FOOD AND DRUG ADMINISTRATION (FDA)

Center for Drug Evaluation and Research (CDER)

Pharmacy Compounding Advisory Committee (PCAC) MeetingJune 8, 2022

AGENDA

During the **morning session**, the committee will discuss two bulk drug substances nominated for inclusion on the list of bulk drug substances that can be used to compound drug products in accordance with section 503A of the FD&C Act: enclomiphene citrate and glutathione.

June 8, 2022, Al	M Session		
9:30 a.m.	Call to Order	Allen J. Vaida, BSc, PharmD, FASHP Acting Chairperson, PCAC	
9:35 a.m.	Introduction of Committee and Conflict of Interest Statement	Takyiah Stevenson, PharmD Designated Federal Officer, PCAC	
9:45 a.m.	FDA INTRODUCTORY REMARKS	Frances Gail Bormel, RPh, JD Director Office of Compounding Quality and Compliance (OCQC) Office of Compliance (OC), CDER, FDA	
9:55 a.m.	FDA Investigational New Drug (IND)/Expanded Access Presentation	Lori Bickel, JD Regulatory Counsel Division of Medical Policy Development (DMPD) Office of Medical Policy (OMP), CDER, FDA	
10:05 a.m.	SECTION 503A BULK DRUG SUBSTANCES LIST – ENCLOMIPHENE CITRATE		
	FDA PRESENTATION	Madeline Wolfert, MD Physician Pharmacy Compounding Review Team Office of Specialty Medicine (OSM) Office of New Drugs (OND), CDER, FDA	
	Clarifying Questions from the Committee		
	NOMINATOR PRESENTATIONS	Marwa Elsaied, PharmD, RPh and Thomas Masterson III, MD Empower Pharmacy	
	Clarifying Questions from the Committee		
10:40 a.m.	OPEN PUBLIC HEARING		
10:55 a.m.	COMMITTEE DISCUSSION AND VOTE		
11:05 a.m.	Break		

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AGENDA (cont.)

11:15 a.m. SECTION 503A BULK DRUG SUBSTANCES LIST – GLUTATHIONE

FDA PRESENTATION Emily Kneeream, PharmD

Clinical Analyst

Pharmacy Compounding Review Team

OSM, OND, CDER, FDA

Clarifying Questions from the Committee

NOMINATOR PRESENTATION A.J. Day, PharmD

Professional Compounding Centers of America and

National Community Pharmacists Association

Clarifying Questions from the Committee

12:15 p.m. **OPEN PUBLIC HEARING**

12:30 p.m. COMMITTEE DISCUSSION AND VOTE

12:40 p.m. LUNCH

FOOD AND DRUG ADMINISTRATION (FDA)

Center for Drug Evaluation and Research (CDER)

Pharmacy Compounding Advisory Committee (PCAC) Meeting June 8, 2022

AGENDA (cont.)

During the **afternoon session**, the committee will discuss two additional bulk drug substances nominated for inclusion on the list of bulk drug substances that can be used to compound drug products in accordance with section 503A of the FD&C Act: ammonium tetrathiomolybdate and ferric subsulfate. The committee will also discuss one drug being considered for inclusion on the list of drug products that may not be compounded because they have been withdrawn or removed from the market because they have been found to be unsafe or not effective (Withdrawn or Removed List, codified at 21 CFR 216.24): lorcaserin hydrochloride.

June 8, 2022, PM Session

1:25 p.m. SECTION 503A BULK DRUG SUBSTANCES LIST – AMMONIUM TETRATHIOMOLYBDATE

(ATTM)

FDA PRESENTATION Raquel Tapia, MD

Physician

Pharmacy Compounding Review Team

OSM, OND, CDER, FDA

Clarifying Questions from the Committee

NOMINATOR PRESENTATION Mark Rosenberg, MD

Pharmacy Solutions

Clarifying Questions from the Committee

2:25 p.m. **OPEN PUBLIC HEARING**

2:40 p.m. COMMITTEE DISCUSSION AND VOTE

2:50 p.m. **BREAK**

3:05 p.m. SECTION 503A BULK DRUG SUBSTANCES LIST – FERRIC SUBSULFATE

FDA PRESENTATION Anam Tariq, DO, MHS

Physician

Pharmacy Compounding Review Team

OSM, OND, CDER, FDA

Clarifying Questions from the Committee

NOMINATOR PRESENTATION

Clarifying Questions from the Committee

3:45 p.m. **OPEN PUBLIC HEARING**

4:00 p.m. COMMITTEE DISCUSSION AND VOTE

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

Pharmacy Compounding Advisory Committee (PCAC) Meeting June 8, 2022

AGENDA (cont.)

4:10 p.m.	BREAK	
4:20 p.m.	Conflict of Interest Statement	Takyiah Stevenson, PharmD Designated Federal Officer, PCAC
4:25 p.m.	WITHDRAWN OR REMOVED LIST PROCESS	Gabrielle Cosel, MSc Director Division of Compounding Policy and Outreach OCQC, OC, CDER, FDA
4:35 p.m.	DRUGS TO BE CONSIDERED FOR THE WITHDRAWN OR REMOVED LIST – LORCASERIN HYDROCHLORIDE	
	FDA PRESENTATION	Marianne San Antonio, DO Physician Pharmacy Compounding Review Team OSM, OND, CDER, FDA
	Clarifying Questions from the Committee	
4:50 p.m.	OPEN PUBLIC HEARING	
5:05 p.m.	COMMITTEE DISCUSSION AND VOTE	
5:15 p.m.	ADJOURNMENT	