The voice of microbiology for 30 years: 1991 – 2021

Cleaning & Disinfection:
A roadmap to compliance and how to make sense of it all

2 x ½ virtual meetings (pm) Via Zoom - 15th & 16th September 2021

Attend and receive the latest updated:
Guide to Disinfectants & their use in the Pharmaceutical Industry 2021

KEY RE: TIMES BLUE – BST (British Summer Time) / **GREEN – EST (Eastern Standard Time)**

WEDNESDAY 15th SEPTEMBER 2021

13.00 – 13.10
08.00 – 08.10 **Welcome**
Rachel Kirkham – Director, Global Technical Consulting Contamination Control & Validation, Ecolab

13.10 – 13.55
08.10 – 08.55 **Upcoming regulatory changes and their impact for cleaning & disinfection: EudraLex draft Annex 1 v.12 & USP draft chapter <1072>**

- Brief summary of the history of these publications and their revisions to date
- Key changes in the proposed draft regulations relating to cleaning and disinfection, versus current industry practice

Laura Guardi – Associate QA Director, AstraZeneca

13.55 – 14.00
08.55 – 09.00 **Q&A session & speaker change-over**

14.00 – 14.45
09.00 – 09.45 **Update on current methods**
This presentation will cover key considerations when designing an efficacy study, including the selection of organisms and surface materials, alignment to in-house cleaning practices and pros and cons of the standard test methods available.
Kim Morwood – MD, MGS Laboratories

14.45 – 15.00
09.45 – 10.00 **Q&A session, virtual break, & speaker change-over**

15.00 – 15.45
10.00 – 10.45 **Disinfectant Workshop: Factors to consider when designing a disinfectant efficacy test and interpretation of results**
Delegates will be provided with a set of data and asked to develop a rationale for a disinfectant regime and interpretation of results
Laura Guardi – Associate QA Director, AstraZeneca & Rachel Kirkham – Director, Global Technical Consulting Contamination Control & Validation, Ecolab

15.45 – 16.00
10.45 – 11.00 **Q&A session, round up and close of virtual day 1**

THURSDAY 16th SEPTEMBER 2021

13.00 – 13.10
08.00 – 08.10 **Welcome**
Rachel Kirkham – Director, Global Technical Consulting Contamination Control & Validation, Ecolab

13.10 – 13.55
08.10 – 08.55 **Final phase of disinfectant efficacy – the field trials**
After completion of the laboratory studies, the final phase of introducing a new disinfectant is the field trial. This remains the only part of disinfectant qualification for which no standard has been written. This session will cover:

- The aims of the field trial
- Designing the study
- Selecting cleanrooms
- Setting acceptance criteria
- Interpreting microbial counts and statistical review
- Assessing recovered microorganisms
- When to repeat the trial
- This session includes case study data

Dr Tim Sandle – Head of Microbiology, Risk Management, & Sterility Assurance, BPL

13.55 – 14.00
08.55 – 09.00 **Q&A session & speaker change-over**

14.00 – 14.45
09.00 – 09.45 **Where can disinfectants go wrong and the consequences: life lessons!**
Delegates will be taken through 3 real-life scenarios from presenters who had to deal with them, followed by an open discussion session where you too can share your experiences: Lessons learnt
Lesson 1 – Disinfectant efficacy validation studies
Lesson 2 – Cleaning & disinfection: What happens when no-one is looking?
Lesson 3 – You think your operators are following the SOPs? Think again!
Lesson 4 - See no evil, hear no evil, speak no evil – when everyone collectively fails to see the obvious"

14.45 – 15.00
09.45 – 10.00 **Q&A session, virtual break, & speaker change-over**

15.00 – 15.45
10.00 – 10.45 **Implementing a new disinfectant regime: commissioning to first use**
Introducing a new cleaning and disinfection regime can be a daunting step, with lots of elements to consider to ensure a compliant implementation. This presentation provides a detailed account taking you through the stages of a new cleaning and disinfection project.

- Planning the project and completing the change control
- Reviewing and approving the supplier
- Selecting the appropriate agents for your facility
- Planning and executing the relevant validation steps
- Introducing quality control measures for the new products
- Designing the rotational regime aligned to the contamination control risks
- The final stage where the cleaning and disinfection stage is implemented, including the additional cleaning steps for start-up of a new facility.

Helen Gates – Global Technical Consultant, Ecolab

15.45 – 16.00
10.45 – 11.00 **Q&A session, summary and close of virtual day two**

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BOOKING FORM

Cleaning & Disinfection:
A roadmap to compliance and how to make sense of it all

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Attend and receive the latest updated:

Guide to Disinfectants & their use in the Pharmaceutical Industry 2021

Company: _____

Address: _____

Contact name if different from the delegate: _____

Tel: _____ Email: _____

DELEGATE 1

Surname: _____ First Name: _____

Job Title: _____ Email: _____

DELEGATE 2

Surname: _____ First Name: _____

Job Title: _____ Email: _____

Cleaning & Disinfection: A roadmap to compliance
15th/16th September 2021 (2 x 1/2-day meetings via Zoom)

Attend and receive the updated: Guide to Disinfectants & their use in the Pharmaceutical Industry

FEES

Member Fees **Euro / Dollar**
Member Delegate - £450/ *€538 /*$688

Non-Member Fees **Euro / Dollar**
Non-Member Delegate - £650/ *€768 / *$912

NHS Member Fees
Member Delegate - £250

NHS Non-Member Fees
Non-Member Delegate - £350

NOTE: * fee is higher to cover conversion rates

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PAYMENT METHODS – please tick relevant box

☐ Please raise an invoice to cover the delegate fee(s) £/€/$ _____

☐ UK BACS Sort code 60 19 28 Account 80843867 £/€/$ _____

☐ Wire Transfer Natwest Bank, 118 High Street, Slough, Berkshire SL1 1JH
SWIFT (BIC) NWB KGB2L Account 80843867
IBAN GB64 NWBK 6019 2880 843 867
£/€/$_____

☐ Please quote company approved purchase order no. _____ For £/€//$ _____

☐ I/we wish to pay by credit card (Pharmig will contact you for details)

Please note: Due to these unusual times, and the need for Pharmig to ensure fees are received in a timely manner before the event, it will have the right to cancel a delegate's attendance unless alternative arrangements have been agreed in advance of the event(s) by both parties.

INFORMATION ON FEES & PAYMENTS

Where possible fees must be paid in advance in order to attend the virtual conference. This can be done via credit card payment OR by an approved company purchase order number

INFORMATION ON ZOOM SET UP

Once you have sent back your registration form and payment (please see info of fees and payments above), Pharmig will add you to the Zoom list and will send out an email which will include the Zoom will include the Zoom link and additional information.
For those not familiar with Zoom – we will do a quick tour around the screen (it's all very straightforward) before starting the session on the afternoon of the 15th September 2021.

Please note: all information addressed by the speakers are of their own/ their company opinions and viewpoints. Pharmig is not responsible for any content presented at the meeting.
Pharmig also has the right to change the programme at any time due to unforeseen circumstances.

By registering for these events, I accept the processing of my Personal Data. Pharmig will use my data for the processing of this order for which I hereby declare to agree that my personal data is stored and processed. Pharmig will only send information in relation to this order or similar events/publications/training courses etc. Pharmig may send your name and company only to other companies attending the same event in the form of an attendee list. Your full personal data will not be disclosed to third parties (see also privacy policy at https://www.pharmig.org.uk/en/privacy-policy/). You can ask for the modification, correction or deletion of my data at any time via an email to maxine@pharmig.org.uk