



EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

12 July 2018
EMA/613008/2018
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: ciprofloxacin hydrochloride / hydrocortisone

Procedure no.: PSUSA/00000774/201711



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
Ciprobay HC, körvatilgad suspensioonina	not available	318300	ALCON CUSI, S.A	EE
Ciflox, øredråber, suspension	not available	30805	NOVARTIS HEALTHCARE A/S	DK
CIPROXIN* HC OTIC, ear drops, suspension	not available	MA 088/06101	NOVARTIS PHARMACEUTICALS UK LIMITED	MT
Ciproxin-Hydrocortison 2 mg/ml + 10 mg/ml, eyrnadropar, dreifa	not available	980196 (IS)	NOVARTIS HEALTHCARE A/S	IS
CIPROXINA 2 mg/ml + 10 mg/ml gotas óticas en suspensión	ES/H/0175/001	63.285	NOVARTIS FARMACÉUTICA S.A.	ES
MEDIFLOX 2 mg/ml + 10 mg/ml gocce auricolari, sospensione.	ES/H/0175/001	035271016	NOVARTIS FARMA S.P.A.	IT
Ciproxin-Hydrocortison 2 mg/ml + 10 mg/ml öronddropar, suspension	not available	13316	NOVARTIS FINLAND OY	FI
Ciproxin-Hydrocortison 2 mg/ml + 10 mg/ml korvatipat, suspensio	not available	13316	NOVARTIS FINLAND OY	FI