Duxes Pharma Series



China Pharma Manufacturing & Supply Chain Management Summit 2010

25-27 August, 2010 Swissôtel Beijing Hong Kong Macau Center, Beijing, China



- ◆ Interpretation of the latest requirements & standards of pharma manufacturing
- Optimizing manufacturing planning to drive operation excellence
- Clarifying the significant contribution of engineering on pharma manufacturing
- Identifying how to develop an efficient and cost-effective supply chain management system
- Exploring the innovative solutions to optimize quality assurance systems
- Learning how to build the relationship between waste management and sustainable development
- Digging the real demand of Chinese pharma manufacturing market

Media Partners

























Background

Growing at a CAGR of around 6%, the global pharmaceutical market is forecasted to reach US\$ 1043.4 Billion by 2012. However, it is expected that the growth will slow down in near future due to patent expiration of key drugs and increased prevalence of generics. It is therefore more important than ever that pharma manufacturers should optimize their manufacturing line for cost reduction while following latest standards, like CGMP, QbD and PAT concepts etc.

A large untapped population and strong economic growth in major countries is expected to make Asia-Pacific the most lucrative pharmaceutical market in future. Growing with 20% per year China is expected to be the third largest pharma market by 2013. However, the lack of manufacturing technologies and streamlined and consistent global supply network still are key hindrances for many local companies' development. To help national pharma manufacturers enter the global market, currently Chinese government is drafting new GMP standards which is expected to be released in this first half year. Making a thorough understanding of this vital policy and the pharma market itself is critical for doing successful business in China.

The summit will provide you a unique platform for first hand information of newly released policies by face to face contact with competent authorities. World-leading pharma manufacturers will share with you their successful experience and lessons through keynote speeches and real-life case study. In addition, the summit will be a valuable opportunity for your networking with business partners and potential clients.

Who Should Attend?

By industry:

- · Pharmaceutical companies
- Raw material providers: Ingredients, API, PI etc.
- Equipment providers
- · Engineering companies
- Service Providers
- Consulting Firm
- Law firm

By job title:

- President/Vice President
- CEO/Managing Director/Partner
- General Manager/Deputy General Manager
- Plant Manager
- Operation Manager/Director
- Manufacturing Manager/Director

- QA & QC Manager
- Technical Director
- BD Manager
- Marketing Manager/Director
- Sales Manager/Director
- Regional Manager/Director

PRE-EVENT CONFERENCE

Wednesday, 25th August, 2010

Pharma Packaging Special

- 14:00 Deep Understanding of Latest
 Development of Regulations &
 Criteria on the Pharmaceutical
 Packaging
 - ♦ Implementation of the strict requirements on material and processing quality
 - ♦ Regulations & polices on pharmaceutical packaging

Director, NICPBP SFDA

- 14:45 Implementing Sustainable
 Development into Packaging
 Processes
 - ♦Strategies for improving and optimizing your ecological packaging production
 - ♦ Systematic process optimization for effective packaging management

Packaging Plants Project Manager Bayer Schering Pharma

- 15:30 Tea Break
- 16:00 Insight into Great Contribution of Pharma Packaging to Pharma Manufacturing
 - ◆Development trend of the pharma packaging technologies
 - ◆Deep analysis of the current situation of pharma packaging market

Director
Pharma Packaging
Schott

16:45 Customized Packaging Solutions by Uhlmann

Werner Florczak Head of Customized Packaging Systems Uhlmann Pac-Systeme GmbH & Co. KG, Germany

17:30 Close Pre-event Conference

DAY ONE

Thursday, 26th August, 2010

08:30 Sign In

Policies & Regulations

08:45 Implementation of new GMP

- ♦ Comprehensive understanding of the details of new GMP
- ♦ Comparison of new GMP and previous one

SFDA

Outlook & Development

09:25 Swift Reaction to Latest
Development of Requirements &
Standards of Pharma Manufacturing

- ♦Insight into Latest Development of cGMP & ISPE guidelines
- ♦Comparison of the new GMP and cGMP

Director FDA & ISPE

10:05 Tea Break

Manufacturing Strategy

Manufacturing Planning

10:20 Optimized Manufacturing Planning to Avoid Risk

- ◆Preparation for additional growth avoiding undue risk
- ♦How to balance excess capacity and insufficient capacity situations

Stephen Reich Risk Management Principal Global Quality, Pfizer

Efficiency & Cost Reduction

11:00 Cost Reduction Strategies for Efficient manufacturing & Operations

- ♦ Make efficient cost reduction strategies
- ♦Find new solutions to implement cost reduction strategies

Yin Xudong President AstraZeneca China

Panel Discussion

11:40 Insight into the Crucial for Cost Optimization

- ♦Identifying the solutions from pharma manufactures ,equipment & service providers to implement cost reduction strategies
- Understanding cost reduction strategies to help you thrive in the industry

12:20 Luncheon

Engineering

14:00 Facility Construction - Solid Basis for Pharma Manufacturing

- Improving the quality of the engineering services
- ♦ Upgrading engineering qualification & certification services

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Supply Chain Management

Pharma Supply Chain

14:40 Optimizing Supply Chain Management

- ♦Improving real-time visibility across the enterprise to ensure product quality, improve profitability and response time
- ♦Eliminating data silos in order to identify unrealized business benefits and minimize risk throughout the production and distribution cycle

Director Supply Chain Project Management Novartis

15:20 Advancing Information System for Effectiveness & Efficiency of Pharma Supply Chain Management

- ♦Implementing information system to ensure supply chain management efficiently
- Understanding the role played by information system in supply chain management

16:00 Tea Break

Raw Materials & Ingredients

16:15 Upgraded Sourcing Criteria to Optimize Supply Chain Management

- New souring criteria from leading pharma manufacturers
- ♦The importance of new criteria to supply chain management

Dittmar Nerger Head of Strategic Sourcing Bayer healthcare

Panel Discussion

- 16:55 How to Implement the Track & Trace
 System of Drug Security Effectively in
 Supply Chain Management
 - ♦Identification the key success factors in a track & trace system of drug security
 - ◆Cooperation and interoperable collaboration among pharma manufacturers government officials, regulators and technology providers

Logistics

17:35 Advanced Logistics Solutions to Ensure in-time Supply Chain Operation

- ♦The innovation solutions of the logistics makes supply chain efficiently
- ♦Enhancing the role played by logistics system in supply chain

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18:15 Close Day One



Friday, 27th August, 2010

Technology & Industrialization 08:45 Identifying Mature Technologies for

Industrialized Manufacturing ◆The current development situation of pharmaceuticals manufacturing

♦ The bottlenecks of industrialization of pharmaceuticals manufacturing

Violetta Georgescu-Kyburz CEO Kenta

Quality Assurance

Quality Management

09:25 Optimizing the Quality Management System to Enhance Pharma Manufacturing Processes

- ♦QMS plays a key role in the pharma manufacturing
- Evaluating and upgrading the quality management system

10:05 Tea Break

System & Processing

10:20 Advanced Processing System - Assurance of Quality Control

- Implementing advanced processing system
- ♦Advancing processing system to ensure effective quality control

Dr. Harald Stahl Senior Pharmaceutical Technologist GEA Pharma Systems

11:00 Automation Enhancing the Performance of Pharmaceutical Manufacturing

- ♦The development of the pharma manufacturing automation
- ♦ Automation will play an increasing role in pharma manufacturing

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11:40 Information System to Ensure Efficient Quality Management

- ♦Information System plays an increasing role in manufacturing
- ♦How to implement the information system efficiently

12:20 Luncheon

Equipment Update

14:00 Insight into the Safe and Efficient Preparation Equipments and Quality Assurance

- ♦The development trend of the preparation equipment
- ♦ More efficient & safer equipments are needed

14:40 Optimized Disinfection Solutions for Qualified Pharma Manufacturing

- ♦ Tailor-made equipments to different pharmaceutical manufacturers
- ♦Innovation of the disinfection solutions to ensure pharma manufacturing safer Open For Sponsor



15:20 Batch and Continuous Reparative Chromatography for Large Scale Production of API's and Biological

♦ Application to antibiotics, Chiral and non chiral API's, Monoclonal Antibodies and Proteins

Yvan Ruland Technology Manager NOVASEP Asia

16:00 Tea Break

Testing & Certification

16:15 Quality Assurance with the Help of Testing Instruments

- ◆Testing instruments ensure quality control operation efficiently
- ◆Updated manufacturing standards need new testing solutions

16:55 Strengthening Comprehensive Validation & Certification - Essential Ways to Ensure Quality

- ♦Understanding the importance of successful cleaning validation
- ♦Upgrading certification to ensure production timely

Waste Management & Sustainability

17:35 Efficient Waste Management System to Reach Sustainable Production Goal

- ◆The importance of waste management system to achieve sustainable manufacturing
- ♦Inspecting the solution to optimize waste management system

Robert Colucci Senior Director Energy & Sustainability Merck & Co., Inc

18:15 Water Treatment Solutions for Green Pharma Manufacturing

- Increasing the efficiency of water use in manufacturing
- ♦ Implementing water cycle management solutions

Michel Buser Managing Director ecoSign

19:00 Close the Conference

Sponsorship and Exhibition Stand Opportunities

Enhance your profile by taking one or more of these marketing options

Taking a sponsorship option at China Pharma Manufacturing & Supply Chain Management Summit 2010 gives you a portfolio of opportunities to reinforce the strength of your brand and enhance awareness about your company and products in front of a precisely targeted audience of decision makers on the event site as well as thousands of top executives through our powerful marketing coverage and sales reach.

What options are still available?

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- Exhibitions
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What benefits will sponsorship bring you?

- · Increased brand exposure
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- 200 words business profile and a hyperlink to your website
- Logo placement on the Summit's backdrop and on all event-related materials

To receive details about sponsorship, please contact:

Peter Xu

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Fax: +86 21 5258 8011 E-mail: pxu@duxes.cn

REGISTRATION FORM

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Please fax back the regis	tration form after complet	ting at +86 21 5258 8011				
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Should you be unable to attend, a substitute delegate is welcome at no extra charge. A charge of 50% of the registration fee, plus 10% administrative charge will be billed for cancellations received in writing at least four weeks prior to the event. Alternatively, you may choose a credit voucher for the full value of the registration price, which may be deemed for future Duxes events. Duxes regrets that no cancellations will be accepted within four weeks prior to the event start date and payments will not be refunded and invoiced sums will be payable in full.

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