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

National Cancer Institute

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Introduction

The National Cancer Institute (NCI) and the [COOP GROUP NAME] would like your opinions about a new NCI clinical trial for [TRIAL DESCRIPTION].

They have developed a **brief online survey** tool to quickly and easily solicit feedback from physicians and research staff in the field to learn any potential issues with opening and accruing to this trial.

The PDF attachment to your email invitation provides an overview of the [TRIAL NAME] trial's concept. After reviewing this brief document we ask that you take 1 minute to answer this short survey.

Your comments will help us plan in advance for any concerns about the [TRIAL NAME] trial identified from this survey. 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 top 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.

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 (0000-0000-000). 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

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Trial Summary Sheet

IMPORTANT:

Please review the 2-page trial description attached to the email you received regarding this survey.

You can open a copy of the document here.

(This document will open in a new tab.)

I have reviewed the trial description and am ready to begin -->

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

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	Opt out of survey
Please reply to all questions from the perspective of y	<u>vour</u> site.
Please indicate which best describes your site:	
 My practice is located within an academic med 	tical center
 My practice is located within an NCI-Designate 	ed Cancer Center
 My practice is located within a community hosp hospitals) 	pital (i.e., non-academic, medical center
IWe are a free-standing private practice	
Other:	
1a. What best describes the size of your communit	ty hospital compared to others?
 We are a small-size community hospital (fev 	wer than 100 beds)
 We are a mid-size community hospital (between 	
 We are a large-size community hospital (mo 	
2. Please indicate which best describes your site's af	ffiliation(s): (Please check all that apply)
CCOP	
MB-CCOP	
NCCCP	
ALLIANCE	
ECOG-ACRIN	
SWOG	
NRG	
COG	
□ EORTC	
COGNO (Cancer Australia)	
NCIC	
Other:	
Which category best describes your role at your pro	actice?
Physician	
Staff member/other	
PC - PER PASSAGE (\$100 \$100 \$100 \$100 \$100 \$100 \$100 \$10	

Attachment_C10_prospectivesurvey

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

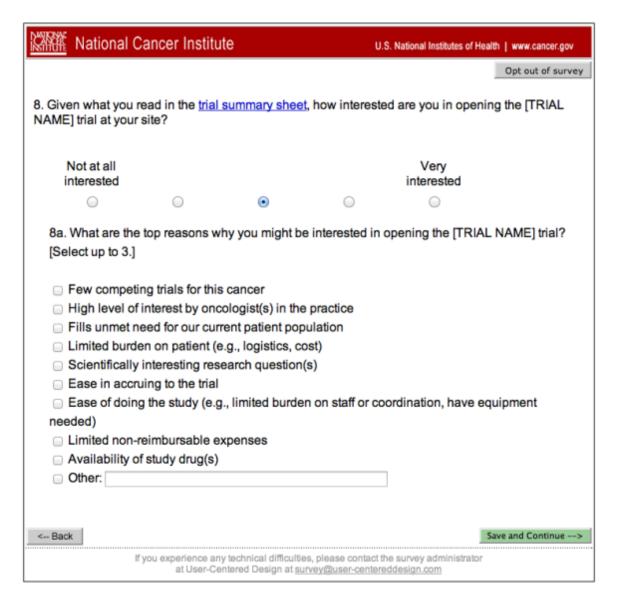
If user selects "physician" to Q3, continue below. If user selects "Staff member/other", continue HERE.

Mational Ca	ancer Institu	ite	U	.S. National Institutes of Health www.c	ancer.g
				Opt out	of su
What type of onco	logy best des	cribes your exp	ertise?		
Medical oncol	oav				
Surgical onco					
Radiation onc	7 7				
Gynecologic of	STATE OF THE STATE				
Pediatric onco	2000				
[VARIABLE]					
Other:					
The [TRIAL NAMI entifically interest RQ1: [INSERT R	ing it is to you	L.]. [For each,] Please tell us h	ow
ia. How scientific	ally interestin	g is this researc	ch question to	you?	
Not at all				Very	
interesting				interesting	
0	0	0	0	0	
Please elaborate	Č.				1
				9	
RQ2: [INSERT R	Q2 FROM TR	IAL SUMMAR	Y SHEET]		
b. How scientific	ally interestin	g is this research	ch question to	you?	
Not at all				Very	
interesting				interesting	
0	0	0	0	0	
Please elaborate					
					1
				9	1
					-
					i.
				Save and Co	

					Opt out of surve
. Assuming the [TF ndings will have or				hat impact do you	believe the
Low impact				High impact	
0	0	0	0	0	
The potential be to open and cor	enefits of this	trial for patient		e effort and reso	ources required
The potential be to open and cor Strongly	enefits of this	trial for patient		Strongly	ources required
The potential be to open and cor	enefits of this	trial for patient			ources required
to open and cor Strongly	enefits of this	trial for patient		Strongly	ources required

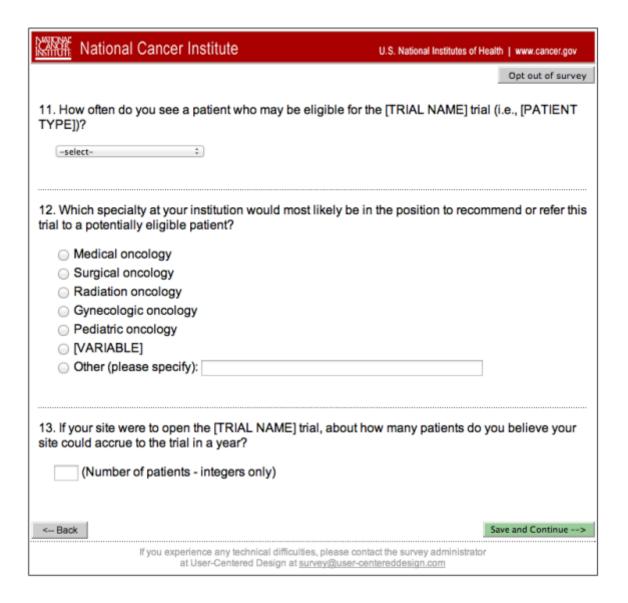
National (Cancer Institu	ite	U.S	6. National Institutes of Hea	ith www.cancer.gov
8. Given what you NAME] trial at your		summary shee	et, how intereste	d are you in openir	Opt out of survey
Not at all interested	•	0	0	Very interested	
8a. What are the [Select up to 3.]		hy you are not	that interested i	n opening the [TRI	AL NAME] trial?
☐ Too many co ☐ Limited inter ☐ Too difficult high refusals) ☐ Financial co	st to our site wo	for this cancer st(s) here nts (e.g., rando	mization, screer	ning many to idention	
Does not maNot scientific	ally interesting	population enough	den on staff coo	ordination required,	equipment
needed) Other:	our de la contraction de la co	, to.g., but	our on our, oo		oquipmont.
< Back			ies, piease contact th	e survey administrator	save and Continue>

Note: This 8a appears if the user selects either of the two leftmost radio buttons for $\mathsf{Q8}.$



Note: This Q8a appears if the user selects any of the three rightmost radio buttons for Q9.

		ents listed below, p NAME] trial at you		l us whether	or not it will n	nake it
	ill this require un this trial at	ement make it to d	pen or	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 2	2 FROM TRIA	L SUMMARY SHE	ET]	0	0	0
[INSERT ITEM I	N FROM TRIA	L SUMMARY SH	EET]	0	0	0
[INSERT ITEM	1 FROM TRIA	L SUMMARY SHE	ET]	0	0	0
[INSERT ITEM :	3 FROM TRIA	L SUMMARY SHE	ET]	0	0	0
[INSERT ITEM 4	FROM TRIA	L SUMMARY SHE	ET]	0	0	0
Overall, how diff Not at all difficult	ficult do you b	elieve the [TRIAL I	NAME] tr	ial will be to Very d		site?
0	0	0	0	0)	



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			C	pt out of survey
	For each of the items below, please tell us whether or no rue patients to the [TRIAL NAME] trial.	t you think it v	will make it di	fficult to
	How difficult will 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)	0	0	0
	Cost to the patient (e.g., insurance, reimbursement)			0
	Patients declining to enroll (e.g., unwilling to randomize, prefer one study arm)	0	0	0
- 11	Explaining the trial's details to a patient, including consenting	0	0	0
	Getting patients referred to the trial			
	Inclusion/exclusion criteria of the trial			0
	Please elaborate – or provide additional reasons – why your corue patients at your site:	ou believe thi	s trial might t	pe difficult to
			_	
<	Back			and Continue>
	If you experience any technical difficulties, please contr at User-Centered Design at <u>survey@user-cer</u>			

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		Opt out of survey
	w many similar trials are open at your site that tion as the [TRIAL NAME] trial? (Integers onl	
	Number of similar trials open a	t your site from each sponsor
	# competing trials that are sponsored by NO	CI (including cooperative group trials)
	# competing trials that are sponsored by ph	armaceutical/biotech industry
	# competing trials that are investigator initia	ted trials from academic medical centers
[TRIAI	-SPECIFIC QUESTION 1 HERE]	
[TRIAI	-SPECIFIC QUESTION 2 HERE]	
< Bac		Save and Continue>
	If you experience any technical difficulties, at User-Centered Design at <u>surve</u>	



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Survey Complete	Opt out of surve
Thank you for compl	eting this survey!
Your answers have been s	ubmitted anonymously.
We appreciate your feedb Your comments will help ensure that we pla	
If you would like a summary of the findings afte 61164thflrlab@	
You may now clos	e this window.
If you experience any technical difficulties, part User-Centered Design at surve	

END OF SURVEY

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	Opt out of survey						
Please reply to all questions from the perspective of	your site.						
1. Please indicate which best describes your site:							
 My practice is located within an academic med 	dical center						
 My practice is located within an NCI-Designate 	ed Cancer Center						
 My practice is located within a community hospital (i.e., non-academic, medical center hospitals) 							
 I/We are a free-standing private practice 							
Other:							
1a. What best describes the size of your communi							
 We are a small-size community hospital (fe 							
 We are a mid-size community hospital (bets) 							
 We are a large-size community hospital (me 	ore than 250 beds)						
2. Please indicate which best describes your site's a CCOP MB-CCOP NCCCP ALLIANCE ECOG-ACRIN SWOG NRG COG COG COG COG COG COG COGNO (Cancer Australia) NCIC COther:	ffiliation(s): (Please check all that apply)						
Which category best describes your role at your property of the state of the s	ractice?						
O Physician							
Staff member/other							
< Back	Save and Continue>						

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

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	Opt out of survey
S1. What category best describes your role within your pract	tice? (Select one)
Research Nurse	
 Site Administrator / Manager 	
○ Coordinator	
CRA (non-nurse)	
Regulatory Specialist	
Data Manager	
Other:	
< Back	Save and Continue>
lf you experience any technical difficulties, please cont at User-Centered Design at <u>survey@user-ce</u>	

紹維 National C	ancer Institute)	U	S. National Institutes of Health www	.cancer.gov
				Opt o	ut of survey
62. Given what you at your site will be in				sted do you believe the onc	cologists
Not at all interested				Very interested	
0	•	0	0	0	
S2a. What are th NAME] trial? [Se		hy your site m	ight not be int	erested in opening the [TR	IAL
Does not mat Too difficult to high refusals) Financial cos Concerns abo Too difficult to needed) Too many con	of accrue patients of to our site wou out drug available conduct the stu	opulation s (e.g., randon ld be too grea ility udy (e.g., burd r this cancer (s) here	it (e.g., non-rei	ning many to identify one; a mbursable expenses) ordination required, equipm	
3. How much do y The potential be to open and cor	enefits of this tr	ial for patient		ne effort and resources re	quired
Strongly Disagree				Strongly Agree	
0	0	0	0	0	
Please elaborate or	n your response	above:			
c Back				Save and C	Continue>
If			s, please contact rvey@user-center	he survey administrator addesign.com	

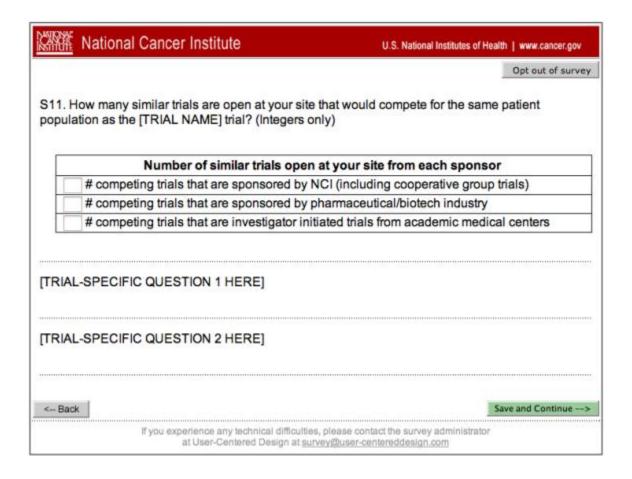
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National C	ancer Institut	te	į	J.S. National Institutes of Health www.cancer.gov
				Opt out of surve
S2. Given what you at your site will be in				sted do you believe the oncologists
Not at all interested				Very interested
0	0	•	0	0
S2a. What are th trial? [Select up t		vhy your site m	ight be intere	sted in opening the [TRIAL NAME]
Limited burde Fills unmet no Few competin Ease in accru Ease of doing needed) High level of Availability of	eed for our curr ng trials for this uing to the trial	.g., logistics, co ent patient pop cancer ., limited burder blogist(s) in the	n on staff or c	oo <mark>rdination, have equipment</mark>
Other:	ou agree or dis	agree with this	statement?	
The potential be to open and cor			s are worth t	the effort and resources required
Strongly Disagree				Strongly Agree
0	0	0	0	0
Please elaborate or	n your response	above:		
< Back		rtechnical difficultie		Save and Continue — the survey administrator reddesign.com

	Not make it difficult to open/ run the trial	Make it somewhat difficult to open/run the trial	Make it yery difficult to open/run the trial	
ARY SHEET]	0	0	0	
[INSERT ITEM N FROM TRIAL SUMMARY SHEET]				
[INSERT ITEM 1 FROM TRIAL SUMMARY SHEET]				
[INSERT ITEM 4 FROM TRIAL SUMMARY SHEET]				
[INSERT ITEM 3 FROM TRIAL SUMMARY SHEET]			0	
[TRIAL NAME] tr	al will be to Very d		site?	
[TRIAL NAME] tri			site?	
	Very d	ifficult		
	ARY SHEET]	make it difficult to open/run the trial ARY SHEET] ARY SHEET] ARY SHEET]	make it difficult to open/run the trial ARY SHEET] ARY SHEET] ARY SHEET] ARY SHEET]	

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	Opt out of survey			
S6. How often does your practice see a patient who may be [PATIENT TYPE])?	eligible for the [TRIAL NAME] trial (i.e.,			
	in the position to many mond or refer			
S7. Which specialty at your institution would most likely be it this trial to a potentially eligible patient?	in the position to recommend or refer			
Medical oncology				
Surgical oncology				
 Radiation oncology 				
 Gynecologic oncology 				
 Pediatric oncology 				
○ [VARIABLE]				
Other (please specify):				
S8. If your site were to open the [TRIAL NAME] trial, about how many patients do you believe your site could accrue to the trial in a year? [Number of patients - integers only)				
< Back	Save and Continue>			
If you experience any technical difficulties, please con at User-Centered Design at <u>survey@user-ce</u>				

ur site? Not at all difficult			Very di	fficult		
\circ	0	0 0	0)		
\$10. For each of the items below, please tell us whether or not you think it will make it difficult to ccrue patients to the [TRIAL NAME] trial.						
How diffic	cult will this issue ma patients to this tri		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 patient	ts referred to the trial		0		0	
Cost to the patient (e.g., insurance, reimbursement)			0	0	0	
Explaining the consenting	trial's details to a pati	ent, including	0	0	0	
Inclusion/exclu	usion criteria of the tria	l	0		0	
Burden on pat time)	ient to participate in th	e trial (e.g., logistics,	0	0	0	
Patients declir prefer one stud	ning to enroll (e.g., unv dy arm)	villing to randomize,	0	0	0	
Please elabora accrue patients	ite – or provide additionate:	nal reasons – why yo	ou believe thi	s trial might t	pe difficult to	



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		Opt out of survey
S12.1	Do you have any final comments about the [TRIAL	NAME] trial that you would like to share?
< Ba	ck	Submit Survey>
	If you experience any technical difficulties, please of at User-Centered Design at survey@user	

