

<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/1/06/363/001	Sprycel	20 mg	Film-coated tablet	Oral use	bottle (HDPE)	60 tablets
EU/1/06/363/002	Sprycel	50 mg	Film-coated tablet	Oral use	bottle (HDPE)	60 tablets
EU/1/06/363/003	Sprycel	70 mg	Film-coated tablet	Oral use	bottle (HDPE)	60 tablets
EU/1/06/363/004	Sprycel	20 mg	Film-coated tablet	Oral use	blister (alu/alu)	56 tablets
EU/1/06/363/005	Sprycel	50 mg	Film-coated tablet	Oral use	blister (alu/alu)	56 tablets
EU/1/06/363/006	Sprycel	70 mg	Film-coated tablet	Oral use	blister (alu/alu)	56 tablets
EU/1/06/363/007	Sprycel	20 mg	Film-coated tablet	Oral use	blister (alu/alu)	60 x 1 tablets (unit dose)
EU/1/06/363/008	Sprycel	50 mg	Film-coated tablet	Oral use	blister (alu/alu)	60 x 1 tablets (unit dose)
EU/1/06/363/009	Sprycel	70 mg	Film-coated tablet	Oral use	blister (alu/alu)	60 x 1 tablets (unit dose)
EU/1/06/363/010	Sprycel	100 mg	Film-coated tablet	Oral use	bottle (HDPE)	30 tablets
EU/1/06/363/011	Sprycel	100 mg	Film-coated tablet	Oral use	blister (alu/alu)	30 x 1 tablets (unit dose)
EU/1/06/363/012	Sprycel	80 mg	Film-coated tablet	Oral use	bottle (HDPE)	30 tablets
EU/1/06/363/013	Sprycel	80 mg	Film-coated tablet	Oral use	blister (alu/alu)	30 x 1 tablets (unit dose)
EU/1/06/363/014	Sprycel	140 mg	Film-coated tablet	Oral use	bottle (HDPE)	30 tablets
EU/1/06/363/015	Sprycel	140 mg	Film-coated tablet	Oral use	blister (alu/alu)	30 x 1 tablets (unit dose)
EU/1/06/363/016	Sprycel	10 mg/ml	Powder for oral suspension	Oral use	bottle (HDPE)	1 bottle + 1 oral syringe + 1 press-in bottle adapter