

Coalition Against Major Diseases and FDA 2015 Annual Scientific Workshop

October 15, 2015

FDA White Oak Campus

CO-SPONSORED BY: Critical Path Institute U.S. Food and Drug Administration

Overview and Objectives

The Coalition Against Major Diseases (CAMD) is a public-private-partnership aimed at creating new tools and methods that can be applied to increase the efficiency of the development process of new treatments for Alzheimer's disease (AD) and Parkinson's disease (PD). The annual meeting brings together members from the pharmaceutical industry, academic key opinion leaders, NIA, FDA, EMA and advocacy groups. The objectives of the meeting are: understand accomplishments of CAMD scientific projects, discuss how these tools are currently or will be applied in drug development, obtain commitment for sharing information/data to begin quantifying benefits of these tools, and facilitate robust and open discussion among all parties of drug development in Alzheimer's and Parkinson's diseases. Experts in the fields of Alzheimer's disease and Parkinson's disease, and leaders of the patient stakeholder community will deliver keynote presentations and regulatory science will be prominently featured throughout the meeting.

Click here for the 2015 Annual Meeting Minutes.

Agenda

7:30-8:30 am	Continental Breakfast
8:30-9:00 am	Welcoming Remarks Martha Brumfield, CEO (Critical Path Institute) Janet Woodcock, Director (Center for Drug Evaluation and Research, FDA) Diane Stephenson & Stephen Arneric, Executive Co-Directors (CAMD)
9:00-9:20 am	Keynote Address Manuel Haas (EMA) Coalition Against Major Diseases and FDA 2015 Annual Scientific Workshop

9:20-9:40 am	Regulatory Perspectives
7.40-7.40 am	ShaAvrée Buckman-Garner (FDA)
	FDA Biomarker Learnings and the Future
9:40-9:55 am	BREAK
SESSION I: Exciting Dev Mark Gordon, Chair	elopments in CAMD Working Groups
9:55-10:10 am	Meeting the Needs of the Parkinson's Community Steve Ford (Parkinson's UK)
7.00 10110 um	
10:10-10:25 am	Computational Modeling for AD
10.10-10.25 am	Julie Stone (Merck) & Klaus Romero (C-Path)
	Where has CAMD come and where do we need to go?
	How can we achieve better understanding of disease progression and efficient clinical populations?
10:25-10:40 am	AD Hippocampal Volume Team Derek Hill (Ixico)
	AD CSF Biomarkers Team
10:40-10:55 am	Robert Dean (Lilly)
10:55-11:40 am	Regulatory Panel Discussion: Regulatory Innovation Now and in the Future Moderator: <i>Richard MeibachJim Kaiser (FDA), Eric Bastings (FDA), Keiju Motohast</i> <i>(FDA), Sandra Kweder (FDA), Maria Isaac (EMA), Vikrma Sinha (FDA), Chris Lept</i>
	LUNCH & AWARDS
11:40-12:25 pm	
SESSION II: Strategies for Successful I	Implementation of Biomarkers in Clinical Trials IntegrationLooking to the Future i) Data SharingWhat Can Be Learned from ALS? Melanie Leitner (Biogen)
SESSION II: Strategies for Successful I CAMD Data Sharing and Peter Loupos, Chair (Sanof	IntegrationLooking to the Future i) Data SharingWhat Can Be Learned from ALS?
SESSION II: Strategies for Successful I CAMD Data Sharing and Peter Loupos, Chair (Sanof 12:25-12:40 pm	IntegrationLooking to the Future i) Data SharingWhat Can Be Learned from ALS? Melanie Leitner (Biogen) PPMI Paving the Way for Defining Prodromal PD
SESSION II: Strategies for Successful I CAMD Data Sharing and Peter Loupos, Chair (Sanof 12:25-12:40 pm 12:40-12:55 pm	IntegrationLooking to the Future i) Data SharingWhat Can Be Learned from ALS? Melanie Leitner (Biogen) PPMI Paving the Way for Defining Prodromal PD Ken Marek (MNI) Data sharingSuccess Story from Multiple Sclerosis (MSOAC)
SESSION II: Strategies for Successful I CAMD Data Sharing and Peter Loupos, Chair (Sanof 12:25-12:40 pm 12:40-12:55 pm 12:55-1:10 pm	IntegrationLooking to the Future i) Data SharingWhat Can Be Learned from ALS? Melanie Leitner (Biogen) PPMI Paving the Way for Defining Prodromal PD Ken Marek (MNI) Data sharingSuccess Story from Multiple Sclerosis (MSOAC) Jesse Cedarbaum (Biogen)
SESSION II: Strategies for Successful I CAMD Data Sharing and Peter Loupos, Chair (Sanof 12:25-12:40 pm 12:40-12:55 pm 12:55-1:10 pm 1:10-1:25 pm	IntegrationLooking to the Future Data SharingWhat Can Be Learned from ALS? Melanie Leitner (Biogen) PPMI Paving the Way for Defining Prodromal PD Ken Marek (MNI) Data sharingSuccess Story from Multiple Sclerosis (MSOAC) Jesse Cedarbaum (Biogen) Panel Discussion on Prospective Directions for CAMDFocus on Data
SESSION II: Strategies for Successful I CAMD Data Sharing and Peter Loupos, Chair (Sanof 12:25-12:40 pm 12:40-12:55 pm 12:55-1:10 pm	IntegrationLooking to the Future ii) Data SharingWhat Can Be Learned from ALS? Melanie Leitner (Biogen) PPMI Paving the Way for Defining Prodromal PD Ken Marek (MNI) Data sharingSuccess Story from Multiple Sclerosis (MSOAC) Jesse Cedarbaum (Biogen) Panel Discussion on Prospective Directions for CAMDFocus on Data Melanie Leitner, Jesse Cedarbaum, Ken Marek, Paul Maruff (Cogstate, AIBL)

Integrated Focus Sessions

1:50-2:30 pm	Session III: Modeling How can we achieve better understanding of disease progression and efficient clinical populations? Vikram Sinha (FDA Co-chair) & Klaus Romero (CAMD Co-chair)
2:30-3:10 pm	Session IV: Biomarkers Chris Leptak (FDA Co-chair) & Richard Meibach (CAMD Co-chair)
3:10-3:50 pm	Session V: Digital Biomarker Technologies Medical Device Regulatory Decision Points Defining Context of Use and Challenges to Deploying Wearables and Digitial Technologies Peter Como (FDA Co-chair) & Jesse Cedarbaum (CAMD Co-chair)
3:50-4:05 pm	BREAK
4:05-4:35 pm	KEY RECOMMENDATIONS: SESSIONS III-V
4:35-4:45 pm	Wrap-up and Looking Ahead Diane Stephenson & Stephen Arneric