

# Title: Prospective outcomes of management of third molar extractions via a large multicenter study

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## PROTOCOL SIGNATURE PAGE

I agree to conduct this clinical study in accordance with the design and specific provisions of this protocol and will only make changes in the protocol after notifying the sponsor.

I understand that I may terminate or suspend enrollment of the study at any time if it becomes necessary to protect the best interests of the study subjects. This study may be terminated by American Association of Oral and Maxillofacial Surgeons (AAOMS), with or without cause.

I agree to personally conduct or supervise this investigation and to ensure that all associates, colleagues, and employees assisting in the conduct of this study are informed about their obligations in meeting these commitments.

I will conduct the study in accordance with Good Clinical Practice, the Declaration of Helsinki, and the moral, ethical and scientific principles that justify medical research. The study will be conducted in accordance with all relevant laws and regulations relating to clinical studies and the protection of patients.

I agree to maintain adequate and accurate records and to make those records available for audit and inspection in accordance with relevant regulatory requirements.

I will ensure that the requirements relating to Institutional Review Board (IRB) review and approval are met. I will provide AAOMS with any material which is to be provided to obtain IRB approval. I agree to promptly report to the AAOMS any changes in the research activity and all unanticipated problems involving risks to human subjects or others. Additionally, I will not make any changes in the research without AAOMS or IRB approval, except where necessary to ensure the safety of study participants.

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Name

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Signature

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Date