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 – BREAK
11:15AM

11:15 – Open Public Comment

12:00PM Moderator: Cara Welch, Senior Advisor, Office of Dietary Supplement Programs, Center for Food Safety and Applied Nutrition, FDA

12:00 – LUNCH
1:15PM

1:15 – Panel 2: What Process Should Be Used to Develop the List of Pre-DSHEA Ingredients?

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

2:50 – BREAK
3:15PM

3:15 – Open Public Comment

4:45PM Moderator: Cara Welch, Senior Advisor, Office of Dietary Supplement Programs, Center for Food Safety and Applied Nutrition, FDA

4:45 PM Wrap-Up
5:00PM ADJOURN