25 June 2026
EMADOC-1700519818-3369884
Human Medicines Division

List of nationally authorised medicinal products

Active substance(s): bisoprolol

EURD list No. PSUSA/00000419/202509

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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
Concor 5 mg Filmtabletten	not available	1-18587	MERCK GESELLSCHAFT MBH	AT
Concor 10 mg Filmtabletten	not available	1-18586	MERCK GESELLSCHAFT MBH	AT
Concor 10 mg Filmtabletten	not available	1-18586	MERCK GESELLSCHAFT MBH	AT
Concor 5 mg Filmtabletten	not available	1-18587	MERCK GESELLSCHAFT MBH	AT
Bisoprolol Viatris 10 mg Filmtabletten	not available	1-22023	VIATRIS LIMITED	AT
Bisoprolol Viatris 5 mg Filmtabletten	not available	1-22024	VIATRIS LIMITED	AT
Concor Cor 1,25 mg Filmtabletten	SE/H/0184/001	1-23302	MERCK GESELLSCHAFT MBH	AT
Concor Cor 1,25 mg Filmtabletten	SE/H/0184/001	1-23302	MERCK GESELLSCHAFT MBH	AT
Concor Cor 1,25 mg Filmtabletten	SE/H/0184/001	1-23302	MERCK GESELLSCHAFT MBH	AT
Concor Cor 1,25 mg Filmtabletten	SE/H/0184/001	1-23302	MERCK GESELLSCHAFT MBH	AT
Concor Cor 1,25 mg Filmtabletten	SE/H/0184/001	1-23302	MERCK GESELLSCHAFT MBH	AT
Concor Cor 1,25 mg Filmtabletten	SE/H/0184/001	1-23302	MERCK GESELLSCHAFT MBH	AT
Concor Cor 1,25 mg Filmtabletten	SE/H/0184/001	1-23302	MERCK GESELLSCHAFT MBH	AT
Concor Cor 1,25 mg Filmtabletten	SE/H/0184/001	1-23302	MERCK GESELLSCHAFT MBH	AT
Concor Cor 1,25 mg Filmtabletten	SE/H/0184/001	1-23302	MERCK GESELLSCHAFT MBH	AT
Concor Cor 1,25 mg Filmtabletten	SE/H/0184/001	1-23302	MERCK GESELLSCHAFT MBH	AT
Concor Cor 1,25 mg Filmtabletten	SE/H/0184/001	1-23302	MERCK GESELLSCHAFT MBH	AT
Concor Cor 1,25 mg	SE/H/0184/001	1-23302	MERCK GESELLSCHAFT MBH	AT

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
Filmtabletten				
Concor Cor 1,25 mg Filmtabletten	SE/H/0184/001	1-23302	MERCK GESELLSCHAFT MBH	AT
Concor Cor 1,25 mg Filmtabletten	SE/H/0184/001	1-23302	MERCK GESELLSCHAFT MBH	AT
Concor Cor 1,25 mg Filmtabletten	SE/H/0184/001	1-23302	MERCK GESELLSCHAFT MBH	AT
Concor Cor 1,25 mg Filmtabletten	SE/H/0184/001	1-23302	MERCK GESELLSCHAFT MBH	AT
Concor Cor 2,5 mg Filmtabletten	SE/H/0184/002	1-23303	MERCK GESELLSCHAFT MBH	AT
Concor Cor 2,5 mg Filmtabletten	SE/H/0184/002	1-23303	MERCK GESELLSCHAFT MBH	AT
Concor Cor 2,5 mg Filmtabletten	SE/H/0184/002	1-23303	MERCK GESELLSCHAFT MBH	AT
Concor Cor 2,5 mg Filmtabletten	SE/H/0184/002	1-23303	MERCK GESELLSCHAFT MBH	AT
Concor Cor 2,5 mg Filmtabletten	SE/H/0184/002	1-23303	MERCK GESELLSCHAFT MBH	AT
Concor Cor 2,5 mg Filmtabletten	SE/H/0184/002	1-23303	MERCK GESELLSCHAFT MBH	AT
Concor Cor 2,5 mg Filmtabletten	SE/H/0184/002	1-23303	MERCK GESELLSCHAFT MBH	AT
Concor Cor 2,5 mg Filmtabletten	SE/H/0184/002	1-23303	MERCK GESELLSCHAFT MBH	AT
Concor Cor 2,5 mg Filmtabletten	SE/H/0184/002	1-23303	MERCK GESELLSCHAFT MBH	AT
Concor Cor 2,5 mg Filmtabletten	SE/H/0184/002	1-23303	MERCK GESELLSCHAFT MBH	AT
Concor Cor 2,5 mg Filmtabletten	SE/H/0184/002	1-23303	MERCK GESELLSCHAFT MBH	AT
Concor Cor 2,5 mg Filmtabletten	SE/H/0184/002	1-23303	MERCK GESELLSCHAFT MBH	AT
Concor Cor 2,5 mg Filmtabletten	SE/H/0184/002	1-23303	MERCK GESELLSCHAFT MBH	AT

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
Concor Cor 2,5 mg Filmtabletten	SE/H/0184/002	1-23303	MERCK GESELLSCHAFT MBH	AT
Concor Cor 2,5 mg Filmtabletten	SE/H/0184/002	1-23303	MERCK GESELLSCHAFT MBH	AT
Concor Cor 2,5 mg Filmtabletten	SE/H/0184/002	1-23303	MERCK GESELLSCHAFT MBH	AT
Concor Cor 3,75 mg Filmtabletten	SE/H/0184/003	1-23304	MERCK GESELLSCHAFT MBH	AT
Concor Cor 3,75 mg Filmtabletten	SE/H/0184/003	1-23304	MERCK GESELLSCHAFT MBH	AT
Concor Cor 3,75 mg Filmtabletten	SE/H/0184/003	1-23304	MERCK GESELLSCHAFT MBH	AT
Concor Cor 3,75 mg Filmtabletten	SE/H/0184/003	1-23304	MERCK GESELLSCHAFT MBH	AT
Concor Cor 3,75 mg Filmtabletten	SE/H/0184/003	1-23304	MERCK GESELLSCHAFT MBH	AT
Concor Cor 3,75 mg Filmtabletten	SE/H/0184/003	1-23304	MERCK GESELLSCHAFT MBH	AT
Concor Cor 3,75 mg Filmtabletten	SE/H/0184/003	1-23304	MERCK GESELLSCHAFT MBH	AT
Concor Cor 3,75 mg Filmtabletten	SE/H/0184/003	1-23304	MERCK GESELLSCHAFT MBH	AT
Concor Cor 3,75 mg Filmtabletten	SE/H/0184/003	1-23304	MERCK GESELLSCHAFT MBH	AT
Concor Cor 3,75 mg Filmtabletten	SE/H/0184/003	1-23304	MERCK GESELLSCHAFT MBH	AT
Concor Cor 3,75 mg Filmtabletten	SE/H/0184/003	1-23304	MERCK GESELLSCHAFT MBH	AT
Concor Cor 3,75 mg Filmtabletten	SE/H/0184/003	1-23304	MERCK GESELLSCHAFT MBH	AT
Concor Cor 3,75 mg Filmtabletten	SE/H/0184/003	1-23304	MERCK GESELLSCHAFT MBH	AT
Concor Cor 3,75 mg Filmtabletten	SE/H/0184/003	1-23304	MERCK GESELLSCHAFT MBH	AT
Concor Cor 3,75 mg	SE/H/0184/003	1-23304	MERCK GESELLSCHAFT MBH	AT

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
Filmtabletten				
Concor Cor 3,75 mg Filmtabletten	SE/H/0184/003	1-23304	MERCK GESELLSCHAFT MBH	AT
Concor Cor 5 mg Filmtabletten	SE/H/0184/004	1-23305	MERCK GESELLSCHAFT MBH	AT
Concor Cor 5 mg Filmtabletten	SE/H/0184/004	1-23305	MERCK GESELLSCHAFT MBH	AT
Concor Cor 5 mg Filmtabletten	SE/H/0184/004	1-23305	MERCK GESELLSCHAFT MBH	AT
Concor Cor 5 mg Filmtabletten	SE/H/0184/004	1-23305	MERCK GESELLSCHAFT MBH	AT
Concor Cor 5 mg Filmtabletten	SE/H/0184/004	1-23305	MERCK GESELLSCHAFT MBH	AT
Concor Cor 5 mg Filmtabletten	SE/H/0184/004	1-23305	MERCK GESELLSCHAFT MBH	AT
Concor Cor 5 mg Filmtabletten	SE/H/0184/004	1-23305	MERCK GESELLSCHAFT MBH	AT
Concor Cor 5 mg Filmtabletten	SE/H/0184/004	1-23305	MERCK GESELLSCHAFT MBH	AT
Concor Cor 5 mg Filmtabletten	SE/H/0184/004	1-23305	MERCK GESELLSCHAFT MBH	AT
Concor Cor 5 mg Filmtabletten	SE/H/0184/004	1-23305	MERCK GESELLSCHAFT MBH	AT
Concor Cor 5 mg Filmtabletten	SE/H/0184/004	1-23305	MERCK GESELLSCHAFT MBH	AT
Concor Cor 5 mg Filmtabletten	SE/H/0184/004	1-23305	MERCK GESELLSCHAFT MBH	AT
Concor Cor 5 mg Filmtabletten	SE/H/0184/004	1-23305	MERCK GESELLSCHAFT MBH	AT
Concor Cor 5 mg Filmtabletten	SE/H/0184/004	1-23305	MERCK GESELLSCHAFT MBH	AT
Concor Cor 5 mg Filmtabletten	SE/H/0184/004	1-23305	MERCK GESELLSCHAFT MBH	AT
Concor Cor 5 mg Filmtabletten	SE/H/0184/004	1-23305	MERCK GESELLSCHAFT MBH	AT

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
Concor Cor 7,5 mg Filmtabletten	SE/H/0184/005	1-23306	MERCK GESELLSCHAFT MBH	AT
Concor Cor 7,5 mg Filmtabletten	SE/H/0184/005	1-23306	MERCK GESELLSCHAFT MBH	AT
Concor Cor 7,5 mg Filmtabletten	SE/H/0184/005	1-23306	MERCK GESELLSCHAFT MBH	AT
Concor Cor 7,5 mg Filmtabletten	SE/H/0184/005	1-23306	MERCK GESELLSCHAFT MBH	AT
Concor Cor 7,5 mg Filmtabletten	SE/H/0184/005	1-23306	MERCK GESELLSCHAFT MBH	AT
Concor Cor 7,5 mg Filmtabletten	SE/H/0184/005	1-23306	MERCK GESELLSCHAFT MBH	AT
Concor Cor 7,5 mg Filmtabletten	SE/H/0184/005	1-23306	MERCK GESELLSCHAFT MBH	AT
Concor Cor 7,5 mg Filmtabletten	SE/H/0184/005	1-23306	MERCK GESELLSCHAFT MBH	AT
Concor Cor 7,5 mg Filmtabletten	SE/H/0184/005	1-23306	MERCK GESELLSCHAFT MBH	AT
Concor Cor 7,5 mg Filmtabletten	SE/H/0184/005	1-23306	MERCK GESELLSCHAFT MBH	AT
Concor Cor 7,5 mg Filmtabletten	SE/H/0184/005	1-23306	MERCK GESELLSCHAFT MBH	AT
Concor Cor 7,5 mg Filmtabletten	SE/H/0184/005	1-23306	MERCK GESELLSCHAFT MBH	AT
Concor Cor 7,5 mg Filmtabletten	SE/H/0184/005	1-23306	MERCK GESELLSCHAFT MBH	AT
Concor Cor 7,5 mg Filmtabletten	SE/H/0184/005	1-23306	MERCK GESELLSCHAFT MBH	AT
Concor Cor 7,5 mg Filmtabletten	SE/H/0184/005	1-23306	MERCK GESELLSCHAFT MBH	AT
Concor Cor 7,5 mg Filmtabletten	SE/H/0184/005	1-23306	MERCK GESELLSCHAFT MBH	AT
Concor Cor 10 mg Filmtabletten	SE/H/0184/006	1-23307	MERCK GESELLSCHAFT MBH	AT
Concor Cor 10 mg	SE/H/0184/006	1-23307	MERCK GESELLSCHAFT MBH	AT

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
Filmtabletten				
Concor Cor 10 mg Filmtabletten	SE/H/0184/006	1-23307	MERCK GESELLSCHAFT MBH	AT
Concor Cor 10 mg Filmtabletten	SE/H/0184/006	1-23307	MERCK GESELLSCHAFT MBH	AT
Concor Cor 10 mg Filmtabletten	SE/H/0184/006	1-23307	MERCK GESELLSCHAFT MBH	AT
Concor Cor 10 mg Filmtabletten	SE/H/0184/006	1-23307	MERCK GESELLSCHAFT MBH	AT
Concor Cor 10 mg Filmtabletten	SE/H/0184/006	1-23307	MERCK GESELLSCHAFT MBH	AT
Concor Cor 10 mg Filmtabletten	SE/H/0184/006	1-23307	MERCK GESELLSCHAFT MBH	AT
Concor Cor 10 mg Filmtabletten	SE/H/0184/006	1-23307	MERCK GESELLSCHAFT MBH	AT
Concor Cor 10 mg Filmtabletten	SE/H/0184/006	1-23307	MERCK GESELLSCHAFT MBH	AT
Concor Cor 10 mg Filmtabletten	SE/H/0184/006	1-23307	MERCK GESELLSCHAFT MBH	AT
Concor Cor 10 mg Filmtabletten	SE/H/0184/006	1-23307	MERCK GESELLSCHAFT MBH	AT
Concor Cor 10 mg Filmtabletten	SE/H/0184/006	1-23307	MERCK GESELLSCHAFT MBH	AT
Concor Cor 10 mg Filmtabletten	SE/H/0184/006	1-23307	MERCK GESELLSCHAFT MBH	AT
Concor Cor 10 mg Filmtabletten	SE/H/0184/006	1-23307	MERCK GESELLSCHAFT MBH	AT
Concor Cor 10 mg Filmtabletten	SE/H/0184/006	1-23307	MERCK GESELLSCHAFT MBH	AT
Concor 1,25 mg Filmtabletten	not available	138509	MERCK GESELLSCHAFT MBH	AT
Concor 1,25 mg Filmtabletten	not available	138509	MERCK GESELLSCHAFT MBH	AT
Concor 1,25 mg Filmtabletten	not available	138509	MERCK GESELLSCHAFT MBH	AT

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
Concor 1,25 mg Filmtabletten	not available	138509	MERCK GESELLSCHAFT MBH	AT
Concor 1,25 mg Filmtabletten	not available	138509	MERCK GESELLSCHAFT MBH	AT
Concor 1,25 mg Filmtabletten	not available	138509	MERCK GESELLSCHAFT MBH	AT
Concor 1,25 mg Filmtabletten	not available	138509	MERCK GESELLSCHAFT MBH	AT
Concor 2,5 mg Filmtabletten	not available	138510	MERCK GESELLSCHAFT MBH	AT
Concor 2,5 mg Filmtabletten	not available	138510	MERCK GESELLSCHAFT MBH	AT
Concor 2,5 mg Filmtabletten	not available	138510	MERCK GESELLSCHAFT MBH	AT
Concor 2,5 mg Filmtabletten	not available	138510	MERCK GESELLSCHAFT MBH	AT
Concor 2,5 mg Filmtabletten	not available	138510	MERCK GESELLSCHAFT MBH	AT
Concor 2,5 mg Filmtabletten	not available	138510	MERCK GESELLSCHAFT MBH	AT
Concor 2,5 mg Filmtabletten	not available	138510	MERCK GESELLSCHAFT MBH	AT
Concor 1,25 mg Filmtabletten	not available	138509	MERCK GESELLSCHAFT MBH	AT
Concor 2,5 mg Filmtabletten	not available	138510	MERCK GESELLSCHAFT MBH	AT
Bisoprolol Sandoz 10 mg filmomhulde tabletten	NL/H/0684/006	BE407513	SANDOZ N.V.	BE
Bisoprolol Sandoz 5 mg filmomhulde tabletten	NL/H/0684/004	BE327372	SANDOZ N.V.	BE
Bisoprolol Sandoz 5 mg filmomhulde tabletten	NL/H/0684/004	BE407495	SANDOZ N.V.	BE
Bisoprolol Sandoz 10 mg filmomhulde tabletten	NL/H/0684/006	BE327397	SANDOZ N.V.	BE
Конкор 5 mg	not available	20030463	MERCK, BULGARIA EAD	BG

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
филмирани таблетки				
Конкор 5 mg филмирани таблетки	not available	20030463	MERCK, BULGARIA EAD	BG
Конкор 5 mg филмирани таблетки	not available	20030463	MERCK, BULGARIA EAD	BG
Конкор 5 mg филмирани таблетки	not available	20030463	MERCK, BULGARIA EAD	BG
Конкор 10 mg филмирани таблетки	not available	20030464	MERCK, BULGARIA EAD	BG
Конкор 10 mg филмирани таблетки	not available	20030464	MERCK, BULGARIA EAD	BG
Конкор 10 mg филмирани таблетки	not available	20030464	MERCK, BULGARIA EAD	BG
Конкор 10 mg филмирани таблетки	not available	20030464	MERCK, BULGARIA EAD	BG
Конкор 10 mg филмирани таблетки	not available	20030464	MERCK, BULGARIA EAD	BG
Конкор 5 mg филмирани таблетки	not available	20030463	MERCK, BULGARIA EAD	BG
Конкор КОР 2,5 mg филмирани таблетки	not available	20060181	MERCK, BULGARIA EAD	BG
Конкор КОР 2,5 mg филмирани таблетки	not available	20060181	MERCK, BULGARIA EAD	BG
Конкор КОР 5 mg филмирани таблетки	not available	20060182	MERCK, BULGARIA EAD	BG
Конкор КОР 5 mg филмирани таблетки	not available	20060182	MERCK, BULGARIA EAD	BG
Конкор КОР 2,5 mg филмирани таблетки	not available	20060181	MERCK, BULGARIA EAD	BG
Конкор КОР 5 mg филмирани таблетки	not available	20060182	MERCK, BULGARIA EAD	BG
Concor 10 mg επικαλυμμένα με λεπτό υμένιο δισκία	not available	021708	MERCK A.E.	CY
Concor 5 mg	not available	12863	MERCK A.E.	CY

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
επικαλυμμένα με λεπτό υμένιο δισκία				
Emconcor® 5 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0187/004	22173	MERCK A.E.	CY
Emconcor® 5 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0187/004	22173	MERCK A.E.	CY
Emconcor® 5 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0187/004	22173	MERCK A.E.	CY
Emconcor® 5 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0187/004	22173	MERCK A.E.	CY
Emconcor® 5 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0187/004	22173	MERCK A.E.	CY
Emconcor® 5 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0187/004	22173	MERCK A.E.	CY
Emconcor® 5 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0187/004	22173	MERCK A.E.	CY
Emconcor® 2,5 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0187/002	022172	MERCK A.E.	CY
Emconcor® 2,5 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0187/002	022172	MERCK A.E.	CY
Emconcor® 2,5 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0187/002	022172	MERCK A.E.	CY
Emconcor® 2,5 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0187/002	022172	MERCK A.E.	CY

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
Emconcor® 2,5 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0187/002	022172	MERCK A.E.	CY
Emconcor® 2,5 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0187/002	022172	MERCK A.E.	CY
Emconcor® 2,5 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0187/002	022172	MERCK A.E.	CY
Emconcor® 2,5 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0187/002	022172	MERCK A.E.	CY
Emconcor® 5 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0187/004	22173	MERCK A.E.	CY
Concor COR 1,25 mg potahované tablety	not available	77-503-19-C	MERCK SPOL.S.R.O.	CZ
Concor COR 1,25 mg potahované tablety	not available	77-503-19-C	MERCK SPOL.S.R.O.	CZ
Concor COR 1,25 mg potahované tablety	not available	77-503-19-C	MERCK SPOL.S.R.O.	CZ
Concor COR 7,5 mg potahované tablety	not available	77-504-19-C	MERCK SPOL.S.R.O.	CZ
Concor COR 7,5 mg potahované tablety	not available	77-504-19-C	MERCK SPOL.S.R.O.	CZ
Concor COR 7,5 mg potahované tablety	not available	77-504-19-C	MERCK SPOL.S.R.O.	CZ
Concor COR 5 mg potahované tablety	not available	77/027/01-C	MERCK SPOL.S.R.O.	CZ
Concor COR 5 mg potahované tablety	not available	77/027/01-C	MERCK SPOL.S.R.O.	CZ
Concor COR 2,5 mg potahované tablety	not available	77/026/01-C	MERCK SPOL.S.R.O.	CZ
Concor COR 2,5 mg potahované tablety	not available	77/026/01-C	MERCK SPOL.S.R.O.	CZ

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
Concor COR 10 mg potahované tablety	not available	77/028/01-C	MERCK SPOL.S.R.O.	CZ
Concor COR 10 mg potahované tablety	not available	77/028/01-C	MERCK SPOL.S.R.O.	CZ
Concor COR 10 mg potahované tablety	not available	77/028/01-C	MERCK SPOL.S.R.O.	CZ
Concor COR 2,5 mg potahované tablety	not available	77/026/01-C	MERCK SPOL.S.R.O.	CZ
Concor COR 5 mg potahované tablety	not available	77/027/01-C	MERCK SPOL.S.R.O.	CZ
Concor 5 mg potahované tablety	not available	41/304/89-A/C	MERCK SPOL.S.R.O.	CZ
Concor 10 mg potahované tablety	not available	41/304/89-B/C	MERCK SPOL.S.R.O.	CZ
Concor 10 mg potahované tablety	not available	41/304/89-B/C	MERCK SPOL.S.R.O.	CZ
Concor 5 mg potahované tablety	not available	41/304/89-A/C	MERCK SPOL.S.R.O.	CZ
Bisoprolol PMCS 2,5 mg tablety	not available	41/559/12-C	PRO.MED.CS PRAHA A.S.	CZ
Conaret 5 mg Tabletten	DE/H/6433/004	2204679.00.00	ZENTIVA PHARMA GMBH	DE
Conaret 5 mg Tabletten	DE/H/6433/004	2204679.00.00	ZENTIVA PHARMA GMBH	DE
Conaret 5 mg Tabletten	DE/H/6433/004	2204679.00.00	ZENTIVA PHARMA GMBH	DE
Conaret 5 mg Tabletten	DE/H/6433/004	2204679.00.00	ZENTIVA PHARMA GMBH	DE
Conaret 5 mg Tabletten	DE/H/6433/004	2204679.00.00	ZENTIVA PHARMA GMBH	DE
Conaret 5 mg Tabletten	DE/H/6433/004	2204679.00.00	ZENTIVA PHARMA GMBH	DE
Conaret 5 mg Tabletten	DE/H/6433/004	2204679.00.00	ZENTIVA PHARMA GMBH	DE
Conaret 5 mg Tabletten	DE/H/6433/004	2204679.00.00	ZENTIVA PHARMA GMBH	DE
Conaret 5 mg Tabletten	DE/H/6433/004	2204679.00.00	ZENTIVA PHARMA GMBH	DE
Conaret 5 mg Tabletten	DE/H/6433/004	2204679.00.00	ZENTIVA PHARMA GMBH	DE
Conaret 5 mg Tabletten	DE/H/6433/004	2204679.00.00	ZENTIVA PHARMA GMBH	DE
Conaret 5 mg Tabletten	DE/H/6433/004	2204679.00.00	ZENTIVA PHARMA GMBH	DE
Conaret 5 mg Tabletten	DE/H/6433/004	2204679.00.00	ZENTIVA PHARMA GMBH	DE
Conaret 10 mg Tabletten	DE/H/6433/006	2204681.00.00	ZENTIVA PHARMA GMBH	DE

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
Conaret 10 mg Tabletten	DE/H/6433/006	2204681.00.00	ZENTIVA PHARMA GMBH	DE
Conaret 10 mg Tabletten	DE/H/6433/006	2204681.00.00	ZENTIVA PHARMA GMBH	DE
Conaret 10 mg Tabletten	DE/H/6433/006	2204681.00.00	ZENTIVA PHARMA GMBH	DE
Conaret 10 mg Tabletten	DE/H/6433/006	2204681.00.00	ZENTIVA PHARMA GMBH	DE
Conaret 10 mg Tabletten	DE/H/6433/006	2204681.00.00	ZENTIVA PHARMA GMBH	DE
Conaret 10 mg Tabletten	DE/H/6433/006	2204681.00.00	ZENTIVA PHARMA GMBH	DE
Conaret 10 mg Tabletten	DE/H/6433/006	2204681.00.00	ZENTIVA PHARMA GMBH	DE
Conaret 10 mg Tabletten	DE/H/6433/006	2204681.00.00	ZENTIVA PHARMA GMBH	DE
Conaret 10 mg Tabletten	DE/H/6433/006	2204681.00.00	ZENTIVA PHARMA GMBH	DE
Conaret 10 mg Tabletten	DE/H/6433/006	2204681.00.00	ZENTIVA PHARMA GMBH	DE
Conaret 10 mg Tabletten	DE/H/6433/006	2204681.00.00	ZENTIVA PHARMA GMBH	DE
Conaret 10 mg Tabletten	DE/H/6433/006	2204681.00.00	ZENTIVA PHARMA GMBH	DE
Conaret 5 mg Tabletten	DE/H/6433/004	2204679.00.00	ZENTIVA PHARMA GMBH	DE
Conaret 10 mg Tabletten	DE/H/6433/006	2204681.00.00	ZENTIVA PHARMA GMBH	DE
Concor COR 1,25 mg Filmtabletten	SE/H/0184/001	46660.00.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor COR 1,25 mg Filmtabletten	SE/H/0184/001	46660.00.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor COR 1,25 mg Filmtabletten	SE/H/0184/001	46660.00.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor COR 1,25 mg Filmtabletten	SE/H/0184/001	46660.00.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor COR 1,25 mg Filmtabletten	SE/H/0184/001	46660.00.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor COR 1,25 mg Filmtabletten	SE/H/0184/001	46660.00.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor COR 1,25 mg Filmtabletten	SE/H/0184/001	46660.00.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor COR 1,25 mg Filmtabletten	SE/H/0184/001	46660.00.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor COR 1,25 mg Filmtabletten	SE/H/0184/001	46660.00.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor COR 1,25 mg Filmtabletten	SE/H/0184/001	46660.00.00	MERCK HEALTHCARE GERMANY GMBH	DE

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
Concor COR 10 mg Filmtabletten	SE/H/0184/006	46660.05.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor COR 10 mg Filmtabletten	SE/H/0184/006	46660.05.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor COR 10 mg Filmtabletten	SE/H/0184/006	46660.05.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor COR 10 mg Filmtabletten	SE/H/0184/006	46660.05.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor COR 10 mg Filmtabletten	SE/H/0184/006	46660.05.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor COR 10 mg Filmtabletten	SE/H/0184/006	46660.05.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor COR 10 mg Filmtabletten	SE/H/0184/006	46660.05.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor COR 10 mg Filmtabletten	SE/H/0184/006	46660.05.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor COR 5 mg Filmtabletten	SE/H/0184/004	46660.03.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor COR 5 mg Filmtabletten	SE/H/0184/004	46660.03.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor COR 5 mg Filmtabletten	SE/H/0184/004	46660.03.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor COR 5 mg Filmtabletten	SE/H/0184/004	46660.03.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor COR 5 mg Filmtabletten	SE/H/0184/004	46660.03.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor COR 5 mg Filmtabletten	SE/H/0184/004	46660.03.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor COR 5 mg Filmtabletten	SE/H/0184/004	46660.03.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor COR 5 mg Filmtabletten	SE/H/0184/004	46660.03.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor COR 3,75 mg Filmtabletten	SE/H/0184/003	46660.02.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor COR 3,75 mg	SE/H/0184/003	46660.02.00	MERCK HEALTHCARE	DE

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
Filmtabletten			GERMANY GMBH	
Concor COR 3,75 mg Filmtabletten	SE/H/0184/003	46660.02.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor COR 3,75 mg Filmtabletten	SE/H/0184/003	46660.02.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor COR 3,75 mg Filmtabletten	SE/H/0184/003	46660.02.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor COR 3,75 mg Filmtabletten	SE/H/0184/003	46660.02.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor COR 3,75 mg Filmtabletten	SE/H/0184/003	46660.02.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor COR 3,75 mg Filmtabletten	SE/H/0184/003	46660.02.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor COR 7,5 mg Filmtabletten	SE/H/0184/005	46660.04.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor COR 7,5 mg Filmtabletten	SE/H/0184/005	46660.04.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor COR 7,5 mg Filmtabletten	SE/H/0184/005	46660.04.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor COR 7,5 mg Filmtabletten	SE/H/0184/005	46660.04.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor COR 7,5 mg Filmtabletten	SE/H/0184/005	46660.04.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor COR 7,5 mg Filmtabletten	SE/H/0184/005	46660.04.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor COR 7,5 mg Filmtabletten	SE/H/0184/005	46660.04.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor COR 7,5 mg Filmtabletten	SE/H/0184/005	46660.04.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor COR 2,5 mg Filmtabletten	SE/H/0184/002	46660.01.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor COR 2,5 mg Filmtabletten	SE/H/0184/002	46660.01.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor COR 2,5 mg Filmtabletten	SE/H/0184/002	46660.01.00	MERCK HEALTHCARE GERMANY GMBH	DE

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
Concor COR 2,5 mg Filmtabletten	SE/H/0184/002	46660.01.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor COR 2,5 mg Filmtabletten	SE/H/0184/002	46660.01.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor COR 2,5 mg Filmtabletten	SE/H/0184/002	46660.01.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor COR 2,5 mg Filmtabletten	SE/H/0184/002	46660.01.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor COR 2,5 mg Filmtabletten	SE/H/0184/002	46660.01.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor 10 mg Filmtabletten	not available	6849.01.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor 10 mg Filmtabletten	not available	6849.01.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor 10 mg Filmtabletten	not available	6849.01.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor 10 mg Filmtabletten	not available	6849.01.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor 10 mg Filmtabletten	not available	6849.01.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor 10 mg Filmtabletten	not available	6849.01.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor 10 mg Filmtabletten	not available	6849.01.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor 10 mg Filmtabletten	not available	6849.01.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor 5 mg Filmtabletten	not available	6849.00.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor 5 mg Filmtabletten	not available	6849.00.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor 5 mg Filmtabletten	not available	6849.00.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor 5 mg Filmtabletten	not available	6849.00.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor 5 mg	not available	6849.00.00	MERCK HEALTHCARE	DE

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
Filmtabletten			GERMANY GMBH	
Concor 5 mg Filmtabletten	not available	6849.00.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor 5 mg Filmtabletten	not available	6849.00.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor 5 mg Filmtabletten	not available	6849.00.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor 5 mg Filmtabletten	not available	6849.00.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor 5 mg Filmtabletten	not available	6849.00.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor 10 mg Filmtabletten	not available	6849.01.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor 10 mg Filmtabletten	not available	6849.01.00	MERCK HEALTHCARE GERMANY GMBH	DE
Concor 5 mg Filmtabletten	not available	6849.00.00	MERCK HEALTHCARE GERMANY GMBH	DE
Bisoprolol "Zentiva", tabletter	DE/H/6433/004	68586	ZENTIVA, K.S.	DK
Bisoprolol "Zentiva", tabletter	DE/H/6433/006	68589	ZENTIVA, K.S.	DK
Bisoprolol "Zentiva", tabletter	DE/H/6433/004	68586	ZENTIVA, K.S.	DK
Bisoprolol "Zentiva", tabletter	DE/H/6433/004	68586	ZENTIVA, K.S.	DK
Bisoprolol "Zentiva", tabletter	DE/H/6433/004	68586	ZENTIVA, K.S.	DK
Bisoprolol "Zentiva", tabletter	DE/H/6433/004	68586	ZENTIVA, K.S.	DK
Bisoprolol "Zentiva", tabletter	DE/H/6433/004	68586	ZENTIVA, K.S.	DK
Bisoprolol "Zentiva", tabletter	DE/H/6433/004	68586	ZENTIVA, K.S.	DK
Bisoprolol "Zentiva", tabletter	DE/H/6433/004	68586	ZENTIVA, K.S.	DK

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
Bisoprolol "Zentiva", tabletter	DE/H/6433/004	68586	ZENTIVA, K.S.	DK
Bisoprolol "Zentiva", tabletter	DE/H/6433/004	68586	ZENTIVA, K.S.	DK
Bisoprolol "Zentiva", tabletter	DE/H/6433/004	68586	ZENTIVA, K.S.	DK
Bisoprolol "Zentiva", tabletter	DE/H/6433/004	68586	ZENTIVA, K.S.	DK
Bisoprolol "Zentiva", tabletter	DE/H/6433/004	68586	ZENTIVA, K.S.	DK
Bisoprolol "Zentiva", tabletter	DE/H/6433/004	68586	ZENTIVA, K.S.	DK
Bisoprolol "Zentiva", tabletter	DE/H/6433/006	68589	ZENTIVA, K.S.	DK
Bisoprolol "Zentiva", tabletter	DE/H/6433/006	68589	ZENTIVA, K.S.	DK
Bisoprolol "Zentiva", tabletter	DE/H/6433/006	68589	ZENTIVA, K.S.	DK
Bisoprolol "Zentiva", tabletter	DE/H/6433/006	68589	ZENTIVA, K.S.	DK
Bisoprolol "Zentiva", tabletter	DE/H/6433/006	68589	ZENTIVA, K.S.	DK
Bisoprolol "Zentiva", tabletter	DE/H/6433/006	68589	ZENTIVA, K.S.	DK
Bisoprolol "Zentiva", tabletter	DE/H/6433/006	68589	ZENTIVA, K.S.	DK
Bisoprolol "Zentiva", tabletter	DE/H/6433/006	68589	ZENTIVA, K.S.	DK
Bisoprolol "Zentiva", tabletter	DE/H/6433/006	68589	ZENTIVA, K.S.	DK
Bisoprolol "Zentiva", tabletter	DE/H/6433/006	68589	ZENTIVA, K.S.	DK
Bisoprolol "Zentiva", tabletter	DE/H/6433/006	68589	ZENTIVA, K.S.	DK
Bisoprolol "Zentiva",	DE/H/6433/006	68589	ZENTIVA, K.S.	DK

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
tabletter				
Bisoprolol "Zentiva", tabletter	DE/H/6433/006	68589	ZENTIVA, K.S.	DK
Bisoprolol Zentiva 5 mg tabletid	DE/H/6433/004	1026221	ZENTIVA, K.S.	EE
Bisoprolol Zentiva 5 mg tabletid	DE/H/6433/004	1026221	ZENTIVA, K.S.	EE
Bisoprolol Zentiva 5 mg tabletid	DE/H/6433/004	1026221	ZENTIVA, K.S.	EE
Bisoprolol Zentiva 5 mg tabletid	DE/H/6433/004	1026221	ZENTIVA, K.S.	EE
Bisoprolol Zentiva 5 mg tabletid	DE/H/6433/004	1026221	ZENTIVA, K.S.	EE
Bisoprolol Zentiva 5 mg tabletid	DE/H/6433/004	1026221	ZENTIVA, K.S.	EE
Bisoprolol Zentiva 5 mg tabletid	DE/H/6433/004	1026221	ZENTIVA, K.S.	EE
Bisoprolol Zentiva 5 mg tabletid	DE/H/6433/004	1026221	ZENTIVA, K.S.	EE
Bisoprolol Zentiva 5 mg tabletid	DE/H/6433/004	1026221	ZENTIVA, K.S.	EE
Bisoprolol Zentiva 5 mg tabletid	DE/H/6433/004	1026221	ZENTIVA, K.S.	EE
Bisoprolol Zentiva 5 mg tabletid	DE/H/6433/004	1026221	ZENTIVA, K.S.	EE
Bisoprolol Zentiva 5 mg tabletid	DE/H/6433/004	1026221	ZENTIVA, K.S.	EE
Bisoprolol Zentiva 5 mg tabletid	DE/H/6433/004	1026221	ZENTIVA, K.S.	EE
Bisoprolol Zentiva 10 mg tabletid	DE/H/6433/006	1026321	ZENTIVA, K.S.	EE
Bisoprolol Zentiva 10 mg tabletid	DE/H/6433/006	1026321	ZENTIVA, K.S.	EE
Bisoprolol Zentiva 10 mg tabletid	DE/H/6433/006	1026321	ZENTIVA, K.S.	EE

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
Bisoprolol Zentiva 10 mg tabletid	DE/H/6433/006	1026321	ZENTIVA, K.S.	EE
Bisoprolol Zentiva 10 mg tabletid	DE/H/6433/006	1026321	ZENTIVA, K.S.	EE
Bisoprolol Zentiva 10 mg tabletid	DE/H/6433/006	1026321	ZENTIVA, K.S.	EE
Bisoprolol Zentiva 10 mg tabletid	DE/H/6433/006	1026321	ZENTIVA, K.S.	EE
Bisoprolol Zentiva 10 mg tabletid	DE/H/6433/006	1026321	ZENTIVA, K.S.	EE
Bisoprolol Zentiva 10 mg tabletid	DE/H/6433/006	1026321	ZENTIVA, K.S.	EE
Bisoprolol Zentiva 10 mg tabletid	DE/H/6433/006	1026321	ZENTIVA, K.S.	EE
Bisoprolol Zentiva 10 mg tabletid	DE/H/6433/006	1026321	ZENTIVA, K.S.	EE
Bisoprolol Zentiva 10 mg tabletid	DE/H/6433/006	1026321	ZENTIVA, K.S.	EE
Bisoprolol Zentiva 10 mg tabletid	DE/H/6433/006	1026321	ZENTIVA, K.S.	EE
Bisoprolol Zentiva 5 mg tabletid	DE/H/6433/004	1026221	ZENTIVA, K.S.	EE
Bisoprolol Zentiva 10 mg tabletid	DE/H/6433/006	1026321	ZENTIVA, K.S.	EE
Euradal 5 mg comprimidos recubiertos con película	not available	57.542	LACER S.A.	ES
Euradal 10 mg comprimidos recubiertos con película	not available	57.543	LACER S.A.	ES
Euradal 5 mg comprimidos recubiertos con película	not available	57.542	LACER S.A.	ES
Euradal 10 mg comprimidos recubiertos	not available	57.543	LACER S.A.	ES

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
con película				
Emconcor 10 mg comprimidos recubiertos con película	not available	57.655	MERCK, S.L.	ES
Emconcor 5 mg comprimidos recubiertos con película	not available	57.654	MERCK, S.L.	ES
Emconcor Cor 2,5 mg comprimidos recubiertos con película	SE/H/0184/002	63046	MERCK, S.L.	ES
Emconcor Cor 2,5 mg comprimidos recubiertos con película	SE/H/0184/002	63046	MERCK, S.L.	ES
Emconcor Cor 2,5 mg comprimidos recubiertos con película	SE/H/0184/002	63046	MERCK, S.L.	ES
Emconcor Cor 2,5 mg comprimidos recubiertos con película	SE/H/0184/002	63046	MERCK, S.L.	ES
Emconcor Cor 2,5 mg comprimidos recubiertos con película	SE/H/0184/002	63046	MERCK, S.L.	ES
Emconcor Cor 2,5 mg comprimidos recubiertos con película	SE/H/0184/002	63046	MERCK, S.L.	ES
Emconcor Cor 2,5 mg comprimidos recubiertos con película	SE/H/0184/002	63046	MERCK, S.L.	ES
Emconcor Cor 2,5 mg comprimidos recubiertos con película	SE/H/0184/002	63046	MERCK, S.L.	ES
Emconcor Cor 5 mg comprimidos recubiertos con película	SE/H/0184/004	63048	MERCK, S.L.	ES
Emconcor Cor 5 mg	SE/H/0184/004	63048	MERCK, S.L.	ES

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
comprimidos recubiertos con película				
Emconcor Cor 5 mg comprimidos recubiertos con película	SE/H/0184/004	63048	MERCK, S.L.	ES
Emconcor Cor 5 mg comprimidos recubiertos con película	SE/H/0184/004	63048	MERCK, S.L.	ES
Emconcor Cor 5 mg comprimidos recubiertos con película	SE/H/0184/004	63048	MERCK, S.L.	ES
Emconcor Cor 5 mg comprimidos recubiertos con película	SE/H/0184/004	63048	MERCK, S.L.	ES
Emconcor Cor 5 mg comprimidos recubiertos con película	SE/H/0184/004	63048	MERCK, S.L.	ES
Emconcor Cor 5 mg comprimidos recubiertos con película	SE/H/0184/004	63048	MERCK, S.L.	ES
Emconcor 10 mg tabletti, kalvopäällysteinen	not available	10191	MERCK OY	FI
Emconcor 5 mg tabletti, kalvopäällysteinen	not available	10190	MERCK OY	FI
Emconcor 5 mg filmdragerade tabletter	not available	10190	MERCK OY	FI
Emconcor 5 mg filmdragerade tabletter	not available	10190	MERCK OY	FI
Emconcor 10 mg filmdragerade tabletter	not available	10191	MERCK OY	FI
Emconcor 10 mg filmdragerade tabletter	not available	10191	MERCK OY	FI
Emconcor 10 mg tabletti,	not available	10191	MERCK OY	FI

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
kalvopäällysteinen				
Emconcor 5 mg tabletti, kalvopäällysteinen	not available	10190	MERCK OY	FI
Emconcor CHF 1,25 mg filmdragerade tabletter	SE/H/0184/001	14531	MERCK OY	FI
Emconcor CHF 1,25 mg filmdragerade tabletter	SE/H/0184/001	14531	MERCK OY	FI
Emconcor CHF 1,25 mg filmdragerade tabletter	SE/H/0184/001	14531	MERCK OY	FI
Emconcor CHF 1,25 mg filmdragerade tabletter	SE/H/0184/001	14531	MERCK OY	FI
Emconcor CHF 1,25 mg filmdragerade tabletter	SE/H/0184/001	14531	MERCK OY	FI
Emconcor CHF 1,25 mg filmdragerade tabletter	SE/H/0184/001	14531	MERCK OY	FI
Emconcor CHF 1,25 mg filmdragerade tabletter	SE/H/0184/001	14531	MERCK OY	FI
Emconcor CHF 1,25 mg filmdragerade tabletter	SE/H/0184/001	14531	MERCK OY	FI
Emconcor CHF 1,25 mg filmdragerade tabletter	SE/H/0184/001	14531	MERCK OY	FI
Emconcor CHF 1,25 mg filmdragerade tabletter	SE/H/0184/001	14531	MERCK OY	FI
Emconcor CHF 1,25 mg filmdragerade tabletter	SE/H/0184/001	14531	MERCK OY	FI
Emconcor CHF 1,25 mg filmdragerade tabletter	SE/H/0184/001	14531	MERCK OY	FI
Emconcor CHF 1,25 mg filmdragerade tabletter	SE/H/0184/001	14531	MERCK OY	FI
Emconcor CHF 1,25 mg filmdragerade tabletter	SE/H/0184/001	14531	MERCK OY	FI
Emconcor CHF 1,25 mg filmdragerade tabletter	SE/H/0184/001	14531	MERCK OY	FI
Emconcor CHF 1,25 mg filmdragerade tabletter	SE/H/0184/001	14531	MERCK OY	FI

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
Emconcor CHF 1,25 mg tabletti, kalvopäällysteinen	SE/H/0184/001	14531	MERCK OY	FI
Emconcor CHF 1,25 mg tabletti, kalvopäällysteinen	SE/H/0184/001	14531	MERCK OY	FI
Emconcor CHF 1,25 mg tabletti, kalvopäällysteinen	SE/H/0184/001	14531	MERCK OY	FI
Emconcor CHF 1,25 mg tabletti, kalvopäällysteinen	SE/H/0184/001	14531	MERCK OY	FI
Emconcor CHF 1,25 mg tabletti, kalvopäällysteinen	SE/H/0184/001	14531	MERCK OY	FI
Emconcor CHF 1,25 mg tabletti, kalvopäällysteinen	SE/H/0184/001	14531	MERCK OY	FI
Emconcor CHF 1,25 mg tabletti, kalvopäällysteinen	SE/H/0184/001	14531	MERCK OY	FI
Emconcor CHF 1,25 mg tabletti, kalvopäällysteinen	SE/H/0184/001	14531	MERCK OY	FI
Emconcor CHF 1,25 mg tabletti, kalvopäällysteinen	SE/H/0184/001	14531	MERCK OY	FI
Emconcor CHF 1,25 mg tabletti, kalvopäällysteinen	SE/H/0184/001	14531	MERCK OY	FI
Emconcor CHF 1,25 mg tabletti, kalvopäällysteinen	SE/H/0184/001	14531	MERCK OY	FI
Emconcor CHF 1,25 mg tabletti,	SE/H/0184/001	14531	MERCK OY	FI

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
kalvopäällysteinen				
Emconcor CHF 1,25 mg tabletti, kalvopäällysteinen	SE/H/0184/001	14531	MERCK OY	FI
Emconcor CHF 1,25 mg tabletti, kalvopäällysteinen	SE/H/0184/001	14531	MERCK OY	FI
Emconcor CHF 1,25 mg tabletti, kalvopäällysteinen	SE/H/0184/001	14531	MERCK OY	FI
Emconcor CHF 1,25 mg tabletti, kalvopäällysteinen	SE/H/0184/001	14531	MERCK OY	FI
Emconcor CHF 10 mg tabletti, kalvopäällysteinen	SE/H/0184/006	14536	MERCK OY	FI
Emconcor CHF 10 mg tabletti, kalvopäällysteinen	SE/H/0184/006	14536	MERCK OY	FI
Emconcor CHF 10 mg tabletti, kalvopäällysteinen	SE/H/0184/006	14536	MERCK OY	FI
Emconcor CHF 10 mg tabletti, kalvopäällysteinen	SE/H/0184/006	14536	MERCK OY	FI
Emconcor CHF 10 mg tabletti, kalvopäällysteinen	SE/H/0184/006	14536	MERCK OY	FI
Emconcor CHF 10 mg tabletti, kalvopäällysteinen	SE/H/0184/006	14536	MERCK OY	FI
Emconcor CHF 10 mg tabletti, kalvopäällysteinen	SE/H/0184/006	14536	MERCK OY	FI
Emconcor CHF 10 mg	SE/H/0184/006	14536	MERCK OY	FI

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
tabletti, kalvopäällysteinen				
Emconcor CHF 10 mg tabletti, kalvopäällysteinen	SE/H/0184/006	14536	MERCK OY	FI
Emconcor CHF 10 mg tabletti, kalvopäällysteinen	SE/H/0184/006	14536	MERCK OY	FI
Emconcor CHF 10 mg tabletti, kalvopäällysteinen	SE/H/0184/006	14536	MERCK OY	FI
Emconcor CHF 10 mg tabletti, kalvopäällysteinen	SE/H/0184/006	14536	MERCK OY	FI
Emconcor CHF 10 mg tabletti, kalvopäällysteinen	SE/H/0184/006	14536	MERCK OY	FI
Emconcor CHF 10 mg tabletti, kalvopäällysteinen	SE/H/0184/006	14536	MERCK OY	FI
Emconcor CHF 10 mg tabletti, kalvopäällysteinen	SE/H/0184/006	14536	MERCK OY	FI
Emconcor CHF 10 mg tabletti, kalvopäällysteinen	SE/H/0184/006	14536	MERCK OY	FI
Emconcor CHF 10 mg filmdragerade tabletter	SE/H/0184/006	14536	MERCK OY	FI
Emconcor CHF 10 mg filmdragerade tabletter	SE/H/0184/006	14536	MERCK OY	FI
Emconcor CHF 10 mg filmdragerade tabletter	SE/H/0184/006	14536	MERCK OY	FI
Emconcor CHF 10 mg filmdragerade tabletter	SE/H/0184/006	14536	MERCK OY	FI
Emconcor CHF 10 mg	SE/H/0184/006	14536	MERCK OY	FI

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
filmdragerade tabletter				
Emconcor CHF 10 mg filmdragerade tabletter	SE/H/0184/006	14536	MERCK OY	FI
Emconcor CHF 10 mg filmdragerade tabletter	SE/H/0184/006	14536	MERCK OY	FI
Emconcor CHF 10 mg filmdragerade tabletter	SE/H/0184/006	14536	MERCK OY	FI
Emconcor CHF 10 mg filmdragerade tabletter	SE/H/0184/006	14536	MERCK OY	FI
Emconcor CHF 10 mg filmdragerade tabletter	SE/H/0184/006	14536	MERCK OY	FI
Emconcor CHF 10 mg filmdragerade tabletter	SE/H/0184/006	14536	MERCK OY	FI
Emconcor CHF 10 mg filmdragerade tabletter	SE/H/0184/006	14536	MERCK OY	FI
Emconcor CHF 10 mg filmdragerade tabletter	SE/H/0184/006	14536	MERCK OY	FI
Emconcor CHF 10 mg filmdragerade tabletter	SE/H/0184/006	14536	MERCK OY	FI
Emconcor CHF 10 mg filmdragerade tabletter	SE/H/0184/006	14536	MERCK OY	FI
Emconcor CHF 10 mg filmdragerade tabletter	SE/H/0184/006	14536	MERCK OY	FI
Emconcor CHF 2,5 mg filmdragerade tabletter	SE/H/0184/002	14532	MERCK OY	FI
Emconcor CHF 2,5 mg filmdragerade tabletter	SE/H/0184/002	14532	MERCK OY	FI
Emconcor CHF 2,5 mg filmdragerade tabletter	SE/H/0184/002	14532	MERCK OY	FI
Emconcor CHF 2,5 mg filmdragerade tabletter	SE/H/0184/002	14532	MERCK OY	FI
Emconcor CHF 2,5 mg filmdragerade tabletter	SE/H/0184/002	14532	MERCK OY	FI
Emconcor CHF 2,5 mg filmdragerade tabletter	SE/H/0184/002	14532	MERCK OY	FI

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
Emconcor CHF 2,5 mg filmdragerade tabletter	SE/H/0184/002	14532	MERCK OY	FI
Emconcor CHF 2,5 mg filmdragerade tabletter	SE/H/0184/002	14532	MERCK OY	FI
Emconcor CHF 2,5 mg filmdragerade tabletter	SE/H/0184/002	14532	MERCK OY	FI
Emconcor CHF 2,5 mg filmdragerade tabletter	SE/H/0184/002	14532	MERCK OY	FI
Emconcor CHF 2,5 mg filmdragerade tabletter	SE/H/0184/002	14532	MERCK OY	FI
Emconcor CHF 2,5 mg filmdragerade tabletter	SE/H/0184/002	14532	MERCK OY	FI
Emconcor CHF 2,5 mg filmdragerade tabletter	SE/H/0184/002	14532	MERCK OY	FI
Emconcor CHF 2,5 mg filmdragerade tabletter	SE/H/0184/002	14532	MERCK OY	FI
Emconcor CHF 2,5 mg filmdragerade tabletter	SE/H/0184/002	14532	MERCK OY	FI
Emconcor CHF 2,5 mg filmdragerade tabletter	SE/H/0184/002	14532	MERCK OY	FI
Emconcor CHF 2,5 mg tabletti, kalvopäällysteinen	SE/H/0184/002	14532	MERCK OY	FI
Emconcor CHF 2,5 mg tabletti, kalvopäällysteinen	SE/H/0184/002	14532	MERCK OY	FI
Emconcor CHF 2,5 mg tabletti, kalvopäällysteinen	SE/H/0184/002	14532	MERCK OY	FI
Emconcor CHF 2,5 mg tabletti, kalvopäällysteinen	SE/H/0184/002	14532	MERCK OY	FI
Emconcor CHF 2,5 mg tabletti, kalvopäällysteinen	SE/H/0184/002	14532	MERCK OY	FI

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
Emconcor CHF 2,5 mg tabletti, kalvopäällysteinen	SE/H/0184/002	14532	MERCK OY	FI
Emconcor CHF 2,5 mg tabletti, kalvopäällysteinen	SE/H/0184/002	14532	MERCK OY	FI
Emconcor CHF 2,5 mg tabletti, kalvopäällysteinen	SE/H/0184/002	14532	MERCK OY	FI
Emconcor CHF 2,5 mg tabletti, kalvopäällysteinen	SE/H/0184/002	14532	MERCK OY	FI
Emconcor CHF 2,5 mg tabletti, kalvopäällysteinen	SE/H/0184/002	14532	MERCK OY	FI
Emconcor CHF 2,5 mg tabletti, kalvopäällysteinen	SE/H/0184/002	14532	MERCK OY	FI
Emconcor CHF 2,5 mg tabletti, kalvopäällysteinen	SE/H/0184/002	14532	MERCK OY	FI
Emconcor CHF 2,5 mg tabletti, kalvopäällysteinen	SE/H/0184/002	14532	MERCK OY	FI
Emconcor CHF 2,5 mg tabletti, kalvopäällysteinen	SE/H/0184/002	14532	MERCK OY	FI
Emconcor CHF 2,5 mg tabletti, kalvopäällysteinen	SE/H/0184/002	14532	MERCK OY	FI
Emconcor CHF 2,5 mg tabletti, kalvopäällysteinen	SE/H/0184/002	14532	MERCK OY	FI
Emconcor CHF 3,75 mg tabletti,	SE/H/0184/003	14533	MERCK OY	FI

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
kalvopäällysteinen				
Emconcor CHF 3,75 mg tabletti, kalvopäällysteinen	SE/H/0184/003	14533	MERCK OY	FI
Emconcor CHF 3,75 mg tabletti, kalvopäällysteinen	SE/H/0184/003	14533	MERCK OY	FI
Emconcor CHF 3,75 mg tabletti, kalvopäällysteinen	SE/H/0184/003	14533	MERCK OY	FI
Emconcor CHF 3,75 mg tabletti, kalvopäällysteinen	SE/H/0184/003	14533	MERCK OY	FI
Emconcor CHF 3,75 mg tabletti, kalvopäällysteinen	SE/H/0184/003	14533	MERCK OY	FI
Emconcor CHF 3,75 mg tabletti, kalvopäällysteinen	SE/H/0184/003	14533	MERCK OY	FI
Emconcor CHF 3,75 mg tabletti, kalvopäällysteinen	SE/H/0184/003	14533	MERCK OY	FI
Emconcor CHF 3,75 mg tabletti, kalvopäällysteinen	SE/H/0184/003	14533	MERCK OY	FI
Emconcor CHF 3,75 mg tabletti, kalvopäällysteinen	SE/H/0184/003	14533	MERCK OY	FI
Emconcor CHF 3,75 mg tabletti, kalvopäällysteinen	SE/H/0184/003	14533	MERCK OY	FI
Emconcor CHF 3,75 mg tabletti, kalvopäällysteinen	SE/H/0184/003	14533	MERCK OY	FI
Emconcor CHF 3,75 mg	SE/H/0184/003	14533	MERCK OY	FI

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
tabletti, kalvopäällysteinen				
Emconcor CHF 3,75 mg tabletti, kalvopäällysteinen	SE/H/0184/003	14533	MERCK OY	FI
Emconcor CHF 3,75 mg tabletti, kalvopäällysteinen	SE/H/0184/003	14533	MERCK OY	FI
Emconcor CHF 3,75 mg tabletti, kalvopäällysteinen	SE/H/0184/003	14533	MERCK OY	FI
Emconcor CHF 3,75 mg filmdragerade tabletter	SE/H/0184/003	14533	MERCK OY	FI
Emconcor CHF 3,75 mg filmdragerade tabletter	SE/H/0184/003	14533	MERCK OY	FI
Emconcor CHF 3,75 mg filmdragerade tabletter	SE/H/0184/003	14533	MERCK OY	FI
Emconcor CHF 3,75 mg filmdragerade tabletter	SE/H/0184/003	14533	MERCK OY	FI
Emconcor CHF 3,75 mg filmdragerade tabletter	SE/H/0184/003	14533	MERCK OY	FI
Emconcor CHF 3,75 mg filmdragerade tabletter	SE/H/0184/003	14533	MERCK OY	FI
Emconcor CHF 3,75 mg filmdragerade tabletter	SE/H/0184/003	14533	MERCK OY	FI
Emconcor CHF 3,75 mg filmdragerade tabletter	SE/H/0184/003	14533	MERCK OY	FI
Emconcor CHF 3,75 mg filmdragerade tabletter	SE/H/0184/003	14533	MERCK OY	FI
Emconcor CHF 3,75 mg filmdragerade tabletter	SE/H/0184/003	14533	MERCK OY	FI
Emconcor CHF 3,75 mg filmdragerade tabletter	SE/H/0184/003	14533	MERCK OY	FI
Emconcor CHF 3,75 mg filmdragerade tabletter	SE/H/0184/003	14533	MERCK OY	FI

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
Emconcor CHF 3,75 mg filmdragerade tabletter	SE/H/0184/003	14533	MERCK OY	FI
Emconcor CHF 3,75 mg filmdragerade tabletter	SE/H/0184/003	14533	MERCK OY	FI
Emconcor CHF 3,75 mg filmdragerade tabletter	SE/H/0184/003	14533	MERCK OY	FI
Emconcor CHF 3,75 mg filmdragerade tabletter	SE/H/0184/003	14533	MERCK OY	FI
Emconcor CHF 7,5 mg tabletti, kalvopäällysteinen	SE/H/0184/005	14535	MERCK OY	FI
Emconcor CHF 7,5 mg tabletti, kalvopäällysteinen	SE/H/0184/005	14535	MERCK OY	FI
Emconcor CHF 7,5 mg tabletti, kalvopäällysteinen	SE/H/0184/005	14535	MERCK OY	FI
Emconcor CHF 7,5 mg tabletti, kalvopäällysteinen	SE/H/0184/005	14535	MERCK OY	FI
Emconcor CHF 7,5 mg tabletti, kalvopäällysteinen	SE/H/0184/005	14535	MERCK OY	FI
Emconcor CHF 7,5 mg tabletti, kalvopäällysteinen	SE/H/0184/005	14535	MERCK OY	FI
Emconcor CHF 7,5 mg tabletti, kalvopäällysteinen	SE/H/0184/005	14535	MERCK OY	FI
Emconcor CHF 7,5 mg tabletti, kalvopäällysteinen	SE/H/0184/005	14535	MERCK OY	FI
Emconcor CHF 7,5 mg tabletti, kalvopäällysteinen	SE/H/0184/005	14535	MERCK OY	FI

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
Emconcor CHF 7,5 mg tabletti, kalvopäällysteinen	SE/H/0184/005	14535	MERCK OY	FI
Emconcor CHF 7,5 mg tabletti, kalvopäällysteinen	SE/H/0184/005	14535	MERCK OY	FI
Emconcor CHF 7,5 mg tabletti, kalvopäällysteinen	SE/H/0184/005	14535	MERCK OY	FI
Emconcor CHF 7,5 mg tabletti, kalvopäällysteinen	SE/H/0184/005	14535	MERCK OY	FI
Emconcor CHF 7,5 mg tabletti, kalvopäällysteinen	SE/H/0184/005	14535	MERCK OY	FI
Emconcor CHF 7,5 mg tabletti, kalvopäällysteinen	SE/H/0184/005	14535	MERCK OY	FI
Emconcor CHF 7,5 mg tabletti, kalvopäällysteinen	SE/H/0184/005	14535	MERCK OY	FI
Emconcor CHF 7,5 mg filmdragerade tabletter	SE/H/0184/005	14535	MERCK OY	FI
Emconcor CHF 7,5 mg filmdragerade tabletter	SE/H/0184/005	14535	MERCK OY	FI
Emconcor CHF 7,5 mg filmdragerade tabletter	SE/H/0184/005	14535	MERCK OY	FI
Emconcor CHF 7,5 mg filmdragerade tabletter	SE/H/0184/005	14535	MERCK OY	FI
Emconcor CHF 7,5 mg filmdragerade tabletter	SE/H/0184/005	14535	MERCK OY	FI
Emconcor CHF 7,5 mg filmdragerade tabletter	SE/H/0184/005	14535	MERCK OY	FI
Emconcor CHF 7,5 mg filmdragerade tabletter	SE/H/0184/005	14535	MERCK OY	FI

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
Emconcor CHF 7,5 mg filmdragerade tabletter	SE/H/0184/005	14535	MERCK OY	FI
Emconcor CHF 7,5 mg filmdragerade tabletter	SE/H/0184/005	14535	MERCK OY	FI
Emconcor CHF 7,5 mg filmdragerade tabletter	SE/H/0184/005	14535	MERCK OY	FI
Emconcor CHF 7,5 mg filmdragerade tabletter	SE/H/0184/005	14535	MERCK OY	FI
Emconcor CHF 7,5 mg filmdragerade tabletter	SE/H/0184/005	14535	MERCK OY	FI
Emconcor CHF 7,5 mg filmdragerade tabletter	SE/H/0184/005	14535	MERCK OY	FI
Emconcor CHF 7,5 mg filmdragerade tabletter	SE/H/0184/005	14535	MERCK OY	FI
Emconcor CHF 7,5 mg filmdragerade tabletter	SE/H/0184/005	14535	MERCK OY	FI
Emconcor CHF 7,5 mg filmdragerade tabletter	SE/H/0184/005	14535	MERCK OY	FI
Emconcor CHF 5 mg filmdragerade tabletter	SE/H/0184/004	14534	MERCK OY	FI
Emconcor CHF 5 mg filmdragerade tabletter	SE/H/0184/004	14534	MERCK OY	FI
Emconcor CHF 5 mg filmdragerade tabletter	SE/H/0184/004	14534	MERCK OY	FI
Emconcor CHF 5 mg filmdragerade tabletter	SE/H/0184/004	14534	MERCK OY	FI
Emconcor CHF 5 mg filmdragerade tabletter	SE/H/0184/004	14534	MERCK OY	FI
Emconcor CHF 5 mg filmdragerade tabletter	SE/H/0184/004	14534	MERCK OY	FI
Emconcor CHF 5 mg filmdragerade tabletter	SE/H/0184/004	14534	MERCK OY	FI
Emconcor CHF 5 mg filmdragerade tabletter	SE/H/0184/004	14534	MERCK OY	FI
Emconcor CHF 5 mg	SE/H/0184/004	14534	MERCK OY	FI

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
filmdragerade tabletter				
Emconcor CHF 5 mg filmdragerade tabletter	SE/H/0184/004	14534	MERCK OY	FI
Emconcor CHF 5 mg filmdragerade tabletter	SE/H/0184/004	14534	MERCK OY	FI
Emconcor CHF 5 mg filmdragerade tabletter	SE/H/0184/004	14534	MERCK OY	FI
Emconcor CHF 5 mg filmdragerade tabletter	SE/H/0184/004	14534	MERCK OY	FI
Emconcor CHF 5 mg filmdragerade tabletter	SE/H/0184/004	14534	MERCK OY	FI
Emconcor CHF 5 mg filmdragerade tabletter	SE/H/0184/004	14534	MERCK OY	FI
Emconcor CHF 5 mg filmdragerade tabletter	SE/H/0184/004	14534	MERCK OY	FI
Emconcor CHF 5 mg tabletti, kalvopäällysteinen	SE/H/0184/004	14534	MERCK OY	FI
Emconcor CHF 5 mg tabletti, kalvopäällysteinen	SE/H/0184/004	14534	MERCK OY	FI
Emconcor CHF 5 mg tabletti, kalvopäällysteinen	SE/H/0184/004	14534	MERCK OY	FI
Emconcor CHF 5 mg tabletti, kalvopäällysteinen	SE/H/0184/004	14534	MERCK OY	FI
Emconcor CHF 5 mg tabletti, kalvopäällysteinen	SE/H/0184/004	14534	MERCK OY	FI
Emconcor CHF 5 mg tabletti, kalvopäällysteinen	SE/H/0184/004	14534	MERCK OY	FI
Emconcor CHF 5 mg tabletti,	SE/H/0184/004	14534	MERCK OY	FI

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
kalvopäällysteinen				
Emconcor CHF 5 mg tabletti, kalvopäällysteinen	SE/H/0184/004	14534	MERCK OY	FI
Emconcor CHF 5 mg tabletti, kalvopäällysteinen	SE/H/0184/004	14534	MERCK OY	FI
Emconcor CHF 5 mg tabletti, kalvopäällysteinen	SE/H/0184/004	14534	MERCK OY	FI
Emconcor CHF 5 mg tabletti, kalvopäällysteinen	SE/H/0184/004	14534	MERCK OY	FI
Emconcor CHF 5 mg tabletti, kalvopäällysteinen	SE/H/0184/004	14534	MERCK OY	FI
Emconcor CHF 5 mg tabletti, kalvopäällysteinen	SE/H/0184/004	14534	MERCK OY	FI
Emconcor CHF 5 mg tabletti, kalvopäällysteinen	SE/H/0184/004	14534	MERCK OY	FI
Emconcor CHF 5 mg tabletti, kalvopäällysteinen	SE/H/0184/004	14534	MERCK OY	FI
Emconcor CHF 5 mg tabletti, kalvopäällysteinen	SE/H/0184/004	14534	MERCK OY	FI
CARDENSIEL 10 mg comprimé pelliculé sécable	SE/H/0184/006	6 635 154 8	MERCK SANTÉ S.A.S.	FR
CARDENSIEL 10 mg comprimé pelliculé sécable	SE/H/0184/006	6 635 154 8	MERCK SANTÉ S.A.S.	FR
CARDENSIEL 10 mg	SE/H/0184/006	6 635 154 8	MERCK SANTÉ S.A.S.	FR

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
comprimé pelliculé sécable				
CARDENSIEL 10 mg comprimé pelliculé sécable	SE/H/0184/006	34009 353 539 7 9	MERCK SANTÉ S.A.S.	FR
CARDENSIEL 10 mg comprimé pelliculé sécable	SE/H/0184/006	34009 353 131 8 8	MERCK SANTÉ S.A.S.	FR
CARDENSIEL 10 mg comprimé pelliculé sécable	SE/H/0184/006	34009 353 132 4 9	MERCK SANTÉ S.A.S.	FR
CARDENSIEL 7,5 mg comprimé pelliculé sécable	SE/H/0184/005	34009 353 129 3 8	MERCK SANTÉ S.A.S.	FR
CARDENSIEL 7,5 mg comprimé pelliculé sécable	SE/H/0184/005	34009 353 540 5 1	MERCK SANTÉ S.A.S.	FR
CARDENSIEL 7,5 mg comprimé pelliculé sécable	SE/H/0184/005	6 926 964 7	MERCK SANTÉ S.A.S.	FR
CARDENSIEL 7,5 mg comprimé pelliculé sécable	SE/H/0184/005	6 926 964 7	MERCK SANTÉ S.A.S.	FR
CARDENSIEL 7,5 mg comprimé pelliculé sécable	SE/H/0184/005	34009 353 130 1 0	MERCK SANTÉ S.A.S.	FR
CARDENSIEL 7,5 mg comprimé pelliculé sécable	SE/H/0184/005	6 926 964 7	MERCK SANTÉ S.A.S.	FR
CARDENSIEL 1,25 mg comprimé pelliculé	SE/H/0184/001	34009 353 535 1 1	MERCK SANTÉ S.A.S.	FR
CARDENSIEL 1,25 mg comprimé pelliculé	SE/H/0184/001	34009 352 968 1 8	MERCK SANTÉ S.A.S.	FR
CARDENSIEL 1,25 mg comprimé pelliculé	SE/H/0184/001	6 316 228 2	MERCK SANTÉ S.A.S.	FR

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
CARDENSIEL 1,25 mg comprimé pelliculé	SE/H/0184/001	6 316 228 2	MERCK SANTÉ S.A.S.	FR
CARDENSIEL 1,25 mg comprimé pelliculé	SE/H/0184/001	34009 352 969 8 6	MERCK SANTÉ S.A.S.	FR
CARDENSIEL 1,25 mg comprimé pelliculé	SE/H/0184/001	6 316 228 2	MERCK SANTÉ S.A.S.	FR
CARDENSIEL 2,5 mg comprimé pelliculé sécable	SE/H/0184/002	6 007 757 4	MERCK SANTÉ S.A.S.	FR
CARDENSIEL 2,5 mg comprimé pelliculé sécable	SE/H/0184/002	6 007 757 4	MERCK SANTÉ S.A.S.	FR
CARDENSIEL 2,5 mg comprimé pelliculé sécable	SE/H/0184/002	34009 352 971 2 9	MERCK SANTÉ S.A.S.	FR
CARDENSIEL 2,5 mg comprimé pelliculé sécable	SE/H/0184/002	6 007 757 4	MERCK SANTÉ S.A.S.	FR
CARDENSIEL 2,5 mg comprimé pelliculé sécable	SE/H/0184/002	34009 353 536 8 9	MERCK SANTÉ S.A.S.	FR
CARDENSIEL 2,5 mg comprimé pelliculé sécable	SE/H/0184/002	34009 352 970 6 8	MERCK SANTÉ S.A.S.	FR
CARDENSIEL 3,75 mg comprimé pelliculé sécable	SE/H/0184/003	6 627 924 8	MERCK SANTÉ S.A.S.	FR
CARDENSIEL 3,75 mg comprimé pelliculé sécable	SE/H/0184/003	6 627 924 8	MERCK SANTÉ S.A.S.	FR
CARDENSIEL 3,75 mg comprimé pelliculé sécable	SE/H/0184/003	34009 353 537 4 0	MERCK SANTÉ S.A.S.	FR
CARDENSIEL 3,75 mg comprimé pelliculé	SE/H/0184/003	6 627 924 8	MERCK SANTÉ S.A.S.	FR

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
sécable				
CARDENSIEL 3,75 mg comprimé pelliculé sécable	SE/H/0184/003	34009 352 973 5 8	MERCK SANTÉ S.A.S.	FR
CARDENSIEL 3,75 mg comprimé pelliculé sécable	SE/H/0184/003	34009 352 972 9 7	MERCK SANTÉ S.A.S.	FR
CARDENSIEL 5 mg comprimé pelliculé sécable	SE/H/0184/004	6 180 453 9	MERCK SANTÉ S.A.S.	FR
CARDENSIEL 5 mg comprimé pelliculé sécable	SE/H/0184/004	34009 353 128 7 7	MERCK SANTÉ S.A.S.	FR
CARDENSIEL 5 mg comprimé pelliculé sécable	SE/H/0184/004	34009 353 538 0 1	MERCK SANTÉ S.A.S.	FR
CARDENSIEL 5 mg comprimé pelliculé sécable	SE/H/0184/004	6 180 453 9	MERCK SANTÉ S.A.S.	FR
CARDENSIEL 5 mg comprimé pelliculé sécable	SE/H/0184/004	34009 352 974 1 9	MERCK SANTÉ S.A.S.	FR
CARDENSIEL 5 mg comprimé pelliculé sécable	SE/H/0184/004	6 180 453 9	MERCK SANTÉ S.A.S.	FR
DETENSIEL 10 mg comprimé pelliculé sécable	not available	34009 358 817 5 5	MERCK SANTÉ S.A.S.	FR
DETENSIEL 10 mg comprimé pelliculé sécable	not available	34009 372 245 5 0	MERCK SANTÉ S.A.S.	FR
DETENSIEL 10 mg comprimé pelliculé sécable	not available	34009 328 450 6 4	MERCK SANTÉ S.A.S.	FR
DETENSIEL 10 mg	not available	34009 558 331 9 2	MERCK SANTÉ S.A.S.	FR

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
comprimé pelliculé sécable				
DETENSIEL 10 mg comprimé pelliculé sécable	not available	34009 300 852 4 0	MERCK SANTÉ S.A.S.	FR
DETENSIEL 10 mg comprimé pelliculé sécable	not available	34009 300 852 5 7	MERCK SANTÉ S.A.S.	FR
Concor 10 mg επικαλυμμένα με λεπτό υμένιο δισκία	not available	78137/9-11-2011	MERCK A.E.	GR
Concor 5 mg επικαλυμμένα με λεπτό υμένιο δισκία	not available	36771/9-11-2011	MERCK A.E.	GR
EMCONCOR® 2,5 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0187/002	15255/5-3-2015	MERCK A.E.	GR
EMCONCOR® 2,5 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0187/002	15255/5-3-2015	MERCK A.E.	GR
EMCONCOR® 2,5 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0187/002	15255/5-3-2015	MERCK A.E.	GR
EMCONCOR® 2,5 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0187/002	15255/5-3-2015	MERCK A.E.	GR
EMCONCOR® 2,5 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0187/002	15255/5-3-2015	MERCK A.E.	GR
EMCONCOR® 2,5 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0187/002	15255/5-3-2015	MERCK A.E.	GR
EMCONCOR® 2,5 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0187/002	15255/5-3-2015	MERCK A.E.	GR

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
EMCONCOR® 5 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0187/004	15259/5-3-2015	MERCK A.E.	GR
EMCONCOR® 5 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0187/004	15259/5-3-2015	MERCK A.E.	GR
EMCONCOR® 5 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0187/004	15259/5-3-2015	MERCK A.E.	GR
EMCONCOR® 5 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0187/004	15259/5-3-2015	MERCK A.E.	GR
EMCONCOR® 5 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0187/004	15259/5-3-2015	MERCK A.E.	GR
EMCONCOR® 5 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0187/004	15259/5-3-2015	MERCK A.E.	GR
EMCONCOR® 5 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0187/004	15259/5-3-2015	MERCK A.E.	GR
EMCONCOR® 5 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0187/004	15259/5-3-2015	MERCK A.E.	GR
EMCONCOR® 2,5 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0187/002	15255/5-3-2015	MERCK A.E.	GR
Concor 10 mg filmom obložene tablete	not available	HR-H-747818074	MERCK D.O.O	HR
Concor 10 mg filmom obložene tablete	not available	HR-H-747818074	MERCK D.O.O	HR
Concor 10 mg filmom obložene tablete	not available	HR-H-747818074	MERCK D.O.O	HR
Concor 5 mg filmom obložene tablete	not available	HR-H-148909302	MERCK D.O.O	HR

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
Concor 5 mg filmom obložene tablete	not available	HR-H-148909302	MERCK D.O.O	HR
Concor 5 mg filmom obložene tablete	not available	HR-H-148909302	MERCK D.O.O	HR
Concor 10 mg filmom obložene tablete	not available	HR-H-747818074	MERCK D.O.O	HR
Concor 5 mg filmom obložene tablete	not available	HR-H-148909302	MERCK D.O.O	HR
Concor COR 2,5 mg filmom obložene tablete	SE/H/0184/002	HR-H-061130840	MERCK D.O.O	HR
Concor COR 2,5 mg filmom obložene tablete	SE/H/0184/002	HR-H-061130840	MERCK D.O.O	HR
Concor COR 2,5 mg filmom obložene tablete	SE/H/0184/002	HR-H-061130840	MERCK D.O.O	HR
Concor COR 2,5 mg filmom obložene tablete	SE/H/0184/002	HR-H-061130840	MERCK D.O.O	HR
Concor COR 2,5 mg filmom obložene tablete	SE/H/0184/002	HR-H-061130840	MERCK D.O.O	HR
Concor COR 2,5 mg filmom obložene tablete	SE/H/0184/002	HR-H-061130840	MERCK D.O.O	HR
Concor COR 2,5 mg filmom obložene tablete	SE/H/0184/002	HR-H-061130840	MERCK D.O.O	HR
Concor COR 2,5 mg filmom obložene tablete	SE/H/0184/002	HR-H-061130840	MERCK D.O.O	HR
Concor COR 2,5 mg filmom obložene tablete	SE/H/0184/002	HR-H-061130840	MERCK D.O.O	HR
Concor COR 2,5 mg filmom obložene tablete	SE/H/0184/002	HR-H-061130840	MERCK D.O.O	HR
Concor COR 2,5 mg filmom obložene tablete	SE/H/0184/002	HR-H-061130840	MERCK D.O.O	HR
Concor COR 2,5 mg filmom obložene tablete	SE/H/0184/002	HR-H-061130840	MERCK D.O.O	HR
Concor COR 2,5 mg filmom obložene tablete	SE/H/0184/002	HR-H-061130840	MERCK D.O.O	HR
Concor COR 2,5 mg	SE/H/0184/002	HR-H-061130840	MERCK D.O.O	HR

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
filmom obložene tablete				
Concor COR 2,5 mg filmom obložene tablete	SE/H/0184/002	HR-H-061130840	MERCK D.O.O	HR
Concor COR 2,5 mg filmom obložene tablete	SE/H/0184/002	HR-H-061130840	MERCK D.O.O	HR
Concor COR 1,25 mg filmom obložene tablete	SE/H/0184/001	HR-H-350108325	MERCK D.O.O	HR
Concor COR 1,25 mg filmom obložene tablete	SE/H/0184/001	HR-H-350108325	MERCK D.O.O	HR
Concor COR 1,25 mg filmom obložene tablete	SE/H/0184/001	HR-H-350108325	MERCK D.O.O	HR
Concor COR 1,25 mg filmom obložene tablete	SE/H/0184/001	HR-H-350108325	MERCK D.O.O	HR
Concor COR 1,25 mg filmom obložene tablete	SE/H/0184/001	HR-H-350108325	MERCK D.O.O	HR
Concor COR 1,25 mg filmom obložene tablete	SE/H/0184/001	HR-H-350108325	MERCK D.O.O	HR
Concor COR 1,25 mg filmom obložene tablete	SE/H/0184/001	HR-H-350108325	MERCK D.O.O	HR
Concor COR 1,25 mg filmom obložene tablete	SE/H/0184/001	HR-H-350108325	MERCK D.O.O	HR
Concor COR 1,25 mg filmom obložene tablete	SE/H/0184/001	HR-H-350108325	MERCK D.O.O	HR
Concor COR 1,25 mg filmom obložene tablete	SE/H/0184/001	HR-H-350108325	MERCK D.O.O	HR
Concor COR 1,25 mg filmom obložene tablete	SE/H/0184/001	HR-H-350108325	MERCK D.O.O	HR
Concor COR 1,25 mg filmom obložene tablete	SE/H/0184/001	HR-H-350108325	MERCK D.O.O	HR
Concor COR 1,25 mg filmom obložene tablete	SE/H/0184/001	HR-H-350108325	MERCK D.O.O	HR
Concor COR 1,25 mg filmom obložene tablete	SE/H/0184/001	HR-H-350108325	MERCK D.O.O	HR
Concor COR 1,25 mg filmom obložene tablete	SE/H/0184/001	HR-H-350108325	MERCK D.O.O	HR

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
Concor COR 1,25 mg filmom obložene tablete	SE/H/0184/001	HR-H-350108325	MERCK D.O.O	HR
Concor COR 1,25 mg filmtabletta	not available	OGYI-T-8324/01	MERCK KFT.	HU
Concor COR 1,25 mg filmtabletta	not available	OGYI-T-8324/02	MERCK KFT.	HU
Concor COR 10 mg filmtabletta	not available	OGYI-T-8324/12	MERCK KFT.	HU
Concor COR 10 mg filmtabletta	not available	OGYI-T-8324/11	MERCK KFT.	HU
Concor COR 2,5 mg filmtabletta	not available	OGYI-T-8324/04	MERCK KFT.	HU
Concor COR 2,5 mg filmtabletta	not available	OGYI-T-8324/03	MERCK KFT.	HU
Concor COR 5 mg filmtabletta	not available	OGYI-T-8324/07	MERCK KFT.	HU
Concor COR 5 mg filmtabletta	not available	OGYI-T-8324/08	MERCK KFT.	HU
Concor COR 10 mg filmtabletta	not available	OGYI-T-8324/16	MERCK KFT.	HU
Concor COR 5 mg filmtabletta	not available	OGYI-T-8324/15	MERCK KFT.	HU
Concor COR 2,5 mg filmtabletta	not available	OGYI-T-8324/13	MERCK KFT.	HU
Concor COR 1,25 mg filmtabletta	not available	OGYI-T-8324/14	MERCK KFT.	HU
Concor 10 mg filmtabletta	not available	OGYI-T-4015/10	MERCK KFT.	HU
Concor 5 mg filmtabletta	not available	OGYI-T-4015/07	MERCK KFT.	HU
Concor 5 mg filmtabletta	not available	OGYI-T-4015/08	MERCK KFT.	HU
Concor 10 mg filmtabletta	not available	OGYI-T-4015/09	MERCK KFT.	HU
Concor 10 mg filmtabletta	not available	OGYI-T-4015/06	MERCK KFT.	HU
Concor 10 mg	not available	OGYI-T-4015/05	MERCK KFT.	HU

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
filmtabletta				
Concor 10 mg filmtabletta	not available	OGYI-T-4015/04	MERCK KFT.	HU
Concor 5 mg filmtabletta	not available	OGYI-T-4015/03	MERCK KFT.	HU
Concor 5 mg filmtabletta	not available	OGYI-T-4015/02	MERCK KFT.	HU
Concor 5 mg filmtabletta	not available	OGYI-T-4015/01	MERCK KFT.	HU
Bisoprolol Zentiva 5 mg tablets	DE/H/6433/004	PA1701/010/004	ZENTIVA, K.S.	IE
Bisoprolol Zentiva 10 mg tablets	DE/H/6433/006	PA 1701/010/006	ZENTIVA, K.S.	IE
Bisoprolol Zentiva 5 mg tablets	DE/H/6433/004	PA1701/010/004	ZENTIVA, K.S.	IE
Bisoprolol Zentiva 5 mg tablets	DE/H/6433/004	PA1701/010/004	ZENTIVA, K.S.	IE
Bisoprolol Zentiva 5 mg tablets	DE/H/6433/004	PA1701/010/004	ZENTIVA, K.S.	IE
Bisoprolol Zentiva 5 mg tablets	DE/H/6433/004	PA1701/010/004	ZENTIVA, K.S.	IE
Bisoprolol Zentiva 5 mg tablets	DE/H/6433/004	PA1701/010/004	ZENTIVA, K.S.	IE
Bisoprolol Zentiva 5 mg tablets	DE/H/6433/004	PA1701/010/004	ZENTIVA, K.S.	IE
Bisoprolol Zentiva 5 mg tablets	DE/H/6433/004	PA1701/010/004	ZENTIVA, K.S.	IE
Bisoprolol Zentiva 5 mg tablets	DE/H/6433/004	PA1701/010/004	ZENTIVA, K.S.	IE
Bisoprolol Zentiva 5 mg tablets	DE/H/6433/004	PA1701/010/004	ZENTIVA, K.S.	IE
Bisoprolol Zentiva 5 mg tablets	DE/H/6433/004	PA1701/010/004	ZENTIVA, K.S.	IE
Bisoprolol Zentiva 5 mg tablets	DE/H/6433/004	PA1701/010/004	ZENTIVA, K.S.	IE
Bisoprolol Zentiva 5 mg tablets	DE/H/6433/004	PA1701/010/004	ZENTIVA, K.S.	IE
Bisoprolol Zentiva 5 mg	DE/H/6433/004	PA1701/010/004	ZENTIVA, K.S.	IE

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
tablets				
Bisoprolol Zentiva 10 mg tablets	DE/H/6433/006	PA 1701/010/006	ZENTIVA, K.S.	IE
Bisoprolol Zentiva 10 mg tablets	DE/H/6433/006	PA 1701/010/006	ZENTIVA, K.S.	IE
Bisoprolol Zentiva 10 mg tablets	DE/H/6433/006	PA 1701/010/006	ZENTIVA, K.S.	IE
Bisoprolol Zentiva 10 mg tablets	DE/H/6433/006	PA 1701/010/006	ZENTIVA, K.S.	IE
Bisoprolol Zentiva 10 mg tablets	DE/H/6433/006	PA 1701/010/006	ZENTIVA, K.S.	IE
Bisoprolol Zentiva 10 mg tablets	DE/H/6433/006	PA 1701/010/006	ZENTIVA, K.S.	IE
Bisoprolol Zentiva 10 mg tablets	DE/H/6433/006	PA 1701/010/006	ZENTIVA, K.S.	IE
Bisoprolol Zentiva 10 mg tablets	DE/H/6433/006	PA 1701/010/006	ZENTIVA, K.S.	IE
Bisoprolol Zentiva 10 mg tablets	DE/H/6433/006	PA 1701/010/006	ZENTIVA, K.S.	IE
Bisoprolol Zentiva 10 mg tablets	DE/H/6433/006	PA 1701/010/006	ZENTIVA, K.S.	IE
Bisoprolol Zentiva 10 mg tablets	DE/H/6433/006	PA 1701/010/006	ZENTIVA, K.S.	IE
Bisoprolol Zentiva 10 mg tablets	DE/H/6433/006	PA 1701/010/006	ZENTIVA, K.S.	IE
Bisoprolol Zentiva 10 mg tablets	DE/H/6433/006	PA 1701/010/006	ZENTIVA, K.S.	IE
Bisoprolol Syri Pharma 0.5 mg/ml Oral Solution	UK/H/7058/001	PA22697/019/001	SYRI PHARMA LIMITED	IE
Cardicor 1.25 mg film-coated tablets	SE/H/0184/001	PA 2286/4/1	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 1.25 mg film-coated tablets	SE/H/0184/001	PA 2286/4/1	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 1.25 mg film-coated tablets	SE/H/0184/001	PA 2286/4/1	MERCK SERONO (IRELAND) LIMITED	IE

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
Cardicor 1.25 mg film-coated tablets	SE/H/0184/001	PA 2286/4/1	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 1.25 mg film-coated tablets	SE/H/0184/001	PA 2286/4/1	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 1.25 mg film-coated tablets	SE/H/0184/001	PA 2286/4/1	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 1.25 mg film-coated tablets	SE/H/0184/001	PA 2286/4/1	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 1.25 mg film-coated tablets	SE/H/0184/001	PA 2286/4/1	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 1.25 mg film-coated tablets	SE/H/0184/001	PA 2286/4/1	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 1.25 mg film-coated tablets	SE/H/0184/001	PA 2286/4/1	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 1.25 mg film-coated tablets	SE/H/0184/001	PA 2286/4/1	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 1.25 mg film-coated tablets	SE/H/0184/001	PA 2286/4/1	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 1.25 mg film-coated tablets	SE/H/0184/001	PA 2286/4/1	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 1.25 mg film-coated tablets	SE/H/0184/001	PA 2286/4/1	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 1.25 mg film-coated tablets	SE/H/0184/001	PA 2286/4/1	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 1.25 mg film-coated tablets	SE/H/0184/001	PA 2286/4/1	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 10 mg film-coated tablets	SE/H/0184/006	PA 2286/4/6	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 10 mg film-coated tablets	SE/H/0184/006	PA 2286/4/6	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 10 mg film-coated tablets	SE/H/0184/006	PA 2286/4/6	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 10 mg film-coated tablets	SE/H/0184/006	PA 2286/4/6	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 10 mg film-	SE/H/0184/006	PA 2286/4/6	MERCK SERONO (IRELAND)	IE

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
coated tablets			LIMITED	
Cardicor 10 mg film-coated tablets	SE/H/0184/006	PA 2286/4/6	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 10 mg film-coated tablets	SE/H/0184/006	PA 2286/4/6	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 10 mg film-coated tablets	SE/H/0184/006	PA 2286/4/6	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 10 mg film-coated tablets	SE/H/0184/006	PA 2286/4/6	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 10 mg film-coated tablets	SE/H/0184/006	PA 2286/4/6	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 10 mg film-coated tablets	SE/H/0184/006	PA 2286/4/6	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 10 mg film-coated tablets	SE/H/0184/006	PA 2286/4/6	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 10 mg film-coated tablets	SE/H/0184/006	PA 2286/4/6	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 10 mg film-coated tablets	SE/H/0184/006	PA 2286/4/6	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 10 mg film-coated tablets	SE/H/0184/006	PA 2286/4/6	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 10 mg film-coated tablets	SE/H/0184/006	PA 2286/4/6	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 2.5 mg film-coated tablets	SE/H/0184/002	PA 2286/4/2	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 2.5 mg film-coated tablets	SE/H/0184/002	PA 2286/4/2	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 2.5 mg film-coated tablets	SE/H/0184/002	PA 2286/4/2	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 2.5 mg film-coated tablets	SE/H/0184/002	PA 2286/4/2	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 2.5 mg film-coated tablets	SE/H/0184/002	PA 2286/4/2	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 2.5 mg film-coated tablets	SE/H/0184/002	PA 2286/4/2	MERCK SERONO (IRELAND) LIMITED	IE

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
Cardicor 2.5 mg film-coated tablets	SE/H/0184/002	PA 2286/4/2	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 2.5 mg film-coated tablets	SE/H/0184/002	PA 2286/4/2	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 2.5 mg film-coated tablets	SE/H/0184/002	PA 2286/4/2	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 2.5 mg film-coated tablets	SE/H/0184/002	PA 2286/4/2	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 2.5 mg film-coated tablets	SE/H/0184/002	PA 2286/4/2	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 2.5 mg film-coated tablets	SE/H/0184/002	PA 2286/4/2	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 2.5 mg film-coated tablets	SE/H/0184/002	PA 2286/4/2	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 2.5 mg film-coated tablets	SE/H/0184/002	PA 2286/4/2	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 2.5 mg film-coated tablets	SE/H/0184/002	PA 2286/4/2	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 2.5 mg film-coated tablets	SE/H/0184/002	PA 2286/4/2	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 5 mg film-coated tablets	SE/H/0184/004	PA 2286/4/4	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 5 mg film-coated tablets	SE/H/0184/004	PA 2286/4/4	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 5 mg film-coated tablets	SE/H/0184/004	PA 2286/4/4	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 5 mg film-coated tablets	SE/H/0184/004	PA 2286/4/4	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 5 mg film-coated tablets	SE/H/0184/004	PA 2286/4/4	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 5 mg film-coated tablets	SE/H/0184/004	PA 2286/4/4	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 5 mg film-coated tablets	SE/H/0184/004	PA 2286/4/4	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 5 mg film-	SE/H/0184/004	PA 2286/4/4	MERCK SERONO (IRELAND)	IE

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
coated tablets			LIMITED	
Cardicor 5 mg film-coated tablets	SE/H/0184/004	PA 2286/4/4	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 5 mg film-coated tablets	SE/H/0184/004	PA 2286/4/4	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 5 mg film-coated tablets	SE/H/0184/004	PA 2286/4/4	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 5 mg film-coated tablets	SE/H/0184/004	PA 2286/4/4	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 5 mg film-coated tablets	SE/H/0184/004	PA 2286/4/4	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 5 mg film-coated tablets	SE/H/0184/004	PA 2286/4/4	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 5 mg film-coated tablets	SE/H/0184/004	PA 2286/4/4	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 5 mg film-coated tablets	SE/H/0184/004	PA 2286/4/4	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 3.75 mg film-coated tablets	SE/H/0184/003	PA 2286/4/3	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 3.75 mg film-coated tablets	SE/H/0184/003	PA 2286/4/3	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 3.75 mg film-coated tablets	SE/H/0184/003	PA 2286/4/3	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 3.75 mg film-coated tablets	SE/H/0184/003	PA 2286/4/3	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 3.75 mg film-coated tablets	SE/H/0184/003	PA 2286/4/3	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 3.75 mg film-coated tablets	SE/H/0184/003	PA 2286/4/3	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 3.75 mg film-coated tablets	SE/H/0184/003	PA 2286/4/3	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 3.75 mg film-coated tablets	SE/H/0184/003	PA 2286/4/3	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 3.75 mg film-coated tablets	SE/H/0184/003	PA 2286/4/3	MERCK SERONO (IRELAND) LIMITED	IE

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
Cardicor 3.75 mg film-coated tablets	SE/H/0184/003	PA 2286/4/3	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 3.75 mg film-coated tablets	SE/H/0184/003	PA 2286/4/3	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 3.75 mg film-coated tablets	SE/H/0184/003	PA 2286/4/3	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 3.75 mg film-coated tablets	SE/H/0184/003	PA 2286/4/3	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 3.75 mg film-coated tablets	SE/H/0184/003	PA 2286/4/3	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 3.75 mg film-coated tablets	SE/H/0184/003	PA 2286/4/3	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 3.75 mg film-coated tablets	SE/H/0184/003	PA 2286/4/3	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 7.5 mg film-coated tablets	SE/H/0184/005	PA 2286/4/5	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 7.5 mg film-coated tablets	SE/H/0184/005	PA 2286/4/5	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 7.5 mg film-coated tablets	SE/H/0184/005	PA 2286/4/5	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 7.5 mg film-coated tablets	SE/H/0184/005	PA 2286/4/5	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 7.5 mg film-coated tablets	SE/H/0184/005	PA 2286/4/5	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 7.5 mg film-coated tablets	SE/H/0184/005	PA 2286/4/5	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 7.5 mg film-coated tablets	SE/H/0184/005	PA 2286/4/5	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 7.5 mg film-coated tablets	SE/H/0184/005	PA 2286/4/5	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 7.5 mg film-coated tablets	SE/H/0184/005	PA 2286/4/5	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 7.5 mg film-coated tablets	SE/H/0184/005	PA 2286/4/5	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 7.5 mg film-	SE/H/0184/005	PA 2286/4/5	MERCK SERONO (IRELAND)	IE

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
coated tablets			LIMITED	
Cardicor 7.5 mg film-coated tablets	SE/H/0184/005	PA 2286/4/5	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 7.5 mg film-coated tablets	SE/H/0184/005	PA 2286/4/5	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 7.5 mg film-coated tablets	SE/H/0184/005	PA 2286/4/5	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 7.5 mg film-coated tablets	SE/H/0184/005	PA 2286/4/5	MERCK SERONO (IRELAND) LIMITED	IE
Cardicor 7.5 mg film-coated tablets	SE/H/0184/005	PA 2286/4/5	MERCK SERONO (IRELAND) LIMITED	IE
Bisoprololo Zentiva 5 mg compresse rivestite con film.	IT/H/0173/001	037690017	ZENTIVA ITALIA S.R.L.	IT
Bisoprololo Zentiva 5 mg compresse rivestite con film.	IT/H/0173/001	037690195	ZENTIVA ITALIA S.R.L.	IT
Bisoprololo Zentiva 5 mg compresse rivestite con film.	IT/H/0173/001	037690070	ZENTIVA ITALIA S.R.L.	IT
Bisoprololo Zentiva 5 mg compresse rivestite con film.	IT/H/0173/001	037690106	ZENTIVA ITALIA S.R.L.	IT
Bisoprololo Zentiva 5 mg compresse rivestite con film.	IT/H/0173/001	037690157	ZENTIVA ITALIA S.R.L.	IT
Bisoprololo Zentiva 5 mg compresse rivestite con film.	IT/H/0173/001	037690082	ZENTIVA ITALIA S.R.L.	IT
Bisoprololo Zentiva 5 mg compresse rivestite con film.	IT/H/0173/001	037690029	ZENTIVA ITALIA S.R.L.	IT
Bisoprololo Zentiva 5 mg compresse rivestite con film.	IT/H/0173/001	037690043	ZENTIVA ITALIA S.R.L.	IT

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
Bisoprololo Zentiva 5 mg compresse rivestite con film.	IT/H/0173/001	037690120	ZENTIVA ITALIA S.R.L.	IT
Bisoprololo Zentiva 5 mg compresse rivestite con film.	IT/H/0173/001	037690094	ZENTIVA ITALIA S.R.L.	IT
Bisoprololo Zentiva 5 mg compresse rivestite con film.	IT/H/0173/001	037690031	ZENTIVA ITALIA S.R.L.	IT
Bisoprololo Zentiva 5 mg compresse rivestite con film.	IT/H/0173/001	037690056	ZENTIVA ITALIA S.R.L.	IT
Bisoprololo Zentiva 5 mg compresse rivestite con film.	IT/H/0173/001	037690118	ZENTIVA ITALIA S.R.L.	IT
Bisoprololo Zentiva 5 mg compresse rivestite con film.	IT/H/0173/001	037690169	ZENTIVA ITALIA S.R.L.	IT
Bisoprololo Zentiva 5 mg compresse rivestite con film.	IT/H/0173/001	037690068	ZENTIVA ITALIA S.R.L.	IT
Bisoprololo Zentiva 5 mg compresse rivestite con film.	IT/H/0173/001	037690132	ZENTIVA ITALIA S.R.L.	IT
Bisoprololo Zentiva 5 mg compresse rivestite con film.	IT/H/0173/001	037690144	ZENTIVA ITALIA S.R.L.	IT
Bisoprololo Zentiva Generics 5 mg compresse	DE/H/6433/004	048314355	ZENTIVA ITALIA S.R.L.	IT
Bisoprololo Zentiva Generics 5 mg compresse	DE/H/6433/004	048314367	ZENTIVA ITALIA S.R.L.	IT
Bisoprololo Zentiva Generics 5 mg	DE/H/6433/004	048314379	ZENTIVA ITALIA S.R.L.	IT

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
compresse				
Bisoprololo Zentiva Generics 5 mg compresse	DE/H/6433/004	048314381	ZENTIVA ITALIA S.R.L.	IT
Bisoprololo Zentiva Generics 5 mg compresse	DE/H/6433/004	048314393	ZENTIVA ITALIA S.R.L.	IT
Bisoprololo Zentiva Generics 5 mg compresse	DE/H/6433/004	048314405	ZENTIVA ITALIA S.R.L.	IT
Bisoprololo Zentiva Generics 5 mg compresse	DE/H/6433/004	048314417	ZENTIVA ITALIA S.R.L.	IT
Bisoprololo Zentiva Generics 5 mg compresse	DE/H/6433/004	048314429	ZENTIVA ITALIA S.R.L.	IT
Bisoprololo Zentiva Generics 5 mg compresse	DE/H/6433/004	048314431	ZENTIVA ITALIA S.R.L.	IT
Bisoprololo Zentiva Generics 5 mg compresse	DE/H/6433/004	048314443	ZENTIVA ITALIA S.R.L.	IT
Bisoprololo Zentiva Generics 5 mg compresse	DE/H/6433/004	048314456	ZENTIVA ITALIA S.R.L.	IT
Bisoprololo Zentiva Generics 5 mg compresse	DE/H/6433/004	048314468	ZENTIVA ITALIA S.R.L.	IT
Bisoprololo Zentiva Generics 5 mg compresse	DE/H/6433/004	048314470	ZENTIVA ITALIA S.R.L.	IT
Bisoprololo Zentiva Generics 5 mg compresse	DE/H/6433/004	048314482	ZENTIVA ITALIA S.R.L.	IT
Bisoprololo Zentiva	DE/H/6433/006	048314595	ZENTIVA ITALIA S.R.L.	IT

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
Generics 10 mg compresse				
Bisoprololo Zentiva Generics 10 mg compresse	DE/H/6433/006	048314607	ZENTIVA ITALIA S.R.L.	IT
Bisoprololo Zentiva Generics 10 mg compresse	DE/H/6433/006	048314619	ZENTIVA ITALIA S.R.L.	IT
Bisoprololo Zentiva Generics 10 mg compresse	DE/H/6433/006	048314621	ZENTIVA ITALIA S.R.L.	IT
Bisoprololo Zentiva Generics 10 mg compresse	DE/H/6433/006	048314633	ZENTIVA ITALIA S.R.L.	IT
Bisoprololo Zentiva Generics 10 mg compresse	DE/H/6433/006	048314645	ZENTIVA ITALIA S.R.L.	IT
Bisoprololo Zentiva Generics 10 mg compresse	DE/H/6433/006	048314658	ZENTIVA ITALIA S.R.L.	IT
Bisoprololo Zentiva Generics 10 mg compresse	DE/H/6433/006	048314660	ZENTIVA ITALIA S.R.L.	IT
Bisoprololo Zentiva Generics 10 mg compresse	DE/H/6433/006	048314672	ZENTIVA ITALIA S.R.L.	IT
Bisoprololo Zentiva Generics 10 mg compresse	DE/H/6433/006	048314684	ZENTIVA ITALIA S.R.L.	IT
Bisoprololo Zentiva Generics 10 mg compresse	DE/H/6433/006	048314696	ZENTIVA ITALIA S.R.L.	IT
Bisoprololo Zentiva Generics 10 mg compresse	DE/H/6433/006	048314708	ZENTIVA ITALIA S.R.L.	IT

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
Bisoprololo Zentiva Generics 10 mg compresse	DE/H/6433/006	048314710	ZENTIVA ITALIA S.R.L.	IT
Bisoprololo Zentiva Generics 10 mg compresse	DE/H/6433/006	048314722	ZENTIVA ITALIA S.R.L.	IT
CONCOR 10 mg compresse rivestite con film	not available	026573016	DOMPÉ FARMACEUTICI S.P.A.	IT
SEQUACOR 1,25 mg compresse rivestite con film	SE/H/0184/001	034952010	DOMPÉ FARMACEUTICI S.P.A.	IT
SEQUACOR 1,25 mg compresse rivestite con film	SE/H/0184/001	034952022	DOMPÉ FARMACEUTICI S.P.A.	IT
SEQUACOR 1,25 mg compresse rivestite con film	SE/H/0184/001	034952046	DOMPÉ FARMACEUTICI S.P.A.	IT
SEQUACOR 1,25 mg compresse rivestite con film	SE/H/0184/001	034952061	DOMPÉ FARMACEUTICI S.P.A.	IT
SEQUACOR 1,25 mg compresse rivestite con film	SE/H/0184/001	034952059	DOMPÉ FARMACEUTICI S.P.A.	IT
SEQUACOR 1,25 mg compresse rivestite con film	SE/H/0184/001	034952085	DOMPÉ FARMACEUTICI S.P.A.	IT
SEQUACOR 1,25 mg compresse rivestite con film	SE/H/0184/001	034952491	DOMPÉ FARMACEUTICI S.P.A.	IT
SEQUACOR 1,25 mg compresse rivestite con film	SE/H/0184/001	034952034	DOMPÉ FARMACEUTICI S.P.A.	IT
SEQUACOR 1,25 mg compresse rivestite con	SE/H/0184/001	034952073	DOMPÉ FARMACEUTICI S.P.A.	IT

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
film				
SEQUACOR 10 mg compresse rivestite con film	SE/H/0184/006	034952414	DOMPÉ FARMACEUTICI S.P.A.	IT
SEQUACOR 10 mg compresse rivestite con film	SE/H/0184/006	034952477	DOMPÉ FARMACEUTICI S.P.A.	IT
SEQUACOR 10 mg compresse rivestite con film	SE/H/0184/006	034952453	DOMPÉ FARMACEUTICI S.P.A.	IT
SEQUACOR 10 mg compresse rivestite con film	SE/H/0184/006	034952438	DOMPÉ FARMACEUTICI S.P.A.	IT
SEQUACOR 10 mg compresse rivestite con film	SE/H/0184/006	034952440	DOMPÉ FARMACEUTICI S.P.A.	IT
SEQUACOR 10 mg compresse rivestite con film	SE/H/0184/006	034952426	DOMPÉ FARMACEUTICI S.P.A.	IT
SEQUACOR 10 mg compresse rivestite con film	SE/H/0184/006	034952465	DOMPÉ FARMACEUTICI S.P.A.	IT
SEQUACOR 10 mg compresse rivestite con film	SE/H/0184/006	034952541	DOMPÉ FARMACEUTICI S.P.A.	IT
SEQUACOR 10 mg compresse rivestite con film	SE/H/0184/006	034952489	DOMPÉ FARMACEUTICI S.P.A.	IT
SEQUACOR 2,5 mg compresse rivestite con film	SE/H/0184/002	034952097	DOMPÉ FARMACEUTICI S.P.A.	IT
SEQUACOR 2,5 mg compresse rivestite con film	SE/H/0184/002	034952111	DOMPÉ FARMACEUTICI S.P.A.	IT
SEQUACOR 2,5 mg	SE/H/0184/002	034952109	DOMPÉ FARMACEUTICI	IT

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
compresse rivestite con film			S.P.A.	
SEQUACOR 2,5 mg compresse rivestite con film	SE/H/0184/002	034952150	DOMPÉ FARMACEUTICI S.P.A.	IT
SEQUACOR 2,5 mg compresse rivestite con film	SE/H/0184/002	034952503	DOMPÉ FARMACEUTICI S.P.A.	IT
SEQUACOR 2,5 mg compresse rivestite con film	SE/H/0184/002	034952162	DOMPÉ FARMACEUTICI S.P.A.	IT
SEQUACOR 2,5 mg compresse rivestite con film	SE/H/0184/002	034952147	DOMPÉ FARMACEUTICI S.P.A.	IT
SEQUACOR 2,5 mg compresse rivestite con film	SE/H/0184/002	034952123	DOMPÉ FARMACEUTICI S.P.A.	IT
SEQUACOR 2,5 mg compresse rivestite con film	SE/H/0184/002	034952135	DOMPÉ FARMACEUTICI S.P.A.	IT
SEQUACOR 3,75 mg compresse rivestite con film	SE/H/0184/003	034952186	DOMPÉ FARMACEUTICI S.P.A.	IT
SEQUACOR 3,75 mg compresse rivestite con film	SE/H/0184/003	034952248	DOMPÉ FARMACEUTICI S.P.A.	IT
SEQUACOR 3,75 mg compresse rivestite con film	SE/H/0184/003	034952200	DOMPÉ FARMACEUTICI S.P.A.	IT
SEQUACOR 3,75 mg compresse rivestite con film	SE/H/0184/003	034952515	DOMPÉ FARMACEUTICI S.P.A.	IT
SEQUACOR 3,75 mg compresse rivestite con film	SE/H/0184/003	034952174	DOMPÉ FARMACEUTICI S.P.A.	IT

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
SEQUACOR 3,75 mg compresse rivestite con film	SE/H/0184/003	034952236	DOMPÉ FARMACEUTICI S.P.A.	IT
SEQUACOR 3,75 mg compresse rivestite con film	SE/H/0184/003	034952212	DOMPÉ FARMACEUTICI S.P.A.	IT
SEQUACOR 3,75 mg compresse rivestite con film	SE/H/0184/003	034952198	DOMPÉ FARMACEUTICI S.P.A.	IT
SEQUACOR 3,75 mg compresse rivestite con film	SE/H/0184/003	034952224	DOMPÉ FARMACEUTICI S.P.A.	IT
SEQUACOR 5 mg compresse rivestite con film	SE/H/0184/004	034952275	DOMPÉ FARMACEUTICI S.P.A.	IT
SEQUACOR 5 mg compresse rivestite con film	SE/H/0184/004	034952263	DOMPÉ FARMACEUTICI S.P.A.	IT
SEQUACOR 5 mg compresse rivestite con film	SE/H/0184/004	034952527	DOMPÉ FARMACEUTICI S.P.A.	IT
SEQUACOR 5 mg compresse rivestite con film	SE/H/0184/004	034952301	DOMPÉ FARMACEUTICI S.P.A.	IT
SEQUACOR 5 mg compresse rivestite con film	SE/H/0184/004	034952313	DOMPÉ FARMACEUTICI S.P.A.	IT
SEQUACOR 5 mg compresse rivestite con film	SE/H/0184/004	034952299	DOMPÉ FARMACEUTICI S.P.A.	IT
SEQUACOR 5 mg compresse rivestite con film	SE/H/0184/004	034952287	DOMPÉ FARMACEUTICI S.P.A.	IT
SEQUACOR 5 mg compresse rivestite con	SE/H/0184/004	034952325	DOMPÉ FARMACEUTICI S.P.A.	IT

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
film				
SEQUACOR 5 mg compresse rivestite con film	SE/H/0184/004	034952251	DOMPÉ FARMACEUTICI S.P.A.	IT
SEQUACOR 7,5 mg compresse rivestite con film	SE/H/0184/005	034952349	DOMPÉ FARMACEUTICI S.P.A.	IT
SEQUACOR 7,5 mg compresse rivestite con film	SE/H/0184/005	034952364	DOMPÉ FARMACEUTICI S.P.A.	IT
SEQUACOR 7,5 mg compresse rivestite con film	SE/H/0184/005	034952376	DOMPÉ FARMACEUTICI S.P.A.	IT
SEQUACOR 7,5 mg compresse rivestite con film	SE/H/0184/005	034952352	DOMPÉ FARMACEUTICI S.P.A.	IT
SEQUACOR 7,5 mg compresse rivestite con film	SE/H/0184/005	034952337	DOMPÉ FARMACEUTICI S.P.A.	IT
SEQUACOR 7,5 mg compresse rivestite con film	SE/H/0184/005	034952390	DOMPÉ FARMACEUTICI S.P.A.	IT
SEQUACOR 7,5 mg compresse rivestite con film	SE/H/0184/005	034952402	DOMPÉ FARMACEUTICI S.P.A.	IT
SEQUACOR 7,5 mg compresse rivestite con film	SE/H/0184/005	034952539	DOMPÉ FARMACEUTICI S.P.A.	IT
SEQUACOR 7,5 mg compresse rivestite con film	SE/H/0184/005	034952388	DOMPÉ FARMACEUTICI S.P.A.	IT
Cardicor 5 mg compresse rivestite con film	SE/H/0185/004	034954267	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Cardicor 5 mg	SE/H/0185/004	034954279	RECORDATI INDUSTRIA	IT

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
compresse rivestite con film			CHIMICA E FARMACEUTICA S.P.A.	
Cardicor 5 mg compresse rivestite con film	SE/H/0185/004	034954281	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Cardicor 5 mg compresse rivestite con film	SE/H/0185/004	034954293	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Cardicor 5 mg compresse rivestite con film	SE/H/0185/004	034954305	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Cardicor 5 mg compresse rivestite con film	SE/H/0185/004	034954317	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Cardicor 5 mg compresse rivestite con film	SE/H/0185/004	034954329	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Cardicor 1,25 mg compresse rivestite con film	SE/H/0185/001	034954014	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Cardicor 1,25 mg compresse rivestite con film	SE/H/0185/001	034954038	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Cardicor 1,25 mg compresse rivestite con film	SE/H/0185/001	034954040	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Cardicor 1,25 mg compresse rivestite con film	SE/H/0185/001	034954053	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Cardicor 1,25 mg compresse rivestite con film	SE/H/0185/001	034954065	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Cardicor 1,25 mg compresse rivestite con film	SE/H/0185/001	034954077	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT

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
Cardicor 1,25 mg compresse rivestite con film	SE/H/0185/001	034954089	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Cardicor 2,5 mg compresse rivestite con film	SE/H/0185/002	034954103	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Cardicor 2,5 mg compresse rivestite con film	SE/H/0185/002	034954115	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Cardicor 2,5 mg compresse rivestite con film	SE/H/0185/002	034954127	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Cardicor 2,5 mg compresse rivestite con film	SE/H/0185/002	034954139	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Cardicor 2,5 mg compresse rivestite con film	SE/H/0185/002	034954141	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Cardicor 2,5 mg compresse rivestite con film	SE/H/0185/002	034954154	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Cardicor 2,5 mg compresse rivestite con film	SE/H/0185/002	034954166	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Cardicor 3,75 mg compresse rivestite con film	SE/H/0185/003	034954180	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Cardicor 3,75 mg compresse rivestite con film	SE/H/0185/003	034954192	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Cardicor 3,75 mg compresse rivestite con film	SE/H/0185/003	034954204	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Cardicor 3,75 mg compresse rivestite con	SE/H/0185/003	034954216	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA	IT

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
film			S.P.A.	
Cardicor 3,75 mg compresse rivestite con film	SE/H/0185/003	034954228	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Cardicor 3,75 mg compresse rivestite con film	SE/H/0185/003	034954230	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Cardicor 3,75 mg compresse rivestite con film	SE/H/0185/003	034954242	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Cardicor 7,5 mg compresse rivestite con film	SE/H/0185/005	034954343	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Cardicor 7,5 mg compresse rivestite con film	SE/H/0185/005	034954356	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Cardicor 7,5 mg compresse rivestite con film	SE/H/0185/005	034954368	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Cardicor 7,5 mg compresse rivestite con film	SE/H/0185/005	034954370	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Cardicor 7,5 mg compresse rivestite con film	SE/H/0185/005	034954382	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Cardicor 7,5 mg compresse rivestite con film	SE/H/0185/005	034954394	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Cardicor 7,5 mg compresse rivestite con film	SE/H/0185/005	034954406	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Cardicor 10 mg compresse rivestite con film	SE/H/0185/006	034954420	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Cardicor 10 mg	SE/H/0185/006	034954432	RECORDATI INDUSTRIA	IT

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
compresse rivestite con film			CHIMICA E FARMACEUTICA S.P.A.	
Cardicor 10 mg compresse rivestite con film	SE/H/0185/006	034954444	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Cardicor 10 mg compresse rivestite con film	SE/H/0185/006	034954457	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Cardicor 10 mg compresse rivestite con film	SE/H/0185/006	034954469	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Cardicor 10 mg compresse rivestite con film	SE/H/0185/006	034954471	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Cardicor 10 mg compresse rivestite con film	SE/H/0185/006	034954483	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Cardicor 1,25 mg compresse rivestite con film	SE/H/0185/001	034954026	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Cardicor 2,5 mg compresse rivestite con film	SE/H/0185/002	034954091	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Cardicor 3,75 mg compresse rivestite con film	SE/H/0185/003	034954178	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Cardicor 5 mg compresse rivestite con film	SE/H/0185/004	034954255	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Cardicor 7,5 mg compresse rivestite con film	SE/H/0185/005	034954331	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Cardicor 10 mg compresse rivestite con film	SE/H/0185/006	034954418	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT

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
CONGESCOR 1,25 mg compresse rivestite con film	SE/H/0186/001	034953012	MERCK SERONO S.P.A.	IT
CONGESCOR 1,25 mg compresse rivestite con film	SE/H/0186/001	034953024	MERCK SERONO S.P.A.	IT
CONGESCOR 1,25 mg compresse rivestite con film	SE/H/0186/001	034953036	MERCK SERONO S.P.A.	IT
CONGESCOR 1,25 mg compresse rivestite con film	SE/H/0186/001	034953048	MERCK SERONO S.P.A.	IT
CONGESCOR 1,25 mg compresse rivestite con film	SE/H/0186/001	034953051	MERCK SERONO S.P.A.	IT
CONGESCOR 1,25 mg compresse rivestite con film	SE/H/0186/001	034953063	MERCK SERONO S.P.A.	IT
CONGESCOR 1,25 mg compresse rivestite con film	SE/H/0186/001	034953075	MERCK SERONO S.P.A.	IT
CONGESCOR 1,25 mg compresse rivestite con film	SE/H/0186/001	034953481	MERCK SERONO S.P.A.	IT
CONGESCOR 2,5 mg compresse rivestite con film	SE/H/0186/002	034953087	MERCK SERONO S.P.A.	IT
CONGESCOR 3,75 mg compresse rivestite con film	SE/H/0186/003	034953164	MERCK SERONO S.P.A.	IT
CONGESCOR 5 mg compresse rivestite con film	SE/H/0186/004	034953240	MERCK SERONO S.P.A.	IT
CONGESCOR 7,5 mg compresse rivestite con	SE/H/0186/005	034953327	MERCK SERONO S.P.A.	IT

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
film				
CONGESCOR 2,5 mg compresse rivestite con film	SE/H/0186/002	034953099	MERCK SERONO S.P.A.	IT
CONGESCOR 2,5 mg compresse rivestite con film	SE/H/0186/002	034953101	MERCK SERONO S.P.A.	IT
CONGESCOR 2,5 mg compresse rivestite con film	SE/H/0186/002	034953113	MERCK SERONO S.P.A.	IT
CONGESCOR 2,5 mg compresse rivestite con film	SE/H/0186/002	034953125	MERCK SERONO S.P.A.	IT
CONGESCOR 2,5 mg compresse rivestite con film	SE/H/0186/002	034953137	MERCK SERONO S.P.A.	IT
CONGESCOR 2,5 mg compresse rivestite con film	SE/H/0186/002	034953149	MERCK SERONO S.P.A.	IT
CONGESCOR 2,5 mg compresse rivestite con film	SE/H/0186/002	034953152	MERCK SERONO S.P.A.	IT
CONGESCOR 3,75 mg compresse rivestite con film	SE/H/0186/003	034953176	MERCK SERONO S.P.A.	IT
CONGESCOR 3,75 mg compresse rivestite con film	SE/H/0186/003	034953188	MERCK SERONO S.P.A.	IT
CONGESCOR 3,75 mg compresse rivestite con film	SE/H/0186/003	034953214	MERCK SERONO S.P.A.	IT
CONGESCOR 3,75 mg compresse rivestite con film	SE/H/0186/003	034953202	MERCK SERONO S.P.A.	IT
CONGESCOR 3,75 mg	SE/H/0186/003	034953226	MERCK SERONO S.P.A.	IT

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
compresse rivestite con film				
CONGESCOR 3,75 mg compresse rivestite con film	SE/H/0186/003	034953238	MERCK SERONO S.P.A.	IT
CONGESCOR 5 mg compresse rivestite con film	SE/H/0186/004	034953253	MERCK SERONO S.P.A.	IT
CONGESCOR 5 mg compresse rivestite con film	SE/H/0186/004	034953277	MERCK SERONO S.P.A.	IT
CONGESCOR 5 mg compresse rivestite con film	SE/H/0186/004	034953265	MERCK SERONO S.P.A.	IT
CONGESCOR 5 mg compresse rivestite con film	SE/H/0186/004	034953289	MERCK SERONO S.P.A.	IT
CONGESCOR 5 mg compresse rivestite con film	SE/H/0186/004	034953291	MERCK SERONO S.P.A.	IT
CONGESCOR 5 mg compresse rivestite con film	SE/H/0186/004	034953303	MERCK SERONO S.P.A.	IT
CONGESCOR 5 mg compresse rivestite con film	SE/H/0186/004	034953315	MERCK SERONO S.P.A.	IT
CONGESCOR 7,5 mg compresse rivestite con film	SE/H/0186/005	034953339	MERCK SERONO S.P.A.	IT
CONGESCOR 7,5 mg compresse rivestite con film	SE/H/0186/005	034953380	MERCK SERONO S.P.A.	IT
CONGESCOR 7,5 mg compresse rivestite con film	SE/H/0186/005	034953378	MERCK SERONO S.P.A.	IT

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
CONGESCOR 7,5 mg compresse rivestite con film	SE/H/0186/005	034953392	MERCK SERONO S.P.A.	IT
CONGESCOR 10 mg compresse rivestite con film	SE/H/0186/006	034953416	MERCK SERONO S.P.A.	IT
CONGESCOR 10 mg compresse rivestite con film	SE/H/0186/006	034953428	MERCK SERONO S.P.A.	IT
CONGESCOR 10 mg compresse rivestite con film	SE/H/0186/006	034953430	MERCK SERONO S.P.A.	IT
CONGESCOR 10 mg compresse rivestite con film	SE/H/0186/006	034953467	MERCK SERONO S.P.A.	IT
CONGESCOR 10 mg compresse rivestite con film	SE/H/0186/006	034953455	MERCK SERONO S.P.A.	IT
CONGESCOR 10 mg compresse rivestite con film	SE/H/0186/006	034953442	MERCK SERONO S.P.A.	IT
CONGESCOR 10 mg compresse rivestite con film	SE/H/0186/006	034953479	MERCK SERONO S.P.A.	IT
Bisoprolol fumarate Zentiva 5 mg tabletės	DE/H/6433/004	LT/1/21/4754/007	ZENTIVA, K.S.	LT
Bisoprolol fumarate Zentiva 5 mg tabletės	DE/H/6433/004	LT/1/21/4754/006	ZENTIVA, K.S.	LT
Bisoprolol fumarate Zentiva 5 mg tabletės	DE/H/6433/004	LT/1/21/4754/001	ZENTIVA, K.S.	LT
Bisoprolol fumarate Zentiva 5 mg tabletės	DE/H/6433/004	LT/1/21/4754/002	ZENTIVA, K.S.	LT
Bisoprolol fumarate Zentiva 5 mg tabletės	DE/H/6433/004	LT/1/21/4754/003	ZENTIVA, K.S.	LT
Bisoprolol fumarate	DE/H/6433/004	LT/1/21/4754/004	ZENTIVA, K.S.	LT

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
Zentiva 5 mg tabletės				
Bisoprolol fumarate Zentiva 5 mg tabletės	DE/H/6433/004	LT/1/21/4754/005	ZENTIVA, K.S.	LT
Bisoprolol fumarate Zentiva 10 mg tabletės	DE/H/6433/006	LT/1/21/4755/007	ZENTIVA, K.S.	LT
Bisoprolol fumarate Zentiva 10 mg tabletės	DE/H/6433/006	LT/1/21/4755/001	ZENTIVA, K.S.	LT
Bisoprolol fumarate Zentiva 10 mg tabletės	DE/H/6433/006	LT/1/21/4755/002	ZENTIVA, K.S.	LT
Bisoprolol fumarate Zentiva 10 mg tabletės	DE/H/6433/006	LT/1/21/4755/003	ZENTIVA, K.S.	LT
Bisoprolol fumarate Zentiva 10 mg tabletės	DE/H/6433/006	LT/1/21/4755/004	ZENTIVA, K.S.	LT
Bisoprolol fumarate Zentiva 10 mg tabletės	DE/H/6433/006	LT/1/21/4755/005	ZENTIVA, K.S.	LT
Bisoprolol fumarate Zentiva 10 mg tabletės	DE/H/6433/006	LT/1/21/4755/006	ZENTIVA, K.S.	LT
Bisoprolol fumarate Zentiva 5 mg tabletės	DE/H/6433/004	LT/1/21/4754/001	ZENTIVA, K.S.	LT
Bisoprolol fumarate Zentiva 5 mg tabletės	DE/H/6433/004	LT/1/21/4754/002	ZENTIVA, K.S.	LT
Bisoprolol fumarate Zentiva 5 mg tabletės	DE/H/6433/004	LT/1/21/4754/003	ZENTIVA, K.S.	LT
Bisoprolol fumarate Zentiva 5 mg tabletės	DE/H/6433/004	LT/1/21/4754/004	ZENTIVA, K.S.	LT
Bisoprolol fumarate Zentiva 5 mg tabletės	DE/H/6433/004	LT/1/21/4754/005	ZENTIVA, K.S.	LT
Bisoprolol fumarate Zentiva 5 mg tabletės	DE/H/6433/004	LT/1/21/4754/006	ZENTIVA, K.S.	LT
Bisoprolol fumarate Zentiva 5 mg tabletės	DE/H/6433/004	LT/1/21/4754/007	ZENTIVA, K.S.	LT
Bisoprolol fumarate Zentiva 10 mg tabletės	DE/H/6433/006	LT/1/21/4755/001	ZENTIVA, K.S.	LT
Bisoprolol fumarate Zentiva 10 mg tabletės	DE/H/6433/006	LT/1/21/4755/002	ZENTIVA, K.S.	LT

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
Bisoprolol fumarate Zentiva 10 mg tabletės	DE/H/6433/006	LT/1/21/4755/003	ZENTIVA, K.S.	LT
Bisoprolol fumarate Zentiva 10 mg tabletės	DE/H/6433/006	LT/1/21/4755/004	ZENTIVA, K.S.	LT
Bisoprolol fumarate Zentiva 10 mg tabletės	DE/H/6433/006	LT/1/21/4755/005	ZENTIVA, K.S.	LT
Bisoprolol fumarate Zentiva 10 mg tabletės	DE/H/6433/006	LT/1/21/4755/006	ZENTIVA, K.S.	LT
Bisoprolol fumarate Zentiva 10 mg tabletės	DE/H/6433/006	LT/1/21/4755/007	ZENTIVA, K.S.	LT
Concor COR 2,5 mg Filmtabletten	SE/H/0184/002	2005068803	MERCK HEALTHCARE GERMANY GMBH	LU
Concor COR 2,5 mg Filmtabletten	SE/H/0184/002	2005068803	MERCK HEALTHCARE GERMANY GMBH	LU
Concor COR 2,5 mg Filmtabletten	SE/H/0184/002	2005068803	MERCK HEALTHCARE GERMANY GMBH	LU
Concor 10 mg Filmtabletten	not available	2008089996	MERCK HEALTHCARE GERMANY GMBH	LU
Concor 5 mg Filmtabletten	not available	2008089995	MERCK HEALTHCARE GERMANY GMBH	LU
Concor 5 mg Filmtabletten	not available	2008089995	MERCK HEALTHCARE GERMANY GMBH	LU
Concor 10 mg Filmtabletten	not available	2008089996	MERCK HEALTHCARE GERMANY GMBH	LU
Bisoprolol Zentiva 10 mg tabletes	DE/H/6433/006	21-0066	ZENTIVA, K.S.	LV
Bisoprolol Zentiva 10 mg tabletes	DE/H/6433/006	21-0066	ZENTIVA, K.S.	LV
Bisoprolol Zentiva 10 mg tabletes	DE/H/6433/006	21-0066	ZENTIVA, K.S.	LV
Bisoprolol Zentiva 10 mg tabletes	DE/H/6433/006	21-0066	ZENTIVA, K.S.	LV
Bisoprolol Zentiva 10 mg tabletes	DE/H/6433/006	21-0066	ZENTIVA, K.S.	LV
Bisoprolol Zentiva 10 mg	DE/H/6433/006	21-0066	ZENTIVA, K.S.	LV

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
tabletes				
Bisoprolol Zentiva 10 mg tabletes	DE/H/6433/006	21-0066	ZENTIVA, K.S.	LV
Bisoprolol Zentiva 10 mg tabletes	DE/H/6433/006	21-0066	ZENTIVA, K.S.	LV
Bisoprolol Zentiva 10 mg tabletes	DE/H/6433/006	21-0066	ZENTIVA, K.S.	LV
Bisoprolol Zentiva 10 mg tabletes	DE/H/6433/006	21-0066	ZENTIVA, K.S.	LV
Bisoprolol Zentiva 10 mg tabletes	DE/H/6433/006	21-0066	ZENTIVA, K.S.	LV
Bisoprolol Zentiva 10 mg tabletes	DE/H/6433/006	21-0066	ZENTIVA, K.S.	LV
Bisoprolol Zentiva 10 mg tabletes	DE/H/6433/006	21-0066	ZENTIVA, K.S.	LV
Bisoprolol Zentiva 10 mg tabletes	DE/H/6433/006	21-0066	ZENTIVA, K.S.	LV
Concor COR 2,5 mg apvalkotās tabletes	not available	02-0168	MERCK SERONO SIA	LV
Concor COR 5 mg apvalkotās tabletes	not available	02-0169	MERCK SERONO SIA	LV
Concor COR 2,5 mg apvalkotās tabletes	not available	02-0168	MERCK SERONO SIA	LV
Concor COR 5 mg apvalkotās tabletes	not available	02-0169	MERCK SERONO SIA	LV
Concor 10 mg apvalkotās tabletes	not available	99-0011	MERCK SERONO SIA	LV
Concor 5 mg apvalkotās tabletes	not available	99-0010	MERCK SERONO SIA	LV
Concor 5 mg apvalkotās tabletes	not available	99-0010	MERCK SERONO SIA	LV
Concor 5 mg apvalkotās tabletes	not available	99-0010	MERCK SERONO SIA	LV
Concor 5 mg apvalkotās tabletes	not available	99-0010	MERCK SERONO SIA	LV

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
Concor 10 mg apvalkotās tabletes	not available	99-0011	MERCK SERONO SIA	LV
Concor 10 mg apvalkotās tabletes	not available	99-0011	MERCK SERONO SIA	LV
Concor 10 mg apvalkotās tabletes	not available	99-0011	MERCK SERONO SIA	LV
Bisoprolol Syri Pharma 0,5 mg/ml, drank	NL/H/5906/001	RVG 132206	SYRI PHARMA LIMITED	NL
EMCOR DECO 7,5 mg filmomhulde tabletten	SE/H/0184/005	RVG 24506	MERCK B.V.	NL
EMCOR DECO 7,5 mg filmomhulde tabletten	SE/H/0184/005	RVG 24506	MERCK B.V.	NL
EMCOR DECO 7,5 mg filmomhulde tabletten	SE/H/0184/005	RVG 24506	MERCK B.V.	NL
EMCOR DECO 7,5 mg filmomhulde tabletten	SE/H/0184/005	RVG 24506	MERCK B.V.	NL
EMCOR DECO 7,5 mg filmomhulde tabletten	SE/H/0184/005	RVG 24506	MERCK B.V.	NL
EMCOR DECO 7,5 mg filmomhulde tabletten	SE/H/0184/005	RVG 24506	MERCK B.V.	NL
EMCOR DECO 7,5 mg filmomhulde tabletten	SE/H/0184/005	RVG 24506	MERCK B.V.	NL
EMCOR DECO 7,5 mg filmomhulde tabletten	SE/H/0184/005	RVG 24506	MERCK B.V.	NL
EMCOR DECO 7,5 mg filmomhulde tabletten	SE/H/0184/005	RVG 24506	MERCK B.V.	NL
EMCOR DECO 7,5 mg filmomhulde tabletten	SE/H/0184/005	RVG 24506	MERCK B.V.	NL
EMCOR DECO 7,5 mg filmomhulde tabletten	SE/H/0184/005	RVG 24506	MERCK B.V.	NL
EMCOR DECO 7,5 mg filmomhulde tabletten	SE/H/0184/005	RVG 24506	MERCK B.V.	NL
EMCOR DECO 7,5 mg filmomhulde tabletten	SE/H/0184/005	RVG 24506	MERCK B.V.	NL
EMCOR DECO 7,5 mg	SE/H/0184/005	RVG 24506	MERCK B.V.	NL

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
filmomhulde tabletten				
EMCOR DECO 7,5 mg filmomhulde tabletten	SE/H/0184/005	RVG 24506	MERCK B.V.	NL
EMCOR DECO 7,5 mg filmomhulde tabletten	SE/H/0184/005	RVG 24506	MERCK B.V.	NL
EMCOR DECO 2,5 mg filmomhulde tabletten	SE/H/0184/002	RVG 24503	MERCK B.V.	NL
EMCOR DECO 2,5 mg filmomhulde tabletten	SE/H/0184/002	RVG 24503	MERCK B.V.	NL
EMCOR DECO 2,5 mg filmomhulde tabletten	SE/H/0184/002	RVG 24503	MERCK B.V.	NL
EMCOR DECO 2,5 mg filmomhulde tabletten	SE/H/0184/002	RVG 24503	MERCK B.V.	NL
EMCOR DECO 2,5 mg filmomhulde tabletten	SE/H/0184/002	RVG 24503	MERCK B.V.	NL
EMCOR DECO 2,5 mg filmomhulde tabletten	SE/H/0184/002	RVG 24503	MERCK B.V.	NL
EMCOR DECO 2,5 mg filmomhulde tabletten	SE/H/0184/002	RVG 24503	MERCK B.V.	NL
EMCOR DECO 2,5 mg filmomhulde tabletten	SE/H/0184/002	RVG 24503	MERCK B.V.	NL
EMCOR DECO 2,5 mg filmomhulde tabletten	SE/H/0184/002	RVG 24503	MERCK B.V.	NL
EMCOR DECO 2,5 mg filmomhulde tabletten	SE/H/0184/002	RVG 24503	MERCK B.V.	NL
EMCOR DECO 2,5 mg filmomhulde tabletten	SE/H/0184/002	RVG 24503	MERCK B.V.	NL
EMCOR DECO 2,5 mg filmomhulde tabletten	SE/H/0184/002	RVG 24503	MERCK B.V.	NL
EMCOR DECO 2,5 mg filmomhulde tabletten	SE/H/0184/002	RVG 24503	MERCK B.V.	NL
EMCOR DECO 2,5 mg filmomhulde tabletten	SE/H/0184/002	RVG 24503	MERCK B.V.	NL
EMCOR DECO 2,5 mg filmomhulde tabletten	SE/H/0184/002	RVG 24503	MERCK B.V.	NL

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
EMCOR DECO 2,5 mg filmomhulde tabletten	SE/H/0184/002	RVG 24503	MERCK B.V.	NL
Bisoprolol Zentiva 5 mg tabletter	DE/H/6433/004	22-15047	ZENTIVA, K.S.	NO
Bisoprolol Zentiva 10 mg tabletter	DE/H/6433/006	22-15048	ZENTIVA, K.S.	NO
Bisoprolol Zentiva 5 mg tabletter	DE/H/6433/004	22-15047	ZENTIVA, K.S.	NO
Bisoprolol Zentiva 5 mg tabletter	DE/H/6433/004	22-15047	ZENTIVA, K.S.	NO
Bisoprolol Zentiva 5 mg tabletter	DE/H/6433/004	22-15047	ZENTIVA, K.S.	NO
Bisoprolol Zentiva 5 mg tabletter	DE/H/6433/004	22-15047	ZENTIVA, K.S.	NO
Bisoprolol Zentiva 5 mg tabletter	DE/H/6433/004	22-15047	ZENTIVA, K.S.	NO
Bisoprolol Zentiva 5 mg tabletter	DE/H/6433/004	22-15047	ZENTIVA, K.S.	NO
Bisoprolol Zentiva 5 mg tabletter	DE/H/6433/004	22-15047	ZENTIVA, K.S.	NO
Bisoprolol Zentiva 5 mg tabletter	DE/H/6433/004	22-15047	ZENTIVA, K.S.	NO
Bisoprolol Zentiva 5 mg tabletter	DE/H/6433/004	22-15047	ZENTIVA, K.S.	NO
Bisoprolol Zentiva 5 mg tabletter	DE/H/6433/004	22-15047	ZENTIVA, K.S.	NO
Bisoprolol Zentiva 5 mg tabletter	DE/H/6433/004	22-15047	ZENTIVA, K.S.	NO
Bisoprolol Zentiva 5 mg tabletter	DE/H/6433/004	22-15047	ZENTIVA, K.S.	NO
Bisoprolol Zentiva 5 mg tabletter	DE/H/6433/004	22-15047	ZENTIVA, K.S.	NO
Bisoprolol Zentiva 10 mg tabletter	DE/H/6433/006	22-15048	ZENTIVA, K.S.	NO
Bisoprolol Zentiva 10 mg	DE/H/6433/006	22-15048	ZENTIVA, K.S.	NO

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
tabletter				
Bisoprolol Zentiva 10 mg tabletter	DE/H/6433/006	22-15048	ZENTIVA, K.S.	NO
Bisoprolol Zentiva 10 mg tabletter	DE/H/6433/006	22-15048	ZENTIVA, K.S.	NO
Bisoprolol Zentiva 10 mg tabletter	DE/H/6433/006	22-15048	ZENTIVA, K.S.	NO
Bisoprolol Zentiva 10 mg tabletter	DE/H/6433/006	22-15048	ZENTIVA, K.S.	NO
Bisoprolol Zentiva 10 mg tabletter	DE/H/6433/006	22-15048	ZENTIVA, K.S.	NO
Bisoprolol Zentiva 10 mg tabletter	DE/H/6433/006	22-15048	ZENTIVA, K.S.	NO
Bisoprolol Zentiva 10 mg tabletter	DE/H/6433/006	22-15048	ZENTIVA, K.S.	NO
Bisoprolol Zentiva 10 mg tabletter	DE/H/6433/006	22-15048	ZENTIVA, K.S.	NO
Bisoprolol Zentiva 10 mg tabletter	DE/H/6433/006	22-15048	ZENTIVA, K.S.	NO
Bisoprolol Zentiva 10 mg tabletter	DE/H/6433/006	22-15048	ZENTIVA, K.S.	NO
Bisoprolol Zentiva 10 mg tabletter	DE/H/6433/006	22-15048	ZENTIVA, K.S.	NO
Emconcor 5 mg tabletter, filmdrasjerte	not available	02-1326	MERCK AB	NO
Emconcor 5 mg tabletter, filmdrasjerte	not available	02-1326	MERCK AB	NO
Emconcor CHF 1,25 mg tablett, filmdrasjert	not available	99-7737	MERCK AB	NO
Emconcor CHF 1,25 mg tablett, filmdrasjert	not available	99-7737	MERCK AB	NO
Emconcor CHF 1,25 mg tablett, filmdrasjert	not available	99-7737	MERCK AB	NO
Emconcor CHF 1,25 mg tablett, filmdrasjert	not available	99-7737	MERCK AB	NO

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
Emconcor CHF 1,25 mg tablett, filmdrasjert	not available	99-7737	MERCK AB	NO
Emconcor CHF 1,25 mg tablett, filmdrasjert	not available	99-7737	MERCK AB	NO
Emconcor CHF 1,25 mg tablett, filmdrasjert	not available	99-7737	MERCK AB	NO
Emconcor CHF 2,5 mg tablett, filmdrasjert	not available	99-7738	MERCK AB	NO
Emconcor CHF 2,5 mg tablett, filmdrasjert	not available	99-7738	MERCK AB	NO
Emconcor CHF 2,5 mg tablett, filmdrasjert	not available	99-7738	MERCK AB	NO
Emconcor CHF 2,5 mg tablett, filmdrasjert	not available	99-7738	MERCK AB	NO
Emconcor CHF 2,5 mg tablett, filmdrasjert	not available	99-7738	MERCK AB	NO
Emconcor CHF 2,5 mg tablett, filmdrasjert	not available	99-7738	MERCK AB	NO
Emconcor CHF 2,5 mg tablett, filmdrasjert	not available	99-7738	MERCK AB	NO
Emconcor CHF 3,75 mg tablett, filmdrasjert	not available	99-7739	MERCK AB	NO
Emconcor CHF 3,75 mg tablett, filmdrasjert	not available	99-7739	MERCK AB	NO
Emconcor CHF 3,75 mg tablett, filmdrasjert	not available	99-7739	MERCK AB	NO
Emconcor CHF 3,75 mg tablett, filmdrasjert	not available	99-7739	MERCK AB	NO
Emconcor CHF 3,75 mg tablett, filmdrasjert	not available	99-7739	MERCK AB	NO
Emconcor CHF 3,75 mg tablett, filmdrasjert	not available	99-7739	MERCK AB	NO
Emconcor CHF 3,75 mg tablett, filmdrasjert	not available	99-7739	MERCK AB	NO
Emconcor CHF 5 mg	not available	99-7740	MERCK AB	NO

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
tablett, filmdrasjert				
Emconcor CHF 5 mg tablett, filmdrasjert	not available	99-7740	MERCK AB	NO
Emconcor CHF 5 mg tablett, filmdrasjert	not available	99-7740	MERCK AB	NO
Emconcor CHF 5 mg tablett, filmdrasjert	not available	99-7740	MERCK AB	NO
Emconcor CHF 5 mg tablett, filmdrasjert	not available	99-7740	MERCK AB	NO
Emconcor CHF 5 mg tablett, filmdrasjert	not available	99-7740	MERCK AB	NO
Emconcor CHF 5 mg tablett, filmdrasjert	not available	99-7740	MERCK AB	NO
Emconcor CHF 10 mg tablett, filmdrasjert	not available	99-7742	MERCK AB	NO
Emconcor CHF 10 mg tablett, filmdrasjert	not available	99-7742	MERCK AB	NO
Emconcor CHF 10 mg tablett, filmdrasjert	not available	99-7742	MERCK AB	NO
Emconcor CHF 10 mg tablett, filmdrasjert	not available	99-7742	MERCK AB	NO
Emconcor CHF 10 mg tablett, filmdrasjert	not available	99-7742	MERCK AB	NO
Emconcor CHF 10 mg tablett, filmdrasjert	not available	99-7742	MERCK AB	NO
Emconcor CHF 10 mg tablett, filmdrasjert	not available	99-7742	MERCK AB	NO
Emconcor CHF 1,25 mg tablett, filmdrasjert	not available	99-7737	MERCK AB	NO
Emconcor CHF 10 mg tablett, filmdrasjert	not available	99-7742	MERCK AB	NO
Emconcor CHF 2,5 mg tablett, filmdrasjert	not available	99-7738	MERCK AB	NO
Emconcor CHF 3,75 mg tablett, filmdrasjert	not available	99-7739	MERCK AB	NO

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
Emconcor CHF 5 mg tablett, filmdrasjert	not available	99-7740	MERCK AB	NO
Concor 5, 5 mg tabletki powlekane	not available	R/3734	MERCK SP.Z O.O.	PL
Concor 5, 5 mg tabletki powlekane	not available	R/3734	MERCK SP.Z O.O.	PL
Concor 10, 10 mg tabletki powlekane	not available	R/3735	MERCK SP.Z O.O.	PL
Concor 10, 10 mg tabletki powlekane	not available	R/3735	MERCK SP.Z O.O.	PL
Concor 10, 10 mg tabletki powlekane	not available	R/3735	MERCK SP.Z O.O.	PL
Concor 5, 5 mg tabletki powlekane	not available	R/3734	MERCK SP.Z O.O.	PL
Bisopromerck 5, 5 mg tabletki powlekane	not available	4894	MERCK SP.Z O.O.	PL
Bisopromerck 10, 10 mg tabletki powlekane	not available	4895	MERCK SP.Z O.O.	PL
Bisopromerck 10, 10 mg tabletki powlekane	not available	4895	MERCK SP.Z O.O.	PL
Bisopromerck 5, 5 mg tabletki powlekane	not available	4894	MERCK SP.Z O.O.	PL
Concor Cor 3,75, 3,75 mg tabletki powlekane	not available	8591	MERCK SP.Z O.O.	PL
Concor Cor 3,75, 3,75 mg tabletki powlekane	not available	8591	MERCK SP.Z O.O.	PL
Concor Cor 2,5, 2,5 mg tabletki powlekane	not available	8590	MERCK SP.Z O.O.	PL
Concor Cor 2,5, 2,5 mg tabletki powlekane	not available	8590	MERCK SP.Z O.O.	PL
Concor Cor 7,5, 7,5 mg tabletki powlekane	not available	8593	MERCK SP.Z O.O.	PL
Concor Cor 7,5, 7,5 mg tabletki powlekane	not available	8593	MERCK SP.Z O.O.	PL
Concor Cor 1,25, 1,25	not available	8589	MERCK SP.Z O.O.	PL

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 tabletki powlekane				
Concor Cor 1,25, 1,25 mg tabletki powlekane	not available	8589	MERCK SP.Z O.O.	PL
Concor Cor 5, 5 mg tabletki powlekane	not available	8592	MERCK SP.Z O.O.	PL
Concor Cor 5, 5 mg tabletki powlekane	not available	8592	MERCK SP.Z O.O.	PL
Concor Cor 10, 10 mg tabletki powlekane	not available	8594	MERCK SP.Z O.O.	PL
Concor Cor 10, 10 mg tabletki powlekane	not available	8594	MERCK SP.Z O.O.	PL
Concor Cor 1,25, 1,25 mg tabletki powlekane	not available	8589	MERCK SP.Z O.O.	PL
Concor Cor 10, 10 mg tabletki powlekane	not available	8594	MERCK SP.Z O.O.	PL
Concor Cor 2,5, 2,5 mg tabletki powlekane	not available	8590	MERCK SP.Z O.O.	PL
Concor Cor 3,75, 3,75 mg tabletki powlekane	not available	8591	MERCK SP.Z O.O.	PL
Concor Cor 5, 5 mg tabletki powlekane	not available	8592	MERCK SP.Z O.O.	PL
Concor Cor 7,5, 7,5 mg tabletki powlekane	not available	8593	MERCK SP.Z O.O.	PL
Bisoprolol Zentiva 5 mg comprimidos	DE/H/6433/004	5817101	ZENTIVA PORTUGAL, LDA	PT
Bisoprolol Zentiva 5 mg comprimidos	DE/H/6433/004	5817077	ZENTIVA PORTUGAL, LDA	PT
Bisoprolol Zentiva 10 mg comprimidos	DE/H/6433/006	5817119	ZENTIVA PORTUGAL, LDA	PT
Bisoprolol Zentiva 10 mg comprimidos	DE/H/6433/006	5817127	ZENTIVA PORTUGAL, LDA	PT
Bisoprolol Aquizent 5 mg comprimidos	not available	5906631	ZENTIVA, K.S.	PT
Bisoprolol Aquizent 10 mg comprimidos	not available	5906649	ZENTIVA, K.S.	PT

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
Concor 10 mg comprimido revestido	not available	5869532	MERCK, S.A.	PT
Concor 5 mg comprimido revestido	not available	5869540	MERCK, S.A.	PT
Concor 10 mg comprimido revestido	not available	8776468	MERCK, S.A.	PT
Concor 10 mg comprimido revestido	not available	8776450	MERCK, S.A.	PT
Concor 5 mg comprimido revestido	not available	8776476	MERCK, S.A.	PT
Concor 5 mg comprimido revestido	not available	8776435	MERCK, S.A.	PT
Concor IC 2,5 mg comprimidos revestidos por película	SE/H/0184/002	SE/H/0184/002	MERCK, S.A.	PT
Concor IC 2,5 mg comprimidos revestidos por película	SE/H/0184/002	SE/H/0184/002	MERCK, S.A.	PT
Concor IC 2,5 mg comprimidos revestidos por película	SE/H/0184/002	3031085	MERCK, S.A.	PT
Concor IC 2,5 mg comprimidos revestidos por película	SE/H/0184/002	SE/H/0184/002	MERCK, S.A.	PT
Concor IC 2,5 mg comprimidos revestidos por película	SE/H/0184/002	SE/H/0184/002	MERCK, S.A.	PT
Concor IC 2,5 mg comprimidos revestidos por película	SE/H/0184/002	3034980	MERCK, S.A.	PT
Concor IC 2,5 mg comprimidos revestidos por película	SE/H/0184/002	3030780	MERCK, S.A.	PT
Concor IC 2,5 mg comprimidos revestidos	SE/H/0184/002	3030988	MERCK, S.A.	PT

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
por película				
Concor IC 2,5 mg comprimidos revestidos por película	SE/H/0184/002	3030889	MERCK, S.A.	PT
Concor IC 2,5 mg comprimidos revestidos por película	SE/H/0184/002	SE/H/0184/002	MERCK, S.A.	PT
Concor IC 2,5 mg comprimidos revestidos por película	SE/H/0184/002	SE/H/0184/002	MERCK, S.A.	PT
Concor IC 2,5 mg comprimidos revestidos por película	SE/H/0184/002	SE/H/0184/002	MERCK, S.A.	PT
Concor IC 2,5 mg comprimidos revestidos por película	SE/H/0184/002	SE/H/0184/002	MERCK, S.A.	PT
Concor IC 2,5 mg comprimidos revestidos por película	SE/H/0184/002	3031382	MERCK, S.A.	PT
Concor IC 2,5 mg comprimidos revestidos por película	SE/H/0184/002	3031184	MERCK, S.A.	PT
Concor IC 2,5 mg comprimidos revestidos por película	SE/H/0184/002	3031283	MERCK, S.A.	PT
CONCOR COR 2,5 mg comprimate filmate	not available	6094/2014/05	MERCK ROMANIA S.R.L.	RO
CONCOR COR 10 mg comprimate filmate	not available	6096/2014/03	MERCK ROMANIA S.R.L.	RO
CONCOR COR 10 mg comprimate filmate	not available	6096/2014/04	MERCK ROMANIA S.R.L.	RO
CONCOR COR 10 mg comprimate filmate	not available	6096/2014/01	MERCK ROMANIA S.R.L.	RO
CONCOR COR 10 mg comprimate filmate	not available	6096/2014/02	MERCK ROMANIA S.R.L.	RO

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
CONCOR COR 2,5 mg comprimate filmate	not available	6094/2014/03	MERCK ROMANIA S.R.L.	RO
CONCOR COR 2,5 mg comprimate filmate	not available	6094/2014/02	MERCK ROMANIA S.R.L.	RO
CONCOR COR 2,5 mg comprimate filmate	not available	6094/2014/01	MERCK ROMANIA S.R.L.	RO
CONCOR COR 2,5 mg comprimate filmate	not available	6094/2014/04	MERCK ROMANIA S.R.L.	RO
CONCOR COR 5 mg comprimate filmate	not available	6095/2014/01	MERCK ROMANIA S.R.L.	RO
CONCOR COR 5 mg comprimate filmate	not available	6095/2014/04	MERCK ROMANIA S.R.L.	RO
CONCOR COR 5 mg comprimate filmate	not available	6095/2014/03	MERCK ROMANIA S.R.L.	RO
CONCOR COR 5 mg comprimate filmate	not available	6095/2014/02	MERCK ROMANIA S.R.L.	RO
Concor 10 mg comprimate filmate	not available	6252/2014/01	MERCK ROMANIA S.R.L.	RO
Concor 5 mg comprimate filmate	not available	6251/2014/01	MERCK ROMANIA S.R.L.	RO
Concor 5 mg comprimate filmate	not available	6251/2014/02	MERCK ROMANIA S.R.L.	RO
Concor 10 mg comprimate filmate	not available	6252/2014/02	MERCK ROMANIA S.R.L.	RO
Bisoprolol Zentiva 5 mg tabletter	DE/H/6433/004	63962	ZENTIVA, K.S.	SE
Bisoprolol Zentiva 10 mg tabletter	DE/H/6433/006	63963	ZENTIVA, K.S.	SE
Bisoprolol Zentiva 5 mg tabletter	DE/H/6433/004	63962	ZENTIVA, K.S.	SE
Bisoprolol Zentiva 5 mg tabletter	DE/H/6433/004	63962	ZENTIVA, K.S.	SE
Bisoprolol Zentiva 5 mg tabletter	DE/H/6433/004	63962	ZENTIVA, K.S.	SE
Bisoprolol Zentiva 5 mg	DE/H/6433/004	63962	ZENTIVA, K.S.	SE

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
tabletter				
Bisoprolol Zentiva 5 mg tabletter	DE/H/6433/004	63962	ZENTIVA, K.S.	SE
Bisoprolol Zentiva 5 mg tabletter	DE/H/6433/004	63962	ZENTIVA, K.S.	SE
Bisoprolol Zentiva 5 mg tabletter	DE/H/6433/004	63962	ZENTIVA, K.S.	SE
Bisoprolol Zentiva 5 mg tabletter	DE/H/6433/004	63962	ZENTIVA, K.S.	SE
Bisoprolol Zentiva 5 mg tabletter	DE/H/6433/004	63962	ZENTIVA, K.S.	SE
Bisoprolol Zentiva 5 mg tabletter	DE/H/6433/004	63962	ZENTIVA, K.S.	SE
Bisoprolol Zentiva 5 mg tabletter	DE/H/6433/004	63962	ZENTIVA, K.S.	SE
Bisoprolol Zentiva 5 mg tabletter	DE/H/6433/004	63962	ZENTIVA, K.S.	SE
Bisoprolol Zentiva 5 mg tabletter	DE/H/6433/004	63962	ZENTIVA, K.S.	SE
Bisoprolol Zentiva 10 mg tabletter	DE/H/6433/006	63963	ZENTIVA, K.S.	SE
Bisoprolol Zentiva 10 mg tabletter	DE/H/6433/006	63963	ZENTIVA, K.S.	SE
Bisoprolol Zentiva 10 mg tabletter	DE/H/6433/006	63963	ZENTIVA, K.S.	SE
Bisoprolol Zentiva 10 mg tabletter	DE/H/6433/006	63963	ZENTIVA, K.S.	SE
Bisoprolol Zentiva 10 mg tabletter	DE/H/6433/006	63963	ZENTIVA, K.S.	SE
Bisoprolol Zentiva 10 mg tabletter	DE/H/6433/006	63963	ZENTIVA, K.S.	SE
Bisoprolol Zentiva 10 mg tabletter	DE/H/6433/006	63963	ZENTIVA, K.S.	SE
Bisoprolol Zentiva 10 mg tabletter	DE/H/6433/006	63963	ZENTIVA, K.S.	SE

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
Bisoprolol Zentiva 10 mg tabletter	DE/H/6433/006	63963	ZENTIVA, K.S.	SE
Bisoprolol Zentiva 10 mg tabletter	DE/H/6433/006	63963	ZENTIVA, K.S.	SE
Bisoprolol Zentiva 10 mg tabletter	DE/H/6433/006	63963	ZENTIVA, K.S.	SE
Bisoprolol Zentiva 10 mg tabletter	DE/H/6433/006	63963	ZENTIVA, K.S.	SE
Bisoprolol Zentiva 10 mg tabletter	DE/H/6433/006	63963	ZENTIVA, K.S.	SE
Emconcor 5 mg filmdragerade tabletter	not available	11202	MERCK AB	SE
Emconcor 5 mg filmdragerade tabletter	not available	11202	MERCK AB	SE
Emconcor 5 mg filmdragerade tabletter	not available	11202	MERCK AB	SE
Emconcor CHF 1,25 mg filmdragerade tabletter	SE/H/0184/001	15371	MERCK AB	SE
Emconcor CHF 1,25 mg filmdragerade tabletter	SE/H/0184/001	15371	MERCK AB	SE
Emconcor CHF 1,25 mg filmdragerade tabletter	SE/H/0184/001	15371	MERCK AB	SE
Emconcor CHF 1,25 mg filmdragerade tabletter	SE/H/0184/001	15371	MERCK AB	SE
Emconcor CHF 1,25 mg filmdragerade tabletter	SE/H/0184/001	15371	MERCK AB	SE
Emconcor CHF 1,25 mg filmdragerade tabletter	SE/H/0184/001	15371	MERCK AB	SE
Emconcor CHF 1,25 mg filmdragerade tabletter	SE/H/0184/001	15371	MERCK AB	SE
Emconcor CHF 1,25 mg filmdragerade tabletter	SE/H/0184/001	15371	MERCK AB	SE
Emconcor CHF 1,25 mg filmdragerade tabletter	SE/H/0184/001	15371	MERCK AB	SE
Emconcor CHF 1,25 mg	SE/H/0184/001	15371	MERCK AB	SE

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
filmdragerade tabletter				
Emconcor CHF 1,25 mg filmdragerade tabletter	SE/H/0184/001	15371	MERCK AB	SE
Emconcor CHF 1,25 mg filmdragerade tabletter	SE/H/0184/001	15371	MERCK AB	SE
Emconcor CHF 1,25 mg filmdragerade tabletter	SE/H/0184/001	15371	MERCK AB	SE
Emconcor CHF 1,25 mg filmdragerade tabletter	SE/H/0184/001	15371	MERCK AB	SE
Emconcor CHF 1,25 mg filmdragerade tabletter	SE/H/0184/001	15371	MERCK AB	SE
Emconcor CHF 1,25 mg filmdragerade tabletter	SE/H/0184/001	15371	MERCK AB	SE
Emconcor CHF 10 mg filmdragerade tabletter	SE/H/0184/006	15231	MERCK AB	SE
Emconcor CHF 10 mg filmdragerade tabletter	SE/H/0184/006	15231	MERCK AB	SE
Emconcor CHF 10 mg filmdragerade tabletter	SE/H/0184/006	15231	MERCK AB	SE
Emconcor CHF 10 mg filmdragerade tabletter	SE/H/0184/006	15231	MERCK AB	SE
Emconcor CHF 10 mg filmdragerade tabletter	SE/H/0184/006	15231	MERCK AB	SE
Emconcor CHF 10 mg filmdragerade tabletter	SE/H/0184/006	15231	MERCK AB	SE
Emconcor CHF 10 mg filmdragerade tabletter	SE/H/0184/006	15231	MERCK AB	SE
Emconcor CHF 10 mg filmdragerade tabletter	SE/H/0184/006	15231	MERCK AB	SE
Emconcor CHF 10 mg filmdragerade tabletter	SE/H/0184/006	15231	MERCK AB	SE
Emconcor CHF 10 mg filmdragerade tabletter	SE/H/0184/006	15231	MERCK AB	SE
Emconcor CHF 10 mg filmdragerade tabletter	SE/H/0184/006	15231	MERCK AB	SE

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
Emconcor CHF 10 mg filmdragerade tabletter	SE/H/0184/006	15231	MERCK AB	SE
Emconcor CHF 10 mg filmdragerade tabletter	SE/H/0184/006	15231	MERCK AB	SE
Emconcor CHF 10 mg filmdragerade tabletter	SE/H/0184/006	15231	MERCK AB	SE
Emconcor CHF 10 mg filmdragerade tabletter	SE/H/0184/006	15231	MERCK AB	SE
Emconcor CHF 10 mg filmdragerade tabletter	SE/H/0184/006	15231	MERCK AB	SE
Emconcor CHF 2,5 mg filmdragerade tabletter	SE/H/0184/002	15229	MERCK AB	SE
Emconcor CHF 2,5 mg filmdragerade tabletter	SE/H/0184/002	15229	MERCK AB	SE
Emconcor CHF 2,5 mg filmdragerade tabletter	SE/H/0184/002	15229	MERCK AB	SE
Emconcor CHF 2,5 mg filmdragerade tabletter	SE/H/0184/002	15229	MERCK AB	SE
Emconcor CHF 2,5 mg filmdragerade tabletter	SE/H/0184/002	15229	MERCK AB	SE
Emconcor CHF 2,5 mg filmdragerade tabletter	SE/H/0184/002	15229	MERCK AB	SE
Emconcor CHF 2,5 mg filmdragerade tabletter	SE/H/0184/002	15229	MERCK AB	SE
Emconcor CHF 2,5 mg filmdragerade tabletter	SE/H/0184/002	15229	MERCK AB	SE
Emconcor CHF 2,5 mg filmdragerade tabletter	SE/H/0184/002	15229	MERCK AB	SE
Emconcor CHF 2,5 mg filmdragerade tabletter	SE/H/0184/002	15229	MERCK AB	SE
Emconcor CHF 2,5 mg filmdragerade tabletter	SE/H/0184/002	15229	MERCK AB	SE
Emconcor CHF 2,5 mg filmdragerade tabletter	SE/H/0184/002	15229	MERCK AB	SE
Emconcor CHF 2,5 mg	SE/H/0184/002	15229	MERCK AB	SE

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
filmdragerade tabletter				
Emconcor CHF 2,5 mg filmdragerade tabletter	SE/H/0184/002	15229	MERCK AB	SE
Emconcor CHF 2,5 mg filmdragerade tabletter	SE/H/0184/002	15229	MERCK AB	SE
Emconcor CHF 2,5 mg filmdragerade tabletter	SE/H/0184/002	15229	MERCK AB	SE
Emconcor CHF 3,75 mg filmdragerade tabletter	SE/H/0184/003	15372	MERCK AB	SE
Emconcor CHF 3,75 mg filmdragerade tabletter	SE/H/0184/003	15372	MERCK AB	SE
Emconcor CHF 3,75 mg filmdragerade tabletter	SE/H/0184/003	15372	MERCK AB	SE
Emconcor CHF 3,75 mg filmdragerade tabletter	SE/H/0184/003	15372	MERCK AB	SE
Emconcor CHF 3,75 mg filmdragerade tabletter	SE/H/0184/003	15372	MERCK AB	SE
Emconcor CHF 3,75 mg filmdragerade tabletter	SE/H/0184/003	15372	MERCK AB	SE
Emconcor CHF 3,75 mg filmdragerade tabletter	SE/H/0184/003	15372	MERCK AB	SE
Emconcor CHF 3,75 mg filmdragerade tabletter	SE/H/0184/003	15372	MERCK AB	SE
Emconcor CHF 3,75 mg filmdragerade tabletter	SE/H/0184/003	15372	MERCK AB	SE
Emconcor CHF 3,75 mg filmdragerade tabletter	SE/H/0184/003	15372	MERCK AB	SE
Emconcor CHF 3,75 mg filmdragerade tabletter	SE/H/0184/003	15372	MERCK AB	SE
Emconcor CHF 3,75 mg filmdragerade tabletter	SE/H/0184/003	15372	MERCK AB	SE
Emconcor CHF 3,75 mg filmdragerade tabletter	SE/H/0184/003	15372	MERCK AB	SE
Emconcor CHF 3,75 mg filmdragerade tabletter	SE/H/0184/003	15372	MERCK AB	SE

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**
Emconcor CHF 3,75 mg filmdragerade tabletter	SE/H/0184/003	15372	MERCK AB	SE
Emconcor CHF 3,75 mg filmdragerade tabletter	SE/H/0184/003	15372	MERCK AB	SE
Emconcor CHF 5 mg filmdragerade tabletter	SE/H/0184/004	15373	MERCK AB	SE
Emconcor CHF 5 mg filmdragerade tabletter	SE/H/0184/004	15373	MERCK AB	SE
Emconcor CHF 5 mg filmdragerade tabletter	SE/H/0184/004	15373	MERCK AB	SE
Emconcor CHF 5 mg filmdragerade tabletter	SE/H/0184/004	15373	MERCK AB	SE
Emconcor CHF 5 mg filmdragerade tabletter	SE/H/0184/004	15373	MERCK AB	SE
Emconcor CHF 5 mg filmdragerade tabletter	SE/H/0184/004	15373	MERCK AB	SE
Emconcor CHF 5 mg filmdragerade tabletter	SE/H/0184/004	15373	MERCK AB	SE
Emconcor CHF 5 mg filmdragerade tabletter	SE/H/0184/004	15373	MERCK AB	SE
Emconcor CHF 5 mg filmdragerade tabletter	SE/H/0184/004	15373	MERCK AB	SE
Emconcor CHF 5 mg filmdragerade tabletter	SE/H/0184/004	15373	MERCK AB	SE
Emconcor CHF 5 mg filmdragerade tabletter	SE/H/0184/004	15373	MERCK AB	SE
Emconcor CHF 5 mg filmdragerade tabletter	SE/H/0184/004	15373	MERCK AB	SE
Emconcor CHF 5 mg filmdragerade tabletter	SE/H/0184/004	15373	MERCK AB	SE
Emconcor CHF 5 mg filmdragerade tabletter	SE/H/0184/004	15373	MERCK AB	SE
Emconcor CHF 5 mg filmdragerade tabletter	SE/H/0184/004	15373	MERCK AB	SE
Emconcor CHF 5 mg	SE/H/0184/004	15373	MERCK AB	SE

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
filmdragerade tabletter				
Emconcor CHF 7.5 mg film-coated tablets	SE/H/0184/005	15230	MERCK AB	SE
Emconcor CHF 7.5 mg film-coated tablets	SE/H/0184/005	15230	MERCK AB	SE
Emconcor CHF 7.5 mg film-coated tablets	SE/H/0184/005	15230	MERCK AB	SE
Emconcor CHF 7.5 mg film-coated tablets	SE/H/0184/005	15230	MERCK AB	SE
Emconcor CHF 7.5 mg film-coated tablets	SE/H/0184/005	15230	MERCK AB	SE
Emconcor CHF 7.5 mg film-coated tablets	SE/H/0184/005	15230	MERCK AB	SE
Emconcor CHF 7.5 mg film-coated tablets	SE/H/0184/005	15230	MERCK AB	SE
Emconcor CHF 7.5 mg film-coated tablets	SE/H/0184/005	15230	MERCK AB	SE
Emconcor CHF 7.5 mg film-coated tablets	SE/H/0184/005	15230	MERCK AB	SE
Emconcor CHF 7.5 mg film-coated tablets	SE/H/0184/005	15230	MERCK AB	SE
Emconcor CHF 7.5 mg film-coated tablets	SE/H/0184/005	15230	MERCK AB	SE
Emconcor CHF 7.5 mg film-coated tablets	SE/H/0184/005	15230	MERCK AB	SE
Emconcor CHF 7.5 mg film-coated tablets	SE/H/0184/005	15230	MERCK AB	SE
Emconcor CHF 7.5 mg film-coated tablets	SE/H/0184/005	15230	MERCK AB	SE
Emconcor CHF 7.5 mg film-coated tablets	SE/H/0184/005	15230	MERCK AB	SE
Emconcor CHF 7.5 mg film-coated tablets	SE/H/0184/005	15230	MERCK AB	SE
Cardicor 2.5 mg film-coated tablets	SE/H/0185/002	15232	MERCK AB	SE

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
Cardicor 2.5 mg film-coated tablets	SE/H/0185/002	15232	MERCK AB	SE
Cardicor 2.5 mg film-coated tablets	SE/H/0185/002	15232	MERCK AB	SE
Cardicor 2.5 mg film-coated tablets	SE/H/0185/002	15232	MERCK AB	SE
Cardicor 2.5 mg film-coated tablets	SE/H/0185/002	15232	MERCK AB	SE
Cardicor 2.5 mg film-coated tablets	SE/H/0185/002	15232	MERCK AB	SE
Cardicor 2.5 mg film-coated tablets	SE/H/0185/002	15232	MERCK AB	SE
Cardicor 2.5 mg film-coated tablets	SE/H/0185/002	15232	MERCK AB	SE
Cardicor 1.25 mg film-coated tablets	SE/H/0185/001	15374	MERCK AB	SE
Cardicor 1.25 mg film-coated tablets	SE/H/0185/001	15374	MERCK AB	SE
Cardicor 1.25 mg film-coated tablets	SE/H/0185/001	15374	MERCK AB	SE
Cardicor 1.25 mg film-coated tablets	SE/H/0185/001	15374	MERCK AB	SE
Cardicor 1.25 mg film-coated tablets	SE/H/0185/001	15374	MERCK AB	SE
Cardicor 1.25 mg film-coated tablets	SE/H/0185/001	15374	MERCK AB	SE
Cardicor 1.25 mg film-coated tablets	SE/H/0185/001	15374	MERCK AB	SE
Cardicor 1.25 mg film-coated tablets	SE/H/0185/001	15374	MERCK AB	SE
Cardicor 3.75 mg film-coated tablets	SE/H/0185/003	15375	MERCK AB	SE
Cardicor 3.75 mg film-coated tablets	SE/H/0185/003	15375	MERCK AB	SE
Cardicor 3.75 mg film-	SE/H/0185/003	15375	MERCK AB	SE

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
coated tablets				
Cardicor 3.75 mg film-coated tablets	SE/H/0185/003	15375	MERCK AB	SE
Cardicor 3.75 mg film-coated tablets	SE/H/0185/003	15375	MERCK AB	SE
Cardicor 3.75 mg film-coated tablets	SE/H/0185/003	15375	MERCK AB	SE
Cardicor 3.75 mg film-coated tablets	SE/H/0185/003	15375	MERCK AB	SE
Cardicor 3.75 mg film-coated tablets	SE/H/0185/003	15375	MERCK AB	SE
Cardicor 5 mg film-coated tablets	SE/H/0185/004	15376	MERCK AB	SE
Cardicor 5 mg film-coated tablets	SE/H/0185/004	15376	MERCK AB	SE
Cardicor 5 mg film-coated tablets	SE/H/0185/004	15376	MERCK AB	SE
Cardicor 5 mg film-coated tablets	SE/H/0185/004	15376	MERCK AB	SE
Cardicor 5 mg film-coated tablets	SE/H/0185/004	15376	MERCK AB	SE
Cardicor 5 mg film-coated tablets	SE/H/0185/004	15376	MERCK AB	SE
Cardicor 5 mg film-coated tablets	SE/H/0185/004	15376	MERCK AB	SE
Cardicor 5 mg film-coated tablets	SE/H/0185/004	15376	MERCK AB	SE
Cardicor 7.5 mg film-coated tablets	SE/H/0185/005	15233	MERCK AB	SE
Cardicor 7.5 mg film-coated tablets	SE/H/0185/005	15233	MERCK AB	SE
Cardicor 7.5 mg film-coated tablets	SE/H/0185/005	15233	MERCK AB	SE
Cardicor 7.5 mg film-coated tablets	SE/H/0185/005	15233	MERCK AB	SE

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
Cardicor 7.5 mg film-coated tablets	SE/H/0185/005	15233	MERCK AB	SE
Cardicor 7.5 mg film-coated tablets	SE/H/0185/005	15233	MERCK AB	SE
Cardicor 7.5 mg film-coated tablets	SE/H/0185/005	15233	MERCK AB	SE
Cardicor 7.5 mg film-coated tablets	SE/H/0185/005	15233	MERCK AB	SE
Cardicor 10 mg film-coated tablets	SE/H/0185/006	15234	MERCK AB	SE
Cardicor 10 mg film-coated tablets	SE/H/0185/006	15234	MERCK AB	SE
Cardicor 10 mg film-coated tablets	SE/H/0185/006	15234	MERCK AB	SE
Cardicor 10 mg film-coated tablets	SE/H/0185/006	15234	MERCK AB	SE
Cardicor 10 mg film-coated tablets	SE/H/0185/006	15234	MERCK AB	SE
Cardicor 10 mg film-coated tablets	SE/H/0185/006	15234	MERCK AB	SE
Cardicor 10 mg film-coated tablets	SE/H/0185/006	15234	MERCK AB	SE
Cardicor 2.5 mg film-coated tablets	SE/H/0185/002	15232	MERCK AB	SE
Cardicor 1.25 mg film-coated tablets	SE/H/0185/001	15374	MERCK AB	SE
Cardicor 10 mg film-coated tablets	SE/H/0185/006	15234	MERCK AB	SE
Cardicor 5 mg film-coated tablets	SE/H/0185/004	15376	MERCK AB	SE
Cardicor 3.75 mg film-coated tablets	SE/H/0185/003	15375	MERCK AB	SE
Cardicor 7.5 mg film-coated tablets	SE/H/0185/005	15233	MERCK AB	SE
Bisoprolol Merck 3.75	SE/H/0187/003	15381	MERCK AB	SE

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 film-coated tablets				
Bisoprolol Merck 3.75 mg film-coated tablets	SE/H/0187/003	15381	MERCK AB	SE
Bisoprolol Merck 3.75 mg film-coated tablets	SE/H/0187/003	15381	MERCK AB	SE
Bisoprolol Merck 3.75 mg film-coated tablets	SE/H/0187/003	15381	MERCK AB	SE
Bisoprolol Merck 3.75 mg film-coated tablets	SE/H/0187/003	15381	MERCK AB	SE
Bisoprolol Merck 3.75 mg film-coated tablets	SE/H/0187/003	15381	MERCK AB	SE
Bisoprolol Merck 3.75 mg film-coated tablets	SE/H/0187/003	15381	MERCK AB	SE
Bisoprolol Merck 5 mg film-coated tablets	SE/H/0187/004	15382	MERCK AB	SE
Bisoprolol Merck 5 mg film-coated tablets	SE/H/0187/004	15382	MERCK AB	SE
Bisoprolol Merck 5 mg film-coated tablets	SE/H/0187/004	15382	MERCK AB	SE
Bisoprolol Merck 5 mg film-coated tablets	SE/H/0187/004	15382	MERCK AB	SE
Bisoprolol Merck 5 mg film-coated tablets	SE/H/0187/004	15382	MERCK AB	SE
Bisoprolol Merck 5 mg film-coated tablets	SE/H/0187/004	15382	MERCK AB	SE
Bisoprolol Merck 5 mg film-coated tablets	SE/H/0187/004	15382	MERCK AB	SE
Bisoprolol Merck 10 mg film-coated tablets	SE/H/0187/006	15240	MERCK AB	SE
Bisoprolol Merck 10 mg film-coated tablets	SE/H/0187/006	15240	MERCK AB	SE
Bisoprolol Merck 10 mg film-coated tablets	SE/H/0187/006	15240	MERCK AB	SE
Bisoprolol Merck 10 mg film-coated tablets	SE/H/0187/006	15240	MERCK AB	SE

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
Bisoprolol Merck 10 mg film-coated tablets	SE/H/0187/006	15240	MERCK AB	SE
Bisoprolol Merck 10 mg film-coated tablets	SE/H/0187/006	15240	MERCK AB	SE
Bisoprolol Merck 10 mg film-coated tablets	SE/H/0187/006	15240	MERCK AB	SE
Bisoprolol Merck 7.5 mg film-coated tablets	SE/H/0187/005	15239	MERCK AB	SE
Bisoprolol Merck 7.5 mg film-coated tablets	SE/H/0187/005	15239	MERCK AB	SE
Bisoprolol Merck 7.5 mg film-coated tablets	SE/H/0187/005	15239	MERCK AB	SE
Bisoprolol Merck 7.5 mg film-coated tablets	SE/H/0187/005	15239	MERCK AB	SE
Bisoprolol Merck 7.5 mg film-coated tablets	SE/H/0187/005	15239	MERCK AB	SE
Bisoprolol Merck 7.5 mg film-coated tablets	SE/H/0187/005	15239	MERCK AB	SE
Bisoprolol Merck 7.5 mg film-coated tablets	SE/H/0187/005	15239	MERCK AB	SE
Bisoprolol Merck 2.5 mg film-coated tablets	SE/H/0187/002	15238	MERCK AB	SE
Bisoprolol Merck 2.5 mg film-coated tablets	SE/H/0187/002	15238	MERCK AB	SE
Bisoprolol Merck 2.5 mg film-coated tablets	SE/H/0187/002	15238	MERCK AB	SE
Bisoprolol Merck 2.5 mg film-coated tablets	SE/H/0187/002	15238	MERCK AB	SE
Bisoprolol Merck 2.5 mg film-coated tablets	SE/H/0187/002	15238	MERCK AB	SE
Bisoprolol Merck 2.5 mg film-coated tablets	SE/H/0187/002	15238	MERCK AB	SE
Bisoprolol Merck 2.5 mg film-coated tablets	SE/H/0187/002	15238	MERCK AB	SE
Bisoprolol Merck 1.25	SE/H/0187/001	15380	MERCK AB	SE

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 film-coated tablets				
Bisoprolol Merck 1.25 mg film-coated tablets	SE/H/0187/001	15380	MERCK AB	SE
Bisoprolol Merck 1.25 mg film-coated tablets	SE/H/0187/001	15380	MERCK AB	SE
Bisoprolol Merck 1.25 mg film-coated tablets	SE/H/0187/001	15380	MERCK AB	SE
Bisoprolol Merck 1.25 mg film-coated tablets	SE/H/0187/001	15380	MERCK AB	SE
Bisoprolol Merck 1.25 mg film-coated tablets	SE/H/0187/001	15380	MERCK AB	SE
Bisoprolol Merck 1.25 mg film-coated tablets	SE/H/0187/001	15380	MERCK AB	SE
Bisoprolol Merck 3.75 mg film-coated tablets	SE/H/0187/003	15381	MERCK AB	SE
Bisoprolol Merck 5 mg film-coated tablets	SE/H/0187/004	15382	MERCK AB	SE
Bisoprolol Merck 1.25 mg film-coated tablets	SE/H/0187/001	15380	MERCK AB	SE
Bisoprolol Merck 10 mg film-coated tablets	SE/H/0187/006	15240	MERCK AB	SE
Bisoprolol Merck 2.5 mg film-coated tablets	SE/H/0187/002	15238	MERCK AB	SE
Bisoprolol Merck 7.5 mg film-coated tablets	SE/H/0187/005	15239	MERCK AB	SE
Libracor 2.5 mg film-coated tablets	SE/H/0186/002	15235	MERCK AB	SE
Libracor 2.5 mg film-coated tablets	SE/H/0186/002	15235	MERCK AB	SE
Libracor 2.5 mg film-coated tablets	SE/H/0186/002	15235	MERCK AB	SE
Libracor 2.5 mg film-coated tablets	SE/H/0186/002	15235	MERCK AB	SE
Libracor 2.5 mg film-coated tablets	SE/H/0186/002	15235	MERCK AB	SE

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
Libracor 2.5 mg film-coated tablets	SE/H/0186/002	15235	MERCK AB	SE
Libracor 2.5 mg film-coated tablets	SE/H/0186/002	15235	MERCK AB	SE
Libracor 2.5 mg film-coated tablets	SE/H/0186/002	15235	MERCK AB	SE
Libracor 5 mg film-coated tablets	SE/H/0186/004	15379	MERCK AB	SE
Libracor 5 mg film-coated tablets	SE/H/0186/004	15379	MERCK AB	SE
Libracor 5 mg film-coated tablets	SE/H/0186/004	15379	MERCK AB	SE
Libracor 5 mg film-coated tablets	SE/H/0186/004	15379	MERCK AB	SE
Libracor 5 mg film-coated tablets	SE/H/0186/004	15379	MERCK AB	SE
Libracor 5 mg film-coated tablets	SE/H/0186/004	15379	MERCK AB	SE
Libracor 5 mg film-coated tablets	SE/H/0186/004	15379	MERCK AB	SE
Libracor 5 mg film-coated tablets	SE/H/0186/004	15379	MERCK AB	SE
Libracor 3.75 mg film-coated tablets	SE/H/0186/003	15378	MERCK AB	SE
Libracor 3.75 mg film-coated tablets	SE/H/0186/003	15378	MERCK AB	SE
Libracor 3.75 mg film-coated tablets	SE/H/0186/003	15378	MERCK AB	SE
Libracor 3.75 mg film-coated tablets	SE/H/0186/003	15378	MERCK AB	SE
Libracor 3.75 mg film-coated tablets	SE/H/0186/003	15378	MERCK AB	SE
Libracor 3.75 mg film-coated tablets	SE/H/0186/003	15378	MERCK AB	SE
Libracor 3.75 mg film-	SE/H/0186/003	15378	MERCK AB	SE

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
coated tablets				
Libracor 3.75 mg film-coated tablets	SE/H/0186/003	15378	MERCK AB	SE
Libracor 7.5 mg film-coated tablets	SE/H/0186/005	15236	MERCK AB	SE
Libracor 7.5 mg film-coated tablets	SE/H/0186/005	15236	MERCK AB	SE
Libracor 7.5 mg film-coated tablets	SE/H/0186/005	15236	MERCK AB	SE
Libracor 7.5 mg film-coated tablets	SE/H/0186/005	15236	MERCK AB	SE
Libracor 7.5 mg film-coated tablets	SE/H/0186/005	15236	MERCK AB	SE
Libracor 7.5 mg film-coated tablets	SE/H/0186/005	15236	MERCK AB	SE
Libracor 7.5 mg film-coated tablets	SE/H/0186/005	15236	MERCK AB	SE
Libracor 7.5 mg film-coated tablets	SE/H/0186/005	15236	MERCK AB	SE
Libracor 10 mg film-coated tablets	SE/H/0186/006	15237	MERCK AB	SE
Libracor 10 mg film-coated tablets	SE/H/0186/006	15237	MERCK AB	SE
Libracor 10 mg film-coated tablets	SE/H/0186/006	15237	MERCK AB	SE
Libracor 10 mg film-coated tablets	SE/H/0186/006	15237	MERCK AB	SE
Libracor 10 mg film-coated tablets	SE/H/0186/006	15237	MERCK AB	SE
Libracor 10 mg film-coated tablets	SE/H/0186/006	15237	MERCK AB	SE
Libracor 10 mg film-coated tablets	SE/H/0186/006	15237	MERCK AB	SE
Libracor 10 mg film-coated tablets	SE/H/0186/006	15237	MERCK AB	SE

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
Libracor 1.25 mg film-coated tablets	SE/H/0186/001	15377	MERCK AB	SE
Libracor 1.25 mg film-coated tablets	SE/H/0186/001	15377	MERCK AB	SE
Libracor 1.25 mg film-coated tablets	SE/H/0186/001	15377	MERCK AB	SE
Libracor 1.25 mg film-coated tablets	SE/H/0186/001	15377	MERCK AB	SE
Libracor 1.25 mg film-coated tablets	SE/H/0186/001	15377	MERCK AB	SE
Libracor 1.25 mg film-coated tablets	SE/H/0186/001	15377	MERCK AB	SE
Libracor 1.25 mg film-coated tablets	SE/H/0186/001	15377	MERCK AB	SE
Libracor 1.25 mg film-coated tablets	SE/H/0186/001	15377	MERCK AB	SE
Libracor 1.25 mg film-coated tablets	SE/H/0186/001	15377	MERCK AB	SE
Libracor 10 mg film-coated tablets	SE/H/0186/006	15237	MERCK AB	SE
Libracor 2.5 mg film-coated tablets	SE/H/0186/002	15235	MERCK AB	SE
Libracor 3.75 mg film-coated tablets	SE/H/0186/003	15378	MERCK AB	SE
Libracor 5 mg film-coated tablets	SE/H/0186/004	15379	MERCK AB	SE
Libracor 7.5 mg film-coated tablets	SE/H/0186/005	15236	MERCK AB	SE
Concor COR 1,25 mg filmsko obložene tablete	not available	H/02/00413/005	MERCK D.O.O.	SI
Concor COR 10 mg filmsko obložene tablete	not available	H/02/00413/004	MERCK D.O.O.	SI
Concor COR 2,5 mg filmsko obložene tablete	not available	H/02/00413/001	MERCK D.O.O.	SI
Concor COR 5 mg	not available	H/02/00413/002	MERCK D.O.O.	SI

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
filmsko obložene tablete				
Concor COR 7,5 mg filmsko obložene tablete	not available	H/02/00413/003	MERCK D.O.O.	SI
Byol 1,25 mg filmsko obložene tablete	NL/H/0684/001	H/10/00320/001	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Byol 1,25 mg filmsko obložene tablete	NL/H/0684/001	H/10/00320/002	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Byol 1,25 mg filmsko obložene tablete	NL/H/0684/001	H/10/00320/003	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Byol 1,25 mg filmsko obložene tablete	NL/H/0684/001	H/10/00320/004	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Byol 1,25 mg filmsko obložene tablete	NL/H/0684/001	H/10/00320/005	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Byol 1,25 mg filmsko obložene tablete	NL/H/0684/001	H/10/00320/006	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Byol 1,25 mg filmsko obložene tablete	NL/H/0684/001	H/10/00320/007	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Byol 1,25 mg filmsko obložene tablete	NL/H/0684/001	H/10/00320/008	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Byol 1,25 mg filmsko obložene tablete	NL/H/0684/001	H/10/00320/009	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Byol 1,25 mg filmsko obložene tablete	NL/H/0684/001	H/10/00320/010	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Byol 1,25 mg filmsko obložene tablete	NL/H/0684/001	H/10/00320/011	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Byol 1,25 mg filmsko obložene tablete	NL/H/0684/001	H/10/00320/012	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Byol 1,25 mg filmsko obložene tablete	NL/H/0684/001	H/10/00320/013	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Byol 1,25 mg filmsko obložene tablete	NL/H/0684/001	H/10/00320/014	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Byol 1,25 mg filmsko obložene tablete	NL/H/0684/001	H/10/00320/015	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Byol 1,25 mg filmsko obložene tablete	NL/H/0684/001	H/10/00320/016	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI

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
Byol 1,25 mg filmsko obložene tablete	NL/H/0684/001	H/10/00320/017	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Byol 1,25 mg filmsko obložene tablete	NL/H/0684/001	H/10/00320/018	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Byol 1,25 mg filmsko obložene tablete	NL/H/0684/001	H/10/00320/019	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Byol 1,25 mg filmsko obložene tablete	NL/H/0684/001	H/10/00320/020	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Byol 1,25 mg filmsko obložene tablete	NL/H/0684/001	H/10/00320/021	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Concor 10 mg filmsko obložene tablete	not available	H/98/00412/002	MERCK D.O.O.	SI
Concor 5 mg filmsko obložene tablete	not available	H/98/00412/001	MERCK D.O.O.	SI
Conaret 5 mg tablety	DE/H/6433/004	41/0098/021-S	ZENTIVA, K.S.	SK
Conaret 5 mg tablety	DE/H/6433/004	41/0098/021-S	ZENTIVA, K.S.	SK
Conaret 5 mg tablety	DE/H/6433/004	41/0098/021-S	ZENTIVA, K.S.	SK
Conaret 5 mg tablety	DE/H/6433/004	41/0098/021-S	ZENTIVA, K.S.	SK
Conaret 5 mg tablety	DE/H/6433/004	41/0098/021-S	ZENTIVA, K.S.	SK
Conaret 5 mg tablety	DE/H/6433/004	41/0098/021-S	ZENTIVA, K.S.	SK
Conaret 5 mg tablety	DE/H/6433/004	41/0098/021-S	ZENTIVA, K.S.	SK
Conaret 5 mg tablety	DE/H/6433/004	41/0098/021-S	ZENTIVA, K.S.	SK
Conaret 5 mg tablety	DE/H/6433/004	41/0098/021-S	ZENTIVA, K.S.	SK
Conaret 5 mg tablety	DE/H/6433/004	41/0098/021-S	ZENTIVA, K.S.	SK
Conaret 5 mg tablety	DE/H/6433/004	41/0098/021-S	ZENTIVA, K.S.	SK
Conaret 5 mg tablety	DE/H/6433/004	41/0098/021-S	ZENTIVA, K.S.	SK
Conaret 5 mg tablety	DE/H/6433/004	41/0098/021-S	ZENTIVA, K.S.	SK
Conaret 10 mg tablety	DE/H/6433/006	41/0100/21-S	ZENTIVA, K.S.	SK
Conaret 10 mg tablety	DE/H/6433/006	41/0100/21-S	ZENTIVA, K.S.	SK
Conaret 10 mg tablety	DE/H/6433/006	41/0100/21-S	ZENTIVA, K.S.	SK
Conaret 10 mg tablety	DE/H/6433/006	41/0100/21-S	ZENTIVA, K.S.	SK
Conaret 10 mg tablety	DE/H/6433/006	41/0100/21-S	ZENTIVA, K.S.	SK
Conaret 10 mg tablety	DE/H/6433/006	41/0100/21-S	ZENTIVA, K.S.	SK
Conaret 10 mg tablety	DE/H/6433/006	41/0100/21-S	ZENTIVA, K.S.	SK

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
Conaret 10 mg tablety	DE/H/6433/006	41/0100/21-S	ZENTIVA, K.S.	SK
Conaret 10 mg tablety	DE/H/6433/006	41/0100/21-S	ZENTIVA, K.S.	SK
Conaret 10 mg tablety	DE/H/6433/006	41/0100/21-S	ZENTIVA, K.S.	SK
Conaret 10 mg tablety	DE/H/6433/006	41/0100/21-S	ZENTIVA, K.S.	SK
Conaret 10 mg tablety	DE/H/6433/006	41/0100/21-S	ZENTIVA, K.S.	SK
Conaret 10 mg tablety	DE/H/6433/006	41/0100/21-S	ZENTIVA, K.S.	SK
Conaret 5 mg tablety	DE/H/6433/004	41/0098/021-S	ZENTIVA, K.S.	SK
Conaret 10 mg tablety	DE/H/6433/006	41/0100/21-S	ZENTIVA, K.S.	SK
Bisomerck 5 mg filmom obalené tablety	not available	41/0054/03-S	MERCK SPOL.S.R.O.	SK
Concor 5 mg filmom obalené tablety	not available	41/0304/89-S	MERCK SPOL.S.R.O.	SK
Concor 5 mg filmom obalené tablety	not available	41/0304/89-S	MERCK SPOL.S.R.O.	SK
Concor 10 mg filmom obalené tablety	not available	41/0176/19-S	MERCK SPOL.S.R.O.	SK
Concor 10 mg filmom obalené tablety	not available	41/0176/19-S	MERCK SPOL.S.R.O.	SK
Concor 10 mg filmom obalené tablety	not available	41/0176/19-S	MERCK SPOL.S.R.O.	SK
Concor 5 mg filmom obalené tablety	not available	41/0304/89-S	MERCK SPOL.S.R.O.	SK
Concor COR 5 mg filmom obalené tablety	not available	41/0011/02-S	MERCK SPOL.S.R.O.	SK
Concor COR 10 mg filmom obalené tablety	not available	41/0012/02-S	MERCK SPOL.S.R.O.	SK
Concor COR 2,5 mg filmom obalené tablety	not available	41/0010/02-S	MERCK SPOL.S.R.O.	SK
Concor COR 2,5 mg filmom obalené tablety	not available	41/0010/02-S	MERCK SPOL.S.R.O.	SK
Concor COR 10 mg filmom obalené tablety	not available	41/0012/02-S	MERCK SPOL.S.R.O.	SK
Concor COR 5 mg filmom obalené tablety	not available	41/0011/02-S	MERCK SPOL.S.R.O.	SK
Concor COR 2,5 mg	not available	41/0010/02-S	MERCK SPOL.S.R.O.	SK

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
filmom obalené tablety				
Bisoprolol Fumarate 0.5mg/ml Oral Solution	UK/H/	PL 39307/0140	SYRI LIMITED T/A THAME LABORATORIES	XI
Cardicor 10 mg film-coated tablets	SE/H/0184/006	PL 11648/0076	MERCK SERONO LIMITED	XI
Cardicor 10 mg film-coated tablets	SE/H/0184/006	PL 11648/0076	MERCK SERONO LIMITED	XI
Cardicor 10 mg film-coated tablets	SE/H/0184/006	PL 11648/0076	MERCK SERONO LIMITED	XI
Cardicor 10 mg film-coated tablets	SE/H/0184/006	PL 11648/0076	MERCK SERONO LIMITED	XI
Cardicor 10 mg film-coated tablets	SE/H/0184/006	PL 11648/0076	MERCK SERONO LIMITED	XI
Cardicor 10 mg film-coated tablets	SE/H/0184/006	PL 11648/0076	MERCK SERONO LIMITED	XI
Cardicor 10 mg film-coated tablets	SE/H/0184/006	PL 11648/0076	MERCK SERONO LIMITED	XI
Cardicor 10 mg film-coated tablets	SE/H/0184/006	PL 11648/0076	MERCK SERONO LIMITED	XI
Cardicor 10 mg film-coated tablets	SE/H/0184/006	PL 11648/0076	MERCK SERONO LIMITED	XI
Cardicor 10 mg film-coated tablets	SE/H/0184/006	PL 11648/0076	MERCK SERONO LIMITED	XI
Cardicor 10 mg film-coated tablets	SE/H/0184/006	PL 11648/0076	MERCK SERONO LIMITED	XI
Cardicor 10 mg film-coated tablets	SE/H/0184/006	PL 11648/0076	MERCK SERONO LIMITED	XI
Cardicor 10 mg film-coated tablets	SE/H/0184/006	PL 11648/0076	MERCK SERONO LIMITED	XI
Cardicor 10 mg film-coated tablets	SE/H/0184/006	PL 11648/0076	MERCK SERONO LIMITED	XI
Cardicor 10 mg film-coated tablets	SE/H/0184/006	PL 11648/0076	MERCK SERONO LIMITED	XI
Cardicor 10 mg film-coated tablets	SE/H/0184/006	PL 11648/0076	MERCK SERONO LIMITED	XI

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
Cardicor 3.75 mg film-coated tablets	SE/H/0184/003	PL 11648/0073	MERCK SERONO LIMITED	XI
Cardicor 3.75 mg film-coated tablets	SE/H/0184/003	PL 11648/0073	MERCK SERONO LIMITED	XI
Cardicor 3.75 mg film-coated tablets	SE/H/0184/003	PL 11648/0073	MERCK SERONO LIMITED	XI
Cardicor 3.75 mg film-coated tablets	SE/H/0184/003	PL 11648/0073	MERCK SERONO LIMITED	XI
Cardicor 3.75 mg film-coated tablets	SE/H/0184/003	PL 11648/0073	MERCK SERONO LIMITED	XI
Cardicor 3.75 mg film-coated tablets	SE/H/0184/003	PL 11648/0073	MERCK SERONO LIMITED	XI
Cardicor 3.75 mg film-coated tablets	SE/H/0184/003	PL 11648/0073	MERCK SERONO LIMITED	XI
Cardicor 3.75 mg film-coated tablets	SE/H/0184/003	PL 11648/0073	MERCK SERONO LIMITED	XI
Cardicor 3.75 mg film-coated tablets	SE/H/0184/003	PL 11648/0073	MERCK SERONO LIMITED	XI
Cardicor 3.75 mg film-coated tablets	SE/H/0184/003	PL 11648/0073	MERCK SERONO LIMITED	XI
Cardicor 3.75 mg film-coated tablets	SE/H/0184/003	PL 11648/0073	MERCK SERONO LIMITED	XI
Cardicor 3.75 mg film-coated tablets	SE/H/0184/003	PL 11648/0073	MERCK SERONO LIMITED	XI
Cardicor 3.75 mg film-coated tablets	SE/H/0184/003	PL 11648/0073	MERCK SERONO LIMITED	XI
Cardicor 3.75 mg film-coated tablets	SE/H/0184/003	PL 11648/0073	MERCK SERONO LIMITED	XI
Cardicor 3.75 mg film-coated tablets	SE/H/0184/003	PL 11648/0073	MERCK SERONO LIMITED	XI
Cardicor 3.75 mg film-coated tablets	SE/H/0184/003	PL 11648/0073	MERCK SERONO LIMITED	XI
Cardicor 7.5 mg film-coated tablets	SE/H/0184/005	PL 11648/0075	MERCK SERONO LIMITED	XI
Cardicor 7.5 mg film-	SE/H/0184/005	PL 11648/0075	MERCK SERONO LIMITED	XI

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
coated tablets				
Cardicor 7.5 mg film-coated tablets	SE/H/0184/005	PL 11648/0075	MERCK SERONO LIMITED	XI
Cardicor 7.5 mg film-coated tablets	SE/H/0184/005	PL 11648/0075	MERCK SERONO LIMITED	XI
Cardicor 7.5 mg film-coated tablets	SE/H/0184/005	PL 11648/0075	MERCK SERONO LIMITED	XI
Cardicor 7.5 mg film-coated tablets	SE/H/0184/005	PL 11648/0075	MERCK SERONO LIMITED	XI
Cardicor 7.5 mg film-coated tablets	SE/H/0184/005	PL 11648/0075	MERCK SERONO LIMITED	XI
Cardicor 7.5 mg film-coated tablets	SE/H/0184/005	PL 11648/0075	MERCK SERONO LIMITED	XI
Cardicor 7.5 mg film-coated tablets	SE/H/0184/005	PL 11648/0075	MERCK SERONO LIMITED	XI
Cardicor 7.5 mg film-coated tablets	SE/H/0184/005	PL 11648/0075	MERCK SERONO LIMITED	XI
Cardicor 7.5 mg film-coated tablets	SE/H/0184/005	PL 11648/0075	MERCK SERONO LIMITED	XI
Cardicor 7.5 mg film-coated tablets	SE/H/0184/005	PL 11648/0075	MERCK SERONO LIMITED	XI
Cardicor 7.5 mg film-coated tablets	SE/H/0184/005	PL 11648/0075	MERCK SERONO LIMITED	XI
Cardicor 7.5 mg film-coated tablets	SE/H/0184/005	PL 11648/0075	MERCK SERONO LIMITED	XI
Cardicor 7.5 mg film-coated tablets	SE/H/0184/005	PL 11648/0075	MERCK SERONO LIMITED	XI
Cardicor 7.5 mg film-coated tablets	SE/H/0184/005	PL 11648/0075	MERCK SERONO LIMITED	XI
Cardicor 1.25 mg film-coated tablets	SE/H/0184/001	PL 11648/0071	MERCK SERONO LIMITED	XI
Cardicor 1.25 mg film-coated tablets	SE/H/0184/001	PL 11648/0071	MERCK SERONO LIMITED	XI
Cardicor 1.25 mg film-coated tablets	SE/H/0184/001	PL 11648/0071	MERCK SERONO LIMITED	XI

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
Cardicor 1.25 mg film-coated tablets	SE/H/0184/001	PL 11648/0071	MERCK SERONO LIMITED	XI
Cardicor 1.25 mg film-coated tablets	SE/H/0184/001	PL 11648/0071	MERCK SERONO LIMITED	XI
Cardicor 1.25 mg film-coated tablets	SE/H/0184/001	PL 11648/0071	MERCK SERONO LIMITED	XI
Cardicor 1.25 mg film-coated tablets	SE/H/0184/001	PL 11648/0071	MERCK SERONO LIMITED	XI
Cardicor 1.25 mg film-coated tablets	SE/H/0184/001	PL 11648/0071	MERCK SERONO LIMITED	XI
Cardicor 1.25 mg film-coated tablets	SE/H/0184/001	PL 11648/0071	MERCK SERONO LIMITED	XI
Cardicor 1.25 mg film-coated tablets	SE/H/0184/001	PL 11648/0071	MERCK SERONO LIMITED	XI
Cardicor 1.25 mg film-coated tablets	SE/H/0184/001	PL 11648/0071	MERCK SERONO LIMITED	XI
Cardicor 1.25 mg film-coated tablets	SE/H/0184/001	PL 11648/0071	MERCK SERONO LIMITED	XI
Cardicor 1.25 mg film-coated tablets	SE/H/0184/001	PL 11648/0071	MERCK SERONO LIMITED	XI
Cardicor 1.25 mg film-coated tablets	SE/H/0184/001	PL 11648/0071	MERCK SERONO LIMITED	XI
Cardicor 1.25 mg film-coated tablets	SE/H/0184/001	PL 11648/0071	MERCK SERONO LIMITED	XI
Cardicor 1.25 mg film-coated tablets	SE/H/0184/001	PL 11648/0071	MERCK SERONO LIMITED	XI
Cardicor 2.5 mg film-coated tablets	SE/H/0184/002	PL 11648/0072	MERCK SERONO LIMITED	XI
Cardicor 2.5 mg film-coated tablets	SE/H/0184/002	PL 11648/0072	MERCK SERONO LIMITED	XI
Cardicor 2.5 mg film-coated tablets	SE/H/0184/002	PL 11648/0072	MERCK SERONO LIMITED	XI
Cardicor 2.5 mg film-coated tablets	SE/H/0184/002	PL 11648/0072	MERCK SERONO LIMITED	XI
Cardicor 2.5 mg film-	SE/H/0184/002	PL 11648/0072	MERCK SERONO LIMITED	XI

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
coated tablets				
Cardicor 2.5 mg film-coated tablets	SE/H/0184/002	PL 11648/0072	MERCK SERONO LIMITED	XI
Cardicor 2.5 mg film-coated tablets	SE/H/0184/002	PL 11648/0072	MERCK SERONO LIMITED	XI
Cardicor 2.5 mg film-coated tablets	SE/H/0184/002	PL 11648/0072	MERCK SERONO LIMITED	XI
Cardicor 2.5 mg film-coated tablets	SE/H/0184/002	PL 11648/0072	MERCK SERONO LIMITED	XI
Cardicor 2.5 mg film-coated tablets	SE/H/0184/002	PL 11648/0072	MERCK SERONO LIMITED	XI
Cardicor 2.5 mg film-coated tablets	SE/H/0184/002	PL 11648/0072	MERCK SERONO LIMITED	XI
Cardicor 2.5 mg film-coated tablets	SE/H/0184/002	PL 11648/0072	MERCK SERONO LIMITED	XI
Cardicor 2.5 mg film-coated tablets	SE/H/0184/002	PL 11648/0072	MERCK SERONO LIMITED	XI
Cardicor 2.5 mg film-coated tablets	SE/H/0184/002	PL 11648/0072	MERCK SERONO LIMITED	XI
Cardicor 2.5 mg film-coated tablets	SE/H/0184/002	PL 11648/0072	MERCK SERONO LIMITED	XI
Cardicor 2.5 mg film-coated tablets	SE/H/0184/002	PL 11648/0072	MERCK SERONO LIMITED	XI
Cardicor 5 mg film-coated tablets	SE/H/0184/004	PL 11648/0074	MERCK SERONO LIMITED	XI
Cardicor 5 mg film-coated tablets	SE/H/0184/004	PL 11648/0074	MERCK SERONO LIMITED	XI
Cardicor 5 mg film-coated tablets	SE/H/0184/004	PL 11648/0074	MERCK SERONO LIMITED	XI
Cardicor 5 mg film-coated tablets	SE/H/0184/004	PL 11648/0074	MERCK SERONO LIMITED	XI
Cardicor 5 mg film-coated tablets	SE/H/0184/004	PL 11648/0074	MERCK SERONO LIMITED	XI
Cardicor 5 mg film-coated tablets	SE/H/0184/004	PL 11648/0074	MERCK SERONO LIMITED	XI

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
Cardicor 5 mg film-coated tablets	SE/H/0184/004	PL 11648/0074	MERCK SERONO LIMITED	XI
Cardicor 5 mg film-coated tablets	SE/H/0184/004	PL 11648/0074	MERCK SERONO LIMITED	XI
Cardicor 5 mg film-coated tablets	SE/H/0184/004	PL 11648/0074	MERCK SERONO LIMITED	XI
Cardicor 5 mg film-coated tablets	SE/H/0184/004	PL 11648/0074	MERCK SERONO LIMITED	XI
Cardicor 5 mg film-coated tablets	SE/H/0184/004	PL 11648/0074	MERCK SERONO LIMITED	XI
Cardicor 5 mg film-coated tablets	SE/H/0184/004	PL 11648/0074	MERCK SERONO LIMITED	XI
Cardicor 5 mg film-coated tablets	SE/H/0184/004	PL 11648/0074	MERCK SERONO LIMITED	XI
Cardicor 5 mg film-coated tablets	SE/H/0184/004	PL 11648/0074	MERCK SERONO LIMITED	XI
Cardicor 5 mg film-coated tablets	SE/H/0184/004	PL 11648/0074	MERCK SERONO LIMITED	XI
Cardicor 5 mg film-coated tablets	SE/H/0184/004	PL 11648/0074	MERCK SERONO LIMITED	XI
Bisoprolol 2.5mg film-coated Tablets	not available	PL 44041/0003	NOUMED LIFE SCIENCES	XI
Bisoprolol 10mg film-coated Tablets	not available	PL 44041/0005	NOUMED LIFE SCIENCES	XI
Bisoprolol Fumarate 10 mg Film-coated Tablets	not available	PL 43461/0052	FLAMINGO PHARMA UK LTD	XI
Bisoprolol Fumarate 5 mg Film-coated Tablets	not available	PL 43461/0051	FLAMINGO PHARMA UK LTD	XI