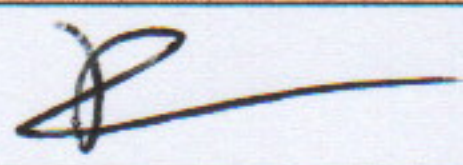


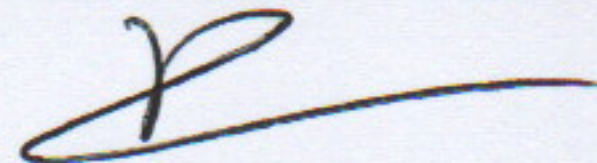
PHARMACY DEPARTMENT

ASSURANCE OF QP CERTIFICATION DOCUMENTATION

TRIAL NAME / PROTOCOL NUMBER		REMAP-CAP
COMPANY CARRYING OUT QP CERTIFICATION	COMPANY NAME	Sanofi
	SPONSOR or MANUFACTURER (please tick)	SPONSOR <input type="checkbox"/> MANUFACTURER <input checked="" type="checkbox"/>
	COMPANY ADDRESS	sanofi R&D
		371, rue du Prof. J.Blaiac
		34184 Montpellier cedex 04
France		
COUNTRY IN WHICH QP CERTIFICATION TAKES PLACE		France

The MANUFACTURER holds copies of the QP Certification documentation and have an appropriate system in place to track these to the drug at site.	SIGN	
	PRINT	Rocio CUADRADO
	DATE	14/MAY/2020
	POSITION	Sanofi Qualified Person

The SPONSOR warrants that any changes to QP Certification will follow all due legal process and will maintain MHRA approval and compliance with cGMP.	SIGN	
	PRINT	
	DATE	
	POSITION	

The MANUFACTURER certifies that all clinical trial material supplied for the study named above is free from Transmissible Spongiform Encephalopathies (TSE).	SIGN	
	PRINT	Rocio CUADRADO
	DATE	14/MAY/2020
	POSITION	Sanofi Qualified Person

***Signatures must be from the Sponsor or Manufacturer (final QP releasing company) not from any other party.**

Once completed, please return to

Email: @nhs.net

Fax: