

Annexure 1

**Patient Consent Form**

**TITLE OF STUDY: “A Comparative study between intracuff Lidocaine and Alkalised Lidocaine for Sedation and Analgesia Requirements in Mechanically Ventilated Patient in UPUMS Saifai, Etawah”**

Investigators:

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**PATIENT INFORMATION:**

We are conducting above mentioned study in this institute. In this study we compare the compare intracuff lidocaine and alkalised Lidocaine for sedation and analgesia requirements in mechanically ventilated patients in ICU. There is no apparent danger due to this study but known complications include numbness, tongue parasthesia, tinnitus, nervousness, twitching of muscles, seizures, hypersensitivity reaction, decreased thrombosis, and platelet aggregation. These complications are readily treatable. Any complication arising due to the study will be treated free of cost, but there is provision for financial compensation if the institute agrees.

Your participation in this study is completely voluntary and you can choose to withdraw from the study at any point without any fear of adverse treatment.

**PART 2: CONSENT**

I have been told/read and understood the above mentioned information about the study participation. I have no further doubts about my participation and my participation is completely voluntary. I give my permission for enrolling me in this study

Name of participant:

Date:

Signature of participant:

Place:

Name & Relation of witness I

Date:

Signature of witness:

Place:

Name and relation of witness II

Date

Signature of witness:

Place: