

Introduction

This workshop aims to enhance participants' understanding of Good Laboratory Practices (GLP) and the registration procedures for generic drugs, traditional medicines, health cosmetic supplements, and products. Featuring expert speakers from the National Pharmaceutical Regulatory Agency (NPRA), the programme equips industry professionals and academics with comprehensive knowledge and practical guidance on NPRA's registration and notification methods. Participants will gain valuable insights directly from regulatory authorities, enabling them to navigate compliance processes effectively and elevate the standards of their research and product development initiatives.











Objectives

- To provide exposure on products registration and notification in the NPRA.
- To encourage the exchange of ideas among researchers and industry players in the field of product development and registration.

Who should attend?

- Regulatory Affairs Executive
- Product Development Executive
- Academician
- Cosmetic Business Operator
- Anyone Who Wanted to Update Their Knowledge On MOH Legal Requirements or Guidelines and Licensing

Content of the workshop

- Quest 3+ System
- Generic Drug and Health Supplement Registration
- Traditional Medicine with Therapuetic Guidelines
- Cosmetic Notification
- Good Laboratory Practice (GLP)
- Regulatory Perspective for Clinical Trial

Registration fee

- RM1,000.00/person
- MNPS members and those who register before October 18th will get 5% discount

QUEST 3+ SYSTEMS, **REGISTRATION OF PHARMACEUTICAL** PRODUCTS, AND **COSMETIC NOTIFICATION** & MEDICINES **ADVERTISEMENT WORKSHOP**

5 - 6 November 2024 **Dorsett Hotel, Putrajaya**

Programme Tentatives

Day 1:5 November 2024

0830 Registration

0900 Opening remark

0905 Keynote 1: Advancing herbal medicine:
 enhancing product quality and safety
 through robust quality control practices

1005 Photo session and morning break

1030 Session 1: Quest 3+ system (NPRA rep.)

1130 Session 2: Medicines advertisement (MOH

rep.)

1230 Lunch break

1400 Session 3: Regulatory perspective of clinical

trial in Malaysia (IMR rep.)

1500 Session 4: Generic product registration

(NPRA rep.)

1600 Session 5: Good laboratory practices (GLP)

overview and preclinical testing (UKM rep.)

1700 Tea time and end of session



0830 Registration

0900 Keynote 2: NMR-based metabolomics approach for quality evaluation of medicinal plants

1000 Morning break

1030 Session 6: Cosmetic product notification (NPRA rep.)

1130 Session 7: Sharing session with industry (Duopharma rep.)

1230 Lunch break

1400 Session 8: Traditional medicine with therapeutic guidelines and registration (NPRA rep.)

1500 Session 9: Health supplement product

registration (NPRA rep.)

1600 Closing ceremony

1700 Tea time and end of session

Speakers

Meet the

Keynote 1

Assoc. Prof. Dr.

Mazlina Mohd Said

Universiti Kebangsaan

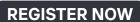
Malaysia



Prof. Dr. Faridah
Abas
Malaysian Natural
Products Society



Mr. Leonard Ariff
Abdul Shatar
Executive Director,
Duopharma Biotech
Berhad





Register before 18 October 2024

For more information or any inquiries reach us at

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