

DEPARTMENT OF HEALTH AND HUMAN SERVICES Public Health Service Food and Drug Administration	Form Approved: OMB No. xxx-xxxx Expiration Date: xx/xx/xx
DISCLOSURE: FINANCIAL INTERESTS AND ARRANGEMENTS OF CLINICAL INVESTIGATORS	
TO BE COMPLETED BY APPLICANT	
The following information concerning <u>PAYED FINANCE</u> , who participated as a clinical investigator in the submitted study <u>GP0000</u> , is submitted in accordance with 21 CFR part 54. The named individual has participated in financial arrangements or holds financial interests that are required to be disclosed as follows:	
Please mark the applicable checkboxes.	
<input type="checkbox"/> any financial arrangement entered into between the sponsor of the covered study and the clinical investigator involved in the conduct of the covered study, whereby the value of the compensation to the clinical investigator for conducting the study could be influenced by the outcome of the study;	
<input type="checkbox"/> any significant payments of other sorts from the sponsor of the covered study such as a grant to fund ongoing research, compensation in the form of equipment, retainer for ongoing consultation, or honoraria;	
<input type="checkbox"/> any proprietary interest in the product tested in the covered study held by the clinical investigator;	
<input checked="" type="checkbox"/> any significant equity interest in the sponsor of the covered study held by the clinical investigator as defined in 21 CFR 54.2(b).	
Details of the individual's disclosable financial arrangements and interests are attached, along with a description of steps taken to minimize the potential bias of clinical study results by any of the disclosed arrangements or interests.	
NAME	TITLE
Johnfin Money, M.B.A.	Chief Financial Officer
FIRM/ORGANIZATION	
Good Pharmaceuticals	
SIGNATURE	DATE
Johnfin Money, M.B.A.	5-1-01
Public reporting burden for this collection of information is estimated to average 4 hours per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing reviewing the collection of information. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden to:	
DHHS Reports Clearance Officer Paperwork Reduction Project (0910-xxxx) Humphrey Building, Room 531-H 200 Independence Ave., SW Washington, DC 20201	
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.	
Please DO NOT RETURN this application to this address.	

DEPARTMENT OF HEALTH AND HUMAN SERVICES Public Health Service Food and Drug Administration		Form Approved: OMB No. xxx-xxxx Expiration Date: xx/xx/xx
CERTIFICATION: FINANCIAL INTERESTS AND ARRANGEMENTS OF CLINICAL INVESTIGATORS		
TO BE COMPLETED BY APPLICANT		
With respect to all covered clinical studies (or specific clinical studies listed below (if appropriate)) submitted in support of this application, I certify to one of the statements below as appropriate. I understand that this certification is made in compliance with 21 CFR part 54 and that for the purposes of this statement, a clinical investigator includes the spouse and each dependent child of the investigator as defined in 21 CFR 54.2(d).		
Please mark the applicable checkbox.		
<input type="checkbox"/> (1) As the sponsor of the submitted studies, I certify that I have not entered into any financial arrangement with the listed clinical investigators (enter names of clinical investigators below or attach list of names to this form) whereby the value of compensation to the investigator could be affected by the outcome of the study as defined in 21 CFR 54.2(a). I also certify that each listed clinical investigator required to disclose to the sponsor whether the investigator had a proprietary interest in this product or a significant equity in the sponsor as defined in 21 CFR 54.2(b) did not disclose any such interests. I further certify that no listed investigator was the recipient of significant payments of other sorts as defined in 21 CFR 54.2(f).		
Clinical Investigators		
<input checked="" type="checkbox"/> (2) As the applicant who is submitting a study or studies sponsored by a firm or party other than the applicant, I certify that based on information obtained from the sponsor or from participating clinical investigators, the listed clinical investigators (attach list of names to this form) did not participate in any financial arrangement with the sponsor of a covered study whereby the value of compensation to the investigator for conducting the study could be affected by the outcome of the study (as defined in 21 CFR 54.2(a)); had no proprietary interest in this product or significant equity interest in the sponsor of the covered study (as defined in 21 CFR 54.2(b)); and was not the recipient of significant payments of other sorts (as defined in 21 CFR 54.2(f)).		
<input type="checkbox"/> (3) As the applicant who is submitting a study or studies sponsored by a firm or party other than the applicant, I certify that I have acted with due diligence to obtain from the listed clinical investigators (attach list of names) or from the sponsor the information required under 54.4 and it was not possible to do so. The reason why this information could not be obtained is attached.		
NAME		TITLE
Johnfin Money, M.B.A.		Chief Financial Officer
FIRM/ORGANIZATION		
Good Pharmaceuticals		
SIGNATURE		DATE
Johnfin Money, M.B.A.		5-1-01
Public reporting burden for this collection of information is estimated to average 1 hour per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing reviewing the collection of information. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden to:		
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