CONSENT FORM

PART I

INFORMATION FOR PARTICIPANTS OF THE STUDY

Instructions - This is the patient information sheet. It should address the participant of this study. Depending upon the nature of the individual project, the details provided to the participant may vary. A separate consent form for the patient/test group and control (drug/procedure or placebo) should be provided as applicable. While formulating this sheet, the investigator must provide the following information as applicable in a simple language in English and Hindi which can be understood by the participant

- Title of the project
- Name of the investigator/guide
- Purpose of this project/study
- Procedure/methods of the study
- Expected duration of the subject participation
- The benefits to be expected from the research to the participant or to others and the post trial responsibilities of the investigator
- Any risks expected from the study to the participant
- Maintenance of confidentiality of records
- Provision of free treatment for research related injury
- Compensation of the participants not only for disability or death resulting from such injury but also for unforeseeable risks.
- Freedom to withdraw from the study at any time during the study period without the loss of benefits that the participant would otherwise be entitled
- Possible current and future uses of the biological material and of the data to be generated from the research and if the material is likely to be used for secondary purposes or would be shared with others, this should be mentioned
- Address and telephone number of the investigator and co-investigator/guide
- The patient information sheet must be duly signed by the investigator

CONSENT FORM

PART II

CONSENT FORM (for the subject)

The advantages and disadvantages of the research in which I am expected to participate and/or for which I have to donate blood/tissue has been explained to me.

I willingly, under no pressure from the researcher (Please ✔)

☐ agree to take part in this research, and agree to participate in all investigations which will help acquire knowledge for the benefit of the mankind,
☐ agree to donate my blood/ tissue
☐ I am agree to use my survey data for research purposes

My consent is explicitly not for disclosing any personal information. For disclosing any such personal information obtained from the investigations conducted on my samples, further consent should be obtained.

I have been informed that the guide/ researchers/ PI and her/his research student will take my prior consent before they draw benefits from research based on my samples.

Signatures:

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Subject Witness Principle Investigator

Mobile number…