N° d'enregistrement :
Ares(2013)3462610

Subject: RE: Possibiity to respond to Revision of EU Commission guidelines on Good Manufacturing Practice for Medicinal Products; Annex 16

Dear Sir/Madam,

With apologies for delayed information, it has been decided Teva has no comments to the EC consultation on the Guidelines on Good Manufacturing Practice for Medicinal Products; Annex 16 Certification by a Qualified Person and Batch Release. Thank you for giving us the extra time.

Yours sincerely,