1 December 2022
EMA/PRAC/918194/2022
Human Medicines Division

List of nationally authorised medicinal products

Active substance(s): sertraline

Procedure No. PSUSA/00002696/202203

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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
Aremis 100 mg comprimidos recubiertos con película	not available	59.734	ESTEVE PHARMACEUTICALS, S.A.	ES
Aremis 50 mg comprimidos recubiertos con película	not available	59.735	ESTEVE PHARMACEUTICALS, S.A.	ES
Besitran 100 mg comprimidos recubiertos con película	NL/H/1732/003	59.718	VIATRIS HEALTHCARE S.L.	ES
Besitran 20 mg/ml concentrado para solución oral	NL/H/1732/004	63.477	VIATRIS HEALTHCARE S.L.	ES
Besitran 50 mg comprimidos recubiertos con película	NL/H/1732/002	59.717	VIATRIS HEALTHCARE S.L.	ES
Dosertra 150 mg compresse rivestite con film	IT/H/0788/001	046561015	BRUNO FARMACEUTICI	IT
Dosertra 200 mg compresse rivestite con film	IT/H/0788/001-002	046561027	BRUNO FARMACEUTICI	IT
Ferbrain 150 mg comprimidos recubiertos con película	ES/H/0559/001	84732	FERRER INTERNACIONAL, S.A.	ES
Ferbrain 200 mg comprimidos recubiertos con película	ES/H/0559/002	84733	FERRER INTERNACIONAL, S.A.	ES
LUSTRAL® 100 mg film coated tablets	NL/H/1732/003	PA 23055/001/002	UPJOHN EESV	IE
LUSTRAL® 100 mg film coated tablets	NL/H/1732/003	PL 50622/0044	UPJOHN UK LIMITED	XI
LUSTRAL® 50 mg film coated tablets	NL/H/1732/002	PA 23055/001/001	UPJOHN EESV	IE
LUSTRAL® 50 mg film coated tablets	NL/H/1732/002	PL 50622/0045	UPJOHN UK 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
Semonic 150 mg comprimidos recubiertos con película	ES/H/0630/001	84190	NEURAXPHARM SPAIN, S.L.U.	ES
Semonic 150 mg comprimidos revestidos por película	ES/H/0630/001	5773577	NEURAXPHARM SPAIN, S.L.U.	PT
Semonic 150 mg comprimidos revestidos por película	ES/H/0630/001	5787015	NEURAXPHARM SPAIN, S.L.U.	PT
Semonic 200 mg comprimidos recubiertos con película	ES/H/0630/002	84191	NEURAXPHARM SPAIN, S.L.U.	ES
Semonic 200 mg comprimidos revestidos por película	ES/H/0630/002	5773601	NEURAXPHARM SPAIN, S.L.U.	PT
Semonic 200 mg comprimidos revestidos por película	ES/H/0630/002	5787148	NEURAXPHARM SPAIN, S.L.U.	PT
Serlain 100 mg comprimés pelliculés	NL/H/1732/003	BE157351	UPJOHN SRL	BE
Serlain 100 mg Filmtabletten	NL/H/1732/003	2011010936	UPJOHN SRL	LU
Serlain 50 mg Filmtabletten	NL/H/1732/002	BE157324	UPJOHN SRL	BE
Serlain 50 mg Filmtabletten	NL/H/1732/002	2011010935	UPJOHN SRL	LU
Sertralin Viatris 100 mg Filmtabletten	NL/H/1736/002	1-30158	UPJOHN EESV	AT
Sertralin Viatris 50 mg Filmtabletten	NL/H/1736/001	1-30157	UPJOHN EESV	AT
Sertraline 100 mg film coated tablets	NL/H/1736/002	PL 50622/0056	UPJOHN UK LIMITED	XI
Sertraline 25 mg Film-coated Tablets	not available	PL 17780/0956	ZENTIVA PHARMA UK LIMITED	XI
Sertraline 25 mg Film-coated Tablets	not available	PL 12762/0553	MERCURY PHARMACEUTICALS LTD.	XI
Sertraline 50 mg film coated tablets	NL/H/1736/001	PL 50622/0057	UPJOHN UK 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
Sertraline Viatris 100 mg, filmomhulde tabletten	NL/H/1736/002	RVG 105382	VIATRIS NETHERLANDS B.V.	NL
Sertraline Viatris 20 mg/ml, concentraat voor oplossing voor oraal gebruik	NL/H/1736/003	RVG 105383	VIATRIS NETHERLANDS B.V.	NL
Sertraline Viatris 25 mg, filmomhulde tabletten	NL/H/1736/004	RVG 106062	VIATRIS NETHERLANDS B.V.	NL
SERTRALINE VIATRIS 25 mg, gélule	not available	NL 30272	VIATRIS SANTE	FR
Sertraline Viatris 50 mg, filmomhulde tabletten	NL/H/1736/001	RVG 105381	VIATRIS NETHERLANDS B.V.	NL
SERTRALINE VIATRIS 50 mg, gélule	not available	NL 30273	VIATRIS SANTE	FR
Sertralin-neuraxpharm 150 mg Filmtabletten	ES/H/0630/001	2205942.00.00	NEURAXPHARM ARZNEIMITTEL GMBH	DE
Sertralin-neuraxpharm 200 mg Filmtabletten	ES/H/0630/002	2205943.00.00	NEURAXPHARM ARZNEIMITTEL GMBH	DE
Tatig 100 mg compresse rivestite con film	NL/H/1736/002	027754050	VIATRIS PHARMA S.R.L.	IT
Tatig 100 mg compresse rivestite con film	NL/H/1736/002	027754098	VIATRIS PHARMA S.R.L.	IT
Tatig 100 mg compresse rivestite con film	NL/H/1736/002	027754403	VIATRIS PHARMA S.R.L.	IT
Tatig 20 mg/ml concentrato per soluzione orale	NL/H/1736/003	027754035	VIATRIS PHARMA S.R.L.	IT
Tatig 50 mg compresse rivestite con film	NL/H/1736/001	027754086	VIATRIS PHARMA S.R.L.	IT
Tatig 50 mg compresse rivestite con film	NL/H/1736/001	027754047	VIATRIS PHARMA S.R.L.	IT
Tatig 50 mg compresse rivestite con film	NL/H/1736/001	027754391	VIATRIS PHARMA S.R.L.	IT
Tresleen® 100 mg Filmtabletten	NL/H/1732/003	1-30163	UPJOHN EESV	AT
Tresleen® 50 mg Filmtabletten	NL/H/1732/002	1-21385	UPJOHN EESV	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
Zoloft 100 mg compresse rivestite con film	NL/H/1732/003	027753045	VIATRIS PHARMA S.R.L.	IT
Zoloft 100 mg compresse rivestite con film	NL/H/1732/003	027753110	VIATRIS PHARMA S.R.L.	IT
Zoloft 100 mg compresse rivestite con film	NL/H/1732/003	027753452	VIATRIS PHARMA S.R.L.	IT
Zoloft 100 mg compresse rivestite con film	NL/H/1732/003	027753300	VIATRIS PHARMA S.R.L.	IT
Zoloft 100 mg compresse rivestite con film	NL/H/1732/003	027753312	VIATRIS PHARMA S.R.L.	IT
Zoloft 100 mg compresse rivestite con film	NL/H/1732/003	027753336	VIATRIS PHARMA S.R.L.	IT
Zoloft 100 mg compresse rivestite con film	NL/H/1732/003	027753348	VIATRIS PHARMA S.R.L.	IT
Zoloft 100 mg compresse rivestite con film	NL/H/1732/003	027753324	VIATRIS PHARMA S.R.L.	IT
Zoloft 100 mg compresse rivestite con film	NL/H/1732/003	027753298	VIATRIS PHARMA S.R.L.	IT
Zoloft 100 mg compresse rivestite con film	NL/H/1732/003	027753351	VIATRIS PHARMA S.R.L.	IT
Zoloft 100 mg compresse rivestite con film	NL/H/1732/003	027753375	VIATRIS PHARMA S.R.L.	IT
Zoloft 100 mg compresse rivestite con film	NL/H/1732/003	027753425	VIATRIS PHARMA S.R.L.	IT
Zoloft 100 mg compresse rivestite con	NL/H/1732/003	027753401	VIATRIS PHARMA S.R.L.	IT

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film				
Zoloft 100 mg compresse rivestite con film	NL/H/1732/003	027753399	VIATRIS PHARMA S.R.L.	IT
Zoloft 100 mg compresse rivestite con film	NL/H/1732/003	027753413	VIATRIS PHARMA S.R.L.	IT
Zoloft 100 mg compresse rivestite con film	NL/H/1732/003	027753387	VIATRIS PHARMA S.R.L.	IT
Zoloft 100 mg compresse rivestite con film	NL/H/1732/003	027753363	VIATRIS PHARMA S.R.L.	IT
Zoloft 100 mg comprimate filmate	NL/H/1732/003	8228/2015/02	UPJOHN EESV	RO
Zoloft 100 mg comprimate filmate	NL/H/1732/003	8228/2015/14	UPJOHN EESV	RO
Zoloft 100 mg comprimate filmate	NL/H/1732/003	8228/2015/16	UPJOHN EESV	RO
Zoloft 100 mg comprimate filmate	NL/H/1732/003	8228/2015/04	UPJOHN EESV	RO
Zoloft 100 mg comprimate filmate	NL/H/1732/003	8228/2015/10	UPJOHN EESV	RO
Zoloft 100 mg comprimate filmate	NL/H/1732/003	8228/2015/08	UPJOHN EESV	RO
Zoloft 100 mg comprimate filmate	NL/H/1732/003	8228/2015/06	UPJOHN EESV	RO
Zoloft 100 mg comprimate filmate	NL/H/1732/003	8228/2015/12	UPJOHN EESV	RO
Zoloft 100 mg comprimate filmate	NL/H/1732/003	8228/2015/03	UPJOHN EESV	RO
Zoloft 100 mg comprimate filmate	NL/H/1732/003	8228/2015/01	UPJOHN EESV	RO
Zoloft 100 mg comprimate filmate	NL/H/1732/003	8228/2015/07	UPJOHN EESV	RO
Zoloft 100 mg comprimate filmate	NL/H/1732/003	8228/2015/11	UPJOHN EESV	RO
Zoloft 100 mg	NL/H/1732/003	8228/2015/09	UPJOHN EESV	RO

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comprimate filmate				
Zoloft 100 mg comprimate filmate	NL/H/1732/003	8228/2015/15	UPJOHN EESV	RO
Zoloft 100 mg comprimate filmate	NL/H/1732/003	8228/2015/05	UPJOHN EESV	RO
Zoloft 100 mg comprimate filmate	NL/H/1732/003	8228/2015/13	UPJOHN EESV	RO
Zoloft 100 mg comprimate filmate	NL/H/1732/003	8228/2015/17	UPJOHN EESV	RO
Zoloft 100 mg comprimidos revestidos por película	NL/H/1732/003	5854484	UPJOHN EESV	PT
Zoloft 100 mg comprimidos revestidos por película	NL/H/1732/003	2182889	UPJOHN EESV	PT
Zoloft 100 mg comprimidos revestidos por película	NL/H/1732/003	2182988	UPJOHN EESV	PT
Zoloft 100 mg filmdragerad tablett	NL/H/1732/003	12567	UPJOHN EESV	SE
Zoloft 100 mg filmdrasjerte tabletter	NL/H/1732/003	00-8204	UPJOHN EESV	NO
ZOLOFT 100 mg filmom obalené tablety	NL/H/1732/003	30/0430/10-S	UPJOHN EESV	SK
Zoloft 100 mg filmsko obložene tablete	NL/H/1732/003	H/94/01718/017	UPJOHN EESV	SI
Zoloft 100 mg filmsko obložene tablete	NL/H/1732/003	H/94/01718/032	UPJOHN EESV	SI
Zoloft 100 mg filmsko obložene tablete	NL/H/1732/003	H/94/01718/022	UPJOHN EESV	SI
Zoloft 100 mg filmsko obložene tablete	NL/H/1732/003	H/94/01718/018	UPJOHN EESV	SI
Zoloft 100 mg filmsko obložene tablete	NL/H/1732/003	H/94/01718/020	UPJOHN EESV	SI
Zoloft 100 mg filmsko obložene tablete	NL/H/1732/003	H/94/01718/025	UPJOHN EESV	SI
Zoloft 100 mg filmsko obložene tablete	NL/H/1732/003	H/94/01718/021	UPJOHN EESV	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
Zoloft 100 mg filmsko obložene tablete	NL/H/1732/003	H/94/01718/028	UPJOHN EESV	SI
Zoloft 100 mg filmsko obložene tablete	NL/H/1732/003	H/94/01718/019	UPJOHN EESV	SI
Zoloft 100 mg filmsko obložene tablete	NL/H/1732/003	H/94/01718/026	UPJOHN EESV	SI
Zoloft 100 mg filmsko obložene tablete	NL/H/1732/003	H/94/01718/024	UPJOHN EESV	SI
Zoloft 100 mg filmsko obložene tablete	NL/H/1732/003	H/94/01718/029	UPJOHN EESV	SI
Zoloft 100 mg filmsko obložene tablete	NL/H/1732/003	H/94/01718/027	UPJOHN EESV	SI
Zoloft 100 mg filmsko obložene tablete	NL/H/1732/003	H/94/01718/023	UPJOHN EESV	SI
Zoloft 100 mg filmsko obložene tablete	NL/H/1732/003	H/94/01718/031	UPJOHN EESV	SI
Zoloft 100 mg filmsko obložene tablete	NL/H/1732/003	H/94/01718/030	UPJOHN EESV	SI
Zoloft 100 mg filmsko obložene tablete	NL/H/1732/003	H/94/01718/035	UPJOHN EESV	SI
Zoloft 100 mg filmuhúðaðar töflur.	NL/H/1732/003	990469	UPJOHN EESV	IS
Zoloft 100 mg kalvopäällysteinen tabletti	NL/H/1732/003	11558	UPJOHN EESV	FI
Zoloft 100 mg potahované tablety	NL/H/1732/003	30/1093/94-B/C	UPJOHN EESV	CZ
Zoloft 100 mg επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/1732/003	14678	UPJOHN HELLAS L.T.D.	CY
Zoloft 100 mg επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/1732/003	43085/23-06-2015	UPJOHN HELLAS L.T.D.	GR
Zoloft 100, filmomhulde tabletten 100 mg	NL/H/1732/003	RVG 105255	VIATRIS NETHERLANDS B.V.	NL
Zoloft 20 mg/ml concentrado para solução oral	NL/H/1732/004	3268083	UPJOHN EESV	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
Zoloft 20 mg/ml concentrat pentru soluţie orală	NL/H/1732/004	8229/2015/01	UPJOHN EESV	RO
Zoloft 20 mg/ml concentrato per soluzione orale	NL/H/1732/004	027753096	VIATRIS PHARMA S.R.L.	IT
Zoloft 20 mg/ml koncentrat till oral lösning	NL/H/1732/004	15504	UPJOHN EESV	SE
Zoloft 20 mg/ml koncentrat za peroralno raztopino	NL/H/1732/004	H/94/01718/033	UPJOHN EESV	SI
Zoloft 20 mg/ml koncentrátum belsőleges oldathoz	NL/H/1732/004	OGYI-T-4342/02	UPJOHN EESV	HU
Zoloft 20 mg/ml, concentraat voor oplossing voor oraal gebruik	NL/H/1732/004	RVG 24641	VIATRIS NETHERLANDS B.V.	NL
Zoloft 25 mg compresse rivestite con film	NL/H/1732/001	027753437	VIATRIS PHARMA S.R.L.	IT
Zoloft 25 mg compresse rivestite con film	NL/H/1732/001	027753134	VIATRIS PHARMA S.R.L.	IT
Zoloft 25 mg compresse rivestite con film	NL/H/1732/001	027753122	VIATRIS PHARMA S.R.L.	IT
Zoloft 25 mg compresse rivestite con film	NL/H/1732/001	027753146	VIATRIS PHARMA S.R.L.	IT
Zoloft 25 mg filmdragerad tablett	NL/H/1732/001	16689	UPJOHN EESV	SE
Zoloft 25 mg filmdrasjerte tabletter.	NL/H/1732/001	00-7908	UPJOHN EESV	NO
Zoloft 25 mg filmuhúðaðar töflur.	NL/H/1732/001	IS/1/01/102/01	UPJOHN EESV	IS
ZOLOFT 25 mg, gélule	not available	34009 355 621 2 8	VIATRIS UP	FR
ZOLOFT 25 mg, gélule	not available	34009 563 118 8 0	VIATRIS UP	FR
ZOLOFT 25 mg, gélule	not available	34009 563 119 4 1	VIATRIS UP	FR
ZOLOFT 25 mg, gélule	not available	34009 355 620 6 7	VIATRIS UP	FR
Zoloft 25, filmomhulde	NL/H/1732/001	RVG 105254	VIATRIS NETHERLANDS 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
tabletten 25 mg				
Zoloft 50 mg apvalkotās tabletes	NL/H/1732/002	97-0311	UPJOHN EESV	LV
Zoloft 50 mg compresse rivestite con film	NL/H/1732/002	027753033	VIATRIS PHARMA S.R.L.	IT
Zoloft 50 mg compresse rivestite con film	NL/H/1732/002	027753108	VIATRIS PHARMA S.R.L.	IT
Zoloft 50 mg compresse rivestite con film	NL/H/1732/002	027753449	VIATRIS PHARMA S.R.L.	IT
Zoloft 50 mg compresse rivestite con film	NL/H/1732/002	027753209	VIATRIS PHARMA S.R.L.	IT
Zoloft 50 mg compresse rivestite con film	NL/H/1732/002	027753286	VIATRIS PHARMA S.R.L.	IT
Zoloft 50 mg compresse rivestite con film	NL/H/1732/002	027753161	VIATRIS PHARMA S.R.L.	IT
Zoloft 50 mg compresse rivestite con film	NL/H/1732/002	027753262	VIATRIS PHARMA S.R.L.	IT
Zoloft 50 mg compresse rivestite con film	NL/H/1732/002	027753274	VIATRIS PHARMA S.R.L.	IT
Zoloft 50 mg compresse rivestite con film	NL/H/1732/002	027753197	VIATRIS PHARMA S.R.L.	IT
Zoloft 50 mg compresse rivestite con film	NL/H/1732/002	027753185	VIATRIS PHARMA S.R.L.	IT
Zoloft 50 mg compresse rivestite con film	NL/H/1732/002	027753211	VIATRIS PHARMA S.R.L.	IT
Zoloft 50 mg compresse rivestite con film	NL/H/1732/002	027753173	VIATRIS PHARMA S.R.L.	IT
Zoloft 50 mg compresse rivestite con film	NL/H/1732/002	027753159	VIATRIS PHARMA S.R.L.	IT
Zoloft 50 mg compresse rivestite con film	NL/H/1732/002	027753247	VIATRIS PHARMA S.R.L.	IT
Zoloft 50 mg compresse rivestite con film	NL/H/1732/002	027753223	VIATRIS PHARMA S.R.L.	IT
Zoloft 50 mg compresse rivestite con film	NL/H/1732/002	027753235	VIATRIS PHARMA S.R.L.	IT
Zoloft 50 mg compresse rivestite con film	NL/H/1732/002	027753250	VIATRIS PHARMA S.R.L.	IT
Zoloft 50 mg	NL/H/1732/002	8227/2015/02	UPJOHN EESV	RO

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comprimate filmate				
Zoloft 50 mg comprimate filmate	NL/H/1732/002	8227/2015/14	UPJOHN EESV	RO
Zoloft 50 mg comprimate filmate	NL/H/1732/002	8227/2015/16	UPJOHN EESV	RO
Zoloft 50 mg comprimate filmate	NL/H/1732/002	8227/2015/10	UPJOHN EESV	RO
Zoloft 50 mg comprimate filmate	NL/H/1732/002	8227/2015/12	UPJOHN EESV	RO
Zoloft 50 mg comprimate filmate	NL/H/1732/002	8227/2015/03	UPJOHN EESV	RO
Zoloft 50 mg comprimate filmate	NL/H/1732/002	8227/2015/15	UPJOHN EESV	RO
Zoloft 50 mg comprimate filmate	NL/H/1732/002	8227/2015/11	UPJOHN EESV	RO
Zoloft 50 mg comprimate filmate	NL/H/1732/002	8227/2015/01	UPJOHN EESV	RO
Zoloft 50 mg comprimate filmate	NL/H/1732/002	8227/2015/13	UPJOHN EESV	RO
Zoloft 50 mg comprimate filmate	NL/H/1732/002	8227/2015/05	UPJOHN EESV	RO
Zoloft 50 mg comprimate filmate	NL/H/1732/002	8227/2015/09	UPJOHN EESV	RO
Zoloft 50 mg comprimate filmate	NL/H/1732/002	8227/2015/07	UPJOHN EESV	RO
Zoloft 50 mg comprimate filmate	NL/H/1732/002	8227/2015/06	UPJOHN EESV	RO
Zoloft 50 mg comprimate filmate	NL/H/1732/002	8227/2015/04	UPJOHN EESV	RO
Zoloft 50 mg comprimate filmate	NL/H/1732/002	8227/2015/08	UPJOHN EESV	RO
Zoloft 50 mg comprimate filmate	NL/H/1732/002	8227/2015/17	UPJOHN EESV	RO
Zoloft 50 mg comprimidos revestidos por película	NL/H/1732/002	2182780	UPJOHN EESV	PT
Zoloft 50 mg comprimidos revestidos	NL/H/1732/002	5854385	UPJOHN EESV	PT

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por película				
Zoloft 50 mg comprimidos revestidos por película	NL/H/1732/002	5830989	UPJOHN EESV	PT
Zoloft 50 mg comprimidos revestidos por película	NL/H/1732/002	2182681	UPJOHN EESV	PT
Zoloft 50 mg filmdragerad tablett	NL/H/1732/002	11557	UPJOHN EESV	FI
Zoloft 50 mg filmdragerad tablett	NL/H/1732/002	12566	UPJOHN EESV	SE
Zoloft 50 mg filmdrasjerte tabletter.	NL/H/1732/002	00-8203	UPJOHN EESV	NO
ZOLOFT 50 mg filmom obalené tablety	NL/H/1732/002	30/0090/96-S	UPJOHN EESV	SK
ZOLOFT 50 mg filmom obložene tablete	not available	HR-H-153369585	UPJOHN EESV	HR
Zoloft 50 mg filmsko obložene tablete	NL/H/1732/002	H/94/01718/002	UPJOHN EESV	SI
Zoloft 50 mg filmsko obložene tablete	NL/H/1732/002	H/94/01718/001	UPJOHN EESV	SI
Zoloft 50 mg filmsko obložene tablete	NL/H/1732/002	H/94/01718/012	UPJOHN EESV	SI
Zoloft 50 mg filmsko obložene tablete	NL/H/1732/002	H/94/01718/006	UPJOHN EESV	SI
Zoloft 50 mg filmsko obložene tablete	NL/H/1732/002	H/94/01718/009	UPJOHN EESV	SI
Zoloft 50 mg filmsko obložene tablete	NL/H/1732/002	H/94/01718/011	UPJOHN EESV	SI
Zoloft 50 mg filmsko obložene tablete	NL/H/1732/002	H/94/01718/004	UPJOHN EESV	SI
Zoloft 50 mg filmsko obložene tablete	NL/H/1732/002	H/94/01718/016	UPJOHN EESV	SI
Zoloft 50 mg filmsko obložene tablete	NL/H/1732/002	H/94/01718/007	UPJOHN EESV	SI
Zoloft 50 mg filmsko obložene tablete	NL/H/1732/002	H/94/01718/015	UPJOHN EESV	SI
Zoloft 50 mg filmsko	NL/H/1732/002	H/94/01718/013	UPJOHN EESV	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
obložene tablete				
Zoloft 50 mg filmsko obložene tablete	NL/H/1732/002	H/94/01718/008	UPJOHN EESV	SI
Zoloft 50 mg filmsko obložene tablete	NL/H/1732/002	H/94/01718/010	UPJOHN EESV	SI
Zoloft 50 mg filmsko obložene tablete	NL/H/1732/002	H/94/01718/014	UPJOHN EESV	SI
Zoloft 50 mg filmsko obložene tablete	NL/H/1732/002	H/94/01718/005	UPJOHN EESV	SI
Zoloft 50 mg filmsko obložene tablete	NL/H/1732/002	H/94/01718/003	UPJOHN EESV	SI
Zoloft 50 mg filmsko obložene tablete	NL/H/1732/002	H/94/01718/034	UPJOHN EESV	SI
Zoloft 50 mg filmtabletta	NL/H/1732/002	OGYI-T-4342/01	UPJOHN EESV	HU
Zoloft 50 mg filmuhúðaðar töflur.	NL/H/1732/002	950077	UPJOHN EESV	IS
Zoloft 50 mg plėvele dengtos tabletės	NL/H/1732/002	LT/1/96/2037/005	UPJOHN EESV	LT
Zoloft 50 mg plėvele dengtos tabletės	NL/H/1732/002	LT/1/96/2037/004	UPJOHN EESV	LT
Zoloft 50 mg plėvele dengtos tabletės	NL/H/1732/002	LT/1/96/2037/008	UPJOHN EESV	LT
Zoloft 50 mg plėvele dengtos tabletės	NL/H/1732/002	LT/1/96/2037/006	UPJOHN EESV	LT
Zoloft 50 mg plėvele dengtos tabletės	NL/H/1732/002	LT/1/96/2037/010	UPJOHN EESV	LT
Zoloft 50 mg plėvele dengtos tabletės	NL/H/1732/002	LT/1/96/2037/013	UPJOHN EESV	LT
Zoloft 50 mg plėvele dengtos tabletės	NL/H/1732/002	LT/1/96/2037/012	UPJOHN EESV	LT
Zoloft 50 mg plėvele dengtos tabletės	NL/H/1732/002	LT/1/96/2037/002	UPJOHN EESV	LT
Zoloft 50 mg plėvele dengtos tabletės	NL/H/1732/002	LT/1/96/2037/016	UPJOHN EESV	LT
Zoloft 50 mg plėvele dengtos tabletės	NL/H/1732/002	LT/1/96/2037/009	UPJOHN EESV	LT
Zoloft 50 mg plėvele dengtos tabletės	NL/H/1732/002	LT/1/96/2037/001	UPJOHN EESV	LT

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Zoloft 50 mg plėvele dengtos tabletės	NL/H/1732/002	LT/1/96/2037/014	UPJOHN EESV	LT
Zoloft 50 mg plėvele dengtos tabletės	NL/H/1732/002	LT/1/96/2037/003	UPJOHN EESV	LT
Zoloft 50 mg plėvele dengtos tabletės	NL/H/1732/002	LT/1/96/2037/007	UPJOHN EESV	LT
Zoloft 50 mg plėvele dengtos tabletės	NL/H/1732/002	LT/1/96/2037/015	UPJOHN EESV	LT
Zoloft 50 mg plėvele dengtos tabletės	NL/H/1732/002	LT/1/96/2037/011	UPJOHN EESV	LT
Zoloft 50 mg plėvele dengtos tabletės	NL/H/1732/002	LT/1/96/2037/017	UPJOHN EESV	LT
Zoloft 50 mg potahované tablety	NL/H/1732/002	30/1093/94-A/C	UPJOHN EESV	CZ
Zoloft 50 mg επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/1732/002	14677	UPJOHN HELLAS L.T.D.	CY
Zoloft 50 mg επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/1732/002	43084/23-06-2015	UPJOHN HELLAS L.T.D.	GR
ZOLOFT 50 mg, gélule	not available	34009 300 379 3 5	VIATRIS UP	FR
ZOLOFT 50 mg, gélule	not available	34009 340 341 9 0	VIATRIS UP	FR
ZOLOFT 50 mg, gélule	not available	34009 340 344 8 0	VIATRIS UP	FR
ZOLOFT 50 mg, gélule	not available	34009 340 342 5 1	VIATRIS UP	FR
ZOLOFT 50 mg, gélule	not available	34009 340 343 1 2	VIATRIS UP	FR
Zoloft 50, filmomhulde tabletten 50 mg	NL/H/1732/002	RVG 16292	VIATRIS NETHERLANDS B.V.	NL
ZOLOFT OC 20 mg/ml koncentrát na perorálny roztok	NL/H/1732/004	30/0207/01-S	UPJOHN EESV	SK
ZOLOFT, 100 mg, tabletki powlekane	NL/H/1732/003	7532	UPJOHN EESV	PL
Zoloft, 50 mg õhukese polümeerikattega tabletid	NL/H/1732/002	122095	UPJOHN EESV	EE
ZOLOFT, 50 mg, tabletki powlekane	NL/H/1732/002	7531	UPJOHN EESV	PL
Zoloft, filmovertrukne	NL/H/1732/001	32146	UPJOHN EESV	DK

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tabletter				
Zoloft, filmovertrukne tabletter	NL/H/1732/002	13471	UPJOHN EESV	DK
Zoloft, filmovertrukne tabletter	NL/H/1732/003	13472	UPJOHN EESV	DK
Zoloft® 100 mg, Filmtabletten	NL/H/1732/003	37076.01.00	VIATRIS PHARMA GMBH	DE
Zoloft® 20 mg/ml Konzentrat zur Herstellung einer Lösung zum Einnehmen	NL/H/1732/004	46312.00.00	VIATRIS PHARMA GMBH	DE
Zoloft® 50 mg, Filmtabletten	NL/H/1732/002	37076.00.00	VIATRIS PHARMA GMBH	DE
Золофт 50 mg филмирани таблетки	NL/H/1732/002	20000514	UPJOHN EESV	BG