EUROPEAN PRE-FILLED SYRINGES SUMMIT 7th-8th Sept 2016, London UK

Innovations in Injectable Device Development

KEY TOPICS

• Meeting demands for change: Change Management with Owen Mumford
• Lessons learned from PFS development and prospects for future applications with Itshak Golan Project Evaluator for The European Commission
• Hear from Senior clinical development manager Florence Schwabenbach from Becton Dickenson (BD) with Expert Insight on Patient Focused Development
• Integrating Quality by Design - Value Adding Principles and Challenges from Novartis
• Exploring Safety Features & Safety Syringes while overcoming hurdles with Roche Pharmaceuticals
• Hear the Latest Initiatives from Mike Hrytsak from Regeneron Pharmaceuticals

Event Sponsor:

Nemera STERISYS

For More Information & Registration:
Mohammad Ahsan +44 (0) 20 3141 0606 mahsan@acieu.net
The Conference

Healthcare systems around the world are trying to move healthcare out of hospitals and into the home. Self-injection continues to offer a promising alternative for the delivery of drugs, prompted by the rise in chronic disease, and rapid growth of biologics. As more sensitive drugs enter the market the need for higher quality, more reliable packaging becomes ever more important. The ongoing challenges remain within the evolving regulatory landscape, packaging material variability and stability logistics in the supply chain. To succeed in this new environment, device manufacturers must rethink a pioneering model for market success. As the industry continues to grow, we shall be looking beyond forthcoming obstacles which will help the industry get equipped for solutions for the future. The programme has been designed to contribute to helping the European biopharma sector step into the future of manufacturing, where the demand is for the prefilled syringes to transition from being an off-the-shelf commodity device, to one in which component materials, functionality, and branding are fully customisable.

Confirmed Topics for Discussion:

- The latest prospects for applications in prefilled technology - anticipating requirements for the future
- Meeting demands for change
- Lessons learned from PFS development and prospects for future applications
- The latest initiatives on patient focused development
- Integrating quality by design: value adding principles
- Exploring safety features while overcoming technical hurdles
- Large volume development of prefilled syringes
- Mitigating risk in the supply chain
- Leading profiles product development activities
- Continual improvement through data

Call For Papers

If you would like to be considered as a speaker at the event for a 30-45 minute presentation, please submit an abstract for consideration to:

Who Will Be Attending?

The two-day conference will bring together the leading executives and experts from across the entire value chain uniting a variety of industry perspectives to discuss solutions and strategies for topical discussions, helping companies of all sizes to develop, prove, and commercialise new and improved processes and technologies for biopharmaceutical manufacture with an emphasis on the competitive landscape.

- Head of Device Development, Head of formulation development, Head of Product Management
- Head of Human Factors, Senior Application Specialists, Senior Packaging Engineers, Medical Devices Senior Director, Packaging, Head of Medical Devices

Commercial & Sponsorship Opportunities

With leading companies and organisations from the aromatic industry attending and speaking at our event, we have the perfect vehicle to provide outstanding exposure to a senior level audience. There are varying sponsorship packages available, including sponsorship of a cocktail reception on the first evening of the event and sponsorship of a networking lunch.

For further details, please contact:

Timothy Rowley-Evans
+44 203 141 0637 / trowleyevans@acieu.net

Registration Is Simple

If you would like to register for this event or wish to find out more information, contact Mohammed Ahsan
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D DAY 1
Wednesday 7th September 2016

08:00 REGISTRATION & COFFEE

09:00 CHAIRMAN’S OPENING REMARKS
Mike Hrytsak
Combination products & Regulatory affairs
Regeneron Pharmaceuticals Inc.

09:15 SESSION ONE
Regulation & Compliance: Meeting Demands for Change
Exploring the shifting landscape for combination products: how to stay competitive in the prefilled syringe space, and the expected impact on the traditional pharmaceutical model in years to come.
Prospects for applications in prefilled technology - anticipating requirements for the future.
Janine Jamieson
Former Assessor
MHRA

Rob Fabien
Senior associate
Hogan Lovells LLP
• Conceptual choices in design - what to look for in the ideal device: platforms, pumps, connectivity, featured functionality that works for the value chain.

Martin Murphy
Medtech Division
Cambridge Consultants
• Change management of drug delivery combination products

Andy Varde
Director of Research & Development
Owen Mumford

10:45 SESSION TWO
Revolutionising Biologics Administration: Safety Approaches to Design
The increasing demand in homecare prompts the need for more innovative safety systems this session looks at the considerations.
• Exploring safety features and safety syringes while overcoming technical hurdles
Hadj Latreche
Combination Products Methods Group Leader
Roche Pharmaceuticals

• Making it safer for healthcare workers
Emmie Galilee
Head of Health and Social Care Services Unit
Health and Safety Executive (HSE)

11:45 MORNING REFRESHMENTS

12:15 SESSION THREE
Integrating Quality by Design
The latest opportunities and challenges within the development & maintenance of the product lifecycle: Value adding process principles for a quality system that merges design factors, material selection issues, technologies, science, alongside integrating it into manufacturing operations.
Prefilled syringes at Novartis - biologic products and related technical and quality issues with respect to the prefilled syringe
Tom Fischer
Senior Quality Manager
Novartis

• Development & enhancement in design & manufacturing
Phillipe Lauwers
Business Unit Director PFS
Nipro Europe

13:00 PANEL DISCUSSION
Lessons Learned in PFS Manufacturing & Mitigating Risk in the Supply Chain
Sharing success and lessons learned from routine production and previous applications and how these factors could further optimise development through to product launch while reducing instability and exploring new opportunities in functionality within the supply chain.
• What are the best integrated manufacturing methods for risk management?
• How can the product development cycle adapt to increase patient compliance?
• Preventing needlestick injury - the latest initiatives
Dawn Smith
HM Principal Specialist Inspector (Occupational Health)
Health and Safety Executive (HSE)

Dr. Rene Dathe
Project Manager Medical Devices
Sanofi

13:45 LUNCH
14:45  SESSION FOUR
The Patient Focused Development Guide – Defining the User
Exploring strategy and technologies that merge innovation and commercialization while incorporating human factors engineering.

- Human factors and usability - what is the most anticipated need for packaging, thus the primary concern for patients?

Florence Schwarzenbach
Senior Clinical Development Manager
Becton Dickenson (BD)

- Pain associated with injecting higher doses from a patient and device interaction perspective

David Blakey PhD
Senior Engineer - R&D Platform Technology & Science
GSK

16:00  AFTERNOON REFRESHMENTS

16:30  SESSION FIVE
Technical Spotlight: Achieving More Flexibility in Manufacturing
The need for flexible solutions to bring products to market faster is crucial. This session explores more efficient process options, to manufacturing methods and additional product requirement solutions.

- Process and efficiency of applying the TRIZ methodology to medical device innovations

Dr. Rene Darthe
Project Manager Medical Devices
Sanofi

Patrick Radermacher
Managing Director
Sterisys Sàrl

17:10  CLOSE OF DAY ONE

Commercial & Sponsorship Opportunities
For information on available commercial opportunities, please contact Tim Rowley Evans:
T. +44 20 3141 0637
F. trowleyevans@racieu.net

DAY 2
Thursday 8th September 2016

08:30  REGISTRATION & COFFEE

09:00  CHAIRMAN’S OPENING REMARKS

Mike Hrytsak
Combination products & Regulatory affairs
Regeneron Pharmaceuticals Inc.

09:05  CONFERENCE PRESENTATION
Bridging the Gap Between Patient, Drugs & Delivery
- The latest steps forward in patient adherence
- Improving medication adherence
- We answer to common questions about how to involve staff and patients in identifying nonadherence and changing behaviors

09:30  SESSION SIX
Advanced Delivery Technologies: Trends, Success Stories and Opportunities. How to Design the Next Successful Self-Use Combination Device
Various drug delivery projects both from a technology and user focus.

Jeremy Kooyman
Medical Device Design & Development
Cambridge Design Partnership

10:15  MORNING REFRESHMENTS

10:45  CONFERENCE PRESENTATION

Adrian Tisserand
Global Category Manager
Nemera

11:25  CONFERENCE PRESENTATION

Bastiaan Delleauw
Head of Business development
Oval Medical Technologies

12:05  LUNCH

13:05  CONFERENCE PRESENTATION
Smart Devices Showcase: Next Generation Injectable Devices & Recent Collaborations

Ian Thompson
Vice President Business Development
Ypsomed AG

14:15  CHAIRMAN’S CLOSING REMARKS

16:00  END OF CONFERENCE & AFTERNOON REFRESHMENTS
European Prefilled Syringes Summit
London, UK

Registration Is Simple

If you would like to register for this event or wish to find out more information, you can contact Mohammed Ahsan using any of the following methods:

+44 (0)20 3141 0606
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Postal Address:
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Terms & Conditions

Payment
Payment must be received within five business days of returning the signed contract. After receiving payment a VAT receipt will be issued. If you do not receive a letter outlining details two weeks prior to the event, please contact the Conference Coordinator at ACI Europe Ltd.

Discounts are available for multiple/group bookings. Please call +44 (0)20 3141 0606 for more information.

Cancellations
Substitutions are welcome up to 24 hours prior to the event. Cancellations must be received in writing no less than 3 weeks prior to the start of the conference; a full credit voucher towards a future ACI conference will be issued. Any cancellation received less than 3 weeks prior to the start of the event shall be deemed to be a breach of this contract by client, and accordingly, no credits will be given. Cancellations must be received in writing by mail or fax three weeks before the conference. Thereafter the full conference fee is payable. If for any reason ACI Europe Ltd decides to amend, cancel or postpone this conference, the conference fee will not be refunded. Furthermore, ACI Europe Ltd will not be responsible for covering airfare, hotel or other costs incurred by registrants. In the event that ACI Europe Ltd cancel or postpone the event, ACI Europe Ltd reserves the right to transfer this booking to another conference to be held in the following twelve months, or to provide a credit of an equivalent amount to another conference within the following twelve months. The construction, validity and performance of this agreement shall be governed in all respects by the laws of England to the exclusive jurisdiction of whose courts the Parties hereby agree to submit.

Accommodation
The cost of accommodation is not included in the event fee. Preferential rates will be arranged with or near the event venue, and all confirmed delegates will be given details of how to book accommodation at this rate in due course.

About ACI

ACI, a UK owned company, has been running successful conferences in the USA since 1999. Headquartered in Chicago with offices all around the States, ACI opened its European head office at the end of 2005 and has expanded rapidly, launching a series of events in key industries including maritime, energy, oil & gas, cosmetics, chemicals & media.

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