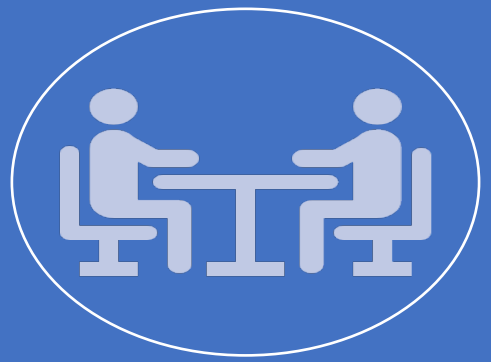
efpia
European Federation of Pharmaceutical Industries and Associations

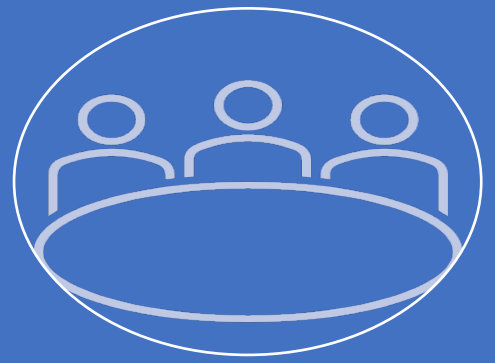
ACT-EU MSP workshop – 22-23 June

Session 2: The Role of Ethics Committees in Clinical Trials

How can medical research ethics committees further contribute to maintaining the EU as an attractive and competitive region for conducting scientific and ethically sound clinical trials?



Providing a space for discussions about ethical issues to take place to encourage best practise sharing.



Facilitating a dialogue including ethics committees and Health Authorities to optimise collaboration

Facilitating common standards and expectations for ethical review including ensuring no overlap with Health Authority assessments. This would include a common understanding of information required for the ethical review, for example what information is required for an eConsent review?

Industry can support through training on innovative approaches used in trials