**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)***

|  |  |  |
| --- | --- | --- |
| **Tick the appropriate box** | New notification  Update of an existing notification | |
| 1. Name and address of the maker |  | |
| 1. Purpose of making | Export  Storing  Export and storing | |
| 1. 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)) |  |
| 1. 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 | Certificate of Member State of making |  |
| (Certificate of Member State of first related act (if any)) |  |
| 1. 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 |  | |
|  | |
|  | |