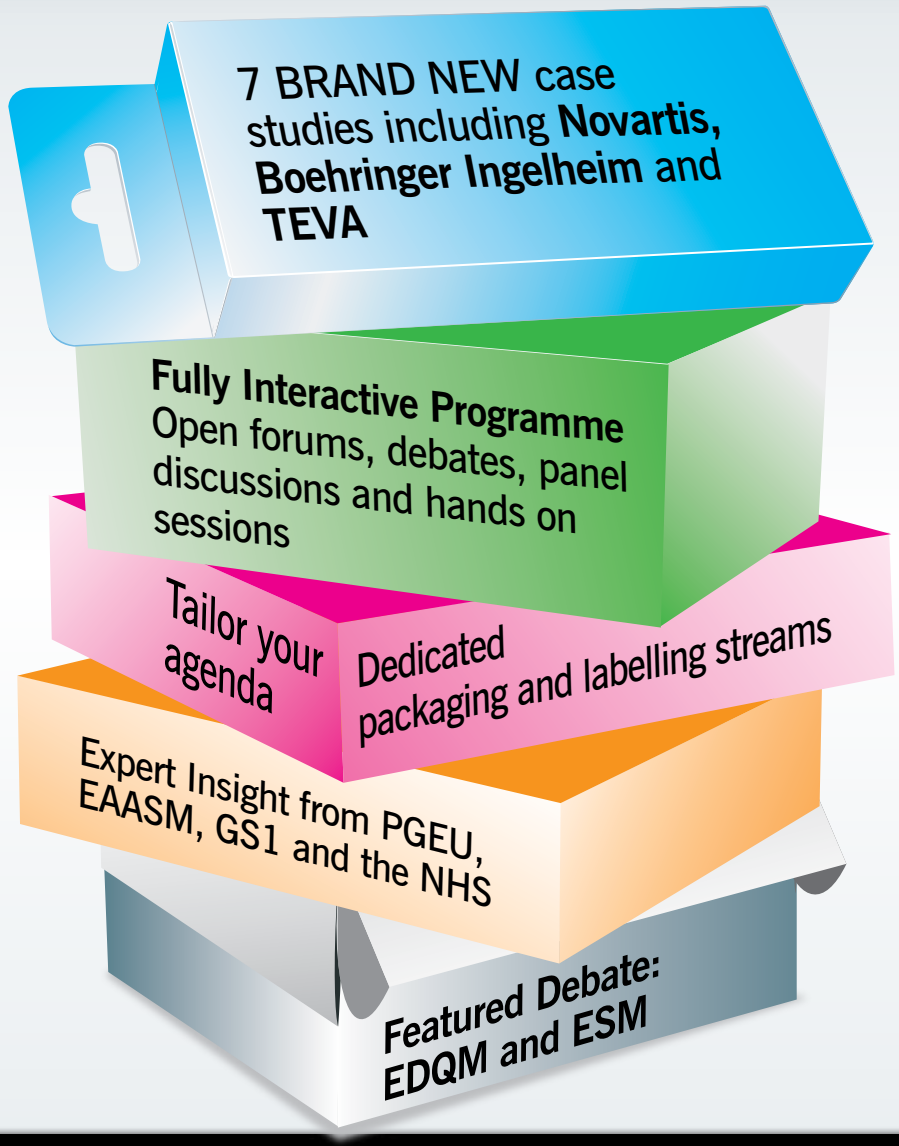
Pharma iQ
a division of IQPC
Presents

Pharmaceutical Packaging and Labelling Summit 2013

Pre-conference day:	Conference:	Location:
26th June 2013	27th-28th June 2013	Ramada Plaza, Basel, Switzerland

Improve Patient Compliance, Prepare for the Falsified Medicines Directive and Maximise Artwork Capabilities while Protecting your Supply Chain



2013's Expert Speaker Panel Features:


Michael Ritter, Global Project Manager Serialization & Product Tracking, **Novartis**


Christoph Krähenbühl, EFPIA


Johannes Schoen, Senior Manager, Anti-Counterfeiting, **Boehringer Ingelheim**


Omar Ali, Pharmacist, **NHS Healthcare Trust**


Eduardo Vidal Blanco, Device & Packaging Development, **Novartis**


John Chave, Secretary General, **PGEU**

Ezri Deshen, Senior Director - Solid Operations, **Teva Pharmaceuticals**

Hans Bigalke, Deputy Head, **EDQM**

Massimo De Carlo, Global Supply Chain Quality System Manager, **F. Hoffmann- La Roche**

"Perfect organisation, great setting and high level of speaker panel. All in all an excellent conference!"
HCPC Delegate 2012

"Very valuable, learned a lot, inspired!"
Novartis Pharma 2012

Not just another packaging conference!

Gain critical insight from industry leaders, key association and regulatory bodies to keep you ahead of the curve.

- Face the challenge of supply chain security head on with cost-effective anti-counterfeit strategies
- Prepare for implementation of mass serialization and the Falsified Medicines Directive
- Maximise patient compliance
- Reduce costs and streamline your artwork capabilities while facing the complex regulatory environment.

www.pharmapackaginglabelling.com

Dear Colleague,

I am delighted to introduce **Pharma IQ's 3rd Annual Pharmaceutical Packaging and Labelling Summit** 2013. Once again we bring together the global leaders from across the field to thrash out the core challenges facing the industry today.
Benchmark against your peers and get cutting edge insight into the latest solutions and best practices for:

- **Implementing track and trace and serialisation** across multiple geographic jurisdictions where strict regulatory requirements differ
- **Effectively preparing for the falsified medicines directive**
- Preventing counterfeit and **ensuring tamper prevention**
- **Capitalising on artwork capabilities** and dealing with the growing complexity while continuing to maximise efficiency
- Cost-effectively implementing methods to **improve patient compliance**
- Ensuring **effective text management to maximise efficiency** while maintaining the highest standards
- Analysing the latest developments in **cost effective "green" packaging** and technology

New for 2013:

- Structured debate: EDQM and EFPIA
- Latest regulatory updates on the EU GDP guidelines
- Dedicated labelling and tamper evidence streams allowing you to tailor to the agenda to suit your needs
- Fully Interactive programme with panel discussions, open forums, roundtable sessions and interactive discussion sessions

Join industry experts, global regulatory influencers and powerful bodies such as **EFPIA, EDQM, GS1** and **PGEU** to discuss the latest advances in the packaging and labelling field. Benchmark with your peers and network with the latest solution providers keeping you at the cutting edge in this ever changing environment.

I look forward to meeting you in Basel.

Kind regards

Anna Fraenkel
Event Director -**Pharma-IQ**
Anna.fraenkel@iqpc.co.uk

2013's Expert Speaker Panel Features:

Ezri Deshen, Senior Director - Solid Operations, **Teva Pharmaceuticals**

Michael Ritter, Global Project Manager Serialization & Product Tracking, **Novartis**

Carolina Sánchez-Céspedes, Head of Packaging**, Lilly, S.A. Spain**

Jochen Weisser, Project Lead**, F. Hoffmann- La Roche**

Christoph Krähenbühl, EFPIA

Johannes Schoen, Senior Manager, Anti-Counterfeiting**, Boehringer Ingelheim**

Stephen Wilkins, Chairman, **Child-Safe Packaging Group**

Horst Kastrup, Senior Regulatory Advisor**, MEDA Pharma**

Omar Ali, Formulary Development Pharmacist, **Surrey & Sussex NHS Trust, UK**

Eduardo Vidal Blanco, Device & Packaging Development, **Novartis**

John Chave, Secretary General, **PGEU**

Michael Niewesteeg, Managing Director, **NVC Netherlands Packaging Centre**

Jim Thomson, Chairman, **EAASM**

Massimo De Carlo, Global Supply Chain Quality System Manager, **F. Hoffmann- La Roche**

Hans Bigalke, Deputy Head, **EDQM**

Andrew Love, VP Capability Development, **Be4ward**

Benoit Defrasne, Business Development Manager, **SICPA**

Christian Hay, Senior Consultant Healthcare, **GS1**

Dirk Hendrik Kneusels, Commercial Director and Business Development**, Laetus GmbH**

Dieter Moessner, Chairman, **German Standards Institute DIN / the European Committee for Standardisation CEN**

Taha Yayci, consultant, **Ministry of Health of Turkey**

Marco Kutscha, General Manager, T.S.P. Germany GmbH, **Delfort Group**

Pharmaceutical Packaging and Labelling Summit

2012 saw representatives from companies such as:

Perigord Group
Meda Pharma GmbH & Co. KG
Reckitt Benkiser.
Roche
Novartis Pharma AG
Mhra
J&J
Pfizer Wyeth
Mylan Inc
GS1
HCPC-Europe
Viropharma Europe
SwissMedic
Blue Sphere Health Ltd
EAHP
Arjowiggins Security
SICPA Holding SA
Bristol-Myers Squibb GmbH
Fresenius Medical Care Deutschland
Eli Lilly
CSL Behring
Spirig Pharma AG
Vifor Pharma
GSK
Genericon Pharma
CSL Behring
Actelion Pharmaceuticals
Ltd.
Orion Corporation
UCB Pharma S.A.
Synthes GmbH
Actavis
SANTO member of Polpharma Group
Coloplast Danmark A/S
GSK Vaccines
K+D AG
Abbott Healthcare Products BV
Abbott Products Operations
NNE Pharmaplan
Drossapharm AG
Intertek Expert Services
Movianto UK Ltd.
Abbott Laboratories
Rolling Optics AB/ ePP
Tervakoski Oy
SYSTECH International
Kessler A.S
SICPA Product Security S.A
B4ward, Former Director Global Pack Management, GSK
SICPA Product Security S.A
Perigord Group
Nordvalls
Atlantic Zeiser
Novo Nordisk A/S
Avery Dennison
Wessex Technology

Testimonials from Packaging and Labelling 2012:

"First conference I have seen with actual implementation of serialisation case studies from the big pharma" – **Janice Kite GS1**

"Perfect organisation, great setting and high level of speaker panel. All in all an excellent conference!" – **HCPC**

"A very Complete and Diverse Panel" – **Roche**

"Fantastic opportunity to network with industry peers" **GSK**

"Very valuable, learned a lot, inspired!" **Novartis Pharma**

Pre-Conference Day: Wednesday June 26th 2013

Our pre-conference day provides a unique working environment where specific specialised groups meet and discuss the hottest topics in a relaxed informal environment.

To make the most out the Pharma IQ packaging and Labelling Summit 2013, get involved with the workshops and take away key learning's that are only developed in the small exclusive environment provided here.

09:00	*WORKSHOP ONE*
	Minimising Costs by Delivering Right-First-time Artwork Across the Globe **Understanding all the capability elements required to deliver accurate artwork in a global pharmaceutical company.** Top takeaways: 1) The challenges facing artwork capabilities, internally and externally - Introduction to SWOT analysis - Sub-group break-outs to develop SWOT for artwork capabilities 2) Introducing model capability - Discussing the elements of a full artwork capability model 3) Defining capability elements - Sub-group break-outs to develop key contents of each capability on a sliding scale of "In Control" through to "World Class" 4) Personal gap analysis - Tailored input into your challenges. The opportunity to give a personal review of where your organisation is and what gaps they might have and receive expert feedback **Andrew Love,** VP Capability Development , **Be4Ward (Former Director Global Pack Management - GlaxoSmithKline)**
13:00	*WORKSHOP TWO*
	Ensuring Manufacturing Readiness: The Delegated Acts are Not a Waiting Game The delegated acts are due for release in 2017 but there is plenty we can prepare for in the mean time. No company, big or small, can be in a position to sit waiting for the specific delegated acts to be released following the falsified medicines directive. This interactive discussion-based workshop session will walk you through the key steps to ensuring you as prepared as you can possibly be before the final release of the delegated acts. **Christoph Krähenbühl, EFPIA**

Day One, Thursday June 27th 2013

08.15	**Registration**
08.50	**Pharma-IQ Welcome**
08.50	**Chairman's Welcome**
	REMAINING REGULATORY COMPLIANT TO ENSURE GLOBAL PRODUCT DELIVERY
09.00	**Best Practices on How to Ensure Pharma Contract Packagers Compliance with US ePedigree Law and EU Directive of Falsified Medicines** - Analysing the multiple regulations impacting contract packagers - Best approaches taken to achieve contract packagers compliance - What decisions need to be taken to ensure compliance? - Outlining and overcoming the challenges faced **Michael Ritter,** Global Project Manager Serialisation & Product Tracking, **Novartis**
09.30	**Implementing E-Pedigree in a Global Company – The Challenges** - Highlighting the latest requirements across the USA - Experiences and lessons learned from current implementation - Our approach to overcoming the challenges of global supply **Ezri Deshen,** Senior Director - Solid Operations, Global Operations, **Teva Pharmaceuticals**
10.00	**Networking Coffee Break**
10.30	**How to Implement Smart Product Protection and Gain Targeting Visibility** - Building smart solutions protecting products and supply chains, beyond compliance requirements - Low invasive authentication and tamper-evident solutions - Value added traceability **Benoit Defrasne,** Business Development Manager, **SICPA**
	THE FALSIFIED MEDICINES DIRECTIVE: COST EFFECTIVLY IMPLEMENTING SERIALISATION
11.00	**EDQM project for an anti-counterfeiting traceability service for medicines: moving towards a real-scale system:** - The eTACT project - Added value of involvement of EDQM in pharmaceutical serialisation - Public governance - Next steps **Hans Bigalke,** Deputy Head**, EDQM**

Day One Continued:

11.30	**Interactive Discussion Session: What are our Expectations for the Delegated Acts?** Interactive! - What will be expected of the pharmaceutical industry? - What do we expect from the specific acts? - The black and white list: ruling products in and out - Best practices for generics: should they be expected to comply? **Panel to feature members of our expert speaker panel including John Chave,** Secretary General, **PGEU**
12.30	**Networking Lunch**
	ENSURING PATIENT SAFETY AND MAXIMISING COMPLIANCE
13.30	**Designing Packaging around Child Safety and Medication for the Elderly** - Child resistant packaging and the latest regulations - Overcoming the challenge of ensuring child safety and maximising elderly access to medication - Best approach for cost effectively implementing child resistant packaging **Stephen Wilkins**, Chairman, **Child-Safe Packaging Group**
14.00	**Interactive Session: the Latest Developments in Pack Design for Patient Safety** Interactive! Topics to be covered in discussion include: - Latest regulations surrounding patient compliance - Cost effective strategies to increase patient safety - Smart packaging, a reality? - Maximising profit by reducing patient error and increasing patient adherence Introduction and Facilitation by: **Eduardo Vidal Blanco**, Device & Packaging Development**, Novartis**
14.30	**TECHNOLOGY SPOTLIGHT SESSION!** **Hear the latest innovations from exciting new service providers as they each give 15 minutes flash sessions on their latest offerings.** *Are you interesting in showcasing your latest innovations in the packaging and labelling space? Please contact me on +44 (0) 207368 9300 or sponsorship@iqpc.co.uk*
15.30	**Networking Coffee Break**
16.00	**Packaging & Labelling on Assessment for Formulary Inclusion - A Payer's Perspective** - Designing packaging for those with chronic disease: Adapting for self medication - Maximising returns by ensuring medication is taken as prescribed - Reducing areas of error and improving patient compliance **Omar Ali,** Formulary Development Pharmacist, **Surrey & Sussex NHS Trust, UK**

Day One Continued:

16.30	**Structured Debate:** **ESM and EDQM Developing the Best Possible Model to meet the FMD** *(to be confirmed)* A chance to look at both the EDQM and ESM advocated plans to meet the Falsified Medicines Directive. A chance to compare the two models side by side and ask those specific questions to both parties at the same time. Interactive! Chairman: (to be confirmed) - **Hans Bigalke,** Deputy Head**, EDQM** - **John Chave,** Secretary General, **PGEU**
IMPLEMENTING RECYCLED PACKAGING AND OVERCOMING ENVIRONMENTAL CHALLENGES	
17.00	**Cost Effectively Driving Sustainability of your Packaging Line** - Minimising packaging and end of life materials to reduce waste - A demonstration of packaging we have made "greener" : old to new - Complying to regualtion on a global scale - Changing the packaging materials: what is available? **Johnson and Johnson** (representative to be confirmed)
17.30	**Chairman's Closing Summary**

"Provided great insight into different opinions on important topics within pharmaceutical packaging"

– Pfizer

Day Two, Friday June 28th 2013

08.00	**Registration & Networking Coffee**
08.20	**Chairman's Day Two Opening** **Michael Niewesteeg,** Managing Director, **NVC Netherlands Packaging Centre**
	PREVENTING ANTI-COUNTERFEIT AND COMPLYING TO LEGISLATION
08.30	**Ensuring the Safety of Your Supply Chain: Mass-Serialisation and Tamper Verification Features** - Facing global regulations: what are the requirements? - Understanding the developments in legislation - Meeting deadlines and complying to varying timelines **Johannes Schoen,** Senior Manager, Anti-Counterfeiting, **Boehringer Ingelheim**
09.00	**Get the most out of Track & Trace: Lessons Learned from Applying Turkish IT'S and the German SecurePharm Pilot** - From serialisation to warehousing: the holistic approach - The importance of being well prepared for aggregation - Best practices or possibilities to maintain high levels of line efficiency - Labelling and packaging in support of seamless track & trace **Dirk Hendrik Kneusels**, Commercial Director and Business Development**, Laetus GmbH**
09.30	**Case Study: Pharmaceutical Track And Trace System in Turkey (ITS)** - Ensuring reliable supply of drugs to patients - Preventing counterfeit, fraud, smuggling and barcode scams - Supporting rational drug use: supplying data to control market - Maintaining current theft and fraud levels at zero - Effective and complex recalls **Taha Yayci,** Consultant**, Ministry of Health of Turkey**
10.00	**GS1 Interactive Workshop Session: An Update on Healthcare Standards** An interactive session allowing you to gain feedback on the GS1standards**.** Analyse their developments and how they have been designed to assist in complying to the heavily regulated area of packaging **Christian Hay,** Senior Consultant Healthcare**, GS1** Interactive!
10.30	**Networking Coffee Break**

Day Two Continued:

	STREAM A COST EFFECTIVE IMPLEMENTATION OF TAMPER PROOF EVIDENCE	STREAM B BEST PRACTICES FOR ADAPTING YOUR LABELLING FOR GLOBAL SUPPLY
11.00	**CEN-Standard Tamper verification features for Medicinal Product Packaging** - Technical solutions for tamper verification features - Integrating Tamper Evidence Into Your Current Processes - Strategising for integrating tamper prevention into your packaging line - Overcoming challenges and reducing timelines - Facing the technical problems of tamper prevention **Dieter Moessner,** Chairman, **German standards Institute DIN / the European Committee for Standardisation CEN**	**Adopting Labelling Process For Global Supply: Driving Standardisation** - Driving Standardisation for braille requirements - Supplying a multilingual market: Streamlining the labelling development - Best approaches for delivering a product to the global market - Multilingual labelling for cost effective manufacture - The mock-up review process supporting economic packaging design **Horst Kastrup,** Senior Regulatory Advisor**, MEDA Pharma**
11.30	**Technologies and Innovations in the Tamper Evidence Space** - Highlighting the latest features that can be implemented to prevent falsification - Cost effective implementation of tamper evidence - Innovations in product security *Facilitator to be confirmed – please visit www.pharmapackaginglabelling.com*	**Effective Text Management: Implementing XML technology** - What is and how does it affect you? - Defining a fundamental change in how you manage information - Defining your template - Example: Our experience of XML *Reserved for a leading provider of XML technology*
12.00	**Panel Discussion: What are the Consequences of Integrating Tamper Evidence into your Packaging Line?** *With input from key experts across our speaker faculty*	**Panel Discussion: Maintaining The Highest Standards and Maximising Efficiency** *With input from key experts across our speaker faculty* Interactive!

Day Two Continued:

12.30	**Networking Lunch**
13:30	**Saving Paper and Saving Money** - Overview of the possibilities in paper materials for pharmaceutical packaging and leaflets - Regulatory considerations & readability guidelines - Critical cost analysis **Marco Kutscha,** General Manager, T.S.P. Germany GmbH - **Delfort Group**
14.00	**Remaining Competitive in the Current Environment: Effectively Implementing New Technologies** • Analysis of the environment: operational implications • Strategic approach • Use of new technologies and infrastructure • How we operate: the human factor **Carolina Sánchez-Céspedes,** Head of Packaging, **Lilly, S.A. Spain**
14.30	**Working to Reduce the Internet Supply of Falsified Medicines** - The Falsified Medicines Directive and the Internet - Collaborative actions to inform patients and counter online crime - The Future of Online Pharmacy - promoting the good guys - corralling the bad guys **Jim Thomson, Chairman, EAASM**
15:00	**ROUNDTABLE SESSIONS:** You can choose to attend one of four informal discussion sessions, depending on your goals and strategies. Join our resident experts for a no holds barred, frank discussion on the topic of your choice **1. Recycled packaging: the future for pharma?** **2. What can we expect from the delegated acts? Are we prepared?** **Horst Kastrup,** Senior Regulatory Advisor**, MEDA Pharma** **3. Globalisation: Facing the challenges of emerging markets head on** **4. Track and trace – What are the next steps?**
16.00	**Networking Coffee Break**

Interactive!

ARTWORK CONTROL SYSTEMS / PACKAGING TECHNOLOGIES	
16.30	**A New Global Artwork Change Management System** - From the artwork errors to the new system requirements - -Key process elements: e.g. regulatory compliance vs supply chain timelines - -Quality control plan for artwork change management - -Learning / next steps conclusion **Massimo De Carlo,** Global Supply Chain Quality System Manager, **F. Hoffmann- La Roche** **Jochen Weisser,** Project Lead, **F. Hoffmann- La Roche**
17:00	**Top Tips For Complexity Management** - How do deal with the ever growing levels of complexity developing in the small orders - How do we set up artwork capabilities? - Maximising efficiency while introducing new variants **Andrew Love**, VP Capability Development, **Be4ward**
17.30	**Chairman's Summary and Close of Conference**

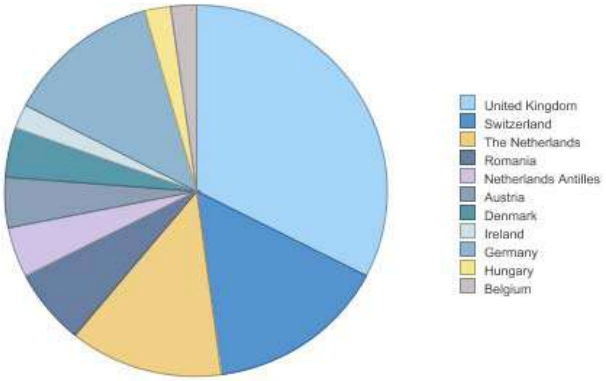
ABOUT THE EVENT

Who Should Attend?

Directors and heads of:

- Packaging
- Labelling
- Regulatory affairs
- Anti-counterfeit
- Supply Chain
- Quality Assurance

2012 Attendees by Country:



2012 saw representatives from companies such as:

Perigord Group
Meda Pharma GmbH & Co. KG
Reckitt Benkiser.
Roche
Novartis Pharma AG
Mhra
J&J
Pfizer Wyeth
Mylan Inc
GS1
HCPC-Europe
Viropharma Europe
SwissMedic
Blue Sphere Health Ltd
EAHP
Arjowiggins Security
SICPA Holding SA
Bristol-Myers Squibb GmbH
Fresenius Medical Care Deutschland

Eli Lilly
CSL Behring
Spirig Pharma AG
Vifor Pharma
GSK
Genericon Pharma
Reckitt Benckiser
CSL Behring
Actelion Pharmaceuticals
Ltd.
Orion Corporation
UCB Pharma S.A.
Synthes GmbH
Actavis
SANTO member of Polpharma Group
Coloplast Danmark A/S
GSK Vaccines
K+D AG
Abbott Healthcare Products BV

Abbott Products Operations
NNE Pharmaplan
Drossapharm AG
Intertek Expert Services
Movianto UK Ltd.
Abbott Laboratories
Rolling Optics AB/ ePP
Tervakoski Oy
SYSTECH International
Kessler A.S
SICPA Product Security S.A
B4ward, Former Director Global Pack Management, GSK
SICPA Product Security S.A
Perigord Group
Nordvalls
Atlantic Zeiser
Novo Nordisk A/S
Avery Dennison
Wessex Technology

Maximise Your Involvement: Sponsorship and Exhibition Opportunities

Packaging and Labelling 2013 is attended by senior officials and decision-makers from industry, bringing together buyers and suppliers in one location.

Focused and high-level, the event will be an excellent platform to initiate new business relationships. With tailored networking, sponsors can achieve the face-to-face contact that overcrowded trade shows cannot deliver.

Exhibiting and Sponsorship options are extensive, and packages can be tailor-made to suit your individual company's needs. Most packages include complimentary entry passes, targeted marketing to industry officials and executives, and bespoke networking opportunities.

Other features of sponsorship include:

- Prominent exhibition space in the main conference networking area
- Participation in comprehensive pre-event marketing campaigns
- Tailored marketing strategies to suit your organisation's size, capabilities and individual requirement

For more information and to discuss the right opportunity, contact us on +44 (0)207 368 9300 or sponsorship@iqpc.co.uk

Sponsors & Exhibitors 2013

Nordvalls is a family owned company established in 1907. We have for over 20 years been producing labels for the pharmaceutical industry on a good GMP-level.

In mid of May this year our new factory dedicated only for Pharma -labels and -leaflets will be up running in full scale. The factory has been lifted to a high GMP level. We can combine silk-screen, flexo and letter press. At Nordvalls we have our own in-house Pre-Press for a safer handling of print media. Safety is very important for us and we have therefor also developed a number of different security solutions against counterfeiting which we gladly would like to discuss with you.

http://www.nordvalls.se/en/

SICPA is a Swiss company, founded in 1927, with headquarters in Lausanne, and is a leading global provider of integrated security and traceability solutions to both, governments and companies of various industry verticals.

The company employs more than 2'500 people and is established on 5 continents, with business activities in most countries worldwide.

In the pharmaceutical and healthcare sector, the customer specific solutions and services delivered by SICPA help companies enhance the visibility and the integrity of their supply chains, while providing value-adding business intelligence and ensuring product protection and regulatory compliance.

SICPA has successfully deployed and is operating product authentication and traceability systems on all continents.

Sponsors & Exhibitors 2013

Laetus GmbH

Laetus is the world leading supplier of innovative and value-adding quality & security inspection systems and complete Track & Trace solutions for the Pharmaceutical, Medical, Cosmetics and FMCG industries.
With 40 years of market experience and our global network of sales, service and project management staff, we help our customers to maintain the highest levels of product quality and production efficiency.
Technological innovation and a strong commitment to our customers and partners have always been our key focus. Many of our product brands such as ARGUS, INSPECT and POLYPHEM have become market standards in code reading, quality assurance and fill inspection.
Laetus is a member of the privately owned Coesia group with 5.000 employees.

delfortgroup is world leader in the design, development, manufacture and promotion of thinner, more sustainable, tailor-made speciality papers. With ongoing product innovations and the ultimate paper solutions, we offer service from a number of locations around the world. Not only are we dedicated to high performance in everything we do, we also have the experience, ability and desire to optimally understand your needs.

delfortgroup is a leading supplier of thinprint paper and supplies standard and tailor-made quality products combined with outstanding service.We know that specialisation and tailor-made products are the key to success. Consequently, our product range is subject to continuous development in line with changing customer needs. State-of-the-art paper machines at all locations enable us to be leaders of technical innovations.

delfortgroup's approach contributes to the reduction of carbon emissions in the value chain. Our paper is produced in using strict environmental management systems and guidelines like ISO14001 & 9001. delfortgroup is committed to care for the environment through sourcing sustainable supplies and efficient production methods.

delfortgroup has several different chains of custody accreditations depending on the type of end product, usage and request of our customers and partners. We aim to provide our customers with innovative quality products and service.

Baumer hhs GmbH

With its headquarters in Krefeld, Baumer hhs GmbH is an internationally operating manufacturer for industrial glue application systems in conjunction with quality assurance systems and camera control systems. Baumer hhs provides its customers with a carefully selected portfolio from the areas of hot-melt and cold glue processing with guns, pumps and pressure tanks, as well as control and monitoring systems for quality assurance in glue application and adhesive application for factory automation. Further information on Baumer hhs, all products and additional services can be found online at www.baumerhhs.com

Pharma Packaging and Labelling Media Partners 2013

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AIPIA Active & Intelligent Packaging Industry Association

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WEBPACKAGING connecting packaging buyers & suppliers

Pharma a division of IQPC iQ

Website: www.pharma-iq.com
Phone: 1-646-454-4559

Delivering quality content and events to enhance your knowledge and strengthen your networks

Pharma IQ, a division of IQPC, provides a forum to address the critical issues facing the Pharmaceutical Industry today. Pharma IQ utilizes workshop, conference and training course formats to facilitate a learning environment for pharmaceutical professionals working in all areas of drug development: from Discovery to Post-Marketing.

Become a member here: www.pharma-iq.com/join.cfm

Pharmaceutical Packaging and Labelling Summit 2013

Pre-conference day: 26th June 2013
Conference: 27th-28th June 2013
Venue: RAMADA PLAZA, Basel, Switzerland

To speed registration, please provide the priority code located on the mailing label or in the box below.

My registration code **PDFW**

Please contact our database manager on +44(0) 207 368 9300 or database@iqpc.co.uk quoting the registration code above to inform us of any changes or to remove your details.

Pharma and Biotech Companies	Pay before 28th March 2013*	Pay before 26th April 2013*	Pay before 31st May 2013*	Standard Price
☐ Conference only	€1,699 SAVE €200	€1,749 SAVE €150	€1,899	€1,899
☐ Conference + 2 Workshops + Conference Recordings	€3,247 SAVE €500	€3,297 SAVE €450	€3,397 SAVE €350	€3,747
☐ Conference + 1 Workshop** + Conference Recordings	€2,698 SAVE €400	€2,748 SAVE €350	€2,848 SAVE €250	€3,098
☐ Conference + 2 Workshops	€2,797 SAVE €400	€2,847 SAVE €350	€2,947 SAVE €250	€3,197
☐ Conference + 1 Workshop**	€2,248 SAVE €300	€2,298 SAVE €250	€2,398 SAVE €150	€2,548
☐ Workshop only [Each]**	€649 SAVE €250	€649 SAVE €250	€649 SAVE €250	€899

Service Provider Companies	Pay before 26th April 2013*	Standard Price
☐ Conference Only	€2299 SAVE €100	€2399
☐ Conference + 2 workshops	€3397 SAVE €300	€3697
☐ Conference + 1 workshop**	€2,798 SAVE €250	€3048
☐ Workshop only [Each]**	€649 SAVE €250	€899

NB: UK Companies will pay UK VAT, all other companies are VAT exempt. VAT Registration #: GB 799225967

*To qualify for discounts, payments must be received by the early bird registration deadline. Early booking discounts are not valid in conjunction with any other offer.

**** Please select your pre-conference choice: Workshop 1** ☐ **Workshop 2** ☐

DELEGATE DETAILS - SIMPLY COMPLETE THIS FORM AND CLICK SUBMIT

Please photocopy for each additional delegate

☐ Mr ☐ Mrs ☐ Miss ☐ Ms ☐ Dr ☐ Other

First Name _____

Family Name _____ Job Title _____

Tel No. _____

Email _____

☐ Yes I would like to receive information about products and services via email

_____

Organisation _____

Nature of business _____

Address _____

Postcode Country _____

Telephone _____

_____

Approving Manager _____

Name of person completing form if different from delegate _____

I agree to IQPC's cancellation, substitution and payment terms

Special dietary requirements: ☐ Vegetarian ☐ Non-dairy ☐ Other (please specify)

Please indicate if you have already registered by: ☐ Phone ☐ Fax ☐ Email ☐ Web

Please note: if you have not received an acknowledgement before the conference, please call us to confirm your booking.

PAYMENT METHOD

Total price for your Organisation: (Add total of all individuals attending):

Card Number: VISA ☐ M/C ☐ AMEX ☐

_____

Exp. Date: _____ _____

Name On Card: _____

_____

City/County/Postcode _____ Cheque enclosed for: € _____ (Made payable to IQPC Ltd.)

(Please quote 20085.003 with remittance advice)

Bank account details (EUR): Account name: International Quality & Productivity Centre Ltd.
Bank: HSBC Bank Plc 67 George Street, Richmond Surrey TW9 1HG, United Kingdom
Account number: 59090618 Sort code: 400515 IBAN: GB98MIDL40051559090618 SWIFT: MIDLGB22

CONFERENCE CODE: 20085.003

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COMPLETE THE FORM AND CLICK SUBMIT

5 WAYS TO REGISTER

PHONE:
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FAX:
+44 (0)20 7368 9301

POST:
YOUR BOOKING FORM TO
IQPC, 129 WILTON ROAD,
VICTORIA, LONDON, SW1V 1JZ

EMAIL:
ENQUIRE@IQPC.CO.UK

WEB:
WWW.PHARMAPACKAGINGLABELLING.COM

TEAM DISCOUNTS*

IQPC recognises the value of learning in teams.
Groups of 3 or more booking at the same time from the same company receive a 10% discount.
5 or more receive a 15% discount.
7 receive a 20% discount.
Only one discount available per person. Team discounts are not applicable in conjunction with another discount.

VENUE & ACCOMMODATION

Venue:
Ramada Plaza, Basel, Switzerland

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