Panel 1: Accelerating Clinical Trials in the Development and Approval of Innovative Medical Products

Moderators:

<u>Laura Esserman, MD, MBA</u> - Professor of Surgery at UCSF <u>Janet Woodcock, MD</u> - Director, Center for Drug Evaluation and Research at FDA

Panelists:

<u>Amy Abernethy, MD, PhD</u> - Principal Deputy Commissioner at FDA

<u>Steven Goodman, MD, MHS, PhD</u> - Associate Dean of Clinical and Translational Research at Stanford

<u>Sandra Horning, MD</u> – Former Chief Medical Officer at Genentech/Roche

<u>Carl Peck, MD</u> - Chairman of NDA Partners

Alfred Sandrock, MD - Chief Medical Officer at Biogen

- What is your top suggestion for an innovation that could accelerate drug development and thereby increase the public's access to effective and cost-effective treatments?
- What is your top suggestion for what we should let go/stop doing and why?

