1 September 2017
EMA/593213/2017
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance(s): furosemide / spironolactone

Procedure No.: PSUSA/00001493/201612

30 Churchill Place ● Canary Wharf ● London E14 5EU ● United Kingdom
Telephone +44 (0)20 3660 6000 **Facsimile** +44 (0)20 3660 5555
Send a question via our website www.ema.europa.eu/contact

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
LASILACTON 20 MG/100 MG-KAPSELN	not available	17.056	SANOFI-AVENTIS GMBH OSTERREICH	AT
LASILACTON 20 MG/100 MG-KAPSELN	not available	17.056	SANOFI-AVENTIS GMBH OSTERREICH	AT
LASILACTON 20 MG/50 MG-KAPSELN	not available	17.055	SANOFI-AVENTIS GMBH OSTERREICH	AT
LASILACTON 20 MG/50 MG-KAPSELN	not available	17.055	SANOFI-AVENTIS GMBH OSTERREICH	AT
LASILACTONE 20MG/50MG CAPSULES	not available	PL 04425/0372	AVENTIS PHARMA LTD	UK
LASILACTONE 20MG/50MG CAPSULES	not available	PL 04425/0372	AVENTIS PHARMA LTD	UK
LASITONE	not available	023770011	SANOFI SPA	IT
OSYROL 100-LASIX	not available	474.01.01	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
OSYROL 100-LASIX	not available	474.01.01	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
OSYROL 100-LASIX	not available	474.01.01	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
OSYROL 100-LASIX	not available	474.01.01	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Osyrol 100-Lasix (100 mg/20 mg Hartkapseln)	not available	474.01.01	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
OSYROL 50-LASIX	not available	474.00.01	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
OSYROL 50-LASIX	not available	474.00.01	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
OSYROL 50-LASIX	not available	474.00.01	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
OSYROL 50-LASIX	not available	474.00.01	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
OSYROL 50-LASIX	not available	474.00.01	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Spiro comp. forte-ratiopharm® 100 mg/20 mg Filmtabletten	not available	2153.01.00	RATIOPHARM GMBH	DE
Spiro comp. forte-ratiopharm® 100 mg/20	not available	0687/05098248	RATIOPHARM GMBH	LU

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
mg Filmtabletten				
SPIROFUR 50 mg + 20 mg capsule rigide	not available	023749056	BRUNO FARMACEUTICI	IT
SPIROFUR 50 mg + 20 mg capsule rigide	not available	023749017	BRUNO FARMACEUTICI	IT