#### Moderna's Respiratory Vaccines: COVID + Flu vaccine (mRNA-1073)

Last program update: November 4, 2021

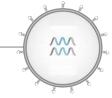
Modality	Program	ID#	Preclinical development	Phase 1	Phase 2	Phase 3	Commercial	Moderna rights
Adults	COVID-19 vaccine	mRNA-1273						Worldwide
		mRNA-1273.351	Beta variant					Worldwide
		mRNA-1273.617	Delta variant					Worldwide
		mRNA-1273.211	Beta variant + w	ild-type				Worldwide
		mRNA-1273.213	Beta + Delta variant					Worldwide
		mRNA-1283	Next generation (	2-5 °C)				Worldwide
<b>→</b>	Flu vaccine	mRNA-1010	Phase 1/2					Worldwide
		mRNA-1020						Worldwide
		mRNA-1030						Worldwide
Prophylactic	COVID + Flu vaccine	mRNA-1073	mRNA-1273 + mRNA-1010					Worldwide
vaccines	Older adults RSV vaccine	mRNA-1345			Phase 2/3 pre	p		Worldwide
Adolescents & Pediatrics	COVID-19 vaccine (adolescents)	mRNA-1273	TeenCOVE					Worldwide
	COVID-19 vaccine (pediatrics)	mRNA-1273	KidCOVE					Worldwide
	Pedialic kyv vaccine	mRNA-1345	Phase 1b					Worldwide
	Pediatric hMPV + PIV3 vaccine	mRNA-1653						Worldwide
	Pediatric RSV + hMPV vaccine	mRNA-1365						Worldwide

## Announcing two new combination vaccines are progressing towards the clinic





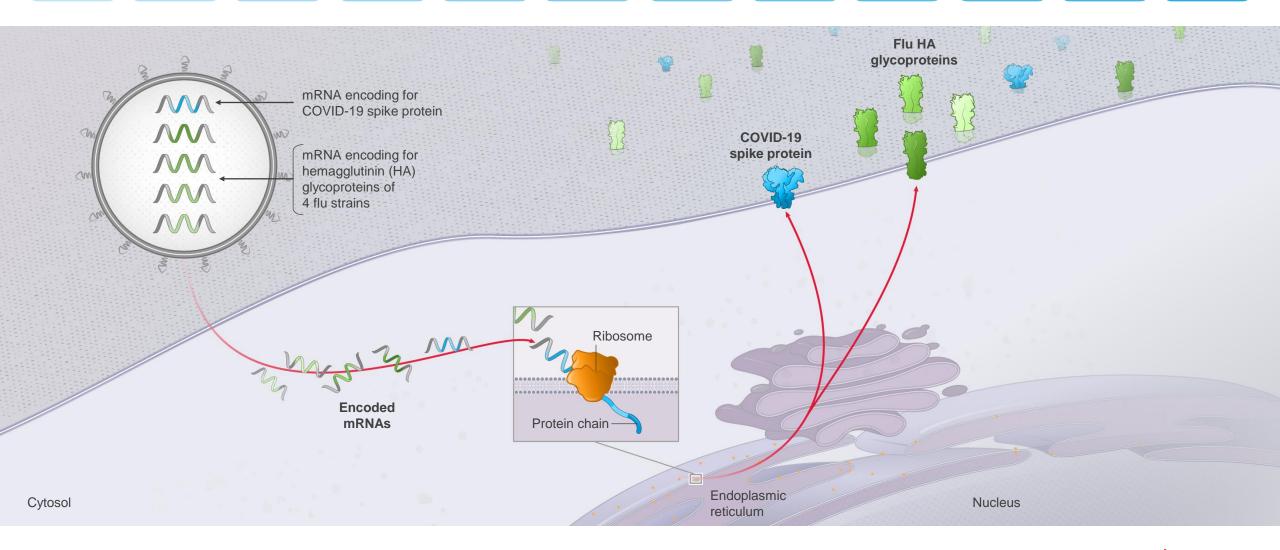




Combination mRNA vaccines against acute respiratory infections



# mRNA-1073 encodes for the COVID-19 spike protein and Flu HA glycoproteins



### We are developing combination vaccines against the respiratory viruses with greatest public health impact

**Developing vaccines** against respiratory viruses with greatest public health impact











PIV3



More viruses...

**INFLUENZA** 

SARS-CoV-2

**RSV** 

**HMPV** 

**Combination vaccines** may lead to:

- Higher compliance
- Better uptake
- Larger benefit to healthcare system

Clinical trials ongoing

**Preparing for** clinical trials







hMPV/PIV3



Pediatric RSV/hMPV

In research



More combinations...



### Forward-looking statements

This presentation contains forward-looking statements within the meaning of the Private Securities Litigation Reform Act of 1995, as amended including, but not limited to, statements concerning potential development candidate applications, development candid ate activities, preclinical and clinical studies, regulatory submissions and approvals, risk management and estimates and forward -looking projections with respect to Moderna or its anticipated future performance or events. In some cases, forward -looking statements can be identified by terminology such as "may," "should," "expects," "intends," "plans," "aims," "anticipates," "believes," "estimate es," "protential," "continue," or the negative of these terms or other comparable terminology, although not all forward -looking statements contain these words. The forward-looking statements in this presentation are neither promises nor guarantees, and you should not place undue reliance on these forward-looking statements because they involve known and unknown risks, uncertainties and other factors, many of which are beyond Moderna's control and which could cause actual results to differ materially from those expressed or implied by these forward - looking statements. These risks, uncertainties and other factors include, among others: preclinical and clinical development is lengthy and uncertain, especially for a new category of medicines such as mRNA, and therefore Moderna's preclinical programs or development candidates may be delayed, terminated, or may never advance to or in the clinic; no mRNA drug has been approved in this new p otential category of medicines; and those described in Moderna's most recent Annual Report on Form 10-K filed with the U.S. Securities and Exchange Commission (SEC) and in subsequent filings any forward-looking statements in this presentation in the event of new information, future developments or otherwise. These forward-looking statements are based on Moderna's current expectations and speak only as of the date hereof.

