13 December 2017
EMA/46974/2018
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance(s): simvastatin

Procedure No.: PSUSA/00002709/201704

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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
Belmalip 10 mg comprimidos recubiertos con película	not available	64.652	TEVA PHARMA S.L.U	ES
Belmalip 10 mg comprimidos recubiertos con película	not available	64.652	TEVA PHARMA S.L.U	ES
Belmalip 20 mg comprimidos recubiertos con película.	not available	64.653	TEVA PHARMA S.L.U	ES
Belmalip 20 mg comprimidos recubiertos con película.	not available	64.653	TEVA PHARMA S.L.U	ES
Belmalip 40 mg comprimidos recubiertos con película.	not available	64.654	TEVA PHARMA S.L.U	ES
Belmalip 40 mg comprimidos recubiertos con película.	not available	64.654	TEVA PHARMA S.L.U	ES
COLEMIN 10 mg comprimidos	not available	59.181	VIFOR PHARMA ESPA&#xFFFD;A, S.L.	ES
COLEMIN 10 mg comprimidos	not available	59.181	VIFOR PHARMA ESPA&#xFFFD;A, S.L.	ES
COLEMIN 20 mg comprimidos	not available	59.180	VIFOR PHARMA ESPA&#xFFFD;A, S.L.	ES
COLEMIN 20 mg comprimidos	not available	59.180	VIFOR PHARMA ESPA&#xFFFD;A, S.L.	ES
COLEMIN FORTE 40 mg comprimidos	not available	62329	VIFOR PHARMA ESPA&#xFFFD;A, S.L.	ES
COLEMIN FORTE 40 mg comprimidos	not available	62329	VIFOR PHARMA ESPA&#xFFFD;A, S.L.	ES
Jabastatina 10 mg comprimidos revestidos por pelテュcula	not available	3487592	JABA RECORDATI, S.A.	PT
Jabastatina 10 mg comprimidos revestidos	not available	3487691	JABA RECORDATI, 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 pelテュcula				
Jabastatina 10 mg comprimidos revestidos por pelテュcula	not available	3487493	JABA RECORDATI, S.A.	PT
Jabastatina 10 mg comprimidos revestidos por pelテュcula	not available	3487592	JABA RECORDATI, S.A.	PT
Jabastatina 10 mg comprimidos revestidos por pelテュcula	not available	3487691	JABA RECORDATI, S.A.	PT
Jabastatina 10 mg comprimidos revestidos por pelテュcula	not available	3487493	JABA RECORDATI, S.A.	PT
Jabastatina 20 mg comprimidos revestidos por pelテュcula	not available	3487998	JABA RECORDATI, S.A.	PT
Jabastatina 20 mg comprimidos revestidos por pelテュcula	not available	3487790	JABA RECORDATI, S.A.	PT
Jabastatina 20 mg comprimidos revestidos por pelテュcula	not available	3487899	JABA RECORDATI, S.A.	PT
Jabastatina 20 mg comprimidos revestidos por pelテュcula	not available	3487998	JABA RECORDATI, S.A.	PT
Jabastatina 20 mg comprimidos revestidos por pelテュcula	not available	3487790	JABA RECORDATI, S.A.	PT
Jabastatina 20 mg comprimidos revestidos por pelテュcula	not available	3487899	JABA RECORDATI, S.A.	PT
Jabastatina 40 mg comprimidos revestidos	not available	4217097	JABA RECORDATI, 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 pel□cula				
Jabastatina 40 mg comprimidos revestidos por pel□cula	not available	4217196	JABA RECORDATI, S.A.	PT
Jabastatina 40 mg comprimidos revestidos por pel□cula	not available	4217097	JABA RECORDATI, S.A.	PT
Jabastatina 40 mg comprimidos revestidos por pel□cula	not available	4217196	JABA RECORDATI, S.A.	PT
Lipcut 30 mg tabletti, kalvopäällysteinen	FI/H/0141/004	16015	SANDOZ A/S	FI
Lipcut 30 mg tabletti, kalvopäällysteinen	FI/H/0141/004	16015	SANDOZ A/S	FI
Lipcut 60 mg tabletti, kalvopäällysteinen	FI/H/0141/007	18449	SANDOZ A/S	FI
Lipcut 60 mg tabletti, kalvopäällysteinen	FI/H/0141/007	18449	SANDOZ A/S	FI
LIPEX® 10 mg filmom obložene tablete	not available	HR-H-509178324	MERCK SHARP & DOHME D.O.O.	HR
LIPEX® 10 mg filmom obložene tablete	not available	HR-H-509178324	MERCK SHARP & DOHME D.O.O.	HR
LIPEX® 20 mg filmom obložene tablete	not available	HR-H-604977393	MERCK SHARP & DOHME D.O.O.	HR
LIPEX® 20 mg filmom obložene tablete	not available	HR-H-604977393	MERCK SHARP & DOHME D.O.O.	HR
LIPEX® 40 mg filmom obložene tablete	not available	HR-H-597927430	MERCK SHARP & DOHME D.O.O.	HR
LIPEX® 40 mg filmom obložene tablete	not available	HR-H-597927430	MERCK SHARP & DOHME D.O.O.	HR
LIPEX® 80 mg filmom obložene tablete	not available	HR-H-164284956	MERCK SHARP & DOHME D.O.O.	HR
LIPEX® 80 mg filmom obložene tablete	not available	HR-H-164284956	MERCK SHARP & DOHME 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
Liponorm 10 mg, compresse rivestite con film	not available	027228016	NEOPHARMED GENTILI S.R.L.	IT
Liponorm 10 mg, compresse rivestite con film	not available	027228016	NEOPHARMED GENTILI S.R.L.	IT
Liponorm 10 mg, compresse rivestite con film	not available	027228016	NEOPHARMED GENTILI S.R.L.	IT
Liponorm 20 mg, compresse rivestite con film	not available	027228028	NEOPHARMED GENTILI S.R.L.	IT
Liponorm 20 mg, compresse rivestite con film	not available	027228079	NEOPHARMED GENTILI S.R.L.	IT
Liponorm 20 mg, compresse rivestite con film	not available	027228028	NEOPHARMED GENTILI S.R.L.	IT
Liponorm 20 mg, compresse rivestite con film	not available	027228079	NEOPHARMED GENTILI S.R.L.	IT
Liponorm 20 mg, compresse rivestite con film	not available	027228028	NEOPHARMED GENTILI S.R.L.	IT
Liponorm 20 mg, compresse rivestite con film	not available	027228079	NEOPHARMED GENTILI S.R.L.	IT
Liponorm 40 mg, compresse rivestite con film	not available	027228030	NEOPHARMED GENTILI S.R.L.	IT
Liponorm 40 mg, compresse rivestite con film	not available	027228081	NEOPHARMED GENTILI S.R.L.	IT
Liponorm 40 mg,	not available	027228030	NEOPHARMED GENTILI	IT

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compresse rivestite con film			S.R.L.	
Liponorm 40 mg, compresse rivestite con film	not available	027228081	NEOPHARMED GENTILI S.R.L.	IT
Liponorm 40 mg, compresse rivestite con film	not available	027228030	NEOPHARMED GENTILI S.R.L.	IT
Liponorm 40 mg, compresse rivestite con film	not available	027228081	NEOPHARMED GENTILI S.R.L.	IT
Medipo 10 mg, compresse rivestite con film	not available	028370017	MEDIOLANUM FARMACEUTICI SPA	IT
Medipo 10 mg, compresse rivestite con film	not available	028370017	MEDIOLANUM FARMACEUTICI SPA	IT
Medipo 20 mg, compresse rivestite con film	not available	028370029	MEDIOLANUM FARMACEUTICI SPA	IT
Medipo 20 mg, compresse rivestite con film	not available	028370106	MEDIOLANUM FARMACEUTICI SPA	IT
Medipo 20 mg, compresse rivestite con film	not available	028370029	MEDIOLANUM FARMACEUTICI SPA	IT
Medipo 20 mg, compresse rivestite con film	not available	028370106	MEDIOLANUM FARMACEUTICI SPA	IT
Medipo 40 mg, compresse rivestite con film	not available	028370043	MEDIOLANUM FARMACEUTICI SPA	IT
Medipo 40 mg, compresse rivestite con	not available	028370118	MEDIOLANUM FARMACEUTICI SPA	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				
Medipo 40 mg, compresse rivestite con film	not available	028370043	MEDIOLANUM FARMACEUTICI SPA	IT
Medipo 40 mg, compresse rivestite con film	not available	028370118	MEDIOLANUM FARMACEUTICI SPA	IT
PANTOK 10 mg comprimidos recubiertos con pelicula	not available	59.182	LACER S.A.	ES
PANTOK 10 mg comprimidos recubiertos con pelicula	not available	59.182	LACER S.A.	ES
PANTOK 20 mg comprimidos recubiertos con pelicula	not available	59.183	LACER S.A.	ES
PANTOK 20 mg comprimidos recubiertos con pelicula	not available	59.183	LACER S.A.	ES
PANTOK FORTE 40 mg comprimidos recubiertos con pelicula	not available	62.330	LACER S.A.	ES
PANTOK FORTE 40 mg comprimidos recubiertos con pelicula	not available	62.330	LACER S.A.	ES
SIMVA BASICS 30 mg Filmtabletten	DE/H/3518/001	86718.00.00	BASICS GMBH	DE
SIMVA BASICS 30 mg Filmtabletten	DE/H/3518/001	86718.00.00	BASICS GMBH	DE
SIMVA BASICS 60 mg Filmtabletten	DE/H/3518/002	86719.00.00	BASICS GMBH	DE
SIMVA BASICS 60 mg Filmtabletten	DE/H/3518/002	86719.00.00	BASICS GMBH	DE
Simva TAD® 30 mg	not available	77051.00.00	TAD 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
Filmtabletten				
Simva TAD® 30 mg Filmtabletten	not available	77051.00.00	TAD PHARMA GMBH	DE
Simva TAD® 60 mg Filmtabletten	not available	77052.00.00	TAD PHARMA GMBH	DE
Simva TAD® 60 mg Filmtabletten	not available	77052.00.00	TAD PHARMA GMBH	DE
Simvastatin - 1 A Pharma 60 mg Filmtabletten	DE/H/2046/007	62540.00.00	1 A PHARMA GMBH	DE
Simvastatin - 1 A Pharma 60 mg Filmtabletten	DE/H/2046/007	62540.00.00	1 A PHARMA GMBH	DE
Simvastatin 10 mg film-coated tablets	not available	PL 21880/0064	MEDREICH PLC	UK
Simvastatin 10 mg film-coated tablets	not available	PL 11311/0398	TILLOMED LABORATORIES LTD	UK
Simvastatin 10 mg film-coated tablets	not available	PL 45994/0001	IMPERIAL WORLDWIDE LIMITED	UK
Simvastatin 10 mg film-coated tablets	not available	PL 45994/0001	IMPERIAL WORLDWIDE LIMITED	UK
Simvastatin 10 mg film-coated tablets	not available	PL 21880/0064	MEDREICH PLC	UK
Simvastatin 10 mg film-coated tablets	not available	PL 11311/0398	TILLOMED LABORATORIES LTD	UK
Simvastatin 10 mg film-coated tablets	not available	PL 45994/0001	IMPERIAL WORLDWIDE LIMITED	UK
Simvastatin 10mg Tablets	not available	PL 17496/0027	DALKEITH LABORATORIES LIMITED	UK
Simvastatin 10mg Tablets	not available	PL 17496/0027	DALKEITH LABORATORIES LIMITED	UK
Simvastatin 20 mg film-coated tablets	not available	PL 21880/0065	MEDREICH PLC	UK
Simvastatin 20 mg film-	not available	PL 11311/0399	TILLOMED LABORATORIES	UK

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			LTD	
Simvastatin 20 mg film-coated tablets	not available	PL 45994/0002	IMPERIAL WORLDWIDE LIMITED	UK
Simvastatin 20 mg film-coated tablets	not available	PL 45994/0002	IMPERIAL WORLDWIDE LIMITED	UK
Simvastatin 20 mg film-coated tablets	not available	PL 21880/0065	MEDREICH PLC	UK
Simvastatin 20 mg film-coated tablets	not available	PL 11311/0399	TILLOMED LABORATORIES LTD	UK
Simvastatin 20 mg film-coated tablets	not available	PL 45994/0002	IMPERIAL WORLDWIDE LIMITED	UK
Simvastatin 40 mg film-coated tablets	not available	PL 21880/0066	MEDREICH PLC	UK
Simvastatin 40 mg film-coated tablets	not available	PL11311/0400	TILLOMED LABORATORIES LTD	UK
Simvastatin 40 mg film-coated tablets	not available	PL 45994/0003	IMPERIAL WORLDWIDE LIMITED	UK
Simvastatin 40 mg film-coated tablets	not available	PL 45994/0003	IMPERIAL WORLDWIDE LIMITED	UK
Simvastatin 40 mg film-coated tablets	not available	PL 21880/0066	MEDREICH PLC	UK
Simvastatin 40 mg film-coated tablets	not available	PL11311/0400	TILLOMED LABORATORIES LTD	UK
Simvastatin 40 mg film-coated tablets	not available	PL 45994/0003	IMPERIAL WORLDWIDE LIMITED	UK
Simvastatin 5 mg film-coated tablets	DE/H/2045/001	PL 04416/0677	SANDOZ LTD	UK
Simvastatin 5 mg film-coated tablets	DE/H/2045/001	PL 04416/0677	SANDOZ LTD	UK
Simvastatin 80 mg film-coated tablets	not available	PL 20416/0362	CRESCENT PHARMA LIMITED	UK
Simvastatin 80 mg film-coated tablets	not available	PL 21880/0067	MEDREICH PLC	UK
Simvastatin 80 mg film-	not available	PL11311/0401	TILLOMED LABORATORIES	UK

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			LTD	
Simvastatin 80 mg film-coated tablets	not available	PL 20416/0362	CRESCENT PHARMA LIMITED	UK
Simvastatin 80 mg film-coated tablets	not available	PL 21880/0067	MEDREICH PLC	UK
Simvastatin 80 mg film-coated tablets	not available	PL11311/0401	TILLOMED LABORATORIES LTD	UK
Simvastatin Aurobindo 40 mg apvalkotās tabletes	NL/H/1311/004	09 - 0221	AUROBINDO PHARMA LIMITED	LV
Simvastatin Aurobindo 40 mg apvalkotās tabletes	NL/H/1311/004	09 - 0221	AUROBINDO PHARMA LIMITED	LV
Simvastatin Aurobindo 80 mg apvalkotās tabletes	NL/H/1311/005	09 - 0222	AUROBINDO PHARMA LIMITED	LV
Simvastatin Aurobindo 80 mg apvalkotās tabletes	NL/H/1311/005	09 - 0222	AUROBINDO PHARMA LIMITED	LV
Simvastatin Hexal 30 mg - Filmtabletten	FI/H/0141/004	1-24282	HEXAL PHARMA GMBH	AT
Simvastatin Hexal 30 mg - Filmtabletten	FI/H/0141/004	1-24282	HEXAL PHARMA GMBH	AT
Simvastatin Sandoz 60 mg Filmtabletten	DE/H/2045/007	52536.06.00	HEXAL AG	DE
Simvastatin Sandoz 60 mg Filmtabletten	DE/H/2045/007	52536.06.00	HEXAL AG	DE
Simvastatin Teva 5 mg filmdragerade tabletter	FR/H/0323/001	25274	TEVA SWEDEN AB	SE
Simvastatin Teva 5 mg filmdragerade tabletter	FR/H/0323/001	25274	TEVA SWEDEN AB	SE
SIMVASTATINA ratiopharm 5 mg Compresse rivestite con	FR/H/0458/001	041511054/M	TEVA 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
film				
SIMVASTATINA ratiopharm 5 mg Compresse rivestite con film	FR/H/0458/001	041511041/M	TEVA ITALIA S.R.L.	IT
SIMVASTATINA ratiopharm 5 mg Compresse rivestite con film	FR/H/0458/001	041511015/M	TEVA ITALIA S.R.L.	IT
SIMVASTATINA ratiopharm 5 mg Compresse rivestite con film	FR/H/0458/001	041511039/M	TEVA ITALIA S.R.L.	IT
SIMVASTATINA ratiopharm 5 mg Compresse rivestite con film	FR/H/0458/001	041511066/M	TEVA ITALIA S.R.L.	IT
SIMVASTATINA ratiopharm 5 mg Compresse rivestite con film	FR/H/0458/001	041511078/M	TEVA ITALIA S.R.L.	IT
SIMVASTATINA ratiopharm 5 mg Compresse rivestite con film	FR/H/0458/001	041511054/M	TEVA ITALIA S.R.L.	IT
SIMVASTATINA ratiopharm 5 mg Compresse rivestite con film	FR/H/0458/001	041511041/M	TEVA ITALIA S.R.L.	IT
SIMVASTATINA ratiopharm 5 mg Compresse rivestite con film	FR/H/0458/001	041511015/M	TEVA ITALIA S.R.L.	IT
SIMVASTATINA	FR/H/0458/001	041511039/M	TEVA 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
ratiopharm 5 mg Compresse rivestite con film				
SIMVASTATINA ratiopharm 5 mg Compresse rivestite con film	FR/H/0458/001	041511066/M	TEVA ITALIA S.R.L.	IT
SIMVASTATINA ratiopharm 5 mg Compresse rivestite con film	FR/H/0458/001	041511027/M	TEVA ITALIA S.R.L.	IT
SIMVASTATINA ratiopharm 5 mg Compresse rivestite con film	FR/H/0458/001	041511078/M	TEVA ITALIA S.R.L.	IT
SIMVASTATINA ratiopharm 5 mg Compresse rivestite con film	FR/H/0458/001	041511027/M	TEVA ITALIA S.R.L.	IT
SIMVASTATINA ratiopharm 80 mg Compresse rivestite con film	FR/H/0458/005	041511344/M	TEVA ITALIA S.R.L.	IT
SIMVASTATINA ratiopharm 80 mg Compresse rivestite con film	FR/H/0458/005	041511320/M	TEVA ITALIA S.R.L.	IT
SIMVASTATINA ratiopharm 80 mg Compresse rivestite con film	FR/H/0458/005	041511371/M	TEVA ITALIA S.R.L.	IT
SIMVASTATINA ratiopharm 80 mg Compresse rivestite con	FR/H/0458/005	041511357/M	TEVA ITALIA S.R.L.	IT

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film				
SIMVASTATINA ratiopharm 80 mg Compresse rivestite con film	FR/H/0458/005	041511332/M	TEVA ITALIA S.R.L.	IT
SIMVASTATINA ratiopharm 80 mg Compresse rivestite con film	FR/H/0458/005	041511369/M	TEVA ITALIA S.R.L.	IT
SIMVASTATINA ratiopharm 80 mg Compresse rivestite con film	FR/H/0458/005	041511383/M	TEVA ITALIA S.R.L.	IT
SIMVASTATINA ratiopharm 80 mg Compresse rivestite con film	FR/H/0458/005	041511344/M	TEVA ITALIA S.R.L.	IT
SIMVASTATINA ratiopharm 80 mg Compresse rivestite con film	FR/H/0458/005	041511320/M	TEVA ITALIA S.R.L.	IT
SIMVASTATINA ratiopharm 80 mg Compresse rivestite con film	FR/H/0458/005	041511371/M	TEVA ITALIA S.R.L.	IT
SIMVASTATINA ratiopharm 80 mg Compresse rivestite con film	FR/H/0458/005	041511357/M	TEVA ITALIA S.R.L.	IT
SIMVASTATINA ratiopharm 80 mg Compresse rivestite con film	FR/H/0458/005	041511369/M	TEVA ITALIA S.R.L.	IT
SIMVASTATINA	FR/H/0458/005	041511383/M	TEVA ITALIA S.R.L.	IT

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ratiopharm 80 mg Compresse rivestite con film				
Simvastatina Teva Italia 10 mg compresse rivestite con film	FR/H/0459/002	041581202	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 10 mg compresse rivestite con film	FR/H/0459/002	041581190	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 10 mg compresse rivestite con film	FR/H/0459/002	041581164	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 10 mg compresse rivestite con film	FR/H/0459/002	041581149	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 10 mg compresse rivestite con film	FR/H/0459/002	041581188	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 10 mg compresse rivestite con film	FR/H/0459/002	041581214	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 10 mg compresse rivestite con film	FR/H/0459/002	041581176	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 10 mg compresse rivestite con film	FR/H/0459/002	041581152	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 10 mg compresse rivestite con film	FR/H/0459/002	041581125	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 10 mg compresse rivestite con film	FR/H/0459/002	041581113	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia	FR/H/0459/002	041581137	TEVA ITALIA S.R.L.	IT

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10 mg compresse rivestite con film				
Simvastatina Teva Italia 20 mg compresse rivestite con film	FR/H/0459/003	041581315	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 20 mg compresse rivestite con film	FR/H/0459/003	041581265	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 20 mg compresse rivestite con film	FR/H/0459/003	041581327	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 20 mg compresse rivestite con film	FR/H/0459/003	041581303	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 20 mg compresse rivestite con film	FR/H/0459/003	041581253	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 20 mg compresse rivestite con film	FR/H/0459/003	041581240	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 20 mg compresse rivestite con film	FR/H/0459/003	041581226	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 20 mg compresse rivestite con film	FR/H/0459/003	041581277	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 20 mg compresse rivestite con film	FR/H/0459/003	041581289	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 20 mg compresse rivestite con film	FR/H/0459/003	041581291	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 20 mg compresse	FR/H/0459/003	041581238	TEVA ITALIA S.R.L.	IT

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rivestite con film				
Simvastatina Teva Italia 40 mg compresse rivestite con film	FR/H/0459/004	041581341	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 40 mg compresse rivestite con film	FR/H/0459/004	041581392	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 40 mg compresse rivestite con film	FR/H/0459/004	041581380	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 40 mg compresse rivestite con film	FR/H/0459/004	041581404	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 40 mg compresse rivestite con film	FR/H/0459/004	041581428	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 40 mg compresse rivestite con film	FR/H/0459/004	041581378	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 40 mg compresse rivestite con film	FR/H/0459/004	041581366	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 40 mg compresse rivestite con film	FR/H/0459/004	041581430	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 40 mg compresse rivestite con film	FR/H/0459/004	041581354	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 40 mg compresse rivestite con film	FR/H/0459/004	041581416	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 40 mg compresse rivestite con film	FR/H/0459/004	041581339	TEVA 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
Simvastatina Teva Italia 5 mg compresse rivestite con film	FR/H/0459/001	041581051	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 5 mg compresse rivestite con film	FR/H/0459/001	041581036	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 5 mg compresse rivestite con film	FR/H/0459/001	041581075	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 5 mg compresse rivestite con film	FR/H/0459/001	041581087	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 5 mg compresse rivestite con film	FR/H/0459/001	041581063	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 5 mg compresse rivestite con film	FR/H/0459/001	041581048	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 5 mg compresse rivestite con film	FR/H/0459/001	041581012	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 5 mg compresse rivestite con film	FR/H/0459/001	041581024	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 5 mg compresse rivestite con film	FR/H/0459/001	041581101	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 5 mg compresse rivestite con film	FR/H/0459/001	041581099	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 5 mg compresse rivestite con film	FR/H/0459/001	041581051	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia	FR/H/0459/001	041581036	TEVA 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
5 mg compresse rivestite con film				
Simvastatina Teva Italia 5 mg compresse rivestite con film	FR/H/0459/001	041581075	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 5 mg compresse rivestite con film	FR/H/0459/001	041581087	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 5 mg compresse rivestite con film	FR/H/0459/001	041581063	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 5 mg compresse rivestite con film	FR/H/0459/001	041581048	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 5 mg compresse rivestite con film	FR/H/0459/001	041581012	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 5 mg compresse rivestite con film	FR/H/0459/001	041581024	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 5 mg compresse rivestite con film	FR/H/0459/001	041581101	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 5 mg compresse rivestite con film	FR/H/0459/001	041581099	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 80 mg compresse rivestite con film	FR/H/0459/005	041581531	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 80 mg compresse rivestite con film	FR/H/0459/005	041581529	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 80 mg compresse	FR/H/0459/005	041581467	TEVA 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
rivestite con film				
Simvastatina Teva Italia 80 mg compresse rivestite con film	FR/H/0459/005	041581493	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 80 mg compresse rivestite con film	FR/H/0459/005	041581505	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 80 mg compresse rivestite con film	FR/H/0459/005	041581479	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 80 mg compresse rivestite con film	FR/H/0459/005	041581442	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 80 mg compresse rivestite con film	FR/H/0459/005	041581481	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 80 mg compresse rivestite con film	FR/H/0459/005	041581517	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 80 mg compresse rivestite con film	FR/H/0459/005	041581455	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 80 mg compresse rivestite con film	FR/H/0459/005	041581531	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 80 mg compresse rivestite con film	FR/H/0459/005	041581529	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 80 mg compresse rivestite con film	FR/H/0459/005	041581467	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 80 mg compresse rivestite con film	FR/H/0459/005	041581493	TEVA 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
Simvastatina Teva Italia 80 mg compresse rivestite con film	FR/H/0459/005	041581505	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 80 mg compresse rivestite con film	FR/H/0459/005	041581479	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 80 mg compresse rivestite con film	FR/H/0459/005	041581442	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 80 mg compresse rivestite con film	FR/H/0459/005	041581481	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 80 mg compresse rivestite con film	FR/H/0459/005	041581517	TEVA ITALIA S.R.L.	IT
Simvastatina Teva Italia 80 mg compresse rivestite con film	FR/H/0459/005	041581455	TEVA ITALIA S.R.L.	IT
Simvastatin-CT 30 mg Filmtabletten	not available	86023.00.00	ABZ-PHARMA GMBH	DE
Simvastatin-CT 30 mg Filmtabletten	not available	86023.00.00	ABZ-PHARMA GMBH	DE
Simvastatin-CT 60 mg Filmtabletten	not available	86024.00.00	ABZ-PHARMA GMBH	DE
Simvastatin-CT 60 mg Filmtabletten	not available	86024.00.00	ABZ-PHARMA GMBH	DE
SIMVASTATINE ISOMED 10 mg, comprimé pelliculé sécable	not available	NL29313	TEVA SANTÉ	FR
SIMVASTATINE RATIOPHARM 10 mg, comprimé pelliculé sécable	not available	NL29314	RATIOPHARM GMBH	FR
SIMVASTATINE	not available	NL29314	RATIOPHARM GMBH	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
RATIOPHARM 10 mg, comprimé pelliculé sécable				
Simvastatine Sandoz 60 mg, filmomhulde tabletten	FI/H/0141/007	RVG 32196	SANDOZ B.V.	NL
Simvastatine Sandoz 60 mg, filmomhulde tabletten	FI/H/0141/007	RVG 32196	SANDOZ B.V.	NL
Simvastatine Teva 10 mg comprimés pelliculés	FR/H/0323/002	BE310843	TEVA PHARMA BELGIUM N.V./S.A	BE
SIMVASTATINE TEVA 10 mg, comprimé pelliculé sécable	FR/H/0323/002	NL29366	TEVA SANTÉ	FR
SIMVASTATINE TEVA 10 mg, comprimé pelliculé sécable	FR/H/0323/002	NL29366	TEVA SANTÉ	FR
imvastatine Teva 40 mg comprimés pelliculés	FR/H/0323/004	BE310877	TEVA PHARMA BELGIUM N.V./S.A	BE
Simvastatine Teva 40 mg filmomhulde tabletten	FR/H/0323/004	BE310877	TEVA PHARMA BELGIUM N.V./S.A	BE
Simvastatine Teva 40 mg filmomhulde tabletten	FR/H/0323/004	BE310877	TEVA PHARMA BELGIUM N.V./S.A	BE
SIMVASTATINE TEVA 40 mg FILMTABLETTEN	FR/H/0323/004	BE310877	TEVA PHARMA BELGIUM N.V./S.A	BE
SIMVASTATINE TEVA 80 mg, comprimé pelliculé sécable	FR/H/0323/005	NL29369	TEVA SANTÉ	FR
SIMVASTATINE TEVA 80 mg, comprimé pelliculé sécable	FR/H/0323/005	NL29369	TEVA SANTÉ	FR
Simvastatin-	not available	85907.00.00	RATIOPHARM 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
ratiopharm® 30 mg Filmtabletten				
Simvastatin-ratiopharm® 30 mg Filmtabletten	not available	85907.00.00	RATIOPHARM GMBH	DE
Simvastatin-ratiopharm® 60 mg Filmtabletten	not available	85908.00.00	RATIOPHARM GMBH	DE
Simvastatin-ratiopharm® 60 mg Filmtabletten	not available	85908.00.00	RATIOPHARM GMBH	DE
Simvastatinum 123ratio, 20 mg, tabletki powlekane	FR/H/0323/003	14378	123RATIO SP. Z O.O.	PL
Simvastatinum 123ratio, 20 mg, tabletki powlekane	FR/H/0323/004	14378	123RATIO SP. Z O.O.	PL
Sinvacor 10 mg compresse rivestite con film	UK/H/0687/002	027209016	MSD ITALIA S.R.L.	IT
SINVACOR 10 mg filmsko obložene tablete	not available	H/99/01426/001	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
SINVACOR 10 mg filmsko obložene tablete	not available	H/99/01426/001	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Sinvacor 20 mg compresse rivestite con film	UK/H/0687/003	027209028	MSD ITALIA S.R.L.	IT
Sinvacor 20 mg compresse rivestite con film	UK/H/0687/003	027209105	MSD ITALIA S.R.L.	IT
SINVACOR 20 mg filmsko obložene tablete	not available	H/99/01426/002	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA	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
			D.O.O.	
SINVACOR 20 mg filmsko obložene tablete	not available	H/99/01426/003	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
SINVACOR 20 mg filmsko obložene tablete	not available	H/99/01426/002	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
SINVACOR 20 mg filmsko obložene tablete	not available	H/99/01426/003	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Sinvacor 40 mg compresse rivestite con film	UK/H/0687/004	027209117	MSD ITALIA S.R.L.	IT
Sinvacor 40 mg compresse rivestite con film	UK/H/0687/004	027209042	MSD ITALIA S.R.L.	IT
SINVACOR FORTE 40 mg filmsko obložene tablete	not available	H/99/01426/004	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
SINVACOR FORTE 40 mg filmsko obložene tablete	not available	H/99/01426/005	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
SINVACOR FORTE 40 mg filmsko obložene tablete	not available	H/99/01426/004	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
SINVACOR FORTE 40 mg filmsko obložene tablete	not available	H/99/01426/005	MERCK SHARP & DOHME INOVATIVNA ZDRAVILA D.O.O.	SI
Sinvastatina Mylan 10 mg comprimidos revestidos por película	01/H/0177/001	3884780	MYLAN, LDA	PT
Sinvastatina Mylan 10 mg comprimidos revestidos por película	01/H/0177/001	3884889	MYLAN, LDA	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
Sinvastatina Mylan 10 mg comprimidos revestidos por película	01/H/0177/001	3884780	MYLAN, LDA	PT
Sinvastatina Mylan 10 mg comprimidos revestidos por película	01/H/0177/001	3884889	MYLAN, LDA	PT
Sinvastatina Mylan 20 mg comprimidos revestidos por película	01/H/0177/002	3884988	MYLAN, LDA	PT
Sinvastatina Mylan 20 mg comprimidos revestidos por película	01/H/0177/002	3885084	MYLAN, LDA	PT
Sinvastatina Mylan 20 mg comprimidos revestidos por película	01/H/0177/002	4026589	MYLAN, LDA	PT
Sinvastatina Mylan 20 mg comprimidos revestidos por película	01/H/0177/002	3884988	MYLAN, LDA	PT
Sinvastatina Mylan 20 mg comprimidos revestidos por película	01/H/0177/002	3885084	MYLAN, LDA	PT
Sinvastatina Mylan 20 mg comprimidos revestidos por película	01/H/0177/002	4026589	MYLAN, LDA	PT
Sinvastatina Mylan 40 mg comprimidos revestidos por película	01/H/0177/003	3885183	MYLAN, LDA	PT
Sinvastatina Mylan 40 mg comprimidos revestidos por película	01/H/0177/003	3885282	MYLAN, LDA	PT
Sinvastatina Mylan 40 mg comprimidos revestidos por película	01/H/0177/003	3885183	MYLAN, LDA	PT
Sinvastatina Mylan 40	01/H/0177/003	3885282	MYLAN, LDA	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
mg comprimidos revestidos por película				
SIVASTIN 10 mg, compresse rivestite con film	UK/H/687/002	027208014	SIGMA-TAU INDUSTRIE FARMACEUTICHE RIUNITE S.P.A.	IT
SIVASTIN 10 mg, compresse rivestite con film	UK/H/687/002	027208014	SIGMA-TAU INDUSTRIE FARMACEUTICHE RIUNITE S.P.A.	IT
SIVASTIN 20 mg, compresse rivestite con film	UK/H/687/003	027208026	SIGMA-TAU INDUSTRIE FARMACEUTICHE RIUNITE S.P.A.	IT
SIVASTIN 20 mg, compresse rivestite con film	UK/H/687/003	027208103	SIGMA-TAU INDUSTRIE FARMACEUTICHE RIUNITE S.P.A.	IT
SIVASTIN 20 mg, compresse rivestite con film	UK/H/687/003	027208026	SIGMA-TAU INDUSTRIE FARMACEUTICHE RIUNITE S.P.A.	IT
SIVASTIN 20 mg, compresse rivestite con film	UK/H/687/003	027208103	SIGMA-TAU INDUSTRIE FARMACEUTICHE RIUNITE S.P.A.	IT
SIVASTIN 40 mg, compresse rivestite con film	UK/H/687/004	027208040	SIGMA-TAU INDUSTRIE FARMACEUTICHE RIUNITE S.P.A.	IT
SIVASTIN 40 mg, compresse rivestite con film	UK/H/687/004	027208115	SIGMA-TAU INDUSTRIE FARMACEUTICHE RIUNITE S.P.A.	IT
SIVASTIN 40 mg, compresse rivestite con film	UK/H/687/004	027208040	SIGMA-TAU INDUSTRIE FARMACEUTICHE RIUNITE S.P.A.	IT
SIVASTIN 40 mg, compresse rivestite con film	UK/H/687/004	027208115	SIGMA-TAU INDUSTRIE FARMACEUTICHE RIUNITE S.P.A.	IT
Zocor ®40 mg, film-coated tablets.	UK/H/0687/004	PL0025/0243	MERCK SHARP & DOHME LTD.	UK

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
Zocor ®80 mg, film-coated tablets.	UK/H/0687/005	PL0025/0366	MERCK SHARP & DOHME LTD.	UK
ZOCOR 10 mg	not available	31/0155/92-C/S	MERCK SHARP & DOHME BV	SK
ZOCOR 10 mg	not available	31/0155/92-C/S	MERCK SHARP & DOHME BV	SK
ZOCOR 10 mg comprimidos recubiertos con película	UK/H/0687/002	58.845	MERCK SHARP & DOHME DE ESPAÑA, S.A	ES
Zocor 10 mg filmtabletta	not available	OGYI-T-4005/01	MSD PHARMA HUNGARY KFT.	HU
Zocor 10 mg filmtabletta	not available	OGYI-T-4005/01	MSD PHARMA HUNGARY KFT.	HU
ZOCOR 10 mg tabletter, filmdrasjerte	UK/H/0687/002	7907	MERCK SHARP & DOHME BV	NO
Zocor 10 mg tabletti, kalvopäällysteinen	UK/H/0687/002	10660	MERCK SHARP & DOHME BV	FI
Zocor 10 mg, compresse rivestite con film	not available	027216011	NEOPHARMED GENTILI S.R.L.	IT
Zocor 10 mg, compresse rivestite con film	not available	027216011	NEOPHARMED GENTILI S.R.L.	IT
Zocor 10 mg, compresse rivestite con film	not available	027216011	NEOPHARMED GENTILI S.R.L.	IT
Zocor 10 mg, film coated tablets	UK/H/0687/002	PA 1286/22/2	MERCK SHARP & DOHME IRELAND (HUMAN HEALTH) LTD	IE
Zocor 10 mg, film-coated tablets	not available	12353	MERCK SHARP & DOHME BV	CY
Zocor 10 mg, film-coated tablets	not available	12353	MERCK SHARP & DOHME BV	CY
Zocor 10 mg, film-coated tablets	not available	MA058/00101	MERCK SHARP & DOHME LTD.	MT
Zocor 10 mg, film-coated tablets	not available	MA058/00101	MERCK SHARP & DOHME LTD.	MT
ZOCOR 10 mg, filmdragerade tabletter	UK/H/0687/002	10660	MERCK SHARP & DOHME BV	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
ZOCOR 10 mg, filmomhulde tabletten	UK/H/0687/002	RVG 13193	MERCK SHARP & DOHME BV	NL
ZOCOR 10 mg, επικαλυμμένα με λεπτό υμένιο δισκία	UK/H/0687/002	28198	VIANEX S.A.	GR
ZOCOR 10, 10 mg, tabletki powlekane	not available	R/3659	MSD POLSKA SP. Z O.O.	PL
ZOCOR 10, 10 mg, tabletki powlekane	not available	R/3659	MSD POLSKA SP. Z O.O.	PL
ZOCOR 20 mg	not available	31/0041/13-S	MERCK SHARP & DOHME BV	SK
ZOCOR 20 mg	not available	31/0041/13-S	MERCK SHARP & DOHME BV	SK
ZOCOR 20 mg comprimate filmate	not available	6795/2014/01	MERCK SHARP & DOHME ROMANIA SRL	RO
ZOCOR 20 mg comprimate filmate	not available	6795/2014/02	MERCK SHARP & DOHME ROMANIA SRL	RO
ZOCOR 20 mg comprimate filmate	not available	6795/2014/01	MERCK SHARP & DOHME ROMANIA SRL	RO
ZOCOR 20 mg comprimate filmate	not available	6795/2014/02	MERCK SHARP & DOHME ROMANIA SRL	RO
ZOCOR 20 mg comprimidos recubiertos con película	UK/H/0687/003	58.846	MERCK SHARP & DOHME DE ESPAÑA, S.A	ES
ZOCOR 20 mg comprimidos revestidos por película	UK/H/0687/003	8768754	MERCK SHARP & DOHME, LDA.	PT
ZOCOR 20 mg comprimidos revestidos por película	UK/H/0687/003	8768739	MERCK SHARP & DOHME, LDA.	PT
ZOCOR 20 mg comprimidos revestidos por película	UK/H/0687/003	8768721	MERCK SHARP & DOHME, LDA.	PT
Zocor 20 mg filmtabletta	not available	OGYI-T-4005/02	MSD PHARMA HUNGARY KFT.	HU
Zocor 20 mg filmtabletta	not available	OGYI-T-4005/02	MSD PHARMA HUNGARY	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
			KFT.	
ZOCOR 20 mg Filmtabletten	UK/H/0687/003	BE147235	MSD BELGIUM BVBA/SPRL	BE
ZOCOR 20 mg tabletter, filmdrasjerte	UK/H/0687/003	7908	MERCK SHARP & DOHME BV	NO
Zocor 20 mg tabletti, kalvopäällysteinen	UK/H/0687/003	10661	MERCK SHARP & DOHME BV	FI
Zocor 20 mg, compresse rivestite con film	not available	027216023	NEOPHARMED GENTILI S.R.L.	IT
Zocor 20 mg, compresse rivestite con film	not available	027216098	NEOPHARMED GENTILI S.R.L.	IT
Zocor 20 mg, compresse rivestite con film	not available	027216023	NEOPHARMED GENTILI S.R.L.	IT
Zocor 20 mg, compresse rivestite con film	not available	027216098	NEOPHARMED GENTILI S.R.L.	IT
Zocor 20 mg, compresse rivestite con film	not available	027216023	NEOPHARMED GENTILI S.R.L.	IT
Zocor 20 mg, compresse rivestite con film	not available	027216098	NEOPHARMED GENTILI S.R.L.	IT
ZOCOR 20 mg, comprimé pelliculé sécable	UK/H/0687/003	34009 377 623-8 0	MSD FRANCE	FR
ZOCOR 20 mg, comprimé pelliculé sécable	UK/H/0687/003	34009 566 392-3 6	MSD FRANCE	FR
ZOCOR 20 mg, comprimé pelliculé sécable	UK/H/0687/003	34009 377 491-4 5	MSD FRANCE	FR
ZOCOR 20 mg, comprimé pelliculé sécable	UK/H/0687/003	34009 330 954-8 2	MSD FRANCE	FR
ZOCOR 20 mg, comprimé pelliculé sécable	UK/H/0687/003	34009 377 624-4 1	MSD FRANCE	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
ZOCOR 20 mg, comprimé pelliculé sécable	UK/H/0687/003	34009 367 668-9 1	MSD FRANCE	FR
ZOCOR 20 mg, comprimé pelliculé sécable	UK/H/0687/003	34009 556 648-5 7	MSD FRANCE	FR
ZOCOR 20 mg, comprimé pelliculé sécable	UK/H/0687/003	34009 342 518-3 2	MSD FRANCE	FR
Zocor 20 mg, film coated tablets	UK/H/0687/003	PA 1286/22/3	MERCK SHARP & DOHME IRELAND (HUMAN HEALTH) LTD	IE
Zocor 20 mg, film-coated tablets	not available	16156	MERCK SHARP & DOHME BV	CY
Zocor 20 mg, film-coated tablets	not available	16156	MERCK SHARP & DOHME BV	CY
Zocor 20 mg, film-coated tablets	not available	MA058/00102	MERCK SHARP & DOHME LTD.	MT
Zocor 20 mg, film-coated tablets	not available	MA058/00102	MERCK SHARP & DOHME LTD.	MT
ZOCOR 20 mg, filmdragerade tabletter	UK/H/0687/003	10661	MERCK SHARP & DOHME BV	FI
ZOCOR 20 mg, filmomhulde tabletten	UK/H/0687/003	RVG 13194	MERCK SHARP & DOHME BV	NL
ZOCOR 20 mg, επικαλυμμένα με λεπτό υμένιο δισκία	UK/H/0687/003	28197	VIANEX S.A.	GR
ZOCOR 20, 20 mg, tabletki powlekane	not available	R/3660	MSD POLSKA SP. Z O.O.	PL
ZOCOR 20, 20 mg, tabletki powlekane	not available	R/3660	MSD POLSKA SP. Z O.O.	PL
ZOCOR 40 mg	not available	31/0042/13-S	MERCK SHARP & DOHME BV	SK
ZOCOR 40 mg	not available	31/0042/13-S	MERCK SHARP & DOHME BV	SK
ZOCOR 40 mg	UK/H/0687/004	BE190206	MSD BELGIUM BVBA/SPRL	BE

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
comprimés pelliculés				
ZOCOR 40 mg comprimidos revestidos por película	UK/H/0687/004	9768762	MERCK SHARP & DOHME, LDA.	PT
ZOCOR 40 mg comprimidos revestidos por película	UK/H/0687/004	9768770	MERCK SHARP & DOHME, LDA.	PT
ZOCOR 40 mg Filmtabletten	UK/H/0687/004	BE190206	MSD BELGIUM BVBA/SPRL	BE
ZOCOR 40 mg tabletter, filmdrasjerte	UK/H/0687/004	94-3357	MERCK SHARP & DOHME BV	NO
Zocor 40 mg tabletti, kalvopäällysteinen	UK/H/0687/004	12198	MERCK SHARP & DOHME BV	FI
Zocor 40 mg, compresse rivestite con film	not available	027216035	NEOPHARMED GENTILI S.R.L.	IT
Zocor 40 mg, compresse rivestite con film	not available	027216100	NEOPHARMED GENTILI S.R.L.	IT
Zocor 40 mg, compresse rivestite con film	not available	027216035	NEOPHARMED GENTILI S.R.L.	IT
Zocor 40 mg, compresse rivestite con film	not available	027216100	NEOPHARMED GENTILI S.R.L.	IT
Zocor 40 mg, compresse rivestite con film	not available	027216035	NEOPHARMED GENTILI S.R.L.	IT
Zocor 40 mg, compresse rivestite con film	not available	027216100	NEOPHARMED GENTILI S.R.L.	IT
ZOCOR 40 mg, comprimé pelliculé	UK/H/0687/004	34009 361 104-6 5	MSD FRANCE	FR
ZOCOR 40 mg, comprimé pelliculé	UK/H/0687/004	34009 361 106-9 4	MSD FRANCE	FR
ZOCOR 40 mg, comprimé pelliculé	UK/H/0687/004	34009 563 036-1 8	MSD FRANCE	FR
ZOCOR 40 mg, comprimé pelliculé	UK/H/0687/004	34009 564 622-1 6	MSD FRANCE	FR
ZOCOR 40 mg,	UK/H/0687/004	34009 377 625-0 2	MSD FRANCE	FR

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comprimé pelliculé				
ZOCOR 40 mg, comprimé pelliculé	UK/H/0687/004	34009 361 105-2 6	MSD FRANCE	FR
ZOCOR 40 mg, comprimé pelliculé	UK/H/0687/004	34009 356 188-0 1	MSD FRANCE	FR
ZOCOR 40 mg, comprimé pelliculé	UK/H/0687/004	34009 356 190-5 1	MSD FRANCE	FR
ZOCOR 40 mg, comprimé pelliculé	UK/H/0687/004	34009 564 623-8 4	MSD FRANCE	FR
ZOCOR 40 mg, comprimé pelliculé	UK/H/0687/004	34009 563 037-8 6	MSD FRANCE	FR
ZOCOR 40 mg, comprimé pelliculé	UK/H/0687/004	34009 377 492-0 6	MSD FRANCE	FR
ZOCOR 40 mg, comprimé pelliculé	UK/H/0687/004	34009 356 189-7 9	MSD FRANCE	FR
ZOCOR 40 mg, comprimé pelliculé	UK/H/0687/004	34009 377 626-7 0	MSD FRANCE	FR
Zocor 40 mg, film coated tablets	UK/H/0687/004	PA 1286/22/4	MERCK SHARP & DOHME IRELAND (HUMAN HEALTH) LTD	IE
Zocor 40 mg, film-coated tablets	not available	16898	MERCK SHARP & DOHME BV	CY
Zocor 40 mg, film-coated tablets	not available	16898	MERCK SHARP & DOHME BV	CY
Zocor 40 mg, film-coated tablets	not available	MA058/00103	MERCK SHARP & DOHME LTD.	MT
Zocor 40 mg, film-coated tablets	not available	MA058/00103	MERCK SHARP & DOHME LTD.	MT
ZOCOR 40 mg, filmdragerade tabletter	UK/H/0687/004	12198	MERCK SHARP & DOHME BV	FI
ZOCOR 40 mg, filmomhulde tabletten	UK/H/0687/004	RVG 13195	MERCK SHARP & DOHME BV	NL
ZOCOR 40 mg, επικαλυμμένα με λεπτό	UK/H/0687/004	28196	VIANEX S.A.	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
υμένιο δισκία				
ZOCOR 40, 40 mg, tabletki powlekane	not available	7691	MSD POLSKA SP. Z O.O.	PL
ZOCOR 40, 40 mg, tabletki powlekane	not available	7691	MSD POLSKA SP. Z O.O.	PL
ZOCOR 80 mg	not available	31/0043/13-S	MERCK SHARP & DOHME BV	SK
ZOCOR 80 mg	not available	31/0043/13-S	MERCK SHARP & DOHME BV	SK
ZOCOR 80 mg filmdragerade tabletter	UK/H/0687/005	13311	MERCK SHARP & DOHME BV	FI
ZOCOR 80 mg tabletter, filmdrasjerte	UK/H/0687/005	97-3651	MERCK SHARP & DOHME BV	NO
Zocor 80 mg tabletti, kalvopäällysteinen	UK/H/0687/005	13311	MERCK SHARP & DOHME BV	FI
ZOCOR 80, 80 mg, tabletki powlekane	not available	R/8466	MSD POLSKA SP. Z O.O.	PL
ZOCOR 80, 80 mg, tabletki powlekane	not available	R/8466	MSD POLSKA SP. Z O.O.	PL
ZOCOR FORTE 40 mg comprimidos recubiertos con película	UK/H/0687/004	61.497	MERCK SHARP & DOHME DE ESPAÑA, S.A	ES
Zocor Forte 40 mg filmtabletta	not available	OGYI-T-4005/03	MSD PHARMA HUNGARY KFT.	HU
Zocor Forte 40 mg filmtabletta	not available	OGYI-T-4005/03	MSD PHARMA HUNGARY KFT.	HU
ZOCOR FORTE, 40 mg õhukese polümeerikattega tabletid	not available	169797	MERCK SHARP & DOHME OU	EE
ZOCOR FORTE, 40 mg õhukese polümeerikattega tabletid	not available	169797	MERCK SHARP & DOHME OU	EE
ZOCOR, 10 mg õhukese polümeerikattega	not available	088794	MERCK SHARP & DOHME OU	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
tabletid				
ZOCOR, 10 mg õhukese polümeerikattega tabletid	not available	088794	MERCK SHARP & DOHME OU	EE
ZOCOR, 20 mg õhukese polümeerikattega tabletid	not available	088894	MERCK SHARP & DOHME OU	EE
ZOCOR, 20 mg õhukese polümeerikattega tabletid	not available	088894	MERCK SHARP & DOHME OU	EE
Zocor, filmovertrukne tabletter	UK/H/0687/002	13313	MERCK SHARP & DOHME BV	DK
Zocor, filmovertrukne tabletter	UK/H/0687/004	17195	MERCK SHARP & DOHME BV	DK
Zocor, filmovertrukne tabletter	UK/H/0687/005	19300	MERCK SHARP & DOHME BV	DK
Zocor, filmovertrukne tabletter	UK/H/0687/003	13314	MERCK SHARP & DOHME BV	DK
ZOCOR® 10 mg Filmtabletten	UK/H/0687/002	35406.01.00	MSD SHARP & DOHME GMBH	DE
ZOCOR® 10 mg potahované tablety	not available	31/155/92-A/C	MERCK SHARP & DOHME BV	CZ
ZOCOR® 10 mg potahované tablety	not available	31/155/92-A/C	MERCK SHARP & DOHME BV	CZ
Zocor® 10 mg, film-coated tablets.	UK/H/0687/002	PL0025/0241	MERCK SHARP & DOHME LTD.	UK
ZOCOR® 20 mg comprimés pelliculés	UK/H/0687/003	BE147235	MSD BELGIUM BVBA/SPRL	BE
ZOCOR® 20 mg comprimés pelliculés	UK/H/0687/003	2011031037	MSD BELGIUM BVBA/SPRL	LU
ZOCOR® 20 mg filmomhulde tabletten	UK/H/0687/003	BE147235	MSD BELGIUM BVBA/SPRL	BE
ZOCOR® 20 mg Filmtabletten	UK/H/0687/003	35406.02.00	MSD SHARP & DOHME 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
ZOCOR® 20 mg potahované tablety	not available	31/155/92-B/C	MERCK SHARP & DOHME BV	CZ
ZOCOR® 20 mg potahované tablety	not available	31/155/92-B/C	MERCK SHARP & DOHME BV	CZ
Zocor® 20 mg, film-coated tablets.	UK/H/0687/003	PL0025/0242	MERCK SHARP & DOHME LTD.	UK
ZOCOR® 40 mg comprimés pelliculés	UK/H/0687/004	2011031036	MSD BELGIUM BVBA/SPRL	LU
ZOCOR® 40 mg filmomhulde tabletten	UK/H/0687/004	BE190206	MSD BELGIUM BVBA/SPRL	BE
ZOCOR® FORTE 40 mg Filmtabletten	UK/H/0687/004	35406.03.00	MSD SHARP & DOHME GMBH	DE
ZOCOR® FORTE 40 mg potahované tablety	not available	31/155/92-C/C	MERCK SHARP & DOHME BV	CZ
ZOCOR® FORTE 40 mg potahované tablety	not available	31/155/92-C/C	MERCK SHARP & DOHME BV	CZ
Zocord 10 mg filmdragerade tabletter	UK/H/0687/002	10772	MERCK SHARP & DOHME BV	SE
Zocord 20 mg filmdragerade tabletter	UK/H/0687/003	10773	MERCK SHARP & DOHME BV	SE
Zocord 20 mg Filmtabletten	UK/H/0687/003	1-19502	MERCK SHARP & DOHME GES.M.B.H.	AT
Zocord 40 mg filmdragerade tabletter	UK/H/0687/004	12462	MERCK SHARP & DOHME BV	SE
Zocord 40 mg Filmtabletten	UK/H/0687/004	1-21714	MERCK SHARP & DOHME GES.M.B.H.	AT
Zocord 80 mg Filmtabletten	UK/H/0687/005	1-23653	MERCK SHARP & DOHME GES.M.B.H.	AT
ЗОКОР 10 mg филмирани таблетки	not available	20010478	MERCK SHARP & DOHME BULGARIA EOOD	BG
ЗОКОР 10 mg филмирани таблетки	not available	20010478	MERCK SHARP & DOHME BULGARIA EOOD	BG
ЗОКОР 20 mg филмирани таблетки	not available	9600024	MERCK SHARP & DOHME BULGARIA EOOD	BG

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ЗОКОР 20 mg филмирани таблетки	not available	9600024	MERCK SHARP & DOHME BULGARIA EOOD	BG
ЗОКОР ФОРТЕ 40 mg филмирани таблетки	not available	9900075	MERCK SHARP & DOHME BULGARIA EOOD	BG
ЗОКОР ФОРТЕ 40 mg филмирани таблетки	not available	9900075	MERCK SHARP & DOHME BULGARIA EOOD	BG