FDA Public Meeting:
To Discuss Development of a List of Pre-DSHEA Dietary Ingredients

Tuesday, October 3, 2017; 8:00 am – 5:00 pm

Center for Food Safety and Applied Nutrition
U.S. Food and Drug Administration
Wiley Auditorium
5001 Campus Drive
College Park, MD 20740

No. FDA-2017-N-4625

AGENDA

8:00AM **Registration**

8:30AM **Greeting & Housekeeping Items**
Cara Welch*, Senior Advisor, Office of Dietary Supplement Programs, Center for Food Safety and Applied Nutrition, FDA*

8:35AM **Welcome and Opening Remarks**
Dr. Stephen Ostroff, *Deputy Commissioner for Foods & Veterinary Medicine, FDA*

8:45AM **Overview of Compiling an Authoritative List of Dietary Ingredients**
Steve Tave, *Director, Office of Dietary Supplement Programs, Center for Food Safety and Applied Nutrition, FDA*

9:15 – 10:30AM **Panel 1: What Evidence is Necessary to Show that an Ingredient was Marketed Before October 15, 1994?**
Moderator: Cara Welch, *Senior Advisor, Office of Dietary Supplement Programs, Center for Food Safety and Applied Nutrition, FDA*
Panelists:
Joe Betz, Ph.D., *Director, Office of Dietary Supplements Analytical Methods and Reference Materials Program, Office of Dietary Supplements, National Institutes of Health*
Dr. Pieter Cohen, *Associate Professor of Medicine, Harvard Medical School*
Loren Israelsen, *President, United Natural Products Alliance*
Duffy Mackay, N.D., *Senior Vice President, Scientific & Regulatory Affairs, Council for Responsible Nutrition*
Michael McGuffin, *President, American Herbal Products Association*

10:30 – 10:50 AM **Q&A Session**
Moderator: Cara Welch, *Senior Advisor, Office of Dietary Supplement Programs, Center for Food Safety and Applied Nutrition, FDA*

10:50 – 11:15AM **BREAK**

11:15 – 12:00PM **<u>Open Public Comment</u>**
Moderator: Cara Welch, *Senior Advisor, Office of Dietary Supplement Programs, Center for Food Safety and Applied Nutrition, FDA*

12:00 – 1:15PM **LUNCH**

1:15 – 2:30PM **<u>Panel 2: What Process Should Be Used to Develop the List of Pre-DSHEA Ingredients?</u>**
Moderator: Robert Durkin, *Deputy Director, Office of Dietary Supplement Programs, Center for Food Safety and Applied Nutrition, FDA*
Panelists:
Chuck Bell, *Programs Director, Consumers Union*
Daniel Fabricant, Ph.D., *Executive Director and CEO, Natural Products Association*
Laura MacCleery, *Director of Regulatory Affairs, Center for Science in the Public Interest*
Stephanie Scarmo, Ph.D., M.P.H., *Officer, Health Care Products Project, The Pew Charitable Trusts*
Jay Sirois, Ph.D., *Senior Director, Regulatory & Scientific Affairs, Consumer Healthcare Products Association*

2:30 – 2:50PM **<u>Q&A Session</u>**
Moderator: Robert Durkin, *Deputy Director, Office of Dietary Supplement Programs, Center for Food Safety and Applied Nutrition, FDA*

2:50 – 3:15PM **BREAK**

3:15 – 4:45PM **<u>Open Public Comment</u>**
Moderator: Cara Welch, *Senior Advisor, Office of Dietary Supplement Programs, Center for Food Safety and Applied Nutrition, FDA*

4:45 PM **<u>Wrap-Up</u>**

5:00PM **ADJOURN**