22 May 2025
EMA/139612/2025
Human Medicines Division

List of nationally authorised medicinal products

Active substance(s): diclofenac (systemic formulations)

Procedure No. PSUSA/00001048/202409

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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
Voltaren ® 100 mg - Zäpfchen für Erwachsene	not available	1-16308	NOVARTIS PHARMA GMBH	AT
Voltaren ® 100 mg - Zäpfchen für Erwachsene	not available	1-16308	NOVARTIS PHARMA GMBH	AT
Voltaren ® 100 mg - Zäpfchen für Erwachsene	not available	1-16308	NOVARTIS PHARMA GMBH	AT
Voltaren ® 100 mg - Zäpfchen für Erwachsene	not available	1-16308	NOVARTIS PHARMA GMBH	AT
Voltaren ® 100 mg - Zäpfchen für Erwachsene	not available	1-16308	NOVARTIS PHARMA GMBH	AT
Voltaren ® 50 mg - Filmtabletten	not available	1-16506	NOVARTIS PHARMA GMBH	AT
Voltaren ® 50 mg - Filmtabletten	not available	1-16506	NOVARTIS PHARMA GMBH	AT
Voltaren ® 50 mg - Filmtabletten	not available	1-16506	NOVARTIS PHARMA GMBH	AT
Voltaren ® 50 mg - Filmtabletten	not available	1-16506	NOVARTIS PHARMA GMBH	AT
Voltaren ® 50 mg - Filmtabletten	not available	1-16506	NOVARTIS PHARMA GMBH	AT
Voltaren ® 50 mg - Zäpfchen für Erwachsene	not available	1-15868	NOVARTIS PHARMA GMBH	AT
Voltaren ® 50 mg - Zäpfchen für Erwachsene	not available	1-15868	NOVARTIS PHARMA GMBH	AT
Voltaren ® 50 mg - Zäpfchen für Erwachsene	not available	1-15868	NOVARTIS PHARMA GMBH	AT
Voltaren ® 50 mg - Zäpfchen für Erwachsene	not available	1-15868	NOVARTIS PHARMA GMBH	AT
Voltaren ® 50 mg - Zäpfchen für Erwachsene	not available	1-15868	NOVARTIS PHARMA GMBH	AT
Voltaren® 75 mg/3 ml - Injektionslösung	not available	1-15915	NOVARTIS PHARMA GMBH	AT
Voltaren® 75 mg/3 ml - Injektionslösung	not available	1-15915	NOVARTIS PHARMA GMBH	AT
Voltaren® 75 mg/3 ml - Injektionslösung	not available	1-15915	NOVARTIS PHARMA GMBH	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
Voltaren® 75 mg/3 ml - Injektionslösung	not available	1-15915	NOVARTIS PHARMA GMBH	AT
Voltaren® 75 mg/3 ml - Injektionslösung	not available	1-15915	NOVARTIS PHARMA GMBH	AT
Voltaren® rapid 50 mg - Dragees	not available	1-19098	NOVARTIS PHARMA GMBH	AT
Voltaren® rapid 50 mg - Dragees	not available	1-19098	NOVARTIS PHARMA GMBH	AT
Voltaren® rapid 50 mg - Dragees	not available	1-19098	NOVARTIS PHARMA GMBH	AT
Voltaren® rapid 50 mg - Dragees	not available	1-19098	NOVARTIS PHARMA GMBH	AT
Voltaren® rapid 50 mg - Dragees	not available	1-19098	NOVARTIS PHARMA GMBH	AT
Voltaren® retard 100 mg - Filmtabletten	not available	1-16856	NOVARTIS PHARMA GMBH	AT
Voltaren® retard 100 mg - Filmtabletten	not available	1-16856	NOVARTIS PHARMA GMBH	AT
Voltaren® retard 100 mg - Filmtabletten	not available	1-16856	NOVARTIS PHARMA GMBH	AT
Voltaren® retard 100 mg - Filmtabletten	not available	1-16856	NOVARTIS PHARMA GMBH	AT
Voltaren® retard 100 mg - Filmtabletten	not available	1-16856	NOVARTIS PHARMA GMBH	AT
Cataflam 50 mg omhulde tabletten	not available	BE147402	NOVARTIS PHARMA N.V.	BE
Cataflam 50 mg omhulde tabletten	not available	BE147402	NOVARTIS PHARMA N.V.	BE
Cataflam 50 mg omhulde tabletten	not available	BE147402	NOVARTIS PHARMA N.V.	BE
Cataflam 50 mg omhulde tabletten	not available	BE147402	NOVARTIS PHARMA N.V.	BE
Cataflam 50 mg omhulde tabletten	not available	BE147402	NOVARTIS PHARMA N.V.	BE
Cataflam 50 mg, comprimés enrobés	not available	BE147402	NOVARTIS PHARMA N.V.	BE
Cataflam 50 mg, comprimés enrobés	not available	BE147402	NOVARTIS PHARMA N.V.	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Cataflam 50 mg, comprimés enrobés	not available	BE147402	NOVARTIS PHARMA N.V.	BE
Cataflam 50 mg, comprimés enrobés	not available	BE147402	NOVARTIS PHARMA N.V.	BE
Cataflam 50 mg, comprimés enrobés	not available	BE147402	NOVARTIS PHARMA N.V.	BE
Cataflam 50 mg, überzogene Tabletten	not available	BE147402	NOVARTIS PHARMA N.V.	BE
Cataflam 50 mg, überzogene Tabletten	not available	BE147402	NOVARTIS PHARMA N.V.	BE
Cataflam 50 mg, überzogene Tabletten	not available	BE147402	NOVARTIS PHARMA N.V.	BE
Cataflam 50 mg, überzogene Tabletten	not available	BE147402	NOVARTIS PHARMA N.V.	BE
Cataflam 50 mg, überzogene Tabletten	not available	BE147402	NOVARTIS PHARMA N.V.	BE
Motifene 75 mg gélules à libération modifiée	not available	PL 22824/0014	GLENWOOD GMBH	BE
Voltaren 100 mg suppositoires.	not available	BE109261	NOVARTIS PHARMA N.V.	BE
Voltaren 100 mg suppositoires.	not available	BE109261	NOVARTIS PHARMA N.V.	BE
Voltaren 100 mg suppositoires.	not available	BE109261	NOVARTIS PHARMA N.V.	BE
Voltaren 100 mg suppositoires.	not available	BE109261	NOVARTIS PHARMA N.V.	BE
Voltaren 100 mg suppositoires.	not available	BE109261	NOVARTIS PHARMA N.V.	BE
Voltaren 100 mg Zäpfchen	not available	BE109261	NOVARTIS PHARMA N.V.	BE
Voltaren 100 mg Zäpfchen	not available	BE109261	NOVARTIS PHARMA N.V.	BE
Voltaren 100 mg Zäpfchen	not available	BE109261	NOVARTIS PHARMA N.V.	BE
Voltaren 100 mg Zäpfchen	not available	BE109261	NOVARTIS PHARMA N.V.	BE
Voltaren 100 mg Zäpfchen	not available	BE109261	NOVARTIS PHARMA N.V.	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Voltaren 100 mg zetpillen.	not available	BE109261	NOVARTIS PHARMA N.V.	BE
Voltaren 100 mg zetpillen.	not available	BE109261	NOVARTIS PHARMA N.V.	BE
Voltaren 100 mg zetpillen.	not available	BE109261	NOVARTIS PHARMA N.V.	BE
Voltaren 100 mg zetpillen.	not available	BE109261	NOVARTIS PHARMA N.V.	BE
Voltaren 100 mg zetpillen.	not available	BE109261	NOVARTIS PHARMA N.V.	BE
Voltaren 25 mg comprimés gastro-résistants.	not available	BE095916	NOVARTIS PHARMA N.V.	BE
Voltaren 25 mg comprimés gastro-résistants.	not available	BE095916	NOVARTIS PHARMA N.V.	BE
Voltaren 25 mg comprimés gastro-résistants.	not available	BE095916	NOVARTIS PHARMA N.V.	BE
Voltaren 25 mg comprimés gastro-résistants.	not available	BE095916	NOVARTIS PHARMA N.V.	BE
Voltaren 25 mg comprimés gastro-résistants.	not available	BE095916	NOVARTIS PHARMA N.V.	BE
Voltaren 25 mg maagsapresistente tabletten.	not available	BE095916	NOVARTIS PHARMA N.V.	BE
Voltaren 25 mg maagsapresistente tabletten.	not available	BE095916	NOVARTIS PHARMA N.V.	BE
Voltaren 25 mg maagsapresistente tabletten.	not available	BE095916	NOVARTIS PHARMA N.V.	BE
Voltaren 25 mg maagsapresistente tabletten.	not available	BE095916	NOVARTIS PHARMA N.V.	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Voltaren 25 mg maagsapresistente tabletten.	not available	BE095916	NOVARTIS PHARMA N.V.	BE
Voltaren 25 mg magensaftresistente Tabletten	not available	BE095916	NOVARTIS PHARMA N.V.	BE
Voltaren 25 mg magensaftresistente Tabletten	not available	BE095916	NOVARTIS PHARMA N.V.	BE
Voltaren 25 mg magensaftresistente Tabletten	not available	BE095916	NOVARTIS PHARMA N.V.	BE
Voltaren 25 mg magensaftresistente Tabletten	not available	BE095916	NOVARTIS PHARMA N.V.	BE
Voltaren 25 mg magensaftresistente Tabletten	not available	BE095916	NOVARTIS PHARMA N.V.	BE
Voltaren 50 mg comprimés gastro-résistants.	not available	BE121116	NOVARTIS PHARMA N.V.	BE
Voltaren 50 mg comprimés gastro-résistants.	not available	BE121116	NOVARTIS PHARMA N.V.	BE
Voltaren 50 mg comprimés gastro-résistants.	not available	BE121116	NOVARTIS PHARMA N.V.	BE
Voltaren 50 mg comprimés gastro-résistants.	not available	BE121116	NOVARTIS PHARMA N.V.	BE
Voltaren 50 mg comprimés gastro-résistants.	not available	BE121116	NOVARTIS PHARMA N.V.	BE
Voltaren 50 mg maagsapresistente tabletten.	not available	BE121116	NOVARTIS PHARMA N.V.	BE
Voltaren 50 mg maagsapresistente tabletten.	not available	BE121116	NOVARTIS PHARMA N.V.	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Voltaren 50 mg maagsapresistente tabletten.	not available	BE121116	NOVARTIS PHARMA N.V.	BE
Voltaren 50 mg maagsapresistente tabletten.	not available	BE121116	NOVARTIS PHARMA N.V.	BE
Voltaren 50 mg maagsapresistente tabletten.	not available	BE121116	NOVARTIS PHARMA N.V.	BE
Voltaren 50 mg magensaftresistente Tabletten	not available	BE121116	NOVARTIS PHARMA N.V.	BE
Voltaren 50 mg magensaftresistente Tabletten	not available	BE121116	NOVARTIS PHARMA N.V.	BE
Voltaren 50 mg magensaftresistente Tabletten	not available	BE121116	NOVARTIS PHARMA N.V.	BE
Voltaren 50 mg magensaftresistente Tabletten	not available	BE121116	NOVARTIS PHARMA N.V.	BE
Voltaren 50 mg magensaftresistente Tabletten	not available	BE121116	NOVARTIS PHARMA N.V.	BE
Voltaren 75 mg/3 ml Injektionslösung	not available	BE113206	NOVARTIS PHARMA N.V.	BE
Voltaren 75 mg/3 ml Injektionslösung	not available	BE113206	NOVARTIS PHARMA N.V.	BE
Voltaren 75 mg/3 ml Injektionslösung	not available	BE113206	NOVARTIS PHARMA N.V.	BE
Voltaren 75 mg/3 ml Injektionslösung	not available	BE113206	NOVARTIS PHARMA N.V.	BE
Voltaren 75 mg/3 ml Injektionslösung	not available	BE113206	NOVARTIS PHARMA N.V.	BE
Voltaren 75 mg/3 ml oplossing voor injectie.	not available	BE113206	NOVARTIS PHARMA N.V.	BE
Voltaren 75 mg/3 ml oplossing voor injectie.	not available	BE113206	NOVARTIS PHARMA N.V.	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Voltaren 75 mg/3 ml oplossing voor injectie.	not available	BE113206	NOVARTIS PHARMA N.V.	BE
Voltaren 75 mg/3 ml oplossing voor injectie.	not available	BE113206	NOVARTIS PHARMA N.V.	BE
Voltaren 75 mg/3 ml oplossing voor injectie.	not available	BE113206	NOVARTIS PHARMA N.V.	BE
Voltaren 75 mg/3 ml solution injectable.	not available	BE113206	NOVARTIS PHARMA N.V.	BE
Voltaren 75 mg/3 ml solution injectable.	not available	BE113206	NOVARTIS PHARMA N.V.	BE
Voltaren 75 mg/3 ml solution injectable.	not available	BE113206	NOVARTIS PHARMA N.V.	BE
Voltaren 75 mg/3 ml solution injectable.	not available	BE113206	NOVARTIS PHARMA N.V.	BE
Voltaren 75 mg/3 ml solution injectable.	not available	BE113206	NOVARTIS PHARMA N.V.	BE
Voltaren Retard 100 mg comprimés à libération prolongée.	not available	BE122071	NOVARTIS PHARMA N.V.	BE
Voltaren Retard 100 mg comprimés à libération prolongée.	not available	BE122071	NOVARTIS PHARMA N.V.	BE
Voltaren Retard 100 mg comprimés à libération prolongée.	not available	BE122071	NOVARTIS PHARMA N.V.	BE
Voltaren Retard 100 mg comprimés à libération prolongée.	not available	BE122071	NOVARTIS PHARMA N.V.	BE
Voltaren Retard 100 mg comprimés à libération prolongée.	not available	BE122071	NOVARTIS PHARMA N.V.	BE
Voltaren Retard 100 mg Retardtabletten	not available	BE122071	NOVARTIS PHARMA N.V.	BE
Voltaren Retard 100 mg Retardtabletten	not available	BE122071	NOVARTIS PHARMA N.V.	BE
Voltaren Retard 100 mg Retardtabletten	not available	BE122071	NOVARTIS PHARMA N.V.	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Voltaren Retard 100 mg Retardtabletten	not available	BE122071	NOVARTIS PHARMA N.V.	BE
Voltaren Retard 100 mg Retardtabletten	not available	BE122071	NOVARTIS PHARMA N.V.	BE
Voltaren Retard 100 mg tabletten met verlengde afgifte.	not available	BE122071	NOVARTIS PHARMA N.V.	BE
Voltaren Retard 100 mg tabletten met verlengde afgifte.	not available	BE122071	NOVARTIS PHARMA N.V.	BE
Voltaren Retard 100 mg tabletten met verlengde afgifte.	not available	BE122071	NOVARTIS PHARMA N.V.	BE
Voltaren Retard 100 mg tabletten met verlengde afgifte.	not available	BE122071	NOVARTIS PHARMA N.V.	BE
Voltaren Retard 100 mg tabletten met verlengde afgifte.	not available	BE122071	NOVARTIS PHARMA N.V.	BE
Voltaren Retard 75 mg comprimés à libération prolongée.	not available	BE165471	NOVARTIS PHARMA N.V.	BE
Voltaren Retard 75 mg comprimés à libération prolongée.	not available	BE165471	NOVARTIS PHARMA N.V.	BE
Voltaren Retard 75 mg comprimés à libération prolongée.	not available	BE165471	NOVARTIS PHARMA N.V.	BE
Voltaren Retard 75 mg comprimés à libération prolongée.	not available	BE165471	NOVARTIS PHARMA N.V.	BE
Voltaren Retard 75 mg comprimés à libération prolongée.	not available	BE165471	NOVARTIS PHARMA N.V.	BE
Voltaren Retard 75 mg Retardtabletten	not available	BE165471	NOVARTIS PHARMA N.V.	BE
Voltaren Retard 75 mg Retardtabletten	not available	BE165471	NOVARTIS PHARMA N.V.	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Voltaren Retard 75 mg Retardtabletten	not available	BE165471	NOVARTIS PHARMA N.V.	BE
Voltaren Retard 75 mg Retardtabletten	not available	BE165471	NOVARTIS PHARMA N.V.	BE
Voltaren Retard 75 mg Retardtabletten	not available	BE165471	NOVARTIS PHARMA N.V.	BE
Voltaren Retard 75 mg tabletten met verlengde afgifte.	not available	BE165471	NOVARTIS PHARMA N.V.	BE
Voltaren Retard 75 mg tabletten met verlengde afgifte.	not available	BE165471	NOVARTIS PHARMA N.V.	BE
Voltaren Retard 75 mg tabletten met verlengde afgifte.	not available	BE165471	NOVARTIS PHARMA N.V.	BE
Voltaren Retard 75 mg tabletten met verlengde afgifte.	not available	BE165471	NOVARTIS PHARMA N.V.	BE
Voltaren Retard 75 mg tabletten met verlengde afgifte.	not available	BE165471	NOVARTIS PHARMA N.V.	BE
Волтарен 100 mg супозитории	not available	20020739	NOVARTIS EUROPHARM LIMITED	BG
Волтарен 100 mg супозитории	not available	20020739	NOVARTIS EUROPHARM LIMITED	BG
Волтарен 100 mg супозитории	not available	20020739	NOVARTIS EUROPHARM LIMITED	BG
Волтарен 100 mg супозитории	not available	20020739	NOVARTIS EUROPHARM LIMITED	BG
Волтарен 100 mg супозитории	not available	20020739	NOVARTIS EUROPHARM LIMITED	BG
Волтарен 25 mg стомашно-устойчиви таблетки	not available	20020736	NOVARTIS PHARMA GMBH	BG
Волтарен 25 mg стомашно-устойчиви таблетки	not available	20020736	NOVARTIS PHARMA GMBH	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
Волтарен 25 mg стомашно-устойчиви таблетки	not available	20020736	NOVARTIS PHARMA GMBH	BG
Волтарен 25 mg стомашно-устойчиви таблетки	not available	20020736	NOVARTIS PHARMA GMBH	BG
Волтарен 25 mg стомашно-устойчиви таблетки	not available	20020736	NOVARTIS PHARMA GMBH	BG
Волтарен 50 mg стомашно-устойчиви таблетки	not available	20020737	NOVARTIS EUROPHARM LIMITED	BG
Волтарен 50 mg стомашно-устойчиви таблетки	not available	20020737	NOVARTIS EUROPHARM LIMITED	BG
Волтарен 50 mg стомашно-устойчиви таблетки	not available	20020737	NOVARTIS EUROPHARM LIMITED	BG
Волтарен 50 mg стомашно-устойчиви таблетки	not available	20020737	NOVARTIS EUROPHARM LIMITED	BG
Волтарен 50 mg стомашно-устойчиви таблетки	not available	20020737	NOVARTIS EUROPHARM LIMITED	BG
Волтарен 50 mg супозитории	not available	20020738	NOVARTIS EUROPHARM LIMITED	BG
Волтарен 50 mg супозитории	not available	20020738	NOVARTIS EUROPHARM LIMITED	BG
Волтарен 50 mg супозитории	not available	20020738	NOVARTIS EUROPHARM LIMITED	BG
Волтарен 50 mg супозитории	not available	20020738	NOVARTIS EUROPHARM LIMITED	BG
Волтарен 50 mg супозитории	not available	20020738	NOVARTIS EUROPHARM LIMITED	BG
Волтарен Ретард 100 mg таблетки с удължено освобождаване	not available	20020740	NOVARTIS EUROPHARM LIMITED	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
Волтарен Ретард 100 mg таблетки с удължено освобождаване	not available	20020740	NOVARTIS EUROPHARM LIMITED	BG
Волтарен Ретард 100 mg таблетки с удължено освобождаване	not available	20020740	NOVARTIS EUROPHARM LIMITED	BG
Волтарен Ретард 100 mg таблетки с удължено освобождаване	not available	20020740	NOVARTIS EUROPHARM LIMITED	BG
Волтарен Ретард 100 mg таблетки с удължено освобождаване	not available	20020740	NOVARTIS EUROPHARM LIMITED	BG
CATAFLAM Sugar Coated Tablets 50mg	not available	18492	NOVARTIS IRELAND LIMITED	CY
CATAFLAM Sugar Coated Tablets 50mg	not available	18492	NOVARTIS IRELAND LIMITED	CY
CATAFLAM Sugar Coated Tablets 50mg	not available	18492	NOVARTIS IRELAND LIMITED	CY
CATAFLAM Sugar Coated Tablets 50mg	not available	18492	NOVARTIS IRELAND LIMITED	CY
CATAFLAM Sugar Coated Tablets 50mg	not available	18492	NOVARTIS IRELAND LIMITED	CY
Voltaren®	not available	18443	NOVARTIS IRELAND LIMITED	CY
Voltaren®	not available	18434	NOVARTIS IRELAND LIMITED	CY
Voltaren®	not available	18444	NOVARTIS IRELAND LIMITED	CY
Voltaren®	not available	18459	NOVARTIS IRELAND LIMITED	CY
Voltaren®	not available	18455	NOVARTIS IRELAND LIMITED	CY
Voltaren®	not available	18400	NOVARTIS IRELAND LIMITED	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
Voltaren®	not available	18457	NOVARTIS IRELAND LIMITED	CY
Voltaren®	not available	18443	NOVARTIS IRELAND LIMITED	CY
Voltaren®	not available	18434	NOVARTIS IRELAND LIMITED	CY
Voltaren®	not available	18444	NOVARTIS IRELAND LIMITED	CY
Voltaren®	not available	18459	NOVARTIS IRELAND LIMITED	CY
Voltaren®	not available	18455	NOVARTIS IRELAND LIMITED	CY
Voltaren®	not available	18400	NOVARTIS IRELAND LIMITED	CY
Voltaren®	not available	18457	NOVARTIS IRELAND LIMITED	CY
Voltaren®	not available	18443	NOVARTIS IRELAND LIMITED	CY
Voltaren®	not available	18434	NOVARTIS IRELAND LIMITED	CY
Voltaren®	not available	18444	NOVARTIS IRELAND LIMITED	CY
Voltaren®	not available	18459	NOVARTIS IRELAND LIMITED	CY
Voltaren®	not available	18455	NOVARTIS IRELAND LIMITED	CY
Voltaren®	not available	18400	NOVARTIS IRELAND LIMITED	CY
Voltaren®	not available	18457	NOVARTIS IRELAND LIMITED	CY
Voltaren®	not available	18443	NOVARTIS IRELAND LIMITED	CY
Voltaren®	not available	18434	NOVARTIS IRELAND LIMITED	CY
Voltaren®	not available	18444	NOVARTIS IRELAND LIMITED	CY
Voltaren®	not available	18459	NOVARTIS IRELAND LIMITED	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
Voltaren®	not available	18455	NOVARTIS IRELAND LIMITED	CY
Voltaren®	not available	18400	NOVARTIS IRELAND LIMITED	CY
Voltaren®	not available	18457	NOVARTIS IRELAND LIMITED	CY
Voltaren®	not available	18443	NOVARTIS IRELAND LIMITED	CY
Voltaren®	not available	18434	NOVARTIS IRELAND LIMITED	CY
Voltaren®	not available	18444	NOVARTIS IRELAND LIMITED	CY
Voltaren®	not available	18459	NOVARTIS IRELAND LIMITED	CY
Voltaren®	not available	18455	NOVARTIS IRELAND LIMITED	CY
Voltaren®	not available	18400	NOVARTIS IRELAND LIMITED	CY
Voltaren®	not available	18457	NOVARTIS IRELAND LIMITED	CY
Voltaren® for Children Suppositories 12.5mg	not available	18454	NOVARTIS IRELAND LIMITED	CY
Voltaren® for Children Suppositories 12.5mg	not available	18454	NOVARTIS IRELAND LIMITED	CY
Voltaren® for Children Suppositories 12.5mg	not available	18454	NOVARTIS IRELAND LIMITED	CY
Voltaren® for Children Suppositories 12.5mg	not available	18454	NOVARTIS IRELAND LIMITED	CY
Voltaren® for Children Suppositories 12.5mg	not available	18454	NOVARTIS IRELAND LIMITED	CY
Flector 50 mg granule pro perorální roztok v sáčku	not available	29/514/99-C	IBSA SLOVAKIA S.R.O.	CZ
UNO 150 mg tablety s prodlouženým uvolňováním	not available	29/216/00-C	RATIOPHARM GMBH	CZ
VOLTAREN 50 mg enterosolventní tablety	not available	29/294/91-C	NOVARTIS S.R.O.,	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
VOLTAREN 50 mg enterosolventní tablety	not available	29/294/91-C	NOVARTIS S.R.O.,	CZ
VOLTAREN 50 mg enterosolventní tablety	not available	29/294/91-C	NOVARTIS S.R.O.,	CZ
VOLTAREN 50 mg enterosolventní tablety	not available	29/294/91-C	NOVARTIS S.R.O.,	CZ
VOLTAREN 50 mg enterosolventní tablety	not available	29/294/91-C	NOVARTIS S.R.O.,	CZ
Voltaren Actigo Extra 25 mg obalené tablety	not available	29/549/00-C	HALEON CZECH REPUBLIC S.R.O.	CZ
Voltaren Rapid 25 mg měkké tobolky	not available	29/173/11-C	HALEON CZECH REPUBLIC S.R.O.	CZ
VOLTAREN RAPID 50 mg obalené tablety	not available	29/098/91-S/C	NOVARTIS S.R.O.,	CZ
VOLTAREN RAPID 50 mg obalené tablety	not available	29/098/91-S/C	NOVARTIS S.R.O.,	CZ
VOLTAREN RAPID 50 mg obalené tablety	not available	29/098/91-S/C	NOVARTIS S.R.O.,	CZ
VOLTAREN RAPID 50 mg obalené tablety	not available	29/098/91-S/C	NOVARTIS S.R.O.,	CZ
VOLTAREN RAPID 50 mg obalené tablety	not available	29/098/91-S/C	NOVARTIS S.R.O.,	CZ
VOLTAREN RETARD 100 mg tablety s prodlouženým uvolňováním	not available	29/247/80-C	NOVARTIS S.R.O.,	CZ
VOLTAREN RETARD 100 mg tablety s prodlouženým uvolňováním	not available	29/247/80-C	NOVARTIS S.R.O.,	CZ
VOLTAREN RETARD 100 mg tablety s prodlouženým uvolňováním	not available	29/247/80-C	NOVARTIS S.R.O.,	CZ
VOLTAREN RETARD 100 mg tablety s prodlouženým uvolňováním	not available	29/247/80-C	NOVARTIS S.R.O.,	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
VOLTAREN RETARD 100 mg tablety s prodlouženým uvolňováním	not available	29/247/80-C	NOVARTIS S.R.O.,	CZ
Diclo-Divido long: Hartkapseln, retardiert Wirkstoff: Diclofenac-Natrium 100 mg	not available	13847.00.00	GLENWOOD GMBH	DE
Diclo-Divido: Hartkapseln, retardiert Wirkstoff: Diclofenac-Natrium 75 mg	not available	13847.01.00	GLENWOOD GMBH	DE
Diclofenac Zentiva 25 mg Filmtabletten	not available	74529.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	95765.00.00	MICRO LABS GMBH	DE
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	95765.00.00	MICRO LABS GMBH	DE
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	95765.00.00	MICRO LABS GMBH	DE
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	95765.00.00	MICRO LABS GMBH	DE
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	95765.00.00	MICRO LABS GMBH	DE
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	95765.00.00	MICRO LABS GMBH	DE
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	95765.00.00	MICRO LABS GMBH	DE
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	95765.00.00	MICRO LABS 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
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	95765.00.00	MICRO LABS GMBH	DE
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	95765.00.00	MICRO LABS GMBH	DE
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	95765.00.00	MICRO LABS GMBH	DE
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	95765.00.00	MICRO LABS GMBH	DE
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	95765.00.00	MICRO LABS GMBH	DE
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	95765.00.00	MICRO LABS GMBH	DE
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	95765.00.00	MICRO LABS GMBH	DE
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	95765.00.00	MICRO LABS GMBH	DE
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	95765.00.00	MICRO LABS GMBH	DE
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	95765.00.00	MICRO LABS GMBH	DE
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	95765.00.00	MICRO LABS GMBH	DE
Diclo-ratiopharm® bei Schmerzen und Fieber 25 mg Filmtabletten	not available	74527.00.00	RATIOPHARM GMBH	DE
Voltaren Dolo 25 mg überzogene Tablette	not available	64958.00.00	HALEON GERMANY 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
Voltaren Dolo 25 mg überzogene Tablette	not available	43151.00.00	HALEON GERMANY GMBH	DE
Voltaren Dolo Liquid 25 mg Weichkapsel	not available	82502.00.00	HALEON GERMANY GMBH	DE
Voltaren® 100 mg retard Retardtabletten	not available	14651.00.01	NOVARTIS PHARMA GMBH	DE
Voltaren® 100 mg retard Retardtabletten	not available	14651.00.01	NOVARTIS PHARMA GMBH	DE
Voltaren® 100 mg retard Retardtabletten	not available	14651.00.01	NOVARTIS PHARMA GMBH	DE
Voltaren® 100 mg retard Retardtabletten	not available	14651.00.01	NOVARTIS PHARMA GMBH	DE
Voltaren® 100 mg retard Retardtabletten	not available	14651.00.01	NOVARTIS PHARMA GMBH	DE
Voltaren® 100 mg retard Retardtabletten	not available	14651.00.01	NOVARTIS PHARMA GMBH	DE
Voltaren® 100 mg Zäpfchen	not available	14651.00.02	NOVARTIS PHARMA GMBH	DE
Voltaren® 100 mg Zäpfchen	not available	14651.00.02	NOVARTIS PHARMA GMBH	DE
Voltaren® 100 mg Zäpfchen	not available	14651.00.02	NOVARTIS PHARMA GMBH	DE
Voltaren® 100 mg Zäpfchen	not available	14651.00.02	NOVARTIS PHARMA GMBH	DE
Voltaren® 100 mg Zäpfchen	not available	14651.00.02	NOVARTIS PHARMA GMBH	DE
Voltaren® 100 mg Zäpfchen	not available	14651.00.02	NOVARTIS PHARMA GMBH	DE
Voltaren® 25 mg Zäpfchen	not available	520.00.00	NOVARTIS PHARMA GMBH	DE
Voltaren® 25 mg Zäpfchen	not available	520.00.00	NOVARTIS PHARMA GMBH	DE
Voltaren® 25 mg Zäpfchen	not available	520.00.00	NOVARTIS PHARMA GMBH	DE
Voltaren® 25 mg Zäpfchen	not available	520.00.00	NOVARTIS PHARMA 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
Voltaren® 25 mg Zäpfchen	not available	520.00.00	NOVARTIS PHARMA GMBH	DE
Voltaren® 25 mg Zäpfchen	not available	520.00.00	NOVARTIS PHARMA GMBH	DE
Voltaren® 50 mg Zäpfchen	not available	6164405.00.00	NOVARTIS PHARMA GMBH	DE
Voltaren® 50 mg Zäpfchen	not available	6164405.00.00	NOVARTIS PHARMA GMBH	DE
Voltaren® 50 mg Zäpfchen	not available	6164405.00.00	NOVARTIS PHARMA GMBH	DE
Voltaren® 50 mg Zäpfchen	not available	6164405.00.00	NOVARTIS PHARMA GMBH	DE
Voltaren® 50 mg Zäpfchen	not available	6164405.00.00	NOVARTIS PHARMA GMBH	DE
Voltaren® 50 mg Zäpfchen	not available	6164405.00.00	NOVARTIS PHARMA GMBH	DE
Voltaren® Dispers 46,5 mg Tabletten zur Herstellung einer Suspension zum Einnehmen	not available	6164351.00.00	NOVARTIS PHARMA GMBH	DE
Voltaren® Dispers 46,5 mg Tabletten zur Herstellung einer Suspension zum Einnehmen	not available	6164351.00.00	NOVARTIS PHARMA GMBH	DE
Voltaren® Dispers 46,5 mg Tabletten zur Herstellung einer Suspension zum Einnehmen	not available	6164351.00.00	NOVARTIS PHARMA GMBH	DE
Voltaren® Dispers 46,5 mg Tabletten zur Herstellung einer Suspension zum Einnehmen	not available	6164351.00.00	NOVARTIS PHARMA 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
Voltaren® Dispers 46,5 mg Tabletten zur Herstellung einer Suspension zum Einnehmen	not available	6164351.00.00	NOVARTIS PHARMA GMBH	DE
Voltaren® Dispers 46,5 mg Tabletten zur Herstellung einer Suspension zum Einnehmen	not available	6164351.00.00	NOVARTIS PHARMA GMBH	DE
Voltaren® Resinat 145,6 mg Diclofenac-Colestyramin (entsprechend 75 mg Diclofenac-Natrium) Hartkapseln	not available	17982.00.00	NOVARTIS PHARMA GMBH	DE
Voltaren® Resinat 145,6 mg Diclofenac-Colestyramin (entsprechend 75 mg Diclofenac-Natrium) Hartkapseln	not available	17982.00.00	NOVARTIS PHARMA GMBH	DE
Voltaren® Resinat 145,6 mg Diclofenac-Colestyramin (entsprechend 75 mg Diclofenac-Natrium) Hartkapseln	not available	17982.00.00	NOVARTIS PHARMA GMBH	DE
Voltaren® Resinat 145,6 mg Diclofenac-Colestyramin (entsprechend 75 mg Diclofenac-Natrium) Hartkapseln	not available	17982.00.00	NOVARTIS PHARMA 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
Voltaren® Resinat 145,6 mg Diclofenac-Colestyramin (entsprechend 75 mg Diclofenac-Natrium) Hartkapseln	not available	17982.00.00	NOVARTIS PHARMA GMBH	DE
Voltaren Akti, 12,5 mg õhukese polümeerikattega tabletid	not available	434104	HALEON HUNGARY KFT.	EE
Voltaren Akti, 12,5 mg õhukese polümeerikattega tabletid	not available	434104	HALEON HUNGARY KFT.	EE
Akis 25 mg solución inyectable en jeringa precargada	HU/H/0796/004	83369	IBSA FARMACEUTICI ITALIA	ES
Akis 50 mg solución inyectable en jeringa precargada	HU/H/0796/005	83370	IBSA FARMACEUTICI ITALIA	ES
Akis 75 mg solución inyectable en jeringa precargada	HU/H/0796/006	83372	IBSA FARMACEUTICI ITALIA	ES
Dolotren 100 mg supositorios	not available	57.203	FAES FARMA, S.A.	ES
Dolotren 46,5 mg comprimidos dispersables	not available	60.068	FAES FARMA, S.A.	ES
Dolotren 50 mg comprimidos gastrorresistentes.	not available	57.202	FAES FARMA, S.A.	ES
Dolotren 75 mg solución inyectable	not available	57.204	FAES FARMA, S.A.	ES
Dolotren retard 100 mg cápsulas duras de liberación prolongada	not available	58.435	FAES FARMA, S.A.	ES
Dolo-Voltarén 46,5 mg comprimidos dispersables	not available	61.440	NOVARTIS FARMACÉUTICA S.A.	ES

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Dolo-Voltarén 46,5 mg comprimidos dispersables	not available	61.440	NOVARTIS FARMACÉUTICA S.A.	ES
Dolo-Voltarén 46,5 mg comprimidos dispersables	not available	61.440	NOVARTIS FARMACÉUTICA S.A.	ES
Dolo-Voltarén 46,5 mg comprimidos dispersables	not available	61.440	NOVARTIS FARMACÉUTICA S.A.	ES
Dolo-Voltarén 46,5 mg comprimidos dispersables	not available	61.440	NOVARTIS FARMACÉUTICA S.A.	ES
Voltarén 100 mg supositorios	not available	55.004	NOVARTIS FARMACÉUTICA S.A.	ES
Voltarén 100 mg supositorios	not available	55.004	NOVARTIS FARMACÉUTICA S.A.	ES
Voltarén 100 mg supositorios	not available	55.004	NOVARTIS FARMACÉUTICA S.A.	ES
Voltarén 100 mg supositorios	not available	55.004	NOVARTIS FARMACÉUTICA S.A.	ES
Voltarén 100 mg supositorios	not available	55.004	NOVARTIS FARMACÉUTICA S.A.	ES
Voltarén 50 mg comprimidos gastrorresistentes	not available	55.005	NOVARTIS FARMACÉUTICA S.A.	ES
Voltarén 50 mg comprimidos gastrorresistentes	not available	55.005	NOVARTIS FARMACÉUTICA S.A.	ES
Voltarén 50 mg comprimidos gastrorresistentes	not available	55.005	NOVARTIS FARMACÉUTICA S.A.	ES
Voltarén 50 mg comprimidos gastrorresistentes	not available	55.005	NOVARTIS FARMACÉUTICA S.A.	ES
Voltarén 50 mg comprimidos gastrorresistentes	not available	55.005	NOVARTIS FARMACÉUTICA S.A.	ES

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Voltarén 75 mg solución inyectable	not available	55.010	NOVARTIS FARMACÉUTICA S.A.	ES
Voltarén 75 mg solución inyectable	not available	55.010	NOVARTIS FARMACÉUTICA S.A.	ES
Voltarén 75 mg solución inyectable	not available	55.010	NOVARTIS FARMACÉUTICA S.A.	ES
Voltarén 75 mg solución inyectable	not available	55.010	NOVARTIS FARMACÉUTICA S.A.	ES
Voltarén 75 mg solución inyectable	not available	55.010	NOVARTIS FARMACÉUTICA S.A.	ES
Voltarén Retard 100 mg comprimidos de liberación modificada	not available	56.562	NOVARTIS FARMACÉUTICA S.A.	ES
Voltarén Retard 100 mg comprimidos de liberación modificada	not available	56.562	NOVARTIS FARMACÉUTICA S.A.	ES
Voltarén Retard 100 mg comprimidos de liberación modificada	not available	56.562	NOVARTIS FARMACÉUTICA S.A.	ES
Voltarén Retard 100 mg comprimidos de liberación modificada	not available	56.562	NOVARTIS FARMACÉUTICA S.A.	ES
Voltarén Retard 100 mg comprimidos de liberación modificada	not available	56.562	NOVARTIS FARMACÉUTICA S.A.	ES
Voltarén Retard 75 mg comprimidos de liberación modificada	not available	62.024	NOVARTIS FARMACÉUTICA S.A.	ES
Voltarén Retard 75 mg comprimidos de liberación modificada	not available	62.024	NOVARTIS FARMACÉUTICA S.A.	ES
Voltarén Retard 75 mg comprimidos de liberación modificada	not available	62.024	NOVARTIS FARMACÉUTICA S.A.	ES
Voltarén Retard 75 mg comprimidos de liberación modificada	not available	62.024	NOVARTIS FARMACÉUTICA S.A.	ES

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Voltarén Retard 75 mg comprimidos de liberación modificada	not available	62.024	NOVARTIS FARMACÉUTICA S.A.	ES
Voltaren 100 mg peräpuikot	not available	8090	NOVARTIS FINLAND OY	FI
Voltaren 100 mg peräpuikot	not available	8090	NOVARTIS FINLAND OY	FI
Voltaren 100 mg peräpuikot	not available	8090	NOVARTIS FINLAND OY	FI
Voltaren 100 mg peräpuikot	not available	8090	NOVARTIS FINLAND OY	FI
Voltaren 100 mg peräpuikot	not available	8090	NOVARTIS FINLAND OY	FI
Voltaren 100 mg peräpuikot	not available	8090	NOVARTIS FINLAND OY	FI
Voltaren 100 mg suppositorier	not available	8090	NOVARTIS FINLAND OY	FI
Voltaren 100 mg suppositorier	not available	8090	NOVARTIS FINLAND OY	FI
Voltaren 100 mg suppositorier	not available	8090	NOVARTIS FINLAND OY	FI
Voltaren 100 mg suppositorier	not available	8090	NOVARTIS FINLAND OY	FI
Voltaren 100 mg suppositorier	not available	8090	NOVARTIS FINLAND OY	FI
Voltaren 100 mg suppositorier	not available	8090	NOVARTIS FINLAND OY	FI
Voltaren 25 mg/ml injektio-/infuusioneste, liuos	not available	8629	NOVARTIS FINLAND OY	FI
Voltaren 25 mg/ml injektio-/infuusioneste, liuos	not available	8629	NOVARTIS FINLAND OY	FI
Voltaren 25 mg/ml injektio-/infuusioneste, liuos	not available	8629	NOVARTIS FINLAND 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
Voltaren 25 mg/ml injektio-/infuusioneste, liuos	not available	8629	NOVARTIS FINLAND OY	FI
Voltaren 25 mg/ml injektio-/infuusioneste, liuos	not available	8629	NOVARTIS FINLAND OY	FI
Voltaren 25 mg/ml injektio-/infuusioneste, liuos	not available	8629	NOVARTIS FINLAND OY	FI
Voltaren 25 mg/ml injektions-/infusionsvätska, lösning	not available	8629	NOVARTIS FINLAND OY	FI
Voltaren 25 mg/ml injektions-/infusionsvätska, lösning	not available	8629	NOVARTIS FINLAND OY	FI
Voltaren 25 mg/ml injektions-/infusionsvätska, lösning	not available	8629	NOVARTIS FINLAND OY	FI
Voltaren 25 mg/ml injektions-/infusionsvätska, lösning	not available	8629	NOVARTIS FINLAND OY	FI
Voltaren 25 mg/ml injektions-/infusionsvätska, lösning	not available	8629	NOVARTIS FINLAND OY	FI
Voltaren 25 mg/ml injektions-/infusionsvätska, lösning	not available	8629	NOVARTIS FINLAND OY	FI
Voltaren 50 mg enterotabletit	not available	7906	NOVARTIS FINLAND OY	FI
Voltaren 50 mg enterotabletit	not available	7906	NOVARTIS FINLAND OY	FI
Voltaren 50 mg enterotabletit	not available	7906	NOVARTIS FINLAND OY	FI
Voltaren 50 mg enterotabletit	not available	7906	NOVARTIS FINLAND OY	FI
Voltaren 50 mg enterotabletit	not available	7906	NOVARTIS FINLAND 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
Voltaren 50 mg enterotabletit	not available	7906	NOVARTIS FINLAND OY	FI
Voltaren 50 mg enterotabletter	not available	7906	NOVARTIS FINLAND OY	FI
Voltaren 50 mg enterotabletter	not available	7906	NOVARTIS FINLAND OY	FI
Voltaren 50 mg enterotabletter	not available	7906	NOVARTIS FINLAND OY	FI
Voltaren 50 mg enterotabletter	not available	7906	NOVARTIS FINLAND OY	FI
Voltaren 50 mg enterotabletter	not available	7906	NOVARTIS FINLAND OY	FI
Voltaren 50 mg enterotabletter	not available	7906	NOVARTIS FINLAND OY	FI
Voltaren Rapid 50 mg dragerade tabletter	not available	10203	NOVARTIS FINLAND OY	FI
Voltaren Rapid 50 mg dragerade tabletter	not available	10203	NOVARTIS FINLAND OY	FI
Voltaren Rapid 50 mg dragerade tabletter	not available	10203	NOVARTIS FINLAND OY	FI
Voltaren Rapid 50 mg dragerade tabletter	not available	10203	NOVARTIS FINLAND OY	FI
Voltaren Rapid 50 mg dragerade tabletter	not available	10203	NOVARTIS FINLAND OY	FI
Voltaren Rapid 50 mg päällystetyt tabletit	not available	10203	NOVARTIS FINLAND OY	FI
Voltaren Rapid 50 mg päällystetyt tabletit	not available	10203	NOVARTIS FINLAND OY	FI
Voltaren Rapid 50 mg päällystetyt tabletit	not available	10203	NOVARTIS FINLAND OY	FI
Voltaren Rapid 50 mg päällystetyt tabletit	not available	10203	NOVARTIS FINLAND OY	FI
Voltaren Rapid 50 mg päällystetyt tabletit	not available	10203	NOVARTIS FINLAND OY	FI
Voltaren Retard 100 mg depottabletit	not available	9228	NOVARTIS FINLAND OY	FI
Voltaren Retard 100 mg depottabletit	not available	9228	NOVARTIS FINLAND 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
Voltaren Retard 100 mg depottabletit	not available	9228	NOVARTIS FINLAND OY	FI
Voltaren Retard 100 mg depottabletit	not available	9228	NOVARTIS FINLAND OY	FI
Voltaren Retard 100 mg depottabletit	not available	9228	NOVARTIS FINLAND OY	FI
Voltaren Retard 100 mg depottabletit	not available	9228	NOVARTIS FINLAND OY	FI
Voltaren Retard 100 mg depottabletter	not available	9228	NOVARTIS FINLAND OY	FI
Voltaren Retard 100 mg depottabletter	not available	9228	NOVARTIS FINLAND OY	FI
Voltaren Retard 100 mg depottabletter	not available	9228	NOVARTIS FINLAND OY	FI
Voltaren Retard 100 mg depottabletter	not available	9228	NOVARTIS FINLAND OY	FI
Voltaren Retard 100 mg depottabletter	not available	9228	NOVARTIS FINLAND OY	FI
Voltaren Retard 100 mg depottabletter	not available	9228	NOVARTIS FINLAND OY	FI
Voltaren Retard 75 mg depottabletit	not available	10919	NOVARTIS FINLAND OY	FI
Voltaren Retard 75 mg depottabletit	not available	10919	NOVARTIS FINLAND OY	FI
Voltaren Retard 75 mg depottabletit	not available	10919	NOVARTIS FINLAND OY	FI
Voltaren Retard 75 mg depottabletit	not available	10919	NOVARTIS FINLAND OY	FI
Voltaren Retard 75 mg depottabletit	not available	10919	NOVARTIS FINLAND OY	FI
Voltaren Retard 75 mg depottabletit	not available	10919	NOVARTIS FINLAND OY	FI
Voltaren Retard 75 mg depottabletter	not available	10919	NOVARTIS FINLAND OY	FI
Voltaren Retard 75 mg depottabletter	not available	10919	NOVARTIS FINLAND OY	FI
Voltaren Retard 75 mg depottabletter	not available	10919	NOVARTIS FINLAND 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
Voltaren Retard 75 mg depottabletter	not available	10919	NOVARTIS FINLAND OY	FI
Voltaren Retard 75 mg depottabletter	not available	10919	NOVARTIS FINLAND OY	FI
Voltaren Retard 75 mg depottabletter	not available	10919	NOVARTIS FINLAND OY	FI
AKIS 25 mg/ml, solution injectable	HU/H/0796/001	34009 301 865 2 7	IBSA PHARMA SAS	FR
AKIS 25 mg/ml, solution injectable	HU/H/0796/001	34009 301 865 3 4	IBSA PHARMA SAS	FR
AKIS 25 mg/ml, solution injectable	HU/H/0796/001	34009 301 865 4 1	IBSA PHARMA SAS	FR
AKIS 25 mg/ml, solution injectable en seringue préremplie	HU/H/0796/004	34009 301 866 2 6	IBSA PHARMA SAS	FR
AKIS 25 mg/ml, solution injectable en seringue préremplie	HU/H/0796/004	34009 301 866 3 3	IBSA PHARMA SAS	FR
AKIS 25 mg/ml, solution injectable en seringue préremplie	HU/H/0796/004	34009 301 866 4 0	IBSA PHARMA SAS	FR
AKIS 50 mg/ml, solution injectable	HU/H/0796/002	34009 301 865 5 8	IBSA PHARMA SAS	FR
AKIS 50 mg/ml, solution injectable	HU/H/0796/002	34009 301 865 6 5	IBSA PHARMA SAS	FR
AKIS 50 mg/ml, solution injectable	HU/H/0796/002	34009 301 865 7 2	IBSA PHARMA SAS	FR
AKIS 50 mg/ml, solution injectable en seringue préremplie	HU/H/0796/005	34009 301 866 5 7	IBSA PHARMA SAS	FR
AKIS 50 mg/ml, solution injectable en seringue préremplie	HU/H/0796/005	34009 301 866 8 8	IBSA PHARMA SAS	FR
AKIS 50 mg/ml, solution injectable en seringue préremplie	HU/H/0796/005	34009 301 866 6 4	IBSA PHARMA SAS	FR
AKIS 75 mg/ml, solution injectable	HU/H/0796/003	34009 301 865 8 9	IBSA PHARMA SAS	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
AKIS 75 mg/ml, solution injectable	HU/H/0796/003	34009 301 865 9 6	IBSA PHARMA SAS	FR
AKIS 75 mg/ml, solution injectable	HU/H/0796/003	34009 301 866 0 2	IBSA PHARMA SAS	FR
AKIS 75 mg/ml, solution injectable en seringue préremplie	HU/H/0796/006	34009 301 866 9 5	IBSA PHARMA SAS	FR
AKIS 75 mg/ml, solution injectable en seringue préremplie	HU/H/0796/006	34009 301 867 0 1	IBSA PHARMA SAS	FR
AKIS 75 mg/ml, solution injectable en seringue préremplie	HU/H/0796/006	34009 301 867 1 8	IBSA PHARMA SAS	FR
FLECTOR 25 mg, granulés pour solution buvable en sachet-dose	not available	34009 352 615 1 9	IBSA PHARMA SAS	FR
FLECTOR 25 mg, granulés pour solution buvable en sachet-dose	not available	34009 352 641 2 1	IBSA PHARMA SAS	FR
FLECTOR 25 mg, granulés pour solution buvable en sachet-dose	not available	34009 352 616 8 7	IBSA PHARMA SAS	FR
FLECTOR 50 mg, comprimé	not available	34009 301 460 6 4	IBSA PHARMA SAS	FR
FLECTOR 50 mg, granulés pour solution buvable en sachet-dose	not available	34009 352 617 4 8	IBSA PHARMA SAS	FR
FLECTOR 50 mg, granulés pour solution buvable en sachet-dose	not available	34009 352 642 9 9	IBSA PHARMA SAS	FR
FLECTOR 50 mg, granulés pour solution buvable en sachet-dose	not available	34009 352 618 0 9	IBSA PHARMA SAS	FR
VAFOTENA LP 75 mg, comprimé pelliculé à libération prolongée	DE/H/4552/001	34009 302 406 9 4	MICRO LABS GMBH	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
VAFOTENA LP 75 mg, comprimé pelliculé à libération prolongée	DE/H/4552/001	34009 302 407 0 0	MICRO LABS GMBH	FR
VAFOTENA LP 75 mg, comprimé pelliculé à libération prolongée	DE/H/4552/001	34009 302 407 1 7	MICRO LABS GMBH	FR
VAFOTENA LP 75 mg, comprimé pelliculé à libération prolongée	DE/H/4552/001	34009 302 407 2 4	MICRO LABS GMBH	FR
VAFOTENA LP 75 mg, comprimé pelliculé à libération prolongée	DE/H/4552/001	34009 302 407 3 1	MICRO LABS GMBH	FR
VAFOTENA LP 75 mg, comprimé pelliculé à libération prolongée	DE/H/4552/001	34009 302 407 4 8	MICRO LABS GMBH	FR
VAFOTENA LP 75 mg, comprimé pelliculé à libération prolongée	DE/H/4552/001	34009 302 407 5 5	MICRO LABS GMBH	FR
VAFOTENA LP 75 mg, comprimé pelliculé à libération prolongée	DE/H/4552/001	34009 302 407 7 9	MICRO LABS GMBH	FR
VAFOTENA LP 75 mg, comprimé pelliculé à libération prolongée	DE/H/4552/001	DE/H/4552/001	MICRO LABS GMBH	FR
VAFOTENA LP 75 mg, comprimé pelliculé à libération prolongée	DE/H/4552/001	34009 302 407 8 6	MICRO LABS GMBH	FR
VAFOTENA LP 75 mg, comprimé pelliculé à libération prolongée	DE/H/4552/001	34009 302 407 9 3	MICRO LABS GMBH	FR
VAFOTENA LP 75 mg, comprimé pelliculé à libération prolongée	DE/H/4552/001	34009 302 408 0 9	MICRO LABS GMBH	FR
VAFOTENA LP 75 mg, comprimé pelliculé à libération prolongée	DE/H/4552/001	34009 302 408 1 6	MICRO LABS GMBH	FR
VAFOTENA LP 75 mg, comprimé pelliculé à libération prolongée	DE/H/4552/001	34009 302 408 3 0	MICRO LABS GMBH	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
VAFOTENA LP 75 mg, comprimé pelliculé à libération prolongée	DE/H/4552/001	34009 302 408 4 7	MICRO LABS GMBH	FR
VAFOTENA LP 75 mg, comprimé pelliculé à libération prolongée	DE/H/4552/001	34009 302 408 5 4	MICRO LABS GMBH	FR
VAFOTENA LP 75 mg, comprimé pelliculé à libération prolongée	DE/H/4552/001	34009 302 408 6 1	MICRO LABS GMBH	FR
VAFOTENA LP 75 mg, comprimé pelliculé à libération prolongée	DE/H/4552/001	34009 550 851 7 1	MICRO LABS GMBH	FR
VAFOTENA LP 75 mg, comprimé pelliculé à libération prolongée	DE/H/4552/001	34009 550 851 8 8	MICRO LABS GMBH	FR
VOLTARENE 100 mg, suppositoire	not available	34009 322 143 4 1	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE 100 mg, suppositoire	not available	34009 322 143 4 1	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE 100 mg, suppositoire	not available	34009 322 143 4 1	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE 100 mg, suppositoire	not available	34009 322 143 4 1	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE 100 mg, suppositoire	not available	34009 322 143 4 1	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE 25 mg, comprimé enrobé gastro-résistant	not available	34009 318 952 9 9	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE 25 mg, comprimé enrobé gastro-résistant	not available	34009 319 787 1 8	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE 25 mg, comprimé enrobé gastro-résistant	not available	34009 338 144 5 8	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE 25 mg, comprimé enrobé gastro-résistant	not available	34009 338 145 1 9	NOVARTIS PHARMA S.A.S.	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
VOLTARENE 25 mg, comprimé enrobé gastro-résistant	not available	34009 318 952 9 9	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE 25 mg, comprimé enrobé gastro-résistant	not available	34009 319 787 1 8	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE 25 mg, comprimé enrobé gastro-résistant	not available	34009 338 144 5 8	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE 25 mg, comprimé enrobé gastro-résistant	not available	34009 338 145 1 9	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE 25 mg, comprimé enrobé gastro-résistant	not available	34009 318 952 9 9	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE 25 mg, comprimé enrobé gastro-résistant	not available	34009 319 787 1 8	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE 25 mg, comprimé enrobé gastro-résistant	not available	34009 338 144 5 8	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE 25 mg, comprimé enrobé gastro-résistant	not available	34009 338 145 1 9	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE 25 mg, comprimé enrobé gastro-résistant	not available	34009 318 952 9 9	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE 25 mg, comprimé enrobé gastro-résistant	not available	34009 319 787 1 8	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE 25 mg, comprimé enrobé gastro-résistant	not available	34009 338 144 5 8	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE 25 mg, comprimé enrobé gastro-résistant	not available	34009 338 145 1 9	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE 25 mg, comprimé enrobé gastro-résistant	not available	34009 318 952 9 9	NOVARTIS PHARMA S.A.S.	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
VOLTARENE 25 mg, comprimé enrobé gastro-résistant	not available	34009 319 787 1 8	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE 25 mg, comprimé enrobé gastro-résistant	not available	34009 338 144 5 8	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE 25 mg, comprimé enrobé gastro-résistant	not available	34009 338 145 1 9	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE 50 mg, comprimé enrobé gastro-résistant	not available	34009 323 511 7 6	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE 50 mg, comprimé enrobé gastro-résistant	not available	34009 323 511 7 6	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE 50 mg, comprimé enrobé gastro-résistant	not available	34009 323 511 7 6	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE 50 mg, comprimé enrobé gastro-résistant	not available	34009 323 511 7 6	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE 50 mg, comprimé enrobé gastro-résistant	not available	34009 323 511 7 6	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE LP 100 mg, comprimé enrobé à libération prolongée	not available	34009 324 604 9 6	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE LP 100 mg, comprimé enrobé à libération prolongée	not available	34009 324 605 5 7	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE LP 100 mg, comprimé enrobé à libération prolongée	not available	34009 330 701 2 0	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE LP 100 mg, comprimé enrobé à libération prolongée	not available	34009 330 702 9 8	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE LP 100 mg, comprimé enrobé à libération prolongée	not available	34009 324 604 9 6	NOVARTIS PHARMA S.A.S.	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
VOLTARENE LP 100 mg, comprimé enrobé à libération prolongée	not available	34009 324 605 5 7	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE LP 100 mg, comprimé enrobé à libération prolongée	not available	34009 330 701 2 0	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE LP 100 mg, comprimé enrobé à libération prolongée	not available	34009 330 702 9 8	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE LP 100 mg, comprimé enrobé à libération prolongée	not available	34009 324 604 9 6	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE LP 100 mg, comprimé enrobé à libération prolongée	not available	34009 324 605 5 7	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE LP 100 mg, comprimé enrobé à libération prolongée	not available	34009 330 701 2 0	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE LP 100 mg, comprimé enrobé à libération prolongée	not available	34009 330 702 9 8	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE LP 100 mg, comprimé enrobé à libération prolongée	not available	34009 324 604 9 6	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE LP 100 mg, comprimé enrobé à libération prolongée	not available	34009 324 605 5 7	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE LP 100 mg, comprimé enrobé à libération prolongée	not available	34009 330 701 2 0	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE LP 100 mg, comprimé enrobé à libération prolongée	not available	34009 330 702 9 8	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE LP 100 mg, comprimé enrobé à libération prolongée	not available	34009 324 604 9 6	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE LP 100 mg, comprimé enrobé à libération prolongée	not available	34009 324 605 5 7	NOVARTIS PHARMA S.A.S.	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
VOLTARENE LP 100 mg, comprimé enrobé à libération prolongée	not available	34009 330 701 2 0	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE LP 100 mg, comprimé enrobé à libération prolongée	not available	34009 330 702 9 8	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE LP 75 mg, comprimé enrobé à libération prolongée	not available	34009 335 919 6 0	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE LP 75 mg, comprimé enrobé à libération prolongée	not available	34009 335 920 4 2	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE LP 75 mg, comprimé enrobé à libération prolongée	not available	34009 335 934 5 2	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE LP 75 mg, comprimé enrobé à libération prolongée	not available	34009 335 935 1 3	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE LP 75 mg, comprimé enrobé à libération prolongée	not available	34009 345 956 1 5	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE LP 75 mg, comprimé enrobé à libération prolongée	not available	34009 335 919 6 0	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE LP 75 mg, comprimé enrobé à libération prolongée	not available	34009 335 920 4 2	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE LP 75 mg, comprimé enrobé à libération prolongée	not available	34009 335 934 5 2	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE LP 75 mg, comprimé enrobé à libération prolongée	not available	34009 335 935 1 3	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE LP 75 mg, comprimé enrobé à libération prolongée	not available	34009 345 956 1 5	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE LP 75 mg, comprimé enrobé à libération prolongée	not available	34009 335 919 6 0	NOVARTIS PHARMA S.A.S.	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
VOLTARENE LP 75 mg, comprimé enrobé à libération prolongée	not available	34009 335 920 4 2	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE LP 75 mg, comprimé enrobé à libération prolongée	not available	34009 335 934 5 2	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE LP 75 mg, comprimé enrobé à libération prolongée	not available	34009 335 935 1 3	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE LP 75 mg, comprimé enrobé à libération prolongée	not available	34009 345 956 1 5	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE LP 75 mg, comprimé enrobé à libération prolongée	not available	34009 335 919 6 0	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE LP 75 mg, comprimé enrobé à libération prolongée	not available	34009 335 920 4 2	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE LP 75 mg, comprimé enrobé à libération prolongée	not available	34009 335 934 5 2	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE LP 75 mg, comprimé enrobé à libération prolongée	not available	34009 335 935 1 3	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE LP 75 mg, comprimé enrobé à libération prolongée	not available	34009 345 956 1 5	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE LP 75 mg, comprimé enrobé à libération prolongée	not available	34009 335 919 6 0	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE LP 75 mg, comprimé enrobé à libération prolongée	not available	34009 335 920 4 2	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE LP 75 mg, comprimé enrobé à libération prolongée	not available	34009 335 934 5 2	NOVARTIS PHARMA S.A.S.	FR
VOLTARENE LP 75 mg, comprimé enrobé à libération prolongée	not available	34009 335 935 1 3	NOVARTIS PHARMA S.A.S.	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
VOLTARENE LP 75 mg, comprimé enrobé à libération prolongée	not available	34009 345 956 1 5	NOVARTIS PHARMA S.A.S.	FR
Voltaren 100 mg δισκία παρατεταμένης αποδέσμευσης	not available	122880301	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren 100 mg δισκία παρατεταμένης αποδέσμευσης	not available	122880301	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren 100 mg δισκία παρατεταμένης αποδέσμευσης	not available	122880301	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren 100 mg δισκία παρατεταμένης αποδέσμευσης	not available	122880301	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren 100 mg δισκία παρατεταμένης αποδέσμευσης	not available	122880301	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren 100 mg δισκία παρατεταμένης αποδέσμευσης	not available	122880301	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren 50 mg γαστροανθεκτικά δισκία	not available	122880201	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren 50 mg γαστροανθεκτικά δισκία	not available	122880202	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren 50 mg γαστροανθεκτικά δισκία	not available	122880201	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren 50 mg γαστροανθεκτικά δισκία	not available	122880202	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren 50 mg γαστροανθεκτικά δισκία	not available	122880201	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren 50 mg γαστροανθεκτικά δισκία	not available	122880202	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren 50 mg γαστροανθεκτικά δισκία	not available	122880201	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren 50 mg γαστροανθεκτικά δισκία	not available	122880202	NOVARTIS (HELLAS) S.A.C.I.	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
Voltaren 50 mg γαστροανθεκτικά δισκία	not available	122880201	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren 50 mg γαστροανθεκτικά δισκία	not available	122880202	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren 50 mg γαστροανθεκτικά δισκία	not available	122880201	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren 50 mg γαστροανθεκτικά δισκία	not available	122880202	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren 50 mg διασπειρόμενα δισκία	not available	122880901	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren 50 mg διασπειρόμενα δισκία	not available	122880901	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren 50 mg διασπειρόμενα δισκία	not available	122880901	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren 50 mg διασπειρόμενα δισκία	not available	122880901	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren 50 mg διασπειρόμενα δισκία	not available	122880901	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren 50 mg διασπειρόμενα δισκία	not available	122880901	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren 50 mg υπόθετα	not available	122880501	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren 50 mg υπόθετα	not available	122880501	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren 50 mg υπόθετα	not available	122880501	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren 50 mg υπόθετα	not available	122880501	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren 50 mg υπόθετα	not available	122880501	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren 50 mg υπόθετα	not available	122880501	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren 75 mg δισκία παρατεταμένης αποδέσμευσης	not available	122880801	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren 75 mg δισκία παρατεταμένης αποδέσμευσης	not available	122880802	NOVARTIS (HELLAS) S.A.C.I.	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
Voltaren 75 mg δισκία παρατεταμένης αποδέσμευσης	not available	122880801	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren 75 mg δισκία παρατεταμένης αποδέσμευσης	not available	122880802	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren 75 mg δισκία παρατεταμένης αποδέσμευσης	not available	122880801	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren 75 mg δισκία παρατεταμένης αποδέσμευσης	not available	122880802	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren 75 mg δισκία παρατεταμένης αποδέσμευσης	not available	122880801	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren 75 mg δισκία παρατεταμένης αποδέσμευσης	not available	122880802	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren 75 mg δισκία παρατεταμένης αποδέσμευσης	not available	122880801	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren 75 mg δισκία παρατεταμένης αποδέσμευσης	not available	122880802	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren 75 mg δισκία παρατεταμένης αποδέσμευσης	not available	122880801	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren 75 mg δισκία παρατεταμένης αποδέσμευσης	not available	122880802	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren 75 mg/3 ml ενέσιμο διάλυμα	not available	122880401	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren 75 mg/3 ml ενέσιμο διάλυμα	not available	122880401	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren 75 mg/3 ml ενέσιμο διάλυμα	not available	122880401	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren 75 mg/3 ml ενέσιμο διάλυμα	not available	122880401	NOVARTIS (HELLAS) S.A.C.I.	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
Voltaren 75 mg/3 ml ενέσιμο διάλυμα	not available	122880401	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren 75 mg/3 ml ενέσιμο διάλυμα	not available	122880401	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren fast 50 mg κόνις για πόσιμο διάλυμα	not available	251250401	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren fast 50 mg κόνις για πόσιμο διάλυμα	not available	251250402	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren fast 50 mg κόνις για πόσιμο διάλυμα	not available	251250403	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren fast 50 mg κόνις για πόσιμο διάλυμα	not available	251250401	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren fast 50 mg κόνις για πόσιμο διάλυμα	not available	251250402	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren fast 50 mg κόνις για πόσιμο διάλυμα	not available	251250403	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren fast 50 mg κόνις για πόσιμο διάλυμα	not available	251250401	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren fast 50 mg κόνις για πόσιμο διάλυμα	not available	251250402	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren fast 50 mg κόνις για πόσιμο διάλυμα	not available	251250403	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren fast 50 mg κόνις για πόσιμο διάλυμα	not available	251250401	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren fast 50 mg κόνις για πόσιμο διάλυμα	not available	251250402	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren fast 50 mg κόνις για πόσιμο διάλυμα	not available	251250403	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren fast 50 mg κόνις για πόσιμο διάλυμα	not available	251250401	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren fast 50 mg κόνις για πόσιμο διάλυμα	not available	251250402	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltaren fast 50 mg κόνις για πόσιμο διάλυμα	not available	251250403	NOVARTIS (HELLAS) S.A.C.I.	GR
Voltarol Acti-Go 12,5 mg επικαλυμμένα με λεπτό υμένιο δισκία	not available	321460301	HALEON HELLAS SINGLE MEMBER SOCIETE ANONYME	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
Voltarol Acti-Go 12,5 mg επικαλυμμένα με λεπτό υμένιο δισκία	not available	321460302	HALEON HELLAS SINGLE MEMBER SOCIETE ANONYME	GR
Voltarol Acti-Go 25 mg επικαλυμμένα δισκία	not available	3214601	HALEON HELLAS SINGLE MEMBER SOCIETE ANONYME	GR
Voltarol Acti-Go 50 mg επικαλυμμένα δισκία	not available	3214602	HALEON HELLAS SINGLE MEMBER SOCIETE ANONYME	GR
Cataflam 15 mg/ml belsőleges szuszpenziós cseppek	not available	OGYI-T-5573/03	NOVARTIS HUNGÁRIA KFT.	HU
Cataflam 15 mg/ml belsőleges szuszpenziós cseppek	not available	OGYI-T-5573/03	NOVARTIS HUNGÁRIA KFT.	HU
Cataflam 15 mg/ml belsőleges szuszpenziós cseppek	not available	OGYI-T-5573/03	NOVARTIS HUNGÁRIA KFT.	HU
Cataflam 15 mg/ml belsőleges szuszpenziós cseppek	not available	OGYI-T-5573/03	NOVARTIS HUNGÁRIA KFT.	HU
Cataflam 15 mg/ml belsőleges szuszpenziós cseppek	not available	OGYI-T-5573/03	NOVARTIS HUNGÁRIA KFT.	HU
Cataflam 50 mg bevont tabletta	not available	OGYI-T-5573/18	NOVARTIS HUNGÁRIA KFT.	HU
Cataflam 50 mg bevont tabletta	not available	OGYI-T-5573/02	NOVARTIS HUNGÁRIA KFT.	HU
Cataflam 50 mg bevont tabletta	not available	OGYI-T-5573/18	NOVARTIS HUNGÁRIA KFT.	HU
Cataflam 50 mg bevont tabletta	not available	OGYI-T-5573/02	NOVARTIS HUNGÁRIA KFT.	HU
Cataflam 50 mg bevont tabletta	not available	OGYI-T-5573/18	NOVARTIS HUNGÁRIA KFT.	HU
Cataflam 50 mg bevont tabletta	not available	OGYI-T-5573/02	NOVARTIS HUNGÁRIA KFT.	HU
Cataflam 50 mg bevont tabletta	not available	OGYI-T-5573/18	NOVARTIS HUNGÁRIA KFT.	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
Cataflam 50 mg bevont tabletta	not available	OGYI-T-5573/02	NOVARTIS HUNGÁRIA KFT.	HU
Cataflam 50 mg bevont tabletta	not available	OGYI-T-5573/18	NOVARTIS HUNGÁRIA KFT.	HU
Cataflam 50 mg bevont tabletta	not available	OGYI-T-5573/02	NOVARTIS HUNGÁRIA KFT.	HU
Cataflam Dolo 25 mg bevont tabletta	not available	OGYI-T-5573/06	HALEON HUNGARY KFT.	HU
Cataflam Dolo 25 mg bevont tabletta	not available	OGYI-T-5573/04	HALEON HUNGARY KFT.	HU
Cataflam Dolo 25 mg bevont tabletta	not available	OGYI-T-5573/05	HALEON HUNGARY KFT.	HU
Cataflam Dolo 25 mg bevont tabletta	not available	OGYI-T-5573/15	HALEON HUNGARY KFT.	HU
Cataflam Dolo Rapid 25 mg lágy kapszula	not available	OGYI-T-5573/09	HALEON HUNGARY KFT.	HU
Cataflam Dolo Rapid 25 mg lágy kapszula	not available	OGYI-T-5573/10	HALEON HUNGARY KFT.	HU
Cataflam Dolo Rapid 25 mg lágy kapszula	not available	OGYI-T-5573/11	HALEON HUNGARY KFT.	HU
Cataflam Dolo Rapid 25 mg lágy kapszula	not available	OGYI-T-5573/12	HALEON HUNGARY KFT.	HU
Cataflam Dolo Rapid 25 mg lágy kapszula	not available	OGYI-T-5573/13	HALEON HUNGARY KFT.	HU
Cataflam Dolo Rapid 25 mg lágy kapszula	not available	OGYI-T-5573/14	HALEON HUNGARY KFT.	HU
Cataflam Dolo Rapid 25 mg lágy kapszula	not available	OGYI-T-5573/16	HALEON HUNGARY KFT.	HU
Cataflam Dolo Rapid 25 mg lágy kapszula	not available	OGYI-T-5573/17	HALEON HUNGARY KFT.	HU
Cataflam-V 50 mg tabletta	not available	OGYI-T-5573/01	NOVARTIS HUNGÁRIA KFT.	HU
Cataflam-V 50 mg tabletta	not available	OGYI-T-5573/01	NOVARTIS HUNGÁRIA KFT.	HU
Cataflam-V 50 mg tabletta	not available	OGYI-T-5573/01	NOVARTIS HUNGÁRIA KFT.	HU
Cataflam-V 50 mg tabletta	not available	OGYI-T-5573/01	NOVARTIS HUNGÁRIA KFT.	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
Cataflam-V 50 mg tabletta	not available	OGYI-T-5573/01	NOVARTIS HUNGÁRIA KFT.	HU
Diclac 150 mg retard tabletta	not available	OGYI-T-4000/05	SANDOZ HUNGÁRIA KFT	HU
Diclac 150 mg retard tabletta	not available	OGYI-T-4000/06	SANDOZ HUNGÁRIA KFT	HU
Diclac 75 mg retard tabletta	not available	OGYI-T-4000/08	SANDOZ HUNGÁRIA KFT	HU
Diclac 75 mg retard tabletta	not available	OGYI-T-4000/07	SANDOZ HUNGÁRIA KFT	HU
Diclac 75 mg retard tabletta	not available	OGYI-T-4000/03	SANDOZ HUNGÁRIA KFT	HU
Diclac 75 mg retard tabletta	not available	OGYI-T-4000/04	SANDOZ HUNGÁRIA KFT	HU
Flector Dolo Rapid 12,5 mg lágy kapszula	CZ/H/0928/001	OGYI-T-23250/01	IBSA PHARMA KFT	HU
Flector Dolo Rapid 12,5 mg lágy kapszula	CZ/H/0928/001	OGYI-T-23250/02	IBSA PHARMA KFT	HU
Flector Dolo Rapid 12,5 mg lágy kapszula	CZ/H/0928/001	OGYI-T-23250/03	IBSA PHARMA KFT	HU
Flector Dolo Rapid 12,5 mg lágy kapszula	CZ/H/0928/001	OGYI-T-23250/04	IBSA PHARMA KFT	HU
Flector RAPID 50 mg granulátum	not available	OGYI-T-05033/14	IBSA PHARMA KFT	HU
Flector RAPID 50 mg granulátum	not available	OGYI-T-05033/03	IBSA PHARMA KFT	HU
Flector RAPID 50 mg granulátum	not available	OGYI-T-05033/04	IBSA PHARMA KFT	HU
Flector Rapiven 25 mg/ml oldatos injekció előretöltött fecskendőben	HU/H/0796/004	OGYI-T-23318/10	IBSA PHARMA KFT	HU
Flector Rapiven 50 mg/ml oldatos injekció előretöltött fecskendőben	HU/H/0796/005	OGYI-T-23318/11	IBSA PHARMA KFT	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
Flector Rapiven 75 mg/ml oldatos injekció eloretöltött fecskendoben	HU/H/0796/006	OGYI-T-23318/12	IBSA PHARMA KFT	HU
Voltaren 100 mg retard filmtabletta	not available	OGYI-T-5572/02	NOVARTIS HUNGÁRIA KFT.	HU
Voltaren 100 mg retard filmtabletta	not available	OGYI-T-5572/02	NOVARTIS HUNGÁRIA KFT.	HU
Voltaren 100 mg retard filmtabletta	not available	OGYI-T-5572/02	NOVARTIS HUNGÁRIA KFT.	HU
Voltaren 100 mg retard filmtabletta	not available	OGYI-T-5572/02	NOVARTIS HUNGÁRIA KFT.	HU
Voltaren 100 mg retard filmtabletta	not available	OGYI-T-5572/02	NOVARTIS HUNGÁRIA KFT.	HU
Voltaren 25 mg gyomornedv-ellenálló filmtabletta	not available	OGYI-T-5572/03	NOVARTIS HUNGÁRIA KFT.	HU
Voltaren 25 mg gyomornedv-ellenálló filmtabletta	not available	OGYI-T-5572/03	NOVARTIS HUNGÁRIA KFT.	HU
Voltaren 25 mg gyomornedv-ellenálló filmtabletta	not available	OGYI-T-5572/03	NOVARTIS HUNGÁRIA KFT.	HU
Voltaren 25 mg gyomornedv-ellenálló filmtabletta	not available	OGYI-T-5572/03	NOVARTIS HUNGÁRIA KFT.	HU
Voltaren 25 mg gyomornedv-ellenálló filmtabletta	not available	OGYI-T-5572/03	NOVARTIS HUNGÁRIA KFT.	HU
Voltaren 50 mg gyomornedv-ellenálló filmtabletta	not available	OGYI-T-5572/04	NOVARTIS HUNGÁRIA KFT.	HU
Voltaren 50 mg gyomornedv-ellenálló filmtabletta	not available	OGYI-T-5572/04	NOVARTIS HUNGÁRIA KFT.	HU
Voltaren 50 mg gyomornedv-ellenálló filmtabletta	not available	OGYI-T-5572/04	NOVARTIS HUNGÁRIA KFT.	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
Voltaren 50 mg gyomornedv-ellenálló filmtabletta	not available	OGYI-T-5572/04	NOVARTIS HUNGÁRIA KFT.	HU
Voltaren 50 mg gyomornedv-ellenálló filmtabletta	not available	OGYI-T-5572/04	NOVARTIS HUNGÁRIA KFT.	HU
Voltaren 50 mg végbélkúp	not available	OGYI-T-5572/05	NOVARTIS HUNGÁRIA KFT.	HU
Voltaren 50 mg végbélkúp	not available	OGYI-T-5572/05	NOVARTIS HUNGÁRIA KFT.	HU
Voltaren 50 mg végbélkúp	not available	OGYI-T-5572/05	NOVARTIS HUNGÁRIA KFT.	HU
Voltaren 50 mg végbélkúp	not available	OGYI-T-5572/05	NOVARTIS HUNGÁRIA KFT.	HU
Voltaren 50 mg végbélkúp	not available	OGYI-T-5572/05	NOVARTIS HUNGÁRIA KFT.	HU
Voltaren 75 mg retard filmtabletta	not available	OGYI-T-5572/01	NOVARTIS HUNGÁRIA KFT.	HU
Voltaren 75 mg retard filmtabletta	not available	OGYI-T-5572/01	NOVARTIS HUNGÁRIA KFT.	HU
Voltaren 75 mg retard filmtabletta	not available	OGYI-T-5572/01	NOVARTIS HUNGÁRIA KFT.	HU
Voltaren 75 mg retard filmtabletta	not available	OGYI-T-5572/01	NOVARTIS HUNGÁRIA KFT.	HU
Voltaren 75 mg retard filmtabletta	not available	OGYI-T-5572/01	NOVARTIS HUNGÁRIA KFT.	HU
Voltaren 75 mg/3 ml oldatos injekció	not available	OGYI-T-5572/06	NOVARTIS HUNGÁRIA KFT.	HU
Voltaren 75 mg/3 ml oldatos injekció	not available	OGYI-T-5572/06	NOVARTIS HUNGÁRIA KFT.	HU
Voltaren 75 mg/3 ml oldatos injekció	not available	OGYI-T-5572/06	NOVARTIS HUNGÁRIA KFT.	HU
Voltaren 75 mg/3 ml oldatos injekció	not available	OGYI-T-5572/06	NOVARTIS HUNGÁRIA KFT.	HU
Voltaren 75 mg/3 ml oldatos injekció	not available	OGYI-T-5572/06	NOVARTIS HUNGÁRIA KFT.	HU
Voltaren Dolo 12,5 mg lágy kapszula	not available	OGYI-T-5572/22	HALEON HUNGARY KFT.	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
Voltaren Dolo 12,5 mg lágy kapszula	not available	OGYI-T-5572/21	HALEON HUNGARY KFT.	HU
Voltaren Dolo 25 mg bevont tabletta	not available	OGYI-T-5572/23	HALEON HUNGARY KFT.	HU
Voltaren Dolo 25 mg bevont tabletta	not available	OGYI-T-5572/25	HALEON HUNGARY KFT.	HU
Voltaren Dolo 25 mg bevont tabletta	not available	OGYI-T-5572/24	HALEON HUNGARY KFT.	HU
Voltaren Dolo Rapid 25 mg lágy kapszula	not available	OGYI-T-5572/76	HALEON HUNGARY KFT.	HU
Voltaren Dolo Rapid 25 mg lágy kapszula	not available	OGYI-T-5572/77	HALEON HUNGARY KFT.	HU
Voltaren Dolo Rapid 25 mg lágy kapszula	not available	OGYI-T-5572/30	HALEON HUNGARY KFT.	HU
Voltaren Dolo Rapid 25 mg lágy kapszula	not available	OGYI-T-5572/29	HALEON HUNGARY KFT.	HU
Voltaren Dolo Rapid 25 mg lágy kapszula	not available	OGYI-T-5572/26	HALEON HUNGARY KFT.	HU
Voltaren Dolo Rapid 25 mg lágy kapszula	not available	OGYI-T-5572/31	HALEON HUNGARY KFT.	HU
Voltaren Dolo Rapid 25 mg lágy kapszula	not available	OGYI-T-5572/28	HALEON HUNGARY KFT.	HU
Voltaren Dolo Rapid 25 mg lágy kapszula	not available	OGYI-T-5572/27	HALEON HUNGARY KFT.	HU
Cataflam 50mg Coated Tablets	not available	PA0896/004/001	NOVARTIS IRELAND LIMITED	IE
Cataflam 50mg Coated Tablets	not available	PA0896/004/001	NOVARTIS IRELAND LIMITED	IE
Cataflam 50mg Coated Tablets	not available	PA0896/004/001	NOVARTIS IRELAND LIMITED	IE
Cataflam 50mg Coated Tablets	not available	PA0896/004/001	NOVARTIS IRELAND LIMITED	IE
Cataflam 50mg Coated Tablets	not available	PA0896/004/001	NOVARTIS IRELAND LIMITED	IE
Diclac 75 mg Prolonged-release Tablet	not available	PA0711/009/003	ROWEX LTD	IE
Voltarol 50mg Gastro-Resistant Tablets	not available	PA0896/034/001	NOVARTIS IRELAND LIMITED	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
Voltarol 50mg Gastro-Resistant Tablets	not available	PA0896/034/001	NOVARTIS IRELAND LIMITED	IE
Voltarol 50mg Gastro-Resistant Tablets	not available	PA0896/034/001	NOVARTIS IRELAND LIMITED	IE
Voltarol 50mg Gastro-Resistant Tablets	not available	PA0896/034/001	NOVARTIS IRELAND LIMITED	IE
Voltarol 50mg Gastro-Resistant Tablets	not available	PA0896/034/001	NOVARTIS IRELAND LIMITED	IE
Voltarol Retard 100 mg Film-coated Prolonged Release Tablets	not available	PA0896/034/002	NOVARTIS IRELAND LIMITED	IE
Voltarol Retard 100 mg Film-coated Prolonged Release Tablets	not available	PA0896/034/002	NOVARTIS IRELAND LIMITED	IE
Voltarol Retard 100 mg Film-coated Prolonged Release Tablets	not available	PA0896/034/002	NOVARTIS IRELAND LIMITED	IE
Voltarol Retard 100 mg Film-coated Prolonged Release Tablets	not available	PA0896/034/002	NOVARTIS IRELAND LIMITED	IE
Voltarol Retard 100 mg Film-coated Prolonged Release Tablets	not available	PA0896/034/002	NOVARTIS IRELAND LIMITED	IE
Voltarol Retard 75mg film-coated prolonged-release tablets.	not available	PA 0896/034/003	NOVARTIS IRELAND LIMITED	IE
Voltarol Retard 75mg film-coated prolonged-release tablets.	not available	PA 0896/034/003	NOVARTIS IRELAND LIMITED	IE
Voltarol Retard 75mg film-coated prolonged-release tablets.	not available	PA 0896/034/003	NOVARTIS IRELAND LIMITED	IE
Voltarol Retard 75mg film-coated prolonged-release tablets.	not available	PA 0896/034/003	NOVARTIS IRELAND LIMITED	IE
Voltarol Retard 75mg film-coated prolonged-release tablets.	not available	PA 0896/034/003	NOVARTIS IRELAND LIMITED	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
Modifenac 75 mg hart hylki með breyttan losunarhraða	not available	980422	TEVA B.V	IS
DEALGIC 100 mg capsule rigide a rilascio prolungato	not available	028943025	FARMA GROUP S.R.L.	IT
DEALGIC 75 mg capsule rigide a rilascio prolungato	not available	028943013	FARMA GROUP S.R.L.	IT
DICLOFENAC Alfasigma 150 mg capsule a rilascio prolungato	not available	033612058	ALFASIGMA S.P.A.	IT
DICLOFENAC Alfasigma 50 mg compresse	not available	033612033	ALFASIGMA S.P.A.	IT
DICLOFENAC Alfasigma 75 mg/3 ml soluzione iniettabile per uso intramuscolare	not available	033612096	ALFASIGMA S.P.A.	IT
DICLOREUM 100 MG COMPRESSE A RILASCIO PROLUNGATO	not available	024515088	ALFASIGMA S.P.A.	IT
DICLOREUM 150 mg capsule rigide a rilascio prolungato	not available	024515114	ALFASIGMA S.P.A.	IT
DICLOREUM 50 MG COMPRESSE GASTRORESISTENTI	not available	024515049	ALFASIGMA S.P.A.	IT
DICLOREUM 50 MG GRANULATO PER SOLUZIONE ORALE	not available	024515138	ALFASIGMA S.P.A.	IT
DICLOREUM 75 MG/3 ML SOLUZIONE INIETTABILE PER USO INTRAMUSCOLARE	not available	024515076	ALFASIGMA S.P.A.	IT
DICLOREUM DOLORE 25 mg granulato per soluzione orale	not available	028618015	ALFASIGMA 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
DICLOREUM DOLORE 25 mg granulato per soluzione orale	not available	028618041	ALFASIGMA S.P.A.	IT
FLECTOR DOLORE 25 mg granulato per soluzione orale	not available	028617037	IBSA FARMACEUTICI ITALIA	IT
FLECTOR DOLORE 25 mg granulato per soluzione orale	not available	028617052	IBSA FARMACEUTICI ITALIA	IT
FLECTORFLAM 50 mg granulato per soluzione orale	not available	036058030	IBSA FARMACEUTICI ITALIA	IT
FLECTORGO 12,5 mg capsule molli	CZ/H/0928/001	044608014	IBSA FARMACEUTICI ITALIA	IT
FLECTORGO 12,5 mg capsule molli	CZ/H/0928/001	044608026	IBSA FARMACEUTICI ITALIA	IT
FLECTORGO 12,5 mg capsule molli	CZ/H/0928/001	044608038	IBSA FARMACEUTICI ITALIA	IT
FLECTORGO 12,5 mg capsule molli	CZ/H/0928/001	044608040	IBSA FARMACEUTICI ITALIA	IT
FlectorIn 25 mg/ml soluzione iniettabile	HU/H/0796/001	040528198	IBSA FARMACEUTICI ITALIA	IT
FlectorIn 25 mg/ml soluzione iniettabile	HU/H/0796/001	040528200	IBSA FARMACEUTICI ITALIA	IT
FlectorIn 25 mg/ml soluzione iniettabile	HU/H/0796/001	040528212	IBSA FARMACEUTICI ITALIA	IT
FlectorIn 25 mg/ml soluzione iniettabile in siringa preriempita	HU/H/0796/004	040528287	IBSA FARMACEUTICI ITALIA	IT
FlectorIn 25 mg/ml soluzione iniettabile in siringa preriempita	HU/H/0796/004	040528299	IBSA FARMACEUTICI ITALIA	IT
FlectorIn 25 mg/ml soluzione iniettabile in siringa preriempita	HU/H/0796/004	040528301	IBSA FARMACEUTICI ITALIA	IT
FlectorIn 50 mg/ml soluzione iniettabile	HU/H/0796/002	040528224	IBSA FARMACEUTICI ITALIA	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
FlectorIn 50 mg/ml soluzione iniettabile	HU/H/0796/002	040528236	IBSA FARMACEUTICI ITALIA	IT
FlectorIn 50 mg/ml soluzione iniettabile	HU/H/0796/002	040528248	IBSA FARMACEUTICI ITALIA	IT
FlectorIn 50 mg/ml soluzione iniettabile in siringa preriempita	HU/H/0796/005	040528313	IBSA FARMACEUTICI ITALIA	IT
FlectorIn 50 mg/ml soluzione iniettabile in siringa preriempita	HU/H/0796/005	040528325	IBSA FARMACEUTICI ITALIA	IT
FlectorIn 50 mg/ml soluzione iniettabile in siringa preriempita	HU/H/0796/005	040528337	IBSA FARMACEUTICI ITALIA	IT
FlectorIn 75 mg/ml soluzione iniettabile	HU/H/0796/003	040528251	IBSA FARMACEUTICI ITALIA	IT
FlectorIn 75 mg/ml soluzione iniettabile	HU/H/0796/003	040528263	IBSA FARMACEUTICI ITALIA	IT
FlectorIn 75 mg/ml soluzione iniettabile	HU/H/0796/003	040528275	IBSA FARMACEUTICI ITALIA	IT
FlectorIn 75 mg/ml soluzione iniettabile in siringa preriempita	HU/H/0796/006	040528349	IBSA FARMACEUTICI ITALIA	IT
FlectorIn 75 mg/ml soluzione iniettabile in siringa preriempita	HU/H/0796/006	040528352	IBSA FARMACEUTICI ITALIA	IT
FlectorIn 75 mg/ml soluzione iniettabile in siringa preriempita	HU/H/0796/006	040528364	IBSA FARMACEUTICI ITALIA	IT
FLOGOFENAC 100 mg capsule rigide a rilascio prolungato	not available	025536020	A. MENARINI - INDUSTRIE FARMACEUTICHE RIUNITE - S.R.L.	IT
Itamifast 25 mg compresse rivestite con film	IT/H/0352/001	041736012	FIDIA FARMACEUTICI S.P.A	IT
Itamifast 25 mg compresse rivestite con film	IT/H/0352/001	041736024	FIDIA FARMACEUTICI 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
Itamifast 50 mg compresse rivestite con film	IT/H/0352/002	041736036	FIDIA FARMACEUTICI S.P.A	IT
Itamifast 50 mg compresse rivestite con film	IT/H/0352/002	041736048	FIDIA FARMACEUTICI S.P.A	IT
SAFIDOL 25 mg compresse rivestite con film	IT/H/0483/001	041735010	SAFI MEDICAL CARE S.R.L	IT
SAFIDOL 25 mg compresse rivestite con film	IT/H/0483/001	041735022	SAFI MEDICAL CARE S.R.L	IT
TRAULEN 100 mg compresse a rilascio prolungato	not available	033420.023	OP PHARMA S.R.L.	IT
TRAULEN 75 mg/3 ml soluzione iniettabile per uso intramuscolare	not available	033420.047	OP PHARMA S.R.L.	IT
VOLTAREN 100 mg compresse a rilascio prolungato	not available	023181035	NOVARTIS FARMA S.P.A.	IT
VOLTAREN 100 mg compresse a rilascio prolungato	not available	023181035	NOVARTIS FARMA S.P.A.	IT
VOLTAREN 100 mg compresse a rilascio prolungato	not available	023181035	NOVARTIS FARMA S.P.A.	IT
VOLTAREN 100 mg compresse a rilascio prolungato	not available	023181035	NOVARTIS FARMA S.P.A.	IT
VOLTAREN 100 mg compresse a rilascio prolungato	not available	023181035	NOVARTIS FARMA S.P.A.	IT
VOLTAREN 100 mg supposte	not available	023181023	NOVARTIS FARMA S.P.A.	IT
VOLTAREN 100 mg supposte	not available	023181023	NOVARTIS FARMA 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
VOLTAREN 100 mg supposte	not available	023181023	NOVARTIS FARMA S.P.A.	IT
VOLTAREN 100 mg supposte	not available	023181023	NOVARTIS FARMA S.P.A.	IT
VOLTAREN 100 mg supposte	not available	023181023	NOVARTIS FARMA S.P.A.	IT
VOLTAREN 50 mg compresse gastroresistenti	not available	023181011	NOVARTIS FARMA S.P.A.	IT
VOLTAREN 50 mg compresse gastroresistenti	not available	023181011	NOVARTIS FARMA S.P.A.	IT
VOLTAREN 50 mg compresse gastroresistenti	not available	023181011	NOVARTIS FARMA S.P.A.	IT
VOLTAREN 50 mg compresse gastroresistenti	not available	023181011	NOVARTIS FARMA S.P.A.	IT
VOLTAREN 50 mg compresse gastroresistenti	not available	023181011	NOVARTIS FARMA S.P.A.	IT
VOLTAREN 50 mg compresse solubili	not available	023181086	NOVARTIS FARMA S.P.A.	IT
VOLTAREN 50 mg compresse solubili	not available	023181086	NOVARTIS FARMA S.P.A.	IT
VOLTAREN 50 mg compresse solubili	not available	023181086	NOVARTIS FARMA S.P.A.	IT
VOLTAREN 50 mg compresse solubili	not available	023181086	NOVARTIS FARMA S.P.A.	IT
VOLTAREN 50 mg compresse solubili	not available	023181086	NOVARTIS FARMA S.P.A.	IT
VOLTAREN 75 mg compresse a rilascio prolungato	not available	023181074	NOVARTIS FARMA S.P.A.	IT
VOLTAREN 75 mg compresse a rilascio prolungato	not available	023181074	NOVARTIS FARMA 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
VOLTAREN 75 mg compresse a rilascio prolungato	not available	023181074	NOVARTIS FARMA S.P.A.	IT
VOLTAREN 75 mg compresse a rilascio prolungato	not available	023181074	NOVARTIS FARMA S.P.A.	IT
VOLTAREN 75 mg compresse a rilascio prolungato	not available	023181074	NOVARTIS FARMA S.P.A.	IT
VOLTAREN 75 mg/3 ml soluzione iniettabile per uso intramuscolare	not available	023181047	NOVARTIS FARMA S.P.A.	IT
VOLTAREN 75 mg/3 ml soluzione iniettabile per uso intramuscolare	not available	023181047	NOVARTIS FARMA S.P.A.	IT
VOLTAREN 75 mg/3 ml soluzione iniettabile per uso intramuscolare	not available	023181047	NOVARTIS FARMA S.P.A.	IT
VOLTAREN 75 mg/3 ml soluzione iniettabile per uso intramuscolare	not available	023181047	NOVARTIS FARMA S.P.A.	IT
VOLTAREN 75 mg/3 ml soluzione iniettabile per uso intramuscolare	not available	023181047	NOVARTIS FARMA S.P.A.	IT
VOLTFAST 50 mg compresse rivestite.	not available	028945020	NOVARTIS FARMA S.P.A.	IT
VOLTFAST 50 mg compresse rivestite.	not available	028945020	NOVARTIS FARMA S.P.A.	IT
VOLTFAST 50 mg compresse rivestite.	not available	028945020	NOVARTIS FARMA S.P.A.	IT
VOLTFAST 50 mg compresse rivestite.	not available	028945020	NOVARTIS FARMA S.P.A.	IT
VOLTFAST 50 mg compresse rivestite.	not available	028945020	NOVARTIS FARMA S.P.A.	IT
VOLTFAST 50 mg granulato per soluzione orale	not available	028945032	NOVARTIS FARMA 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
VOLTFAST 50 mg granulato per soluzione orale	not available	028945032	NOVARTIS FARMA S.P.A.	IT
VOLTFAST 50 mg granulato per soluzione orale	not available	028945032	NOVARTIS FARMA S.P.A.	IT
VOLTFAST 50 mg granulato per soluzione orale	not available	028945032	NOVARTIS FARMA S.P.A.	IT
VOLTFAST 50 mg granulato per soluzione orale	not available	028945032	NOVARTIS FARMA S.P.A.	IT
Voltaren Akti 12,5 mg plėvele dengtos tabletės	not available	LT/1/94/0948/003	HALEON HUNGARY KFT.	LT
Voltaren Akti 25 mg minkštosios kapsulės	not available	LT/1/94/0948/005	HALEON HUNGARY KFT.	LT
Voltaren Akti 25 mg minkštosios kapsulės	not available	LT/1/94/0948/004	HALEON HUNGARY KFT.	LT
Cataflam 50 mg, comprimés enrobés	not available	2001107175	NOVARTIS PHARMA N.V.	LU
Cataflam 50 mg, comprimés enrobés	not available	2001107175	NOVARTIS PHARMA N.V.	LU
Cataflam 50 mg, comprimés enrobés	not available	2001107175	NOVARTIS PHARMA N.V.	LU
Cataflam 50 mg, comprimés enrobés	not available	2001107175	NOVARTIS PHARMA N.V.	LU
Cataflam 50 mg, überzogene Tabletten	not available	2001107175	NOVARTIS PHARMA N.V.	LU
Cataflam 50 mg, überzogene Tabletten	not available	2001107175	NOVARTIS PHARMA N.V.	LU
Cataflam 50 mg, überzogene Tabletten	not available	2001107175	NOVARTIS PHARMA N.V.	LU
Cataflam 50 mg, überzogene Tabletten	not available	2001107175	NOVARTIS PHARMA N.V.	LU
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	2017040080	MICRO LABS GMBH	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
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	2017040080	MICRO LABS GMBH	LU
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	2017040080	MICRO LABS GMBH	LU
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	2017040080	MICRO LABS GMBH	LU
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	2017040080	MICRO LABS GMBH	LU
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	2017040080	MICRO LABS GMBH	LU
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	2017040080	MICRO LABS GMBH	LU
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	2017040080	MICRO LABS GMBH	LU
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	2017040080	MICRO LABS GMBH	LU
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	2017040080	MICRO LABS GMBH	LU
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	2017040080	MICRO LABS GMBH	LU
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	2017040080	MICRO LABS GMBH	LU
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	2017040080	MICRO LABS GMBH	LU
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	2017040080	MICRO LABS GMBH	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
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	2017040080	MICRO LABS GMBH	LU
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	2017040080	MICRO LABS GMBH	LU
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	2017040080	MICRO LABS GMBH	LU
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	2017040080	MICRO LABS GMBH	LU
Diclofenac-Natrium Micro Labs 75 mg Retardtabletten	DE/H/4552/001	2017040080	MICRO LABS GMBH	LU
Motifene 75 mg gélules à libération modifiée	not available	BE 176671	GLENWOOD GMBH	LU
Voltaren 100 mg suppositoires.	not available	2008049767	NOVARTIS PHARMA N.V.	LU
Voltaren 100 mg suppositoires.	not available	2008049767	NOVARTIS PHARMA N.V.	LU
Voltaren 100 mg suppositoires.	not available	2008049767	NOVARTIS PHARMA N.V.	LU
Voltaren 100 mg suppositoires.	not available	2008049767	NOVARTIS PHARMA N.V.	LU
Voltaren 100 mg Zäpfchen	not available	2008049767	NOVARTIS PHARMA N.V.	LU
Voltaren 100 mg Zäpfchen	not available	2008049767	NOVARTIS PHARMA N.V.	LU
Voltaren 100 mg Zäpfchen	not available	2008049767	NOVARTIS PHARMA N.V.	LU
Voltaren 100 mg Zäpfchen	not available	2008049767	NOVARTIS PHARMA N.V.	LU
Voltaren 50 mg comprimés gastro-résistants.	not available	2002100205	NOVARTIS PHARMA N.V.	LU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Voltaren 50 mg comprimés gastro-résistants.	not available	2002100205	NOVARTIS PHARMA N.V.	LU
Voltaren 50 mg comprimés gastro-résistants.	not available	2002100205	NOVARTIS PHARMA N.V.	LU
Voltaren 50 mg comprimés gastro-résistants.	not available	2002100205	NOVARTIS PHARMA N.V.	LU
Voltaren 50 mg magensaftresistente Tabletten	not available	2002100205	NOVARTIS PHARMA N.V.	LU
Voltaren 50 mg magensaftresistente Tabletten	not available	2002100205	NOVARTIS PHARMA N.V.	LU
Voltaren 50 mg magensaftresistente Tabletten	not available	2002100205	NOVARTIS PHARMA N.V.	LU
Voltaren 50 mg magensaftresistente Tabletten	not available	2002100205	NOVARTIS PHARMA N.V.	LU
Voltaren Retard 100 mg comprimés à libération prolongée.	not available	2008049766	NOVARTIS PHARMA N.V.	LU
Voltaren Retard 100 mg comprimés à libération prolongée.	not available	2008049766	NOVARTIS PHARMA N.V.	LU
Voltaren Retard 100 mg comprimés à libération prolongée.	not available	2008049766	NOVARTIS PHARMA N.V.	LU
Voltaren Retard 100 mg comprimés à libération prolongée.	not available	2008049766	NOVARTIS PHARMA N.V.	LU
Voltaren Retard 100 mg Retardtabletten	not available	2008049766	NOVARTIS PHARMA N.V.	LU
Voltaren Retard 100 mg Retardtabletten	not available	2008049766	NOVARTIS PHARMA N.V.	LU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Voltaren Retard 100 mg Retardtabletten	not available	2008049766	NOVARTIS PHARMA N.V.	LU
Voltaren Retard 100 mg Retardtabletten	not available	2008049766	NOVARTIS PHARMA N.V.	LU
Voltaren Akti 12,5 mg apvalkotās tabletes	not available	04 – 0291	HALEON HUNGARY KFT.	LV
Catafast 50 mg powder for oral solution.	not available	MA1249/00203	NOVARTIS IRELAND LIMITED	MT
Catafast 50 mg powder for oral solution.	not available	MA1249/00203	NOVARTIS IRELAND LIMITED	MT
Catafast 50 mg powder for oral solution.	not available	MA1249/00203	NOVARTIS IRELAND LIMITED	MT
Catafast 50 mg powder for oral solution.	not available	MA1249/00203	NOVARTIS IRELAND LIMITED	MT
Catafast 50 mg powder for oral solution.	not available	MA1249/00203	NOVARTIS IRELAND LIMITED	MT
CATAFLAM 25 mg sugar-coated tablets.	not available	MA1249/00201	NOVARTIS IRELAND LIMITED	MT
CATAFLAM 25 mg sugar-coated tablets.	not available	MA1249/00201	NOVARTIS IRELAND LIMITED	MT
CATAFLAM 25 mg sugar-coated tablets.	not available	MA1249/00201	NOVARTIS IRELAND LIMITED	MT
CATAFLAM 25 mg sugar-coated tablets.	not available	MA1249/00201	NOVARTIS IRELAND LIMITED	MT
CATAFLAM 25 mg sugar-coated tablets.	not available	MA1249/00201	NOVARTIS IRELAND LIMITED	MT
CATAFLAM 50 mg sugar-coated tablets.	not available	MA1249/00202	NOVARTIS IRELAND LIMITED	MT
CATAFLAM 50 mg sugar-coated tablets.	not available	MA1249/00202	NOVARTIS IRELAND LIMITED	MT
CATAFLAM 50 mg sugar-coated tablets.	not available	MA1249/00202	NOVARTIS IRELAND LIMITED	MT
CATAFLAM 50 mg sugar-coated tablets.	not available	MA1249/00202	NOVARTIS IRELAND LIMITED	MT
CATAFLAM 50 mg sugar-coated tablets.	not available	MA1249/00202	NOVARTIS IRELAND LIMITED	MT
VOLTAREN 100 mg suppositories.	not available	MA1249/00703	NOVARTIS IRELAND LIMITED	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
VOLTAREN 100 mg suppositories.	not available	MA1249/00703	NOVARTIS IRELAND LIMITED	MT
VOLTAREN 100 mg suppositories.	not available	MA1249/00703	NOVARTIS IRELAND LIMITED	MT
VOLTAREN 100 mg suppositories.	not available	MA1249/00703	NOVARTIS IRELAND LIMITED	MT
VOLTAREN 100 mg suppositories.	not available	MA1249/00703	NOVARTIS IRELAND LIMITED	MT
VOLTAREN 25 mg gastro-resistant tablets.	not available	MA1249/00704	NOVARTIS IRELAND LIMITED	MT
VOLTAREN 25 mg gastro-resistant tablets.	not available	MA1249/00704	NOVARTIS IRELAND LIMITED	MT
VOLTAREN 25 mg gastro-resistant tablets.	not available	MA1249/00704	NOVARTIS IRELAND LIMITED	MT
VOLTAREN 25 mg gastro-resistant tablets.	not available	MA1249/00704	NOVARTIS IRELAND LIMITED	MT
VOLTAREN 25 mg gastro-resistant tablets.	not available	MA1249/00704	NOVARTIS IRELAND LIMITED	MT
VOLTAREN 50 mg gastro-resistant tablets.	not available	MA1249/00705	NOVARTIS IRELAND LIMITED	MT
VOLTAREN 50 mg gastro-resistant tablets.	not available	MA1249/00705	NOVARTIS IRELAND LIMITED	MT
VOLTAREN 50 mg gastro-resistant tablets.	not available	MA1249/00705	NOVARTIS IRELAND LIMITED	MT
VOLTAREN 50 mg gastro-resistant tablets.	not available	MA1249/00705	NOVARTIS IRELAND LIMITED	MT
VOLTAREN 50 mg gastro-resistant tablets.	not available	MA1249/00705	NOVARTIS IRELAND LIMITED	MT
VOLTAREN 50 mg suppositories.	not available	MA1249/00702	NOVARTIS IRELAND LIMITED	MT
VOLTAREN 50 mg suppositories.	not available	MA1249/00702	NOVARTIS IRELAND LIMITED	MT
VOLTAREN 50 mg suppositories.	not available	MA1249/00702	NOVARTIS IRELAND LIMITED	MT
VOLTAREN 50 mg suppositories.	not available	MA1249/00702	NOVARTIS IRELAND LIMITED	MT
VOLTAREN 50 mg suppositories.	not available	MA1249/00702	NOVARTIS IRELAND LIMITED	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
VOLTAREN 75mg/3mL solution for injection.	not available	MA1249/00706	NOVARTIS IRELAND LIMITED	MT
VOLTAREN 75mg/3mL solution for injection.	not available	MA1249/00706	NOVARTIS IRELAND LIMITED	MT
VOLTAREN 75mg/3mL solution for injection.	not available	MA1249/00706	NOVARTIS IRELAND LIMITED	MT
VOLTAREN 75mg/3mL solution for injection.	not available	MA1249/00706	NOVARTIS IRELAND LIMITED	MT
VOLTAREN 75mg/3mL solution for injection.	not available	MA1249/00706	NOVARTIS IRELAND LIMITED	MT
VOLTAREN D 50mg dispersible tablets.	not available	MA1249/00707	NOVARTIS IRELAND LIMITED	MT
VOLTAREN D 50mg dispersible tablets.	not available	MA1249/00707	NOVARTIS IRELAND LIMITED	MT
VOLTAREN D 50mg dispersible tablets.	not available	MA1249/00707	NOVARTIS IRELAND LIMITED	MT
VOLTAREN D 50mg dispersible tablets.	not available	MA1249/00707	NOVARTIS IRELAND LIMITED	MT
VOLTAREN D 50mg dispersible tablets.	not available	MA1249/00707	NOVARTIS IRELAND LIMITED	MT
VOLTAREN Retard 100	not available	MA1249/00709	NOVARTIS IRELAND LIMITED	MT
VOLTAREN Retard 100	not available	MA1249/00709	NOVARTIS IRELAND LIMITED	MT
VOLTAREN Retard 100	not available	MA1249/00709	NOVARTIS IRELAND LIMITED	MT
VOLTAREN Retard 100	not available	MA1249/00709	NOVARTIS IRELAND LIMITED	MT
VOLTAREN Retard 100	not available	MA1249/00709	NOVARTIS IRELAND LIMITED	MT
VOLTAREN SR 75	not available	MA1249/00708	NOVARTIS IRELAND LIMITED	MT
VOLTAREN SR 75	not available	MA1249/00708	NOVARTIS IRELAND LIMITED	MT
VOLTAREN SR 75	not available	MA1249/00708	NOVARTIS IRELAND LIMITED	MT
VOLTAREN SR 75	not available	MA1249/00708	NOVARTIS IRELAND LIMITED	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
VOLTAREN SR 75	not available	MA1249/00708	NOVARTIS IRELAND LIMITED	MT
VOLTAREN12.5 mg suppositories.	not available	MA1249/00701	NOVARTIS IRELAND LIMITED	MT
VOLTAREN12.5 mg suppositories.	not available	MA1249/00701	NOVARTIS IRELAND LIMITED	MT
VOLTAREN12.5 mg suppositories.	not available	MA1249/00701	NOVARTIS IRELAND LIMITED	MT
VOLTAREN12.5 mg suppositories.	not available	MA1249/00701	NOVARTIS IRELAND LIMITED	MT
VOLTAREN12.5 mg suppositories.	not available	MA1249/00701	NOVARTIS IRELAND LIMITED	MT
Cataflam 50, omhulde tabletten 50 mg	not available	RVG 13245	NOVARTIS PHARMA B.V.	NL
Cataflam 50, omhulde tabletten 50 mg	not available	RVG 13245	NOVARTIS PHARMA B.V.	NL
Cataflam 50, omhulde tabletten 50 mg	not available	RVG 13245	NOVARTIS PHARMA B.V.	NL
Cataflam 50, omhulde tabletten 50 mg	not available	RVG 13245	NOVARTIS PHARMA B.V.	NL
Cataflam 50, omhulde tabletten 50 mg	not available	RVG 13245	NOVARTIS PHARMA B.V.	NL
Cataflam 50, omhulde tabletten 50 mg	not available	RVG 13245	NOVARTIS PHARMA B.V.	NL
Voltaren 75 mg/3 ml, oplossing voor injectie 25 mg/ml	not available	RVG 07460	NOVARTIS PHARMA B.V.	NL
Voltaren 75 mg/3 ml, oplossing voor injectie 25 mg/ml	not available	RVG 07460	NOVARTIS PHARMA B.V.	NL
Voltaren 75 mg/3 ml, oplossing voor injectie 25 mg/ml	not available	RVG 07460	NOVARTIS PHARMA B.V.	NL
Voltaren 75 mg/3 ml, oplossing voor injectie 25 mg/ml	not available	RVG 07460	NOVARTIS PHARMA 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
Voltaren 75 mg/3 ml, oplossing voor injectie 25 mg/ml	not available	RVG 07460	NOVARTIS PHARMA B.V.	NL
Voltaren 75 mg/3 ml, oplossing voor injectie 25 mg/ml	not available	RVG 07460	NOVARTIS PHARMA B.V.	NL
Voltaren K 12,5 mg, filmomhulde tabletten	not available	RVG 20982	HALEON NETHERLANDS B.V.	NL
Voltaren K 25 mg, omhulde tabletten.	not available	RVG 13244	HALEON NETHERLANDS B.V.	NL
CATAFLAM 50 mg drasjerte tabletter	not available	8287	NOVARTIS NORGE AS	NO
CATAFLAM 50 mg drasjerte tabletter	not available	8287	NOVARTIS NORGE AS	NO
CATAFLAM 50 mg drasjerte tabletter	not available	8287	NOVARTIS NORGE AS	NO
CATAFLAM 50 mg drasjerte tabletter	not available	8287	NOVARTIS NORGE AS	NO
CATAFLAM 50 mg drasjerte tabletter	not available	8287	NOVARTIS NORGE AS	NO
CATAFLAM 50 mg drasjerte tabletter	not available	8287	NOVARTIS NORGE AS	NO
Modifenac 75 mg kapsler med modifisert frisetting, harde	not available	96-3618	TEVA B.V	NO
Voltaren 100 mg stikkpiller	not available	7784	NOVARTIS NORGE AS	NO
Voltaren 100 mg stikkpiller	not available	7784	NOVARTIS NORGE AS	NO
Voltaren 100 mg stikkpiller	not available	7784	NOVARTIS NORGE AS	NO
Voltaren 100 mg stikkpiller	not available	7784	NOVARTIS NORGE AS	NO
Voltaren 100 mg stikkpiller	not available	7784	NOVARTIS NORGE AS	NO
Voltaren 100 mg stikkpiller	not available	7784	NOVARTIS 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
Voltaren 25 mg enterotabletter	not available	7780	NOVARTIS NORGE AS	NO
Voltaren 25 mg enterotabletter	not available	7780	NOVARTIS NORGE AS	NO
Voltaren 25 mg enterotabletter	not available	7780	NOVARTIS NORGE AS	NO
Voltaren 25 mg enterotabletter	not available	7780	NOVARTIS NORGE AS	NO
Voltaren 25 mg enterotabletter	not available	7780	NOVARTIS NORGE AS	NO
Voltaren 25 mg enterotabletter	not available	7780	NOVARTIS NORGE AS	NO
Voltaren 25 mg stikkpiller	not available	7782	NOVARTIS NORGE AS	NO
Voltaren 25 mg stikkpiller	not available	7782	NOVARTIS NORGE AS	NO
Voltaren 25 mg stikkpiller	not available	7782	NOVARTIS NORGE AS	NO
Voltaren 25 mg stikkpiller	not available	7782	NOVARTIS NORGE AS	NO
Voltaren 25 mg stikkpiller	not available	7782	NOVARTIS NORGE AS	NO
Voltaren 25 mg stikkpiller	not available	7782	NOVARTIS NORGE AS	NO
Voltaren 25 mg/ml injeksjonsvæske / konsentrat til infusjonsvæske, oppløsning	not available	7377	NOVARTIS NORGE AS	NO
Voltaren 25 mg/ml injeksjonsvæske / konsentrat til infusjonsvæske, oppløsning	not available	7377	NOVARTIS 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
Voltaren 25 mg/ml injeksjonsvæske / konsentrat til infusjonsvæske, oppløsning	not available	7377	NOVARTIS NORGE AS	NO
Voltaren 25 mg/ml injeksjonsvæske / konsentrat til infusjonsvæske, oppløsning	not available	7377	NOVARTIS NORGE AS	NO
Voltaren 25 mg/ml injeksjonsvæske / konsentrat til infusjonsvæske, oppløsning	not available	7377	NOVARTIS NORGE AS	NO
Voltaren 25 mg/ml injeksjonsvæske / konsentrat til infusjonsvæske, oppløsning	not available	7377	NOVARTIS NORGE AS	NO
Voltaren 50 mg enterotabletter	not available	7781	NOVARTIS NORGE AS	NO
Voltaren 50 mg enterotabletter	not available	7781	NOVARTIS NORGE AS	NO
Voltaren 50 mg enterotabletter	not available	7781	NOVARTIS NORGE AS	NO
Voltaren 50 mg enterotabletter	not available	7781	NOVARTIS NORGE AS	NO
Voltaren 50 mg enterotabletter	not available	7781	NOVARTIS NORGE AS	NO
Voltaren 50 mg enterotabletter	not available	7781	NOVARTIS NORGE AS	NO
Voltaren 50 mg stikkpiller	not available	7783	NOVARTIS NORGE AS	NO
Voltaren 50 mg stikkpiller	not available	7783	NOVARTIS NORGE AS	NO
Voltaren 50 mg stikkpiller	not available	7783	NOVARTIS 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
Voltaren 50 mg stikkpiller	not available	7783	NOVARTIS NORGE AS	NO
Voltaren 50 mg stikkpiller	not available	7783	NOVARTIS NORGE AS	NO
Voltaren 50 mg stikkpiller	not available	7783	NOVARTIS NORGE AS	NO
AKIS, 25 mg/ml, roztwór do wstrzykiwań	HU/H/0796/001	24766	IBSA FARMACEUTICI ITALIA	PL
AKIS, 25 mg/ml, roztwór do wstrzykiwań w ampułko-strzykawce	HU/H/0796/004	24769	IBSA FARMACEUTICI ITALIA	PL
AKIS, 50 mg/ml, roztwór do wstrzykiwań	HU/H/0796/002	24767	IBSA FARMACEUTICI ITALIA	PL
AKIS, 50 mg/ml, roztwór do wstrzykiwań w ampułko-strzykawce	HU/H/0796/005	24770	IBSA FARMACEUTICI ITALIA	PL
AKIS, 75 mg/ml, roztwór do wstrzykiwań	HU/H/0796/003	24768	IBSA FARMACEUTICI ITALIA	PL
AKIS, 75 mg/ml, roztwór do wstrzykiwań w ampułko-strzykawce	HU/H/0796/006	24771	IBSA FARMACEUTICI ITALIA	PL
DICLAC 150 DUO, 150 MG, TABLETKI O ZMODYFIKOWANYM UWALNIANIU	not available	9578	SANDOZ GMBH	PL
DICLAC 75 DUO, 75 MG, TABLETKI O ZMODYFIKOWANYM UWALNIANIU	not available	9577	SANDOZ GMBH	PL
Flectorgo, 12,5 mg, kapsułki, miękkie	CZ/H/0928/001	25425	IBSA FARMACEUTICI ITALIA	PL
Voltaren Acti Forte, 25 mg, tabletki powlekane	not available	16489	HALEON POLAND SP. Z O.O.	PL
Voltaren Acti, 12,5 mg, tabletki powlekane	not available	9462	GLAXOSMITHKLINE CONSUMER HEALTHCARE SP. Z O.O.	PL
Voltaren Express Forte, 25 mg, kapsułki miękkie	not available	17890	HALEON POLAND SP. Z O.O.	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
Voltaren Express, 12,5 mg, kapsułki miękkie	not available	16009	HALEON POLAND SP. Z O.O.	PL
Voltaren SR 100, 100 mg, tabletki o przedłużonym uwalnianiu	not available	R/1207	NOVARTIS POLAND SP. Z O. O.	PL
Voltaren SR 100, 100 mg, tabletki o przedłużonym uwalnianiu	not available	R/1207	NOVARTIS POLAND SP. Z O. O.	PL
Voltaren SR 100, 100 mg, tabletki o przedłużonym uwalnianiu	not available	R/1207	NOVARTIS POLAND SP. Z O. O.	PL
Voltaren SR 100, 100 mg, tabletki o przedłużonym uwalnianiu	not available	R/1207	NOVARTIS POLAND SP. Z O. O.	PL
Voltaren SR 100, 100 mg, tabletki o przedłużonym uwalnianiu	not available	R/1207	NOVARTIS POLAND SP. Z O. O.	PL
Voltaren SR 75, 75 mg, tabletki o przedłużonym uwalnianiu	not available	R/6637	NOVARTIS POLAND SP. Z O. O.	PL
Voltaren SR 75, 75 mg, tabletki o przedłużonym uwalnianiu	not available	R/6637	NOVARTIS POLAND SP. Z O. O.	PL
Voltaren SR 75, 75 mg, tabletki o przedłużonym uwalnianiu	not available	R/6637	NOVARTIS POLAND SP. Z O. O.	PL
Voltaren SR 75, 75 mg, tabletki o przedłużonym uwalnianiu	not available	R/6637	NOVARTIS POLAND SP. Z O. O.	PL
Voltaren SR 75, 75 mg, tabletki o przedłużonym uwalnianiu	not available	R/6637	NOVARTIS POLAND SP. Z O. O.	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
Voltaren, 25 mg/ml, roztwór do wstrzykiwań lub do sporządzania roztworu do infuzji	not available	R/1205	NOVARTIS POLAND SP. Z O. O.	PL
Voltaren, 25 mg/ml, roztwór do wstrzykiwań lub do sporządzania roztworu do infuzji	not available	R/1205	NOVARTIS POLAND SP. Z O. O.	PL
Voltaren, 25 mg/ml, roztwór do wstrzykiwań lub do sporządzania roztworu do infuzji	not available	R/1205	NOVARTIS POLAND SP. Z O. O.	PL
Voltaren, 25 mg/ml, roztwór do wstrzykiwań lub do sporządzania roztworu do infuzji	not available	R/1205	NOVARTIS POLAND SP. Z O. O.	PL
Voltaren, 25 mg/ml, roztwór do wstrzykiwań lub do sporządzania roztworu do infuzji	not available	R/1205	NOVARTIS POLAND SP. Z O. O.	PL
Voltaren 100 mg supositórios	not available	9447037	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 100 mg supositórios	not available	9447037	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 100 mg supositórios	not available	9447037	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 100 mg supositórios	not available	9447037	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 100 mg supositórios	not available	9447037	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 100 mg supositórios	not available	9447037	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS 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
Voltaren 12,5 12,5 mg comprimidos revestidos por película	not available	5047246	HALEON PORTUGAL, LDA	PT
Voltaren 12,5 12,5 mg comprimidos revestidos por película	not available	5047253	HALEON PORTUGAL, LDA	PT
Voltaren 12,5 12,5 mg comprimidos revestidos por película	not available	5185533	HALEON PORTUGAL, LDA	PT
Voltaren 12,5 12,5 mg comprimidos revestidos por película	not available	3852589	HALEON PORTUGAL, LDA	PT
Voltaren 12,5 12,5 mg comprimidos revestidos por película	not available	3852688	HALEON PORTUGAL, LDA	PT
Voltaren 12,5 12,5 mg comprimidos revestidos por película	not available	5185525	HALEON PORTUGAL, LDA	PT
Voltaren 25, 25 mg, cápsulas moles	not available	5204466	HALEON PORTUGAL, LDA	PT
Voltaren 25, 25 mg, cápsulas moles	not available	5204508	HALEON PORTUGAL, LDA	PT
Voltaren 25, 25 mg, cápsulas moles	not available	5204458	HALEON PORTUGAL, LDA	PT
Voltaren 25, 25 mg, cápsulas moles	not available	5204474	HALEON PORTUGAL, LDA	PT
Voltaren 50 mg comprimidos gastrorresistentes	not available	4636098	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 50 mg comprimidos gastrorresistentes	not available	4636197	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 50 mg comprimidos gastrorresistentes	not available	9427815	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 50 mg comprimidos gastrorresistentes	not available	9427831	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS 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
Voltaren 50 mg comprimidos gastrorresistentes	not available	4636098	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 50 mg comprimidos gastrorresistentes	not available	4636197	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 50 mg comprimidos gastrorresistentes	not available	9427815	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 50 mg comprimidos gastrorresistentes	not available	9427831	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 50 mg comprimidos gastrorresistentes	not available	4636098	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 50 mg comprimidos gastrorresistentes	not available	4636197	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 50 mg comprimidos gastrorresistentes	not available	9427815	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 50 mg comprimidos gastrorresistentes	not available	9427831	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 50 mg comprimidos gastrorresistentes	not available	4636098	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 50 mg comprimidos gastrorresistentes	not available	4636197	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 50 mg comprimidos gastrorresistentes	not available	9427815	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 50 mg comprimidos gastrorresistentes	not available	9427831	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 50 mg comprimidos gastrorresistentes	not available	4636098	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS 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
Voltaren 50 mg comprimidos gastrorresistentes	not available	4636197	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 50 mg comprimidos gastrorresistentes	not available	9427815	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 50 mg comprimidos gastrorresistentes	not available	9427831	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 50 mg comprimidos gastrorresistentes	not available	4636098	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 50 mg comprimidos gastrorresistentes	not available	4636197	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 50 mg comprimidos gastrorresistentes	not available	9427815	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 50 mg comprimidos gastrorresistentes	not available	9427831	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 75 75 mg comprimidos de libertação prolongada	not available	4726394	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 75 75 mg comprimidos de libertação prolongada	not available	4726493	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 75 75 mg comprimidos de libertação prolongada	not available	9427856	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 75 75 mg comprimidos de libertação prolongada	not available	9427864	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 75 75 mg comprimidos de libertação prolongada	not available	4726394	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 75 75 mg comprimidos de libertação prolongada	not available	4726493	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS 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
Voltaren 75 75 mg comprimidos de libertação prolongada	not available	9427856	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 75 75 mg comprimidos de libertação prolongada	not available	9427864	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 75 75 mg comprimidos de libertação prolongada	not available	4726394	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 75 75 mg comprimidos de libertação prolongada	not available	4726493	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 75 75 mg comprimidos de libertação prolongada	not available	9427856	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 75 75 mg comprimidos de libertação prolongada	not available	9427864	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 75 75 mg comprimidos de libertação prolongada	not available	4726394	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 75 75 mg comprimidos de libertação prolongada	not available	4726493	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 75 75 mg comprimidos de libertação prolongada	not available	9427856	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 75 75 mg comprimidos de libertação prolongada	not available	9427864	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 75 75 mg comprimidos de libertação prolongada	not available	4726394	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 75 75 mg comprimidos de libertação prolongada	not available	4726493	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 75 75 mg comprimidos de libertação prolongada	not available	9427856	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS 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
Voltaren 75 75 mg comprimidos de libertação prolongada	not available	9427864	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 75 75 mg comprimidos de libertação prolongada	not available	4726394	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 75 75 mg comprimidos de libertação prolongada	not available	4726493	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 75 75 mg comprimidos de libertação prolongada	not available	9427856	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 75 75 mg comprimidos de libertação prolongada	not available	9427864	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 75 mg/3 ml solução injetável	not available	4636593	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 75 mg/3 ml solução injetável	not available	8446906	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 75 mg/3 ml solução injetável	not available	4636593	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 75 mg/3 ml solução injetável	not available	8446906	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 75 mg/3 ml solução injetável	not available	4636593	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 75 mg/3 ml solução injetável	not available	8446906	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 75 mg/3 ml solução injetável	not available	4636593	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 75 mg/3 ml solução injetável	not available	8446906	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS 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
Voltaren 75 mg/3 ml solução injetável	not available	4636593	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 75 mg/3 ml solução injetável	not available	8446906	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 75 mg/3 ml solução injetável	not available	4636593	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren 75 mg/3 ml solução injetável	not available	8446906	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
VOLTAREN Rapid 50 mg comprimidos revestidos	not available	2134492	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
VOLTAREN Rapid 50 mg comprimidos revestidos	not available	2134591	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
VOLTAREN Rapid 50 mg comprimidos revestidos	not available	2134690	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
VOLTAREN Rapid 50 mg comprimidos revestidos	not available	4636494	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
VOLTAREN Rapid 50 mg comprimidos revestidos	not available	2134492	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
VOLTAREN Rapid 50 mg comprimidos revestidos	not available	2134591	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
VOLTAREN Rapid 50 mg comprimidos revestidos	not available	2134690	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
VOLTAREN Rapid 50 mg comprimidos revestidos	not available	4636494	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
VOLTAREN Rapid 50 mg comprimidos revestidos	not available	2134492	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS 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
VOLTAREN Rapid 50 mg comprimidos revestidos	not available	2134591	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
VOLTAREN Rapid 50 mg comprimidos revestidos	not available	2134690	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
VOLTAREN Rapid 50 mg comprimidos revestidos	not available	4636494	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
VOLTAREN Rapid 50 mg comprimidos revestidos	not available	2134492	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
VOLTAREN Rapid 50 mg comprimidos revestidos	not available	2134591	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
VOLTAREN Rapid 50 mg comprimidos revestidos	not available	2134690	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
VOLTAREN Rapid 50 mg comprimidos revestidos	not available	4636494	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
VOLTAREN Rapid 50 mg comprimidos revestidos	not available	2134492	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
VOLTAREN Rapid 50 mg comprimidos revestidos	not available	2134591	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
VOLTAREN Rapid 50 mg comprimidos revestidos	not available	2134690	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
VOLTAREN Rapid 50 mg comprimidos revestidos	not available	4636494	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
VOLTAREN Rapid 50 mg comprimidos revestidos	not available	2134492	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
VOLTAREN Rapid 50 mg comprimidos revestidos	not available	2134591	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS 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
VOLTAREN Rapid 50 mg comprimidos revestidos	not available	2134690	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
VOLTAREN Rapid 50 mg comprimidos revestidos	not available	4636494	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren Retard 100 mg comprimidos de libertação prolongada	not available	9427823	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren Retard 100 mg comprimidos de libertação prolongada	not available	9427823	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren Retard 100 mg comprimidos de libertação prolongada	not available	9427823	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren Retard 100 mg comprimidos de libertação prolongada	not available	9427823	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren Retard 100 mg comprimidos de libertação prolongada	not available	9427823	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Voltaren Retard 100 mg comprimidos de libertação prolongada	not available	9427823	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Dicloreum 150 mg capsule cu eliberare prelungită	not available	8293/2015/01	ALFASIGMA S.P.A.	RO
Voltaren 25 mg enterotabletter.	not available	9737	NOVARTIS SVERIGE AB	SE
Voltaren 25 mg enterotabletter.	not available	9737	NOVARTIS SVERIGE AB	SE
Voltaren 25 mg enterotabletter.	not available	9737	NOVARTIS SVERIGE AB	SE
Voltaren 25 mg enterotabletter.	not available	9737	NOVARTIS SVERIGE AB	SE
Voltaren 25 mg enterotabletter.	not available	9737	NOVARTIS SVERIGE 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
Voltaren 25 mg enterotabletter.	not available	9737	NOVARTIS SVERIGE AB	SE
Voltaren 25 mg/ml, injektionsvätska, lösning.	not available	09940	NOVARTIS SVERIGE AB	SE
Voltaren 25 mg/ml, injektionsvätska, lösning.	not available	09940	NOVARTIS SVERIGE AB	SE
Voltaren 25 mg/ml, injektionsvätska, lösning.	not available	09940	NOVARTIS SVERIGE AB	SE
Voltaren 25 mg/ml, injektionsvätska, lösning.	not available	09940	NOVARTIS SVERIGE AB	SE
Voltaren 25 mg/ml, injektionsvätska, lösning.	not available	09940	NOVARTIS SVERIGE AB	SE
Voltaren 25 mg/ml, injektionsvätska, lösning.	not available	09940	NOVARTIS SVERIGE AB	SE
Voltaren 50 mg enterotabletter	not available	9738	NOVARTIS SVERIGE AB	SE
Voltaren 50 mg enterotabletter	not available	9738	NOVARTIS SVERIGE AB	SE
Voltaren 50 mg enterotabletter	not available	9738	NOVARTIS SVERIGE AB	SE
Voltaren 50 mg enterotabletter	not available	9738	NOVARTIS SVERIGE AB	SE
Voltaren 50 mg enterotabletter	not available	9738	NOVARTIS SVERIGE AB	SE
Voltaren 50 mg enterotabletter	not available	9738	NOVARTIS SVERIGE AB	SE
Voltaren suppositorier 100 mg	not available	9818	NOVARTIS SVERIGE AB	SE
Voltaren suppositorier 100 mg	not available	9818	NOVARTIS SVERIGE AB	SE
Voltaren suppositorier 100 mg	not available	9818	NOVARTIS SVERIGE 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
Voltaren suppositorier 100 mg	not available	9818	NOVARTIS SVERIGE AB	SE
Voltaren suppositorier 100 mg	not available	9818	NOVARTIS SVERIGE AB	SE
Voltaren suppositorier 100 mg	not available	9818	NOVARTIS SVERIGE AB	SE
Voltaren suppositorier 25 mg	not available	12840	NOVARTIS SVERIGE AB	SE
Voltaren suppositorier 25 mg	not available	12840	NOVARTIS SVERIGE AB	SE
Voltaren suppositorier 25 mg	not available	12840	NOVARTIS SVERIGE AB	SE
Voltaren suppositorier 25 mg	not available	12840	NOVARTIS SVERIGE AB	SE
Voltaren suppositorier 25 mg	not available	12840	NOVARTIS SVERIGE AB	SE
Voltaren suppositorier 25 mg	not available	12840	NOVARTIS SVERIGE AB	SE
Voltaren suppositorier 50 mg	not available	9941	NOVARTIS SVERIGE AB	SE
Voltaren suppositorier 50 mg	not available	9941	NOVARTIS SVERIGE AB	SE
Voltaren suppositorier 50 mg	not available	9941	NOVARTIS SVERIGE AB	SE
Voltaren suppositorier 50 mg	not available	9941	NOVARTIS SVERIGE AB	SE
Voltaren suppositorier 50 mg	not available	9941	NOVARTIS SVERIGE AB	SE
Voltaren suppositorier 50 mg	not available	9941	NOVARTIS SVERIGE AB	SE
Voltaren T 12,5 mg filmdragerade tabletter	not available	16081	HALEON DENMARK APS	SE
Voltaren T 25 mg mjuka kapslar.	not available	41904	HALEON DENMARK APS	SE
Voltaren T 25 mg, dragerade tabletter	not available	12427	HALEON DENMARK APS	SE
Voltaren T 50 mg, dragerade tabletter.	not available	11242	NOVARTIS SVERIGE 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
Voltaren T 50 mg, dragerade tabletter.	not available	11242	NOVARTIS SVERIGE AB	SE
Voltaren T 50 mg, dragerade tabletter.	not available	11242	NOVARTIS SVERIGE AB	SE
Voltaren T 50 mg, dragerade tabletter.	not available	11242	NOVARTIS SVERIGE AB	SE
Voltaren T 50 mg, dragerade tabletter.	not available	11242	NOVARTIS SVERIGE AB	SE
Voltaren 12,5 mg svečke	not available	H/94/01506/001	NOVARTIS EUROPHARM LIMITED	SI
Voltaren 12,5 mg svečke	not available	H/94/01506/001	NOVARTIS EUROPHARM LIMITED	SI
Voltaren 12,5 mg svečke	not available	H/94/01506/001	NOVARTIS EUROPHARM LIMITED	SI
Voltaren 12,5 mg svečke	not available	H/94/01506/001	NOVARTIS EUROPHARM LIMITED	SI
Voltaren 12,5 mg svečke	not available	H/94/01506/001	NOVARTIS EUROPHARM LIMITED	SI
Voltaren 25 mg svečke	not available	H/94/01506/002	NOVARTIS EUROPHARM LIMITED	SI
Voltaren 25 mg svečke	not available	H/94/01506/002	NOVARTIS EUROPHARM LIMITED	SI
Voltaren 25 mg svečke	not available	H/94/01506/002	NOVARTIS EUROPHARM LIMITED	SI
Voltaren 25 mg svečke	not available	H/94/01506/002	NOVARTIS EUROPHARM LIMITED	SI
Voltaren 25 mg svečke	not available	H/94/01506/002	NOVARTIS EUROPHARM LIMITED	SI
Voltaren Dolo 12,5 mg filmsko obložene tablete	not available	H/09/01954/001	HALEON HUNGARY KFT.	SI
Voltaren Dolo 12,5 mg filmsko obložene tablete	not available	H/09/01954/002	HALEON HUNGARY KFT.	SI
Voltaren Dolo 12,5 mg filmsko obložene tablete	not available	H/09/01954/003	HALEON HUNGARY KFT.	SI
Voltaren Dolo 12,5 mg filmsko obložene tablete	not available	H/09/01954/004	HALEON HUNGARY KFT.	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
AKIS Plus 25 mg/ml injekčný roztok naplnený v injekčnej striekačke	HU/H/0796/004	29/0331/17-S	IBSA SLOVAKIA S.R.O.	SK
AKIS Plus 50 mg/ml injekčný roztok naplnený v injekčnej striekačke	HU/H/0796/005	29/0332/17-S	IBSA SLOVAKIA S.R.O.	SK
AKIS Plus 75 mg/ml injekčný roztok naplnený v injekčnej striekačke	HU/H/0796/006	29/0333/17-S	IBSA SLOVAKIA S.R.O.	SK
Flector 12,5 mg	CZ/H/0928/001	29/0249/17-S	IBSA SLOVAKIA S.R.O.	SK
Flector Rapid 50 mg	not available	29/0906/96-S	IBSA SLOVAKIA S.R.O.	SK
Voltaren 100 100 mg čapíky	not available	29/0184/80-CS	NOVARTIS SLOVAKIA S.R.O.	SK
Voltaren 100 100 mg čapíky	not available	29/0184/80-CS	NOVARTIS SLOVAKIA S.R.O.	SK
Voltaren 100 100 mg čapíky	not available	29/0184/80-CS	NOVARTIS SLOVAKIA S.R.O.	SK
Voltaren 100 100 mg čapíky	not available	29/0184/80-CS	NOVARTIS SLOVAKIA S.R.O.	SK
Voltaren 100 100 mg čapíky	not available	29/0184/80-CS	NOVARTIS SLOVAKIA S.R.O.	SK
Voltaren 100 100 mg čapíky	not available	29/0184/80-CS	NOVARTIS SLOVAKIA S.R.O.	SK
Voltaren 50 50 mg gastrorezistentné tablety	not available	29/0294/91-CS	NOVARTIS SLOVAKIA S.R.O.	SK
Voltaren 50 50 mg gastrorezistentné tablety	not available	29/0294/91-CS	NOVARTIS SLOVAKIA S.R.O.	SK
Voltaren 50 50 mg gastrorezistentné tablety	not available	29/0294/91-CS	NOVARTIS SLOVAKIA S.R.O.	SK
Voltaren 50 50 mg gastrorezistentné tablety	not available	29/0294/91-CS	NOVARTIS SLOVAKIA S.R.O.	SK
Voltaren 50 50 mg gastrorezistentné tablety	not available	29/0294/91-CS	NOVARTIS SLOVAKIA S.R.O.	SK
Voltaren 50 50 mg gastrorezistentné tablety	not available	29/0294/91-CS	NOVARTIS SLOVAKIA S.R.O.	SK
Voltaren Actigo Extra 25 mg obalené tablety	not available	29/0227/01-S	HALEON CZECH REPUBLIC S.R.O.	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
Voltaren Rapid 25 mg mäkké kapsuly	not available	29/0709/10-S	HALEON CZECH REPUBLIC S.R.O.	SK
Voltaren Rapid 50 mg obalené tablety	not available	29/0098/91-S	NOVARTIS SLOVAKIA S.R.O.	SK
Voltaren Rapid 50 mg obalené tablety	not available	29/0098/91-S	NOVARTIS SLOVAKIA S.R.O.	SK
Voltaren Rapid 50 mg obalené tablety	not available	29/0098/91-S	NOVARTIS SLOVAKIA S.R.O.	SK
Voltaren Rapid 50 mg obalené tablety	not available	29/0098/91-S	NOVARTIS SLOVAKIA S.R.O.	SK
Voltaren Rapid 50 mg obalené tablety	not available	29/0098/91-S	NOVARTIS SLOVAKIA S.R.O.	SK
Voltaren Rapid 50 mg obalené tablety	not available	29/0098/91-S	NOVARTIS SLOVAKIA S.R.O.	SK
Voltaren Rapid 50 mg obalené tablety	not available	29/0098/91-S	NOVARTIS SLOVAKIA S.R.O.	SK
Voltaren Rapid 50 mg obalené tablety	not available	29/0098/91-S	NOVARTIS SLOVAKIA S.R.O.	SK
Voltaren Rapid 50 mg obalené tablety	not available	29/0098/91-S	NOVARTIS SLOVAKIA S.R.O.	SK
Voltaren Rapid 50 mg obalené tablety	not available	29/0098/91-S	NOVARTIS SLOVAKIA S.R.O.	SK
Voltaren Retard 100 100 mg tablety s predĺženým uvoľňovaním	not available	29/0247/80-CS	NOVARTIS SLOVAKIA S.R.O.	SK
Voltaren Retard 100 100 mg tablety s predĺženým uvoľňovaním	not available	29/0247/80-CS	NOVARTIS SLOVAKIA S.R.O.	SK
Voltaren Retard 100 100 mg tablety s predĺženým uvoľňovaním	not available	29/0247/80-CS	NOVARTIS SLOVAKIA S.R.O.	SK
Voltaren Retard 100 100 mg tablety s predĺženým uvoľňovaním	not available	29/0247/80-CS	NOVARTIS SLOVAKIA S.R.O.	SK
Voltaren Retard 100 100 mg tablety s predĺženým uvoľňovaním	not available	29/0247/80-CS	NOVARTIS SLOVAKIA S.R.O.	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
Voltaren Retard 100 100 mg tablety s predĺženým uvoľňovaním	not available	29/0247/80-CS	NOVARTIS SLOVAKIA S.R.O.	SK
Voltaren Retard 100 100 mg tablety s predĺženým uvoľňovaním	not available	29/0247/80-CS	NOVARTIS SLOVAKIA S.R.O.	SK
Voltaren Retard 100 100 mg tablety s predĺženým uvoľňovaním	not available	29/0247/80-CS	NOVARTIS SLOVAKIA S.R.O.	SK
Voltaren Retard 100 100 mg tablety s predĺženým uvoľňovaním	not available	29/0247/80-CS	NOVARTIS SLOVAKIA S.R.O.	SK
Voltaren Retard 100 100 mg tablety s predĺženým uvoľňovaním	not available	29/0247/80-CS	NOVARTIS SLOVAKIA S.R.O.	SK
Voltaren Retard 100 100 mg tablety s predĺženým uvoľňovaním	not available	29/0247/80-CS	NOVARTIS SLOVAKIA S.R.O.	SK
Voltaren Retard 100 100 mg tablety s predĺženým uvoľňovaním	not available	29/0247/80-CS	NOVARTIS SLOVAKIA S.R.O.	SK
AKIS 25 mg/ml solution for injection in prefilled syringe	HU/H/0796/004	PL 21039/0043	IBSA FARMACEUTICI ITALIA	XI
AKIS 50 mg/ml solution for injection in prefilled syringe	HU/H/0796/005	PL 21039/0044	IBSA FARMACEUTICI ITALIA	XI
AKIS 75 mg/ml solution for injection in prefilled syringe	HU/H/0796/006	PL 21039/0045	IBSA FARMACEUTICI ITALIA	XI
Diclofenac potassium 12.5mg tablets	not available	PL 36687/0214	TORRENT PHARMA (UK) LTD.	XI
Diclofenac Potassium 25 mg Tablets	not available	PL 20075/0682	ACCORD HEALTHCARE LIMITED	XI
Diclofenac Potassium 50 mg Tablets	not available	PL 20075/0683	ACCORD HEALTHCARE LIMITED	XI
Diclofenac Sodium 25 mg Tablets	not available	PL 33414/0038	CHELONIA HEALTHCARE LIMITED	XI

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
Diclofenac Sodium 50 mg Tablets	not available	PL 33414/0143	CHELONIA HEALTHCARE LIMITED	XI
Diclofenac Tablets 25 mg	not available	PL 04416/0361	SANDOZ LTD	XI
Diclofenac Tablets BP 25mg	not available	PL 20416/0219	CRESCENT PHARMA LIMITED	XI
Diclofenac Tablets BP 50mg	not available	PL 20416/0220	CRESCENT PHARMA LIMITED	XI
Diclomax Retard.	not available	PL 27827/0005	GALEN LIMITED	XI
Diclovol 75mg SR Tablets	not available	PL 04416/0645	SANDOZ LTD	XI
Econac SR 75 mg Tablets	not available	PL 12762/0100	MERCURY PHARMACEUTICALS LTD.	XI
Econac SR 75 mg Tablets	not available	PL 12762/0100	MERCURY PHARMACEUTICALS LTD.	XI
Econac SR 75 mg Tablets	not available	PL 12762/0100	MERCURY PHARMACEUTICALS LTD.	XI
Econac SR 75 mg Tablets	not available	PL 12762/0100	MERCURY PHARMACEUTICALS LTD.	XI
Econac SR 75 mg Tablets	not available	PL 12762/0100	MERCURY PHARMACEUTICALS LTD.	XI
Econac SR 75 mg Tablets	not available	PL 12762/0100	MERCURY PHARMACEUTICALS LTD.	XI
Econac SR 75 mg Tablets	not available	PL 12762/0100	MERCURY PHARMACEUTICALS LTD.	XI
Econac SR 75 mg Tablets	not available	PL 12762/0100	MERCURY PHARMACEUTICALS LTD.	XI
Econac SR 75 mg Tablets	not available	PL 12762/0100	MERCURY PHARMACEUTICALS LTD.	XI
Econac SR 75 mg Tablets	not available	PL 12762/0100	MERCURY PHARMACEUTICALS LTD.	XI
Econac SR 75 mg Tablets	not available	PL 12762/0100	MERCURY PHARMACEUTICALS LTD.	XI
Econac SR 75 mg Tablets	not available	PL 12762/0100	MERCURY PHARMACEUTICALS LTD.	XI
Rhumalgan CR 100	not available	PL 04416/0243	SANDOZ LTD	XI
Rhumalgan CR 75	not available	PL 04416/0242	SANDOZ LTD	XI

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
Rhumalgan SR 75 mg Modified Release Capsules	not available	PL 04416/0737	SANDOZ LTD	XI
Rhumalgan XL 100 mg modified-release capsules.	not available	PL 04416/0738	SANDOZ LTD	XI
Voltarol® Ampoules	not available	PL 00101/0466	NOVARTIS PHARMACEUTICALS UK LIMITED	XI
Voltarol® Ampoules	not available	PL 00101/0466	NOVARTIS PHARMACEUTICALS UK LIMITED	XI
Voltarol® Ampoules	not available	PL 00101/0466	NOVARTIS PHARMACEUTICALS UK LIMITED	XI
Voltarol® Ampoules	not available	PL 00101/0466	NOVARTIS PHARMACEUTICALS UK LIMITED	XI
Voltarol® Rapid Tablets 50mg	not available	PL 00101/0482	NOVARTIS PHARMACEUTICALS UK LIMITED	XI
Voltarol® Rapid Tablets 50mg	not available	PL 00101/0482	NOVARTIS PHARMACEUTICALS UK LIMITED	XI
Voltarol® Rapid Tablets 50mg	not available	PL 00101/0482	NOVARTIS PHARMACEUTICALS UK LIMITED	XI
Voltarol® Rapid Tablets 50mg	not available	PL 00101/0482	NOVARTIS PHARMACEUTICALS UK LIMITED	XI
Voltarol® Suppositories 100mg	not available	PL 00101/0475	NOVARTIS PHARMACEUTICALS UK LIMITED	XI
Voltarol® Suppositories 100mg	not available	PL 00101/0475	NOVARTIS PHARMACEUTICALS UK LIMITED	XI
Voltarol® Suppositories 100mg	not available	PL 00101/0475	NOVARTIS PHARMACEUTICALS UK LIMITED	XI

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
Voltarol® Suppositories 100mg	not available	PL 00101/0475	NOVARTIS PHARMACEUTICALS UK LIMITED	XI
Voltarol® Suppositories 12.5mg	not available	PL 00101/0472	NOVARTIS PHARMACEUTICALS UK LIMITED	XI
Voltarol® Suppositories 12.5mg	not available	PL 00101/0472	NOVARTIS PHARMACEUTICALS UK LIMITED	XI
Voltarol® Suppositories 12.5mg	not available	PL 00101/0472	NOVARTIS PHARMACEUTICALS UK LIMITED	XI
Voltarol® Suppositories 12.5mg	not available	PL 00101/0472	NOVARTIS PHARMACEUTICALS UK LIMITED	XI
Voltarol® Suppositories 25mg	not available	PL 00101/0473	NOVARTIS PHARMACEUTICALS UK LIMITED	XI
Voltarol® Suppositories 25mg	not available	PL 00101/0473	NOVARTIS PHARMACEUTICALS UK LIMITED	XI
Voltarol® Suppositories 25mg	not available	PL 00101/0473	NOVARTIS PHARMACEUTICALS UK LIMITED	XI
Voltarol® Suppositories 25mg	not available	PL 00101/0473	NOVARTIS PHARMACEUTICALS UK LIMITED	XI
Voltarol® Suppositories 50mg	not available	PL 00101/0474	NOVARTIS PHARMACEUTICALS UK LIMITED	XI
Voltarol® Suppositories 50mg	not available	PL 00101/0474	NOVARTIS PHARMACEUTICALS UK LIMITED	XI
Voltarol® Suppositories 50mg	not available	PL 00101/0474	NOVARTIS PHARMACEUTICALS UK LIMITED	XI
Voltarol® Suppositories 50mg	not available	PL 00101/0474	NOVARTIS PHARMACEUTICALS UK LIMITED	XI