

Clinical Trials Directive Re-opening: Concrete orientations to increase the European biomedical research competitiveness for the benefit of patients

An exchange of views to prepare the forthcoming reopening of the Clinical Trials Directive 2001/20/CE

**European Parliament, June 22, 2011 – Room: “Salon des Membres”
12.30 – 14.00**

Chair: Pr. Philippe JUVIN, MD, PhD

MEP, Group of the European People's Party

Member of the Committee on the Environment, Public Health and Food Safety

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| 12:15 | Registration |
| 12:30 – 12:35 | Welcome by Pr. Philippe Juvin, MD, PhD, MEP |
| 12:35 – 12:40 | Consultation state of play by Andrzej Rys, Director Health systems and products, DG SANCO |
| 12:40 – 13:00 | How to better harmonise/rationalise the technico scientific administrative process of Clinical Trials? |
| 13:00 – 13:20 | How to adapt the regulatory oversight to a clinical Trials risk based approach? |
| 13:20 – 13:40 | How to facilitate access to information and capacity building for patients? |
| 13:40 – 14:00 | Conclusions & Next Steps by Pr. Philippe Juvin |

participating organisations include:



*lunch is kindly offered by sanofi