| **<EMA procedure number>** **<MA (EU) number>** | **(Invented) name** | **Strength** | **Pharmaceutical form** | **Route of administration** | **Immediate Packaging** | **<Content (concentration)>** | **Pack size** |
| --- | --- | --- | --- | --- | --- | --- | --- |
|  |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |

**NOTE:**

**This is a sample English Annex A template only.**

The English Annex A template is not provided as it is prepared by the European Medicines Agency. Translations of Annex A must be in line and consistent with the adopted English product information annexes of the product concerned.