Technology Transfer

9 & 10 May, 7 & 8 November 2011
Window Conference Venue, London UK

This 2-day interactive workshop is intended to provide an overview of the challenges that must be overcome in transferring technology and to highlight some proven techniques for overcoming those challenges.

Some of the benefits to be derived from this workshop include:

- An appreciation of where TT fits in the life cycle of a pharmaceutical product.
- An appreciation of the importance of planning and project management in TT.
- An overview of the potential complexity of technology and product transfer.
- An outline of proven best practice in technology transfer.
- An opportunity to discuss current issues and challenges in technology transfer with peers and with an expert faculty.

Course Leader:

Chris Barnett is an independent GMP and compliance consultancy. Chris has a talent for coaching and explaining complex regulations in a straight-forward manner.

An expert in quality management, technology transfer and new product introduction, Chris specialises in Quality Management, Technology Transfer and New Product Introduction.

Chris has a background as an analytical chemist and over 20 years experience in QA roles, with the Wellcome Foundation and GlaxoSmithKline. He has had QA management posts in India and Mexico, as well as new product introduction project management roles in pharma manufacturing. Chris graduated from Cambridge in natural sciences, with an MSc in analytical chemistry, and subsequently an MBA from Greenwich University.

Comments from previous attendees:

"Very good, informative, gave a good overview of all aspects of TT"

"Good with experienced lecturer, lots of questions could be answered"

"Good overview of what a transfer should be"

www.pharma-training-courses.com

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Who should attend?
Pharmaceutical technologists, Quality Assurance staff, Regulatory staff, Engineering staff, Project Managers and anybody else involved in the transfer of products or technology from one location to another.

PROGRAMME

Start time:
Coffee and registration will be available from 8.30am on Monday 22 November, course proper will commence at 9.00am. We anticipate the course will finish at 5.00pm on Tuesday 23 November 2010.

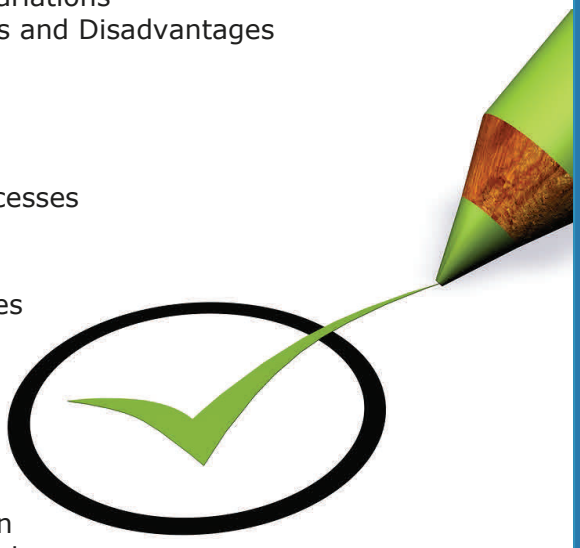
Day 1
Why Transfer Technology?

- Pharmaceutical Product Life Cycle
  - Development
  - Innovative
  - Mature
  - Generic
- Product Transfers and Technology Transfers
  - IP
  - Technology
  - Techniques
  - Regulatory Considerations
- Registration Procedures
  - Europe
  - Non-European
- Variation Procedures
  - Types of Variations
  - Advantages and Disadvantages

Evening reception

Day 2
Technical Challenges

- Manufacturing processes
- Validation
- Stability
- Packaging processes



Logistical Challenges

- Label Changes
  - Text
  - Physical
- Timing
  - Project Plan
  - Phase in/out

Human challenges

- Push versus Pull
- Team membership and team roles
- Cross-Cultural Issues
- Progress Reporting and breaking bad news

Venue

Window Conference Venue 13 Windsor Street, London, N1 8QG convenient for central London, in a pleasant informal setting.

The course does not include accommodation, hotel details are available from our website www.pharma-training-courses.com

Numbers are limited to give participants the opportunity for thorough discussion of the issues to be covered by the programmes and one-to-one consultation with speaker(s)

COURSE PROGRAMME 2011

Planning for Commercial Launch: *29 & 30 March 2011*
GMP Auditor Training*: New Jersey USA - 11 & 12 April 2011*
London - 9 & 10 May, 7 & 8 November 2011
How to Audit API Manufacturers: *New Jersey USA - 13 April 2011*
London - 11 May, 9 November 2011
Supply Chain Management in Pharma/Biotech: *5 & 6 May 2011*
Technology Transfer: *London - 9 & 10 May, 7 & 8 November 2011*
Integrated Tablet Formulation Development: *New Jersey USA - 7 & 8 April*
London - 9 & 10 June, 24 & 25 November 2011
Tablet Process Development, Validation and the application of QbD:
New Jersey USA - 11 & 12 April, London - 13 & 14 June, 28 & 29 November
Pharmacokinetics in Drug Development - an Integrated Approach:
9 & 10 June 2011
An Introduction to LC/MS: *19 September 2011*
Quantitative Analysis: *20 & 21 September 2011*
Writing effective SOPs in a GMP Environment: *13 & 14 October 2011*
OOS investigations in a GMP Environment: *18 & 19 October 2011*
Stability Testing in Pharmaceutical Development: *16 & 17 May 2011,*
12 & 13 December 2011
Introduction to Photostability: *14 December 2011*
HPLC Analytical Method Development and Validation: *22 & 23 November*
Oral Solid Dosage Manufacturing Technology: *28 November 2011*
Development and Manufacture of Effervescent Tablets: *30 November 2011*
Pharmaceutical Packaging - an Introductory Course: *14 December 2011*

Check out the benefits, content, details, dates and times of our range of training programmes:

We deliver a range of expert programmes in pharmaceutical development, quality assurance and regulatory topics, plus a new range of industry awareness courses. We employ speakers/trainers with a high degree of expertise, completely up to date with industry trends.

We run our programmes in a variety of locations, throughout the year.

For 5 or more staff requiring training it may be beneficial to run a course in-house.

The benefits of running a course in-house:

- Up to 70% savings on delegate fees
- Save on travel or accommodation costs
- Customised content to meet your requirements
- Big print savings on course material - especially with larger groups
- Courses arranged for large groups up to 24 staff
- Tutorials available for small groups of 2 or 3 staff
- Meet course speakers in advance to discuss design and content

Contact **Judy Callanan** Ph: 0044 20 7193 7703, Fax: 0044 20 7681 3582
Email: judy@pharma-training-courses.com

REGISTRATION FEES

Technology Transfer:

9 & 10 May 2011, London ☐

7 & 10 November 2011, London ☐

2 day course £1180.00 + VAT £236.00 **Total £1416.00** ☐

Discounted rate for registering and paying 8 weeks prior to commencement of course £1062.00 + VAT £232.40 **Total £1294.40** ☐

Send completed registration form to:

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Nottingham NG1 1GF
United Kingdom

Fax: 0044 (0)20 7681 3582

Email: info@pharma-training-courses.com

Title (Mr/Mrs/Ms/Dr/Prof): First Name:

Surname:

Company:

Job Title:

Address:

Post Code: Country:

Tel: Fax:

Email address:

Signature:

Method of Payment:

☐ Cheque (**Please make payable to "Pharmaceutical Assurance Services")**

☐ Bank transfer

☐ Credit/Debit Card (If paying by Credit Card please register online)

Online Registration is available on our website:
www.pharma-training-courses.com

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Terms and Conditions

Delegate fees: Fees for this programme or suite of programmes are shown overleaf. Delegate fees are inclusive of course documentation, refreshments and lunch. Payment of the registration fee must be paid within 14 days of commencement of the course. Upon receipt of payment, a proof of payment will be sent to you.

Cancellation Policy: Full refunds less a handling fee of £100 will be made for cancellations received in writing within 28 days of the commencement of the course. Refunds of 50% will be made for cancellations received in writing between 28 and 7 days prior to the commencement of the course. Regrettably no refunds will be made after 7 days prior to commencement of the course. Substitutions can be made at any time.

Liability: PharmaTraining Ltd reserves the right to change the programme, speakers, date or venue without notice or cancel the event. If cancellation occurs delegates will be notified as soon as possible and will receive full refund of fees paid. PharmaTraining Ltd will not be responsible for any airfare, accommodation or other travel costs incurred.

Data Protection

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