



EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

12 January 2017
EMA/45667/2017
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance(s): benazepril / hydrochlorothiazide

Procedure No.: PSUSA/00000314/201605



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
BRIAZIDE 10 mg/12,5 mg, comprimé pelliculé sécable	not available	34009 335 310 1 0	MEDA PHARMA SAS	FR
BRIAZIDE 10 mg/12,5 mg, comprimé pelliculé sécable	not available	34009 335 311 8 8	MEDA PHARMA SAS	FR
BRIAZIDE 10 mg/12,5 mg, comprimé pelliculé sécable	not available	34009 372 388 0 9	MEDA PHARMA SAS	FR
BRIAZIDE 10 mg/12,5 mg, comprimé pelliculé sécable	not available	34009 372 389 7 7	MEDA PHARMA SAS	FR
BRIAZIDE 10 mg/12,5 mg, comprimé pelliculé sécable	not available	34009 372 390 5 9	MEDA PHARMA SAS	FR
CIBADREX 10 mg + 12.5 mg compresse rivestite con film	not available	028037024	MEDA PHARMA S.P.A.	IT
CIBADREX 10 mg + 12.5 mg compresse rivestite con film	not available	028037048	MEDA PHARMA S.P.A.	IT
Cibadrex 10 mg/12,5 mg Filmdabletten	not available	25958.01.00	MEDA PHARMA GMBH & CO. KG	DE
CIBADREX 10 mg/12,5 mg, comprimé pelliculé sécable	not available	335 312-4	MEDA PHARMA SAS	FR
CIBADREX 10 mg/12,5 mg, comprimé pelliculé sécable	not available	371 559-6	MEDA PHARMA SAS	FR
CIBADREX 10 mg/12,5 mg, comprimé pelliculé sécable	not available	335 313-0	MEDA PHARMA SAS	FR
CIBADREX 10 mg/12,5 mg, comprimé pelliculé sécable	not available	371 560-4	MEDA PHARMA SAS	FR
CIBADREX 10 mg/12,5 mg, comprimé pelliculé sécable	not available	371 561-0	MEDA PHARMA SAS	FR
CIBADREX 20 mg + 25 mg compresse rivestite con film	not available	028037036	MEDA PHARMA S.P.A.	IT
Cibadrex 20 mg/25 mg film-coated tablets / Цибадрекс 20 mg/25 mg филмирани таблетки	not available	20050055	NOVARTIS PHARMA GMBH	BG
Cibadrex 20 mg/25mg Filmdabletten	not available	25958.02.00	MEDA PHARMA GMBH & CO. KG	DE
CIBADREX 5 mg + 6,25 mg compresse rivestite con film	not available	028037012	MEDA PHARMA S.P.A.	IT
Cibadrex 5 mg/6,25 mg Filmdabletten	not available	25958.00.00	MEDA PHARMA GMBH & CO. KG	DE

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
Cibadrex® 10 mg / 12,5 mg comprimidos recubiertos	not available	65860	MEDA PHARMA S.L.	ES
Cibadrex® 10 mg/12,5 mg Filmtabletten	not available	2005038699	MEDA PHARMA GMBH & CO. KG	LU
Cibadrex® 20 mg / 25 mg comprimidos recubiertos	not available	65859	MEDA PHARMA S.L.	ES
Cibadrex® επικαλυμμένα με λεπτό υμένιο δισκία των (10+12,5) mg	not available	25627/01-04-2013	MEDA PHARMACEUTICALS S.A.	GR
Cibadrex® επικαλυμμένα με λεπτό υμένιο δισκία των (20+25) mg	not available	25628/01-04-2013	MEDA PHARMACEUTICALS S.A.	GR
Cibadrex® επικαλυμμένα με λεπτό υμένιο δισκία των (5+6,25) mg	not available	70576/12/01-04-2013	MEDA PHARMACEUTICALS S.A.	GR
Lotensin HCT 10 mg/12.5 mg filmtabletta	not available	OGYI-T-4506/03	MEDA PHARMA HUNGARY KFT.	HU
Lotensin HCT 10 mg/12.5 mg filmtabletta	not available	OGYI-T-4506/04	MEDA PHARMA HUNGARY KFT.	HU
Lotensin HCT 5 mg/6,25 mg filmtabletta	not available	OGYI-T-4506/01	MEDA PHARMA HUNGARY KFT.	HU
Lotensin HCT 5 mg/6,25 mg filmtabletta	not available	OGYI-T-4506/02	MEDA PHARMA HUNGARY KFT.	HU
TENSADIUR "10mg + 12,5 mg compresse rivestite con film" 14 compresse	not available	028211023	MEDA PHARMA S.P.A.	IT
TENSADIUR "10mg + 12,5 mg compresse rivestite con film" 28 compresse	not available	028211047	MEDA PHARMA S.P.A.	IT
TENSADIUR "20 mg + 25 mg compresse rivestite con film" 14 compresse	not available	028211035	MEDA PHARMA S.P.A.	IT
TENSADIUR "5 mg + 6,25 mg compresse rivestite con film" 14 compresse	not available	028211011	MEDA PHARMA S.P.A.	IT
ZINADIUR 10 mg + 12,5 mg compresse rivestite con film	not available	028193011	ERREKAPPA EUROTHERAPICI SPA	IT

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
ZINADIUR 10 mg + 12,5 mg compresse rivestite con film	not available	028193047	ERREKAPPA EUROTHERAPICI SPA	IT
ZINADIUR 20 mg + 25 mg compresse rivestite con film	not available	028193035	ERREKAPPA EUROTHERAPICI SPA	IT
ZINADIUR 5 mg + 6,25 mg compresse rivestite con film	not available	028193023	ERREKAPPA EUROTHERAPICI SPA	IT