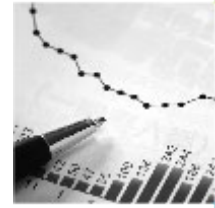


Identifying Clinical Trials and Trial Results for Systematic Reviews & Other Evidence Syntheses: Making the Most of Trials Registers, Regulatory Agency Sources & Other Novel Resources

Session 1

TIMETABLE

- Trainers:** Julie Glanville, Independent Consultant in Information Retrieval, Glanville.info, York, UK
Carol Lefebvre, Independent Information Consultant, Lefebvre Associates Ltd, Oxford, UK
- 15.00** Welcome and introduction to the day
- 15.05** Why do we need to know about clinical trials and ongoing research? (**Julie**)
- 15.25** What are trials registers? Background and brief history (**Carol**)
- 16.00** Key trials and trials results registers (**Julie**):
- Focus on ClinicalTrials.gov, the WHO ICTRP and the presenters' trial register website
- 17.00** Break (5 minutes)
- 17.05** Questions / Discussion
- 17.30** Close



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Session 2

TIMETABLE

- Trainers:** Julie Glanville, Independent Consultant in Information Retrieval, Glanville.info, York, UK
Carol Lefebvre, Independent Information Consultant, Lefebvre Associates Ltd, Oxford, UK
- 15.00** Clinical study reports and obtaining trial information from regulatory agencies: the (European) EMA and the (US) FDA (**Carol**)
- 15.45** Other resources to identify unpublished trial data: preprints, HTA sources and other trial initiatives such as Crossmark (**Julie**)
- 16.15** Record management issues: downloading results (**Julie**)
- 16.35** Record management issues: documenting and reporting searches (**Carol**)
- 17.00** Break (5 minutes)
- 17.05** Questions / Discussion
- 17.30** Close