

SECTION I: APPLICATION INFORMATION				
1. Study Protocol Code	1.1 Reference Number:			
	1.2 UV-IRB CODE:	1.2 UV-IRB CODE:		
	1.3 Study Code (spons	1.3 Study Code (sponsored study only):		
2. Date of Initial	Click here to enter a c	late.		
Submission		Γ		
3. Resubmission History	Date Submitted	Type of Submission	Version	Version Date
Resubmissions – responses to		□ Resubmission		
initial review recommendations		□ Amendment		
or submission of studies with investigator-initiated changes				
prior to ethics approval		Resubmission		
Amendments – responses to		□ Amendment		
initial review recommendations				
or submission of studies with		Resubmission		
investigator-initiated changes after ethics approval		□ Amendment		
		Resubmission		
		□ Amendment		
		Resubmission		
		□ Amendment		
			Ear spansored study, copy t	he version code (if any). For
				ave these columns blank.
4. Study Category	• 4.1 Research involving human participants			
	\Box 4.1.1 Research involving Indigenous Herbal Medicine or People – Apply			
			mission on Indigenous	
	□4.1.2 Clinical Tr	ial intended for Mark	eting Registration – Ap	ply simultaneously
	with the Food and Drug Administration © 4.2 Research involving animals - STOP.			
	Please contact the University of the Visayas, Institutional Animal Care and Use Committee			
	C 4.3 Research involving non-human participants -			
	Please apply for UV-IRB Exempt for Review			
	C 4.4 Research involving biological and hazardous materials - STOP.			
	Please contact the National Committee of Biosafety of the Philippines			
5. Category of	● 5.1 UV Faculty/Staff ○ 5.2 UV Undergraduate Student			
Investigator	© 5.3 UV Graduate © 5.3.1 Masteral Thesis © 5.3.2 Doctoral Dissertation			
	© 5.4 Non-UV: (NOTE	: This category requi	res completion of PAF	RT



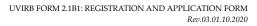
	BENEFICENCE			
6.	Purpose of study	© 6.1 Academic Requirement (Thesis, Dissertation, Training Requirement)		
		© 6.2 Institutional Requirement (Work-Related)		
		© 6.3 Multi-Institutional or Multi-Country Collaboration		
		6.4 Others (indicate): Click here to enter text.		
7.	Study Title	Click here to enter text.		
8.	Study Duration (Data Gathering, Analysis and Reporting)	(in months) Click here to enter text.		
9.	Type of study:	9.1 Social and Behavioral Researches, specifically (choose one):		
		 9.1.1 Research on Economics, Business, Leadership, Management, Organizational Psychology, Operations, and Process 9.1.2 Research on Law, Penology, Criminal and Political Science, and Public Administration 		
		□9.1.3 Research on Media and Communication		
		□9.1.4 Research on Pedagogy and Educational Sciences		
		9.1.5 Research on Sociology, Anthropology, Ethnology, Social Psychology, History, and Socio-Cultural		
		Geography \Box 9.1.6 Research on Urban studies (social aspect of planning and development)		
		\square 9.1.7 Social and Educational Experiments		
		\square 9.2 Technical Researches and Data Science , specifically (choose one):		
		\square 9.2.1 Data Mining and Review of Records		
		\square 9.2.2 Researches on Informatics, Information Technology and Computer Science		
		\Box 9.3.4 Research on Technology, Invention, Innovation and Design		
		9.3 Health Non-Clinical Trial, specifically (choose one):		
		□9.3.1 In vitro study		
		\Box 9.3.2 Herbal Research, and Complementary and Alternative Medicine Research		
		9.2.3 Epidemiological study		
		9.2.4 Clinical Psychology and Special Education		
		9.4 Experimental Studies, specifically (choose one):		
		\Box 9.4.1 Clinical Trial (drug or pharmaceutical trials, diagnostic trials, trials on devices, and other therapy trials)		
		therapy trials)		
		\Box 9.4.1.1 intended for marketing registration – Phase: $\bigcirc 1 \bigcirc 2 \bigcirc 3 \bigcirc 4$		
		□9.4.1.2 Not interfaced for marketing registration		
		9.4.3 Social and Educational Experiments		
		□9.5 Humanities:		
		9.5.1 Visual and Performing Arts, and Composition		
		9.5.2 Language and Literature		
		\Box 9.5.3 Philosophy		
		\Box 9.6 Others, please indicate:		



BENEFICENCE			
10. Use of special	□10.1 Children (under 18)	\Box 10.12 Cognitively and emotionally	
populations or	□10.2 Indigenous People	impaired persons	
vulnerable groups	□10.3 Elderly	\Box 10.13 Institutionalized people	
□10.0 Not applicable	□ 10.4 People on Welfare/Social Assistance	(Hospital, Facility, Homes, Hospice,	
	□10.5 Poor and Unemployed	Rehabilitation)	
	10.6 Patients in Emergency Care	\Box 10.14 Members of the armed forces	
	□10.7 Homeless Persons	or police	
	□10.8 Refugees or Displaced Persons	\Box 10.15 Physically disabled	
	□10.9 Patients with Incurable Diseases	Politically powerless	
	□10.10 Prisoners	\Box 10.16 Students, employees and	
	□10.11 Pregnant Women and Fetus	subordinates \Box 10.17 Junior members of hierarchial	
	□10.11 AIDS/HIV+ subjects	groups	
		10.18 Others (indicate):	
11. Endorsing	C 11.1 Callega of Dusiness and		
College/Unit/	© 11.1 College of Business and	O 11.9 College of Maritime Studies	
Institution	© 11.2 College of Arts and Sciences	© 11.10 College of Medicine	
© 11.0 Not applicable	© 11.3 College of Engineering and	© 11.11 College of Education	
	© 11.4 College of Computer Science	11.12 College of Law	
	© 11.5 College of Dentistry	11.13 GS- Education	
	© 11.6 College of Nursing	11.13 GS- Business	
	© 11.7 College of Pharmacy	11.13 GS- Nursing	
	© 11.8 College of Criminal Justice Education	O 11.13 UV Staff:	
		© 11.14 Non-UV:	
12. Funding Agency:	12.1 (NAME):		
C 12.0 Not Applicable	 12.1 Higher Education Institution 12.2 PHL Government Agency/Office/Entity 12.3 Multilateral Agency (UN agencies and other Intergovernmental Agencies) 12.4 Private Company or Non-Governmental Organization (NGO) 		
	C 12.5 Others (indicate):		
13. Study Budget (For	NOTE: This refers to line item amounts. However, if a separate budget sheet is		
funded researches	available, just indicate total amount and attach budget sheet		
only)			
□13.0 Not Applicable	-	-	
14. Previous ethics	14.1 Name of Institutional Review Board or Ethics Review Committee:		
approval or clearance	14.2 Date of Ethics Approval:		
issued by other sites	14.3 Date of Expiration of Ethics Approval:		
14.0 Not Applicable			
15. Proponent			



16. Proponent Address			
17. Proponent			
Telephone:			
18. Proponent Facsimile:			
19. Proponent Mobile:			
20. Proponent Email:			
21. Other Ongoing	21.1 Title:	21.3 Title:	
studies	21.1.1 UVIRB Code (if applicable):	21.3.1 UVIRB Code (if applicable):	
	21.2 Title:	21.4 Title:	
	21.2.1 UVIRB Code (if applicable):	21.4.1 UVIRB Code (if applicable):	
22. Declaration of	22.1 I have personal/family financial in	terest in the results of the study	
Conflict of Interest of	NATURE:	· · · · · · · · · · · · · · · · · · ·	
Proponent	\Box 22.2 I Have proprietary interest in the research for which this application is being		
\Box 22.0 I have no conflict	made (patent, trademark, copyright, licensing)		
of interest in any form	NATURE:		
(financial, proprietary,	22.3 I have significant financial Interest	s as defined in US 45 CFR Part 94 (Note: This	
professional) with	category is only for applications for which	this regulation may apply. For information,	
sponsor, the study, Co-	refer to http://www.ecfr.gov)		
Investigators, or the site	NATURE		
23. Other investigators	Co-Investigator: Task description:		
with corresponding			
task description (add			
additional rows as			
applicable) \Box 22.0 Not Applicable			
\Box 23.0 Not Applicable			
24. Submitted by:		1	





DENTITICENCE		
	Study designation	
25. Proponent signature		



SECTION II: SCIENTIFIC/TECHNICAL REVIEW APPROVAL ENDORSEMENT This section should be signed by the Chair/Head of the Scientific/Technical Review Committee/Office that reviewed the scientific soundness of the study and issued the appropriate approval. Alternatively, results of Scientific/Technical Review disposition may be appended to this application, instead of completing this section, provided that the information required below had been appropriately addressed. STUDY PROTOCOL TITLE: <with Version Number and Date> Proponent: <Title, Name, Surname> I confirm that the (NAME OF SCIENTIFIC/TECHNICAL REVIEW COMMITTEE/OFFICE) has reviewed and approved the following study protocol-related information: Objectives/Expected output supported by literature review; overall research design; sampling design, sample size, Inclusion/exclusion/ withdrawal criteria; data collection plan and specimen collection, processing, and storage as applicable; data analysis plan including statistical design/framework, as applicable. Issuing Committee/Office/College: Head of Committee/Office <Title, Name, Surname> (Chair, Thesis/Dissertation Committee): Signature: Date of Signature: SECTION III: INSTITUTIONAL/DEPARTMENTAL ENDORSEMENT This section should be signed by the head of unit (administrative authority legally empowered to sign on behalf the unit such as Dean, Director, and the like) of the Principal Investigator. This section is required only for initial submission, provided there are no changes in study protocol information below. I confirm that I have read this Application and that the research will be implemented under the oversight of this Department/Institution in accordance with the conditions of approval by the University of the Visayas Institutional Review Board. I also confirm that the Principal Investigator has a regular appointment/is a bonafide student in this institution. Head of unit (Dean or **Research Coordinator):** Date of Signature: Signature:



SECTION IV: AUTHORIZA	SECTION IV: AUTHORIZATION AND ACKNOWLEDGEMENT OF REVIEW			
This section should be completed by the signatory official who can sign on behalf of the institution that has oversight				
on the research site, IF the rese	on the research site, IF the research site is OUTSIDE the scope of authority of UV and the PI is non UV personnel . If			
not applicable, put N/A in all fie	not applicable, put N/A in all fields. This section is required only for initial submission, provided there are no changes			
in study protocol information b	elow.			
STUDY PROTOCOL TITLE:				
Proponent:	<title, name,="" surname=""></title,>			
This is to certify that the <name of="" research="" site="">:</name>				
1) Has no local Institutional Review Board/ Ethics Review Committee; and				
2) Authorizes and acknowledges the University of the Visayas Institutional Review Board (UVIRB), located at the 2 nd				
Floor Admin Building, UV Main Campus, Corner D. Jakosalem and Colon St. Cebu City, 6000 Philippines, to perform				
the ethical review of the abovementioned study protocol in accordance with international ethical standards and				
national regulatory requirements, and oversee the conduct of the research study which includes progress				
monitoring, adverse event monitoring, and site visits.				
Name of Research Site				
Address of Research Site				
Signatory Official	<title, name,="" surname=""></title,>			
Position of Official				
Signature		Date of Signature:		