# Working Group 3:

How to establish communication between Enpr-EMA networks and industry

# Working Group 5:

Sharing Good Practice within Enpr-EMA and with Industry Partners

Susan Tansey
Acting Chair WG 3 and WG 5

## Enpr-EMA WG 3

#### Role:

**Develop recommendations for how Enpr-EMA can:** 

- facilitate communication between industry and networks
- provide industry with easy access to information about capacities of individual Enpr-EMA networks
- increase the visibility of individual networks
- make contact with a range of industry partners (big Pharma, SMEs, biotechs, CROs etc.

## Enpr-EMA WG 5

### Role:

- Gather examples of network involvement in good practice for the development and implementation of clinical trials in children and young people
- Gather examples of good practice from industry colleagues working with Enpr-EMA networks
- Develop proposals to disseminate examples of good practice to Enpr-EMA members and industry partners.

## Enpr-EMA WG 3 and WG 5

Following discussion at the first meeting of WG 5 it was decided to propose merging the two WGs for the reasons below:

- many of the members of WG 5 are also in WG 3
- the remit was similar and it was felt that sending out a survey would be the best way to get some initial information on both examples of best practice (WG 5) but also on the current communication status between networks and industry (WG 3)

## Enpr-EMA WG 3 and WG 5

#### **Industry** members:

Ron Portman
Charles A Thompson
Matthew McClure
Martine Dehlinger-Kremer
Klaus Hartmann
Lynda Wight

#### **Network members:**

Susan Tansey (Acting Chair)
Pirkko Lepola (Steering group)
Jenny Preston (Steering group)
Pamela Dicks (Steering group
Nicola Ruperto
Andy Pearson
Christina Peters
Richard Trompeter

### Enpr-EMA WGs 3 and 5

Key Tasks, Timeframes and Outputs (1) Progress so far

Draft questions for a survey monkey style questionnaire to collect good practice examples from both network members and Industry colleagues. Planned date: end of September 2013

Actual date: 9th December 2013

Finalise survey monkey questionnaire and distribute.

Planned date: mid-October 2013 Actual date: 18<sup>th</sup> December 2013

Deadline for responses.

Planned date: end of November 2013

Actual date: 13th January 2014

Proposed extended date: 14th February 2014

### Enpr-EMA WGs 3 and 5

Key Tasks, Timeframes and Outputs (2)

Next steps

Organise teleconference to discuss and summarise responses.

Planned date: week commencing 2nd December 2013

Revised planned date: 3<sup>rd</sup>/4<sup>th</sup> March 2014

Send draft report of survey monkey to EnprEMA.

Planned date: week commencing 9th December 2013

Revised planned date: 30<sup>th</sup> March 2014

Develop a dissemination strategy to distribute and share good practice. Some ideas would include distributing via newsletters, EnprEMA website, Meeting Presentations, Webinars.

Planned date: January 2013

Revised planned date: April 2014

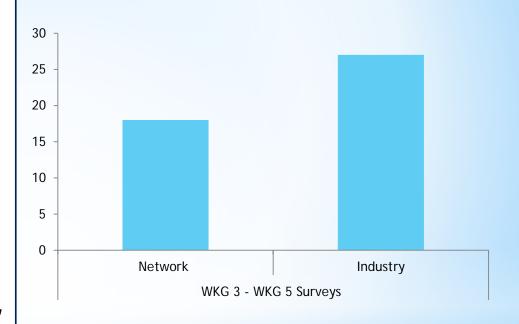
## Initial results of Industry and Network surveys

Responses received: Total 45

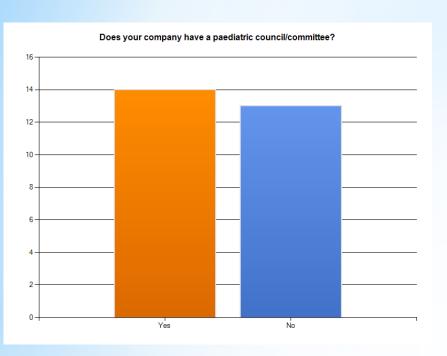
Industry responses (27):

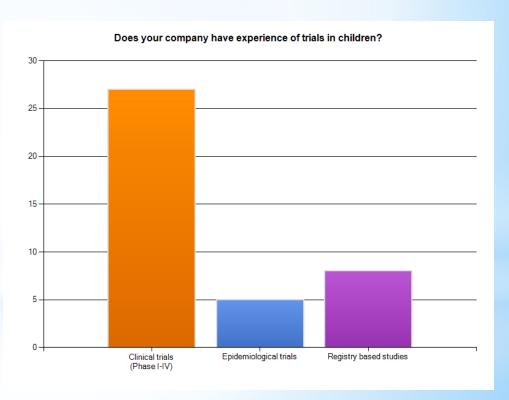
CROs: PRA, Quintiles, Covance, INC, Premier Research and Illingworth

Pharmaceutical/Biotechs: BMS (5), Astra Zeneca(3), Novartis (3), Janssen(2), Aegerion, Alios, Bayer (2), Kowa Research, Mitsubishi, Proveca and Vertex.

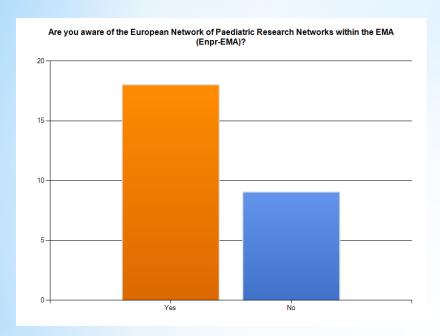


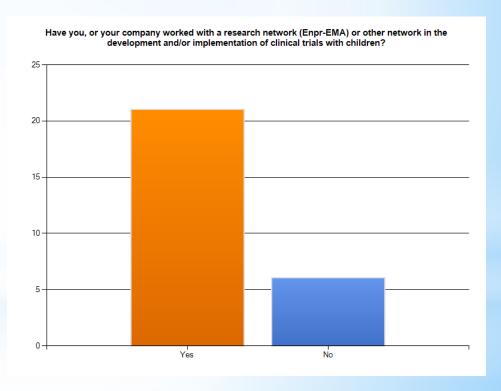
### Initial results of Industry survey

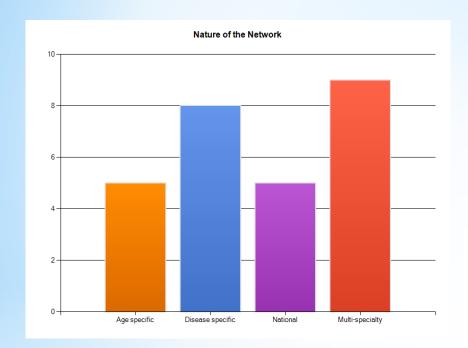


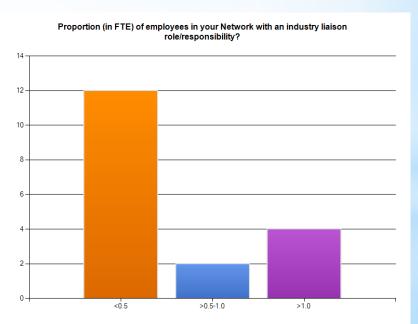


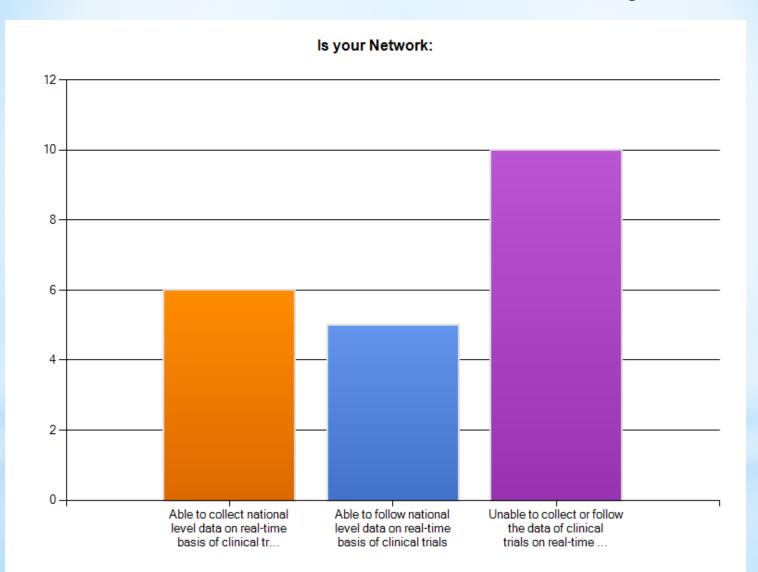
### Initial results of Industry survey

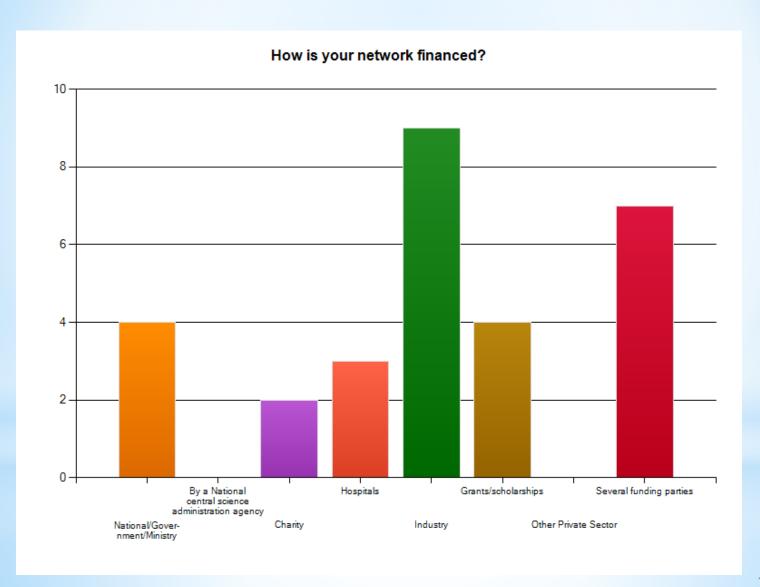


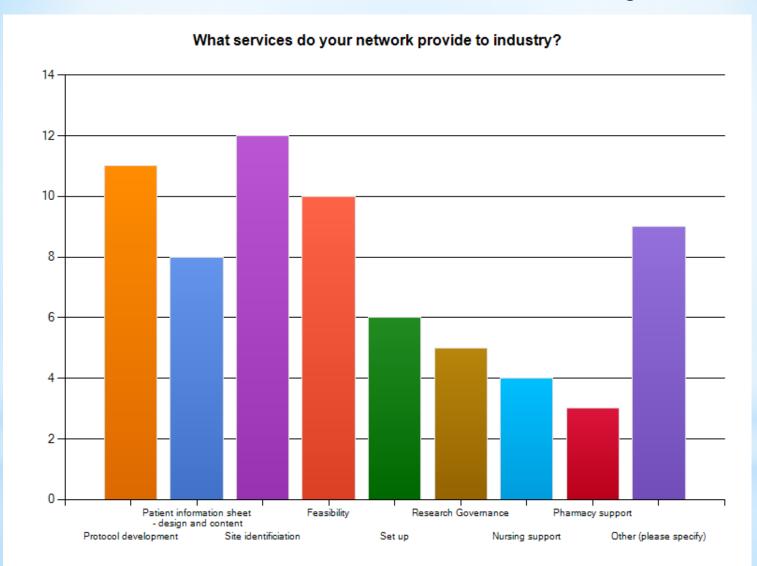












### Summary

### Next steps:

- Extend deadline for survey
- Full analysis of surveys
- Draft report to EnprEMA
- Dissemination strategy for best practice examples
- Communication strategy