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| **NATIONAL MARKETING AUTHORISATION** | |
| **MA number** |  |
| **Product name** |  |
| **Pharmaceutical form** |  |
| **Strength** |  |
| **MAH\*** |  |
| **Indication\*\*** |  |
| **Manufacturer responsible for batch release** |  |
| **Manufacturer responsible for quality control** |  |
| **Manufacturer of the medicinal product** |  |
| **Manufacturer responsible for packaging (primary, secondary)** |  |
| **Manufacturer of the active substance** |  |
| **Legal status** |  |
| **Type of immediate packaging** |  |

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| **SÚKL code\*\*\*** | **Pack size** | **SÚKL code\*\*\*** | **Pack size** |
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| **MRP/DCP MARKETING AUTHORISATION** | | | |
| **MA number** |  | | |
| **Product name** |  | | |
| **Pharmaceutical form** |  | | |
| **Strength** |  | | |
| **MAH\*** |  | | |
| **Indication\*\*** |  | | |
| **Manufacturer responsible for batch release** |  | | |
| **Manufacturer responsible for quality control** |  | | |
| **Manufacturer of the medicinal product** |  | | |
| **Manufacturer responsible for packaging (primary, secondary)** |  | | |
| **Manufacturer of the active substance** |  | | |
| **Legal status** |  | | |
| **Type of immediate packaging** |  | | |
| **SÚKL code\*\*\*** | **Pack size** | **SÚKL code\*\*\*** | **Pack size** |
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| \* MAH must be identical. | | | |
| \*\* The last version of SmPC should be provided. | |  |  |
| \*\*\*Please state SÚKL code assigned as part of the national MA, if the code is to be maintained for the relevant product presentation. If product presentation from MR/DC procedure is to be maintained and is not part of national MA, new SÚKL code will be assigned. | | | |