14 November 2018
EMA/854912/2018
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: Magnesium sulphate heptahydrate, sodium sulphate anhydrous, potassium sulphate

Procedure no.: EMEA/H/N/PSR/S/0016

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Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Izinova	FR/H/0511/001/DC	NL41696	Ipsen Pharma	France (RMS)
Eziclen	FR/H/0511/001/DC	BE434323	Ipsen NV	Belgium
Eziclen	FR/H/0511/001/DC	61/106/13-C	Ipsen Pharma	Czech Republic
Eziclen	FR/H/0511/001/DC	808713	Ipsen Pharma	Estonia
Eziclen	FR/H/0511/001/DC	86203.00.00	Ipsen Pharma GmbH	Germany
Eziclen	FR/H/0511/001/DC	300760101	Ipsen epe	Greece
Izinova	FR/H/0511/001/DC	041807013/M	Ipsen S.p.A.	Italy
Eziclen	FR/H/0511/001/DC	13-0037	Ipsen Pharma	Latvia
Eziclen	FR/H/0511/001/DC	LT/1/13/3275/001	Ipsen Pharma	Lithuania
Eziclen	FR/H/0511/001/DC	2013080288	Ipsen NV	Luxembourg
Eziclen	FR/H/0511/001/DC	RVG 110863	Ipsen Farmaceutica BV	Netherlands
Eziclen	FR/H/0511/001/DC	21110	Ipsen Pharma	Poland
Eziclen	FR/H/0511/001/DC	5573555	Ipsen Portugal-Produtos Farmaceuticos, S.A.	Portugal
Eziclen	FR/H/0511/001/DC	5417/2013/01	Ipsen Pharma	Romania
Eziclen	FR/H/0511/001/DC	77754	Ipsen Pharma S.A.	Spain
Izinova	FR/H/0511/001/DC	PL 34926/0016	Ipsen Limited	United Kingdom