## Global spotlight 15.2: Key additions for the second half of March 2022



There is one newly added synthesis and seven updates to living evidence syntheses that are already included in the public-health measures parts of the COVID-END inventory of 'best' evidence syntheses\*, as well as two newly added syntheses and eight updates to living evidence syntheses that are already included in the clinical management parts of the inventory.

\*COVID-END assigns 'best' status to evidence syntheses based on an assessment of how up-to-date they are (i.e., the date of the last search, with priority given to living reviews), quality (using the AMSTAR tool), and whether there is an evidence profile available (e.g., GRADE).

Taxonomy section	Title	Type of synthesis	Criteria for best evidence synthesis		
			Date of last search	Quality (AMSTA R) rating	Evidence profile (e.g., GRADE) available
Public-health measures	Evidence suggests that face coverings may reduce the transmission of SARS-CoV-2 with no serious harms, while medical masks appear to have higher efficacy than fabric masks [Review of studies of mainly low quality]	Newly added living rapid review	2021-07-15	4/9	No
Public-health measures	[BioNTech/Pfizer against variants of concern] BNT162b2 [Pfizer] vaccine may prevent infection from the Omicron variant of concern up to 44 days and may provide limited protection up to 60 days after the second dose; it may also prevent symptomatic infection up to 63 days after the second dose, and may provide limited protection up to 90 days after the second dose (other variants are also included in the report)	Update to living rapid review	2022-03-16	7/9	Yes
Public-health measures	[BioNTech/Pfizer against variants of concern] Three doses of BNT162b2 [Pfizer] vaccine may prevent infection from the Omicron variant of concern up to 30 days after the third dose, it may prevent infection up to 60 days, and it may provide limited protection after 90 days of the third dose; it may also provide strong protection against severe, critical, or fatal disease produced by the Omicron variant of concern (other variants are also included in the report)	Update to living rapid review	2022-03-16	7/9	Yes
Public-health measures	[Johnson & Johnson against variants of concern] Johnson & Johnson/AD26.COV2.S vaccine may provide limited protection from infection from the Omicron variant of concern up to 60 days after the second dose (other variants are also included in the report)	Update to living rapid review	2022-03-16	7/9	Yes
Public-health measures	[Johnson & Johnson against variants of concern] One dose of Johnson & Johnson/AD26.COV2.S vaccine followed by a second dose of an mRNA vaccine may prevent infection from the Omicron variant of concern at least 7 days after the second dose (other variants are also included in the report)	Update to living rapid review	2022-03-16	7/9	Yes

Public-health	[Moderna against variants of concern] mRNA-1273	Update to living	2022-03-16	7/9	Yes
measures	[Moderna] vaccine may	rapid review	2022-03-10	119	100
	provide limited protection for infection from the	·I · · · · · ·			
	Omicron variant up to 44 days, 60 days and 90 days				
	after the second dose, and it may prevent				
	symptomatic infection up to 30 days after the second				
	dose (other variants are also included in the report)				
Public-health	[Moderna against variants of concern] Three doses of	Update to living	2022-03-16	7/9	Yes
measures	Moderna vaccine may prevent infection from the	rapid review			
	Omicron variant of concern up to 30 days, and up to				
	60 days after the third dose, and it may prevent severe,				
	critical, and fatal disease death up to 42 days after the				
	third dose (other variants are also included in the				
	<u>report)</u>				
Public-health	[Oxford/AstraZeneca against variants of concern]	Update to living	2022-03-16	7/9	Yes
measures	ChAdOx1 [AstraZeneca] vaccine may prevent	rapid review			
	infection caused by the Omicron variant of concern at				
	up to 60 days after the second dose (other variants are				
	also included in the report)	NT 1 11 1	2022 02 10	40/44	37
Clinical management of COVID-19 and	[Ciclesonide] In COVID-19 outpatients, ciclesonide	Newly added living review	2022-03-18	10/11	Yes
pandemic-related	<u>may make little or no difference in hospitalization or</u> <u>death while it may slightly increase adverse events; its</u>	living review			
health issues	effects on other outcomes are uncertain				
Clinical management	The effects and cost-effectiveness of home	Newly added	2021-06-09	4/9	No
of COVID-19 and	monitoring using pulse oximetry in people with	living rapid review	2021-00-09	4/ 9	110
pandemic-related	<u>COVID-19 symptoms are currently uncertain</u>	inving rapid review			
health issues	, <u></u>				
Clinical management	[Aspirin] Among hospitalized patients, aspirin	Update to living	2022-03-18	10/11	Yes
of COVID-19 and	probably slightly reduces mortality and slightly	review			
pandemic-related	increases clinical improvement				
health issues Clinical management	Compared to therapeutic anticoagulant therapy,	The data to limite a	2022-03-18	10/11	Yes
of COVID-19 and	prophylactic anticoagulants in patients hospitalized	Update to living review	2022-03-16	10/11	1 68
pandemic-related	with COVID-19 may make little or no difference in	ic vic w			
health issues	mortality, and may probably slightly increases clinical				
	improvement; its safety outcomes are uncertain				
Clinical management	[Favipiravir] In COVID-19 hospitalized patients,	Update to living	2022-03-18	10/11	Yes
of COVID-19 and	favipiravir may make little or no difference in	review			
pandemic-related	mortality, clinical improvement and disease				
health issues	progression, while it may not have an effect on viral negative conversion among COVID-19 outpatients;				
	its safety outcomes and other effects on outpatients.				
	are uncertain				
Clinical management	[Hydroxychloroquine] In hospitalized patients,	Update to living	2022-03-18	10/11	Yes
of COVID-19 and	hydroxychloroquine probably does not have an effect	review			
pandemic-related	on mortality and clinical improvement, and it may				
health issues	make little or no difference in disease progression,				
	while it may increase adverse events; in outpatients, it may not have an effect on mortality, and it probably				
	makes little or no difference in viral negative				
	<u>conversion</u>				
Clinical management	In hospitalized patients, adding convalescent plasma	Update to living	2022-03-18	10/11	Yes
of COVID-19 and	to standard care probably does not have an effect on	review			
pandemic-related	mortality at 28 days and clinical improvement, while it				
health issues	may not have an effect on disease progression, and it				
	<u>may slightly increase serious adverse events; in</u>				
	outpatients, convalescent plasma may slightly reduce hospitalization or death and may not increase adverse				
	nospitalization of ucatif and may not increase adverse				

	events, while its effects on other outcomes are				
	uncertain				
Clinical management	[Lopinavir + ritonavir] Adding lopinavir + ritonavir to	Update to living	2022-03-18	10/11	Yes
of COVID-19 and	standard care probably makes little or no difference	review		-	
pandemic-related	on mortality, it may not have a substantial effect on				
health issues	clinical improvement, viral conversion or disease				
	progression, whereas it may slightly increase adverse				
	events				
Clinical management	[REGEN-COV2] In hospitalized COVID-19 patients,	Update to living	2022-03-18	10/11	Yes
of COVID-19 and	REGEN-COV2 (casirimab + imdevimab) may slightly	review			
pandemic-related	reduce mortality and disease progression, and it				
health issues	probably slightly increases clinical improvement, while				
	no information on safety outcomes is available; in				
	outpatients, it may slightly reduce hospitalization or				
	death and it may not increase serious adverse events,				
	while its effects on other outcomes are uncertain				
Clinical management	[Tofacitinib] Using tofacitinib in hospitalized patients	Update to living	2022-03-18	10/11	Yes
of COVID-19 and	may slightly reduce mortality, and it may slightly	review			
pandemic-related	increase clinical improvement, while it may also				
health issues	slightly increase severe adverse events				