YHEC
York Health Economics Consortium

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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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15.45	Other resources to identify unpublished trial data: preprints, HTA sources and other trial initiatives such as Crossmark **(Julie)**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