For Patients who are Smokers with any Type of Cancer

EAQ171CD Available Through ECOG-ACRIN Cancer Research Group

Implementing a Virtual Tobacco Treatment in Community Oncology Practices: “Smoke Free Support Study 2.0”

Patient Population
See Section 3.0 for Complete Details for Participant Selection

Selection of NCORP Site Staff Participants: The CCDR Leader and site PI from each participating site will identify ~10 multidisciplinary staff members to complete brief surveys (~15 minutes) and participate in focus group interviews (~45-60 minutes; conducted ~24 months post site activation/after 23 month survey). Site staff may vary, but is envisioned as the CCDR Leader, site PI, 2-3 oncology nurses, 2-3 medical oncologists, and 2-3 additional staff members
- Eligible NCORP site staff will be English-speaking and employed at the NCORP site for at least 3 months and able to provide informed consent to participate (see protocol for additional details)

Selection of Patients:
Eligibility Step 0 and 1:
- Age ≥ 18 years
- Must present with any type of cancer with a date of diagnosis within the past 4 months (recurrence, diagnosed within the last 4 months, of tumors in patients with past cancer diagnoses/patients with a new primary cancer, diagnosed within the last 4 months, who have been treated previously for other types of cancer/in situ cancers, diagnosed within the past 4 months, will all be considered eligible)
- Must be a current smoker defined per protocol
- Must be fluent in both written and spoken English/ written and spoken Spanish
- Must have telephone, web, and email access
- Patients with an ECOG PS of 3 or above, or deemed medically unable to participate by investigators/oncology clinicians are excluded
- Patients with no intention of receiving their cancer care/monitoring at an NCORP site are excluded

Eligibility Step 2 (Randomization):
- Must have completed Baseline Survey (EASEE-PRO) within 30 days of the informed consent date (Step 1)

Treatment Plan
See Section 5.0 for Complete Methodology Details

- Site staff to review new patient lists/schedules to identify all patients who are adult, newly diagnosed, and who are potential current smokers (per protocol)
  - Patients meeting these criteria should be entered into the study-specific screening log in REDCap, which will assign the next available unique screening ID code (sociodemographics and eligibility criteria should be entered in the screening log)
- Then, eligible patients are approached about the study, and eligibility is confirmed
- Patient must verbally agree to be contacted by staff to view an informational video about the study; then, the patient will be registered in OPEN (Screening Step 0)
- Once the patient’s information is in OPEN, the informational video will be sent to the email address provided by EASEE-PRO
- Site staff will then follow-up with all patients who received the informational video to assess their interest in participating in the study (it is recommended site staff make at least 3 outreach attempts)
- Interested, eligible patients will be given an IC; once the patient has signed the IC, they will be registered to OPEN Step 1 and will be prompted to complete the Baseline Survey in EASEE-PRO
- After completion of the survey, the patient will be eligible to register to Step 2, which will randomize them to a treatment arm (see protocol for details):
  - **Arm A- Enhanced Usual Care:** reflects the NCCN Smoking Cessation Guidelines (i.e., assessment of smoking status, provision of quitting advice, referral the NCI Smokers’ Quitline)
- **Arm B- Virtual Intervention Treatment:** offered up to 11 counseling sessions over approximately 6 months and up to 12 weeks of free combination Nicotine Replacement Therapy (see protocol for details/counseling topics)

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Please Enroll Your Eligible Patients!
Note: Please refer to the protocol for NCORP Site Staff Schema.