

**FOOD AND DRUG ADMINISTRATION (FDA)  
Center for Biologics Evaluation and Research (CBER)  
76<sup>th</sup> Meeting of the Cellular, Tissue, and Gene Therapies  
Advisory Committee (CTGTAC)  
October 31, 2023  
DRAFT AGENDA**

The committee will meet in open session to discuss and make recommendations on biologics license application (BLA) 125787 from Vertex Pharmaceuticals, Inc. for exagamglogene autotemcel (exa-cel). The applicant has requested an indication for the treatment of sickle cell disease in patients 12 years and older with recurrent vaso-occlusive crises.

Time EDT	Presentation/Presenter
9:00 a.m.	<p><b><u>Opening Remarks: Call to Order and Welcome (5 Min)</u></b></p> <p>Tabassum (Taby) Ahsan, Ph.D., Acting Chairperson, CTGTAC Vice President, Cell Therapy Operations City of Hope, Duarte, CA</p> <p><b><u>Administrative Announcements, Roll Call, Introduction of Committee, Conflict of Interest Statement (20 Min)</u></b></p> <p>LCDR Cicely Reese, Pharm.D., Designated Federal Officer Division of Scientific Advisors and Consultants (DSAC), OM, CBER, FDA</p>
9:25 a.m.	<p><b><u>FDA Introduction (5 Min)</u></b></p> <p><b>Introductory Remarks</b></p> <ul style="list-style-type: none"> <li>• <b>Nicole Verdun, M.D.</b> Director Office of Therapeutic Products (OTP), CBER, FDA</li> </ul>
9:30 a.m.	<p><b><u>Guest Speaker Presentations (55 Min including Q &amp; A)</u></b></p> <p><b>Genetic Editing</b></p> <ul style="list-style-type: none"> <li>• <b>Fyodor Urnov, Ph.D. (20 Min)</b> Professor, Department of Molecular and Cell Biology University of California, Berkeley Director of Technology and Translation Innovative Genomics Institute Berkeley, California</li> </ul>

**FOOD AND DRUG ADMINISTRATION (FDA)  
Center for Biologics Evaluation and Research (CBER)  
76<sup>th</sup> Meeting of the Cellular, Tissue, and Gene Therapies  
Advisory Committee (CTGTAC)  
October 31, 2023  
DRAFT AGENDA**

	<p><b>Off-Targets of Genetic Editing</b></p> <ul style="list-style-type: none"> <li>• <b>Daniel E. Bauer, Ph.D. (20 Min)</b> Principal Investigator and Staff Physician Dana-Farber/Boston Children’s Cancer and Blood Disorders Center Boston Children’s Hospital Boston, MA</li> </ul> <p><b>Q &amp; A (15 Min)</b></p>
10:25 a.m.	<b>BREAK (10 min.)</b>
10:35 a.m.	<p><b><u>Applicant Presentations</u> (75 Min including Q &amp; A)</b></p> <p><b>Exa-cel for the Treatment of Sickle Cell Disease (SCD) in Patients ≥ 12 Years With Recurrent Vaso-Occlusive Crises (VOCs).</b></p> <p>Introduction                      <b>Stephanie Krogmeier, Ph.D.</b> Vice President, Global Regulatory Affairs Vertex Pharmaceuticals Incorporated</p> <p>Unmet Need                         <b>Alexis Thompson, M.D., M.P.H.</b> Division Chief, Hematology Children’s Hospital of Philadelphia</p> <p>Efficacy                              <b>William Hobbs, M.D., Ph.D.</b> Vice President Clinical Development, Hematology Vertex Pharmaceuticals Incorporated</p> <p>Non-Clinical Safety               <b>David Altshuler, M.D., Ph.D.</b> Executive Vice President and Chief Scientific Officer, Vertex Pharmaceuticals Incorporated</p> <p>Safety                                 <b>Christopher Simard, M.D.</b> Vice President, Global Patient Safety Vertex Pharmaceuticals Incorporated</p>

**FOOD AND DRUG ADMINISTRATION (FDA)**  
**Center for Biologics Evaluation and Research (CBER)**  
**76<sup>th</sup> Meeting of the Cellular, Tissue, and Gene Therapies**  
**Advisory Committee (CTGTAC)**  
**October 31, 2023**  
**DRAFT AGENDA**

	<p>Clinical Perspective      <b>Haydar Frangoul, M.D.</b>  Medical Director, Pediatric Hematology/Oncology  Tristar Centennial Medical Center</p> <p><b>Q &amp; A (15 Min)</b></p>
<b>11:50 a.m.</b>	<b>LUNCH (45 Min)</b>
<b>12:35 p.m.</b>	<b><u>Open Public Hearing (60 Min)</u></b>
<b>1:35 p.m.</b>	<b>BREAK (10 Min)</b>
<b>1:45 p.m.</b>	<p><b><u>FDA Presentations (75 Min including Q &amp; A)</u></b></p> <p><b>BLA 125787 Exagamglogene autotemcel (exa-cel)</b></p> <ul style="list-style-type: none"> <li>• <b>Karl Kasamon, M.D.</b>  Reviewer, Officer of Clinical Evaluation  Division of Hematology, Benign Hematology  Branch, OTP, CBER. FDA</li> <li>• <b>Komudi Singh, Ph.D.</b>  Bioinformatics Reviewer, Office of Cellular  Therapy and Human Tissue  OTP, CBER, FDA</li> </ul> <p><b>Q &amp; A (15 Min)</b></p>
<b>3:00 p.m.</b>	<p><b><u>Committee Discussion (110 min)</u></b></p> <ul style="list-style-type: none"> <li>• Discussion Question</li> </ul>
<b>4:50 p.m.</b>	<p><b>Closing Remarks (10 Min)</b></p> <ul style="list-style-type: none"> <li>• <b>Nicole Verdun</b></li> </ul>
<b>5:00 p.m.</b>	<b>ADJOURNMENT</b>