## Updated4 April 2023

## **UN Medical Directors**



No additional safety concerns have been identified the use of heterologous schedule (either within the primary series or the booster dose).

Note that specific guidance or the administration of Moderna is provided in Annex A

Summary table 1:

ary table in			
DOSE #1	DOSE #2 <sup>13</sup>	DOSE #3/BOOSTER#1 <sup>14</sup>	DOSE#4 / BOOSTER#2 <sup>15</sup>
mRNA*	mRNA*	vectored	mRNA
mRNA*	vectored	mRNA	mRNA
vectored	vectored	mRNA	mRNA
vectored	mRNA*	mRNA	mRNA
Inactivated	inactivated	mRNAor vectored	mRNAor vectored
Inactivated	mRNAor vectored	mRNAor vectored	mRNAor vectored
non WHO EUL	non WHO EUL	After primary vaccination with WHO EU®	After primary vaccination ar booster dose with WHO E®
	DOSE #1  mRNA*  mRNA*  vectored  vectored  Inactivated  Inactivated	mRNA* mRNA* vectored vectored vectored vectored mRNA* Inactivated inactivated Inactivated mRNAor vectored	DOSE #1 DOSE #2 <sup>13</sup> DOSE #3/BOOSTER#1 <sup>14</sup> mRNA* wectored mRNA  vectored vectored mRNA  vectored mRNA  vectored mRNA  inactivated mRNA  mR



Annex A: UNMD recommendations for utilization of Moderna vaccine for the UN System-wide vaccination programme

## Disclaimer

- This guidance does not imply that Moderna is the only option for booster doses within the context of the COVID 19 vaccination series but refers specifically to situations where the Moderna vaccine is an available option.
- This advice is based on currently available evidence on the possible use of Moderna vaccine either as a primary SAR&COV2 immunization tool or as aximatch second dose of a primary immunization cycle (heterologous vaccination) or as a booster dose of a completed primary immunization cycle.
- Primarydoses can be monovalent/ancestral strain or bivalent ancestral/Omicron validated
- Booster doses cabe monovalent/ancestral strain or bivalent ancestral/Omicron variant
- Homologous vaccination (i.e., same vaccine) schedules are considered standard practice based on substantial safety, immunogenicity, and efficacy data available for each CO9/Laccinevith WHO EUL approval.
- Heterologous vaccination (i.e., mixed vaccines) with any two WHO EUL approved vaccines<sup>18</sup> is considered a
  complete primary series.<sup>19</sup>
- Where scientific evidence of above uses of Moderna vaccine has been judged weak or unavailable error peer review process has informed the technical recommendation.
- Advice regarding additional dosing for nb/HO EUL COVID vaccines is less well established and should be based on an individual risk assessment.
- As new information becomes available, the recommendations presented in this document will be adjusted accordingly.
- See table 2 below for details.

## Rules for use of Moderna

- Primary schedule age 12-17 and 18 and above: administerdose#1 and dose#2sfull dose monovalent1(00 μg) or bivalent original/Omicron BA!5
- Primary schedule age 6-11: administerdose#1 and dose#2shalf dose monovalent (5µg)
- Moderna as 2<sup>nd</sup> dose (and 3<sup>rd</sup> dose for selected immunocompromised individuals) **should be administered-8** weeks after the initial dose (week interval recommended for mRNA) primary vaccination to further increase effectiveness and further duce rare risk of myocardiffs.
- First booster doses age 12-17 and 18 and above: monovalent or bivalental dose \$0 µg) given 4 to 6 months (or less depending on local health authority policies) at the recompletion of the last dose.
- Subsequent booster doses for high risk grou4(pl)5(etET@0g00004]TJE) ca



Non-WHO	Sputnik,	Commence a new primary schedule at least 4 weeks after the last do
EUL	Soberana,	administered.
approved	Abdala	
vaccines		