

FECC TECHNICAL WORKSHOP 15 MAY, 2018



CHALLENGES AND TOOLS FOR *PHARMACEUTICAL EXCIPIENTS SUPPLY CHAIN*

When

15 May 2018, 10:00-17:00

Where

Fecc offices, Rue du Luxembourg 16b, B-1000 Brussels

Target audience

Professionals working in the supply chain of pharmaceutical starting materials.

Introduction

The current medicinal products regulatory framework in Europe sets additional legal requirements to ensure the quality and traceability of pharmaceutical starting materials. GMP and GDP as well as supplier audits and appropriate oversight on the supply chain became challenging than in the past. In order to achieve this compliance suppliers, distributors and pharmaceutical companies share responsibilities that need to be properly managed through the entire supply chain.

With this workshop distributors, as the link in the supply chain, will be informed about existing tools helping to manage challenges under the starting materials legislation.



Programme

10:00-10:15	Registration
10:15-10:30	Opening and welcome
10:30-11:15	GMP and GDP for pharmaceutical excipients along the supply chain <i>Dr. Mathias Brenken, former Dow Chemical</i>
11:15-12:00	Quality Risk Assessment for pharmaceutical Excipients – Applicability in the entire supply chain <i>Kaat Bracquine, Lonza,</i>
12:00-12:45	Expectations of a pharma company for suppliers' contribution to the excipients risk assessment <i>TBC</i>
12:45-13:30	<i>Lunch break</i>
13:30-14:15	Challenges for Quality Agreements between Manufacturer, Distributor and User of Excipients <i>Astrid Stockrahm-Uhling, DFE Pharma</i>
14:15-15:00	Challenges under the current excipient legislation with working with distributors <i>TBC</i>
15:00-15:15	<i>Coffee break</i>
15:15-16:00	Distributors' challenges in the supply chain of excipients <i>Catherine Martin, Univar</i>
16:00-16:45	Current challenges on APIs distribution <i>François Vandeweyer- APIC</i>
16:45-17:00	Closure

For registration

Contact Mrs. Irantzu Garmendia: iga@fecc.org