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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.  |