OMB No.: 0925-0766 Expiration Date: 04/30/2023

\* 1. Company Information

Public reporting burden for this collection of information is estimated to average 2 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 20892-7974, ATTN: PRA (0925-0766). Do not return the completed form to this address.

## 2022 NCI SBIR Development Center Program with FDA CARE - Connecting Awardees with Regulatory Experts

Company Name				
Contact Person				
Title/Role				
Email				
* 2. Please indicate the extent to whi CARE Program helped our team	ch you agree wit Strongly Agree	h the follow Agree	ing. Participation	on in the Strongly Disagree
Learn and/or confirm the FDA Center (CBER, CDER, CDRH) or Office that will regulate our technology	0	$\circ$	0	0
Begin developing the regulatory strategy for our technology	$\bigcirc$	$\bigcirc$	$\circ$	$\bigcirc$
Plan the next regulatory step(s) for our technology	$\bigcirc$		$\bigcirc$	$\circ$
Clarify and/or confirm the appropriate regulatory path for our technology	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\bigcirc$
Gain a better understanding of the process to contact FDA for a meeting request	$\bigcirc$	$\bigcirc$	$\bigcirc$	$\circ$
Find information on FDA's website related to a particular regulatory topic	$\bigcirc$		$\circ$	
Understand how to contact an FDA office that provides free resource assistance to industry (i.e., MATTB, SBIA, or DICE)	0	$\bigcirc$	$\circ$	$\circ$

Program, our team plans to				
	Strongly Agree	Agree	Disagree	Strongly Disagree
Contact an FDA office via phone or email that provides free resource assistance to industry (i.e., MATTB, SBIA, or DICE)	$\bigcirc$			$\circ$
Watch an FDA webinar(s) on a particular regulatory topic		$\bigcirc$	$\bigcirc$	
Read an FDA guidance(s) on a particular regulatory topic				
Access an FDA website to learn more on a particular regulatory topic		$\bigcirc$	$\bigcirc$	$\bigcirc$
Submit a meeting request to FDA to discuss regulatory strategy (e.g., INTERACT meeting, pre-IND meeting, pre-submission meeting, etc.)	$\circ$	$\bigcirc$		$\circ$
* 4. Please indicate the extent to which	ch you agree wit	h the follow	ing. Our team	
				Strongly
	Strongly Agree	Agree	Disagree	
Found the CARE Program useful	Strongly Agree	Agree	Disagree	Strongly
Found the CARE Program useful  Found the NCI SBIR FDA resources website useful <a href="https://sbir.cancer.gov/resources/fda-resources">https://sbir.cancer.gov/resources/fda-resources</a>	Strongly Agree	Agree	Disagree	Strongly
Found the NCI SBIR FDA resources website useful <a href="https://sbir.cancer.gov/resources/fda-">https://sbir.cancer.gov/resources/fda-</a>	Strongly Agree	Agree	Disagree	Strongly
Found the NCI SBIR FDA resources website useful <a href="https://sbir.cancer.gov/resources/fda-resources">https://sbir.cancer.gov/resources/fda-resources</a> Received information from the CARE Program that will affect our future	Strongly Agree	Agree	Disagree	Strongly