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Public reporting burden for this collection of information is estimated to average 5 minutes per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing the collection of information. An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a current valid OMB control number. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden to NIH Project Clearance Branch, 6705 Rockledge Drive, MSC 7974, Bethesda, MD 20892-7974, ATTN: PRA (0925-0648). Do not return the completed form to this address.

ClinicalTrials.gov Train-the-Trainer Feedback Survey

Please rate the usefulness of the following educational formats for the workshop:

	Very Useful (1)	Somewhat Useful (2)	Neither Useful nor Useless (3)	Not Useful (4)	Useless (5)	I don't know (6)
Lecture (1)	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
Interactive Discussion (2)	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
Hands-on Time (3)	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>

Comments/clarifications about the format of the workshop:

How appropriate was the length of the workshop?

- Too long (1)
- Just right (2)
- Not long enough (3)

Comments/clarifications about the length of the workshop:

How much novel information and/or skills did you learn about each section that you did not know before?

	A lot of new info/skills (1)	Some new info/skills (2)	No new info/skills, I already knew this (3)	I don't know (4)
Overview of the Clinical Trial Disclosure Landscape (1)	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
NIH Office of Extramural Research (OER) Perspective (2)	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
Managing Registration and Results Reporting at and Academic Institution (3)	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
Participant Flow Module (4)	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
Baseline Characteristics Module (5)	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
Protocol Registration and Results System Overview (6)	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
PRS Guided Tutorials Overview (7)	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
Outcome Measures Module (8)	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
Adverse Events Module (9)	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>

Comments/clarifications about what you learned:

Comments/clarifications about your confidence in helping colleagues with their questions:

What did you like most or find most helpful about the Train-the-Trainer Workshop?

What did you like least or find least helpful about the Train-the-Trainer Workshop? How could the Workshop improve?

Do you have any questions or concerns that were not addressed as a part of the Train-the-Trainer Workshop? How could they be better addressed?

What additional resources and information would be useful to you, your institution, or your colleagues for reporting to ClinicalTrials.gov
