

Section 2. Subject of the substantial modification

<p>Please provide a short rationale of this substantial modification</p>	
<p>Is this substantial modification likely to have an impact on subjects participating in the performance study? (Select all that apply)</p> <ul style="list-style-type: none">Rights of subjectsSafety of subjectsHealth of subjectsOtherNo impact on the subjects	
<p>Do you consider this substantial modification will likely have an impact on generated clinical data? (Select all that apply)</p> <ul style="list-style-type: none">Robustness of clinical data generated by the performance studyReliability of clinical data generated by the investigationOtherNo impact on clinical data	

Please use the template in [MDCG 2022-X](#) named “appendix of documents to attach” to identify clearly which documents are being proposed for modification with this substantial modification.

I hereby certify that the information and documentation submitted with this substantial modification is correct in detail and all the information requested has been supplied. The investigated device complies with the applicable general safety and performance requirements, apart from those covered by the investigation and that every precaution has been taken to protect the health and safety of the patient and/or user. I confirm that all the performance study information collected for this notification, has been done in compliance with the European data protection legislation (GDPR).

Date

Name

Position
