

## **Commitment to Continuing Education**

Paladin Medical, Inc. is committed to continuing education, both as a lecturer, author and student.

### **Presentations** Selected Examples 2007-2002

*Navigating Standards & Regulations for Medical Textiles*, IFAI Medical Textile Symposium, October 31, 2006

*GLPs for Medical Device Sponsors and Combination Products*, CRL invitational conference, August 25, 2005

*Biomaterials Qualification and Selection for Spinal Implants*, MDM Conference, November 1, 2005

*Practical Tips for Regulatory Compliance: Top Guidance and Website Help*, AusBiotech in Sydney Australia, April 22, 2004, and Medical Alley, March 13, 2004

*Regulatory Changes Impacting Medical Device Development: The Good, the Bad and the Ugly*, U of MN Annual Medical Device Design Conference, April 25, 2003

*Surface Analysis: Tales of Woe & Lessons From the Past for the Future*, MDM Minneapolis, October 22-24, 2002 and in Anaheim, CA, February 2003

*GLP Compliance for Combination Products*, Barnett International, November 18, 2002

*GLP Training for Sponsors: Corporate Forum (Workshop)*, Society for Biomaterials, April 24, 2002

*Customized GLP Training for Sponsor*, AnokaRamsey County College Education Grant, January 25, 2002

### **Continuous Service Improvements** Selected Examples 2007-2002

NASS, Austin, TX October 23-27, 2007

*Developments in Medical Device Testing: Sterilization and Preclinical Testing*, NAMSA May 17, 2007

*Investigational Testing of Medical Devices in Canada*, Advamed, September 7, 2006

AAOS, Chicago, IL March 21-25-2007

*7<sup>th</sup> World Congress of Biomaterials*, Sydney, Australia, May 17-21, 2004

*Litigation Exposures in Clinical Trials*, Medical Alley, January 21, 2004

*HIPPA Update*, Medical Alley, September 26, 2002, Minneapolis, MN

*Data Management Process for Clinical Trials*, Medical Alley, August 21, 2002, Minneapolis, MN

*Good Laboratory Practices – Professional Training Course*, IQPC, July 24-25, 2002, Philadelphia, PA

*21 CFR Part 11– Electronic Records; Electronic Signatures*, Medical Alley, April 5, 2002, Mpls. MN

**Published Articles** Selected Examples

Duncan, Elaine. "123: Development and Regulation of Medical Technology." *Industrial Fabric Products Review*, November 2006.

Duncan, Elaine, et al. "Standards." Chapter 10 of *Biomaterials Science: 2nd Edition*, Elsevier, Inc. 2004.

Duncan, E. "Good Laboratory Practices for Testing Biomaterials." *Biomaterials Forum*, September-October 2001.

Duncan, E. "Combination Products and Design Control." *SurFACTS in Biomaterials*, Spring, 2001.

Duncan, E. "Combination Products Can Create Combination Compliance Headaches." *SurFACTS in Biomaterials*, Vol. 4, Issue 2. Summer 1999.

Duncan, E. *Regulatory Update for SurFACTS Readers*, Vol. 3, Issue 3. 1998, p. 7.

Duncan, E. "Potential Impact of Regulatory Changes on Medical Devices." *New Venture Review Minnesota*, Vol. 1. Sept/Oct 1988, pp. 395-397.

Duncan, E. "The Value of a Risk Management Approach to Clinical Trial Design and Management." *Critical Reviews in Biomedical Engineering*, Vol. 25, Issue 2. 1997, pp. 84-85.

Duncan, E. "Custom Devices: Until FDA Finalizes a Guidance, What Can You Do?" *The Validation Consultant*, Vol. 4 No. 9. September 1997.

Duncan, E. "What is a Biomaterial?" *MDDI Magazine*, Canon Communications, Inc. January 1990, p. 138.

Duncan, E. "Biostability, Stability and Controlled Instability" *MDDI Magazine*, Canon Communications, Inc. March 1990, p. 94.