EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

11 July 2019
EMA/408178/2019
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: naltrexone

Procedure no.: PSUSA/00002117/201811

Official address Domenico Scarlattilaan 6 ● 1083 HS Amsterdam ● The Netherlands
Address for visits and deliveries Refer to www.ema.europa.eu/how-to-find-us
Send us a question Go to www.ema.europa.eu/contact **Telephone** +31 (0)88 781 6000 An agency of the European Union

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
Antaxon 100 mg belsőleges oldat	not available	OGYI-T-7922/03	ZAMBON S.P.A.	HU
Antaxon 50 mg belsőleges oldat	not available	OGYI-T-7922/02	ZAMBON S.P.A.	HU
Antaxon 50 mg kemény kapszula	not available	OGYI-T-7922/01	ZAMBON S.P.A.	HU
ANTAXONE 100 mg/20 ml soluzione orale	not available	025855040	ZAMBON ITALIA S.R.L.	IT
ANTAXONE 50 mg capsule rigide	not available	025855077	ZAMBON ITALIA S.R.L.	IT
ANTAXONE 50 mg/10 ml soluzione orale	not available	025855065	ZAMBON ITALIA S.R.L.	IT
NALOREX 50 mg compresse rivestite con film	not available	025969039	L. MOLTENI AND C. DEI F.LLI ALITTI SOCIETÀ DI ESERCIZIO S.P.A.	IT
Revia 50 mg filmsko obložene tablete	not available	H/98/01345/001	CHIESI PHARMACEUTICALS GMBH	SI
REVIA 50 mg, comprimé pelliculé sécable	not available	34009 398 972 1 9	BRISTOL-MYERS SQUIBB	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
REVIA 50 mg, comprimé pelliculé sécable	not available	34009 341 751-6 9	BRISTOL-MYERS SQUIBB	FR
ReVia® 50 mg Filmtabletten	not available	1-18743	CHIESI PHARMACEUTICALS GMBH	AT