13 May 2016
EMA/351661/2016
Procedure Management and Committees Support

List of nationally authorised medicinal products

Active substance: bisoprolol

Procedure no.: PSUSA/00000419/201509

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Product full name (in authorisation country)	**MRP / DCP Authorisation number**	**National Authorisation Number**	**MAH of product in the member state**	**Member State where product is authorised**	**Legal basis**
Bisoprolol AUROBINDO 5 mg, comprimé pelliculé sécable	UK/H/1684/001	223 277-2	AUROBINDO PHARMA FRANCE SARL	FR	Hybrid application (Article 10(3) of Directive No 2001/83/EC)
Bisoprolol AUROBINDO 5 mg, comprimé pelliculé sécable	UK/H/1684/001	223 278-9	AUROBINDO PHARMA FRANCE SARL	FR	Hybrid application (Article 10(3) of Directive No 2001/83/EC)
Bisoprolol AUROBINDO 5 mg, comprimé pelliculé sécable	UK/H/1684/001	582 637-7	AUROBINDO PHARMA FRANCE SARL	FR	Hybrid application (Article 10(3) of Directive No 2001/83/EC)
Concor COR 1,25 mg filmom obložene tablete	SE/H/0184/001	HR-H-350108325-04	MERCK D.O.O	HR	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor COR 1,25 mg filmom obložene tablete	SE/H/0184/001	HR-H-350108325-05	MERCK D.O.O	HR	Full application (Article 8(3) of Directive No 2001/83/EC)

Product full name (in authorisation country)	**MRP / DCP Authorisation number**	**National Authorisation Number**	**MAH of product in the member state**	**Member State where product is authorised**	**Legal basis**
Concor COR 1,25 mg filmom obložene tablete	SE/H/0184/001	HR-H-350108325-06	MERCK D.O.O	HR	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor COR 1,25 mg filmom obložene tablete	SE/H/0184/001	HR-H-350108325-07	MERCK D.O.O	HR	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor COR 2,5 mg filmom obložene tablete	SE/H/0184/002	HR-H-061130840-08	MERCK D.O.O	HR	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor COR 2,5 mg filmom obložene tablete	SE/H/0184/002	HR-H-061130840-01	MERCK D.O.O	HR	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor COR 2,5 mg filmom obložene tablete	SE/H/0184/002	HR-H-061130840-02	MERCK D.O.O	HR	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor COR 2,5 mg filmom obložene tablete	SE/H/0184/002	HR-H-061130840-03	MERCK D.O.O	HR	Full application (Article 8(3) of Directive No 2001/83/EC)

Product full name (in authorisation country)	**MRP / DCP Authorisation number**	**National Authorisation Number**	**MAH of product in the member state**	**Member State where product is authorised**	**Legal basis**
Concor COR 2,5 mg filmom obložene tablete	SE/H/0184/002	HR-H-061130840-04	MERCK D.O.O	HR	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor COR 2,5 mg filmom obložene tablete	SE/H/0184/002	HR-H-061130840-05	MERCK D.O.O	HR	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor COR 2,5 mg filmom obložene tablete	SE/H/0184/002	HR-H-061130840-06	MERCK D.O.O	HR	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor COR 2,5 mg filmom obložene tablete	SE/H/0184/002	HR-H-061130840-07	MERCK D.O.O	HR	Full application (Article 8(3) of Directive No 2001/83/EC)
Emconcor 10 mg Filmtabletten	not available	BE138363	MERCK N.V.-S.A.	BE	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor 10 mg film-coated tablets	not available	21708	MERCK A.E.	CY	Full application (Article 8(3) of Directive No 2001/83/EC)

Product full name (in authorisation country)	**MRP / DCP Authorisation number**	**National Authorisation Number**	**MAH of product in the member state**	**Member State where product is authorised**	**Legal basis**
Bisoprolol Merck 3.75 mg film-coated tablets	SE/H/0187/003	15381	MERCK KGAA	SE	Full application (Article 8(3) of Directive No 2001/83/EC)
Bisoprolol Merck 5 mg film-coated tablets	SE/H/0187/004	15382	MERCK KGAA	SE	Full application (Article 8(3) of Directive No 2001/83/EC)
Cardicor 2.5 mg film-coated tablets	SE/H/0185/002	15232	MERCK KGAA	SE	Full application (Article 8(3) of Directive No 2001/83/EC)
EMCONCOR® 5 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0187/004	15259/5-3-2015	MERCK A.E.	GR	Full application (Article 8(3) of Directive No 2001/83/EC)
EMCONCOR® 2,5 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0187/002	15255/5-3-2015	MERCK A.E.	GR	Full application (Article 8(3) of Directive No 2001/83/EC)
Emconcor Minor 2,5 mg comprimés pelliculés	SE/H/0184/002	BE472746	MERCK N.V.-S.A.	BE	Full application (Article 8(3) of Directive No 2001/83/EC)

Product full name (in authorisation country)	**MRP / DCP Authorisation number**	**National Authorisation Number**	**MAH of product in the member state**	**Member State where product is authorised**	**Legal basis**
Конкор КОР 2.5 mg филмирани таблетки	not available	20060181	MERCK KGAA	BG	Full application (Article 8(3) of Directive No 2001/83/EC)
Конкор КОР 5 mg филмирани таблетки	not available	20060182	MERCK KGAA	BG	Full application (Article 8(3) of Directive No 2001/83/EC)
Конкор 10 mg филмирани таблетки	not available	II-7679/27.06.2003	MERCK KGAA	BG	Full application (Article 8(3) of Directive No 2001/83/EC)
Конкор 5 mg филмирани таблетки	not available	II-7678/27.06.2003	MERCK KGAA	BG	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor 5 mg film-coated tablets	not available	12863	MERCK A.E.	CY	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor COR 10 mg potahované tablety	not available	77/028/01-C	MERCK KGAA	CZ	Full application (Article 8(3) of Directive No 2001/83/EC)

Product full name (in authorisation country)	**MRP / DCP Authorisation number**	**National Authorisation Number**	**MAH of product in the member state**	**Member State where product is authorised**	**Legal basis**
Concor COR 2,5 mg potahované tablety	not available	77/026/01-C	MERCK KGAA	CZ	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor COR 5 mg potahované tablety	not available	77/027/01-C	MERCK KGAA	CZ	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor 10, potahované tablety	not available	41/304/89-B/C	MERCK KGAA	CZ	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor 5, potahované tablety	not available	41/304/89-A/C	MERCK KGAA	CZ	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor 10 mg Filmtabletten	not available	6849.01.00	MERCK SERONO GMBH	DE	Full application (Article 8(3) of Directive No 2001/83/EC)
Emconcor Cor 2,5 mg comprimidos recubiertos con película	SE/H/0184/002	63046	MERCK KGAA	ES	Full application (Article 8(3) of Directive No 2001/83/EC)

Product full name (in authorisation country)	**MRP / DCP Authorisation number**	**National Authorisation Number**	**MAH of product in the member state**	**Member State where product is authorised**	**Legal basis**
Emconcor Cor 5 mg comprimidos recubiertos con película	SE/H/0184/004	63048	MERCK KGAA	ES	Full application (Article 8(3) of Directive No 2001/83/EC)
Emconcor 10 mg comprimidos recubiertos con película	not available	57.655	MERCK, S.L.	ES	Full application (Article 8(3) of Directive No 2001/83/EC)
Emconcor 5 mg comprimidos recubiertos con película	not available	57.654	MERCK, S.L.	ES	Full application (Article 8(3) of Directive No 2001/83/EC)
Emconcor CHF 1,25 mg tabletti, kalvopäällysteinen	SE/H/0184/001	14531	MERCK OY	FI	Full application (Article 8(3) of Directive No 2001/83/EC)
EMCONCOR CHF 10 mg tabletti, kalvopäällysteinen	SE/H/0184/006	14536	MERCK OY	FI	Full application (Article 8(3) of Directive No 2001/83/EC)
Emconcor CHF 2,5 mg tabletti, kalvopäällysteinen	SE/H/0184/002	14532	MERCK OY	FI	Full application (Article 8(3) of Directive No 2001/83/EC)

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Emconcor CHF 3,75 mg tabletti, kalvopäällysteinen	SE/H/0184/003	14533	MERCK OY	FI	Full application (Article 8(3) of Directive No 2001/83/EC)
Emconcor CHF 5 mg tabletti, kalvopäällysteinen	SE/H/0184/004	14534	MERCK OY	FI	Full application (Article 8(3) of Directive No 2001/83/EC)
Emconcor CHF 7,5 mg tabletti, kalvopäällysteinen	SE/H/0184/005	14535	MERCK OY	FI	Full application (Article 8(3) of Directive No 2001/83/EC)
Emconcor 10 mg tabletti, kalvopäällysteinen	not available	10191	MERCK KGAA	FI	Full application (Article 8(3) of Directive No 2001/83/EC)
Emconcor 5 mg tabletti, kalvopäällysteinen	not available	10190	MERCK KGAA	FI	Full application (Article 8(3) of Directive No 2001/83/EC)
Congescor 1.25 mg film-coated tablets	not available	PL 11648/0035	MERCK SERONO LTD.	UK	Full application (Article 8(3) of Directive No 2001/83/EC)

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Cardicor 1.25 mg film-coated tablets	SE/H/0184/001	PL 11648/0071	MERCK SERONO LTD.	UK	Full application (Article 8(3) of Directive No 2001/83/EC)
Cardicor 10 mg film-coated tablets	SE/H/0184/006	PL 11648/0076	MERCK SERONO LTD.	UK	Full application (Article 8(3) of Directive No 2001/83/EC)
Congescor 2.5 mg film-coated tablets	not available	PL 11648/0036	MERCK SERONO LTD.	UK	Full application (Article 8(3) of Directive No 2001/83/EC)
Cardicor 2.5 mg film-coated tablets	SE/H/0184/002	PL 11648/0072	MERCK SERONO LTD.	UK	Full application (Article 8(3) of Directive No 2001/83/EC)
Cardicor 3.75 mg film-coated tablets	SE/H/0184/003	PL 11648/0073	MERCK SERONO LTD.	UK	Full application (Article 8(3) of Directive No 2001/83/EC)
Cardicor 5 mg film-coated tablets	SE/H/0184/004	PL 11648/0074	MERCK SERONO LTD.	UK	Full application (Article 8(3) of Directive No 2001/83/EC)

Product full name (in authorisation country)	**MRP / DCP Authorisation number**	**National Authorisation Number**	**MAH of product in the member state**	**Member State where product is authorised**	**Legal basis**
Emcor 10 mg film coated tablets	not available	PL 11648/0070	MERCK SERONO LTD.	UK	Full application (Article 8(3) of Directive No 2001/83/EC)
Emcor LS 5 mg film-coated tablets	not available	PL 11648/0069	MERCK SERONO LTD.	UK	Full application (Article 8(3) of Directive No 2001/83/EC)
CONCOR ® 10 mg επικαλυμμένα με λεπτό υμένιο δισκία	not available	78137/09-11-2011	MERCK A.E.	GR	Full application (Article 8(3) of Directive No 2001/83/EC)
CONCOR ® 5 mg επικαλυμμένα με λεπτό υμένιο δισκία	not available	36771/09-11-2011	MERCK A.E.	GR	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor COR 1.25 mg filmtabletta	not available	OGYI-T-8324/01	MERCK KFT.	HU	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor COR 1,25 mg filmtabletta	not available	OGYI-T-8324/02	MERCK KFT.	HU	Full application (Article 8(3) of Directive No 2001/83/EC)

Product full name (in authorisation country)	**MRP / DCP Authorisation number**	**National Authorisation Number**	**MAH of product in the member state**	**Member State where product is authorised**	**Legal basis**
Concor COR 10 mg filmtabletta	not available	OGYI-T-8324/12	MERCK KFT.	HU	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor COR 10 mg filmtabletta	not available	OGYI-T-8324/11	MERCK KFT.	HU	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor COR 2,5 mg filmtabletta	not available	OGYI-T-8324/04	MERCK KFT.	HU	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor COR 2,5 mg filmtabletta	not available	OGYI-T-8324/03	MERCK KFT.	HU	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor 10 mg filmtabletta	not available	OGYI-T-4015/06	MERCK KFT.	HU	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor 10 mg filmtabletta	not available	OGYI-T-4015/05	MERCK KFT.	HU	Full application (Article 8(3) of Directive No 2001/83/EC)

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Concor 10 mg filmtabletta	not available	OGYI-T-4015/04	MERCK KFT.	HU	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor 5 mg filmtabletta	not available	OGYI-T-4015/03	MERCK KFT.	HU	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor 5 mg filmtabletta	not available	OGYI-T-4015/02	MERCK KFT.	HU	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor 5 mg filmtabletta	not available	OGYI-T-4015/01	MERCK KFT.	HU	Full application (Article 8(3) of Directive No 2001/83/EC)
Cardicor 1.25 mg film-coated tablets	SE/H/0184/001	PA 654/7/1	MERCK SERONO LTD.	IE	Full application (Article 8(3) of Directive No 2001/83/EC)
Cardicor 10 mg film-coated tablets	SE/H/0184/006	PA 654/7/6	MERCK SERONO LTD.	IE	Full application (Article 8(3) of Directive No 2001/83/EC)

Product full name (in authorisation country)	**MRP / DCP Authorisation number**	**National Authorisation Number**	**MAH of product in the member state**	**Member State where product is authorised**	**Legal basis**
Cardicor 2.5 mg film-coated tablets	SE/H/0184/002	PA 654/7/2	MERCK SERONO LTD.	IE	Full application (Article 8(3) of Directive No 2001/83/EC)
Cardicor 3.75 mg film-coated tablets	SE/H/0184/003	PA 654/7/3	MERCK SERONO LTD.	IE	Full application (Article 8(3) of Directive No 2001/83/EC)
Cardicor 5 mg film-coated tablets	SE/H/0184/004	PA 654/7/4	MERCK SERONO LTD.	IE	Full application (Article 8(3) of Directive No 2001/83/EC)
Cardicor 7.5 mg film-coated tablets	SE/H/0184/005	PA 654/7/5	MERCK SERONO LTD.	IE	Full application (Article 8(3) of Directive No 2001/83/EC)
Emcor 10 mg film-coated tablets	not available	PA 654/12/2	MERCK SERONO LTD.	IE	Full application (Article 8(3) of Directive No 2001/83/EC)
Emcor 5 mg film-coated tablets	not available	PA 654/12/1	MERCK SERONO LTD.	IE	Full application (Article 8(3) of Directive No 2001/83/EC)

Product full name (in authorisation country)	**MRP / DCP Authorisation number**	**National Authorisation Number**	**MAH of product in the member state**	**Member State where product is authorised**	**Legal basis**
Concor COR 2,5 mg apvalkotās tabletes	not available	02-0168	MERCK KGAA	LV	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor COR 5 mg apvalkotās tabletes	not available	02-0169	MERCK KGAA	LV	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor 10 mg apvalkotās tabletes	not available	99-0011	MERCK KGAA	LV	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor 5 mg apvalkotās tabletes	not available	99-0010	MERCK KGAA	LV	Full application (Article 8(3) of Directive No 2001/83/EC)
EMCORDECO 2,5, filmomhulde tabletten 2,5 mg	SE/H/0184/002	RVG 24503	MERCK B.V.	NL	Full application (Article 8(3) of Directive No 2001/83/EC)
EMCORDECO 7,5, filmomhulde tabletten 7,5 mg	SE/H/0184/005	RVG 24506	MERCK B.V.	NL	Full application (Article 8(3) of Directive No 2001/83/EC)

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Emcor 5, filmomhulde tabletten 5 mg	not available	RVG 12408	MERCK B.V.	NL	Full application (Article 8(3) of Directive No 2001/83/EC)
Emconcor CHF 1,25 mg tablett, filmdrasjert	SE/H/0184/001	99-7737	MERCK KGAA	NO	Full application (Article 8(3) of Directive No 2001/83/EC)
Emconcor CHF 10 mg tablett, filmdrasjert	SE/H/0184/006	99-7742	MERCK KGAA	NO	Full application (Article 8(3) of Directive No 2001/83/EC)
Emconcor CHF 2.5 mg tablett, filmdrasjert	SE/H/0184/002	99-7738	MERCK KGAA	NO	Full application (Article 8(3) of Directive No 2001/83/EC)
Emconcor CHF 3.75 mg tablett, filmdrasjert	SE/H/0184/003	99-7739	MERCK KGAA	NO	Full application (Article 8(3) of Directive No 2001/83/EC)
Emconcor CHF 5 mg tablett, filmdrasjert	SE/H/0184/004	99-7740	MERCK KGAA	NO	Full application (Article 8(3) of Directive No 2001/83/EC)

Product full name (in authorisation country)	**MRP / DCP Authorisation number**	**National Authorisation Number**	**MAH of product in the member state**	**Member State where product is authorised**	**Legal basis**
Emconcor 5 mg film-coated tablets	not available	02-1326	MERCK KGAA	NO	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor Cor 1 ,25, 1,25 mg, tabletki powlekane	not available	8589	MERCK KGAA	PL	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor Cor 10, 10 mg, tabletki powlekane	not available	8594	MERCK KGAA	PL	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor Cor 2,5, 2,5 mg, tabletki powlekane	not available	8590	MERCK KGAA	PL	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor Cor 3,75, 3,75 mg, tabletki powlekane	not available	8591	MERCK KGAA	PL	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor Cor 5, 5 mg, tabletki powlekane	not available	8592	MERCK KGAA	PL	Full application (Article 8(3) of Directive No 2001/83/EC)

Product full name (in authorisation country)	**MRP / DCP Authorisation number**	**National Authorisation Number**	**MAH of product in the member state**	**Member State where product is authorised**	**Legal basis**
Concor Cor 7,5, 7,5 mg, tabletki powlekane	not available	8593	MERCK KGAA	PL	Full application (Article 8(3) of Directive No 2001/83/EC)
Bisopromerck 10; 10 mg tabletki powlekane	not available	4895	MERCK KGAA	PL	Generic application (Article 10(1)
Concor, 10 mg tabletki powlekane	not available	R/3735	MERCK KGAA	PL	Full application (Article 8(3) of Directive No 2001/83/EC)
Bisopromerck 5; 5 mg tabletki powlekane	not available	4894	MERCK KGAA	PL	Generic application (Article 10(1)
Concor, 5 mg tabletki powlekane	not available	R/3734	MERCK KGAA	PL	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor IC 2,5 mg comprimidos revestidos por película	SE/H/0184/002	3031283	MERCK, S.A.	PT	Full application (Article 8(3) of Directive No 2001/83/EC)

Product full name (in authorisation country)	**MRP / DCP Authorisation number**	**National Authorisation Number**	**MAH of product in the member state**	**Member State where product is authorised**	**Legal basis**
Concor 10 mg comprimido revestido	not available	8776468	MERCK, S.A.	PT	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor 10 mg comprimido revestido	not available	8776450	MERCK, S.A.	PT	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor 5 mg comprimido revestido	not available	8776476	MERCK, S.A.	PT	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor 5 mg comprimido revestido	not available	8776435	MERCK, S.A.	PT	Full application (Article 8(3) of Directive No 2001/83/EC)
CONCOR COR 10 mg comprimate filmate	not available	6096/2014/03	MERCK KGAA	RO	Full application (Article 8(3) of Directive No 2001/83/EC)
CONCOR COR 10 mg comprimate filmate	not available	6096/2014/04	MERCK KGAA	RO	Full application (Article 8(3) of Directive No 2001/83/EC)

Product full name (in authorisation country)	**MRP / DCP Authorisation number**	**National Authorisation Number**	**MAH of product in the member state**	**Member State where product is authorised**	**Legal basis**
CONCOR COR 10 mg comprimate filmate	not available	6096/2014/01	MERCK KGAA	RO	Full application (Article 8(3) of Directive No 2001/83/EC)
CONCOR COR 10 mg comprimate filmate	not available	6096/2014/02	MERCK KGAA	RO	Full application (Article 8(3) of Directive No 2001/83/EC)
CONCOR COR 2,5 mg comprimate filmate	not available	6094/2014/03	MERCK KGAA	RO	Full application (Article 8(3) of Directive No 2001/83/EC)
CONCOR COR 2,5 mg comprimate filmate	not available	6094/2014/02	MERCK KGAA	RO	Full application (Article 8(3) of Directive No 2001/83/EC)
CONCOR COR 2,5 mg comprimate filmate	not available	6094/2014/01	MERCK KGAA	RO	Full application (Article 8(3) of Directive No 2001/83/EC)
CONCOR COR 2,5 mg comprimate filmate	not available	6094/2014/04	MERCK KGAA	RO	Full application (Article 8(3) of Directive No 2001/83/EC)

Product full name (in authorisation country)	**MRP / DCP Authorisation number**	**National Authorisation Number**	**MAH of product in the member state**	**Member State where product is authorised**	**Legal basis**
CONCOR COR 5 mg comprimate filmate	not available	6095/2014/01	MERCK KGAA	RO	Full application (Article 8(3) of Directive No 2001/83/EC)
CONCOR COR 5 mg comprimate filmate	not available	6095/2014/04	MERCK KGAA	RO	Full application (Article 8(3) of Directive No 2001/83/EC)
CONCOR COR 5 mg comprimate filmate	not available	6095/2014/03	MERCK KGAA	RO	Full application (Article 8(3) of Directive No 2001/83/EC)
CONCOR COR 5 mg comprimate filmate	not available	6095/2014/02	MERCK KGAA	RO	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor 10 mg comprimate filmate	not available	6252/2014/01	MERCK KGAA	RO	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor 5 mg comprimate filmate	not available	6251/2014/01	MERCK KGAA	RO	Full application (Article 8(3) of Directive No 2001/83/EC)

Product full name (in authorisation country)	**MRP / DCP Authorisation number**	**National Authorisation Number**	**MAH of product in the member state**	**Member State where product is authorised**	**Legal basis**
CARDENSIEL 1,25 mg, comprimé pelliculé	SE/H/0184/001	24 901	MERCK SANTÉ S.A.S.	FR	Full application (Article 8(3) of Directive No 2001/83/EC)
CARDENSIEL 10 mg, comprimé pelliculé sécable	SE/H/0184/006	24 906	MERCK SANTÉ S.A.S.	FR	Full application (Article 8(3) of Directive No 2001/83/EC)
CARDENSIEL 2,5 mg, comprimé pelliculé sécable	SE/H/0184/002	24902	MERCK SANTÉ S.A.S.	FR	Full application (Article 8(3) of Directive No 2001/83/EC)
CARDENSIEL 3,75 mg, comprimé pelliculé sécable	SE/H/0184/003	24 903	MERCK SANTÉ S.A.S.	FR	Full application (Article 8(3) of Directive No 2001/83/EC)
CARDENSIEL 5 mg, comprimé pelliculé sécable	SE/H/0184/004	24 904	MERCK SANTÉ S.A.S.	FR	Full application (Article 8(3) of Directive No 2001/83/EC)
CARDENSIEL 7,5 mg, comprimé pelliculé sécable	SE/H/0184/005	24 905	MERCK SANTÉ S.A.S.	FR	Full application (Article 8(3) of Directive No 2001/83/EC)

Product full name (in authorisation country)	**MRP / DCP Authorisation number**	**National Authorisation Number**	**MAH of product in the member state**	**Member State where product is authorised**	**Legal basis**
Bisoprolol Merck 1.25 mg film-coated tablets	SE/H/0187/001	15380	MERCK KGAA	SE	Full application (Article 8(3) of Directive No 2001/83/EC)
Libracor 1.25 mg film-coated tablets	SE/H/0186/001	15377	MERCK KGAA	SE	Full application (Article 8(3) of Directive No 2001/83/EC)
Emconcor CHF 1,25 mg filmdragerade tabletter	SE/H/0184/001	15371	MERCK KGAA	SE	Full application (Article 8(3) of Directive No 2001/83/EC)
Cardicor 1.25 mg film-coated tablets	SE/H/0185/001	15374	MERCK KGAA	SE	Full application (Article 8(3) of Directive No 2001/83/EC)
Emconcor CHF 10 mg filmdragerade tabletter	SE/H/0184/006	15231	MERCK KGAA	SE	Full application (Article 8(3) of Directive No 2001/83/EC)
Bisoprolol Merck 10 mg film-coated tablets	SE/H/0187/006	15240	MERCK KGAA	SE	Full application (Article 8(3) of Directive No 2001/83/EC)

Product full name (in authorisation country)	**MRP / DCP Authorisation number**	**National Authorisation Number**	**MAH of product in the member state**	**Member State where product is authorised**	**Legal basis**
Libracor 10 mg film-coated tablets	SE/H/0186/006	15237	MERCK KGAA	SE	Full application (Article 8(3) of Directive No 2001/83/EC)
Cardicor 10 mg film-coated tablets	SE/H/0185/006	15234	MERCK KGAA	SE	Full application (Article 8(3) of Directive No 2001/83/EC)
Emconcor CHF 2,5 mg filmdragerade tabletter	SE/H/0184/002	15229	MERCK KGAA	SE	Full application (Article 8(3) of Directive No 2001/83/EC)
Bisoprolol Merck 2.5 mg film-coated tablets	SE/H/0187/002	15238	MERCK KGAA	SE	Full application (Article 8(3) of Directive No 2001/83/EC)
Cardicor 5 mg film-coated tablets	SE/H/0185/004	15376	MERCK KGAA	SE	Full application (Article 8(3) of Directive No 2001/83/EC)
Libracor 2.5 mg film-coated tablets	SE/H/0186/002	15235	MERCK KGAA	SE	Full application (Article 8(3) of Directive No 2001/83/EC)

Product full name (in authorisation country)	**MRP / DCP Authorisation number**	**National Authorisation Number**	**MAH of product in the member state**	**Member State where product is authorised**	**Legal basis**
Cardicor 3.75 mg film-coated tablets	SE/H/0185/003	15375	MERCK KGAA	SE	Full application (Article 8(3) of Directive No 2001/83/EC)
Emconcor CHF 3,75 mg filmdragerade tabletter	SE/H/0184/003	15372	MERCK KGAA	SE	Full application (Article 8(3) of Directive No 2001/83/EC)
Libracor 3.75 mg film-coated tablets	SE/H/0186/003	15378	MERCK KGAA	SE	Full application (Article 8(3) of Directive No 2001/83/EC)
Emconcor CHF 5 mg filmdragerade tabletter	SE/H/0184/004	15373	MERCK KGAA	SE	Full application (Article 8(3) of Directive No 2001/83/EC)
Libracor 5 mg film-coated tablets	SE/H/0186/004	15379	MERCK KGAA	SE	Full application (Article 8(3) of Directive No 2001/83/EC)
Libracor 7.5 mg film-coated tablets	SE/H/0186/005	15236	MERCK KGAA	SE	Full application (Article 8(3) of Directive No 2001/83/EC)

Product full name (in authorisation country)	**MRP / DCP Authorisation number**	**National Authorisation Number**	**MAH of product in the member state**	**Member State where product is authorised**	**Legal basis**
Emconcor CHF 7,5 mg filmdragerade tabletter	SE/H/0184/005	15230	MERCK KGAA	SE	Full application (Article 8(3) of Directive No 2001/83/EC)
Cardicor 7.5 mg film-coated tablets	SE/H/0185/005	15233	MERCK KGAA	SE	Full application (Article 8(3) of Directive No 2001/83/EC)
Bisoprolol Merck 7.5 mg film-coated tablets	SE/H/0187/005	15239	MERCK KGAA	SE	Full application (Article 8(3) of Directive No 2001/83/EC)
Emconcor 5 mg filmdragerade tabletter	not available	11202	MERCK KGAA	SE	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor COR 10 mg	not available	41/0012/02-S	MERCK KGAA	SK	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor COR 5 mg	not available	41/0011/02-S	MERCK KGAA	SK	Full application (Article 8(3) of Directive No 2001/83/EC)

Product full name (in authorisation country)	**MRP / DCP Authorisation number**	**National Authorisation Number**	**MAH of product in the member state**	**Member State where product is authorised**	**Legal basis**
Concor 10	not available	41/0304/89-S	MERCK KGAA	SK	Full application (Article 8(3) of Directive No 2001/83/EC)
Bisomerck 10	not available	41/0055/03-S	MERCK KGAA	SK	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor 5	not available	41/0304/89-S	MERCK KGAA	SK	Full application (Article 8(3) of Directive No 2001/83/EC)
Bisomerck 5	not available	41/0054/03-S	MERCK KGAA	SK	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor COR 2,5 mg	not available	41/0010/02-S	MERCK KGAA	SK	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor COR 1,25 mg filmsko obložene tablete	not available	5363-I-555/12	MERCK D.O.O.	SI	Full application (Article 8(3) of Directive No 2001/83/EC)

Product full name (in authorisation country)	**MRP / DCP Authorisation number**	**National Authorisation Number**	**MAH of product in the member state**	**Member State where product is authorised**	**Legal basis**
Concor COR 10 mg filmsko obložene tablete	not available	5363-I-1803/13	MERCK D.O.O.	SI	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor COR 2,5 mg filmsko obložene tablete	not available	5363-I-26/13	MERCK D.O.O.	SI	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor COR 5 mg filmsko obložene tablete	not available	5363-I-558/12	MERCK D.O.O.	SI	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor COR 7,5 mg filmsko obložene tablete	not available	5363-I-1185/12	MERCK D.O.O.	SI	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor 10 mg filmsko obložene tablete	not available	5363-I-554/12	MERCK D.O.O.	SI	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor 5 mg filmsko obložene tablete	not available	5363-I-27/13	MERCK D.O.O.	SI	Full application (Article 8(3) of Directive No 2001/83/EC)

Product full name (in authorisation country)	**MRP / DCP Authorisation number**	**National Authorisation Number**	**MAH of product in the member state**	**Member State where product is authorised**	**Legal basis**
CONGESCOR 1,25 mg, compresse film-rivestite	SE/H/0186/001	034953024/M	DAIICHI SANKYO ITALIA S.P.A	IT	Informed consent application (Article 10c of Directive No 2001/83/EC)
CONGESCOR 2,5 mg, compresse film-rivestite	SE/H/0186/002	034953099/M	DAIICHI SANKYO ITALIA S.P.A	IT	Informed consent application (Article 10c of Directive No 2001/83/EC)
CONGESCOR 3,75 mg, compresse film-rivestite	SE/H/0186/003	034953176/M	DAIICHI SANKYO ITALIA S.P.A	IT	Informed consent application (Article 10c of Directive No 2001/83/EC)
CONGESCOR 5 mg, compresse film-rivestite	SE/H/0186/004	034953253/M	DAIICHI SANKYO ITALIA S.P.A	IT	Informed consent application (Article 10c of Directive No 2001/83/EC)

Product full name (in authorisation country)	**MRP / DCP Authorisation number**	**National Authorisation Number**	**MAH of product in the member state**	**Member State where product is authorised**	**Legal basis**
CONGESCOR 7,5 mg, compresse film-rivestite	SE/H/0186/005	034953339/M	DAIICHI SANKYO ITALIA S.P.A	IT	Informed consent application (Article 10c of Directive No 2001/83/EC)
CONGESCOR 10 mg, compresse film-rivestite	SE/H/0186/006	034953416/M	DAIICHI SANKYO ITALIA S.P.A	IT	Informed consent application (Article 10c of Directive No 2001/83/EC)
Concor Cor 1,25 mg Filmtabletten	SE/H/0184/001	1-23302	MERCK GESELLSCHAFT MBH	AT	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor Cor 10 mg Filmtabletten	SE/H/0184/006	1-23307	MERCK GESELLSCHAFT MBH	AT	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor Cor 2,5 mg Filmtabletten	SE/H/0184/002	1-23303	MERCK GESELLSCHAFT MBH	AT	Full application (Article 8(3) of Directive No 2001/83/EC)

Product full name (in authorisation country)	**MRP / DCP Authorisation number**	**National Authorisation Number**	**MAH of product in the member state**	**Member State where product is authorised**	**Legal basis**
Concor Cor 3,75 mg Filmtabletten	SE/H/0184/003	1-23304	MERCK GESELLSCHAFT MBH	AT	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor Cor 5 mg Filmtabletten	SE/H/0184/004	1-23305	MERCK GESELLSCHAFT MBH	AT	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor Cor 7,5 mg Filmtabletten	SE/H/0184/005	1-23306	MERCK GESELLSCHAFT MBH	AT	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor 10 mg - Filmtabletten	not available	1-18586	MERCK GESELLSCHAFT MBH	AT	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor 5 mg - Filmtabletten	not available	1-18587	MERCK GESELLSCHAFT MBH	AT	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor COR 1,25 mg Filmtabletten	SE/H/0184/001	46660.00.00	MERCK SERONO GMBH	DE	Full application (Article 8(3) of Directive No 2001/83/EC)

Product full name (in authorisation country)	**MRP / DCP Authorisation number**	**National Authorisation Number**	**MAH of product in the member state**	**Member State where product is authorised**	**Legal basis**
Concor COR 10 mg Filmtabletten	SE/H/0184/006	46660.00.05	MERCK SERONO GMBH	DE	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor COR 2,5 mg Filmtabletten	SE/H/0184/002	46660.00.01	MERCK SERONO GMBH	DE	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor COR 3,75 mg Filmtabletten	SE/H/0184/003	46660.00.02	MERCK SERONO GMBH	DE	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor COR 5 mg Filmtabletten	SE/H/0184/004	46660.00.03	MERCK SERONO GMBH	DE	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor COR 7,5 mg Filmtabletten	SE/H/0184/005	46660.00.04	MERCK SERONO GMBH	DE	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor 5 mg Filmtabletten	not available	6849.00.00	MERCK SERONO GMBH	DE	Full application (Article 8(3) of Directive No 2001/83/EC)

Product full name (in authorisation country)	**MRP / DCP Authorisation number**	**National Authorisation Number**	**MAH of product in the member state**	**Member State where product is authorised**	**Legal basis**
Cardicor 2,5 mg compresse rivestite con film	SE/H/0185/002	034954091/M	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT	Full application (Article 8(3) of Directive No 2001/83/EC)
Cardicor 3,75 mg compresse rivestite con film	SE/H/0185/003	034954175/M	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT	Full application (Article 8(3) of Directive No 2001/83/EC)
Cardicor 5 mg compresse rivestite con film	SE/H/0185/004	034954255	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT	Full application (Article 8(3) of Directive No 2001/83/EC)
Cardicor 7,5 mg compresse rivestite con film	SE/H/0185/005	034954331	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT	Full application (Article 8(3) of Directive No 2001/83/EC)
Emconcor Mitis 5 mg filmomhulde tabletten	not available	BE155346	MERCK N.V.-S.A.	BE	Full application (Article 8(3) of Directive No 2001/83/EC)

Product full name (in authorisation country)	**MRP / DCP Authorisation number**	**National Authorisation Number**	**MAH of product in the member state**	**Member State where product is authorised**	**Legal basis**
Sequacor 1,25 mg compresse rivestite con film	SE/H/184/001-006	034952010/M	BRACCO SPA	IT	Full application (Article 8(3) of Directive No 2001/83/EC)
Sequacor 2,5 mg compresse rivestite con film	SE/H/184/001-006	034952097/M	BRACCO SPA	IT	Full application (Article 8(3) of Directive No 2001/83/EC)
Sequacor 3.75 mg compresse rivestite con film	SE/H/184/001-006	034952174/M	BRACCO SPA	IT	Full application (Article 8(3) of Directive No 2001/83/EC)
Sequacor 5 mg compresse rivestite con film	SE/H/184/001-006	034952251/M	BRACCO SPA	IT	Full application (Article 8(3) of Directive No 2001/83/EC)
Sequacor 7,5 mg compresse rivestite con film	SE/H/184/001-006	034952337/M	BRACCO SPA	IT	Full application (Article 8(3) of Directive No 2001/83/EC)
Sequacor 10 mg compresse rivestite con film	SE/H/184/001-006	034952414/M	BRACCO SPA	IT	Full application (Article 8(3) of Directive No 2001/83/EC)

Product full name (in authorisation country)	**MRP / DCP Authorisation number**	**National Authorisation Number**	**MAH of product in the member state**	**Member State where product is authorised**	**Legal basis**
Isoten 5 mg filmomhulde tabletten	SE/H/185/04	BE 210314	S.A. MEDA PHARMA N.V.	BE	Full application (Article 8(3) of Directive No 2001/83/EC)
Isoten 10 mg filmomhulde tabletten	SE/H/185/06	BE 210332	S.A. MEDA PHARMA N.V.	BE	Full application (Article 8(3) of Directive No 2001/83/EC)
CARDIOCOR 1,25 mg, comprimé pelliculé	SE/H/185/001	353 152- 5	MEDA PHARMA SAS	FR	Full application (Article 8(3) of Directive No 2001/83/EC)
CARDIOCOR 2,5 mg, comprimé pelliculé sécable	SE/H/185/002	353 154-8	MEDA PHARMA SAS	FR	Full application (Article 8(3) of Directive No 2001/83/EC)
CARDIOCOR 5 mg, comprimé pelliculé sécable	SE/H/185/004	353 158-3	MEDA PHARMA SAS	FR	Full application (Article 8(3) of Directive No 2001/83/EC)

Product full name (in authorisation country)	**MRP / DCP Authorisation number**	**National Authorisation Number**	**MAH of product in the member state**	**Member State where product is authorised**	**Legal basis**
BISOPROLOL BIOGARAN® 10 mg, comprimé pelliculé sécable	not available	27752	BIOGARAN	FR	Informed consent application (Article 10c of Directive No 2001/83/EC)
Bisoprolol Apotex 5 mg filmomhulde tabletten	not available	BE218626	APOTEX NV	BE	
Bisoprolol Apotex 10 mg filmomhulde tabletten	not available	BE218635	APOTEX NV	BE	
Bisoprolol 'Arcana' 5 mg - Filmtabletten		1-22024	Arcana Arzneimittel GmbH	AT	Full application (Article 8(3) of Directive No 2001/83/EC)
Bisoprolol 'Arcana' 10 mg - Filmtabletten		1-22023	Arcana Arzneimittel GmbH	AT	Full application (Article 8(3) of Directive No 2001/83/EC)
CONCOR 10mg compresse		026573016	Bracco SpA	IT	Full application (Article 8(3) of Directive No 2001/83/EC)

Product full name (in authorisation country)	**MRP / DCP Authorisation number**	**National Authorisation Number**	**MAH of product in the member state**	**Member State where product is authorised**	**Legal basis**
Bisoprolol AUROBINDO 5 mg, comprimé pelliculé sécable	UK/H/1684/001	223 270-8	AUROBINDO PHARMA FRANCE SARL	FR	Hybrid application (Article 10(3) of Directive No 2001/83/EC)
Bisoprolol AUROBINDO 5 mg, comprimé pelliculé sécable	UK/H/1684/001	223 271-4	AUROBINDO PHARMA FRANCE SARL	FR	Hybrid application (Article 10(3) of Directive No 2001/83/EC)
Bisoprolol AUROBINDO 5 mg, comprimé pelliculé sécable	UK/H/1684/001	223 272-0	AUROBINDO PHARMA FRANCE SARL	FR	Hybrid application (Article 10(3) of Directive No 2001/83/EC)
Bisoprolol AUROBINDO 5 mg, comprimé pelliculé sécable	UK/H/1684/001	223 273-7	AUROBINDO PHARMA FRANCE SARL	FR	Hybrid application (Article 10(3) of Directive No 2001/83/EC)
Bisoprolol AUROBINDO 5 mg, comprimé pelliculé sécable	UK/H/1684/001	223 274-3	AUROBINDO PHARMA FRANCE SARL	FR	Hybrid application (Article 10(3) of Directive No 2001/83/EC)

Product full name (in authorisation country)	**MRP / DCP Authorisation number**	**National Authorisation Number**	**MAH of product in the member state**	**Member State where product is authorised**	**Legal basis**
Bisoprolol AUROBINDO 5 mg, comprimé pelliculé sécable	UK/H/1684/001	223 276-6	AUROBINDO PHARMA FRANCE SARL	FR	Hybrid application (Article 10(3) of Directive No 2001/83/EC)
Concor COR	SE/H/0184/002	2005068803	MERCK SERONO GMBH	LU	Full application (Article 8(3) of Directive No 2001/83/EC)
Isoten 2,5 mg	SE/H/0185/002	BE210196	Meda PHARMA, Belgium	BE	Full application (Article 8(3) of Directive No 2001/83/EC)
Cardicor 1,25 mg compresse rivestite con film	SE/H/0185/001	034954014/M	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT	Full application (Article 8(3) of Directive No 2001/83/EC)
Bisprololfumaraat Deco Mylan 2,5 mg	SE/H/0185/002	RVG 24509	Mylan B.V. Netherlands	NL	Full application (Article 8(3) of Directive No 2001/83/EC)

Product full name (in authorisation country)	**MRP / DCP Authorisation number**	**National Authorisation Number**	**MAH of product in the member state**	**Member State where product is authorised**	**Legal basis**
Bisprololfumaraat Deco Mylan 10 mg	SE/H/0185/006	RVG 24513	Mylan B.V. Netherlands	NL	Full application (Article 8(3) of Directive No 2001/83/EC)
Bisoce, 1,25 mg	SE/H/0186/001	353 139-9, 353 140-7, 353 541-1	Mylan, France	FR	Full application (Article 8(3) of Directive No 2001/83/EC)
Bisoce, 2,5 mg	SE/H/0186/002	353 542-8	Mylan, France	FR	Full application (Article 8(3) of Directive No 2001/83/EC)
Bisoce, 3,75 mg	SE/H/0186/003	353 144-2, 353 145-9, 353 543-4	Mylan, France	FR	Full application (Article 8(3) of Directive No 2001/83/EC)
Bisoce, 5mg	SE/H/0186/004	353 544-0, 353 146-5	Mylan, France	FR	Full application (Article 8(3) of Directive No 2001/83/EC)
Bisoce, 7,5 mg	SE/H/0186/005	353 545-7, 353 148-8	Mylan, France	FR	Full application (Article 8(3) of Directive No 2001/83/EC)

Product full name (in authorisation country)	**MRP / DCP Authorisation number**	**National Authorisation Number**	**MAH of product in the member state**	**Member State where product is authorised**	**Legal basis**
Bisoce, 10mg	SE/H/0186/006	353 546-3, 353 150-2	Mylan, France	FR	
Emconcor 2,5mg	SE/H/0187/002	22172	CG Papaloisou	CY	Full application (Article 8(3) of Directive No 2001/83/EC)
Emconcor 5 mg	SE/H/0187/004	22173	CG Papaloisou	CY	Full application (Article 8(3) of Directive No 2001/83/EC)
BISOPROLOL BGR 1,25 mg comprimé pelliculé sécable	SE/H/187/01	353 548-6, 352 954-0	BIOGARAN	FR	Informed consent application (Article 10c of Directive No 2001/83/EC)
BISOPROLOL BGR 2,5 mg comprimé pelliculé sécable	SE/H/0187/002	353 549-2, 352 956-3	BIOGARAN	FR	Informed consent application (Article 10c of Directive No 2001/83/EC)

Product full name (in authorisation country)	**MRP / DCP Authorisation number**	**National Authorisation Number**	**MAH of product in the member state**	**Member State where product is authorised**	**Legal basis**
BISOPROLOL BGR 1,25 mg comprimé pelliculé	SE/H/187/04	NL 24922	BIOGARAN	FR	Informed consent application (Article 10c of Directive No 2001/83/EC)
BISOPROLOL BGR 7,5 mg comprimé pelliculé sécable	SE/H/187/05	353 552-3, 352 964-6	BIOGARAN	FR	Informed consent application (Article 10c of Directive No 2001/83/EC)
BISOPROLOL BGR 10 mg comprimé pelliculé sécable	SE/H/187/05	NL 24924	BIOGARAN	FR	Informed consent application (Article 10c of Directive No 2001/83/EC)
DETENSIEL 1		14214	MERCK SANTÉ S.A.S.	FR	Full application (Article 8(3) of Directive No 2001/83/EC)

Product full name (in authorisation country)	**MRP / DCP Authorisation number**	**National Authorisation Number**	**MAH of product in the member state**	**Member State where product is authorised**	**Legal basis**
Bisoprolol Mylan		NL 27 756	Mylan, France	FR	Full application (Article 8(3) of Directive No 2001/83/EC)
Bisoprolol Zentiva 10 mg		361252-5, 361253-1, 564665-2, 372049-1	Sanofi-Aventis, France	FR	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor COR 1,25 mg filmtabletta	not available	OGYI-T-8324/14	MERCK KFT.	HU	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor COR 2,5 mg filmtabletta	not available	OGYI-T-8324/13	MERCK KFT.	HU	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor COR 10 mg filmtabletta	not available	OGYI-T-8324/16	MERCK KFT.	HU	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor 10 mg filmtabletta	not available	OGYI-T-4015/09	MERCK KFT.	HU	Full application (Article 8(3) of Directive No 2001/83/EC)

Product full name (in authorisation country)	**MRP / DCP Authorisation number**	**National Authorisation Number**	**MAH of product in the member state**	**Member State where product is authorised**	**Legal basis**
Concor 10 mg filmtabletta	not available	OGYI-T-4015/10	MERCK KFT.	HU	Full application (Article 8(3) of Directive No 2001/83/EC)
Emcolol 5mg		PA 577/38/1	MERCK SERONO LTD.	IE	Full application (Article 8(3) of Directive No 2001/83/EC)
Emcolol 10 mg		PA 577/38/2	MERCK SERONO LTD.	IE	Full application (Article 8(3) of Directive No 2001/83/EC)
Bisoprolol 5mg		RVG 25233	Mylan B.V. Netherlands	NL	Full application (Article 8(3) of Directive No 2001/83/EC)
Bisoprolol 10 mg		RVG 12409	Mylan B.V. Netherlands	NL	Full application (Article 8(3) of Directive No 2001/83/EC)
Concor 5 mg	not available	1/11/86	MERCK, S.A.	PT	Full application (Article 8(3) of Directive No 2001/83/EC)

Product full name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised	Legal basis
Concor 10 mg	not available	1/11/86	MERCK, S.A.	PT	Full application (Article 8(3) of Directive No 2001/83/EC)
Bisoprololo Zentiva	IT/H/0173/001/MRP	037690056 037690068 037690017 037690070 037690029 037690031 037690195 037690082 037690043 037690132 037690144 037690094 037690157 037690106 037690118 037690169 037690120 037690171 037690183	Zentiva Italia SRL	IT	Hybrid application (Article 10(3) of Directive No 2001/83/EC)

Product full name (in authorisation country)	**MRP / DCP Authorisation number**	**National Authorisation Number**	**MAH of product in the member state**	**Member State where product is authorised**	**Legal basis**
Cardicor 7.5mg Film-coated Tablets	SE/H/0184/001/IA/048	PL 11648/0075	MERCK SERONO LTD.	UK	Full application (Article 8(3) of Directive No 2001/83/EC)
Cardicor 10 mg film-coated tablets	SE/H/0185/006	034953471/M	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT	Full application (Article 8(3) of Directive No 2001/83/EC)
Bisoprolol Sandoz 5 mg		0659742, 0659756, 0659773, 0659791, 0659806, 0659823, 0659837, 0659841, 0659854, 0659868, 0659871, 0659885, 0659899, 0660144, 0660158, 0660161, 0660175, 0660189, 0660192, 0660208, 2012090024	SANDOZ N.V.	LU	Hybrid application (Article 10(3) of Directive No 2001/83/EC)

Product full name (in authorisation country)	**MRP / DCP Authorisation number**	**National Authorisation Number**	**MAH of product in the member state**	**Member State where product is authorised**	**Legal basis**
Bisoprolol Sandoz 10 mg		0659904, 0659918, 0659921, 0659935, 0659949, 0659952, 0659966, 0659983, 0659997, 0660001, 0660015, 0660029, 0660032, 0660046, 0660063, 0660077, 0660081, 0660094, 0660113, 0660127, 0660131, 2012090025	SANDOZ N.V.	LU	Hybrid application (Article 10(3) of Directive No 2001/83/EC)