

Eric Stephan Kastango

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PROFESSIONAL EXPERIENCE

CLINICALIQ, LLC and CriticalPoint, LLC

Principal, Florham Park, New Jersey

December 1999 – Present

Initiated and developed all aspects of company operations whose mission is to become leading provider of quality aseptic process and business development strategies for the pharmaceutical, medical device and healthcare industry. Areas of expertise include aseptic processing (pharmacy (USP Chapters <795> and <797> and pharmaceutical industry-based), medical device manufacturing, regulatory and accreditation compliance auditing, and the development and implementation of evidence-based quality systems. Client portfolio includes:

Baxter Healthcare (US, Australia, Canada, China, Ireland, Japan, New Zealand and Puerto Rico)

Express Scripts, Inc.

Coram Healthcare

SoluNet, LLC

MedStar Health System

Richie's Pharmacy

B. Braun Manufacturing Division

B. Braun/CAPS

Plum Creek Pharmaceuticals, Inc.

Child Health Corporation of America

Greater New York Hospital Association

US Department of Veterans Affairs, VISN 3, VISN 6

Pharmacy OneSource

Home Healthcare Laboratory of America

Tyco/Mallinckrodt, Inc.

American Society of Health-System Pharmacists

University of California, Irvine Medical Center

Shands at the University of Florida Hospital

ForHealth Technologies, Inc.

Baxa Corporation

The Cleveland Clinic

University of Texas, MD Anderson Cancer Center

Seattle Children's Regional Hospital and Medical Center

CORAM HEALTHCARE CORPORATION

Vice President, Pharmacy Services, Whippany, New Jersey

January 1999 – December 1999

- Responsible for all aspects of clinical services, policy and procedures, clinical initiatives, staff training, field support of 88 branches in 44 states. Responsible for implementing standardized clean room operating guidelines through Computer-Based Training Modules (Good Compounding Practices), physical plant design requirements and policy and procedures. Implemented Home Healthcare Laboratory of America Clinical Laboratory Service Model, Primary Care Management Model with Vice President of Nursing Services, Aminoglycoside Patient Assessment Tool and Process and Calcium Phosphate TPN Stability Process.

COGNITIVE DESIGN ASSOCIATES, Inc

Principal, Upper Saddle River, New Jersey

April 1997 – January 2004

- Initiated and developed all aspects of company operations whose mission is to become leading provider of customized educational strategies for performance improvement in the healthcare industry. CDA facilitates the achievement of identified organizational goals through a performance consulting methodology. Responsible for the development, implementation, and support of customized Interactive Computer-Assisted Learning Modules (ICALMs) for employee competency validation. Additional service offerings include process design, service model revision, clinical and operational policy/procedure development, competency tool development, and instructional design for the hospital and homecare markets.

MILLERS HOMECARE SERVICES, Inc.

Executive Vice-President, Hawthorne, New Jersey
August 1996 – May 1997

- Responsible for all aspects of business operations for Millers Homecare Services, which included the following divisions: Home Medical Equipment and Respiratory, Rehabilitation Technology Services and Infusion and Clinical Services. Annual sales for 1996 were 8.5 million dollars. Responsible for closing a one million dollar supply management contract with the Visiting Nurse Association of Northern New Jersey that provided nursing and patient surgical/medical supplies. Position eliminated due to a potential acquisition.

BAXTER COMPASS, a business unit of Baxter Healthcare Corporation, IV Systems Division.

Compass Facility Manager, Edison, New Jersey
January 1994-August 1996

- Responsible for the start-up and day-to-day operation of a US Food and Drug Administration registered parenteral nutrition and cardioplegia solution manufacturing operation. Operations supported six area hospitals. Areas of responsibilities included compliance with cGMPs, staff development and supervision, purchasing and inventory management, budget and finance management and all aspects of Quality System Management. Facility underwent two FDA site audits without the issuance of any 483's.

HMSS, Inc.

Branch Operations Manager, Wayne, New Jersey
September 1993-January 1994

- Responsible for the day-to-day operation of a home infusion therapy facility. Areas of responsibilities included: development and supervision of three department managers, budget and finance management. Coram, Inc purchased company in March 1994. Position was to be eliminated.

CAREMARK, Inc.

Staff Pharmacist, March 1984-August 1985
Manager of Pharmacy Services, August 1985-September 1986 and January 1990 -January 1993,
Branch Manager, January 1993-August 1993

- Participated in the team management of a branch staff of eighty (80) encompassing pharmacy, nursing and patient services in Totowa, New Jersey. Branch generated sales of forty-one (41) millions dollars and a profit of eleven (11) million dollars in 1992.
- Responsible for the management of the Manhattan Clinical Service Center that provided home infusion and nursing services for over one hundred and fifty (150) HIV patients in New York and New Jersey. The center employed twenty-two (22) clinical and service specialists.
- Responsible for the development and implementation of a clinical pharmacy-training program for the Women's Health Initiative, and trained one hundred and fifteen (115) clinical pharmacists in tocolytic therapy.
- Co-authored and edited a physician sponsored Investigational New Drug Application (IND) for L-glutamine and its use in the home TPN patients. The Food and Drug Administration approved IND 37,331 in May 1991. A paper was published in the Journal of Parenteral and Enteral Nutrition.

TOKOS MEDICAL CORPORATION

Area Clinical Pharmacist, Edison, New Jersey
January 1989-January 1990

- Responsible for managing all women's health (tocolytic and home infusion) patients out of four regional centers from Maine to New Jersey and out to Ohio.
- Responsible for establishing local provider relationships with independent home infusion operations in order to serve infusion patients.
- Designed a trifold-marketing brochure detailing the home infusion services provided by Tokos Medical Corporation. Brochure was implemented nationwide in 1989.

NEW ENGLAND CRITICAL CARE, Inc

Manager of Pharmacy Services, January 1987-September 1987
Branch Operations Manager, September 1987-January 1989

- Responsible for the day-to-day management of an operation that experienced a two hundred percent (200%) increase in patient census with an associated increase in gross sales from six million to eleven million dollars in 1988
- Responsible for obtaining outpatient licensure for the first homecare blood bank/transfusion service in New Jersey
- Participated in several corporate groups that focused on enhancements to clinical operating policy and procedures, computerized pharmacy operating systems and employee incentive programs

EDUCATION

Massachusetts College of Pharmacy and Allied Health Sciences Boston, Massachusetts	Bachelor of Science, Pharmacy March 1983
University of Phoenix Phoenix, Arizona	Masters, Business Administration March 2001 (Honors)
University of Tennessee, Department of Pharmaceutical Sciences Memphis, Tennessee <i>The Comprehensive Industrial Course in the Preparation of Parenteral Products</i>	Certificate Program March 2007
BD Six Sigma Program, Certified Green Belt	In Process July and September 2008
Purdue University, West Lafayette, Indiana	65 hours live training Certification in Nuclear Pharmacy May 2009

LICENSURE

Registered Pharmacist, State of New Jersey
Registered Pharmacist, State of New York (Inactive)

APPOINTMENTS

The United States Pharmacopoeia, Council of Experts

Elected member of the Sterile Compounding Committee (USP Chapter <797>), 2005-2010

Pharmacy Purchasing and Products Magazine

Editorial Board 2004-present

APPOINTMENTS, continued

International Journal of Pharmaceutical Compounding

Sterile-Product Preparation Section Editor, 2000-2005

Contributing Author

American Society of Health-System Pharmacists

-Vice Chairman, Council of Educational Affairs, 2002-2003

-Member, Committee for the Home, Ambulatory, and Chronic Care Practitioners, 2002-present

-Member, Council on Educational Affairs, 2001-2002

-Chair, Committee on Communications, 2000-2001

-Professional Practice Committee Member, 1999-2000

-Home Care Practitioners Section Member, 1999

Cleanrooms Magazine

Editorial Board 2007-present

National Home Infusion Association

-Standards Committee Member, 2002-2005

Visiting Nurse Services of New York (VNSNY)

-Medical and High-Tech Advisory Board, 1991-1999

FACULTY

Instructor, Baxa Star Center, Englewood, CO

September 2007-Present

Instructor, APEx Sterile Training Program, Bethlehem, PA

June 2005-December 2006

AWARDS

Eagle Scout, Boys Scouts of America, June 1977

Fellow, American Society of Health-System Pharmacists, March 2001

PROFESSIONAL ASSOCIATIONS/MEMBERSHIPS

American Society of Health-System Pharmacists

May 1983-Present

National Home Infusion Association

April 1997-May 2006

PDA

December 1999-Present

Institute for Environmental Sciences and Technology

December 1999 -Present

Association for Professionals in Infection Control and Epidemiology

November 2007-Present

The American Society for Quality

November 2003-Present

JOURNAL PUBLICATIONS

Kastango, ES. The Top Ten Gaps in USP Chapter <797> Compliance. *Pharmacy Purchasing and Products Magazine*. TBD

Kastango ES. Spread the Word: Aseptic Technique Prevents Infection. *Pharmacy Purchasing and Products Magazine*. April 2009; 6-9

Okeke CC, Newton DW, Kastango ES, and Allen LV. Basics of Compounding: United States Pharmacopeia General Chapter <797> Pharmaceutical Compounding-Sterile Preparations, Part 10: First Revision: The Main Changes, Events and Rationale. *Int J Pharm Compd*. 2008; 12:530-536.

Kastango, ES. Compounded Sterile Preparations Continue to Harm or Kill Patients. *Journal of Pharmacy Practice and Research*. 2008; 3:173-174.

Kastango, ES, Wagner JT, Kastango KB, Kastango NE, and Wagner TJ. Generation of particulate matter during handling of needle and syringe packaging. *Am J Health-Syst Pharm*. 2008; 65:1443-50.

Kastango, ES. USP Chapter <797> and Environmental Monitoring. *Clinical Microbiology Newsletter*.30:14, 2008

Kastango, ES. Microbial Air-Sampling Equipment, Part 1: Meeting United States Pharmacopeia Chapter <797> Standard. *Int J Pharm Compd*. 2008; 12:216-268.

Kastango ES. The revised USP Chapter <797> in review. *Cleanrooms Magazine*, April 2008.

Kastango ES. The updated <797> in Review: How the Changes to USP Chapter Impact Your Practice. *Pharmacy Purchasing and Products Magazine*. February 2008; 18-22

Kastango ES. USP <797>: The First Step to Compliance. *Pharmacy Purchasing and Products Magazine*. July 2007, 10-11.

Kastango, ES. Quality Assurance and <797> Compliance: Fact versus Fiction-Which are your environmental monitoring and sterility testing programs based on? *Pharmacy Purchasing and Products Magazine*. May 2007, 20-24.

Kastango ES. So What Have You Done Today to Improve Patient Safety? *Pharmacy Purchasing and Products Magazine*. Special Issue: Cleanrooms & Compounding 2006: 1

Kastango ES. A Review of USP's Updates to <797>: Don't Just Know About It. *Pharmacy Purchasing and Products Magazine*. Special Issue: Cleanrooms & Compounding 2006; 3:8-10.

Kastango ES. Putting the Science back into the "Art and Science" of Compounding. *Int J Pharm Compd*. 2006; 10:263-268.

Trissel LA, Zhang Y, Douglass K and Kastango ES. Extended Stability of Oxytocin in Common Infusion Solutions. *Int J Pharm Compd*. 2006; 10:156-158.

Kastango ES. Pharmacy Compounding Urban Legends. *Int J Pharm Compd*. 2006; 10:28-31.

Kastango ES, Faylor, K. The Importance of Environmental Monitoring, Part II: Surface Testing. *Pharmacy Purchasing and Products Magazine*. 2005; 2: 24-26.

Kastango ES. USP <797>: Making the case for improving environmental controls in pharmaceutical compounding. Supplement to *Infusion*. 2005; 11(4):S1-S12

Kastango ES. A Blueprint for Implementing USP Chapter <797>, Pharmaceutical Considerations: Sterile Preparations, *Am J Health-Syst Pharm*. 2005; 62:1271-88.

JOURNAL PUBLICATIONS, continued

Kastango ES. The Importance of Environmental Monitoring, Part I: Air Quality. *Pharmacy Purchasing and Products Magazine*. 2005; 4:4-5.

Kastango ES. A Practical Guide to Aseptic Technique Verification: Policies and Procedures that Meet USP Chapter <797> Requirements. *Pharmacy Purchasing and Products Magazine*, 2005; 3:16-18.

Wagner J, Kastango ES. Understanding Cleanroom Design. *Pharmacy Purchasing and Products Magazine*. 2005; 2: 21-27.

Kastango ES. Using Automated Compounding Devices (ACDs) in the Practice of Pharmacy, *Int J Pharm Compd*. 2005; 9:15-21.

Kastango ES. Using Equipment and Products to Successfully Meet USP Chapter <797> Standards. *Pharmacy Purchasing and Products Magazine*. 2004; 1: 12-17.

Kastango ES, Bradshaw BD. USP Chapter 797: Establishing a practice standard for the profession of pharmacy. *Am J Health-Syst Pharm*. 2004; 61:1928-38

Kastango ES. Quality-Control Analytical Methods: USP Chapter <797>Compounded Sterile Preparations Sterility Requirements and Their Relationship to Beyond-Use Dating. *Int J Pharm Compd*. 2004; 8: 393-397.

Kastango ES. The ASHP Discussion Guide for Compounding Sterile Preparations: Summary and Implementation of USP Chapter 797. Published by ASHP in collaboration with Baxter Healthcare. April 2004.

Kastango ES, Trissel LA, Bradshaw B. An Ounce of Prevention: Controlling Hazards in Extemporaneous Compounding Practices. *Int J Pharm Compd*. 2003; 5: 401-416

Kastango ES, Bradshaw B. Sterile Product Compounding: Developing Quality-Based Standards for Pharmacy Practice. *Infusion*. 2002; 8: 23-27

Kastango, ES. The Cost of Quality for Sterile Products, *Int J Pharm Compd*. 2002; 6: 404-407

Stranz M, Kastango ES. A Review of pH and Osmolarity. *Int J Pharm Compd*. 2002; 6: 216-220

Kastango ES, Hadaway, L. New Perspectives on Vancomycin Use in Homecare, Part 2. *Int J Pharm Compd*. 2002; 6: 55-57

Kastango ES, Hadaway L. New Perspectives on Vancomycin Use in Homecare, Part 1. *Int J Pharm Compd*. 2001; 5: 465-469

Kastango ES. Compounding Containment Devices: Buyer Be Aware. *Int J Pharm Compd*. 2001; 5: 384-388

Kastango ES, Douglass K. Quality Assurance for Sterile Products. *Int J Pharm Compd*. 2001; 5: 246-253

Kastango, ES, DeMarco S. Pharmacy Cleanroom Project Management Considerations: An Experience-Based Perspective. *Int J Pharm Compd*. 2001; 5: 221-225

Douglass K, Kastango ES. Consolidation of Pharmacy Compounding Services: An Alternative to Outsourcing. *Int J Pharm Compd*. 2001; 5: 140-144

Kastango ES. Sterile-Product Preparations: Mix or Buy? *Int J Pharm Compd*. 2001; 5:59-63

Kastango ES, Douglass, K. Improving the Management, Operations and Cost Effectiveness of Sterile-Product Compounding – Improve the ability to predict employee efficiency, cost of service and product wastage while simultaneously improving quality. *Int J Pharm Compd*. 1999; 3:253-258

JOURNAL PUBLICATIONS, continued

Kastango ES. Circadian Rhythm Chemotherapy for Renal Cell Cancer. *Alternative Site Care Case Study, CADD-Plus Ambulatory Infusion Pumps*, Number 11, July 1993, Pharmacia Deltec Inc., St. Paul, Minnesota.

Kastango ES. Tocolytic Therapy. *Caremark Pharmacy Report*, Vol. 4, 2. December 1991

REFERENCES AND BOOKS

Kastango ES, St. John KH, Weber, DJ. (2009) Chapter 61-Pharmacy. The APIC Text of Infection Control and Epidemiology. Washington, DC.

Kastango ES. (2008). Chapter 14-Finished Preparation Release Checks and Tests. In E. Clyde Buchanan & Philip J. Schneider (Eds.), Compounding Sterile Preparations, 3rd edition. Bethesda, MD. ASHP.

Kastango ES. (2008). Chapter 24-Environmental Quality and Control. In E. Clyde Buchanan & Philip J. Schneider (Eds.), Compounding Sterile Preparations, 3rd edition. Bethesda, MD. ASHP.

Kastango ES. (2008). Chapter 26- Personnel Training and Competency Evaluation of Garbing, Aseptic Work Practices. In E. Clyde Buchanan & Philip J. Schneider (Eds.), Compounding Sterile Preparations, 3rd edition. Bethesda, MD. ASHP.

Kastango ES. (2008). Chapter 13- Verification of Compounding Accuracy and Sterility. In E. Clyde Buchanan & Philip J. Schneider (Eds.), Compounding Sterile Preparations, 3rd edition. Bethesda, MD. ASHP.

Kastango ES. (2004). Chapter 21-End-Preparation Evaluation. In E. Clyde Buchanan & Philip J. Schneider (Eds.), Compounding Sterile Preparations, 2nd edition. Bethesda, MD. ASHP.

Kastango ES. (2004). Chapter 20-Process Validation. In E. Clyde Buchanan & Philip J. Schneider (Eds.), Compounding Sterile Preparations, 2nd edition. Bethesda, MD. ASHP.

Kastango ES. (2004). Chapter 18-Environmental Monitoring. In E. Clyde Buchanan & Philip J. Schneider (Eds.), Compounding Sterile Preparations, 2nd edition. Bethesda, MD. ASHP.

ELECTRONICS PUBLICATIONS

CriticalPoint Virtual Compounder™, 2007

Editor and Author of three (3) lessons in CriticalPoint's Sterile thirty-two (35) 1-hour ACPE approved Compounding eLearning Modules, 2007.

Clinical IQ, LLC (www.clinicaliq.com) and CriticalPoint (www.criticalpoint.info) websites

Kastango, ES. (2005). *Compounding Sterile Preparation, 2nd ed: A Multimedia Learning Tool CD-ROM*. American Society of Health-System Pharmacists, Bethesda, MD.

Kastango ES (2004). *The Comprehensive Sterile Compounding Compliance Gap Analysis and Risk Assessment Tool*. Available online: www.797complianceadvisor.com. American Society of Health-System Pharmacists, Bethesda, MD.

Kastango, ES. (2004). *Sterile Product Preparation: A Multimedia Learning Tool CD-ROM, 2nd edition*. American Society of Health-System Pharmacists, Bethesda, MD.

Kastango, ES. (2002). *Sterile Product Preparation: A Multimedia Learning Tool CD-ROM*. American Society of Health-System Pharmacists, Bethesda, MD.

Critical Issues in Homecare Compounding. Training Video, Clintec Nutrition Division, Baxter Healthcare, Inc., Deerfield, Illinois. 1997.

INTERNATIONAL PRESENTATIONS

Keynote speaker and presenter at “Updates in Sterile Compounding and Parenteral Nutrition” International Symposium, King Fahad Medical City (KFMC), Riyadh, Saudi Arabia, June 21-22, 2008.

International 7 City Speaking Tour in Japan, China, and Australia to inform/educate Key Healthcare and Political agencies/Pharmacists on the current US USP 797 experience and sterile compounding regulations”, Baxter Healthcare, April 4-18, 2008.

Keynote speaker and “Regulation of Aseptic Dispensing in Hospital Pharmacies: Federal, State, or...?” Society of Hospital Pharmacist of Australia (SHPA) 28th Federal Conference 2007, Sydney, NSW.

“Applying Aseptic Compounding Standards to protect patients and employee: Advances in Targeted Therapies: Where Next & How?” 5th Annual Hospital Pharmacists’ Association, Ireland (HPAI) meeting, Dublin, Ireland, January 18, 2007.

International 3 City Tour in Ireland to discuss current hospital aseptic compounding practices in the Republic of Ireland. Baxter Healthcare Ltd, January 15-20, 2007

“Practices and Requirements for the Compounding of Sterile Pharmaceuticals (CSPs)” Cleanrooms Europe 2006, Brno, Czech Republic, October 26, 2006

“USP Chapter <797>: A Compounding Patient Safety Roadmap Achieving Compliance”. Colegio de Farmaceuticos 2006 Convencion, Rio Grande, PR, August 25, 2006.

International 6 City Speaking Tour in Australia and New Zealand to inform/educate Key Healthcare and Political agencies/people on the current US USP 797 experience, Baxter Healthcare ANZ, August 7-14, 2006.

NATIONAL AND LOCAL PRESENTATIONS

“The Top 10 Gaps in USP 797 Compliance” Pharmacy OneSource Webinar, September 2, 2009

“Understanding the Environmental Sampling Requirements of USP <797>.” 28th Annual Meeting, Southwest Association of Clinical Microbiology, Ft. Worth, TX, September 3, 2009

“USP <797> Update 2009.” 2009 ASHP Summer Meeting, Rosemont, IL, June 17, 2009.

“Seven Deadly Myths of USP Chapter 797” Pharmacy OneSource Webinar, March, 2009

“USP <797>: Overcoming Challenges on the Road to Compliance.” 2009 NHIA Annual Conference & Exposition, Baltimore, MD, March 1, 2009.

“USP <797> Update 2008.” 2007 ASHP Summer Meeting, San Francisco, CA, June 10, 2008.

“Practical Application of USP to Pharmaceutical Compounding and Dispensing.” The United States Pharmacopeia Chapter <797> Workshops, Bethesda, MD, March and May 2008.

“USP <797>: New Requirements for Compounding Sterile Preparations.” NHIA 2008 Annual Conference, Phoenix, AZ, March 12, 2008.

“Understanding Pharmaceutical Compounding-USP Chapter <797>.” Virginia Association of Hematologists and Oncologists. Irvington, VA, March 7, 2008.

“USP 797 Basics.” BD Medical Surgical, National Sales Training Program, Charlotte, NC and Salt Lake City, UT, February and March 2008.

“Understanding the USP process and the recent changes to USP Chapter.” 2007 ASHP Midyear Clinical Meeting, Las Vegas, NV, December 5, 2007.

NATIONAL AND LOCAL PRESENTATIONS, continued

“USP <797> Update 2007.” 2007 ASHP Summer Meeting, San Francisco, CA, June 27, 2007.

“A Practical and Evidence-based approach to Surveying Pharmacies for Compliance to USP <795> and <797>.” NABP 103rd Annual Meeting, Portland, OR, May 19, 2007.

“USP <797> Update” ESTECH 2007 Conference, Institute of Environmental Sciences and Technology, Bloomingdale, IL, May 1, 2007.

“USP <797>: Understanding the past, present, and future of sterile compounding regulations”. Texas Society of Health-System Pharmacists’ 59th Annual Seminar, San Antonio, TX, April 22, 2007.

“Achieving USP 797 Compliance”. American Pharmacists Association 2007 Conference, Atlanta, GA, March 2007.

“Handling Hazardous Drugs: Implementation Strategies for a Safer Work Environment: Short-term and long-term compliance activities for USP 797 and the NIOSH Alert.” 2006 ASHP Midyear Clinical Meeting, Anaheim, CA, December 6, 2006.

“USP <797> Updates” California Society of Health-System Pharmacists Seminar, Sacramento, CA,

“Tuning UP Your Plan for USP 797, Parts I and II” 2006 ASHP Summer Meeting, Orlando, FL, June 28, 2006.

“USP <797>: Understanding the Air, Surface and Personnel Sampling Monitoring Requirements.” 2006 ASHP Midyear Clinical Meeting, Anaheim, CA, December 5, 2006.

“USP Chapter <797>: Surveying for Sterile Compounding Compliance”. NABP 102nd Annual Meeting Compliance Officer Programming, San Francisco, CA, April 11, 2006

“Extemporaneous Sterile Compounding for Preparing Clinical Supplies Roundtable: Environmental Controls Required During Extemporaneous Sterile Pharmaceutical Compounding.” 2006 American Association of Pharmaceutical Scientists (AAPS) Annual Meeting and Exposition, San Antonio, TX. November 1, 2006

“Tuning up your plan for USP 797, Part I and Part II” 2006 ASHP Annual Meeting, Orlando, FL June 28, 2005.

“USP 797 Webinar” USP sponsored web-based conference, Bethesda, MD, June 7, 2006.

“USP 797 Update: Next Steps” Tenet Healthcare 2006 Quality Summit. Dallas TX. February 1, 2006.

“Automating the Compounding of Sterile Products - A USP <797> Perspective” ForHealth Technology ASHP 39th MCM Exhibitor’s Theater. Las Vegas, NV December 6, 2005

“What’s Next for USP <797> Compliance” Baxa ASHP 39th MCM Exhibitor’s Theater. Las Vegas, NV December 5, 2005

“USP Chapter <797>: Steps to Compliance in 2006” 40th ASHP Midyear Clinical Meeting, Las Vegas, NV, December 4, 2005

“USP Chapter <797> and Infection Control”. Northern Chapter Monthly Meeting, Association for Professionals in Infection Control and Epidemiology, Montclair, NJ, October 26, 2005.

“Implementing Your USP 797 Plan 6: Issues Around Outsourcing” 2005 ASHP Annual Meeting, Boston, MA June 15, 2005

“Implementing Your USP 797 Plan 1: Environmental Monitoring, Infection Control, and Process Validation of Media Fills” 2005 ASHP Annual Meeting, Boston, MA, June 13, 2005

NATIONAL AND LOCAL PRESENTATIONS, continued

“Using USP 797 to Improve Patient Care and Healthcare Processes”. 2005 INS Annual Meeting, Ft. Lauderdale, FL, May 14, 2005.

“USP Chapter <797>: Reducing Sterile Compounding Risk”. 2005 NHIA Meeting,

“Meeting USP <797>: Buyer Beware: Avoid costly mistakes in complying with the requirements of USP Chapter” Baxa ASHP MCM Exhibitor’s Theater. Orlando, FL, December 7, 2004

“USP Chapter <797> Standards: An Expert Panel Offers Practical Strategies for Implementation” 39th ASHP Midyear Clinical Meeting. Orlando, FL, December 6, 2004.

“USP Chapter <797>, The New Standard of Practice for Compounding Sterile Preparations VHA Gulf States Pharmacy Directors Meeting. Cranbury, NJ, November 18, 2004

“USP <797> Pharmaceutical Compounding-Sterile Preparations: What is it and what does it mean to pharmacy practice?”. 2004 NABP/AACP District II Meeting, Chester, WV October 22, 2004

“Practical Strategies in meeting the requirements of Chapter USP <797>” The United States Pharmacopeia Chapter <797> Workshop, Bethesda, MD, August, November 2004 and May and August 2005.

“Quality Assurance and Compounded Prescriptions” The Ohio Northern University’s Raabe College of Pharmacy 14th Symposia, Ada, Ohio, July 2004.

“Your homecare practice and meeting the new USP Guidelines, The New Standard of Practice for Compounding Sterile Preparations.” Annual Education Conference, Pharmacy Society of Wisconsin, Madison, Wisconsin, April 2004.

“Understanding USP Chapter <797> Standards for Compounded Sterile Preparations.” Greater New York Hospital Association Pharmacy Directors Meeting, Flushing, NY, April 2004.

“USP Chapter <797>, Meeting the Challenges in Homecare.” Child Health Corporation of America, Homecare Directors Forum Meeting, Phoenix, AZ, April 2004.

“USP Chapter <797>, Meeting the Challenges in Hospitals.” Child Health Corporation of America, Hospital Pharmacy Directors Forum Meeting, San Antonio, TX. April 2004.

“USP <797> Pharmaceutical Compounding-Sterile Preparations: What is it and what does it mean to hospital pharmacy practice?” New Jersey Society of Health-System Pharmacists Spring Meeting, East Rutherford, NJ. April 2004.

“Sterile Preparations: A Full Day Workshop on Meeting New USP Guidelines, Regulations, and Standards of Practice for Compounding Sterile Preparations”, NHIA 13th Annual Conference, Las Vegas, Nevada. March 2004.

“Meeting the Future Challenges in Sterile Product Compounding, an overview of USP Chapter <797>”, NHIA Audio-teleconference: October 2003

“NHIA Pharmacy Compounding Conference, Regulatory Issue Focus: Sterile Product Preparation” NHIA 12th Annual Conference, San Antonio, Texas. February 2003.

“Designing Robust Sterile Compounding Conditions” NHIA Regional Conference: Quality and Compounding-Meeting the New Challenge in California, Sacramento, California. September 2002.

“Sterile Product Preparations-Mix or Buy?” Maryland Society of Health-System Pharmacist, March Meeting. Columbia, Maryland. March 2002.

NATIONAL AND LOCAL PRESENTATIONS, continued

“Impact of Accreditation: ACHC” NHIA Clinical Track Conference Program: NHIA 11th Annual Conference, Las Vegas, Nevada. February 2002.

“Pharmacy Compounding Errors: Learn the Lessons Before They Occur.” Expert Panel Presentation. International Academy of Compounding Pharmacists. IACP National Meeting, Houston, Texas, October 2001.

“Nursing and Pharmacy Collaboration in Alternate Care Setting” NHIA Clinical Track Conference Program: NHIA 9th Annual Conference, Cincinnati, Ohio. May 2000.

“1999 National Clinical Meeting, Maximizing Divisional Synergies: The Clinical Approach to Customer Service”. Phoenix, Arizona, Coram Healthcare Corporation. Program Organizer, Director, and Presenter. October 1999.

“Pharmacologic Management of Preterm Labor.” Various conferences nationwide. (1994-1998)

“Infusion Pharmacy Services: Improving Management, Operations, and Cost-Effectiveness.” NHIA Pre-Conference Program: NHIA 7th Annual Conference, Pittsburgh, Pennsylvania. May 1998