8 March 2018
EMA/113527/2018
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: hydrochlorothiazide moexipril

Procedure no.: PSUSA/00002082/201706

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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
Femipres Plus 15 mg/25 mg compresse rivestite con film	IT/H/321/001	033907027	UCB PHARMA SPA (MILAN IT)	IT
ENULID 15 mg/25 mg compresse rivestite con film	IT/H/0321/001	033908029	MEDA PHARMA S.P.A.	IT
ENULID 15 mg/25 mg compresse rivestite con film	IT/H/0321/001	033908029	MEDA PHARMA S.P.A.	IT
Fempress plus 15 mg/25 mg	DE/H/3397/001	42479.01.00	PUREN PHARMA GMBH & CO. KG	DE
ENULID 15 mg/25 mg compresse rivestite con film	IT/H/0321/001	033908029	MEDA PHARMA S.P.A.	IT
ENULID 15 mg/25 mg compresse rivestite con film	IT/H/0321/001	033908029	MEDA PHARMA S.P.A.	IT