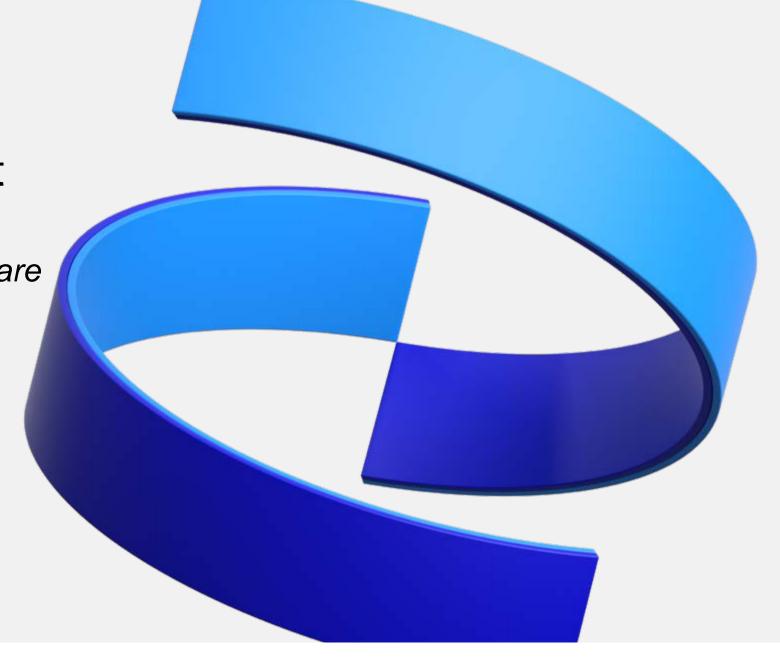
Advancing Our mRNA Strategic Development

40th Annual JP Morgan HealthCare Conference

Albert Bourla, DVM, PhD CEO & Chairman January 10, 2022





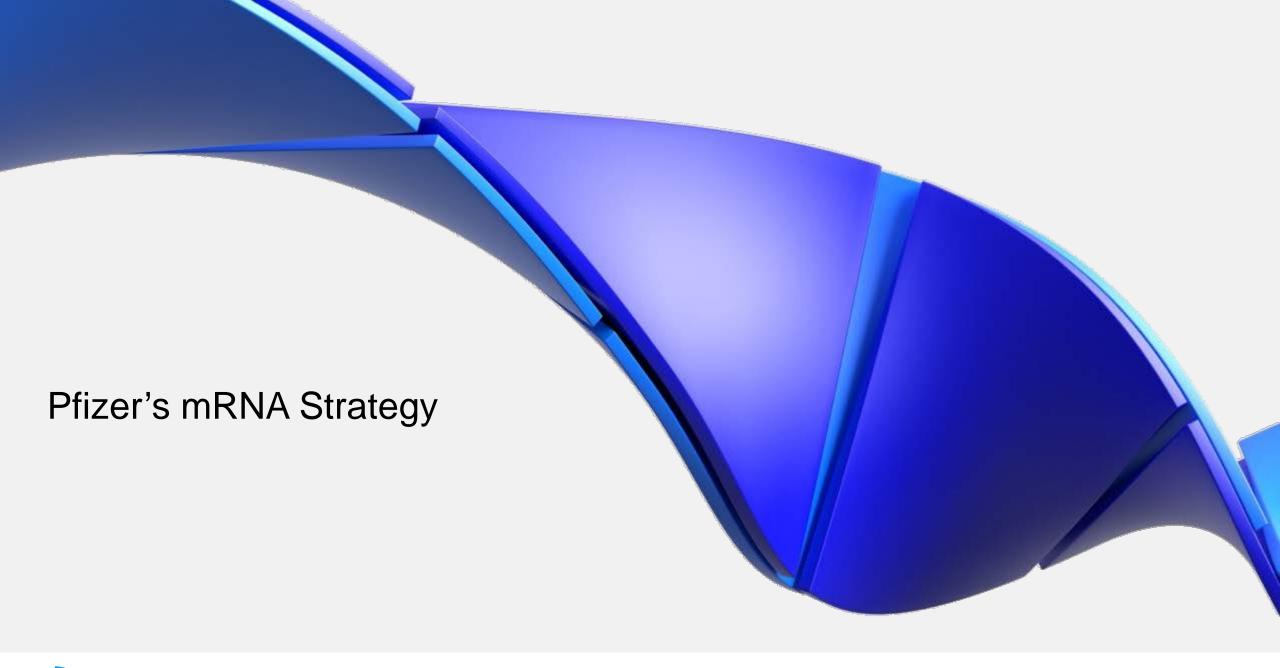
Forward-Looking Statements and Other Notices

This presentation includes forward-looking statements about, among other topics, Pfizer's mRNA strategy and pipeline portfolio; Pfizer's collaboration agreement with BioNTech to co-develop a shingles mRNA vaccine; Pfizer's collaboration with Beam Therapeutics for base editing for certain rare diseases; Pfizer's option agreement with Acuitas to non-exclusively license LNP technology; Pfizer's research collaboration with Codex DNA for its synthetic DNA technology; our efforts to respond to COVID-19, including Comirnaty; our anticipated future operating and financial performance, business plans and prospects; our ability to successfully capitalize on growth opportunities and prospects; and other statements about our business, operations and financial results that are each subject to substantial risks and uncertainties that could cause actual results to differ materially from those expressed or implied by such statements. Among other things, statements regarding revenue and earnings per share growth; the development or commercial potential of our product pipeline, in-line products, product candidates and additional indications, including expected clinical trial protocols, the timing of the initiation and progress of clinical trials and data read-outs from trials; the timing for the submission of applications for and receipt of regulatory approvals; expected breakthrough, best or first-in-class or blockbuster status of our medicines or vaccines; and the anticipated benefits of our lightspeed approach to research and development of certain areas of our product pipeline, are forward looking and are estimates that are subject to change and clinical trial and regulatory success. These statements are subject to risks, uncertainties and other factors that may cause actual results to differ materially from past results, future plans and projected future results.

Additional information regarding these and other factors can be found in Pfizer's Annual Report on Form 10-K for the fiscal year ended December 31, 2020 and in our subsequent reports on Form 10-Q, including in the sections thereof captioned "Risk Factors" and "Forward-Looking Information and Factors That May Affect Future Results", as well as in our subsequent reports on Form 8-K, all of which are filed with the U.S. Securities and Exchange Commission and available at www.sec.gov and <a href="https://www.se

Potential risks and uncertainties also include the impact of COVID-19 on our sales and operations, including impacts on employees, manufacturing, supply chain, research and development, and clinical trials. The forward-looking statements in this presentation speak only as of the original date of the presentation and we undertake no obligation to update or revise any of these statements. This presentation is intended for the investor community only; it is not intended to promote the products referenced herein or otherwise influence healthcare prescribing decisions. All trademarks in this presentation are the property of their respective owners.





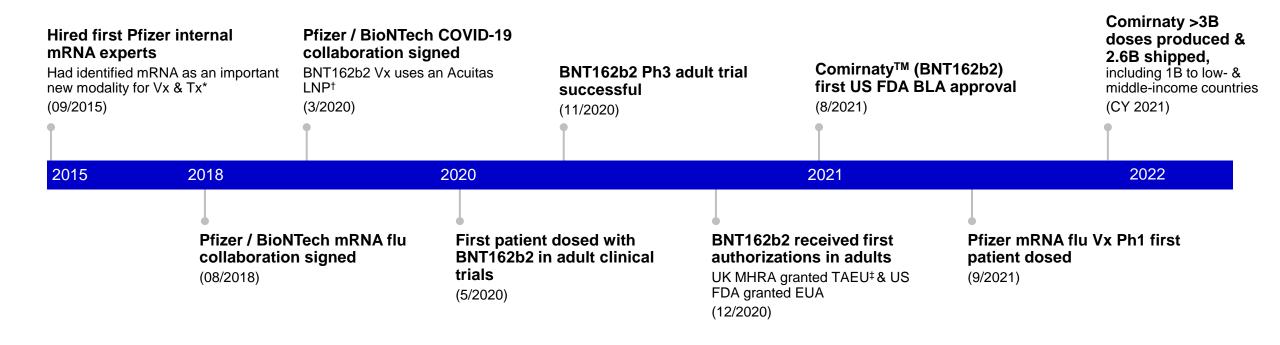


mRNA is expected to be one of our sustainable business drivers going forward

- Adds another potential treatment modality to our armamentarium for vaccines and therapeutics, across our therapeutic areas
- Our Lightspeed methodology & internal cultural changes resulted in the acceleration of the platform's development in recent years
- Looking forward, we aim to continue to harness the power of mRNA to address the greatest patient challenges and contribute substantially to Pfizer's growth targets in the years ahead



mRNA - Where we've come from

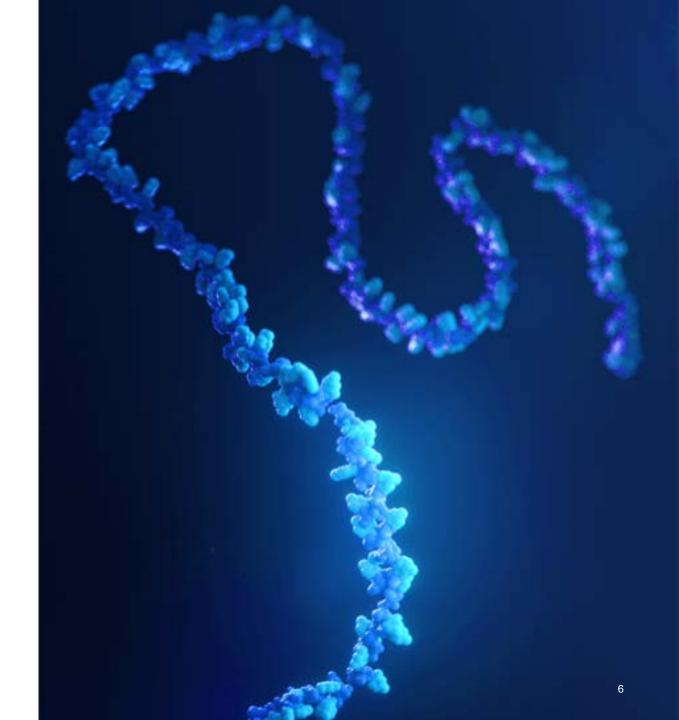


"Vx = Vaccine, Tx = Treatment, †LNP= cationic Lipid NanoParticle, †TAEU = UK Temporary Authorization for Emergency Use. EUA = Emergency Use Authorization, BLA = Biologics License Application, CY = Calendar Year.



Our vision of mRNA leadership

Harness the power of mRNA to address the greatest unmet needs for patients with breakthrough medicines that contribute substantially to Pfizer's growth targets by 2025 and beyond





Pfizer's mRNA Strategy Roadmap

Harness the power of mRNA to address the greatest unmet needs for patients with breakthrough medicines

Invest to Strengthen Core Franchise

Grow Prophylactic Vaccines*

Pursue Additional TAs with Strongest Benefit/Risk

Explore opportunities in larger indications





Flu and Shingles**



Rare Disease

Base editing in partnership with Beam Therapeutics



(e.g., immuno-tolerance)





Oncology

Internal effort leveraging cancer vaccine experience



Internal Medicine

(e.g., next-gen base editing)

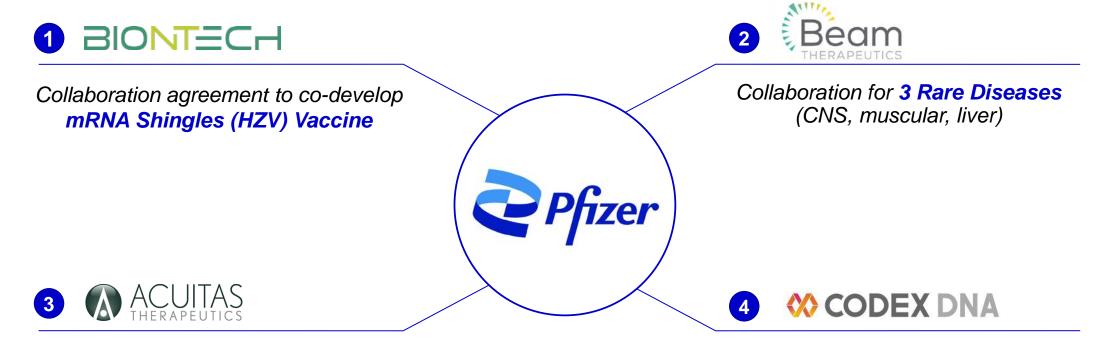
TA = Therapeutic Area; I&I = Inflammation and Immunology



^{*}Programs are currently investigational

^{**}Programs in collaboration with BioNTech

To advance our mRNA strategy, we are pursuing four agreements aligned with our strategic priorities



Collaboration & Option to non-exclusively license **LNP technology** for up to 10 targets

Research collaboration / license for synthetic DNA technology

CNS = central nervous system; HZV = Herpes Zoster Virus (aka Varicella Zoster Virus or VZV) which causes Chicken Pox as a primary infection and later (usually many years later) can cause Shingles after dormancy at the nerve endings; LNP = cationic Lipid NanoParticles



New collaborations established to advance mRNA programs in vaccines and rare diseases

Focus Area





Partnership



Agreement to co-develop mRNA Shingles (VZV) Vaccine



Collaboration for 3 Rare Diseases (CNS, muscular, liver)

- Strategic Benefits
- Candidates to use Pfizer's HZV antigen sequences & certain of BNT's mRNA technology
- Potential for well tolerated, highly efficacious vaccine that could be manufactured at scale
- Third vaccines collaboration between the companies since 2018

- Jointly advance Rare Disease programs using Beam's base editing technology to perform targeted editing of individual nucleotides
- Base Editing does not involve double strand breaks and is potentially more accurate and safer than CRISPR-Cas9* approaches



Formulation and manufacturing partnerships to accelerate efforts in core disease areas

Focus Area



Formulation

Access improved nanoparticle structure to optimize stability, targeting & efficacy



Manufacturing

Produce high-quality medicines at scale with efficient production processes

Partnership



Collaboration & Option to nonexclusively license LNP technology for up to 10 targets



Research collaboration / license for synthetic DNA technology

Strategic Benefits

- Proven LNP* technology used in the Comirnaty COVID-19 Vaccine could be applied to wider range of programs
- Shown to efficiently deliver mRNA to target **cells** while minimizing side effects

- Codex DNA's synthetic DNA technology allows enzymatic synthesis of DNA
- Moving from biological to synthetic DNA assembly may allow significant time savings at front-end of the mRNA production process



Pfizer is uniquely positioned to be a leader in mRNA



Deep understanding of disease within our five core therapeutic areas

- Vaccine expertise & infrastructure built through development of Prevnar 13 and 20, **RSV*** and Comirnaty
- Depth in gene therapy & rare disease with three AAV**-based gene therapies in Phase 3

Demonstrated ability to progress & manufacture promising therapies at "lightspeed"

Operational Excellence

- Industry-leading manufacturing expertise with >3B doses of Comirnaty manufactured in 2021
- Implemented lightspeed cultural changes to accelerate promising new therapies





Our approach to collaborating with the ecosystem to co-create innovation



Pfizer Ventures

Monitors early-stage companies for investment opportunities aligned with our mRNA strategy



RNA Incubator

Establishes academic collaborations and initiates exploratory projects through Pfizer's Center for Therapeutic Innovation



Business Development

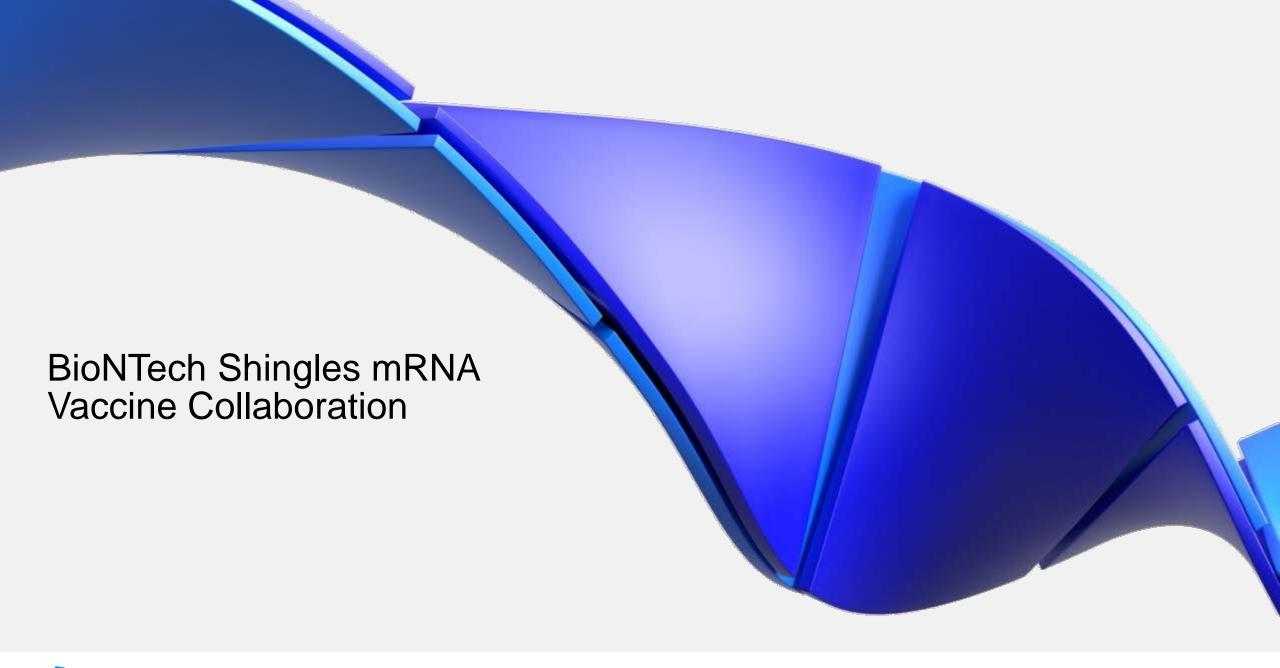
Dedicated team tasked with identifying M&A and partnership opportunities with mature companies that fit our enterprise-wide priorities



mRNA Centers of Excellence

Internal COE comprising research nodes focused on our mRNA priorities (Vaccines, Rare Disease, Oncology, Medicinal Sciences)







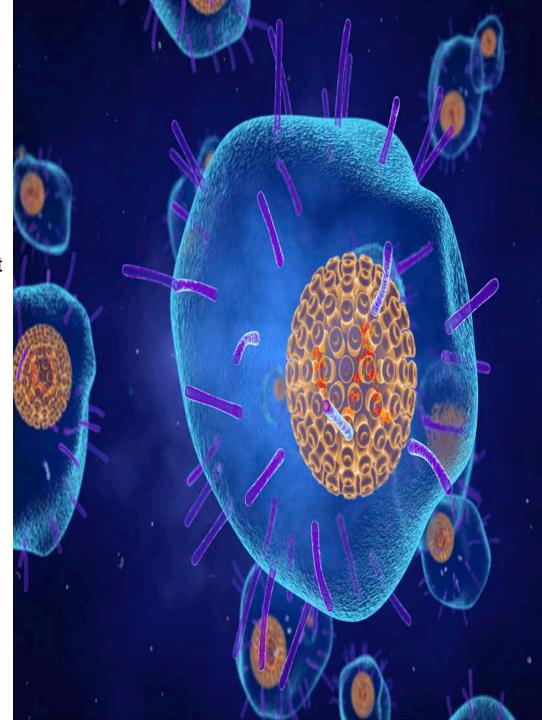
What is Shingles / Herpes Zoster?

Painful disease with potentially severe & long-lasting effects in older & immunocompromised adults

- Reactivation of virus that causes Chicken Pox which has been dormant in nerve endings since childhood infection
- 1 in 3 US adults will experience Shingles at least once in their lifetime with most attacks lasting 3-5 weeks
 - Most cases occur in adults >60 and those who are immunocompromised
- Presents usually as unilateral rash; post herpetic neuralgia (PHN) is a major complication defined as pain persisting to 90 days or more
- PHN is the third most common cause of chronic neuropathic pain in the US with an estimated 500K yearly cases
- ACIP* recommends Shingles vaccination for adults >50

Date Source: Shingles, National Institute on Aging, accessed at https://www.nia.nih.gov/health/shingles Illustration source: Getty Images *ACIP = US CDC's Advisory Committee on Immunization Practices





Opportunity for an Improved Shingles Vaccine Using mRNA Technology

Builds upon prior Pfizer-BioNTech collaboration for COVID-19

Potential mRNA VZV* Vaccine Could Allow: Maintaining High Efficacy + Reduced Side Effects + Robust, Scalable Manufacturing

- Leverages Pfizer's depth in Vaccines (\$6.6B in 2020 revenues,+2% op)
- Uses same expertise & infrastructure as Comirnaty with opportunity for:
 - High Efficacy (≈Current Vaccine)
 - Potential for reduced number of doses (1 vs. 2)
 - Reduced Vaccine Side Effects
 - Currently licensed adjuvanted recombinant vaccine has high efficacy but also rates of side effects that we think can be improved
 - Robust, Scalable Manufacturing
 - Produced >3B doses of Comirnaty in CY[†] 2021 and have production capacity of 4B doses in CY[†] 2022



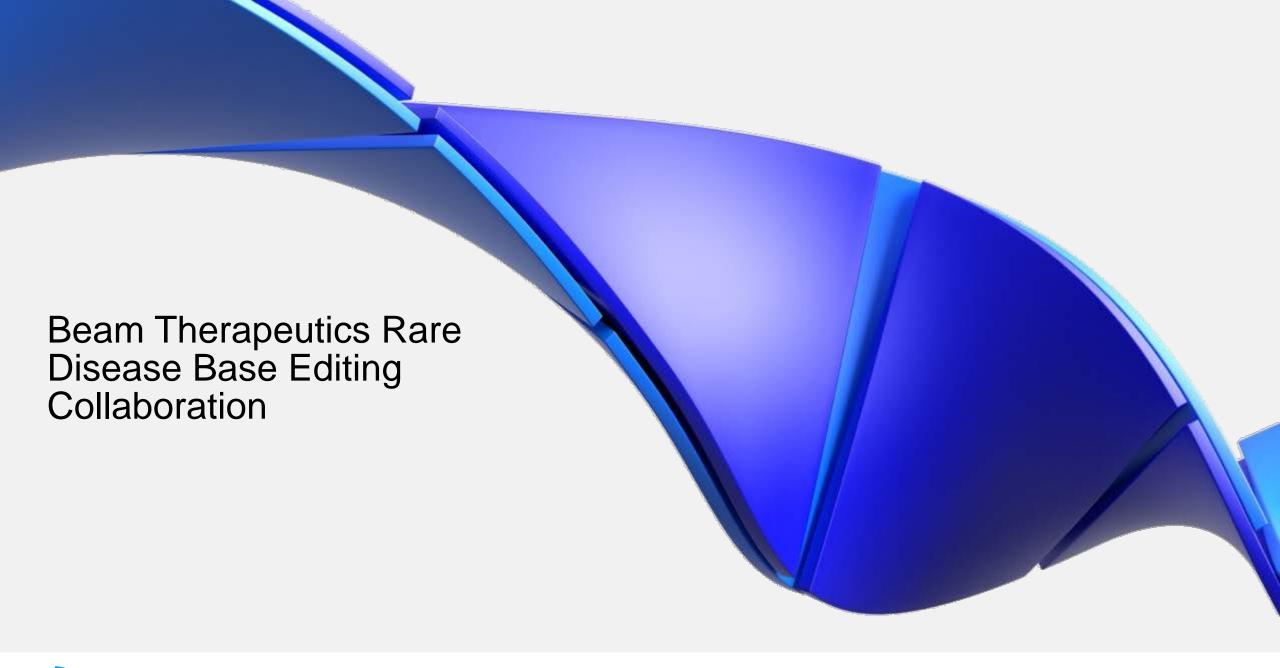
Currently Licensed Adjuvanted Recombinant Vaccine

Currently Licensed Vaccine – Effective but we believe tolerability & capacity-constrained supply can be improved

- Vaccine is highly effective (up to >90% in preventing attacks*) according to data from their label, but
- Large portions of prevalent and incident populations of all adults >50 years and immunocompromised adults >18 years remain unvaccinated
 - The populations total nearly 400M in major markets[†], with only a small portion having been vaccinated to date
- Most patients experience side effects for 2-3 days post-vaccination (2 doses)
- Multi-step reconstitution process with adjuvant

*Source: Shingrix US PI, accessed at https://gskpro.com/content/dam/global/hcpportal/en_US/Prescribing_Information/Shingrix/pdf/SHINGRIX.PDF †Pfizer calculation. Major Markets include US, EU, UK, CAN, AUST, NZ, JPN, SK. ‡Gastro-intestinal includes nausea / vomiting / diarrhea / pain.





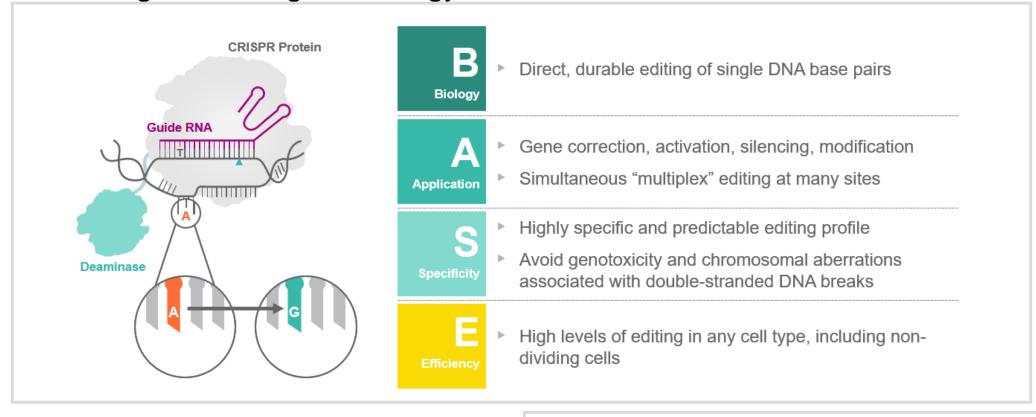


Beam's Base Editing Platform for Rare Diseases Advances Pfizer's Path to Leadership in mRNA

- Leverages Pfizer's depth in gene therapy & rare diseases
 - Pfizer has 3 AAV-based* gene therapies for Rare Diseases in Phase 3 trials and 12 in pre-clinical development with 1-2 INDs expected annually
 - Pfizer Rare Disease Therapeutic Area has \$2.9B 2020 revenues (+29% op) with marketed product strength in hematology, endocrinology, and cardiology
- Demonstrates Pfizer's commitment to mRNA / LNP Technology Across Therapeutic Areas
 - Delivery of mRNA encoding base editing machinery by LNP⁺ is the method of choice for gene editing
 - mRNA delivery facilitates transient expression of editing activity
 - Ability to redose to achieve desired effect level if necessary
 - Progress made in targeting LNP+ to tissues beyond the liver
- Newer generations of gene editing may be more suitable for in vivo therapeutic development
 - CRISPR-Cas9[‡] editing now clinically validated for effecting gene knockdowns
 - Beam's Base Editing does not use double strand breaks and non-homologous end-joining, and may be more accurate and efficient
 - Leverages a non-viral delivery platform



Base editing is a highly differentiated, potentially best-in-class gene editing technology



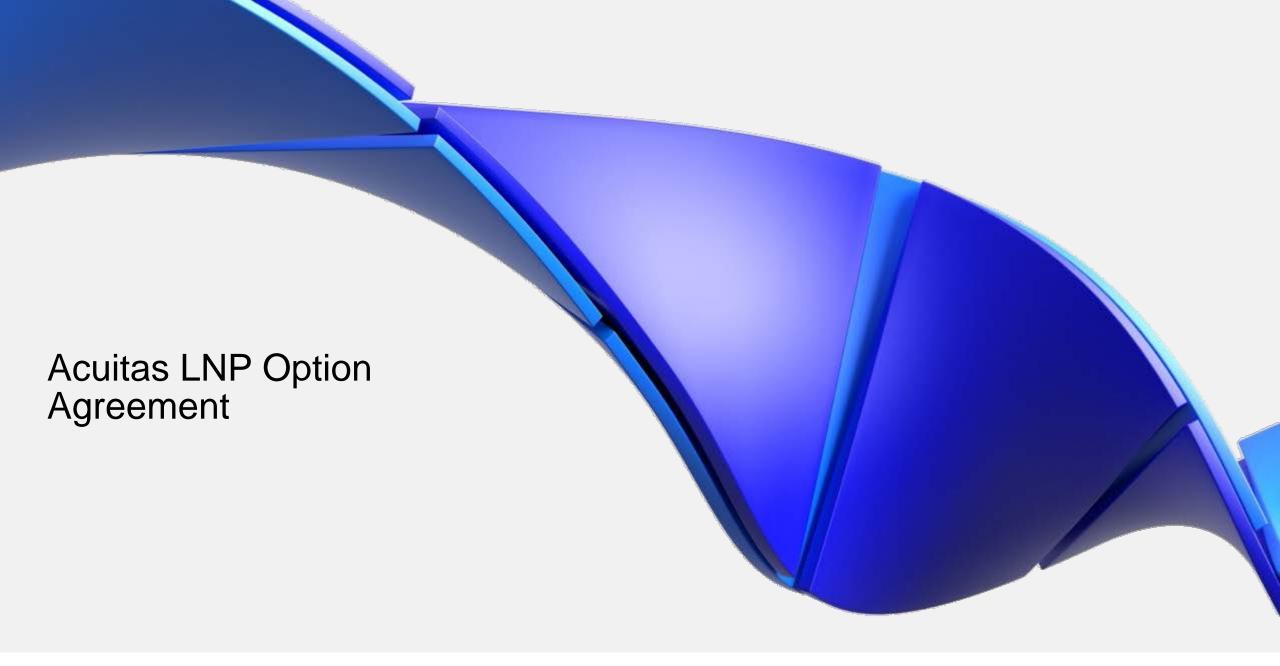
ABE = Adenine Base Editor, DSPC = DiStearoylPhosphatidylCholine, PEG= PolyEthylene Glycol



Therapeutic consists of BE mRNA, guide RNA and LNP

1. mRNA for ABE
2. guide RNA 1
3. Amino lipid
4. Cholesterol
5. DSPC
6. PEG-lipid

Lipid components
60 nm





Acuitas LNP Collaboration

Acuitas Collaboration Terms

- Acuitas is one of the global leaders in development of LNPs
- Allows Pfizer to explore LNPs for a variety of vaccines or therapeutics and exercise option for up to 10 different Vx and Tx
- Acuitas' proven LNP technology is used in the Comirnaty COVID-19 Vx, and shown to efficiently deliver mRNA to target cells while minimizing side effects

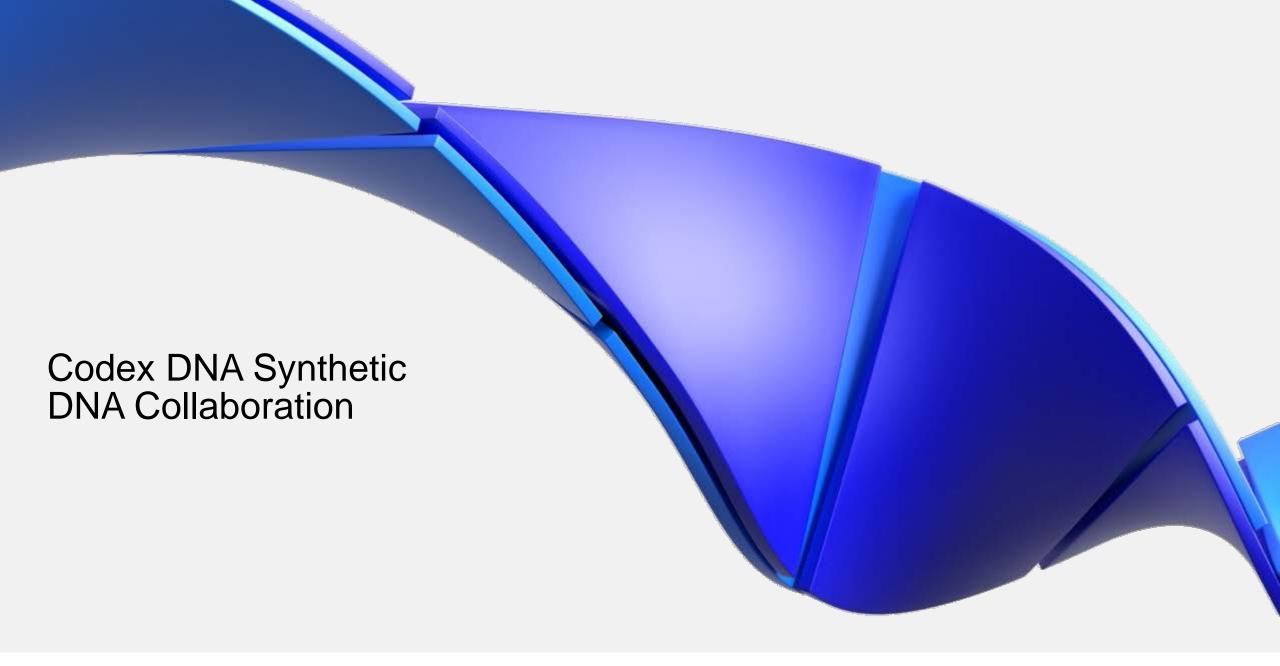
Why are LNPs important for mRNA delivery?

- Engineered LNPs allow efficient intracellular delivery of mRNA vaccines/medicines to the cytoplasm
 - "Naked" mRNA can be significantly degraded or attacked by the immune system before reaching desired cells
 - mRNA has a hard time passing through the cell wall unaided
 - Each specific LNP's properties help determine tissue tropism, reactogenicity, and immunogenicity



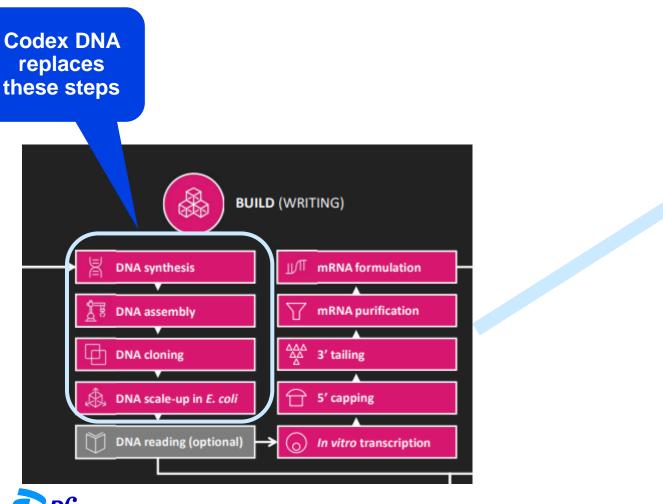


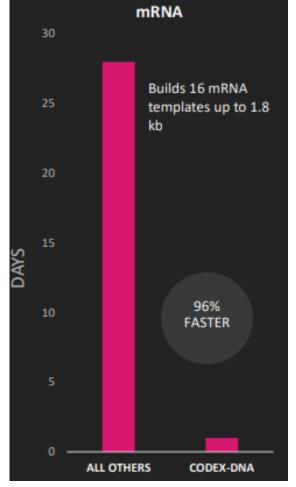






Codex DNA's technology may allow significant time savings at front-end of the mRNA production process by moving from biologic to synthetic DNA assembly, potentially reducing time from several weeks to days





mRNA Collaborations - Key Takeaways

New BioNTech, Beam, Acuitas and Codex DNA collaborations allow Pfizer to continue to advance its mRNA strategy

- These strategic collaborations provide many of the key enabling technologies needed to deepen and expand Pfizer's mRNA development
 - Beam expands the use of mRNA into Rare Disease with the opportunity for precise gene editing
 - BioNTech shingles collaboration adds a potential third mRNA viral vaccine candidate
 - Acuitas LNP collaboration / option / license provides potential LNPs for additional future vaccine & therapeutics candidates
- Platform Capabilities:
 - Codex DNA's synthetic DNA assembly could allow for reduced production cycle times.

Bottom Line: These collaborations provide many of the key enabling technologies needed to deepen and expand Pfizer's mRNA strategic development



