

SWOG Lead Oncology Research Professional (ORP) Training:

[SWOG Lead ORP Workshop](#) - Enroll to the entire workshop or the individual course components via the links below. (Login at <https://swog.exphosted.com/> prior to clicking on the course links.)

Introductory Modules

- [SWOG Lead Oncology Research Professional Workshop Introduction](#) (10 mins)
- [An Introduction to SWOG and the NCI](#) (45 mins)

Leadership and Site Operations Modules

- [Leadership Perspective](#) (17 mins)
- [NCI Trials - Site Operations Perspective](#) (NEW / forthcoming)

SWOG and NCI Systems Module

- [SWOG and NCI Systems Overview Training](#) (14 mins)

Data Management and Resource Access Modules

- [Data Management and Access Module Introduction for Lead ORPs](#) (1 min)
- [CTSUS Website Overview](#) (9 mins)
- [SWOG Website Overview](#) (4 mins)
- [SWOG ORP \(CRA\) Workbench Overview](#) (4 mins)
- [iMedidata RAVE Access for Lead ORPs](#) (6 mins)
- [SWOG Specimen Tracking System for Lead ORPs](#) (8 mins)
- [Central Monitoring](#) (5 mins)

Funding and Site Payments Modules

- [NCTN and NCORP Study Funding and Payment Distribution](#) (20 mins)
- [Open Funding](#) (10 mins)
- [National Coverage Analysis Overview](#) (4 mins)

Study Activation and Management Modules

- [Delegation of Tasks Log](#) (2 mins)
- [Adverse Events Training for Lead ORPs](#) (8 mins)
- [Dose Modifications Training for Lead ORPs](#) (1 min)
- [Serious Adverse Event Reporting Training for Lead ORPs](#) (30 mins)
- [SAE Reporting - Specific Exceptions to Expedited Reporting for Lead ORPs](#) (4 mins)
- [Data Entry in Difficult Situations Training for Lead ORPs](#) (2 mins)
- [Record Retention Training for Lead ORPs](#) (3 mins)

Reports and Tools for Data Quality Modules

- [Expectations and Expectation Reports for Lead ORPs](#) (10 mins)
- [Vital Status Expectations Training for Lead ORPs](#) (5 mins)
- [Specimen Expectations for Lead ORPs](#) (8 mins)
- [Institution Performance Review \(IPR\) for Lead ORPs](#) (10 mins)
- [Query Reports Training for Lead ORPs](#) (4 mins)
- [CTSUS reports - Data Quality Portal \(DQP\) Training for Lead ORPs](#) (7 mins)

Regulatory Module

- [Regulatory Expectations from a QA Perspective - Training for Lead ORPs](#) (13 mins)

Quality Assurance Modules

- [Audits and Quality Assurance Program - Training for Lead ORPs](#) (20 mins)
- [Protocol Deviations vs. Deficiencies Training for Lead ORPs](#) (4 mins)
- [When is my Institution's next Audit Due? - Training for Lead ORPs](#) (2 mins)

Policy 50 Description and Responsibilities of a SWOG Lead Oncology Research Professional

Additional Training Resources:

- [SWOG Regulatory Workshop](https://swog.exphosted.com/) (Login at <https://swog.exphosted.com/>)
 - [Session 1: Introduction](#)
 - [Session 2: Regulatory 101 Panel Discussion](#)
 - [Session 3: Understanding the PMB](#)
 - [Session 4: Know Your IRBs Panel Discussion](#)
 - [Session 5: SWOG Perspectives and Common Challenges Panel Discussion](#)
 - [Session 6: Past Accomplishments & Hope for the Future](#)
- [SWOG Advanced Practice Provider Clinical Research Workshop](https://classlms.org/) (Login at <https://classlms.org/>)
- [\(Re\)Igniting Leadership](https://swog.exphosted.com/) (Login at <https://swog.exphosted.com/>)
- [Compiled Researcher Resources Spreadsheet](https://ctsu.cancer.gov/) (login at <https://ctsu.cancer.gov/>)

Helpful Webpages:

- [Lead ORP Workbench | SWOG](#)
- [SWOG Protocol Development Tracking Reports and Dashboards](#) (login at [SWOG.org](https://swog.org))
- [SWOG ORP \(CRA\) Workbench](#) (login at [ORP Workbench](#))
 - [SWOG CRA Manual for Oncology Research Professionals](#) (login at [ORP Workbench](#))
 - [Current ORP \(CRA\) Newsletter & Newsletter Archive](#) login at [ORP Workbench](#)
- [SWOG Quality Assurance & Audits](#), includes essential guidelines.
 - [SWOG Best Practices Document](#)
 - [SWOG Quality Assurance Audit Guidelines](#)
 - [SWOG Regulatory Guidance](#)
 - [NCI-CTMB Guidelines for Monitoring of Clinical Trials](#)
 - [Serious Adverse Events | SWOG](#)
 - [NCI Guidelines for Investigators: Adverse Event Reporting Requirements for DCTD \(CTEP and CIP\) INDs and IDEs](#)
- [Biospecimen Collection and Submission Procedures | SWOG](#)
- [SWOG Oncology Research Professionals](#) (login at [SWOG.org](https://swog.org))
- [Frequently Asked Questions | SWOG](#)
- [Clinical Research Resources | SWOG](#)
- [SWOG Member Directory | SWOG](#) (login at [SWOG.org](https://swog.org))
- [SWOG Publications Database | SWOG](#)
- [SWOG Cancer Research Network Meetings | SWOG](#)
- [The Hope Foundation Funding Opportunities - SWOG Cancer Research](#)

Feasibility Assessment Tool: [Clinical Trial Review Guide](#) (login at [SWOG.org](https://swog.org))

Resources provided by NCI Cancer Trials Support Unit (CTSU):

- [NCI CTSU Researcher Resources](https://ctsu.cancer.gov/) (login at <https://ctsu.cancer.gov/>)
 - [NCI CTSU Glossary and Acronyms](https://ctsu.cancer.gov/) (login at <https://ctsu.cancer.gov/>)

NCI Pharmaceutical Management Branch (PMB) Agent Management Application:

- [NCI PMB, CTEP Inventory Management System: AURORA Application Training](#)
- [NCI CTEP AURORA Document Access Training Course](#)

NCI CIRB Resources for Institutions: [Institutions | NCICIRB](#)