A 3-DAY INTRODUCTORY WORKSHOP IN POPULATION PK DATA ANALYSIS

A HANDS-ON COURSE USING NONMEM®

Thursday, May 14 – Saturday, May 16, 2009
Buffalo, NY

WORKSHOP SYNOPSIS

This introductory population PK training workshop has been designed to provide the necessary information to successfully implement population pharmacokinetic methodology in a drug development program. The material is structured to impart both the theoretical and practical aspects of the population approach and is versatile so that participants with diverse backgrounds and areas of expertise may benefit. Examples of the use of population PK studies in drug development programs, especially those from the published scientific literature, will be presented whenever possible to provide specific details of various implementations and better illustrate essential aspects of population PK methods. Emphasis will be placed on compliance with the FDA’s Guidance for Industry on Population PK and the EMEA’s Guideline on Reporting the Results of Population PK Analyses; participants will gain an appreciation for the importance of protocol compliance, the essentials of accurate and sufficient data collection, and learn how to proactively plan in order to maximize study effectiveness.

The workshop content is provided as a combination of formal lectures, review of data, code, and data analysis results, and hands-on exercises. Participants will be provided with a computer terminal where they will be afforded the opportunity to practice coding control streams, running various models, and evaluating the results. A thorough examination of an example dataset, from development of the structural model, through covariate analysis, and model refinement will be covered. Overall, this workshop will provide the audience with a comprehensive understanding of the population PK approach to data analysis, its usefulness and added value in drug development, as well as when and where to employ population PK methods and sparse sampling within a given development program. The format is designed to be both comprehensive and interactive.

LEARNING OBJECTIVES

Following the workshop, the participant should be able to:

1. Understand the conceptual basis and rationale for the population approach to data analysis
2. Understand where and when population methods may be optimally applied to PK and PK/PD analyses during the drug development process
3. Understand and describe the potential benefits and advantages to implementing a population strategy
4. Identify the critical logistic and practical issues involved in study design, protocol development, case report form development, overall planning, and efficient execution for population PK studies
5. Describe the critical documentation standards for population PK reports intended for submission to the FDA
6. Write, execute, and de-bug basic NONMEM® control streams for simple structural PK models
7. Outline the requirements and format for basic NONMEM® datasets
8. Understand, identify, and code some basic functional forms for covariate-parameter relationships
9. Perform covariate analysis using a forward selection followed by backward elimination approach
10. Understand the importance of exploratory data analysis (EDA) and the interpretation of standard goodness-of-fit diagnostic plots
11. Understand the basis for model selection strategies and discriminate between model candidates on the basis of both quantitative and qualitative factors
12. Have insight into potential model refinement issues

COURSE INSTRUCTION

The workshop is organized and taught by experienced pharmacometricians from Cognigen Corporation and the University at Buffalo School of Pharmacy and Pharmaceutical Sciences, a pioneer and global leader in the field of pharmacodynamics and pharmacokinetics. Cognigen Corporation has been providing clinical pharmacology consulting services, including population PK/PD modeling and simulation to the global pharmaceutical industry for over 15 years. Cognigen specializes in performing innovative data management and analyses to generate and communicate the knowledge required for time-sensitive decision-making and regulatory review. Course faculty will include: Jill Fiedler-Kelly, Alan Forrest, and David Jaworowicz.

This session precedes a 3-day separate course in the concepts and applications of Pharmacokinetic/Pharmacodynamic Modeling coordinated by Dr. William J. Jusko. For information see: http://pharmsci.buffalo.edu/symposia or contact rrurben@buffalo.edu.

Jill Fiedler-Kelly
**REGISTRATION FORM: INTRODUCTORY NONMEM® WORKSHOP**

**Name:**

**Organization:**

**Address:**

**City:**

**Postal Code:**

**Telephone:**

**Fax:**

**E-mail:**

**For credit card payment:**

**Credit card number:**

**Signature:**

**Expiration Date:**

Kindly return to: PK/PD MODELING – NONMEM Workshop, Dept. of Pharmaceutics, School of Pharmacy, University at Buffalo, 519 Hochstetter Hall, Buffalo, NY 14260; phone: 716 645 2842, x. 540; fax: 716 645 3693; e-mail Rita Urben at rrurben@buffalo.edu.