

EUROPEAN PATIENTS' FORUM SPRING CONFERENCE Empowerment, Information, Sustainability

Renaissance Hotel, Brussels: 20/21 March, 2007.

Tuesday, 20th March, 2007.

12.30 to 14.00 Registration and Buffet Lunch

14.00 to 14.10 Welcome and outline of the Meeting Programme by Mr Anders Olauson, President, European Patients' Forum

14.10 to 14.30 Keynote Address by Mr Bernard Merkel, Health and Consumer Protection Directorate

The Empowered Patient: A benefit or a liability?

14.30 to 14.50 The Regulatory Perspective – Mr. Noel Wathion, The European Medicines Agency What role should the EMEA perform with regard to the provision of information on medicines?

14.50 to 15.10 The Physician's Perspective –Dr. Daniel Mart, President of CPME Should the healthcare profession be the gatekeeper of all information?

15.10 to 15.30 The Economic Perspective -Mr. David Mc Daid, London School of Economics Is there a financial case to be made for an informed patient?

15.30 to 15.50 Tea Break

European Patients' Forum, "Park Leopold" Rue Wiertz 50/28, B1050 Brussels Conference organizer's phone number: +32 497 19 38 99

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- **15.50 to 16.10** The Politician's Perspective Ms Dagmar Roth-Behrendt, MEP Can information be controlled by legislation nationally or internationally?
- 16.10 to 16.30 The Commission's Perspective Mr Martin Terberger, Head of Unit,

 Pharmaceuticals, Consumer Goods, Enterprise and Industry

 Directorate- General.

 By whom and to whom should information be provided?
- 16.30 to 16.50 How can we ensure the sustainability of patient organizations and who will be funding them in 2017? What part will corporate social responsibility play? Mr Richard Bergstrom, Director of Swedish Association of the Pharmaceutical Industry. Followed by Patient Perspective, Mr Anders Olauson.
- 16.50 to 17.30 General Discussion with Speakers comprising the panel
- **19.30 to 22.00** Conference Dinner, Renaissance Hotel



Wednesday, 21st March, 2007.

Break out into 3 Breakout Groups 09.15 to 12.45

It is suggested that each breakout group should last for 90 minutes with a coffee break at 10.45 until 11.15. Delegates may then attend another breakout group from 11.15 until 12.45.

GROUP A

Who really represents patients?

Patient organisations may be experts in their condition but rarely represent more then 2% of those affected by the condition. How can they speak for all patients? What is the role of consumer bodies? Are they legitimate representatives? How can increased solidarity be achieved across the European Patients' Movement? Should healthcare professionals be involved? Who validates the information provided by patient organisations?

| Facilitator: Ms Mel Read | Rapporteur: Mr Mike O'Donovan |
|--|-------------------------------|
| Member of European Parliament 1989-2004 | Treasurer, European Patients' |
| President, European Cervical Cancer Assoc. | Forum |
| Honorary Chair, Health First Europe | |
| Member of GlaxoSmithKline Health Advisory | |
| Board | |

GROUP B

Is ignorance bliss or is an informed patient a sound economic investment?

What is the evidence that an informed and educated patient uses healthcare services more effectively and more efficiently? Or does the more informed and empowered patient cost more by demanding the latest and most costly drugs and treatments? Should health be viewed as an investment or a cost?

| Facilitator: Mr Jorgo Chatzimarkakis | Rapporteur: Mr Jean Georges |
|--------------------------------------|-------------------------------------|
| Member of the European Parliament | The Board, European Patients' Forum |

GROUP C

Are patient organisations viable and sustainable? If so, by whom?

Many national and supranational patient organisations have become highly effective in their respective disease areas. But many are extremely fragile financially. How sustainable are they in the medium to long term? Competition is fierce for the finite funding available. How realistic is it for patient groups to devote such many resources



to fundraising to ensure their continued existence? Is this a good and legitimate use of such resources? Is there too much duplication within a disease area resulting in wastage? What may be classified as legitimate funding sources? What role should the pharmaceutical industry continue to play? Who else would provide funding? What should be the Commission's position?

| Facilitator: Ms Celine van Doosselaere | Rapporteur: Mr Rodney Elgie |
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| European Health Management Assoc | GAMIAN Europe |

12.45 to 14.00 Lunch

Plenary Session: How do we manage the future?

14.00 to 14.30 Feedback from each Rapporteur

14.30 to 14.55 The prevailing environment in 2017. How well informed will patients be ten years from now and from what sources will information be available?
<u>Dr. Ilona Kickbusch, Senior Health Policy Advisor and WHO Consultant.</u>

14.55 to 15.20 The Patients' Perspective

Ms Susanna Palkonen, European Federation of
Allergy and Airways Disease Patients' Associations

15.20 to 16.00 General facilitated discussion

16.00 to 16.15 Conclusions and close of the meeting

Mr Anders Olauson, President European Patients'Forum

This conference has been made possible thanks to an unrestricted educational grant from GlaxoSmithKline and Pfizer Public Affairs Europe.