30 November 2017
EMA/802792/2017
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: sertraline

Procedure no.: PSUSA/00002696/201703

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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
Tresleen® 50 mg Filmtabletten	NL/H/1732/002	1-21385	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	AT
Tresleen® 100 mg Filmtabletten	NL/H/1732/003	1-30163	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	AT
Tresleen® 50 mg Filmtabletten	NL/H/1732/002	1-21385	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	AT
Tresleen® 100 mg Filmtabletten	NL/H/1732/003	1-30163	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	AT
Gladem® 50 mg - Filmtabletten	not available	1-21401	BOEHRINGER INGELHEIM RCV GMBH & CO KG	AT
Sertralin Pfizer 100 mg Filmtabletten	NL/H/1736/002	1-30158	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	AT
Sertralin Pfizer 50 mg Filmtabletten	NL/H/1736/001	1-30157	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	AT
Serlain 100 mg Filmtabletten	NL/H/1732/003	BE157351	PFIZER S.A. (BELGIUM)	BE
Serlain 50 mg Filmtabletten	NL/H/1732/002	BE157324	PFIZER S.A. (BELGIUM)	BE
Serlain 20 mg / ml Konzentrat zur Herstellung einer Lösung zum Einnehmen	NL/H/1732/004	BE226755	PFIZER S.A. (BELGIUM)	BE
Serlain 50 mg filmomhulde tabletten	NL/H/1732/002	BE157324	PFIZER S.A. (BELGIUM)	BE
Serlain 100 mg filmomhulde tabletten	NL/H/1732/003	BE157351	PFIZER S.A. (BELGIUM)	BE
Serlain 20 mg/ml concentraat voor drank	NL/H/1732/004	BE226755	PFIZER S.A. (BELGIUM)	BE
Serlain 50 mg comprimés pelliculés	NL/H/1732/002	BE157324	PFIZER S.A. (BELGIUM)	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
Serlain 20 mg/ml solution à diluer pour solution buvable	NL/H/1732/004	BE226755	PFIZER S.A. (BELGIUM)	BE
Serlain 100 mg comprimés pelliculés	NL/H/1732/003	BE157351	PFIZER S.A. (BELGIUM)	BE
Serlain 100 mg Filmtabletten	NL/H/1732/003	BE157351	PFIZER S.A. (BELGIUM)	BE
Serlain 50 mg Filmtabletten	NL/H/1732/002	BE157324	PFIZER S.A. (BELGIUM)	BE
Serlain 20 mg / ml Konzentrat zur Herstellung einer Lösung zum Einnehmen	NL/H/1732/004	BE226755	PFIZER S.A. (BELGIUM)	BE
Serlain 50 mg filmomhulde tabletten	NL/H/1732/002	BE157324	PFIZER S.A. (BELGIUM)	BE
Serlain 100 mg filmomhulde tabletten	NL/H/1732/003	BE157351	PFIZER S.A. (BELGIUM)	BE
Serlain 20 mg/ml concentraat voor drank	NL/H/1732/004	BE226755	PFIZER S.A. (BELGIUM)	BE
Serlain 50 mg comprimés pelliculés	NL/H/1732/002	BE157324	PFIZER S.A. (BELGIUM)	BE
Serlain 20 mg/ml solution à diluer pour solution buvable	NL/H/1732/004	BE226755	PFIZER S.A. (BELGIUM)	BE
Serlain 100 mg comprimés pelliculés	NL/H/1732/003	BE157351	PFIZER S.A. (BELGIUM)	BE
Золофт 50 mg филмирани таблетки	NL/H/1732/002	20000514	PFIZER EUROPE MA EEIG	BG
Золофт 50 mg филмирани таблетки	NL/H/1732/002	20000514	PFIZER EUROPE MA EEIG	BG
Zoloft, 50 mg επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/1732/002	14677/ 5.4.1999	PFIZER HELLAS, A.E.	CY
Zoloft, 100 mg επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/1732/003	14678/ 5.4.1999	PFIZER HELLAS, A.E.	CY

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zoloft, 50 mg επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/1732/002	14677/ 5.4.1999	PFIZER HELLAS, A.E.	CY
Zoloft, 100 mg επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/1732/003	14678/ 5.4.1999	PFIZER HELLAS, A.E.	CY
Zoloft 100 mg potahované tablety	NL/H/1732/003	30/1093/94-B/C	PFIZER, SPOL. S R.O.	CZ
Zoloft 50 mg potahované tablety	NL/H/1732/002	30/1093/94-A/C	PFIZER, SPOL. S R.O.	CZ
Zoloft 100 mg potahované tablety	NL/H/1732/003	30/1093/94-B/C	PFIZER, SPOL. S R.O.	CZ
Zoloft 50 mg potahované tablety	NL/H/1732/002	30/1093/94-A/C	PFIZER, SPOL. S R.O.	CZ
Zoloft® 20 mg/ml Konzentrat zur Herstellung einer Lösung zum Einnehmen	NL/H/1732/004	46312.00.00	PFIZER PHARMA PFE GMBH	DE
Zoloft® 50 mg, Filmtabletten	NL/H/1732/002	37076.00.00	PFIZER PHARMA PFE GMBH	DE
Zoloft® 100 mg, Filmtabletten	NL/H/1732/003	37076.01.00	PFIZER PHARMA PFE GMBH	DE
Zoloft® 20 mg/ml Konzentrat zur Herstellung einer Lösung zum Einnehmen	NL/H/1732/004	46312.00.00	PFIZER PHARMA PFE GMBH	DE
Zoloft® 50 mg, Filmtabletten	NL/H/1732/002	37076.00.00	PFIZER PHARMA PFE GMBH	DE
Zoloft® 100 mg, Filmtabletten	NL/H/1732/003	37076.01.00	PFIZER PHARMA PFE GMBH	DE
Zoloft, filmovertrukne tabletter	NL/H/1732/001	32146	PFIZER APS	DK
Zoloft, filmovertrukne tabletter	NL/H/1732/002	13471	PFIZER APS	DK
Zoloft, filmovertrukne tabletter	NL/H/1732/003	13472	PFIZER APS	DK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zoloft, koncentrat til oral opløsning	NL/H/1732/004	30610	PFIZER APS	DK
Zoloft, filmovertrukne tabletter	NL/H/1732/001	32146	PFIZER APS	DK
Zoloft, filmovertrukne tabletter	NL/H/1732/002	13471	PFIZER APS	DK
Zoloft, filmovertrukne tabletter	NL/H/1732/003	13472	PFIZER APS	DK
Zoloft, koncentrat til oral opløsning	NL/H/1732/004	30610	PFIZER APS	DK
Zoloft, 50 mg õhukese polümeerikattega tabletid	NL/H/1732/002	122095	PFIZER EUROPE MA EEIG	EE
Zoloft, 50 mg õhukese polümeerikattega tabletid	NL/H/1732/002	122095	PFIZER EUROPE MA EEIG	EE
Besitran 50 mg comprimidos recubiertos con película	NL/H/1732/002	59.717	PFIZER, S.L.	ES
Besitran 100 mg comprimidos recubiertos con película	NL/H/1732/003	59.718	PFIZER, S.L.	ES
Besitran 20 mg/ml concentrado para solución oral	NL/H/1732/004	63.477	PFIZER, S.L.	ES
Besitran 50 mg comprimidos recubiertos con película	NL/H/1732/002	59.717	PFIZER, S.L.	ES
Besitran 100 mg comprimidos recubiertos con película	NL/H/1732/003	59.718	PFIZER, S.L.	ES
Besitran 20 mg/ml concentrado para solución oral	NL/H/1732/004	63.477	PFIZER, S.L.	ES
AREMIS 100 mg comprimidos recubiertos con película	not available	59.734	LABORATORIOS DR. ESTEVE S.A.	ES

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
AREMIS 50 mg comprimidos recubiertos con película	not available	59.735	LABORATORIOS DR. ESTEVE S.A.	ES
Zoloft 50 mg tabletti, kalvopäällysteinen	NL/H/1732/002	11557	PFIZER OY	FI
Zoloft 100 mg tabletti, kalvopäällysteinen	NL/H/1732/003	11558	PFIZER OY	FI
Zoloft 100 mg filmdragerade tabletter	NL/H/1732/003	11558	PFIZER OY	FI
Zoloft 50 mg filmdragerade tabletter	NL/H/1732/002	11557	PFIZER OY	FI
Zoloft 50 mg tabletti, kalvopäällysteinen	NL/H/1732/002	11557	PFIZER OY	FI
Zoloft 100 mg tabletti, kalvopäällysteinen	NL/H/1732/003	11558	PFIZER OY	FI
Zoloft 100 mg filmdragerade tabletter	NL/H/1732/003	11558	PFIZER OY	FI
Zoloft 50 mg filmdragerade tabletter	NL/H/1732/002	11557	PFIZER OY	FI
ZOLOFT 25 mg, gélule	not available	34009 355 621 2 8	PFIZER HOLDING FRANCE (S.C.A.)	FR
ZOLOFT 25 mg, gélule	not available	34009 563 118 8 0	PFIZER HOLDING FRANCE (S.C.A.)	FR
ZOLOFT 25 mg, gélule	not available	34009 563 119 4 1	PFIZER HOLDING FRANCE (S.C.A.)	FR
ZOLOFT 50 mg, gélule	not available	34009 300 379 3 5	PFIZER HOLDING FRANCE	FR
ZOLOFT 25 mg, gélule	not available	34009 355 620 6 7	PFIZER HOLDING FRANCE	FR
ZOLOFT 50 mg, gélule	not available	34009 340 341 9 0	PFIZER HOLDING FRANCE (S.C.A.)	FR
ZOLOFT 50 mg, gélule	not available	34009 340 344 8 0	PFIZER HOLDING FRANCE (S.C.A.)	FR
ZOLOFT 50 mg, gélule	not available	34009 340 342 5 1	PFIZER HOLDING FRANCE (S.C.A.)	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
ZOLOFT 50 mg, gélule	not available	34009 340 343 1 2	PFIZER HOLDING FRANCE (S.C.A.)	FR
Zoloft, 50 mg επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/1732/002	43084/23-06-2015	PFIZER HELLAS, A.E.	GR
Zoloft, 100 mg επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/1732/003	43085/23-06-2015	PFIZER HELLAS, A.E.	GR
Sertraline/Generics 100 mg, επικαλυμμένα με λεπτό υμένιο δισκία	EL/H/0177/002	12399/12-02-2013	GENERICS [UK] LIMITED	GR
Zoloft, 50 mg επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/1732/002	43084/23-06-2015	PFIZER HELLAS, A.E.	GR
Zoloft, 100 mg επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/1732/003	43085/23-06-2015	PFIZER HELLAS, A.E.	GR
ZOLOFT 50 mg filmom obložene tablete	not available	HR-H-153369585	PFIZER CROATIA D.O.O.	HR
Zoloft 50 mg filmtabletta	NL/H/1732/002	OGYI-T-4342/01	PFIZER KFT.	HU
Zoloft 20 mg/ml koncentrátum belsőleges oldathoz	NL/H/1732/004	OGYI-T-4342/02	PFIZER KFT.	HU
Zoloft 50 mg filmtabletta	NL/H/1732/002	OGYI-T-4342/01	PFIZER KFT.	HU
Zoloft 20 mg/ml koncentrátum belsőleges oldathoz	NL/H/1732/004	OGYI-T-4342/02	PFIZER KFT.	HU
LUSTRAL® 50 mg film coated tablets	NL/H/1732/002	PA 822/1/4	PFIZER HEALTHCARE IRELAND	IE
LUSTRAL® 100 mg film coated tablets	NL/H/1732/003	PA 822/1/5	PFIZER HEALTHCARE IRELAND	IE
LUSTRAL® 50 mg film coated tablets	NL/H/1732/002	PA 822/1/4	PFIZER HEALTHCARE IRELAND	IE
LUSTRAL® 100 mg film coated tablets	NL/H/1732/003	PA 822/1/5	PFIZER HEALTHCARE IRELAND	IE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zoloft 20 mg/ml mixtúruþykkni, lausn	NL/H/1732/004	990043 (IS)	PFIZER APS	IS
Zoloft 100 mg filmuhúðaðar töflur	NL/H/1732/003	990469 (IS)	PFIZER APS	IS
Zoloft 25 mg filmuhúðaðar töflur	NL/H/1732/001	IS/1/01/102/01	PFIZER APS	IS
Zoloft 50 mg filmuhúðaðar töflur	NL/H/1732/002	950077 (IS)	PFIZER APS	IS
Zoloft 20 mg/ml mixtúruþykkni, lausn	NL/H/1732/004	990043 (IS)	PFIZER APS	IS
Zoloft 100 mg filmuhúðaðar töflur	NL/H/1732/003	990469 (IS)	PFIZER APS	IS
Zoloft 25 mg filmuhúðaðar töflur	NL/H/1732/001	IS/1/01/102/01	PFIZER APS	IS
Zoloft 50 mg filmuhúðaðar töflur	NL/H/1732/002	950077 (IS)	PFIZER APS	IS
Zoloft compresse rivestite con film 25 mg	NL/H/1732/001	027753437	PFIZER ITALIA S.R.L.	IT
Zoloft concentrato per soluzione orale 20 mg/ml	NL/H/1732/004	027753096	PFIZER ITALIA S.R.L.	IT
Zoloft compresse rivestite con film 50 mg	NL/H/1732/002	027753108	PFIZER ITALIA S.R.L.	IT
Zoloft compresse rivestite con film 25 mg	NL/H/1732/001	027753437	PFIZER ITALIA S.R.L.	IT
Zoloft concentrato per soluzione orale 20 mg/ml	NL/H/1732/004	027753096	PFIZER ITALIA S.R.L.	IT
Zoloft compresse rivestite con film 50 mg	NL/H/1732/002	027753108	PFIZER ITALIA S.R.L.	IT
Tatig concentrato per soluzione orale 20 mg/ml	NL/H/1736/003	027754035	PFIZER ITALIA S.R.L.	IT
Tatig compresse rivestite con film 50 mg	NL/H/1736/001	027754086	PFIZER ITALIA S.R.L.	IT
Tatig compresse rivestite con film 50 mg	NL/H/1736/001	027754047	PFIZER ITALIA S.R.L.	IT
Tatig compresse rivestite con film 100 mg	NL/H/1736/002	027754050	PFIZER 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
Tatig compresse rivestite con film 100 mg	NL/H/1736/002	027754098	PFIZER ITALIA S.R.L.	IT
Zoloft 50 mg plėvele dengtos tabletės	NL/H/1732/002	LT/1/96/2037/005	PFIZER LIMITED	LT
Zoloft 50 mg plėvele dengtos tabletės	NL/H/1732/002	LT/1/96/2037/004	PFIZER LIMITED	LT
Zoloft 50 mg plėvele dengtos tabletės	NL/H/1732/002	LT/1/96/2037/008	PFIZER LIMITED	LT
Zoloft 50 mg plėvele dengtos tabletės	NL/H/1732/002	LT/1/96/2037/006	PFIZER LIMITED	LT
Zoloft 50 mg plėvele dengtos tabletės	NL/H/1732/002	LT/1/96/2037/010	PFIZER LIMITED	LT
Zoloft 50 mg plėvele dengtos tabletės	NL/H/1732/002	LT/1/96/2037/013	PFIZER LIMITED	LT
Zoloft 50 mg plėvele dengtos tabletės	NL/H/1732/002	LT/1/96/2037/012	PFIZER LIMITED	LT
Zoloft 50 mg plėvele dengtos tabletės	NL/H/1732/002	LT/1/96/2037/002	PFIZER LIMITED	LT
Zoloft 50 mg plėvele dengtos tabletės	NL/H/1732/002	LT/1/96/2037/016	PFIZER LIMITED	LT
Zoloft 50 mg plėvele dengtos tabletės	NL/H/1732/002	LT/1/96/2037/009	PFIZER LIMITED	LT
Zoloft 50 mg plėvele dengtos tabletės	NL/H/1732/002	LT/1/96/2037/001	PFIZER LIMITED	LT
Zoloft 50 mg plėvele dengtos tabletės	NL/H/1732/002	LT/1/96/2037/014	PFIZER LIMITED	LT
Zoloft 50 mg plėvele dengtos tabletės	NL/H/1732/002	LT/1/96/2037/003	PFIZER LIMITED	LT
Zoloft 50 mg plėvele dengtos tabletės	NL/H/1732/002	LT/1/96/2037/007	PFIZER LIMITED	LT
Zoloft 50 mg plėvele dengtos tabletės	NL/H/1732/002	LT/1/96/2037/015	PFIZER LIMITED	LT
Zoloft 50 mg plėvele dengtos tabletės	NL/H/1732/002	LT/1/96/2037/011	PFIZER LIMITED	LT
Zoloft 50 mg plėvele dengtos tabletės	NL/H/1732/002	LT/1/96/2037/005	PFIZER LIMITED	LT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zoloft 50 mg plėvele dengtos tabletės	NL/H/1732/002	LT/1/96/2037/004	PFIZER LIMITED	LT
Zoloft 50 mg plėvele dengtos tabletės	NL/H/1732/002	LT/1/96/2037/008	PFIZER LIMITED	LT
Zoloft 50 mg plėvele dengtos tabletės	NL/H/1732/002	LT/1/96/2037/006	PFIZER LIMITED	LT
Zoloft 50 mg plėvele dengtos tabletės	NL/H/1732/002	LT/1/96/2037/010	PFIZER LIMITED	LT
Zoloft 50 mg plėvele dengtos tabletės	NL/H/1732/002	LT/1/96/2037/013	PFIZER LIMITED	LT
Zoloft 50 mg plėvele dengtos tabletės	NL/H/1732/002	LT/1/96/2037/012	PFIZER LIMITED	LT
Zoloft 50 mg plėvele dengtos tabletės	NL/H/1732/002	LT/1/96/2037/002	PFIZER LIMITED	LT
Zoloft 50 mg plėvele dengtos tabletės	NL/H/1732/002	LT/1/96/2037/016	PFIZER LIMITED	LT
Zoloft 50 mg plėvele dengtos tabletės	NL/H/1732/002	LT/1/96/2037/009	PFIZER LIMITED	LT
Zoloft 50 mg plėvele dengtos tabletės	NL/H/1732/002	LT/1/96/2037/001	PFIZER LIMITED	LT
Zoloft 50 mg plėvele dengtos tabletės	NL/H/1732/002	LT/1/96/2037/014	PFIZER LIMITED	LT
Zoloft 50 mg plėvele dengtos tabletės	NL/H/1732/002	LT/1/96/2037/003	PFIZER LIMITED	LT
Zoloft 50 mg plėvele dengtos tabletės	NL/H/1732/002	LT/1/96/2037/007	PFIZER LIMITED	LT
Zoloft 50 mg plėvele dengtos tabletės	NL/H/1732/002	LT/1/96/2037/015	PFIZER LIMITED	LT
Zoloft 50 mg plėvele dengtos tabletės	NL/H/1732/002	LT/1/96/2037/011	PFIZER LIMITED	LT
Serlain 100 mg Filmtabletten	NL/H/1732/003	2011010936	PFIZER S.A. (BELGIUM)	LU
Serlain 50 mg Filmtabletten	NL/H/1732/002	2011010935	PFIZER S.A. (BELGIUM)	LU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Serlain 20 mg / ml Konzentrat zur Herstellung einer Lösung zum Einnehmen	NL/H/1732/004	2011010938	PFIZER S.A. (BELGIUM)	LU
Serlain 100 mg comprimés pelliculés	NL/H/1732/003	2011010936	PFIZER S.A. (BELGIUM)	LU
Serlain 50 mg comprimés pelliculés	NL/H/1732/002	2011010935	PFIZER S.A. (BELGIUM)	LU
Serlain 20 mg/ml solution à diluer pour solution buvable	NL/H/1732/004	2011010938	PFIZER S.A. (BELGIUM)	LU
Serlain 100 mg Filmtabletten	NL/H/1732/003	2011010936	PFIZER S.A. (BELGIUM)	LU
Serlain 50 mg Filmtabletten	NL/H/1732/002	2011010935	PFIZER S.A. (BELGIUM)	LU
Serlain 20 mg / ml Konzentrat zur Herstellung einer Lösung zum Einnehmen	NL/H/1732/004	2011010938	PFIZER S.A. (BELGIUM)	LU
Serlain 100 mg comprimés pelliculés	NL/H/1732/003	2011010936	PFIZER S.A. (BELGIUM)	LU
Serlain 50 mg comprimés pelliculés	NL/H/1732/002	2011010935	PFIZER S.A. (BELGIUM)	LU
Serlain 20 mg/ml solution à diluer pour solution buvable	NL/H/1732/004	2011010938	PFIZER S.A. (BELGIUM)	LU
Zoloft 50 mg apvalkotās tabletes	NL/H/1732/002	97-0311	PFIZER LIMITED	LV
Zoloft 50 mg apvalkotās tabletes	NL/H/1732/002	97-0311	PFIZER LIMITED	LV
Lustral 100 mg film coated tablets	NL/H/1732/003	MA 505/03202	PFIZER HELLAS, A.E.	MT
Lustral 50 mg film coated tablets	NL/H/1732/002	MA 505/03201	PFIZER HELLAS, A.E.	MT
Lustral 100 mg film coated tablets	NL/H/1732/003	MA 505/03202	PFIZER HELLAS, A.E.	MT

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
Lustral 50 mg film coated tablets	NL/H/1732/002	MA 505/03201	PFIZER HELLAS, A.E.	MT
Zoloft 20 mg/ml, concentraat voor oplossing voor oraal gebruik	NL/H/1732/004	RVG 24641	PFIZER B.V.	NL
Zoloft 50, filmomhulde tabletten 50 mg	NL/H/1732/002	RVG 16292	PFIZER B.V.	NL
Zoloft 25, filmomhulde tabletten 25 mg	NL/H/1732/001	RVG 105254	PFIZER B.V.	NL
Zoloft 100, filmomhulde tabletten 100 mg	NL/H/1732/003	RVG 105255	PFIZER B.V.	NL
Zoloft 20 mg/ml, concentraat voor oplossing voor oraal gebruik	NL/H/1732/004	RVG 24641	PFIZER B.V.	NL
Zoloft 50, filmomhulde tabletten 50 mg	NL/H/1732/002	RVG 16292	PFIZER B.V.	NL
Zoloft 25, filmomhulde tabletten 25 mg	NL/H/1732/001	RVG 105254	PFIZER B.V.	NL
Zoloft 100, filmomhulde tabletten 100 mg	NL/H/1732/003	RVG 105255	PFIZER B.V.	NL
Sertraline Pfizer 100 mg, filmomhulde tabletten	NL/H/1736/002	RVG 105382	PFIZER B.V.	NL
Sertraline Pfizer 50 mg, filmomhulde tabletten	NL/H/1736/001	RVG 105381	PFIZER B.V.	NL
Sertraline Pfizer 20 mg/ml, concentraat voor oplossing voor oraal gebruik	NL/H/1736/003	RVG 105383	PFIZER B.V.	NL
Sertraline Pfizer 25 mg, filmomhulde tabletten	NL/H/1736/004	RVG 106062	PFIZER B.V.	NL
Zoloft 50 mg filmdrasjerte tabletter	NL/H/1732/002	00-8203	PFIZER AS	NO
Zoloft 100 mg filmdrasjerte tabletter	NL/H/1732/003	00-8204	PFIZER AS	NO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zoloft 25 mg filmdrasjerte tabletter	NL/H/1732/001	00-7908	PFIZER AS	NO
Zoloft 20 mg/ml konsentrat til mikstur, oppløsning	NL/H/1732/004	99-6843	PFIZER AS	NO
Zoloft 50 mg filmdrasjerte tabletter	NL/H/1732/002	00-8203	PFIZER AS	NO
Zoloft 100 mg filmdrasjerte tabletter	NL/H/1732/003	00-8204	PFIZER AS	NO
Zoloft 25 mg filmdrasjerte tabletter	NL/H/1732/001	00-7908	PFIZER AS	NO
Zoloft 20 mg/ml konsentrat til mikstur, oppløsning	NL/H/1732/004	99-6843	PFIZER AS	NO
ZOLOFT, 50 mg, tabletki powlekane	NL/H/1732/002	7531	PFIZER EUROPE MA EEIG	PL
ZOLOFT, 100 mg, tabletki powlekane	NL/H/1732/003	7532	PFIZER EUROPE MA EEIG	PL
ZOLOFT, 50 mg, tabletki powlekane	NL/H/1732/002	7531	PFIZER EUROPE MA EEIG	PL
ZOLOFT, 100 mg, tabletki powlekane	NL/H/1732/003	7532	PFIZER EUROPE MA EEIG	PL
Zoloft 50 mg comprimidos revestidos por película	NL/H/1732/002	2182780	LABORATÓRIOS PFIZER, LDA.	PT
Zoloft 50 mg comprimidos revestidos por película	NL/H/1732/002	5854385	LABORATÓRIOS PFIZER, LDA.	PT
Zoloft 50 mg comprimidos revestidos por película	NL/H/1732/002	5830989	LABORATÓRIOS PFIZER, LDA.	PT
Zoloft 50 mg comprimidos revestidos por película	NL/H/1732/002	2182681	LABORATÓRIOS PFIZER, 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
Zoloft 100 mg comprimidos revestidos por película	NL/H/1732/003	5854484	LABORATÓRIOS PFIZER, LDA.	PT
Zoloft 100 mg comprimidos revestidos por película	NL/H/1732/003	2182889	LABORATÓRIOS PFIZER, LDA.	PT
Zoloft 100 mg comprimidos revestidos por película	NL/H/1732/003	2182988	LABORATÓRIOS PFIZER, LDA.	PT
Zoloft 20 mg/ml concentrado para solução oral	NL/H/1732/004	3268083	LABORATÓRIOS PFIZER, LDA.	PT
Zoloft 50 mg comprimidos revestidos por película	NL/H/1732/002	2182780	LABORATÓRIOS PFIZER, LDA.	PT
Zoloft 50 mg comprimidos revestidos por película	NL/H/1732/002	5854385	LABORATÓRIOS PFIZER, LDA.	PT
Zoloft 50 mg comprimidos revestidos por película	NL/H/1732/002	5830989	LABORATÓRIOS PFIZER, LDA.	PT
Zoloft 50 mg comprimidos revestidos por película	NL/H/1732/002	2182681	LABORATÓRIOS PFIZER, LDA.	PT
Zoloft 100 mg comprimidos revestidos por película	NL/H/1732/003	5854484	LABORATÓRIOS PFIZER, LDA.	PT
Zoloft 100 mg comprimidos revestidos por película	NL/H/1732/003	2182889	LABORATÓRIOS PFIZER, LDA.	PT
Zoloft 100 mg comprimidos revestidos por película	NL/H/1732/003	2182988	LABORATÓRIOS PFIZER, LDA.	PT
Zoloft 20 mg/ml concentrado para solução oral	NL/H/1732/004	3268083	LABORATÓRIOS PFIZER, 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
Zoloft 50 mg comprimate filmate	NL/H/1732/002	8227/2015/02	PFIZER EUROPE MA EEIG	RO
Zoloft 50 mg comprimate filmate	NL/H/1732/002	8227/2015/14	PFIZER EUROPE MA EEIG	RO
Zoloft 100 mg comprimate filmate	NL/H/1732/003	8228/2015/02	PFIZER EUROPE MA EEIG	RO
Zoloft 100 mg comprimate filmate	NL/H/1732/003	8228/2015/14	PFIZER EUROPE MA EEIG	RO
Zoloft 100 mg comprimate filmate	NL/H/1732/003	8228/2015/16	PFIZER EUROPE MA EEIG	RO
Zoloft 50 mg comprimate filmate	NL/H/1732/002	8227/2015/16	PFIZER EUROPE MA EEIG	RO
Zoloft 100 mg comprimate filmate	NL/H/1732/003	8228/2015/04	PFIZER EUROPE MA EEIG	RO
Zoloft 100 mg comprimate filmate	NL/H/1732/003	8228/2015/10	PFIZER EUROPE MA EEIG	RO
Zoloft 50 mg comprimate filmate	NL/H/1732/002	8227/2015/10	PFIZER EUROPE MA EEIG	RO
Zoloft 100 mg comprimate filmate	NL/H/1732/003	8228/2015/08	PFIZER EUROPE MA EEIG	RO
Zoloft 50 mg comprimate filmate	NL/H/1732/002	8227/2015/12	PFIZER EUROPE MA EEIG	RO
Zoloft 100 mg comprimate filmate	NL/H/1732/003	8228/2015/06	PFIZER EUROPE MA EEIG	RO
Zoloft 50 mg comprimate filmate	NL/H/1732/002	8227/2015/03	PFIZER EUROPE MA EEIG	RO
Zoloft 100 mg comprimate filmate	NL/H/1732/003	8228/2015/12	PFIZER EUROPE MA EEIG	RO
Zoloft 50 mg comprimate filmate	NL/H/1732/002	8227/2015/15	PFIZER EUROPE MA EEIG	RO
Zoloft 100 mg comprimate filmate	NL/H/1732/003	8228/2015/03	PFIZER EUROPE MA EEIG	RO
Zoloft 50 mg comprimate filmate	NL/H/1732/002	8227/2015/11	PFIZER EUROPE MA EEIG	RO
Zoloft 50 mg comprimate filmate	NL/H/1732/002	8227/2015/01	PFIZER EUROPE MA EEIG	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zoloft 100 mg comprimate filmate	NL/H/1732/003	8228/2015/01	PFIZER EUROPE MA EEIG	RO
Zoloft 50 mg comprimate filmate	NL/H/1732/002	8227/2015/13	PFIZER EUROPE MA EEIG	RO
Zoloft 100 mg comprimate filmate	NL/H/1732/003	8228/2015/07	PFIZER EUROPE MA EEIG	RO
Zoloft 100 mg comprimate filmate	NL/H/1732/003	8228/2015/11	PFIZER EUROPE MA EEIG	RO
Zoloft 100 mg comprimate filmate	NL/H/1732/003	8228/2015/09	PFIZER EUROPE MA EEIG	RO
Zoloft 100 mg comprimate filmate	NL/H/1732/003	8228/2015/15	PFIZER EUROPE MA EEIG	RO
Zoloft 100 mg comprimate filmate	NL/H/1732/003	8228/2015/05	PFIZER EUROPE MA EEIG	RO
Zoloft 20 mg/ml concentrat pentru soluţie orală	NL/H/1732/004	8229/2015/01	PFIZER EUROPE MA EEIG	RO
Zoloft 100 mg comprimate filmate	NL/H/1732/003	8228/2015/13	PFIZER EUROPE MA EEIG	RO
Zoloft 50 mg comprimate filmate	NL/H/1732/002	8227/2015/05	PFIZER EUROPE MA EEIG	RO
Zoloft 50 mg comprimate filmate	NL/H/1732/002	8227/2015/09	PFIZER EUROPE MA EEIG	RO
Zoloft 50 mg comprimate filmate	NL/H/1732/002	8227/2015/07	PFIZER EUROPE MA EEIG	RO
Zoloft 50 mg comprimate filmate	NL/H/1732/002	8227/2015/06	PFIZER EUROPE MA EEIG	RO
Zoloft 50 mg comprimate filmate	NL/H/1732/002	8227/2015/04	PFIZER EUROPE MA EEIG	RO
Zoloft 50 mg comprimate filmate	NL/H/1732/002	8227/2015/08	PFIZER EUROPE MA EEIG	RO
Zoloft 50 mg comprimate filmate	NL/H/1732/002	8227/2015/02	PFIZER EUROPE MA EEIG	RO
Zoloft 50 mg comprimate filmate	NL/H/1732/002	8227/2015/14	PFIZER EUROPE MA EEIG	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zoloft 100 mg comprimate filmate	NL/H/1732/003	8228/2015/02	PFIZER EUROPE MA EEIG	RO
Zoloft 100 mg comprimate filmate	NL/H/1732/003	8228/2015/14	PFIZER EUROPE MA EEIG	RO
Zoloft 100 mg comprimate filmate	NL/H/1732/003	8228/2015/16	PFIZER EUROPE MA EEIG	RO
Zoloft 50 mg comprimate filmate	NL/H/1732/002	8227/2015/16	PFIZER EUROPE MA EEIG	RO
Zoloft 100 mg comprimate filmate	NL/H/1732/003	8228/2015/04	PFIZER EUROPE MA EEIG	RO
Zoloft 100 mg comprimate filmate	NL/H/1732/003	8228/2015/10	PFIZER EUROPE MA EEIG	RO
Zoloft 50 mg comprimate filmate	NL/H/1732/002	8227/2015/10	PFIZER EUROPE MA EEIG	RO
Zoloft 100 mg comprimate filmate	NL/H/1732/003	8228/2015/08	PFIZER EUROPE MA EEIG	RO
Zoloft 50 mg comprimate filmate	NL/H/1732/002	8227/2015/12	PFIZER EUROPE MA EEIG	RO
Zoloft 100 mg comprimate filmate	NL/H/1732/003	8228/2015/06	PFIZER EUROPE MA EEIG	RO
Zoloft 50 mg comprimate filmate	NL/H/1732/002	8227/2015/03	PFIZER EUROPE MA EEIG	RO
Zoloft 100 mg comprimate filmate	NL/H/1732/003	8228/2015/12	PFIZER EUROPE MA EEIG	RO
Zoloft 50 mg comprimate filmate	NL/H/1732/002	8227/2015/15	PFIZER EUROPE MA EEIG	RO
Zoloft 100 mg comprimate filmate	NL/H/1732/003	8228/2015/03	PFIZER EUROPE MA EEIG	RO
Zoloft 50 mg comprimate filmate	NL/H/1732/002	8227/2015/11	PFIZER EUROPE MA EEIG	RO
Zoloft 50 mg comprimate filmate	NL/H/1732/002	8227/2015/01	PFIZER EUROPE MA EEIG	RO
Zoloft 100 mg comprimate filmate	NL/H/1732/003	8228/2015/01	PFIZER EUROPE MA EEIG	RO
Zoloft 50 mg comprimate filmate	NL/H/1732/002	8227/2015/13	PFIZER EUROPE MA EEIG	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zoloft 100 mg comprimate filmate	NL/H/1732/003	8228/2015/07	PFIZER EUROPE MA EEIG	RO
Zoloft 100 mg comprimate filmate	NL/H/1732/003	8228/2015/11	PFIZER EUROPE MA EEIG	RO
Zoloft 100 mg comprimate filmate	NL/H/1732/003	8228/2015/09	PFIZER EUROPE MA EEIG	RO
Zoloft 100 mg comprimate filmate	NL/H/1732/003	8228/2015/15	PFIZER EUROPE MA EEIG	RO
Zoloft 100 mg comprimate filmate	NL/H/1732/003	8228/2015/05	PFIZER EUROPE MA EEIG	RO
Zoloft 20 mg/ml concentrat pentru soluţie orală	NL/H/1732/004	8229/2015/01	PFIZER EUROPE MA EEIG	RO
Zoloft 100 mg comprimate filmate	NL/H/1732/003	8228/2015/13	PFIZER EUROPE MA EEIG	RO
Zoloft 50 mg comprimate filmate	NL/H/1732/002	8227/2015/05	PFIZER EUROPE MA EEIG	RO
Zoloft 50 mg comprimate filmate	NL/H/1732/002	8227/2015/09	PFIZER EUROPE MA EEIG	RO
Zoloft 50 mg comprimate filmate	NL/H/1732/002	8227/2015/07	PFIZER EUROPE MA EEIG	RO
Zoloft 50 mg comprimate filmate	NL/H/1732/002	8227/2015/06	PFIZER EUROPE MA EEIG	RO
Zoloft 50 mg comprimate filmate	NL/H/1732/002	8227/2015/04	PFIZER EUROPE MA EEIG	RO
Zoloft 50 mg comprimate filmate	NL/H/1732/002	8227/2015/08	PFIZER EUROPE MA EEIG	RO
Zoloft 100 mg filmdragerad tablett	NL/H/1732/003	12567	PFIZER AB	SE
Zoloft 50 mg filmdragerad tablett	NL/H/1732/002	12566	PFIZER AB	SE
Zoloft 25 mg filmdragerad tablett	NL/H/1732/001	16689	PFIZER AB	SE
Zoloft 20 mg/ml koncentrat till oral lösning	NL/H/1732/004	15504	PFIZER AB	SE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zoloft 100 mg filmdragerad tablett	NL/H/1732/003	12567	PFIZER AB	SE
Zoloft 50 mg filmdragerad tablett	NL/H/1732/002	12566	PFIZER AB	SE
Zoloft 25 mg filmdragerad tablett	NL/H/1732/001	16689	PFIZER AB	SE
Zoloft 20 mg/ml koncentrat till oral lösning	NL/H/1732/004	15504	PFIZER AB	SE
Zoloft 100 mg filmsko obložene tablete	NL/H/1732/003	H/94/01718/017	PFIZER LUXEMBOURG SARL	SI
Zoloft 100 mg filmsko obložene tablete	NL/H/1732/003	H/94/01718/032	PFIZER LUXEMBOURG SARL	SI
Zoloft 100 mg filmsko obložene tablete	NL/H/1732/003	H/94/01718/022	PFIZER LUXEMBOURG SARL	SI
Zoloft 100 mg filmsko obložene tablete	NL/H/1732/003	H/94/01718/018	PFIZER LUXEMBOURG SARL	SI
Zoloft 100 mg filmsko obložene tablete	NL/H/1732/003	H/94/01718/020	PFIZER LUXEMBOURG SARL	SI
Zoloft 100 mg filmsko obložene tablete	NL/H/1732/003	H/94/01718/025	PFIZER LUXEMBOURG SARL	SI
Zoloft 100 mg filmsko obložene tablete	NL/H/1732/003	H/94/01718/021	PFIZER LUXEMBOURG SARL	SI
Zoloft 100 mg filmsko obložene tablete	NL/H/1732/003	H/94/01718/028	PFIZER LUXEMBOURG SARL	SI
Zoloft 100 mg filmsko obložene tablete	NL/H/1732/003	H/94/01718/019	PFIZER LUXEMBOURG SARL	SI
Zoloft 50 mg filmsko obložene tablete	NL/H/1732/002	H/94/01718/002	PFIZER LUXEMBOURG SARL	SI
Zoloft 100 mg filmsko obložene tablete	NL/H/1732/003	H/94/01718/026	PFIZER LUXEMBOURG SARL	SI
Zoloft 100 mg filmsko obložene tablete	NL/H/1732/003	H/94/01718/024	PFIZER LUXEMBOURG SARL	SI
Zoloft 100 mg filmsko obložene tablete	NL/H/1732/003	H/94/01718/029	PFIZER LUXEMBOURG SARL	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/94/01718/027	PFIZER LUXEMBOURG SARL	SI
Zoloft 100 mg filmsko obložene tablete	NL/H/1732/003	H/94/01718/023	PFIZER LUXEMBOURG SARL	SI
Zoloft 100 mg filmsko obložene tablete	NL/H/1732/003	H/94/01718/031	PFIZER LUXEMBOURG SARL	SI
Zoloft 100 mg filmsko obložene tablete	NL/H/1732/003	H/94/01718/030	PFIZER LUXEMBOURG SARL	SI
Zoloft 50 mg filmsko obložene tablete	NL/H/1732/002	H/94/01718/001	PFIZER LUXEMBOURG SARL	SI
Zoloft 50 mg filmsko obložene tablete	NL/H/1732/002	H/94/01718/012	PFIZER LUXEMBOURG SARL	SI
Zoloft 50 mg filmsko obložene tablete	NL/H/1732/002	H/94/01718/006	PFIZER LUXEMBOURG SARL	SI
Zoloft 50 mg filmsko obložene tablete	NL/H/1732/002	H/94/01718/009	PFIZER LUXEMBOURG SARL	SI
Zoloft 50 mg filmsko obložene tablete	NL/H/1732/002	H/94/01718/011	PFIZER LUXEMBOURG SARL	SI
Zoloft 50 mg filmsko obložene tablete	NL/H/1732/002	H/94/01718/004	PFIZER LUXEMBOURG SARL	SI
Zoloft 50 mg filmsko obložene tablete	NL/H/1732/002	H/94/01718/016	PFIZER LUXEMBOURG SARL	SI
Zoloft 50 mg filmsko obložene tablete	NL/H/1732/002	H/94/01718/007	PFIZER LUXEMBOURG SARL	SI
Zoloft 50 mg filmsko obložene tablete	NL/H/1732/002	H/94/01718/015	PFIZER LUXEMBOURG SARL	SI
Zoloft 50 mg filmsko obložene tablete	NL/H/1732/002	H/94/01718/013	PFIZER LUXEMBOURG SARL	SI
Zoloft 50 mg filmsko obložene tablete	NL/H/1732/002	H/94/01718/008	PFIZER LUXEMBOURG SARL	SI
Zoloft 50 mg filmsko obložene tablete	NL/H/1732/002	H/94/01718/010	PFIZER LUXEMBOURG SARL	SI
Zoloft 50 mg filmsko obložene tablete	NL/H/1732/002	H/94/01718/014	PFIZER LUXEMBOURG SARL	SI
Zoloft 50 mg filmsko obložene tablete	NL/H/1732/002	H/94/01718/005	PFIZER LUXEMBOURG SARL	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 50 mg filmsko obložene tablete	NL/H/1732/002	H/94/01718/003	PFIZER LUXEMBOURG SARL	SI
Zoloft 20 mg/ml koncentrat za peroralno raztopino	NL/H/1732/004	H/94/01718/033	PFIZER LUXEMBOURG SARL	SI
Zoloft 100 mg filmsko obložene tablete	NL/H/1732/003	H/94/01718/017	PFIZER LUXEMBOURG SARL	SI
Zoloft 100 mg filmsko obložene tablete	NL/H/1732/003	H/94/01718/032	PFIZER LUXEMBOURG SARL	SI
Zoloft 100 mg filmsko obložene tablete	NL/H/1732/003	H/94/01718/022	PFIZER LUXEMBOURG SARL	SI
Zoloft 100 mg filmsko obložene tablete	NL/H/1732/003	H/94/01718/018	PFIZER LUXEMBOURG SARL	SI
Zoloft 100 mg filmsko obložene tablete	NL/H/1732/003	H/94/01718/020	PFIZER LUXEMBOURG SARL	SI
Zoloft 100 mg filmsko obložene tablete	NL/H/1732/003	H/94/01718/025	PFIZER LUXEMBOURG SARL	SI
Zoloft 100 mg filmsko obložene tablete	NL/H/1732/003	H/94/01718/021	PFIZER LUXEMBOURG SARL	SI
Zoloft 100 mg filmsko obložene tablete	NL/H/1732/003	H/94/01718/028	PFIZER LUXEMBOURG SARL	SI
Zoloft 100 mg filmsko obložene tablete	NL/H/1732/003	H/94/01718/019	PFIZER LUXEMBOURG SARL	SI
Zoloft 50 mg filmsko obložene tablete	NL/H/1732/002	H/94/01718/002	PFIZER LUXEMBOURG SARL	SI
Zoloft 100 mg filmsko obložene tablete	NL/H/1732/003	H/94/01718/026	PFIZER LUXEMBOURG SARL	SI
Zoloft 100 mg filmsko obložene tablete	NL/H/1732/003	H/94/01718/024	PFIZER LUXEMBOURG SARL	SI
Zoloft 100 mg filmsko obložene tablete	NL/H/1732/003	H/94/01718/029	PFIZER LUXEMBOURG SARL	SI
Zoloft 100 mg filmsko obložene tablete	NL/H/1732/003	H/94/01718/027	PFIZER LUXEMBOURG SARL	SI
Zoloft 100 mg filmsko obložene tablete	NL/H/1732/003	H/94/01718/023	PFIZER LUXEMBOURG SARL	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/94/01718/031	PFIZER LUXEMBOURG SARL	SI
Zoloft 100 mg filmsko obložene tablete	NL/H/1732/003	H/94/01718/030	PFIZER LUXEMBOURG SARL	SI
Zoloft 50 mg filmsko obložene tablete	NL/H/1732/002	H/94/01718/001	PFIZER LUXEMBOURG SARL	SI
Zoloft 50 mg filmsko obložene tablete	NL/H/1732/002	H/94/01718/012	PFIZER LUXEMBOURG SARL	SI
Zoloft 50 mg filmsko obložene tablete	NL/H/1732/002	H/94/01718/006	PFIZER LUXEMBOURG SARL	SI
Zoloft 50 mg filmsko obložene tablete	NL/H/1732/002	H/94/01718/009	PFIZER LUXEMBOURG SARL	SI
Zoloft 50 mg filmsko obložene tablete	NL/H/1732/002	H/94/01718/011	PFIZER LUXEMBOURG SARL	SI
Zoloft 50 mg filmsko obložene tablete	NL/H/1732/002	H/94/01718/004	PFIZER LUXEMBOURG SARL	SI
Zoloft 50 mg filmsko obložene tablete	NL/H/1732/002	H/94/01718/016	PFIZER LUXEMBOURG SARL	SI
Zoloft 50 mg filmsko obložene tablete	NL/H/1732/002	H/94/01718/007	PFIZER LUXEMBOURG SARL	SI
Zoloft 50 mg filmsko obložene tablete	NL/H/1732/002	H/94/01718/015	PFIZER LUXEMBOURG SARL	SI
Zoloft 50 mg filmsko obložene tablete	NL/H/1732/002	H/94/01718/013	PFIZER LUXEMBOURG SARL	SI
Zoloft 50 mg filmsko obložene tablete	NL/H/1732/002	H/94/01718/008	PFIZER LUXEMBOURG SARL	SI
Zoloft 50 mg filmsko obložene tablete	NL/H/1732/002	H/94/01718/010	PFIZER LUXEMBOURG SARL	SI
Zoloft 50 mg filmsko obložene tablete	NL/H/1732/002	H/94/01718/014	PFIZER LUXEMBOURG SARL	SI
Zoloft 50 mg filmsko obložene tablete	NL/H/1732/002	H/94/01718/005	PFIZER LUXEMBOURG SARL	SI
Zoloft 50 mg filmsko obložene tablete	NL/H/1732/002	H/94/01718/003	PFIZER LUXEMBOURG SARL	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 20 mg/ml koncentrat za peroralno raztopino	NL/H/1732/004	H/94/01718/033	PFIZER LUXEMBOURG SARL	SI
ZOLOFT OC 20 mg/ml koncentrát na perorálny roztok	NL/H/1732/004	30/0207/01–S	PFIZER EUROPE MA EEIG	SK
ZOLOFT 100 mg filmom obalené tablety	NL/H/1732/003	30/0430/10–S	PFIZER EUROPE MA EEIG	SK
ZOLOFT 50 mg filmom obalené tablety	NL/H/1732/002	30/0090/96–S	PFIZER EUROPE MA EEIG	SK
ZOLOFT OC 20 mg/ml koncentrát na perorálny roztok	NL/H/1732/004	30/0207/01–S	PFIZER EUROPE MA EEIG	SK
ZOLOFT 100 mg filmom obalené tablety	NL/H/1732/003	30/0430/10–S	PFIZER EUROPE MA EEIG	SK
ZOLOFT 50 mg filmom obalené tablety	NL/H/1732/002	30/0090/96–S	PFIZER EUROPE MA EEIG	SK
LUSTRAL® 50 mg film coated tablets	NL/H/1732/002	PL 00057/0308	PFIZER LIMITED	UK
LUSTRAL® 100 mg film coated tablets	NL/H/1732/003	PL 0057/0309	PFIZER LIMITED	UK
Sertraline 50 mg Film-coated Tablets	not available	PL 44041/0031	NOUMED LIFE SCIENCES	UK
Sertraline 100mg Film-coated Tablets	not available	PL 44041/0032	NOUMED LIFE SCIENCES	UK
LUSTRAL® 50 mg film coated tablets	NL/H/1732/002	PL 00057/0308	PFIZER LIMITED	UK
LUSTRAL® 100 mg film coated tablets	NL/H/1732/003	PL 0057/0309	PFIZER LIMITED	UK
Sertraline 50 mg Tablets	not available	PL 21880/0097	MEDREICH PLC	UK
Sertraline 100mg Tablets	not available	PL 21880/0098	MEDREICH PLC	UK
Sertraline 50 mg film-coated tablet	not available	PL 11311/0491	TILLOMED LABORATORIES LTD	UK
Sertraline 100 mg film-coated tablet	not available	PL 11311/0492	TILLOMED LABORATORIES 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
Sertraline 100 mg film coated tablets	NL/H/1736/002	PL 00057/1205	PFIZER LIMITED	UK
Sertraline 50 mg film coated tablets	NL/H/1736/001	PL 00057/1204	PFIZER LIMITED	UK