# **INSTITUTIONAL ETHICS COMMITTEE**

National Ethics Committee Registry for Biomedical and Health Research (NECRBHR), Department of Health Research (DHR) Regd. No: EC/NEW/INST/2020/687

iecvspm@gmail.com

#### VSPM'S DENTAL COLLEGE & RESEARCH CENTRE, NAGPUR-440019

Digdoh Hills, Hingna Road, Nagpur-440 019 Ph: (07104) 665000,665301 Fax: (07104) 665011, Email: <a href="mailto:vspmdcrcnagpur@gmail.com">vspmdcrcnagpur@gmail.com</a> Website: <a href="mailto:https://vspmdcrc.edu.in">https://vspmdcrc.edu.in</a>

Chairperson: Dr. R. K Yeltiwar

Member Secretary: Dr. Alka Dive

#### Application form for permission of Research Project/Dissertation

(N.B. to be submitted in triplicate, one copy will be returned to the department after approval) (N.B. To be preserved by IEC & departments for minimum 3 years)

	Dr. R. K Yeltiwar			Dr. Alka Dive	
	Chairperson			Member Secretary	
	Date of approval by IEC				
	Date of resubmission to IEC if applicable			<u>.                                      </u>	
	Date of IEC meeting			Dept:	
	Date of Receipt by IEC			Sr.No	
	(For IEC Office Use Only)				
	vi) Signature of the Dean (with seal)	:			
	Department involved (with seal)				
	v) Signature of HODs of other	:			
	Investigator (with seal)				
	iv) Signature of HOD of Principal	:			
	. ) C. A. CHOD CD 1				
	iii) Signature of Guide	:			
	ii) Signature of Co-Investigator	:			
5.	i) Signature of Principal Investigator	:			
	•	•			
4.	Title of Research Project/ Dissertation				
٥.	Department Department	•			
3.	Name of Guide and				
2.	Name of Co-Investigator	:			
	(UG/Intern/PG/Staff)	:			
1.	Name of Principal Investigator				

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6. Place where research work carried out

		At VSPM'S DCRC Campus	a)		
	b)	Outside VSPM'S DCRC	b)		
7.	Tin	ne period required for completion	:		
	of I	Research project & its analysis			
8.	Ris	k factors for the patient (give details)	:		
		rocedural			
		dverse drug reaction (ADRs)			
		vasive investigations xplain the measures to control the			
		ove risk factors			
9.	Det	tails of financial burden involved &	:		
	Ho	w it will be met			
10.	. Wh	nether research project is sponsored	:		
11.	. An	y other relevant information	:		
Ple	ease	tick relevant enclosures			
1) Three copies of Research protocol on A-4 size paper only					
	2)	Patient information sheet			
	3)	Informed consent form in English/Marathi(I	CF)		
	4)	Case record form (CRF)			
	,	Article publication matter			
	6)	Permission letters of other institute if any			
-		are that I shall follow national and internation ting the above research project.	onal Good Clinical Practice (GCF) guidelines in		
			Signature of Principal Investigator		
No	to. D	binging Investigator has to submit COMDITI	NON DEPORT of the research project to IEC in		

Note: Principal Investigator has to submit COMPLETION REPORT of the research project to IEC in prescribed format on completion of the project in hard copy to office and soft copy to <a href="mailto:iecvspm@gmail.com">iecvspm@gmail.com</a>.

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#### COMPLETION REPORT OF RESEARCH PROJECT

1.	Name of the Department	:	
2.	Name of the Principal Investigator		
	(UG/Intern/PG/Staff) :		
3.	Title of the Project	:	
4.	Collaborating Institutions, if any	:	
5.	Date of Approval by IEC and IEC		
	number	:	
6.	Duration	:	
7.	Date of Completion :		
8.	Clinical trial registration details		
	(if any)	:	
9.	Conclusion and Scope for future	:	
	work		
10.	Academic publication/presentation	:	
	with complete details		
11.	Patents taken, if any :		
12.	Products developed, if any	:	
P	rincipal Investigator	Name of Guide (if applicable)	Head of the Department
	Signature with date	Signature with date	Signature with date