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In reply please
refer to:

Your reference: EMA/308793/2020

Executive Director Guido Rasi
European Medicines Agency
Domenico Scarlattilaan 6
NL-1083 HS Amsterdam

27 October 2020

Dear Professor Rasi,

Thank you for your letter EMA/308793/2020 outlining the conditions for making available reports of suspected adverse reactions by EMA to WHO/UMC in accordance with Article 28c(1), 2nd paragraph of Regulation (EC) No 726/2004. I acknowledge receipt and confirm that we accept the modalities and conditions detailed in the letter for the continued transfer of data between the European Medicines Agency (EMA) and the Uppsala Monitoring Centre (UMC).

I would also like to thank you for the continued collaboration on this important project and look forward to future EMA's continued cooperation with WHO and UMC.

Yours sincerely,

Emer Cooke
Director
Regulation and Prequalification