

<u>MA (EU) number</u>	<u>(Invented) name</u>	<u>Strength</u>	<u>Pharmaceutical form</u>	<u>Route of administration</u>	<u>Immediate Packaging</u>	<u>Pack size</u>
EU/1/22/1714/001	Pombiliti	105 mg	Powder for concentrate for solution for infusion	Intravenous use	vial (glass)	1 vial
EU/1/22/1714/002	Pombiliti	105 mg	Powder for concentrate for solution for infusion	Intravenous use	vial (glass)	10 vials
EU/1/22/1714/003	Pombiliti	105 mg	Powder for concentrate for solution for infusion	Intravenous use	vial (glass)	25 vials

**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.