

Patient Perspectives on the Impact of Rare Diseases: Bridging the Commonalities

Public Meeting

April 29, 2019

FDA White Oak Campus

10903 New Hampshire Avenue, Building 31, the Great Room (Room 1503-C)

Silver Spring, Maryland 20993

Agenda

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| 12:15 PM – 1:00 PM | Registration |
| 1:00 PM – 1:10 PM | Opening Remarks
Speaker: Janet Maynard, MD, MHS, Director, Office of Orphan Products Development, FDA |
| 1:10 PM – 1:20 PM | Meeting Overview
Moderator: Andrea Furia-Helms, MPH, Director, Patient Affairs Staff, FDA |
| 1:20 PM – 2:40 PM | First Session
Panel Discussion
Facilitated Group Discussion
Facilitator: Susan Chittooran, MSW, Patient Affairs Staff, FDA |
| 2:40 PM – 2:55 PM | Remarks by FDA's Principal Deputy Commissioner and Acting Chief Information Officer
Speaker: Amy Abernethy, MD, PhD |
| 2:55 PM – 3:15 PM | Break |
| 3:15 PM – 4:15 PM | Second Session
Panel Discussion
Facilitated Group Discussion
Facilitator: Susan Chittooran, MSW, Patient Affairs Staff, FDA |
| 4:15 PM – 4:50 PM | Open Public Comment |
| 4:50 PM – 5:00 PM | Closing Remarks
Speaker: Janet Maynard, MD, MHS, Director, Office of Orphan Products Development, FDA |