

**FOOD AND DRUG ADMINISTRATION**  
**Center for Biologics Evaluation and Research (CBER)**  
**67<sup>th</sup> Meeting of the Cellular, Tissue, and Gene Therapies Advisory Committee (CTGTAC)**  
**October 12, 2017**

**FDA White Oak Conference Center, Room 1503**  
**Silver Spring, Maryland**

**DRAFT AGENDA**

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**Topic:** The committees will discuss Voretigene Neparvovec, Spark Therapeutics, BLA 125610, for the treatment of patients with vision loss due to confirmed biallelic RPE65 mutation-associated retinal dystrophy

8:30 a.m.	Welcome and Introduction of Members	<b>Barry Byrne, M.D. Ph.D.</b> Acting Chair, CTGTAC
	Conflicts of Interest Statement	<b>Prabhakara Atreya, Ph.D.</b> Designated Federal Officer, CTGTAC
8:50 a.m.	FDA Introduction	<b>Wilson W. Bryan, M.D.</b> Director, Office of Tissues and Advanced Therapies (OTAT), CBER
	Sponsor Presentation(s)	
9:00 a.m.	Introduction	<b>Kathryn High, M.D.</b> President and Head of Res. & Dev.
	Unmet Need	<b>Mark Pennessi, M.D. Ph.D.</b> Associate Prof. Ophthal. Genetics Oregon Health and Sci. University
	Efficacy	<b>Kathleen Reape, M.D.</b> Head, Clinical Res. & Dev.
	Safety	<b>Deborah Kelley, M.D.</b> Head, Pharmacovigilance
	Clinical Perspective	<b>Albert Maguire, M.D.</b> Clin. Associate, Div. Ped. Ophthal. Children's Hospital of Philadelphia

## FDA Presentation

10:00 a.m. BLA125610 Voretigene Neparvovec  
Spark Therapeutics, Inc.

**Yao-Yao Zhu, M.D., Ph.D.**  
Medical Officer, OTAT/CBER

10:45 a.m. Break

11:00 a.m. Q & A

11:15 a.m. Open Public Hearing

12:15 p.m. Lunch

1:15 p.m. Q&A

1:30 p.m. Committee Discussion

2:45 p.m. Break

3:00 p.m. Committee Discussion

5:00 p.m. Adjourn Meeting