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 **(Julie)**
15.05	Why do we need to know about clinical trials and ongoing research? **(Julie)**
15.25	What are trial registers? Background and brief history **(Carol)**
16.00	Key trials and trials results registers part 1 **(Julie):** ▪ Focus on ClinicalTrials.gov
16.30	Break
16.40	Key trials and trials results registers part 2 **(Julie):** ▪ the WHO ICTRP, CENTRAL and other resources
17.10	Questions / Discussion
17.30	Close

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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15.45	Other resources to identify unpublished trial data: preprints, HTA sources and other trial initiatives such as Crossmark **(Julie)**
16.30	Break
16.40	Record management issues: downloading results **(Julie)**
17.00	Record management issues: documenting and reporting searches **(Carol)**
17.25	Questions / Discussion
17.45	Close