DEPARTMENT OF HEALTH AND HUMAN SERVICES
Food and Drug Administration

CERTIFICATION: FINANCIAL INTERESTS AND ARRANGEMENTS OF CLINICAL INVESTIGATORS

Form Approved: OMB No. 0910-0396 Expiration Date: December 31, 2015

With respect to all covered clinical studies (or spessor support of this application, I certify to one of t certification is made in compliance with 21 CFR investigator includes the spouse and each dependent	the statements belo part 54 and that for	ow as appropriate. I understand that this r the purposes of this statement, a clinica
Please mar	k the applicable check b	ox.
(1) As the sponsor of the submitted studies, with the listed clinical investigators (enter this form) whereby the value of compensa study as defined in 21 CFR 54.2(a). I also to the sponsor whether the investigator has the sponsor as defined in 21 CFR 54.2(b listed investigator was the recipient of sign	names of clinical in ation to the investiga o certify that each lis ad a proprietary inte b) did not disclose a	vestigators below or attach list of names to tor could be affected by the outcome of the ted clinical investigator required to discloss rest in this product or a significant equity is any such interests. I further certify that n
Clinical Investigators		
(2) As the applicant who is submitting a st applicant, I certify that based on informa investigators, the listed clinical investigator financial arrangement with the sponsor o investigator for conducting the study cou CFR 54.2(a)); had no proprietary interest the covered study (as defined in 21 CFR other sorts (as defined in 21 CFR 54.2(f)).	ation obtained from ors (attach list of na of a covered study v Id be affected by th t in this product or s 54.2(b)); and was	the sponsor or from participating clinical mes to this form) did not participate in an whereby the value of compensation to the ne outcome of the study (as defined in 2 significant equity interest in the sponsor of
(3) As the applicant who is submitting a st applicant, I certify that I have acted with (attach list of names) or from the sponsor do so. The reason why this information co	n due diligence to c the information req	obtain from the listed clinical investigator uired under 54.4 and it was not possible t
NAME	TITLE	
FIRM/ORGANIZATION		
SIGNATURE		
SIGNATURE		DATE (mm/dd/yyyy)
L		

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."

or any other aspect of this collection of information to the address to the right: