Effect of probe material (metallic vs plastic) on peri-implant probing: a randomized clinical trial

NCT ID not yet assigned

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INFORMED CONSENT OF THE STUDY PARTICIPANT

Study Title: *"Effect of Probe Material (Metallic vs. Plastic) on Peri-implant Probing: A Randomized Clinical Trial."*

Mr./Ms. ..... as the participant with ID number .....or as the legal representative, family member, etc. of the participant, Mr./Ms. .....with ID number .....

I DECLARE that Dr. / Dr. ..... ,
(First name and two surnames of the investigator providing the information)
has given me sufficient information about the study and has provided me with the corresponding Participant Information Sheet.

I have understood the explanations given to me in clear and simple language, and the clinician who attended me has allowed me to make all the remarks I considered necessary and has clarified all the questions I raised.

I also understand that, at any moment and without the need to give any explanation, I may revoke the consent I now give and I may withdraw from the study whenever I wish, without having to provide reasons and without any consequences.

Therefore, I state that I am satisfied with the information received and that I give my consent to participate in this study.

Signature	Signature	Signature
Investigator	Participant	Participant's legal representative

Hospitalet de Llobregat, ... day of ....., 20 .....

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INFORMED CONSENT OF THE STUDY PARTICIPANT

WITHDRAWAL OF CONSENT

Mr./Ms. ..... as a participant with ID number .....or as the legal representative, family member, etc. of the participant, Mr./Ms. ..... with ID number .....

I DECLARE that:

Dr./ Dr. .....

has correctly informed me about the study:.....

.....

I WITHDRAW my consent.

Hospitalet de Llobregat, ... day of ....., 20 .....

Signature	Signature	Signature
Investigator	Participant	Participant's legal representative

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