

## PARTICIPANT INFORMED CONSENT FORM (PICF)

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.

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(Signatures / Left Thumb Impression)

Date:  
Place:

Name of the Participant: \_\_\_\_\_

Son / Daughter / Spouse of: \_\_\_\_\_

Complete postal address: \_\_\_\_\_

This is to certify that the above consent has been obtained in my presence.

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Signatures of the Principal Investigator

Date:  
Place:

1) Witness – 1

2) Witness – 2

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Signatures

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Signatures

Name:

Name:

Address:

Address:

**NB Three copies should be made, for (1) patient, (2) researcher, (2) Institution**  
**(Investigators are advised to prepare the translation in simple understandable Hindi on their own.)**