



EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

27 November 2025
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: sulprostone

Procedure no.: PSUSA/00002828/202504

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
Nalador		RVG 08996	Teofarma S.r.l.	Kingdom of the Netherlands
Nalador-500		6714.00.00	Teofarma S.r.l.	Federal Republic of Germany
Nalador		1-17577	Teofarma S.r.l.	Republic of Austria
NALADOR		34009 550 796 1 3	Teofarma S.r.l.	French Republic
Nalador 500			Teofarma S.r.l.	Portuguese Republic
Nalador		025998030	Teofarma S.r.l.	Italian Republic
Nalador		1-17577	Teofarma S.r.l.	Republic of Austria
Nalador		OGYI-T-4024/02	Teofarma S.r.l.	Republic of Hungary
Nalador		RVG 08996	Teofarma S.r.l.	Kingdom of the Netherlands
NALADOR		34009 555 252 0 2	Teofarma S.r.l.	French Republic
Nalador		OGYI-T-4024/01	Teofarma S.r.l.	Republic of Hungary