

Offer USA-37:

for training courses in industrial pharmacy for manufacturers and inspectors in developing countries.

Michael H. Anisfeld from Globepharm Consulting in Chicago, USA, has offered to Pharmabridge to teach the following industrial pharmacy courses in developing countries:

- Preparing for, and Passing an FDA/EU Inspection (1 day)
- FDA cGMPs for Finished Dosage Form Manufacturers (1 day)
- FDA cGMPs for Manufacturers of Sterile and/or Biotechnology Products (1 day)
- FDA cGMPs for Manufacturers of Active Pharmaceutical Ingredients (APIs) (1 day)
- EU GMPs for Finished Dosage Form Manufacturers (1 day)
- USA and EU GMPs for Sterile Product Manufacture (1 day)
- International GMPs for Cosmetic and Complementary Medicines Manufacturers (1 day)
- International GMPs for Engineers, Architects and Facility Designers (1 day)
- International GMPs for Pharmaceutical Development Scientists (1 day)
  
- Managing the QC Lab in a GMP Compliant Manner (1 day)
- Fundamentals and Essentials of Validation (2 days)
- Good Documentation Practices (1 day)
- Good Distribution Practices – Securing the Supply Chain (1 day)
- Effective Investigations and Corrective Actions (1 day or 1½ days with case studies)
- Effective Quality Assurance Auditing (1 day or 1½ days with case studies)
- Fundamentals and Essentials of Pharmaceutical Water Systems (1 day)
- GMP Week: A week of intensive GMP training for every department in the company (5 days)
  
- Industrial Pharmacy – a post graduate course into the manufacture and quality testing of pharmaceutical (20 days) – *this course is particularly suited to Universities/Colleges*
  
- How To Perform An Effective GMP Inspection (2 days)  
– *course only available to members of government inspection agencies*

These courses are designed for 25 – 50 people in attendance; more than 50 is possible, but spoils the intense interactive nature of the courses.

Detailed course descriptions of all these courses can be found at Mr Anisfeld's website at <http://www.globepharm.org/seminars.html>

Michael Anisfeld is prepared to tailor courses to local needs with a duration of 5 to 20 days.

The best local organizers would be the local pharmaceutical industry associations, and/or/with Ministries of Health, because these courses would also be ideal to train government GMP inspectors.

Michael Anisfeld does not expect to be paid for his time, but expects the local organizers to cover his travel (prepaid ticket) and living expenses at a business class hotel, ex-Chicago, USA.

Obligations on receiving party:

**Organization**

Local organization of everything relating to the course with a minimum of 25 participants per course

Whoever takes over the organization, a combination of participants from industry and regulatory agencies is preferred.

**Financial obligation**

All organizational expenses and return air ticket (full fare economy) out of Chicago and local accommodation and living expenses (food, local travel, etc.) for Mr Anisfeld to be covered either by the organizer (local manufacturers association or Ministry of Health) or through course registration fees.

Anybody interested in the offer should initially contact

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Pharmabridge coordinator  
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GMP training carried out under Pharmabridge so far:

Sri Lanka:

2005: 14-16 June: 1x2 +2x1 days with 54,43 and 43 participants respectively

2012: 12-16 November: 1x2 + 3x1 days, 141 people from both government and private sectors

Nigeria:

2005:

12-13 December: How to perform GMP Inspection (20 inspectors: 10 from NAFDAC, 10 from Pharmacy Council of Nigeria)

14-15 December: Current Trends and Expectations in GMP with 33 participants from industry

2006:

4-5 December: Validation, Water systems attended by 13 participants from Pharmacists Council of Nigeria

5-7 December: Validation, Water systems attended by 34 participants from industry (from 16 companies)

2007:

11-12 June (day 1: Good documentation. practices (41 participants); day 2:Managing a QC Lab in a GMP Way (38 participants). Most participants attended both days

Mongolia:

2007: 5 days (including visits to manufacturers) with 50 to 65 participants from Ministry of Health

Ghana:

2007, 07-11 May: 5 days on 5 GMP topics; 60 participants from regulatory body and industry

7<sup>th</sup> May, 2007            International (EU & USA) GMPs For Cosmetics And Drug Manufacturers

8 <sup>th</sup> May, 2007	Managing QC Laboratories In a GMP Compliant Manner
9 <sup>th</sup> May, 2007	Good Documentation Practices
10 <sup>th</sup> May, 2007	Fundamentals and Essentials of Validation
11 <sup>th</sup> May, 2007	Fundamentals and Essentials of Pharmaceutical Water Systems

Ghana

2008, 19-23 May: 5 days on 5 GMP topics; 61 participants from regulatory body and industry

19 <sup>th</sup> May 2008	Passing an FDA Inspection: A - Z
20 <sup>th</sup> May 2008	Design and Engineering of GMP Compliant Pharmaceutical Manufacturing Facilities
21 <sup>st</sup> May 2008	Performing Effective GMP Inspections: Strategies and Tactics
22 <sup>nd</sup> May 2008	GMPs For Senior Directors and Executive Management
22-23 May 2008	Effective Investigations and Corrective Actions (CAPA) Proper Investigation Of Quality Events

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