

<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/1/17/1248/001	Darunavir Krka d.d.	400 mg	Film-coated tablet	Oral use	bottle (HDPE)	30 tablets
EU/1/17/1248/002	Darunavir Krka d.d.	400 mg	Film-coated tablet	Oral use	bottle (HDPE)	60 (2 x 30) tablets
EU/1/17/1248/003	Darunavir Krka d.d.	400 mg	Film-coated tablet	Oral use	bottle (HDPE)	90 (3 x 30) tablets
EU/1/17/1248/004	Darunavir Krka d.d.	400 mg	Film-coated tablet	Oral use	bottle (HDPE)	180 (6 x 30) tablets
EU/1/17/1248/005	Darunavir Krka d.d.	600 mg	Film-coated tablet	Oral use	bottle (HDPE)	30 tablets
EU/1/17/1248/006	Darunavir Krka d.d.	600 mg	Film-coated tablet	Oral use	bottle (HDPE)	60 (2 x 30) tablets
EU/1/17/1248/007	Darunavir Krka d.d.	600 mg	Film-coated tablet	Oral use	bottle (HDPE)	90 (3 x 30) tablets
EU/1/17/1248/008	Darunavir Krka d.d.	600 mg	Film-coated tablet	Oral use	bottle (HDPE)	180 (6 x 30) tablets
EU/1/17/1248/009	Darunavir Krka d.d.	800 mg	Film-coated tablet	Oral use	bottle (HDPE)	30 tablets
EU/1/17/1248/010	Darunavir Krka d.d.	800 mg	Film-coated tablet	Oral use	bottle (HDPE)	90 (3 x 30) tablets