

Joint EFGCP / DIA / EMA  
Better Medicines for Children Conference 2014 on

# Explore Ways to Enhance Collaboration Between Key Players

30<sup>th</sup> September & 1<sup>st</sup> October 2014  
EMA Headquarters, London, United Kingdom

*Organised by*



[conferences@efgcp.eu](mailto:conferences@efgcp.eu) – [www.efgcp.eu](http://www.efgcp.eu)

## Conference Rationale

The EU paediatric regulation is now in force since 2007. Drug development is no longer possible without considering children. Furthermore, companies developing medicines need to consider the paediatric requirements early in the development. This legislation has transformed paediatric drug development from a topic discussed by a few interested paediatricians, clinical pharmacologists and regulators to an issue that is broadly known within pharmaceutical industry and regulatory authorities, and to a lesser degree in the clinical world. More than 1000 PIP decisions are now published on the EMA website, and virtually everybody within pharmaceutical industry has heard of paediatric investigation plans and waivers. The EU Commission has published a 5-year report, and a 10-year report will be submitted by the Commission to the European Parliament and Council in 2017.

The aim of this conference is to discuss on a high level how the EU paediatric regulation is working and how it contributes to children's health. This will include a discussion on the preparedness for the 10-year report; strategic thoughts within the EMA on how to streamline paediatric development and a session dedicated to paediatric oncology.

As always, experts from all involved parties will be present, and on day 1 participants will discuss more specialized and hot topic issues in four breakout sessions. This will allow participants to discuss face-to-face with all stakeholders, which otherwise usually occurs by email or phone. Questions on any topic relating to the Agency's activities can be submitted before the conference to [paediatrics@efgcp.eu](mailto:paediatrics@efgcp.eu), and will be answered by Peter Karolyi from the Paediatric Medicines office at the EMA.

## Programme Committee

<b>Gesine Bejeuhr</b>	vfa Research-based Pharmaceutical Companies, DIA Paediatric Group, Germany
<b>Emilie Desfontaine</b>	European Medicines Agency (EMA)
<b>Thorsten Olski</b>	European Medicines Agency (EMA)
<b>Jytte Lyngvig</b>	Drug Information Association (DIA), Switzerland
<b>Klaus Rose</b>	klausrose Consulting & EFGCP Children's Medicines Working Party, Switzerland
<b>Paolo Tomasi</b>	European Medicines Agency (EMA)
<b>Mette Due Theilade Thomsen</b>	Novo Nordisk, Denmark

## Faculty

<b>Peter Adamson</b>	The Children's Hospital of Philadelphia, United States
<b>Kate Beaujeux</b>	MedImmune, United Kingdom
<b>Gesine Bejeuhr</b>	vfa Research-based Pharmaceutical Companies, DIA Paediatric Group, Germany
<b>Gerlind Bode</b>	International Confederation of Childhood Cancer Parent Organizations (ICCCPO), Germany
<b>Christina Bucci-Rechtweg</b>	Novartis Pharmaceuticals, Switzerland
<b>Alexander Cvetkovich-Muntañola</b>	INC Research, Spain
<b>Emilie Desfontaine</b>	European Medicines Agency (EMA)
<b>Irmgard Eichler</b>	European Medicines Agency (EMA)
<b>Oliver Gross</b>	University Medicine Göttingen, Germany
<b>Ralf Herold</b>	European Medicines Agency (EMA)
<b>Peter Karolyi</b>	European Medicines Agency (EMA)
<b>Dirk Mentzer</b>	Paediatric Committee (PDCO), European Medicines Agency (EMA) & Paul Ehrlich Institut, Germany
<b>Koenraad Norga</b>	Paediatric Committee (PDCO), European Medicines Agency (EMA) & Antwerp University Hospital (UZA), Belgium
<b>Thorsten Olski</b>	European Medicines Agency (EMA)

<b>Andy Pearson</b>	Institute of Cancer Research, United Kingdom
<b>Solange Rohou</b>	AstraZeneca, United Kingdom
<b>Klaus Rose</b>	klausrose Consulting & EFGCP Children's Medicines Working Party, Switzerland
<b>Agnes Saint-Raymond</b>	European Medicines Agency (EMA)
<b>Mette Due Theilade Thomsen</b>	Novo Nordisk, Denmark
<b>Paolo Tomasi</b>	European Medicines Agency (EMA)
<b>Gilles Vassal</b>	Gustave Roussy, France
<b>Philip Walson</b>	Georg-August-University Medical School, Göttingen, Germany

## Conference Language

The language of the conference will be English.

## Conference Venue

### EMA Headquarters (NEW ADDRESS!)

Churchill Place 30, Canary Wharf  
London E14 5AB, United Kingdom  
Website: [www.ema.europa.eu](http://www.ema.europa.eu)

## Registration & Information

E-mail [conferences@efgcp.eu](mailto:conferences@efgcp.eu) or visit [www.efgcp.eu](http://www.efgcp.eu)

# Programme

Tuesday 30<sup>th</sup> September

08:00 Registration & Welcome Coffee

08:45 **Welcome**

*Emilie Desfontaine, European Medicines Agency (EMA) & Klaus Rose, klausrose Consulting & EFGCP Children's Medicines Working Party, Switzerland*

## SESSION 1 SETTING THE SCENE

Chairperson: *Emilie Desfontaine, European Medicines Agency (EMA)*

09:00 **Preparedness for the 10-year report and perspectives of the PDCO chairman**

*Dirk Mentzer, Chairman of the Paediatric Committee (PDCO), European Medicines Agency (EMA) & Paul Ehrlich Institut, Germany*

09:30 **The Drug Developers' Perspective**

*Christina Bucci-Rechtweg, Novartis Pharmaceuticals, Switzerland*

10:00 Discussion

10:30 Coffee Break

11:00 **Parallel break-out groups**

**Group 1 Pre-competitive paediatric collaboration between pharma companies and the role of EnprEMA?**

Chair: *Gesine Bejeuhr, vfa Research-based Pharmaceutical Companies, DIA Paediatric Group, Germany*

Introduction: *Irmgard Eichler, European Medicines Agency (EMA)*

**Group 2 Practical challenges of paediatrics clinical trials**

Chair: *Mette Due Theilade Thomsen, Novo Nordisk, Denmark*

Introduction: *Alexander Cvetkovich-Muntañola, INC Research, Spain*

**Group 3 Practical challenges with PIPs**

Chair: *Klaus Rose, klausrose Consulting & EFGCP Children's Medicines Working Party, Switzerland*

Introduction: *Kate Beaujeux, MedImmune, United Kingdom*

**Group 4 International framework for paediatric drug development**

Chair: *Emilie Desfontaine, European Medicines Agency (EMA)*

Introduction: *Agnes Saint-Raymond, European Medicines Agency (EMA)*

13:00 Lunch

## SESSION 2

### GROUP REPORTS AND FEEDBACK FROM EMA

Chairpersons: *Mette Due Theilade Thomsen, Novo Nordisk, Denmark & Klaus Rose, klausrose Consulting & EFGCP Children's Medicines Working Party, Switzerland*

- 14:00 **Feedback from the 4 parallel breakout groups**  
*Rapporteurs of the Groups*
- 15:00 **One stop-shop: a new EMA approach**  
*Paolo Tomasi, European Medicines Agency (EMA)*
- 15:30 **Report & Tentative answers on questions collected prior to the conference (send questions to [paediatrics@efgcp.eu](mailto:paediatrics@efgcp.eu)) / Update on paediatric procedures / PIP Format and content guideline revision**  
*Peter Karolyi, European Medicines Agency (EMA)*
- 16:00 **Panel & General Discussion**
- 16:50 **Conclusions**
- 17:00 **End of Day 1**
- 18:30 **Social Event**
- Key note speech: *Professor Philip Walson, Board Certified in Paediatrics, Clinical Pharmacology and Medical Toxicology; Visiting Professor, Department of Laboratory Medicine at Georg-August-University Medical School, Goettingen, Germany*

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## Wednesday 1<sup>st</sup> October

08:30 Welcome coffee

## SESSION 3

### ENHANCED COLLABORATION BETWEEN KEY PLAYERS

Chairpersons: *Ralf Herold, European Medicines Agency (EMA) & Gesine Bejeuhr, vfa Research-based Pharmaceutical Companies, DIA Paediatric Group, Germany*

- 09:00 **Paediatric Oncology: The EU Perspective**  
*Gilles Vassal, Gustave Roussy, France*
- 09:25 **Paediatric Oncology: The US Perspective**  
*Peter Adamson, The Children's Hospital of Philadelphia, United States*
- 09:50 **Paediatric Oncology: The UK ICR Perspective**  
*Andy Pearson, Institute of Cancer Research, United Kingdom*
- 10:15 **Paediatric Oncology: The Parent of Patient Perspective**  
*Gerlind Bode, International Confederation of Childhood Cancer Parent Organizations (ICCCPO), Germany*
- 10:40 Coffee Break

- 11:00 **EMA / PDCO Representative**  
*Koenraad Norga, Chairman of the Paediatric Committee (PDCO), European Medicines Agency (EMA) & Antwerp University Hospital (UZA), Belgium*
- 11:35 **Allport Syndrom: Challenges of clinical research and its translation into clinical practice**  
*Oliver Gross, University Medicine Göttingen, Germany*
- 12:00 **Panel Discussion**  
*Speakers of the session*
- 13:00 Lunch

## **SESSION 4**

### **INTERNATIONAL DEVELOPMENT IN PAEDIATRICS**

Chairperson: *Thorsten Olski, European Medicines Agency (EMA)*

- 14:00 **ICH E 11/ ICH E 6**  
*Dirk Mentzer, Paediatric Committee (PDCO), European Medicines Agency (EMA) & Paul Ehrlich Institut, Germany*
- 14:20 **ICH E 11/ ICH E 6**  
*Solange Rohou, AstraZeneca, United Kingdom*
- 14:40 **TTIP (Transatlantic Trade and Investment Partnership)**  
*Speaker invited*
- 15:00 **Panel & General Discussion**
- 15:50 **Conference Wrap Up**  
*Thorsten Olski, European Medicines Agency (EMA) & Mette Due Theilade Thomsen, Novo Nordisk, Denmark*
- 16:00 **End of Day 2**