Introduction

This workshop aims to enhance participants' understanding of Good Laboratory Practices (GLP) and the registration procedures for generic drugs, traditional medicines, health supplements, and cosmetic 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

Day 2 : 6 November 2024

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

REGISTER NOW

Register before **18 October 2024**

For more information or any inquiries reach us at

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Meet the Speakers

Keynote 1

Assoc. Prof. Dr. Mazlina Mohd Said
Universiti Kebangsaan Malaysia

Keynote 2

Prof. Dr. Faridah Abas
Malaysian Natural Products Society

Industry Talk

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