## Performance study – Substantial modification of performance study under In Vitro Medical Device Regulation.

## **Notification form Version**

## Section 1. Identification of the performance study

Please provide the performance study ID (CIV-ID)	
Does this substantial modification relate	Yes
to a performance study that is currently suspended/stopped?	No
How many patients have been recruited	in the performance study
Worldwide	
Europe	
In the Member State you are submit	ting this substantial modification
Select the Member State where this pe	rformance study is ongoing:

## Section 2. Subject of the substantial modification

Please provide a short rationale					
of this substantial modification					
Is this substantial modification lik	ely to have an impact on subjects participating in the				
performance study? (Select all th					
Rights of subjects					
<b>G,</b>					
Safety of subjects					
Health of subjects					
Other					
Other					
No impact on the subjects	No impact on the subjects				
Do you consider this substantial r	nodification will likely have an impact on generated clinical data?				
(Select all that apply)	mountained with the city mate an impact on generated chinical data.				
Robustness of clinical performance data generated by the performance study					
Reliability of clinical performance data generated by the performance study					
Othor					
Other					
No impact on clinical data					
, , , , , , , , , , , , , , , , , , ,					

Please use the template in 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 device for performance complies with the applicable general safety and performance requirements, apart from those covered by the performance study 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			
D. attitus			