EUROPEAN
COMMISSION

Brussels, 27.4.2023
COM(2023) 222 final

ANNEX 2

**ANNEX**

**to the**

**Proposal for a Regulation of the European Parliament and of the Council**

**on the unitary supplementary certificate for medicinal products, and amending Regulation (EU) 2017/1001, Regulation (EC) No 1901/2006 as well as Regulation (EU) No 608/2013**

{SEC(2023) 172 final} - {SWD(2023) 117 final} - {SWD(2023) 118 final} - {SWD(2023) 119 final}

**ANNEX II**

Standard form for notification pursuant to Article 5(3), points (b) and (c).

| | | | |
|---|---|---|---|
| Tick the appropriate box | | □ New notification<br>□ Update of an existing notification | |
| (a) | Name and address of the maker | … | |
| (b) | Purpose of making | □ Export<br>□ Storing<br>□ Export and storing | |
| (c) | Member State in which making is to take place and Member State in which first related act (if any) prior to making is to take place | Member State of making | |
| | | (Member State of first related act (if any)) | |
| (d) | Number of unitary certificate having effect in the Member State of making and number of certificate granted in Member State of first related act (if any) prior to making | Unitary certificate having effect in the Member State of making | |
| | | (Certificate of Member State of first related act (if any)) | |
| (e) | For medicinal products to be exported to third countries, reference number of marketing authorisation, or the equivalent of such authorisation, in each third country of export | | |
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