

中国医药产品国际化专题论坛

---美国药品法规政策及仿制药和新药注册审批过程

U.S. FDA Regulatory Policies & How Does FDA Approve Generic and Innovative Drugs



中国 郑州

Zhengzhou, China
September 20-21, 2009

Conference Information and Pharmaceutical Job Opportunities in China

SAPA-GP and **Frontage Laboratories, Inc.** are co-sponsoring a workshop in Zhengzhou China on September 20 - 21, 2009. The goal of the workshop is to provide educational information about latest U.S. FDA regulatory policies for generic and new drugs approval in order to help Chinese Pharmaceutical companies launch their products in the international market. Workshop will be given by current and previous FDA officials and experienced cross-border industrial practitioners and specialists.



The invited speakers include:

Dr. Ethan Stier, Team Leader, Division of Bioequivalence II, Office of Generic Drugs, Center for Drug Evaluation and Research (CDER), FDA

Dr. Jonca Bull, Former Office Director (Office V), Office of New Drugs, Center for Drug Evaluation and Research (CDER), FDA. Vice President, Regulatory Affairs, Novartis Pharmaceuticals Corporation

Dr. Brenda Uratnai, Beijing Office, FDA

Dr. Derek Zhang, VP Regulatory Affairs and Clinical Pharmacology, Frontage Laboratories, Inc. Former Senior Clinical Pharmacology Reviewer, CDER, FDA

Mr. Ronald Connolly, SVP Operations, Frontage Laboratories, Inc.

Mr. Robert Edwards, CEO, BOCA Pharmaceuticals

They will interact with a wide range of participants such as company executives, policy-makers, and professionals from across the industry, the government, and the academia.

During the workshop, Chinese pharmaceutical companies with international ambition will recruit US-trained professionals with pharmaceutical industry experience who have the intention to work in China. Candidates with experience in Regulatory Affairs, QA, QC, DMPK, CMC, Preclinical or Clinical R&D are highly recommended to meet with the employers at the workshop, or to submit their resumes to Ms. Wang at hwang@frontagelab.com not later than August 15th, 2009.

美中药协大费城分会与美国方达医药技术公司将于 2009 年 9 月 20-21 在中国郑州共同举办中国医药产品国际化专题论坛。论坛旨在为中国医药工业提供美国 FDA 药品法规政策及仿制药和新药注册审批过程的相关信息，帮助中国医药产品打入国际市场。论坛特邀多位美国 FDA 医药产品审批官员及药物研发、申报和市场流通专家执讲。执讲人将与来自各界的参会者，如制药公司高层领导、政府官员和决策者、高校及其它专业人士进行充分互动。

会议期间，致力于国际化拓展的各大中国制药公司将对有意到中国工作并具备美国教育及药业工作背景的业界精英展开招聘活动，尤其欢迎有 **Regulatory Affairs, QA, QC, DMPK, CMC, Preclinical or Clinical R&D** 经验者。有意者可报名参会或在 2009 年 8 月 15 日前，将电子版简历提前发送给王女士，hwang@frontagelab.com。



Sponsors:



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