



## **Regeneron and Colorado Center for Personalized Medicine Announce Major New Human Genetics Research Collaboration**

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***Regeneron Genetics Center will sequence 450,000 informed and consented patients***

***Sequencing data will be paired with de-identified health records to aid in genomic medicine, drug discovery and personalized medicine approaches***

Regeneron Pharmaceuticals, Inc. (NASDAQ: REGN) and the Colorado Center for Personalized Medicine (CCPM) at the University of Colorado Anschutz Medical Campus today announced a large-scale research collaboration designed to advance the field of human genetics and precision medicine through the sharing of 450,000 DNA samples and corresponding health records from de-identified, consented patient participants in the expansive UHealth system. The Regeneron Genetics Center (RGC), a wholly owned subsidiary of Regeneron, has entered into the collaboration with CCPM and will sequence these samples, producing genomic data that can be used to facilitate translational medical research and ultimately enable physicians to make better decisions for their patients.

The CCPM is one of the largest health data warehouses in the United States and a pioneer in the use of a secure cloud platform with more than 8.7 million de-identified patient records. The five-year old research enterprise is also one of the first and largest programs in the country to integrate personalized genomic information with clinical data via a research biobank. CCPM physicians will validate any genetic findings from the RGC data in their CLIA-certified lab, enabling the return of clinically-actionable results to patients.

"This collaboration will take an already notable program at the CCPM and expand the depth and breadth of its capabilities, allowing us to give more back to our patient participants than ever before," said Kathleen Barnes, Ph.D., Professor and Director of CCPM at the University of Colorado Anschutz Medical Campus. "We have made tremendous strides with our work in pharmacogenomics, but having access to such a large genomic dataset that enables the return of clinically actionable results will be transformative. Our collaboration with the RGC will lead to an optimization of patient care, using personalized results to better inform clinical decision making, and potentially leading to new ways of diagnosing, preventing and treating illnesses."

The RGC has built one of the world's largest genetics databases, pairing the sequenced exomes and de-identified electronic health records of more than one million people, through collaborations with more than 80 global healthcare and academic institutions. Building upon Regeneron's strengths in genetics-driven drug discovery, the information secured from this initiative will allow for the elucidation, on a large scale, of genetic factors that cause or influence a range of human diseases.

"We're excited to collaborate with the CCPM and UHealth to further expand the RGC's large-scale genomics initiatives," said Aris Baras, M.D., Senior Vice President at Regeneron and Head of the Regeneron Genetics Center. "In the search for new and improved medicines, as well as the advancement of validated and improved risk scores in medicine, both scale and quality of data matter. This partnership opens up new doors for meaningful discovery, strengthens Regeneron's ability to speed and improve the drug development process, and allows us to work alongside other leaders in the advancement of genomic and precision medicine."

The impressive scope of this effort is also thanks to the wide-ranging footprint of UHealth. Large numbers of the nonprofit health care system's patients have consented to biobank participation, more than half of whom live outside of the metro Denver area and in neighboring states like Nebraska and Wyoming.

"Our patients are already benefitting from the remarkable work of the CCPM which is allowing providers to use genomics to make more accurate diagnoses and precisely tailor treatment to individual patients," said Richard Zane, M.D., UHealth Chief Innovation Officer, who is also the Professor and Chair of emergency medicine at the University of Colorado School of Medicine. "This partnership will help drive the health care discoveries of tomorrow and realize the full potential of precision medicine. We so appreciate our patients who have consented to participate and without whom discovery would not be possible."

### **About The RGC-Colorado Center for Personalized Medicine Research Collaboration**

The RGC and CCPM will create a comprehensive, large-scale genomics database comprising approximately 450,000 fully annotated, whole exome sequences for discovery research and to support patient care at UHealth. The goal of the partnership is to leverage de-identified genetic information from consenting patients with large sets of corresponding medical data that will ultimately build a warehouse of information useful for the discovery of genetic factors that could inform clinical care. For the newly-established CCPM, this enterprise will exponentially expand biobank research capabilities.

### **About the University of Colorado Anschutz Medical Campus**

The University of Colorado Anschutz Medical Campus is a world-class medical destination at the forefront of transformative science, medicine, education, and patient care. The campus encompasses the University of Colorado health professional schools, more than 60 centers and institutes, and two nationally ranked hospitals that treat more than 2 million adult and pediatric patients each year. Innovative, interconnected and highly collaborative, together we deliver life-changing treatments, patient care, professional training, and conduct world-renowned research powered by more than \$550 million in research awards. For more information, visit [www.cuanschutz.edu](http://www.cuanschutz.edu).

### **About Regeneron**

Regeneron (NASDAQ: REGN) is a leading biotechnology company that invents life-transforming medicines for people with serious diseases. Founded

and led for over 30 years by physician-scientists, our unique ability to repeatedly and consistently translate science into medicine has led to seven FDA-approved treatments and numerous product candidates in development, all of which were homegrown in our laboratories. Our medicines and pipeline are designed to help patients with eye diseases, allergic and inflammatory diseases, cancer, cardiovascular and metabolic diseases, pain, infectious diseases and rare diseases.

Regeneron is accelerating and improving the traditional drug development process through our proprietary *VelociSuite*<sup>®</sup> technologies, such as *VelocImmune*<sup>®</sup> which uses unique genetically-humanized mice to produce optimized fully-human antibodies and bispecific antibodies, and through ambitious research initiatives such as the Regeneron Genetics Center, which is conducting one of the largest genetics sequencing efforts in the world.

For additional information about the company, please visit [www.regeneron.com](http://www.regeneron.com) or follow [@Regeneron](https://twitter.com/Regeneron) on Twitter.

#### **Regeneron Forward-Looking Statements and Use of Digital Media**

*This press release includes forward-looking statements that involve risks and uncertainties relating to future events and the future performance of Regeneron Pharmaceuticals, Inc. ("Regeneron" or the "Company"), and actual events or results may differ materially from these forward-looking statements. Words such as "anticipate," "expect," "intend," "plan," "believe," "seek," "estimate," variations of such words, and similar expressions are intended to identify such forward-looking statements, although not all forward-looking statements contain these identifying words. These statements concern, and these risks and uncertainties include, among others, the impact of SARS-CoV-2 (the virus that has caused the COVID-19 pandemic) on Regeneron's business and its employees, collaborators, suppliers, and other third parties on which Regeneron relies, Regeneron's and its collaborators' ability to continue to conduct research and clinical programs, Regeneron's ability to manage its supply chain, net product sales of products marketed by Regeneron and/or its collaborators (collectively, "Regeneron's Products"), and the global economy; the nature, timing, and possible success and therapeutic applications of Regeneron's Products and Regeneron's product candidates and research and clinical programs now underway or planned, as well as the use of human genetics in Regeneron's research programs; the extent to which the results from the research and development programs conducted by Regeneron and/or its collaborators (such as Regeneron's collaboration with the Colorado Center for Personalized Medicine at the University of Colorado Anschutz Medical Campus discussed in this press release) may lead to advancement of product candidates to clinical trials or therapeutic applications; the likelihood, timing, and scope of possible regulatory approval and commercial launch of Regeneron's product candidates and new indications for Regeneron's Products; unforeseen safety issues resulting from the administration of Regeneron's Products and product candidates in patients, including serious complications or side effects in connection with the use of Regeneron's Products and product candidates in clinical trials; determinations by regulatory and administrative governmental authorities which may delay or restrict Regeneron's ability to continue to develop or commercialize Regeneron's Products and product candidates; ongoing regulatory obligations and oversight impacting Regeneron's Products, research and clinical programs, and business, including those relating to patient privacy and the use of genomic data as discussed in this press release; uncertainty of market acceptance and commercial success of Regeneron's Products and product candidates and the impact of studies (whether conducted by Regeneron or others and whether mandated or voluntary) on the commercial success of Regeneron's Products and product candidates; the availability and extent of reimbursement of Regeneron's Products from third-party payers, including private payer healthcare and insurance programs, health maintenance organizations, pharmacy benefit management companies, and government programs such as Medicare and Medicaid; coverage and reimbursement determinations by such payers and new policies and procedures adopted by such payers; competing drugs and product candidates that may be superior to Regeneron's Products and product candidates; the ability of Regeneron to manufacture and manage supply chains for multiple products and product candidates; the ability of Regeneron's collaborators, suppliers, or other third parties (as applicable) to perform manufacturing, filling, finishing, packaging, labeling, distribution, and other steps related to Regeneron's Products and product candidates; unanticipated expenses; the costs of developing, producing, and selling products; the ability of Regeneron to meet any of its financial projections or guidance and changes to the assumptions underlying those projections or guidance; the potential for any license or collaboration agreement, including Regeneron's agreements with Sanofi, Bayer, and Teva Pharmaceutical Industries Ltd. (or their respective affiliated companies, as applicable) to be cancelled or terminated without any further product success; and risks associated with intellectual property of other parties and pending or future litigation relating thereto (including without limitation the patent litigation and other related proceedings relating to Dupixent<sup>®</sup> (dupilumab) and Praluent<sup>®</sup> (alirocumab)), other litigation and other proceedings and government investigations relating to the Company and/or its operations, the ultimate outcome of any such proceedings and investigations, and the impact any of the foregoing may have on Regeneron's business, prospects, operating results, and financial condition. A more complete description of these and other material risks can be found in Regeneron's filings with the U.S. Securities and Exchange Commission, including its Form 10-K for the year ended December 31, 2019 and its Form 10-Q for the quarterly period ended March 31, 2020. Any forward-looking statements are made based on management's current beliefs and judgment, and the reader is cautioned not to rely on any forward-looking statements made by Regeneron. Regeneron does not undertake any obligation to update publicly any forward-looking statement, including without limitation any financial projection or guidance, whether as a result of new information, future events, or otherwise.*

*Regeneron uses its media and investor relations website and social media outlets to publish important information about the Company, including information that may be deemed material to investors. Financial and other information about Regeneron is routinely posted and is accessible on Regeneron's media and investor relations website (<http://newsroom.regeneron.com>) and its Twitter feed (<http://twitter.com/regeneron>).*

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