

Fourth Quarter and Full-Year 2023 Financial Results and Business Update

April 24, 2024

CureVac Speakers









Head of Intellectual Property









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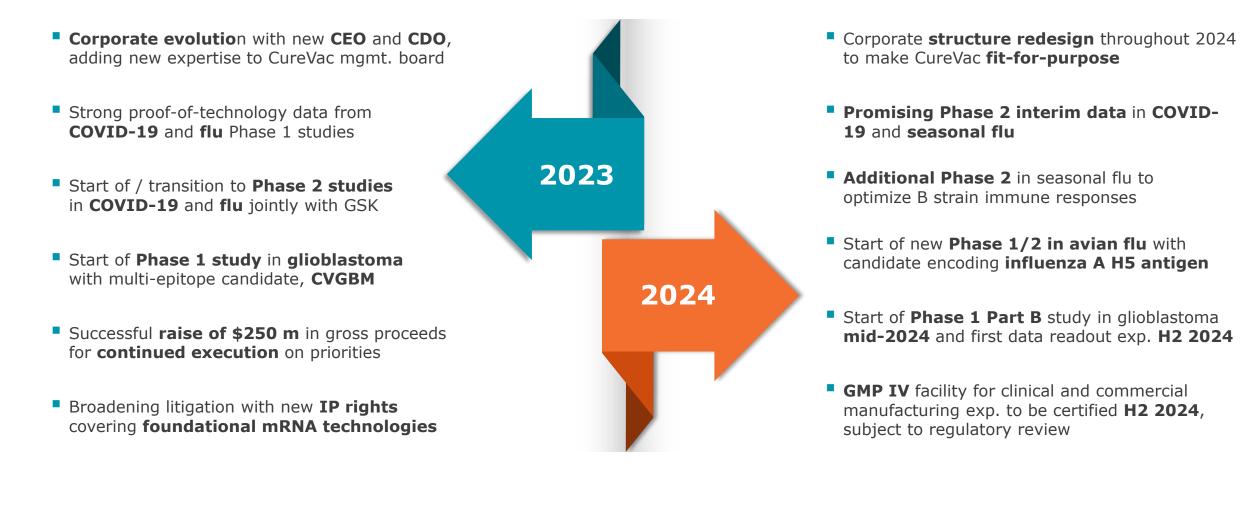
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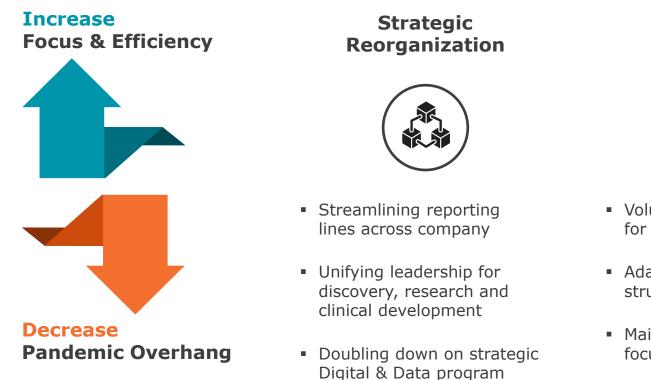


Corporate Development	Oncology Partnerships	Prophylactic Vaccines		
 Organizational redesign initiated with voluntary leaver program for 150 positions Together with GSK, wind down of 	 Strategic collaboration with world- leading cancer center MD Anderson for development of novel cancer vaccines 	 Promising interim Phase 2 data from seasonal flu and COVID-19 programs jointly developed with GSK Start of combined Phase 1/2 study in avian flu within broad infectious disease collaboration with GSK 		
Pandemic Preparedness Agreement after consultation with German government	Oncology Pipeline			
Thaminda Ramanayake to strengthen CureVac Mgmt. Team as Chief Business Officer	Phase 1 Part A dose-escalation in patients with resected glioblastoma fully recruited			
Finance	 Successful safety data review by DSMB confirmed no dose-limiting toxicities across tested doses 			
€402.5 million ✓ Cash position as of December 31, 2023 providing runway into Q4 2025	 Recommendation for 100 µg as preferred dose in subsequent Phase 1 dose- confirmation Part B 	The RNA Printer [®] achieved next regulatory milestone with Framework License to support oncology strategy		









Targeted Rightsizing



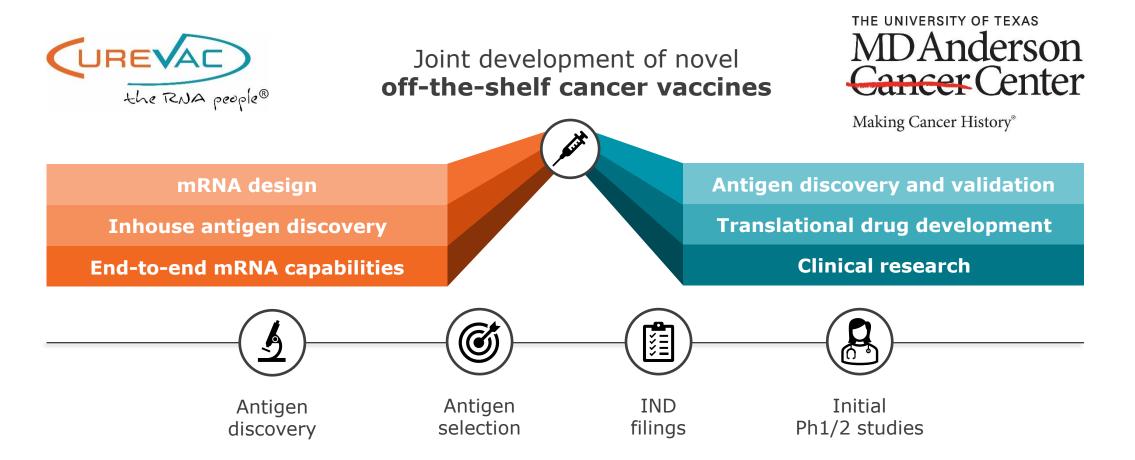
- Voluntary leaver program for 150 positions initiated
- Adapting 'pandemic' structures across company
- Maintaining strong focus on R&D activities

Financial Discipline



- Decreasing OPEX across departments
- Financial discipline and lean budgets
- Extending cash runway into Q4 2025





Harnessing combined expertise in the development of novel off-the-shelf cancer vaccines





Thaminda Ramanayake

Joining CureVac from his position as CBO at Affini-T Therapeutics

- Extensive experience from established biopharma companies such as Sanofi, BioMarin Pharmaceuticals and Amgen
- Strong track record of successful clinical collaborations, M&A, asset in-licensing and strategic financing initiatives across multiple therapeutic areas
- Strong technical experience advancing corporate strategy, business development and strengthening strategic partnerships and alliances
- Strong science background with master's degree in immunology and a bachelor's degree in cellular, molecular and systems biology
- Vast network of Pharma and biotech organizations and community with a focus on the U.S and Europe



			Collaborator	Preclinical development	Phase 1 development	Phase 2 development	Phase 3 developmen
	COVID-19	CV0601 / CV0701	GSK				
Prophylactic	Seasonal influenza	Multivalent candidate	GSK				
Vaccines	H5-pandemic influenza	Monovalent candidate	GSK	NE	W		
	Infectious diseases	3 undisclosed targets	GSK				
	Resected glioblastoma	CVGBM					
Oncology Und indi Und	Undisclosed indications	Off-the-shelf cancer vaccines					
	Undisclosed indications	Off-the-shelf cancer vaccines	MD Anderson	NEW			
	Undisclosed indications	Personalized cancer vaccines					
	Gene editing	Cas9 enzyme					
Molecular Therapies	Ocular diseases	Undisclosed target					
	Therapeutic antibodies	Undisclosed target	Genmab				



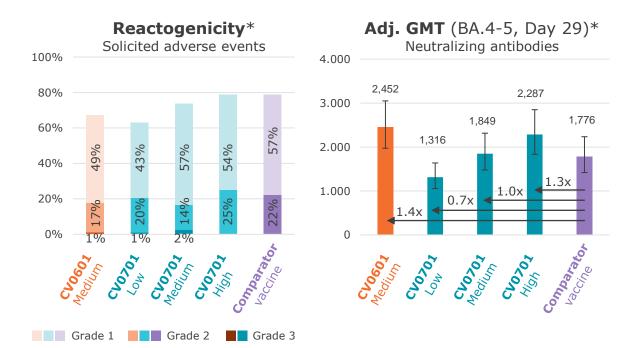
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Seasonal Flu Phase 2 Part

- Candidate from **Phase 1 part** of Phase 1/2 study
- Candidate encodes WHO-recommended strains
- Licensed age-appropriate comparator vaccines

COVID-19 Phase 2

- CV0601, monovalent candidate, encoding BA.4-5
- CV0701, bivalent candidate, encoding BA.4-5 and WT
- Licensed bivalent mRNA comparator vaccine



Influenza A strains

 GMTs consistently numerically exceeded GMTs of comparator vaccines across all doses and age groups

Influenza B strains

- GMTs lower than GMTs of comparator vaccines
- Optimizations to improve immune responses to be tested in additional Phase 2 study





Scalable solutions for indications with high prevalence of **shared antigens**

- Phase 1 study in glioblastoma testing shared antigens with known immunogenicity
- Potentially immunogenic shared antigens identified in collaboration with myNEO
- New Candidate selected for clinical development in 2025 based on proprietary shared antigens
- Collaboration with MD Anderson in solid and hematological cancers

Personalized Cancer Vaccines

Individualized vaccines for broad range of solid cancers

- Further evolution of technology from Frame Cancer Therapeutics acquisition
- Frame technology further directed to specific identification of novel classes of personalized antigens
- Framework License for The RNA Printer[®], critically enabling fast and flexible access to personalized cancer vaccines candidates

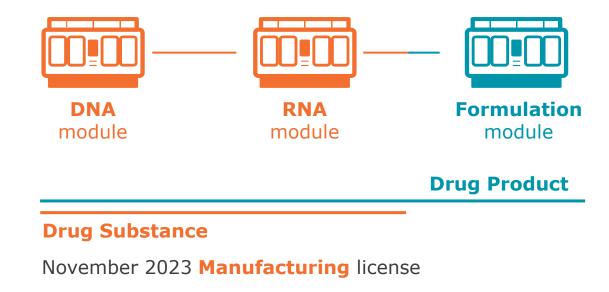


Phase 1 Clinical Study in Patients with Resected Glioblastoma



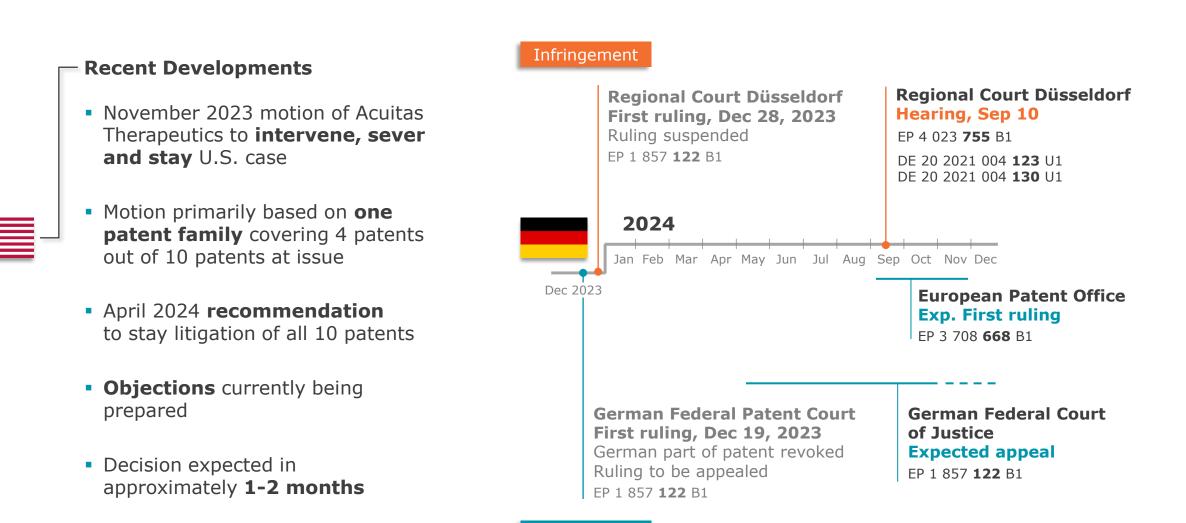




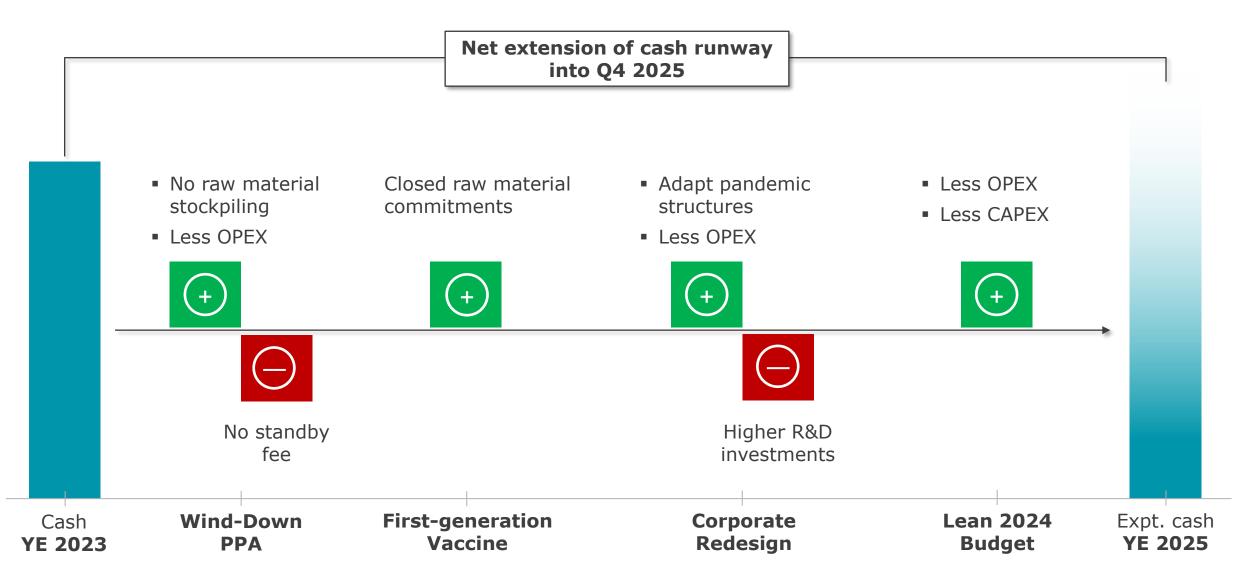


December 2023 Framework license











	December 31, 2023	December 31, 2022
(in € millions)		
Cash and Cash Equivalents	402.5	495.8

Three months ended December 31,

Twelve month ended December 31,

(in € millions)	2023	2022	2023	2022
Revenue	22.6	11.7	53.8	67.4
Cost of Sales, Operating Expenses & Other Operating Income	-110.6	-133.2	-328.0	-316.9
Operating Result	-88.0	-121.5	-274.2	-249.5
Financial Result	1.5	-7.2	14.2	0.3
Pre-Tax Loss	-86.5	-128.7	-260.0	-249.2





Comprehensive **organizational redesign** initiated to adapt residual pandemic infrastructure and improve **operational efficiency** and **agility**



Strong cash position of €402.5 million at year-end 2023, supported in 2024 by stronger financial discipline, **extending cash runway** into the **fourth quarter of 2025**



Significant progress in infectious diseases marked by **new avian flu study** and **promising Phase 2 data** from COVID-19 and seasonal flu programs



Strong advancements of oncology growth driver with **MD Anderson collaboration** and progress **of Phase 1 glioblastoma** study



Thank you for your attention

CureVac www.curevac.com