

**FOOD AND DRUG ADMINISTRATION (FDA)  
Center for Drug Evaluation and Research (CDER)**

*Antimicrobial Drugs Advisory Committee (AMDAC) Meeting  
FDA White Oak Campus, Building 31 Conference Center, the Great Room (Rm. 1503)  
10903 New Hampshire Avenue, Silver Spring, Maryland  
July 26, 2018*

**AGENDA**

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*The committee will discuss new drug application (NDA) 210607, tafenoquine tablet, 100 milligram (mg), sponsored by 60 Degrees Pharmaceuticals, LLC, for the proposed indication of prevention of malaria in adults for up to 6 months of continuous dosing.*

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8:30 a.m.	Call to Order and Introduction of Committee	<b>Lindsey Baden, MD</b> Chairperson, AMDAC
8:40 a.m.	Conflict of Interest Statement	<b>Kalyani Bhatt, BS, MS</b> Acting Designated Federal Officer, AMDAC
8:45 a.m.	FDA Opening Remarks	<b>Yuliya Yasinskaya, MD</b> Clinical Team Leader Division of Anti-Infective Products (DAIP) Office of Antimicrobial Products (OAP) Office of New Drugs (OND), CDER, FDA
8:55 a.m.	<b>APPLICANT PRESENTATIONS</b>	<b>60 Degrees Pharmaceuticals, LLC</b>
	Overview and Background Development of ARAKODA™	<b>Geoffrey Dow, PhD</b> Chief Scientific Officer and CEO 60 Degrees Pharmaceuticals, LLC
	Unmet Medical Need for ARAKODA™ Military and Civilian Travelers	<b>Stephen Toovey, MD, PhD</b> Infectious and Tropical Disease Physician Pegasus Research, Switzerland
		<b>Mark Reid, MBA</b> ADF Veteran Principal Consultant Clinical Network Services, Pty Ltd, Australia
	Efficacy	<b>Jonathan Berman, MD, PhD</b> Senior Vice President for Clinical Affairs Fast-Track Drugs & Biologics, LLC
	Safety	<b>Bryan Smith, MD</b> Chief Medical Officer 60 Degrees Pharmaceuticals, LLC
	Neuropsychiatric Safety	<b>Geoffrey Dow, PhD</b>
	Future ARAKODA™ Use	<b>Stephen Toovey, MD, PhD</b>

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**AGENDA (cont.)**

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10:25 a.m. Clarifying Questions

10:40 a.m. **BREAK**

10:50 a.m. **FDA PRESENTATIONS**

Presentation of Clinical Efficacy

**Xianbin Li, PhD**

Statistical Reviewer

Division of Biometrics IV

Office of Biostatistics

Office of Translational Sciences, CDER, FDA

Presentation of Nonclinical Pharmacology  
and Toxicology

**Owen McMaster, PhD**

Pharmacology/Toxicology Reviewer

DAIP, OAP, OND, CDER, FDA

Presentation of Clinical Safety

**Sheral Patel, MD**

Medical Officer

DAIP, OAP, OND, CDER, FDA

12:20 p.m. Clarifying Questions

12:35 p.m. **LUNCH**

1:30 p.m. **OPEN PUBLIC HEARING**

2:30 p.m. **BREAK**

2:40 p.m. Questions to the Committee/Committee Discussion

4:00 p.m. **ADJOURNMENT**