11 January 2018
EMA/415684/2018
Human Medicines Evaluation Division

List of nationally authorised medicinal

Active substance(s): candesartan, candesartan / hydrochlorothiazide

Procedure No.: PSUSA/00000527/201704

30 Churchill Place ● Canary Wharf ● London E14 5EU ● United Kingdom
Telephone +44 (0)20 3660 6000 **Facsimile** +44 (0)20 3660 5525
Send a question via our website www.ema.europa.eu/contact

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
Amias 16 mg Tablets	not available	PL 16189/0004	TAKEDA UK LTD	UK
Amias 2 mg Tablets	not available	PL 16189/0001	TAKEDA UK LTD	UK
Amias 32 mg Tablets	not available	PL 16189/0007	TAKEDA UK LTD	UK
Amias 4 mg Tablets	not available	PL 16189/0002	TAKEDA UK LTD	UK
Amias 8 mg Tablets	not available	PL 16189/0003	TAKEDA UK LTD	UK
Atacand 16 mg – Tabletten	UK/H/0197/004	1-22289	ASTRAZENECA OSTERREICH GMBH	AT
Atacand 16 mg comprimate	UK/H/0197/004	6233/2014/15	ASTRAZENECA AB	RO
Atacand 16 mg comprimate	UK/H/0197/004	6233/2014/05	ASTRAZENECA AB	RO
Atacand 16 mg comprimate	UK/H/0197/004	6233/2014/02	ASTRAZENECA AB	RO
Atacand 16 mg comprimate	UK/H/0197/004	6233/2014/10	ASTRAZENECA AB	RO
Atacand 16 mg comprimate	UK/H/0197/004	6233/2014/13	ASTRAZENECA AB	RO
Atacand 16 mg comprimate	UK/H/0197/004	6233/2014/04	ASTRAZENECA AB	RO
Atacand 16 mg comprimate	UK/H/0197/004	6233/2014/16	ASTRAZENECA AB	RO
Atacand 16 mg comprimate	UK/H/0197/004	6233/2014/14	ASTRAZENECA AB	RO
Atacand 16 mg comprimate	UK/H/0197/004	6233/2014/11	ASTRAZENECA AB	RO
Atacand 16 mg comprimate	UK/H/0197/004	6233/2014/03	ASTRAZENECA AB	RO
Atacand 16 mg comprimate	UK/H/0197/004	6233/2014/01	ASTRAZENECA AB	RO
Atacand 16 mg comprimate	UK/H/0197/004	6233/2014/06	ASTRAZENECA AB	RO
Atacand 16 mg comprimate	UK/H/0197/004	6233/2014/09	ASTRAZENECA AB	RO
Atacand 16 mg comprimate	UK/H/0197/004	6233/2014/18	ASTRAZENECA AB	RO
Atacand 16 mg	UK/H/0197/004	6233/2014/07	ASTRAZENECA AB	RO

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
comprimate				
Atacand 16 mg comprimate	UK/H/0197/004	6233/2014/17	ASTRAZENECA AB	RO
Atacand 16 mg comprimate	UK/H/0197/004	6233/2014/12	ASTRAZENECA AB	RO
Atacand 16 mg comprimate	UK/H/0197/004	6233/2014/08	ASTRAZENECA AB	RO
Atacand 16 mg comprimidos	UK/H/0197/004	61.894	ASTRAZENECA FARMACÉUTICA SPAIN, S.A.	ES
Atacand 16 mg comprimidos	UK/H/0197/004	2697282	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Atacand 16 mg comprimidos	UK/H/0197/004	3198785	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Atacand 16 mg comprimidos	UK/H/0197/004	2697787	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Atacand 16 mg comprimidos	UK/H/0197/004	2697183	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Atacand 16 mg comprimidos	UK/H/0197/004	2696987	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Atacand 16 mg comprimidos	UK/H/0197/004	2697589	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Atacand 16 mg comprimidos	UK/H/0197/004	2697985	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Atacand 16 mg comprimidos	UK/H/0197/004	2697381	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Atacand 16 mg comprimidos	UK/H/0197/004	2697480	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Atacand 16 mg comprimidos	UK/H/0197/004	2697084	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Atacand 16 mg comprimidos	UK/H/0197/004	2696888	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Atacand 16 mg comprimidos	UK/H/0197/004	2697688	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Atacand 16 mg	UK/H/0197/004	2697886	ASTRAZENECA PRODUTOS	PT

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
comprimidos			FARMACEUTICOS LDA	
ATACAND 16 mg tablete	not available	UP/I-530-09/12-02/216	ASTRAZENECA D.O.O.	HR
Atacand 16 mg tablete	UK/H/0197/004	H/00/00228/037	ASTRAZENECA UK LIMITED	SI
Atacand 16 mg tabletes	UK/H/0197/004	99-0119	ASTRAZENECA AB	LV
Atacand 16 mg tabletės	UK/H/0197/004	LT/1/99/0689/057	ASTRAZENECA AB	LT
Atacand 16 mg tabletės	UK/H/0197/004	LT/1/99/0689/015	ASTRAZENECA AB	LT
Atacand 16 mg tabletės	UK/H/0197/004	LT/1/99/0689/054	ASTRAZENECA AB	LT
Atacand 16 mg tabletės	UK/H/0197/004	LT/1/99/0689/018	ASTRAZENECA AB	LT
Atacand 16 mg tabletės	UK/H/0197/004	LT/1/99/0689/019	ASTRAZENECA AB	LT
Atacand 16 mg tabletės	UK/H/0197/004	LT/1/99/0689/003	ASTRAZENECA AB	LT
Atacand 16 mg tabletės	UK/H/0197/004	LT/1/99/0689/051	ASTRAZENECA AB	LT
Atacand 16 mg tabletės	UK/H/0197/004	LT/1/99/0689/049	ASTRAZENECA AB	LT
Atacand 16 mg tabletės	UK/H/0197/004	LT/1/99/0689/056	ASTRAZENECA AB	LT
Atacand 16 mg tabletės	UK/H/0197/004	LT/1/99/0689/016	ASTRAZENECA AB	LT
Atacand 16 mg tabletės	UK/H/0197/004	LT/1/99/0689/055	ASTRAZENECA AB	LT
Atacand 16 mg tabletės	UK/H/0197/004	LT/1/99/0689/058	ASTRAZENECA AB	LT
Atacand 16 mg tabletės	UK/H/0197/004	LT/1/99/0689/059	ASTRAZENECA AB	LT
Atacand 16 mg tabletės	UK/H/0197/004	LT/1/99/0689/060	ASTRAZENECA AB	LT
Atacand 16 mg tabletės	UK/H/0197/004	LT/1/99/0689/053	ASTRAZENECA AB	LT
Atacand 16 mg tabletės	UK/H/0197/004	LT/1/99/0689/075	ASTRAZENECA AB	LT
Atacand 16 mg tabletės	UK/H/0197/004	LT/1/99/0689/052	ASTRAZENECA AB	LT
Atacand 16 mg tabletės	UK/H/0197/004	LT/1/99/0689/017	ASTRAZENECA AB	LT
Atacand 16 mg tabletės	UK/H/0197/004	LT/1/99/0689/050	ASTRAZENECA AB	LT
ATACAND 16 MG TABLETS	UK/H/0197/004	17748	ASTRAZENECA AB	CY
Atacand 16 mg tablets	UK/H/0197/004	970/30/4	ASTRAZENECA UK LIMITED	IE
Atacand 16 mg tablets	UK/H/0197/004	MA046/00602	ASTRAZENECA AB	MT
Atacand 16 mg tablets	UK/H/0197/004	PL 17901/0099	ASTRAZENECA UK LIMITED	UK
Atacand 16 mg tablets.	UK/H/0197/004	PL 15661/0008	ASTRAZENECA UK LIMITED	UK
Atacand 16 mg tabletter	UK/H/0197/004	18974	ASTRAZENECA A/S	DK
Atacand 16 mg tabletter	UK/H/0197/004	96-2907	ASTRAZENECA AS	NO
Atacand 16 mg tabletter	UK/H/0197/004	13727	ASTRAZENECA AB	SE
Atacand 16 mg tabletti	UK/H/0197/004	12920	ASTRAZENECA OY	FI

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
Atacand 16 mg tablety	UK/H/0197/004	58/0153/99-S	ASTRAZENECA AB	SK
Atacand 16 mg töflur	UK/H/0197/004	970005	ASTRAZENECA A/S	IS
ATACAND 16 mg, comprimé sécable	UK/H/0197/004	NL 22850	ASTRAZENECA S.A.S.	FR
Atacand 16 mg, comprimés	UK/H/0197/004	BE190531	ASTRAZENECA S.A. / N.V.	BE
Atacand 16 mg, comprimés	UK/H/0197/004	BE190592	ASTRAZENECA S.A. / N.V.	BE
Atacand 16 mg, comprimés	UK/H/0197/004	2010010676	ASTRAZENECA S.A. / N.V.	LU
Atacand 16, tabletten 16 mg	UK/H/0197/004	RVG 21706	ASTRAZENECA BV	NL
Atacand 32 mg – Tabletten	UK/H/0197/005	1-25535	ASTRAZENECA OSTERREICH GMBH	AT
Atacand 32 mg comprimidos	UK/H/0197/005	66.300	ASTRAZENECA FARMACÉUTICA SPAIN, S.A.	ES
Atacand 32 mg comprimidos	UK/H/0197/005	5177589	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Atacand 32 mg comprimidos	UK/H/0197/005	5177480	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
ATACAND 32 mg tablete	not available	UP/I-530-09/12-02/217	ASTRAZENECA D.O.O.	HR
Atacand 32 mg tabletės	UK/H/0197/005	LT/1/99/0689/072	ASTRAZENECA AB	LT
Atacand 32 mg tabletės	UK/H/0197/005	LT/1/99/0689/071	ASTRAZENECA AB	LT
Atacand 32 mg tabletės	UK/H/0197/005	LT/1/99/0689/061	ASTRAZENECA AB	LT
Atacand 32 mg tabletės	UK/H/0197/005	LT/1/99/0689/069	ASTRAZENECA AB	LT
Atacand 32 mg tabletės	UK/H/0197/005	LT/1/99/0689/021	ASTRAZENECA AB	LT
Atacand 32 mg tabletės	UK/H/0197/005	LT/1/99/0689/004	ASTRAZENECA AB	LT
Atacand 32 mg tabletės	UK/H/0197/005	LT/1/99/0689/063	ASTRAZENECA AB	LT
Atacand 32 mg tabletės	UK/H/0197/005	LT/1/99/0689/067	ASTRAZENECA AB	LT
Atacand 32 mg tabletės	UK/H/0197/005	LT/1/99/0689/023	ASTRAZENECA AB	LT
Atacand 32 mg tabletės	UK/H/0197/005	LT/1/99/0689/068	ASTRAZENECA AB	LT
Atacand 32 mg tabletės	UK/H/0197/005	LT/1/99/0689/064	ASTRAZENECA AB	LT
Atacand 32 mg tabletės	UK/H/0197/005	LT/1/99/0689/070	ASTRAZENECA AB	LT
Atacand 32 mg tabletės	UK/H/0197/005	LT/1/99/0689/066	ASTRAZENECA AB	LT

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
Atacand 32 mg tabletės	UK/H/0197/005	LT/1/99/0689/020	ASTRAZENECA AB	LT
Atacand 32 mg tabletės	UK/H/0197/005	LT/1/99/0689/024	ASTRAZENECA AB	LT
Atacand 32 mg tabletės	UK/H/0197/005	LT/1/99/0689/065	ASTRAZENECA AB	LT
Atacand 32 mg tabletės	UK/H/0197/005	LT/1/99/0689/076	ASTRAZENECA AB	LT
Atacand 32 mg tabletės	UK/H/0197/005	LT/1/99/0689/062	ASTRAZENECA AB	LT
Atacand 32 mg tabletės	UK/H/0197/005	LT/1/99/0689/022	ASTRAZENECA AB	LT
Atacand 32 mg tablets	UK/H/0197/005	20090	ASTRAZENECA AB	CY
Atacand 32 mg tablets	UK/H/0197/005	PL 17901/0210	ASTRAZENECA UK LIMITED	UK
Atacand 32 mg tabletter	UK/H/0197/005	36493	ASTRAZENECA A/S	DK
Atacand 32 mg tabletter	UK/H/0197/005	04-2641	ASTRAZENECA AS	NO
Atacand 32 mg tabletter	UK/H/0197/005	20864	ASTRAZENECA AB	SE
Atacand 32 mg tabletti	UK/H/0197/005	19355	ASTRAZENECA OY	FI
Atacand 32 mg tablety	UK/H/0197/005	58/0018/05-S	ASTRAZENECA AB	SK
Atacand 32 mg töflur	UK/H/0197/005	IS/1/04/134/01	ASTRAZENECA A/S	IS
Atacand 32 mg, comprimate	UK/H/0197/005	6234/2014/17	ASTRAZENECA AB	RO
Atacand 32 mg, comprimate	UK/H/0197/005	6234/2014/09	ASTRAZENECA AB	RO
Atacand 32 mg, comprimate	UK/H/0197/005	6234/2014/10	ASTRAZENECA AB	RO
Atacand 32 mg, comprimate	UK/H/0197/005	6234/2014/11	ASTRAZENECA AB	RO
Atacand 32 mg, comprimate	UK/H/0197/005	6234/2014/01	ASTRAZENECA AB	RO
Atacand 32 mg, comprimate	UK/H/0197/005	6234/2014/16	ASTRAZENECA AB	RO
Atacand 32 mg, comprimate	UK/H/0197/005	6234/2014/03	ASTRAZENECA AB	RO
Atacand 32 mg, comprimate	UK/H/0197/005	6234/2014/05	ASTRAZENECA AB	RO
Atacand 32 mg, comprimate	UK/H/0197/005	6234/2014/07	ASTRAZENECA AB	RO
Atacand 32 mg, comprimate	UK/H/0197/005	6234/2014/15	ASTRAZENECA AB	RO

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
Atacand 32 mg, comprimate	UK/H/0197/005	6234/2014/18	ASTRAZENECA AB	RO
Atacand 32 mg, comprimate	UK/H/0197/005	6234/2014/12	ASTRAZENECA AB	RO
Atacand 32 mg, comprimate	UK/H/0197/005	6234/2014/04	ASTRAZENECA AB	RO
Atacand 32 mg, comprimate	UK/H/0197/005	6234/2014/06	ASTRAZENECA AB	RO
Atacand 32 mg, comprimate	UK/H/0197/005	6234/2014/08	ASTRAZENECA AB	RO
Atacand 32 mg, comprimate	UK/H/0197/005	6234/2014/13	ASTRAZENECA AB	RO
Atacand 32 mg, comprimate	UK/H/0197/005	6234/2014/02	ASTRAZENECA AB	RO
Atacand 32 mg, comprimate	UK/H/0197/005	6234/2014/14	ASTRAZENECA AB	RO
ATACAND 32 mg, comprimé sécable	UK/H/0197/005	NL 30321	ASTRAZENECA S.A.S.	FR
Atacand 32 mg, comprimés	UK/H/0197/005	BE267924	ASTRAZENECA S.A. / N.V.	BE
Atacand 32 mg, comprimés	UK/H/0197/005	BE267906	ASTRAZENECA S.A. / N.V.	BE
Atacand 32 mg, comprimés	UK/H/0197/005	0173/10010677	ASTRAZENECA S.A. / N.V.	LU
Atacand 32, tabletten 32 mg	UK/H/0197/005	RVG 30755	ASTRAZENECA BV	NL
Atacand 4 mg – Tabletten	UK/H/0197/002	1-22287	ASTRAZENECA OSTERREICH GMBH	AT
Atacand 4 mg comprimidos	UK/H/0197/002	61.892	ASTRAZENECA FARMACÉUTICA SPAIN, S.A.	ES
Atacand 4 mg comprimidos	UK/H/0197/002	2694883	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Atacand 4 mg comprimidos	UK/H/0197/002	2695385	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT

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
Atacand 4 mg comprimidos	UK/H/0197/002	2694784	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Atacand 4 mg comprimidos	UK/H/0197/002	2695286	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Atacand 4 mg comprimidos	UK/H/0197/002	2694982	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Atacand 4 mg comprimidos	UK/H/0197/002	2695088	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Atacand 4 mg comprimidos	UK/H/0197/002	2695187	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Atacand 4 mg comprimidos	UK/H/0197/002	2694685	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Atacand 4 mg comprimidos	UK/H/0197/002	2694586	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Atacand 4 mg comprimidos	UK/H/0197/002	3198587	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Atacand 4 mg comprimidos	UK/H/0197/002	2694487	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Atacand 4 mg comprimidos	UK/H/0197/002	2695484	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Atacand 4 mg comprimidos	UK/H/0197/002	2695583	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
ATACAND 4 mg tablete	not available	UP/I-530-09/12-02/214	ASTRAZENECA D.O.O.	HR
Atacand 4 mg tablete	UK/H/0197/002	H/00/00228/001	ASTRAZENECA UK LIMITED	SI
Atacand 4 mg tabletės	UK/H/0197/002	LT/1/99/0689/006	ASTRAZENECA AB	LT
Atacand 4 mg tabletės	UK/H/0197/002	LT/1/99/0689/031	ASTRAZENECA AB	LT
Atacand 4 mg tabletės	UK/H/0197/002	LT/1/99/0689/008	ASTRAZENECA AB	LT
Atacand 4 mg tabletės	UK/H/0197/002	LT/1/99/0689/034	ASTRAZENECA AB	LT
Atacand 4 mg tabletės	UK/H/0197/002	LT/1/99/0689/001	ASTRAZENECA AB	LT
Atacand 4 mg tabletės	UK/H/0197/002	LT/1/99/0689/033	ASTRAZENECA AB	LT
Atacand 4 mg tabletės	UK/H/0197/002	LT/1/99/0689/036	ASTRAZENECA AB	LT
Atacand 4 mg tabletės	UK/H/0197/002	LT/1/99/0689/027	ASTRAZENECA AB	LT
Atacand 4 mg tabletės	UK/H/0197/002	LT/1/99/0689/035	ASTRAZENECA AB	LT
Atacand 4 mg tabletės	UK/H/0197/002	LT/1/99/0689/030	ASTRAZENECA AB	LT

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
Atacand 4 mg tabletės	UK/H/0197/002	LT/1/99/0689/026	ASTRAZENECA AB	LT
Atacand 4 mg tabletės	UK/H/0197/002	LT/1/99/0689/007	ASTRAZENECA AB	LT
Atacand 4 mg tabletės	UK/H/0197/002	LT/1/99/0689/005	ASTRAZENECA AB	LT
Atacand 4 mg tabletės	UK/H/0197/002	LT/1/99/0689/025	ASTRAZENECA AB	LT
Atacand 4 mg tabletės	UK/H/0197/002	LT/1/99/0689/029	ASTRAZENECA AB	LT
Atacand 4 mg tabletės	UK/H/0197/002	LT/1/99/0689/032	ASTRAZENECA AB	LT
Atacand 4 mg tabletės	UK/H/0197/002	LT/1/99/0689/009	ASTRAZENECA AB	LT
Atacand 4 mg tabletės	UK/H/0197/002	LT/1/99/0689/073	ASTRAZENECA AB	LT
Atacand 4 mg tabletės	UK/H/0197/002	LT/1/99/0689/028	ASTRAZENECA AB	LT
Atacand 4 mg tablets	UK/H/0197/002	17746	ASTRAZENECA AB	CY
Atacand 4 mg tablets	UK/H/0197/002	970/30/2	ASTRAZENECA UK LIMITED	IE
Atacand 4 mg tablets	UK/H/0197/002	PL 17901/0101	ASTRAZENECA UK LIMITED	UK
Atacand 4 mg tablets.	UK/H/0197/002	PL 00017/0383	ASTRAZENECA UK LIMITED	UK
Atacand 4 mg tabletter	UK/H/0197/002	18972	ASTRAZENECA A/S	DK
Atacand 4 mg tabletter	UK/H/0197/002	96-2905	ASTRAZENECA AS	NO
Atacand 4 mg tabletter	UK/H/0197/002	13725	ASTRAZENECA AB	SE
Atacand 4 mg tabletti	UK/H/0197/002	12918	ASTRAZENECA OY	FI
Atacand 4 mg töflur	UK/H/0197/002	970003	ASTRAZENECA A/S	IS
ATACAND 4 mg, comprimé sécable	UK/H/0197/002	NL 22848	ASTRAZENECA S.A.S.	FR
Atacand 4 mg, comprimés	UK/H/0197/002	BE190574	ASTRAZENECA S.A. / N.V.	BE
Atacand 4 mg, comprimés	UK/H/0197/002	BE190644	ASTRAZENECA S.A. / N.V.	BE
Atacand 4 mg, comprimés	UK/H/0197/002	2010010673	ASTRAZENECA S.A. / N.V.	LU
Atacand 4, tabletten 4 mg	UK/H/0197/002	RVG 21704	ASTRAZENECA BV	NL
Atacand 8 mg – Tabletten	UK/H/0197/003	1-22288	ASTRAZENECA OSTERREICH GMBH	AT
Atacand 8 mg comprimate	UK/H/0197/003	6232/2014/18	ASTRAZENECA AB	RO
Atacand 8 mg comprimate	UK/H/0197/003	6232/2014/17	ASTRAZENECA AB	RO

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
Atacand 8 mg comprimate	UK/H/0197/003	6232/2014/11	ASTRAZENECA AB	RO
Atacand 8 mg comprimate	UK/H/0197/003	6232/2014/10	ASTRAZENECA AB	RO
Atacand 8 mg comprimate	UK/H/0197/003	6232/2014/01	ASTRAZENECA AB	RO
Atacand 8 mg comprimate	UK/H/0197/003	6232/2014/16	ASTRAZENECA AB	RO
Atacand 8 mg comprimate	UK/H/0197/003	6232/2014/05	ASTRAZENECA AB	RO
Atacand 8 mg comprimate	UK/H/0197/003	6232/2014/13	ASTRAZENECA AB	RO
Atacand 8 mg comprimate	UK/H/0197/003	6232/2014/09	ASTRAZENECA AB	RO
Atacand 8 mg comprimate	UK/H/0197/003	6232/2014/06	ASTRAZENECA AB	RO
Atacand 8 mg comprimate	UK/H/0197/003	6232/2014/02	ASTRAZENECA AB	RO
Atacand 8 mg comprimate	UK/H/0197/003	6232/2014/15	ASTRAZENECA AB	RO
Atacand 8 mg comprimate	UK/H/0197/003	6232/2014/08	ASTRAZENECA AB	RO
Atacand 8 mg comprimate	UK/H/0197/003	6232/2014/12	ASTRAZENECA AB	RO
Atacand 8 mg comprimate	UK/H/0197/003	6232/2014/03	ASTRAZENECA AB	RO
Atacand 8 mg comprimate	UK/H/0197/003	6232/2014/07	ASTRAZENECA AB	RO
Atacand 8 mg comprimate	UK/H/0197/003	6232/2014/04	ASTRAZENECA AB	RO
Atacand 8 mg comprimate	UK/H/0197/003	6232/2014/14	ASTRAZENECA AB	RO
Atacand 8 mg comprimidos	UK/H/0197/003	61.893	ASTRAZENECA FARMACÉUTICA SPAIN, S.A.	ES

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
Atacand 8 mg comprimidos	UK/H/0197/003	3198686	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Atacand 8 mg comprimidos	UK/H/0197/003	2695989	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Atacand 8 mg comprimidos	UK/H/0197/003	2695781	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Atacand 8 mg comprimidos	UK/H/0197/003	2695880	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Atacand 8 mg comprimidos	UK/H/0197/003	2696383	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Atacand 8 mg comprimidos	UK/H/0197/003	2695682	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Atacand 8 mg comprimidos	UK/H/0197/003	2696284	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Atacand 8 mg comprimidos	UK/H/0197/003	2696482	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Atacand 8 mg comprimidos	UK/H/0197/003	2696185	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Atacand 8 mg comprimidos	UK/H/0197/003	2696086	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Atacand 8 mg comprimidos	UK/H/0197/003	2696581	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Atacand 8 mg comprimidos	UK/H/0197/003	2696680	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Atacand 8 mg comprimidos	UK/H/0197/003	2696789	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
ATACAND 8 mg tablete	not available	UP/I-530-09/12-02/215	ASTRAZENECA D.O.O.	HR
Atacand 8 mg tablete	UK/H/0197/003	H/00/00228/019	ASTRAZENECA UK LIMITED	SI
Atacand 8 mg tabletes	UK/H/0197/003	99-0118	ASTRAZENECA AB	LV
Atacand 8 mg tabletės	UK/H/0197/003	LT/1/99/0689/011	ASTRAZENECA AB	LT
Atacand 8 mg tabletės	UK/H/0197/003	LT/1/99/0689/046	ASTRAZENECA AB	LT
Atacand 8 mg tabletės	UK/H/0197/003	LT/1/99/0689/002	ASTRAZENECA AB	LT
Atacand 8 mg tabletės	UK/H/0197/003	LT/1/99/0689/039	ASTRAZENECA AB	LT
Atacand 8 mg tabletės	UK/H/0197/003	LT/1/99/0689/048	ASTRAZENECA AB	LT

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
Atacand 8 mg tabletės	UK/H/0197/003	LT/1/99/0689/047	ASTRAZENECA AB	LT
Atacand 8 mg tabletės	UK/H/0197/003	LT/1/99/0689/045	ASTRAZENECA AB	LT
Atacand 8 mg tabletės	UK/H/0197/003	LT/1/99/0689/041	ASTRAZENECA AB	LT
Atacand 8 mg tabletės	UK/H/0197/003	LT/1/99/0689/043	ASTRAZENECA AB	LT
Atacand 8 mg tabletės	UK/H/0197/003	LT/1/99/0689/074	ASTRAZENECA AB	LT
Atacand 8 mg tabletės	UK/H/0197/003	LT/1/99/0689/037	ASTRAZENECA AB	LT
Atacand 8 mg tabletės	UK/H/0197/003	LT/1/99/0689/038	ASTRAZENECA AB	LT
Atacand 8 mg tabletės	UK/H/0197/003	LT/1/99/0689/042	ASTRAZENECA AB	LT
Atacand 8 mg tabletės	UK/H/0197/003	LT/1/99/0689/044	ASTRAZENECA AB	LT
Atacand 8 mg tabletės	UK/H/0197/003	LT/1/99/0689/013	ASTRAZENECA AB	LT
Atacand 8 mg tabletės	UK/H/0197/003	LT/1/99/0689/010	ASTRAZENECA AB	LT
Atacand 8 mg tabletės	UK/H/0197/003	LT/1/99/0689/014	ASTRAZENECA AB	LT
Atacand 8 mg tabletės	UK/H/0197/003	LT/1/99/0689/040	ASTRAZENECA AB	LT
Atacand 8 mg tabletės	UK/H/0197/003	LT/1/99/0689/012	ASTRAZENECA AB	LT
Atacand 8 mg tablets	UK/H/0197/003	17747	ASTRAZENECA AB	CY
Atacand 8 mg tablets	UK/H/0197/003	970/30/3	ASTRAZENECA UK LIMITED	IE
Atacand 8 mg tablets	UK/H/0197/003	MA046/00601	ASTRAZENECA AB	MT
Atacand 8 mg tablets	UK/H/0197/003	PL 17901/0102	ASTRAZENECA UK LIMITED	UK
Atacand 8 mg tabletter	UK/H/0197/003	18973	ASTRAZENECA A/S	DK
Atacand 8 mg tabletter	UK/H/0197/003	96-2906	ASTRAZENECA AS	NO
Atacand 8 mg tabletter	UK/H/0197/003	13726	ASTRAZENECA AB	SE
Atacand 8 mg tabletti	UK/H/0197/003	12919	ASTRAZENECA OY	FI
Atacand 8 mg tablety	UK/H/0197/003	58/0152/99-S	ASTRAZENECA AB	SK
Atacand 8 mg töflur	UK/H/0197/003	970004	ASTRAZENECA A/S	IS
ATACAND 8 mg, comprimé sécable	UK/H/0197/003	NL22849	ASTRAZENECA S.A.S.	FR
Atacand 8 mg, comprimés	UK/H/0197/003	BE190556	ASTRAZENECA S.A. / N.V.	BE
Atacand 8 mg, comprimés	UK/H/0197/003	BE190617	ASTRAZENECA S.A. / N.V.	BE
Atacand 8 mg, comprimés	UK/H/0197/003	2010010675	ASTRAZENECA S.A. / N.V.	LU
Atacand 8, tabletten 8 mg	UK/H/0197/003	RVG 21705	ASTRAZENECA BV	NL

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
Atacand Plus 16 mg/12,5 mg comprimate	SE/H/0162/002	3853/2011/01	ASTRAZENECA AB	RO
Atacand Plus 16 mg/12,5 mg comprimate	SE/H/0162/002	3853/2011/02	ASTRAZENECA AB	RO
Atacand Plus 16 mg/12,5 mg comprimate	SE/H/0162/002	3853/2011/06	ASTRAZENECA AB	RO
Atacand Plus 16 mg/12,5 mg comprimate	SE/H/0162/002	3853/2011/12	ASTRAZENECA AB	RO
Atacand Plus 16 mg/12,5 mg comprimate	SE/H/0162/002	3853/2011/05	ASTRAZENECA AB	RO
Atacand Plus 16 mg/12,5 mg comprimate	SE/H/0162/002	3853/2011/10	ASTRAZENECA AB	RO
Atacand Plus 16 mg/12,5 mg comprimate	SE/H/0162/002	3853/2011/09	ASTRAZENECA AB	RO
Atacand Plus 16 mg/12,5 mg comprimate	SE/H/0162/002	3853/2011/08	ASTRAZENECA AB	RO
Atacand Plus 16 mg/12,5 mg comprimate	SE/H/0162/002	3853/2011/13	ASTRAZENECA AB	RO
Atacand Plus 16 mg/12,5 mg comprimate	SE/H/0162/002	3853/2011/03	ASTRAZENECA AB	RO
Atacand Plus 16 mg/12,5 mg comprimate	SE/H/0162/002	3853/2011/04	ASTRAZENECA AB	RO
Atacand Plus 16 mg/12,5 mg comprimate	SE/H/0162/002	3853/2011/07	ASTRAZENECA AB	RO
Atacand Plus 16 mg/12,5 mg comprimate	SE/H/0162/002	3853/2011/11	ASTRAZENECA AB	RO
Atacand Plus 16 mg/12,5 mg comprimate	SE/H/0162/002	3853/2011/15	ASTRAZENECA AB	RO
Atacand Plus 16 mg/12,5 mg comprimate	SE/H/0162/002	3853/2011/17	ASTRAZENECA AB	RO
Atacand Plus 16 mg/12,5 mg comprimate	SE/H/0162/002	3853/2011/18	ASTRAZENECA AB	RO
Atacand Plus 16 mg/12,5 mg comprimate	SE/H/0162/002	3853/2011/14	ASTRAZENECA AB	RO

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
Atacand Plus 16 mg/12,5 mg comprimate	SE/H/0162/002	3853/2011/16	ASTRAZENECA AB	RO
Atacand Plus 16 mg/12,5 mg comprimidos	SE/H/0162/002	63.209	ASTRAZENECA FARMACÉUTICA SPAIN, S.A.	ES
ATACAND Plus 16 mg/12,5 mg tablete	not available	UP/I-530-09/12-02/218	ASTRAZENECA D.O.O.	HR
Atacand Plus 16 mg/12,5 mg tablete	SE/H/0162/002	H/04/00229/001	ASTRAZENECA UK LIMITED	SI
Atacand Plus 16 mg/12,5 mg tabletes	SE/H/0162/002	11-0413	ASTRAZENECA AB	LV
Atacand Plus 16 mg/12,5 mg Tabletten	SE/H/0162/002	1-23588	ASTRAZENECA OSTERREICH GMBH	AT
Atacand Plus 16 mg/12,5 mg tabletten	SE/H/0162/002	BE213071	ASTRAZENECA S.A. / N.V.	BE
Atacand Plus 16 mg/12,5 mg tabletten	SE/H/0162/002	BE213062	ASTRAZENECA S.A. / N.V.	BE
Atacand Plus 16 mg/12,5 mg tabletter	SE/H/0162/002	99-2649	ASTRAZENECA AS	NO
Atacand Plus 16 mg/12,5 mg tabletter	SE/H/0162/002	15487	ASTRAZENECA AB	SE
Atacand Plus 16 mg/12,5 mg tabletti	SE/H/0162/002	15139	ASTRAZENECA OY	FI
Atacand Plus 16 mg/12,5 mg töflur	SE/H/0162/002	990122	ASTRAZENECA A/S	IS
Atacand Plus 16 mg/12,5 mg, comprimés	SE/H/0162/002	BE213062	ASTRAZENECA S.A. / N.V.	BE
Atacand Plus 16 mg/12,5 mg, comprimés	SE/H/0162/002	BE213071	ASTRAZENECA S.A. / N.V.	BE
Atacand Plus 16 mg/12,5 mg, comprimés	SE/H/0162/002	2005038733	ASTRAZENECA S.A. / N.V.	LU
Atacand Plus 16/12,5 mg tablety	SE/H/0162/002	58/0258/01-S	ASTRAZENECA AB	SK
Atacand Plus 16/12,5, tabletten 16/12,5 mg	SE/H/0162/002	RVG 24995	ASTRAZENECA BV	NL

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
Atacand Plus 16/12.5 mg tablets	SE/H/0162/002	19464	ASTRAZENECA AB	CY
Atacand Plus 16/12.5 mg tablets	SE/H/0162/002	970/31/2	ASTRAZENECA UK LIMITED	IE
Atacand Plus 16/12.5 mg tablets	SE/H/0162/002	MA046/01701	ASTRAZENECA AB	MT
Atacand Plus 32 mg/12,5 mg comprimidos	SE/H/0162/003	71.062	ASTRAZENECA FARMACÉUTICA SPAIN, S.A.	ES
Atacand Plus 32 mg/12,5 mg Tabletten	SE/H/0162/003	1-28289	ASTRAZENECA OSTERREICH GMBH	AT
Atacand Plus 32 mg/12,5 mg tabletten	SE/H/0162/003	BE345055	ASTRAZENECA S.A. / N.V.	BE
Atacand Plus 32 mg/12,5 mg tabletten	SE/H/0162/003	BE345064	ASTRAZENECA S.A. / N.V.	BE
Atacand Plus 32 mg/12,5 mg tabletter	SE/H/0162/003	08/6166	ASTRAZENECA AS	NO
Atacand Plus 32 mg/12,5 mg tabletter	SE/H/0162/003	27695	ASTRAZENECA AB	SE
Atacand Plus 32 mg/12,5 mg tabletti	SE/H/0162/003	25243	ASTRAZENECA OY	FI
Atacand Plus 32 mg/12,5 mg tabletti	SE/H/0162/003	25244	ASTRAZENECA OY	FI
Atacand Plus 32 mg/12,5 mg tablety	SE/H/0162/003	58/0318/10-S	ASTRAZENECA AB	SK
Atacand Plus 32 mg/12,5 mg töflur	SE/H/0162/003	IS/1/09/004/01	ASTRAZENECA A/S	IS
Atacand Plus 32 mg/12,5 mg, comprimés	SE/H/0162/003	BE345064	ASTRAZENECA S.A. / N.V.	BE
Atacand Plus 32 mg/12,5 mg, comprimés	SE/H/0162/003	BE345055	ASTRAZENECA S.A. / N.V.	BE
Atacand Plus 32 mg/12,5 mg, comprimés	SE/H/0162/003	2010030001	ASTRAZENECA S.A. / N.V.	LU
Atacand Plus 32 mg/12,5 mg, comprimés	SE/H/0162/003	0173/10030001	ASTRAZENECA S.A. / N.V.	LU

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
Atacand Plus 32 mg/25 mg	SE/H/0162/004	58/0319/10-S	ASTRAZENECA AB	SK
Atacand Plus 32 mg/25 mg comprimate	SE/H/0162/004	3855/2011/12	ASTRAZENECA AB	RO
Atacand Plus 32 mg/25 mg comprimate	SE/H/0162/004	3855/2011/04	ASTRAZENECA AB	RO
Atacand Plus 32 mg/25 mg comprimate	SE/H/0162/004	3855/2011/06	ASTRAZENECA AB	RO
Atacand Plus 32 mg/25 mg comprimate	SE/H/0162/004	3855/2011/10	ASTRAZENECA AB	RO
Atacand Plus 32 mg/25 mg comprimate	SE/H/0162/004	3855/2011/09	ASTRAZENECA AB	RO
Atacand Plus 32 mg/25 mg comprimate	SE/H/0162/004	3855/2011/13	ASTRAZENECA AB	RO
Atacand Plus 32 mg/25 mg comprimate	SE/H/0162/004	3855/2011/05	ASTRAZENECA AB	RO
Atacand Plus 32 mg/25 mg comprimate	SE/H/0162/004	3855/2011/08	ASTRAZENECA AB	RO
Atacand Plus 32 mg/25 mg comprimate	SE/H/0162/004	3855/2011/01	ASTRAZENECA AB	RO
Atacand Plus 32 mg/25 mg comprimate	SE/H/0162/004	3855/2011/14	ASTRAZENECA AB	RO
Atacand Plus 32 mg/25 mg comprimate	SE/H/0162/004	3855/2011/11	ASTRAZENECA AB	RO
Atacand Plus 32 mg/25 mg comprimate	SE/H/0162/004	3855/2011/03	ASTRAZENECA AB	RO
Atacand Plus 32 mg/25 mg comprimate	SE/H/0162/004	3855/2011/07	ASTRAZENECA AB	RO
Atacand Plus 32 mg/25 mg comprimate	SE/H/0162/004	3855/2011/02	ASTRAZENECA AB	RO
Atacand Plus 32 mg/25 mg comprimate	SE/H/0162/004	3855/2011/15	ASTRAZENECA AB	RO
Atacand Plus 32 mg/25 mg comprimate	SE/H/0162/004	3855/2011/16	ASTRAZENECA AB	RO

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
Atacand Plus 32 mg/25 mg Tabletten	SE/H/0162/004	1-28290	ASTRAZENECA OSTERREICH GMBH	AT
Atacand Plus 32 mg/25 mg tabletten	SE/H/0162/004	BE345073	ASTRAZENECA S.A. / N.V.	BE
Atacand Plus 32 mg/25 mg tabletter	SE/H/0162/004	08/6167	ASTRAZENECA AS	NO
Atacand Plus 32 mg/25 mg tabletter	SE/H/0162/004	27696	ASTRAZENECA AB	SE
Atacand Plus 32 mg/25 mg tabletti	SE/H/0162/004	25244	ASTRAZENECA OY	FI
Atacand Plus 32 mg/25 mg töflur	SE/H/0162/004	IS/1/09/004/02	ASTRAZENECA A/S	IS
Atacand Plus 32 mg/25 mg, comprimés	SE/H/0162/004	BE345073	ASTRAZENECA S.A. / N.V.	BE
Atacand Plus 32 mg/25 mg, comprimés	SE/H/0162/004	2010030002	ASTRAZENECA S.A. / N.V.	LU
Atacand Plus 32 mg/25 mg, comprimés	SE/H/0162/004	0173/10030002	ASTRAZENECA S.A. / N.V.	LU
Atacand Plus 8 mg/12,5 mg tabletten	SE/H/0162/001	BE203603	ASTRAZENECA S.A. / N.V.	BE
Atacand Plus 8 mg/12,5 mg tabletten	SE/H/0162/001	BE203585	ASTRAZENECA S.A. / N.V.	BE
Atacand Plus 8 mg/12,5 mg tabletter	SE/H/0162/001	14274	ASTRAZENECA AB	SE
Atacand Plus 8 mg/12,5 mg tabletti	SE/H/0162/001	13751	ASTRAZENECA OY	FI
Atacand Plus 8 mg/12,5 mg, comprimés	SE/H/0162/001	BE203585	ASTRAZENECA S.A. / N.V.	BE
Atacand Plus 8 mg/12,5 mg, comprimés	SE/H/0162/001	BE203603	ASTRAZENECA S.A. / N.V.	BE
Atacand Plus 8 mg/12,5 mg, comprimés	SE/H/0162/001	2005038732	ASTRAZENECA S.A. / N.V.	LU
Atacand Plus 8/12,5, tabletten 8/12,5 mg	SE/H/0162/001	RVG 23317	ASTRAZENECA BV	NL

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
Atacand Plus 8/12.5 mg tablets	SE/H/0162/001	PA 970/31/1	ASTRAZENECA UK LIMITED	IE
Atacand Plus Forte 32 mg/25 mg comprimidos	SE/H/0162/004	71.068	ASTRAZENECA FARMACÉUTICA SPAIN, S.A.	ES
Atacand Plus mite 8 mg/12,5 mg Tabletten	SE/H/0162/001	1-22882	ASTRAZENECA OSTERREICH GMBH	AT
Atacand Plus Mite 8 mg/12,5 mg tabletter	SE/H/0162/001	98-2451	ASTRAZENECA AS	NO
Atacand Plus, 16/12,5 mg tabletid	SE/H/0162/002	454804	ASTRAZENECA AB	EE
Atacand Plus, 32/12,5 mg tabletid	SE/H/0162/003	674910	ASTRAZENECA AB	EE
Atacand Plus, 32/25 mg tabletid	SE/H/0162/004	675010	ASTRAZENECA AB	EE
Atacand Plus® 16/12,5 mg δισκία	SE/H/0162/002	44501/15-7-2008	ASTRAZENECA S.A	GR
Atacand Plus® 32/12,5 mg δισκία	not available	70102/07-10-2009	ASTRAZENECA S.A	GR
Atacand Plus® 32/25 mg δισκία	SE/H/0162/004	70103/07-10-2009	ASTRAZENECA S.A	GR
Atacand Plus® 8/12,5 mg Δισκία	SE/H/0162/001	44500/18-11-2008	ASTRAZENECA S.A	GR
Atacand Zid 16 mg/12,5 mg, tabletter	SE/H/0162/002	31691	ASTRAZENECA A/S	DK
Atacand Zid 32 mg/12,5 mg, tabletter	SE/H/0162/003	43645	ASTRAZENECA A/S	DK
Atacand Zid 32 mg/25 mg, tabletter	SE/H/0162/004	43646	ASTRAZENECA A/S	DK
Atacand Zid 8 mg/12,5 mg tabletter	SE/H/0162/001	30284	ASTRAZENECA A/S	DK
Atacand, 16 mg tabletid	UK/H/0197/004	207898	ASTRAZENECA AB	EE
Atacand, 16 mg tabletki	UK/H/0197/004	4301	ASTRAZENECA AB	PL
Atacand, 16 mg tabletten	UK/H/0197/004	BE190531	ASTRAZENECA S.A. / N.V.	BE
Atacand, 16 mg tabletten	UK/H/0197/004	BE190592	ASTRAZENECA S.A. / N.V.	BE

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
Atacand, 32 mg tabletid	UK/H/0197/005	511306	ASTRAZENECA AB	EE
Atacand, 32 mg tabletten	UK/H/0197/005	BE267906	ASTRAZENECA S.A. / N.V.	BE
Atacand, 32 mg tabletten	UK/H/0197/005	BE267924	ASTRAZENECA S.A. / N.V.	BE
Atacand, 4 mg tabletten	UK/H/0197/002	BE190574	ASTRAZENECA S.A. / N.V.	BE
Atacand, 4 mg tabletten	UK/H/0197/002	BE190644	ASTRAZENECA S.A. / N.V.	BE
Atacand, 8 mg tabletki	UK/H/0197/003	4300	ASTRAZENECA AB	PL
Atacand, 8 mg tabletten	UK/H/0197/003	BE190556	ASTRAZENECA S.A. / N.V.	BE
Atacand, 8 mg tabletten	UK/H/0197/003	BE190617	ASTRAZENECA S.A. / N.V.	BE
Atacand® 16 mg Tabletten	UK/H/0197/004	41261.03.00	ASTRAZENECA GMBH	DE
Atacand® 4 mg Tabletten	UK/H/0197/002	41261.01.00	ASTRAZENECA GMBH	DE
Atacand® 8 mg Tabletten	not available	41261.02.00	ASTRAZENECA GMBH	DE
Atacand® PLUS 16 mg/12,5 mg Tabletten	SE/H/0162/002	44433.01.00	ASTRAZENECA GMBH	DE
Atacand® PLUS 32 mg/12,5 mg Tabletten	not available	74088.00.00	ASTRAZENECA GMBH	DE
Atacand® PLUS 8 mg/12,5 mg Tabletten	SE/H/0162/001	44433.00.00	ASTRAZENECA GMBH	DE
Atacand® PLUS forte 32 mg/25 mg Tabletten	SE/H/0162/004	74089.00.00	ASTRAZENECA GMBH	DE
Atacand® PROTECT 32 mg Tabletten	UK/H/0197/005	60029.00.00	ASTRAZENECA GMBH	DE
Atacand® Δισκία, 16 mg	UK/H/0197/004	8925/13-06-2014	ASTRAZENECA S.A	GR
Atacand® Δισκία, 32 mg	UK/H/0197/005	8926/4-2-2013	ASTRAZENECA S.A	GR
Atacand® Δισκία, 4 mg	UK/H/0197/002	8923/4-2-2013	ASTRAZENECA S.A	GR
Atacand® Δισκία, 8 mg	UK/H/0197/003	8924/4-2-2013	ASTRAZENECA S.A	GR
BLOPRESID 16 mg/12,5 mg compresse	SE/H/0163/002	034187118	TAKEDA ITALIA S.P.A.	IT
BLOPRESID 16 mg/12,5 mg compresse	SE/H/0163/002	034187120	TAKEDA ITALIA S.P.A.	IT
BLOPRESID 16 mg/12,5 mg compresse	SE/H/0163/002	034187132	TAKEDA ITALIA S.P.A.	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
BLOPRESID 16 mg/12,5 mg compresse	SE/H/0163/002	034187144	TAKEDA ITALIA S.P.A.	IT
BLOPRESID 16 mg/12,5 mg compresse	SE/H/0163/002	034187157	TAKEDA ITALIA S.P.A.	IT
BLOPRESID 16 mg/12,5 mg compresse	SE/H/0163/002	034187169	TAKEDA ITALIA S.P.A.	IT
BLOPRESID 16 mg/12,5 mg compresse	SE/H/0163/002	034187171	TAKEDA ITALIA S.P.A.	IT
BLOPRESID 16 mg/12,5 mg compresse	SE/H/0163/002	034187183	TAKEDA ITALIA S.P.A.	IT
BLOPRESID 16 mg/12,5 mg compresse	SE/H/0163/002	034187195	TAKEDA ITALIA S.P.A.	IT
BLOPRESID 32 mg/12,5 mg compresse	SE/H/0163/003	034187207	TAKEDA ITALIA S.P.A.	IT
BLOPRESID 32 mg/12,5 mg compresse	SE/H/0163/003	034187219	TAKEDA ITALIA S.P.A.	IT
BLOPRESID 32 mg/12,5 mg compresse	SE/H/0163/003	034187221	TAKEDA ITALIA S.P.A.	IT
BLOPRESID 32 mg/12,5 mg compresse	SE/H/0163/003	034187233	TAKEDA ITALIA S.P.A.	IT
BLOPRESID 32 mg/12,5 mg compresse	SE/H/0163/003	034187245	TAKEDA ITALIA S.P.A.	IT
BLOPRESID 32 mg/12,5 mg compresse	SE/H/0163/003	034187258	TAKEDA ITALIA S.P.A.	IT
BLOPRESID 32 mg/12,5 mg compresse	SE/H/0163/003	034187260	TAKEDA ITALIA S.P.A.	IT
BLOPRESID 32 mg/12,5 mg compresse	SE/H/0163/003	034187272	TAKEDA ITALIA S.P.A.	IT
BLOPRESID 32 mg/12,5 mg compresse	SE/H/0163/003	034187284	TAKEDA ITALIA S.P.A.	IT
BLOPRESID 32 mg/25 mg compresse	SE/H/0163/004	034187296	TAKEDA ITALIA S.P.A.	IT
BLOPRESID 32 mg/25 mg compresse	SE/H/0163/004	034187308	TAKEDA ITALIA S.P.A.	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
BLOPRESID 32 mg/25 mg compresse	SE/H/0163/004	034187310	TAKEDA ITALIA S.P.A.	IT
BLOPRESID 32 mg/25 mg compresse	SE/H/0163/004	034187322	TAKEDA ITALIA S.P.A.	IT
BLOPRESID 32 mg/25 mg compresse	SE/H/0163/004	034187334	TAKEDA ITALIA S.P.A.	IT
BLOPRESID 32 mg/25 mg compresse	SE/H/0163/004	034187346	TAKEDA ITALIA S.P.A.	IT
BLOPRESID 32 mg/25 mg compresse	SE/H/0163/004	034187359	TAKEDA ITALIA S.P.A.	IT
BLOPRESID 32 mg/25 mg compresse	SE/H/0163/004	034187361	TAKEDA ITALIA S.P.A.	IT
BLOPRESID 32 mg/25 mg compresse	SE/H/0163/004	034187373	TAKEDA ITALIA S.P.A.	IT
BLOPRESID 8 mg/12,5 mg compresse	SE/H/0163/001	034187017	TAKEDA ITALIA S.P.A.	IT
BLOPRESID 8 mg/12,5 mg compresse	SE/H/0163/001	034187029	TAKEDA ITALIA S.P.A.	IT
BLOPRESID 8 mg/12,5 mg compresse	SE/H/0163/001	034187031	TAKEDA ITALIA S.P.A.	IT
BLOPRESID 8 mg/12,5 mg compresse	SE/H/0163/001	034187043	TAKEDA ITALIA S.P.A.	IT
BLOPRESID 8 mg/12,5 mg compresse	SE/H/0163/001	034187056	TAKEDA ITALIA S.P.A.	IT
BLOPRESID 8 mg/12,5 mg compresse	SE/H/0163/001	034187068	TAKEDA ITALIA S.P.A.	IT
BLOPRESID 8 mg/12,5 mg compresse	SE/H/0163/001	034187070	TAKEDA ITALIA S.P.A.	IT
BLOPRESID 8 mg/12,5 mg compresse	SE/H/0163/001	034187082	TAKEDA ITALIA S.P.A.	IT
BLOPRESID 8 mg/12,5 mg compresse	SE/H/0163/001	034187094	TAKEDA ITALIA S.P.A.	IT
BLOPRESID 8 mg/12,5 mg compresse	SE/H/0163/001	034187106	TAKEDA ITALIA S.P.A.	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
Blopress 16 mg – Tabletten	UK/H/0198/004	1-22236	TAKEDA PHARMA GES.M.B.H.	AT
Blopress 16 mg + 12,5 mg comprimidos	SE/H/0163/002	3197696	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Blopress 16 mg + 12,5 mg comprimidos	SE/H/0163/002	3197795	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Blopress 16 mg + 12,5 mg comprimidos	SE/H/0163/002	3197894	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Blopress 16 mg + 12,5 mg comprimidos	SE/H/0163/002	3197993	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Blopress 16 mg + 12,5 mg comprimidos	SE/H/0163/002	3198090	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Blopress 16 mg + 12,5 mg comprimidos	SE/H/0163/002	3198199	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Blopress 16 mg + 12,5 mg comprimidos	SE/H/0163/002	3198298	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Blopress 16 mg + 12,5 mg comprimidos	SE/H/0163/002	3198397	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Blopress 16 mg + 12,5 mg comprimidos	SE/H/0163/002	3198496	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
BLOPRESS 16 mg compresse	UK/H/0198/004	033451232	TAKEDA ITALIA S.P.A.	IT
BLOPRESS 16 mg compresse	UK/H/0198/004	033451244	TAKEDA ITALIA S.P.A.	IT
BLOPRESS 16 mg compresse	UK/H/0198/004	033451257	TAKEDA ITALIA S.P.A.	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
BLOPRESS 16 mg compresse	UK/H/0198/004	033451269	TAKEDA ITALIA S.P.A.	IT
BLOPRESS 16 mg compresse	UK/H/0198/004	033451271	TAKEDA ITALIA S.P.A.	IT
BLOPRESS 16 mg compresse	UK/H/0198/004	033451283	TAKEDA ITALIA S.P.A.	IT
BLOPRESS 16 mg compresse	UK/H/0198/004	033451295	TAKEDA ITALIA S.P.A.	IT
BLOPRESS 16 mg compresse	UK/H/0198/004	033451307	TAKEDA ITALIA S.P.A.	IT
BLOPRESS 16 mg compresse	UK/H/0198/004	033451319	TAKEDA ITALIA S.P.A.	IT
BLOPRESS 16 mg compresse	UK/H/0198/004	033451321	TAKEDA ITALIA S.P.A.	IT
Blopress 16 mg comprimidos	UK/H/0198/004	61.997	TAKEDA FARMACÉUTICA ESPAÑA S.A	ES
Blopress 16 mg Comprimidos	UK/H/0198/004	2700284	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Blopress 16 mg Comprimidos	UK/H/0198/004	2700383	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Blopress 16 mg Comprimidos	UK/H/0198/004	2700482	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Blopress 16 mg Comprimidos	UK/H/0198/004	2700581	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Blopress 16 mg Comprimidos	UK/H/0198/004	2700680	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Blopress 16 mg Comprimidos	UK/H/0198/004	2700789	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Blopress 16 mg	UK/H/0198/004	2700888	LUSOMEDICAMENTA –	PT

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
Comprimidos			SOCIEDADE TÉCNICA FARMACÊUTICA, SA	
Blopress 16 mg Comprimidos	UK/H/0198/004	2700987	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Blopress 16 mg Comprimidos	UK/H/0198/004	2701084	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Blopress 16 mg Comprimidos	UK/H/0198/004	2701183	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Blopress 16 mg Tablets	UK/H/0198/004	PA 1547/1/4	TAKEDA UK LTD	IE
Blopress 16 mg Tablets	UK/H/0198/004	PL 15475/0025	TAKEDA PHARMA A/S	UK
Blopress 16® mg Tabletten	UK/H/0198/004	41294.03.00	TAKEDA GMBH (KONSTANZ)	DE
Blopress 2 mg compresse	UK/H/0198/001	033451016	TAKEDA ITALIA S.P.A.	IT
BLOPRESS 2 mg compresse	UK/H/0198/001	033451028	TAKEDA ITALIA S.P.A.	IT
Blopress 2 mg Comprimidos	UK/H/0198/001	2698082	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Blopress 2 mg Comprimidos	UK/H/0198/001	2698181	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Blopress 2 mg Tablets	UK/H/0198/001	PA 1547/1/1	TAKEDA UK LTD	IE
Blopress 2 mg Tablets	UK/H/0198/001	PL 15475/0022	TAKEDA PHARMA A/S	UK
Blopress 32 mg – Tabletten	UK/H/0198/005	1-25719	TAKEDA PHARMA GES.M.B.H.	AT
Blopress 32 mg + 12,5 mg comprimidos	SE/H/0163/003	5193008	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Blopress 32 mg + 12,5 mg comprimidos	SE/H/0163/003	5193016	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT

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
Blopress 32 mg + 25 mg comprimidos	SE/H/0163/004	5193024	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Blopress 32 mg + 25 mg comprimidos	SE/H/0163/004	5193032	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
BLOPRESS 32 mg compresse	UK/H/0198/005	033451333	TAKEDA ITALIA S.P.A.	IT
BLOPRESS 32 mg compresse	UK/H/0198/005	033451345	TAKEDA ITALIA S.P.A.	IT
BLOPRESS 32 mg compresse	UK/H/0198/005	033451358	TAKEDA ITALIA S.P.A.	IT
BLOPRESS 32 mg compresse	UK/H/0198/005	033451360	TAKEDA ITALIA S.P.A.	IT
BLOPRESS 32 mg compresse	UK/H/0198/005	033451372	TAKEDA ITALIA S.P.A.	IT
BLOPRESS 32 mg compresse	UK/H/0198/005	033451384	TAKEDA ITALIA S.P.A.	IT
BLOPRESS 32 mg compresse	UK/H/0198/005	033451396	TAKEDA ITALIA S.P.A.	IT
BLOPRESS 32 mg compresse	UK/H/0198/005	033451408	TAKEDA ITALIA S.P.A.	IT
BLOPRESS 32 mg compresse	UK/H/0198/005	033451410	TAKEDA ITALIA S.P.A.	IT
Blopress 32 mg comprimidos	UK/H/0198/005	66.439	TAKEDA FARMACÉUTICA ESPAÑA S.A	ES
Blopress 32 mg Comprimidos	UK/H/0198/005	5854187	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Blopress 32 mg Comprimidos	UK/H/0198/005	5177688	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Blopress 32 mg Comprimidos	UK/H/0198/005	5177787	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT

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
Blopress 32 mg Tablets	UK/H/0198/005	PA 1547/1/5	TAKEDA UK LTD	IE
Blopress 32 mg Tablets	UK/H/0198/005	PL 15475/0026	TAKEDA PHARMA A/S	UK
Blopress 32 mg Tablets	UK/H/0198/005	PL 15475/0026	TAKEDA PHARMA A/S	UK
Blopress 32® mg Tabletten	UK/H/0198/005	41294.04.00	TAKEDA GMBH (KONSTANZ)	DE
Blopress 4 mg – Tabletten	UK/H/0198/002	1-22235	TAKEDA PHARMA GES.M.B.H.	AT
BLOPRESS 4 mg compresse	UK/H/0198/002	033451030	TAKEDA ITALIA S.P.A.	IT
BLOPRESS 4 mg compresse	UK/H/0198/002	033451042	TAKEDA ITALIA S.P.A.	IT
BLOPRESS 4 mg compresse	UK/H/0198/002	033451055	TAKEDA ITALIA S.P.A.	IT
BLOPRESS 4 mg compresse	UK/H/0198/002	033451067	TAKEDA ITALIA S.P.A.	IT
BLOPRESS 4 mg compresse	UK/H/0198/002	033451079	TAKEDA ITALIA S.P.A.	IT
BLOPRESS 4 mg compresse	UK/H/0198/002	033451081	TAKEDA ITALIA S.P.A.	IT
BLOPRESS 4 mg compresse	UK/H/0198/002	033451093	TAKEDA ITALIA S.P.A.	IT
BLOPRESS 4 mg compresse	UK/H/0198/002	033451105	TAKEDA ITALIA S.P.A.	IT
BLOPRESS 4 mg compresse	UK/H/0198/002	033451117	TAKEDA ITALIA S.P.A.	IT
BLOPRESS 4 mg compresse	UK/H/0198/002	033451129	TAKEDA ITALIA S.P.A.	IT
Blopress 4 mg comprimidos	UK/H/0198/002	61.995	TAKEDA FARMACÉUTICA ESPAÑA S.A	ES
Blopress 4 mg Comprimidos	UK/H/0198/002	2698280	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Blopress 4 mg Comprimidos	UK/H/0198/002	2698389	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA	PT

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
			FARMACÊUTICA, SA	
Blopress 4 mg Comprimidos	UK/H/0198/002	2698488	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Blopress 4 mg Comprimidos	UK/H/0198/002	2698587	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Blopress 4 mg Comprimidos	UK/H/0198/002	2698686	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Blopress 4 mg Comprimidos	UK/H/0198/002	2698785	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Blopress 4 mg Comprimidos	UK/H/0198/002	2698884	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Blopress 4 mg Comprimidos	UK/H/0198/002	2698983	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Blopress 4 mg Comprimidos	UK/H/0198/002	2699080	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Blopress 4 mg Comprimidos	UK/H/0198/002	2699189	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Blopress 4 mg Tablets	UK/H/0198/002	PA 1547/1/2	TAKEDA UK LTD	IE
Blopress 4 mg Tablets	UK/H/0198/002	PL 15475/0023	TAKEDA PHARMA A/S	UK
Blopress 4® mg Tabletten	UK/H/0198/002	41294.01.00	TAKEDA GMBH (KONSTANZ)	DE
Blopress 8 mg – Tabletten	UK/H/0198/003	1-22233	TAKEDA PHARMA GES.M.B.H.	AT
Blopress 8 mg + 12,5 mg comprimidos	SE/H/0163/001	2843282	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Blopress 8 mg + 12,5	SE/H/0163/001	2843381	LUSOMEDICAMENTA –	PT

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
mg comprimidos			SOCIEDADE TÉCNICA FARMACÊUTICA, SA	
Blopress 8 mg + 12,5 mg comprimidos	SE/H/0163/001	2843480	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Blopress 8 mg + 12,5 mg comprimidos	SE/H/0163/001	2843589	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Blopress 8 mg + 12,5 mg comprimidos	SE/H/0163/001	2843688	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Blopress 8 mg + 12,5 mg comprimidos	SE/H/0163/001	2843787	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Blopress 8 mg + 12,5 mg comprimidos	SE/H/0163/001	2843886	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Blopress 8 mg + 12,5 mg comprimidos	SE/H/0163/001	2843985	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Blopress 8 mg + 12,5 mg comprimidos	SE/H/0163/001	2844082	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Blopress 8 mg + 12,5 mg comprimidos	SE/H/0163/001	2844181	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
BLOPRESS 8 mg compresse	UK/H/0198/003	033451131	TAKEDA ITALIA S.P.A.	IT
BLOPRESS 8 mg compresse	UK/H/0198/003	033451143	TAKEDA ITALIA S.P.A.	IT
BLOPRESS 8 mg compresse	UK/H/0198/003	033451156	TAKEDA ITALIA S.P.A.	IT
BLOPRESS 8 mg compresse	UK/H/0198/003	033451168	TAKEDA ITALIA S.P.A.	IT
BLOPRESS 8 mg	UK/H/0198/003	033451170	TAKEDA ITALIA S.P.A.	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
compresse				
BLOPRESS 8 mg compresse	UK/H/0198/003	033451182	TAKEDA ITALIA S.P.A.	IT
BLOPRESS 8 mg compresse	UK/H/0198/003	033451194	TAKEDA ITALIA S.P.A.	IT
BLOPRESS 8 mg compresse	UK/H/0198/003	033451206	TAKEDA ITALIA S.P.A.	IT
BLOPRESS 8 mg compresse	UK/H/0198/003	033451218	TAKEDA ITALIA S.P.A.	IT
BLOPRESS 8 mg compresse	UK/H/0198/003	033451220	TAKEDA ITALIA S.P.A.	IT
Blopress 8 mg comprimidos	UK/H/0198/003	61.996	TAKEDA FARMACÉUTICA ESPAÑA S.A	ES
Blopress 8 mg Comprimidos	UK/H/0198/003	2699387	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Blopress 8 mg Comprimidos	UK/H/0198/003	2699486	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Blopress 8 mg Comprimidos	UK/H/0198/003	2699585	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Blopress 8 mg Comprimidos	UK/H/0198/003	2699684	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Blopress 8 mg Comprimidos	UK/H/0198/003	2699783	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Blopress 8 mg Comprimidos	UK/H/0198/003	2699882	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Blopress 8 mg Comprimidos	UK/H/0198/003	2699981	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Blopress 8 mg	UK/H/0198/003	2700086	LUSOMEDICAMENTA –	PT

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
Comprimidos			SOCIEDADE TÉCNICA FARMACÊUTICA, SA	
Blopress 8 mg Comprimidos	UK/H/0198/003	2700185	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Blopress 8 mg Comprimidos	UK/H/0198/003	2699288	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Blopress 8 mg Tablets	UK/H/0198/003	PA 1547/1/3	TAKEDA UK LTD	IE
Blopress 8 mg Tablets	UK/H/0198/003	PL 15475/0024	TAKEDA PHARMA A/S	UK
Blopress 8® mg Tabletten	UK/H/0198/003	41294.02.00	TAKEDA GMBH (KONSTANZ)	DE
Blopress Comp 16 mg/12,5 mg tablett	SE/H/0163/002	15479	TAKEDA PHARMA A/S	SE
Blopress Comp 32 mg/12,5 mg tablett	SE/H/0163/003	27705	TAKEDA PHARMA A/S	SE
Blopress Comp 32 mg/25 mg tablett	SE/H/0163/004	27706	TAKEDA PHARMA A/S	SE
Blopress Comp 8 mg/12,5 mg tablett	SE/H/0163/001	14283	TAKEDA PHARMA A/S	SE
Blopress Forte 32 mg /25 mg comprimidos	SE/H/0163/004	71028	TAKEDA FARMACÉUTICA ESPAÑA S.A	ES
Blopress Plus 16 mg/12,5 mg comprimidos	SE/H/0163/002	66.050	TAKEDA FARMACÉUTICA ESPAÑA S.A	ES
Blopress Plus 16 mg/12.5 mg tablets	SE/H/0163/002	PA 1547/002/002	TAKEDA UK LTD	IE
Blopress Plus 16 mg/12,5 mg – Tabletten	SE/H/0163/002	1-23584	TAKEDA PHARMA GES.M.B.H.	AT
Blopress Plus 32 mg /12,5 mg comprimidos	SE/H/0163/003	71026	TAKEDA FARMACÉUTICA ESPAÑA S.A	ES
Blopress Plus 32 mg/12,5 mg – Tabletten	SE/H/0163/003	1-28291	TAKEDA PHARMA GES.M.B.H.	AT

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
Blopress Plus 32 mg/12.5 mg tablets	SE/H/0163/003	PA 1547/2/3	TAKEDA UK LTD	IE
Blopress Plus 32 mg/25 mg – Tabletten	SE/H/0163/004	1-28292	TAKEDA PHARMA GES.M.B.H.	AT
Blopress Plus 32 mg/25 mg tablets	SE/H/0163/004	PA 1547/2/4	TAKEDA UK LTD	IE
Blopress Plus 8 mg/12.5 mg tablets	SE/H/0163/001	PA 1547/2/1	TAKEDA UK LTD	IE
Blopress Plus 8 mg/12,5 mg – Tabletten	SE/H/0163/001	1-22844	TAKEDA PHARMA GES.M.B.H.	AT
Blopress® 16 mg Plus 12,5 mg Tabletten	SE/H/0163/002	44475.01.00	TAKEDA GMBH (KONSTANZ)	DE
Blopress® 32 mg Plus 12,5 mg Tabletten	SE/H/0163/003	74307.00.00	TAKEDA GMBH (KONSTANZ)	DE
Blopress® 8 mg Plus 12,5 mg Tabletten	SE/H/0163/001	44475.00.00	TAKEDA GMBH (KONSTANZ)	DE
Blopress® forte 32 mg Plus 25 mg Tabletten	SE/H/0163/004	74308.00.00	TAKEDA GMBH (KONSTANZ)	DE
Candea HCT 8 mg/12,5 mg tablete	DE/H/1828/001	H/10/00331/001	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Candea HCT 8 mg/12,5 mg tablete	DE/H/1828/001	H/10/00331/002	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Candea HCT 8 mg/12,5 mg tablete	DE/H/1828/001	H/10/00331/003	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Candea HCT 8 mg/12,5 mg tablete	DE/H/1828/001	H/10/00331/005	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Candea HCT 8 mg/12,5 mg tablete	DE/H/1828/001	H/10/00331/006	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Candea HCT 8 mg/12,5 mg tablete	DE/H/1828/001	H/10/00331/008	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Candea HCT 8 mg/12,5 mg tablete	DE/H/1828/001	H/10/00331/009	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Candea HCT 8 mg/12,5	DE/H/1828/001	H/10/00331/010	LEK PHARMACEUTICALS	SI

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
mg tablete			D.D. LJUBLJANA	
Candea HCT 8 mg/12,5 mg tablete	DE/H/1828/001	H/10/00331/011	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Candea HCT 8 mg/12,5 mg tablete	DE/H/1828/001	H/10/00331/012	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Candea HCT 8 mg/12,5 mg tablete	DE/H/1828/001	H/10/00331/013	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Candea HCT 8 mg/12,5 mg tablete	DE/H/1828/001	H/10/00331/014	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Candea HCT 8 mg/12,5 mg tablete	DE/H/1828/001	H/10/00331/016	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Candea HCT 8 mg/12,5 mg tablete	DE/H/1828/001	H/10/00331/017	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Candea HCT 8 mg/12,5 mg tablete	DE/H/1828/001	H/10/00331/018	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Candea HCT 8 mg/12,5 mg tablete	DE/H/1828/001	H/10/00331/019	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Candea HCT 8 mg/12,5 mg tablete	DE/H/1828/001	H/10/00331/021	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Candea HCT 8 mg/12,5 mg tablete	DE/H/1828/001	H/10/00331/022	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Candea HCT 8 mg/12,5 mg tablete	DE/H/1828/001	H/10/00331/023	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Candea HCT 8 mg/12,5 mg tablete	DE/H/1828/001	H/10/00331/024	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Candea HCT 8 mg/12,5 mg tablete	DE/H/1828/001	H/10/00331/004	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Candea HCT 8 mg/12,5 mg tablete	DE/H/1828/001	H/10/00331/007	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Candea HCT 8 mg/12,5 mg tablete	DE/H/1828/001	H/10/00331/015	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Candea HCT 8 mg/12,5 mg tablete	DE/H/1828/001	H/10/00331/020	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
CANDEPRES, 32 MG,	DE/H/1097/004	16104	SANDOZ GMBH	PL

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
TABLETKI				
CANDESARTAN ACTAVIS 32 mg tabletes	DK/H/1838/004	11-0255	ACTAVIS GROUP PTC EHF.	LV
Candesartan Actavis, 4 mg tabletid	DK/H/1838/001	735911	ACTAVIS GROUP PTC EHF.	EE
Candesartan cilexetil 2 mg tablets	PT/H/0888/001	MA807/06601	AUROBINDO PHARMA (MALTA) LIMITED	MT
Candesartan cilexetil 32 mg tablets	PT/H/0888/005	MA807/06605	AUROBINDO PHARMA (MALTA) LIMITED	MT
Candesartan cilexetil 4 mg tablets	PT/H/0888/002	MA807/06602	AUROBINDO PHARMA (MALTA) LIMITED	MT
Candesartan HCT Actavis 32 mg/25 mg tabletes	DK/H/1839/004	14-0065	ACTAVIS GROUP PTC EHF.	LV
Candesartan HCT Actavis, 8 mg/12,5 mg tabletid	DK/H/1839/001	736111	ACTAVIS GROUP PTC EHF.	EE
Candesartan/Hydrochlorothiazide Orion 8 mg/12,5 mg tabletter	FI/H/0833/001	10-7705	ORION CORPORATION	NO
Candpress Comp 8 mg/12,5 mg töflur	DK/H/1839/001	IS/1/11/054/01	ACTAVIS GROUP PTC EHF.	IS
COKENZEN 16 mg/12,5 mg, comprimé sécable	SE/H/0163/002	34009 355 572 1 6	TAKEDA FRANCE S.A.S.	FR
COKENZEN 16 mg/12,5 mg, comprimé sécable	SE/H/0163/002	34009 371 417 7 2	TAKEDA FRANCE S.A.S.	FR
COKENZEN 16 mg/12,5 mg, comprimé sécable	SE/H/0163/002	34009 355 573 8 4	TAKEDA FRANCE S.A.S.	FR
COKENZEN 16 mg/12,5 mg, comprimé sécable	SE/H/0163/002	34009 371 418 3 3	TAKEDA FRANCE S.A.S.	FR
COKENZEN 16 mg/12,5 mg, comprimé sécable	SE/H/0163/002	34009 371 420 8 3	TAKEDA FRANCE S.A.S.	FR
COKENZEN 16 mg/12,5 mg, comprimé sécable	SE/H/0163/002	34009 371 421 4 4	TAKEDA FRANCE S.A.S.	FR
COKENZEN 16 mg/12,5	SE/H/0163/002	34009 562 942 9 3	TAKEDA FRANCE S.A.S.	FR

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
mg, comprimé sécable				
COKENZEN 16 mg/12,5 mg, comprimé sécable	SE/H/0163/002	34009 562 943 5 0	TAKEDA FRANCE S.A.S.	FR
COKENZEN 16 mg/12,5 mg, comprimé sécable	SE/H/0163/002	34009 562 944 1 1	TAKEDA FRANCE S.A.S.	FR
COKENZEN 16 mg/12,5 mg, comprimé sécable	SE/H/0163/002	34009 355 571 5 5	TAKEDA FRANCE S.A.S.	FR
COKENZEN 8 mg/12,5 mg, comprimé sécable	SE/H/0163/001	34009 353 932 0 3	TAKEDA FRANCE S.A.S.	FR
COKENZEN 8 mg/12,5 mg, comprimé sécable	SE/H/0163/001	34009 371 409 4 2	TAKEDA FRANCE S.A.S.	FR
COKENZEN 8 mg/12,5 mg, comprimé sécable	SE/H/0163/001	34009 353 933 7 1	TAKEDA FRANCE S.A.S.	FR
COKENZEN 8 mg/12,5 mg, comprimé sécable	SE/H/0163/001	34009 371 410 2 4	TAKEDA FRANCE S.A.S.	FR
COKENZEN 8 mg/12,5 mg, comprimé sécable	SE/H/0163/001	34009 371 411 9 2	TAKEDA FRANCE S.A.S.	FR
COKENZEN 8 mg/12,5 mg, comprimé sécable	SE/H/0163/001	34009 371 412 5 3	TAKEDA FRANCE S.A.S.	FR
COKENZEN 8 mg/12,5 mg, comprimé sécable	SE/H/0163/001	34009 562 419 4 1	TAKEDA FRANCE S.A.S.	FR
COKENZEN 8 mg/12,5 mg, comprimé sécable	SE/H/0163/001	34009 562 420 2 3	TAKEDA FRANCE S.A.S.	FR
COKENZEN 8 mg/12,5 mg, comprimé sécable	SE/H/0163/001	34009 562 421 9 1	TAKEDA FRANCE S.A.S.	FR
Hytacand 16 mg, 16 mg/12,5 mg comprimidos	SE/H/0162/002	3212685	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Hytacand 16 mg, 16 mg/12,5 mg comprimidos	SE/H/0162/002	3212388	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Hytacand 16 mg, 16 mg/12,5 mg comprimidos	SE/H/0162/002	3212289	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT

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
Hytacand 16 mg, 16 mg/12,5 mg comprimidos	SE/H/0162/002	3212784	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Hytacand 16 mg, 16 mg/12,5 mg comprimidos	SE/H/0162/002	3212081	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Hytacand 16 mg, 16 mg/12,5 mg comprimidos	SE/H/0162/002	3212586	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Hytacand 16 mg, 16 mg/12,5 mg comprimidos	SE/H/0162/002	3212487	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Hytacand 16 mg, 16 mg/12,5 mg comprimidos	SE/H/0162/002	3211984	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Hytacand 16 mg, 16 mg/12,5 mg comprimidos	SE/H/0162/002	3212180	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Hytacand 16 mg, 16 mg/12,5 mg comprimidos	SE/H/0162/002	3212883	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
HYTACAND 16 mg/12,5 mg comprimé	SE/H/0162/002	NL 24952	ASTRAZENECA S.A.S.	FR
Hytacand 32 mg/12,5 mg comprimidos	SE/H/0162/003	5187729	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA.	PT
Hytacand 32 mg/12,5 mg comprimidos	SE/H/0162/003/DC	5187729	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA.	PT
Hytacand 32 mg/12,5 mg comprimidos	SE/H/0162/003	5187711	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Hytacand 32 mg/25 mg comprimidos	SE/H/0162/004	5187745	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Hytacand 32 mg/25 mg comprimidos	SE/H/0162/004	5187737	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
HYTACAND 8 mg/12,5 mg comprimé	SE/H/0162/001	NL 23494	ASTRAZENECA S.A.S.	FR

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
Hytacand 8 mg/12,5 mg comprimidos	SE/H/0162/001	2845386	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Hytacand 8 mg/12,5 mg comprimidos	SE/H/0162/001	2844587	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Hytacand 8 mg/12,5 mg comprimidos	SE/H/0162/001	2845188	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Hytacand 8 mg/12,5 mg comprimidos	SE/H/0162/001	2844389	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Hytacand 8 mg/12,5 mg comprimidos	SE/H/0162/001	2845089	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Hytacand 8 mg/12,5 mg comprimidos	SE/H/0162/001	2844488	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Hytacand 8 mg/12,5 mg comprimidos	SE/H/0162/001	2844280	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Hytacand 8 mg/12,5 mg comprimidos	SE/H/0162/001	2845287	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Hytacand 8 mg/12,5 mg comprimidos	SE/H/0162/001	2844686	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Hytacand 8 mg/12,5 mg comprimidos	SE/H/0162/001	2844884	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Hytacand 8 mg/12,5 mg comprimidos	SE/H/0162/001	2844785	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Hytacand 8 mg/12,5 mg comprimidos	SE/H/0162/001	2844983	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Kenzen 16 mg, comprimé sécable	UK/H/0198/004	350 856-1	TAKEDA FRANCE S.A.S.	FR
Kenzen 16 mg, comprimé sécable	UK/H/0198/004	350 857-8	TAKEDA FRANCE S.A.S.	FR
Kenzen 16 mg, comprimé sécable	UK/H/0198/004	372 042-7	TAKEDA FRANCE S.A.S.	FR
Kenzen 16 mg, comprimé sécable	UK/H/0198/004	350 858-4	TAKEDA FRANCE S.A.S.	FR
Kenzen 16 mg, comprimé sécable	UK/H/0198/004	372 043-3	TAKEDA FRANCE S.A.S.	FR

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
Kenzen 16 mg, comprimé sécable	UK/H/0198/004	372 045-6	TAKEDA FRANCE S.A.S.	FR
Kenzen 16 mg, comprimé sécable	UK/H/0198/004	372 046-2	TAKEDA FRANCE S.A.S.	FR
Kenzen 16 mg, comprimé sécable	UK/H/0198/004	350 859-0	TAKEDA FRANCE S.A.S.	FR
Kenzen 16 mg, comprimé sécable	UK/H/0198/004	561 617-7	TAKEDA FRANCE S.A.S.	FR
Kenzen 16 mg, comprimé sécable	UK/H/0198/004	561 618-3	TAKEDA FRANCE S.A.S.	FR
Kenzen 32 mg, comprimé sécable	UK/H/0198/005	368 699-5	TAKEDA FRANCE S.A.S.	FR
Kenzen 32 mg, comprimé sécable	UK/H/0198/005	368 700-3	TAKEDA FRANCE S.A.S.	FR
Kenzen 32 mg, comprimé sécable	UK/H/0198/005	368 702-6	TAKEDA FRANCE S.A.S.	FR
Kenzen 32 mg, comprimé sécable	UK/H/0198/005	368 703-2	TAKEDA FRANCE S.A.S.	FR
Kenzen 32 mg, comprimé sécable	UK/H/0198/005	368 704-9	TAKEDA FRANCE S.A.S.	FR
Kenzen 32 mg, comprimé sécable	UK/H/0198/005	368 705-5	TAKEDA FRANCE S.A.S.	FR
Kenzen 32 mg, comprimé sécable	UK/H/0198/005	368 706-1	TAKEDA FRANCE S.A.S.	FR
Kenzen 32 mg, comprimé sécable	UK/H/0198/005	368 707-8	TAKEDA FRANCE S.A.S.	FR
Kenzen 32 mg, comprimé sécable	UK/H/0198/005	369 509-5	TAKEDA FRANCE S.A.S.	FR
Kenzen 32 mg, comprimé sécable	UK/H/0198/005	368 708-4	TAKEDA FRANCE S.A.S.	FR
Kenzen 32 mg, comprimé sécable	UK/H/0198/005	368 709-0	TAKEDA FRANCE S.A.S.	FR
Kenzen 32 mg, comprimé sécable	UK/H/0198/005	368 710-9	TAKEDA FRANCE S.A.S.	FR

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
Kenzen 32 mg, comprimé sécable	UK/H/0198/005	368 711-5	TAKEDA FRANCE S.A.S.	FR
Kenzen 32 mg, comprimé sécable	UK/H/0198/005	567 125-9	TAKEDA FRANCE S.A.S.	FR
Kenzen 32 mg, comprimé sécable	UK/H/0198/005	567 126-5	TAKEDA FRANCE S.A.S.	FR
Kenzen 32 mg, comprimé sécable	UK/H/0198/005	567 127-1	TAKEDA FRANCE S.A.S.	FR
Kenzen 32 mg, comprimé sécable	UK/H/0198/005	567 128-8	TAKEDA FRANCE S.A.S.	FR
Kenzen 32 mg, comprimé sécable	UK/H/0198/005	567 129-4	TAKEDA FRANCE S.A.S.	FR
Kenzen 32 mg, comprimé sécable	UK/H/0198/005	567 130-2	TAKEDA FRANCE S.A.S.	FR
Kenzen 32 mg, comprimé sécable	UK/H/0198/005	566 817-4	TAKEDA FRANCE S.A.S.	FR
Kenzen 32 mg, comprimé sécable	UK/H/0198/005	566 818-0	TAKEDA FRANCE S.A.S.	FR
Kenzen 32 mg, comprimé sécable	UK/H/0198/005	567 131-9	TAKEDA FRANCE S.A.S.	FR
Kenzen 32 mg, comprimé sécable	UK/H/0198/005	567 132-5	TAKEDA FRANCE S.A.S.	FR
Kenzen 32 mg, comprimé sécable	UK/H/0198/005	566 819-7	TAKEDA FRANCE S.A.S.	FR
Kenzen 32 mg, comprimé sécable	UK/H/0198/005	566 820-5	TAKEDA FRANCE S.A.S.	FR
Kenzen 32 mg, comprimé sécable	UK/H/0198/005	566 821-1	TAKEDA FRANCE S.A.S.	FR
Kenzen 32 mg, comprimé sécable	UK/H/0198/005	566 822-8	TAKEDA FRANCE S.A.S.	FR
Kenzen 32 mg, comprimé sécable	UK/H/0198/005	567 133-1	TAKEDA FRANCE S.A.S.	FR
Kenzen 32 mg, comprimé sécable	UK/H/0198/005	567 134-8	TAKEDA FRANCE S.A.S.	FR

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
Kenzen 32 mg, comprimé sécable	UK/H/0198/005	369 510-3	TAKEDA FRANCE S.A.S.	FR
Kenzen 4 mg, comprimé sécable	UK/H/0198/002	345 693-0	TAKEDA FRANCE S.A.S.	FR
Kenzen 4 mg, comprimé sécable	UK/H/0198/002	345 694-7	TAKEDA FRANCE S.A.S.	FR
Kenzen 4 mg, comprimé sécable	UK/H/0198/002	372 033-8	TAKEDA FRANCE S.A.S.	FR
Kenzen 4 mg, comprimé sécable	UK/H/0198/002	345 695-3	TAKEDA FRANCE S.A.S.	FR
Kenzen 4 mg, comprimé sécable	UK/H/0198/002	372 034-4	TAKEDA FRANCE S.A.S.	FR
Kenzen 4 mg, comprimé sécable	UK/H/0198/002	372 035-0	TAKEDA FRANCE S.A.S.	FR
Kenzen 4 mg, comprimé sécable	UK/H/0198/002	372 036-7	TAKEDA FRANCE S.A.S.	FR
Kenzen 4 mg, comprimé sécable	UK/H/0198/002	345 697-6	TAKEDA FRANCE S.A.S.	FR
Kenzen 4 mg, comprimé sécable	UK/H/0198/002	345 698-2	TAKEDA FRANCE S.A.S.	FR
Kenzen 4 mg, comprimé sécable	UK/H/0198/002	560 915-4	TAKEDA FRANCE S.A.S.	FR
Kenzen 8 mg, comprimé sécable	UK/H/0198/003	345 699-9	TAKEDA FRANCE S.A.S.	FR
Kenzen 8 mg, comprimé sécable	UK/H/0198/003	345 700-7	TAKEDA FRANCE S.A.S.	FR
Kenzen 8 mg, comprimé sécable	UK/H/0198/003	372 037-3	TAKEDA FRANCE S.A.S.	FR
Kenzen 8 mg, comprimé sécable	UK/H/0198/003	345 701-3	TAKEDA FRANCE S.A.S.	FR
Kenzen 8 mg, comprimé sécable	UK/H/0198/003	372 039-6	TAKEDA FRANCE S.A.S.	FR
Kenzen 8 mg, comprimé sécable	UK/H/0198/003	372 040-4	TAKEDA FRANCE S.A.S.	FR

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
Kenzen 8 mg, comprimé sécable	UK/H/0198/003	372 041-0	TAKEDA FRANCE S.A.S.	FR
Kenzen 8 mg, comprimé sécable	UK/H/0198/003	345 703-6	TAKEDA FRANCE S.A.S.	FR
Kenzen 8 mg, comprimé sécable	UK/H/0198/003	345 704-2	TAKEDA FRANCE S.A.S.	FR
Kenzen 8 mg, comprimé sécable	UK/H/0198/003	560 916-0	TAKEDA FRANCE S.A.S.	FR
Parapres 16 mg comprimidos	not available	62.008	ALMIRALL, S.A.	ES
PARAPRES 32 mg Comprimidos	not available	66.729	ALMIRALL, S.A.	ES
PARAPRES 4 mg Comprimidos	not available	62.006	ALMIRALL, S.A.	ES
PARAPRES 8 mg Comprimidos	not available	62.007	ALMIRALL, S.A.	ES
Parapres Comp 32 mg/12,5 mg tablett	SE/H/0197/002	27713	TAKEDA PHARMA A/S	SE
Parapres Comp 32 mg/25 mg tablett	SE/H/0197/003	27714	TAKEDA PHARMA A/S	SE
Parapres Comp Forte 16 mg/12,5 mg tablett	SE/H/0197/001	15797	TAKEDA PHARMA A/S	SE
Parapres Plus 16 mg/12,5 mg comprimidos	SE/H/0197/001	63.175	ALMIRALL, S.A.	ES
Parapres Plus 32 mg/12,5 mg comprimidos	SE/H/0197/002	71.021	ALMIRALL, S.A.	ES
Parapres Plus Forte 32 mg/25 mg comprimidos	SE/H/0197/003	71.022	ALMIRALL, S.A.	ES
RATACAND 16 mg compresse	UK/H/0197/004	033577368	ASTRAZENECA S.P.A.	IT
RATACAND 16 mg compresse	UK/H/0197/004	033577305	ASTRAZENECA S.P.A.	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
RATACAND 16 mg compresse	UK/H/0197/004	033577281	ASTRAZENECA S.P.A.	IT
RATACAND 16 mg compresse	UK/H/0197/004	033577356	ASTRAZENECA S.P.A.	IT
RATACAND 16 mg compresse	UK/H/0197/004	033577370	ASTRAZENECA S.P.A.	IT
RATACAND 16 mg compresse	UK/H/0197/004	033577293	ASTRAZENECA S.P.A.	IT
RATACAND 16 mg compresse	UK/H/0197/004	033577331	ASTRAZENECA S.P.A.	IT
RATACAND 16 mg compresse	UK/H/0197/004	033577317	ASTRAZENECA S.P.A.	IT
RATACAND 16 mg compresse	UK/H/0197/004	033577329	ASTRAZENECA S.P.A.	IT
RATACAND 16 mg compresse	UK/H/0197/004	033577343	ASTRAZENECA S.P.A.	IT
RATACAND 16 mg compresse	UK/H/0197/004	033577279	ASTRAZENECA S.P.A.	IT
RATACAND 16 mg compresse	UK/H/0197/004	033577382	ASTRAZENECA S.P.A.	IT
RATACAND 16 mg compresse	UK/H/0197/004	033577418	ASTRAZENECA S.P.A.	IT
RATACAND 16 mg compresse	UK/H/0197/004	033577622	ASTRAZENECA S.P.A.	IT
RATACAND 16 mg compresse	UK/H/0197/004	033577673	ASTRAZENECA S.P.A.	IT
RATACAND 16 mg compresse	UK/H/0197/004	033577610	ASTRAZENECA S.P.A.	IT
RATACAND 16 mg compresse	UK/H/0197/004	033577735	ASTRAZENECA S.P.A.	IT
RATACAND 16 mg compresse	UK/H/0197/004	033577685	ASTRAZENECA S.P.A.	IT
RATACAND 32 mg compresse	UK/H/0197/005	033577519	ASTRAZENECA S.P.A.	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
RATACAND 32 mg compresse	UK/H/0197/005	033577471	ASTRAZENECA S.P.A.	IT
RATACAND 32 mg compresse	UK/H/0197/005	033577533	ASTRAZENECA S.P.A.	IT
RATACAND 32 mg compresse	UK/H/0197/005	033577483	ASTRAZENECA S.P.A.	IT
RATACAND 32 mg compresse	UK/H/0197/005	033577545	ASTRAZENECA S.P.A.	IT
RATACAND 32 mg compresse	UK/H/0197/005	033577420	ASTRAZENECA S.P.A.	IT
RATACAND 32 mg compresse	UK/H/0197/005	033577469	ASTRAZENECA S.P.A.	IT
RATACAND 32 mg compresse	UK/H/0197/005	033577507	ASTRAZENECA S.P.A.	IT
RATACAND 32 mg compresse	UK/H/0197/005	033577457	ASTRAZENECA S.P.A.	IT
RATACAND 32 mg compresse	UK/H/0197/005	033577521	ASTRAZENECA S.P.A.	IT
RATACAND 32 mg compresse	UK/H/0197/005	033577432	ASTRAZENECA S.P.A.	IT
RATACAND 32 mg compresse	UK/H/0197/005	033577444	ASTRAZENECA S.P.A.	IT
RATACAND 32 mg compresse	UK/H/0197/005	033577495	ASTRAZENECA S.P.A.	IT
RATACAND 32 mg compresse	UK/H/0197/005	033577697	ASTRAZENECA S.P.A.	IT
RATACAND 32 mg compresse	UK/H/0197/005	033577747	ASTRAZENECA S.P.A.	IT
RATACAND 32 mg compresse	UK/H/0197/005	033577560	ASTRAZENECA S.P.A.	IT
RATACAND 32 mg compresse	UK/H/0197/005	033577709	ASTRAZENECA S.P.A.	IT
RATACAND 32 mg compresse	UK/H/0197/005	033577558	ASTRAZENECA S.P.A.	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
RATACAND 4 mg compresse	UK/H/0197/002	033577077	ASTRAZENECA S.P.A.	IT
RATACAND 4 mg compresse	UK/H/0197/002	033577394	ASTRAZENECA S.P.A.	IT
RATACAND 4 mg compresse	UK/H/0197/002	033577089	ASTRAZENECA S.P.A.	IT
RATACAND 4 mg compresse	UK/H/0197/002	033577115	ASTRAZENECA S.P.A.	IT
RATACAND 4 mg compresse	UK/H/0197/002	033577053	ASTRAZENECA S.P.A.	IT
RATACAND 4 mg compresse	UK/H/0197/002	033577127	ASTRAZENECA S.P.A.	IT
RATACAND 4 mg compresse	UK/H/0197/002	033577038	ASTRAZENECA S.P.A.	IT
RATACAND 4 mg compresse	UK/H/0197/002	033577065	ASTRAZENECA S.P.A.	IT
RATACAND 4 mg compresse	UK/H/0197/002	033577139	ASTRAZENECA S.P.A.	IT
RATACAND 4 mg compresse	UK/H/0197/002	033577141	ASTRAZENECA S.P.A.	IT
RATACAND 4 mg compresse	UK/H/0197/002	033577103	ASTRAZENECA S.P.A.	IT
RATACAND 4 mg compresse	UK/H/0197/002	033577091	ASTRAZENECA S.P.A.	IT
RATACAND 4 mg compresse	UK/H/0197/002	033577040	ASTRAZENECA S.P.A.	IT
RATACAND 4 mg compresse	UK/H/0197/002	033577646	ASTRAZENECA S.P.A.	IT
RATACAND 4 mg compresse	UK/H/0197/002	033577634	ASTRAZENECA S.P.A.	IT
RATACAND 4 mg compresse	UK/H/0197/002	033577584	ASTRAZENECA S.P.A.	IT
RATACAND 4 mg compresse	UK/H/0197/002	033577572	ASTRAZENECA S.P.A.	IT

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RATACAND 4 mg compresse	UK/H/0197/002	033577711	ASTRAZENECA S.P.A.	IT
RATACAND 8 mg compresse	UK/H/0197/003	033577216	ASTRAZENECA S.P.A.	IT
RATACAND 8 mg compresse	UK/H/0197/003	033577242	ASTRAZENECA S.P.A.	IT
RATACAND 8 mg compresse	UK/H/0197/003	033577192	ASTRAZENECA S.P.A.	IT
RATACAND 8 mg compresse	UK/H/0197/003	033577230	ASTRAZENECA S.P.A.	IT
RATACAND 8 mg compresse	UK/H/0197/003	033577180	ASTRAZENECA S.P.A.	IT
RATACAND 8 mg compresse	UK/H/0197/003	033577228	ASTRAZENECA S.P.A.	IT
RATACAND 8 mg compresse	UK/H/0197/003	033577267	ASTRAZENECA S.P.A.	IT
RATACAND 8 mg compresse	UK/H/0197/003	033577154	ASTRAZENECA S.P.A.	IT
RATACAND 8 mg compresse	UK/H/0197/003	033577406	ASTRAZENECA S.P.A.	IT
RATACAND 8 mg compresse	UK/H/0197/003	033577178	ASTRAZENECA S.P.A.	IT
RATACAND 8 mg compresse	UK/H/0197/003	033577204	ASTRAZENECA S.P.A.	IT
RATACAND 8 mg compresse	UK/H/0197/003	033577255	ASTRAZENECA S.P.A.	IT
RATACAND 8 mg compresse	UK/H/0197/003	033577166	ASTRAZENECA S.P.A.	IT
RATACAND 8 mg compresse	UK/H/0197/003	033577661	ASTRAZENECA S.P.A.	IT
RATACAND 8 mg compresse	UK/H/0197/003	033577608	ASTRAZENECA S.P.A.	IT
RATACAND 8 mg compresse	UK/H/0197/003	033577659	ASTRAZENECA S.P.A.	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
RATACAND 8 mg compresse	UK/H/0197/003	033577723	ASTRAZENECA S.P.A.	IT
RATACAND 8 mg compresse	UK/H/0197/003	033577596	ASTRAZENECA S.P.A.	IT
Ratacand Plus 16 mg/12,5 mg compresse	SE/H/0162/002	034186142	ASTRAZENECA S.P.A.	IT
Ratacand Plus 16 mg/12,5 mg compresse	SE/H/0162/002	034186155	ASTRAZENECA S.P.A.	IT
Ratacand Plus 16 mg/12,5 mg compresse	SE/H/0162/002	034186181	ASTRAZENECA S.P.A.	IT
Ratacand Plus 16 mg/12,5 mg compresse	SE/H/0162/002	034186243	ASTRAZENECA S.P.A.	IT
Ratacand Plus 16 mg/12,5 mg compresse	SE/H/0162/002	034186229	ASTRAZENECA S.P.A.	IT
Ratacand Plus 16 mg/12,5 mg compresse	SE/H/0162/002	034186179	ASTRAZENECA S.P.A.	IT
Ratacand Plus 16 mg/12,5 mg compresse	SE/H/0162/002	034186-130	ASTRAZENECA S.P.A.	IT
Ratacand Plus 16 mg/12,5 mg compresse	SE/H/0162/002	034186231	ASTRAZENECA S.P.A.	IT
Ratacand Plus 16 mg/12,5 mg compresse	SE/H/0162/002	034186193	ASTRAZENECA S.P.A.	IT
Ratacand Plus 16 mg/12,5 mg compresse	SE/H/0162/002	034186205	ASTRAZENECA S.P.A.	IT
Ratacand Plus 16 mg/12,5 mg compresse	SE/H/0162/002	034186167	ASTRAZENECA S.P.A.	IT
Ratacand Plus 16 mg/12,5 mg compresse	SE/H/0162/002	034186217	ASTRAZENECA S.P.A.	IT
Ratacand Plus 16 mg/12,5 mg compresse	SE/H/0162/002	034186256	ASTRAZENECA S.P.A.	IT
Ratacand Plus 16 mg/12,5 mg compresse	SE/H/0162/002	034186270	ASTRAZENECA S.P.A.	IT
Ratacand Plus 16 mg/12,5 mg compresse	SE/H/0162/002	034186268	ASTRAZENECA S.P.A.	IT

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Ratacand Plus 32 mg/12,5 mg compresse	SE/H/0162/003	034186282	ASTRAZENECA S.P.A.	IT
Ratacand Plus 32 mg/12,5 mg compresse	SE/H/0162/003	034186421	ASTRAZENECA S.P.A.	IT
Ratacand Plus 32 mg/12,5 mg compresse	SE/H/0162/003	034186294	ASTRAZENECA S.P.A.	IT
Ratacand Plus 32 mg/12,5 mg compresse	SE/H/0162/003	034186306	ASTRAZENECA S.P.A.	IT
Ratacand Plus 32 mg/12,5 mg compresse	SE/H/0162/003	034186320	ASTRAZENECA S.P.A.	IT
Ratacand Plus 32 mg/12,5 mg compresse	SE/H/0162/003	034186332	ASTRAZENECA S.P.A.	IT
Ratacand Plus 32 mg/12,5 mg compresse	SE/H/0162/003	034186344	ASTRAZENECA S.P.A.	IT
Ratacand Plus 32 mg/12,5 mg compresse	SE/H/0162/003	034186383	ASTRAZENECA S.P.A.	IT
Ratacand Plus 32 mg/12,5 mg compresse	SE/H/0162/003	034186371	ASTRAZENECA S.P.A.	IT
Ratacand Plus 32 mg/12,5 mg compresse	SE/H/0162/003	034186357	ASTRAZENECA S.P.A.	IT
Ratacand Plus 32 mg/12,5 mg compresse	SE/H/0162/003	034186395	ASTRAZENECA S.P.A.	IT
Ratacand Plus 32 mg/12,5 mg compresse	SE/H/0162/003	034186369	ASTRAZENECA S.P.A.	IT
Ratacand Plus 32 mg/12,5 mg compresse	SE/H/0162/003	034186407	ASTRAZENECA S.P.A.	IT
Ratacand Plus 32 mg/12,5 mg compresse	SE/H/0162/003	034186419	ASTRAZENECA S.P.A.	IT
Ratacand Plus 32 mg/12,5 mg compresse	SE/H/0162/003	034186318	ASTRAZENECA S.P.A.	IT
Ratacand Plus 32 mg/25 mg compresse	SE/H/0162/004	034186433	ASTRAZENECA S.P.A.	IT
Ratacand Plus 32 mg/25 mg compresse	SE/H/0162/004	034186445	ASTRAZENECA S.P.A.	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
Ratacand Plus 32 mg/25 mg compresse	SE/H/0162/004	034186458	ASTRAZENECA S.P.A.	IT
Ratacand Plus 32 mg/25 mg compresse	SE/H/0162/004	034186496	ASTRAZENECA S.P.A.	IT
Ratacand Plus 32 mg/25 mg compresse	SE/H/0162/004	034186508	ASTRAZENECA S.P.A.	IT
Ratacand Plus 32 mg/25 mg compresse	SE/H/0162/004	034186484	ASTRAZENECA S.P.A.	IT
Ratacand Plus 32 mg/25 mg compresse	SE/H/0162/004	034186460	ASTRAZENECA S.P.A.	IT
Ratacand Plus 32 mg/25 mg compresse	SE/H/0162/004	034186472	ASTRAZENECA S.P.A.	IT
Ratacand Plus 32 mg/25 mg compresse	SE/H/0162/004	034186510	ASTRAZENECA S.P.A.	IT
Ratacand Plus 32 mg/25 mg compresse	SE/H/0162/004	034186561	ASTRAZENECA S.P.A.	IT
Ratacand Plus 32 mg/25 mg compresse	SE/H/0162/004	034186559	ASTRAZENECA S.P.A.	IT
Ratacand Plus 32 mg/25 mg compresse	SE/H/0162/004	034186546	ASTRAZENECA S.P.A.	IT
Ratacand Plus 32 mg/25 mg compresse	SE/H/0162/004	034186534	ASTRAZENECA S.P.A.	IT
Ratacand Plus 32 mg/25 mg compresse	SE/H/0162/004	034186522	ASTRAZENECA S.P.A.	IT
Ratacand Plus 8 mg/12,5 mg compresse	SE/H/0162/001	034186128	ASTRAZENECA S.P.A.	IT
Ratacand Plus 8 mg/12,5 mg compresse	SE/H/0162/001	034186104	ASTRAZENECA S.P.A.	IT
Ratacand Plus 8 mg/12,5 mg compresse	SE/H/0162/001	034186041	ASTRAZENECA S.P.A.	IT
Ratacand Plus 8 mg/12,5 mg compresse	SE/H/0162/001	034186080	ASTRAZENECA S.P.A.	IT
Ratacand Plus 8 mg/12,5 mg compresse	SE/H/0162/001	034186039	ASTRAZENECA S.P.A.	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
Ratacand Plus 8 mg/12,5 mg compresse	SE/H/0162/001	034186027	ASTRAZENECA S.P.A.	IT
Ratacand Plus 8 mg/12,5 mg compresse	SE/H/0162/001	034186092	ASTRAZENECA S.P.A.	IT
Ratacand Plus 8 mg/12,5 mg compresse	SE/H/0162/001	034186078	ASTRAZENECA S.P.A.	IT
Ratacand Plus 8 mg/12,5 mg compresse	SE/H/0162/001	034186054	ASTRAZENECA S.P.A.	IT
Ratacand Plus 8 mg/12,5 mg compresse	SE/H/0162/001	034186066	ASTRAZENECA S.P.A.	IT
Ratacand Plus 8 mg/12,5 mg compresse	SE/H/0162/001	034186015	ASTRAZENECA S.P.A.	IT
Ratacand Plus 8 mg/12,5 mg compresse	SE/H/0162/001	034186116	ASTRAZENECA S.P.A.	IT
Xaleec Combi 8 mg/12,5 mg tablety	DE/H/1828/001	58/346/10-C	SANDOZ S.R.O.	CZ
Атаканд 16 mg таблетки	UK/H/0197/004	20060846	ASTRAZENECA AB	BG
Атаканд 8 mg таблетки	UK/H/0197/003	20060845	ASTRAZENECA AB	BG
Кандесартан HCT Ауробиндо 8 mg/12,5 mg таблетки	PT/H/1135/001	20160028	AUROBINDO PHARMA (MALTA) LIMITED	BG
Кандесартан Актавис 32 mg таблетки	DK/H/1838/004	20110314	ACTAVIS GROUP PTC EHF.	BG