Bibliography Advisory Committee for Pharmaceutical Science April 13-14, 2004

- Hussain, Ajaz S., "FDA's Initiative on Pharmaceutical Quality Systems for the 21st Century and the Debate on Delivered Dose Uniformity for Inhalers", prepared for submission to *Respiratory Drug Delivery IX*, p. 1-12 [Not yet published]
- Schadt, Randall, "Process Analytical Technology Changing the Validation Paradigm", *American Pharmaceutical Review*, Vol. 7, Issue 1 (2004): 58-61
- 3. Cooley, Rick E., Egan, J. Carmel, "The Impact of Process Analytical Technology (PAT) on Pharmaceutical Manufacturing", *American Pharmaceutical Review*, Vol. 7, Issue 1 (2004): 62-68
- 4. Arrivo, Steven M., "The role of PAT in Research and Development", *American Pharmaceutical Review*, Vol. 6, Issue 2 (2003): 46-53
- 5. Tothfalusi, Laszlo, Endrenyi, Laszlo, "Limits for the Scaled Average Bioequivalence of Highly Variable Drugs and Drug Products" *Pharmaceutical Research*, Vol. 20, No. 3, (March 2003): 382-389
- Lionberger, Robert A., Lawrence X, Yu, et. al., "Development of Bioequivalence Methods for Topical Dermatological Drug Products", prepared for submission to *Pharmaceutical Research, p. 1-32* [Not yet published]