DEPARTMENT OF HEALTH AND HUMAN SERVICES
Food and Drug Administration

Food and Drug Administration Exp

TO DE COMPLETED DY ADDITION

Form Approved: OMB No. 0910-0396 Expiration Date: April 30, 2022

DISCLOSURE: FINANCIAL INTERESTS AND ARRANGEMENTS OF CLINICAL INVESTIGATORS

The following information concerning	, who participated
as a clinical investigator in the submitted study	
is submitted in accordance with	21 CFR part 54. The
clinical study named individual has participated in financial arrangements or holds fina required to be disclosed as follows:	ancial interests that are
Please mark the applicable check boxes.	
any financial arrangement entered into between the sponsor of the cove investigator involved in the conduct of the covered study, whereby the vato the clinical investigator for conducting the study could be influenced study;	alue of the compensation
any significant payments of other sorts made on or after February 2, 19 the covered study, such as a grant to fund ongoing research, compequipment, retainer for ongoing consultation, or honoraria;	
 any proprietary interest in the product tested in the covered studinvestigator; 	dy held by the clinical
any significant equity interest, as defined in 21 CFR 54.2(b), held by the sponsor of the covered study.	he clinical investigator in
Details of the individual's disclosable financial arrangements and interests a description of steps taken to minimize the potential bias of clinical stud disclosed arrangements or interests.	
NAME TITLE	
FIRM/ORGANIZATION	
SIGNATURE Date (mm/c	dd/yyyy)

This section applies only to the requirements of the Paperwork Reduction Act of 1995.

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. Public reporting burden for this collection of information is estimated to average 5 hours per response, including time for reviewing instructions, searching existing data sources, gathering and maintaining the necessary data, and completing and reviewing the collection of information. Send comments regarding this burden estimate or any other aspect of this collection of information to the address to the right:

Do NOT send your completed form to the PRA Staff email address below.

Department of Health and Human Services Food and Drug Administration Office of Operations PRAStaff@fda.hhs.gov

"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 number."