	Time interval	VE	95% CI		
Severe	Adults ≥16 yr	≥7 days after d2	92	(75-100)	. epaideon					
COVID19	[Dagan <sup>B</sup> ]									
I										

\*SARS-CoV-2 infection irrespective of symptoms. °90% Credible interval. <sup>®</sup>No preprint. Peer reviewed publication in the New England Journal of Medicine. <sup>\*</sup>No preprint. Correspondence of the Lancet. <sup>I</sup>Median age at first vaccination dose (interquartile range). <sup>o</sup>Median age. D1, dose1; D2, dose 2; HCW, Health Care Workers; LCTF, Long Term Care Facility residents.