## PARTICIPANT INFORMED CONSENT FORM (PICF) (English)

Protocol / Study number :	
Participant identification number for this trial:	
Title of project:	

Name of Principal Investigator: \_\_\_\_\_\_ Tel.No(s).\_\_\_\_\_ The contents of the information sheet dated that was provided have been read carefully by me / explained in detail to me, in a language that I comprehend, and I have fully understood the contents. I confirm that I have had the opportunity to ask questions.

The nature and purpose of the study and its potential risks / benefits and expected duration of the study, and other relevant details of the study have been explained to me in detail. I understand that my participation is voluntary and that I am free to withdraw at any time, without giving any reason, without my medical care or legal right being affected.

I understand that the information collected about me from my participation in this research and sections of any of my medical notes may be looked at by responsible individuals from AIIMS. I give permission for these individuals to have access to my records.

I agree to take part in the above study.

	Date:
(Signatures / Left Thumb Impression)	Place:
Name of the Participant: Son / Daughter / Spouse of: Complete postal address:	
This is to certify that the above consent has be	en obtained in my presence.
Signatures of the Principal Investigator	Date: Place:
1) Witness – 1	2) Witness – 2
Signatures	Signatures
Name:	Name:
Address: <u>NB Three copies should be made, for (1) pa</u>	Address: tient, (2) researcher, (2) Institution
(Investigators are advised to prepare the transla	ation in simple understandable Hindi on their own.)