

## PARTICIPANT/LAR INFORMED CONSENT FORM (PICF)

Protocol / Study number: \_\_\_\_\_

Participant identification number for this trial: \_\_\_\_\_

**Title of project:** "Thromboelastography-guided blood product transfusion in cirrhosis patients with acute variceal bleeding: a Randomized Controlled Trial".

**Name of Principal Investigator:** Dr Shalimar, Phone number- 09868397211

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: 14-12-2020

Place: DELHI

(Signatures / Left Thumb Impression)

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.

\_\_\_\_\_

Signatures of the Principal Investigator

Date: 14-12-2020

Place: DELHI

2) Witness - 2

Signatures

Address:

\_\_\_\_\_

1) Witness - 1

Signatures

Name \_\_\_\_\_

Address: \_\_\_\_\_