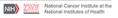
OMB #0925-xxxx

Expiration Date: xx/xx/xxxx



Team Driven. Cancer Therapy Focused.

Experimental Therapeutics Clinical Trials Network



Introduction

The National Cancer Institute (NCI) would like to know about your experiences with the Experimental Therapeutics Clinical Trials Network (ETCTN) over the past grant year (April 2014 - March 2015).

Your input will help NCI assess and refine the ETCTN processes and identify areas for improvement.

The survey will ask questions about ETCTN processes and activities and trial portfolio. It should take approximately 15 minutes to complete.

Your responses are confidential and all results will be reported in the aggregate. Your participation and responses will have no bearing on your ETCTN grant, or any future interactions with NCI.

We thank you for your assistance!

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

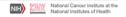


If you experience any technical difficulties, please contact the survey administrator, User-Centered Design at survey@user-centereddesign.com



Team Driven. Cancer Therapy Focused.

Experimental Therapeutics Clinical Trials Network



OMB# 0925-0046-09 Exp. Date: 05/31/2016

Privacy Statement and Consent

Collection of this information is authorized by The Public Health Service Act, Section 411 (42 USC 285a). Participation is voluntary and there are no penalties for not participating or withdrawing from the study at any time. The information collected in this study will be kept private to the extent provided by law. Names and other identifiers will not appear in any report of the study. Information provided will be combined for all study participants and reported as summaries. You are being contacted by User-Centered Design, Inc. via email to complete this instrument so that we can gain feedback from you on your experiences and recommendations regarding the implementation of the ETCTN program.

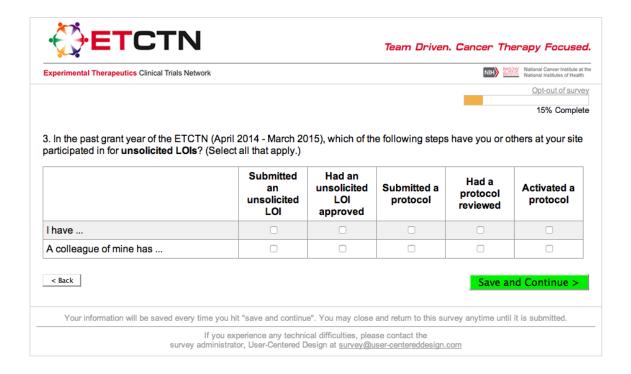
Please click the "Next" button if you consent to taking this survey.

Public reporting burden for this collection of information is estimated to average 15 minutes 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 (0925-0046-09). Do not return the completed form to this address.



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

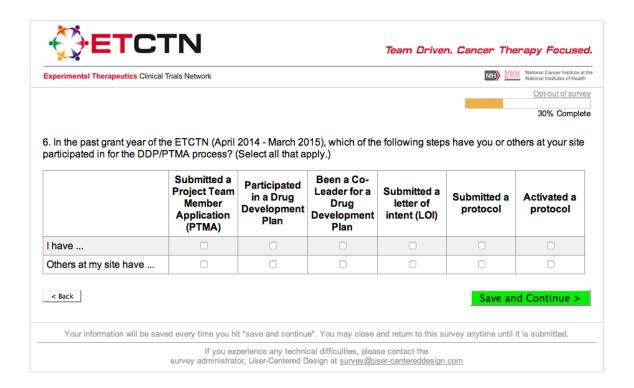
perimental Therapeutics Clinical Trials Network					NIH Natio	nal Cancer Institute a nal Institutes of Heal
					<u>(</u>	Opt-out of surv
				_		5% Comple
Unsolicited Letter of Intent (LOI) Process for	ETCTN Trials					
This set of questions asks you about your opinio	on of the <i>unsolicit</i> e	e d letter of	intent (LOI)	process.		
How aware are you of the unsolicited LOI pro-	cess used in the E	TCTN?				
Not very aware			ery vare			
0 0 0	0		0			
ransparency of the unsolicited LOI submission	Not very satisfied	0			Very satisfied	Don't know
ranguages of the uncellaited LOI submission	satisfied				satisfied	know
rocess	0	0	0	0	0	0
ransparency of the unsolicited LOI review proce		0	0	0	0	0
fficiency of the unsolicited LOI process	0	0	0	0	0	0
ommunication received from NCI about the nsolicited LOI process	0	\circ	0	0	0	0
umber of drugs available to investigators to evelop quality LOIs	0	0	0	0	0	0
larity of which drugs are available for unsolicite Ols	d	\circ	0	0	0	0
a. Please describe any changes NCI could make	e to increase your	satisfactio	n with the u	nsolicited	LOI process.	



Note: 3a appears conditionally, populated with any items selected for "I have..." in 03.

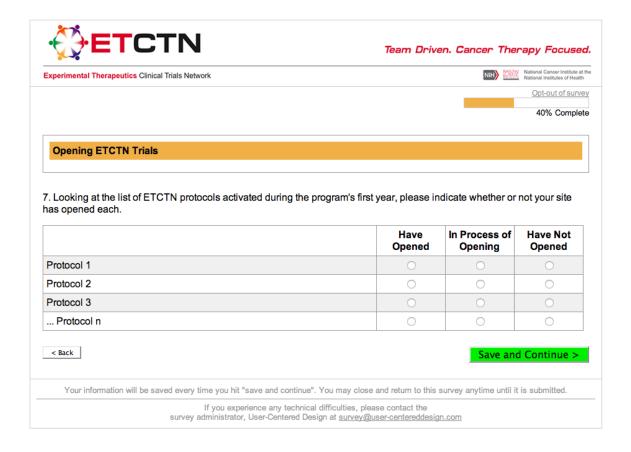
					National Cancer Institute National Institutes of He Opt-out of Sul
					20% Comp
a. What is your level of satisfaction for these phases of the	unsolicited LOI prod	cess that y	ou were in	olved with	1?
	Not very satisfied				Very satisfied
Submitted an unsolicited LOI	0	0	0	0	0
Ol review process	0	0	0	0	0
Submitted a protocol	0	0	0	0	0
Protocol review process	0	0	0	0	0
Activated a protocol	0	0	0	0	0

perimental Therapeutics Clinical Trials Network				NIH Natio	onal Cancer Institutional Institutes of H
				(Opt-out of su
					25% Comp
Project Team Member Applications and Drug Developmer	nt Plan to De	velop ETCT	N Trials		
this set of questions asks you about your opinion of the Drug L	Development	Project Teal	m Member	Application	DDP/
PTMA) process used to submit LOIs and protocols.					
And the Development PTMA	(DDD/DT144)		ETOTNO		
How aware are you of the Drug Development PTMA process ((DDP/PTMA)	used in the	EICIN?		
,					
Not very aware		/ery ware			
Not very		•			
Not very aware		ware			
Not very aware	a	ware			
Not very aware	a	ware	A process	?	
Not very aware	a	ware	A process	Strongly agree	Don't know
Not very aware How much do you agree or disagree with the following statem	ents about th	ware	•	Strongly	
Not very aware How much do you agree or disagree with the following statem The DDP/PTMA process	ents about th Strongly disagree	e DDP/PTM	Agree	Strongly agree	know
Not very aware How much do you agree or disagree with the following statem The DDP/PTMA process reamlines the development of early phase trials.	ents about th Strongly disagree	e DDP/PTM Disagree	Agree	Strongly agree	know
Not very aware How much do you agree or disagree with the following statem The DDP/PTMA process Teamlines the development of early phase trials. Incourages multidisciplinary teams.	ents about th Strongly disagree	e DDP/PTM Disagree	Agree	Strongly agree	know
Not very aware How much do you agree or disagree with the following statem The DDP/PTMA process reamlines the development of early phase trials. accourages multidisciplinary teams. a fair process to develop early phase trials. as reduced my workload in comparison to other early phase	ents about th Strongly disagree	e DDP/PTM Disagree	Agree	Strongly agree	know
Not very aware How much do you agree or disagree with the following statem The DDP/PTMA process reamlines the development of early phase trials. Incourages multidisciplinary teams. It a fair process to develop early phase trials. Its reduced my workload in comparison to other early phase all processes.	ents about th Strongly disagree	e DDP/PTM Disagree	Agree	Strongly agree	know

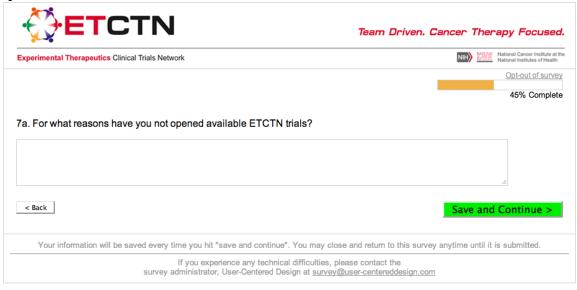


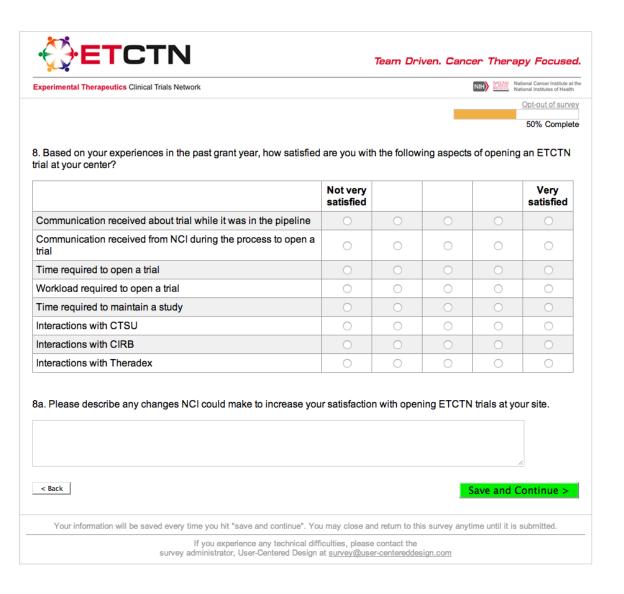
Note: 6a and 6b appear conditionally; 6a is populated with any items selected for "I have..." in Q6. 6b appears if 6a appears.

What is your level of satisfaction for each phase of the DDP/PTMA process in which you were involved? Not very satisfied Very satisfied	satisfied satisfied PTMA submission process Development of drug plan Co-leading drug development plan Letter of intent (LOI) submission Protocol submission	TMA process				
What is your level of satisfaction for each phase of the DDP/PTMA process in which you were involved? Not very satisfied Very satisfied	a. What is your level of satisfaction for each phase of the DDP/PTMA process in which you were involved? Not very satisfied Very satisfied PTMA submission process Oevelopment of drug plan Oevelopment of drug development plan Oevelopment (LOI) submission Oevelopment (LOI) submission Oevelopment of drug development plan Oevelopment plan Oevelop	TMA process	to orbitals or			
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satisfied satisfied MA submission process evelopment of drug plan beleading drug development plan tter of intent (LOI) submission cotocol submission cotocol activation	satisfied satisfied PTMA submission process Development of drug plan Co-leading drug development plan Letter of intent (LOI) submission Protocol submission Protocol activation		in which y	ou were inv	olved?	
MA submission process evelopment of drug plan p-leading drug development plan tter of intent (LOI) submission ptocol submission ptocol activation	PTMA submission process Development of drug plan Co-leading drug development plan Letter of intent (LOI) submission Protocol submission Protocol activation					
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otocol activation	Protocol activation	0	0	0	0	0
		0	0	0	0	0
Please describe any changes NCI could make to increase your satisfaction with the DDD/DTMA process	b. Please describe any changes NCI could make to increase your satisfaction with the DDP/PTMA process.	0	0	0	0	0
riease describe any changes Not could make to increase your satisfaction with the DDFF Livia process.		0	0	0	0	0
Please describe any changes NOI could make to increase yo	< Back		satisfied	satisfied	satisfied O O O O O O O O O O O O O O O O O O O	satisfied



Note: 7a appears conditionally if "have not opened" is selected for any protocol in Q7.





ETCTN protocols are intended to be innovative and are not to be duplicative of research funded by industry or other sources. Iow is a list of ETCTN protocols that were activated during the past grant year. For each protocol, please indicate whethout you consider the trial to be answering an important scientific question? Not important scientific question Important scientific question Don't know rotocol 1 Ontocol 2 Ontocol 3 Ontocol 3 Ontocol 3 Ontocol 3 Ontocol 4 Ontocol 5 Ontocol 5 Ontocol 6 Ontocol 7 Ontocol 7 Ontocol 8 Ontocol 8 Ontocol 9 Ontocol	•					NIH Nation	nal Cancer Institute nal Institutes of Hea
ETCTN protocols are intended to be innovative and are not to be duplicative of research funded by industry or other sources. Blow is a list of ETCTN protocols that were activated during the past grant year. For each protocol, please indicate whether not you consider the trial to be answering an important scientific question? Not						0	pt-out of sur
elow is a list of ETCTN protocols that were activated during the past grant year. For each protocol, please indicate whether not you consider the trial to be answering an important scientific question? Not important scientific question Important scientific question Don't know							55% Compl
important scientific question Protocol 1 Protocol 2 Protocol 3 Protocol n OLooking at the list of trials above, to what degree are you satisfied with the overall portfolio of activated protocols in the TCTN? Not satisfied Very satisfied	ETCTN Trial Portfolio						
Protocol 1 Protocol 2 Protocol 3 Protocol n Coloring at the list of trials above, to what degree are you satisfied with the overall portfolio of activated protocols in the TCTN? Not satisfied Very satisfied	esources. elow is a list of ETCTN protocols that were act	ivated during the pa mportant scientific of Not important	st grant ye			, please indica	ate whether
Protocol 2 Protocol 3 Protocol n O Looking at the list of trials above, to what degree are you satisfied with the overall portfolio of activated protocols in the TCTN? Not satisfied Very satisfied							know
Protocol 3 Protocol n Protoc	Protocol 1	0	0	0	0	0	0
D. Looking at the list of trials above, to what degree are you satisfied with the overall portfolio of activated protocols in the TCTN? Not satisfied Very satisfied	Protocol 2	0	0	0	0	0	0
0. Looking at the list of trials above, to what degree are you satisfied with the overall portfolio of activated protocols in the TCTN? Not satisfied Very satisfied	Protocol 3	0	\circ	0	0	0	0
Not satisfied Very satisfied O O O O O O O O O O O O O O O O O O	Protocol n	0	\circ	\circ	0	0	\circ
	TCTN? Not satisfied		v	ery isfied	tfolio of ac	tivated protoco	ols in the

			NIH)	National Cancer Institute at National Institutes of Health
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ect to the ETC	TN nortfolio	of trials		
	TT portiono	or trials.		
Not satisfied				Very satisfied
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er O	0	0	0	0
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			Save and	Continue >
	Not satisfied O O O O O O O O O O O O O O O O O O	Not satisfied O O O O O O O O O O O O O O O O O O O	satisfied O	ect to the ETCTN portfolio of trials. Not

Experimental Therapeutics Clinical Trials Network		NIH)	National Cancer Institutes of
			Opt-out of s
			65% Com
2. What changes would you like to see in the emphasis of	different scientific areas in trials	in the ETCTN p	ortfolio?
	Less emphasis	About the same emphasis	More emphasis
Angiogenesis	0	0	0
Immunotherapy	0	0	0
Molecular characterizations of tumor features	0	0	0
Organ dysfunction	0	0	0
Organ dysfunction	0	0	0
Pharmacokinetics	0	0	0
Signal transduction	0	0	0
Organ dysfunction Pharmacokinetics Signal transduction 12a. If other, please describe:	0	0	
			Z,
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If you experience any tech	nical difficulties, please contact the		

experimental Therapeutics Clinical Trials Network						tional Cancer Instit tional Institutes of I
						Opt-out of s
						70% Com
Network Readiness and Satisfaction with	th the ETCTN					
 Overall, how ready do you believe each or rant year in April 2014? 	of the groups below we	ere to partic	ipate in the	ETCTN at	the beginn	ing of th
rant year III April 2014:						
	Not				Very	Don't
	ready				read	know
Your center		0	0	0	read	know
	ready	0	0	0		
	ready				0	0
The ETCTN overall	ready	0	0	0	0	0
The ETCTN overall	ready	0	0	0	0	0
The ETCTN overall	ready of the groups below is	0	0	0	Very	O Don't
The ETCTN overall 4. Overall, how ready do you believe each o	ready of the groups below is Not ready	now to par	ticipate in th	e ETCTN?	Very	Don't know
The ETCTN overall 4. Overall, how ready do you believe each of the second of the seco	of the groups below is Not ready	now to par	ticipate in th	e ETCTN?	Very read	Don't know
Your center The ETCTN overall 4. Overall, how ready do you believe each of the center Your center The ETCTN overall	ready of the groups below is Not ready	now to par	ticipate in th	e ETCTN?	Very	Don't know
The ETCTN overall 4. Overall, how ready do you believe each of the second of the seco	of the groups below is Not ready	now to par	ticipate in th	e ETCTN?	Very read	Don't know

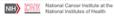
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Opt-out of survey

80% Complete

16. Below is a list of potential benefits to be achieved via the ETCTN network.

Please indicate if you think that each of the potential benefits is:

- More likely to occur with ETCTN trials,
- More likely to occur with other investigator-initiated trials at your center OR
- Equally likely to occur with other investigator-initiated trials and with ETCTN trials

	More Likely to Occur With ETCTN Trials at my Center	More Likely to Occur With Other Investigator- Initiated Trials at my Center	Likely to Occur Equally With Other Investigator- Initiated Trials AND ETCTN Trials
Obtain IRB approval quickly.	0	0	0
Activate trials quickly at my center.	0	0	0
Participate in scientifically important trials.	0	0	0
Access drugs early in their development.	0	0	0
Participate in trials that will help advance my career.	0	0	0
Enroll patients to trials at the projected accrual rate.	0	0	0
Provide my patients access to a wide variety of early phase clinical trial options.	0	0	0
Collaborate with investigators at other centers.	0	0	0
Collaborate with researchers from other disciplines.	0	0	0

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Social Network Collaborations

We would like to learn more about your work with other researchers that are involved in the ETCTN.

17. Please look at the list of ETCTN organization members below and indicate a "yes" for those you had direct collaboration with over the past year (either developing protocols or opening trials).

ETCTN Lead Academic Organizations and Affiliates	Yes	No
Translational Genomics Research Institute	0	0
City of Hope Comprehensive Cancer Center	0	0
UC Davis Comprehensive Cancer Center	0	0
USC Norris Comprehensive Cancer Center	0	0
University of Colorado Cancer Center - Anschutz Cancer Pavilion	0	0
Yale Cancer Center	0	0
H. Lee Moffitt Cancer Center and Research Institute	0	0
Emory University/Winship Cancer Institute	0	0
University of Chicago	0	0
Johns Hopkins University/Sidney Kimmel Comprehensive Cancer Center	0	0
National Cancer Institute Developmental Therapeutics Clinic	0	0
University of Maryland Greenbaum Cancer Center	0	0
Dana-Farber Cancer Center	0	0
Massachusetts General Hospital	0	0
Wayne State University/Karmanos Cancer Institute	0	0
Mayo Clinic Rochester	0	0
Washington University	0	0
Rutgers University - Cancer Institute of New Jersey	0	0
Roswell Park Cancer Institute	0	0
Duke University	0	0
UNC Chapel Hill	0	0
Case Western Reserve University	0	0
Cleveland Clinic Foundation	0	0
Ohio State University Comprehensive Cancer Center	0	0
Fox Chase Cancer Center	0	0
University of Pittsburgh Cancer Institute	0	0
Vanderbilt-Ingram Cancer Center	0	0
University of Texas MD Anderson Cancer Center	0	0
Virginia Commonwealth University	0	0
University Wisconsin Carbone Cancer Center	0	0
British Columbia Cancer Agency	0	0
Juravinski Cancer Center	0	0
University Health Network/Princess Margaret Cancer Center	0	0

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