



EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

10 July 2025
EMADOC-1700519818-2315708
Human Medicines Division

List of nationally authorised medicinal products

Active substance(s): lornoxicam

EURD list No. PSUSA/00001911/202412

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

© European Medicines Agency, 2025. Reproduction is authorised provided the source is acknowledged.



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
Xefo 4 mg - Filmtabletten	DK/H/0123/001	1-21851	TAKEDA AUSTRIA GMBH	AT
Xefo 8 mg - Filmtabletten	DK/H/0123/002	1-21852	TAKEDA AUSTRIA GMBH	AT
Xefo 8 mg Pulver und Lösungsmittel zur Herstellung einer Injektionslösung	DK/H/0123/005	1-22673	TAKEDA AUSTRIA GMBH	AT
Xefo Rapid 8 mg - Filmtabletten	DK/H/0123/006	1-24965	TAKEDA AUSTRIA GMBH	AT
КСЕФО 8 mg прах и разтворител за инжекционен разтвор (4mg/ml, 2 ml)	DK/H/0123/005	II-12721/29.03.2011	TAKEDA AUSTRIA GMBH	BG
Ксефо Репид 8 mg филмирани таблетки	DK/H/0123/006	II-12722/29.03.2011	TAKEDA AUSTRIA GMBH	BG
Lornoxicam "Takeda", filmovertrukne tabletter	DK/H/0137/001	19246	TAKEDA PHARMA A/S	DK
Lornoxicam "Takeda", filmovertrukne tabletter	DK/H/0137/002	19247	TAKEDA PHARMA A/S	DK
Lornoxicam "Takeda", pulver og solvens til injektionsvæske, opløsning	DK/H/0137/005	30643	TAKEDA PHARMA A/S	DK
Xefo Rapid, filmovertrukne tabletter	DK/H/0123/006	31688	TAKEDA PHARMA A/S	DK
Xefo, filmovertrukne tabletter	DK/H/0123/001	15784	TAKEDA PHARMA A/S	DK
Xefo, filmovertrukne tabletter	DK/H/0123/002	15785	TAKEDA PHARMA A/S	DK
Xefo, pulver og solvens til injektionsvæske, opløsning	DK/H/0123/005	18241	TAKEDA PHARMA A/S	DK
Xefo 4 mg επικαλυμμένο με λεπτό υμένιο δισκίο	DK/H/0123/001	92082/07-09-2023	TAKEDA HELLAS S.A.	GR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Xefo 8 mg κόνις και διαλύτης για ενέσιμο διάλυμα	DK/H/0123/005	92084/07-09-2023	TAKEDA HELLAS S.A.	GR
Xefo Rapid 8 mg επικαλυμμένο με λεπτό υμένιο δισκίο	DK/H/0123/006	92083/07-09-2023	TAKEDA HELLAS S.A.	GR
Xefo 4 mg filmdabletta	DK/H/0123/001	OGYI-T-8545/12	TAKEDA AUSTRIA GMBH	HU
Xefo 4 mg filmdabletta	DK/H/0123/001	OGYI-T-8545/08	TAKEDA AUSTRIA GMBH	HU
Xefo 4 mg filmdabletta	DK/H/0123/001	OGYI-T-8545/09	TAKEDA AUSTRIA GMBH	HU
Xefo 4 mg filmdabletta	DK/H/0123/001	OGYI-T-8545/10	TAKEDA AUSTRIA GMBH	HU
Xefo 4 mg filmdabletta	DK/H/0123/001	OGYI-T-8545/11	TAKEDA AUSTRIA GMBH	HU
Xefo 8 mg filmdabletta	DK/H/0123/002	OGYI-T-8545/01	TAKEDA AUSTRIA GMBH	HU
Xefo 8 mg filmdabletta	DK/H/0123/002	OGYI-T-8545/02	TAKEDA AUSTRIA GMBH	HU
Xefo 8 mg filmdabletta	DK/H/0123/002	OGYI-T-8545/03	TAKEDA AUSTRIA GMBH	HU
Xefo 8 mg filmdabletta	DK/H/0123/002	OGYI-T-8545/04	TAKEDA AUSTRIA GMBH	HU
Xefo 8 mg filmdabletta	DK/H/0123/002	OGYI-T-8545/05	TAKEDA AUSTRIA GMBH	HU
Xefo Rapid 8 mg filmdabletta	DK/H/0123/006	OGYI-T-8545/06	TAKEDA AUSTRIA GMBH	HU
Xefo Rapid 8 mg filmdabletta	DK/H/0123/006	OGYI-T-8545/07	TAKEDA AUSTRIA GMBH	HU
xefo 4 mg plevele dengtos tabletes	DK/H/0123/001	LT/1/98/2432/003	TAKEDA AUSTRIA GMBH	LT
xefo 4 mg plèvele dengtos tabletės	DK/H/0123/001	LT/1/98/2432/001	TAKEDA AUSTRIA GMBH	LT
xefo 4 mg plèvele dengtos tabletės	DK/H/0123/001	LT/1/98/2432/002	TAKEDA AUSTRIA GMBH	LT
xefo 4 mg plèvele dengtos tabletės	DK/H/0123/001	LT/1/98/2432/004	TAKEDA AUSTRIA GMBH	LT
xefo 4 mg plèvele dengtos tabletės	DK/H/0123/001	LT/1/98/2432/005	TAKEDA AUSTRIA GMBH	LT
xefo 8 mg milteliai ir tirpiklis injekciniam tirpalui	DK/H/0123/005	LT/1/98/2432/015	TAKEDA AUSTRIA GMBH	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
xefo 8 mg milteliai ir tirpiklis injekciniam tirpalui	DK/H/0123/005	LT/1/98/2432/016	TAKEDA AUSTRIA GMBH	LT
xefo 8 mg milteliai ir tirpiklis injekciniam tirpalui	DK/H/0123/005	LT/1/98/2432/017	TAKEDA AUSTRIA GMBH	LT
xefo 8 mg milteliai ir tirpiklis injekciniam tirpalui	DK/H/0123/005	LT/1/98/2432/018	TAKEDA AUSTRIA GMBH	LT
xefo 8 mg plevele dengtos tabletės	DK/H/0123/002	LT/1/98/2432/012	TAKEDA AUSTRIA GMBH	LT
xefo 8 mg plėvele dengtos tabletės	DK/H/0123/002	LT/1/98/2432/008	TAKEDA AUSTRIA GMBH	LT
xefo 8 mg plėvele dengtos tabletės	DK/H/0123/002	LT/1/98/2432/009	TAKEDA AUSTRIA GMBH	LT
xefo 8 mg plėvele dengtos tabletės	DK/H/0123/002	LT/1/98/2432/010	TAKEDA AUSTRIA GMBH	LT
xefo 8 mg plėvele dengtos tabletės	DK/H/0123/002	LT/1/98/2432/011	TAKEDA AUSTRIA GMBH	LT
Xefo 4 mg apvalkotās tabletes	DK/H/0123/001	99-0202	TAKEDA AUSTRIA GMBH	LV
Xefo Rapid 8 mg tabletki powlekane	DK/H/0123/006	15430	TAKEDA PHARMA SP.Z.O.O.	PL
Xefo Rapid 8 mg comprimate filmate	DK/H/0123/006	3732/2011/01	TAKEDA AUSTRIA GMBH	RO
Xefo Rapid 8 mg comprimate filmate	DK/H/0123/006	3732/2011/02	TAKEDA AUSTRIA GMBH	RO
Xefo Rapid 8 mg comprimate filmate	DK/H/0123/006	3732/2011/03	TAKEDA AUSTRIA GMBH	RO
Xefo Rapid 8 mg comprimate filmate	DK/H/0123/006	3732/2011/04	TAKEDA AUSTRIA GMBH	RO
Xefo Rapid 8 mg comprimate filmate	DK/H/0123/006	3732/2011/05	TAKEDA AUSTRIA GMBH	RO
Xefo Rapid 8 mg comprimate filmate	DK/H/0123/006	3732/2011/06	TAKEDA AUSTRIA GMBH	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
Xefo Rapid 8 mg comprimate filmate	DK/H/0123/006	3732/2011/07	TAKEDA AUSTRIA GMBH	RO
Xefo 4 mg filmom obalené tablety	DK/H/0123/001	29/0596/08-S	TAKEDA PHARMA A/S	SK
Xefo 4 mg/ml prášok a rozpúšťadlo na injekčný roztok	DK/H/0123/005	29/0598/08-S	TAKEDA PHARMA A/S	SK
Xefo 8 mg filmom obalené tablety	DK/H/0123/002	29/0597/08-S	TAKEDA PHARMA A/S	SK
Xefo Rapid filmom obalené tablety	DK/H/0123/006	29/0599/08-S	TAKEDA PHARMA A/S	SK