

# Tip of the Month



## Documentation of Protocol Training

Prior to any research-related procedures being initiated, initial protocol-specific training must be completed and documented for all research personnel listed on the Delegation of Authority (DOA).

Initial protocol-specific training will either be conducted by the Sponsor or Principal Investigator (PI), dependent on the research trial. Subsequent training for research personnel must be conducted by the PI, sub-investigator, clinical research coordinator or other appropriate research personnel who were in attendance at the initial protocol-specific training, or has otherwise been previously trained.

Protocol amendments may require additional training depending on the scope and nature of the changes.

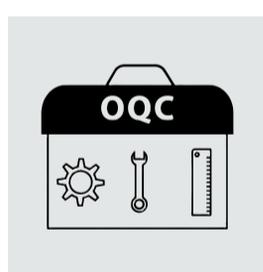
Re-training of research personnel should occur if the protocol is amended with operational aspects that affect study procedures such as:

- Eligibility criteria
- Treatment parameters
- Changes to cohorts or study arms
- Safety or efficacy assessments
- Data collection and reporting



[Training Log Template](#)

[UUSOP-03: Protocol Training](#)



### OQC Toolkit

Explore templates used to assist research teams with conducting, tracking and/or monitoring research activities. Templates are available for research teams to download and adapt for their own use.

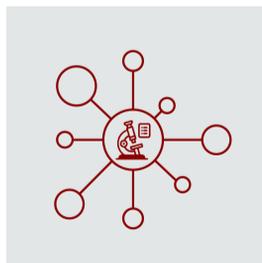
[OQC TOOLKIT](#)



### Tip of the Month Archive

Explore previous research compliance reminders and information.

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### Research Quality Compliance Network

A network for the research community within the University. Quarterly events are held on compliance and research related topics.

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### Contact Us

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