FDA Proposed Regulation
Current Good Manufacturing Practices (CGMPs)
Dietary Ingredients and Dietary Supplements
Public Stakeholder Meeting
April 29, 2003, College Park, MD

9:00-9:10 AM	**Welcome and Opening Remarks** Virginia Wilkening, Deputy Director, Office of Nutritional Products, Labeling, and Dietary Supplements, CFSAN/FDA, College Park, MD Peter Vardon, Economist, Division of Market Studies FDA/CFSAN, College Park, MD
9:10-9:40 AM	**Background and Proposal Highlights** Karen Strauss, Consumer Safety Officer, FDA/CFSAN, College Park, MD
9:40- 9:55 AM	Questions and Answers
9:55-10:10 AM	**Proposed Production and Process Controls** Sara J. Dent Acosta, Consumer Safety Officer FDA San Diego Resident Post, San Diego, CA
10:10-10:25 AM	Questions and Answers
10:25-10:40 AM	Morning Break
10:40-11:00 AM	**Proposed Laboratory Operations** Steven Musser, Lead Scientist for Chemistry FDA/CFSAN, College Park, MD
11:00-11:15 AM	Questions and Answers
11:15-11:20 AM	**Public Comment Period and Next Steps** Karen Strauss, Consumer Safety Officer
11:20-11:35 AM	**Economic Impact Analysis** Peter Vardon, Economist, Division of Market Studies FDA/CFSAN, College Park, MD
11:35-12:00 Noon	Questions and Answers
12:00-1:30 PM	Lunch (on your own—see restaurant guide)
1:30-1:50 PM	**Regulatory Flexibility Act and How to Comment** Richard Williams, Director, Division of Market Studies FDA/CFSAN, College Park, MD Marie Falcone, Small Business Representative, FDA Central Region
1:50-2:00 PM	Questions and Answers

2:00-2:30 PM	**Small Business Questions on Proposed Requirements** Richard Williams, Karen Strauss, Sara Dent Acosta, and Peter Vardon
2:30-3:45 PM	**Breakout Sessions**
3:45-5:00 PM	**Breakout Session Summaries and Discussion**