12 April 2018
EMA/227734/2018
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance(s): buprenorphine

Procedure No.: PSUSA/00000459/201707

Digitally signed by Chirtoacă Iurie
Date: 2023.08.01 17:29:16 EEST
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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
Natzon 2 mg Sublingual Tablets	UK/H/1827/002	PL 20117/0119	MORNINGSIDE HEALTHCARE LTD	UK
Natzon 8 mg Sublingual Tablets	UK/H/1827/003	PL 20117/0120	MORNINGSIDE HEALTHCARE LTD	UK
Nimedol 0,4 mg sublingvalne tablete	PT/H/1288/001	HR-H-221990733	PLIVA HRVATSKA D.O.O.	HR
Nimedol 2 mg sublingvalne tablete	PT/H/1288/002	HR-H-056604005	PLIVA HRVATSKA D.O.O.	HR
Nimedol 8 mg sublingvalne tablete	PT/H/1288/003	HR-H-481991441	PLIVA HRVATSKA D.O.O.	HR
Noprex 35 mikrogramov/h transdermálna náplasť	DE/H/5080/001	65/0161/16-S	GLENMARK PHARMACEUTICALS S.R.O.	SK
Noprex 35 mikrogramů/h transdermální náplast	DE/H/5080/001	65/280/16-C	GLENMARK PHARMACEUTICALS S.R.O.	CZ
Noprex 52,5 mikrogramů/h transdermální náplast	DE/H/5080/002	65/281/16-C	GLENMARK PHARMACEUTICALS S.R.O.	CZ
Noprex 52.5 mikrogramov/h transdermálna náplasť	DE/H/5080/002	65/0162/16-S	GLENMARK PHARMACEUTICALS S.R.O.	SK
Noprex 70 mikrogramov/h transdermálna náplasť	DE/H/5080/003	65/0163/16-S	GLENMARK PHARMACEUTICALS S.R.O.	SK
Noprex 70 mikrogramů/h transdermální náplast	DE/H/5080/003	65/282/16-C	GLENMARK PHARMACEUTICALS S.R.O.	CZ
Norfinox, 52,5 mikrograma/godzinę, system transdermalny, plaster	DE/H/4515/002	23434	SANDOZ GMBH	PL
Norfinox, 70 mikrogramów/godzinę, system transdermalny, plaster	DE/H/4515/003	23435	SANDOZ GMBH	PL
Norfinox,35 mikrogramów/godzinę, system transdermalny, plaster	DE/H/4515/001	23433	SANDOZ GMBH	PL
Norspan 40 mikrog/t depotlaastari	DK/H/0718/007	31551	MUNDIPHARMA OY	FI