

**Building Effective Partnerships  
FDA and Stakeholders Public Meeting**

**Stanford Law School 8529**

'00 MAR 28 P1:56

**March 23, 2000**

**Agenda**

**I. Welcome** Mark Barnett, Moderator

**II. Opening Remarks** Jane Henney, M.D.

**III. Presentations of Leveraging Models**

- Tobias Massa, Ph.D.  
Eli Lilly and Company
- Erica Jones  
California Medical Review, Inc.
- Charles E. Sizer, Ph.D.  
National Center for Food Safety and Technology

**IV. Discussion of Proposed Leveraging Initiatives**

- Safety Review for New Products  
*Safety Assurance in Clinical Trials*
- Assuring Industry Compliance with Safety Regulations  
*Gene Therapy, Human Cellular and Tissue Based Products*
- Patient/Consumer Education on the Safe Use of Products  
*Risk Management*
- Safety-related Research  
*NCTR Chip Technology*
- Additional Comments

**V. Closing Remarks** Jane Henney, M.D.

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## CUSTOMER SURVEY

**NAME OF EVENT: Leveraging-Collaborating With Stakeholders**

**DATE** \_\_\_\_\_

**DID YOU GIVE/RECEIVE THE INFORMATION YOU WANTED/NEEDED/EXPECTED?** YES \_\_\_ NO \_\_\_

**IS THE DIALOGUE USEFUL TO YOU?** YES \_\_\_ NO \_\_\_

**WERE MATERIALS USEFUL TO YOU?** YES \_\_\_ NO \_\_\_

**WERE THE LISTENERS PROFESSIONAL, KNOWLEDGEABLE AND ABLE TO ANSWER YOUR QUESTIONS?**

YES \_\_\_ NO \_\_\_

**WOULD YOU PARTICIPATE IN ANOTHER SIMILAR EVENT?** YES \_\_\_ NO \_\_\_

**HOW CAN WE IMPROVE? COMMENTS/SUGGESTIONS:**

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