

SPONSOR INITIATED CLINICAL TRIAL INFORMATION FORM 1. $Please\ submit\ this\ form\ to\ the\ Grants\ \&\ Contracts\ Coordinator,\ Providence\ Research\ c/o\ \underline{jvalenzuela@providencehealth.bc.ca}.$ 2. For all other funding, please use UBC Research Project Information Form (RPIF). Research Ethics Board (REB)# Date Received: Contact Information Research Institute Principal Investigator Name: Academic Rank Faculty: Telephone: Email: Department: Address: Division: Study Coordinator/Nurse Email: Name: Facsimile: Telephone: Address: Sponsor/Legal Agreement Contact Company/ Organization Contact Name: Email: Telephone: Facsimile: Address: B. **Project Details** Study Title: Protocol/Study Nickname: Type of Study: Phase:

Does th	ne Project involve:						
	Clinical Study Drug	No	Yes	Certificate/Application #			
	Clinical Study Device	No	Yes	Certificate/Application #			
i)	Funding Source/Sponsor:						
ii)	CRO (if applicable):						
The account for this study will be held at:							
Pro	ovidence Health Care Research Institute		-	Γhe University of British Colu	mbia		
Indicate Institution (UBC, RI or formally affiliated institution) where research activity for the project will be undertaken. Select all that apply:							
	UBC Vancouver Campus UBC Okanagan Campus BC Cancer Agency BC Centre for Disease Control Providence Health Care Research Institute BC Mental Health & Addictions Research Institute						
				_			
∐ Chi	ild & Family Research Institute	Nomen's Heal	th Research Institute	e ∐ Vancouver Co	pastal Health Research Institute		
Otl	her:						
C.	Conflict of Interest						
Are you	aware of any conflicts of interest that may	have a bearing	g on this project?	No - Please proceed to Se			
				Yes - Please check applica	ible boxes below.		
		F	Principal Investigato	r Co-Investigator(s)	Student(s)		
	Seat on Board of Directors						
	Seat on Scientific Advisory Board						
	Any Role within the Company						
	Shares in Sponsor Company						
	License/Option Agreement						
	Non-Disclosure Agreement						
	Consulting Agreement						
L	Other Conflicts of Interest:						
D.	Additional Information						
i) Will you be using any confidential materials or information in the project? No Yes - Please specify below							
Source							
of Materia	al						
Matura							
Nature of							
Materia	al						

ii)	Are you conducting any research for another collaborator or sponsor that might No Yes - F overlap with this project?	lease describ	e below		
iii)	Will any employees of the collaborator or sponsor be participating in the project?	No	Yes		
	If YES, will they be participating on site at PHC or UBC ?	No	Yes		
iv)	Will part of this study be subcontracted by the Principal Investigator to a third party external to the institution? (Eg., a non-institutional pharmacy or lab?). Do not include study functions subcontracted by the Sponsor (ie. central lab arranged by Sponsor that is to be used by all sites participating in the study).	No	Yes		
	If YES, please advise what part(s) of the study will be subcontracted and the name of the third party(ies):				
v)	Will any personnel not employed by the institution (UBC or PHC) be part of the research team? If YES, who is the employer of these individuals?	No	Yes		
	If YES, how are the non-institutional personnel insured?				
vi) vii)	Does any investigator plan to publish or present the results of this study? Has any investigator or other personnel involved in the study been debarred or investigated by the FDA or any	No	Yes		
·	other regulatory authority for debarment action?	No	Yes		
viii)	Is there any investigator involved in the study that does not have CMPA coverage?	No	Yes		
	If YES, please provide additional details:				
ix)	Is there any other information you wish to provide (eg., regarding timelines, study start up meetings, etc.)?				
E.	Signatures				
	ordance with institutional policies, holders of research and trust accounts must be members of the permanent academ ened for lecturers or research associates, if allowed by the sponsor, and at the specific request of the Dean.	ic staff. Acco	unts ma		
Princip	oal Investigator. I understand that Indirect Costs must be included in the budget as per <u>UBC Policy #LR2</u> .				
Signa	ignature: Date:				
	Please submit this form, with a copy of the study protocol and draft agreement, electronically to: Providence Research c/o jvalenzuela@providencehealth.bc.ca				