26 November 2020
EMA/646224/2020
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: racecadotril

Procedure no.: PSUSA/00002602/202003

Official address Domenico Scarlattilaan 6 ● 1083 HS Amsterdam ● The Netherlands
Address for visits and deliveries Refer to www.ema.europa.eu/how-to-find-us
Send us a question Go to www.ema.europa.eu/contact **Telephone** +31 (0)88 781 6000 An agency of the European Union

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
Hidrasec 10 mg Granulat zur Herstellung einer Suspension zum Einnehmen	SE/H/1342/002	1-30818	BIOPROJET EUROPE LTD	AT
Hidrasec 10 mg Granulat zur Herstellung einer Suspension zum Einnehmen	SE/H/1342/002	1-30818	BIOPROJET EUROPE LTD	AT
Hidrasec 10 mg Granulat zur Herstellung einer Suspension zum Einnehmen	SE/H/1342/002	1-30818	BIOPROJET EUROPE LTD	AT
Hidrasec 10 mg Granulat zur Herstellung einer Suspension zum Einnehmen	SE/H/1342/002	1-30818	BIOPROJET EUROPE LTD	AT
Hidrasec 10 mg Granulat zur Herstellung einer Suspension zum Einnehmen	SE/H/1342/002	1-30818	BIOPROJET EUROPE LTD	AT
Hidrasec 30 mg Granulat zur Herstellung einer Suspension zum Einnehmen	SE/H/1342/001	1-30819	BIOPROJET EUROPE LTD	AT
Hidrasec 30 mg Granulat zur Herstellung einer Suspension zum Einnehmen	SE/H/1342/001	1-30819	BIOPROJET EUROPE LTD	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
Hidrasec 30 mg Granulat zur Herstellung einer Suspension zum Einnehmen	SE/H/1342/001	1-30819	BIOPROJET EUROPE LTD	AT
Hidrasec 30 mg Granulat zur Herstellung einer Suspension zum Einnehmen	SE/H/1342/001	1-30819	BIOPROJET EUROPE LTD	AT
Hidrasec 30 mg Granulat zur Herstellung einer Suspension zum Einnehmen	SE/H/1342/001	1-30819	BIOPROJET EUROPE LTD	AT
Hidrasec 100 mg Hartkapseln	SE/H/1342/003	1-30820	BIOPROJET EUROPE LTD	AT
Hidrasec 100 mg Hartkapseln	SE/H/1342/003	1-30820	BIOPROJET EUROPE LTD	AT
Hidrasec 100 mg Hartkapseln	SE/H/1342/003	1-30820	BIOPROJET EUROPE LTD	AT
Hidrasec 100 mg Hartkapseln	SE/H/1342/003	1-30820	BIOPROJET EUROPE LTD	AT
Vaprino® 100 mg Hartkapseln	AT/H/0542/001	136194	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
Vaprino® 100 mg Hartkapseln	AT/H/0542/001	136194	SANOFI-AVENTIS GMBH OSTERREICH	AT
Hidrasec 10 mg Granulat zur Herstellung einer Suspension zum Einnehmen	SE/H/1342/002	1-30818	BIOPROJET EUROPE LTD	AT
Hidrasec 30 mg Granulat zur Herstellung einer Suspension zum Einnehmen	SE/H/1342/001	1-30819	BIOPROJET EUROPE LTD	AT
Hidrasec 100 mg Hartkapseln	SE/H/1342/003	1-30820	BIOPROJET EUROPE LTD	AT
Tiorfix Baby 10 mg granulés pour suspension buvable	SE/H/1342/001	BE400723	BIOPROJET EUROPE LTD	BE
Tiorfix Baby 10 mg granulés pour suspension buvable	SE/H/1342/001	BE400723	BIOPROJET EUROPE LTD	BE
Tiorfix Baby 10 mg granulés pour suspension buvable	SE/H/1342/001	BE400723	BIOPROJET EUROPE LTD	BE
Tiorfix Baby 10 mg granulés pour suspension buvable	SE/H/1342/001	BE400723	BIOPROJET EUROPE LTD	BE
Tiorfix Baby 10 mg granulés pour suspension buvable	SE/H/1342/001	BE400723	BIOPROJET EUROPE LTD	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**
Tiorfix Junior 30 mg granulés pour suspension buvable	SE/H/1342/002	BE400732	BIOPROJET EUROPE LTD	BE
Tiorfix Junior 30 mg granulés pour suspension buvable	SE/H/1342/002	BE400732	BIOPROJET EUROPE LTD	BE
Tiorfix Junior 30 mg granulés pour suspension buvable	SE/H/1342/002	BE400732	BIOPROJET EUROPE LTD	BE
Tiorfix Junior 30 mg granulés pour suspension buvable	SE/H/1342/002	BE400732	BIOPROJET EUROPE LTD	BE
Tiorfix Junior 30 mg granulés pour suspension buvable	SE/H/1342/002	BE400732	BIOPROJET EUROPE LTD	BE
Tiorfix 100 mg gélules	SE/H/1342/003	BE400741	BIOPROJET EUROPE LTD	BE
Tiorfix 100 mg gélules	SE/H/1342/003	BE400741	BIOPROJET EUROPE LTD	BE
Tiorfix 100 mg gélules	SE/H/1342/003	BE400741	BIOPROJET EUROPE LTD	BE
Tiorfix 100 mg gélules	SE/H/1342/003	BE400741	BIOPROJET EUROPE LTD	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
Tiorfix Baby 10 mg, granulaat voor orale suspensie	SE/H/1342/001	BE400723	BIOPROJET EUROPE LTD	BE
Tiorfix Baby 10 mg, granulaat voor orale suspensie	SE/H/1342/001	BE400723	BIOPROJET EUROPE LTD	BE
Tiorfix Baby 10 mg, granulaat voor orale suspensie	SE/H/1342/001	BE400723	BIOPROJET EUROPE LTD	BE
Tiorfix Baby 10 mg, granulaat voor orale suspensie	SE/H/1342/001	BE400723	BIOPROJET EUROPE LTD	BE
Tiorfix Baby 10 mg, granulaat voor orale suspensie	SE/H/1342/001	BE400723	BIOPROJET EUROPE LTD	BE
Tiorfix Junior 30 mg granulaat voor orale suspensie	SE/H/1342/002	BE400732	BIOPROJET EUROPE LTD	BE
Tiorfix Junior 30 mg granulaat voor orale suspensie	SE/H/1342/002	BE400732	BIOPROJET EUROPE LTD	BE
Tiorfix Junior 30 mg granulaat voor orale suspensie	SE/H/1342/002	BE400732	BIOPROJET EUROPE LTD	BE
Tiorfix Junior 30 mg granulaat voor orale suspensie	SE/H/1342/002	BE400732	BIOPROJET EUROPE LTD	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
Tiorfix Junior 30 mg granulaat voor orale suspensie	SE/H/1342/002	BE400732	BIOPROJET EUROPE LTD	BE
Tiorfix 100 mg harde capsules	SE/H/1342/003	BE400741	BIOPROJET EUROPE LTD	BE
Tiorfix 100 mg harde capsules	SE/H/1342/003	BE400741	BIOPROJET EUROPE LTD	BE
Tiorfix 100 mg harde capsules	SE/H/1342/003	BE400741	BIOPROJET EUROPE LTD	BE
Tiorfix 100 mg harde capsules	SE/H/1342/003	BE400741	BIOPROJET EUROPE LTD	BE
Vaprino® 100 mg Hartkapseln	AT/H/0542/001	BE494160	SANOFI BELGIUM	BE
Vaprino 100 mg, gélules	AT/H/0542/001	BE494160	SANOFI BELGIUM	BE
Vaprino 100 mg harde capsules	AT/H/0542/001	BE494160	SANOFI BELGIUM	BE
Tiorfix 175 mg comprimés pelliculés	FR/H/0623/001	BE534497	BIOPROJET PHARMA	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
Tiorfix 175 mg Filmtabletten	FR/H/0623/001	BE534497	BIOPROJET EUROPE LTD	BE
Tiorfix 175 mg filmomhulde tabletten	FR/H/0623/001	BE534497	BIOPROJET EUROPE LTD	BE
Tiorfix Junior 30 mg Granulat zur Herstellung einer Suspension zum Einnehmen	SE/H/1342/002	BE400732	BIOPROJET EUROPE LTD	BE
Tiorfix Junior 30 mg Granulat zur Herstellung einer Suspension zum Einnehmen	SE/H/1342/002	BE400732	BIOPROJET EUROPE LTD	BE
Tiorfix Junior 30 mg Granulat zur Herstellung einer Suspension zum Einnehmen	SE/H/1342/002	BE400732	BIOPROJET EUROPE LTD	BE
Tiorfix Junior 30 mg Granulat zur Herstellung einer Suspension zum Einnehmen	SE/H/1342/002	BE400732	BIOPROJET EUROPE LTD	BE
Tiorfix Junior 30 mg Granulat zur Herstellung einer Suspension zum Einnehmen	SE/H/1342/002	BE400732	BIOPROJET EUROPE LTD	BE
Tiorfix Junior 30 mg Granulat zur Herstellung einer Suspension zum	SE/H/1342/002	BE400732	BIOPROJET EUROPE LTD	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
Einnehmen				
Tiorfix 100 mg Hartkapsel	SE/H/1342/003	BE400741	BIOPROJET EUROPE LTD	BE
Tiorfix 100 mg Hartkapsel	SE/H/1342/003	BE400741	BIOPROJET EUROPE LTD	BE
Tiorfix 100 mg Hartkapsel	SE/H/1342/003	BE400741	BIOPROJET EUROPE LTD	BE
Tiorfix 100 mg Hartkapsel	SE/H/1342/003	BE400741	BIOPROJET EUROPE LTD	BE
Tiorfix 100 mg Hartkapsel	SE/H/1342/003	BE400741	BIOPROJET EUROPE LTD	BE
Tiorfix Baby 10 mg Granulat zur Herstellung einer Suspension zum Einnehmen	SE/H/1342/001	BE400723	BIOPROJET EUROPE LTD	BE
Tiorfix Baby 10 mg Granulat zur Herstellung einer Suspension zum Einnehmen	SE/H/1342/001	BE400723	BIOPROJET EUROPE LTD	BE
Tiorfix Baby 10 mg Granulat zur Herstellung einer Suspension zum	SE/H/1342/001	BE400723	BIOPROJET EUROPE LTD	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
Einnehmen				
Tiorfix Baby 10 mg Granulat zur Herstellung einer Suspension zum Einnehmen	SE/H/1342/001	BE400723	BIOPROJET EUROPE LTD	BE
Tiorfix Baby 10 mg Granulat zur Herstellung einer Suspension zum Einnehmen	SE/H/1342/001	BE400723	BIOPROJET EUROPE LTD	BE
Tiorfix Baby 10 mg Granulat zur Herstellung einer Suspension zum Einnehmen	SE/H/1342/001	BE400723	BIOPROJET EUROPE LTD	BE
Tiorfix Baby 10 mg granulés pour suspension buvable	SE/H/1342/001	BE400723	BIOPROJET EUROPE LTD	BE
Tiorfix Junior 30 mg granulés pour suspension buvable	SE/H/1342/002	BE400732	BIOPROJET EUROPE LTD	BE
Tiorfix 100 mg gélules	SE/H/1342/003	BE400741	BIOPROJET EUROPE LTD	BE
Tiorfix Baby 10 mg, granulaat voor orale suspensie	SE/H/1342/001	BE400723	BIOPROJET EUROPE LTD	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
Tiorfix Junior 30 mg granulaat voor orale suspensie	SE/H/1342/002	BE400732	BIOPROJET EUROPE LTD	BE
Tiorfix 100 mg harde capsules	SE/H/1342/003	BE400741	BIOPROJET EUROPE LTD	BE
Хидрасек 10 mg гранули за перорална суспензия	not available	II-12465/14.03.2011	BIOPROJET PHARMA	BG
Хидрасек 10 mg гранули за перорална суспензия	not available	II-12465/14.03.2011	BIOPROJET PHARMA	BG
Хидрасек 30 mg гранули за перорална суспензия	not available	II-12466/14.03.2011	BIOPROJET PHARMA	BG
Хидрасек 30 mg гранули за перорална суспензия	not available	II-12466/14.03.2011	BIOPROJET PHARMA	BG
Хидрасек 30 mg гранули за перорална суспензия	not available	II-12466/14.03.2011	BIOPROJET PHARMA	BG
Хидрасек 100 mg твърди капсули	not available	II-12464/14.03.2011	BIOPROJET PHARMA	BG
Хидрасек 100 mg твърди капсули	not available	II-12464/14.03.2011	BIOPROJET PHARMA	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
Hidrasec pro kojence 10 mg granule pro perorální suspenzi	SE/H/1342/002	49/554/11-C	BIOPROJET EUROPE LTD	CZ
Hidrasec pro kojence 10 mg granule pro perorální suspenzi	SE/H/1342/002	49/554/11-C	BIOPROJET EUROPE LTD	CZ
Hidrasec pro kojence 10 mg granule pro perorální suspenzi	SE/H/1342/002	49/554/11-C	BIOPROJET EUROPE LTD	CZ
Hidrasec pro kojence 10 mg granule pro perorální suspenzi	SE/H/1342/002	49/554/11-C	BIOPROJET EUROPE LTD	CZ
Hidrasec pro kojence 10 mg granule pro perorální suspenzi	SE/H/1342/002	49/554/11-C	BIOPROJET EUROPE LTD	CZ
Hidrasec pro děti 30 mg granule pro perorální suspenzi	SE/H/1342/001	49/555/11-C	BIOPROJET EUROPE LTD	CZ
Hidrasec pro děti 30 mg granule pro perorální suspenzi	SE/H/1342/001	49/555/11-C	BIOPROJET EUROPE LTD	CZ
Hidrasec pro děti 30 mg granule pro perorální suspenzi	SE/H/1342/001	49/555/11-C	BIOPROJET EUROPE LTD	CZ
Hidrasec pro děti 30 mg granule pro perorální suspenzi	SE/H/1342/001	49/555/11-C	BIOPROJET EUROPE LTD	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
Hidrasec pro děti 30 mg granule pro perorální suspenzi	SE/H/1342/001	49/555/11-C	BIOPROJET EUROPE LTD	CZ
Hidrasec 100 mg tvrdé tobolky	SE/H/1342/003	49/556/11-C	BIOPROJET EUROPE LTD	CZ
Hidrasec 100 mg tvrdé tobolky	SE/H/1342/003	49/556/11-C	BIOPROJET EUROPE LTD	CZ
Hidrasec 100 mg tvrdé tobolky	SE/H/1342/003	49/556/11-C	BIOPROJET EUROPE LTD	CZ
Hidrasec 100 mg tvrdé tobolky	SE/H/1342/003	49/556/11-C	BIOPROJET EUROPE LTD	CZ
Hidrasec pro kojence 10 mg granule pro perorální suspenzi	SE/H/1342/002	49/554/11-C	BIOPROJET EUROPE LTD	CZ
Hidrasec pro děti 30 mg granule pro perorální suspenzi	SE/H/1342/001	49/555/11-C	BIOPROJET EUROPE LTD	CZ
Hidrasec 100 mg tvrdé tobolky	SE/H/1342/003	49/556/11-C	BIOPROJET EUROPE LTD	CZ
VAPRINO® GEGEN AKUTEN DURCHFALL 100 MG HARTKAPSELN	not available	85316.00.00	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
TIORFAN 10 mg, Granulat zur Herstellung einer Suspension zum Einnehmen.	SE/H/1342/002	59563.00.00	BIOPROJET EUROPE LTD	DE
TIORFAN 10 mg, Granulat zur Herstellung einer Suspension zum Einnehmen.	SE/H/1342/002	59563.00.00	BIOPROJET EUROPE LTD	DE
TIORFAN 10 mg, Granulat zur Herstellung einer Suspension zum Einnehmen.	SE/H/1342/002	59563.00.00	BIOPROJET EUROPE LTD	DE
TIORFAN 10 mg, Granulat zur Herstellung einer Suspension zum Einnehmen.	SE/H/1342/002	59563.00.00	BIOPROJET EUROPE LTD	DE
TIORFAN 10 mg, Granulat zur Herstellung einer Suspension zum Einnehmen.	SE/H/1342/002	59563.00.00	BIOPROJET EUROPE LTD	DE
TIORFAN 30 mg, Granulat zur Herstellung einer Suspension zum Einnehmen.	SE/H/1342/001	59564.00.00	BIOPROJET EUROPE LTD	DE
TIORFAN 30 mg, Granulat zur Herstellung einer Suspension zum Einnehmen.	SE/H/1342/001	59564.00.00	BIOPROJET EUROPE LTD	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
TIORFAN 30 mg, Granulat zur Herstellung einer Suspension zum Einnehmen.	SE/H/1342/001	59564.00.00	BIOPROJET EUROPE LTD	DE
TIORFAN 30 mg, Granulat zur Herstellung einer Suspension zum Einnehmen.	SE/H/1342/001	59564.00.00	BIOPROJET EUROPE LTD	DE
TIORFAN 30 mg, Granulat zur Herstellung einer Suspension zum Einnehmen.	SE/H/1342/001	59564.00.00	BIOPROJET EUROPE LTD	DE
TIORFAN 100 mg Hartkapseln	SE/H/1342/003	62751.00.00	BIOPROJET EUROPE LTD	DE
TIORFAN 100 mg Hartkapseln	SE/H/1342/003	62751.00.00	BIOPROJET EUROPE LTD	DE
TIORFAN 100 mg Hartkapseln	SE/H/1342/003	62751.00.00	BIOPROJET EUROPE LTD	DE
TIORFAN 100 mg Hartkapseln	SE/H/1342/003	62751.00.00	BIOPROJET EUROPE LTD	DE
VAPRINO® GEGEN AKUTEN DURCHFALL 100 MG HARTKAPSELN	not available	85316.00.00	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
TIORFAN 10 mg, Granulat zur Herstellung einer Suspension zum Einnehmen.	SE/H/1342/002	59563.00.00	BIOPROJET EUROPE LTD	DE
TIORFAN 30 mg, Granulat zur Herstellung einer Suspension zum Einnehmen.	SE/H/1342/001	59564.00.00	BIOPROJET EUROPE LTD	DE
TIORFAN 100 mg Hartkapseln	SE/H/1342/003	62751.00.00	BIOPROJET EUROPE LTD	DE
Hidrasec, granulat til oral suspension	SE/H/1342/002	48809	BIOPROJET EUROPE LTD	DK
Hidrasec, granulat til oral suspension	SE/H/1342/002	48809	BIOPROJET EUROPE LTD	DK
Hidrasec, granulat til oral suspension	SE/H/1342/002	48809	BIOPROJET EUROPE LTD	DK
Hidrasec, granulat til oral suspension	SE/H/1342/002	48809	BIOPROJET EUROPE LTD	DK
Hidrasec, granulat til oral suspension	SE/H/1342/002	48809	BIOPROJET EUROPE LTD	DK
Hidrasec, granulat til oral suspension	SE/H/1342/001	48810	BIOPROJET EUROPE LTD	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
Hidrasec, granulat til oral suspension	SE/H/1342/001	48810	BIOPROJET EUROPE LTD	DK
Hidrasec, granulat til oral suspension	SE/H/1342/001	48810	BIOPROJET EUROPE LTD	DK
Hidrasec, granulat til oral suspension	SE/H/1342/001	48810	BIOPROJET EUROPE LTD	DK
Hidrasec, granulat til oral suspension	SE/H/1342/001	48810	BIOPROJET EUROPE LTD	DK
Hidrasec, hårde kapsler	SE/H/1342/003	48811	BIOPROJET EUROPE LTD	DK
Hidrasec, hårde kapsler	SE/H/1342/003	48811	BIOPROJET EUROPE LTD	DK
Hidrasec, hårde kapsler	SE/H/1342/003	48811	BIOPROJET EUROPE LTD	DK
Hidrasec, hårde kapsler	SE/H/1342/003	48811	BIOPROJET EUROPE LTD	DK
Hidrasec, granulat til oral suspension	SE/H/1342/002	48809	BIOPROJET EUROPE LTD	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
Hidrasec, granulat til oral suspension	SE/H/1342/001	48810	BIOPROJET EUROPE LTD	DK
Hidrasec, hårde kapsler	SE/H/1342/003	48811	BIOPROJET EUROPE LTD	DK
Hidrasec, 10 mg suukaudse suspensiooni graanulid	SE/H/1342/002	758111	BIOPROJET EUROPE LTD	EE
Hidrasec, 10 mg suukaudse suspensiooni graanulid	SE/H/1342/002	758111	BIOPROJET EUROPE LTD	EE
Hidrasec, 10 mg suukaudse suspensiooni graanulid	SE/H/1342/002	758111	BIOPROJET EUROPE LTD	EE
Hidrasec, 10 mg suukaudse suspensiooni graanulid	SE/H/1342/002	758111	BIOPROJET EUROPE LTD	EE
Hidrasec, 10 mg suukaudse suspensiooni graanulid	SE/H/1342/002	758111	BIOPROJET EUROPE LTD	EE
Hidrasec, 100 mg kõvakapslid	SE/H/1342/003	758211	BIOPROJET EUROPE LTD	EE
Hidrasec, 100 mg kõvakapslid	SE/H/1342/003	758211	BIOPROJET EUROPE LTD	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
Hidrasec, 100 mg kõvakapslid	SE/H/1342/003	758211	BIOPROJET EUROPE LTD	EE
Hidrasec, 30 mg suukaudse suspensiooni graanulid	SE/H/1342/001	758011	BIOPROJET EUROPE LTD	EE
Hidrasec, 30 mg suukaudse suspensiooni graanulid	SE/H/1342/001	758011	BIOPROJET EUROPE LTD	EE
Hidrasec, 30 mg suukaudse suspensiooni graanulid	SE/H/1342/001	758011	BIOPROJET EUROPE LTD	EE
Hidrasec, 30 mg suukaudse suspensiooni graanulid	SE/H/1342/001	758011	BIOPROJET EUROPE LTD	EE
Hidrasec, 30 mg suukaudse suspensiooni graanulid	SE/H/1342/001	758011	BIOPROJET EUROPE LTD	EE
Hidrasec, 100 mg kõvakapslid	SE/H/1342/003	758211	BIOPROJET EUROPE LTD	EE
Hidrasec, 10 mg suukaudse suspensiooni graanulid	SE/H/1342/002	758111	BIOPROJET EUROPE LTD	EE
Hidrasec, 30 mg suukaudse suspensiooni graanulid	SE/H/1342/001	758011	BIOPROJET EUROPE LTD	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
Hidrasec, 100 mg kõvakapslid	SE/H/1342/003	758211	BIOPROJET EUROPE LTD	EE
HIDRASEC 100 mg cápsulas duras	not available	77809	BIOPROJET FERRER SL	ES
TIORFAN LACTANTES 10 mg granulado para suspensión oral	SE/H/1342/002	64.816	BIOPROJET FERRER SL	ES
TIORFAN NIÑOS 30 mg granulado para suspensión oral	SE/H/1342/001	64.809	BIOPROJET FERRER SL	ES
TIORFAN 100 mg cápsulas duras	SE/H/1342/003	63.286	BIOPROJET FERRER SL	ES
Hidrasec 10 mg rakeet oraalisuspensiota varten	SE/H/1342/002	29702	BIOPROJET EUROPE LTD	FI
Hidrasec 10 mg rakeet oraalisuspensiota varten	SE/H/1342/002	29702	BIOPROJET EUROPE LTD	FI
Hidrasec 10 mg rakeet oraalisuspensiota varten	SE/H/1342/002	29702	BIOPROJET EUROPE LTD	FI
Hidrasec 10 mg rakeet oraalisuspensiota varten	SE/H/1342/002	29702	BIOPROJET EUROPE LTD	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
Hidrasec 10 mg rakeet oraalisuspensiota varten	SE/H/1342/002	29702	BIOPROJET EUROPE LTD	FI
Hidrasec 30 mg rakeet oraalisuspensiota varten	SE/H/1342/001	29703	BIOPROJET EUROPE LTD	FI
Hidrasec 30 mg rakeet oraalisuspensiota varten	SE/H/1342/001	29703	BIOPROJET EUROPE LTD	FI
Hidrasec 30 mg rakeet oraalisuspensiota varten	SE/H/1342/001	29703	BIOPROJET EUROPE LTD	FI
Hidrasec 30 mg rakeet oraalisuspensiota varten	SE/H/1342/001	29703	BIOPROJET EUROPE LTD	FI
Hidrasec 30 mg rakeet oraalisuspensiota varten	SE/H/1342/001	29703	BIOPROJET EUROPE LTD	FI
Hidrasec 100 mg kapseli, kova	SE/H/1342/003	29704	BIOPROJET EUROPE LTD	FI
Hidrasec 100 mg kapseli, kova	SE/H/1342/003	29704	BIOPROJET EUROPE LTD	FI
Hidrasec 100 mg kapseli, kova	SE/H/1342/003	29704	BIOPROJET EUROPE LTD	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
Hidrasec 100 mg kapseli, kova	SE/H/1342/003	29704	BIOPROJET EUROPE LTD	FI
Hidrasec 10 mg rakeet oraalisuspensiota varten	SE/H/1342/002	29702	BIOPROJET EUROPE LTD	FI
Hidrasec 30 mg rakeet oraalisuspensiota varten	SE/H/1342/001	29703	BIOPROJET EUROPE LTD	FI
Hidrasec 100 mg kapseli, kova	SE/H/1342/003	29704	BIOPROJET EUROPE LTD	FI
TIORFAN 10 mg NOURRISSONS, poudre orale en sachet-dose	not available	34009 352 111 3 2	BIOPROJET PHARMA	FR
TIORFAN 30 mg ENFANTS, poudre orale en sachet-dose	not available	34009 352 114-2 2	BIOPROJET PHARMA	FR
TIORFAN 100mg, gélule	not available	34009 334 967 7 7	BIOPROJET PHARMA	FR
TIORFAST 100mg, gélule	not available	34009 389 616 1 4	BIOPROJET PHARMA	FR
TIORFANOR 175 mg, comprimé pelliculé	FR/H/0623/001	34009 382 003 4 8	BIOPROJET PHARMA	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
DIARFIX 100 mg, gélule	not available	34009 492 159 9 7	BIOPROJET PHARMA	FR
TIORFAN 4 mg/mL NOURRISSONS ET ENFANTS, suspension buvable	not available	34009 301 738 4 8	BIOPROJET PHARMA	FR
TIORFAN 4 mg/mL NOURRISSONS ET ENFANTS, suspension buvable	not available	34009 301 738 5 5	BIOPROJET PHARMA	FR
HIDRASEC ΓΙΑ ΠΑΙΔΙΑ 30 mg, Κοκκία για πόσιμο εναιώρημα (σε συσκευασία μιας δόσης)	SE/H/1342/002	2641502	FERRER-GALENICA S.A.	GR
HIDRASEC ΓΙΑ ΒΡΕΦΗ 10 mg, Κοκκία για πόσιμο εναιώρημα (σε συσκευασία μιας δόσης)	SE/H/1342/001	2641501	FERRER-GALENICA S.A.	GR
HIDRASEC 100 mg καψάκια σκληρά	SE/H/1342/003	2641503	FERRER-GALENICA S.A.	GR
Hidrasec Baby 10 mg granulátum belsőleges szuszpenzióhoz	SE/H/1342/002	OGYI-T-22076/02	BIOPROJET EUROPE LTD	HU
Hidrasec Baby 10 mg granulátum belsőleges szuszpenzióhoz	SE/H/1342/002	OGYI-T-22076/02	BIOPROJET EUROPE LTD	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
Hidrasec Baby 10 mg granulátum belsőleges szuszpenzióhoz	SE/H/1342/002	OGYI-T-22076/02	BIOPROJET EUROPE LTD	HU
Hidrasec Baby 10 mg granulátum belsőleges szuszpenzióhoz	SE/H/1342/002	OGYI-T-22076/02	BIOPROJET EUROPE LTD	HU
Hidrasec Baby 10 mg granulátum belsőleges szuszpenzióhoz	SE/H/1342/002	OGYI-T-22076/02	BIOPROJET EUROPE LTD	HU
Hidrasec Junior 30 mg granulátum belsőleges szuszpenzióhoz	SE/H/1342/001	OGYI-T-22076/01	BIOPROJET EUROPE LTD	HU
Hidrasec Junior 30 mg granulátum belsőleges szuszpenzióhoz	SE/H/1342/001	OGYI-T-22076/01	BIOPROJET EUROPE LTD	HU
Hidrasec Junior 30 mg granulátum belsőleges szuszpenzióhoz	SE/H/1342/001	OGYI-T-22076/01	BIOPROJET EUROPE LTD	HU
Hidrasec Junior 30 mg granulátum belsőleges szuszpenzióhoz	SE/H/1342/001	OGYI-T-22076/01	BIOPROJET EUROPE LTD	HU
Hidrasec Junior 30 mg granulátum belsőleges szuszpenzióhoz	SE/H/1342/001	OGYI-T-22076/01	BIOPROJET EUROPE LTD	HU
Hidrasec 100 mg kemény kapszula	SE/H/1342/003	OGYI-T-22076/03	BIOPROJET EUROPE LTD	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
Hidrasec 100 mg kemény kapszula	SE/H/1342/003	OGYI-T-22076/03	BIOPROJET EUROPE LTD	HU
Hidrasec 100 mg kemény kapszula	SE/H/1342/003	OGYI-T-22076/03	BIOPROJET EUROPE LTD	HU
Hidrasec 100 mg kemény kapszula	SE/H/1342/003	OGYI-T-22076/03	BIOPROJET EUROPE LTD	HU
Hidrasec Baby 10 mg granulátum belsőleges szuszpenzióhoz	SE/H/1342/002	OGYI-T-22076/02	BIOPROJET EUROPE LTD	HU
Hidrasec Junior 30 mg granulátum belsőleges szuszpenzióhoz	SE/H/1342/001	OGYI-T-22076/01	BIOPROJET EUROPE LTD	HU
Hidrasec 100 mg kemény kapszula	SE/H/1342/003	OGYI-T-22076/03	BIOPROJET EUROPE LTD	HU
Hidrasec Infants 10mg Granules for Oral Suspension.	SE/H/1342/002	PA1714/001/002	BIOPROJET EUROPE LTD	IE
Hidrasec Infants 10mg Granules for Oral Suspension.	SE/H/1342/002	PA1714/001/002	BIOPROJET EUROPE LTD	IE
Hidrasec Infants 10mg Granules for Oral Suspension.	SE/H/1342/002	PA1714/001/002	BIOPROJET EUROPE 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
Hidrasec Infants 10mg Granules for Oral Suspension.	SE/H/1342/002	PA1714/001/002	BIOPROJET EUROPE LTD	IE
Hidrasec Infants 10mg Granules for Oral Suspension.	SE/H/1342/002	PA1714/001/002	BIOPROJET EUROPE LTD	IE
Hidrasec Children 30mg Granules for Oral Suspension	SE/H/1342/001	PA1714/001/003	BIOPROJET EUROPE LTD	IE
Hidrasec Children 30mg Granules for Oral Suspension	SE/H/1342/001	PA1714/001/003	BIOPROJET EUROPE LTD	IE
Hidrasec Children 30mg Granules for Oral Suspension	SE/H/1342/001	PA1714/001/003	BIOPROJET EUROPE LTD	IE
Hidrasec Children 30mg Granules for Oral Suspension	SE/H/1342/001	PA1714/001/003	BIOPROJET EUROPE LTD	IE
Hidrasec Children 30mg Granules for Oral Suspension	SE/H/1342/001	PA1714/001/003	BIOPROJET EUROPE LTD	IE
Hidrasec 100 mg hard capsules	SE/H/1342/003	PA1714/001/001	BIOPROJET EUROPE LTD	IE
Hidrasec 100 mg hard capsules	SE/H/1342/003	PA1714/001/001	BIOPROJET EUROPE 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
Hidrasec 100 mg hard capsules	SE/H/1342/003	PA1714/001/001	BIOPROJET EUROPE LTD	IE
Hidrasec 100 mg hard capsules	SE/H/1342/003	PA1714/001/001	BIOPROJET EUROPE LTD	IE
Hidrasec Infants 10mg Granules for Oral Suspension.	SE/H/1342/002	PA1714/001/002	BIOPROJET EUROPE LTD	IE
Hidrasec Children 30mg Granules for Oral Suspension	SE/H/1342/001	PA1714/001/003	BIOPROJET EUROPE LTD	IE
Hidrasec 100 mg hard capsules	SE/H/1342/003	PA1714/001/001	BIOPROJET EUROPE LTD	IE
TIORFIX 100 mg capsule rigide	SE/H/1342/003	037518178	BIOPROJET EUROPE LTD	IT
Tiorfanor 175 mg compresse rivestite con film	FR/H/0623/001	046299018	BIOPROJET PHARMA	IT
TIORFIX 10 mg, granulato per sospensione orale PRIMA INFANZIA	SE/H/1342/002	037518115	BIOPROJET EUROPE LTD	IT
TIORFIX 10 mg, granulato per sospensione orale PRIMA INFANZIA	SE/H/1342/002	037518127	BIOPROJET EUROPE LTD	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
TIORFIX 10 mg, granulato per sospensione orale PRIMA INFANZIA	SE/H/1342/002	037518139	BIOPROJET EUROPE LTD	IT
TIORFIX 10 mg, granulato per sospensione orale PRIMA INFANZIA	SE/H/1342/002	037518141	BIOPROJET EUROPE LTD	IT
TIORFIX 10 mg, granulato per sospensione orale PRIMA INFANZIA	SE/H/1342/002	037518154	BIOPROJET EUROPE LTD	IT
TIORFIX 10 mg, granulato per sospensione orale PRIMA INFANZIA	SE/H/1342/002	037518166	BIOPROJET EUROPE LTD	IT
TIORFIX 30 mg, granulato per sospensione orale BAMBINI	SE/H/1342/001	037518053	BIOPROJET EUROPE LTD	IT
TIORFIX 30 mg, granulato per sospensione orale BAMBINI	SE/H/1342/001	037518065	BIOPROJET EUROPE LTD	IT
TIORFIX 30 mg, granulato per sospensione orale BAMBINI	SE/H/1342/001	037518077	BIOPROJET EUROPE LTD	IT
TIORFIX 30 mg, granulato per sospensione orale BAMBINI	SE/H/1342/001	037518089	BIOPROJET EUROPE LTD	IT
TIORFIX 30 mg, granulato per sospensione orale BAMBINI	SE/H/1342/001	037518091	BIOPROJET EUROPE LTD	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
TIORFIX 30 mg, granulato per sospensione orale BAMBINI	SE/H/1342/001	037518103	BIOPROJET EUROPE LTD	IT
TIORFIX 100 mg capsule rigide	SE/H/1342/003	037518014	BIOPROJET EUROPE LTD	IT
TIORFIX 100 mg capsule rigide	SE/H/1342/003	037518026	BIOPROJET EUROPE LTD	IT
TIORFIX 100 mg capsule rigide	SE/H/1342/003	037518038	BIOPROJET EUROPE LTD	IT
TIORFIX 100 mg capsule rigide	SE/H/1342/003	037518040	BIOPROJET EUROPE LTD	IT
Hidrasec 100 mg kietosios kapsulės	SE/H/1342/003	LT/1/12/2804/013	BIOPROJET EUROPE LTD	LT
Hidrasec 100 mg kietosios kapsulės	SE/H/1342/003	LT/1/12/2804/017	BIOPROJET EUROPE LTD	LT
Hidrasec 100 mg kietosios kapsulės	SE/H/1342/003	LT/1/12/2804/014	BIOPROJET EUROPE LTD	LT
Hidrasec 10 mg granulės geriamajai suspensijai	SE/H/1342/002	LT/1/12/2804/001	BIOPROJET EUROPE LTD	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
Hidrasec 10 mg granulės geriamajai suspensijai	SE/H/1342/002	LT/1/12/2804/002	BIOPROJET EUROPE LTD	LT
Hidrasec 10 mg granulės geriamajai suspensijai	SE/H/1342/002	LT/1/12/2804/003	BIOPROJET EUROPE LTD	LT
Hidrasec 10 mg granulės geriamajai suspensijai	SE/H/1342/002	LT/1/12/2804/004	BIOPROJET EUROPE LTD	LT
Hidrasec 10 mg granulės geriamajai suspensijai	SE/H/1342/002	LT/1/12/2804/005	BIOPROJET EUROPE LTD	LT
Hidrasec 10 mg granulės geriamajai suspensijai	SE/H/1342/002	LT/1/12/2804/006	BIOPROJET EUROPE LTD	LT
Hidrasec 30 mg granulės geriamajai suspensijai	SE/H/1342/001	LT/1/12/2804/007	BIOPROJET EUROPE LTD	LT
Hidrasec 30 mg granulės geriamajai suspensijai	SE/H/1342/001	LT/1/12/2804/009	BIOPROJET EUROPE LTD	LT
Hidrasec 30 mg granulės geriamajai suspensijai	SE/H/1342/001	LT/1/12/2804/010	BIOPROJET EUROPE LTD	LT
Hidrasec 30 mg granulės geriamajai suspensijai	SE/H/1342/001	LT/1/12/2804/011	BIOPROJET EUROPE LTD	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
Hidrasec 30 mg granulės geriamajai suspensijai	SE/H/1342/001	LT/1/12/2804/012	BIOPROJET EUROPE LTD	LT
Hidrasec 30 mg granulės geriamajai suspensijai	SE/H/1342/001	LT/1/12/2804/008	BIOPROJET EUROPE LTD	LT
Hidrasec 100 mg kietosios kapsulės	SE/H/1342/003	LT/1/12/2804/015	BIOPROJET EUROPE LTD	LT
Hidrasec 100 mg kietosios kapsulės	SE/H/1342/003	LT/1/12/2804/016	BIOPROJET EUROPE LTD	LT
Tiorfix Baby 10 mg granulés pour suspension buvable	SE/H/1342/002	2012010049	BIOPROJET PHARMA	LU
Tiorfix Baby 10 mg granulés pour suspension buvable	SE/H/1342/002	2012010049	BIOPROJET PHARMA	LU
Tiorfix Baby 10 mg granulés pour suspension buvable	SE/H/1342/002	2012010049	BIOPROJET PHARMA	LU
Tiorfix Baby 10 mg granulés pour suspension buvable	SE/H/1342/002	2012010049	BIOPROJET PHARMA	LU
Tiorfix Baby 10 mg granulés pour suspension buvable	SE/H/1342/002	2012010049	BIOPROJET PHARMA	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
Tiorfix Junior 30 mg granulés pour suspension buvable	SE/H/1342/001	2012010050	BIOPROJET PHARMA	LU
Tiorfix Junior 30 mg granulés pour suspension buvable	SE/H/1342/001	2012010050	BIOPROJET PHARMA	LU
Tiorfix Junior 30 mg granulés pour suspension buvable	SE/H/1342/001	2012010050	BIOPROJET PHARMA	LU
Tiorfix Junior 30 mg granulés pour suspension buvable	SE/H/1342/001	2012010050	BIOPROJET PHARMA	LU
Tiorfix Junior 30 mg granulés pour suspension buvable	SE/H/1342/001	2012010050	BIOPROJET PHARMA	LU
Tiorfix 100 mg gélules	ES/H/0122/003	2012010048	BIOPROJET PHARMA	LU
Tiorfix 100 mg gélules	ES/H/0122/003	2012010048	BIOPROJET PHARMA	LU
Tiorfix 100 mg gélules	ES/H/0122/003	2012010048	BIOPROJET PHARMA	LU
Tiorfix 100 mg gélules	ES/H/0122/003	2012010048	BIOPROJET PHARMA	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
Tiorfix Junior 30 mg granulés pour suspension buvable	SE/H/1342/001	2012010050	BIOPROJET PHARMA	LU
Tiorfix Baby 10 mg granulés pour suspension buvable	SE/H/1342/002	2012010049	BIOPROJET PHARMA	LU
Tiorfix 100 mg gélules	ES/H/0122/003	2012010048	BIOPROJET PHARMA	LU
Hidrasec 10 mg granulas iekšķīgi lietojamas suspensijas pagatavošanai	SE/H/1342/002	11-0392	BIOPROJET EUROPE LTD	LV
Hidrasec 10 mg granulas iekšķīgi lietojamas suspensijas pagatavošanai	SE/H/1342/002	11-0392	BIOPROJET EUROPE LTD	LV
Hidrasec 10 mg granulas iekšķīgi lietojamas suspensijas pagatavošanai	SE/H/1342/002	11-0392	BIOPROJET EUROPE LTD	LV
Hidrasec 10 mg granulas iekšķīgi lietojamas suspensijas pagatavošanai	SE/H/1342/002	11-0392	BIOPROJET EUROPE LTD	LV
Hidrasec 10 mg granulas iekšķīgi lietojamas suspensijas pagatavošanai	SE/H/1342/002	11-0392	BIOPROJET EUROPE LTD	LV
Hidrasec 30 mg granulas iekšķīgi lietojamas suspensijas pagatavošanai	SE/H/1342/001	11-0393	BIOPROJET EUROPE LTD	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
Hidrasec 30 mg granulas iekšķīgi lietojamas suspensijas pagatavošanai	SE/H/1342/001	11-0393	BIOPROJET EUROPE LTD	LV
Hidrasec 30 mg granulas iekšķīgi lietojamas suspensijas pagatavošanai	SE/H/1342/001	11-0393	BIOPROJET EUROPE LTD	LV
Hidrasec 30 mg granulas iekšķīgi lietojamas suspensijas pagatavošanai	SE/H/1342/001	11-0393	BIOPROJET EUROPE LTD	LV
Hidrasec 30 mg granulas iekšķīgi lietojamas suspensijas pagatavošanai	SE/H/1342/001	11-0393	BIOPROJET EUROPE LTD	LV
Hidrasec 100 mg cietās kapsulas	SE/H/1342/003	11-0494	BIOPROJET EUROPE LTD	LV
Hidrasec 100 mg cietās kapsulas	SE/H/1342/003	11-0494	BIOPROJET EUROPE LTD	LV
Hidrasec 100 mg cietās kapsulas	SE/H/1342/003	11-0495	BIOPROJET EUROPE LTD	LV
Hidrasec 100 mg cietās kapsulas	SE/H/1342/003	11-0494	BIOPROJET EUROPE LTD	LV
Hidrasec 100 mg cietās kapsulas	SE/H/1342/003	11-0495	BIOPROJET EUROPE LTD	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
Hidrasec 10 mg granulas iekšķīgi lietojamas suspensijas pagatavošanai	SE/H/1342/002	11-0392	BIOPROJET EUROPE LTD	LV
Hidrasec 30 mg granulas iekšķīgi lietojamas suspensijas pagatavošanai	SE/H/1342/001	11-0393	BIOPROJET EUROPE LTD	LV
Hidrasec Baby 10 mg, granulaat voor orale suspensie.	SE/H/1342/002	RVG 109470	BIOPROJET EUROPE LTD	NL
Hidrasec Baby 10 mg, granulaat voor orale suspensie.	SE/H/1342/002	RVG 109470	BIOPROJET EUROPE LTD	NL
Hidrasec Baby 10 mg, granulaat voor orale suspensie.	SE/H/1342/002	RVG 109470	BIOPROJET EUROPE LTD	NL
Hidrasec Baby 10 mg, granulaat voor orale suspensie.	SE/H/1342/002	RVG 109470	BIOPROJET EUROPE LTD	NL
Hidrasec Baby 10 mg, granulaat voor orale suspensie.	SE/H/1342/002	RVG 109470	BIOPROJET EUROPE LTD	NL
Hidrasec Junior 30 mg, granulaat voor orale suspensie.	SE/H/1342/001	RVG 109466	BIOPROJET EUROPE LTD	NL
Hidrasec Junior 30 mg, granulaat voor orale suspensie.	SE/H/1342/001	RVG 109466	BIOPROJET EUROPE LTD	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
Hidrasec Junior 30 mg, granulaat voor orale suspensie.	SE/H/1342/001	RVG 109466	BIOPROJET EUROPE LTD	NL
Hidrasec Junior 30 mg, granulaat voor orale suspensie.	SE/H/1342/001	RVG 109466	BIOPROJET EUROPE LTD	NL
Hidrasec Junior 30 mg, granulaat voor orale suspensie.	SE/H/1342/001	RVG 109466	BIOPROJET EUROPE LTD	NL
Hidrasec 100 mg, capsules hard	SE/H/1342/003	RVG 109471	BIOPROJET EUROPE LTD	NL
Hidrasec 100 mg, capsules hard	SE/H/1342/003	RVG 109471	BIOPROJET EUROPE LTD	NL
Hidrasec 100 mg, capsules hard	SE/H/1342/003	RVG 109471	BIOPROJET EUROPE LTD	NL
Hidrasec 100 mg, capsules hard	SE/H/1342/003	RVG 109471	BIOPROJET EUROPE LTD	NL
Hidrasec Baby 10 mg, granulaat voor orale suspensie.	SE/H/1342/002	RVG 109470	BIOPROJET EUROPE LTD	NL
Hidrasec Junior 30 mg, granulaat voor orale suspensie.	SE/H/1342/001	RVG 109466	BIOPROJET EUROPE LTD	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
Hidrasec 100 mg, capsules hard	SE/H/1342/003	RVG 109471	BIOPROJET EUROPE LTD	NL
Hidrasec 10 mg granulat til mikstur, suspensjon	SE/H/1342/002	11-8261	BIOPROJET EUROPE LTD	NO
Hidrasec 10 mg granulat til mikstur, suspensjon	SE/H/1342/002	11-8261	BIOPROJET EUROPE LTD	NO
Hidrasec 10 mg granulat til mikstur, suspensjon	SE/H/1342/002	11-8261	BIOPROJET EUROPE LTD	NO
Hidrasec 10 mg granulat til mikstur, suspensjon	SE/H/1342/002	11-8261	BIOPROJET EUROPE LTD	NO
Hidrasec 10 mg granulat til mikstur, suspensjon	SE/H/1342/002	11-8261	BIOPROJET EUROPE LTD	NO
Hidrasec 30 mg granulat til mikstur, suspensjon	SE/H/1342/001	11-8260	BIOPROJET EUROPE LTD	NO
Hidrasec 30 mg granulat til mikstur, suspensjon	SE/H/1342/001	11-8260	BIOPROJET EUROPE LTD	NO
Hidrasec 30 mg granulat til mikstur, suspensjon	SE/H/1342/001	11-8260	BIOPROJET EUROPE LTD	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
Hidrasec 30 mg granulat til mikstur, suspensjon	SE/H/1342/001	11-8260	BIOPROJET EUROPE LTD	NO
Hidrasec 30 mg granulat til mikstur, suspensjon	SE/H/1342/001	11-8260	BIOPROJET EUROPE LTD	NO
Hidrasec 10 mg granulat til mikstur, suspensjon	SE/H/1342/002	11-8261	BIOPROJET EUROPE LTD	NO
Hidrasec 30 mg granulat til mikstur, suspensjon	SE/H/1342/001	11-8260	BIOPROJET EUROPE LTD	NO
Hidrasec 10 mg, 10 mg, granulat do sporządzania zawiesiny doustnej	SE/H/1342/002	18916	BIOPROJET EUROPE LTD	PL
Hidrasec 10 mg, 10 mg, granulat do sporządzania zawiesiny doustnej	SE/H/1342/002	18916	BIOPROJET EUROPE LTD	PL
Hidrasec 10 mg, 10 mg, granulat do sporządzania zawiesiny doustnej	SE/H/1342/002	18916	BIOPROJET EUROPE LTD	PL
Hidrasec 10 mg, 10 mg, granulat do sporządzania zawiesiny doustnej	SE/H/1342/002	18916	BIOPROJET EUROPE LTD	PL
Hidrasec 10 mg, 10 mg, granulat do sporządzania zawiesiny doustnej	SE/H/1342/002	18916	BIOPROJET EUROPE LTD	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
Hidrasec 30 mg, 30 mg, granulat do sporządzania zawiesiny doustnej	SE/H/1342/001	18917	BIOPROJET EUROPE LTD	PL
Hidrasec 30 mg, 30 mg, granulat do sporządzania zawiesiny doustnej	SE/H/1342/001	18917	BIOPROJET EUROPE LTD	PL
Hidrasec 30 mg, 30 mg, granulat do sporządzania zawiesiny doustnej	SE/H/1342/001	18917	BIOPROJET EUROPE LTD	PL
Hidrasec 30 mg, 30 mg, granulat do sporządzania zawiesiny doustnej	SE/H/1342/001	18917	BIOPROJET EUROPE LTD	PL
Hidrasec 30 mg, 30 mg, granulat do sporządzania zawiesiny doustnej	SE/H/1342/001	18917	BIOPROJET EUROPE LTD	PL
Tiorfan, 100 mg, kapsułki, twarde	SE/H/1342/003	18918	BIOPROJET EUROPE LTD	PL
Tiorfan, 100 mg, kapsułki, twarde	SE/H/1342/003	18918	BIOPROJET EUROPE LTD	PL
Tiorfan, 100 mg, kapsułki, twarde	SE/H/1342/003	18918	BIOPROJET EUROPE LTD	PL
Tiorfan, 100 mg, kapsułki, twarde	SE/H/1342/003	18918	BIOPROJET EUROPE LTD	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
Hidrasec 30 mg, 30 mg, granulat do sporządzania zawiesiny doustnej	SE/H/1342/001	18917	BIOPROJET EUROPE LTD	PL
Hidrasec 10 mg, 10 mg, granulat do sporządzania zawiesiny doustnej	SE/H/1342/002	18916	BIOPROJET EUROPE LTD	PL
Tiorfan, 100 mg, kapsułki, twarde	SE/H/1342/003	18918	BIOPROJET EUROPE LTD	PL
TIORFAN 100 mg cápsulas	SE/H/1342/003	5756689	FERRER INTERNACIONAL, S.A.	PT
TIORFAN 100 mg cápsulas	SE/H/1342/003	5756788	FERRER INTERNACIONAL, S.A.	PT
TIORFAN INFANTIL 30 mg granulado para suspensão oral	SE/H/1342/001	5109186	FERRER INTERNACIONAL, S.A.	PT
TIORFAN INFANTIL 30 mg granulado para suspensão oral	SE/H/1342/001	5109285	FERRER INTERNACIONAL, S.A.	PT
TIORFAN INFANTIL 30 mg granulado para suspensão oral	SE/H/1342/001	5109384	FERRER INTERNACIONAL, S.A.	PT
TIORFAN LACTENTE 10 mg granulado para suspensão oral	SE/H/1342/002	5109087	FERRER INTERNACIONAL, S.A.	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
TIORFAN LACTENTE 10 mg granulado para suspensão oral	SE/H/1342/002	5108980	FERRER INTERNACIONAL, S.A.	PT
TIORFAN LACTENTE 10 mg granulado para suspensão oral	SE/H/1342/002	5108881	FERRER INTERNACIONAL, S.A.	PT
TIORFAN 100 mg hard capsules	ES/H/0122/003	46035	BIOPROJET EUROPE LTD	SE
TIORFAN 100 mg hard capsules	ES/H/0122/003	46035	BIOPROJET EUROPE LTD	SE
TIORFAN 100 mg hard capsules	ES/H/0122/003	46035	BIOPROJET EUROPE LTD	SE
TIORFAN 100 mg hard capsules	ES/H/0122/003	46035	BIOPROJET EUROPE LTD	SE
Hidrasec 10 mg granulat till oral suspension	ES/H/0122/002	46034	BIOPROJET EUROPE LTD	SE
Hidrasec 10 mg granulat till oral suspension	ES/H/0122/002	46034	BIOPROJET EUROPE LTD	SE
Hidrasec 10 mg granulat till oral suspension	ES/H/0122/002	46034	BIOPROJET EUROPE LTD	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
Hidrasec 10 mg granulat till oral suspension	ES/H/0122/002	46034	BIOPROJET EUROPE LTD	SE
Hidrasec 10 mg granulat till oral suspension	ES/H/0122/002	46034	BIOPROJET EUROPE LTD	SE
Hidrasec 30 mg granulat till oral suspension	ES/H/0122/001	46033	BIOPROJET EUROPE LTD	SE
Hidrasec 30 mg granulat till oral suspension	ES/H/0122/001	46033	BIOPROJET EUROPE LTD	SE
Hidrasec 30 mg granulat till oral suspension	ES/H/0122/001	46033	BIOPROJET EUROPE LTD	SE
Hidrasec 30 mg granulat till oral suspension	ES/H/0122/001	46033	BIOPROJET EUROPE LTD	SE
Hidrasec 30 mg granulat till oral suspension	ES/H/0122/001	46033	BIOPROJET EUROPE LTD	SE
Hidrasec 100 mg hårda kapslar	ES/H/0122/003	46035	BIOPROJET EUROPE LTD	SE
Hidrasec 100 mg hårda kapslar	ES/H/0122/003	46035	BIOPROJET EUROPE LTD	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
Hidrasec 100 mg hårda kapslar	ES/H/0122/003	46035	BIOPROJET EUROPE LTD	SE
Hidrasec 100 mg hårda kapslar	ES/H/0122/003	46035	BIOPROJET EUROPE LTD	SE
Hidrasec 100 mg hårda kapslar	ES/H/0122/003	46035	BIOPROJET EUROPE LTD	SE
Hidrasec 10 mg granulat till oral suspension	ES/H/0122/002	46034	BIOPROJET EUROPE LTD	SE
Hidrasec 30 mg granulat till oral suspension	ES/H/0122/001	46033	BIOPROJET EUROPE LTD	SE
TIORFAN 100 mg hard capsules	ES/H/0122/003	46035	BIOPROJET EUROPE LTD	SE
Hidrasec za dojenčke 10 mg zrnca za peroralno suspenzijo	SE/H/1342/002	H/12/00735/006	BIOPROJET EUROPE LTD	SI
Hidrasec za dojenčke 10 mg zrnca za peroralno suspenzijo	SE/H/1342/002	H/12/00735/008	BIOPROJET EUROPE LTD	SI
Hidrasec za dojenčke 10 mg zrnca za peroralno suspenzijo	SE/H/1342/002	H/12/00735/009	BIOPROJET EUROPE LTD	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
Hidrasec za dojenčke 10 mg zrnca za peroralno suspenzijo	SE/H/1342/002	H/12/00735/010	BIOPROJET EUROPE LTD	SI
Hidrasec za dojenčke 10 mg zrnca za peroralno suspenzijo	SE/H/1342/002	H/12/00735/011	BIOPROJET EUROPE LTD	SI
Hidrasec za otroke 30 mg zrnca za peroralno suspenzijo	SE/H/1342/001	H/12/00735/013	BIOPROJET EUROPE LTD	SI
Hidrasec za otroke 30 mg zrnca za peroralno suspenzijo	SE/H/1342/001	H/12/00735/014	BIOPROJET EUROPE LTD	SI
Hidrasec za otroke 30 mg zrnca za peroralno suspenzijo	SE/H/1342/001	H/12/00735/015	BIOPROJET EUROPE LTD	SI
Hidrasec za otroke 30 mg zrnca za peroralno suspenzijo	SE/H/1342/001	H/12/00735/016	BIOPROJET EUROPE LTD	SI
Hidrasec za otroke 30 mg zrnca za peroralno suspenzijo	SE/H/1342/001	H/12/00735/017	BIOPROJET EUROPE LTD	SI
Hidrasec 100 mg trde kapsule	SE/H/1342/003	H/12/00735/003	BIOPROJET EUROPE LTD	SI
Hidrasec 100 mg trde kapsule	SE/H/1342/003	H/12/00735/004	BIOPROJET EUROPE LTD	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
Hidrasec 100 mg trde kapsule	SE/H/1342/003	H/12/00735/005	BIOPROJET EUROPE LTD	SI
Hidrasec 100 mg trde kapsule	SE/H/1342/003	H/12/00735/002	BIOPROJET EUROPE LTD	SI
Hidrasec za otroke 30 mg zrnca za peroralno suspenzijo	SE/H/1342/001	H/12/00735/012	BIOPROJET EUROPE LTD	SI
Hidrasec za dojencke 10 mg zrnca za peroralno suspenzijo	SE/H/1342/002	H/12/00735/007	BIOPROJET EUROPE LTD	SI
Hidrasec 100 mg trde kapsule	SE/H/1342/003	H/12/00735/001	BIOPROJET EUROPE LTD	SI
Hidrasec pre dojcatá 10 mg, granulát na perorálnu suspenziu	SE/H/1342/002	49/0581/11-S	BIOPROJET EUROPE LTD	SK
Hidrasec pre dojčatá 10 mg, granulát na perorálnu suspenziu	SE/H/1342/002	49/0581/11-S	BIOPROJET EUROPE LTD	SK
Hidrasec pre dojčatá 10 mg, granulát na perorálnu suspenziu	SE/H/1342/002	49/0581/11-S	BIOPROJET EUROPE LTD	SK
Hidrasec pre dojčatá 10 mg, granulát na perorálnu suspenziu	SE/H/1342/002	49/0581/11-S	BIOPROJET EUROPE LTD	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
Hidrasec pre dojčatá 10 mg, granulát na perorálnu suspenziu	SE/H/1342/002	49/0581/11-S	BIOPROJET EUROPE LTD	SK
Hidrasec pre deti 30 mg, granulát na perorálnu suspenziu	SE/H/1342/001	49/0580/11-S	BIOPROJET EUROPE LTD	SK
Hidrasec pre deti 30 mg, granulát na perorálnu suspenziu	SE/H/1342/001	49/0580/11-S	BIOPROJET EUROPE LTD	SK
Hidrasec pre deti 30 mg, granulát na perorálnu suspenziu	SE/H/1342/001	49/0580/11-S	BIOPROJET EUROPE LTD	SK
Hidrasec pre deti 30 mg, granulát na perorálnu suspenziu	SE/H/1342/001	49/0580/11-S	BIOPROJET EUROPE LTD	SK
Hidrasec pre deti 30 mg, granulát na perorálnu suspenziu	SE/H/1342/001	49/0580/11-S	BIOPROJET EUROPE LTD	SK
Hidrasec 100 mg tvrdé kapsuly	SE/H/1342/003	49/0582/11-S	BIOPROJET EUROPE LTD	SK
Hidrasec 100 mg tvrdé kapsuly	SE/H/1342/003	49/0582/11-S	BIOPROJET EUROPE LTD	SK
Hidrasec 100 mg tvrdé kapsuly	SE/H/1342/003	49/0582/11-S	BIOPROJET EUROPE LTD	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
Hidrasec 100 mg tvrdé kapsuly	SE/H/1342/003	49/0582/11-S	BIOPROJET EUROPE LTD	SK
Hidrasec pre dojčatá 10 mg, granulát na perorálnu suspenziu	SE/H/1342/002	49/0581/11-S	BIOPROJET EUROPE LTD	SK
Hidrasec pre deti 30 mg, granulát na perorálnu suspenziu	SE/H/1342/001	49/0580/11-S	BIOPROJET EUROPE LTD	SK
Hidrasec 100 mg tvrdé kapsuly	SE/H/1342/003	49/0582/11-S	BIOPROJET EUROPE LTD	SK
Vaprino 100 mg capsules, hard	UK/H/6676/001	PL 04425/0729	AVENTIS PHARMA LTD	UK
Vaprino 100 mg capsules, hard	UK/H/6676/001	PL 04425/0729	AVENTIS PHARMA LTD	UK
HIDRASEC INFANTS 10 mg, Granules for oral suspension	SE/H/1342/002	PL 39418/0001	BIOPROJET EUROPE LTD	UK
HIDRASEC INFANTS 10 mg, Granules for oral suspension	SE/H/1342/002	PL 39418/0001	BIOPROJET EUROPE LTD	UK
HIDRASEC INFANTS 10 mg, Granules for oral suspension	SE/H/1342/002	PL 39418/0001	BIOPROJET EUROPE 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
HIDRASEC INFANTS 10 mg, Granules for oral suspension	SE/H/1342/002	PL 39418/0001	BIOPROJET EUROPE LTD	UK
HIDRASEC INFANTS 10 mg, Granules for oral suspension	SE/H/1342/002	PL 39418/0001	BIOPROJET EUROPE LTD	UK
HIDRASEC CHILDREN 30 mg, Granules for oral suspension	SE/H/1342/001	PL 39418/0002	BIOPROJET EUROPE LTD	UK
HIDRASEC CHILDREN 30 mg, Granules for oral suspension	SE/H/1342/001	PL 39418/0002	BIOPROJET EUROPE LTD	UK
HIDRASEC CHILDREN 30 mg, Granules for oral suspension	SE/H/1342/001	PL 39418/0002	BIOPROJET EUROPE LTD	UK
HIDRASEC CHILDREN 30 mg, Granules for oral suspension	SE/H/1342/001	PL 39418/0002	BIOPROJET EUROPE LTD	UK
HIDRASEC CHILDREN 30 mg, Granules for oral suspension	SE/H/1342/001	PL 39418/0002	BIOPROJET EUROPE LTD	UK
HIDRASEC 100 mg Hard Capsules	SE/H/1342/003	PL 39418/0003	BIOPROJET EUROPE LTD	UK
HIDRASEC 100 mg Hard Capsules	SE/H/1342/003	PL 39418/0003	BIOPROJET EUROPE 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
HIDRASEC 100 mg Hard Capsules	SE/H/1342/003	PL 39418/0003	BIOPROJET EUROPE LTD	UK
HIDRASEC 100 mg Hard Capsules	SE/H/1342/003	PL 39418/0003	BIOPROJET EUROPE LTD	UK
Vaprino 100 mg capsules, hard	UK/H/6676/001	PL 04425/0729	AVENTIS PHARMA LTD	UK
Vaprino 100 mg capsules, hard	UK/H/6676/001	PL 04425/0729	AVENTIS PHARMA LTD	UK
HIDRASEC INFANTS 10 mg, Granules for oral suspension	SE/H/1342/002	PL 39418/0001	BIOPROJET EUROPE LTD	UK
HIDRASEC CHILDREN 30 mg, Granules for oral suspension	SE/H/1342/001	PL 39418/0002	BIOPROJET EUROPE LTD	UK
HIDRASEC 100 mg Hard Capsules	SE/H/1342/003	PL 39418/0003	BIOPROJET EUROPE LTD	UK