

SWOG

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Policy Memorandum No. 46
Subject: Job Description of Principal Investigator
Departments Affected: All

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JOB DESCRIPTION OF A PRINCIPAL INVESTIGATOR

General Responsibilities

The Principal Investigator (PI) will assume responsibility for the overall scope of his/her institutional program, including, but not limited to, ensuring that the institution meets the procedures and standards set forth by SWOG, notifying SWOG of important changes in member relationships, agreeing to quality assurance audits, and communicating all policies (and any changes in policy) on the conduct of clinical research to clinical researchers and staff members. The PI is further charged with promoting SWOG activities within the institution, ensuring investigators and trainees are provided the opportunity to learn about SWOG and participate in SWOG activities, as well as identifying institutional research that could be translated to cooperative group science and assisting in bringing those ideas forth within SWOG.

Qualifications

The Principal Investigator must be a licensed physician and be qualified to personally supervise all aspects of clinical trial conduct. Principal Investigators must have a minimum of three years clinical faculty experience in conducting clinical trials. The Principal Investigator should possess effective communication skills and the ability to work as part of a multidisciplinary team to provide leadership in the development of the institution's program both scientifically (protocol development and coordination, publications, committee leadership and participation) and by accrual contributions.

If a Principal Investigator is designated by an institution that does not meet the above minimum requirements, it is recommended that a senior Co-Principal Investigator (at least at an Associate Professor or equivalent level) be designated to provide additional support and mentorship.

Specific Responsibilities

1. Understands and upholds the policies and procedures of the Group and assist with, and participates in communications as necessary with all clinical researchers and staff members in ensuring the same.
2. Ensures that specialists from relevant oncology disciplines, as well as appropriate data management and nursing resources, are available within the institution to support the activities of SWOG.
3. Regularly monitors institutional accrual and ensures the institution meets minimum accrual goals set forth by SWOG to maintain membership.
4. Monitors performance of affiliate sites and provides support to those sites as needed.
5. Directs the use of funds received by SWOG to support SWOG activities.

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6. Assumes patient follow-up responsibility from investigators at his/her institution or affiliates in accordance with Group Policy No. 30.
7. Principal Investigator is expected to be available during the exit interview for a recap of the audit findings and to provide clarification of issues raised by the auditors.
8. Ensures that deficiencies in institutional performance reviews and quality assurance audits are addressed and corrected.
9. Approves all new investigator membership nominations at his/her institution and affiliates.
10. Appoints individuals from his/her institution as authors on Group publications in accordance with Group Policy No. 24, as requested.
11. It is required that the Principal Investigator attends at least one semi-annual Group meeting every two years.
12. All Principal Investigators from Member institutions in good standing will serve as a member of the Board of Governors.
13. All Principal Investigators from the Community Clinical Oncology Program (CCOP) that choose SWOG as their primary research base will serve as a member of the Board of Governors.
14. When resigning as the Principal Investigator, the Principal Investigator is responsible for ensuring that the Group Chair is informed of this decision and that a new Principal Investigator is named.