30 October 2025
EMADOC-1700519818-2594276
Human Medicines Division

List of nationally authorised medicinal products

Active substance(s): alprazolam

EURD list No. PSUSA/00000109/202503

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
Alpraz 0,5 mg comprimés	not available	BE165751	LABORATOIRES SMB S.A.	BE
Alpraz 0,5 mg comprimés	not available	BE165751	LABORATOIRES SMB S.A.	BE
Alpraz 0,5 mg comprimés	not available	BE165751	LABORATOIRES SMB S.A.	BE
Alpraz 0,5 mg comprimés	not available	BE165751	LABORATOIRES SMB S.A.	BE
Alpraz 0,5 mg comprimés	not available	BE165751	LABORATOIRES SMB S.A.	BE
Alpraz 0,5 mg comprimés	not available	BE165751	LABORATOIRES SMB S.A.	BE
Alpraz 0,5 mg comprimés	not available	BE165751	LABORATOIRES SMB S.A.	BE
Alpraz 0,5 mg comprimés	not available	1994120670	LABORATOIRES SMB S.A.	LU
Alpraz 0,5 mg comprimés	not available	1994120670	LABORATOIRES SMB S.A.	LU
Alpraz 0,5 mg comprimés	not available	1994120670	LABORATOIRES SMB S.A.	LU
Alpraz 0,5 mg comprimés	not available	1994120670	LABORATOIRES SMB S.A.	LU
Alpraz 0,5 mg comprimés	not available	1994120670	LABORATOIRES SMB S.A.	LU
Alpraz 0,5 mg comprimés	not available	1994120670	LABORATOIRES SMB S.A.	LU
Alpraz 0,5 mg comprimés	not available	1994120670	LABORATOIRES SMB S.A.	LU
Alpraz 0,5 mg tabletten	not available	BE165751	LABORATOIRES SMB S.A.	BE
Alpraz 0,5 mg Tabletten	not available	BE165751	LABORATOIRES SMB S.A.	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
Alpraz 0,5 mg tabletten	not available	BE165751	LABORATOIRES SMB S.A.	BE
Alpraz 0,5 mg Tabletten	not available	BE165751	LABORATOIRES SMB S.A.	BE
Alpraz 0,5 mg tabletten	not available	BE165751	LABORATOIRES SMB S.A.	BE
Alpraz 0,5 mg Tabletten	not available	BE165751	LABORATOIRES SMB S.A.	BE
Alpraz 0,5 mg tabletten	not available	BE165751	LABORATOIRES SMB S.A.	BE
Alpraz 0,5 mg Tabletten	not available	BE165751	LABORATOIRES SMB S.A.	BE
Alpraz 0,5 mg tabletten	not available	BE165751	LABORATOIRES SMB S.A.	BE
Alpraz 0,5 mg Tabletten	not available	BE165751	LABORATOIRES SMB S.A.	BE
Alpraz 0,5 mg tabletten	not available	BE165751	LABORATOIRES SMB S.A.	BE
Alpraz 0,5 mg Tabletten	not available	BE165751	LABORATOIRES SMB S.A.	BE
Alpraz 0,5 mg tabletten	not available	BE165751	LABORATOIRES SMB S.A.	BE
Alpraz 0,5 mg Tabletten	not available	BE165751	LABORATOIRES SMB S.A.	BE
Alpraz 0,5 mg Tabletten	not available	1994120670	LABORATOIRES SMB S.A.	LU
Alpraz 0,5 mg Tabletten	not available	1994120670	LABORATOIRES SMB S.A.	LU
Alpraz 0,5 mg Tabletten	not available	1994120670	LABORATOIRES SMB S.A.	LU
Alpraz 0,5 mg Tabletten	not available	1994120670	LABORATOIRES SMB S.A.	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
Alpraz 0,5 mg Tabletten	not available	1994120670	LABORATOIRES SMB S.A.	LU
Alpraz 0,5 mg Tabletten	not available	1994120670	LABORATOIRES SMB S.A.	LU
Alpraz 0,5 mg Tabletten	not available	1994120670	LABORATOIRES SMB S.A.	LU
Alpraz 1 mg comprimés	not available	BE165767	LABORATOIRES SMB S.A.	BE
Alpraz 1 mg comprimés	not available	BE165767	LABORATOIRES SMB S.A.	BE
Alpraz 1 mg comprimés	not available	BE165767	LABORATOIRES SMB S.A.	BE
Alpraz 1 mg comprimés	not available	BE165767	LABORATOIRES SMB S.A.	BE
Alpraz 1 mg comprimés	not available	BE165767	LABORATOIRES SMB S.A.	BE
Alpraz 1 mg comprimés	not available	BE165767	LABORATOIRES SMB S.A.	BE
Alpraz 1 mg comprimés	not available	BE165767	LABORATOIRES SMB S.A.	BE
Alpraz 1 mg comprimés	not available	1994120671	LABORATOIRES SMB S.A.	LU
Alpraz 1 mg comprimés	not available	1994120671	LABORATOIRES SMB S.A.	LU
Alpraz 1 mg comprimés	not available	1994120671	LABORATOIRES SMB S.A.	LU
Alpraz 1 mg comprimés	not available	1994120671	LABORATOIRES SMB S.A.	LU
Alpraz 1 mg comprimés	not available	1994120671	LABORATOIRES SMB S.A.	LU
Alpraz 1 mg comprimés	not available	1994120671	LABORATOIRES SMB S.A.	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
Alpraz 1 mg comprimés	not available	1994120671	LABORATOIRES SMB S.A.	LU
Alpraz 1 mg tabletten	not available	BE165767	LABORATOIRES SMB S.A.	BE
Alpraz 1 mg Tabletten	not available	BE165767	LABORATOIRES SMB S.A.	BE
Alpraz 1 mg tabletten	not available	BE165767	LABORATOIRES SMB S.A.	BE
Alpraz 1 mg Tabletten	not available	BE165767	LABORATOIRES SMB S.A.	BE
Alpraz 1 mg tabletten	not available	BE165767	LABORATOIRES SMB S.A.	BE
Alpraz 1 mg Tabletten	not available	BE165767	LABORATOIRES SMB S.A.	BE
Alpraz 1 mg tabletten	not available	BE165767	LABORATOIRES SMB S.A.	BE
Alpraz 1 mg Tabletten	not available	BE165767	LABORATOIRES SMB S.A.	BE
Alpraz 1 mg tabletten	not available	BE165767	LABORATOIRES SMB S.A.	BE
Alpraz 1 mg Tabletten	not available	BE165767	LABORATOIRES SMB S.A.	BE
Alpraz 1 mg tabletten	not available	BE165767	LABORATOIRES SMB S.A.	BE
Alpraz 1 mg Tabletten	not available	BE165767	LABORATOIRES SMB S.A.	BE
Alpraz 1 mg tabletten	not available	BE165767	LABORATOIRES SMB S.A.	BE
Alpraz 1 mg Tabletten	not available	BE165767	LABORATOIRES SMB S.A.	BE
Alpraz 1 mg Tabletten	not available	1994120671	LABORATOIRES SMB S.A.	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
Alpraz 1 mg Tabletten	not available	1994120671	LABORATOIRES SMB S.A.	LU
Alpraz 1 mg Tabletten	not available	1994120671	LABORATOIRES SMB S.A.	LU
Alpraz 1 mg Tabletten	not available	1994120671	LABORATOIRES SMB S.A.	LU
Alpraz 1 mg Tabletten	not available	1994120671	LABORATOIRES SMB S.A.	LU
Alpraz 1 mg Tabletten	not available	1994120671	LABORATOIRES SMB S.A.	LU
Alpraz 1 mg Tabletten	not available	1994120671	LABORATOIRES SMB S.A.	LU
Alprazolam ”Upjohn”, tabletter	not available	14213	UPJOHN EESV	DK
Alprazolam ”Upjohn”, tabletter	not available	14630	UPJOHN EESV	DK
Alprazolam ”Upjohn”, tabletter	not available	14214	UPJOHN EESV	DK
Alprazolam ”Upjohn”, tabletter	not available	14213	UPJOHN EESV	DK
Alprazolam ”Upjohn”, tabletter	not available	14630	UPJOHN EESV	DK
Alprazolam ”Upjohn”, tabletter	not available	14214	UPJOHN EESV	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
Alprazolam ”Upjohn”, tabletter	not available	14213	UPJOHN EESV	DK
Alprazolam ”Upjohn”, tabletter	not available	14630	UPJOHN EESV	DK
Alprazolam ”Upjohn”, tabletter	not available	14214	UPJOHN EESV	DK
Alprazolam ”Upjohn”, tabletter	not available	14213	UPJOHN EESV	DK
Alprazolam ”Upjohn”, tabletter	not available	14630	UPJOHN EESV	DK
Alprazolam ”Upjohn”, tabletter	not available	14214	UPJOHN EESV	DK
Alprazolam ”Upjohn”, tabletter	not available	14213	UPJOHN EESV	DK
Alprazolam ”Upjohn”, tabletter	not available	14630	UPJOHN EESV	DK
Alprazolam ”Upjohn”, tabletter	not available	14214	UPJOHN EESV	DK
Alprazolam AbZ 0,25 mg	NL/H/0338/001	54833.00.00	ABZ-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
Tabletten				
Alprazolam AbZ 0,25 mg Tabletten	NL/H/0338/001	54833.00.00	ABZ-PHARMA GMBH	DE
Alprazolam AbZ 0,25 mg Tabletten	NL/H/0338/001	54833.00.00	ABZ-PHARMA GMBH	DE
Alprazolam AbZ 0,25 mg Tabletten	NL/H/0338/001	54833.00.00	ABZ-PHARMA GMBH	DE
Alprazolam AbZ 0,25 mg Tabletten	NL/H/0338/001	54833.00.00	ABZ-PHARMA GMBH	DE
Alprazolam AbZ 0,25 mg Tabletten	NL/H/0338/001	54833.00.00	ABZ-PHARMA GMBH	DE
Alprazolam AbZ 0,25 mg Tabletten	NL/H/0338/001	54833.00.00	ABZ-PHARMA GMBH	DE
Alprazolam AbZ 0,25 mg Tabletten	NL/H/0338/001	54833.00.00	ABZ-PHARMA GMBH	DE
Alprazolam AbZ 0,25 mg Tabletten	NL/H/0338/001	54833.00.00	ABZ-PHARMA GMBH	DE
Alprazolam AbZ 0,25 mg Tabletten	NL/H/0338/001	54833.00.00	ABZ-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
Alprazolam AbZ 0,25 mg Tabletten	NL/H/0338/001	54833.00.00	ABZ-PHARMA GMBH	DE
Alprazolam AbZ 0,25 mg Tabletten	NL/H/0338/001	54833.00.00	ABZ-PHARMA GMBH	DE
Alprazolam AbZ 0,25 mg Tabletten	NL/H/0338/001	54833.00.00	ABZ-PHARMA GMBH	DE
Alprazolam AbZ 0,25 mg Tabletten	NL/H/0338/001	54833.00.00	ABZ-PHARMA GMBH	DE
Alprazolam AbZ 0,25 mg Tabletten	NL/H/0338/001	54833.00.00	ABZ-PHARMA GMBH	DE
Alprazolam AbZ 0,25 mg Tabletten	NL/H/0338/001	54833.00.00	ABZ-PHARMA GMBH	DE
Alprazolam AbZ 0,25 mg Tabletten	NL/H/0338/001	54833.00.00	ABZ-PHARMA GMBH	DE
Alprazolam AbZ 0,25 mg Tabletten	NL/H/0338/001	54833.00.00	ABZ-PHARMA GMBH	DE
Alprazolam AbZ 0,25 mg Tabletten	NL/H/0338/001	54833.00.00	ABZ-PHARMA GMBH	DE
Alprazolam AbZ 0,25 mg	NL/H/0338/001	54833.00.00	ABZ-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
Tabletten				
Alprazolam AbZ 0,25 mg Tabletten	NL/H/0338/001	54833.00.00	ABZ-PHARMA GMBH	DE
Alprazolam AbZ 0,25 mg Tabletten	NL/H/0338/001	54833.00.00	ABZ-PHARMA GMBH	DE
Alprazolam AbZ 0,25 mg Tabletten	NL/H/0338/001	54833.00.00	ABZ-PHARMA GMBH	DE
Alprazolam AbZ 0,25 mg Tabletten	NL/H/0338/001	54833.00.00	ABZ-PHARMA GMBH	DE
Alprazolam AbZ 0,25 mg Tabletten	NL/H/0338/001	54833.00.00	ABZ-PHARMA GMBH	DE
Alprazolam AbZ 0,25 mg Tabletten	NL/H/0338/001	54833.00.00	ABZ-PHARMA GMBH	DE
Alprazolam AbZ 0,25 mg Tabletten	NL/H/0338/001	54833.00.00	ABZ-PHARMA GMBH	DE
Alprazolam AbZ 0,25 mg Tabletten	NL/H/0338/001	54833.00.00	ABZ-PHARMA GMBH	DE
Alprazolam AbZ 0,25 mg Tabletten	NL/H/0338/001	54833.00.00	ABZ-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
Alprazolam AbZ 0,25 mg Tabletten	NL/H/0338/001	54833.00.00	ABZ-PHARMA GMBH	DE
Alprazolam AbZ 0,25 mg Tabletten	NL/H/0338/001	54833.00.00	ABZ-PHARMA GMBH	DE
Alprazolam AbZ 0,25 mg Tabletten	NL/H/0338/001	54833.00.00	ABZ-PHARMA GMBH	DE
Alprazolam AbZ 0,25 mg Tabletten	NL/H/0338/001	54833.00.00	ABZ-PHARMA GMBH	DE
Alprazolam IBSA 0,25 mg drank in verpakking voor eenmalig gebruik	BE/H/0344/001	BE660580	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,25 mg drank in verpakking voor eenmalig gebruik	BE/H/0344/001	BE660580	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,25 mg drank in verpakking voor eenmalig gebruik	BE/H/0344/001	BE660580	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,25 mg drank in verpakking voor eenmalig gebruik	BE/H/0344/001	BE660580	IBSA FARMACEUTICI ITALIA	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
Alprazolam IBSA 0,25 mg drank in verpakking voor eenmalig gebruik	BE/H/0344/001	BE660580	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,25 mg drank in verpakking voor eenmalig gebruik	BE/H/0344/001	BE660580	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,25 mg drank in verpakking voor eenmalig gebruik	BE/H/0344/001	BE660580	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,25 mg drank in verpakking voor eenmalig gebruik	BE/H/0344/001	BE660580	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,25 mg drank in verpakking voor eenmalig gebruik	BE/H/0344/001	BE660580	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,25 mg drank in verpakking voor eenmalig gebruik	BE/H/0344/001	BE660580	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,25 mg drank in verpakking voor eenmalig gebruik	BE/H/0344/001	BE660580	IBSA FARMACEUTICI ITALIA	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
Alprazolam IBSA 0,25 mg drank in verpakking voor eenmalig gebruik	BE/H/0344/001	BE660580	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,25 mg drank in verpakking voor eenmalig gebruik	BE/H/0344/001	BE660580	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,25 mg drank in verpakking voor eenmalig gebruik	BE/H/0344/001	BE660580	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,25 mg solution buvable en récipient unidose	BE/H/0344/001	BE660580	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,25 mg solution buvable en récipient unidose	BE/H/0344/001	BE660580	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,25 mg solution buvable en récipient unidose	BE/H/0344/001	BE660580	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,25 mg solution buvable en récipient unidose	BE/H/0344/001	BE660580	IBSA FARMACEUTICI ITALIA	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
Alprazolam IBSA 0,25 mg solution buvable en récipient unidose	BE/H/0344/001	BE660580	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,25 mg solution buvable en récipient unidose	BE/H/0344/001	BE660580	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,25 mg solution buvable en récipient unidose	BE/H/0344/001	BE660580	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,25 mg solution buvable en récipient unidose	BE/H/0344/001	BE660580	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,25 mg solution buvable en récipient unidose	BE/H/0344/001	BE660580	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,25 mg solution buvable en récipient unidose	BE/H/0344/001	BE660580	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,25 mg solution buvable en récipient unidose	BE/H/0344/001	BE660580	IBSA FARMACEUTICI ITALIA	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
Alprazolam IBSA 0,25 mg solution buvable en récipient unidose	BE/H/0344/001	BE660580	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,25 mg solution buvable en récipient unidose	BE/H/0344/001	BE660580	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,25 mg solution buvable en récipient unidose	BE/H/0344/001	BE660580	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,25 mg solution buvable en récipient unidose	BE/H/0344/001	2022110229	IBSA FARMACEUTICI ITALIA	LU
Alprazolam IBSA 0,25 mg solution buvable en récipient unidose	BE/H/0344/001	2022110229	IBSA FARMACEUTICI ITALIA	LU
Alprazolam IBSA 0,25 mg solution buvable en récipient unidose	BE/H/0344/001	2022110229	IBSA FARMACEUTICI ITALIA	LU
Alprazolam IBSA 0,25 mg solution buvable en récipient unidose	BE/H/0344/001	2022110229	IBSA FARMACEUTICI ITALIA	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**
Alprazolam IBSA 0,25 mg solution buvable en récipient unidose	BE/H/0344/001	2022110229	IBSA FARMACEUTICI ITALIA	LU
Alprazolam IBSA 0,25 mg solution buvable en récipient unidose	BE/H/0344/001	2022110229	IBSA FARMACEUTICI ITALIA	LU
Alprazolam IBSA 0,25 mg solution buvable en récipient unidose	BE/H/0344/001	2022110229	IBSA FARMACEUTICI ITALIA	LU
Alprazolam IBSA 0,25 mg solution buvable en récipient unidose	BE/H/0344/001	2022110229	IBSA FARMACEUTICI ITALIA	LU
Alprazolam IBSA 0,25 mg solution buvable en récipient unidose	BE/H/0344/001	2022110229	IBSA FARMACEUTICI ITALIA	LU
Alprazolam IBSA 0,25 mg solution buvable en récipient unidose	BE/H/0344/001	2022110229	IBSA FARMACEUTICI ITALIA	LU
Alprazolam IBSA 0,25 mg solution buvable en récipient unidose	BE/H/0344/001	2022110229	IBSA FARMACEUTICI ITALIA	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
Alprazolam IBSA 0,25 mg solution buvable en récipient unidose	BE/H/0344/001	2022110229	IBSA FARMACEUTICI ITALIA	LU
Alprazolam IBSA 0,25 mg solution buvable en récipient unidose	BE/H/0344/001	2022110229	IBSA FARMACEUTICI ITALIA	LU
Alprazolam IBSA 0,25 mg solution buvable en récipient unidose	BE/H/0344/001	2022110229	IBSA FARMACEUTICI ITALIA	LU
Alprazolam IBSA 0,25 mg soluzione orale in contenitore monodose	BE/H/0344/001	049508017	IBSA FARMACEUTICI ITALIA	IT
Alprazolam IBSA 0,25 mg soluzione orale in contenitore monodose	BE/H/0344/001	049508029	IBSA FARMACEUTICI ITALIA	IT
Alprazolam IBSA 0,25 mg soluzione orale in contenitore monodose	BE/H/0344/001	049508017	IBSA FARMACEUTICI ITALIA	IT
Alprazolam IBSA 0,25 mg soluzione orale in contenitore monodose	BE/H/0344/001	049508029	IBSA FARMACEUTICI ITALIA	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**
Alprazolam IBSA 0,25 mg soluzione orale in contenitore monodose	BE/H/0344/001	049508017	IBSA FARMACEUTICI ITALIA	IT
Alprazolam IBSA 0,25 mg soluzione orale in contenitore monodose	BE/H/0344/001	049508029	IBSA FARMACEUTICI ITALIA	IT
Alprazolam IBSA 0,25 mg soluzione orale in contenitore monodose	BE/H/0344/001	049508017	IBSA FARMACEUTICI ITALIA	IT
Alprazolam IBSA 0,25 mg soluzione orale in contenitore monodose	BE/H/0344/001	049508029	IBSA FARMACEUTICI ITALIA	IT
Alprazolam IBSA 0,25 mg soluzione orale in contenitore monodose	BE/H/0344/001	049508017	IBSA FARMACEUTICI ITALIA	IT
Alprazolam IBSA 0,25 mg soluzione orale in contenitore monodose	BE/H/0344/001	049508029	IBSA FARMACEUTICI ITALIA	IT
Alprazolam IBSA 0,25 mg soluzione orale in contenitore monodose	BE/H/0344/001	049508017	IBSA FARMACEUTICI ITALIA	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
Alprazolam IBSA 0,25 mg soluzione orale in contenitore monodose	BE/H/0344/001	049508029	IBSA FARMACEUTICI ITALIA	IT
Alprazolam IBSA 0,25 mg soluzione orale in contenitore monodose	BE/H/0344/001	049508017	IBSA FARMACEUTICI ITALIA	IT
Alprazolam IBSA 0,25 mg soluzione orale in contenitore monodose	BE/H/0344/001	049508029	IBSA FARMACEUTICI ITALIA	IT
Alprazolam IBSA 0,25 Milligramm Lösung zum Einnehmen im Einzeldosisbehältnis	BE/H/0344/001	BE660580	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,25 Milligramm Lösung zum Einnehmen im Einzeldosisbehältnis	BE/H/0344/001	BE660580	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,25 Milligramm Lösung zum Einnehmen im Einzeldosisbehältnis	BE/H/0344/001	BE660580	IBSA FARMACEUTICI ITALIA	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**
Alprazolam IBSA 0,25 Milligramm Lösung zum Einnehmen im Einzeldosisbehältnis	BE/H/0344/001	BE660580	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,25 Milligramm Lösung zum Einnehmen im Einzeldosisbehältnis	BE/H/0344/001	BE660580	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,25 Milligramm Lösung zum Einnehmen im Einzeldosisbehältnis	BE/H/0344/001	BE660580	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,25 Milligramm Lösung zum Einnehmen im Einzeldosisbehältnis	BE/H/0344/001	BE660580	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,25 Milligramm Lösung zum Einnehmen im Einzeldosisbehältnis	BE/H/0344/001	BE660580	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,25 Milligramm Lösung zum	BE/H/0344/001	BE660580	IBSA FARMACEUTICI ITALIA	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
Einnehmen im Einzeldosisbehältnis				
Alprazolam IBSA 0,25 Milligramm Lösung zum Einnehmen im Einzeldosisbehältnis	BE/H/0344/001	BE660580	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,25 Milligramm Lösung zum Einnehmen im Einzeldosisbehältnis	BE/H/0344/001	BE660580	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,25 Milligramm Lösung zum Einnehmen im Einzeldosisbehältnis	BE/H/0344/001	BE660580	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,25 Milligramm Lösung zum Einnehmen im Einzeldosisbehältnis	BE/H/0344/001	BE660580	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,25 Milligramm Lösung zum Einnehmen im Einzeldosisbehältnis	BE/H/0344/001	BE660580	IBSA FARMACEUTICI ITALIA	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
Alprazolam IBSA 0,50 mg drank in verpakking voor eenmalig gebruik	BE/H/0344/002	BE660581	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,50 mg drank in verpakking voor eenmalig gebruik	BE/H/0344/002	BE660581	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,50 mg drank in verpakking voor eenmalig gebruik	BE/H/0344/002	BE660581	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,50 mg drank in verpakking voor eenmalig gebruik	BE/H/0344/002	BE660581	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,50 mg drank in verpakking voor eenmalig gebruik	BE/H/0344/002	BE660581	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,50 mg drank in verpakking voor eenmalig gebruik	BE/H/0344/002	BE660581	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,50 mg drank in verpakking voor eenmalig gebruik	BE/H/0344/002	BE660581	IBSA FARMACEUTICI ITALIA	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
Alprazolam IBSA 0,50 mg drank in verpakking voor eenmalig gebruik	BE/H/0344/002	BE660581	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,50 mg drank in verpakking voor eenmalig gebruik	BE/H/0344/002	BE660581	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,50 mg drank in verpakking voor eenmalig gebruik	BE/H/0344/002	BE660581	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,50 mg drank in verpakking voor eenmalig gebruik	BE/H/0344/002	BE660581	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,50 mg drank in verpakking voor eenmalig gebruik	BE/H/0344/002	BE660581	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,50 mg drank in verpakking voor eenmalig gebruik	BE/H/0344/002	BE660581	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,50 mg drank in verpakking voor eenmalig gebruik	BE/H/0344/002	BE660581	IBSA FARMACEUTICI ITALIA	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
Alprazolam IBSA 0,50 mg solution buvable en récipient unidose	BE/H/0344/002	BE660581	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,50 mg solution buvable en récipient unidose	BE/H/0344/002	BE660581	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,50 mg solution buvable en récipient unidose	BE/H/0344/002	BE660581	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,50 mg solution buvable en récipient unidose	BE/H/0344/002	BE660581	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,50 mg solution buvable en récipient unidose	BE/H/0344/002	BE660581	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,50 mg solution buvable en récipient unidose	BE/H/0344/002	BE660581	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,50 mg solution buvable en récipient unidose	BE/H/0344/002	BE660581	IBSA FARMACEUTICI ITALIA	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**
Alprazolam IBSA 0,50 mg solution buvable en récipient unidose	BE/H/0344/002	BE660581	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,50 mg solution buvable en récipient unidose	BE/H/0344/002	BE660581	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,50 mg solution buvable en récipient unidose	BE/H/0344/002	BE660581	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,50 mg solution buvable en récipient unidose	BE/H/0344/002	BE660581	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,50 mg solution buvable en récipient unidose	BE/H/0344/002	BE660581	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,50 mg solution buvable en récipient unidose	BE/H/0344/002	BE660581	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,50 mg solution buvable en récipient unidose	BE/H/0344/002	BE660581	IBSA FARMACEUTICI ITALIA	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
Alprazolam IBSA 0,50 mg solution buvable en récipient unidose	BE/H/0344/002	2022110230	IBSA FARMACEUTICI ITALIA	LU
Alprazolam IBSA 0,50 mg solution buvable en récipient unidose	BE/H/0344/002	2022110230	IBSA FARMACEUTICI ITALIA	LU
Alprazolam IBSA 0,50 mg solution buvable en récipient unidose	BE/H/0344/002	2022110230	IBSA FARMACEUTICI ITALIA	LU
Alprazolam IBSA 0,50 mg solution buvable en récipient unidose	BE/H/0344/002	2022110230	IBSA FARMACEUTICI ITALIA	LU
Alprazolam IBSA 0,50 mg solution buvable en récipient unidose	BE/H/0344/002	2022110230	IBSA FARMACEUTICI ITALIA	LU
Alprazolam IBSA 0,50 mg solution buvable en récipient unidose	BE/H/0344/002	2022110230	IBSA FARMACEUTICI ITALIA	LU
Alprazolam IBSA 0,50 mg solution buvable en récipient unidose	BE/H/0344/002	2022110230	IBSA FARMACEUTICI ITALIA	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
Alprazolam IBSA 0,50 mg solution buvable en récipient unidose	BE/H/0344/002	2022110230	IBSA FARMACEUTICI ITALIA	LU
Alprazolam IBSA 0,50 mg solution buvable en récipient unidose	BE/H/0344/002	2022110230	IBSA FARMACEUTICI ITALIA	LU
Alprazolam IBSA 0,50 mg solution buvable en récipient unidose	BE/H/0344/002	2022110230	IBSA FARMACEUTICI ITALIA	LU
Alprazolam IBSA 0,50 mg solution buvable en récipient unidose	BE/H/0344/002	2022110230	IBSA FARMACEUTICI ITALIA	LU
Alprazolam IBSA 0,50 mg solution buvable en récipient unidose	BE/H/0344/002	2022110230	IBSA FARMACEUTICI ITALIA	LU
Alprazolam IBSA 0,50 mg solution buvable en récipient unidose	BE/H/0344/002	2022110230	IBSA FARMACEUTICI ITALIA	LU
Alprazolam IBSA 0,50 mg solution buvable en récipient unidose	BE/H/0344/002	2022110230	IBSA FARMACEUTICI ITALIA	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**
Alprazolam IBSA 0,50 mg soluzione orale in contenitore monodose	BE/H/0344/002	049508031	IBSA FARMACEUTICI ITALIA	IT
Alprazolam IBSA 0,50 mg soluzione orale in contenitore monodose	BE/H/0344/002	049508043	IBSA FARMACEUTICI ITALIA	IT
Alprazolam IBSA 0,50 mg soluzione orale in contenitore monodose	BE/H/0344/002	049508031	IBSA FARMACEUTICI ITALIA	IT
Alprazolam IBSA 0,50 mg soluzione orale in contenitore monodose	BE/H/0344/002	049508043	IBSA FARMACEUTICI ITALIA	IT
Alprazolam IBSA 0,50 mg soluzione orale in contenitore monodose	BE/H/0344/002	049508031	IBSA FARMACEUTICI ITALIA	IT
Alprazolam IBSA 0,50 mg soluzione orale in contenitore monodose	BE/H/0344/002	049508043	IBSA FARMACEUTICI ITALIA	IT
Alprazolam IBSA 0,50 mg soluzione orale in contenitore monodose	BE/H/0344/002	049508031	IBSA FARMACEUTICI ITALIA	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
Alprazolam IBSA 0,50 mg soluzione orale in contenitore monodose	BE/H/0344/002	049508043	IBSA FARMACEUTICI ITALIA	IT
Alprazolam IBSA 0,50 mg soluzione orale in contenitore monodose	BE/H/0344/002	049508031	IBSA FARMACEUTICI ITALIA	IT
Alprazolam IBSA 0,50 mg soluzione orale in contenitore monodose	BE/H/0344/002	049508043	IBSA FARMACEUTICI ITALIA	IT
Alprazolam IBSA 0,50 mg soluzione orale in contenitore monodose	BE/H/0344/002	049508031	IBSA FARMACEUTICI ITALIA	IT
Alprazolam IBSA 0,50 mg soluzione orale in contenitore monodose	BE/H/0344/002	049508043	IBSA FARMACEUTICI ITALIA	IT
Alprazolam IBSA 0,50 mg soluzione orale in contenitore monodose	BE/H/0344/002	049508031	IBSA FARMACEUTICI ITALIA	IT
Alprazolam IBSA 0,50 mg soluzione orale in contenitore monodose	BE/H/0344/002	049508043	IBSA FARMACEUTICI ITALIA	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
Alprazolam IBSA 0,50 Milligramm Lösung zum Einnehmen im Einzeldosisbehältnis	BE/H/0344/002	BE660581	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,50 Milligramm Lösung zum Einnehmen im Einzeldosisbehältnis	BE/H/0344/002	BE660581	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,50 Milligramm Lösung zum Einnehmen im Einzeldosisbehältnis	BE/H/0344/002	BE660581	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,50 Milligramm Lösung zum Einnehmen im Einzeldosisbehältnis	BE/H/0344/002	BE660581	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,50 Milligramm Lösung zum Einnehmen im Einzeldosisbehältnis	BE/H/0344/002	BE660581	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,50 Milligramm Lösung zum	BE/H/0344/002	BE660581	IBSA FARMACEUTICI ITALIA	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
Einnehmen im Einzeldosisbehältnis				
Alprazolam IBSA 0,50 Milligramm Lösung zum Einnehmen im Einzeldosisbehältnis	BE/H/0344/002	BE660581	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,50 Milligramm Lösung zum Einnehmen im Einzeldosisbehältnis	BE/H/0344/002	BE660581	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,50 Milligramm Lösung zum Einnehmen im Einzeldosisbehältnis	BE/H/0344/002	BE660581	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,50 Milligramm Lösung zum Einnehmen im Einzeldosisbehältnis	BE/H/0344/002	BE660581	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,50 Milligramm Lösung zum Einnehmen im Einzeldosisbehältnis	BE/H/0344/002	BE660581	IBSA FARMACEUTICI ITALIA	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
Alprazolam IBSA 0,50 Milligramm Lösung zum Einnehmen im Einzeldosisbehältnis	BE/H/0344/002	BE660581	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,50 Milligramm Lösung zum Einnehmen im Einzeldosisbehältnis	BE/H/0344/002	BE660581	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,50 Milligramm Lösung zum Einnehmen im Einzeldosisbehältnis	BE/H/0344/002	BE660581	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,75 mg drank in verpakking voor eenmalig gebruik	BE/H/0344/003	BE660582	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,75 mg drank in verpakking voor eenmalig gebruik	BE/H/0344/003	BE660582	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,75 mg drank in verpakking voor eenmalig gebruik	BE/H/0344/003	BE660582	IBSA FARMACEUTICI ITALIA	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
Alprazolam IBSA 0,75 mg drank in verpakking voor eenmalig gebruik	BE/H/0344/003	BE660582	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,75 mg drank in verpakking voor eenmalig gebruik	BE/H/0344/003	BE660582	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,75 mg drank in verpakking voor eenmalig gebruik	BE/H/0344/003	BE660582	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,75 mg drank in verpakking voor eenmalig gebruik	BE/H/0344/003	BE660582	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,75 mg drank in verpakking voor eenmalig gebruik	BE/H/0344/003	BE660582	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,75 mg drank in verpakking voor eenmalig gebruik	BE/H/0344/003	BE660582	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,75 mg drank in verpakking voor eenmalig gebruik	BE/H/0344/003	BE660582	IBSA FARMACEUTICI ITALIA	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
Alprazolam IBSA 0,75 mg drank in verpakking voor eenmalig gebruik	BE/H/0344/003	BE660582	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,75 mg drank in verpakking voor eenmalig gebruik	BE/H/0344/003	BE660582	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,75 mg drank in verpakking voor eenmalig gebruik	BE/H/0344/003	BE660582	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,75 mg drank in verpakking voor eenmalig gebruik	BE/H/0344/003	BE660582	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,75 mg solution buvable en récipient unidose	BE/H/0344/003	BE660582	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,75 mg solution buvable en récipient unidose	BE/H/0344/003	BE660582	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,75 mg solution buvable en récipient unidose	BE/H/0344/003	BE660582	IBSA FARMACEUTICI ITALIA	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
Alprazolam IBSA 0,75 mg solution buvable en récipient unidose	BE/H/0344/003	BE660582	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,75 mg solution buvable en récipient unidose	BE/H/0344/003	BE660582	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,75 mg solution buvable en récipient unidose	BE/H/0344/003	BE660582	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,75 mg solution buvable en récipient unidose	BE/H/0344/003	BE660582	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,75 mg solution buvable en récipient unidose	BE/H/0344/003	BE660582	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,75 mg solution buvable en récipient unidose	BE/H/0344/003	BE660582	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,75 mg solution buvable en récipient unidose	BE/H/0344/003	BE660582	IBSA FARMACEUTICI ITALIA	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
Alprazolam IBSA 0,75 mg solution buvable en récipient unidose	BE/H/0344/003	BE660582	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,75 mg solution buvable en récipient unidose	BE/H/0344/003	BE660582	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,75 mg solution buvable en récipient unidose	BE/H/0344/003	BE660582	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,75 mg solution buvable en récipient unidose	BE/H/0344/003	BE660582	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,75 mg solution buvable en récipient unidose	BE/H/0344/003	2022110231	IBSA FARMACEUTICI ITALIA	LU
Alprazolam IBSA 0,75 mg solution buvable en récipient unidose	BE/H/0344/003	2022110231	IBSA FARMACEUTICI ITALIA	LU
Alprazolam IBSA 0,75 mg solution buvable en récipient unidose	BE/H/0344/003	2022110231	IBSA FARMACEUTICI ITALIA	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
Alprazolam IBSA 0,75 mg solution buvable en récipient unidose	BE/H/0344/003	2022110231	IBSA FARMACEUTICI ITALIA	LU
Alprazolam IBSA 0,75 mg solution buvable en récipient unidose	BE/H/0344/003	2022110231	IBSA FARMACEUTICI ITALIA	LU
Alprazolam IBSA 0,75 mg solution buvable en récipient unidose	BE/H/0344/003	2022110231	IBSA FARMACEUTICI ITALIA	LU
Alprazolam IBSA 0,75 mg solution buvable en récipient unidose	BE/H/0344/003	2022110231	IBSA FARMACEUTICI ITALIA	LU
Alprazolam IBSA 0,75 mg solution buvable en récipient unidose	BE/H/0344/003	2022110231	IBSA FARMACEUTICI ITALIA	LU
Alprazolam IBSA 0,75 mg solution buvable en récipient unidose	BE/H/0344/003	2022110231	IBSA FARMACEUTICI ITALIA	LU
Alprazolam IBSA 0,75 mg solution buvable en récipient unidose	BE/H/0344/003	2022110231	IBSA FARMACEUTICI ITALIA	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
Alprazolam IBSA 0,75 mg solution buvable en récipient unidose	BE/H/0344/003	2022110231	IBSA FARMACEUTICI ITALIA	LU
Alprazolam IBSA 0,75 mg solution buvable en récipient unidose	BE/H/0344/003	2022110231	IBSA FARMACEUTICI ITALIA	LU
Alprazolam IBSA 0,75 mg solution buvable en récipient unidose	BE/H/0344/003	2022110231	IBSA FARMACEUTICI ITALIA	LU
Alprazolam IBSA 0,75 mg solution buvable en récipient unidose	BE/H/0344/003	2022110231	IBSA FARMACEUTICI ITALIA	LU
Alprazolam IBSA 0,75 mg soluzione orale in contenitore monodose	BE/H/0344/003	049508056	IBSA FARMACEUTICI ITALIA	IT
Alprazolam IBSA 0,75 mg soluzione orale in contenitore monodose	BE/H/0344/003	049508068	IBSA FARMACEUTICI ITALIA	IT
Alprazolam IBSA 0,75 mg soluzione orale in contenitore monodose	BE/H/0344/003	049508056	IBSA FARMACEUTICI ITALIA	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**
Alprazolam IBSA 0,75 mg soluzione orale in contenitore monodose	BE/H/0344/003	049508068	IBSA FARMACEUTICI ITALIA	IT
Alprazolam IBSA 0,75 mg soluzione orale in contenitore monodose	BE/H/0344/003	049508056	IBSA FARMACEUTICI ITALIA	IT
Alprazolam IBSA 0,75 mg soluzione orale in contenitore monodose	BE/H/0344/003	049508068	IBSA FARMACEUTICI ITALIA	IT
Alprazolam IBSA 0,75 mg soluzione orale in contenitore monodose	BE/H/0344/003	049508056	IBSA FARMACEUTICI ITALIA	IT
Alprazolam IBSA 0,75 mg soluzione orale in contenitore monodose	BE/H/0344/003	049508068	IBSA FARMACEUTICI ITALIA	IT
Alprazolam IBSA 0,75 mg soluzione orale in contenitore monodose	BE/H/0344/003	049508056	IBSA FARMACEUTICI ITALIA	IT
Alprazolam IBSA 0,75 mg soluzione orale in contenitore monodose	BE/H/0344/003	049508068	IBSA FARMACEUTICI ITALIA	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
Alprazolam IBSA 0,75 mg soluzione orale in contenitore monodose	BE/H/0344/003	049508056	IBSA FARMACEUTICI ITALIA	IT
Alprazolam IBSA 0,75 mg soluzione orale in contenitore monodose	BE/H/0344/003	049508068	IBSA FARMACEUTICI ITALIA	IT
Alprazolam IBSA 0,75 mg soluzione orale in contenitore monodose	BE/H/0344/003	049508056	IBSA FARMACEUTICI ITALIA	IT
Alprazolam IBSA 0,75 mg soluzione orale in contenitore monodose	BE/H/0344/003	049508068	IBSA FARMACEUTICI ITALIA	IT
Alprazolam IBSA 0,75 Milligramm Lösung zum Einnehmen im Einzeldosisbehältnis	BE/H/0344/003	BE660582	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,75 Milligramm Lösung zum Einnehmen im Einzeldosisbehältnis	BE/H/0344/003	BE660582	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,75 Milligramm Lösung zum	BE/H/0344/003	BE660582	IBSA FARMACEUTICI ITALIA	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
Einnehmen im Einzeldosisbehältnis				
Alprazolam IBSA 0,75 Milligramm Lösung zum Einnehmen im Einzeldosisbehältnis	BE/H/0344/003	BE660582	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,75 Milligramm Lösung zum Einnehmen im Einzeldosisbehältnis	BE/H/0344/003	BE660582	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,75 Milligramm Lösung zum Einnehmen im Einzeldosisbehältnis	BE/H/0344/003	BE660582	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,75 Milligramm Lösung zum Einnehmen im Einzeldosisbehältnis	BE/H/0344/003	BE660582	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,75 Milligramm Lösung zum Einnehmen im Einzeldosisbehältnis	BE/H/0344/003	BE660582	IBSA FARMACEUTICI ITALIA	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**
Alprazolam IBSA 0,75 Milligramm Lösung zum Einnehmen im Einzeldosisbehältnis	BE/H/0344/003	BE660582	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,75 Milligramm Lösung zum Einnehmen im Einzeldosisbehältnis	BE/H/0344/003	BE660582	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,75 Milligramm Lösung zum Einnehmen im Einzeldosisbehältnis	BE/H/0344/003	BE660582	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,75 Milligramm Lösung zum Einnehmen im Einzeldosisbehältnis	BE/H/0344/003	BE660582	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,75 Milligramm Lösung zum Einnehmen im Einzeldosisbehältnis	BE/H/0344/003	BE660582	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 0,75 Milligramm Lösung zum	BE/H/0344/003	BE660582	IBSA FARMACEUTICI ITALIA	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
Einnehmen im Einzeldosisbehältnis				
Alprazolam IBSA 1 mg drank in verpakking voor eenmalig gebruik	BE/H/0344/004	BE660583	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 1 mg drank in verpakking voor eenmalig gebruik	BE/H/0344/004	BE660583	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 1 mg drank in verpakking voor eenmalig gebruik	BE/H/0344/004	BE660583	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 1 mg drank in verpakking voor eenmalig gebruik	BE/H/0344/004	BE660583	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 1 mg drank in verpakking voor eenmalig gebruik	BE/H/0344/004	BE660583	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 1 mg drank in verpakking voor eenmalig gebruik	BE/H/0344/004	BE660583	IBSA FARMACEUTICI ITALIA	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
Alprazolam IBSA 1 mg drank in verpakking voor eenmalig gebruik	BE/H/0344/004	BE660583	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 1 mg drank in verpakking voor eenmalig gebruik	BE/H/0344/004	BE660583	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 1 mg drank in verpakking voor eenmalig gebruik	BE/H/0344/004	BE660583	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 1 mg drank in verpakking voor eenmalig gebruik	BE/H/0344/004	BE660583	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 1 mg drank in verpakking voor eenmalig gebruik	BE/H/0344/004	BE660583	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 1 mg drank in verpakking voor eenmalig gebruik	BE/H/0344/004	BE660583	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 1 mg drank in verpakking voor eenmalig gebruik	BE/H/0344/004	BE660583	IBSA FARMACEUTICI ITALIA	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
Alprazolam IBSA 1 mg drank in verpakking voor eenmalig gebruik	BE/H/0344/004	BE660583	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 1 mg solution buvable en récipient unidose	BE/H/0344/004	BE660583	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 1 mg solution buvable en récipient unidose	BE/H/0344/004	BE660583	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 1 mg solution buvable en récipient unidose	BE/H/0344/004	BE660583	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 1 mg solution buvable en récipient unidose	BE/H/0344/004	BE660583	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 1 mg solution buvable en récipient unidose	BE/H/0344/004	BE660583	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 1 mg solution buvable en récipient unidose	BE/H/0344/004	BE660583	IBSA FARMACEUTICI ITALIA	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**
Alprazolam IBSA 1 mg solution buvable en récipient unidose	BE/H/0344/004	BE660583	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 1 mg solution buvable en récipient unidose	BE/H/0344/004	BE660583	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 1 mg solution buvable en récipient unidose	BE/H/0344/004	BE660583	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 1 mg solution buvable en récipient unidose	BE/H/0344/004	BE660583	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 1 mg solution buvable en récipient unidose	BE/H/0344/004	BE660583	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 1 mg solution buvable en récipient unidose	BE/H/0344/004	BE660583	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 1 mg solution buvable en récipient unidose	BE/H/0344/004	BE660583	IBSA FARMACEUTICI ITALIA	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
Alprazolam IBSA 1 mg solution buvable en récipient unidose	BE/H/0344/004	BE660583	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 1 mg solution buvable en récipient unidose	BE/H/0344/004	2022110232	IBSA FARMACEUTICI ITALIA	LU
Alprazolam IBSA 1 mg solution buvable en récipient unidose	BE/H/0344/004	2022110232	IBSA FARMACEUTICI ITALIA	LU
Alprazolam IBSA 1 mg solution buvable en récipient unidose	BE/H/0344/004	2022110232	IBSA FARMACEUTICI ITALIA	LU
Alprazolam IBSA 1 mg solution buvable en récipient unidose	BE/H/0344/004	2022110232	IBSA FARMACEUTICI ITALIA	LU
Alprazolam IBSA 1 mg solution buvable en récipient unidose	BE/H/0344/004	2022110232	IBSA FARMACEUTICI ITALIA	LU
Alprazolam IBSA 1 mg solution buvable en récipient unidose	BE/H/0344/004	2022110232	IBSA FARMACEUTICI ITALIA	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**
Alprazolam IBSA 1 mg solution buvable en récipient unidose	BE/H/0344/004	2022110232	IBSA FARMACEUTICI ITALIA	LU
Alprazolam IBSA 1 mg solution buvable en récipient unidose	BE/H/0344/004	2022110232	IBSA FARMACEUTICI ITALIA	LU
Alprazolam IBSA 1 mg solution buvable en récipient unidose	BE/H/0344/004	2022110232	IBSA FARMACEUTICI ITALIA	LU
Alprazolam IBSA 1 mg solution buvable en récipient unidose	BE/H/0344/004	2022110232	IBSA FARMACEUTICI ITALIA	LU
Alprazolam IBSA 1 mg solution buvable en récipient unidose	BE/H/0344/004	2022110232	IBSA FARMACEUTICI ITALIA	LU
Alprazolam IBSA 1 mg solution buvable en récipient unidose	BE/H/0344/004	2022110232	IBSA FARMACEUTICI ITALIA	LU
Alprazolam IBSA 1 mg solution buvable en récipient unidose	BE/H/0344/004	2022110232	IBSA FARMACEUTICI ITALIA	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
Alprazolam IBSA 1 mg solution buvable en récipient unidose	BE/H/0344/004	2022110232	IBSA FARMACEUTICI ITALIA	LU
Alprazolam IBSA 1 mg soluzione orale in contenitore monodose	BE/H/0344/004	049508070	IBSA FARMACEUTICI ITALIA	IT
Alprazolam IBSA 1 mg soluzione orale in contenitore monodose	BE/H/0344/004	049508082	IBSA FARMACEUTICI ITALIA	IT
Alprazolam IBSA 1 mg soluzione orale in contenitore monodose	BE/H/0344/004	049508070	IBSA FARMACEUTICI ITALIA	IT
Alprazolam IBSA 1 mg soluzione orale in contenitore monodose	BE/H/0344/004	049508082	IBSA FARMACEUTICI ITALIA	IT
Alprazolam IBSA 1 mg soluzione orale in contenitore monodose	BE/H/0344/004	049508070	IBSA FARMACEUTICI ITALIA	IT
Alprazolam IBSA 1 mg soluzione orale in contenitore monodose	BE/H/0344/004	049508082	IBSA FARMACEUTICI ITALIA	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
Alprazolam IBSA 1 mg soluzione orale in contenitore monodose	BE/H/0344/004	049508070	IBSA FARMACEUTICI ITALIA	IT
Alprazolam IBSA 1 mg soluzione orale in contenitore monodose	BE/H/0344/004	049508082	IBSA FARMACEUTICI ITALIA	IT
Alprazolam IBSA 1 mg soluzione orale in contenitore monodose	BE/H/0344/004	049508070	IBSA FARMACEUTICI ITALIA	IT
Alprazolam IBSA 1 mg soluzione orale in contenitore monodose	BE/H/0344/004	049508082	IBSA FARMACEUTICI ITALIA	IT
Alprazolam IBSA 1 mg soluzione orale in contenitore monodose	BE/H/0344/004	049508070	IBSA FARMACEUTICI ITALIA	IT
Alprazolam IBSA 1 mg soluzione orale in contenitore monodose	BE/H/0344/004	049508082	IBSA FARMACEUTICI ITALIA	IT
Alprazolam IBSA 1 mg soluzione orale in contenitore monodose	BE/H/0344/004	049508070	IBSA FARMACEUTICI ITALIA	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
Alprazolam IBSA 1 mg soluzione orale in contenitore monodose	BE/H/0344/004	049508082	IBSA FARMACEUTICI ITALIA	IT
Alprazolam IBSA 1 Milligramm Lösung zum Einnehmen im Einzeldosisbehältnis	BE/H/0344/004	BE660583	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 1 Milligramm Lösung zum Einnehmen im Einzeldosisbehältnis	BE/H/0344/004	BE660583	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 1 Milligramm Lösung zum Einnehmen im Einzeldosisbehältnis	BE/H/0344/004	BE660583	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 1 Milligramm Lösung zum Einnehmen im Einzeldosisbehältnis	BE/H/0344/004	BE660583	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 1 Milligramm Lösung zum Einnehmen im	BE/H/0344/004	BE660583	IBSA FARMACEUTICI ITALIA	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
Einzeldosisbehältnis				
Alprazolam IBSA 1 Milligramm Lösung zum Einnehmen im Einzeldosisbehältnis	BE/H/0344/004	BE660583	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 1 Milligramm Lösung zum Einnehmen im Einzeldosisbehältnis	BE/H/0344/004	BE660583	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 1 Milligramm Lösung zum Einnehmen im Einzeldosisbehältnis	BE/H/0344/004	BE660583	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 1 Milligramm Lösung zum Einnehmen im Einzeldosisbehältnis	BE/H/0344/004	BE660583	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 1 Milligramm Lösung zum Einnehmen im Einzeldosisbehältnis	BE/H/0344/004	BE660583	IBSA FARMACEUTICI ITALIA	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
Alprazolam IBSA 1 Milligramm Lösung zum Einnehmen im Einzeldosisbehältnis	BE/H/0344/004	BE660583	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 1 Milligramm Lösung zum Einnehmen im Einzeldosisbehältnis	BE/H/0344/004	BE660583	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 1 Milligramm Lösung zum Einnehmen im Einzeldosisbehältnis	BE/H/0344/004	BE660583	IBSA FARMACEUTICI ITALIA	BE
Alprazolam IBSA 1 Milligramm Lösung zum Einnehmen im Einzeldosisbehältnis	BE/H/0344/004	BE660583	IBSA FARMACEUTICI ITALIA	BE
ALPRAZOLAM VIATRIS 0,25 mg compresse	not available	028644019	MYLAN S.P.A.	IT
ALPRAZOLAM VIATRIS 0,25 mg compresse	not available	028644019	MYLAN S.P.A.	IT
ALPRAZOLAM VIATRIS	not available	028644019	MYLAN S.P.A.	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
0,25 mg compresse				
ALPRAZOLAM VIATRIS 0,25 mg compresse	not available	028644019	MYLAN S.P.A.	IT
ALPRAZOLAM VIATRIS 0,25 mg compresse	not available	028644019	MYLAN S.P.A.	IT
ALPRAZOLAM VIATRIS 0,25 mg compresse	not available	028644019	MYLAN S.P.A.	IT
ALPRAZOLAM VIATRIS 0,25 mg compresse	not available	028644019	MYLAN S.P.A.	IT
ALPRAZOLAM VIATRIS 0,50 mg compresse	not available	028644021	MYLAN S.P.A.	IT
ALPRAZOLAM VIATRIS 0,50 mg compresse	not available	028644021	MYLAN S.P.A.	IT
ALPRAZOLAM VIATRIS 0,50 mg compresse	not available	028644021	MYLAN S.P.A.	IT
ALPRAZOLAM VIATRIS 0,50 mg compresse	not available	028644021	MYLAN S.P.A.	IT
ALPRAZOLAM VIATRIS 0,50 mg compresse	not available	028644021	MYLAN S.P.A.	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
ALPRAZOLAM VIATRIS 0,50 mg compresse	not available	028644021	MYLAN S.P.A.	IT
ALPRAZOLAM VIATRIS 0,50 mg compresse	not available	028644021	MYLAN S.P.A.	IT
ALPRAZOLAM VIATRIS 0,75 mg/ml gocce orali, soluzione	not available	028644084	MYLAN S.P.A.	IT
ALPRAZOLAM VIATRIS 0,75 mg/ml gocce orali, soluzione	not available	028644084	MYLAN S.P.A.	IT
ALPRAZOLAM VIATRIS 0,75 mg/ml gocce orali, soluzione	not available	028644084	MYLAN S.P.A.	IT
ALPRAZOLAM VIATRIS 0,75 mg/ml gocce orali, soluzione	not available	028644084	MYLAN S.P.A.	IT
ALPRAZOLAM VIATRIS 0,75 mg/ml gocce orali, soluzione	not available	028644084	MYLAN S.P.A.	IT
ALPRAZOLAM VIATRIS 0,75 mg/ml gocce orali,	not available	028644084	MYLAN S.P.A.	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
soluzione				
ALPRAZOLAM VIATRIS 0,75 mg/ml gocce orali, soluzione	not available	028644084	MYLAN S.P.A.	IT
ALPRAZOLAM VIATRIS 1 mg compresse	not available	028644033	MYLAN S.P.A.	IT
ALPRAZOLAM VIATRIS 1 mg compresse	not available	028644033	MYLAN S.P.A.	IT
ALPRAZOLAM VIATRIS 1 mg compresse	not available	028644033	MYLAN S.P.A.	IT
ALPRAZOLAM VIATRIS 1 mg compresse	not available	028644033	MYLAN S.P.A.	IT
ALPRAZOLAM VIATRIS 1 mg compresse	not available	028644033	MYLAN S.P.A.	IT
ALPRAZOLAM VIATRIS 1 mg compresse	not available	028644033	MYLAN S.P.A.	IT
ALPRAZOLAM VIATRIS 1 mg compresse	not available	028644033	MYLAN S.P.A.	IT
Alprox, tabletter	FI/H/0752/003	47141	ORION CORPORATION	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
Alprox, tabletter	FI/H/0752/003	47141	ORION CORPORATION	DK
Alprox, tabletter	FI/H/0752/003	47141	ORION CORPORATION	DK
Alprox, tabletter	FI/H/0752/003	47141	ORION CORPORATION	DK
Tafil 0,25 mg töflur.	not available	802634	VIATRIS APS	IS
Tafil 0,25 mg töflur.	not available	802634	VIATRIS APS	IS
Tafil 0,25 mg töflur.	not available	802634	VIATRIS APS	IS
Tafil 0,25 mg töflur.	not available	802634	VIATRIS APS	IS
Tafil 0,25 mg töflur.	not available	802634	VIATRIS APS	IS
Tafil 0,25 mg töflur.	not available	802634	VIATRIS APS	IS
Tafil 0,25 mg töflur.	not available	802634	VIATRIS APS	IS
Tafil 0,5 mg töflur.	not available	802633	VIATRIS APS	IS
Tafil 0,5 mg töflur.	not available	802633	VIATRIS APS	IS
Tafil 0,5 mg töflur.	not available	802633	VIATRIS APS	IS
Tafil 0,5 mg töflur.	not available	802633	VIATRIS APS	IS
Tafil 0,5 mg töflur.	not available	802633	VIATRIS APS	IS
Tafil 0,5 mg töflur.	not available	802633	VIATRIS APS	IS

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
Tafil 0,5 mg töflur.	not available	802633	VIATRIS APS	IS
Tafil 1 mg töflur.	not available	880115	VIATRIS APS	IS
Tafil 1 mg töflur.	not available	880115	VIATRIS APS	IS
Tafil 1 mg töflur.	not available	880115	VIATRIS APS	IS
Tafil 1 mg töflur.	not available	880115	VIATRIS APS	IS
Tafil 1 mg töflur.	not available	880115	VIATRIS APS	IS
Tafil 1 mg töflur.	not available	880115	VIATRIS APS	IS
Tafil 1 mg töflur.	not available	880115	VIATRIS APS	IS
Tafil Retard 0,5 mg forðatöflur.	not available	940124	VIATRIS APS	IS
Tafil Retard 1 mg forðatöflur.	not available	940123	VIATRIS APS	IS
Tafil Retard 2 mg forðatöflur.	not available	940122	VIATRIS APS	IS
Tafil Retard 3 mg forðatöflur.	not available	940121	VIATRIS APS	IS
Tafil Retard,	not available	15826	VIATRIS 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
depottabletter				
Tafil Retard, depottabletter	not available	15825	VIATRIS APS	DK
Tafil Retard, depottabletter	not available	15827	VIATRIS APS	DK
Tafil Retard, depottabletter	not available	15828	VIATRIS APS	DK
Tafil, tabletter	not available	10403	VIATRIS APS	DK
Tafil, tabletter	not available	10405	VIATRIS APS	DK
Tafil, tabletter	not available	10404	VIATRIS APS	DK
Tafil, tabletter	not available	13776	VIATRIS APS	DK
Tafil, tabletter	not available	10403	VIATRIS APS	DK
Tafil, tabletter	not available	10405	VIATRIS APS	DK
Tafil, tabletter	not available	10404	VIATRIS APS	DK
Tafil, tabletter	not available	13776	VIATRIS APS	DK
Tafil, tabletter	not available	10403	VIATRIS APS	DK
Tafil, tabletter	not available	10405	VIATRIS 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
Tafil, tabletter	not available	10404	VIATRIS APS	DK
Tafil, tabletter	not available	13776	VIATRIS APS	DK
Tafil® 0,5 mg Tabletten	not available	29946.00.00	VIATRIS PHARMA GMBH	DE
Tafil® 1,0 mg Tabletten	not available	29946.01.00	VIATRIS PHARMA GMBH	DE
TRANKIMAZIN 0,25 mg comprimidos	not available	56027	VIATRIS HEALTHCARE S.L.	ES
TRANKIMAZIN 0,50 mg comprimidos	not available	56026	VIATRIS HEALTHCARE S.L.	ES
TRANKIMAZIN 0,75 mg/ml gotas orales en solución	not available	64079	VIATRIS HEALTHCARE S.L.	ES
TRANKIMAZIN 1 mg comprimidos	not available	56025	VIATRIS HEALTHCARE S.L.	ES
TRANKIMAZIN 2 mg comprimidos	not available	58784	VIATRIS HEALTHCARE S.L.	ES
TRANKIMAZIN Retard 0,5 mg comprimidos	not available	61152	VIATRIS HEALTHCARE S.L.	ES
TRANKIMAZIN Retard 0,5 mg comprimidos	not available	61152	VIATRIS HEALTHCARE S.L.	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
TRANKIMAZIN Retard 1 mg comprimidos	not available	61151	VIATRIS HEALTHCARE S.L.	ES
TRANKIMAZIN Retard 1 mg comprimidos	not available	61151	VIATRIS HEALTHCARE S.L.	ES
TRANKIMAZIN Retard 2 mg comprimidos	not available	61153	VIATRIS HEALTHCARE S.L.	ES
TRANKIMAZIN Retard 2 mg comprimidos	not available	61153	VIATRIS HEALTHCARE S.L.	ES
TRANKIMAZIN Retard 3 mg comprimidos	not available	61154	VIATRIS HEALTHCARE S.L.	ES
TRANKIMAZIN Retard 3 mg comprimidos	not available	61154	VIATRIS HEALTHCARE S.L.	ES
VALEANS® 0,25 mg capsule rigide	not available	025941 028	NEOPHARMED GENTILI SPA	IT
VALEANS® 0,25 mg capsule rigide	not available	025941 028	NEOPHARMED GENTILI SPA	IT
VALEANS® 0,25 mg capsule rigide	not available	025941 028	NEOPHARMED GENTILI SPA	IT
VALEANS® 0,25 mg	not available	025941 028	NEOPHARMED GENTILI 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
capsule rigide				
VALEANS® 0,25 mg capsule rigide	not available	025941 028	NEOPHARMED GENTILI SPA	IT
VALEANS® 0,25 mg capsule rigide	not available	025941 028	NEOPHARMED GENTILI SPA	IT
VALEANS® 0,25 mg capsule rigide	not available	025941 028	NEOPHARMED GENTILI SPA	IT
VALEANS® 0,50 mg capsule rigide	not available	025941 016	NEOPHARMED GENTILI SPA	IT
VALEANS® 0,50 mg capsule rigide	not available	025941 016	NEOPHARMED GENTILI SPA	IT
VALEANS® 0,50 mg capsule rigide	not available	025941 016	NEOPHARMED GENTILI SPA	IT
VALEANS® 0,50 mg capsule rigide	not available	025941 016	NEOPHARMED GENTILI SPA	IT
VALEANS® 0,50 mg capsule rigide	not available	025941 016	NEOPHARMED GENTILI SPA	IT
VALEANS® 0,50 mg capsule rigide	not available	025941 016	NEOPHARMED GENTILI 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
VALEANS® 0,50 mg capsule rigide	not available	025941 016	NEOPHARMED GENTILI SPA	IT
VALEANS® 1 mg capsule rigide	not available	025941 042	NEOPHARMED GENTILI SPA	IT
VALEANS® 1 mg capsule rigide	not available	025941 042	NEOPHARMED GENTILI SPA	IT
VALEANS® 1 mg capsule rigide	not available	025941 042	NEOPHARMED GENTILI SPA	IT
VALEANS® 1 mg capsule rigide	not available	025941 042	NEOPHARMED GENTILI SPA	IT
VALEANS® 1 mg capsule rigide	not available	025941 042	NEOPHARMED GENTILI SPA	IT
VALEANS® 1 mg capsule rigide	not available	025941 042	NEOPHARMED GENTILI SPA	IT
VALEANS® 1 mg capsule rigide	not available	025941 042	NEOPHARMED GENTILI SPA	IT
VALEANS®0,75 mg/ml gocce orali, soluzione	not available	025941 055	NEOPHARMED GENTILI SPA	IT
VALEANS®0,75 mg/ml	not available	025941 055	NEOPHARMED GENTILI 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
gocce orali, soluzione				
VALEANS®0,75 mg/ml gocce orali, soluzione	not available	025941 055	NEOPHARMED GENTILI SPA	IT
VALEANS®0,75 mg/ml gocce orali, soluzione	not available	025941 055	NEOPHARMED GENTILI SPA	IT
VALEANS®0,75 mg/ml gocce orali, soluzione	not available	025941 055	NEOPHARMED GENTILI SPA	IT
VALEANS®0,75 mg/ml gocce orali, soluzione	not available	025941 055	NEOPHARMED GENTILI SPA	IT
VALEANS®0,75 mg/ml gocce orali, soluzione	not available	025941 055	NEOPHARMED GENTILI SPA	IT
XANAX	not available	23876/26.06.95	VIATRIS HELLAS LTD.	GR
XANAX	not available	23877/26.06.95	VIATRIS HELLAS LTD.	GR
XANAX	not available	40778/22.12.97	VIATRIS HELLAS LTD.	GR
XANAX	not available	23875/26.06.95	VIATRIS HELLAS LTD.	GR
XANAX	not available	23876/26.06.95	VIATRIS HELLAS LTD.	GR
XANAX	not available	23877/26.06.95	VIATRIS HELLAS LTD.	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
XANAX	not available	40778/22.12.97	VIATRIS HELLAS LTD.	GR
XANAX	not available	23875/26.06.95	VIATRIS HELLAS LTD.	GR
XANAX	not available	23876/26.06.95	VIATRIS HELLAS LTD.	GR
XANAX	not available	23877/26.06.95	VIATRIS HELLAS LTD.	GR
XANAX	not available	40778/22.12.97	VIATRIS HELLAS LTD.	GR
XANAX	not available	23875/26.06.95	VIATRIS HELLAS LTD.	GR
XANAX	not available	23876/26.06.95	VIATRIS HELLAS LTD.	GR
XANAX	not available	23877/26.06.95	VIATRIS HELLAS LTD.	GR
XANAX	not available	40778/22.12.97	VIATRIS HELLAS LTD.	GR
XANAX	not available	23875/26.06.95	VIATRIS HELLAS LTD.	GR
XANAX	not available	23876/26.06.95	VIATRIS HELLAS LTD.	GR
XANAX	not available	23877/26.06.95	VIATRIS HELLAS LTD.	GR
XANAX	not available	40778/22.12.97	VIATRIS HELLAS LTD.	GR
XANAX	not available	23875/26.06.95	VIATRIS HELLAS LTD.	GR
XANAX 0,25 mg compresse	not available	025980057	VIATRIS PHARMA 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
Xanax 0,25 mg comprimate	not available	7758/2015/01	UPJOHN EESV	RO
XANAX 0,25 mg comprimés	not available	BE120994	VIATRIS HEALTHCARE	BE
XANAX 0,25 mg comprimés	not available	BE120994	VIATRIS HEALTHCARE	BE
XANAX 0,25 mg comprimés	not available	BE120994	VIATRIS HEALTHCARE	BE
XANAX 0,25 mg comprimés	not available	BE120994	VIATRIS HEALTHCARE	BE
XANAX 0,25 mg comprimés	not available	2007079357	VIATRIS HEALTHCARE	LU
XANAX 0,25 mg comprimés	not available	2007079357	VIATRIS HEALTHCARE	LU
XANAX 0,25 mg comprimés	not available	2007079357	VIATRIS HEALTHCARE	LU
XANAX 0,25 mg comprimés	not available	2007079357	VIATRIS HEALTHCARE	LU
Xanax 0,25 mg	not available	4578795	VIATRIS HEALTHCARE 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
comprimidos				
Xanax 0,25 mg comprimidos	not available	8573345	VIATRIS HEALTHCARE LDA.	PT
Xanax 0,25 mg comprimidos	not available	4578787	VIATRIS HEALTHCARE LDA.	PT
Xanax 0,25 mg comprimidos	not available	5787692	VIATRIS HEALTHCARE LDA.	PT
Xanax 0,25 mg comprimidos	not available	9573345	VIATRIS HEALTHCARE LDA.	PT
Xanax 0,25 mg tablete	not available	HR-H-204001816	UPJOHN EESV	HR
Xanax 0,25 mg tablete	not available	HR-H-204001816	UPJOHN EESV	HR
Xanax 0,25 mg tablete	not available	H/92/01672/001	UPJOHN EESV	SI
Xanax 0,25 mg tablete	not available	H/92/01672/001	UPJOHN EESV	SI
Xanax 0,25 mg tablete	not available	H/92/01672/001	UPJOHN EESV	SI
Xanax 0,25 mg tablete	not available	H/92/01672/001	UPJOHN EESV	SI
Xanax 0,25 mg tablete	not available	H/92/01672/001	UPJOHN EESV	SI
XANAX 0,25 mg tabletės	not available	LT/1/96/2950/002	UPJOHN EESV	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
XANAX 0,25 mg tabletės	not available	LT/1/96/2950/001	UPJOHN EESV	LT
Xanax 0,25 mg tabletid	not available	205998	UPJOHN EESV	EE
XANAX 0,25 mg tabletta	not available	OGYI-T-4617/05	VIATRIS HEALTHCARE LTD	HU
XANAX 0,25 mg tabletta	not available	OGYI-T-4617/04	VIATRIS HEALTHCARE LTD	HU
XANAX 0,25 mg tabletta	not available	OGYI-T-4617/05	VIATRIS HEALTHCARE LTD	HU
XANAX 0,25 mg tabletta	not available	OGYI-T-4617/04	VIATRIS HEALTHCARE LTD	HU
XANAX 0,25 mg Tabletten	not available	BE120994	VIATRIS HEALTHCARE	BE
XANAX 0,25 mg tabletten	not available	BE120994	VIATRIS HEALTHCARE	BE
XANAX 0,25 mg Tabletten	not available	BE120994	VIATRIS HEALTHCARE	BE
XANAX 0,25 mg tabletten	not available	BE120994	VIATRIS HEALTHCARE	BE
XANAX 0,25 mg Tabletten	not available	BE120994	VIATRIS HEALTHCARE	BE
XANAX 0,25 mg tabletten	not available	BE120994	VIATRIS HEALTHCARE	BE
XANAX 0,25 mg Tabletten	not available	BE120994	VIATRIS HEALTHCARE	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
XANAX 0,25 mg tabletten	not available	BE120994	VIATRIS HEALTHCARE	BE
XANAX 0,25 mg Tabletten	not available	2007079357	VIATRIS HEALTHCARE	LU
XANAX 0,25 mg Tabletten	not available	2007079357	VIATRIS HEALTHCARE	LU
XANAX 0,25 mg Tabletten	not available	2007079357	VIATRIS HEALTHCARE	LU
XANAX 0,25 mg Tabletten	not available	2007079357	VIATRIS HEALTHCARE	LU
XANAX 0,25 mg tablety	not available	70/214/85-A/C	UPJOHN EESV	CZ
XANAX 0,25 mg tablety	not available	70/214/85-A/C	UPJOHN EESV	CZ
XANAX 0,25 mg tablety	not available	70/214/85-A/C	UPJOHN EESV	CZ
XANAX 0,25 mg tablety	not available	70/0214/85-CS	UPJOHN EESV	SK
XANAX 0,25 mg tablety	not available	70/0214/85-CS	UPJOHN EESV	SK
XANAX 0,25 mg tablety	not available	70/0214/85-CS	UPJOHN EESV	SK
XANAX 0,25 mg, comprimé sécable	not available	34009 554 218 3 2	VIATRIS UP	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
XANAX 0,25 mg, comprimé sécable	not available	34009 326 444 9 0	VIATRIS UP	FR
XANAX 0,25 mg, comprimé sécable	not available	34009 554 218 3 2	VIATRIS UP	FR
XANAX 0,25 mg, comprimé sécable	not available	34009 326 444 9 0	VIATRIS UP	FR
XANAX 0,25 mg, comprimé sécable	not available	34009 554 218 3 2	VIATRIS UP	FR
XANAX 0,25 mg, comprimé sécable	not available	34009 326 444 9 0	VIATRIS UP	FR
XANAX 0,25 mg, comprimé sécable	not available	34009 554 218 3 2	VIATRIS UP	FR
XANAX 0,25 mg, comprimé sécable	not available	34009 326 444 9 0	VIATRIS UP	FR
Xanax 0,25, tabletten 0,25 mg	not available	RVG 14409	VIATRIS HEALTHCARE LTD	NL
XANAX 0,5 mg compresse a rilascio prolungato	not available	025980172	VIATRIS PHARMA 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
XANAX 0,5 mg compresse a rilascio prolungato	not available	025980160	VIATRIS PHARMA S.R.L.	IT
XANAX 0,5 mg compresse a rilascio prolungato	not available	025980145	VIATRIS PHARMA S.R.L.	IT
XANAX 0,5 mg compresse a rilascio prolungato	not available	025980158	VIATRIS PHARMA S.R.L.	IT
XANAX 0,5 mg compresse a rilascio prolungato	not available	025980133	VIATRIS PHARMA S.R.L.	IT
Xanax 0,5 mg comprimate	not available	7759/2015/01	UPJOHN EESV	RO
XANAX 0,5 mg comprimés	not available	BE120933	VIATRIS HEALTHCARE	BE
XANAX 0,5 mg comprimés	not available	BE120933	VIATRIS HEALTHCARE	BE
XANAX 0,5 mg comprimés	not available	BE120933	VIATRIS HEALTHCARE	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
XANAX 0,5 mg comprimés	not available	BE120933	VIATRIS HEALTHCARE	BE
XANAX 0,5 mg comprimés	not available	2007079358	VIATRIS HEALTHCARE	LU
XANAX 0,5 mg comprimés	not available	2007079358	VIATRIS HEALTHCARE	LU
XANAX 0,5 mg comprimés	not available	2007079358	VIATRIS HEALTHCARE	LU
XANAX 0,5 mg comprimés	not available	2007079358	VIATRIS HEALTHCARE	LU
Xanax 0,5 mg comprimidos	not available	4578993	VIATRIS HEALTHCARE LDA.	PT
Xanax 0,5 mg comprimidos	not available	4578886	VIATRIS HEALTHCARE LDA.	PT
Xanax 0,5 mg comprimidos	not available	4578894	VIATRIS HEALTHCARE LDA.	PT
Xanax 0,5 mg comprimidos	not available	5787791	VIATRIS HEALTHCARE LDA.	PT
Xanax 0,5 mg	not available	4578985	VIATRIS HEALTHCARE 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
comprimidos				
Xanax 0,5 mg tablete	not available	HR-H-365377512	UPJOHN EESV	HR
Xanax 0,5 mg tablete	not available	HR-H-365377512	UPJOHN EESV	HR
Xanax 0,5 mg tablete	not available	H/92/01672/002	UPJOHN EESV	SI
Xanax 0,5 mg tablete	not available	H/92/01672/002	UPJOHN EESV	SI
Xanax 0,5 mg tablete	not available	H/92/01672/002	UPJOHN EESV	SI
Xanax 0,5 mg tablete	not available	H/92/01672/002	UPJOHN EESV	SI
Xanax 0,5 mg tablete	not available	H/92/01672/002	UPJOHN EESV	SI
XANAX 0,5 mg tabletes	not available	97-0166	UPJOHN EESV	LV
XANAX 0,5 mg tabletės	not available	LT/1/96/2950/004	UPJOHN EESV	LT
XANAX 0,5 mg tabletės	not available	LT/1/96/2950/003	UPJOHN EESV	LT
Xanax 0,5 mg tabletid	not available	239398	UPJOHN EESV	EE
XANAX 0,5 mg tabletta	not available	OGYI-T-4617/06	VIATRIS HEALTHCARE LTD	HU
XANAX 0,5 mg tabletta	not available	OGYI-T-4617/07	VIATRIS HEALTHCARE LTD	HU
XANAX 0,5 mg tabletta	not available	OGYI-T-4617/06	VIATRIS HEALTHCARE LTD	HU
XANAX 0,5 mg tabletta	not available	OGYI-T-4617/07	VIATRIS HEALTHCARE LTD	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
XANAX 0,5 mg Tabletten	not available	BE120933	VIATRIS HEALTHCARE	BE
XANAX 0,5 mg tabletten	not available	BE120933	VIATRIS HEALTHCARE	BE
XANAX 0,5 mg Tabletten	not available	BE120933	VIATRIS HEALTHCARE	BE
XANAX 0,5 mg tabletten	not available	BE120933	VIATRIS HEALTHCARE	BE
XANAX 0,5 mg Tabletten	not available	BE120933	VIATRIS HEALTHCARE	BE
XANAX 0,5 mg tabletten	not available	BE120933	VIATRIS HEALTHCARE	BE
XANAX 0,5 mg Tabletten	not available	BE120933	VIATRIS HEALTHCARE	BE
XANAX 0,5 mg tabletten	not available	BE120933	VIATRIS HEALTHCARE	BE
XANAX 0,5 mg Tabletten	not available	2007079358	VIATRIS HEALTHCARE	LU
XANAX 0,5 mg Tabletten	not available	2007079358	VIATRIS HEALTHCARE	LU
XANAX 0,5 mg Tabletten	not available	2007079358	VIATRIS HEALTHCARE	LU
XANAX 0,5 mg Tabletten	not available	2007079358	VIATRIS HEALTHCARE	LU
XANAX 0,5 mg tablety	not available	70/214/85-B/C	UPJOHN EESV	CZ
XANAX 0,5 mg tablety	not available	70/214/85-B/C	UPJOHN EESV	CZ
XANAX 0,5 mg tablety	not available	70/214/85-B/C	UPJOHN EESV	CZ
XANAX 0,5 mg tablety	not available	70/0051/17-S	UPJOHN EESV	SK

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
XANAX 0,5 mg tablety	not available	70/0051/17-S	UPJOHN EESV	SK
XANAX 0,5 mg tablety	not available	70/0051/17-S	UPJOHN EESV	SK
Xanax 0,5, tabletten 0,5 mg	not available	RVG 14410	VIATRIS HEALTHCARE LTD	NL
XANAX 0,50 mg compresse	not available	025980069	VIATRIS PHARMA S.R.L.	IT
XANAX 0,50 mg, comprimé sécable	not available	34009 326 445 5 1	VIATRIS UP	FR
XANAX 0,50 mg, comprimé sécable	not available	34009 554 220 8 2	VIATRIS UP	FR
XANAX 0,50 mg, comprimé sécable	not available	34009 326 445 5 1	VIATRIS UP	FR
XANAX 0,50 mg, comprimé sécable	not available	34009 554 220 8 2	VIATRIS UP	FR
XANAX 0,50 mg, comprimé sécable	not available	34009 326 445 5 1	VIATRIS UP	FR
XANAX 0,50 mg, comprimé sécable	not available	34009 554 220 8 2	VIATRIS UP	FR
XANAX 0,50 mg,	not available	34009 326 445 5 1	VIATRIS UP	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
comprimé sécable				
XANAX 0,50 mg, comprimé sécable	not available	34009 554 220 8 2	VIATRIS UP	FR
XANAX 0,75 mg/ml druppels voor oraal gebruik, oplossing	not available	BE187311	VIATRIS HEALTHCARE	BE
XANAX 0,75 mg/ml druppels voor oraal gebruik, oplossing	not available	BE187311	VIATRIS HEALTHCARE	BE
XANAX 0,75 mg/ml druppels voor oraal gebruik, oplossing	not available	BE187311	VIATRIS HEALTHCARE	BE
XANAX 0,75 mg/ml druppels voor oraal gebruik, oplossing	not available	BE187311	VIATRIS HEALTHCARE	BE
XANAX 0,75 mg/ml gocce orali, soluzione	not available	025980083	VIATRIS PHARMA S.R.L.	IT
XANAX 0,75 mg/ml solution buvable en gouttes	not available	BE187311	VIATRIS HEALTHCARE	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
XANAX 0,75 mg/ml solution buvable en gouttes	not available	BE187311	VIATRIS HEALTHCARE	BE
XANAX 0,75 mg/ml solution buvable en gouttes	not available	BE187311	VIATRIS HEALTHCARE	BE
XANAX 0,75 mg/ml solution buvable en gouttes	not available	BE187311	VIATRIS HEALTHCARE	BE
XANAX 0,75 mg/ml solution buvable en gouttes	not available	2007079356	VIATRIS HEALTHCARE	LU
XANAX 0,75 mg/ml solution buvable en gouttes	not available	2007079356	VIATRIS HEALTHCARE	LU
XANAX 0,75 mg/ml solution buvable en gouttes	not available	2007079356	VIATRIS HEALTHCARE	LU
XANAX 0,75 mg/ml solution buvable en gouttes	not available	2007079356	VIATRIS HEALTHCARE	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
XANAX 0,75 mg/ml Tropfen zum Einnehmen, Lösung	not available	BE187311	VIATRIS HEALTHCARE	BE
XANAX 0,75 mg/ml Tropfen zum Einnehmen, Lösung	not available	BE187311	VIATRIS HEALTHCARE	BE
XANAX 0,75 mg/ml Tropfen zum Einnehmen, Lösung	not available	BE187311	VIATRIS HEALTHCARE	BE
XANAX 0,75 mg/ml Tropfen zum Einnehmen, Lösung	not available	BE187311	VIATRIS HEALTHCARE	BE
XANAX 0,75 mg/ml Tropfen zum Einnehmen, Lösung	not available	2007079356	VIATRIS HEALTHCARE	LU
XANAX 0,75 mg/ml Tropfen zum Einnehmen, Lösung	not available	2007079356	VIATRIS HEALTHCARE	LU
XANAX 0,75 mg/ml Tropfen zum Einnehmen, Lösung	not available	2007079356	VIATRIS HEALTHCARE	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
XANAX 0,75 mg/ml Tropfen zum Einnehmen, Lösung	not available	2007079356	VIATRIS HEALTHCARE	LU
XANAX 1 mg compresse	not available	025980071	VIATRIS PHARMA S.R.L.	IT
XANAX 1 mg compresse a rilascio prolungato	not available	025980196	VIATRIS PHARMA S.R.L.	IT
XANAX 1 mg compresse a rilascio prolungato	not available	025980184	VIATRIS PHARMA S.R.L.	IT
XANAX 1 mg compresse a rilascio prolungato	not available	025980210	VIATRIS PHARMA S.R.L.	IT
XANAX 1 mg compresse a rilascio prolungato	not available	025980246	VIATRIS PHARMA S.R.L.	IT
XANAX 1 mg compresse a rilascio prolungato	not available	025980234	VIATRIS PHARMA S.R.L.	IT
XANAX 1 mg compresse a rilascio prolungato	not available	025980208	VIATRIS PHARMA S.R.L.	IT
XANAX 1 mg compresse a rilascio prolungato	not available	025980222	VIATRIS PHARMA S.R.L.	IT
Xanax 1 mg comprimate	not available	7760/2015/01	UPJOHN EESV	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
XANAX 1 mg comprimés	not available	BE135046	VIATRIS HEALTHCARE	BE
XANAX 1 mg comprimés	not available	BE135046	VIATRIS HEALTHCARE	BE
XANAX 1 mg comprimés	not available	BE135046	VIATRIS HEALTHCARE	BE
XANAX 1 mg comprimés	not available	BE135046	VIATRIS HEALTHCARE	BE
XANAX 1 mg comprimés	not available	2007079359	VIATRIS HEALTHCARE	LU
XANAX 1 mg comprimés	not available	2007079359	VIATRIS HEALTHCARE	LU
XANAX 1 mg comprimés	not available	2007079359	VIATRIS HEALTHCARE	LU
XANAX 1 mg comprimés	not available	2007079359	VIATRIS HEALTHCARE	LU
Xanax 1 mg comprimidos	not available	5787890	VIATRIS HEALTHCARE LDA.	PT
Xanax 1 mg comprimidos	not available	4579199	VIATRIS HEALTHCARE LDA.	PT
Xanax 1 mg comprimidos	not available	4579181	VIATRIS HEALTHCARE LDA.	PT
Xanax 1 mg comprimidos	not available	4579090	VIATRIS HEALTHCARE LDA.	PT
Xanax 1 mg comprimidos	not available	4579082	VIATRIS HEALTHCARE LDA.	PT
Xanax 1 mg tablete	not available	H/92/01672/003	UPJOHN EESV	SI
Xanax 1 mg tablete	not available	H/92/01672/003	UPJOHN EESV	SI
Xanax 1 mg tablete	not available	H/92/01672/003	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
Xanax 1 mg tablete	not available	H/92/01672/003	UPJOHN EESV	SI
Xanax 1 mg tablete	not available	H/92/01672/003	UPJOHN EESV	SI
XANAX 1 mg tabletes	not available	97- 0165	UPJOHN EESV	LV
XANAX 1 mg tabletės	not available	LT/1/96/2950/005	UPJOHN EESV	LT
XANAX 1 mg tabletės	not available	LT/1/96/2950/006	UPJOHN EESV	LT
Xanax 1 mg tabletid	not available	239498	UPJOHN EESV	EE
Xanax 1 mg Tablets	not available	PA 23055/010/001	UPJOHN EESV	IE
XANAX 1 mg tabletta	not available	OGYI-T-4617/08	VIATRIS HEALTHCARE LTD	HU
XANAX 1 mg tabletta	not available	OGYI-T-4617/08	VIATRIS HEALTHCARE LTD	HU
XANAX 1 mg tabletten	not available	BE135046	VIATRIS HEALTHCARE	BE
XANAX 1 mg Tabletten	not available	BE135046	VIATRIS HEALTHCARE	BE
XANAX 1 mg tabletten	not available	BE135046	VIATRIS HEALTHCARE	BE
XANAX 1 mg Tabletten	not available	BE135046	VIATRIS HEALTHCARE	BE
XANAX 1 mg tabletten	not available	BE135046	VIATRIS HEALTHCARE	BE
XANAX 1 mg Tabletten	not available	BE135046	VIATRIS HEALTHCARE	BE
XANAX 1 mg tabletten	not available	BE135046	VIATRIS HEALTHCARE	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
XANAX 1 mg Tabletten	not available	BE135046	VIATRIS HEALTHCARE	BE
XANAX 1 mg Tabletten	not available	2007079359	VIATRIS HEALTHCARE	LU
XANAX 1 mg Tabletten	not available	2007079359	VIATRIS HEALTHCARE	LU
XANAX 1 mg Tabletten	not available	2007079359	VIATRIS HEALTHCARE	LU
XANAX 1 mg Tabletten	not available	2007079359	VIATRIS HEALTHCARE	LU
XANAX 1 mg tablety	not available	70/214/85-C/C	UPJOHN EESV	CZ
XANAX 1 mg tablety	not available	70/214/85-C/C	UPJOHN EESV	CZ
XANAX 1 mg tablety	not available	70/214/85-C/C	UPJOHN EESV	CZ
XANAX 1 mg tablety	not available	70/0052/17-S	UPJOHN EESV	SK
XANAX 1 mg tablety	not available	70/0052/17-S	UPJOHN EESV	SK
XANAX 1 mg tablety	not available	70/0052/17-S	UPJOHN EESV	SK
XANAX 2 mg compresse a rilascio prolungato	not available	025980273	VIATRIS PHARMA S.R.L.	IT
XANAX 2 mg compresse a rilascio prolungato	not available	025980261	VIATRIS PHARMA S.R.L.	IT
XANAX 2 mg compresse a rilascio prolungato	not available	025980259	VIATRIS PHARMA 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
XANAX 2 mg compresse a rilascio prolungato	not available	025980285	VIATRIS PHARMA S.R.L.	IT
XANAX 2 mg compresse a rilascio prolungato	not available	025980297	VIATRIS PHARMA S.R.L.	IT
XANAX 2 mg comprimés	not available	BE147576	VIATRIS HEALTHCARE	BE
XANAX 2 mg comprimés	not available	BE147576	VIATRIS HEALTHCARE	BE
XANAX 2 mg comprimés	not available	BE147576	VIATRIS HEALTHCARE	BE
XANAX 2 mg comprimés	not available	BE147576	VIATRIS HEALTHCARE	BE
XANAX 2 mg comprimés	not available	2007079360	VIATRIS HEALTHCARE	LU
XANAX 2 mg comprimés	not available	2007079360	VIATRIS HEALTHCARE	LU
XANAX 2 mg comprimés	not available	2007079360	VIATRIS HEALTHCARE	LU
XANAX 2 mg comprimés	not available	2007079360	VIATRIS HEALTHCARE	LU
XANAX 2 mg tabletten	not available	BE147576	VIATRIS HEALTHCARE	BE
XANAX 2 mg Tabletten	not available	BE147576	VIATRIS HEALTHCARE	BE
XANAX 2 mg tabletten	not available	BE147576	VIATRIS HEALTHCARE	BE
XANAX 2 mg Tabletten	not available	BE147576	VIATRIS HEALTHCARE	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
XANAX 2 mg tabletten	not available	BE147576	VIATRIS HEALTHCARE	BE
XANAX 2 mg Tabletten	not available	BE147576	VIATRIS HEALTHCARE	BE
XANAX 2 mg tabletten	not available	BE147576	VIATRIS HEALTHCARE	BE
XANAX 2 mg Tabletten	not available	BE147576	VIATRIS HEALTHCARE	BE
XANAX 2 mg Tabletten	not available	2007079360	VIATRIS HEALTHCARE	LU
XANAX 2 mg Tabletten	not available	2007079360	VIATRIS HEALTHCARE	LU
XANAX 2 mg Tabletten	not available	2007079360	VIATRIS HEALTHCARE	LU
XANAX 2 mg Tabletten	not available	2007079360	VIATRIS HEALTHCARE	LU
XANAX 2 mg tablety	not available	70/214/85-D/C	UPJOHN EESV	CZ
XANAX 2 mg tablety	not available	70/214/85-D/C	UPJOHN EESV	CZ
XANAX 2 mg tablety	not available	70/214/85-D/C	UPJOHN EESV	CZ
XANAX 2 mg tablety	not available	70/0053/17-S	UPJOHN EESV	SK
XANAX 2 mg tablety	not available	70/0053/17-S	UPJOHN EESV	SK
XANAX 2 mg tablety	not available	70/0053/17-S	UPJOHN EESV	SK
Xanax 250 microgram Tablets	not available	PA 23055/010/002	UPJOHN EESV	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
Xanax 250 microgram Tablets	not available	MA1396/01201	VIATRIS HELLAS LTD.	MT
Xanax 250 microgram Tablets	not available	MA1396/01201	VIATRIS HELLAS LTD.	MT
Xanax 250 microgram Tablets	not available	MA1396/01201	VIATRIS HELLAS LTD.	MT
Xanax 250 microgram Tablets	not available	MA1396/01201	VIATRIS HELLAS LTD.	MT
Xanax 250 microgram Tablets	not available	PL 50622/0066	UPJOHN UK LIMITED	XI
XANAX 3 mg compresse a rilascio prolungato	not available	025980323	VIATRIS PHARMA S.R.L.	IT
XANAX 3 mg compresse a rilascio prolungato	not available	025980309	VIATRIS PHARMA S.R.L.	IT
XANAX 3 mg compresse a rilascio prolungato	not available	025980311	VIATRIS PHARMA S.R.L.	IT
XANAX 3 mg compresse a rilascio prolungato	not available	025980347	VIATRIS PHARMA S.R.L.	IT
XANAX 3 mg compresse a	not available	025980335	VIATRIS PHARMA 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
rilascio prolungato				
Xanax 500 microgram Tablets	not available	PA 23055/010/003	UPJOHN EESV	IE
Xanax 500 microgram Tablets	not available	MA1396/01202	VIATRIS HELLAS LTD.	MT
Xanax 500 microgram Tablets	not available	MA1396/01202	VIATRIS HELLAS LTD.	MT
Xanax 500 microgram Tablets	not available	MA1396/01202	VIATRIS HELLAS LTD.	MT
Xanax 500 microgram Tablets	not available	MA1396/01202	VIATRIS HELLAS LTD.	MT
Xanax 500 microgram Tablets	not available	PL 50622/0067	UPJOHN UK LIMITED	XI
XANAX Retard 0,5 mg comprimés à libération prolongée	not available	BE173223	VIATRIS HEALTHCARE	BE
XANAX Retard 0,5 mg comprimés à libération prolongée	not available	BE173223	VIATRIS HEALTHCARE	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
XANAX Retard 0,5 mg comprimés à libération prolongée	not available	BE173223	VIATRIS HEALTHCARE	BE
XANAX Retard 0,5 mg comprimés à libération prolongée	not available	BE173223	VIATRIS HEALTHCARE	BE
XANAX Retard 0,5 mg comprimés à libération prolongée	not available	2007079361	VIATRIS HEALTHCARE	LU
XANAX Retard 0,5 mg comprimés à libération prolongée	not available	2007079361	VIATRIS HEALTHCARE	LU
XANAX Retard 0,5 mg comprimés à libération prolongée	not available	2007079361	VIATRIS HEALTHCARE	LU
XANAX Retard 0,5 mg comprimés à libération prolongée	not available	2007079361	VIATRIS HEALTHCARE	LU
XANAX Retard 0,5 mg Retardtabletten	not available	BE173223	VIATRIS HEALTHCARE	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
XANAX Retard 0,5 mg Retardtabletten	not available	BE173223	VIATRIS HEALTHCARE	BE
XANAX Retard 0,5 mg Retardtabletten	not available	BE173223	VIATRIS HEALTHCARE	BE
XANAX Retard 0,5 mg Retardtabletten	not available	BE173223	VIATRIS HEALTHCARE	BE
XANAX Retard 0,5 mg Retardtabletten	not available	2007079361	VIATRIS HEALTHCARE	LU
XANAX Retard 0,5 mg Retardtabletten	not available	2007079361	VIATRIS HEALTHCARE	LU
XANAX Retard 0,5 mg Retardtabletten	not available	2007079361	VIATRIS HEALTHCARE	LU
XANAX Retard 0,5 mg Retardtabletten	not available	2007079361	VIATRIS HEALTHCARE	LU
XANAX Retard 0,5 mg tabletten met verlengde afgifte	not available	BE173223	VIATRIS HEALTHCARE	BE
XANAX Retard 0,5 mg tabletten met verlengde afgifte	not available	BE173223	VIATRIS HEALTHCARE	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
XANAX Retard 0,5 mg tabletten met verlengde afgifte	not available	BE173223	VIATRIS HEALTHCARE	BE
XANAX Retard 0,5 mg tabletten met verlengde afgifte	not available	BE173223	VIATRIS HEALTHCARE	BE
Xanax Retard 0,5, tabletten met gereguleerde afgifte 0,5 mg	not available	RVG 19886	VIATRIS HEALTHCARE LTD	NL
Xanax Retard 0,5, tabletten met gereguleerde afgifte 0,5 mg	not available	RVG 19886	VIATRIS HEALTHCARE LTD	NL
Xanax Retard 0,5, tabletten met gereguleerde afgifte 0,5 mg	not available	RVG 19886	VIATRIS HEALTHCARE LTD	NL
Xanax Retard 0,5, tabletten met gereguleerde afgifte 0,5 mg	not available	RVG 19886	VIATRIS HEALTHCARE LTD	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
Xanax Retard 0,5, tabletten met gereguleerde afgifte 0,5 mg	not available	RVG 19886	VIATRIS HEALTHCARE LTD	NL
Xanax Retard 0,5, tabletten met gereguleerde afgifte 0,5 mg	not available	RVG 19886	VIATRIS HEALTHCARE LTD	NL
Xanax Retard 0,5, tabletten met gereguleerde afgifte 0,5 mg	not available	RVG 19886	VIATRIS HEALTHCARE LTD	NL
XANAX Retard 1 mg comprimés à libération prolongée	not available	BE173232	VIATRIS HEALTHCARE	BE
XANAX Retard 1 mg comprimés à libération prolongée	not available	BE173232	VIATRIS HEALTHCARE	BE
XANAX Retard 1 mg comprimés à libération prolongée	not available	BE173232	VIATRIS HEALTHCARE	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
XANAX Retard 1 mg comprimés à libération prolongée	not available	BE173232	VIATRIS HEALTHCARE	BE
XANAX Retard 1 mg comprimés à libération prolongée	not available	2007079362	VIATRIS HEALTHCARE	LU
XANAX Retard 1 mg comprimés à libération prolongée	not available	2007079362	VIATRIS HEALTHCARE	LU
XANAX Retard 1 mg comprimés à libération prolongée	not available	2007079362	VIATRIS HEALTHCARE	LU
XANAX Retard 1 mg comprimés à libération prolongée	not available	2007079362	VIATRIS HEALTHCARE	LU
XANAX Retard 1 mg Retardtabletten	not available	BE173232	VIATRIS HEALTHCARE	BE
XANAX Retard 1 mg Retardtabletten	not available	BE173232	VIATRIS HEALTHCARE	BE
XANAX Retard 1 mg	not available	BE173232	VIATRIS HEALTHCARE	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
Retardtabletten				
XANAX Retard 1 mg Retardtabletten	not available	BE173232	VIATRIS HEALTHCARE	BE
XANAX Retard 1 mg Retardtabletten	not available	2007079362	VIATRIS HEALTHCARE	LU
XANAX Retard 1 mg Retardtabletten	not available	2007079362	VIATRIS HEALTHCARE	LU
XANAX Retard 1 mg Retardtabletten	not available	2007079362	VIATRIS HEALTHCARE	LU
XANAX Retard 1 mg Retardtabletten	not available	2007079362	VIATRIS HEALTHCARE	LU
XANAX Retard 1 mg tabletten met verlengde afgifte	not available	BE173232	VIATRIS HEALTHCARE	BE
XANAX Retard 1 mg tabletten met verlengde afgifte	not available	BE173232	VIATRIS HEALTHCARE	BE
XANAX Retard 1 mg tabletten met verlengde afgifte	not available	BE173232	VIATRIS HEALTHCARE	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
XANAX Retard 1 mg tabletten met verlengde afgifte	not available	BE173232	VIATRIS HEALTHCARE	BE
Xanax Retard 1, tabletten met gereguleerde afgifte 1 mg	not available	RVG 19887	VIATRIS HEALTHCARE LTD	NL
Xanax Retard 1, tabletten met gereguleerde afgifte 1 mg	not available	RVG 19887	VIATRIS HEALTHCARE LTD	NL
Xanax Retard 1, tabletten met gereguleerde afgifte 1 mg	not available	RVG 19887	VIATRIS HEALTHCARE LTD	NL
Xanax Retard 1, tabletten met gereguleerde afgifte 1 mg	not available	RVG 19887	VIATRIS HEALTHCARE LTD	NL
Xanax Retard 1, tabletten met gereguleerde afgifte 1 mg	not available	RVG 19887	VIATRIS HEALTHCARE LTD	NL
Xanax Retard 1, tabletten met gereguleerde afgifte 1 mg	not available	RVG 19887	VIATRIS HEALTHCARE LTD	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
Xanax Retard 1, tabletten met gereguleerde afgifte 1 mg	not available	RVG 19887	VIATRIS HEALTHCARE LTD	NL
XANAX Retard 2 mg comprimés à libération prolongée	not available	BE173241	VIATRIS HEALTHCARE	BE
XANAX Retard 2 mg comprimés à libération prolongée	not available	BE173241	VIATRIS HEALTHCARE	BE
XANAX Retard 2 mg comprimés à libération prolongée	not available	BE173241	VIATRIS HEALTHCARE	BE
XANAX Retard 2 mg comprimés à libération prolongée	not available	BE173241	VIATRIS HEALTHCARE	BE
XANAX Retard 2 mg comprimés à libération prolongée	not available	2007079363	VIATRIS HEALTHCARE	LU
XANAX Retard 2 mg comprimés à libération prolongée	not available	2007079363	VIATRIS HEALTHCARE	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
XANAX Retard 2 mg comprimés à libération prolongée	not available	2007079363	VIATRIS HEALTHCARE	LU
XANAX Retard 2 mg comprimés à libération prolongée	not available	2007079363	VIATRIS HEALTHCARE	LU
XANAX Retard 2 mg Retardtabletten	not available	BE173241	VIATRIS HEALTHCARE	BE
XANAX Retard 2 mg Retardtabletten	not available	BE173241	VIATRIS HEALTHCARE	BE
XANAX Retard 2 mg Retardtabletten	not available	BE173241	VIATRIS HEALTHCARE	BE
XANAX Retard 2 mg Retardtabletten	not available	BE173241	VIATRIS HEALTHCARE	BE
XANAX Retard 2 mg Retardtabletten	not available	2007079363	VIATRIS HEALTHCARE	LU
XANAX Retard 2 mg Retardtabletten	not available	2007079363	VIATRIS HEALTHCARE	LU
XANAX Retard 2 mg Retardtabletten	not available	2007079363	VIATRIS HEALTHCARE	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
XANAX Retard 2 mg Retardtabletten	not available	2007079363	VIATRIS HEALTHCARE	LU
XANAX Retard 2 mg tabletten met verlengde afgifte	not available	BE173241	VIATRIS HEALTHCARE	BE
XANAX Retard 2 mg tabletten met verlengde afgifte	not available	BE173241	VIATRIS HEALTHCARE	BE
XANAX Retard 2 mg tabletten met verlengde afgifte	not available	BE173241	VIATRIS HEALTHCARE	BE
XANAX Retard 2 mg tabletten met verlengde afgifte	not available	BE173241	VIATRIS HEALTHCARE	BE
Xanax Retard 2, tabletten met gereguleerde afgifte 2 mg	not available	RVG 19888	VIATRIS HEALTHCARE LTD	NL
Xanax Retard 2, tabletten met gereguleerde afgifte 2 mg	not available	RVG 19888	VIATRIS HEALTHCARE LTD	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
Xanax Retard 2, tabletten met gereguleerde afgifte 2 mg	not available	RVG 19888	VIATRIS HEALTHCARE LTD	NL
Xanax Retard 2, tabletten met gereguleerde afgifte 2 mg	not available	RVG 19888	VIATRIS HEALTHCARE LTD	NL
Xanax Retard 2, tabletten met gereguleerde afgifte 2 mg	not available	RVG 19888	VIATRIS HEALTHCARE LTD	NL
Xanax Retard 2, tabletten met gereguleerde afgifte 2 mg	not available	RVG 19888	VIATRIS HEALTHCARE LTD	NL
Xanax Retard 2, tabletten met gereguleerde afgifte 2 mg	not available	RVG 19888	VIATRIS HEALTHCARE LTD	NL
XANAX Retard 3 mg comprimés à libération prolongée	not available	BE173257	VIATRIS HEALTHCARE	BE
XANAX Retard 3 mg comprimés à libération prolongée	not available	BE173257	VIATRIS HEALTHCARE	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
XANAX Retard 3 mg comprimés à libération prolongée	not available	BE173257	VIATRIS HEALTHCARE	BE
XANAX Retard 3 mg comprimés à libération prolongée	not available	BE173257	VIATRIS HEALTHCARE	BE
XANAX Retard 3 mg comprimés à libération prolongée	not available	2007079364	VIATRIS HEALTHCARE	LU
XANAX Retard 3 mg comprimés à libération prolongée	not available	2007079364	VIATRIS HEALTHCARE	LU
XANAX Retard 3 mg comprimés à libération prolongée	not available	2007079364	VIATRIS HEALTHCARE	LU
XANAX Retard 3 mg comprimés à libération prolongée	not available	2007079364	VIATRIS HEALTHCARE	LU
XANAX Retard 3 mg Retardtabletten	not available	BE173257	VIATRIS HEALTHCARE	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
XANAX Retard 3 mg Retardtabletten	not available	BE173257	VIATRIS HEALTHCARE	BE
XANAX Retard 3 mg Retardtabletten	not available	BE173257	VIATRIS HEALTHCARE	BE
XANAX Retard 3 mg Retardtabletten	not available	BE173257	VIATRIS HEALTHCARE	BE
XANAX Retard 3 mg Retardtabletten	not available	2007079364	VIATRIS HEALTHCARE	LU
XANAX Retard 3 mg Retardtabletten	not available	2007079364	VIATRIS HEALTHCARE	LU
XANAX Retard 3 mg Retardtabletten	not available	2007079364	VIATRIS HEALTHCARE	LU
XANAX Retard 3 mg Retardtabletten	not available	2007079364	VIATRIS HEALTHCARE	LU
XANAX Retard 3 mg tabletten met verlengde afgifte	not available	BE173257	VIATRIS HEALTHCARE	BE
XANAX Retard 3 mg tabletten met verlengde afgifte	not available	BE173257	VIATRIS HEALTHCARE	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
XANAX Retard 3 mg tabletten met verlengde afgifte	not available	BE173257	VIATRIS HEALTHCARE	BE
XANAX Retard 3 mg tabletten met verlengde afgifte	not available	BE173257	VIATRIS HEALTHCARE	BE
XANAX SR 0,5 mg retard tabletta	not available	OGYI-T-4617/01	VIATRIS HEALTHCARE LIMITED	HU
Xanax SR 0,5 mg tablete s podaljšanim sproščanjem	not available	H/92/01672/004	UPJOHN EESV	SI
XANAX SR 0,5 mg tablete s produljenim oslobađanjem	not available	HR-H-219339638	UPJOHN EESV	HR
XANAX SR 0,5 mg tablety s predlženým uvolnovaním	not available	70/0132/95-S	UPJOHN EESV	SK
XANAX SR 0,5 mg tablety s predlženým uvolnovaním	not available	70/0132/95-S	UPJOHN EESV	SK
XANAX SR 0,5 mg tablety s predlženým	not available	70/0132/95-S	UPJOHN EESV	SK

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
uvolnovaním				
XANAX SR 0,5 mg tablety s prodlouženým uvolňováním	not available	70/735/95-A/C	UPJOHN EESV	CZ
XANAX SR 1 mg retard tabletta	not available	OGYI-T-4617/02	VIATRIS HEALTHCARE LIMITED	HU
Xanax SR 1 mg tablete s podaljšanim sproščanjem	not available	H/92/01672/005	UPJOHN EESV	SI
XANAX SR 1 mg tablete s produljenim oslobađanjem	not available	HR-H-466930724	UPJOHN EESV	HR
XANAX SR 1 mg tablety s predlženým uvolnovaním	not available	70/0054/17-S	UPJOHN EESV	SK
XANAX SR 1 mg tablety s predlženým uvolnovaním	not available	70/0054/17-S	UPJOHN EESV	SK
XANAX SR 1 mg tablety s predlženým uvolnovaním	not available	70/0054/17-S	UPJOHN EESV	SK
XANAX SR 1 mg tablety s prodlouženým uvolňováním	not available	70/735/95-B/C	UPJOHN EESV	CZ

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
XANAX SR 2 mg retard tabletta	not available	OGYI-T-4617/03	VIATRIS HEALTHCARE LIMITED	HU
Xanax SR 2 mg tablete s podaljšanim sproščanjem	not available	H/92/01672/006	UPJOHN EESV	SI
XANAX SR 2 mg tablete s produljenim oslobađanjem	not available	HR-H-020745058	UPJOHN EESV	HR
XANAX SR 2 mg tablety s predlženým uvolnovaním	not available	70/0055/17-S	UPJOHN EESV	SK
XANAX SR 2 mg tablety s predlženým uvolnovaním	not available	70/0055/17-S	UPJOHN EESV	SK
XANAX SR 2 mg tablety s predlženým uvolnovaním	not available	70/0055/17-S	UPJOHN EESV	SK
XANAX SR 2 mg tablety s prodlouženým uvolňováním	not available	70/735/95-C/C	UPJOHN EESV	CZ
XANAX SR 3 mg tablety s predlženým uvolnovaním	not available	70/0056/17-S	UPJOHN EESV	SK
XANAX SR 3 mg tablety s predlženým uvolnovaním	not available	70/0056/17-S	UPJOHN EESV	SK

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
XANAX SR 3 mg tablety s predlženým uvolnovaním	not available	70/0056/17-S	UPJOHN EESV	SK
XANAX Sublingualis 0,5 mg nyelvalatti tabletta	not available	OGYI-T-4617/10	VIATRIS HEALTHCARE LTD	HU
XANAX Sublingualis 0,5 mg nyelvalatti tabletta	not available	OGYI-T-4617/11	VIATRIS HEALTHCARE LTD	HU
XANAX Sublingualis 0,5 mg nyelvalatti tabletta	not available	OGYI-T-4617/09	VIATRIS HEALTHCARE LTD	HU
XANAX Sublingualis 0,5 mg nyelvalatti tabletta	not available	OGYI-T-4617/12	VIATRIS HEALTHCARE LTD	HU
XANAX Sublingualis 0,5 mg nyelvalatti tabletta	not available	OGYI-T-4617/13	VIATRIS HEALTHCARE LTD	HU
XANAX Sublingualis 0,5 mg nyelvalatti tabletta	not available	OGYI-T-4617/10	VIATRIS HEALTHCARE LTD	HU
XANAX Sublingualis 0,5 mg nyelvalatti tabletta	not available	OGYI-T-4617/11	VIATRIS HEALTHCARE LTD	HU
XANAX Sublingualis 0,5 mg nyelvalatti tabletta	not available	OGYI-T-4617/09	VIATRIS HEALTHCARE LTD	HU
XANAX Sublingualis 0,5	not available	OGYI-T-4617/12	VIATRIS HEALTHCARE LTD	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
mg nyelvalatti tabletta				
XANAX Sublingualis 0,5 mg nyelvalatti tabletta	not available	OGYI-T-4617/13	VIATRIS HEALTHCARE LTD	HU
Xanax XR 0,5 mg comprimidos de libertação modificada	not available	5816087	UPJOHN EESV	PT
Xanax XR 0,5 mg comprimidos de libertação modificada	not available	2381085	UPJOHN EESV	PT
Xanax XR 0,5 mg comprimidos de libertação modificada	not available	4579389	UPJOHN EESV	PT
Xanax XR 0,5 mg comprimidos de libertação modificada	not available	4579280	UPJOHN EESV	PT
XANAX XR 0,5 mg pailginto atpalaidavimo tabletės	not available	LT/1/96/2951/001	UPJOHN EESV	LT
Xanax XR 0,5 mg toimeainet prolongeeritult	not available	239798	UPJOHN EESV	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
vabastavad tabletid				
Xanax XR 1 mg comprimidos de libertação modificada	not available	5816186	UPJOHN EESV	PT
Xanax XR 1 mg comprimidos de libertação modificada	not available	4579488	UPJOHN EESV	PT
Xanax XR 1 mg comprimidos de libertação modificada	not available	2381184	UPJOHN EESV	PT
Xanax XR 1 mg comprimidos de libertação modificada	not available	4579587	UPJOHN EESV	PT
XANAX XR 1 mg pailginto atpalaidavimo tabletės	not available	LT/1/96/2951/002	UPJOHN EESV	LT
Xanax XR 1 mg toimeainet prolongeeritult vabastavad tabletid	not available	239698	UPJOHN EESV	EE
Xanax XR 2 mg comprimidos de	not available	4579785	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
libertação modificada				
Xanax XR 2 mg comprimidos de libertação modificada	not available	5816285	UPJOHN EESV	PT
Xanax XR 2 mg comprimidos de libertação modificada	not available	4579686	UPJOHN EESV	PT
Xanax XR 2 mg comprimidos de libertação modificada	not available	2381283	UPJOHN EESV	PT
Xanax XR 3 mg comprimidos de libertação modificada	not available	2381382	UPJOHN EESV	PT
Xanax XR 3 mg comprimidos de libertação modificada	not available	4579983	UPJOHN EESV	PT
Xanax XR 3 mg comprimidos de libertação modificada	not available	4579884	UPJOHN EESV	PT
Xanax XR 3 mg comprimidos de	not available	5816384	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
libertação modificada				
Xanax, 0,25 mg, tabletki	not available	R/0364	UPJOHN EESV	PL
Xanax, 0,25 mg, tabletki	not available	R/0364	UPJOHN EESV	PL
Xanax, 0,25 mg, tabletki	not available	R/0364	UPJOHN EESV	PL
Xanax, 0,25 mg, tabletki	not available	R/0364	UPJOHN EESV	PL
Xanax, 0,25 mg, tabletki	not available	R/0364	UPJOHN EESV	PL
Xanax, 0,25 mg, tabletki	not available	R/0364	UPJOHN EESV	PL
Xanax, 0,25 mg, tabletki	not available	R/0364	UPJOHN EESV	PL
Xanax, 0,25 mg, tabletki	not available	R/0364	UPJOHN EESV	PL
Xanax, 0,5 mg, tabletki	not available	R/0365	UPJOHN EESV	PL
Xanax, 0,5 mg, tabletki	not available	R/0365	UPJOHN EESV	PL
Xanax, 0,5 mg, tabletki	not available	R/0365	UPJOHN EESV	PL
Xanax, 0,5 mg, tabletki	not available	R/0365	UPJOHN EESV	PL
Xanax, 0,5 mg, tabletki	not available	R/0365	UPJOHN EESV	PL
Xanax, 0,5 mg, tabletki	not available	R/0365	UPJOHN EESV	PL
Xanax, 0,5 mg, tabletki	not available	R/0365	UPJOHN EESV	PL

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
Xanax, 0,5 mg, tabletki	not available	R/0365	UPJOHN EESV	PL
Xanax, 1 mg, tabletki	not available	R/0366	UPJOHN EESV	PL
Xanax, 1 mg, tabletki	not available	R/0366	UPJOHN EESV	PL
Xanax, 1 mg, tabletki	not available	R/0366	UPJOHN EESV	PL
Xanax, 1 mg, tabletki	not available	R/0366	UPJOHN EESV	PL
Xanax, 1 mg, tabletki	not available	R/0366	UPJOHN EESV	PL
Xanax, 1 mg, tabletki	not available	R/0366	UPJOHN EESV	PL
Xanax, 1 mg, tabletki	not available	R/0366	UPJOHN EESV	PL
Xanax, 1 mg, tabletki	not available	R/0366	UPJOHN EESV	PL
Xanax, 2 mg, tabletki	not available	R/0367	UPJOHN EESV	PL
Xanax, 2 mg, tabletki	not available	R/0367	UPJOHN EESV	PL
Xanax, 2 mg, tabletki	not available	R/0367	UPJOHN EESV	PL
Xanax, 2 mg, tabletki	not available	R/0367	UPJOHN EESV	PL
Xanax, 2 mg, tabletki	not available	R/0367	UPJOHN EESV	PL
Xanax, 2 mg, tabletki	not available	R/0367	UPJOHN EESV	PL
Xanax, 2 mg, tabletki	not available	R/0367	UPJOHN EESV	PL

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
Xanax, 2 mg, tabletki	not available	R/0367	UPJOHN EESV	PL
XANOR 0,25 mg tablett	not available	9384	UPJOHN EESV	FI
XANOR 0,25 mg tablett	not available	9384	UPJOHN EESV	FI
XANOR 0,25 mg tablett	not available	9384	UPJOHN EESV	FI
XANOR 0,25 mg tablett	not available	9384	UPJOHN EESV	FI
XANOR 0,25 mg tablett	not available	9384	UPJOHN EESV	FI
XANOR 0,25 mg tablett	not available	9384	UPJOHN EESV	FI
XANOR 0,25 mg tablett	not available	9384	UPJOHN EESV	FI
Xanor 0,25 mg tabletter	not available	8026	UPJOHN EESV	NO
Xanor 0,25 mg tabletter	not available	10176	UPJOHN EESV	SE
XANOR 0,25 mg tabletti	not available	9384	UPJOHN EESV	FI
XANOR 0,25 mg tabletti	not available	9384	UPJOHN EESV	FI
XANOR 0,25 mg tabletti	not available	9384	UPJOHN EESV	FI
XANOR 0,25 mg tabletti	not available	9384	UPJOHN EESV	FI
XANOR 0,25 mg tabletti	not available	9384	UPJOHN EESV	FI
XANOR 0,25 mg tabletti	not available	9384	UPJOHN EESV	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
XANOR 0,25 mg tabletti	not available	9384	UPJOHN EESV	FI
XANOR 0,5 mg tablett	not available	9385	UPJOHN EESV	FI
XANOR 0,5 mg tablett	not available	9385	UPJOHN EESV	FI
XANOR 0,5 mg tablett	not available	9385	UPJOHN EESV	FI
XANOR 0,5 mg tablett	not available	9385	UPJOHN EESV	FI
XANOR 0,5 mg tablett	not available	9385	UPJOHN EESV	FI
XANOR 0,5 mg tablett	not available	9385	UPJOHN EESV	FI
XANOR 0,5 mg tablett	not available	9385	UPJOHN EESV	FI
Xanor 0,5 mg tabletter	not available	8027	UPJOHN EESV	NO
Xanor 0,5 mg tabletter	not available	10177	UPJOHN EESV	SE
XANOR 0,5 mg tabletti	not available	9385	UPJOHN EESV	FI
XANOR 0,5 mg tabletti	not available	9385	UPJOHN EESV	FI
XANOR 0,5 mg tabletti	not available	9385	UPJOHN EESV	FI
XANOR 0,5 mg tabletti	not available	9385	UPJOHN EESV	FI
XANOR 0,5 mg tabletti	not available	9385	UPJOHN EESV	FI
XANOR 0,5 mg tabletti	not available	9385	UPJOHN EESV	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
XANOR 0,5 mg tabletti	not available	9385	UPJOHN EESV	FI
XANOR 1 mg tablett	not available	9657	UPJOHN EESV	FI
XANOR 1 mg tablett	not available	9657	UPJOHN EESV	FI
XANOR 1 mg tablett	not available	9657	UPJOHN EESV	FI
XANOR 1 mg tablett	not available	9657	UPJOHN EESV	FI
XANOR 1 mg tablett	not available	9657	UPJOHN EESV	FI
XANOR 1 mg tablett	not available	9657	UPJOHN EESV	FI
XANOR 1 mg tablett	not available	9657	UPJOHN EESV	FI
Xanor 1 mg tabletter	not available	8028	UPJOHN EESV	NO
Xanor 1 mg tabletter	not available	10382	UPJOHN EESV	SE
XANOR 1 mg tabletti	not available	9657	UPJOHN EESV	FI
XANOR 1 mg tabletti	not available	9657	UPJOHN EESV	FI
XANOR 1 mg tabletti	not available	9657	UPJOHN EESV	FI
XANOR 1 mg tabletti	not available	9657	UPJOHN EESV	FI
XANOR 1 mg tabletti	not available	9657	UPJOHN EESV	FI
XANOR 1 mg tabletti	not available	9657	UPJOHN EESV	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
XANOR 1 mg tabletti	not available	9657	UPJOHN EESV	FI
XANOR 2 mg tablett	not available	10729	UPJOHN EESV	FI
XANOR 2 mg tablett	not available	10729	UPJOHN EESV	FI
XANOR 2 mg tablett	not available	10729	UPJOHN EESV	FI
XANOR 2 mg tablett	not available	10729	UPJOHN EESV	FI
XANOR 2 mg tablett	not available	10729	UPJOHN EESV	FI
XANOR 2 mg tablett	not available	10729	UPJOHN EESV	FI
XANOR 2 mg tablett	not available	10729	UPJOHN EESV	FI
Xanor 2 mg tabletter	not available	8029	UPJOHN EESV	NO
Xanor 2 mg tabletter	not available	11424	UPJOHN EESV	SE
XANOR 2 mg tabletti	not available	10729	UPJOHN EESV	FI
XANOR 2 mg tabletti	not available	10729	UPJOHN EESV	FI
XANOR 2 mg tabletti	not available	10729	UPJOHN EESV	FI
XANOR 2 mg tabletti	not available	10729	UPJOHN EESV	FI
XANOR 2 mg tabletti	not available	10729	UPJOHN EESV	FI
XANOR 2 mg tabletti	not available	10729	UPJOHN EESV	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
XANOR 2 mg tabletti	not available	10729	UPJOHN EESV	FI
XANOR DEPOT 0,5 mg depottablett	not available	11534	UPJOHN EESV	FI
XANOR DEPOT 0,5 mg depottablett	not available	11534	UPJOHN EESV	FI
XANOR DEPOT 0,5 mg depottablett	not available	11534	UPJOHN EESV	FI
XANOR DEPOT 0,5 mg depottablett	not available	11534	UPJOHN EESV	FI
Xanor Depot 0,5 mg depottabletter	not available	95-2110	UPJOHN EESV	NO
Xanor Depot 0,5 mg depottabletter	not available	95-2110	UPJOHN EESV	NO
Xanor Depot 0,5 mg depottabletter	not available	95-2110	UPJOHN EESV	NO
Xanor Depot 0,5 mg depottabletter	not available	95-2110	UPJOHN EESV	NO
Xanor Depot 0,5 mg depottabletter	not available	95-2110	UPJOHN EESV	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
Xanor Depot 0,5 mg depottabletter	not available	95-2110	UPJOHN EESV	NO
Xanor Depot 0,5 mg depottabletter	not available	95-2110	UPJOHN EESV	NO
Xanor Depot 0,5 mg depottabletter	not available	12757	UPJOHN EESV	SE
Xanor Depot 0,5 mg depottabletter	not available	12757	UPJOHN EESV	SE
Xanor Depot 0,5 mg depottabletter	not available	12757	UPJOHN EESV	SE
Xanor Depot 0,5 mg depottabletter	not available	12757	UPJOHN EESV	SE
Xanor Depot 0,5 mg depottabletter	not available	12757	UPJOHN EESV	SE
Xanor Depot 0,5 mg depottabletter	not available	12757	UPJOHN EESV	SE
Xanor Depot 0,5 mg depottabletter	not available	12757	UPJOHN EESV	SE
XANOR DEPOT 0,5 mg	not available	11534	UPJOHN EESV	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
depottabletti				
XANOR DEPOT 0,5 mg depottabletti	not available	11534	UPJOHN EESV	FI
XANOR DEPOT 0,5 mg depottabletti	not available	11534	UPJOHN EESV	FI
XANOR DEPOT 0,5 mg depottabletti	not available	11534	UPJOHN EESV	FI
XANOR DEPOT 1 mg depottablett	not available	11535	UPJOHN EESV	FI
XANOR DEPOT 1 mg depottablett	not available	11535	UPJOHN EESV	FI
XANOR DEPOT 1 mg depottablett	not available	11535	UPJOHN EESV	FI
XANOR DEPOT 1 mg depottablett	not available	11535	UPJOHN EESV	FI
Xanor Depot 1 mg depottabletter	not available	12758	UPJOHN EESV	SE
Xanor Depot 1 mg depottabletter	not available	12758	UPJOHN EESV	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
Xanor Depot 1 mg depottabletter	not available	12758	UPJOHN EESV	SE
Xanor Depot 1 mg depottabletter	not available	12758	UPJOHN EESV	SE
Xanor Depot 1 mg depottabletter	not available	12758	UPJOHN EESV	SE
Xanor Depot 1 mg depottabletter	not available	12758	UPJOHN EESV	SE
Xanor Depot 1 mg depottabletter	not available	12758	UPJOHN EESV	SE
XANOR DEPOT 1 mg depottabletti	not available	11535	UPJOHN EESV	FI
XANOR DEPOT 1 mg depottabletti	not available	11535	UPJOHN EESV	FI
XANOR DEPOT 1 mg depottabletti	not available	11535	UPJOHN EESV	FI
XANOR DEPOT 1 mg depottabletti	not available	11535	UPJOHN EESV	FI
XANOR DEPOT 2 mg	not available	11536	UPJOHN EESV	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
depottablett				
XANOR DEPOT 2 mg depottablett	not available	11536	UPJOHN EESV	FI
XANOR DEPOT 2 mg depottablett	not available	11536	UPJOHN EESV	FI
XANOR DEPOT 2 mg depottablett	not available	11536	UPJOHN EESV	FI
Xanor Depot 2 mg depottabletter	not available	12759	UPJOHN EESV	SE
Xanor Depot 2 mg depottabletter	not available	12759	UPJOHN EESV	SE
Xanor Depot 2 mg depottabletter	not available	12759	UPJOHN EESV	SE
Xanor Depot 2 mg depottabletter	not available	12759	UPJOHN EESV	SE
Xanor Depot 2 mg depottabletter	not available	12759	UPJOHN EESV	SE
Xanor Depot 2 mg depottabletter	not available	12759	UPJOHN EESV	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
Xanor Depot 2 mg depottabletter	not available	12759	UPJOHN EESV	SE
XANOR DEPOT 2 mg depottabletti	not available	11536	UPJOHN EESV	FI
XANOR DEPOT 2 mg depottabletti	not available	11536	UPJOHN EESV	FI
XANOR DEPOT 2 mg depottabletti	not available	11536	UPJOHN EESV	FI
XANOR DEPOT 2 mg depottabletti	not available	11536	UPJOHN EESV	FI
Xanor Depot 3 mg depottabletter	not available	95-2113	UPJOHN EESV	NO
Xanor Depot 3 mg depottabletter	not available	95-2113	UPJOHN EESV	NO
Xanor Depot 3 mg depottabletter	not available	95-2113	UPJOHN EESV	NO
Xanor Depot 3 mg depottabletter	not available	95-2113	UPJOHN EESV	NO
Xanor Depot 3 mg	not available	95-2113	UPJOHN EESV	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
depottabletter				
Xanor Depot 3 mg depottabletter	not available	95-2113	UPJOHN EESV	NO
Xanor Depot 3 mg depottabletter	not available	95-2113	UPJOHN EESV	NO
Xanor® 0,5 mg – Tabletten	not available	1-18639	UPJOHN EESV	AT
Xanor® 0,5 mg – Tabletten	not available	1-18639	UPJOHN EESV	AT
Xanor® 0,5 mg – Tabletten	not available	1-18639	UPJOHN EESV	AT
Xanor® 0,5 mg – Tabletten	not available	1-18639	UPJOHN EESV	AT
Xanor® 0,5 mg – Tabletten	not available	1-18639	UPJOHN EESV	AT
Xanor® 0,5 mg – Tabletten	not available	1-18639	UPJOHN EESV	AT
Xanor® 0,5 mg – Tabletten	not available	1-18639	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
Xanor® 1 mg – Tabletten	not available	1-18638	UPJOHN EESV	AT
Xanor® 1 mg – Tabletten	not available	1-18638	UPJOHN EESV	AT
Xanor® 1 mg – Tabletten	not available	1-18638	UPJOHN EESV	AT
Xanor® 1 mg – Tabletten	not available	1-18638	UPJOHN EESV	AT
Xanor® 1 mg – Tabletten	not available	1-18638	UPJOHN EESV	AT
Xanor® 1 mg – Tabletten	not available	1-18638	UPJOHN EESV	AT
Xanor® 1 mg – Tabletten	not available	1-18638	UPJOHN EESV	AT
КСАНАКС 0,25 mg таблетки	not available	20020166	UPJOHN EESV	BG
КСАНАКС 0,5 mg таблетки	not available	20020165	UPJOHN EESV	BG