

WEBINAR: PRINCIPLES OF CLEANING VALIDATION & ITS REGULATORY REQUIREMENTS

20 JANUARY 2022 (THU)
9.30 AM - 12.00 PM



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MAIN CONTENT

- To understand the requirement of Cleaning Validation in GMP
- To understand the best practices in Cleaning Validation
- To understand how the Risk Assessment should be carried out for Cleaning Programme
- Case study on Cleaning Validation Programme
- To understand the expectation from GMP Inspector during a routine audit
- Sharing of the common observations by pharmaceutical manufacturers

RECOMMENDED FOR

QA, QC, Production, Technical Services, Engineering, Project Management, Regulatory Affairs

VENUE

Zoom Platform

CONTACT US:

ispe.malaysia@gmail.com

REGISTRATION FEES

ISPE Members	: RM 250
Non ISPE Members	: RM 500
Foreign Participants	: USD 200

PAYMENT DETAILS

Bank transfer/deposit:

Account Name : **ISPE Malaysia**
Bank : United Overseas Bank (M)
Berhad
Account Number : **1623023449**
(Swift Code : UOVBMKYL)

Cheque:

Payable to : **ISPE Malaysia**
Address : Suite 8.01, Level 8
Menara Binjai, No. 2, Jalan Binjai
50450 Kuala Lumpur

CLOSING DATE

18 JANUARY 2022

**E-Certificate of Attendance will be provided*

ISPE MALAYSIA PRESENTS

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INTRODUCTION

An introductory yet comprehensive overview of cleaning validation principles and requirements that are crucial for ensuring the effectiveness of cleaning as well as compliance with GMP requirements.

AGENDA

- 09.30 am – 10.30 am **Risk Based Cleaning Validation, Maximum Safe Carry Over & Process Automation**
Vivien Santillan, Regional Director, Asia Novatek International
Dr. David Vincent, CEO VTI Life Sciences
- 10.30 am – 11.00 am **Case Study for Cleaning Validation Program**
Vivien Santillan, Regional Director, Asia Novatek International
Dr. David Vincent, CEO VTI Life Sciences
- 11.00 am – 12.00 pm **Regulator's Expectation on Cleaning Validation Programme**
Mr. Mohd Nasrul Mohamad Noor, Head of Good Manufacturing Practice (GMP) Section, NPRA

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