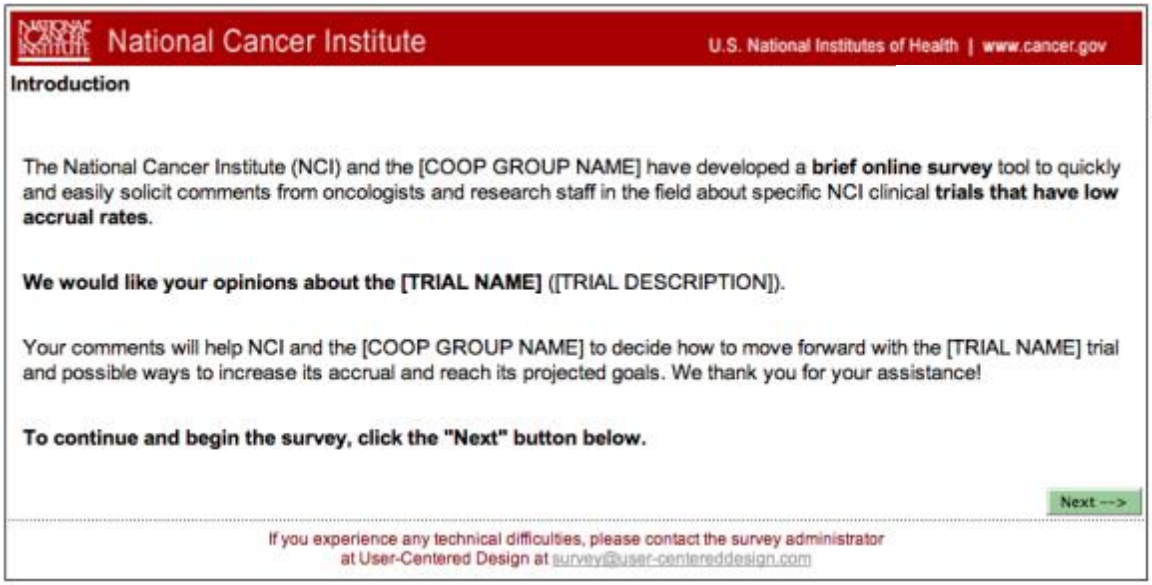
OMB #0925-xxxx
Expiration Date: xx/xx/xxxx



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Introduction

The National Cancer Institute (NCI) and the [COOP GROUP NAME] have developed a **brief online survey** tool to quickly and easily solicit comments from oncologists and research staff in the field about specific NCI clinical **trials that have low accrual rates**.

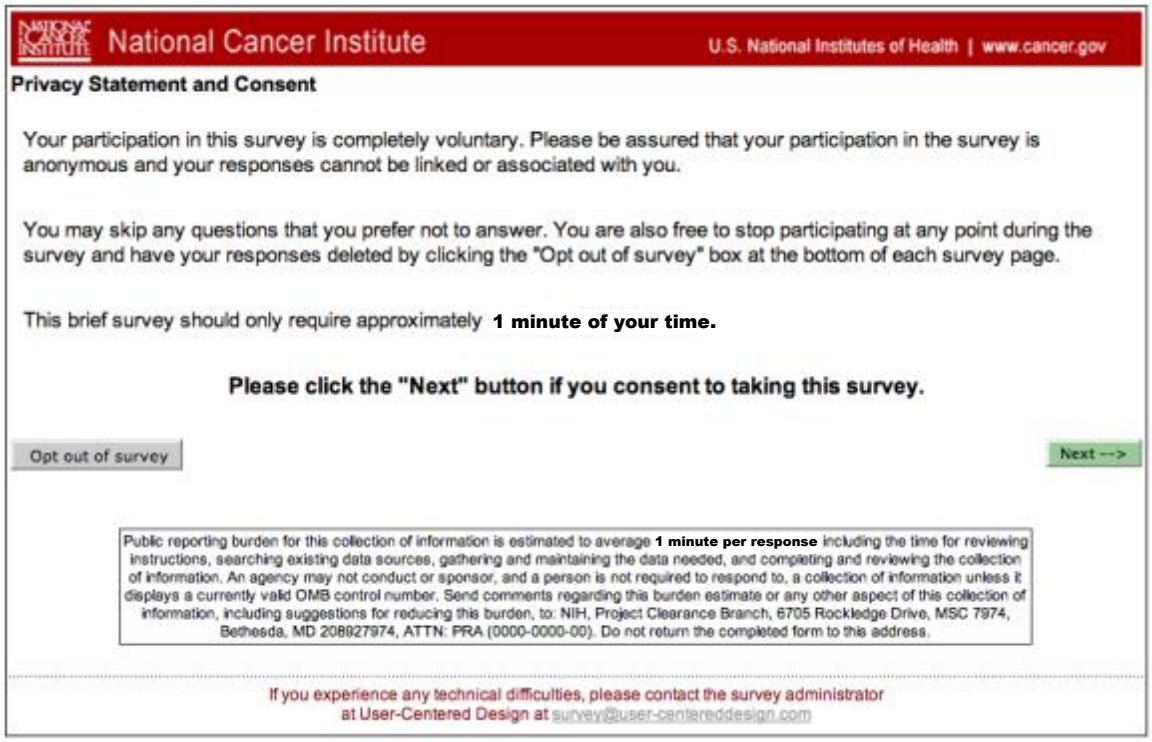
We would like your opinions about the [TRIAL NAME] ([TRIAL DESCRIPTION]).

Your comments will help NCI and the [COOP GROUP NAME] to decide how to move forward with the [TRIAL NAME] trial and possible ways to increase its accrual and reach its projected goals. We thank you for your assistance!

To continue and begin the survey, click the "Next" button below.

Next -->

If you experience any technical difficulties, please contact the survey administrator at User-Centered Design at survey@user-centereddesign.com



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Privacy Statement and Consent

Your participation in this survey is completely voluntary. Please be assured that your participation in the survey is anonymous and your responses cannot be linked or associated with you.

You may skip any questions that you prefer not to answer. You are also free to stop participating at any point during the survey and have your responses deleted by clicking the "Opt out of survey" box at the bottom of each survey page.

This brief survey should only require approximately **1 minute of your time.**

Please click the "Next" button if you consent to taking this survey.

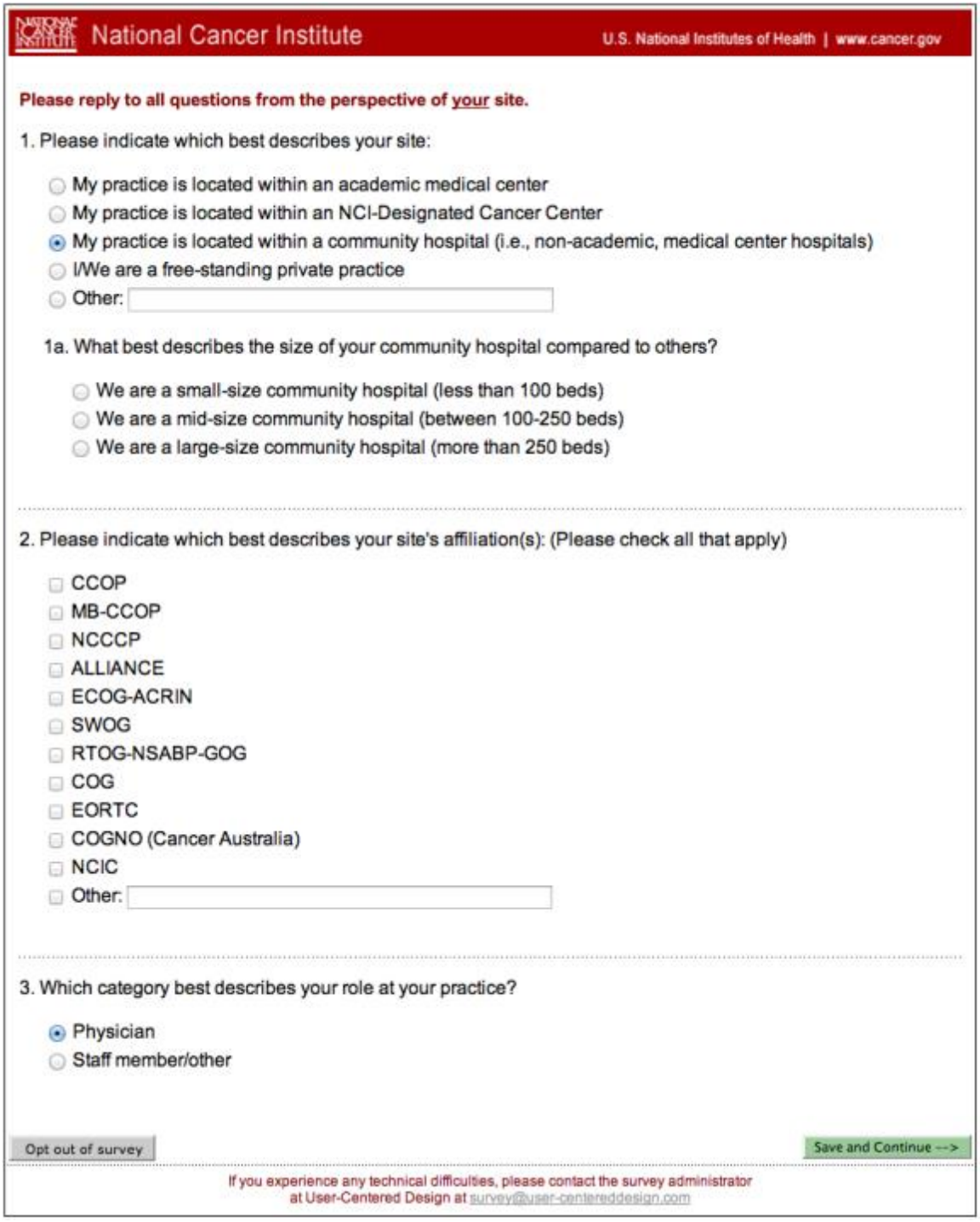
Opt out of survey

Next -->

Public reporting burden for this collection of information is estimated to average **1 minute 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 currently 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 208927974, ATTN: PRA (0000-0000-00). Do not return the completed form to this address.

If you experience any technical difficulties, please contact the survey administrator at User-Centered Design at survey@user-centereddesign.com

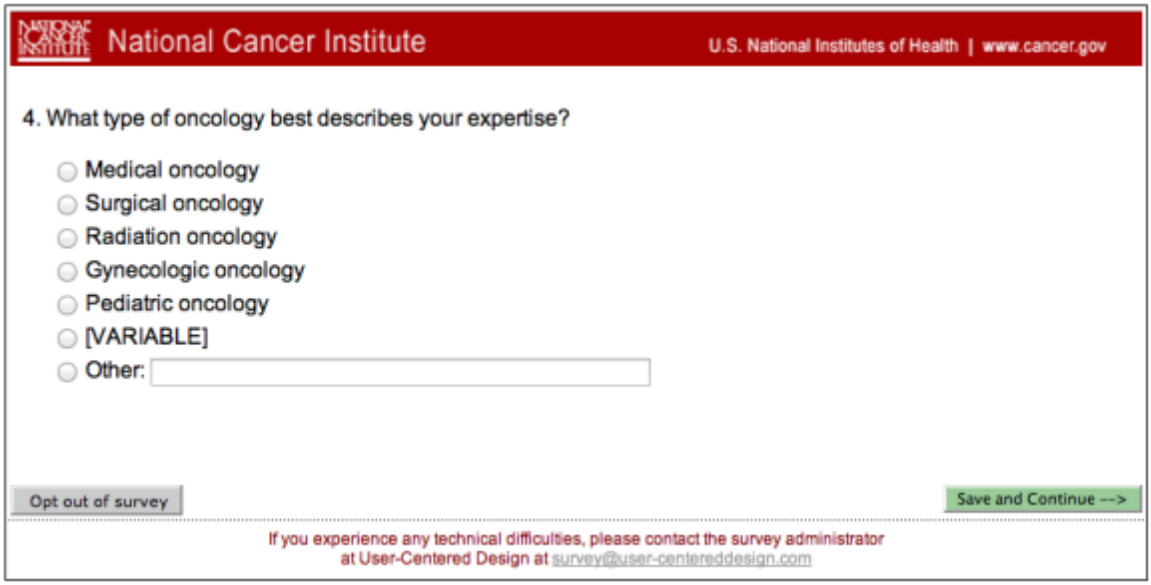
Attachment_C11_lowaccruingsurvey



Note: Q1a appears only if "My practice is located within a community hospital" is selected for Q1.

If user selects "physician" to Q3, continue below. If user selects "Staff member/other", continue HERE.

Attachment_C11_lowaccruingsurvey



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5. The [TRIAL NAME] trial has [INSERT #] research question[s]. [For each,] Please tell us how scientifically interesting it is to you.

RQ1: [INSERT RQ1 FROM CONCEPT SHEET]

5a1. How scientifically interesting is this research question to you?

Not at all interesting				Very interesting
○	○	○	○	○

5a2. Which statement best matches your opinion?

- ○ RQ1 is more interesting to me than when it first opened.
- ○ RQ1 is less interesting to me than when it first opened.
- ○ RQ1 is about the same level interest to me as when it first opened.

Please elaborate:

RQ2: [INSERT RQ2 FROM CONCEPT SHEET]

5b1. How scientifically interesting is this research question to you?

Not at all interesting				Very interesting
○	○	○	○	○

5b2. Which statement best matches your opinion?

- ○ RQ2 is more interesting to me than when it first opened.
- ○ RQ2 is less interesting to me than when it first opened.
- ○ RQ2 is about the same level interest to me as when it first opened.

Please elaborate:

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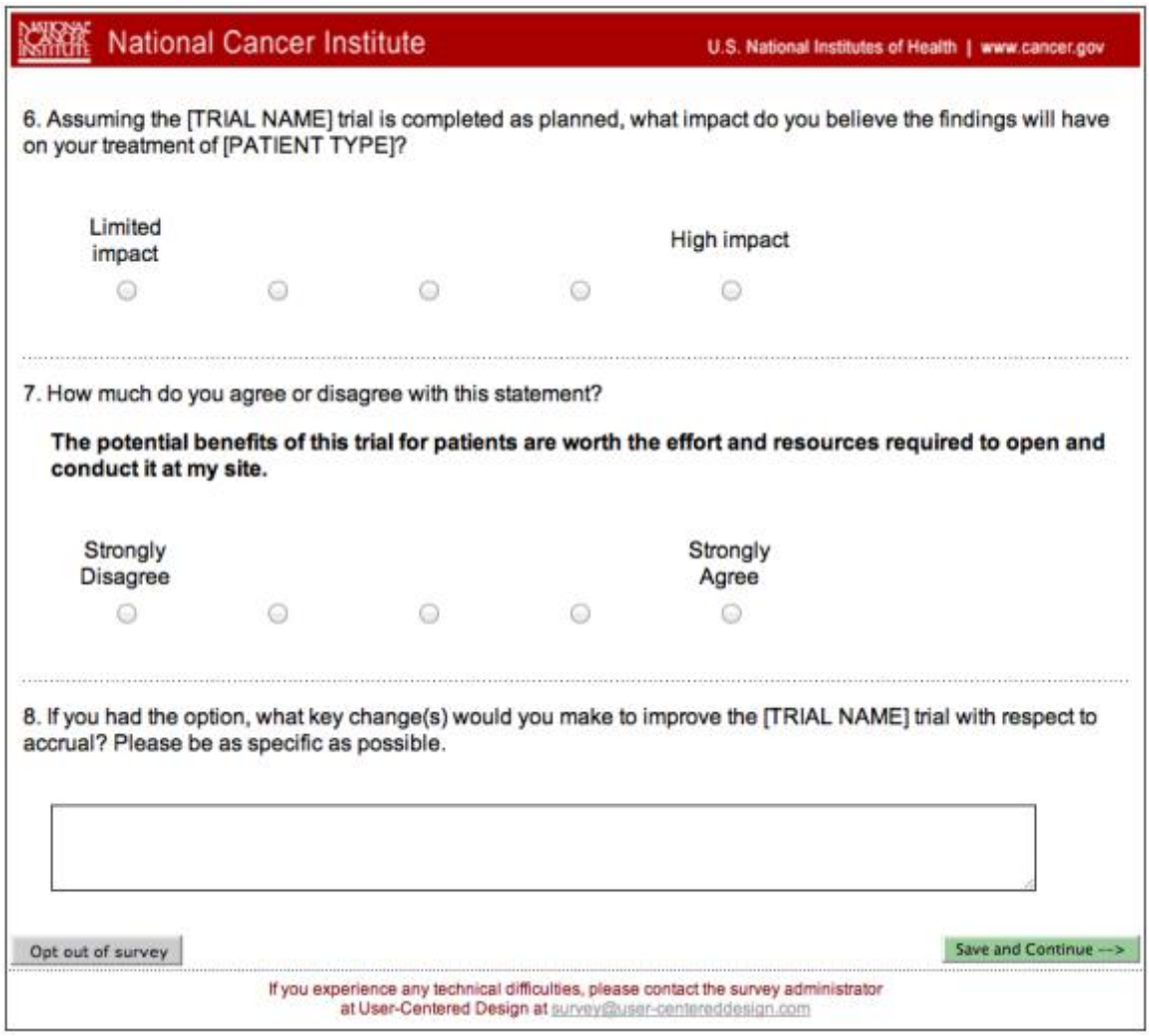

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9. For each of the trial's requirements listed below, please tell us whether or not it does (or would) make it difficult to open or run the [TRIAL NAME] trial at your site.

How difficult does (or would) this requirement make it to open or run this trial at your site?	Not make it difficult to open/ run the trial	Make it somewhat difficult to open/ run the trial	Make it very difficult to open/run the trial
[INSERT ITEM 1 FROM TRIAL SUMMARY SHEET]	○	○	○
[INSERT ITEM 2 FROM TRIAL SUMMARY SHEET]	○	○	○
[INSERT ITEM 3 FROM TRIAL SUMMARY SHEET]	○	○	○
[INSERT ITEM N FROM TRIAL SUMMARY SHEET]	○	○	○

Opt out of survey | Save and Continue -->

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10. How often do you see a patient who may be eligible for the [TRIAL NAME] trial (i.e., [PATIENT TYPE])?

- ○ Daily
- ○ Weekly
- ○ Monthly
- ○ Every few months
- ○ A couple times per year or less

11. Which specialty at your institution would most likely be in the position to recommend or refer this trial to a potentially eligible patient?

- ○ Medical oncology
- ○ Surgical oncology
- ○ Radiation oncology
- ○ Gynecologic oncology
- ○ Pediatric oncology
- ○ [VARIABLE]
- ○ Other (please specify): _____

Opt out of survey | Save and Continue -->

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12. For each of the items below, please tell us whether or not it does (or would) make it difficult to accrue patients to the [TRIAL NAME] trial.

How difficult does (or would) this issue make it to accrue patients to this trial?	Not make it difficult to accrue patients to the trial	Make it somewhat difficult to accrue patients to the trial	Make it very difficult to accrue patients to the trial
Burden on patient to participate in the trial (e.g., logistics, time)	○	○	○
Inclusion/exclusion criteria of the study	○	○	○
Patients declining to enroll (e.g., unwilling to randomize, prefer one study arm)	○	○	○
Explaining the trial's details to a patient, including consenting	○	○	○
Getting patients referred to the trial	○	○	○
Cost to the patient (e.g., insurance, reimbursement)	○	○	○

Please elaborate – or provide additional reasons – why you believe this trial might be difficult to accrue patients at your site:

Opt out of survey Save and Continue -->

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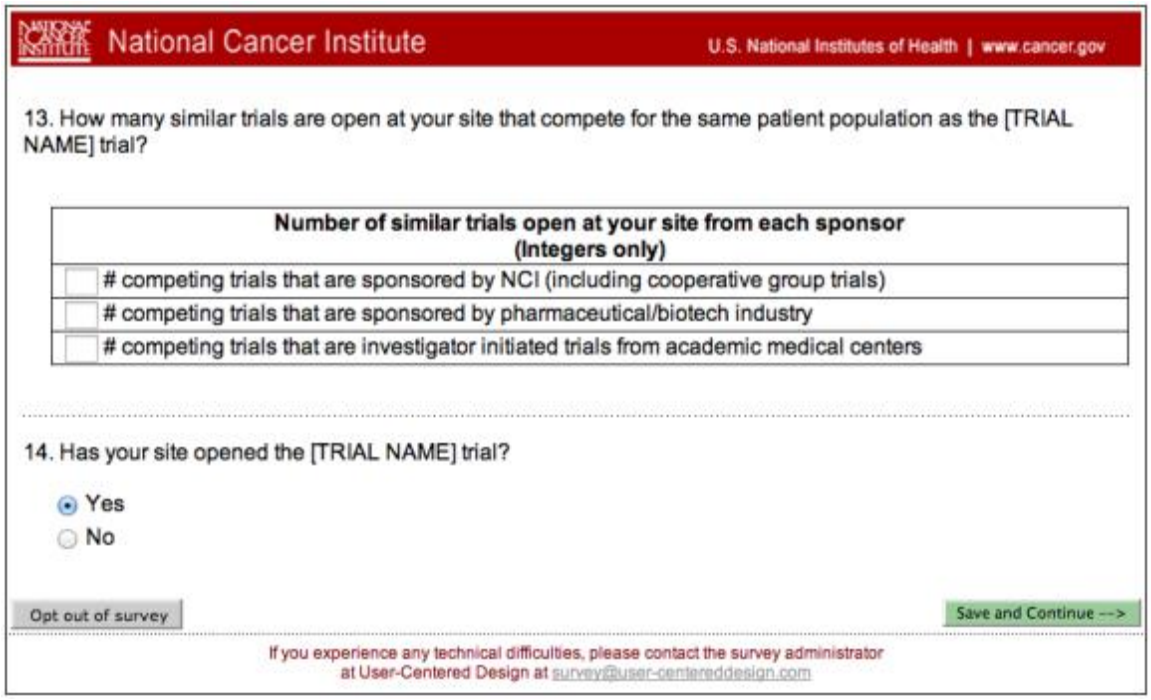
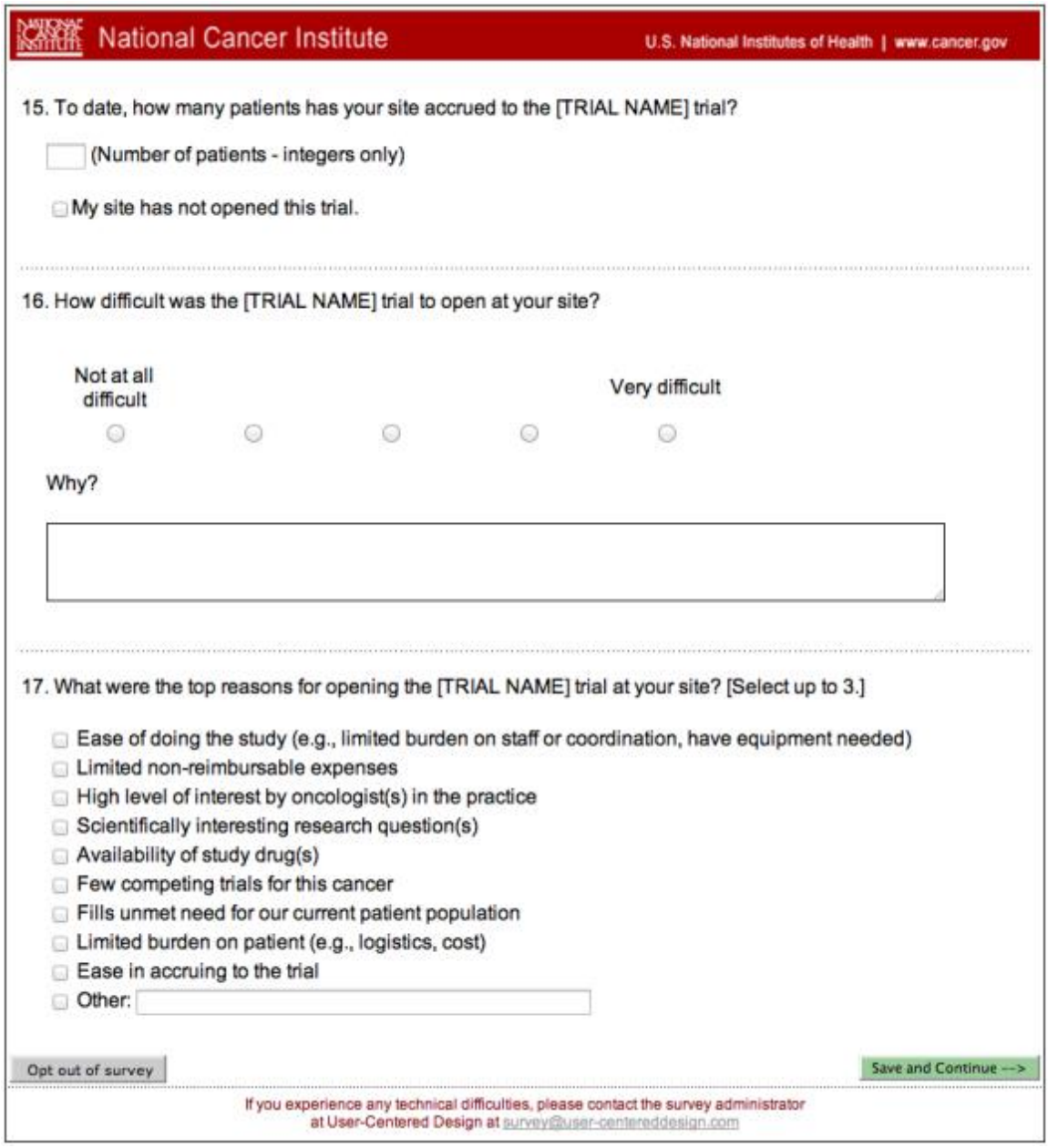


If user selects "Yes" to Q14, continue below. If user selects "No", continue HERE.



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15. To date, how many patients has your site accrued to the [TRIAL NAME] trial?

( ) (Number of patients - integers only)

☐ My site has not opened this trial.

16. How difficult was the [TRIAL NAME] trial to open at your site?

Not at all difficult				Very difficult
○	○	○	○	○

Why?

17. What were the top reasons for opening the [TRIAL NAME] trial at your site? [Select up to 3.]

- ☐ Ease of doing the study (e.g., limited burden on staff or coordination, have equipment needed)
- ☐ Limited non-reimbursable expenses
- ☐ High level of interest by oncologist(s) in the practice
- ☐ Scientifically interesting research question(s)
- ☐ Availability of study drug(s)
- ☐ Few competing trials for this cancer
- ☐ Fills unmet need for our current patient population
- ☐ Limited burden on patient (e.g., logistics, cost)
- ☐ Ease in accruing to the trial
- ☐ Other:

Opt out of survey Save and Continue -->

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END OF SURVEY

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13. How many similar trials are open at your site that compete for the same patient population as the [TRIAL NAME] trial?

Number of similar trials open at your site from each sponsor (Integers only)
☐ # competing trials that are sponsored by NCI (including cooperative group trials)
☐ # competing trials that are sponsored by pharmaceutical/biotech industry
☐ # competing trials that are investigator initiated trials from academic medical centers

14. Has your site opened the [TRIAL NAME] trial?

- ○ Yes
- ◉ No

Opt out of survey Save and Continue -->

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15. Given what you read in the trial summary sheet, how interested are you in opening the [TRIAL NAME] trial at your site?

Not at all interested				Very interested
○	○	○	○	○

16. What are the top reasons your site has not opened the [TRIAL NAME] trial?

- ☐ Not scientifically interesting enough
- ☐ Does not match our patient population
- ☐ Limited interest by oncologist(s) here
- ☐ Too many competing trials for this cancer
- ☐ Too great of a burden on patients (e.g., logistics, cost)
- ☐ Too difficult to conduct the study (e.g., burden on staff, coordination required, equipment needed)
- ☐ Financial cost to our site would be too great (e.g., non-reimbursable expenses)
- ☐ Too difficult to accrue patients (e.g., randomization, screening many to identify one; anticipate high refusals)
- ☐ Concerns about drug availability
- ☐ Other:

Opt out of survey Save and Continue -->

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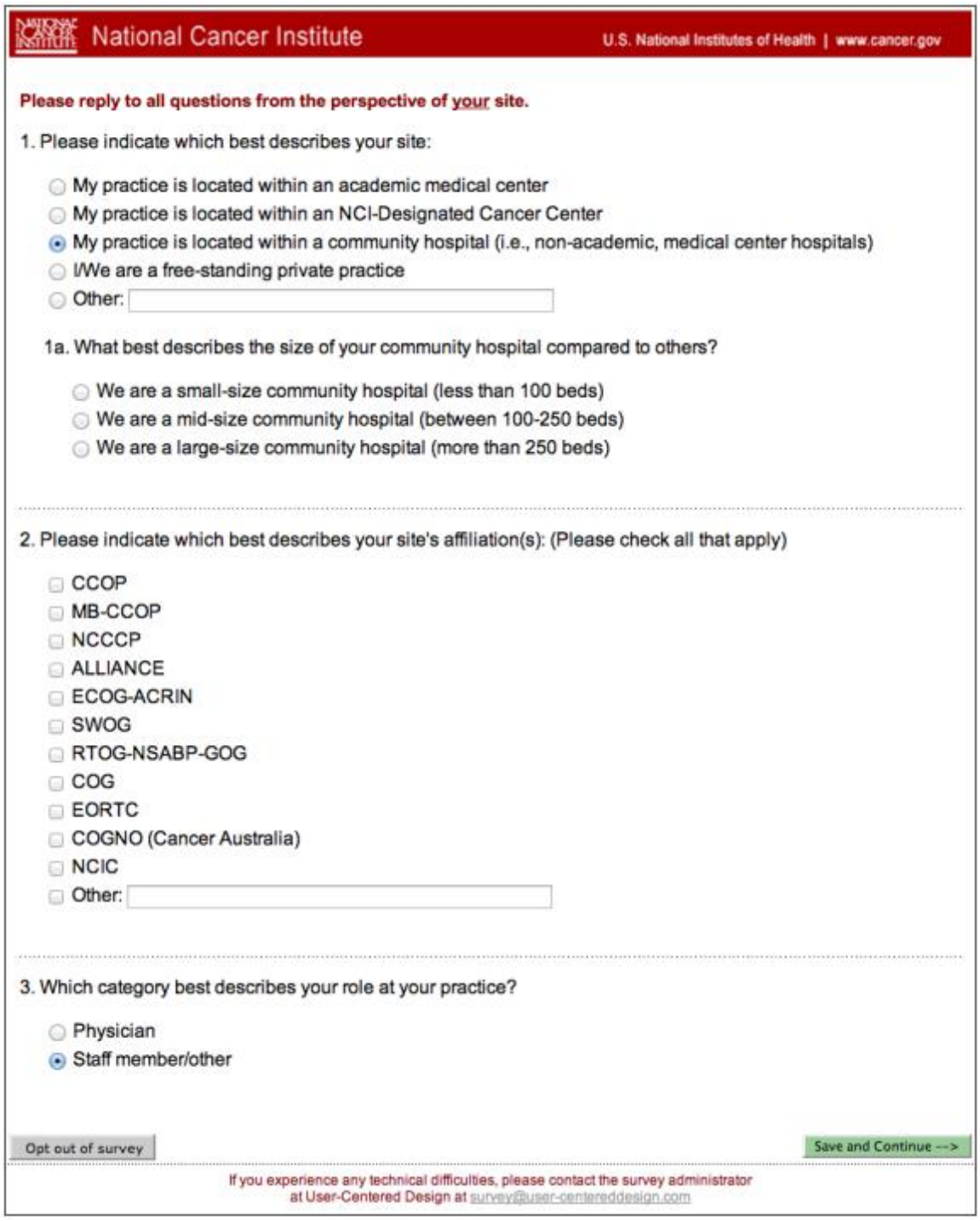
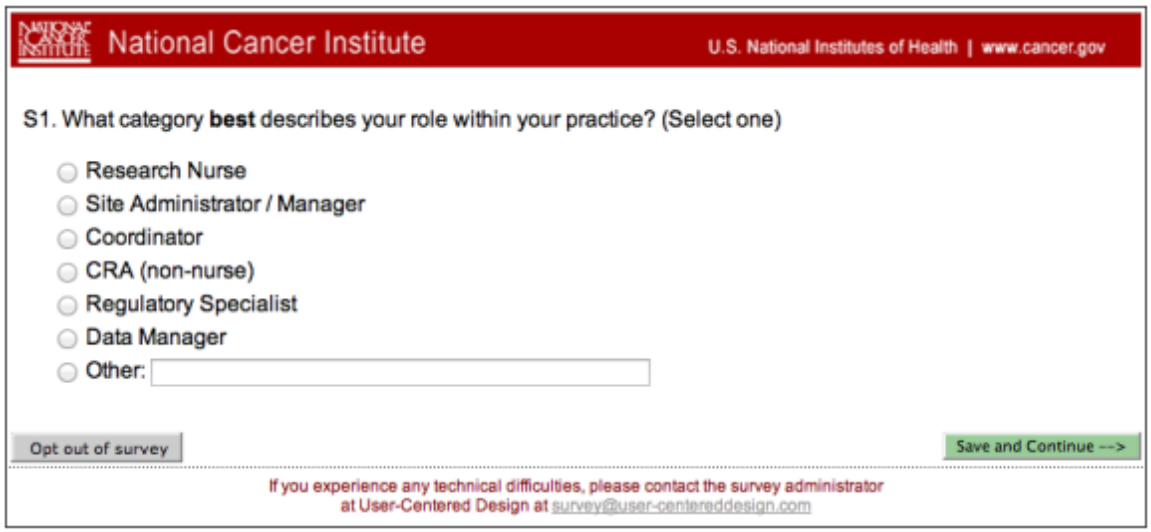


Note: Q1a appears only if "My practice is located within a community hospital" is selected for Q1.

Attachment_C11_lowaccruingsurvey



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The [TRIAL NAME] trial was opened by the [COOP GROUP NAME] in [INSERT MONTH, YEAR]. To date, [INSERT #] patients have been accrued to the trial. We expected a significantly more robust accrual than we have experienced. As such, NCI and [COOP GROUP NAME] are interested in learning from the field what issues are limiting accrual to [TRIAL NAME] and possible ways to increase its accrual and reach its projected goals.

S2. How much do you agree or disagree with this statement?

The potential benefits of this trial for patients are worth the effort and resources required to open and conduct it at my site.

Strongly Disagree				Strongly Agree
○	○	○	○	○

S3. For each of the trial's requirements listed below, please tell us whether or not it does (or would) make it difficult to open or run the [TRIAL NAME] trial at your site.

How difficult does (or would) this requirement make it to open or run this trial at your site?	Not make it difficult to open/ run the trial	Make it somewhat difficult to open/ run the trial	Make it very difficult to open/run the trial
[INSERT ITEM 1 FROM TRIAL SUMMARY SHEET]	○	○	○
[INSERT ITEM 2 FROM TRIAL SUMMARY SHEET]	○	○	○
[INSERT ITEM 3 FROM TRIAL SUMMARY SHEET]	○	○	○
[INSERT ITEM N FROM TRIAL SUMMARY SHEET]	○	○	○

Opt out of survey | Submit Survey -->

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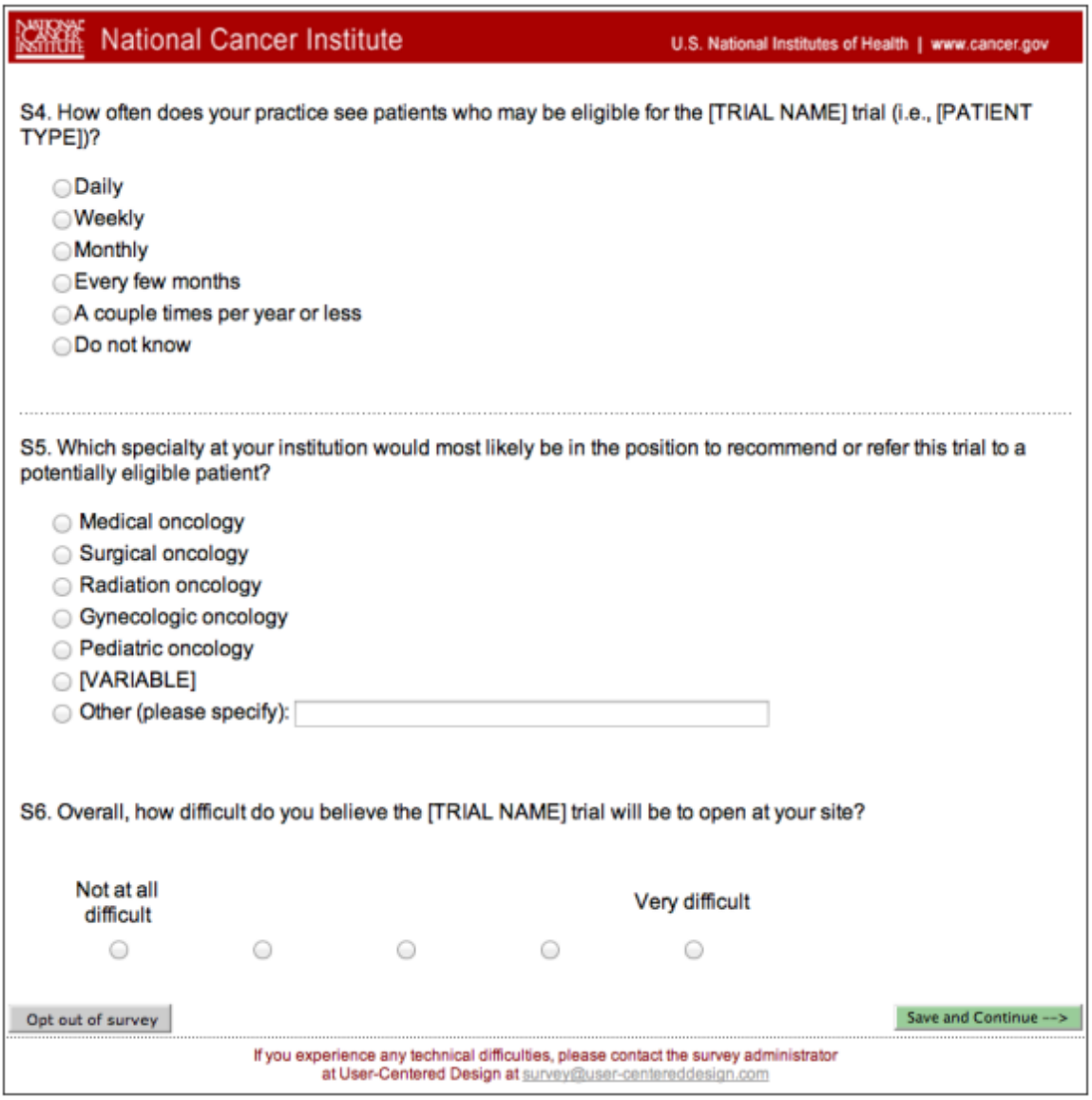

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S4. How often does your practice see patients who may be eligible for the [TRIAL NAME] trial (i.e., [PATIENT TYPE])?

- Daily
- Weekly
- Monthly
- Every few months
- A couple times per year or less
- Do not know

S5. Which specialty at your institution would most likely be in the position to recommend or refer this trial to a potentially eligible patient?

- Medical oncology
- Surgical oncology
- Radiation oncology
- Gynecologic oncology
- Pediatric oncology
- [VARIABLE]
- Other (please specify):

S6. Overall, how difficult do you believe the [TRIAL NAME] trial will be to open at your site?

Not at all difficult				Very difficult
○	○	○	○	○

Opt out of survey Save and Continue -->

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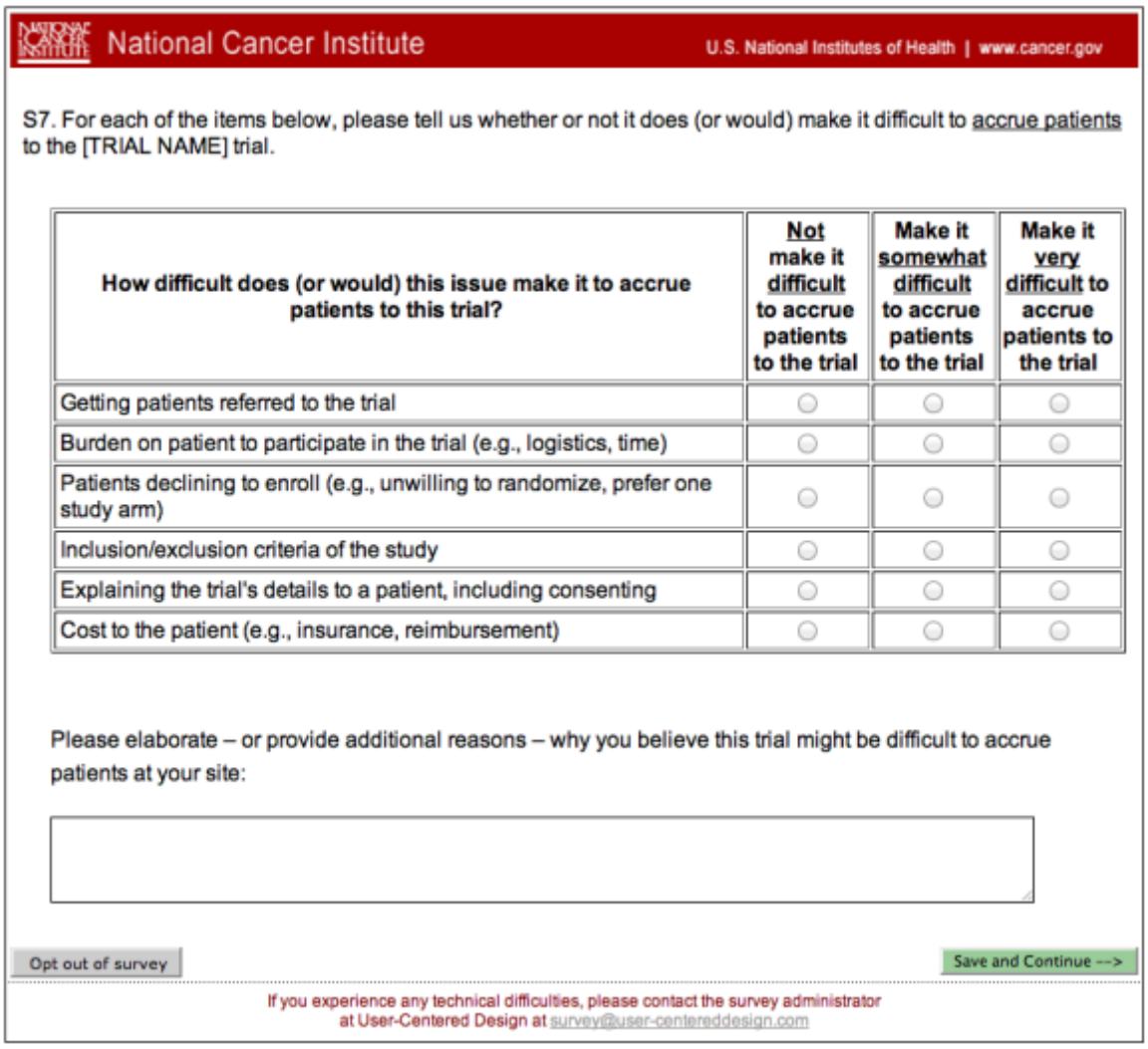
Attachment_C11_lowaccruingsurvey



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S7. For each of the items below, please tell us whether or not it does (or would) make it difficult to accrue patients to the [TRIAL NAME] trial.

How difficult does (or would) this issue make it to accrue patients to this trial?	Not make it difficult to accrue patients to the trial	Make it somewhat difficult to accrue patients to the trial	Make it very difficult to accrue patients to the trial
Getting patients referred to the trial	○	○	○
Burden on patient to participate in the trial (e.g., logistics, time)	○	○	○
Patients declining to enroll (e.g., unwilling to randomize, prefer one study arm)	○	○	○
Inclusion/exclusion criteria of the study	○	○	○
Explaining the trial's details to a patient, including consenting	○	○	○
Cost to the patient (e.g., insurance, reimbursement)	○	○	○

Please elaborate – or provide additional reasons – why you believe this trial might be difficult to accrue patients at your site:

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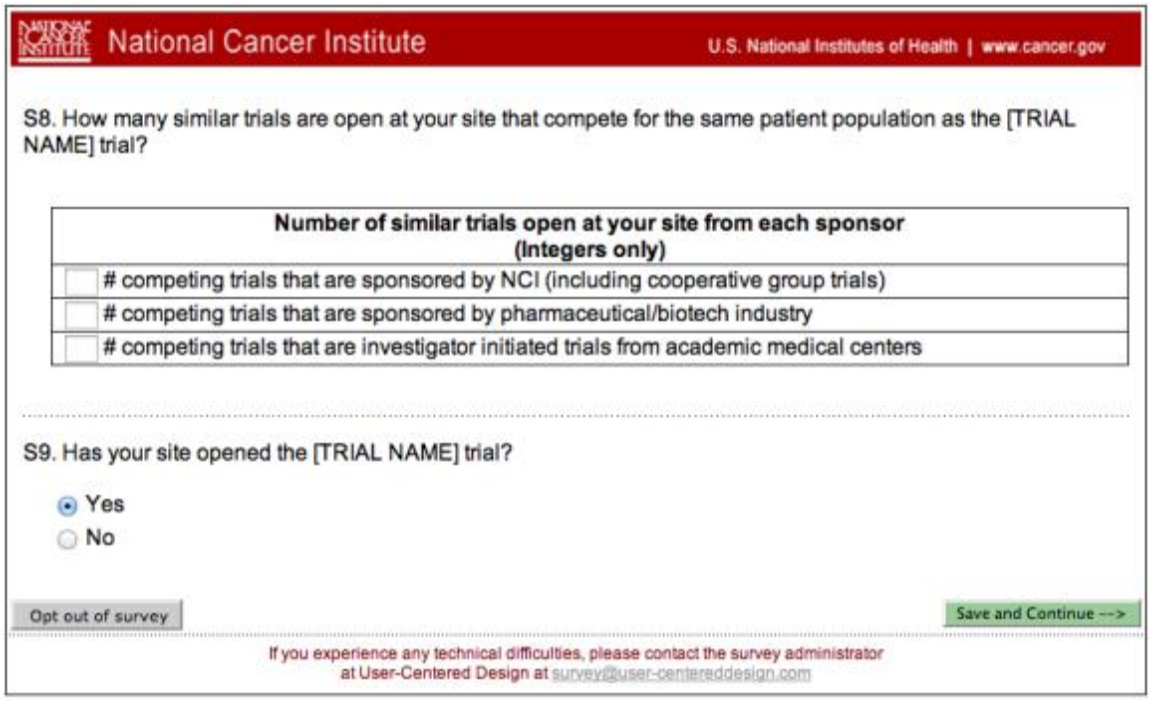
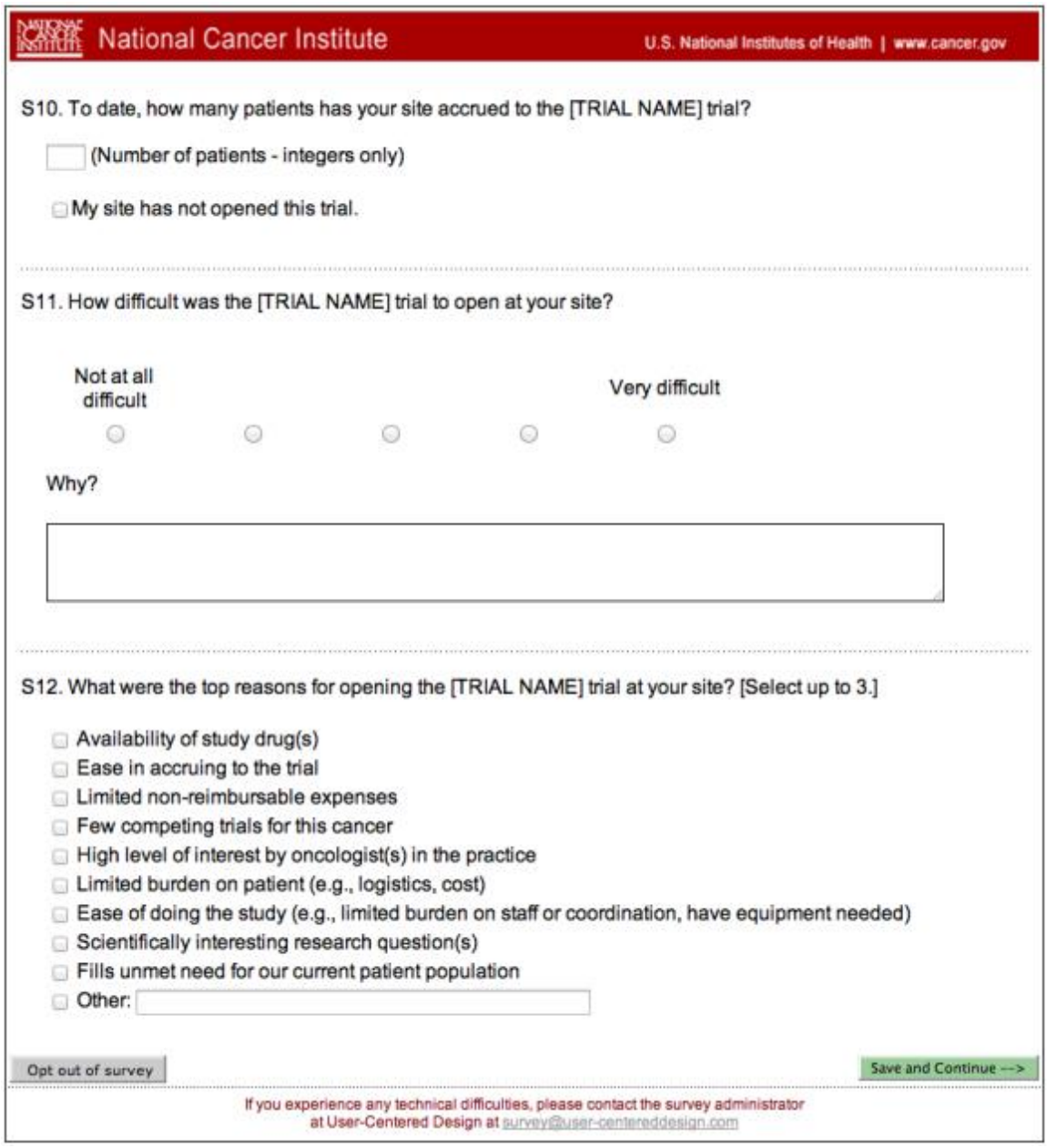


If user selects "Yes" to QS9, continue below. Otherwise continue HERE.

Attachment_C11_lowaccruingsurvey



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S10. To date, how many patients has your site accrued to the [TRIAL NAME] trial?

(Number of patients - integers only)

My site has not opened this trial.

S11. How difficult was the [TRIAL NAME] trial to open at your site?

Not at all difficult — Very difficult

Why?

S12. What were the top reasons for opening the [TRIAL NAME] trial at your site? [Select up to 3.]

- Availability of study drug(s)
- Ease in accruing to the trial
- Limited non-reimbursable expenses
- Few competing trials for this cancer
- High level of interest by oncologist(s) in the practice
- Limited burden on patient (e.g., logistics, cost)
- Ease of doing the study (e.g., limited burden on staff or coordination, have equipment needed)
- Scientifically interesting research question(s)
- Fills unmet need for our current patient population
- Other:

Opt out of survey

Save and Continue -->

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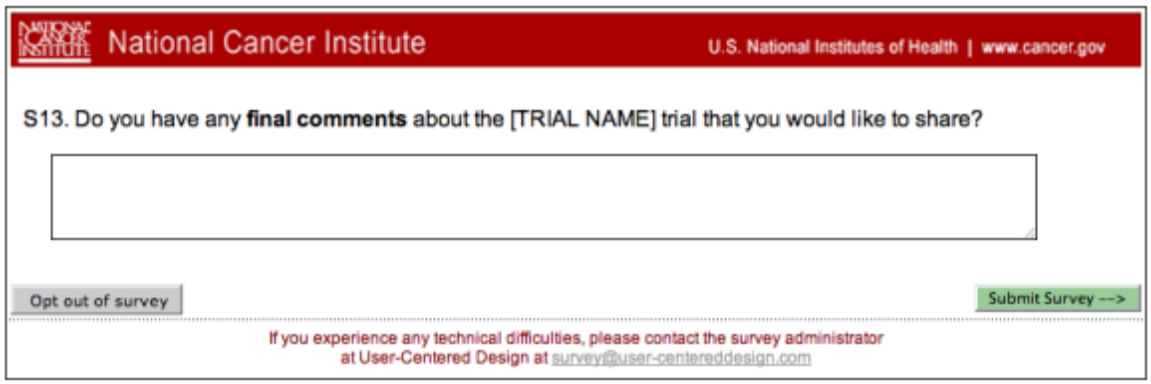

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S8. How many similar trials are open at your site that compete for the same patient population as the [TRIAL NAME] trial?

Number of similar trials open at your site from each sponsor (Integers only)
competing trials that are sponsored by NCI (including cooperative group trials)
competing trials that are sponsored by pharmaceutical/biotech industry
competing trials that are investigator initiated trials from academic medical centers

S9. Has your site opened the [TRIAL NAME] trial?

- ○ Yes
- ◉ No

Opt out of survey Save and Continue -->

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S10. Given what you read in the trial summary sheet, how interested are you in opening the [TRIAL NAME] trial at your site?

Not at all interested				Very interested
○	○	○	○	○

S11. What are the top reasons your site has not opened the [TRIAL NAME] trial?

- ☐ Not scientifically interesting enough
- ☐ Does not match our patient population
- ☐ Limited interest by oncologist(s) here
- ☐ Too many competing trials for this cancer
- ☐ Too great of a burden on patients (e.g., logistics, cost)
- ☐ Too difficult to conduct the study (e.g., burden on staff, coordination required, equipment needed)
- ☐ Financial cost to our site would be too great (e.g., non-reimbursable expenses)
- ☐ Too difficult to accrue patients (e.g., randomization, screening many to identify one; anticipate high refusals)
- ☐ Concerns about drug availability
- ☐ Other:

Opt out of survey Save and Continue -->

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If you experience any technical difficulties, please contact the survey administrator at User-Centered Design at survey@user-centereddesign.com

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