REGULATION (EU) 2019/933 OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL

of 20 May 2019

amending Regulation (EC) No 469/2009 concerning the supplementary protection certificate for medicinal products

Standard form for notification pursuant to points (b) and (c) of Article 5(2)

<table>
<tr><td>Tick the appropriate box</td><td colspan="2">☐ New notification
☐ Update of an existing notification</td></tr>
<tr><td>(a) Name and address of the maker</td><td colspan="2"></td></tr>
<tr><td>(b) Purpose of making</td><td colspan="2">☐ Export
☐ Storing
☐ Export and storing</td></tr>
<tr><td rowspan="2">(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</td><td>Member State of making</td><td></td></tr>
<tr><td>(Member State of first related act (if any))</td><td></td></tr>
<tr><td rowspan="2">(d) Number of certificate granted in the Member State of making and number of certificate granted in Member State of first related act (if any) prior to making</td><td>Certificate of Member State of making</td><td></td></tr>
<tr><td>(Certificate of Member State of first related act (if any))</td><td></td></tr>
<tr><td rowspan="3">(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</td><td colspan="2"></td></tr>
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