

**Panelists for the Part 15 Public Hearing –
Electronic Submission of Regulatory Information, and Creating an Electronic
Platform for Enhanced Information Management**

	Panelists	Title
1.	Janet Woodcock, M.D. (Presiding Officer)	Deputy Commissioner for Operations, FDA
2.	Randall Lutter, Ph.D.	Associate Commissioner for Policy and Planning, Office of the Commissioner, FDA
3.	Randy Levin, M.D.	Director for Health and Regulatory Data Standards, Office of Critical Path Programs, Office of the Commissioner, FDA
4.	Armando Oliva, M.D.	CAPT, U.S. Public Health Service, Deputy Director for Bioinformatics, Office of Critical Path Programs, FDA
5.	Kevin Fain, J.D.	Associate Chief Counsel for Drugs, Office of the Chief Counsel, FDA
6.	Lana Skirboll, M.D.	Director, Office of Policy and Planning, NIH
7.	Ken Buetow, Ph.D.	Associate Director for Bioinformatics and Information Technology, NCI, NIH