

INDUSTRIAL TRAINING AWARDING CERTIFICATE OF MANUFACTURING ASSISTANT

BACKGROUND

Pharmaceutical industry is a well-regulated industry. It is a major source of pharmaceutical products supplying various types of medicines to the public. This industry is required to be operated by following stringent Good Manufacturing Practice in order for the medicines to be manufactured, registered and commercialised. Having a well-trained manufacturing assistant is an added advantage to this industry complimenting its aggressive nature.

OBJECTIVES

1. To expose the trainee to the fundamental aspects of Good Manufacturing Practice (GMP) of pharmaceutical industry.
2. To train the trainee on manufacturing and In-Process Quality Control (IPQC) of pharmaceutical dosage forms abiding Good Manufacturing Practice (GMP).

ENTRY REQUIREMENT

DIPLOMA level of any science-based and not limited to mechanical, chemical and pharmaceutical engineering.

MAXIMUM NO. OF PARTICIPANTS PER INTAKE & DURATION OF TRAINING

Seven (7) participants per intake. Intake is on middle of April, June, August, October, December. Duration per intake is 40 days.

CONTACT INFORMATION

Assoc. Prof. Dr. Farahidah Mohamed, Acting CEO, IKOP Sdn. Bhd. (Pilot Plant Manufacturing of Pharmaceutical, Kulliyah of Pharmacy, International Islamic University, 25200 Kuantan).
farah@ikop.com.my

DETAIL COST OF ACTIVITIES

As depicted in the next page.

A GMP Manufacturer of Shari'ah-Compliant Pharmaceuticals

IKOP Sdn Bhd

Kulliyah of Pharmacy, International Islamic University Malaysia,
Jalan Sultan Ahmad Shah, 25200 Kuantan, Pahang DarulMakmur.

Tel. +60 9 571 6684 Fax. +60 9 571 6541, Website: www.ikop.com.my E-mail info@ikop.com.my

Activities	Materials	Equipments	Trainer's Fees	Student's allowance
1. 5 hr lectures on :	Training materials=		5 hrs lectures + assessment time =	RM10/day = 10 x 40days = RM400
a. Personnel and material flow in a pharmaceutical industry	1. 5 hrs Lectures notes = RM250		=	
b. Introduction to GMP	2. Assessment post-lectures = RM250		RM 150 x 5 hrs = 750	
c. Fundamental aspects of Standard Operating Procedure for manufacturing & IPQC test	3. Hands-on training modules = RM300			
d. Basic information on SOPs of manufacturing equipment				
e. Basic information on IPQC equipments and testing				
2. Dispensing of raw materials	Raw materials =		Hands-on instructions and briefings =	
3. Manufacturing & IPQC test of solid dosage forms (tablet/capsule)	1. For manufacturing of solid dosage forms = RM500	1. For solid line equipment = RM800 (4 weeks)	1. For solid line = RM200 (4 weeks)	
4. Manufacturing & IPQC test of semi-solid or liquid for external use.	2. For manufacturing of semisolid = RM400	2. For Semisolid equipment= RM400 (2 weeks)	2. For semisolid line = RM100 (2 weeks)	
Sub-total fees per pax	RM1,700	RM1,200	RM 1,050	RM400
GRAND TOTAL FEES/pax	RM 4,350			

A GMP Manufacturer of Shari'ah-Compliant Pharmaceuticals

IKOP Sdn Bhd

Kulliyah of Pharmacy, International Islamic University Malaysia,
Jalan Sultan Ahmad Shah, 25200 Kuantan, Pahang DarulMakmur.

Tel. +60 9 571 6684 Fax. +60 9 571 6541, Website: www.ikop.com.my E-mail info@ikop.com.my