ClinicalTrials.gov: Results Reporting, Unique Evidence, and the Role of the Medical Librarian

Training materials:

• ClinicalTrials.gov How to Read a Study Record [https://clinicaltrials.gov/ct2/help/how-read-study]
• ClinicalTrials.gov train the trainer materials [https://clinicaltrials.gov/ct2/manage-recs/present#ResultsTrainTrainer]
• ClinicalTrials.gov additional resources [https://clinicaltrials.gov/ct2/manage-recs/resources]

Additional ClinicalTrials.gov Resources:

• Final Rule Information [https://prsinfo.clinicaltrials.gov]
  • Final Rule Webinar Series
  • Applicable Clinical Trial Checklist and Elaboration (ACT Checklist)
  • Frequently Asked Questions
  • Planned Updates to PRS
  • “Coming Soon”

• Submit Studies [https://clinicaltrials.gov/ct2/manage-recs]
  • FDAAA 801 Requirements
  • Frequently Asked Questions
  • Training Materials – recorded presentations, example study records
  • Support Materials

Links to the readings, articles and tutorials referenced in today’s webinar:


Gøtzsche PC. Why we need easy access to all data from all clinical trials and how to accomplish it. Trials. 2011 Nov 23;12:249. doi: 10.1186/1745-6215-12-249.


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