LETTER OF ANNOUNCEMENT
European Commission Q&A Version 19

17th January 2022

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.19)".

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 (1.29) and the revision of Q&As 1.22, 5.8 and 7.19.

Should you have any questions, please do not hesitate to contact our Helpdesk.

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