

# Understanding the Poison Prevention Packaging Act (PPPA)

## March 24, 2004

8:00 – 8:30	Registration (Hearing Room, 420)
8:30 – 8:35	Introduction – Suzanne Barone, Directorate for Health Sciences
8:35 – 8:45	Welcome – Commissioner Mary Gall
8:45 – 9:00	Overview of the Commission – Jacqueline Elder, Assistant Executive Director, Office of Hazard Identification and Reduction
9:00 – 9:40	PPPA – Overview of the Act and Its Implementing Regulations – Lowell Martin, Office of General Counsel
9:40 – 10:00	PPPA Petitions for Exemption – Suzanne Barone, Directorate for Health Sciences
10:00 – 10:15	Break
10:15 – 11:00	Compliance Issues – Michelle Gillice, Office of Compliance and Geri Smith, Office of Compliance
11:00 – 11:45	Testing Issues – Suzanne Barone, Directorate for Health Sciences and Geri Smith, Office of Compliance
11:45 – 12:00	Child-Resistant Packaging Booklet – John Boja, Directorate for Health Sciences
12:00 – 1:30	Lunch on your own
1:30 – 2:30	Small Group Discussions 1  Session 1A - Prescription Drugs and Physician Samples (Hearing Room, 420) – Suzanne Barone, Directorate for Health Sciences and Geri Smith, Office of Compliance  Session 1B - Federal Hazardous Substances Act Labeling Requirements (Room 410) – Mary Toro, Office of Compliance and Michelle Gillice, Office of Compliance
2:30 – 2:45	Break
2:45 – 3:45	Small Group Discussions 2  Session 2A - Unit Packaging (Hearing Room, 420) – Suzanne Barone, Directorate for Health Sciences and Geri Smith, Office of Compliance  Session 2B - Poisoning Incident Data Sources (Room 410) – Tom Schroeder, Directorate for Epidemiology, John Boja, Directorate for Health Sciences, Robin Ingle, Directorate for Epidemiology
3:45 – 4:30	Wrap up, Questions, and Evaluation (Hearing Room, 420)