

**Panelists at Part 15 Public Hearing –  
 Conduct of Emergency Clinical Research  
 October 11, 2006**

	<b>Panelists</b>	<b>Title</b>
1.	Jeffrey Shuren, MD, JD (Presiding Officer)	Assistant Commissioner for Policy, Office of the Commissioner, FDA
2.	Michael Carome, MD	Associate Director for Regulatory Affairs, Office for Human Research Protections
3.	Sara Goldkind, MD, MA	Senior Bioethicist, Office of Critical Path Programs, Office of the Commissioner, FDA
4.	Bonnie M. Lee	Associate Director for Human Subject Protection Policy, Good Clinical Practice Program, Office of the Commissioner, FDA
5.	Joanne Less, PhD	Associate Director for Clinical Research and Government Affairs, Center for Devices and Radiological Health, FDA
6.	Catherine Lorraine, JD	Director, Policy Development and Coordination Staff, Office of Policy, Office of the Commissioner, FDA
7.	Diane Maloney, JD	Associate Director for Policy, Center for Biologics Evaluation and Research, FDA
8.	Robert Temple, MD	Associate Director for Medical Policy, Center for Drug Evaluation and Research, FDA
9.	Denise Zavagno, JD	Associate Chief Counsel for Biologics, Office of the Chief Counsel, Office of the Commissioner, FDA