LETTER OF ANNOUNCEMENT
European Commission Q&A Version 17

11th March 2020

Dear all,

EMVO would like to kindly inform you that the European Commission has published a new version of the Questions & Answers document on "Safety Features for Medicinal Products for Human Use (V.17)".

The document is available on EMVO's website in the Knowledge Database section – Documents Overview. The Q&A document can be consulted here.

Please note that the modifications from the previous version have been highlighted for your convenience. The changes include the addition of a new Q&A (5.11) and the revision of Q&A 1.22.

In the event of any questions or uncertainty, please do not hesitate to contact our Helpdesk:

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