29 May 2019
EMA/397212/2019
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: levofloxacin (except for the centrally authorised product)

Procedure no.: PSUSA/00001854/201810

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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
Levomer, 5 mg/ml, krople do oczu, roztwór	PT/H/1495/001/DC	23616	ADAMED	PL
LEXAVON 5 mg/ml οφθαλμικές σταγόνες, διάλυμα	PT/H/0821/001	82071/28-09-2017	RAFARM SA.	GR
Lexavon eye drops solution/Rafarm 5mg/mL	PT/H/821/01/DC	021948	RAFARM SA.	CY
Levofloxacina Rafarm 5 mg/ml colírio, solução	PT/H/0821/001	5601257	RAFARM SA.	PT
Oftaquix 5 mg/ml - silmätipat	MRP UK/H/464/001	16992	SANTEN OY	FI
Oftaquix 5 mg/ml ögondroppar	MRP UK/H/464/001	16992	SANTEN OY	FI
Oftaquix 5 mg/ml ögondroppar, lösning	FI/H/0989/001	17994	SANTEN OY	SE
Oftaquix, øjendråber, opløsning	UK/H/0464/001	33160	SANTEN OY	DK
Oftaquix 5 mg/ml oční kapky, roztok	UK/H/0464/001	64/175/05-C	SANTEN OY	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
Oftaquix 5 mg/ml augndropar, lausn	FI/H/0989/001	IS/1/02/003/01	SANTEN OY	IS
Oftaquix 5 mg/ml silmatilgad, lahus	UK/H/0464/001	472305	SANTEN OY	EE
Oftaquix 5 mg/ml Augentropfen, Lösung	UK/H/0464/001	53538.00.00	SANTEN OY	DE
Oftaquix 5 mg/ml oldatos szemcsepp	MRP UK/H/464/001	OGYI-T-10257/01	SANTEN OY	HU
Oftaquix 5 mg/ml collirio, soluzione	FI/H/0989/001	035728017	SANTEN OY	IT
Oftaquix 5 mg/ml acu pilieni, šķīdums	FI/H/0989/001	05-0196	SANTEN OY	LV
OFTAQUIX 5 mg/ml akių lašai (tirpalas)	FI/H/0989/001	LT/1/05/0325/001	SANTEN OY	LT
Oftaquix 5 mg/ml colírio	MRP FI/H/989/01	4041786	SANTEN OY	PT
Oftaquix 5 mg/ml očná roztoková instilácia	FI/H/0989/001	64/0180/05-S	SANTEN OY	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
Oftaquix 5 mg/ml eye drops	FI/H/0989/001	PL 16058/0006	SANTEN OY	UK
Oftaquix 5 mg/ml Augentropfen	FI/H/0989/001	NO.AMM 2002050011, NO. NAT. 0318046	SANTEN OY	LU
Oftaquix 5 mg/ml Augentropfen	MRP UK/H/464/001	1-24655	SANTEN OY	AT
OFTAQUIX 5 mg/ml picături oftalmice, soluţie	MRP FI/H/989/01	4609/2012/01	SANTEN OY	RO
FLOXISTILL 5 mg/ml collirio soluzione	PT/H/1360/001	044071013	BRUSCHETTINI S.R.L	IT
Levofloxacina Bruschettini 5 mg/ml colírio, solução	PT/H/1360/001/DC	5666540	BRUSCHETTINI S.R.L	PT
TAVANIC 5 MG/ML SOLUTION FOR INFUSION	DE/H/5119/003	19282	SANOFI-AVENTIS CYPRUS LTD	CY
TAVANIC 5 MG/ML SOLUTION FOR INFUSION	DE/H/5119/003	19282	SANOFI-AVENTIS CYPRUS LTD	CY
TAVANIC 5 MG/ML SOLUTION FOR INFUSION	DE/H/5119/003	19282	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
TAVANIC 5 MG/ML SOLUTION FOR INFUSION	DE/H/5119/003	19282	SANOFI-AVENTIS CYPRUS LTD	CY
TAVANIC 5 MG/ML SOLUTION FOR INFUSION	DE/H/5119/003	19282	SANOFI-AVENTIS CYPRUS LTD	CY
TAVANIC 5 MG/ML, SOLUTION POUR PERFUSION	DE/H/5119/003	34009 561 899 2 2	SANOFI-AVENTIS FRANCE	FR
TAVANIC 500 MG FILMDRAGERAD TABLETT	DE/H/5119/002	14136	SANOFI AB	SE
TAVANIC 500 MG FILMDRAGERAD TABLETT	DE/H/5119/002	14136	SANOFI AB	SE
TAVANIC 500 MG FILMDRAGERAD TABLETT	DE/H/5119/002	14136	SANOFI AB	SE
TAVANIC 500 MG, COMPRIME	DE/H/5119/002	34009 349 655 6 2	SANOFI-AVENTIS FRANCE	FR
TAVANIC 500 MG FILMDRAGERAD TABLETT	DE/H/5119/002	14136	SANOFI AB	SE
TAVANIC 500 MG FILMDRAGERAD TABLETT	DE/H/5119/002	14136	SANOFI AB	SE

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
TAVANIC 500 MG FILMDRAGERAD TABLETT	DE/H/5119/002	14136	SANOFI AB	SE
Tavanic 250 mg plėvele dengtos tabletės	DE/H/5119/001	LT R 02/7796/6	UAB SANOFI-AVENTIS LIETUVA	LT
TAVANIC 500 MG FILMDRAGERAD TABLETT	DE/H/5119/002	14136	SANOFI AB	SE
TAVANIC	DE/H/5119/003	8766	SANOFI-AVENTIS DEUTSCHLAND GMBH	PL
TAVANIC 5 MG/ML, SOLUTION POUR PERFUSION	DE/H/5119/003	34009 561 898 6 1	SANOFI-AVENTIS FRANCE	FR
TAVANIC 5 MG/ML, SOLUTION POUR PERFUSION	DE/H/5119/003	34009561 901 7 1	SANOFI-AVENTIS FRANCE	FR
TAVANIC 5 MG/ML, SOLUTION POUR PERFUSION	DE/H/5119/003	34009 561 900 0 3	SANOFI-AVENTIS FRANCE	FR
TAVANIC 500 MG, COMPRIME	DE/H/5119/002	34009 349 656 2 3	SANOFI-AVENTIS FRANCE	FR
TAVANIC 500 MG, COMPRIME	DE/H/5119/002	34009 561 904 6 1	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
TAVANIC 500 MG, COMPRIME	DE/H/5119/002	34009 348 996 4 5	SANOFI-AVENTIS FRANCE	FR
TAVANIC 500 MG, COMPRIME	DE/H/5119/002	34009 574 907 9 9	SANOFI-AVENTIS FRANCE	FR
Tavanic 500 mg plėvele dengtos tabletės	DE/H/5119/002	LT R 02/7797/6	UAB SANOFI-AVENTIS LIETUVA	LT
Tavanic 500 mg plėvele dengtos tabletės	DE/H/5119/002	LT R 02/7797/6	UAB SANOFI-AVENTIS LIETUVA	LT
Tavanic 250 mg plėvele dengtos tabletės	DE/H/5119/001	LT R 02/7796/6	UAB SANOFI-AVENTIS LIETUVA	LT
Tavanic 500 mg plėvele dengtos tabletės	DE/H/5119/002	LT R 02/7797/6	UAB SANOFI-AVENTIS LIETUVA	LT
Tavanic 500 mg plėvele dengtos tabletės	DE/H/5119/002	LT R 02/7797/6	UAB SANOFI-AVENTIS LIETUVA	LT
Tavanic 500 mg plėvele dengtos tabletės	DE/H/5119/002	LT R 02/7797/6	UAB SANOFI-AVENTIS LIETUVA	LT
Tavanic 250 mg plėvele dengtos tabletės	DE/H/5119/001	LT R 02/7796/6	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
Tavanic 250 mg plėvele dengtos tabletės	DE/H/5119/001	LT R 02/7796/6	UAB SANOFI-AVENTIS LIETUVA	LT
Tavanic 250 mg plėvele dengtos tabletės	DE/H/5119/001	LT R 02/7796/6	UAB SANOFI-AVENTIS LIETUVA	LT
Tavanic 500 mg plėvele dengtos tabletės	DE/H/5119/002	LT R 02/7797/6	UAB SANOFI-AVENTIS LIETUVA	LT
Tavanic 500 mg plėvele dengtos tabletės	DE/H/5119/002	LT R 02/7797/6	UAB SANOFI-AVENTIS LIETUVA	LT
Tavanic 250 mg plėvele dengtos tabletės	DE/H/5119/001	LT R 02/7796/6	UAB SANOFI-AVENTIS LIETUVA	LT
Tavanic 250 mg plėvele dengtos tabletės	DE/H/5119/001	LT R 02/7796/6	UAB SANOFI-AVENTIS LIETUVA	LT
TAVANIC 5 MG/ML RAZTOPINA ZA INFUNDIRANJE	DE/H/5119/003	H/00/01496/015	SANOFI-AVENTIS D.O.O.	SI
TAVANIC 500 MG FILMSKO OBLOZENE TABLETE	DE/H/5119/002	H/00/01496/008	SANOFI-AVENTIS D.O.O.	SI
TAVANIC 500 MG FILMTABLETTA	DE/H/5119/002	OGYI-T-6837/07	SANOFI-AVENTIS ZRT	HU

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
TAVANIC 500 MG FILMTABLETTA	DE/H/5119/002	OGYI-T-6837/14	SANOFI-AVENTIS ZRT	HU
TAVANIC 500 MG FILMTABLETTA	DE/H/5119/002	OGYI-T-6837/06	SANOFI-AVENTIS ZRT	HU
TAVANIC	DE/H/5119/002	6160/17-02-99	SANOFI-AVENTIS AEBE	GR
TAVANIC	DE/H/5119/003	6161/17-02-99	SANOFI-AVENTIS AEBE	GR
TAVANIC	DE/H/5119/002	6160/17-02-99	SANOFI-AVENTIS AEBE	GR
TAVANIC	DE/H/5119/002	6160/17-02-99	SANOFI-AVENTIS AEBE	GR
TAVANIC	DE/H/5119/002	6160/17-02-99	SANOFI-AVENTIS AEBE	GR
Таваник 500mg филмирани таблетки	DE/H/5119/002	20020443	SANOFI BULGARIA EOOD	BG
Tavanic, 250 mg õhukese polümeerikattega tabletid	DE/H/5119/001	424203	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
TAVANIC, 5 MG/ML INFUSIOONILAHUS	DE/H/5119/003	424003	SANOFI-AVENTIS ESTONIA OÜ	EE
Таваник 500mg филмирани таблетки	DE/H/5119/002	20020443	SANOFI BULGARIA EOOD	BG
Таваник 500mg филмирани таблетки	DE/H/5119/002	20020443	SANOFI BULGARIA EOOD	BG
Таваник 500mg филмирани таблетки	DE/H/5119/002	20020443	SANOFI BULGARIA EOOD	BG
Таваник 500mg филмирани таблетки	DE/H/5119/002	20020443	SANOFI BULGARIA EOOD	BG
Таваник 500mg филмирани таблетки	DE/H/5119/002	20020443	SANOFI BULGARIA EOOD	BG
Таваник 500mg филмирани таблетки	DE/H/5119/002	20020443	SANOFI BULGARIA EOOD	BG
TAVANIC, 500 MG OHUKESE POLUMEERIKATTEGA TABLETID	DE/H/5119/002	424103	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
TAVANIC, 250 MG OHUKESE POLUMEERIKATTEGA TABLETID	DE/H/5119/001	424203	SANOFI-AVENTIS ESTONIA OÜ	EE
TAVANIC, 250 MG OHUKESE POLUMEERIKATTEGA TABLETID	DE/H/5119/001	424203	SANOFI-AVENTIS ESTONIA OÜ	EE
TAVANIC, 250 MG OHUKESE POLUMEERIKATTEGA TABLETID	DE/H/5119/001	424203	SANOFI-AVENTIS ESTONIA OÜ	EE
TAVANIC, 250 MG OHUKESE POLUMEERIKATTEGA TABLETID	DE/H/5119/001	424203	SANOFI-AVENTIS ESTONIA OÜ	EE
TAVANIC, 250 MG OHUKESE POLUMEERIKATTEGA TABLETID	DE/H/5119/001	424203	SANOFI-AVENTIS ESTONIA OÜ	EE
TAVANIC, 500 MG OHUKESE POLUMEERIKATTEGA TABLETID	DE/H/5119/002	424103	SANOFI-AVENTIS ESTONIA OÜ	EE
TAVANIC, 500 MG OHUKESE POLUMEERIKATTEGA TABLETID	DE/H/5119/002	424103	SANOFI-AVENTIS ESTONIA OÜ	EE
TAVANIC, 500 MG OHUKESE POLUMEERIKATTEGA TABLETID	DE/H/5119/002	424103	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
TAVANIC, 500 MG OHUKESE POLUMEERIKATTEGA TABLETID	DE/H/5119/002	424103	SANOFI-AVENTIS ESTONIA OÜ	EE
TAVANIC, 250 MG OHUKESE POLUMEERIKATTEGA TABLETID	DE/H/5119/001	424203	SANOFI-AVENTIS ESTONIA OÜ	EE
TAVANIC, 500 MG OHUKESE POLUMEERIKATTEGA TABLETID	DE/H/5119/002	424103	SANOFI-AVENTIS ESTONIA OÜ	EE
TAVANIC, 500 MG OHUKESE POLUMEERIKATTEGA TABLETID	DE/H/5119/002	424103	SANOFI-AVENTIS ESTONIA OÜ	EE
TAVANIC, 5 MG/ML INFUSIOONILAHUS	DE/H/5119/003	424003	SANOFI-AVENTIS ESTONIA OÜ	EE
TAVANIC, 5 MG/ML INFUSIOONILAHUS	DE/H/5119/003	424003	SANOFI-AVENTIS ESTONIA OÜ	EE
TAVANIC, 5 MG/ML INFUSIOONILAHUS	DE/H/5119/003	424003	SANOFI-AVENTIS ESTONIA OÜ	EE
TAVANIC, 5 MG/ML INFUSIOONILAHUS	DE/H/5119/003	424003	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
TAVANIC	DE/H/5119/002	19283	SANOFI-AVENTIS CYPRUS LTD	CY
TAVANIC	DE/H/5119/002	19283	SANOFI-AVENTIS CYPRUS LTD	CY
TAVANIC	DE/H/5119/002	19283	SANOFI-AVENTIS CYPRUS LTD	CY
TAVANIC	DE/H/5119/002	19283	SANOFI-AVENTIS CYPRUS LTD	CY
TAVANIC	DE/H/5119/002	19283	SANOFI-AVENTIS CYPRUS LTD	CY
TAVANIC	DE/H/5119/002	19283	SANOFI-AVENTIS CYPRUS LTD	CY
TAVANIC	DE/H/5119/002	19283	SANOFI-AVENTIS CYPRUS LTD	CY
TAVANIC 250 MG FILMDRAGERAD TABLETT	DE/H/5119/001	14135	SANOFI AB	SE
TAVANIC 250 MG FILMDRAGERAD TABLETT	DE/H/5119/001	14135	SANOFI AB	SE

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
TAVANIC 250 MG FILMDRAGERAD TABLETT	DE/H/5119/001	14135	SANOFI AB	SE
TAVANIC 250 MG FILMDRAGERAD TABLETT	DE/H/5119/001	14135	SANOFI AB	SE
TAVANIC 250 MG FILMDRAGERAD TABLETT	DE/H/5119/001	14135	SANOFI AB	SE
TAVANIC 250 MG FILMDRAGERAD TABLETT	DE/H/5119/001	14135	SANOFI AB	SE
TAVANIC 250 MG FILMDRAGERAD TABLETT	DE/H/5119/001	14135	SANOFI AB	SE
TAVANIC 5 MG/ML SOLUTION FOR INFUSION	DE/H/5119/003	14134	SANOFI AB	SE
TAVANIC 5 MG/ML SOLUTION FOR INFUSION	DE/H/5119/003	14134	SANOFI AB	SE
TAVANIC 5 MG/ML SOLUTION FOR INFUSION	DE/H/5119/003	14134	SANOFI AB	SE
TAVANIC 5 MG/ML SOLUTION FOR INFUSION	DE/H/5119/003	14134	SANOFI AB	SE

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
TAVANIC 5 MG/ML SOLUTION FOR INFUSION	DE/H/5119/003	14134	SANOFI AB	SE
TAVANIC 500 MG FILM-COATED TABLETS	DE/H/5119/002	MA 082/03502	SANOFI MALTA LTD	MT
TAVANIC 500 MG FILM-COATED TABLETS	DE/H/5119/002	MA 082/03502	SANOFI MALTA LTD	MT
TAVANIC 500 MG FILM-COATED TABLETS	DE/H/5119/002	MA 082/03502	SANOFI MALTA LTD	MT
TAVANIC 500 MG FILM-COATED TABLETS	DE/H/5119/002	MA 082/03502	SANOFI MALTA LTD	MT
TAVANIC 500 MG FILM-COATED TABLETS	DE/H/5119/002	MA 082/03502	SANOFI MALTA LTD	MT
TAVANIC 500 MG FILM-COATED TABLETS	DE/H/5119/002	MA 082/03502	SANOFI MALTA LTD	MT
TAVANIC 500 MG FILM-COATED TABLETS	DE/H/5119/002	MA 082/03502	SANOFI MALTA LTD	MT
TAVANIC I.V.	DE/H/5119/003	42/176/01-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
TAVANIC I.V.	DE/H/5119/003	42/176/01-C	SANOFI-AVENTIS SRO	CZ
TAVANIC I.V.	DE/H/5119/003	42/176/01-C	SANOFI-AVENTIS SRO	CZ
TAVANIC I.V.	DE/H/5119/003	42/176/01-C	SANOFI-AVENTIS SRO	CZ
TAVANIC I.V.	DE/H/5119/003	42/176/01-C	SANOFI-AVENTIS SRO	CZ
Tavanic 5 mg/ml solution pour perfusion	DE/H/5119/003	BE192875	SANOFI BELGIUM	BE
Tavanic 5 mg/ml solution pour perfusion	DE/H/5119/003	BE230237	SANOFI BELGIUM	BE
Tavanic 5 mg/ml solution pour perfusion	DE/H/5119/003	BE192875	SANOFI BELGIUM	BE
Tavanic 5 mg/ml solution pour perfusion	DE/H/5119/003	BE192875	SANOFI BELGIUM	BE
Tavanic 5 mg/ml solution pour perfusion	DE/H/5119/003	BE230237	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
TAVANIC 500 mg comprimidos revestidos por película	DE/H/5119/002	3088382	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
TAVANIC 500 mg comprimidos revestidos por película	DE/H/5119/002	2628584	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Tavanic 500 mg film-coated tablets	DE/H/5119/002	PA 540/77/3	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
TAVANIC 500 mg comprimidos revestidos por película	DE/H/5119/002	3088481	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
TAVANIC 500 mg comprimidos revestidos por película	DE/H/5119/002	2628881	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
TAVANIC 500 mg comprimidos revestidos por película	DE/H/5119/002	2628485	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Tavanic 250 mg kalvopäällysteiset tabletit	DE/H/5119/001	12946	SANOFI OY	FI
TAVANIC 500 mg comprimidos revestidos por película	DE/H/5119/002	3088283	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Tavanic 250 mg kalvopäällysteiset tabletit	DE/H/5119/001	12946	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
Tavanic 500 mg film-coated tablets	DE/H/5119/002	PA 540/77/3	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
TAVANIC 500 mg comprimidos revestidos por película	DE/H/5119/002	2628683	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Tavanic 500 mg kalvopäällysteiset tabletit	DE/H/5119/002	12947	SANOFI OY	FI
Tavanic Infusionsflasche	DE/H/5119/003	1-22321	SANOFI-AVENTIS GMBH OSTERREICH	AT
Tavanic 500 mg film-coated tablets	DE/H/5119/002	PA 540/77/3	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
TAVANIC 5 mg/ml solução para perfusão	DE/H/5119/003	3088580	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Tavanic 500 mg kalvopäällysteiset tabletit	DE/H/5119/002	12947	SANOFI OY	FI
Tavanic® 500 mg Filmtabletten	DE/H/5119/002	1-22320	SANOFI-AVENTIS GMBH OSTERREICH	AT
Tavanic Infusionsflasche	DE/H/5119/003	1-22321	SANOFI-AVENTIS GMBH OSTERREICH	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
Tavanic® 500 mg Filmtabletten	DE/H/5119/002	1-22320	SANOFI-AVENTIS GMBH OSTERREICH	AT
Tavanic 500 mg kalvopäällysteiset tabletit	DE/H/5119/002	12947	SANOFI OY	FI
Tavanic® 500 mg Filmtabletten	DE/H/5119/002	1-22320	SANOFI-AVENTIS GMBH OSTERREICH	AT
Tavanic® 500 mg Filmtabletten	DE/H/5119/002	1-22320	SANOFI-AVENTIS GMBH OSTERREICH	AT
Tavanic Infusionsflasche	DE/H/5119/003	1-22321	SANOFI-AVENTIS GMBH OSTERREICH	AT
TAVANIC 5 mg/ml solução para perfusão	DE/H/5119/003	2628980	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Tavanic 500 mg kalvopäällysteiset tabletit	DE/H/5119/002	12947	SANOFI OY	FI
Tavanic Infusionsflasche	DE/H/5119/003	1-22321	SANOFI-AVENTIS GMBH OSTERREICH	AT
Tavanic® 500 mg Filmtabletten	DE/H/5119/002	1-22320	SANOFI-AVENTIS GMBH OSTERREICH	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
Tavanic 500 mg kalvopäällysteiset tabletit	DE/H/5119/002	12947	SANOFI OY	FI
TAVANIC 5 mg/ml solução para perfusão	DE/H/5119/003	3088788	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Tavanic® 500 mg Filmtabletten	DE/H/5119/002	1-22320	SANOFI-AVENTIS GMBH OSTERREICH	AT
Tavanic® 500 mg Filmtabletten	DE/H/5119/002	1-22320	SANOFI-AVENTIS GMBH OSTERREICH	AT
Tavanic 5 mg/ml solution for infusion	DE/H/5119/003	PA 540/77/001	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
Tavanic Infusionsflasche	DE/H/5119/003	1-22321	SANOFI-AVENTIS GMBH OSTERREICH	AT
Tavanic® 250 mg Filmtabletten	DE/H/5119/001	1-22319	SANOFI-AVENTIS GMBH OSTERREICH	AT
Tavanic 500 mg kalvopäällysteiset tabletit	DE/H/5119/002	12947	SANOFI OY	FI
TAVANIC 5 mg/ml solução para perfusão	DE/H/5119/003	3088689	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
TAVANIC 500 MG COMPRESSE RIVESTITE CON FILM	DE/H/5119/002	033634054	SANOFI S.P.A	IT
Tavanic® 250 mg Filmtabletten	DE/H/5119/001	1-22319	SANOFI-AVENTIS GMBH OSTERREICH	AT
Tavanic 5 mg/ml oplossing voor infusie	DE/H/5119/003	BE192875	SANOFI BELGIUM	BE
Tavanic 500 mg film-coated tablets	DE/H/5119/002	PA 540/77/3	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
Tavanic 500 mg filmomhulde tabletten	DE/H/5119/002	BE192884	SANOFI BELGIUM	BE
Tavanic 500 mg filmomhulde tabletten	DE/H/5119/002	BE192884	SANOFI BELGIUM	BE
Tavanic 500 mg kalvopäällysteiset tabletit	DE/H/5119/002	12947	SANOFI OY	FI
Tavanic 250 mg comprimé pelliculé	DE/H/5119/001	0247087	SANOFI BELGIUM	LU
Tavanic 5 mg/ml oplossing voor infusie	DE/H/5119/003	BE192875	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
Tavanic 500 mg film-coated tablets	DE/H/5119/002	PA 540/77/3	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
Tavanic 5 mg/ml solución para perfusión	DE/H/5119/003	62066	SANOFI-AVENTIS, S.A.	ES
Tavanic 5 mg/ml solution for infusion	DE/H/5119/003	PA 540/77/001	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
Tavanic 250 mg comprimé pelliculé	DE/H/5119/001	0247056	SANOFI BELGIUM	LU
Tavanic 500 mg film-coated tablets	DE/H/5119/002	PA 540/77/3	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
Tavanic® 250 mg Filmtabletten	DE/H/5119/001	1-22319	SANOFI-AVENTIS GMBH OSTERREICH	AT
TAVANIC 500 MG COMPRESSE RIVESTITE CON FILM	DE/H/5119/002	033634041	SANOFI S.P.A	IT
Tavanic® 250 mg Filmtabletten	DE/H/5119/001	1-22319	SANOFI-AVENTIS GMBH OSTERREICH	AT
Tavanic 500 mg film-coated tablets	DE/H/5119/002	PA 540/77/3	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	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
Tavanic® 250 mg Filmtabletten	DE/H/5119/001	1-22319	SANOFI-AVENTIS GMBH OSTERREICH	AT
Tavanic® 250 mg Filmtabletten	DE/H/5119/001	1-22319	SANOFI-AVENTIS GMBH OSTERREICH	AT
Tavanic 250 mg Filmtabletten	DE/H/5119/001	1-22319	SANOFI-AVENTIS GMBH OSTERREICH	AT
Tavanic 250 mg comprimé pelliculé	DE/H/5119/001	0247073	SANOFI BELGIUM	LU
Tavanic 250 mg comprimé pelliculé	DE/H/5119/001	0247915	SANOFI BELGIUM	LU
Tavanic IV, oplossing voor infusie 5 mg/ml	DE/H/5119/003	RVG 21810	SANOFI-AVENTIS NETHERLANDS B.V.	NL
TAVANIC 500 MG COMPRESSE RIVESTITE CON FILM	DE/H/5119/002	033634039	SANOFI S.P.A	IT
TAVANIC 5 mg/ml solução para perfusão	DE/H/5119/003	2978682	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Tavanic IV, oplossing voor infusie 5 mg/ml	DE/H/5119/003	RVG 21810	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
Tavanic 250 mg kalvopäällysteiset tabletit	DE/H/5119/001	12946	SANOFI OY	FI
Tavanic 5 mg/ml oplossing voor infusie	DE/H/5119/003	BE230237	SANOFI BELGIUM	BE
Tavanic 250 mg film-coated tablets	DE/H/5119/001	PA 540/77/2	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
TAVANIC 250 MG FILM-COATED TABLETS	DE/H/5119/001	PA 540/77/2	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
Tavanic 250 mg kalvopäällysteiset tabletit	DE/H/5119/001	12946	SANOFI OY	FI
TAVANIC 250 MG FILM-COATED TABLETS	DE/H/5119/001	PA 540/77/2	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
TAVANIC 250 MG FILM-COATED TABLETS	DE/H/5119/001	PA 540/77/2	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
Tavanic 500 mg filmomhulde tabletten	DE/H/5119/002	BE192884	SANOFI BELGIUM	BE
Tavanic 500 mg filmomhulde tabletten	DE/H/5119/002	BE192884	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
TAVANIC 250 MG FILM-COATED TABLETS	DE/H/5119/001	PA 540/77/2	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
Tavanic 250 mg kalvopäällysteiset tabletit	DE/H/5119/001	12946	SANOFI OY	FI
Tavanic 500 mg filmomhulde tabletten	DE/H/5119/002	BE192884	SANOFI BELGIUM	BE
Tavanic 250 mg kalvopäällysteiset tabletit	DE/H/5119/001	12946	SANOFI OY	FI
Tavanic 250 mg kalvopäällysteiset tabletit	DE/H/5119/001	12946	SANOFI OY	FI
TAVANIC 250 MG FILM-COATED TABLETS	DE/H/5119/001	PA 540/77/2	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
Tavanic 500 mg filmomhulde tabletten	DE/H/5119/002	BE192884	SANOFI BELGIUM	BE
TAVANIC 250 MG FILM-COATED TABLETS	DE/H/5119/001	PA 540/77/2	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
Tavanic 5 mg/ml solution pour perfusion	DE/H/5119/003	0247141	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
Tavanic 5 mg/ml Infusionslösung	DE/H/5119/003	41384.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Tavanic 5 mg/ml Infusionslösung	DE/H/5119/003	41384.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Tavanic 500 mg comprimé pelliculé	DE/H/5119/002	0247137	SANOFI BELGIUM	LU
Tavanic 5 mg/ml Infusionslösung	DE/H/5119/003	41384.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Tavanic 500 mg comprimé pelliculé	DE/H/5119/002	0247106	SANOFI BELGIUM	LU
Tavanic 5 mg/ml solution pour perfusion	DE/H/5119/003	0305701	SANOFI BELGIUM	LU
Tavanic 5 mg/ml Infusionslösung	DE/H/5119/003	41384.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Tavanic 500 mg comprimé pelliculé	DE/H/5119/002	0247091	SANOFI BELGIUM	LU
Tavanic 500 mg comprimé pelliculé	DE/H/5119/002	0247123	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
Tavanic® 250 mg Filmtabletten	DE/H/5119/001	41382.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Tavanic® 250 mg Filmtabletten	DE/H/5119/001	41382.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Tavanic® 250 mg Filmtabletten	DE/H/5119/001	41382.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Tavanic® 250 mg Filmtabletten	DE/H/5119/001	41382.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Tavanic® 250 mg Filmtabletten	DE/H/5119/001	41382.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
TAVANIC 250 mg comprimidos revestidos por película	DE/H/5119/001	2628386	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
TAVANIC 250 mg comprimidos revestidos por película	DE/H/5119/001	2627982	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
TAVANIC 5 MG/ML SOLUZIONE PER INFUSIONE	DE/H/5119/003	033634080	SANOFI S.P.A	IT
TAVANIC 5 MG/ML SOLUZIONE PER INFUSIONE	DE/H/5119/003	033634066	SANOFI 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
TAVANIC 250 MG	DE/H/5119/001	033634015	SANOFI S.P.A	IT
Tavanic 250 mg filmomhulde tabletten	DE/H/5119/001	BE192866	SANOFI BELGIUM	BE
Tavanic® 500 mg Filmtabletten	DE/H/5119/002	41382.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Tavanic 250 mg filmomhulde tabletten	DE/H/5119/001	BE192866	SANOFI BELGIUM	BE
Tavanic 5 mg/ml oplossing voor infusie	DE/H/5119/003	BE192875	SANOFI BELGIUM	BE
TAVANIC 5 MG/ML INFUUSIONESTE, LIUOS	DE/H/5119/003	12948	SANOFI OY	FI
TAVANIC 5 MG/ML INFUUSIONESTE, LIUOS	DE/H/5119/003	12948	SANOFI OY	FI
TAVANIC 5 MG/ML INFUUSIONESTE, LIUOS	DE/H/5119/003	12948	SANOFI OY	FI
Tavanic 250 mg compresse rivestite con film	DE/H/5119/001	033634027	SANOFI 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
Tavanic 250 mg filmomhulde tabletten	DE/H/5119/001	BE192866	SANOFI BELGIUM	BE
TAVANIC 250 mg comprimidos revestidos por película	DE/H/5119/001	2628188	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Tavanic 500 mg comprimidos recubiertos con película	DE/H/5119/002	62065	SANOFI-AVENTIS, S.A.	ES
Tavanic 250 mg filmomhulde tabletten	DE/H/5119/001	BE192866	SANOFI BELGIUM	BE
Tavanic® 500 mg Filmtabletten	DE/H/5119/002	41382.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Tavanic 500 mg comprimidos recubiertos con película	DE/H/5119/002	62065	SANOFI-AVENTIS, S.A.	ES
TAVANIC 5 MG/ML SOLUZIONE PER INFUSIONE	DE/H/5119/003	033634078	SANOFI S.P.A	IT
Tavanic 250 mg filmomhulde tabletten	DE/H/5119/001	BE192866	SANOFI BELGIUM	BE
TAVANIC 250 mg comprimidos revestidos por película	DE/H/5119/001	3088085	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
Tavanic® 500 mg Filmtabletten	DE/H/5119/002	41382.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Tavanic® 500 mg Filmtabletten	DE/H/5119/002	41382.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Tavanic 250 mg filmomhulde tabletten	DE/H/5119/001	BE192866	SANOFI BELGIUM	BE
Tavanic 250 mg filmomhulde tabletten	DE/H/5119/001	BE192866	SANOFI BELGIUM	BE
TAVANIC 250 mg comprimidos revestidos por película	DE/H/5119/001	2628089	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
TAVANIC 250 mg comprimidos revestidos por película	DE/H/5119/001	3484680	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
TAVANIC 250 mg comprimidos revestidos por película	DE/H/5119/001	3088184	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
TAVANIC 5 MG/ML INFUUSIONESTE, LIUOS	DE/H/5119/003	12948	SANOFI OY	FI
Tavanic 5 mg/ml oplossing voor infusie	DE/H/5119/003	BE230237	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
TAVANIC 5 MG/ML INFUUSIONESTE, LIUOS	DE/H/5119/003	12948	SANOFI OY	FI
Tavanic 500 mg compresse rivestite con film	DE/H/5119/002	033634092	SANOFI S.P.A	IT
Tavanic 500 mg comprimidos recubiertos con película	DE/H/5119/002	62065	SANOFI-AVENTIS, S.A.	ES
Visuflox 5 mg/ml collirio, soluzione	PT/H/1495/001	044459016	VISUFARMA SPA	IT
Levofloxacin Actavis 5 mg/ml ögondroppar, lösning	SE/H/1715/001	52248	ACTAVIS GROUP PTC EHF.	SE
Levofloxacin Actavis 5 mg/ml silmatilgad, lahus	SE/H/1715/001	905516	ACTAVIS GROUP PTC EHF.	EE
Levofloxacin Actavis 5 mg/ml akių lašai (tirpalas)	SE/H/1715/001	LT/1/16/3901/001	ACTAVIS GROUP PTC EHF.	LT
Levofloxacină Actavis 5 mg/ml picături oftalmice, soluţie	UK/H/5960/001	8907/2016/01	ACTAVIS GROUP PTC EHF.	RO
Levofloxacin Actavis 5 mg/ml acu pilieni, šķīdums	SE/H/1715/001	16-0090	ACTAVIS GROUP PTC EHF.	LV

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
Levofloxacin Actavis 5 mg/ml –silmätipat, liuos	SE/H/1715/001	32926	ACTAVIS GROUP PTC EHF.	FI
OFTAQUIX, 5 mg/ml, krople do oczu, roztwór	not available	10825	SANTEN OY	PL
Oftaquix sine, 5 mg/ml, krople do oczu, roztwór w pojemniku jednodawkowym	not available	17128	SANTEN OY	PL
Levofloxacina Velka Hellas 5 mg/ml colírio, solução	PT/H/1495/001	5691118	VELKA HELLAS S.A.	PT
Oftaquix 5 mg/ml silmätipat, liuos, kerta-annospakkaus	MRP UK/H/0464/002	23157	SANTEN OY	FI
Oftaquix 5 mg/ml ögondroppar, lösning i endosbehållare	MRP UK/H/0464/002	23157	SANTEN OY	FI
Oftaquix 5 mg/ml ögondroppar, lösning i endosbehållare	FI/H/0989/002	25042	SANTEN OY	SE
Oftaquix, øjendråber, opløsning i enkeltdosisbeholder	MRP UK/H/0464/002	40612	SANTEN OY	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
Oftaquix 5 mg/ml augndropar, lausn stakskammtaílát	FI/H/0989/002	IS/1/07/020/01	SANTEN OY	IS
Oftaquix Unit Dose 5 mg/ml eye drops, solution in single-dose container	FI/H/0989/002	PL 16058/0007	SANTEN OY	UK
Oftaquix sine 5 mg/ml Augentropfen, Lösung im Einzeldosisbehältnis	MRP UK/H/0464/002	67435.00.00	SANTEN OY	DE
Oftaquix 5 mg/ml collirio, soluzione, contenitore monodose	FI/H/0989/002	035728031	SANTEN OY	IT
Oftaquix 5 mg/ml collirio, soluzione, contenitore monodose	FI/H/0989/002	035728029	SANTEN OY	IT
Oftaquix 5 mg/ml collirio, soluzione, contenitore monodose	FI/H/0989/002	035728043	SANTEN OY	IT
Oftaquix 5 mg/ml collirio, soluzione, contenitore monodose	FI/H/0989/002	035728056	SANTEN OY	IT
Офтакуикс 5 mg/ml очни капки Oftaquix 5 mg/ml eye drops	not available	20040266	SANTEN OY	BG

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
PRIXAR	not available	033633037	SANOFI SPA	IT
PRIXAR	not available	033633049	SANOFI SPA	IT
Prixar 250 mg compresse rivestite con film	not available	033633013	SANOFI SPA	IT
PRIXAR	not available	033633052	SANOFI SPA	IT
Prixar 250 mg compresse rivestite con film	not available	033633025	SANOFI SPA	IT
Levofloxacin UNIMED PHARMA 5 mg/ml silmatilgad, lahus	HU/H/0449/001	933917	UNIMED PHARMA SPOL.S R.O.	EE
Levofloxacin UNIMED PHARMA 5 mg/ml oldatos szemcsepp	HU/H/0449/001	OGYI-T-23155/01	UNIMED PHARMA SPOL.S R.O.	HU
Levofloxacin UNIMED PHARMA 5 mg/ml acu pilieni, šķīdums	HU/H/0449/001	17-0057	UNIMED PHARMA SPOL.S R.O.	LV
Levofloxacin UNIMED PHARMA 5 mg/ml akių lašai (tirpalas)	HU/H/0449/001	LT/1/17/4044/001	UNIMED PHARMA SPOL.S R.O.	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
Levofloxacin UNIMED PHARMA, 5 mg/ml, krople do oczu, roztwór	HU/H/0449/001	24081	UNIMED PHARMA SPOL.S R.O.	PL
Levofloksacin UNIMED PHARMA 5 mg/ml kapljice za oko, raztopina	HU/H/0449/001	H/17/02355/001	UNIMED PHARMA SPOL.S R.O.	SI
Levofloxacin 5mg/ml Eye Drops Solution	UK/H/5960/001	PL 0142/0946	ACTAVIS UK LTD.	UK
Levofloxacin 5mg/ml Eye Drops Solution	UK/H/5960/001	PL 0142/0946	ACTAVIS UK LTD.	UK
Levofloxacina NTC 5 mg/ml colírio	PT/H/1738/001	PT/H/1738/001	NTC SRL	PT
Levoxacin 250 mg compresse rivestite con film	not available	033940014	GLAXOSMITHKLINE S.P.A.	IT
Levoxacin 250 mg compresse rivestite con film	not available	033940026	GLAXOSMITHKLINE S.P.A.	IT
Levoxacin 500 mg compresse rivestite con film	not available	033940038	GLAXOSMITHKLINE S.P.A.	IT
Levoxacin 500 mg compresse rivestite con film	not available	033940040	GLAXOSMITHKLINE 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
Levoxacin 500 mg compresse rivestite con film	not available	033940053	GLAXOSMITHKLINE S.P.A.	IT
Levoxacin 5 mg/ml soluzione per infusione	not available	033940065	GLAXOSMITHKLINE S.P.A.	IT
Levodrop 5 mg/ml collirio, soluzione	PT/H/0822/001	042149017	ALFA INTES INDUSTRIA TERAPEUTICA SPLENDORE VIA FRATELLI	IT
Levofloxacina Optacare 5 mg/ml colírio	PT/H/0822/001	5593173	NTC SRL	PT
Levofloxacin 5 mg/ml eye drops solution	PT/H/1495/001	PL35533/0069	ASPIRE PHARMA LIMITED	UK
LEVOFLOXACIN/GENERICS 500 mg/100 ml, διάλυμα προς έγχυση	UK/H/5634/002	21707	MYLAN S.A.S	CY
Levofloxacine Mylan 500 mg/100 ml Infusionslösung	UK/H/5634/002	BE355625	MYLAN BVBA/SPRL	BE
Levofloxacine Mylan 500 mg/100 ml solution pour perfusion	UK/H/5634/002	BE355625	MYLAN BVBA/SPRL	BE
Levofloxacină Mylan 500 mg/100 ml soluţie perfuzabilă	UK/H/5634/002	5900/2013/01	MYLAN S.A.S	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
Levofloxacină Mylan 500 mg/100 ml soluţie perfuzabilă	UK/H/5634/002	5900/2013/02	MYLAN S.A.S	RO
Levofloxacină Mylan 500 mg/100 ml soluţie perfuzabilă	UK/H/5634/002	5900/2013/03	MYLAN S.A.S	RO
Levofloxacină Mylan 500 mg/100 ml soluţie perfuzabilă	UK/H/5634/002	5900/2013/04	MYLAN S.A.S	RO
Levofloxacină Mylan 500 mg/100 ml soluţie perfuzabilă	UK/H/5634/002	5900/2013/05	MYLAN S.A.S	RO
Levofloxacină Mylan 500 mg/100 ml soluţie perfuzabilă	UK/H/5634/002	5900/2013/06	MYLAN S.A.S	RO
Levofloxacină Mylan 500 mg/100 ml soluţie perfuzabilă	UK/H/5634/002	5900/2013/07	MYLAN S.A.S	RO
LEVOFLOXACIN/GENERICS 500 mg/100 ml, διάλυμα προς έγχυση	UK/H/5634/001	51161/13-07-2011	GENERICS PHARMA HELLAS LTD	GR
Levofloxacine Mylan 500 mg/100 ml oplossing voor infusie	UK/H/5634/002	BE355625	MYLAN BVBA/SPRL	BE
Levofloxacin Mylan 250 mg/50 ml, solution for infusion	UK/H/5634/001	MA767/00201	MYLAN S.A.S	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
Levofloxacin 250 mg/50 ml, solution for infusion	UK/H/5634/001	PL 04569/1010	GENERICS [UK] LIMITED	UK
Levofloxacin Arcana 500 mg/100 ml Infusionslösung	UK/H/5634/002	1-29240	ARCANA ARZNEIMITTEL GMBH	AT
Levofloxacina Mylan 500 mg/100 ml soluzione per infusione	UK/H/5634/002	040326011	MYLAN S.P.A.	IT
Levofloxacina Mylan 500 mg/100 ml soluzione per infusione	UK/H/5634/002	040326023	MYLAN S.P.A.	IT
Levofloxacina Mylan 500 mg/100 ml soluzione per infusione	UK/H/5634/002	040326035	MYLAN S.P.A.	IT
Levofloxacina Mylan 500 mg/100 ml soluzione per infusione	UK/H/5634/002	040326047	MYLAN S.P.A.	IT
Levofloxacina Mylan 500 mg/100 ml soluzione per infusione	UK/H/5634/002	040326050	MYLAN S.P.A.	IT
Levofloxacina Mylan 500 mg/100 ml soluzione per infusione	UK/H/5634/002	040326062	MYLAN S.P.A.	IT
Levofloxacina Mylan 500 mg/100 ml soluzione per infusione	UK/H/5634/002	040326074	MYLAN 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
Levofloxacina Mylan 5 mg/ mL solução para perfusão	UK/H/5634/002	5270624	MYLAN, LDA	PT
Levofloxacina Mylan 5 mg/ml solução para perfusão	UK/H/5634/002	5270632	MYLAN, LDA	PT
Levofloxacin Mylan 500 mg/100 ml infuzní roztok	UK/H/5634/002	42/894/09-C	MYLAN S.A.S	CZ
Levofloxacin Mylan 500 mg/100 ml, solution for infusion	UK/H/5634/002	MA767/00202	MYLAN S.A.S	MT
Levofloxacin 500 mg/100 ml, solution for infusion	UK/H/5634/002	PL 04569/1011	GENERICS [UK] LIMITED	UK
Levofloxacine Mylan 5 mg/ml, oplossing voor infusie	UK/H/5634/001	RVG 104110	MYLAN B.V.	NL
TAVANIC 250 mg comprimate filmate		942/2008/01-04	Terapia SA	RO
TAVANIC 500 mg comprimate filmate		943/2008/01-02-03	Terapia SA	RO
Tavanic i.v. 250 mg soluţie perfuzabilă		1016/2008/01	Terapia SA	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
Tavanic i.v. 500 mg soluţie perfuzabilă		1017/2008/01	Terapia SA	RO