

INFORMED CONSENT FORM

TITLE OF THE PROPOSED STUDY:

“To estimate the prevalence of hsCRP levels in acute ischemic stroke and subtypes”

Name of the Patient.....

Age/Sex.....

Registration no......

INFORMED CONSENT

(Permission to participate in research)

1. I have been fully informed about the procedure to be conducted in this study and I authorize Dr. Ish Anand and the team for doing this work on me.
2. I confirm that *I have read and understood the information sheet* for the above study and have had the opportunity to ask questions.
3. I understand that *my participation in the study is voluntary and that I am free to withdraw at any time*, without giving any reason, without my medical care or legal rights being affected.
4. *I agree not to restrict the use of any data* or results that arise from this study provided such a use is only for scientific purposes.
5. I understand *that my treatment will not be affected if I refuse to participate*. I have been informed that I can discontinue my participation at any time.
6. *My identity will be kept confidential*. I agree that I will not demand any limits on the use of the findings of the study.

7. I authorize access to my medical records as explained in this information and consent form.

8. I give my *free* and *full consent* to participate in this study.

➤ **Patient's name:**..... Signature.....
Date: Place:

➤ **Witness's/Relative's name**..... Signature.....
Date: Place:

➤ **Principal Investigator-cum-Guide**
Dr. Ish Anand Signature.....
Date: Place:

➤ **Thesis Scholar-cum-Co-investigator**
Dr. Davinder Singh Rana Signature.....
Date: Place:

➤ **Co-investigator-cum-Co-guide**
1. Dr. (Col) P.K Sethi Signature.....
Date: Place:

2. Dr. Anuradha Batra Signature.....
Date: Place:

3. Dr. Seema Bhargava Signature.....
Date: Place:

If this consent process has been done in a language other than that on this written form with the assistance of a translator, indicate;

Language.....

Translator's name

Signature.....

Date:

Place: