17 October 2024
EMA/459995/2024
Human Medicines Division

List of nationally authorised medicinal products

Active substance(s): mesalazine

Procedure No. PSUSA/00001990/202402

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Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
ARGONAL 2 g/50 ml sospensione rettale	not available	034254033	DICOFARM S.P.A.	IT
ARGONAL 4 g/100 ml sospensione rettale	not available	034254045	DICOFARM S.P.A.	IT
ARGONAL 500 mg supposte	not available	034254058	DICOFARM S.P.A.	IT
ARGONAL 500 mg supposte	not available	034254058	DICOFARM S.P.A.	IT
ARGONAL 800 mg compresse gastroresistenti	not available	034254021	DICOFARM S.P.A.	IT
ARGONAL 800 mg compresse gastroresistenti	not available	034254060	DICOFARM S.P.A.	IT
ARGONAL 400mg COMPRESSE GASTRORESISTENTI	not available	034254019	DICOFARM S.P.A.	IT
Asacol	not available	121388/08-12-2021	TILLOTTS PHARMA ITALY, S.R.L.,	GR
Asacol	not available	121388/08-12-2021	TILLOTTS PHARMA ITALY, S.R.L.,	GR
Asacol	not available	121388/08-12-2021	TILLOTTS PHARMA ITALY, S.R.L.,	GR
Asacol	not available	121388/08-12-2021	TILLOTTS PHARMA ITALY, S.R.L.,	GR
Asacol 500 mg žvakutės	not available	LT/1/05/0283/003	TILLOTTS PHARMA AB	LT
Asacol 1 g čípky	SE/H/2039/001	29/487/19-C	TILLOTTS PHARMA GMBH	CZ
Asacol 1 g čípky	SE/H/2039/001	29/487/19-C	TILLOTTS PHARMA GMBH	CZ

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Asacol 1 g endaþarmsdreifa	not available	970314 (IS)	TILLOTTS PHARMA AB	IS
Asacol 1 g endaþarmsstílar	SE/H/2039/001	IS/1/21/017/01	TILLOTTS PHARMA AB	IS
Asacol 1 g endaþarmsstílar	SE/H/2039/001	IS/1/21/017/01	TILLOTTS PHARMA AB	IS
Asacol 1 g peräpuikot	SE/H/2039/001	37770	TILLOTTS PHARMA AB	FI
Asacol 1 g peräpuikot	SE/H/2039/001	37770	TILLOTTS PHARMA AB	FI
Asacol 1 g peräpuikot	SE/H/2039/001	37770	TILLOTTS PHARMA AB	FI
Asacol 1 g peräpuikot	SE/H/2039/001	37770	TILLOTTS PHARMA AB	FI
Asacol 1 g peräpuikot	SE/H/2039/001	37770	TILLOTTS PHARMA AB	FI
Asacol 1 g peräpuikot	SE/H/2039/001	37770	TILLOTTS PHARMA AB	FI
Asacol 1 g peräpuikot	SE/H/2039/001	37770	TILLOTTS PHARMA AB	FI
Asacol 1 g peräpuikot	SE/H/2039/001	37770	TILLOTTS PHARMA AB	FI
Asacol 1 g peräpuikot	SE/H/2039/001	37770	TILLOTTS PHARMA AB	FI
Asacol 1 g peräpuikot	SE/H/2039/001	37770	TILLOTTS PHARMA AB	FI
Asacol 1 g peräruiske	not available	12432	TILLOTTS PHARMA AB	FI
Asacol 1 g rektalsuspension	not available	13251	TILLOTTS PHARMA AB	SE

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
ASACOL 1 g rektalvæske, suspensjon	not available	96-3269	TILLOTTS PHARMA AB	NO
ASACOL 1 g rektalvätska	not available	12432	TILLOTTS PHARMA AB	FI
ASACOL 1 g schiuma rettale	not available	026416267	GIULIANI SPA	IT
ASACOL 1 g schiuma rettale	not available	026416279	GIULIANI SPA	IT
Asacol 1 g stikkpiller	SE/H/2039/001	20-13255	TILLOTTS PHARMA AB	NO
Asacol 1 g stikkpiller	SE/H/2039/001	20-13255	TILLOTTS PHARMA AB	NO
Asacol 1 g supositorios	SE/H/2039/001	85816	TILLOTTS PHARMA SPAIN S.A.	ES
Asacol 1 g supositorios	SE/H/2039/001	85816	TILLOTTS PHARMA SPAIN S.A.	ES
Asacol 1 g suppositorier	SE/H/2039/001	63888	TILLOTTS PHARMA AB	DK
Asacol 1 g suppositorier	SE/H/2039/001	63888	TILLOTTS PHARMA AB	DK
Asacol 1 g suppositorier	SE/H/2039/001	37770	TILLOTTS PHARMA AB	FI
Asacol 1 g suppositorier	SE/H/2039/001	37770	TILLOTTS PHARMA AB	FI
Asacol 1 g suppositorier	SE/H/2039/001	37770	TILLOTTS PHARMA AB	FI
Asacol 1 g suppositorier	SE/H/2039/001	37770	TILLOTTS PHARMA AB	FI
Asacol 1 g suppositorier	SE/H/2039/001	37770	TILLOTTS PHARMA AB	FI

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Asacol 1 g suppositorier	SE/H/2039/001	37770	TILLOTTS PHARMA AB	FI
Asacol 1 g suppositorier	SE/H/2039/001	37770	TILLOTTS PHARMA AB	FI
Asacol 1 g suppositorier	SE/H/2039/001	37770	TILLOTTS PHARMA AB	FI
Asacol 1 g suppositorier	SE/H/2039/001	37770	TILLOTTS PHARMA AB	FI
Asacol 1 g suppositorier	SE/H/2039/001	37770	TILLOTTS PHARMA AB	FI
Asacol 1 g suppositorier	SE/H/2039/001	60180	TILLOTTS PHARMA AB	SE
Asacol 1 g suppositorier	SE/H/2039/001	60180	TILLOTTS PHARMA AB	SE
Asacol 1 g Zäpfchen	SE/H/2039/001	2205811.00.00	TILLOTTS PHARMA GMBH	DE
Asacol 1 g Zäpfchen	SE/H/2039/001	2205811.00.00	TILLOTTS PHARMA GMBH	DE
Asacol 1600 mg comprimidos gastrorresistentes	not available	86315	TILLOTTS PHARMA SPAIN S.A.	ES
Asacol 1600 mg comprimidos gastrorresistentes	not available	86315	TILLOTTS PHARMA SPAIN S.A.	ES
Asacol 1600 mg modifikuoto atpalaidavimo tabletės	SE/H/1654/001	LT/1/18/4201/001	TILLOTTS PHARMA AB	LT
Asacol 1600 mg modifikuoto atpalaidavimo tabletės	SE/H/1654/001	LT/1/18/4201/002	TILLOTTS PHARMA AB	LT
Asacol 1600 mg modifikuoto atpalaidavimo tabletės	SE/H/1654/001	LT/1/18/4201/003	TILLOTTS PHARMA AB	LT
Asacol 1600 mg säädellysti vapauttavat tabletit	SE/H/1654/001	34959	TILLOTTS PHARMA AB	FI

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Asacol 1600 mg säädellysti vapauttavat tabletit	SE/H/1654/001	34959	TILLOTTS PHARMA AB	FI
Asacol 1600 mg Tabletten mit veränderter Wirkstofffreisetzung	SE/H/1654/001	2202543.00.00	TILLOTTS PHARMA GMBH	DE
Asacol 1600 mg tabletter med modifierad frisättning	SE/H/1654/001	34959	TILLOTTS PHARMA AB	FI
Asacol 1600 mg tabletter med modifierad frisättning	SE/H/1654/001	34959	TILLOTTS PHARMA AB	FI
Asacol 1600 mg tabletter med modifierad frisättning	SE/H/1654/001	34959	TILLOTTS PHARMA AB	FI
Asacol 1600 mg tabletter med modifierad frisättning	SE/H/1654/001	34959	TILLOTTS PHARMA AB	FI
Asacol 1600 mg tabletter med modifierad frisättning	SE/H/1654/001	55999	TILLOTTS PHARMA AB	SE
Asacol 1600 mg tabletter med modifisert frisetting	SE/H/1654/001	17-11557	TILLOTTS PHARMA AB	NO
Asacol 1600 mg tablety s řízeným uvolňováním	SE/H/1654/001	29/034/17-C	TILLOTTS PHARMA GMBH	CZ
Asacol 1600 mg töflur með breyttan losunarhraða	SE/H/1654/001	IS/1/18/021/01	TILLOTTS PHARMA AB	IS
ASACOL 1g supposte	not available	026416305	GIULIANI SPA	IT
ASACOL 2 g granulato per sospensione rettale	not available	026416091	GIULIANI SPA	IT
ASACOL 2 g schiuma rettale	not available	026416216	GIULIANI SPA	IT
ASACOL 2 g/50 ml sospensione rettale	not available	026416141	GIULIANI SPA	IT
ASACOL 4 g in 100 mL enema	not available	19627	TILLOTTS PHARMA GMBH	CY

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
ASACOL 4 g schiuma rettale	not available	026416230	GIULIANI SPA	IT
ASACOL 4 g/100 ml sospensione rettale	not available	026416154	GIULIANI SPA	IT
ASACOL 4 g/50 ml sospensione rettale	not available	026416166	GIULIANI SPA	IT
Asacol 400 mg	not available	30723/03-05-2012	TILLOTTS PHARMA ITALY, S.R.L.,	GR
Asacol 400 mg	not available	30723/03-05-2012	TILLOTTS PHARMA ITALY, S.R.L.,	GR
ASACOL 400 mg capsule a rilascio modificato	not available	026416255	GIULIANI SPA	IT
ASACOL 400 mg compresse gastroresistenti	not available	026416014	GIULIANI SPA	IT
ASACOL 400 mg compresse gastroresistenti	not available	026416329	GIULIANI SPA	IT
ASACOL 400 mg COMPRIMIDOS GASTRORRESISTENTES	not available	57726	TILLOTTS PHARMA SPAIN S.A.	ES
Asacol 400 mg enterosolventní tablety	not available	29/169/97-C	TILLOTTS PHARMA GMBH	CZ
ASACOL 400 mg enterotabletit	not available	11344	TILLOTTS PHARMA AB	FI
Asacol 400 mg enterotablett	not available	7603	TILLOTTS PHARMA AB	NO
Asacol 400 mg enterotabletter	not available	11344	TILLOTTS PHARMA AB	FI
Asacol 400 mg enterotabletter	not available	11580	TILLOTTS PHARMA AB	SE
Asacol 400 mg enterotabletter	not available	11580	TILLOTTS PHARMA AB	SE

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
ASACOL 400 mg Gastro-resistant Tablets	not available	19504	TILLOTTS PHARMA GMBH	CY
ASACOL 400 mg Gastro-resistant Tablets	not available	19504	TILLOTTS PHARMA GMBH	CY
Asacol 400 mg ilgstošās darbības tabletes	not available	04-0435	TILLOTTS PHARMA AB	LV
Asacol 400 mg magensaftresistente Tabletten	not available	7135.00.00	TILLOTTS PHARMA GMBH	DE
Asacol 400 mg magensaftresistente Tabletten	not available	7135.00.00	TILLOTTS PHARMA GMBH	DE
Asacol 400 mg magensaftresistente Tabletten	not available	7135.00.00	TILLOTTS PHARMA GMBH	DE
Asacol 400 mg modifikuoto atpalaidavimo tabletės	not available	LT/1/05/0283/001	TILLOTTS PHARMA AB	LT
Asacol 400 mg modifikuoto atpalaidavimo tabletės	not available	LT/1/05/0283/002	TILLOTTS PHARMA AB	LT
Asacol 400 mg sýruþolnar töflur.	not available	970315	TILLOTTS PHARMA AB	IS
ASACOL 400 tablet, maagsapresistente tabletten 400 mg	not available	RVG 11737	PHARMAAND GMBH	NL
Asacol 400mg MR Tablets	not available	PL 41042/0053	ABBVIE LTD (UK)	XI
Asacol 400mg, comprimido gastrorresistente	not available	8676817	TILLOTTS PHARMA GMBH	PT
Asacol 400mg, comprimido gastrorresistente	not available	8676809	TILLOTTS PHARMA GMBH	PT
Asacol 4g SINGLE DOSE	not available	121389/08-12-2021	TILLOTTS PHARMA ITALY, S.R.L.,	GR
Asacol 4g SINGLE DOSE	not available	121389/08-12-2021	TILLOTTS PHARMA ITALY, S.R.L.,	GR

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Asacol 500 mg stílar	not available	970313 (IS)	TILLOTTS PHARMA AB	IS
Asacol 500 mg supozitoriji	not available	04-0434	TILLOTTS PHARMA AB	LV
Asacol 500 mg suppositorier	not available	15998	TILLOTTS PHARMA AB	DK
Asacol 500 mg suppositorier	not available	11562	TILLOTTS PHARMA AB	FI
Asacol 500 mg suppositorier	not available	11562	TILLOTTS PHARMA AB	FI
Asacol 500 mg suppositorier	not available	11562	TILLOTTS PHARMA AB	FI
Asacol 500 mg suppositorier	not available	11562	TILLOTTS PHARMA AB	FI
Asacol 500 mg suppositorier	not available	12093	TILLOTTS PHARMA AB	SE
Asacol 500 mg suppositorier	not available	12093	TILLOTTS PHARMA AB	SE
ASACOL 500 mg supposte	not available	026416127	GIULIANI SPA	IT
ASACOL 500 mg supposte	not available	026416139	GIULIANI SPA	IT
Asacol 500 mg, stikkpille	not available	94-1742	TILLOTTS PHARMA AB	NO
Asacol 500mg, supositório	not available	2137180	TILLOTTS PHARMA GMBH	PT
Asacol 800 mg	not available	123556/08-12-2021	TILLOTTS PHARMA ITALY, S.R.L.,	GR
Asacol 800 mg	not available	123556/08-12-2021	TILLOTTS PHARMA ITALY, S.R.L.,	GR

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
ASACOL 800 mg compresse gastroresistenti	not available	026416242	GIULIANI SPA	IT
ASACOL 800 mg compresse gastroresistenti	not available	026416317	GIULIANI SPA	IT
ASACOL 800 mg compresse gastroresistenti	not available	026416293	GIULIANI SPA	IT
Asacol 800 mg comprimido gastrorresistente.	not available	5179627	TILLOTTS PHARMA GMBH	PT
ASACOL 800 mg comprimidos gastrorresistentes	not available	78448	TILLOTTS PHARMA SPAIN S.A.	ES
Asacol 800 mg enterosolventní tablety	not available	29/091/08-C	TILLOTTS PHARMA GMBH	CZ
Asacol 800 mg enterotabletit	not available	16882	TILLOTTS PHARMA AB	FI
Asacol 800 mg enterotablett	not available	01-11015	TILLOTTS PHARMA AB	NO
Asacol 800 mg enterotabletter	not available	16882	TILLOTTS PHARMA AB	FI
Asacol 800 mg enterotabletter	not available	17894	TILLOTTS PHARMA AB	SE
Asacol 800 mg ilgstošās darbības tabletes	not available	05-0531	TILLOTTS PHARMA AB	LV
Asacol 800 mg magensaftresistente Tabletten	not available	89227.00.00	TILLOTTS PHARMA GMBH	DE
Asacol 800 mg modifikuoto atpalaidavimo tabletės	not available	LT/1/05/0283/004	TILLOTTS PHARMA AB	LT
Asacol 800 mg modifikuoto atpalaidavimo tabletės	not available	LT/1/05/0283/005	TILLOTTS PHARMA AB	LT
Asacol 800 mg sýruþolnar töflur	not available	IS/1/02/116/01	TILLOTTS PHARMA AB	IS

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
ASACOL 800 tablet, maagsapresistente tabletten 800 mg	not available	RVG 18636	PHARMAAND GMBH	NL
Asacol 800mg MR Tablets	not available	PL 41042/0054	ABBVIE LTD (UK)	XI
ASACOL čípky	not available	29/619/99-C	TILLOTTS PHARMA GMBH	CZ
ASACOL ENEMA 4 G rektální suspenze	not available	29/174/95-C	TILLOTTS PHARMA GMBH	CZ
Asacol, 400 mg toimeainet modifitseeritult vabastavad tabletid	not available	452604	TILLOTTS PHARMA AB	EE
Asacol, 500 mg rektaalsuposiidid	not available	452704	TILLOTTS PHARMA AB	EE
Asacol, 800 mg toimeainet modifitseeritult vabastavad tabletid	not available	501605	TILLOTTS PHARMA AB	EE
Asacol, enterotabletter	not available	33464	TILLOTTS PHARMA AB	DK
Asacol, enterotabletter	not available	13014	TILLOTTS PHARMA AB	DK
Asacol, rektalvæske, suspension, enkeltdosisbeholder	not available	18493	TILLOTTS PHARMA AB	DK
ASACOL® 800 mg gastrorezistentne tablete	not available	H/04/00214/002	TILLOTTS PHARMA GMBH	SI
ASACOL® 800 mg gastrorezistentne tablete	not available	H/04/00214/003	TILLOTTS PHARMA GMBH	SI
ASACOL® 400 mg gastrorezistentne tablete	not available	H/04/00214/001	TILLOTTS PHARMA GMBH	SI

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Asacolon 1000 mg Suppositories	SE/H/2039/001	PA2018/005/001	TILLOTTS PHARMA GMBH	IE
Asacolon 1000 mg Suppositories	SE/H/2039/001	PA2018/005/001	TILLOTTS PHARMA GMBH	IE
Asacolon 1600 mg modified-release tablets	SE/H/1654/001	PA 2018/004/001	TILLOTTS PHARMA GMBH	IE
Asacolon 400mg Gastro-resistant Tablets	not available	PA2018/001/001	TILLOTTS PHARMA GMBH	IE
Asacolon 400mg Gastro-resistant Tablets	not available	PA2018/001/001	TILLOTTS PHARMA GMBH	IE
Asacolon 500 mg Suppositories	not available	PA2018/001/003	TILLOTTS PHARMA GMBH	IE
Asacolon 800mg Gastro-resistant Tablets	not available	PA2018/001/002	TILLOTTS PHARMA GMBH	IE
ASALEX 1,5 g granulato per sospensione rettale	not available	027122112	CHIESI FARMACEUTICI S.P.A.	IT
ASALEX 2 g/60 ml sospensione rettale	not available	027122062	CHIESI FARMACEUTICI S.P.A.	IT
ASALEX 4 g/60 ml sospensione rettale	not available	027122098	CHIESI FARMACEUTICI S.P.A.	IT
ASALEX 400 mg compresse gastroresistenti a rilascio modificato	not available	027122100	CHIESI FARMACEUTICI S.P.A.	IT
ASALEX 800 mg compresse gastroresistenti a rilascio modificato	not available	027122124	CHIESI FARMACEUTICI S.P.A.	IT
Asamax 2 g/50 ml sospensione rettale	not available	034298036	CA.DI.GROUP S.P.A	IT
Asamax 250 250 mg tabletki dojelitowe	not available	10 840	ASTELLAS PHARMA SP. Z O.O.	PL

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Asamax 250, 250 mg, czopki	not available	10 838	ASTELLAS PHARMA SP. Z O.O.	PL
Asamax 4 g/100 ml sospensione rettale	not available	034298048	CA.DI.GROUP S.P.A	IT
Asamax 400 mg compresse gastroresistenti	not available	034298012	CA.DI.GROUP S.P.A	IT
Asamax 500 500 mg tabletki dojelitowe	not available	10 841	ASTELLAS PHARMA SP. Z O.O.	PL
Asamax 500 mg supposte	not available	034298051	CA.DI.GROUP S.P.A	IT
Asamax 500, 500 mg, czopki	not available	10 839	ASTELLAS PHARMA SP. Z O.O.	PL
Asamax 800 mg compresse gastroresistenti	not available	034298063	CA.DI.GROUP S.P.A	IT
Asamax 800 mg compresse gastroresistenti	not available	034298024	CA.DI.GROUP S.P.A	IT
Asamovon 1600 mg comprimés à libération modifiée	SE/H/1654/001	BE535777	TILLOTTS PHARMA GMBH	BE
Asamovon 1600 mg comprimés à libération modifiée	SE/H/1654/001	BE535777	TILLOTTS PHARMA GMBH	BE
Asamovon 1600 mg comprimés à libération modifiée	SE/H/1654/001	BE535777	TILLOTTS PHARMA GMBH	BE
Asamovon 1600 mg tabletten met gereguleerde afgifte	SE/H/1654/001	BE535777	TILLOTTS PHARMA GMBH	BE
Asamovon 1600 mg tabletten met gereguleerde afgifte	SE/H/1654/001	BE535777	TILLOTTS PHARMA GMBH	BE
Asamovon 1600 mg tabletten met gereguleerde	SE/H/1654/001	BE535777	TILLOTTS PHARMA GMBH	BE

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
afgifte				
Asamovon 1600 mg Tabletten mit veränderter Wirkstofffreisetzung	SE/H/1654/001	BE535777	TILLOTTS PHARMA GMBH	BE
Asamovon 1600 mg Tabletten mit veränderter Wirkstofffreisetzung	SE/H/1654/001	BE535777	TILLOTTS PHARMA GMBH	BE
Asamovon 1600 mg Tabletten mit veränderter Wirkstofffreisetzung	SE/H/1654/001	BE535777	TILLOTTS PHARMA GMBH	BE
ASAVIXIN 500 mg compresse rivestite	not available	035356017	VIATRIS ITALIA S.R.L.	IT
Azzavix 1 g comprimido gastrorresistente	not available	5833645	FAES FARMA PORTUGAL, S.A.	PT
Azzavix 1 g comprimido gastrorresistente	not available	5721337	FAES FARMA PORTUGAL, S.A.	PT
Azzavix 1 g comprimido gastrorresistente	not available	5721311	FAES FARMA PORTUGAL, S.A.	PT
Azzavix 1 g comprimido gastrorresistente	not available	5721329	FAES FARMA PORTUGAL, S.A.	PT
Azzavix 1 g comprimido gastrorresistente	not available	5774005	FAES FARMA PORTUGAL, S.A.	PT
Azzavix 1000 mg comprimidos gastrorresistentes	ES/H/0587/002	85724	FAES FARMA, S.A.	ES
Azzavix 1000 mg enterosolventní tablety	ES/H/0588/002	29/446/21-C	FAES FARMA, S.A.	CZ
Azzavix 1000 mg gastrorezistentné tablety	ES/H/0588/002	29/0125/22-S	FAES FARMA, S.A.	SK

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Azzavix 1000 mg maagsapresistente tabletten	ES/H/0856/002	RVG 130146	FAES FARMA, S.A.	NL
Azzavix 1000 mg maagsapresistente tabletten	ES/H/0856/002	RVG 130146	FAES FARMA, S.A.	NL
Azzavix 1000 mg magensaftresistente Tabletten	ES/H/0587/002	140412	ASTRO-PHARMA GMBH	AT
Azzavix 1000 mg supositórios	not available	5833652	FAES FARMA PORTUGAL, S.A.	PT
Azzavix 1000 mg supositórios	not available	5758479	FAES FARMA PORTUGAL, S.A.	PT
Azzavix 1000 mg supositórios	not available	5758453	FAES FARMA PORTUGAL, S.A.	PT
Azzavix 1000 mg supositórios	not available	5758461	FAES FARMA PORTUGAL, S.A.	PT
Azzavix 1000 mg Zäpfchen	PT/H/2281/002	140243	FAES FARMA, S.A.	AT
Azzavix 500 mg čípky	ES/H/0588/003	29/312/21-C	FAES FARMA, S.A.	CZ
Azzavix 500 mg comprimidos gastrorresistentes	ES/H/0587/001	84676	FAES FARMA, S.A.	ES
Azzavix 500 mg comprimidos gastrorresistentes	not available	5650452	FAES FARMA PORTUGAL, S.A.	PT
Azzavix 500 mg comprimidos gastrorresistentes	not available	5650445	FAES FARMA PORTUGAL, S.A.	PT
Azzavix 500 mg comprimidos gastrorresistentes	not available	5708243	FAES FARMA PORTUGAL, S.A.	PT
Azzavix 500 mg gastrorezistentné tablety	ES/H/0588/001	29/0205/20-S	FAES FARMA, S.A.	SK
Azzavix 500 mg maagsapresistente tabletten	ES/H/0856/001	RVG 130143	FAES FARMA, S.A.	NL

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Azzavix 500 mg maagsapresistente tabletten	ES/H/0856/001	RVG 130143	FAES FARMA, S.A.	NL
Azzavix 500 mg magensaftresistente Tabletten	ES/H/0588/001	139270	ASTRO-PHARMA GMBH	AT
Azzavix 500 mg supositórios	not available	5713425	FAES FARMA PORTUGAL, S.A.	PT
Azzavix 500 mg supositórios	not available	5705157	FAES FARMA PORTUGAL, S.A.	PT
Azzavix 500 mg supositórios	not available	5705165	FAES FARMA PORTUGAL, S.A.	PT
Azzavix 500 mg supozitoare	ES/H/0588/003	1472/2022/01	FAES FARMA, S.A.	RO
Azzavix 500 mg supozitoare	ES/H/0588/003	1472/2022/02	FAES FARMA, S.A.	RO
Azzavix 500 mg supozitoare	ES/H/0588/003	1472/2022/03	FAES FARMA, S.A.	RO
Azzavix 500 mg supozitoare	ES/H/0588/003	1472/2022/04	FAES FARMA, S.A.	RO
Azzavix 500 mg supozitoare	ES/H/0588/003	1472/2022/05	FAES FARMA, S.A.	RO
Azzavix 500 mg Zäpfchen	PT/H/2281/001	140244	FAES FARMA, S.A.	AT
Belenix 1000 mg supositórios	PT/H/2838/002/DC	NOT APPLICABLE	FAES FARMA PORTUGAL, S.A.	PT
Belenix 1000 mg supositórios	PT/H/2838/002/DC	NOT APPLICABLE	FAES FARMA PORTUGAL, S.A.	PT
Belenix 1000 mg supositórios	PT/H/2838/002/DC	NOT APPLICABLE	FAES FARMA PORTUGAL, S.A.	PT
Belenix 500 mg supositórios	PT/H/2838/001/DC	NOT APPLICABLE	FAES FARMA PORTUGAL, S.A.	PT

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Belenix 500 mg supositórios	PT/H/2838/001/DC	NOT APPLICABLE	FAES FARMA PORTUGAL, S.A.	PT
Blenix 1000 mg comprimidos gastrorresistentes	ES/H/0856/002	89105	FAES FARMA, S.A.	ES
Blenix 1000 mg comprimidos gastrorresistentes	ES/H/0856/002	89105	FAES FARMA, S.A.	ES
Blenix 500 mg comprimidos gastrorresistentes	ES/H/0856/001	89106	FAES FARMA, S.A.	ES
Blenix 500 mg comprimidos gastrorresistentes	ES/H/0856/001	89106	FAES FARMA, S.A.	ES
Claversal 1 g comprimidos gastrorresistentes	not available	83.162	FAES FARMA, S.A.	ES
Claversal 1000 mg Zäpfchen	PT/H/2281/002	2203439.00.00	FAES FARMA, S.A.	DE
Claversal 1g espuma rectal.	not available	61.335	FAES FARMA, S.A.	ES
Claversal 250 mg Zäpfchen	not available	1-18370	MERCK GESELLSCHAFT MBH	AT
Claversal 500 mg - Filmtabletten	not available	1-18369	MERCK GESELLSCHAFT MBH	AT
CLAVERSAL 500 mg compresse rivestite	not available	027308016	TEOFARMA S.R.L.	IT
CLAVERSAL 500 mg comprimés gastro-résistants	not available	BE213534	TRUVION HEALTHCARE N.V	BE
CLAVERSAL 500 mg comprimés gastro-résistants	not available	2001040038	TRUVION HEALTHCARE N.V	LU
Claversal 500 mg comprimidos gastrorresistentes	not available	58.101	FAES FARMA, S.A.	ES
Claversal 500 mg Filmtabletten	not available	1-18369	MERCK GESELLSCHAFT MBH	AT

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
CLAVERSAL 500 mg maagsapresistente tabletten	not available	BE213534	TRUVION HEALTHCARE N.V	BE
CLAVERSAL 500 mg magensaftresistente Tabletten	not available	BE213534	TRUVION HEALTHCARE N.V	BE
Claversal 500 mg supositorios	not available	60.607	FAES FARMA, S.A.	ES
CLAVERSAL 500 mg Suppositoires	not available	BE144882	TRUVION HEALTHCARE N.V	BE
CLAVERSAL 500 mg suppositoires	not available	2000065845	TRUVION HEALTHCARE N.V	LU
Claversal 500 mg Zäpfchen	not available	1-18371	MERCK GESELLSCHAFT MBH	AT
CLAVERSAL 500 mg Zäpfchen	not available	BE144882	TRUVION HEALTHCARE N.V	BE
CLAVERSAL 500 mg zetpillen	not available	BE144882	TRUVION HEALTHCARE N.V	BE
CLAVERSAL FOAM 1 g pro Dosis, Rektalschaum	not available	BE169592	TRUVION HEALTHCARE N.V	BE
CLAVERSAL Foam 1g/dose, mousse rectale	not available	BE169592	TRUVION HEALTHCARE N.V	BE
CLAVERSAL Foam 1g/dose, mousse rectale	not available	1995070317	TRUVION HEALTHCARE N.V	LU
CLAVERSAL Foam 1g/dosis, schuim voor rectaal gebruik	not available	BE169592	TRUVION HEALTHCARE N.V	BE
Claversal® 250 mg Zäpfchen	not available	5697.00.00	RECORDATI PHARMA GMBH	DE
Claversal® 4 g/60 g Klysmen, Rektalsuspension	not available	40157.00.00	RECORDATI PHARMA GMBH	DE
Claversal® 500 mg Tabletten	not available	5697.01.01	RECORDATI PHARMA GMBH	DE

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Claversal® 500 mg Tabletten	not available	5697.01.01	RECORDATI PHARMA GMBH	DE
Claversal® 500 mg Zäpfchen	not available	5697.01.00	RECORDATI PHARMA GMBH	DE
Claversal® 500 mg Zäpfchen	not available	5697.01.00	RECORDATI PHARMA GMBH	DE
Claversal® Micropellets 1,5 g, magensaftresistentes Granulat	not available	42197.00.01	RECORDATI PHARMA GMBH	DE
Claversal® Rektalschaum 1 g	not available	42197.00.00	RECORDATI PHARMA GMBH	DE
Cletrovaproct 1 g supposte	DE/H/5682/001	050523012	DR. FALK PHARMA G.M.B.H.	IT
Cletrovaproct 1 g supposte	DE/H/5682/001	050523024	DR. FALK PHARMA G.M.B.H.	IT
Cletrovaproct 1 g supposte	DE/H/5682/001	050523036	DR. FALK PHARMA G.M.B.H.	IT
Cletrovaproct 1 g supposte	DE/H/5682/001	050523048	DR. FALK PHARMA G.M.B.H.	IT
Cletrovaproct 1 g supposte	DE/H/5682/001	050523051	DR. FALK PHARMA G.M.B.H.	IT
Cletrovaproct 1 g supposte	DE/H/5682/001	050523063	DR. FALK PHARMA G.M.B.H.	IT
Cletrovaproct 1 g supposte	DE/H/5682/001	050523075	DR. FALK PHARMA G.M.B.H.	IT
Colitofalk 1 g suppositoires	DE/H/5682/001	BE371612	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1 g suppositoires	DE/H/5682/001	BE371612	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1 g suppositoires	DE/H/5682/001	BE371612	DR. FALK PHARMA G.M.B.H.	BE

Product full name (in authorisation country)	**MRP/DCP or CP Authorisation number**	**National Authorisation Number**	**MAH of product in the member state**	**Member State where product is authorised**
Colitofalk 1 g suppositoires	DE/H/5682/001	BE371612	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1 g suppositoires	DE/H/5682/001	BE371612	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1 g suppositoires	DE/H/5682/001	BE371612	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1 g suppositoires	DE/H/5682/001	BE371612	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1 g suppositories	DE/H/5682/001	2011040012	DR. FALK PHARMA G.M.B.H.	LU
Colitofalk 1 g Zäpfchen	DE/H/5682/001	BE371612	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1 g Zäpfchen	DE/H/5682/001	BE371612	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1 g Zäpfchen	DE/H/5682/001	BE371612	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1 g Zäpfchen	DE/H/5682/001	BE371612	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1 g zetpillen	DE/H/5682/001	BE371612	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1 g zetpillen	DE/H/5682/001	BE371612	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1 g zetpillen	DE/H/5682/001	BE371612	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1 g zetpillen	DE/H/5682/001	BE371612	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1 g zetpillen	DE/H/5682/001	BE371612	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1 g zetpillen	DE/H/5682/001	BE371612	DR. FALK PHARMA G.M.B.H.	BE

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Colitofalk 1 g zetpillen	DE/H/5682/001	BE371612	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1,5 g granulaat met verlengde afgifte	DE/H/0363/004	BE347155	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1,5 g granulaat met verlengde afgifte	DE/H/0363/004	BE347155	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1,5 g granulaat met verlengde afgifte	DE/H/0363/004	BE347155	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1,5 g granulaat met verlengde afgifte	DE/H/0363/004	BE347155	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1,5 g granulaat met verlengde afgifte	DE/H/0363/004	BE347155	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1,5 g granulaat met verlengde afgifte	DE/H/0363/004	BE347155	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1,5 g granulaat met verlengde afgifte	DE/H/0363/004	BE347155	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1,5 g granulaat met verlengde afgifte	DE/H/0363/004	BE347155	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1,5 g granulaat met verlengde afgifte	DE/H/0363/004	BE347155	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1,5 g granulaat met verlengde afgifte	DE/H/0363/004	BE347155	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1,5 g granulés á libération prolongée	DE/H/0363/004	2010020010	DR. FALK PHARMA G.M.B.H.	LU
Colitofalk 1,5 g granulés à libération prolongée	DE/H/0363/004	BE347155	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1,5 g granulés à libération prolongée	DE/H/0363/004	BE347155	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1,5 g granulés à libération prolongée	DE/H/0363/004	BE347155	DR. FALK PHARMA G.M.B.H.	BE

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Colitofalk 1,5 g granulés à libération prolongée	DE/H/0363/004	BE347155	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1,5 g granulés à libération prolongée	DE/H/0363/004	BE347155	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1,5 g granulés à libération prolongée	DE/H/0363/004	BE347155	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1,5 g granulés à libération prolongée	DE/H/0363/004	BE347155	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1,5 g granulés à libération prolongée	DE/H/0363/004	BE347155	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1,5 g granulés à libération prolongée	DE/H/0363/004	BE347155	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1,5 g granulés à libération prolongée	DE/H/0363/004	BE347155	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1,5 g Retardgranulat	DE/H/0363/004	BE347155	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1,5 g Retardgranulat	DE/H/0363/004	BE347155	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1,5 g Retardgranulat	DE/H/0363/004	BE347155	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1,5 g Retardgranulat	DE/H/0363/004	BE347155	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1,5 g Retardgranulat	DE/H/0363/004	BE347155	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1,5 g Retardgranulat	DE/H/0363/004	BE347155	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1,5 g Retardgranulat	DE/H/0363/004	BE347155	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1,5 g Retardgranulat	DE/H/0363/004	BE347155	DR. FALK PHARMA G.M.B.H.	BE

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Colitofalk 1,5 g Retardgranulat	DE/H/0363/004	BE347155	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1,5 g Retardgranulat	DE/H/0363/004	BE347155	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1000 mg granulaat met verlengde afgifte	DE/H/0363/002	BE243494	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1000 mg granulaat met verlengde afgifte	DE/H/0363/002	BE243494	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1000 mg granulaat met verlengde afgifte	DE/H/0363/002	BE243494	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1000 mg granulaat met verlengde afgifte	DE/H/0363/002	BE243494	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1000 mg granulaat met verlengde afgifte	DE/H/0363/002	BE243494	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1000 mg granulés à libération prolongée	DE/H/0363/002	BE243494	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1000 mg granulés à libération prolongée	DE/H/0363/002	BE243494	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1000 mg granulés à libération prolongée	DE/H/0363/002	BE243494	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1000 mg granulés à libération prolongée	DE/H/0363/002	BE243494	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1000 mg granulés à libération prolongée	DE/H/0363/002	BE243494	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1000 mg granulés à libération prolongée	DE/H/0363/002	2003040027	DR. FALK PHARMA G.M.B.H.	LU
Colitofalk 1000 mg Retardgranulat	DE/H/0363/002	BE243494	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1000 mg Retardgranulat	DE/H/0363/002	BE243494	DR. FALK PHARMA G.M.B.H.	BE

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Colitofalk 1000 mg Retardgranulat	DE/H/0363/002	BE243494	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1000 mg Retardgranulat	DE/H/0363/002	BE243494	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1000 mg Retardgranulat	DE/H/0363/002	BE243494	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1g Zäpfchen	DE/H/5682/001	BE371612	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1g Zäpfchen	DE/H/5682/001	BE371612	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 1g Zäpfchen	DE/H/5682/001	BE371612	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 2 g Rektalsuspension	not available	BE157272	DR. FALK PHARMA BENELUX B.V.	BE
Colitofalk 2 g suspension rectale	not available	BE157272	DR. FALK PHARMA BENELUX B.V.	BE
Colitofalk 2 g suspension rectale	not available	2011051156	DR. FALK PHARMA BENELUX B.V.	LU
Colitofalk 2g suspensie voor rectaal gebruik	not available	BE157272	DR. FALK PHARMA BENELUX B.V.	BE
Colitofalk 3 g granulaat met verlengde afgifte	DE/H/0363/007	BE398027	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 3 g granulaat met verlengde afgifte	DE/H/0363/007	BE398027	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 3 g granulaat met verlengde afgifte	DE/H/0363/007	BE398027	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 3 g granulaat met verlengde afgifte	DE/H/0363/007	BE398027	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 3 g granulaat met verlengde afgifte	DE/H/0363/007	BE398027	DR. FALK PHARMA G.M.B.H.	BE

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Colitofalk 3 g granulaat met verlengde afgifte	DE/H/0363/007	BE398027	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 3 g granulaat met verlengde afgifte	DE/H/0363/007	BE398027	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 3 g granulaat met verlengde afgifte	DE/H/0363/007	BE398027	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 3 g granulés à libération prolongée	DE/H/0363/007	BE398027	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 3 g granulés à libération prolongée	DE/H/0363/007	BE398027	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 3 g granulés à libération prolongée	DE/H/0363/007	BE398027	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 3 g granulés à libération prolongée	DE/H/0363/007	BE398027	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 3 g granulés à libération prolongée	DE/H/0363/007	BE398027	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 3 g granulés à libération prolongée	DE/H/0363/007	BE398027	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 3 g granulés à libération prolongée	DE/H/0363/007	BE398027	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 3 g granulés à libération prolongée	DE/H/0363/007	BE398027	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 3 g granulés à libération prolongée	DE/H/0363/007	2012030024	DR. FALK PHARMA G.M.B.H.	LU
Colitofalk 3 g Retardgranulat	DE/H/0363/007	BE398027	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 3 g Retardgranulat	DE/H/0363/007	BE398027	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 3 g Retardgranulat	DE/H/0363/007	BE398027	DR. FALK PHARMA G.M.B.H.	BE

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Colitofalk 3 g Retardgranulat	DE/H/0363/007	BE398027	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 3 g Retardgranulat	DE/H/0363/007	BE398027	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 3 g Retardgranulat	DE/H/0363/007	BE398027	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 3 g Retardgranulat	DE/H/0363/007	BE398027	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 3 g Retardgranulat	DE/H/0363/007	BE398027	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 4 g Rektalsuspension	not available	BE145275	DR. FALK PHARMA BENELUX B.V.	BE
Colitofalk 4 g Rektalsuspension	not available	BE145275	DR. FALK PHARMA BENELUX B.V.	BE
Colitofalk 4 g suspensie voor rectaal gebruik	not available	BE145275	DR. FALK PHARMA BENELUX B.V.	BE
Colitofalk 4 g suspensie voor rectaal gebruik	not available	BE145275	DR. FALK PHARMA BENELUX B.V.	BE
Colitofalk 4 g suspension rectale	not available	BE145275	DR. FALK PHARMA BENELUX B.V.	BE
Colitofalk 4 g suspension rectale	not available	BE145275	DR. FALK PHARMA BENELUX B.V.	BE
Colitofalk 4 g suspension rectale	not available	2011051157	DR. FALK PHARMA BENELUX B.V.	LU
Colitofalk 500 mg granulaat met verlengde afgifte	DE/H/0363/001	BE243485	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 500 mg granulaat met verlengde afgifte	DE/H/0363/001	BE243485	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 500 mg granulaat met verlengde afgifte	DE/H/0363/001	BE243485	DR. FALK PHARMA G.M.B.H.	BE

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Colitofalk 500 mg granulés à libération prolongée	DE/H/0363/001	BE243485	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 500 mg granulés à libération prolongée	DE/H/0363/001	BE243485	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 500 mg granulés à libération prolongée	DE/H/0363/001	BE243485	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 500 mg granulés à libération prolongée	DE/H/0363/001	2003040026	DR. FALK PHARMA G.M.B.H.	LU
Colitofalk 500 mg Retardgranulat	DE/H/0363/001	BE243485	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 500 mg Retardgranulat	DE/H/0363/001	BE243485	DR. FALK PHARMA G.M.B.H.	BE
Colitofalk 500 mg Retardgranulat	DE/H/0363/001	BE243485	DR. FALK PHARMA G.M.B.H.	BE
Cosinix 500 mg gyomornedv-ellenálló tabletta	ES/H/0588/001	OGYI-T-23771/01	FAES FARMA S.A.	HU
CROHNAX 250 mg czopki	not available	10740	FARMINA SP. Z O.O.	PL
CROHNAX, 1000 mg, czopki	not available	22519	FARMINA SP. Z O.O.	PL
CROHNAX, 1000 mg, czopki	not available	22519	FARMINA SP. Z O.O.	PL
CROHNAX, 1000 mg, czopki	not available	22519	FARMINA SP. Z O.O.	PL
CROHNAX, 1000 mg, czopki	not available	22519	FARMINA SP. Z O.O.	PL
CROHNAX, 500 mg, czopki	not available	22518	FARMINA SP. Z O.O.	PL
Falk-Asa 500mg magensaftresistente Tablette	DE/H/0363/006	53157.01.00	DR. FALK PHARMA G.M.B.H.	DE

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Falk-Asa 500mg Suppositorien	not available	53157.01.01	DR. FALK PHARMA G.M.B.H.	DE
FIVASA 1 g, suppositoire	SE/H/2039/001	34009 302 222 0 1	TILLOTTS PHARMA GMBH	FR
FIVASA 1 g, suppositoire	SE/H/2039/001	34009 302 222 1 8	TILLOTTS PHARMA GMBH	FR
FIVASA 1 g, suppositoire	SE/H/2039/001	34009 302 222 2 5	TILLOTTS PHARMA GMBH	FR
FIVASA 1 g, suppositoire	SE/H/2039/001	34009 550 795 7 6	TILLOTTS PHARMA GMBH	FR
FIVASA 1 g, suppositoire	SE/H/2039/001	34009 550 795 8 3	TILLOTTS PHARMA GMBH	FR
FIVASA 1 g, suppositoire	SE/H/2039/001	34009 302 222 0 1	TILLOTTS PHARMA GMBH	FR
FIVASA 1 g, suppositoire	SE/H/2039/001	34009 302 222 1 8	TILLOTTS PHARMA GMBH	FR
FIVASA 1 g, suppositoire	SE/H/2039/001	34009 302 222 2 5	TILLOTTS PHARMA GMBH	FR
FIVASA 1 g, suppositoire	SE/H/2039/001	34009 550 795 7 6	TILLOTTS PHARMA GMBH	FR
FIVASA 1 g, suppositoire	SE/H/2039/001	34009 550 795 8 3	TILLOTTS PHARMA GMBH	FR
FIVASA 1600 mg, comprimé gastro-résistant	not available	34009 301 998 6 2	TILLOTTS PHARMA GMBH	FR
FIVASA 1600 mg, comprimé gastro-résistant	not available	34009 301 998 7 9	TILLOTTS PHARMA GMBH	FR
FIVASA 1600 mg, comprimé gastro-résistant	not available	34009 301 998 8 6	TILLOTTS PHARMA GMBH	FR
FIVASA 400 mg, comprimé enrobé gastrorésistant	not available	34009 347 861 8 1	TILLOTTS PHARMA GMBH	FR

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
FIVASA 500 mg, suppositoire	not available	34009 356 683 1 8	TILLOTTS PHARMA GMBH	FR
FIVASA 500 mg, suppositoire	not available	34009 358 811 7 5	TILLOTTS PHARMA GMBH	FR
FIVASA 500 mg, suppositoire	not available	34009 358 810 0 7	TILLOTTS PHARMA GMBH	FR
FIVASA 800 mg, comprimé enrobé gastrorésistant	not available	34009 358 567 9 1	TILLOTTS PHARMA GMBH	FR
FIVASA 800 mg, comprimé enrobé gastrorésistant	not available	34009 358 568 5 2	TILLOTTS PHARMA GMBH	FR
Galtasa 1000 mg gastro-resistant tablets	ES/H/0588/002	PA 0864/001/003	FAES FARMA, S.A.	IE
Galtasa 500 mg gastro-resistant tablets.	ES/H/0588/001	PA 0864/001/001	FAES FARMA, S.A.	IE
Galtasa 500 mg suppositories	ES/H/0588/003	PA 0864/001/002	FAES FARMA, S.A.	IE
Kiudro 1 g zetpil	SE/H/2039/001	RVG 126165	TILLOTTS PHARMA GMBH	NL
Kiudro 1 g zetpil	SE/H/2039/001	RVG 126165	TILLOTTS PHARMA GMBH	NL
Kiudro 1 g Υπόθετα	SE/H/2039/001	327720101	TILLOTTS PHARMA GMBH	GR
Kiudro 1 g Υπόθετα	SE/H/2039/001	327720102	TILLOTTS PHARMA GMBH	GR
Kiudro 1 g Υπόθετα	SE/H/2039/001	327720103	TILLOTTS PHARMA GMBH	GR
Kiudro 1 g Υπόθετα	SE/H/2039/001	327720104	TILLOTTS PHARMA GMBH	GR
Kiudro 1 g Υπόθετα	SE/H/2039/001	327720105	TILLOTTS PHARMA GMBH	GR

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Kiudro 1 g Υπόθετα	SE/H/2039/001	327720101	TILLOTTS PHARMA GMBH	GR
Kiudro 1 g Υπόθετα	SE/H/2039/001	327720102	TILLOTTS PHARMA GMBH	GR
Kiudro 1 g Υπόθετα	SE/H/2039/001	327720103	TILLOTTS PHARMA GMBH	GR
Kiudro 1 g Υπόθετα	SE/H/2039/001	327720104	TILLOTTS PHARMA GMBH	GR
Kiudro 1 g Υπόθετα	SE/H/2039/001	327720105	TILLOTTS PHARMA GMBH	GR
Lextrasa 400mg Enteric-Coated Tablets Mesalazine 400mg Enteric-Coated Tablets	not available	PL 48053/0020	ALFASIGMA S.P.A.	XI
MAZALVIX 500 mg, comprimé gastro-résistant	ES/H/0856/001	3400930274620	LABORATOIRES MAYOLY SPINDLER	FR
MAZALVIX 500 mg, comprimé gastro-résistant	ES/H/0856/001	3400930274620	LABORATOIRES MAYOLY SPINDLER	FR
MAZALVIX 1000 mg, comprimé gastro-résistant	ES/H/0856/002/DC	34009 302 747 6 7	LABORATOIRES MAYOLY SPINDLER	FR
MAZALVIX 1000 mg, comprimé gastro-résistant	ES/H/0856/002/DC	34009 302 747 7 4	LABORATOIRES MAYOLY SPINDLER	FR
MAZALVIX 1000 mg, comprimé gastro-résistant	ES/H/0856/002/DC	34009 302 747 6 7	LABORATOIRES MAYOLY SPINDLER	FR
MAZALVIX 1000 mg, comprimé gastro-résistant	ES/H/0856/002/DC	34009 302 747 7 4	LABORATOIRES MAYOLY	FR

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
			SPINDLER	
MAZALVIX 1000 mg, suppositoire	PT/H/2838/002	34009 302 927 0 9	LABORATOIRES MAYOLY SPINDLER	FR
MAZALVIX 1000 mg, suppositoire	PT/H/2838/002	34009 302 927 1 6	LABORATOIRES MAYOLY SPINDLER	FR
MAZALVIX 1000 mg, suppositoire	PT/H/2838/002	34009 302 927 2 3	LABORATOIRES MAYOLY SPINDLER	FR
MAZALVIX 500 mg, suppositoire	PT/H/2838/001	34009 302 926 8 6	LABORATOIRES MAYOLY SPINDLER	FR
MAZALVIX 500 mg, suppositoire	PT/H/2838/001	34009 302 926 9 3	LABORATOIRES MAYOLY SPINDLER	FR
Mecolvix 1000 mg comprimidos gastrorresistentes	ES/H/0588/002	85738	FAES FARMA, S.A.	ES
Mecolvix 1000 mg supositórios	PT/H/2281/002/DC	PT/H/2281/002/DC	LABORATÓRIOS VITÓRIA, SA	PT
Mecolvix 1000 mg supositórios	PT/H/2281/002/DC	PT/H/2281/002/DC	LABORATÓRIOS VITÓRIA, SA	PT
Mecolvix 1000 mg supositórios	PT/H/2281/002/DC	PT/H/2281/002/DC	LABORATÓRIOS VITÓRIA, SA	PT
Mecolvix 1000 mg supositórios	PT/H/2281/002/DC	PT/H/2281/002/DC	LABORATÓRIOS VITÓRIA, SA	PT
Mecolvix 500 mg comprimidos gastrorresistentes	ES/H/0588/001	84675	FAES FARMA, S.A.	ES

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Mecolvix 500 mg supositorios	ES/H/0588/003	88174	FAES FARMA, S.A.	ES
Mecolvix 500 mg supositórios	PT/H/2281/001/DC	PT/H/2281/001/DC	LABORATÓRIOS VITÓRIA, SA	PT
Mecolvix 500 mg supositórios	PT/H/2281/001/DC	PT/H/2281/001/DC	LABORATÓRIOS VITÓRIA, SA	PT
Mecolvix 500 mg supositórios	PT/H/2281/001/DC	PT/H/2281/001/DC	LABORATÓRIOS VITÓRIA, SA	PT
Mecolvix 500 mg supositórios	PT/H/2281/001/DC	PT/H/2281/001/DC	LABORATÓRIOS VITÓRIA, SA	PT
Mecolzin 1000 mg γαστροανθεκτικά δισκία.	ES/H/0588/002	61304/09-06-2022	INNOVIS PHARMA	GR
Mecolzin 500 mg γαστροανθεκτικά δισκία.	ES/H/0588/001	61303/09-06-22	INNOVIS PHARMA	GR
Mecolzine 1000 mg gastro-resistant tablets.	ES/H/0587/002	MA1278/00102	FAES FARMA, S.A.	MT
Mecolzine 1000 mg γαστροανθεκτικά δισκία.	ES/H/0587/002	023468	FAES FARMA, S.A.	CY
Mecolzine 1000 mg υπόθετο	PT/H/2280/002	023298	FAES FARMA, S.A.	CY
Mecolzine 1g suppositories	PT/H/2280/002	MA1278/00302	FAES FARMA, S.A.	MT
Mecolzine 500 mg gastro-resistant tablets.	ES/H/0587/001	MA1278/00101	FAES FARMA, S.A.	MT
Mecolzine 500 mg Suppositories	PT/H/2280/001	MA1278/00301	FAES FARMA S.A.	MT
Mecolzine 500 mg γαστροανθεκτικά δισκία.	ES/H/0587/001	023332	FAES FARMA, S.A.	CY
Mecolzine 500mg υπόθετο	PT/H/2280/001	023297	FAES FARMA, S.A.	CY

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Mesagran 1000 mg Retardgranulat	DE/H/0363/002	1-24830	DR. FALK PHARMA G.M.B.H.	AT
Mesagran 1500 mg Retardgranulat	DE/H/0363/004	1-27730	DR. FALK PHARMA G.M.B.H.	AT
Mesagran 3000 mg Retardgranulat	DE/H/0363/007	1-30482	DR. FALK PHARMA G.M.B.H.	AT
Mesagran 500 mg Retardgranulat	DE/H/0363/001	1-24829	DR. FALK PHARMA G.M.B.H.	AT
Mesalazin "Orion", enterotabletter 1000 mg	ES/H/0587/002	63563	FAES FARMA, S.A.	DK
Mesalazin "Orion", enterotabletter 500 mg	ES/H/0587/001	61716	FAES FARMA S.A.	DK
Mesalazin "Orion", suppositorier 1 g	PT/H/2280/002	62018	FAES FARMA, S.A.	DK
Mesalazin Orion 1 000 mg enterotabletit	ES/H/0587/002	37572	FAES FARMA, S.A.	FI
Mesalazin Orion 1 000 mg enterotabletter	ES/H/0587/002	37572	FAES FARMA, S.A.	FI
Mesalazin Orion 1 000 mg enterotabletter	ES/H/0587/002	59937	FAES FARMA, S.A.	SE
Mesalazin Orion 1 g peräpuikot	PT/H/2280/002	36636	FAES FARMA, S.A.	FI
Mesalazin Orion 1 g stikkpiller	PT/H/2280/002	18-12604	FAES FARMA, S.A.	NO
Mesalazin Orion 1 g suppositorier	PT/H/2280/002	36636	FAES FARMA, S.A.	FI
Mesalazin Orion 1 g suppositorier	PT/H/2280/002	58602	FAES FARMA, S.A.	SE
Mesalazin Orion 1000 mg enterotabletter	ES/H/0587/002	19-13142	FAES FARMA, S.A.	NO

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Mesalazin Orion 500 mg enterotabletit	ES/H/0587/001	36387	FAES FARMA, S.A.	FI
Mesalazin Orion 500 mg enterotabletter	ES/H/0587/001	36387	FAES FARMA, S.A.	FI
Mesalazin Orion 500 mg enterotabletter	ES/H/0587/001	18-12463	FAES FARMA, S.A.	NO
Mesalazin Orion 500 mg enterotabletter	ES/H/0587/001	58289	FAES FARMA, S.A.	SE
Mesalazin Orion 500 mg peräpuikot	PT/H/2280/001	36635	FAES FARMA, S.A.	FI
Mesalazin Orion 500 mg stikkpiller	PT/H/2280/001	18-12603	FAES FARMA, S.A.	NO
Mesalazin Orion 500 mg suppositorier	PT/H/2280/001	36635	FAES FARMA, S.A.	FI
Mesalazin Orion 500 mg suppositorier	PT/H/2280/001	58601	FAES FARMA, S.A.	SE
Mesalazin ''Orion'', suppositorier 500 mg	PT/H/2280/001	62017	FAES FARMA, S.A.	DK
MESALAZINA ALFASIGMA 400 mg comprimate gastrorezistente	not available	12452/2019/02	ALFASIGMA S.P.A.	RO
MESALAZINA ALFASIGMA 400 mg comprimate gastrorezistente	not available	12452/2019/01	ALFASIGMA S.P.A.	RO
MESALAZINA ALFASIGMA 800 mg comprimate gastrorezistente	not available	12453/2019/02	ALFASIGMA S.P.A.	RO
MESALAZINA ALFASIGMA 800 mg comprimate gastrorezistente	not available	12453/2019/03	ALFASIGMA S.P.A.	RO

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
MESALAZINA ALFASIGMA 800 mg comprimate gastrorezistente	not available	12453/2019/01	ALFASIGMA S.P.A.	RO
MESALAZINA DOC 2 g/50 ml sospensione rettale	not available	035386046	DOC GENERICI S.R.L.	IT
MESALAZINA DOC 4 g/100 ml sospensione rettale	not available	035386034	DOC GENERICI S.R.L.	IT
MESALAZINA DOC 400 mg compresse gastroresistenti	not available	035386073	DOC GENERICI S.R.L.	IT
MESALAZINA DOC 400 mg compresse gastroresistenti	not available	035386010	DOC GENERICI S.R.L.	IT
MESALAZINA DOC 500 mg supposte	not available	035386059	DOC GENERICI S.R.L.	IT
MESALAZINA DOC 800 mg compresse gastroresistenti	not available	035386022	DOC GENERICI S.R.L.	IT
MESALAZINA DOC 800 mg compresse gastroresistenti	not available	035386061	DOC GENERICI S.R.L.	IT
MESALAZINA DOROM 400 mg compresse gastroresistenti	not available	034462061	TEVA ITALIA S.R.L.	IT
Mesalazina Dorom 400 mg compresse gastroresistenti	not available	034462010	TEVA ITALIA S.R.L.	IT
MESALAZINA DOROM 800 mg compresse gastroresistenti	not available	034462073	TEVA ITALIA S.R.L.	IT

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
MESALAZINA DOROM 800 mg compresse gastroresistenti	not available	034462022	TEVA ITALIA S.R.L.	IT
MESALAZINA EG 2 g/60 ml gel rettale	not available	029480023	EG S.P.A.	IT
MESALAZINA EG 4 g/60 ml gel rettale	not available	029480035	EG S.P.A.	IT
MESALAZINA EG 400 mg capsule rigide	not available	029480011	EG S.P.A.	IT
MESALAZINA EG 800 mg compresse gastroresistenti	not available	029480050	EG S.P.A.	IT
MESALAZINA EG 800 mg compresse gastroresistenti	not available	029480086	EG S.P.A.	IT
Mesalazina Mylan Generics 400 mg compresse gastroresistenti	not available	033529052	MYLAN S.P.A.	IT
Mesalazina Mylan Generics 800 mg compresse gastroresistenti	not available	033529064	MYLAN S.P.A.	IT
MESALAZINA SANDOZ	not available	034836015	SANDOZ S.P.A.	IT
MESALAZINA SANDOZ	not available	034836027	SANDOZ S.P.A.	IT
Mesalazina Sandoz 4 g sospensione rettale	not available	034836039	SANDOZ S.P.A.	IT
Mesalazine Aurobindo 500 mg, zetpillen	NL/H/0286/002	RVG 24112	AUROBINDO PHARMA B.V.	NL
Mesalazine Aurobindo EC 250 mg, maagsapresistente tabletten.	not available	RVG 27071	AUROBINDO PHARMA B.V.	NL

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Mesalazine Aurobindo EC 500 mg, maagsapresistente tabletten.	not available	RVG 27072	AUROBINDO PHARMA B.V.	NL
MESALAZINE MAYOLY 500 mg, suppositoire	ES/H/0588/003	34009 302 601 3 5	LABORATOIRES MAYOLY SPINDLER	FR
MESALAZINE MAYOLY 500 mg, suppositoire	ES/H/0588/003	34009 302 601 4 2	LABORATOIRES MAYOLY SPINDLER	FR
MESALAZINE MAYOLY 500 mg, suppositoire	ES/H/0588/003	34009 302 601 5 9	LABORATOIRES MAYOLY SPINDLER	FR
MESALAZINE MAYOLY 500 mg, suppositoire	ES/H/0588/003	34009 302 601 6 6	LABORATOIRES MAYOLY SPINDLER	FR
MESALAZINE MAYOLY 500 mg, suppositoire	ES/H/0588/003	34009 550 911 2 7	LABORATOIRES MAYOLY SPINDLER	FR
Mesalazine Nordic 750 mg, maagsapresistente tabletten	not available	RVG 114896	NORDIC GROUP B.V.	NL
Mesalazine Nordic EC 250, maagsapresistente tabletten 250 mg	not available	RVG 27063	NORDIC GROUP B.V.	NL
Mesalazine Nordic EC 500, maagsapresistente tabletten 500 mg	not available	RVG 27064	NORDIC GROUP B.V.	NL
MESALAZINE SANDOZ TABLET EC 250, maagsapresistente tabletten 250 mg	not available	RVG 26915	SANDOZ B.V.	NL

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
MESALAZINE SANDOZ TABLET EC 500, maagsapresistente tabletten 500 mg	not available	RVG 26916	SANDOZ B.V.	NL
Mesalazine Teva 250 mg, zetpillen	not available	RVG 26917	TEVA NEDERLAND B.V.	NL
Mesalazine Teva 500 mg comprimés gastro-résistants.	not available	BE215747	TEVA PHARMA BELGIUM N.V./S.A	BE
Mesalazine Teva 500 mg maagsapresistente tabletten	not available	BE215747	TEVA PHARMA BELGIUM N.V./S.A	BE
MESALAZINE TEVA 500 mg MAGENSAFTRESISTENTE TABLETTEN	not available	BE215747	TEVA PHARMA BELGIUM N.V./S.A	BE
Mesalazine Teva 500 mg, zetpillen	not available	RVG 26918	TEVA NEDERLAND B.V.	NL
Mesalmin 2g/30g Klysmen Rektalsuspension	not available	43095.00.00	DR. FALK PHARMA G.M.B.H.	DE
Mesalmin 500mg magensaftresistente Tabletten	not available	43104.00.00	DR. FALK PHARMA G.M.B.H.	DE
Mesavancol 1200 mg compresse gastroresistenti, a rilascio prolungato.	NL/H/0733/001	037734011	GIULIANI SPA	IT
Mesavancol 1200 mg compresse gastroresistenti, a rilascio prolungato.	NL/H/0733/001	037734023	GIULIANI SPA	IT
Mesavancol 1200 mg maagsapresistent, tabletten met een verlengde afgifte	NL/H/0733/001	33597	GIULIANI SPA	NL

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Mezavant 1 200 mg comprimés gastro-résistants à libération prolongée.	NL/H/0732/001	BE293587	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Mezavant 1 200 mg comprimés gastro-résistants à libération prolongée.	NL/H/0732/001	BE293587	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Mezavant 1 200 mg comprimés gastro-résistants à libération prolongée.	NL/H/0732/001	0485173	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	LU
Mezavant 1 200 mg comprimés gastro-résistants à libération prolongée.	NL/H/0732/001	0485187	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	LU
Mezavant 1 200 mg enterodepottabletter.	NL/H/0732/001	23419	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	SE
Mezavant 1 200 mg enterodepottabletter.	NL/H/0732/001	23419	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	SE
Mezavant 1.200 mg comprimidos de liberación prolongada gastrorresistentes.	NL/H/0732/001	70144	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	ES
Mezavant 1.200 mg comprimidos de liberación	NL/H/0732/001	70144	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND	ES

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
prolongada gastrorresistentes.			BRANCH	
Mezavant 1200 mg comprimidos gastrorresistentes de libertação prolongada.	NL/H/0732/001	5393632	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	PT
Mezavant 1200 mg comprimidos gastrorresistentes de libertação prolongada.	NL/H/0732/001	5393640	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	PT
Mezavant 1200 mg enterodepottabletter	NL/H/0732/001	06-3932	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	NO
Mezavant 1200 mg enterodepottabletter	NL/H/0732/001	06-3932	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	NO
Mezavant 1200 mg maagsapresistente, tabletten met verlengde afgifte.	NL/H/0732/001	BE293587	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Mezavant 1200 mg maagsapresistente, tabletten met verlengde afgifte.	NL/H/0732/001	BE293587	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Mezavant 1200 mg maagsapresistente, tabletten met verlengde afgifte.	NL/H/0732/001	RVG 33600	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	NL

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Mezavant 1200 mg maagsapresistente, tabletten met verlengde afgifte.	NL/H/0732/001	RVG 33600	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	NL
Mezavant 1200 mg magensaftresistente Retardtabletten	NL/H/0732/001	BE293587	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Mezavant 1200 mg magensaftresistente Retardtabletten	NL/H/0732/001	BE293587	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Mezavant 1200 mg magensaftresistente Retardtabletten	NL/H/0732/001	0485173	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	LU
Mezavant 1200 mg magensaftresistente Retardtabletten	NL/H/0732/001	0485187	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	LU
Mezavant 1200 mg magensaftresistente Retardtabletten.	NL/H/0732/001	1-26880	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	AT
Mezavant 1200 mg magensaftresistente Retardtabletten.	NL/H/0732/001	1-26880	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	AT
Mezavant 1200 mg magensaftresistente	NL/H/0732/001	64753.00.00	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND	DE

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Retardtabletten.			BRANCH	
Mezavant 1200 mg magensaftresistente Retardtabletten.	NL/H/0732/001	64753.00.00	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	DE
Mezavant 1200 mg, γαστροανθεκτικά δισκία παρατεταμένης αποδέσμευσης.	NL/H/0732/001	20250	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	CY
Mezavant 1200 mg, γαστροανθεκτικά δισκία παρατεταμένης αποδέσμευσης.	NL/H/0732/001	20250	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	CY
Mezavant 1200 mg, γαστροανθεκτικά δισκία παρατεταμένης αποδέσμευσης.	NL/H/0732/001	129311/21/28-03-2022	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	GR
Mezavant 1200 mg, γαστροανθεκτικά δισκία παρατεταμένης αποδέσμευσης.	NL/H/0732/001	129311/21/28-03-2022	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	GR
Mezavant XL 1200 mg gastro-resistant, prolonged release tablets	NL/H/0732/001	PA23211/004/001	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	IE
Mezavant XL 1200 mg, gastro-resistant, prolonged release tablets.	NL/H/0732/001	PL 16189/0142	TAKEDA UK LTD	XI

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Mezavant XL 1200 mg, gastro-resistant, prolonged release tablets.	NL/H/0732/001	PL 16189/0142	TAKEDA UK LTD	XI
Mezavant XL 1200 mg, gastro-resistant, prolonged-release tablets.	NL/H/0732/001	MA1464/00201	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	MT
Mezavant XL gastro-resistant, prolonged release tablets	NL/H/0732/001	PA23211/004/001	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	IE
Mezavant, enterodepottabletter	NL/H/0732/001	38969	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	DK
Mezavant, enterodepottabletter	NL/H/0732/001	38969	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	DK
Movevit 1000 mg supositórios	PT/H/2280/002/DC	PT/H/2280/002/DC	LABORATÓRIOS VITÓRIA, SA	PT
Movevit 1000 mg supositórios	PT/H/2280/002/DC	PT/H/2280/002/DC	LABORATÓRIOS VITÓRIA, SA	PT
Movevit 1000 mg supositórios	PT/H/2280/002/DC	PT/H/2280/002/DC	LABORATÓRIOS VITÓRIA, SA	PT
Movevit 1000 mg supositórios	PT/H/2280/002/DC	PT/H/2280/002/DC	LABORATÓRIOS VITÓRIA, SA	PT
Movevit 500 mg supositório	PT/H/2280/001	PT/H/2280/001	FAES FARMA PORTUGAL, S.A.	PT

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Movevit 500 mg supositórios	PT/H/2280/001/DC	PT/H/2280/001/DC	LABORATÓRIOS VITÓRIA, SA	PT
Movevit 500 mg supositórios	PT/H/2280/001/DC	PT/H/2280/001/DC	LABORATÓRIOS VITÓRIA, SA	PT
Movevit 500 mg supositórios	PT/H/2280/001/DC	PT/H/2280/001/DC	LABORATÓRIOS VITÓRIA, SA	PT
Octasa 1 g Suppositories	not available	PL 36633/0011	TILLOTTS PHARMA UK, LTD.	XI
Octasa 1600 mg modified-release tablets	SE/H/1654/001	PL 36633/0009	TILLOTTS PHARMA UK, LTD.	XI
Octasa 400 mg Modified Release Tablets	not available	PL 36633/0002	TILLOTTS PHARMA UK, LTD.	XI
Octasa 800 mg Modified Release Tablets	not available	PL 36633/0001	TILLOTTS PHARMA UK, LTD.	XI
Osperzo 1 g, suppositoire	DE/H/5682/001	34009 302 750 7 8	DR. FALK PHARMA G.M.B.H.	FR
Osperzo 1 g, suppositoire	DE/H/5682/001	34009 302 750 8 5	DR. FALK PHARMA G.M.B.H.	FR
Osperzo 1 g, suppositoire	DE/H/5682/001	34009 302 750 9 2	DR. FALK PHARMA G.M.B.H.	FR
Osperzo 1 g, suppositoire	DE/H/5682/001	34009 302 751 0 8	DR. FALK PHARMA G.M.B.H.	FR
Osperzo 1 g, suppositoire	DE/H/5682/001	34009 550 961 4 6	DR. FALK PHARMA G.M.B.H.	FR
Osperzo 1 g, suppositoire	DE/H/5682/001	34009 550 961 5 3	DR. FALK PHARMA G.M.B.H.	FR
Osperzo 1 g, suppositoire	DE/H/5682/001	34009 302 750 6 1	DR. FALK PHARMA G.M.B.H.	FR
OSPERZO 1,5 g, granulés á libération prolongée	DE/H/0363/004	34009 302 646 6 9	DR. FALK PHARMA G.M.B.H.	FR

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
OSPERZO 1,5 g, granulés á libération prolongée	DE/H/0363/004	34009 302 646 7 6	DR. FALK PHARMA G.M.B.H.	FR
OSPERZO 1,5 g, granulés á libération prolongée	DE/H/0363/004	34009 302 646 8 3	DR. FALK PHARMA G.M.B.H.	FR
OSPERZO 1,5 g, granulés á libération prolongée	DE/H/0363/004	34009 302 646 9 0	DR. FALK PHARMA G.M.B.H.	FR
OSPERZO 1,5 g, granulés á libération prolongée	DE/H/0363/004	34009 302 647 0 6	DR. FALK PHARMA G.M.B.H.	FR
OSPERZO 1,5 g, granulés á libération prolongée	DE/H/0363/004	34009 302 647 1 3	DR. FALK PHARMA G.M.B.H.	FR
OSPERZO 1,5 g, granulés á libération prolongée	DE/H/0363/004	34009 550 927 1 1	DR. FALK PHARMA G.M.B.H.	FR
OSPERZO 1,5 g, granulés á libération prolongée	DE/H/0363/004	34009 550 927 2 8	DR. FALK PHARMA G.M.B.H.	FR
OSPERZO 1,5 g, granulés á libération prolongée	DE/H/0363/004	34009 550 927 3 5	DR. FALK PHARMA G.M.B.H.	FR
OSPERZO 1,5 g, granulés á libération prolongée	DE/H/0363/004	34009 550 927 4 2	DR. FALK PHARMA G.M.B.H.	FR
OSPERZO 1000 mg, granulés à libération prolongée	DE/H/0363/002	34009 302 646 3 8	DR. FALK PHARMA G.M.B.H.	FR
OSPERZO 1000 mg, granulés à libération prolongée	DE/H/0363/002	34009 302 646 4 5	DR. FALK PHARMA G.M.B.H.	FR
OSPERZO 1000 mg, granulés à libération prolongée	DE/H/0363/002	34009 302 646 5 2	DR. FALK PHARMA G.M.B.H.	FR
OSPERZO 1000 mg, granulés à libération prolongée	DE/H/0363/002	34009 550 926 9 8	DR. FALK PHARMA G.M.B.H.	FR
OSPERZO 1000 mg, granulés à libération prolongée	DE/H/0363/002	34009 550 927 0 4	DR. FALK PHARMA G.M.B.H.	FR
OSPERZO 3 g, granulés à libération prolongée	DE/H/0363/007	34009 302 647 2 0	DR. FALK PHARMA G.M.B.H.	FR

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
OSPERZO 3 g, granulés à libération prolongée	DE/H/0363/007	34009 302 647 3 7	DR. FALK PHARMA G.M.B.H.	FR
OSPERZO 3 g, granulés à libération prolongée	DE/H/0363/007	34009 302 647 4 4	DR. FALK PHARMA G.M.B.H.	FR
OSPERZO 3 g, granulés à libération prolongée	DE/H/0363/007	34009 302 647 5 1	DR. FALK PHARMA G.M.B.H.	FR
OSPERZO 3 g, granulés à libération prolongée	DE/H/0363/007	34009 550 927 5 9	DR. FALK PHARMA G.M.B.H.	FR
OSPERZO 3 g, granulés à libération prolongée	DE/H/0363/007	34009 550 927 6 6	DR. FALK PHARMA G.M.B.H.	FR
OSPERZO 3 g, granulés à libération prolongée	DE/H/0363/007	34009 550 927 7 3	DR. FALK PHARMA G.M.B.H.	FR
OSPERZO 3 g, granulés à libération prolongée	DE/H/0363/007	34009 550 927 8 0	DR. FALK PHARMA G.M.B.H.	FR
OSPERZO 500 mg, granulés à libération prolongée	DE/H/0363/001	34009 302 646 1 4	DR. FALK PHARMA G.M.B.H.	FR
OSPERZO 500 mg, granulés à libération prolongée	DE/H/0363/001	34009 302 646 2 1	DR. FALK PHARMA G.M.B.H.	FR
OSPERZO 500 mg, granulés à libération prolongée	DE/H/0363/001	34009 550 926 8 1	DR. FALK PHARMA G.M.B.H.	FR
PENTACOL 1,5 g granulato per sospensione rettale	not available	026925038	ALFASIGMA S.P.A.	IT
PENTACOL 1200 mg compresse gastroresistenti a rilascio modificato	not available	026925228	ALFASIGMA S.P.A.	IT
PENTACOL 1200 mg compresse gastroresistenti a rilascio modificato	not available	026925230	ALFASIGMA S.P.A.	IT
PENTACOL 2 g schiuma rettale	not available	026925127	ALFASIGMA S.P.A.	IT

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
PENTACOL 4 g schiuma rettale	not available	026925115	ALFASIGMA S.P.A.	IT
PENTACOL 4 g/100 ml sospensione rettale	not available	026925141	ALFASIGMA S.P.A.	IT
PENTACOL 400 mg compresse gastroresistenti a rilascio modificato	not available	026925040	ALFASIGMA S.P.A.	IT
PENTACOL 500 mg gel rettale	not available	026925180	ALFASIGMA S.P.A.	IT
PENTACOL 500 mg gel rettale	not available	026925065	ALFASIGMA S.P.A.	IT
PENTACOL 500 mg supposte	not available	026925154	ALFASIGMA S.P.A.	IT
PENTACOL 800 mg compresse gastroresistenti a rilascio modificato	not available	026925166	ALFASIGMA S.P.A.	IT
PENTACOL 800 mg compresse gastroresistenti a rilascio modificato	not available	026925053	ALFASIGMA S.P.A.	IT
PENTACOL 800 mg compresse gastroresistenti a rilascio modificato	not available	026925178	ALFASIGMA S.P.A.	IT
Pentasa 1000 mg Supositórios	not available	2177384	FERRING S.A.U.	PT
Pentasa 1g čípky	not available	29 /009/95 - C	FERRING LÉCIVA A.S.	CZ
Pentasa 1g Granulado de libertação modificada	not available	3292687	FERRING S.A.U.	PT
Pentasa 1 g - Klysma	not available	1-19907	FERRING ARZNEIMITTEL	AT

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
			GES.M.B.H.	
Pentasa 1 g - Zäpfchen	not available	1-20055	FERRING ARZNEIMITTEL GES.M.B.H.	AT
Pentasa 1 g čapíky	not available	29/0425/95-S	FERRING LÉCIVA A.S.	SK
Pentasa 1 g čepići	not available	HR-H-729224011	FERRING GMBH	HR
PENTASA 1 g comprimate cu eliberare prelungită	not available	4480/2012/01	FERRING GMBH	RO
Pentasa 1 g comprimés à libération prolongée	not available	BE477306	FERRING N.V.	BE
Pentasa 1 g comprimés à libération prolongée	not available	2016040070	FERRING N.V.	LU
PENTASA 1 g depotgranulat	not available	23857	FERRING LÄÄKKEET OY	FI
PENTASA 1 g depotrakeet	not available	23857	FERRING LÄÄKKEET OY	FI
PENTASA 1 g depottablett	not available	27787	FERRING LÄÄKKEET OY	FI
Pentasa 1 g depottabletter	not available	09-6965	FERRING LEGEMIDLER AS	NO
Pentasa 1 g depottabletter	not available	42950	FERRING LÄKEMEDEL AB	SE
PENTASA 1 g depottabletti	not available	27787	FERRING LÄÄKKEET OY	FI
Pentasa 1 g endaþarmsdreifa.	not available	880163	FERRING LÆGEMIDLER A/S	IS
Pentasa 1 g endaþarmsstílar.	not available	900152 (IS)	FERRING LÆGEMIDLER A/S	IS

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Pentasa 1 g forðatöflur	not available	IS/1/10/135/01	FERRING LÆGEMIDLER A/S	IS
PENTASA 1 g granulato a rilascio prolungato	not available	027130083	FERRING S.P.A.	IT
Pentasa 1 g ilgstošās darbības granulas	not available	10-0562	FERRING GMBH	LV
PENTASA 1 g pailginto atpalaidavimo granulės	not available	LT/1/96/1965/001	FERRING GMBH	LT
PENTASA 1 g pailginto atpalaidavimo granulės	not available	LT/1/96/1965/002	FERRING GMBH	LT
PENTASA 1 g pailginto atpalaidavimo granulės	not available	LT/1/96/1965/008	FERRING GMBH	LT
PENTASA 1 g pailginto atpalaidavimo granulės	not available	LT/1/96/1965/003	FERRING GMBH	LT
PENTASA 1 g peräpuikko	not available	10993	FERRING LÄÄKKEET OY	FI
Pentasa 1 g prolonged-release tablets	not available	PA 1009/006/007	FERRING IRELAND LTD.	IE
Pentasa 1 g rektálna suspenzia	not available	29/0319/00-S	FERRING LÉCIVA A.S.	SK
Pentasa 1 g rektalna suspenzija	not available	HR-H-296920131	FERRING GMBH	HR
PENTASA 1 g rektální suspenze	not available	29/183/00-C	FERRING LÉCIVA A.S.	CZ
Pentasa 1 g rektalsuspension	not available	14430	FERRING LÄKEMEDEL AB	SE
PENTASA 1 g rektalvæske, suspensjon	not available	7402	FERRING LEGEMIDLER AS	NO
Pentasa 1 g retard tabletta	not available	OGYI-T- 4798/05	FERRING MAGYARORSZÁG GYÓGYSZERKERESKEDELMI KFT.	HU

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Pentasa 1 g Retardtabletten	not available	BE477306	FERRING N.V.	BE
Pentasa 1 g Retardtabletten	not available	2016040070	FERRING N.V.	LU
PENTASA 1 g stikkpiller	not available	03-2326	FERRING LEGEMIDLER AS	NO
Pentasa 1 g supozitoriji	not available	97-0616	FERRING GMBH	LV
Pentasa 1 g suppositoires	not available	BE164972	FERRING N.V.	BE
Pentasa 1 g suppositoires	not available	0349/09050317	FERRING N.V.	LU
PENTASA 1 g suppositorium	not available	10993	FERRING LÄÄKKEET OY	FI
PENTASA 1 g supposte	not available	027130069	FERRING S.P.A.	IT
PENTASA 1 g suspensión rectal	not available	60.853	FERRING S.A.U.	ES
Pentasa 1 g tablete s produljenim oslobađanjem	not available	HR-H-589842039	FERRING GMBH	HR
Pentasa 1 g tabletten met verlengde afgifte	not available	BE477306	FERRING N.V.	BE
Pentasa 1 g végbélkúp	not available	OGYI-T- 4798/02	FERRING MAGYARORSZÁG GYÓGYSZERKERESKEDELMI KFT.	HU
Pentasa 1 g végbélszuszpenzió	not available	OGYI-T- 4798/03	FERRING MAGYARORSZÁG GYÓGYSZERKERESKEDELMI KFT.	HU
Pentasa 1 g Zäpfchen	not available	BE164972	FERRING N.V.	BE

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Pentasa 1 g Zäpfchen	not available	0349/09050317	FERRING N.V.	LU
Pentasa 1 g zetpillen	not available	BE164972	FERRING N.V.	BE
PENTASA 1 g žvakutės	not available	LT/1/96/1965/007	FERRING GMBH	LT
PENTASA 1 g, granulés en sachet	not available	3400934748622	FERRING S.A.S.	FR
PENTASA 1 g, granulés en sachet	not available	3400934424601	FERRING S.A.S.	FR
PENTASA 1 g, granulés en sachet	not available	3400934424779	FERRING S.A.S.	FR
PENTASA 1 g, suppositoire	not available	34009 335 028 4 3	FERRING S.A.S.	FR
PENTASA 1 g, suppositoire	not available	34009 301 817 1 3	FERRING S.A.S.	FR
Pentasa 1 g, tabletten met verlengde afgifte	not available	RVG 105712	FERRING B.V.	NL
Pentasa 1 g, toimeainet prolongeeritult vabastavad graanulid	not available	659709	FERRING GMBH	EE
Pentasa 1 g/100 ml Rektalsuspension	not available	BE150683	FERRING N.V.	BE
PENTASA 1 g/100 ml rektalsuspension	not available	9527	FERRING LÄÄKKEET OY	FI
Pentasa 1 g/100 ml Rektalsuspension	not available	2009050315	FERRING N.V.	LU
Pentasa 1 g/100 ml suspensie voor rectaal gebruik	not available	BE150683	FERRING N.V.	BE
Pentasa 1 g/100 ml suspension rectale	not available	BE150683	FERRING N.V.	BE

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Pentasa 1 g/100 ml suspension rectale	not available	2009050315	FERRING N.V.	LU
PENTASA 1 g/100 ml, suspension rectale	not available	34009 301 819 0 4	FERRING S.A.S.	FR
Pentasa 10 mg/ml Rectal Suspension	not available	PA 1009/006/003	FERRING IRELAND LTD.	IE
Pentasa 1000 mg Comprimidos de libertação prolongada	not available	5323845	FERRING S.A.U.	PT
PENTASA 1000 mg Retardtabletten	not available	78793.00.00	FERRING GMBH	DE
Pentasa 1000 mg Supositórios	not available	5868047	FERRING S.A.U.	PT
Pentasa 1000 mg Supositórios	not available	5818745	FERRING S.A.U.	PT
Pentasa 1000 mg supozitoare	not available	8670/2016/01	FERRING GMBH	RO
PENTASA 1000 mg Zäpfchen	not available	16944.00.02	FERRING GMBH	DE
PENTASA 1g compresse a rilascio modificato	not available	027130107	FERRING S.P.A.	IT
Pentasa 1g granulado de liberación prolongada	DK/H/0140/001	62.670	FERRING S.A.U.	ES
Pentasa 1g Granulado de libertação modificada	not available	3292588	FERRING S.A.U.	PT
Pentasa 1g Granulado de libertação modificada	not available	5779210	FERRING S.A.U.	PT
PENTASA 1g Supositorios	not available	60.151	FERRING S.A.U.	ES
Pentasa 1g suppositorier	not available	16690	FERRING LÄKEMEDEL AB	SE

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Pentasa 1g Suppositories	not available	PA 1009/006/002	FERRING IRELAND LTD.	IE
Pentasa 1g zetpillen	not available	RVG 15064	FERRING B.V.	NL
PENTASA 1g, granulés en sachet.	not available	34009 301 817 2 0	FERRING S.A.S.	FR
PENTASA 1g/100 ml peräruiskesuspensio	not available	9527	FERRING LÄÄKKEET OY	FI
Pentasa 1g/100 ml, suspensie voor rectaal gebruik	not available	RVG 11782	FERRING B.V.	NL
PENTASA 2 g depotgranulat	not available	23858	FERRING LÄÄKKEET OY	FI
PENTASA 2 g depotrakeet	not available	23858	FERRING LÄÄKKEET OY	FI
Pentasa 2 g granule cu eliberare prelungită	not available	7178/2014/01	FERRING GMBH	RO
Pentasa 2 g granule cu eliberare prelungită	not available	7178/2014/02	FERRING GMBH	RO
Pentasa 2 g granule s produljenim oslobaĎanjem u vrećici	not available	HR-H-282835727	FERRING GMBH	HR
Pentasa 2 g ilgstošās darbības granulas	not available	10-0563	FERRING GMBH	LV
PENTASA 2 g pailginto atpalaidavimo granulės	not available	LT/1/96/1965/004	FERRING GMBH	LT
PENTASA 2 g pailginto atpalaidavimo granulės	not available	LT/1/96/1965/005	FERRING GMBH	LT
PENTASA 2 g pailginto atpalaidavimo granulės	not available	LT/1/96/1965/009	FERRING GMBH	LT
Pentasa 2 g retard granulátum	not available	OGYI-T- 4798/04	FERRING MAGYARORSZÁG	HU

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
			GYÓGYSZERKERESKEDELMI KFT.	
Pentasa 2 g zrnca s podaljšanim sproščanjem	DK/H/0140/002	H/12/01236/003	FERRING GMBH	SI
Pentasa 2 g zrnca s podaljšanim sproščanjem	DK/H/0140/002	H/12/01236/001	FERRING GMBH	SI
Pentasa 2 g zrnca s podaljšanim sproščanjem	DK/H/0140/002	H/12/01236/002	FERRING GMBH	SI
Pentasa 2 g κοκκία παρατεταμένης αποδέσμευσης	DK/H/0140/002	021943	FERRING HELLAS MEPE	CY
Pentasa 2 g κοκκία παρατεταμένης αποδέσμευσης	DK/H/0140/002	22417/22.03.2018	FERRING HELLAS MEPE	GR
PENTASA 2 g, granulés en sachet-dose	not available	3400936758797	FERRING S.A.S.	FR
Pentasa 2 g, toimeainet prolongeeritult vabastavad graanulid	not available	659809	FERRING GMBH	EE
Pentasa 2000 mg Granulado de libertação modificada	not available	5731542	FERRING S.A.U.	PT
Pentasa 2g granulado de liberación prolongada	not available	77.668	FERRING S.A.U.	ES
PENTASA 4 g depotgranulat	not available	34347	FERRING LÄÄKKEET OY	FI
PENTASA 4 g depotrakeet	not available	34347	FERRING LÄÄKKEET OY	FI
Pentasa 4 g granulado de liberación prolongada	DK/H/0140/003	79555	FERRING S.A.U.	ES
PENTASA 4 g pailginto atpalaidavimo granulės	not available	LT/1/23/5295/001	FERRING GMBH	LT

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Pentasa 4 g retard granulátum	not available	OGYI-T- 4798/06	FERRING MAGYARORSZÁG GYÓGYSZERKERESKEDELMI KFT.	HU
Pentasa 4 g zrnca s podaljšanim sproščanjem	DK/H/0140/003	H/12/01236/007	FERRING GMBH	SI
Pentasa 4 g zrnca s podaljšanim sproščanjem	DK/H/0140/003	H/12/01236/005	FERRING GMBH	SI
Pentasa 4 g zrnca s podaljšanim sproščanjem	DK/H/0140/003	H/12/01236/004	FERRING GMBH	SI
Pentasa 4 g zrnca s podaljšanim sproščanjem	DK/H/0140/003	H/12/01236/006	FERRING GMBH	SI
Pentasa 4 g κοκκία παρατεταμένης αποδέσμευσης	DK/H/140/003	022373	FERRING HELLAS MEPE	CY
Pentasa 4 g κοκκία παρατεταμένης αποδέσμευσης	DK/H/0140/003	16611/05.03.2015	FERRING HELLAS MEPE	GR
Pentasa 4 g, toimeainet prolongeeritult vabastavad graanulid	not available	1151624	FERRING GMBH	EE
PENTASA 4 g/100 ml sospensione rettale.	not available	027130044	FERRING S.P.A.	IT
Pentasa 4000 mg Granulado de libertação modificada	not available	5757935	FERRING S.A.U.	PT
PENTASA 500 mg compresse a rilascio modificato	not available	027130071	FERRING S.P.A.	IT
Pentasa 500 mg comprimate cu eliberare prelungită	not available	9996/2017/01	FERRING GMBH	RO
Pentasa 500 mg comprimés à libération prolongée	not available	BE156055	FERRING N.V.	BE

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Pentasa 500 mg comprimés à libération prolongée	not available	BE230991	FERRING N.V.	BE
Pentasa 500 mg comprimés à libération prolongée	not available	2009050316	FERRING N.V.	LU
Pentasa 500 mg Comprimido de Libertação Prolongada	not available	2178887	FERRING S.A.U.	PT
Pentasa 500 mg Comprimido de Libertação Prolongada	not available	2178986	FERRING S.A.U.	PT
Pentasa 500 mg Comprimido de Libertação Prolongada	not available	5779202	FERRING S.A.U.	PT
PENTASA 500 mg comprimidos de liberación prolongada	not available	60.152	FERRING S.A.U.	ES
PENTASA 500 mg depottablett	not available	10992	FERRING LÄÄKKEET OY	FI
PENTASA 500 mg depottabletter	not available	7762	FERRING LEGEMIDLER AS	NO
Pentasa 500 mg depottabletter	not available	12307	FERRING LÄKEMEDEL AB	SE
PENTASA 500 mg depottabletti	not available	10992	FERRING LÄÄKKEET OY	FI
Pentasa 500 mg forðatöflur.	not available	900151	FERRING LÆGEMIDLER A/S	IS
Pentasa 500 mg ilgstošās darbības tabletes	not available	98-0118	FERRING GMBH	LV
PENTASA 500 mg pailginto atpalaidavimo tabletės	not available	LT/1/96/1965/006	FERRING GMBH	LT

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Pentasa 500 mg prolonged-release tablets	not available	PA 1009/006/005	FERRING IRELAND LTD.	IE
Pentasa 500 mg retard tabletta	not available	OGYI-T- 4798/01	FERRING MAGYARORSZÁG GYÓGYSZERKERESKEDELMI KFT.	HU
Pentasa 500 mg Retardtabletten	not available	BE156055	FERRING N.V.	BE
Pentasa 500 mg Retardtabletten	not available	BE230991	FERRING N.V.	BE
PENTASA 500 mg Retardtabletten	not available	16944.00.01	FERRING GMBH	DE
Pentasa 500 mg Retardtabletten	not available	2009050316	FERRING N.V.	LU
Pentasa 500 mg tablete s podaljšanim sproščanjem	not available	H/12/01972/001	FERRING GMBH	SI
Pentasa 500 mg tablete s produljenim oslobadanjem	not available	HR-H-120226953	FERRING GMBH	HR
Pentasa 500 mg tabletten met verlengde afgifte	not available	BE156055	FERRING N.V.	BE
Pentasa 500 mg tabletten met verlengde afgifte	not available	BE230991	FERRING N.V.	BE
Pentasa 500 mg, tabletten met verlengde afgifte	not available	RVG 14797	FERRING B.V.	NL
Pentasa Compact 1 g, granulaat met verlengde afgifte	not available	RVG 18706	FERRING B.V.	NL
Pentasa Compact 2 g, granulaat met verlengde afgifte	not available	RVG 31379	FERRING B.V.	NL

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Pentasa Compact 4 g, granulaat met verlengde afgifte	not available	RVG 114015	FERRING B.V.	NL
Pentasa Prolong 1g tablety s prodlouženým uvolňováním	not available	29/011/11-C	FERRING LÉCIVA A.S.	CZ
Pentasa Prolong 500 mg tablety s prodlouženým uvolňováním	not available	29/014/95 - C	FERRING LÉCIVA A.S.	CZ
Pentasa retard 1 g - Granulat	not available	1-22009	FERRING ARZNEIMITTEL GES.M.B.H.	AT
Pentasa retard 1 g - Tabletten	not available	1-31050	FERRING ARZNEIMITTEL GES.M.B.H.	AT
Pentasa retard 2 g - Granulat	not available	1-26559	FERRING ARZNEIMITTEL GES.M.B.H.	AT
Pentasa retard 4 g - Granulat	DK/H/0140/003	139055	FERRING ARZNEIMITTEL GES.M.B.H.	AT
Pentasa retard 500 mg - Tabletten	not available	1-20047	FERRING ARZNEIMITTEL GES.M.B.H.	AT
PENTASA Sachet 1 g depotgranulat	not available	98-5186	FERRING LEGEMIDLER AS	NO
Pentasa Sachet 1 g depotgranulat	DK/H/0140/001	14752	FERRING LÄKEMEDEL AB	SE
Pentasa Sachet 1 g forðakyrni	not available	940305	FERRING LÆGEMIDLER A/S	IS

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Pentasa Sachet 1 g granulaat met verlengde afgifte	not available	BE209772	FERRING N.V.	BE
Pentasa Sachet 1 g granulés à libération prolongée	not available	BE209772	FERRING N.V.	BE
Pentasa Sachet 1 g granulés à libération prolongée	not available	0349/08100013	FERRING N.V.	LU
Pentasa Sachet 1 g Retardgranulat	not available	BE209772	FERRING N.V.	BE
Pentasa Sachet 1 g Retardgranulat	not available	0349/08100013	FERRING N.V.	LU
PENTASA Sachet 1000 mg Retardgranulat	DK/H/0140/001	44367.00.00	FERRING GMBH	DE
PENTASA Sachet 1g prolonged release granules	DK/H/0140/001	PL 03194/0075	FERRING PHARMACEUTICALS LTD.	XI
Pentasa Sachet 1g prolonged-release granules	DK/H/0140/001	PA 1009/6/1	FERRING IRELAND LTD.	IE
PENTASA Sachet 2 g depotgranulat	not available	04-2948	FERRING LEGEMIDLER AS	NO
Pentasa Sachet 2 g depotgranulat	DK/H/0140/002	25569	FERRING LÄKEMEDEL AB	SE
Pentasa Sachet 2 g forðakyrni	not available	IS/1/05/104/01	FERRING LÆGEMIDLER A/S	IS
Pentasa Sachet 2 g granulaat met verlengde afgifte	not available	BE343734	FERRING N.V.	BE
Pentasa Sachet 2 g granulát s predlženým uvolnovaním	not available	29/0155/07-S	FERRING LÉCIVA A.S.	SK
Pentasa Sachet 2 g granulés à libération prolongée	not available	BE343734	FERRING N.V.	BE

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Pentasa Sachet 2 g granulés à libération prolongée	not available	2010040049	FERRING N.V.	LU
Pentasa Sachet 2 g Retardgranulat	not available	BE343734	FERRING N.V.	BE
Pentasa Sachet 2 g Retardgranulat	not available	2010040049	FERRING N.V.	LU
Pentasa Sachet 2g granules prodlouženým uvolnováním	not available	29/669/07-C	FERRING LÉCIVA A.S.	CZ
Pentasa Sachet 2g prolonged release granules	DK/H/0140/002	PL 03194/0102	FERRING PHARMACEUTICALS LTD.	XI
Pentasa Sachet 2g prolonged-release granules	DK/H/0140/002	PA 1009/6/6	FERRING IRELAND LTD.	IE
Pentasa Sachet 4 g depotgranulat	not available	14-10423	FERRING LEGEMIDLER AS	NO
Pentasa Sachet 4 g depotgranulat	DK/H/0140/003	50369	FERRING LÄKEMEDEL AB	SE
Pentasa Sachet 4 g forðakyrni.	not available	IS/1/16/037/01	FERRING LÆGEMIDLER A/S	IS
Pentasa Sachet 4 g granulaat met verlengde afgifte	not available	BE506604	FERRING N.V.	BE
Pentasa Sachet 4 g granulát s predĺženým uvoľňovaním	not available	29/0187/17-S	FERRING LÉCIVA A.S.	SK
Pentasa Sachet 4 g granulés à libération prolongée	not available	BE506604	FERRING N.V.	BE
Pentasa Sachet 4 g granulés à libération prolongée	not available	2018010044	FERRING N.V.	LU

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Pentasa Sachet 4 g Retardgranulat	not available	BE506604	FERRING N.V.	BE
PENTASA Sachet 4 g Retardgranulat	DK/H/0140/003	91296.00.00	FERRING GMBH	DE
Pentasa Sachet 4 g Retardgranulat	not available	2018010044	FERRING N.V.	LU
Pentasa Sachet 4 g granule s prodlouženým uvolňováním v sáčku	not available	29/109/16-C	FERRING LÉCIVA A.S.	CZ
PENTASA Sachet 4g prolonged release granules	DK/H/0140/003	PA 1009/006/008	FERRING IRELAND LTD.	IE
PENTASA Sachet 4g prolonged release granules	DK/H/0140/003	PL 03194/0117	FERRING PHARMACEUTICALS LTD.	XI
Pentasa Sachet, depotgranulat, enkeltdosisbeholder	not available	55370	FERRING LÆGEMIDLER A/S	DK
Pentasa Sachet, depotgranulat, enkeltdosisbeholder	not available	16895	FERRING LÆGEMIDLER A/S	DK
Pentasa Sachet, depotgranulat, enkeltdosisbeholder	not available	36822	FERRING LÆGEMIDLER A/S	DK
Pentasa Slow release tablets 1 g tablety s predĺženým uvoľňovaním	not available	29/0532/10-S	FERRING LÉCIVA A.S.	SK
Pentasa Slow release tablets 500 mg tablety s predlženým uvolnovaním	not available	29/0426/95-S	FERRING LÉCIVA A.S.	SK
PENTASA Xtend 2 g Retardgranulat	DK/H/0140/002	67611.00.00	FERRING GMBH	DE
PENTASA, 1 g suposiidid	not available	145396	FERRING GMBH	EE

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
PENTASA, 1 g, czopki	not available	R/6568	FERRING GMBH	PL
PENTASA, 1 g, granulat o przedłużonym uwalnianiu	not available	8553	FERRING GMBH	PL
PENTASA, 1 g, tabletki o przedłużonym uwalnianiu	not available	20099	FERRING GMBH	PL
PENTASA, 1 g/100 ml, zawiesina doodbytnicza	not available	8188	FERRING GMBH	PL
PENTASA, 2 g, granulat o przedłużonym uwalnianiu	not available	20100	FERRING GMBH	PL
PENTASA, 4 g, granulat o przedłużonym uwalnianiu	not available	23149	FERRING GMBH	PL
Pentasa, 500 mg toimeainet prolongeeritult vabastavad tabletid	not available	145496	FERRING GMBH	EE
PENTASA, 500 mg, tabletki o przedłużonym uwalnianiu	not available	R/6621	FERRING GMBH	PL
Pentasa, depottabletter	not available	13655	FERRING LÆGEMIDLER A/S	DK
Pentasa, depottabletter	not available	45626	FERRING LÆGEMIDLER A/S	DK
Pentasa, rektalvæske, suspension, enkeltdosisbeholder	not available	11718	FERRING LÆGEMIDLER A/S	DK
Pentasa, suppositorier	not available	13943	FERRING LÆGEMIDLER A/S	DK
PENTASA® 1 g/100 ml, suspension rectale	not available	3400932934744	FERRING S.A.S.	FR

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
PENTASA® 1g, comprimé	not available	34009 498 584 3 9	FERRING S.A.S.	FR
PENTASA® 1g, suppositoire	not available	3400933502782	FERRING S.A.S.	FR
PENTASA® 500 mg, comprimé	not available	3400933278717	FERRING S.A.S.	FR
PENTASA® 500 mg, comprimé	not available	3400933278885	FERRING S.A.S.	FR
PENTASA® Mesalazine Enema	not available	PL 3194/0027	FERRING PHARMACEUTICALS LTD.	XI
PENTASA® Slow Release Tablets 1g	not available	PL 3194/0108	FERRING PHARMACEUTICALS LTD.	XI
PENTASA® Slow Release Tablets 500mg	not available	PL 3194/0044	FERRING PHARMACEUTICALS LTD.	XI
PENTASA® Suppositories 1g	not available	PL 03194/0045	FERRING PHARMACEUTICALS LTD.	XI
PENTASA1 g comprimidos de liberación prolongada	not available	77022	FERRING S.A.U.	ES
PENTASA1000 mg Rektalsuspension	not available	13913.00.00	FERRING GMBH	DE
Pentasa1000 mg/100 ml Suspensão retal	not available	2233286	FERRING S.A.U.	PT
Pentasa1000 mg/100 ml Suspensão retal	not available	3968682	FERRING S.A.U.	PT
Quintasa Sachet, depotgranulat	DK/H/0140/003	53238	FERRING LÆGEMIDLER A/S	DK

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Quintasa Sachet, depotgranulat	DK/H/0140/001	18936	FERRING LÆGEMIDLER A/S	DK
Quintasa Sachet, depotgranulat	DK/H/0140/002	36657	FERRING LÆGEMIDLER A/S	DK
ROWASA 250 mg, comprimé enrobé gastro-résistant	not available	333 824-8	TEOFARMA S.R.L.	FR
ROWASA 250 mg, comprimé enrobé gastro-résistant	not available	333 825-4	TEOFARMA S.R.L.	FR
ROWASA 250 mg, comprimé enrobé gastro-résistant	not available	333 826-0	TEOFARMA S.R.L.	FR
ROWASA 250 mg, comprimé enrobé gastro-résistant	not available	333 827-7	TEOFARMA S.R.L.	FR
ROWASA 500 mg, comprimé enrobé gastro-résistant	not available	333 820-2	TEOFARMA S.R.L.	FR
ROWASA 500 mg, comprimé enrobé gastro-résistant	not available	333 821-9	TEOFARMA S.R.L.	FR
ROWASA 500 mg, comprimé enrobé gastro-résistant	not available	333 822-5	TEOFARMA S.R.L.	FR
ROWASA 500 mg, comprimé enrobé gastro-résistant	not available	333 823-1	TEOFARMA S.R.L.	FR
ROWASA 500 mg, comprimé enrobé gastro-résistant	not available	560 454-7	TEOFARMA S.R.L.	FR
ROWASA 500 mg, suppositoire	not available	335 035-0	TEOFARMA S.R.L.	FR
ROWASA 500 mg, suppositoire	not available	335 036-7	TEOFARMA S.R.L.	FR
ROWASA 500 mg, suppositoire	not available	335 037-3	TEOFARMA S.R.L.	FR
Salaza, 1000 mg, czopki	PT/H/2281/002	25973	FAES FARMA, S.A.	PL

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Salaza, 1000 mg, tabletki dojelitowe	ES/H/0588/002	26293	FAES FARMA S.A.	PL
Salaza, 500 mg, czopki	PT/H/2281/001	25972	FAES FARMA, S.A.	PL
Salaza, 500 mg, tabletki dojelitowe	ES/H/0588/001	25650	FAES FARMA, S.A.	PL
Salcrozine 1 000 mg čapíky	PT/H/2281/002	29/0073/20-S	FAES FARMA, S.A.	SK
Salcrozine 1000 mg comprimate gastrorezistente	ES/H/0588/002	14426/2022/01	FAES FARMA, S.A.	RO
Salcrozine 1000 mg comprimate gastrorezistente	ES/H/0588/002	14426/2022/02	FAES FARMA, S.A.	RO
Salcrozine 1000 mg gastro-resistant tablets	not available	PL 18945/0006	FAES FARMA, S.A.	XI
Salcrozine 1000 mg gyomornedv-ellenálló tabletta	ES/H/0588/002	OGYI-T-24080/01	FAES FARMA, S.A.	HU
Salcrozine 1000 mg gyomornedv-ellenálló tabletta	ES/H/0588/002	OGYI-T-24080/02	FAES FARMA, S.A.	HU
Salcrozine 500 mg čapíky	PT/H/2281/001	29/0072/20-S	FAES FARMA, S.A.	SK
Salcrozine 500 mg comprimate gastrorezistente	ES/H/0588/001	13552/2020/01	FAES FARMA, S.A.	RO
Salcrozine 500 mg gastro-resistant tablets.	ES/H/0588/001	PL 18945/0002	FAES FARMA, S.A.	XI
Salcrozine 500 mg suppositories	not available	PL 18945/0010	FAES FARMA, S.A.	XI
Salcrozine 500 mg végbélkúp	ES/H/0588/003	OGYI-T-24080/03	FAES FARMA, S.A.	HU
Salcrozine 500 mg végbélkúp	ES/H/0588/003	OGYI-T-24080/04	FAES FARMA, S.A.	HU

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Salcrozine 500 mg végbélkúp	ES/H/0588/003	OGYI-T-24080/05	FAES FARMA, S.A.	HU
Salcrozine 500 mg végbélkúp	ES/H/0588/003	OGYI-T-24080/06	FAES FARMA, S.A.	HU
Salcrozine 500 mg végbélkúp	ES/H/0588/003	OGYI-T-24080/07	FAES FARMA, S.A.	HU
Salcrozine 500 mg Zäpfchen	PT/H/2281/001	2203438.00.00	FAES FARMA, S.A.	DE
Salo 4g/60ml Klysmen Rektalsuspension	DE/H/0363/005	19179.00.00	DR. FALK PHARMA G.M.B.H.	DE
Salofalk	DE/H/0363/005	44750	DR. FALK PHARMA G.M.B.H.	DK
Salofalk	DE/H/5681/001	43313	DR. FALK PHARMA G.M.B.H.	DK
Salofalk	DE/H/5682/001	43551	DR. FALK PHARMA G.M.B.H.	DK
Salofalk – Салофалк 500 mg	not available	9600067	DR. FALK PHARMA G.M.B.H.	BG
Salofalk 1 g čepići	not available	HR-H-096573039	DR. FALK PHARMA G.M.B.H.	HR
Salofalk 1 g čípky	DE/H/5682/001	29/586/10-C	DR. FALK PHARMA G.M.B.H.	CZ
Salofalk 1 g gyomornedv-ellenálló tabletta	DE/H/0363/008	OGYI-T-1612/19	DR. FALK PHARMA G.M.B.H.	HU
Salofalk 1 g peräpuikko	DE/H/5682/001	25173	DR. FALK PHARMA G.M.B.H.	FI
Salofalk 1 g stikkpille	DE/H/5682/001	08-6052	DR. FALK PHARMA G.M.B.H.	NO
Salofalk 1 g supozitoare	DE/H/5682/001	10973/2018/06	DR. FALK PHARMA G.M.B.H.	RO

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Salofalk 1 g supozitoare	DE/H/5682/001	10973/2018/05	DR. FALK PHARMA G.M.B.H.	RO
Salofalk 1 g supozitoare	DE/H/5682/001	10973/2018/04	DR. FALK PHARMA G.M.B.H.	RO
Salofalk 1 g supozitoare	DE/H/5682/001	10973/2018/03	DR. FALK PHARMA G.M.B.H.	RO
Salofalk 1 g supozitoare	DE/H/5682/001	10973/2018/01	DR. FALK PHARMA G.M.B.H.	RO
Salofalk 1 g supozitoare	DE/H/5682/001	10973/2018/07	DR. FALK PHARMA G.M.B.H.	RO
Salofalk 1 g supozitoare	DE/H/5682/001	10973/2018/02	DR. FALK PHARMA G.M.B.H.	RO
Salofalk 1 g supozitoriji	DE/H/5682/001	10-0300	DR. FALK PHARMA G.M.B.H.	LV
Salofalk 1 g suppositorier	DE/H/5682/001	27700	DR. FALK PHARMA G.M.B.H.	SE
Salofalk 1 g žvakutės	DE/H/5682/001	LT/1/10/2095/004	DR. FALK PHARMA G.M.B.H.	LT
Salofalk 1 g žvakutės	DE/H/5682/001	LT/1/10/2095/003	DR. FALK PHARMA G.M.B.H.	LT
Salofalk 1 g žvakutės	DE/H/5682/001	LT/1/10/2095/002	DR. FALK PHARMA G.M.B.H.	LT
Salofalk 1 g žvakutės	DE/H/5682/001	LT/1/10/2095/001	DR. FALK PHARMA G.M.B.H.	LT
Salofalk 1 g žvakutės	DE/H/5682/001	LT/1/10/2095/005	DR. FALK PHARMA G.M.B.H.	LT
Salofalk 1 g žvakutės	DE/H/5682/001	LT/1/10/2095/007	DR. FALK PHARMA G.M.B.H.	LT
Salofalk 1 g žvakutės	DE/H/5682/001	LT/1/10/2095/006	DR. FALK PHARMA G.M.B.H.	LT

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Salofalk 1 g γαστροανθεκτικά δισκία	DE/H/0363/008	023177	DR. FALK PHARMA G.M.B.H.	CY
Salofalk 1 g γαστροανθεκτικά δισκία Μεσαλαζίνη	DE/H/0363/008	1961813	GALENICA SA	GR
Salofalk 1 g υπόθετα	DE/H/5682/001	20945	DR. FALK PHARMA G.M.B.H.	CY
Salofalk 1 g Υπόθετα	DE/H/5682/001	1961811	GALENICA SA	GR
Salofalk 1 g, 1 g, czopki	DE/H/5682/001	17233	DR. FALK PHARMA G.M.B.H.	PL
Salofalk 1 g/dose rektalskum	DE/H/5681/001	08-5945	DR. FALK PHARMA G.M.B.H.	NO
Salofalk 1 g/pulvérisation mousse rectale	DE/H/5681/001	2003070008	DR. FALK PHARMA G.M.B.H.	LU
Salofalk 1 g/Sprühstoß Rektalschaum	DE/H/5681/001	1-24828	DR. FALK PHARMA G.M.B.H.	AT
Salofalk 1 g/εφαρμογή Ορθικός Αφρός.	DE/H/5681/001	1961810	GALENICA SA	GR
Salofalk 1,5 g depotgranulat i dosepose	DE/H/0363/004	07-4951	DR. FALK PHARMA G.M.B.H.	NO
Salofalk 1,5 g depotrakeet	DE/H/0363/004	23501	DR. FALK PHARMA G.M.B.H.	FI
Salofalk 1,5 g enterodepotgranulat	DE/H/0363/004	41316	DR. FALK PHARMA G.M.B.H.	DK
Salofalk 1,5 g granulado de liberación prolongada	DE/H/0363/004	70134	DR. FALK PHARMA G.M.B.H.	ES
Salofalk 1,5 g granulát s predĺženým uvoľňovaním	DE/H/0363/004	29/0383/13-S	DR. FALK PHARMA G.M.B.H.	SK

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Salofalk 1,5 g Granu-Stix, granulaat met verlengde afgifte	DE/H/0363/004	RVG 100059	DR. FALK PHARMA G.M.B.H.	NL
Salofalk 1,5 g ilgstošās darbības granulas	DE/H/0363/004	13-0205	DR. FALK PHARMA G.M.B.H.	LV
Salofalk 1,5 g pailginto atpalaidavimo granulės	DE/H/0363/004	LT/1/24/5316/001	DR. FALK PHARMA G.M.B.H.	LT
Salofalk 1,5 g pailginto atpalaidavimo granulės	DE/H/0363/004	LT/1/24/5316/002	DR. FALK PHARMA G.M.B.H.	LT
Salofalk 1,5 g pailginto atpalaidavimo granulės	DE/H/0363/004	LT/1/24/5316/003	DR. FALK PHARMA G.M.B.H.	LT
Salofalk 1,5 g pailginto atpalaidavimo granulės	DE/H/0363/004	LT/1/24/5316/004	DR. FALK PHARMA G.M.B.H.	LT
Salofalk 1,5 g pailginto atpalaidavimo granulės	DE/H/0363/004	LT/1/24/5316/005	DR. FALK PHARMA G.M.B.H.	LT
Salofalk 1,5 g pailginto atpalaidavimo granulės	DE/H/0363/004	LT/1/24/5316/006	DR. FALK PHARMA G.M.B.H.	LT
Salofalk 1,5 g pailginto atpalaidavimo granulės	DE/H/0363/004	LT/1/24/5316/007	DR. FALK PHARMA G.M.B.H.	LT
Salofalk 1,5 g pailginto atpalaidavimo granulės	DE/H/0363/004	LT/1/24/5316/008	DR. FALK PHARMA G.M.B.H.	LT
Salofalk 1,5 g pailginto atpalaidavimo granulės	DE/H/0363/004	LT/1/24/5316/009	DR. FALK PHARMA G.M.B.H.	LT
Salofalk 1,5 g pailginto atpalaidavimo granulės	DE/H/0363/004	LT/1/24/5316/010	DR. FALK PHARMA G.M.B.H.	LT
Salofalk 1,5 g retard granulátum	DE/H/0363/004	OGYI-T-1612/18	DR. FALK PHARMA G.M.B.H.	HU
Salofalk 1,5 g retard granulátum	DE/H/0363/004	OGYI-T-1612/14	DR. FALK PHARMA G.M.B.H.	HU
Salofalk 1,5 g retard granulátum	DE/H/0363/004	OGYI-T-1612/13	DR. FALK PHARMA G.M.B.H.	HU

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Salofalk 1,5 g retard granulátum	DE/H/0363/004	OGYI-T-1612/12	DR. FALK PHARMA G.M.B.H.	HU
Salofalk 1.5 g prolonged-release granules	DE/H/0363/004	PA 0573/003/007	DR. FALK PHARMA G.M.B.H.	IE
Salofalk 1.5g prolonged-release granules	DE/H/0363/004	PL 08637/0016	DR. FALK PHARMA G.M.B.H.	XI
Salofalk 1000 mg - Zäpfchen	DE/H/5682/001	1-29764	DR. FALK PHARMA G.M.B.H.	AT
Salofalk 1000 mg depotgranulat	DE/H/0363/002	18622	DR. FALK PHARMA G.M.B.H.	SE
Salofalk 1000 mg depotgranulat i dosepose	DE/H/0363/002	02-1275	DR. FALK PHARMA G.M.B.H.	NO
Salofalk 1000 mg depotrakeet	DE/H/0363/002	17474	DR. FALK PHARMA G.M.B.H.	FI
Salofalk 1000 mg enterodepotgranulat	DE/H/0363/002	33924	DR. FALK PHARMA G.M.B.H.	DK
Salofalk 1000 mg granulado de liberación prolongada	DE/H/0363/002	65772	DR. FALK PHARMA G.M.B.H.	ES
Salofalk 1000 mg granulát s predĺženým uvoľňovaním	DE/H/0363/002	29/0382/13-S	DR. FALK PHARMA G.M.B.H.	SK
Salofalk 1000 mg granule s prodlouženým uvolňováním	DE/H/0363/002	29/031/14-C	DR. FALK PHARMA G.M.B.H.	CZ
Salofalk 1000 mg Granu-Stix, granulaat met verlengde afgifte	DE/H/0363/002	RVG 28131	DR. FALK PHARMA G.M.B.H.	NL
Salofalk 1000 mg ilgstošās darbības granulas	DE/H/0363/002	13-0204	DR. FALK PHARMA G.M.B.H.	LV

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Salofalk 1000 mg pailginto atpalaidavimo granulės	DE/H/0363/002	LT/1/24/5316/001	DR. FALK PHARMA G.M.B.H.	LT
Salofalk 1000 mg pailginto atpalaidavimo granulės	DE/H/0363/002	LT/1/24/5315/002	DR. FALK PHARMA G.M.B.H.	LT
Salofalk 1000 mg pailginto atpalaidavimo granulės	DE/H/0363/002	LT/1/24/5315/003	DR. FALK PHARMA G.M.B.H.	LT
Salofalk 1000 mg pailginto atpalaidavimo granulės	DE/H/0363/002	LT/1/24/5315/004	DR. FALK PHARMA G.M.B.H.	LT
Salofalk 1000 mg pailginto atpalaidavimo granulės	DE/H/0363/002	LT/1/24/5315/005	DR. FALK PHARMA G.M.B.H.	LT
Salofalk 1000 mg prolonged release granules	DE/H/0363/002	PA 0573/003/002	DR. FALK PHARMA G.M.B.H.	IE
Salofalk 1000 mg retard granulátum	DE/H/0363/002	OGYI-T-1612/10	DR. FALK PHARMA G.M.B.H.	HU
Salofalk 1000 mg retard granulátum	DE/H/0363/002	OGYI-T-1612/11	DR. FALK PHARMA G.M.B.H.	HU
Salofalk 1000 mg svečke	DE/H/5682/001	H/11/02031/007	DR. FALK PHARMA G.M.B.H.	SI
Salofalk 1000 mg svečke	DE/H/5682/001	H/11/02031/006	DR. FALK PHARMA G.M.B.H.	SI
Salofalk 1000 mg svečke	DE/H/5682/001	H/11/02031/005	DR. FALK PHARMA G.M.B.H.	SI
Salofalk 1000 mg svečke	DE/H/5682/001	H/11/02031/004	DR. FALK PHARMA G.M.B.H.	SI
Salofalk 1000 mg svečke	DE/H/5682/001	H/11/02031/003	DR. FALK PHARMA G.M.B.H.	SI
Salofalk 1000 mg svečke	DE/H/5682/001	H/11/02031/002	DR. FALK PHARMA G.M.B.H.	SI
Salofalk 1000 mg svečke	DE/H/5682/001	H/11/02031/001	DR. FALK PHARMA G.M.B.H.	SI

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Salofalk 1000 mg želučanootporne granule	not available	HR-H-293118992	DR. FALK PHARMA G.M.B.H.	HR
Salofalk 1000 mg zrnca s podaljšanim sproščanjem	DE/H/0363/002	H/04/01392/004	DR. FALK PHARMA G.M.B.H.	SI
Salofalk 1000 mg zrnca s podaljšanim sproščanjem	DE/H/0363/002	H/04/01392/005	DR. FALK PHARMA G.M.B.H.	SI
Salofalk 1000 mg zrnca s podaljšanim sproščanjem	DE/H/0363/002	H/04/01392/006	DR. FALK PHARMA G.M.B.H.	SI
Salofalk 1000 mg zrnca s podaljšanim sproščanjem	DE/H/0363/002	H/04/01392/007	DR. FALK PHARMA G.M.B.H.	SI
Salofalk 1000 mg zrnca s podaljšanim sproščanjem	DE/H/0363/002	H/04/01392/008	DR. FALK PHARMA G.M.B.H.	SI
Salofalk 1000mg prolonged-release granules	DE/H/0363/002	PL 08637/0008	DR. FALK PHARMA G.M.B.H.	XI
Salofalk 1500 mg depotgranulat	DE/H/0363/004	25474	DR. FALK PHARMA G.M.B.H.	SE
Salofalk 1500 mg granule s prodlouženým uvolňováním	DE/H/0363/004	29/032/14-C	DR. FALK PHARMA G.M.B.H.	CZ
Salofalk 1500 mg zrnca s podaljšanim sproščanjem	DE/H/0363/004	H/04/01392/009	DR. FALK PHARMA G.M.B.H.	SI
Salofalk 1500 mg zrnca s podaljšanim sproščanjem	DE/H/0363/004	H/04/01392/010	DR. FALK PHARMA G.M.B.H.	SI
Salofalk 1500 mg zrnca s podaljšanim sproščanjem	DE/H/0363/004	H/04/01392/018	DR. FALK PHARMA G.M.B.H.	SI
Salofalk 1500 mg zrnca s podaljšanim sproščanjem	DE/H/0363/004	H/04/01392/017	DR. FALK PHARMA G.M.B.H.	SI
Salofalk 1500 mg zrnca s podaljšanim sproščanjem	DE/H/0363/004	H/04/01392/016	DR. FALK PHARMA G.M.B.H.	SI
Salofalk 1500 mg zrnca s podaljšanim sproščanjem	DE/H/0363/004	H/04/01392/015	DR. FALK PHARMA G.M.B.H.	SI

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Salofalk 1500 mg zrnca s podaljšanim sproščanjem	DE/H/0363/004	H/04/01392/014	DR. FALK PHARMA G.M.B.H.	SI
Salofalk 1500 mg zrnca s podaljšanim sproščanjem	DE/H/0363/004	H/04/01392/013	DR. FALK PHARMA G.M.B.H.	SI
Salofalk 1500 mg zrnca s podaljšanim sproščanjem	DE/H/0363/004	H/04/01392/012	DR. FALK PHARMA G.M.B.H.	SI
Salofalk 1500 mg zrnca s podaljšanim sproščanjem	DE/H/0363/004	H/04/01392/011	DR. FALK PHARMA G.M.B.H.	SI
Salofalk 1g čapíky	DE/H/5682/001	73/0715/10-S	DR. FALK PHARMA G.M.B.H.	SK
Salofalk 1g gastro-resistant tablets	DE/H/0363/008	PL 08637/0027	DR. FALK PHARMA G.M.B.H.	XI
Salofalk 1g gyomornedv-ellenálló tabletta	DE/H/0363/008	OGYI-T-1612/20	DR. FALK PHARMA G.M.B.H.	HU
Salofalk 1g magensaftresistente Tabletten	DE/H/0363/008	96061.00.00	DR. FALK PHARMA G.M.B.H.	DE
Salofalk 1g Rektalschaum	DE/H/5681/001	65717.00.00	DR. FALK PHARMA G.M.B.H.	DE
Salofalk 1g supositorios	DE/H/5682/001	73372	DR. FALK PHARMA G.M.B.H.	ES
Salofalk 1g supositórios	DE/H/5682/001	5296371	DR. FALK PHARMA G.M.B.H.	PT
Salofalk 1g Suppositorien	DE/H/5682/001	74251.00.00	DR. FALK PHARMA G.M.B.H.	DE
Salofalk 1g suppositories	DE/H/5682/001	PA 573/4/4	DR. FALK PHARMA G.M.B.H.	IE
Salofalk 1g Suppositories	DE/H/5682/001	PL 08637/0018	DR. FALK PHARMA G.M.B.H.	XI
Salofalk 1g végbélkúp	DE/H/5682/001	OGYI-T-1612/05	DR. FALK PHARMA G.M.B.H.	HU

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Salofalk 1g, zetpillen	DE/H/5682/001	RVG 102870	DR. FALK PHARMA G.M.B.H.	NL
Salofalk 1g/actuation rectal foam	DE/H/5681/001	PA 573/4/5	DR. FALK PHARMA G.M.B.H.	IE
Salofalk 1g/actuation Rectal Foam	DE/H/5681/001	PL 08637/0003	DR. FALK PHARMA G.M.B.H.	XI
Salofalk 1g/aplicação Espuma rectal	DE/H/5681/001	4275582	DR. FALK PHARMA G.M.B.H.	PT
Salofalk 1g/puff rectal foam	DE/H/5681/001	27046	DR. FALK PHARMA G.M.B.H.	SE
Salofalk 2 g/30 g, klysma	not available	RVG 15393	DR. FALK PHARMA BENELUX B.V.	NL
Salofalk 2 g/30 ml végbélszuszpenzió	not available	OGYI-T-1612/06	DR. FALK PHARMA G.M.B.H.	HU
Salofalk 2 g/60 g, klysma	not available	RVG 15845	DR. FALK PHARMA BENELUX B.V.	NL
Salofalk 250 čapíky	not available	29/0027/87-S	DR. FALK PHARMA G.M.B.H.	SK
Salofalk 250 mg bélben oldódó tabletta	not available	OGYI-T-1612/01	DR. FALK PHARMA G.M.B.H.	HU
Salofalk 250 mg čepići	not available	HR-H-717303526	DR. FALK PHARMA G.M.B.H.	HR
Salofalk 250 mg comprimate gastrorezistente	not available	11532/2019/02	DR. FALK PHARMA G.M.B.H.	RO
Salofalk 250 mg comprimate gastrorezistente	not available	11532/2019/01	DR. FALK PHARMA G.M.B.H.	RO
SALOFALK 250 mg comprimidos gastrorresistentes	not available	9639906	DR. FALK PHARMA PORTUGAL	PT
SALOFALK 250 mg comprimidos gastrorresistentes	not available	9639914	DR. FALK PHARMA PORTUGAL	PT

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Salofalk 250 mg gastrorezistentne tablete	not available	H/00/02032/001	DR. FALK PHARMA G.M.B.H.	SI
Salofalk 250 mg gastrorezistentne tablete	not available	H/00/02032/002	DR. FALK PHARMA G.M.B.H.	SI
SALOFALK 250 mg supositórios	not available	9639807	DR. FALK PHARMA PORTUGAL	PT
Salofalk 250 mg supozitoare	not available	11534/2019/01	DR. FALK PHARMA G.M.B.H.	RO
Salofalk 250 mg supozitoare	not available	11534/2019/02	DR. FALK PHARMA G.M.B.H.	RO
Salofalk 250 mg svečke	not available	H/00/02032/006	DR. FALK PHARMA G.M.B.H.	SI
Salofalk 250 mg svečke	not available	H/00/02032/005	DR. FALK PHARMA G.M.B.H.	SI
Salofalk 250 mg végbélkúp	not available	OGYI-T-1612/03	DR. FALK PHARMA G.M.B.H.	HU
Salofalk 250 mg γαστροανθεκτικά δισκία	not available	1961801	GALENICA SA	GR
Salofalk 250 mg υπόθετα	not available	1961803	GALENICA SA	GR
Salofalk 250 tablety	not available	29/0026/87-S	DR. FALK PHARMA G.M.B.H.	SK
Salofalk 250, maagsapresistente tabletten 250 mg	not available	RVG 11086	DR. FALK PHARMA BENELUX B.V.	NL
Salofalk 250mg gastro-resistant tablets	not available	PA 573/4/3	DR. FALK PHARMA G.M.B.H.	IE
Salofalk 250mg Suppositorien	not available	2881.00.00	DR. FALK PHARMA G.M.B.H.	DE
Salofalk 250mg suppositories	not available	PA 573/4/2	DR. FALK PHARMA G.M.B.H.	IE

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Salofalk 250mg Tablets	not available	PL 10341/0004	DR. FALK PHARMA UK LTD	XI
Salofalk 250mg, magensaftresistente Tabletten	not available	2881.00.01	DR. FALK PHARMA G.M.B.H.	DE
Salofalk 2g/30g Klysmen	not available	29/0179/92-C/S	DR. FALK PHARMA G.M.B.H.	SK
Salofalk 2g/30ml Klysmen Rektalsuspension	not available	16629.00.00	DR. FALK PHARMA G.M.B.H.	DE
Salofalk 3 g depotgranulat i dosepose	DE/H/0363/007	10-7491	DR. FALK PHARMA G.M.B.H.	NO
Salofalk 3 g depotrakeet	DE/H/0363/007	28672	DR. FALK PHARMA G.M.B.H.	FI
Salofalk 3 g enterodepotgranulat	DE/H/0363/007	46737	DR. FALK PHARMA G.M.B.H.	DK
Salofalk 3 g granulado de liberación prolongada	DE/H/0363/007	74791	DR. FALK PHARMA G.M.B.H.	ES
Salofalk 3 g granulát s predĺženým uvoľňovaním	DE/H/0363/007	29/0384/13-S	DR. FALK PHARMA G.M.B.H.	SK
Salofalk 3 g Granu-Stix, granulaat met verlengde afgifte	DE/H/0363/007	RVG 107302	DR. FALK PHARMA G.M.B.H.	NL
Salofalk 3 g ilgstošās darbības granulas	DE/H/0363/007	13-0206	DR. FALK PHARMA G.M.B.H.	LV
Salofalk 3 g pailginto atpalaidavimo granulės	DE/H/0363/007	LT/1/24/5317/001	DR. FALK PHARMA G.M.B.H.	LT
Salofalk 3 g pailginto atpalaidavimo granulės	DE/H/0363/007	LT/1/24/5317/002	DR. FALK PHARMA G.M.B.H.	LT
Salofalk 3 g pailginto atpalaidavimo granulės	DE/H/0363/007	LT/1/24/5317/003	DR. FALK PHARMA G.M.B.H.	LT
Salofalk 3 g pailginto atpalaidavimo granulės	DE/H/0363/007	LT/1/24/5317/004	DR. FALK PHARMA G.M.B.H.	LT

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Salofalk 3 g pailginto atpalaidavimo granulės	DE/H/0363/007	LT/1/24/5317/005	DR. FALK PHARMA G.M.B.H.	LT
Salofalk 3 g pailginto atpalaidavimo granulės	DE/H/0363/007	LT/1/24/5317/006	DR. FALK PHARMA G.M.B.H.	LT
Salofalk 3 g pailginto atpalaidavimo granulės	DE/H/0363/007	LT/1/24/5317/007	DR. FALK PHARMA G.M.B.H.	LT
Salofalk 3 g pailginto atpalaidavimo granulės	DE/H/0363/007	LT/1/24/5317/008	DR. FALK PHARMA G.M.B.H.	LT
Salofalk 3 g prolonged-release granules	DE/H/0363/007	PA 0573/003/006	DR. FALK PHARMA G.M.B.H.	IE
Salofalk 3 g retard granulátum	DE/H/0363/007	OGYI-T-1612/15	DR. FALK PHARMA G.M.B.H.	HU
Salofalk 3 g retard granulátum	DE/H/0363/007	OGYI-T-1612/16	DR. FALK PHARMA G.M.B.H.	HU
Salofalk 3 g retard granulátum	DE/H/0363/007	OGYI-T-1612/17	DR. FALK PHARMA G.M.B.H.	HU
Salofalk 3000 mg depotgranulat	DE/H/0363/007	44129	DR. FALK PHARMA G.M.B.H.	SE
Salofalk 3000 mg zrnca s podaljšanim sproščanjem	DE/H/0363/007	H/04/01392/019	DR. FALK PHARMA G.M.B.H.	SI
Salofalk 3000 mg zrnca s podaljšanim sproščanjem	DE/H/0363/007	H/04/01392/020	DR. FALK PHARMA G.M.B.H.	SI
Salofalk 3000 mg zrnca s podaljšanim sproščanjem	DE/H/0363/007	H/04/01392/026	DR. FALK PHARMA G.M.B.H.	SI
Salofalk 3000 mg zrnca s podaljšanim sproščanjem	DE/H/0363/007	H/04/01392/025	DR. FALK PHARMA G.M.B.H.	SI
Salofalk 3000 mg zrnca s podaljšanim sproščanjem	DE/H/0363/007	H/04/01392/024	DR. FALK PHARMA G.M.B.H.	SI
Salofalk 3000 mg zrnca s podaljšanim sproščanjem	DE/H/0363/007	H/04/01392/023	DR. FALK PHARMA G.M.B.H.	SI

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Salofalk 3000 mg zrnca s podaljšanim sproščanjem	DE/H/0363/007	H/04/01392/022	DR. FALK PHARMA G.M.B.H.	SI
Salofalk 3000 mg zrnca s podaljšanim sproščanjem	DE/H/0363/007	H/04/01392/021	DR. FALK PHARMA G.M.B.H.	SI
Salofalk 3000 mggranule s prodlouženým uvolňováním	DE/H/0363/007	29/033/14-C	DR. FALK PHARMA G.M.B.H.	CZ
Salofalk 3g prolonged-release granules	DE/H/0363/007	PL 08637/0025	DR. FALK PHARMA G.M.B.H.	XI
Salofalk 4 g - Klysmen	not available	1-19003	DR. FALK PHARMA G.M.B.H.	AT
Salofalk 4 g rektalna suspenzija	not available	H/00/02032/009	DR. FALK PHARMA G.M.B.H.	SI
Salofalk 4 g/60 g, klysma	not available	RVG 11469	DR. FALK PHARMA BENELUX B.V.	NL
Salofalk 4 g/60 ml tiesiosios žarnos suspensija	not available	LT/1/95/0445/005	DR. FALK PHARMA G.M.B.H.	LT
Salofalk 4 g/60 mL enemas	not available	12246	DR. FALK PHARMA G.M.B.H.	CY
Salofalk 4 g/60 ml ορθικό εναιώρημα	not available	1961802	GALENICA SA	GR
Salofalk 4 g/60 ml peräruiskesuspensio	DE/H/0363/005	27285	DR. FALK PHARMA G.M.B.H.	FI
Salofalk 4 g/60 ml rektālā suspensija	not available	99-0935	DR. FALK PHARMA G.M.B.H.	LV
Salofalk 4 g/60 ml rektalvæske, suspensjon	DE/H/0363/005	09-6598	DR. FALK PHARMA G.M.B.H.	NO
Salofalk 4 g/60 ml végbélszuszpenzió	not available	OGYI-T-1612/07	DR. FALK PHARMA G.M.B.H.	HU
Salofalk 4g/60g Klysmen	not available	29/0229/13-S	DR. FALK PHARMA G.M.B.H.	SK

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Salofalk 4g/60ml enema	not available	PA 573/4/1	DR. FALK PHARMA G.M.B.H.	IE
Salofalk 4g/60ml Klysmen Rektalsuspension	not available	2881.00.02	DR. FALK PHARMA G.M.B.H.	DE
Salofalk 4g/60ml rektalsuspension	DE/H/0363/005	42221	DR. FALK PHARMA G.M.B.H.	SE
Salofalk 4g/60ml suspensión rectal	DE/H/0363/005	71881	DR. FALK PHARMA G.M.B.H.	ES
Salofalk 4g/60ml suspenzija za rektum	not available	HR-H-371400988	DR. FALK PHARMA G.M.B.H.	HR
Salofalk 500 čapíky	not available	29/0228/13-S	DR. FALK PHARMA G.M.B.H.	SK
Salofalk 500 mg - Zäpfchen	not available	1-19005	DR. FALK PHARMA G.M.B.H.	AT
Salofalk 500 mg bélben oldódó tabletta	not available	OGYI-T-1612/02	DR. FALK PHARMA G.M.B.H.	HU
Salofalk 500 mg čepići	not available	HR-H-366666368	DR. FALK PHARMA G.M.B.H.	HR
Salofalk 500 mg čípky	not available	29/273/98-C	DR. FALK PHARMA G.M.B.H.	CZ
Salofalk 500 mg comprimate gastrorezistente	not available	11533/2019/02	DR. FALK PHARMA G.M.B.H.	RO
Salofalk 500 mg comprimate gastrorezistente	not available	11533/2019/01	DR. FALK PHARMA G.M.B.H.	RO
SALOFALK 500 mg comprimidos gastrorresistentes	not available	9639930	DR. FALK PHARMA PORTUGAL	PT
Salofalk 500 mg depotgranulat	DE/H/0363/001	18621	DR. FALK PHARMA G.M.B.H.	SE
Salofalk 500 mg depotgranulat i dosepose	DE/H/0363/001	02-1274	DR. FALK PHARMA G.M.B.H.	NO

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Salofalk 500 mg depotrakeet	DE/H/0363/001	17473	DR. FALK PHARMA G.M.B.H.	FI
Salofalk 500 mg enterodepotgranulat	DE/H/0363/001	33923	DR. FALK PHARMA G.M.B.H.	DK
Salofalk 500 mg gastrorezistentne tablete	not available	H/00/02032/003	DR. FALK PHARMA G.M.B.H.	SI
Salofalk 500 mg gastrorezistentne tablete	not available	H/00/02032/004	DR. FALK PHARMA G.M.B.H.	SI
Salofalk 500 mg granulado de liberación prolongada	DE/H/0363/001	65771	DR. FALK PHARMA G.M.B.H.	ES
Salofalk 500 mg granulát s predĺženým uvoľňovaním	DE/H/0363/001	29/0381/13-S	DR. FALK PHARMA G.M.B.H.	SK
Salofalk 500 mg granule s prodlouženým uvolňováním	DE/H/0363/001	29/030/14-C	DR. FALK PHARMA G.M.B.H.	CZ
Salofalk 500 mg Granu-Stix, granulaat met verlengde afgifte	DE/H/0363/001	RVG 28130	DR. FALK PHARMA G.M.B.H.	NL
Salofalk 500 mg ilgstošās darbības granulas	DE/H/0363/001	13-0203	DR. FALK PHARMA G.M.B.H.	LV
Salofalk 500 mg magensaftresistente Tabletten	not available	1-19004	DR. FALK PHARMA G.M.B.H.	AT
Salofalk 500 mg pailginto atpalaidavimo granulės	DE/H/0363/001	LT/1/24/5314/002	DR. FALK PHARMA G.M.B.H.	LT
Salofalk 500 mg pailginto atpalaidavimo granulės	DE/H/0363/001	LT/1/24/5314/001	DR. FALK PHARMA G.M.B.H.	LT
Salofalk 500 mg pailginto atpalaidavimo granulės	DE/H/0363/001	LT/1/24/5314/003	DR. FALK PHARMA G.M.B.H.	LT
Salofalk 500 mg prolonged release granules	DE/H/0363/001	PA 0573/003/001	DR. FALK PHARMA G.M.B.H.	IE

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Salofalk 500 mg retard granulátum	DE/H/0363/001	OGYI-T-1612/08	DR. FALK PHARMA G.M.B.H.	HU
Salofalk 500 mg retard granulátum	DE/H/0363/001	OGYI-T-1612/09	DR. FALK PHARMA G.M.B.H.	HU
Salofalk 500 mg skrandyje neirios tabletės	not available	LT/1/95/0445/002	DR. FALK PHARMA G.M.B.H.	LT
SALOFALK 500 mg Supositórios	not available	9639815	DR. FALK PHARMA PORTUGAL	PT
Salofalk 500 mg supozitoare	not available	11535/2019/01	DR. FALK PHARMA G.M.B.H.	RO
Salofalk 500 mg supozitoare	not available	11535/2019/02	DR. FALK PHARMA G.M.B.H.	RO
Salofalk 500 mg supozitoriji	not available	99-0383	DR. FALK PHARMA G.M.B.H.	LV
Salofalk 500 mg suppositories	not available	13057	DR. FALK PHARMA G.M.B.H.	CY
Salofalk 500 mg svečke	not available	H/00/02032/007	DR. FALK PHARMA G.M.B.H.	SI
Salofalk 500 mg svečke	not available	H/00/02032/008	DR. FALK PHARMA G.M.B.H.	SI
Salofalk 500 mg végbélkúp	not available	OGYI-T-1612/04	DR. FALK PHARMA G.M.B.H.	HU
Salofalk 500 mg zarnās šķīstošā tablete	not available	99-0382	DR. FALK PHARMA G.M.B.H.	LV
Salofalk 500 mg želučanootporne granule	not available	HR-H-842627661	DR. FALK PHARMA G.M.B.H.	HR
Salofalk 500 mg želučanootporne tablete	not available	HR-H-056926689	DR. FALK PHARMA G.M.B.H.	HR
Salofalk 500 mg zrnca s podaljšanim sproščanjem	DE/H/0363/001	H/04/01392/001	DR. FALK PHARMA G.M.B.H.	SI

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Salofalk 500 mg zrnca s podaljšanim sproščanjem	DE/H/0363/001	H/04/01392/002	DR. FALK PHARMA G.M.B.H.	SI
Salofalk 500 mg zrnca s podaljšanim sproščanjem	DE/H/0363/001	H/04/01392/003	DR. FALK PHARMA G.M.B.H.	SI
Salofalk 500 mg žvakutės	not available	LT/1/95/0445/004	DR. FALK PHARMA G.M.B.H.	LT
Salofalk 500 mg γαστροανθεκτικά δισκία	not available	1961804	GALENICA SA	GR
Salofalk 500 mg υπόθετα	not available	1961805	GALENICA SA	GR
Salofalk 500 tablety	not available	29/0227/13-S	DR. FALK PHARMA G.M.B.H.	SK
SALOFALK 500, 500 mg, czopki	not available	4222	DR. FALK PHARMA G.M.B.H.	PL
SALOFALK 500, 500 mg, tabletki dojelitowe	not available	4001	DR. FALK PHARMA G.M.B.H.	PL
Salofalk 500, enterosolventní tablety	not available	29/207/98-C	DR. FALK PHARMA G.M.B.H.	CZ
Salofalk 500, maagsapresistente tabletten 500 mg	not available	RVG 12144	DR. FALK PHARMA BENELUX B.V.	NL
Salofalk 500mg gastro-resistant tablets	DE/H/0363/006	PL 08637/0019	DR. FALK PHARMA G.M.B.H.	XI
Salofalk 500mg prolonged-release granules	DE/H/0363/001	PL 08637/0007	DR. FALK PHARMA G.M.B.H.	XI
Salofalk 500mg Suppositorien Mesalazin	not available	2881.01.00	DR. FALK PHARMA G.M.B.H.	DE
Salofalk 500mg, gastro-resistant tablets	not available	13056	DR. FALK PHARMA G.M.B.H.	CY
Salofalk 500mg, magensaftresistente Tabletten	not available	2881.01.01	DR. FALK PHARMA G.M.B.H.	DE

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Salofalk Enema 2g	not available	PL 10341/0008	DR. FALK PHARMA UK LTD	XI
SALOFALK enema 4 g/60 ml Suspensão retal	not available	8640003	DR. FALK PHARMA PORTUGAL	PT
SALOFALK enema 4 g/60 ml Suspensão retal	not available	5851993	DR. FALK PHARMA PORTUGAL	PT
SALOFALK enema 4 g/60 ml Suspensão retal	not available	4352886	DR. FALK PHARMA PORTUGAL	PT
Salofalk granu stix 1,5 g κοκκία παρατεταμένης αποδέσμευσης	DE/H/0363/004	1961809	GALENICA SA	GR
Salofalk granu stix 1000 mg κοκκία παρατεταμένης αποδέσμευσης	DE/H/0363/002	1961807	GALENICA SA	GR
Salofalk granu stix 3 g κοκκία παρατεταμένης αποδέσμευσης	DE/H/0363/007	1961812	GALENICA SA	GR
Salofalk granu stix 500 mg κοκκία παρατεταμένης αποδέσμευσης	DE/H/0363/001	1961806	GALENICA SA	GR
Salofalk Grânulos, 1,5 g granulado de libertação prolongada	DE/H/0363/004	5132535	DR. FALK PHARMA G.M.B.H.	PT
Salofalk Grânulos, 1,5 g granulado de libertação prolongada	DE/H/0363/004	5132543	DR. FALK PHARMA G.M.B.H.	PT
Salofalk Grânulos, 1000 mg granulado de libertação prolongada	DE/H/0363/002	4213088	DR. FALK PHARMA G.M.B.H.	PT

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Salofalk Grânulos, 1000 mg granulado de libertação prolongada	DE/H/0363/002	4213187	DR. FALK PHARMA G.M.B.H.	PT
Salofalk Grânulos, 1000 mg granulado de libertação prolongada	DE/H/0363/002	4213286	DR. FALK PHARMA G.M.B.H.	PT
Salofalk Grânulos, 1000 mg granulado de libertação prolongada	DE/H/0363/002	5104245	DR. FALK PHARMA G.M.B.H.	PT
Salofalk Grânulos, 1000 mg granulado de libertação prolongada	DE/H/0363/002	5107735	DR. FALK PHARMA G.M.B.H.	PT
Salofalk Grânulos, 3 g granulado de libertação prolongada	DE/H/0363/007	5653464	DR. FALK PHARMA G.M.B.H.	PT
Salofalk Grânulos, 3 g granulado de libertação prolongada	DE/H/0363/007	5398433	DR. FALK PHARMA G.M.B.H.	PT
Salofalk Grânulos, 500 mg granulado de libertação prolongada	DE/H/0363/001	4212783	DR. FALK PHARMA G.M.B.H.	PT
Salofalk Grânulos, 500 mg granulado de libertação prolongada	DE/H/0363/001	4212882	DR. FALK PHARMA G.M.B.H.	PT
Salofalk Grânulos, 500 mg granulado de libertação prolongada	DE/H/0363/001	4212981	DR. FALK PHARMA G.M.B.H.	PT
Salofalk Granu-Stix 1,5 g κοκκία παρατεταμένης αποδέσμευσης	DE/H/0363/004	023925	DR. FALK PHARMA G.M.B.H.	CY

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Salofalk Granu-Stix 1,5g Retardgranulat	DE/H/0363/004	69313.00.00	DR. FALK PHARMA G.M.B.H.	DE
Salofalk Granu-Stix 1000 mg κοκκία παρατεταμένης αποδέσμευσης	DE/H/0363/002	023923	DR. FALK PHARMA G.M.B.H.	CY
Salofalk Granu-Stix 1000mg Retardgranulat	DE/H/0363/002	50315.01.00	DR. FALK PHARMA G.M.B.H.	DE
Salofalk Granu-Stix 3 g κοκκία παρατεταμένης αποδέσμευσης	DE/H/0363/007	023925	DR. FALK PHARMA G.M.B.H.	CY
Salofalk Granu-Stix 3g Retardgranulat	DE/H/0363/007	81709.00.00	DR. FALK PHARMA G.M.B.H.	DE
Salofalk Granu-Stix 500 mg κοκκία παρατεταμένης αποδέσμευσης	DE/H/0363/001	023922	DR. FALK PHARMA G.M.B.H.	CY
Salofalk Granu-Stix 500mg Retardgranulat	DE/H/0363/001	50315.00.00	DR. FALK PHARMA G.M.B.H.	DE
Salofalk Schuim, 1g/dosis schuim voor rectaal gebruik	DE/H/5681/001	RVG 28179	DR. FALK PHARMA G.M.B.H.	NL
Salofalk Suppositories 500mg	not available	PL 10341/0009	DR. FALK PHARMA UK LTD	XI
Salofalk zetpil 250, zetpillen 250 mg	not available	RVG 10115	DR. FALK PHARMA BENELUX B.V.	NL
Salofalk zetpil 500, zetpillen 500 mg	not available	RVG 11836	DR. FALK PHARMA BENELUX B.V.	NL
Salofalk, 1,5 g, granulat o przedłużonym uwalnianiu	DE/H/0363/004	21830	DR. FALK PHARMA G.M.B.H.	PL
Salofalk, 1000 mg, granulat o przedłużonym	DE/H/0363/002	21829	DR. FALK PHARMA G.M.B.H.	PL

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uwalnianiu				
Salofalk, 3 g, granulat o przedłużonym uwalnianiu	DE/H/0363/007	21831	DR. FALK PHARMA G.M.B.H.	PL
SALOFALK, 4 g/60 ml, suspensie rectală	not available	11536/2019/01	DR. FALK PHARMA G.M.B.H.	RO
Salofalk, 4 g/60 ml, zawiesina doodbytnicza	not available	9701	DR. FALK PHARMA G.M.B.H.	PL
Salofalk® 1 g/painallus rektaalivaahto	DE/H/5681/001	25076	DR. FALK PHARMA G.M.B.H.	FI
Salogran-Falk 1000mg Retardgranulat	not available	52808.01.00	DR. FALK PHARMA G.M.B.H.	DE
Salogran-Falk 500mg Retardgranulat	not available	52808.00.00	DR. FALK PHARMA G.M.B.H.	DE
SALOZINAL 500 mg čípky	not available	29/688/12-C	PRO.MED.CS PRAHA A.S.	CZ
Xalazin 500 mg gyomornedv-ellenálló tabletta	not available	OGYI-T-9202/01	ASTELLAS PHARMA KFT	HU
Xalazin 500 mg végbélkúp	not available	OGYI-T-9204/01	ASTELLAS PHARMA KFT	HU
Yaldigo 1 g rektaalsuposiidid	SE/H/2039/001	1020021	TILLOTTS PHARMA AB	EE
Yaldigo 1 g rektaalsuposiidid	SE/H/2039/001	1020021	TILLOTTS PHARMA AB	EE
Yaldigo 1 g supozitoriji	SE/H/2039/001	21-0007	TILLOTTS PHARMA AB	LV
Yaldigo 1 g supozitoriji	SE/H/2039/001	21-0007	TILLOTTS PHARMA AB	LV
Yaldigo 1 g žvakutės	SE/H/2039/001	LT/1/21/4726/001	TILLOTTS PHARMA AB	LT

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Yaldigo 1 g žvakutės	SE/H/2039/001	LT/1/21/4726/002	TILLOTTS PHARMA AB	LT
Yaldigo 1 g žvakutės	SE/H/2039/001	LT/1/21/4726/003	TILLOTTS PHARMA AB	LT
Yaldigo 1 g žvakutės	SE/H/2039/001	LT/1/21/4726/004	TILLOTTS PHARMA AB	LT
Yaldigo 1 g žvakutės	SE/H/2039/001	LT/1/21/4726/005	TILLOTTS PHARMA AB	LT
Yaldigo 1 g žvakutės	SE/H/2039/001	LT/1/21/4726/001	TILLOTTS PHARMA AB	LT
Yaldigo 1 g žvakutės	SE/H/2039/001	LT/1/21/4726/002	TILLOTTS PHARMA AB	LT
Yaldigo 1 g žvakutės	SE/H/2039/001	LT/1/21/4726/003	TILLOTTS PHARMA AB	LT
Yaldigo 1 g žvakutės	SE/H/2039/001	LT/1/21/4726/004	TILLOTTS PHARMA AB	LT
Yaldigo 1 g žvakutės	SE/H/2039/001	LT/1/21/4726/005	TILLOTTS PHARMA AB	LT
Yaldigo 1600 mg modificētās darbības tabletes	SE/H/1654/001	18-0046	TILLOTTS PHARMA AB	LV
Yaldigo 1600 mg Tabletten mit veränderter Wirkstofffreisetzung	SE/H/1654/001	138193	TILLOTTS PHARMA GMBH	AT
Yaldigo 1600 mg δισκία ελεγχόμενης αποδέσμευσης	SE/H/1654/001	95174/02-10-2020	TILLOTTS PHARMA GMBH	GR
Yaldigo 1600 mg δισκία ελεγχόμενης αποδέσμευσης	SE/H/1654/001	95174/02-10-2020	TILLOTTS PHARMA GMBH	GR
Yaldigo 1600 mg δισκία ελεγχόμενης αποδέσμευσης	SE/H/1654/001	95174/02-10-2020	TILLOTTS PHARMA GMBH	GR
Yaldigo 1600 mg, tabletten met gereguleerde afgifte	SE/H/1654/001	RVG 120637	TILLOTTS PHARMA GMBH	NL

Product full name (in authorisation country)	MRP/DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Yaldigo, 1600 mg toimeainet modifitseeritult vabastavad tabletid	SE/H/1654/001	961518	TILLOTTS PHARMA AB	EE
Меколзин 1000 mg стомашно-устойчиви таблетки	ES/H/0587/002	61291	FAES FARMA, S.A.	BG
Меколзин 1000 mg супозитории	PT/H/2280/002	61293	FAES FARMA, S.A.	BG
Меколзин 500 mg стомашно-устойчиви таблетки	ES/H/0587/001	20200054	FAES FARMA, S.A.	BG
Меколзин 500 mg супозитории	PT/H/2280/001	61292	FAES FARMA, S.A.	BG
Пентаза 1 g супозитории	not available	9900188	FERRING GMBH	BG
Пентаза 500 mg таблетки с удължено освобождаване	not available	9900187	FERRING GMBH	BG
Салофалк 1 g супозитории	DE/H/5682/001	20100771	DR. FALK PHARMA G.M.B.H.	BG
Салофалк 1000 mg стомашно-устойчиви гранули с удължено освобождаване	DE/H/0363/002	20130427	DR. FALK PHARMA G.M.B.H.	BG
Салофалк 3 g стомашно-устойчиви гранули с удължено освобождаване	DE/H/0363/007	20130429	DR. FALK PHARMA G.M.B.H.	BG
Салофалк 4g/60 ml клизми	not available	9600070	DR. FALK PHARMA G.M.B.H.	BG
Салофалк 500 mg супозитории	not available	9600068	DR. FALK PHARMA G.M.B.H.	BG