03 October 2024
EMA/268423/2024
Human Medicines Division

List of nationally authorised medicinal products

Active substance(s): ondansetron

Procedure No. PSUSA/00002217/202402

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

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Ondansetron Accord 4 mg Injektions/Infusionslösung in einer Fertigspritze	ES/H/0627/001	140299	ACCORD HEALTHCARE B.V.	AT
Ondansetron Accord 8 mg Injektions/Infusionslösung in einer Fertigspritze	ES/H/0627/002	140300	ACCORD HEALTHCARE B.V.	AT
Ondansetron Accord 4 mg solution injectable/pour perfusion en seringue préremplie	ES/H/0627/001	BE557937	ACCORD HEALTHCARE B.V.	BE
Ondansetron Accord 8 mg solution injectable/pour perfusion en seringue préremplie	ES/H/0627/002	BE557946	ACCORD HEALTHCARE B.V.	BE
Ondansetron Accord 4 mg oplossing voor injectie/infusie in een voorgevulde spuit	ES/H/0627/001	BE557937	ACCORD HEALTHCARE B.V.	BE
Ondansetron Accord 8 mg oplossing voor injectie/infusie in een voorgevulde spuit	ES/H/0627/002	BE557946	ACCORD HEALTHCARE B.V.	BE
Ondansetron Accord 4 mg/2 ml Lösung zur Injektion/Infusion in einer Fertigspritze	ES/H/0627/001	2203436.00.00	ACCORD HEALTHCARE B.V.	DE
Ondansetron Accord 8 mg/4 ml Lösung zur Injektion/Infusion in einer Fertigspritze	ES/H/0627/002	2203919.00.00	ACCORD HEALTHCARE B.V.	DE
Ondansetron "Accordpharma", injektions-/infusionsvæske, opløsning i fyldt injektionssprøjte	ES/H/0627/001	62160	ACCORD HEALTHCARE B.V.	DK
Ondansetron "Accordpharma", injektions-/infusionsvæske, opløsning i fyldt injektionssprøjte	ES/H/0627/002	62245	ACCORD HEALTHCARE B.V.	DK
Ondansetron Accordpharma 4 mg injektio-/infuusioneste,	ES/H/0627/001	36729	ACCORD HEALTHCARE B.V.	FI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
liuos, esitäytetty ruisku				
Ondansetron Accordpharma 8 mg injektio-/infuusioneste, liuos, esitäytetty ruisku	ES/H/0627/002	36875	ACCORD HEALTHCARE B.V.	FI
Ondansetron Accord 4 mg oplossing voor injectie/infusie in een voorgevulde spuit	ES/H/0627/001	RVG 124397	ACCORD HEALTHCARE B.V.	NL
Ondansetron Accord 8 mg oplossing voor injectie/infusie in een voorgevulde spuit	ES/H/0627/002	RVG 124523	ACCORD HEALTHCARE B.V.	NL
Ondansetron Accord 4 mg oplossing voor injectie/infusie in een voorgevulde spuit	ES/H/0627/001	RVG 124397	ACCORD HEALTHCARE B.V.	NL
Ondansetron Accord 4 mg oplossing voor injectie/infusie in een voorgevulde spuit	ES/H/0627/001	RVG 124397	ACCORD HEALTHCARE B.V.	NL
Ondansetron Accord 8 mg oplossing voor injectie/infusie in een voorgevulde spuit	ES/H/0627/002	RVG 124523	ACCORD HEALTHCARE B.V.	NL
Ondansetron Accord 8 mg oplossing voor injectie/infusie in een voorgevulde spuit	ES/H/0627/002	RVG 124523	ACCORD HEALTHCARE B.V.	NL
Ondansetron Accordpharma 4 mg injeksjons-/infusjonsvæske, oppløsning i ferdigfylt sprøyte	ES/H/0627/001	19-12669	ACCORD HEALTHCARE B.V.	NO
Ondansetron Accordpharma 8 mg injeksjons-/infusjonsvæske, oppløsning i ferdigfylt sprøyte	ES/H/0627/002	21-13843	ACCORD HEALTHCARE B.V.	NO
Ondansetron Accordpharma 4 mg injektions-/infusionsvätska, lösning i förfylld spruta	ES/H/0627/001	58696	ACCORD HEALTHCARE B.V.	SE
Ondansetron Accordpharma 4 mg injektions-/infusionsvätska, lösning i förfylld spruta	ES/H/0627/001	58696	ACCORD HEALTHCARE B.V.	SE
Ondansetron Accordpharma 8 mg injektions-/infusionsvätska,	ES/H/0627/002	60474	ACCORD HEALTHCARE B.V.	SE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
lösning i förfylld spruta				
Ondansetron Accordpharma 8 mg injektions-/infusionsvätska, lösning i förfylld spruta	ES/H/0627/002	60474	ACCORD HEALTHCARE B.V.	SE
Ondansetron 2 mg/ml Solution for Injection/Infusion in Prefilled Syringe	ES/H/0627/001	PA2315/159/001	ACCORD HEALTHCARE IRELAND LIMITED	IE
Ondansetron 2 mg/ml Solution for Injection/Infusion in Prefilled Syringe	ES/H/0627/002	PA2315/159/001	ACCORD HEALTHCARE IRELAND LIMITED	IE
Ondansetron Accord, 8 mg, roztwór do wstrzykiwan/do infuzji w ampulko-strzykawce	ES/H/0627/002	25899	ACCORD HEALTHCARE POLSKA SP. Z O.O.	PL
Ondansetron Accord, 4 mg, roztwór do wstrzykiwan/do infuzji w ampulko-strzykawce	ES/H/0627/001	25898	ACCORD HEALTHCARE POLSKA SP. Z O.O.	PL
Ondansetron Accord 4 mg solución inyectable y para perfusión en jeringa precargada	ES/H/0627/001	85235	ACCORD HEALTHCARE S.L.U.	ES
Ondansetron Accord 4 mg solución inyectable y para perfusión en jeringa precargada	ES/H/0627/001	85235	ACCORD HEALTHCARE S.L.U.	ES
Ondansetron Accord 4 mg solución inyectable y para perfusión en jeringa precargada	ES/H/0627/001	85235	ACCORD HEALTHCARE S.L.U.	ES
Ondansetron Accord 8 mg solución inyectable y para perfusión en jeringa precargada	ES/H/0627/002	85236	ACCORD HEALTHCARE S.L.U.	ES
Ondansetron Accord 8 mg solución inyectable y para perfusión en jeringa precargada	ES/H/0627/002	85236	ACCORD HEALTHCARE S.L.U.	ES
Ondansetron Accord 8 mg solución inyectable y para	ES/H/0627/002	85236	ACCORD HEALTHCARE S.L.U.	ES

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
perfusión en jeringa precargada				
Ondansetron Accord 4 mg soluzione iniettabile/per infusione in siringa preriempita	ES/H/0627/001	048015010	ACCORD HEALTHCARE S.L.U.	IT
Ondansetron Accord 4 mg soluzione iniettabile/per infusione in siringa preriempita	ES/H/0627/001	048015022	ACCORD HEALTHCARE S.L.U.	IT
Ondansetron Accord 4 mg soluzione iniettabile/per infusione in siringa preriempita	ES/H/0627/001	048015034	ACCORD HEALTHCARE S.L.U.	IT
Ondansetron Accord 8 mg soluzione iniettabile/per infusione in siringa preriempita	ES/H/0627/002	048015046	ACCORD HEALTHCARE S.L.U.	IT
Ondansetron Accord 8 mg soluzione iniettabile/per infusione in siringa preriempita	ES/H/0627/002	048015059	ACCORD HEALTHCARE S.L.U.	IT
Ondansetron Accord 8 mg soluzione iniettabile/per infusione in siringa preriempita	ES/H/0627/002	048015061	ACCORD HEALTHCARE S.L.U.	IT
Ondansetron B. Braun 2 mg/ml Injektionslösung	DE/H/0805/001	1-27392	B.BRAUN MELSUNGEN AG	AT
Ondansetron B.Braun 2 mg/ml solution injectable	DE/H/0805/001	BE316897	B.BRAUN MELSUNGEN AG	BE
Ondansetron B.Braun 2 mg/ml solution injectable	DE/H/0805/001	BE316881	B.BRAUN MELSUNGEN AG	BE
Ondansetron B. Braun 2 mg/ml oplossing voor injectie	DE/H/0805/001	BE316881	B.BRAUN MELSUNGEN AG	BE
Ondansetron B. Braun 2 mg/ml oplossing voor injectie	DE/H/0805/001	BE316897	B.BRAUN MELSUNGEN AG	BE
Ondansetron B. Braun 2 mg/ml oplossing voor injectie	DE/H/0805/001	BE316881	B.BRAUN MELSUNGEN AG	BE
Ondansetron B.Braun 2 mg/ml solution injectable	DE/H/0805/001	BE316881	B.BRAUN MELSUNGEN AG	BE
Ondansetron B. Braun 2 mg/ml oplossing voor injectie	DE/H/0805/001	BE316897	B.BRAUN MELSUNGEN AG	BE
Ondansetron B.Braun 2 mg/ml solution injectable	DE/H/0805/001	BE316897	B.BRAUN MELSUNGEN AG	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Ondansetron B. Braun 2 mg/ml oplossing voor injectie	DE/H/0805/001	BE421644	B.BRAUN MELSUNGEN AG	BE
Ondansetron B.Braun 2 mg/ml solution injectable	DE/H/0805/001	BE421644	B.BRAUN MELSUNGEN AG	BE
Ondansetron B. Braun 2 mg/ml Injektionslösung	DE/H/0805/001	BE316881	B.BRAUN MELSUNGEN AG	BE
Ondansetron B. Braun 2 mg/ml Injektionslösung	DE/H/0805/001	BE316897	B.BRAUN MELSUNGEN AG	BE
Ondansetron B. Braun 2 mg/ml Injektionslösung	DE/H/0805/001	BE421644	B.BRAUN MELSUNGEN AG	BE
Ondansetron B. Braun 2 mg/ml Injektionslösung	DE/H/0805/001	BE316881	B.BRAUN MELSUNGEN AG	BE
Ondansetron B. Braun 2 mg/ml Injektionslösung	DE/H/0805/001	BE316897	B.BRAUN MELSUNGEN AG	BE
Ондансетрон Б. Браун 0,08 mg/ml инфузионен разтвор	DE/H/0805/002	20160366	B.BRAUN MELSUNGEN AG	BG
Ondansetron B. Braun 2 mg/ml, Injektionslösung	DE/H/0805/001	59057.00.00	B.BRAUN MELSUNGEN AG	DE
Ondansetron B. Braun 0,16 mg/ml Infusionslösung	DE/H/0805/003	94575.00.00	B.BRAUN MELSUNGEN AG	DE
Ondansetron B. Braun 0,08 mg/ml Infusionslösung	DE/H/0805/002	94574.00.00	B.BRAUN MELSUNGEN AG	DE
Ondansetron "B. Braun", injektionsvæske, opløsning	DE/H/0805/001	40588	B.BRAUN MELSUNGEN AG	DK
Ondansetrón B. Braun 2 mg/ml solución inyectable EFG	DE/H/0805/001	69.905	B.BRAUN MELSUNGEN AG	ES
Ondansetrón B. Braun 0,08 mg/ml solución para perfusión	DE/H/0805/002	81830	B.BRAUN MELSUNGEN AG	ES
Ondansetrón B. Braun 0,16 mg/ml solución para perfusión	DE/H/0805/003	81829	B.BRAUN MELSUNGEN AG	ES
Ondansetron B. Braun 2 mg/ml injektioneste, liuos	DE/H/0805/001	23137	B.BRAUN MELSUNGEN AG	FI
Ondansetron B. Braun 2 mg/ml injektionsvätska, lösning	DE/H/0805/001	23137	B.BRAUN MELSUNGEN AG	FI
Ondansetron B. Braun 0,16 mg/ml infuusioneste, liuos	DE/H/0805/003	33148	B.BRAUN MELSUNGEN AG	FI
Ondansetron B. Braun 0,08 mg/ml infuusioneste, liuos	DE/H/0805/002	33147	B.BRAUN MELSUNGEN AG	FI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Ondansetron B. Braun 2 mg/ml soluzione iniettabile	DE/H/0805/001	038128029	B.BRAUN MELSUNGEN AG	IT
Ondansetron B. Braun 2 mg/ml soluzione iniettabile	DE/H/0805/001	038128031	B.BRAUN MELSUNGEN AG	IT
Ondansetron B. Braun 2 mg/ml soluzione iniettabile.	DE/H/0805/001	038128043	B.BRAUN MELSUNGEN AG	IT
Ondansetron B. Braun 2 mg/ml soluzione iniettabile.	DE/H/0805/001	038128056	B.BRAUN MELSUNGEN AG	IT
Ondansetron B. Braun 2 mg/ml soluzione iniettabile	DE/H/0805/001	038128017	B.BRAUN MELSUNGEN AG	IT
Ondansetron B. Braun 0,08 mg/ml soluzione per infusione	DE/H/0805/002	038128068	B.BRAUN MELSUNGEN AG	IT
Ondansetron B. Braun 0,16 mg/ml soluzione per infusione	DE/H/0805/003	038128070	B.BRAUN MELSUNGEN AG	IT
Ondansetron B. Braun 2 mg/ml, Injektionslösung	DE/H/0805/001	0667/08120021	B.BRAUN MELSUNGEN AG	LU
Ondansetron B. Braun 2 mg/ml, oplossing voor injectie	DE/H/0805/001	RVG 34936	B.BRAUN MELSUNGEN AG	NL
Ondansetron B. Braun 0,08 mg/ml, oplossing voor infusie	DE/H/0805/002	RVG 116983	B.BRAUN MELSUNGEN AG	NL
Ondansetron B. Braun 0,16 mg/ml, oplossing voor infusie	DE/H/0805/003	RVG 116996	B.BRAUN MELSUNGEN AG	NL
Ondansetron B. Braun 2 mg/ml roztwór do wstrzykiwań	DE/H/0805/001	15717	B.BRAUN MELSUNGEN AG	PL
Ondansetron B. Braun, 0,08 mg/ml, roztwór do infuzji	DE/H/0805/002	23712	B.BRAUN MELSUNGEN AG	PL
Ondansetron B. Braun, 0,16 mg/ml, roztwór do infuzji	DE/H/0805/003	23713	B.BRAUN MELSUNGEN AG	PL
Ondansetron B. Braun 2 mg/ml injektionsvätska, lösning	DE/H/0805/001	24997	B.BRAUN MELSUNGEN AG	SE
Ondansetron B. Braun 0,08 mg/ml infusionsvätska, lösning	DE/H/0805/002	52598	B.BRAUN MELSUNGEN AG	SE
Ondansetron B. Braun 0,16 mg/ml infusionsvätska, lösning	DE/H/0805/003	52599	B.BRAUN MELSUNGEN AG	SE
Ondansetron B. Braun 2 mg/ml injekčný roztok	DE/H/0805/001	20/0255/07-S	B.BRAUN MELSUNGEN AG	SK
Zofran 4 mg comprimidos recubiertos con película	not available	59.069	BEXAL FARMACÉUTICA S.A.	ES

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zofran 8 mg comprimidos recubiertos con película	not available	59.070	BEXAL FARMACÉUTICA S.A.	ES
Zofran 4 mg solución inyectable	not available	59.071	BEXAL FARMACÉUTICA S.A.	ES
Zofran Zydis 4 mg liofilizado oral	not available	62.944	BEXAL FARMACÉUTICA S.A.	ES
Zofran Zydis 8 mg liofilizado oral	not available	62.945	BEXAL FARMACÉUTICA S.A.	ES
Zofran 4 mg solución inyectable	not available	59.072	BEXAL FARMACÉUTICA S.A.	ES
ONDANSETRON EG 8 mg, comprimé orodispersible	not available	NL32178	EG LABO LABORATOIRES EUROGENERICS - DO NOT USE	FR
Ondansetron Fresenius Kabi	DE/H/0640/003	65540	FRESENIUS KABI AB	DK
Ondansetron "Fresenius Kabi", infusionsvæske, opløsning	DE/H/0640/002	65539	FRESENIUS KABI AB	DK
Ondansetron Fresenius Kabi 0,08 mg/ml infusionvätska, lösning	DE/H/0640/002	38714	FRESENIUS KABI AB	FI
Ondansetron Fresenius Kabi 0,16 mg/ml infusionsvätska, lösning	DE/H/0640/003	38715	FRESENIUS KABI AB	FI
Ondansetron Fresenius Kabi 0.08 mg/ml infuusioneste, liuos	DE/H/0640/002	38714	FRESENIUS KABI AB	FI
Ondansetron Fresenius Kabi 0,16 mg/ml infuusioneste, liuos	DE/H/0640/003	38715	FRESENIUS KABI AB	FI
Ondansetron Fresenius Kabi 0,08 mg/ml innrennslislyf, lausn	DE/H/0640/002/E/001	IS/1/23/032/01	FRESENIUS KABI AB	IS
Ondansetron Fresenius Kabi 0,16 mg/ml innrennslislyf, lausn	DE/H/0640/003/E/001	IS/1/23/032/02	FRESENIUS KABI AB	IS
Ondansetron Fresenius Kabi 0,08 mg/ml infusionsvätska, lösning	DE/H/0640/002	61556	FRESENIUS KABI AB	SE
Ondansetron Fresenius Kabi 0,16 mg/ml infusionsvätska, lösning	DE/H/0640/003	61557	FRESENIUS KABI AB	SE
Ondansetron Kabi 0,08 mg/ml	DE/H/0640/002/E/001	141721	FRESENIUS KABI AUSTRIA	AT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Infusionslösung			GMBH	
Ondansetron Kabi 0,16 mg/ml Infusionslösung	DE/H/0640/003/E/001	141722	FRESENIUS KABI AUSTRIA GMBH	AT
Ondansetron Kabi 0,08 mg/ml raztopina za infundiranje	DE/H/0640/002/E/001	H/23/03019/001	FRESENIUS KABI AUSTRIA GMBH	SI
Ondansetron Kabi 0,08 mg/ml raztopina za infundiranje	DE/H/0640/002/E/001	H/23/03019/002	FRESENIUS KABI AUSTRIA GMBH	SI
Ondansetron Kabi 0,08 mg/ml raztopina za infundiranje	DE/H/0640/002/E/001	H/23/03019/004	FRESENIUS KABI AUSTRIA GMBH	SI
Ondansetron Kabi 0,08 mg/ml raztopina za infundiranje	DE/H/0640/002/E/001	H/23/03019/003	FRESENIUS KABI AUSTRIA GMBH	SI
Ondansetron Kabi 0,08 mg/ml raztopina za infundiranje	DE/H/0640/002/E/001	H/23/03019/005	FRESENIUS KABI AUSTRIA GMBH	SI
Ondansetron Kabi 0,08 mg/ml raztopina za infundiranje	DE/H/0640/002/E/001	H/23/03019/006	FRESENIUS KABI AUSTRIA GMBH	SI
Ondansetron Kabi 0,08 mg/ml raztopina za infundiranje	DE/H/0640/002/E/001	H/23/03019/008	FRESENIUS KABI AUSTRIA GMBH	SI
Ondansetron Kabi 0,08 mg/ml raztopina za infundiranje	DE/H/0640/002/E/001	H/23/03019/007	FRESENIUS KABI AUSTRIA GMBH	SI
Ondansetron Kabi 0,16 mg/ml raztopina za infundiranje	DE/H/0640/003/E/001	H/23/03019/009	FRESENIUS KABI AUSTRIA GMBH	SI
Ondansetron Kabi 0,16 mg/ml raztopina za infundiranje	DE/H/0640/003/E/001	H/23/03019/010	FRESENIUS KABI AUSTRIA GMBH	SI
Ondansetron Kabi 0,16 mg/ml raztopina za infundiranje	DE/H/0640/003/E/001	H/23/03019/012	FRESENIUS KABI AUSTRIA GMBH	SI
Ondansetron Kabi 0,16 mg/ml raztopina za infundiranje	DE/H/0640/003/E/001	H/23/03019/011	FRESENIUS KABI AUSTRIA GMBH	SI
Ondansetron Kabi 0,08 mg/ml Infusionslösung	DE/H/0640/002/E/001	20230128	FRESENIUS KABI BULGARIA EOOD	BG
Ondansetron Kabi 0,16 mg/ml Infusionslösung	DE/H/0640/003/E/001	20230129	FRESENIUS KABI BULGARIA EOOD	BG
Ondanzetron Kabi 0,08 mg/ml otopina za infuziju	DE/H/0640/002/E/001	HR-H-602841677	FRESENIUS KABI D.O.O.	HR
Ondanzetron Kabi 0,16 mg/ml otopina za infuziju	DE/H/0640/003/E/001	HR-H-272776579	FRESENIUS KABI D.O.O.	HR
Ondansetron Kabi 0,08 mg/ml infuzní roztok	DE/H/0640/002	20/515/20-C	FRESENIUS KABI DEUTSCHLAND GMBH	CZ
Ondansetron Kabi 0,16 mg/ml	DE/H/0640/003	20/516/20-C	FRESENIUS KABI	CZ

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
infuzní roztok			DEUTSCHLAND GMBH	
Ondansetron Kabi 0,08 mg/ml Infusionslösung	DE/H/0640/002	7003676.00.00	FRESENIUS KABI DEUTSCHLAND GMBH	DE
Ondansetron Kabi 0,16 mg/ml Infusionslösung	DE/H/0640/003	7003677.00.00	FRESENIUS KABI DEUTSCHLAND GMBH	DE
Ondansetron Kabi 0,08 mg/ml oldatos infúzió	DE/H/0640/002	OGYI-T-20247/05	FRESENIUS KABI DEUTSCHLAND GMBH	HU
Ondansetron Kabi 0,08 mg/ml oldatos infúzió	DE/H/0640/002	OGYI-T-20247/06	FRESENIUS KABI DEUTSCHLAND GMBH	HU
Ondansetron Kabi 0,08 mg/ml oldatos infúzió	DE/H/0640/002	OGYI-T-20247/08	FRESENIUS KABI DEUTSCHLAND GMBH	HU
Ondansetron Kabi 0,08 mg/ml oldatos infúzió	DE/H/0640/002	OGYI-T-20247/07	FRESENIUS KABI DEUTSCHLAND GMBH	HU
Ondansetron Kabi 0,08 mg/ml oldatos infúzió	DE/H/0640/002	OGYI-T-20247/09	FRESENIUS KABI DEUTSCHLAND GMBH	HU
Ondansetron Kabi 0,08 mg/ml oldatos infúzió	DE/H/0640/002	OGYI-T-20247/10	FRESENIUS KABI DEUTSCHLAND GMBH	HU
Ondansetron Kabi 0,08 mg/ml oldatos infúzió	DE/H/0640/002	OGYI-T-20247/12	FRESENIUS KABI DEUTSCHLAND GMBH	HU
Ondansetron Kabi 0,08 mg/ml oldatos infúzió	DE/H/0640/002	OGYI-T-20247/11	FRESENIUS KABI DEUTSCHLAND GMBH	HU
Ondansetron Kabi 0,16 mg/ml oldatos infúzió	DE/H/0640/003	OGYI-T-20247/13	FRESENIUS KABI DEUTSCHLAND GMBH	HU
Ondansetron Kabi 0,16 mg/ml oldatos infúzió	DE/H/0640/003	OGYI-T-20247/14	FRESENIUS KABI DEUTSCHLAND GMBH	HU
Ondansetron Kabi 0,16 mg/ml oldatos infúzió	DE/H/0640/003	OGYI-T-20247/16	FRESENIUS KABI DEUTSCHLAND GMBH	HU
Ondansetron Kabi 0,16 mg/ml oldatos infúzió	DE/H/0640/003	OGYI-T-20247/15	FRESENIUS KABI DEUTSCHLAND GMBH	HU
Ondansetron Kabi 0.08 mg/ml Solution for infusion	DE/H/0640/002	PA2059/080/001	FRESENIUS KABI DEUTSCHLAND GMBH	IE
Ondansetron Kabi 0.16 mg/ml Solution for infusion	DE/H/0640/003	PA2059/080/002	FRESENIUS KABI DEUTSCHLAND GMBH	IE
Ondansetron Kabi, 0,16 mg/ml, roztwór do infuzji	DE/H/0640/003	27697	FRESENIUS KABI DEUTSCHLAND GMBH	PL
Ondansetron Kabi, 0,08 mg/ml, roztwór do infuzji	DE/H/0640/002	27696	FRESENIUS KABI DEUTSCHLAND GMBH	PL
Ondansetron Kabi 0,08 mg/ml,	DE/H/0640/002	20/0185/22-S	FRESENIUS KABI	SK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
infúzny roztok			DEUTSCHLAND GMBH	
Ondansetron Kabi 0,16 mg/ml, infúzny roztok	DE/H/0640/003	20/0186/22-S	FRESENIUS KABI DEUTSCHLAND GMBH	SK
Ondansetron 0.08 mg/ml Solution for infusion	DE/H/0640/002	87730	FRESENIUS KABI ESPAÑA S.A.U.	ES
Ondansetron 0.16 mg/ml Solution for infusion	DE/H/0640/003	87731	FRESENIUS KABI ESPAÑA S.A.U.	ES
Ondansetron/Kabi 0,08 mg/ml διάλυμα για έγχυση	DE/H/0640/002/E/001	023860	FRESENIUS KABI HELLAS ΜΟΝΟΠΡΌΣΩΠΗ Α.Ε.	CY
Ondansetron/Kabi 0,16 mg/ml διάλυμα για έγχυση	DE/H/0640/003/E/001	023861	FRESENIUS KABI HELLAS ΜΟΝΟΠΡΌΣΩΠΗ Α.Ε.	CY
Ondansetron/Kabi 0,16 mg/ml διάλυμα για έγχυση	DE/H/0640/003	73062/05-07-2022	FRESENIUS KABI HELLAS ΜΟΝΟΠΡΌΣΩΠΗ Α.Ε.	GR
Ondansetron/Kabi 0,08 mg/ml διάλυμα για έγχυση	DE/H/0640/002	73061/05-07-2022	FRESENIUS KABI HELLAS ΜΟΝΟΠΡΌΣΩΠΗ Α.Ε.	GR
Ondansetron Kabi 0,08 mg/ml soluzione per infusione	DE/H/0640/002	037389071	FRESENIUS KABI ITALIA S.R.L.	IT
Ondansetron Kabi 0,08 mg/ml soluzione per infusione	DE/H/0640/002	037389083	FRESENIUS KABI ITALIA S.R.L.	IT
Ondansetron Kabi 0,08 mg/ml soluzione per infusione	DE/H/0640/002	037389107	FRESENIUS KABI ITALIA S.R.L.	IT
Ondansetron Kabi 0,08 mg/ml soluzione per infusione	DE/H/0640/002	037389095	FRESENIUS KABI ITALIA S.R.L.	IT
Ondansetron Kabi 0,08 mg/ml soluzione per infusione	DE/H/0640/002	037389119	FRESENIUS KABI ITALIA S.R.L.	IT
Ondansetron Kabi 0,08 mg/ml soluzione per infusione	DE/H/0640/002	037389121	FRESENIUS KABI ITALIA S.R.L.	IT
Ondansetron Kabi 0,08 mg/ml soluzione per infusione	DE/H/0640/002	037389145	FRESENIUS KABI ITALIA S.R.L.	IT
Ondansetron Kabi 0,08 mg/ml soluzione per infusione	DE/H/0640/002	037389133	FRESENIUS KABI ITALIA S.R.L.	IT
Ondansetron Kabi 0,16 mg/ml soluzione per infusione	DE/H/0640/003	037389158	FRESENIUS KABI ITALIA S.R.L.	IT
Ondansetron Kabi 0,16 mg/ml soluzione per infusione	DE/H/0640/003	037389160	FRESENIUS KABI ITALIA S.R.L.	IT
Ondansetron Kabi 0,16 mg/ml soluzione per infusione	DE/H/0640/003	037389184	FRESENIUS KABI ITALIA S.R.L.	IT
Ondansetron Kabi 0,16 mg/ml	DE/H/0640/003	037389172	FRESENIUS KABI ITALIA S.R.L.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
soluzione per infusione				
Ondansetron Kabi 0.08 mg/ml solution for infusion	DE/H/0640/002	PL 08828/0333	FRESENIUS KABI LIMITED	XI
Ondansetron Kabi 0.16 mg/ml solution for infusion	DE/H/0640/003	PL 08828/0334	FRESENIUS KABI LIMITED	XI
Ondansetron Fresenius Kabi 0,08 mg/ml oplossing voor infusie	DE/H/0640/002	RVG 127773	FRESENIUS KABI NEDERLAND B.V.	NL
Ondansetron Fresenius Kabi 0,16 mg/ml oplossing voor infusie	DE/H/0640/003	RVG 127774	FRESENIUS KABI NEDERLAND B.V.	NL
Ondansetron Fresenius Kabi 0,08 mg/ml infusjonsvæske, oppløsning	DE/H/0640/002	20-13798	FRESENIUS KABI NORGE AS OSLO	NO
Ondansetron Fresenius Kabi 0,16 mg/ml infusjonsvæske, oppløsning	DE/H/0640/003	22-14658	FRESENIUS KABI NORGE AS OSLO	NO
Ondansetron Fresenius Kabi 0,08 mg/ml oplossing voor infusie	DE/H/0640/002	BE599795	FRESENIUS KABI NV/SA	BE
Ondansetron Fresenius Kabi 0,08 mg/ml oplossing voor infusie	DE/H/0640/002	BE599804	FRESENIUS KABI NV/SA	BE
Ondansetron Fresenius Kabi 0,16 mg/ml oplossing voor infusie	DE/H/0640/003	BE599813	FRESENIUS KABI NV/SA	BE
Ondansetron Fresenius Kabi 0,16 mg/ml solution pour perfusion	DE/H/0640/003	BE599813	FRESENIUS KABI NV/SA	BE
Ondansetron Fresenius Kabi 0,08 mg/ml solution pour perfusion	DE/H/0640/002	BE599795	FRESENIUS KABI NV/SA	BE
Ondansetron Fresenius Kabi 0,08 mg/ml solution pour perfusion	DE/H/0640/002	BE599804	FRESENIUS KABI NV/SA	BE
Ondansetrom Kabi 0,08 mg/ml solução para perfusão	DE/H/0640/002/E/001	5857172	FRESENIUS KABI PHARMA PORTUGAL, LDA.	PT
Ondansetrom Kabi 0,08 mg/ml	DE/H/0640/002/E/001	5857206	FRESENIUS KABI PHARMA	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
solução para perfusão			PORTUGAL, LDA.	
Ondansetrom Kabi 0,08 mg/ml solução para perfusão	DE/H/0640/002/E/001	5857230	FRESENIUS KABI PHARMA PORTUGAL, LDA.	PT
Ondansetrom Kabi 0,08 mg/ml solução para perfusão	DE/H/0640/002/E/001	5857248	FRESENIUS KABI PHARMA PORTUGAL, LDA.	PT
Ondansetrom Kabi 0,08 mg/ml solução para perfusão	DE/H/0640/002/E/001	5857214	FRESENIUS KABI PHARMA PORTUGAL, LDA.	PT
Ondansetrom Kabi 0,08 mg/ml solução para perfusão	DE/H/0640/002/E/001	5857255	FRESENIUS KABI PHARMA PORTUGAL, LDA.	PT
Ondansetrom Kabi 0,08 mg/ml solução para perfusão	DE/H/0640/002/E/001	5857222	FRESENIUS KABI PHARMA PORTUGAL, LDA.	PT
Ondansetrom Kabi 0,08 mg/ml solução para perfusão	DE/H/0640/002/E/001	5857263	FRESENIUS KABI PHARMA PORTUGAL, LDA.	PT
Ondansetrom Kabi 0,16 mg/ml solução para perfusão	DE/H/0640/003/E/001	5857370	FRESENIUS KABI PHARMA PORTUGAL, LDA.	PT
Ondansetrom Kabi 0,16 mg/ml solução para perfusão	DE/H/0640/003/E/001	5857362	FRESENIUS KABI PHARMA PORTUGAL, LDA.	PT
Ondansetrom Kabi 0,16 mg/ml solução para perfusão	DE/H/0640/003/E/001	5857354	FRESENIUS KABI PHARMA PORTUGAL, LDA.	PT
Ondansetrom Kabi 0,16 mg/ml solução para perfusão	DE/H/0640/003/E/001	5857347	FRESENIUS KABI PHARMA PORTUGAL, LDA.	PT
Ondansetron Kabi 0,08 mg/ml infusioonilahus	DE/H/0640/002/E/001	1106423	FRESENIUS KABI POLSKA SP. Z O.O.	EE
Ondansetron Kabi 0,16 mg/ml infusioonilahus	DE/H/0640/003/E/001	1106023	FRESENIUS KABI POLSKA SP. Z O.O.	EE
Ondansetron Kabi 0,08 mg/ml infuzinis tirpalas	DE/H/0640/002/E/001	LT/1/23/5166/001	FRESENIUS KABI POLSKA SP. Z O.O.	LT
Ondansetron Kabi 0,08 mg/ml infuzinis tirpalas	DE/H/0640/002/E/001	LT/1/23/5166/002	FRESENIUS KABI POLSKA SP. Z O.O.	LT
Ondansetron Kabi 0,08 mg/ml infuzinis tirpalas	DE/H/0640/002/E/001	LT/1/23/5166/003	FRESENIUS KABI POLSKA SP. Z O.O.	LT
Ondansetron Kabi 0,08 mg/ml infuzinis tirpalas	DE/H/0640/002/E/001	LT/1/23/5166/004	FRESENIUS KABI POLSKA SP. Z O.O.	LT
Ondansetron Kabi 0,08 mg/ml infuzinis tirpalas	DE/H/0640/002/E/001	LT/1/23/5166/005	FRESENIUS KABI POLSKA SP. Z O.O.	LT
Ondansetron Kabi 0,08 mg/ml infuzinis tirpalas	DE/H/0640/002/E/001	LT/1/23/5166/006	FRESENIUS KABI POLSKA SP. Z O.O.	LT
Ondansetron Kabi 0,08 mg/ml	DE/H/0640/002/E/001	LT/1/23/5166/008	FRESENIUS KABI POLSKA SP. Z	LT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
infuzinis tirpalas			O.O.	
Ondansetron Kabi 0,08 mg/ml infuzinis tirpalas	DE/H/0640/002/E/001	LT/1/23/5166/007	FRESENIUS KABI POLSKA SP. Z O.O.	LT
Ondansetron Kabi 0,16 mg/ml infuzinis tirpalas	DE/H/0640/003/E/001	LT/1/23/5167/001	FRESENIUS KABI POLSKA SP. Z O.O.	LT
Ondansetron Kabi 0,16 mg/ml infuzinis tirpalas	DE/H/0640/003/E/001	LT/1/23/5167/002	FRESENIUS KABI POLSKA SP. Z O.O.	LT
Ondansetron Kabi 0,16 mg/ml infuzinis tirpalas	DE/H/0640/003/E/001	LT/1/23/5167/004	FRESENIUS KABI POLSKA SP. Z O.O.	LT
Ondansetron Kabi 0,16 mg/ml infuzinis tirpalas	DE/H/0640/003/E/001	LT/1/23/5167/003	FRESENIUS KABI POLSKA SP. Z O.O.	LT
Ondansetron Kabi 0,08 mg/ml šķīdums infūzijām	DE/H/0640/002/E/001	23-0075	FRESENIUS KABI POLSKA SP. Z O.O.	LV
Ondansetron Kabi 0,16 mg/ml šķīdums infūzijām	DE/H/0640/003/E/001	23-0076	FRESENIUS KABI POLSKA SP. Z O.O.	LV
Ondansetron Kabi 0,08 mg/ml soluție perfuzabilă	DE/H/0640/002/E/001	14999/2023/01	FRESENIUS KABI ROMANIA SRL	RO
Ondansetron Kabi 0,08 mg/ml soluție perfuzabilă	DE/H/0640/002/E/001	14999/2023/02	FRESENIUS KABI ROMANIA SRL	RO
Ondansetron Kabi 0,08 mg/ml soluție perfuzabilă	DE/H/0640/002/E/001	14999/2023/04	FRESENIUS KABI ROMANIA SRL	RO
Ondansetron Kabi 0,08 mg/ml soluție perfuzabilă	DE/H/0640/002/E/001	14999/2023/03	FRESENIUS KABI ROMANIA SRL	RO
Ondansetron Kabi 0,08 mg/ml soluție perfuzabilă	DE/H/0640/002/E/001	14999/2023/05	FRESENIUS KABI ROMANIA SRL	RO
Ondansetron Kabi 0,08 mg/ml soluție perfuzabilă	DE/H/0640/002/E/001	14999/2023/06	FRESENIUS KABI ROMANIA SRL	RO
Ondansetron Kabi 0,08 mg/ml soluție perfuzabilă	DE/H/0640/002/E/001	14999/2023/08	FRESENIUS KABI ROMANIA SRL	RO
Ondansetron Kabi 0,08 mg/ml soluție perfuzabilă	DE/H/0640/002/E/001	14999/2023/07	FRESENIUS KABI ROMANIA SRL	RO
Ondansetron Kabi 0,16 mg/ml soluție perfuzabilă	DE/H/0640/003/E/001	15000/2023/01	FRESENIUS KABI ROMANIA SRL	RO
Ondansetron Kabi 0,16 mg/ml soluție perfuzabilă	DE/H/0640/003/E/001	15000/2023/02	FRESENIUS KABI ROMANIA SRL	RO
Ondansetron Kabi 0,16 mg/ml soluție perfuzabilă	DE/H/0640/003/E/001	15000/2023/04	FRESENIUS KABI ROMANIA SRL	RO
Ondansetron Kabi 0,16 mg/ml	DE/H/0640/003/E/001	15000/2023/03	FRESENIUS KABI ROMANIA SRL	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
soluție perfuzabilă				
Ondansetron-hameln 2 mg/ml Injektionslösung Wirkstoff: Ondansetronhydrochlorid	not available	59062.00.00	HAMELN PHARMA GMBH	DE
Ondansetron-hameln 2 mg/ml Injektionslösung	DE/H/0806/001	59061.00.00	HAMELN PHARMA GMBH	DE
Ondansetron "Hameln"	DE/H/0806/001/MR	40589	HAMELN PHARMA GMBH	DK
Ondansetron Hameln 2 mg/ml injektionsvätska, lösning	DE/H/0806/001	24998	HAMELN PHARMA GMBH	SE
Zofran 4 mg Filmtabletten	not available	22008.00.01	HEXAL AG	DE
Zofran 8 mg Filmtabletten	not available	22008.01.01	HEXAL AG	DE
Zofran Lösung 4 mg/5 ml	not available	37196.00.00	HEXAL AG	DE
Zofran® 8 mg Zydis® Lingual Schmelztablette	not available	41349.01.00	HEXAL AG	DE
Zofran 4 mg Zydis Lingual Schmelztablette	not available	41349.00.00	HEXAL AG	DE
Ondansetron 8mg/5ml oral solution	not available	PL 00427/0271	ROSEMONT PHARMACEUTICALS LTD	XI
Zofran 4 mg film-coated tablets	not available	PA0711/327/003	ROWEX LTD	IE
Zofran 8 mg film-coated tablets	not available	PA0711/327/004	ROWEX LTD	IE
Zofran 4mg/2ml Solution for Injection or Infusion	not available	PA0711/327/001	ROWEX LTD	IE
Zofran 4 mg/5 ml Syrup	not available	PA0711/327/002	ROWEX LTD	IE
Zofran Zydis 4 mg oral lyophilisate	not available	PA0711/327/005	ROWEX LTD	IE
Zofran Zydis 8 mg oral lyophilisate	not available	PA0711/327/006	ROWEX LTD	IE
ZOPHREN 8 mg, comprimé pelliculé	not available	34009 335 387 4 3	SANDOZ	FR
ZOPHREN 8 mg, comprimé pelliculé	not available	34009 335 388 0 4	SANDOZ	FR
ZOPHREN 8 mg, comprimé pelliculé	not available	34009 335 389 7 2	SANDOZ	FR
ZOPHREN 8 mg, comprimé	not available	34009 335 390 5 4	SANDOZ	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
pelliculé				
ZOPHREN 8 mg, comprimé pelliculé	not available	34009 335 391 1 5	SANDOZ	FR
ZOPHREN 8 mg, comprimé pelliculé	not available	34009 361 620 4 4	SANDOZ	FR
ZOPHREN 8 mg, comprimé pelliculé	not available	34009 361 641 1 6	SANDOZ	FR
ZOPHREN 8 mg, comprimé pelliculé	not available	34009 361 642 8 4	SANDOZ	FR
ZOPHREN 8 mg, comprimé pelliculé	not available	34009 361 972 8 2	SANDOZ	FR
ZOPHREN 8 mg, comprimé pelliculé	not available	34009 361 973 4 3	SANDOZ	FR
ZOPHREN 8 mg, comprimé pelliculé	not available	34009 556 875 1 1	SANDOZ	FR
ZOPHREN 8 mg, comprimé pelliculé	not available	34009 556 876 8 9	SANDOZ	FR
ZOPHREN 8 mg, comprimé pelliculé	not available	34009 556 877 4 0	SANDOZ	FR
ZOPHREN 8 mg, comprimé pelliculé	not available	34009 556 878 0 1	SANDOZ	FR
ZOPHREN 8 mg, comprimé pelliculé	not available	34009 556 879 7 9	SANDOZ	FR
ZOPHREN 8 mg, comprimé pelliculé	not available	34009 564 739 6 0	SANDOZ	FR
ZOPHREN 8 mg, comprimé pelliculé	not available	34009 564 740 4 2	SANDOZ	FR
ZOPHREN 2 mg/ml, solution injectable en ampoule (IV)	not available	34009 335 392 8 3	SANDOZ	FR
ZOPHREN 2 mg/ml, solution injectable en ampoule (IV)	not available	34009 335 393 4 4	SANDOZ	FR
ZOPHREN 2 mg/ml, solution injectable en ampoule (IV)	not available	34009 550 795 9 0	SANDOZ	FR
ZOPHREN 2 mg/ml, solution injectable en ampoule (IV)	not available	34009 556 831 4 8	SANDOZ	FR
ZOPHREN 2 mg/ml, solution injectable en ampoule (IV)	not available	34009 556 832 0 9	SANDOZ	FR
ZOPHREN 2 mg/ml, solution	not available	34009 556 839 5 7	SANDOZ	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
injectable en ampoule (IV)				
ZOPHREN 2 mg/ml, solution injectable en ampoule (IV)	not available	34009 556 840 3 9	SANDOZ	FR
ZOPHREN 2 mg/ml, solution injectable en ampoule (IV)	not available	34009 556 843 2 9	SANDOZ	FR
ZOPHREN 2 mg/ml, solution injectable en ampoule (IV)	not available	34009 556 844 9 7	SANDOZ	FR
ZOPHREN 4 mg, comprimé pelliculé	not available	34009 335 382 2 4	SANDOZ	FR
ZOPHREN 4 mg, comprimé pelliculé	not available	34009 335 383 9 2	SANDOZ	FR
ZOPHREN 4 mg, comprimé pelliculé	not available	34009 335 384 5 3	SANDOZ	FR
ZOPHREN 4 mg, comprimé pelliculé	not available	34009 335 385 1 4	SANDOZ	FR
ZOPHREN 4 mg, comprimé pelliculé	not available	34009 335 386 8 2	SANDOZ	FR
ZOPHREN 4 mg, comprimé pelliculé	not available	34009 361 636 8 3	SANDOZ	FR
ZOPHREN 4 mg, comprimé pelliculé	not available	34009 361 637 4 4	SANDOZ	FR
ZOPHREN 4 mg, comprimé pelliculé	not available	34009 361 639 7 3	SANDOZ	FR
ZOPHREN 4 mg, comprimé pelliculé	not available	34009 361 640 5 5	SANDOZ	FR
ZOPHREN 4 mg, comprimé pelliculé	not available	34009 556 861 0 1	SANDOZ	FR
ZOPHREN 4 mg, comprimé pelliculé	not available	34009 556 857 3 9	SANDOZ	FR
ZOPHREN 4 mg, comprimé pelliculé	not available	34009 556 859 6 8	SANDOZ	FR
ZOPHREN 4 mg, comprimé pelliculé	not available	34009 556 860 4 0	SANDOZ	FR
ZOPHREN 4 mg, comprimé pelliculé	not available	34009 556 862 7 9	SANDOZ	FR
ZOPHREN 4 mg, comprimé pelliculé	not available	34009 564 736 7 0	SANDOZ	FR
ZOPHREN 4 mg, lyophilisat oral	not available	34009 343 944 6 1	SANDOZ	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
ZOPHREN 4 mg, lyophilisat oral	not available	34009 343 945 2 2	SANDOZ	FR
ZOPHREN 4 mg, lyophilisat oral	not available	34009 343 946 9 0	SANDOZ	FR
ZOPHREN 4 mg, lyophilisat oral	not available	34009 343 947 5 1	SANDOZ	FR
ZOPHREN 8 mg, lyophilisat oral	not available	34009 343 948 1 2	SANDOZ	FR
ZOPHREN 8 mg, lyophilisat oral	not available	34009 343 949 8 0	SANDOZ	FR
ZOPHREN 8 mg, lyophilisat oral	not available	34009 343 950 6 2	SANDOZ	FR
ZOPHREN 8 mg, lyophilisat oral	not available	34009 343 951 2 3	SANDOZ	FR
ZOPHREN 4 mg/5 ml, sirop	not available	34009 341 656 3 4	SANDOZ	FR
ZOPHREN 4 mg, comprimé pelliculé	not available	34009 361 638 0 5	SANDOZ	FR
ZOPHREN 4 mg, comprimé pelliculé	not available	34009 564 737 3 1	SANDOZ	FR
Zofran, frysetørrede tabletter	not available	18515	SANDOZ A/S	DK
Zofran, frysetørrede tabletter	not available	18516	SANDOZ A/S	DK
Zofran	not available	17600	SANDOZ A/S	DK
Zofran Zydis 4 mg kylmäkuivattu tabletti	not available	12874	SANDOZ A/S	FI
Zofran Zydis 8 mg kylmäkuivattu tabletti	not available	12875	SANDOZ A/S	FI
Zofran 4 mg kalvopäällysteinen tabletti	not available	10793	SANDOZ A/S	FI
Zofran 8 mg kalvopäällysteinen tabletti	not available	10794	SANDOZ A/S	FI
Zofran 2 mg/ml injektioneste, liuos	not available	10792	SANDOZ A/S	FI
Zofran 0,8 mg/ml mixtúra, lausn.	not available	IS/1/13/004/01	SANDOZ A/S	IS
Zofran 4 mg filmuhúðaðar töflur	not available	890111	SANDOZ A/S	IS
Zofran 8 mg filmuhúðaðar töflur.	not available	890112	SANDOZ A/S	IS
Zofran 4 mg filmdrasjerte tabletter	not available	7758	SANDOZ A/S	NO
Zofran 4 mg, smeltetabletter	not available	96-3183	SANDOZ A/S	NO
Zofran 8 mg, smeltetabletter	not available	96-3184	SANDOZ A/S	NO
Zofran 0,8 mg/ml, mikstur, oppløsning	not available	95-2485	SANDOZ A/S	NO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zofran injeksjonsvæske 2 mg/ml, oppløsning	not available	7699	SANDOZ A/S	NO
Zofran 8 mg filmdrasjerte tabletter	not available	7700	SANDOZ A/S	NO
Zofran munlöslig 8 mg frystorkad tablett	not available	13586	SANDOZ A/S	SE
Zofran munlöslig 4 mg frystorkad tablett	not available	13585	SANDOZ A/S	SE
Zofran 4 mg filmdragerade tabletter	not available	11493	SANDOZ A/S	SE
Zofran 8 mg filmdragerade tabletter	not available	11494	SANDOZ A/S	SE
Zofran 0,8 mg/ml oral lösning	not available	13433	SANDOZ A/S	SE
Zofran 4 Zydis, smelttabletten 4 mg	not available	RVG 21471	SANDOZ B.V.	NL
Zofran 8 Zydis, smelttabletten 8 mg	not available	RVG 21472	SANDOZ B.V.	NL
Zofran Stroop, stroop 4 mg/5 ml	not available	RVG 19922	SANDOZ B.V.	NL
Zofran 8 mg filmom obložene tablete	not available	HR-H-774416957	SANDOZ D.O.O.	HR
Zofran 0,8 mg/ml Xarope	not available	2504280	SANDOZ FARMACÊUTICA LDA.	PT
ZOFRAN® 8 mg-Filmtabletten	not available	1-19343	SANDOZ GMBH	AT
Zofran - Lösung zum Einnehmen	not available	1-22404	SANDOZ GMBH	AT
Zofran 4 mg - Filmtabletten	not available	1-19341	SANDOZ GMBH	AT
ZOFRAN® Zydis® 4 mg-Tabletten	not available	1-22512	SANDOZ GMBH	AT
ZOFRAN® Zydis® 8 mg-Tabletten	not available	1-22513	SANDOZ GMBH	AT
ZOFRAN® 4 mg-Ampullen	not available	1-19340	SANDOZ GMBH	AT
Zofran 8 mg - Ampullen	not available	1-19342	SANDOZ GMBH	AT
Zofran-Zydis 8 mg lyofilisaat voor oraal gebruik	not available	BE199193	SANDOZ N.V.	BE
Zofran 8 mg oplossing voor injectie	not available	BE150297	SANDOZ N.V.	BE
Zofran 4 mg oplossing voor injectie	not available	BE150272	SANDOZ N.V.	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Avessaron 8 mg oplossing voor injectie	not available	BE150236	SANDOZ N.V.	BE
Avessaron 4 mg oplossing voor injectie	not available	BE150263	SANDOZ N.V.	BE
ЗОФРАН 2 mg/ml инжекционен разтвор	not available	20011016	SANDOZ PHARMACEUTICALS D.D.	BG
ЗОФРАН 4 mg филмирани таблетки	not available	20011017	SANDOZ PHARMACEUTICALS D.D.	BG
ЗОФРАН 8 mg филмирани таблетки	not available	20011018	SANDOZ PHARMACEUTICALS D.D.	BG
Zofron 4mg επικαλυμμένο με λεπτό υμένιο δισκίο	not available	38976/10/18-04-2011	SANDOZ PHARMACEUTICALS D.D.	GR
Zofron Zydis 8mg, δισκίο λυοφιλοποιημένο, από του στόματος	not available	38987/10/18-04-2011	SANDOZ PHARMACEUTICALS D.D.	GR
Zofron 8mg επικαλυμμένο με λεπτό υμένιο δισκίο	not available	38978/10/18-04-2011	SANDOZ PHARMACEUTICALS D.D.	GR
Zofran 4 mg/2 ml injekcinis ar infuzinis tirpalas	not available	LT/1/94/1091/006	SANDOZ PHARMACEUTICALS D.D.	LT
Zofran 4 mg/2 ml injekcinis ar infuzinis tirpalas	not available	LT/1/94/1091/007	SANDOZ PHARMACEUTICALS D.D.	LT
Zofran 4 mg/2 ml injekcinis ar infuzinis tirpalas	not available	LT/1/94/1091/010	SANDOZ PHARMACEUTICALS D.D.	LT
Zofran 8 mg plėvele dengtos tabletės	not available	LT/1/94/1091/004	SANDOZ PHARMACEUTICALS D.D.	LT
Zofran 8 mg plėvele dengtos tabletės	not available	LT/1/94/1091/005	SANDOZ PHARMACEUTICALS D.D.	LT
Zofran 8 mg/4 ml injekcinis ar infuzinis tirpalas	not available	LT/1/94/1091/008	SANDOZ PHARMACEUTICALS D.D.	LT
Zofran 8 mg/4 ml injekcinis ar infuzinis tirpalas	not available	LT/1/94/1091/009	SANDOZ PHARMACEUTICALS D.D.	LT
Zofran 8 mg/4 ml injekcinis ar infuzinis tirpalas	not available	LT/1/94/1091/011	SANDOZ PHARMACEUTICALS D.D.	LT
Zofran 4 mg filmsko obložene tablete	not available	H/92/01707/003	SANDOZ PHARMACEUTICALS D.D.	SI
Zofran 4 mg Compresse orodispersibili	not available	027612098	SANDOZ S.P.A.	IT
Zofran 4 mg Compresse	not available	027612098	SANDOZ S.P.A.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
orodispersibili				
Zofran 4 mg Compresse orodispersibili	not available	027612098	SANDOZ S.P.A.	IT
Zofran 4 mg Compresse rivestite con film	not available	027612011	SANDOZ S.P.A.	IT
Zofran 8 mg Compresse orodispersibili	not available	027612112	SANDOZ S.P.A.	IT
Zofran 40 mg/20 ml Soluzione iniettabile	not available	027612136	SANDOZ S.P.A.	IT
Zofran 8 mg/4 ml Soluzione iniettabile	not available	027612047	SANDOZ S.P.A.	IT
Zofran 8 mg Compresse rivestite con film	not available	027612023	SANDOZ S.P.A.	IT
Zofran 4 mg/2 ml Soluzione iniettabile	not available	027612035	SANDOZ S.P.A.	IT
Zofran 4 mg/2 ml Soluzione iniettabile	not available	027612163	SANDOZ S.P.A.	IT
Zofran 8 mg/4 ml Soluzione iniettabile	not available	027612175	SANDOZ S.P.A.	IT
Zofran 4 mg/5 ml Sciroppo	not available	027612086	SANDOZ S.P.A.	IT
Zofran Zydis 4 mg tablety dispergovatelné v ústech	not available	20/475/99-C	SANDOZ S.R.O.	CZ
Zofran 2mg/ml injekční roztok	not available	20/164/91-C	SANDOZ S.R.O.	CZ
Zofran Zydis 8 mg tablety dispergovatelné v ústech	not available	20/474/99-C	SANDOZ S.R.O.	CZ
ONDANSETRON ZENTIVA 8 mg, comprimé orodispersible	not available	34009 300 878 4 8	ZENTIVA FRANCE	FR
ONDANSETRON ZENTIVA 8 mg, comprimé orodispersible	not available	34009 382 840 3 4	ZENTIVA FRANCE	FR
ONDANSETRON ZENTIVA 8 mg, comprimé orodispersible	not available	34009 382 842 6 3	ZENTIVA FRANCE	FR
ONDANSETRON ZENTIVA 8 mg, comprimé orodispersible	not available	34009 382 843 2 4	ZENTIVA FRANCE	FR
ONDANSETRON ZENTIVA 8 mg, comprimé orodispersible	not available	34009 382 844 9 2	ZENTIVA FRANCE	FR
ONDANSETRON ZENTIVA 8 mg, comprimé orodispersible	not available	34009 300 878 5 5	ZENTIVA FRANCE	FR
ONDANSETRON ZENTIVA 8	not available	34009 300 878 2 4	ZENTIVA FRANCE	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
mg, comprimé orodispersible				
ONDANSETRON ZENTIVA 8 mg, comprimé orodispersible	not available	34009 300 878 3 1	ZENTIVA FRANCE	FR