

Training Course Agenda

PDA EU00007 Extractables and Leachables

CEST

Day 1, 19 October 2023		09:00 – 17:00
09:00	Intro and Attendee Expectations	
09:15	Introduction on Extractables & Leachables (E/L) <ul style="list-style-type: none"> • What is the importance of a good E/L-qualification? • Historical cases of leachables, impacting the quality or the safety of a drug product • Regulatory requirements (FDA, EMA...) for primary packaging 	
10:30	Coffee Break	
10:45	Understanding the Materials, Used in the Manufacture of Pharmaceutical Containers & Closures <ul style="list-style-type: none"> • Types of polymers and their physicochemical properties: examples in medical/pharmaceutical use • Understanding the composition of polymers: Intentionally added & Non-Intentionally added Compounds: Their function and origin • The issues with glass in parenteral applications 	
11:30	The Mechanism of Leaching <ul style="list-style-type: none"> • What are the physicochemical parameters to be considered when trying to understand polymer migration • How do leachables move through a polymer, the diffusion model • Special cases in migration 	
12:00	Lunch Break	
13:00	How to Set-up Extractables & Leachables Studies for Pharmaceutical Container Closure Systems <ul style="list-style-type: none"> • Selecting the right conditions for extraction • How to select the right compounds to monitor in a leachable study • Designing a leachable study 	
14:45	Coffee Break	
15:00	Analytical Techniques to Perform Extractables & Leachables Research <ul style="list-style-type: none"> • The importance of sample preparation: the corner stone in E/L research • What are the target compounds for material research • How does a classification of these compounds assist in finding the right analytical technique • From basic "screening" methodologies to state-of-the-art equipment 	
15:45	How to Perform a Safety Evaluation – Risk Assessment on Extractables & Leachables <ul style="list-style-type: none"> • Toxicology 101 • EMA Guideline on Genotoxic Impurities • ICH M7 (DNA reactive Impurities) and its suggested staged approach • The Threshold Concept of PQRI (OINDP and PDP/ODP) • Examples 	
17:00	End of Training Course Day 1	

Day 2, 20 October 2023		08:30 – 16:00
08:30	Recap Day 1	
08:45	E/L Testing for a Small Volume Parenteral Container Closure systems <ul style="list-style-type: none"> • Glass Syringes: the issues with tungsten, glue residues and silicone oil and glass metals leaching • The Issue with rubbers: the plunger, the needle shield or the tip cap: different approaches needed? • The impact of secondary packaging – option or necessity? • Setting up extractable & leachable studies for a pre-filled Syringe • Biological Drug Products versus Small Molecule Drug Products. • Primary packaging for the lyophilized drug product – modus of interaction with the DP • Critical aspects when designing leachable studies for lyophilized DP • Reactivity Of Leachables: concern for Lyophilized drug Products • Integration of the administration procedure (e.g. IV-set, pump system) in leachables evaluation 	
10:00	Coffee Break	
10:15	E/L Testing for Disposable and Single-Use Systems in Bioproduction <ul style="list-style-type: none"> • How to classify the risk of different single-use systems in the bioproduction process? • Understanding BPSA & BPOG recommendations, and how they can be implemented in the study design • Performing E/L studies on filters: potential approaches 	
11:00	Large Volume Parenterals <ul style="list-style-type: none"> • The challenge in E/L testing for LVP's • Primary packaging for LVP's – critical materials and components • Secondary packaging for LVP: critical points to consider 	
11:30	Lunch Break	
12:30	Qualification of Injection Devices used for Parenteral Administrations <ul style="list-style-type: none"> • Short introduction into Medical Device Regulations (ISO 10993 series) • Difference in Approaches for Medical Devices, compared to Pharmaceutical Packaging • Considerations for Combination Products: how to proceed? 	
14:00	Coffee Break	
14:15	Updates of E/L- Regulations, Standards and Recommendations <ul style="list-style-type: none"> • Pharma Packaging: <ul style="list-style-type: none"> ○ Final PQRI recommendations of the Parenteral Drug Product (DPD) Chemistry group ○ USP <661> & USP<665> chapters: where are we, where could it go to? • (Bio)Pharmaceutical Manufacturing <ul style="list-style-type: none"> ○ The BPOG protocol 	
15:45	Q&A	
16:00	End of Training Course	
The training course will include two coffee breaks and one lunch break		