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This presentation contains "forward-looking statements" of Intellia Therapeutics, Inc. ("Intellia", "we" or "our") within the meaning of the Private Securities Litigation Reform Act of 1995. These forward-looking statements include, but are not limited to, express or implied statements regarding Intellia's beliefs and expectations regarding: its strategy, business plans and focus; its ability to quickly and efficiently realize the scope and potential of its genome-editing technology; its ability to maintain, expand and maximize its intellectual property portfolio and pipeline as well as accelerate clinical validation for its platform; the therapeutic value and development potential of CRISPR/Cas-9 gene editing technologies and therapies; its ability to combine its allogeneic T cell platform and CRISPR-based cell engineering with GEMoaB's universal CAR-T platform; the expected strategic benefits and value of the formation of the new company, resulting collaborations, and use of Cellex's cell therapy manufacturing sites; its expectations on the new company's ability to develop CAR-T Cell therapies for immunoncology and autoimmune diseases and the timing and potential success of such therapies; its ability to optimize the impact of its collaborations on its development programs, including but not limited to its collaboration with BXLS, Cellex, or the new company's heability of BXLS, Cellex, Intellia, and the new company to obtain all regulatory approvals necessary to close the transaction; its and the new company's ability to enter into a license and collaboration agreement and co-development and co-promotion agreement; Intellia's ability to submit an IND or equivalent regulatory filing for NTLA-5001 in mid-2021; and Intellia's ability to nominate its first allogeneic development candidate by the 1H 2022.

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Any forward-looking statements in this presentation are based on management's current expectations and beliefs of future events, and are subject to a number of risks and uncertainties that could cause actual results to differ materially and adversely from those set forth in or implied by such forward-looking statements. These risks and uncertainties include, but are not limited to risks related to Intellia's ability to protect and maintain its intellectual property position; risks related to Intellia's relationship with third parties, including its licensors and licensees; risks related to the ability of Intellia's licensors to protect and maintain their intellectual property position; uncertainties related to the authorization, initiation and conduct of studies and other development requirements for the new company's product candidates; the risk that any one or more of the new company's product candidates will not be successfully developed and commercialized; the risk that the results of preclinical studies or clinical studies will not be predictive of future results in connection with future studies; and the risk that Intellia's collaborations with BXLS, Cellex, or the new company or its other collaborations will not continue or will not be successful. These and other risks and uncertainties are described in greater detail in the section entitled "Risk Factors" in Intellia's most recent annual report on Form 10-K and quarterly report on Form 10-Q filed with the SEC, as well as discussions of potential risks, uncertainties, and other important factors in Intellia's other filings with the Securities and Exchange Commission. Any forward-looking statements contained in this press release represent Intellia's views only as of the date hereof and should not be relied upon as representing its views as of any subsequent date. Intellia explicitly disclaims any obligation to update any forward-looking statements, except as required by law.



Intellia, Cellex and Blackstone Launch New Company to Develop Allogeneic Universal CAR-T Cell Therapies, With \$250M Committed Funding

Concurrent Cellex deal enables expansion and acceleration of Intellia's ex vivo pipeline with expanded manufacturing capabilities

Intellia's Key Goals

Realize significant value by combining CRISPR/Cas9 with Universal CAR-T (UniCAR, RevCAR) technologies

Strengthen Intellia's investment in core areas of strategic focus, including wholly owned allogeneic programs



New Company Combines Advanced CRISPR Engineering From Intellia With a Universal CAR-T Cell Construct from GEMoaB



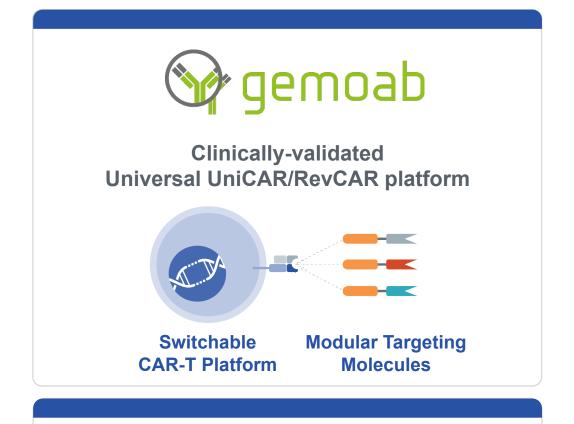
Leading genome editing capabilities with differentiated allogeneic platform



CRISPR/Cas9 Platform



Allogeneic Cell Engineering





Parent company of GEMoaB

Established CMO GMP Cell Therapy Manufacturing



Intellia Receives Significant Value From the New Venture with Blackstone and Cellex, as well as Concurrent Agreement with Cellex

- Rights to 2 Co/Co options in U.S. and key European countries on allogeneic universal CAR-T products
 - Intellia leads U.S. commercialization
- Additional validation of Intellia's proprietary allogeneic platform
- Hold substantial equity stake in new company
- Access to Cellex cell therapy manufacturing site and allogeneic cell donations via a preferred relationship
 - Supports Intellia's wholly owned ex vivo pipeline
 - Expanded capacity to handle additional pipeline growth



Intellia's Proprietary Engineering Platform Provides Flexibility to Expand into Other Cell Therapies

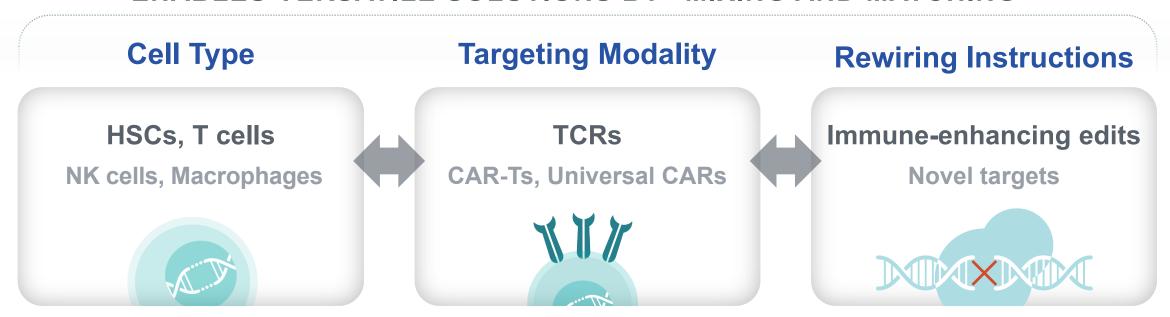
CELL ENGINEERING PLATFORM

Highly efficient sequential editing

Optimal cell performance

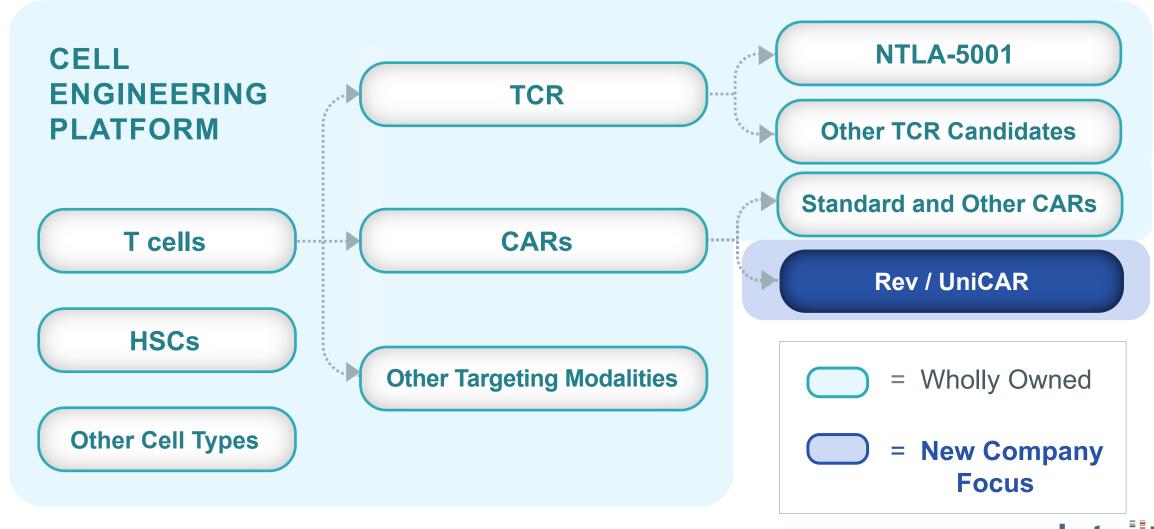
Scalable manufacturing process

ENABLES VERSATILE SOLUTIONS BY "MIXING AND MATCHING"





Intellia Retains Rights Across a Diverse Ex Vivo Landscape



Intellia Gains Access to Important GMP Manufacturing Services from Cellex Under a Preferred Customer Agreement





World-leading manufacturing...

- Manufacturing of intermediates and final products for CAR-T cell products
- Fully automated, audited and inspected GMP facility including Quality Management, Control and Systems
- Experienced and large collection centers for allogeneic cell donations based in Cologne and Dresden
- Worldwide distribution of cell products including Same-Day Delivery for US and Europe

...to support and accelerate key programs.



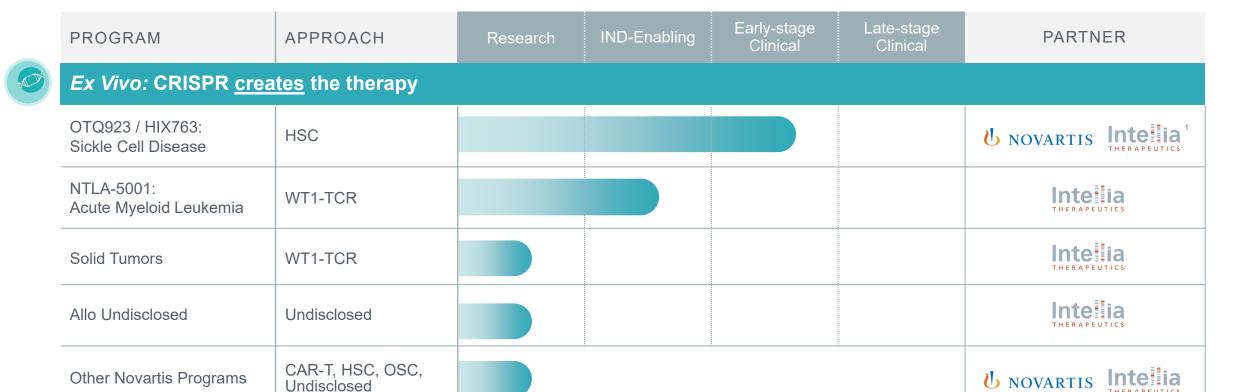
Preferred relationship and dedicated capacity to support Intellia's wholly owned engineered cell therapy programs



High quality and diverse **donor cells** to support allogeneic platform



Expanding and Accelerating Intellia's Ex Vivo Pipeline



- Expect to submit an IND or equivalent regulatory filing for NTLA-5001 in mid-2021
- Plan to nominate first allogeneic development candidate by the 1H 2022



Q&A

Intellia Corporate Update Call

