26 October 2017
EMA/690954/2017
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: fexofenadine

Procedure no.: PSUSA/00001388/201703

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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
TELFAST 30 MG FILM-COATED TABLET	UK/H/0151/005	PL 04425/0162	AVENTIS PHARMA LTD	UK
TELFAST 30 MG FILM-COATED TABLET	UK/H/0151/005	PL 04425/0162	AVENTIS PHARMA LTD	UK
TELFAST 30 MG FILM-COATED TABLET	UK/H/0151/005	PL 04425/0162	AVENTIS PHARMA LTD	UK
TELFAST 30 MG FILM-COATED TABLET	UK/H/0151/005	PL 04425/0162	AVENTIS PHARMA LTD	UK
TELFAST 30 MG FILM-COATED TABLET	UK/H/0151/005	PL 04425/0162	AVENTIS PHARMA LTD	UK
TELFAST 30 MG FILM-COATED TABLET	UK/H/0151/005	PL 04425/0162	AVENTIS PHARMA LTD	UK
TELFAST 30 MG FILM-COATED TABLET	UK/H/0151/005	PL 04425/0162	AVENTIS PHARMA LTD	UK
TELFAST 30 MG FILM-COATED TABLET	UK/H/0151/005	PL 04425/0162	AVENTIS PHARMA LTD	UK
TELFAST 30 MG FILM-COATED TABLET	UK/H/0151/005	PL 04425/0162	AVENTIS PHARMA LTD	UK
TELFAST 30 MG FILM-COATED TABLET	UK/H/0151/005	PL 04425/0162	AVENTIS PHARMA LTD	UK
TELFAST 30 MG FILM-COATED TABLET	UK/H/0151/005	PL 04425/0162	AVENTIS PHARMA LTD	UK
TELFAST 30 MG FILM-COATED TABLET	UK/H/0151/005	PL 04425/0162	AVENTIS PHARMA LTD	UK
ALLEGRA	UK/H/5244/001	51557	SANOFI-AVENTIS DENMARK A/S	DK
ALLEGRA	UK/H/5244/001	51557	SANOFI-AVENTIS DENMARK A/S	DK

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
ALLEGRA	UK/H/5244/001	51557	SANOFI-AVENTIS DENMARK A/S	DK
ALLEGRA	UK/H/5244/001	51557	SANOFI-AVENTIS DENMARK A/S	DK
ALLEGRA	UK/H/5244/001	51557	SANOFI-AVENTIS DENMARK A/S	DK
ALLEGRA	UK/H/5244/001	51557	SANOFI-AVENTIS DENMARK A/S	DK
ALLEGRA	UK/H/5244/001	51557	SANOFI-AVENTIS DENMARK A/S	DK
ALLEGRA	UK/H/5244/001	51557	SANOFI-AVENTIS DENMARK A/S	DK
ALLEGRA	UK/H/5244/001	51557	SANOFI-AVENTIS DENMARK A/S	DK
ALLEGRA 120 MG FILMSKO OBLOZENE TABLETE	not available	H/98/00143/004	SANOFI-AVENTIS D.O.O.	SI
TELFAST 30 MG FILMSKO OBLOZENE TABLETE ZA OTROKE	not available	H/98/00143/003	SANOFI-AVENTIS D.O.O.	SI
TELFAST 180 MG FILMSKO OBLOZENE TABLETE	not available	H/98/00143/002	SANOFI-AVENTIS D.O.O.	SI
TELFAST	not available	18151	SANOFI-AVENTIS CYPRUS LTD	CY
FEXOFENADINE ZENTIVA 120 MG, COMPRIME PELLICULE	not available	345 459-8	SANOFI-AVENTIS FRANCE	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
FEXOFENADINE ZENTIVA 180 MG, COMPRIME PELLICULE	not available	345 462-9	SANOFI-AVENTIS FRANCE	FR
FEXOFENADINE ZENTIVA 120 MG, COMPRIME PELLICULE	not available	348 261-4	SANOFI-AVENTIS FRANCE	FR
FEXOFENADINE ZENTIVA 120 MG, COMPRIME PELLICULE	not available	345 457-5	SANOFI-AVENTIS FRANCE	FR
FEXOFENADINE ZENTIVA 120 MG, COMPRIME PELLICULE	not available	345 458-1	SANOFI-AVENTIS FRANCE	FR
FEXOFENADINE ZENTIVA 120 MG, COMPRIME PELLICULE	not available	345 456-9	SANOFI-AVENTIS FRANCE	FR
FEXOFENADINE ZENTIVA 120 MG, COMPRIME PELLICULE	not available	348 262-0	SANOFI-AVENTIS FRANCE	FR
FEXOFENADINE ZENTIVA 180 MG, COMPRIME PELLICULE	not available	348 263-7	SANOFI-AVENTIS FRANCE	FR
FEXOFENADINE ZENTIVA 180 MG, COMPRIME PELLICULE	not available	348 264-3	SANOFI-AVENTIS FRANCE	FR
FEXOFENADINE ZENTIVA 180 MG, COMPRIME PELLICULE	not available	345 461-2	SANOFI-AVENTIS FRANCE	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
FEXOFENADINE ZENTIVA 180 MG, COMPRIME PELLICULE	not available	345 463-5	SANOFI-AVENTIS FRANCE	FR
FEXOFENADINE ZENTIVA 180 MG, COMPRIME PELLICULE	not available	345 460-6	SANOFI-AVENTIS FRANCE	FR
TELFAST 30 MG, COMPRIME PELLICULE	not available	362 450-5	SANOFI-AVENTIS FRANCE	FR
TELFAST 30 MG, COMPRIME PELLICULE	not available	361 866-3	SANOFI-AVENTIS FRANCE	FR
TELFAST 30 MG, COMPRIME PELLICULE	not available	362 446-8	SANOFI-AVENTIS FRANCE	FR
TELFAST 30 MG, COMPRIME PELLICULE	not available	362 447-4	SANOFI-AVENTIS FRANCE	FR
TELFAST 30 MG, COMPRIME PELLICULE	not available	362 448-0	SANOFI-AVENTIS FRANCE	FR
TELFAST 30 MG, COMPRIME PELLICULE	not available	362 454-0	SANOFI-AVENTIS FRANCE	FR
TELFAST 30 MG, COMPRIME PELLICULE	not available	362 452-8	SANOFI-AVENTIS FRANCE	FR
TELFAST 30 MG, COMPRIME PELLICULE	not available	362 449-7	SANOFI-AVENTIS FRANCE	FR
TELFAST 30 MG, COMPRIME PELLICULE	not available	564 960-4	SANOFI-AVENTIS FRANCE	FR
TELFAST 30 MG, COMPRIME PELLICULE	not available	362 451-1	SANOFI-AVENTIS FRANCE	FR
TELFAST 30 MG, COMPRIME PELLICULE	not available	362 453-4	SANOFI-AVENTIS FRANCE	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
TELFAST 30 MG, COMPRIME PELLICULE	not available	361 868-6	SANOFI-AVENTIS FRANCE	FR
TELFAST 120 COMPRIMIDOS RECUBIERTOS CON PELÍCULA	UK/H/0151/003	699812.0	MARION MERRELL S.A.	ES
TELFAST 120 MG FILMTABLETTA	not available	OGYI-T-6450/02	SANOFI-AVENTIS ZRT	HU
TELFAST 120 MG FILMTABLETTA	not available	OGYI-T-6450/07	SANOFI-AVENTIS ZRT	HU
TELFAST 120 MG FILMTABLETTA	not available	OGYI-T-6450/06	SANOFI-AVENTIS ZRT	HU
TELFAST 120 MG FILMTABLETTA	not available	OGYI-T-6450/01	SANOFI-AVENTIS ZRT	HU
Allegra 120 mg plėvele dengtos tabletės	UK/H/0151/003	LT/1/15/3795/001-009	UAB SANOFI-AVENTIS LIETUVA	LT
Allegra 120 mg plėvele dengtos tabletės	UK/H/0151/003	LT/1/15/3795/003	UAB SANOFI-AVENTIS LIETUVA	LT
Allegra 120 mg plėvele dengtos tabletės	UK/H/0151/003	LT/1/15/3795/001	UAB SANOFI-AVENTIS LIETUVA	LT
Allegra 120 mg plėvele dengtos tabletės	UK/H/0151/003	LT/1/15/3795/001-009	UAB SANOFI-AVENTIS LIETUVA	LT
Allegra 120 mg plėvele dengtos tabletės	UK/H/0151/003	LT/1/15/3795/001-009	UAB SANOFI-AVENTIS LIETUVA	LT
Allegra 120 mg plėvele dengtos tabletės	UK/H/0151/003	LT/1/15/3795/001-009	UAB SANOFI-AVENTIS LIETUVA	LT
Allegra 120 mg plėvele dengtos tabletės	UK/H/0151/003	LT/1/15/3795/006	UAB SANOFI-AVENTIS LIETUVA	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
Allegra 120 mg plėvele dengtos tabletės	UK/H/0151/003	LT/1/15/3795/001-009	UAB SANOFI-AVENTIS LIETUVA	LT
Allegra 120 mg plėvele dengtos tabletės	UK/H/0151/003	LT/1/15/3795/001-009	UAB SANOFI-AVENTIS LIETUVA	LT
Allegra 120 mg plėvele dengtos tabletės	UK/H/0151/003	LT/1/15/3795/001-009	UAB SANOFI-AVENTIS LIETUVA	LT
Allegra 120 mg plėvele dengtos tabletės	UK/H/0151/003	LT/1/15/3795/001-009	UAB SANOFI-AVENTIS LIETUVA	LT
Allegra 120 mg plėvele dengtos tabletės	UK/H/0151/003	LT/1/15/3795/007	UAB SANOFI-AVENTIS LIETUVA	LT
Allegra 120 mg plėvele dengtos tabletės	UK/H/0151/003	LT/1/15/3795/001-009	UAB SANOFI-AVENTIS LIETUVA	LT
Allegra 120 mg plėvele dengtos tabletės	UK/H/0151/003	LT/1/15/3795/001-009	UAB SANOFI-AVENTIS LIETUVA	LT
Allegra 120 mg plėvele dengtos tabletės	UK/H/0151/003	LT/1/15/3795/001-009	UAB SANOFI-AVENTIS LIETUVA	LT
Allegra 120 mg plėvele dengtos tabletės	UK/H/0151/003	LT/1/15/3795/005	UAB SANOFI-AVENTIS LIETUVA	LT
Allegra 120 mg plėvele dengtos tabletės	UK/H/0151/003	LT/1/15/3795/001-009	UAB SANOFI-AVENTIS LIETUVA	LT
Allegra 120 mg plėvele dengtos tabletės	UK/H/0151/003	LT/1/15/3795/001-009	UAB SANOFI-AVENTIS LIETUVA	LT
Allegra 120 mg plėvele dengtos tabletės	UK/H/0151/003	LT/1/15/3795/002	UAB SANOFI-AVENTIS LIETUVA	LT
Allegra 120 mg plėvele dengtos tabletės	UK/H/0151/003	LT/1/15/3795/001-009	UAB SANOFI-AVENTIS LIETUVA	LT
Allegra 120 mg plėvele dengtos tabletės	UK/H/0151/003	LT/1/15/3795/009	UAB SANOFI-AVENTIS LIETUVA	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
Allegra 120 mg plėvele dengtos tabletės	UK/H/0151/003	LT/1/15/3795/004	UAB SANOFI-AVENTIS LIETUVA	LT
Allegra 120 mg plėvele dengtos tabletės	UK/H/0151/003	LT/1/15/3795/001-009	UAB SANOFI-AVENTIS LIETUVA	LT
Allegra 120 mg plėvele dengtos tabletės	UK/H/0151/003	LT/1/15/3795/001-009	UAB SANOFI-AVENTIS LIETUVA	LT
Allegra 120 mg plėvele dengtos tabletės	UK/H/0151/003	LT/1/15/3795/008	UAB SANOFI-AVENTIS LIETUVA	LT
Allegra 120 mg plėvele dengtos tabletės	UK/H/0151/003	LT/1/15/3795/001-009	UAB SANOFI-AVENTIS LIETUVA	LT
Allegra 120 mg plėvele dengtos tabletės	UK/H/0151/003	LT/1/15/3795/003	UAB SANOFI-AVENTIS LIETUVA	LT
Allegra Junior 30 mg filmtabletta	not available	OGYI-T-10513/08	SANOFI-AVENTIS ZRT	HU
FEXALLEGRA 120 mg COMPRESSE RIVESTITE CON FILM	not available	042554042	SANOFI SPA	IT
FEXALLEGRA 120 mg COMPRESSE RIVESTITE CON FILM	not available	042554016	SANOFI SPA	IT
FEXALLEGRA 120 mg COMPRESSE RIVESTITE CON FILM	not available	042554028	SANOFI SPA	IT
FEXALLEGRA 120 mg COMPRESSE RIVESTITE CON FILM	not available	042554030	SANOFI SPA	IT
Telfast 30 mg filmom obalené tablety	not available	24/0079/05-S	SANOFI-AVENTIS SLOVAKIA SRO	SK

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
Telfast 30 mg filmom obalené tablety	not available	24/0079/05-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Telfast 30 mg filmom obalené tablety	not available	24/0079/05-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Telfast 30 mg filmom obalené tablety	not available	24/0079/05-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Telfast 30 mg filmom obalené tablety	not available	24/0079/05-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Telfast 30 mg filmom obalené tablety	not available	24/0079/05-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Telfast 30 mg filmom obalené tablety	not available	24/0079/05-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Telfast 30 mg filmom obalené tablety	not available	24/0079/05-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Telfast 30 mg filmom obalené tablety	not available	24/0079/05-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Telfast 30 mg filmom obalené tablety	not available	24/0079/05-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Telfast 30 mg filmom obalené tablety	not available	24/0079/05-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Telfast, filmovertrukne tabletter 120 mg	UK/H/0151/003	18916	SANOFI-AVENTIS DENMARK A/S	DK
Telfast, filmovertrukne tabletter 120 mg	UK/H/0151/003	18916	SANOFI-AVENTIS DENMARK A/S	DK
Telfast 180 mg filmom obalené tablety	not available	24/0037/99-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Telfast 180 mg filmom obalené tablety	not available	24/0037/99-S	SANOFI-AVENTIS SLOVAKIA SRO	SK

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
Allegra 120 mg potahované tablety	UK/H/0151/003	24/515/15-C	SANOFI-AVENTIS SRO	CZ
Allegra 120 mg potahované tablety	UK/H/0151/003	24/515/15-C	SANOFI-AVENTIS SRO	CZ
Allegra 120 mg potahované tablety	UK/H/0151/003	24/515/15-C	SANOFI-AVENTIS SRO	CZ
Allegra 120 mg potahované tablety	UK/H/0151/003	24/515/15-C	SANOFI-AVENTIS SRO	CZ
Allegra 120 mg potahované tablety	UK/H/0151/003	24/515/15-C	SANOFI-AVENTIS SRO	CZ
Allegra 120 mg potahované tablety	UK/H/0151/003	24/515/15-C	SANOFI-AVENTIS SRO	CZ
Allegra 120 mg potahované tablety	UK/H/0151/003	24/515/15-C	SANOFI-AVENTIS SRO	CZ
Allegra 120 mg potahované tablety	UK/H/0151/003	24/515/15-C	SANOFI-AVENTIS SRO	CZ
Allegra 120 mg potahované tablety	UK/H/0151/003	24/515/15-C	SANOFI-AVENTIS SRO	CZ
Allegra 120 mg potahované tablety	UK/H/0151/003	24/515/15-C	SANOFI-AVENTIS SRO	CZ
Allegra 120 mg potahované tablety	UK/H/0151/003	24/515/15-C	SANOFI-AVENTIS SRO	CZ
Allegra 120 mg potahované tablety	UK/H/0151/003	24/515/15-C	SANOFI-AVENTIS SRO	CZ
Allegra 120 mg potahované tablety	UK/H/0151/003	24/515/15-C	SANOFI-AVENTIS SRO	CZ
Allegra 120 mg potahované tablety	UK/H/0151/003	24/515/15-C	SANOFI-AVENTIS SRO	CZ

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
Allegra 120 mg potahované tablety	UK/H/0151/003	24/515/15-C	SANOFI-AVENTIS SRO	CZ
Allegra 120 mg potahované tablety	UK/H/0151/003	24/515/15-C	SANOFI-AVENTIS SRO	CZ
Allegra 120 mg potahované tablety	UK/H/0151/003	24/515/15-C	SANOFI-AVENTIS SRO	CZ
Allegra 120 mg potahované tablety	UK/H/0151/003	24/515/15-C	SANOFI-AVENTIS SRO	CZ
Allegra 120 mg filmdragerade tabletter	UK/H/0151/003	13650	SANOFI AB	SE
Allegra 120 mg filmdragerade tabletter	UK/H/0151/003	13650	SANOFI AB	SE
Allegra 120 mg filmdragerade tabletter	UK/H/0151/003	13650	SANOFI AB	SE
Allegra 120 mg filmdragerade tabletter	UK/H/0151/003	13650	SANOFI AB	SE
Allegra 120 mg filmdragerade tabletter	UK/H/0151/003	13650	SANOFI AB	SE
Allegra 120 mg filmdragerade tabletter	UK/H/0151/003	13650	SANOFI AB	SE
Allegra 120 mg filmtabletta	not available	OGYI-T-10513/09	SANOFI-AVENTIS ZRT	HU
Allegra 120 mg filmtabletta	not available	OGYI-T-10513/10	SANOFI-AVENTIS ZRT	HU
TELFAST 30 mg comprimés pelliculés	UK/H/0151/005	BE259131	SANOFI BELGIUM	BE
TELFAST 30 mg comprimés pelliculés	UK/H/0151/005	BE259131	SANOFI BELGIUM	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
TELFAST 30 mg comprimés pelliculés	UK/H/0151/005	BE259131	SANOFI BELGIUM	BE
TELFAST 30 mg comprimés pelliculés	UK/H/0151/005	BE259131	SANOFI BELGIUM	BE
TELFAST 30 mg comprimés pelliculés	UK/H/0151/005	BE259131	SANOFI BELGIUM	BE
TELFAST 30 mg comprimés pelliculés	UK/H/0151/005	BE259131	SANOFI BELGIUM	BE
TELFAST 30 mg comprimés pelliculés	UK/H/0151/005	BE259131	SANOFI BELGIUM	BE
TELFAST 30 mg comprimés pelliculés	UK/H/0151/005	BE259131	SANOFI BELGIUM	BE
TELFAST 30 mg comprimés pelliculés	UK/H/0151/005	BE259131	SANOFI BELGIUM	BE
TELFAST 30 mg comprimés pelliculés	UK/H/0151/005	BE259131	SANOFI BELGIUM	BE
TELFAST 30 mg comprimés pelliculés	UK/H/0151/005	BE259131	SANOFI BELGIUM	BE
TELFAST 30 mg comprimés pelliculés	UK/H/0151/005	BE259131	SANOFI BELGIUM	BE
Telfast 120 mg film-coated tablets.	UK/H/0151/003	PL 04425/0157	AVENTIS PHARMA LTD	UK
Telfast 120 mg film-coated tablets.	UK/H/0151/003	PL 04425/0157	AVENTIS PHARMA LTD	UK
Telfast 120 mg film-coated tablets.	UK/H/0151/003	PL 04425/0157	AVENTIS PHARMA LTD	UK
Telfast 120 mg film-coated tablets.	UK/H/0151/003	PL 04425/0157	AVENTIS PHARMA LTD	UK

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
Telfast 120 mg film-coated tablets.	UK/H/0151/003	PL 04425/0157	AVENTIS PHARMA LTD	UK
Telfast 120 mg film-coated tablets.	UK/H/0151/003	PL 04425/0157	AVENTIS PHARMA LTD	UK
Telfast 120 mg film-coated tablets.	UK/H/0151/003	PL 04425/0157	AVENTIS PHARMA LTD	UK
Telfast 120 mg kalvopäällysteiset tabletit	UK/H/0151/003	12882	SANOFI OY	FI
Telfast 120 mg film-coated tablets.	UK/H/0151/003	PL 04425/0157	AVENTIS PHARMA LTD	UK
Telfast 120 mg film-coated tablets.	UK/H/0151/003	PL 04425/0157	AVENTIS PHARMA LTD	UK
Telfast 120 mg kalvopäällysteiset tabletit	UK/H/0151/003	12882	SANOFI OY	FI
TELFAST® 120 MG FILMTABLETTEN	UK/H/0151/003	40858.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
TELFAST® 120 MG FILMTABLETTEN	UK/H/0151/003	40858.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
TELFAST® 120 MG FILMTABLETTEN	UK/H/0151/003	40858.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
TELFAST® 120 MG FILMTABLETTEN	UK/H/0151/003	40858.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
TELFAST® 120 MG FILMTABLETTEN	UK/H/0151/003	40858.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
TELFAST® 120 MG FILMTABLETTEN	UK/H/0151/003	40858.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Telfast 180 mg film-coated tablets	UK/H/0151/004	PL 04425/0158	AVENTIS PHARMA LTD	UK

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
Telfast 180 mg film-coated tablets	UK/H/0151/004	PL 04425/0158	AVENTIS PHARMA LTD	UK
Telfast 180 mg film-coated tablets	UK/H/0151/004	PL 04425/0158	AVENTIS PHARMA LTD	UK
Telfast 180 mg film-coated tablets	UK/H/0151/004	PL 04425/0158	AVENTIS PHARMA LTD	UK
Telfast 180 mg film-coated tablets	UK/H/0151/004	PL 04425/0158	AVENTIS PHARMA LTD	UK
Telfast 180 mg film-coated tablets	UK/H/0151/004	PL 04425/0158	AVENTIS PHARMA LTD	UK
Telfast 180 mg film-coated tablets	UK/H/0151/004	PL 04425/0158	AVENTIS PHARMA LTD	UK
Telfast 180 mg film-coated tablets	UK/H/0151/004	PL 04425/0158	AVENTIS PHARMA LTD	UK
TELFAST 180 MG FILMDRAGERADE TABLETTER	UK/H/0151/004	13651	SANOFI AB	SE
TELFAST 180 MG FILMDRAGERADE TABLETTER	UK/H/0151/004	13651	SANOFI AB	SE
FEOXFENADINA SANOFI 180 MG COMPRIMIDOS RECUBIERTOS CON PELICULA	UK/H/0151/004	61911	SANOFI-AVENTIS, S.A.	ES
Telfast 180 mg film-coated tablets	UK/H/0151/004	PA 540/78/3	SANOFI-AVENTIS IRELAND LTD	IE

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
TELFAST 120 MG COMPRESSE RIVESTITE CON FILM	UK/H/0151/003	033303239	SANOFI SPA	IT
Telfast 120 mg film-coated tablets	UK/H/0151/003	PA 540/78/2	SANOFI-AVENTIS IRELAND LTD	IE
Telfast 120 mg comprimido revestido por película	UK/H/0151/003	2594984	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Telfast 120 mg comprimido revestido por película	UK/H/0151/003	5349964	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
TELFAST 120 MG COMPRESSE RIVESTITE CON FILM	UK/H/0151/003	033303049	SANOFI SPA	IT
TELFAST 120 MG COMPRESSE RIVESTITE CON FILM	UK/H/0151/003	033303227	SANOFI SPA	IT
TELFAST 120 MG COMPRESSE RIVESTITE CON FILM	UK/H/0151/003	033303215	SANOFI SPA	IT
TELFAST 120 MG COMPRESSE RIVESTITE CON FILM	UK/H/0151/003	033303203	SANOFI SPA	IT
TELFAST 120 MG COMPRESSE RIVESTITE CON FILM	UK/H/0151/003	033303177	SANOFI 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
TELFAST 120 MG COMPRESSE RIVESTITE CON FILM	UK/H/0151/003	033303189	SANOFI SPA	IT
TELFAST 120 MG COMPRESSE RIVESTITE CON FILM	UK/H/0151/003	033303191	SANOFI SPA	IT
Telfast 120 mg comprimate filmate	UK/H/0151/003	8315/2015/18	SANOFI ROMANIA SRL	RO
Telfast 120 mg comprimate filmate	UK/H/0151/003	8315/2015/16	SANOFI ROMANIA SRL	RO
Telfast 120 mg comprimate filmate	UK/H/0151/003	8315/2015/06	SANOFI ROMANIA SRL	RO
Telfast 120 mg comprimate filmate	UK/H/0151/003	8315/2015/10	SANOFI ROMANIA SRL	RO
Telfast 120 mg comprimate filmate	UK/H/0151/003	8315/2015/07	SANOFI ROMANIA SRL	RO
Telfast 120 mg comprimate filmate	UK/H/0151/003	8315/2015/17	SANOFI ROMANIA SRL	RO
Telfast 120 mg comprimate filmate	UK/H/0151/003	8315/2015/14	SANOFI ROMANIA SRL	RO
Telfast 120 mg comprimate filmate	UK/H/0151/003	8315/2015/13	SANOFI ROMANIA SRL	RO
Telfast 120 mg comprimate filmate	UK/H/0151/003	8315/2015/02	SANOFI ROMANIA SRL	RO
Telfast 120 mg comprimate filmate	UK/H/0151/003	8315/2015/08	SANOFI ROMANIA SRL	RO
Telfast 120 mg comprimate filmate	UK/H/0151/003	8315/2015/09	SANOFI ROMANIA SRL	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
Telfast 120 mg comprimate filmate	UK/H/0151/003	8315/2015/04	SANOFI ROMANIA SRL	RO
Telfast 120 mg comprimate filmate	UK/H/0151/003	8315/2015/03	SANOFI ROMANIA SRL	RO
Telfast 120 mg comprimate filmate	UK/H/0151/003	8315/2015/12	SANOFI ROMANIA SRL	RO
Telfast 120 mg comprimate filmate	UK/H/0151/003	8315/2015/01	SANOFI ROMANIA SRL	RO
Telfast 120 mg comprimate filmate	UK/H/0151/003	8315/2015/05	SANOFI ROMANIA SRL	RO
Telfast 120 mg comprimate filmate	UK/H/0151/003	8315/2015/11	SANOFI ROMANIA SRL	RO
Telfast 120 mg comprimate filmate	UK/H/0151/003	8315/2015/15	SANOFI ROMANIA SRL	RO
ALLEGRA 120 MG FILMTABLETTA	not available	OGYI-T-10513/04	SANOFI-AVENTIS ZRT	HU
ALLEGRA 120 MG FILMTABLETTA	not available	OGYI-T-10513/03	SANOFI-AVENTIS ZRT	HU
ALLEGRA 120 MG FILMTABLETTA	not available	OGYI-T-10513/05	SANOFI-AVENTIS ZRT	HU
ALLEGRA	not available	4195	SANOFI-AVENTIS DEUTSCHLAND GMBH	PL
Fexofenadina Sanofi 120 mg compresse rivestite con film	not available	033304041	SANOFI SPA	IT
FEXOFENADINA SANOFI	not available	033304039	SANOFI 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
FEXOFENADIN WINTHROP 120MG FILMTABLETTEN	not available	40860.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Fexofenadina Sanofi 120 mg compresse rivestite con film	not available	033304078	SANOFI SPA	IT
FEXOFENADINA SANOFI	not available	033304080	SANOFI SPA	IT
FEXOFENADIN WINTHROP 120MG FILMTABLETTEN	not available	40860.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
FEXOFENADIN WINTHROP 120MG FILMTABLETTEN	not available	40860.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
FEXOFENADIN WINTHROP 120MG FILMTABLETTEN	not available	40860.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
FEXOFENADIN WINTHROP 120MG FILMTABLETTEN	not available	40860.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Fexofenadine hydrochloride 120 mg film-coated tablets.	UK/H/5244/001	PL 04425/0667	AVENTIS PHARMA LTD	UK
Fexofenadine hydrochloride 120 mg film-coated tablets.	UK/H/5244/001	PL 04425/0667	AVENTIS PHARMA LTD	UK
Fexofenadine hydrochloride 120 mg film-coated tablets.	UK/H/5244/001	PL 04425/0667	AVENTIS PHARMA LTD	UK

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
Fexofenadine hydrochloride 120 mg film-coated tablets.	UK/H/5244/001	PL 04425/0667	AVENTIS PHARMA LTD	UK
Fexofenadine hydrochloride 120 mg film-coated tablets.	UK/H/5244/001	PL 04425/0667	AVENTIS PHARMA LTD	UK
Fexofenadine hydrochloride 120 mg film-coated tablets.	UK/H/5244/001	PL 04425/0667	AVENTIS PHARMA LTD	UK
Fexofenadine hydrochloride 120 mg film-coated tablets.	UK/H/5244/001	PL 04425/0667	AVENTIS PHARMA LTD	UK
Fexofenadine hydrochloride 120 mg film-coated tablets.	UK/H/5244/001	PL 04425/0667	AVENTIS PHARMA LTD	UK
Fexofenadine hydrochloride 120 mg film-coated tablets.	UK/H/5244/001	PL 04425/0667	AVENTIS PHARMA LTD	UK
Telfast 30 mg comprimidos revestidos por película	UK/H/0151/005	4920880	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Telfast 30 mg comprimidos revestidos por película	UK/H/0151/005	4921086	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Telfast 30 mg comprimidos revestidos por película	UK/H/0151/005	4920187	SANOFI - 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
Telfast 30 mg comprimidos revestidos por película	UK/H/0151/005	4920484	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Telfast 30 mg comprimidos revestidos por película	UK/H/0151/005	4920286	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Telfast 30 mg comprimidos revestidos por película	UK/H/0151/005	4919981	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Telfast 30 mg comprimidos revestidos por película	UK/H/0151/005	4920088	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Telfast 30 mg comprimidos revestidos por película	UK/H/0151/005	4920781	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Telfast 30 mg comprimidos revestidos por película	UK/H/0151/005	4920583	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Telfast 30 mg comprimidos revestidos por película	UK/H/0151/005	4920989	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Telfast 30 mg comprimidos revestidos por película	UK/H/0151/005	4920682	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Telfast 30 mg comprimidos revestidos por película	UK/H/0151/005	4920385	SANOFI - 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
Telfast 30 mg tabletti, kalvopäällysteinen	UK/H/0151/005	18570	SANOFI OY	FI
Telfast 30 mg tabletti, kalvopäällysteinen	UK/H/0151/005	18570	SANOFI OY	FI
Telfast 30 mg tabletti, kalvopäällysteinen	UK/H/0151/005	18570	SANOFI OY	FI
Telfast 30 mg tabletti, kalvopäällysteinen	UK/H/0151/005	18570	SANOFI OY	FI
Telfast 30 mg tabletti, kalvopäällysteinen	UK/H/0151/005	18570	SANOFI OY	FI
Telfast 30 mg tabletti, kalvopäällysteinen	UK/H/0151/005	18570	SANOFI OY	FI
Telfast 30 mg tabletti, kalvopäällysteinen	UK/H/0151/005	18570	SANOFI OY	FI
Telfast 30 mg tabletti, kalvopäällysteinen	UK/H/0151/005	18570	SANOFI OY	FI
Telfast 30 mg tabletti, kalvopäällysteinen	UK/H/0151/005	18570	SANOFI OY	FI
Telfast 30 mg tabletti, kalvopäällysteinen	UK/H/0151/005	18570	SANOFI OY	FI
Telfast 30 mg tabletti, kalvopäällysteinen	UK/H/0151/005	18570	SANOFI OY	FI
Telfast 30 mg tabletti, kalvopäällysteinen	UK/H/0151/005	18570	SANOFI OY	FI
Telfast, filmovertrukne tabletter 180 mg	UK/H/0151/004	18917	SANOFI-AVENTIS DENMARK A/S	DK
Telfast, filmovertrukne tabletter 180 mg	UK/H/0151/004	18917	SANOFI-AVENTIS DENMARK A/S	DK

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
Telfast, filmovertrukne tabletter 30 mg	UK/H/0151/005	35535	SANOFI-AVENTIS DENMARK A/S	DK
Telfast, filmovertrukne tabletter 180 mg	UK/H/0151/004	18917	SANOFI-AVENTIS DENMARK A/S	DK
Telfast, filmovertrukne tabletter 180 mg	UK/H/0151/004	18917	SANOFI-AVENTIS DENMARK A/S	DK
Telfast, filmovertrukne tabletter 180 mg	UK/H/0151/004	18917	SANOFI-AVENTIS DENMARK A/S	DK
Telfast, filmovertrukne tabletter 180 mg	UK/H/0151/004	18917	SANOFI-AVENTIS DENMARK A/S	DK
Telfast, filmovertrukne tabletter 180 mg	UK/H/0151/004	18917	SANOFI-AVENTIS DENMARK A/S	DK
Telfast, filmovertrukne tabletter 30 mg	UK/H/0151/005	35535	SANOFI-AVENTIS DENMARK A/S	DK
Telfast, filmovertrukne tabletter 30 mg	UK/H/0151/005	35535	SANOFI-AVENTIS DENMARK A/S	DK
Telfast, filmovertrukne tabletter 30 mg	UK/H/0151/005	35535	SANOFI-AVENTIS DENMARK A/S	DK
Telfast, filmovertrukne tabletter 30 mg	UK/H/0151/005	35535	SANOFI-AVENTIS DENMARK A/S	DK
Telfast, filmovertrukne tabletter 180 mg	UK/H/0151/004	18917	SANOFI-AVENTIS DENMARK A/S	DK
Telfast, filmovertrukne tabletter 30 mg	UK/H/0151/005	35535	SANOFI-AVENTIS DENMARK A/S	DK
Telfast, filmovertrukne tabletter 120 mg	UK/H/0151/003	18916	SANOFI-AVENTIS DENMARK A/S	DK
Telfast, filmovertrukne tabletter 120 mg	UK/H/0151/003	18916	SANOFI-AVENTIS DENMARK A/S	DK

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
Telfast, filmovertrukne tabletter 120 mg	UK/H/0151/003	18916	SANOFI-AVENTIS DENMARK A/S	DK
Telfast, filmovertrukne tabletter 120 mg	UK/H/0151/003	18916	SANOFI-AVENTIS DENMARK A/S	DK
Telfast, filmovertrukne tabletter 120 mg	UK/H/0151/003	18916	SANOFI-AVENTIS DENMARK A/S	DK
Telfast, filmovertrukne tabletter 120 mg	UK/H/0151/003	18916	SANOFI-AVENTIS DENMARK A/S	DK
Telfast 30 mg film-coated tablets	UK/H/0151/005	540/78/1	SANOFI-AVENTIS IRELAND LTD	IE
Telfast 30 mg film-coated tablets	UK/H/0151/005	540/78/1	SANOFI-AVENTIS IRELAND LTD	IE
Telfast 30 mg film-coated tablets	UK/H/0151/005	540/78/1	SANOFI-AVENTIS IRELAND LTD	IE
Telfast 30 mg film-coated tablets	UK/H/0151/005	540/78/1	SANOFI-AVENTIS IRELAND LTD	IE
Telfast 30 mg film-coated tablets	UK/H/0151/005	540/78/1	SANOFI-AVENTIS IRELAND LTD	IE
Telfast 30 mg film-coated tablets	UK/H/0151/005	540/78/1	SANOFI-AVENTIS IRELAND LTD	IE
Telfast 30 mg film-coated tablets	UK/H/0151/005	540/78/1	SANOFI-AVENTIS IRELAND LTD	IE
Telfast 30 mg film-coated tablets	UK/H/0151/005	540/78/1	SANOFI-AVENTIS IRELAND LTD	IE
Telfast 30 mg film-coated tablets	UK/H/0151/005	540/78/1	SANOFI-AVENTIS IRELAND LTD	IE
Telfast 30 mg film-coated tablets	UK/H/0151/005	540/78/1	SANOFI-AVENTIS IRELAND LTD	IE

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
Telfast 30 mg film-coated tablets	UK/H/0151/005	540/78/1	SANOFI-AVENTIS IRELAND LTD	IE
Telfast 30 mg film-coated tablets	UK/H/0151/005	540/78/1	SANOFI-AVENTIS IRELAND LTD	IE
TELFAST 30 mg comprimés pelliculés	UK/H/0151/005	0362447	SANOFI BELGIUM	LU
TELFAST 30 mg comprimés pelliculés	UK/H/0151/005	0362501	SANOFI BELGIUM	LU
TELFAST 30 mg comprimés pelliculés	UK/H/0151/005	0362514	SANOFI BELGIUM	LU
TELFAST 30 mg comprimés pelliculés	UK/H/0151/005	0362481	SANOFI BELGIUM	LU
TELFAST 30 mg comprimés pelliculés	UK/H/0151/005	0362416	SANOFI BELGIUM	LU
TELFAST 30 mg comprimés pelliculés	UK/H/0151/005	0362495	SANOFI BELGIUM	LU
TELFAST 30 mg comprimés pelliculés	UK/H/0151/005	0362433	SANOFI BELGIUM	LU
TELFAST 30 mg comprimés pelliculés	UK/H/0151/005	0362402	SANOFI BELGIUM	LU
TELFAST 30 mg comprimés pelliculés	UK/H/0151/005	0362451	SANOFI BELGIUM	LU
TELFAST 30 mg comprimés pelliculés	UK/H/0151/005	0362478	SANOFI BELGIUM	LU
TELFAST 30 mg comprimés pelliculés	UK/H/0151/005	0362464	SANOFI BELGIUM	LU
TELFAST 30 mg comprimés pelliculés	UK/H/0151/005	0362528	SANOFI BELGIUM	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
TELFAST 180 mg comprimés pelliculés	UK/H/0151/004	0244328	SANOFI BELGIUM	LU
TELFAST 180 mg, filmomhulde tabletten	UK/H/0151/004	BE 190303	SANOFI BELGIUM	BE
TELFAST 180 mg, filmomhulde tabletten	UK/H/0151/004	BE 190303	SANOFI BELGIUM	BE
TELFAST 180 mg, filmomhulde tabletten	UK/H/0151/004	BE 190303	SANOFI BELGIUM	BE
TELFAST 180 mg, filmomhulde tabletten	UK/H/0151/004	BE 190303	SANOFI BELGIUM	BE
TELFAST 180 mg, filmomhulde tabletten	UK/H/0151/004	BE 190303	SANOFI BELGIUM	BE
TELFAST 180 mg, filmomhulde tabletten	UK/H/0151/004	BE 190303	SANOFI BELGIUM	BE
TELFAST 180 mg, filmomhulde tabletten	UK/H/0151/004	BE 190303	SANOFI BELGIUM	BE
TELFAST 180 mg, filmomhulde tabletten	UK/H/0151/004	BE 190303	SANOFI BELGIUM	BE
Telfast 30 mg, filmomhulde tabletten	UK/H/0151/005	BE 259131	SANOFI BELGIUM	BE
Telfast 30 mg, filmomhulde tabletten	UK/H/0151/005	BE 259131	SANOFI BELGIUM	BE
Telfast 30 mg, filmomhulde tabletten	UK/H/0151/005	BE 259131	SANOFI BELGIUM	BE
Telfast 30 mg, filmomhulde tabletten	UK/H/0151/005	BE 259131	SANOFI BELGIUM	BE
Telfast 30 mg, filmomhulde tabletten	UK/H/0151/005	BE 259131	SANOFI BELGIUM	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
Telfast 30 mg, filmomhulde tabletten	UK/H/0151/005	BE 259131	SANOFI BELGIUM	BE
Telfast 30 mg, filmomhulde tabletten	UK/H/0151/005	BE 259131	SANOFI BELGIUM	BE
Telfast 30 mg, filmomhulde tabletten	UK/H/0151/005	BE 259131	SANOFI BELGIUM	BE
Telfast 30 mg, filmomhulde tabletten	UK/H/0151/005	BE 259131	SANOFI BELGIUM	BE
Telfast 30 mg, filmomhulde tabletten	UK/H/0151/005	BE 259131	SANOFI BELGIUM	BE
Telfast 30 mg, filmomhulde tabletten	UK/H/0151/005	BE 259131	SANOFI BELGIUM	BE
Telfast 30 mg, filmomhulde tabletten	UK/H/0151/005	BE 259131	SANOFI BELGIUM	BE
Telfast 180 mg comprimido revestido por película	UK/H/0151/004	2595189	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Telfast 180 mg kalvopäällysteiset tabletit	UK/H/0151/004	12884	SANOFI OY	FI
Telfast 180 mg kalvopäällysteiset tabletit	UK/H/0151/004	12884	SANOFI OY	FI
Telfast 180 mg kalvopäällysteiset tabletit	UK/H/0151/004	12884	SANOFI OY	FI
Telfast 180 mg kalvopäällysteiset tabletit	UK/H/0151/004	12884	SANOFI OY	FI
Telfast 180 mg kalvopäällysteiset tabletit	UK/H/0151/004	12884	SANOFI 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
Telfast 180 mg kalvopäällysteiset tabletit	UK/H/0151/004	12884	SANOFI OY	FI
Telfast 180 mg kalvopäällysteiset tabletit	UK/H/0151/004	12884	SANOFI OY	FI
Telfast 180 mg kalvopäällysteiset tabletit	UK/H/0151/004	12884	SANOFI OY	FI
Telfast 120 mg kalvopäällysteiset tabletit	UK/H/0151/003	12882	SANOFI OY	FI
Telfast 120 mg kalvopäällysteiset tabletit	UK/H/0151/003	12882	SANOFI OY	FI
Telfast 120 mg kalvopäällysteiset tabletit	UK/H/0151/003	12882	SANOFI OY	FI
Telfast 120 mg kalvopäällysteiset tabletit	UK/H/0151/003	12882	SANOFI OY	FI
Telfast 120 mg kalvopäällysteiset tabletit	UK/H/0151/003	12882	SANOFI OY	FI
Telfast 120 mg kalvopäällysteiset tabletit	UK/H/0151/003	12882	SANOFI OY	FI
Telfast 120 mg kalvopäällysteiset tabletit	UK/H/0151/003	12882	SANOFI OY	FI
Allegra 120 mg επικαλυμμένα με λεπτό υμένιο δισκία	UK/H/0151/003	14256/15/25-10-2016	SANOFI-AVENTIS AEBE	GR
Allegra 120 mg επικαλυμμένα με λεπτό υμένιο δισκία	UK/H/0151/003	14256/15/25-10-2016	SANOFI-AVENTIS AEBE	GR

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
Allegra 120 mg επικαλυμμένα με λεπτό υμένιο δισκία	UK/H/0151/003	14256/15/25-10-2016	SANOFI-AVENTIS AEBE	GR
Allegra 120 mg επικαλυμμένα με λεπτό υμένιο δισκία	UK/H/0151/003	14256/15/25-10-2016	SANOFI-AVENTIS AEBE	GR
Allegra 120 mg επικαλυμμένα με λεπτό υμένιο δισκία	UK/H/0151/003	14256/15/25-10-2016	SANOFI-AVENTIS AEBE	GR
Allegra 120 mg επικαλυμμένα με λεπτό υμένιο δισκία	UK/H/0151/003	14256/15/25-10-2016	SANOFI-AVENTIS AEBE	GR
Allegra 120 mg επικαλυμμένα με λεπτό υμένιο δισκία	UK/H/0151/003	14256/15/25-10-2016	SANOFI-AVENTIS AEBE	GR
Allegra 120 mg επικαλυμμένα με λεπτό υμένιο δισκία	UK/H/0151/003	14256/15/25-10-2016	SANOFI-AVENTIS AEBE	GR
Allegra 120 mg επικαλυμμένα με λεπτό υμένιο δισκία	UK/H/0151/003	14256/15/25-10-2016	SANOFI-AVENTIS AEBE	GR
Allegra 120 mg επικαλυμμένα με λεπτό υμένιο δισκία	UK/H/0151/003	14256/15/25-10-2016	SANOFI-AVENTIS AEBE	GR
Allegra 120 mg επικαλυμμένα με λεπτό υμένιο δισκία	UK/H/0151/003	14256/15/25-10-2016	SANOFI-AVENTIS AEBE	GR

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
Allegra 120 mg επικαλυμμένα με λεπτό υμένιο δισκία	UK/H/0151/003	14256/15/25-10-2016	SANOFI-AVENTIS AEBE	GR
Allegra 120 mg επικαλυμμένα με λεπτό υμένιο δισκία	UK/H/0151/003	14256/15/25-10-2016	SANOFI-AVENTIS AEBE	GR
Allegra 120 mg επικαλυμμένα με λεπτό υμένιο δισκία	UK/H/0151/003	14256/15/25-10-2016	SANOFI-AVENTIS AEBE	GR
Allegra 120 mg επικαλυμμένα με λεπτό υμένιο δισκία	UK/H/0151/003	14256/15/25-10-2016	SANOFI-AVENTIS AEBE	GR
Allegra 120 mg επικαλυμμένα με λεπτό υμένιο δισκία	UK/H/0151/003	14256/15/25-10-2016	SANOFI-AVENTIS AEBE	GR
Telfast® 30 mg Filmtabletten	UK/H/0151/005	40858.02.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Telfast® 30 mg Filmtabletten	UK/H/0151/005	40858.02.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Telfast® 30 mg Filmtabletten	UK/H/0151/005	40858.02.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Telfast® 30 mg Filmtabletten	UK/H/0151/005	40858.02.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Telfast® 30 mg Filmtabletten	UK/H/0151/005	40858.02.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Telfast® 30 mg Filmtabletten	UK/H/0151/005	40858.02.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	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
Telfast® 30 mg Filmtabletten	UK/H/0151/005	40858.02.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Telfast® 30 mg Filmtabletten	UK/H/0151/005	40858.02.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Telfast® 30 mg Filmtabletten	UK/H/0151/005	40858.02.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Telfast® 30 mg Filmtabletten	UK/H/0151/005	40858.02.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Telfast® 30 mg Filmtabletten	UK/H/0151/005	40858.02.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Telfast® 30 mg Filmtabletten	UK/H/0151/005	40858.02.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Telfast® 180 mg Filmtabletten	UK/H/0151/004	40858.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Telfast® 180 mg Filmtabletten	UK/H/0151/004	40858.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Telfast® 180 mg Filmtabletten	UK/H/0151/004	40858.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Telfast® 180 mg Filmtabletten	UK/H/0151/004	40858.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Telfast® 180 mg Filmtabletten	UK/H/0151/004	40858.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Allegra 120 mg filmom obalené tablety	UK/H/0151/003	24/0473/15-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Allegra 120 mg filmom obalené tablety	UK/H/0151/003	24/0473/15-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Allegra 120 mg filmom obalené tablety	UK/H/0151/003	24/0473/15-S	SANOFI-AVENTIS SLOVAKIA SRO	SK

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
Allegra 120 mg filmom obalené tablety	UK/H/0151/003	24/0473/15-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Allegra 120 mg filmom obalené tablety	UK/H/0151/003	24/0473/15-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Allegra 120 mg filmom obalené tablety	UK/H/0151/003	24/0473/15-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Allegra 120 mg filmom obalené tablety	UK/H/0151/003	24/0473/15-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Allegra 120 mg filmom obalené tablety	UK/H/0151/003	24/0473/15-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Allegra 120 mg filmom obalené tablety	UK/H/0151/003	24/0473/15-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Allegra 120 mg filmom obalené tablety	UK/H/0151/003	24/0473/15-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Allegra 120 mg filmom obalené tablety	UK/H/0151/003	24/0473/15-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Allegra 120 mg filmom obalené tablety	UK/H/0151/003	24/0473/15-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Allegra 120 mg filmom obalené tablety	UK/H/0151/003	24/0473/15-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Allegra 120 mg filmom obalené tablety	UK/H/0151/003	24/0473/15-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Allegra 120 mg filmom obalené tablety	UK/H/0151/003	24/0473/15-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Allegra 120 mg filmom obalené tablety	UK/H/0151/003	24/0473/15-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Allegra 120 mg filmom obalené tablety	UK/H/0151/003	24/0473/15-S	SANOFI-AVENTIS SLOVAKIA SRO	SK

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
Allegra 120 mg filmom obalené tablety	UK/H/0151/003	24/0473/15-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Telfast 120 mg filmuhúðaðar töflur	not available	970359	SANOFI-AVENTIS NORGE AS	IS
Telfast 120 mg filmuhúðaðar töflur	not available	970359	SANOFI-AVENTIS NORGE AS	IS
Telfast 120 mg filmuhúðaðar töflur	not available	970359	SANOFI-AVENTIS NORGE AS	IS
Telfast 120 mg, filmdrasjerte tabletter	not available	96-3616	SANOFI-AVENTIS NORGE AS	NO
Telfast 120 mg, filmdrasjerte tabletter	not available	96-3616	SANOFI-AVENTIS NORGE AS	NO
Telfast 120 mg, filmdrasjerte tabletter	not available	96-3616	SANOFI-AVENTIS NORGE AS	NO
Telfast 120 mg, filmdrasjerte tabletter	not available	96-3616	SANOFI-AVENTIS NORGE AS	NO
Telfast 180 mg, filmdrasjerte tabletter	not available	96-3617	SANOFI-AVENTIS NORGE AS	NO
Telfast 180 mg, filmdrasjerte tabletter	not available	96-3617	SANOFI-AVENTIS NORGE AS	NO
Telfast 180 mg filmuhúðaðar töflur	not available	970360	SANOFI-AVENTIS NORGE AS	IS
Telfast 180 mg filmuhúðaðar töflur	not available	970360	SANOFI-AVENTIS NORGE AS	IS
Telfast 30 mg tabletter, filmdrasjerte	UK/H/0151/005	03-2004	SANOFI-AVENTIS NORGE AS	NO
Telfast 30 mg tabletter, filmdrasjerte	UK/H/0151/005	03-2004	SANOFI-AVENTIS NORGE AS	NO

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
Telfast 30 mg tabletter, filmdrasjerte	UK/H/0151/005	03-2004	SANOFI-AVENTIS NORGE AS	NO
Telfast 30 mg tabletter, filmdrasjerte	UK/H/0151/005	03-2004	SANOFI-AVENTIS NORGE AS	NO
Telfast 30 mg tabletter, filmdrasjerte	UK/H/0151/005	03-2004	SANOFI-AVENTIS NORGE AS	NO
Telfast 30 mg tabletter, filmdrasjerte	UK/H/0151/005	03-2004	SANOFI-AVENTIS NORGE AS	NO
Telfast 30 mg tabletter, filmdrasjerte	UK/H/0151/005	03-2004	SANOFI-AVENTIS NORGE AS	NO
Telfast 30 mg tabletter, filmdrasjerte	UK/H/0151/005	03-2004	SANOFI-AVENTIS NORGE AS	NO
Telfast 30 mg tabletter, filmdrasjerte	UK/H/0151/005	03-2004	SANOFI-AVENTIS NORGE AS	NO
Telfast 30 mg tabletter, filmdrasjerte	UK/H/0151/005	03-2004	SANOFI-AVENTIS NORGE AS	NO
Telfast 30 mg tabletter, filmdrasjerte	UK/H/0151/005	03-2004	SANOFI-AVENTIS NORGE AS	NO
Telfast 30 mg tabletter, filmdrasjerte	UK/H/0151/005	03-2004	SANOFI-AVENTIS NORGE AS	NO
Telfast 30 mg filmuhúðaðar töflur	UK/H/0151/005	IS/1/03/033/01	SANOFI-AVENTIS NORGE AS	IS
Telfast 30 mg filmuhúðaðar töflur	UK/H/0151/005	IS/1/03/033/01	SANOFI-AVENTIS NORGE AS	IS
Telfast 30 mg filmuhúðaðar töflur	UK/H/0151/005	IS/1/03/033/01	SANOFI-AVENTIS NORGE AS	IS
Telfast 30 mg filmuhúðaðar töflur	UK/H/0151/005	IS/1/03/033/01	SANOFI-AVENTIS NORGE AS	IS

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
Telfast 30 mg filmuhúðaðar töflur	UK/H/0151/005	IS/1/03/033/01	SANOFI-AVENTIS NORGE AS	IS
Telfast 30 mg filmuhúðaðar töflur	UK/H/0151/005	IS/1/03/033/01	SANOFI-AVENTIS NORGE AS	IS
Telfast 30 mg filmuhúðaðar töflur	UK/H/0151/005	IS/1/03/033/01	SANOFI-AVENTIS NORGE AS	IS
Telfast 30 mg filmuhúðaðar töflur	UK/H/0151/005	IS/1/03/033/01	SANOFI-AVENTIS NORGE AS	IS
Telfast 30 mg filmuhúðaðar töflur	UK/H/0151/005	IS/1/03/033/01	SANOFI-AVENTIS NORGE AS	IS
Telfast 30 mg filmuhúðaðar töflur	UK/H/0151/005	IS/1/03/033/01	SANOFI-AVENTIS NORGE AS	IS
Telfast 30 mg filmuhúðaðar töflur	UK/H/0151/005	IS/1/03/033/01	SANOFI-AVENTIS NORGE AS	IS
Telfast 30 mg filmuhúðaðar töflur	UK/H/0151/005	IS/1/03/033/01	SANOFI-AVENTIS NORGE AS	IS
ALLEGRA 120 MG	not available	20010553	SANOFI BULGARIA EOOD	BG
ALLEGRA 120 MG	not available	20010553	SANOFI BULGARIA EOOD	BG
Allegra 120 mg apvalkotās tabletes	UK/H/0151/003	15-0215	SANOFI-AVENTIS LATVIA SIA	LV
Allegra 120 mg apvalkotās tabletes	UK/H/0151/003	15-0215	SANOFI-AVENTIS LATVIA SIA	LV
Allegra 120 mg apvalkotās tabletes	UK/H/0151/003	15-0215	SANOFI-AVENTIS LATVIA SIA	LV
Allegra 120 mg õhukese polümeerikattega tabletid	UK/H/0151/003	878415	SANOFI-AVENTIS ESTONIA OÜ	EE

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
Allegra 120 mg õhukese polümeerikattega tabletid	UK/H/0151/003	878415	SANOFI-AVENTIS ESTONIA OÜ	EE
Allegra 120 mg õhukese polümeerikattega tabletid	UK/H/0151/003	878415	SANOFI-AVENTIS ESTONIA OÜ	EE
Allegra 120 mg apvalkotās tabletes	UK/H/0151/003	15-0215	SANOFI-AVENTIS LATVIA SIA	LV
Allegra 120 mg apvalkotās tabletes	UK/H/0151/003	15-0215	SANOFI-AVENTIS LATVIA SIA	LV
Allegra 120 mg apvalkotās tabletes	UK/H/0151/003	15-0215	SANOFI-AVENTIS LATVIA SIA	LV
Allegra 120 mg apvalkotās tabletes	UK/H/0151/003	15-0215	SANOFI-AVENTIS LATVIA SIA	LV
Allegra 120 mg apvalkotās tabletes	UK/H/0151/003	15-0215	SANOFI-AVENTIS LATVIA SIA	LV
Allegra 120 mg apvalkotās tabletes	UK/H/0151/003	15-0215	SANOFI-AVENTIS LATVIA SIA	LV
Allegra 120 mg apvalkotās tabletes	UK/H/0151/003	15-0215	SANOFI-AVENTIS LATVIA SIA	LV
Allegra 120 mg õhukese polümeerikattega tabletid	UK/H/0151/003	878415	SANOFI-AVENTIS ESTONIA OÜ	EE
Allegra 120 mg õhukese polümeerikattega tabletid	UK/H/0151/003	878415	SANOFI-AVENTIS ESTONIA OÜ	EE
Allegra 120 mg õhukese polümeerikattega tabletid	UK/H/0151/003	878415	SANOFI-AVENTIS ESTONIA OÜ	EE
Allegra 120 mg õhukese polümeerikattega tabletid	UK/H/0151/003	878415	SANOFI-AVENTIS ESTONIA OÜ	EE
Allegra 120 mg õhukese polümeerikattega tabletid	UK/H/0151/003	878415	SANOFI-AVENTIS ESTONIA OÜ	EE

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
Allegra 120 mg õhukese polümeerikattega tabletid	UK/H/0151/003	878415	SANOFI-AVENTIS ESTONIA OÜ	EE
Allegra 120 mg Filmtabletten	UK/H/0151/003	1-22153	SANOFI-AVENTIS GMBH OSTERREICH	AT
Allegra 120 mg Filmtabletten	UK/H/0151/003	1-22153	SANOFI-AVENTIS GMBH OSTERREICH	AT
Allegra 120 mg Filmtabletten	UK/H/0151/003	1-22153	SANOFI-AVENTIS GMBH OSTERREICH	AT
Allegra 120 mg Filmtabletten	UK/H/0151/003	1-22153	SANOFI-AVENTIS GMBH OSTERREICH	AT
Allegra 120 mg Filmtabletten	UK/H/0151/003	1-22153	SANOFI-AVENTIS GMBH OSTERREICH	AT
Allegra 120 mg Filmtabletten	UK/H/0151/003	1-22153	SANOFI-AVENTIS GMBH OSTERREICH	AT
Allegra 120 mg Filmtabletten	UK/H/0151/003	1-22153	SANOFI-AVENTIS GMBH OSTERREICH	AT
Allegra 120 mg Filmtabletten	UK/H/0151/003	1-22153	SANOFI-AVENTIS GMBH OSTERREICH	AT
Allegratab 120 mg film-coated tablets	UK/H/5244/001	MA082/07301	SANOFI MALTA LTD	MT
Allegratab 120 mg film-coated tablets	UK/H/5244/001	MA082/07301	SANOFI MALTA LTD	MT
Allegratab 120 mg film-coated tablets	UK/H/5244/001	MA082/07301	SANOFI MALTA LTD	MT
Allegratab 120 mg film-coated tablets	UK/H/5244/001	MA082/07301	SANOFI MALTA LTD	MT
Allegratab 120 mg film-coated tablets	UK/H/5244/001	MA082/07301	SANOFI MALTA LTD	MT

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
Allegratab 120 mg film-coated tablets	UK/H/5244/001	MA082/07301	SANOFI MALTA LTD	MT
Allegratab 120 mg film-coated tablets	UK/H/5244/001	MA082/07301	SANOFI MALTA LTD	MT
Allegratab 120 mg film-coated tablets	UK/H/5244/001	MA082/07301	SANOFI MALTA LTD	MT
Allegratab 120 mg film-coated tablets	UK/H/5244/001	MA082/07301	SANOFI MALTA LTD	MT
TELFAST 180 MG FILM COATED TABLETS	not available	20010554	SANOFI BULGARIA EOOD	BG
TELFAST 180 MG FILM COATED TABLETS	not available	20010554	SANOFI BULGARIA EOOD	BG
Telfast 120 mg filmsko obložene tablete	UK/H/0151/003	H/16/02158/001	SANOFI-AVENTIS D.O.O.	SI
Telfast 120 mg filmsko obložene tablete	UK/H/0151/003	H/16/02158/015	SANOFI-AVENTIS D.O.O.	SI
Telfast 120 mg filmsko obložene tablete	UK/H/0151/003	H/16/02158/002	SANOFI-AVENTIS D.O.O.	SI
Telfast 120 mg filmsko obložene tablete	UK/H/0151/003	H/16/02158/013	SANOFI-AVENTIS D.O.O.	SI
Telfast 120 mg filmsko obložene tablete	UK/H/0151/003	H/16/02158/003	SANOFI-AVENTIS D.O.O.	SI
Telfast 120 mg filmsko obložene tablete	UK/H/0151/003	H/16/02158/007	SANOFI-AVENTIS D.O.O.	SI
Telfast 120 mg filmsko obložene tablete	UK/H/0151/003	H/16/02158/010	SANOFI-AVENTIS D.O.O.	SI
Telfast 120 mg filmsko obložene tablete	UK/H/0151/003	H/16/02158/008	SANOFI-AVENTIS D.O.O.	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
Telfast 120 mg filmsko obložene tablete	UK/H/0151/003	H/16/02158/012	SANOFI-AVENTIS D.O.O.	SI
Telfast 120 mg filmsko obložene tablete	UK/H/0151/003	H/16/02158/011	SANOFI-AVENTIS D.O.O.	SI
Telfast 120 mg filmsko obložene tablete	UK/H/0151/003	H/16/02158/006	SANOFI-AVENTIS D.O.O.	SI
Telfast 120 mg filmsko obložene tablete	UK/H/0151/003	H/16/02158/016	SANOFI-AVENTIS D.O.O.	SI
Telfast 120 mg filmsko obložene tablete	UK/H/0151/003	H/16/02158/014	SANOFI-AVENTIS D.O.O.	SI
Telfast 120 mg filmsko obložene tablete	UK/H/0151/003	H/16/02158/005	SANOFI-AVENTIS D.O.O.	SI
Telfast 120 mg filmsko obložene tablete	UK/H/0151/003	H/16/02158/009	SANOFI-AVENTIS D.O.O.	SI
Telfast 120 mg filmsko obložene tablete	UK/H/0151/003	H/16/02158/017	SANOFI-AVENTIS D.O.O.	SI
Telfast 120 mg filmsko obložene tablete	UK/H/0151/003	H/16/02158/004	SANOFI-AVENTIS D.O.O.	SI
Telfast 30 mg επικαλυμμένα με λεπτό υμένιο δισκία	UK/H/0151/005	20746	SANOFI-AVENTIS CYPRUS LTD	CY
Telfast 30 mg επικαλυμμένα με λεπτό υμένιο δισκία	UK/H/0151/005	20746	SANOFI-AVENTIS CYPRUS LTD	CY
Telfast 30 mg επικαλυμμένα με λεπτό υμένιο δισκία	UK/H/0151/005	20746	SANOFI-AVENTIS CYPRUS LTD	CY

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
Telfast 30 mg επικαλυμμένα με λεπτό υμένιο δισκία	UK/H/0151/005	20746	SANOFI-AVENTIS CYPRUS LTD	CY
Telfast 30 mg επικαλυμμένα με λεπτό υμένιο δισκία	UK/H/0151/005	20746	SANOFI-AVENTIS CYPRUS LTD	CY
Telfast 30 mg επικαλυμμένα με λεπτό υμένιο δισκία	UK/H/0151/005	20746	SANOFI-AVENTIS CYPRUS LTD	CY
Telfast 30 mg επικαλυμμένα με λεπτό υμένιο δισκία	UK/H/0151/005	20746	SANOFI-AVENTIS CYPRUS LTD	CY
Telfast 30 mg επικαλυμμένα με λεπτό υμένιο δισκία	UK/H/0151/005	20746	SANOFI-AVENTIS CYPRUS LTD	CY
Telfast 30 mg επικαλυμμένα με λεπτό υμένιο δισκία	UK/H/0151/005	20746	SANOFI-AVENTIS CYPRUS LTD	CY
Telfast 30 mg επικαλυμμένα με λεπτό υμένιο δισκία	UK/H/0151/005	20746	SANOFI-AVENTIS CYPRUS LTD	CY
Telfast 30 mg επικαλυμμένα με λεπτό υμένιο δισκία	UK/H/0151/005	20746	SANOFI-AVENTIS CYPRUS LTD	CY
Telfast 30 mg επικαλυμμένα με λεπτό υμένιο δισκία	UK/H/0151/005	20746	SANOFI-AVENTIS CYPRUS LTD	CY

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
ALLEGRA FORTE 180 MG FILMTABLETTA	not available	OGYI-T-10513/11	SANOFI-AVENTIS ZRT	HU
ALLEGRA FORTE 180 MG FILMTABLETTA	not available	OGYI-T-10513/06	SANOFI-AVENTIS ZRT	HU
ALLEGRA FORTE 180 MG FILMTABLETTA	not available	OGYI-T-10513/12	SANOFI-AVENTIS ZRT	HU
ALLEGRA FORTE 180 MG FILMTABLETTA	not available	OGYI-T-10513/07	SANOFI-AVENTIS ZRT	HU
TELFAST 180 MG FILMTABLETTA	not available	OGYI-T- 6450/05	SANOFI-AVENTIS ZRT	HU
TELFAST 180 MG FILMTABLETTA	not available	OGYI-T- 6450/04	SANOFI-AVENTIS ZRT	HU
TELFAST 180 MG FILMTABLETTA	not available	OGYI-T-6450/08	SANOFI-AVENTIS ZRT	HU
TELFAST 180 MG FILMTABLETTA	not available	OGYI-T-6450/09	SANOFI-AVENTIS ZRT	HU
Allegra 120 mg filmdragerade tabletter	UK/H/0151/003	13650	SANOFI AB	SE
Allegra 120 mg filmdragerade tabletter	UK/H/0151/003	13650	SANOFI AB	SE
TELFAST JUNIOR 30 MG	not available	RVG 28085	SANOFI-AVENTIS NETHERLANDS B.V.	NL
TELFAST 180	not available	RVG 21625	SANOFI-AVENTIS NETHERLANDS B.V.	NL
ALLEGRA FEXOTABS	not available	RVG 21624	SANOFI-AVENTIS NETHERLANDS B.V.	NL
ALLEGRA FEXOTABS	not available	RVG 21624	SANOFI-AVENTIS NETHERLANDS B.V.	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
ALLEGRA FEXOTABS	not available	RVG 21624	SANOFI-AVENTIS NETHERLANDS B.V.	NL
ALLEGRA FEXOTABS	not available	RVG 21624	SANOFI-AVENTIS NETHERLANDS B.V.	NL
ALLEGRA FEXOTABS	not available	RVG 21624	SANOFI-AVENTIS NETHERLANDS B.V.	NL
ALLEGRA FEXOTABS	not available	RVG21624	SANOFI-AVENTIS NETHERLANDS B.V.	NL
ALLEGRA TAB 120 mg, filmomhulde tabletten	UK/H/0151/003	BE190312	SANOFI BELGIUM	BE
ALLEGRA TAB 120 mg comprimés pelliculés	UK/H/0151/003	0244314	SANOFI BELGIUM	LU
ALLEGRA TAB 120 mg comprimés pelliculés	UK/H/0151/003	0244301	SANOFI BELGIUM	LU
ALLEGRA TAB 120 mg comprimés pelliculés	UK/H/0151/003	BE190312	SANOFI BELGIUM	BE
ALLEGRA TAB 120 mg comprimés pelliculés	UK/H/0151/003	BE190312	SANOFI BELGIUM	BE
ALLEGRA TAB 120 mg, filmomhulde tabletten	UK/H/0151/003	BE190312	SANOFI BELGIUM	BE
ALLEGRA TAB 120 mg comprimés pelliculés	UK/H/0151/003	BE190312	SANOFI BELGIUM	BE
ALLEGRA TAB 120 mg comprimés pelliculés	UK/H/0151/003	BE190312	SANOFI BELGIUM	BE
ALLEGRA TAB 120 mg, filmomhulde tabletten	UK/H/0151/003	BE190312	SANOFI BELGIUM	BE
ALLEGRA TAB 120 mg comprimés pelliculés	UK/H/0151/003	BE190312	SANOFI BELGIUM	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
ALLEGRA TAB 120 mg comprimés pelliculés	UK/H/0151/003	BE190312	SANOFI BELGIUM	BE
ALLEGRA TAB 120 mg, filmomhulde tabletten	UK/H/0151/003	BE190312	SANOFI BELGIUM	BE
ALLEGRA TAB 120 mg, filmomhulde tabletten	UK/H/0151/003	BE190312	SANOFI BELGIUM	BE
ALLEGRA TAB 120 mg, filmomhulde tabletten	UK/H/0151/003	BE190312	SANOFI BELGIUM	BE
ALLEGRA TAB 120 mg, filmomhulde tabletten	UK/H/0151/003	BE190312	SANOFI BELGIUM	BE
ALLEGRA TAB 120 mg comprimés pelliculés	UK/H/0151/003	BE190312	SANOFI BELGIUM	BE
ALLEGRA TAB 120 mg comprimés pelliculés	UK/H/0151/003	BE190312	SANOFI BELGIUM	BE
ALLEGRA TAB 120 mg, filmomhulde tabletten	UK/H/0151/003	BE190312	SANOFI BELGIUM	BE
FEXOFENADINE HYDROCHLORIDE	UK/H/5244/001	PA 0540/170/001	SANOFI-AVENTIS IRELAND LTD	IE
FEXOFENADINE HYDROCHLORIDE	UK/H/5244/001	PA 0540/170/001	SANOFI-AVENTIS IRELAND LTD	IE
FEXOFENADINE HYDROCHLORIDE	UK/H/5244/001	PA 0540/170/001	SANOFI-AVENTIS IRELAND LTD	IE
FEXOFENADINE HYDROCHLORIDE	UK/H/5244/001	PA 0540/170/001	SANOFI-AVENTIS IRELAND LTD	IE
FEXOFENADINE HYDROCHLORIDE	UK/H/5244/001	PA 0540/170/001	SANOFI-AVENTIS IRELAND LTD	IE
FEXOFENADINE HYDROCHLORIDE	UK/H/5244/001	PA 0540/170/001	SANOFI-AVENTIS IRELAND LTD	IE

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**
FEXOFENADINE HYDROCHLORIDE	UK/H/5244/001	PA 0540/170/001	SANOFI-AVENTIS IRELAND LTD	IE
FEXOFENADINE HYDROCHLORIDE	UK/H/5244/001	PA 0540/170/001	SANOFI-AVENTIS IRELAND LTD	IE
FEXOFENADINE HYDROCHLORIDE	UK/H/5244/001	PA 0540/170/001	SANOFI-AVENTIS IRELAND LTD	IE
FEXOFENADIN WINTHROP 180MG FILMTABLETTEN	not available	76707.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
FEXOFENADIN WINTHROP 180MG FILMTABLETTEN	not available	76707.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
FEXOFENADIN WINTHROP 180MG FILMTABLETTEN	not available	76707.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
FEXOFENADIN WINTHROP 180MG FILMTABLETTEN	not available	76707.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
FEXOFENADIN WINTHROP 180MG FILMTABLETTEN	not available	76707.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
TELFAST 180	not available	4194	SANOFI-AVENTIS DEUTSCHLAND GMBH	PL
TELFAST 180	not available	4194	SANOFI-AVENTIS DEUTSCHLAND GMBH	PL
TELFAST 180	not available	4194	SANOFI-AVENTIS DEUTSCHLAND 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
Telfast 30, 30 mg, tabletki powlekane	UK/H/0151/005	17232	SANOFI-AVENTIS DEUTSCHLAND GMBH	PL
Telfast 30, 30 mg, tabletki powlekane	UK/H/0151/005	17232	SANOFI-AVENTIS DEUTSCHLAND GMBH	PL
Telfast 30, 30 mg, tabletki powlekane	UK/H/0151/005	17232	SANOFI-AVENTIS DEUTSCHLAND GMBH	PL
Telfast 30, 30 mg, tabletki powlekane	UK/H/0151/005	17232	SANOFI-AVENTIS DEUTSCHLAND GMBH	PL
Telfast 30, 30 mg, tabletki powlekane	UK/H/0151/005	17232	SANOFI-AVENTIS DEUTSCHLAND GMBH	PL
Telfast 30, 30 mg, tabletki powlekane	UK/H/0151/005	17232	SANOFI-AVENTIS DEUTSCHLAND GMBH	PL
Telfast 30, 30 mg, tabletki powlekane	UK/H/0151/005	17232	SANOFI-AVENTIS DEUTSCHLAND GMBH	PL
Telfast 30, 30 mg, tabletki powlekane	UK/H/0151/005	17232	SANOFI-AVENTIS DEUTSCHLAND GMBH	PL
Telfast 30, 30 mg, tabletki powlekane	UK/H/0151/005	17232	SANOFI-AVENTIS DEUTSCHLAND GMBH	PL
Telfast 30, 30 mg, tabletki powlekane	UK/H/0151/005	17232	SANOFI-AVENTIS DEUTSCHLAND GMBH	PL
Telfast 30, 30 mg, tabletki powlekane	UK/H/0151/005	17232	SANOFI-AVENTIS DEUTSCHLAND GMBH	PL
Telfast 30, 30 mg, tabletki powlekane	UK/H/0151/005	17232	SANOFI-AVENTIS DEUTSCHLAND GMBH	PL
TELFAST 180 mg compresse rivestite con film	UK/H/0151/004	033303254	SANOFI 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
TELFAST 180 mg compresse rivestite con film	UK/H/0151/004	033303292	SANOFI SPA	IT
TELFAST 180 mg compresse rivestite con film	UK/H/0151/004	033303241	SANOFI SPA	IT
TELFAST 180 mg compresse rivestite con film	UK/H/0151/004	033303037	SANOFI SPA	IT
TELFAST 180 mg compresse rivestite con film	UK/H/0151/004	033303280	SANOFI SPA	IT
TELFAST 180 mg compresse rivestite con film	UK/H/0151/004	033303266	SANOFI SPA	IT
TELFAST 120 mg, comprimé pelliculé	not available	348 260-8	SANOFI-AVENTIS FRANCE	FR
TELFAST 120 mg, comprimé pelliculé	not available	348 258-3	SANOFI-AVENTIS FRANCE	FR
TELFAST 120 mg, comprimé pelliculé	not available	345 446-3	SANOFI-AVENTIS FRANCE	FR
TELFAST 120 mg, comprimé pelliculé	not available	345 448-6	SANOFI-AVENTIS FRANCE	FR
TELFAST 120 mg, comprimé pelliculé	not available	345 450-0	SANOFI-AVENTIS FRANCE	FR
TELFAST 120 mg, comprimé pelliculé	not available	345 449-2	SANOFI-AVENTIS FRANCE	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
TELFAST 180 mg, comprimé pelliculé	not available	348 267-2	SANOFI-AVENTIS FRANCE	FR
TELFAST 180 mg, comprimé pelliculé	not available	348 266-6	SANOFI-AVENTIS FRANCE	FR
TELFAST 180 mg, comprimé pelliculé	not available	345 452-3	SANOFI-AVENTIS FRANCE	FR
TELFAST 180 mg, comprimé pelliculé	not available	345 455-2	SANOFI-AVENTIS FRANCE	FR
TELFAST 180 mg, comprimé pelliculé	not available	345 451-7	SANOFI-AVENTIS FRANCE	FR
TELFAST 180 mg, comprimé pelliculé	not available	345 454-6	SANOFI-AVENTIS FRANCE	FR
TELFAST 120MG FILM COATED TABLETS	not available	MA082/03601	SANOFI MALTA LTD	MT
TELFAST 120MG FILM COATED TABLETS	not available	MA082/03601	SANOFI MALTA LTD	MT
TELFAST 120MG FILM COATED TABLETS	not available	MA082/03601	SANOFI MALTA LTD	MT
TELFAST 120MG FILM COATED TABLETS	not available	MA082/03601	SANOFI MALTA LTD	MT
TELFAST 120MG FILM COATED TABLETS	not available	MA082/03601	SANOFI MALTA LTD	MT
TELFAST 120MG FILM COATED TABLETS	not available	MA082/03601	SANOFI MALTA LTD	MT
TELFAST 120MG FILM COATED TABLETS	not available	MA082/03601	SANOFI MALTA LTD	MT
TELFAST 120MG FILM COATED TABLETS	not available	MA082/03601	SANOFI MALTA LTD	MT

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
TELFAST 120MG FILM COATED TABLETS	not available	MA082/03601	SANOFI MALTA LTD	MT
TELFAST 180 MG FILM COATED TABLETS	not available	MA082/03602	SANOFI MALTA LTD	MT
TELFAST 180 MG FILM COATED TABLETS	not available	MA082/03602	SANOFI MALTA LTD	MT
TELFAST 180 MG FILM COATED TABLETS	not available	MA082/03602	SANOFI MALTA LTD	MT
TELFAST 180 MG FILM COATED TABLETS	not available	MA082/03602	SANOFI MALTA LTD	MT
TELFAST 180 MG FILM COATED TABLETS	not available	MA082/03602	SANOFI MALTA LTD	MT
TELFAST 180 MG FILM COATED TABLETS	not available	MA082/03602	SANOFI MALTA LTD	MT
TELFAST 180 MG FILM COATED TABLETS	not available	MA082/03602	SANOFI MALTA LTD	MT
TELFAST 180 MG FILM COATED TABLETS	not available	MA082/03602	SANOFI MALTA LTD	MT
TELFAST 180 MG FILM COATED TABLETS	not available	MA082/03602	SANOFI MALTA LTD	MT