LETTER OF ANNOUNCEMENT
European Commission Q&A Version 20

7th July 2022

EMVO would like to inform you that the European Commission has published a new version of the Questions & Answers (Q&A) document on "**Safety Features for Medicinal Products for Human Use (V.20)**".

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

Please note that the modifications from the previous version have been highlighted for your convenience and include the addition of a new Q&A (5.14).

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

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