



EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

27 January 2022
EMA/148029/2022
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: loperamide, loperamide / simeticone

Procedure no.: PSUSA/00010665/202105



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
Imodium 2mg Kapseln	not available	16.789	JOHNSON & JOHNSON GMBH	AT
Imodium 2mg Kapseln	not available	16.789	JOHNSON & JOHNSON GMBH	AT
Imodium akut 2 mg Schmelztabletten	not available	1-22679	JOHNSON & JOHNSON GMBH	AT
Imodium akut 2 mg Schmelztabletten	not available	1-22679	JOHNSON & JOHNSON GMBH	AT
Imosec 0,2 mg/ml Lösung zum Einnehmen	not available	16.790	JANSSEN-CILAG PHARMA GMBH	AT
Imosec 0,2 mg/ml Lösung zum Einnehmen	not available	16.790	JANSSEN-CILAG PHARMA GMBH	AT
Loperamid Sandoz 2 mg - Kapseln	not available	1-24708	SANDOZ GMBH	AT
Imodium 0,2 mg/ml drank	not available	BE099005	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium 0,2 mg/ml drank	not available	BE099005	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium 0,2 mg/ml Lösung zum Einnehmen	not available	BE099005	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium 0,2 mg/ml Lösung zum Einnehmen	not available	BE099005	JOHNSON & JOHNSON CONSUMER N.V./S.A.	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
Imodium 0,2 mg/ml solution buvable	not available	BE099005	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium 0,2 mg/ml solution buvable	not available	BE099005	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium 2 mg gélules	not available	BE 001215	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium 2 mg gélules	not available	BE 001215	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium 2 mg gélules	not available	BE 001215	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium 2 mg harde capsules	not available	BE 001215	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium 2 mg harde capsules	not available	BE 001215	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium 2 mg harde capsules	not available	BE 001215	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium 2 mg Hartkapseln	not available	BE 001215	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium 2 mg Hartkapseln	not available	BE 001215	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium 2 mg Hartkapseln	not available	BE 001215	JOHNSON & JOHNSON CONSUMER N.V./S.A.	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
Imodium Duo 2 mg/125 mg	DK/H/3047/001	BE294341	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium Duo 2 mg/125 mg	DK/H/3047/001	BE294341	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium Duo 2 mg/125 mg	DK/H/3047/001	BE294341	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium Duo 2 mg/125 mg	DK/H/3047/001	BE294366	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium Duo 2 mg/125 mg	DK/H/3047/001	BE294341	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium Duo 2 mg/125 mg	DK/H/3047/001	BE294366	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium Duo 2 mg/125 mg	DK/H/3047/001	BE294366	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium Duo 2 mg/125 mg	DK/H/3047/001	BE294341	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium Duo 2 mg/125 mg	DK/H/3047/001	BE294341	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium Duo 2 mg/125 mg	DK/H/3047/001	BE294366	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium Duo 2 mg/125 mg	DK/H/3047/001	BE294341	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE

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Imodium Duo 2 mg/125 mg	DK/H/3047/001	BE294366	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium Duo 2 mg/125 mg	DK/H/3047/001	BE294366	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium Duo 2 mg/125 mg	DK/H/3047/001	BE294366	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium Duo 2 mg/125 mg	DK/H/3047/001	BE294366	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium Duo 2 mg/125 mg	DK/H/3047/001	BE294341	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium Duo 2 mg/125 mg	DK/H/3047/001	BE294341	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium Duo 2 mg/125 mg	DK/H/3047/001	BE294366	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium Duo 2 mg/125 mg	DK/H/3047/001	BE294341	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium Duo 2 mg/125 mg	DK/H/3047/001	BE294366	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium Duo 2 mg/125 mg	DK/H/3047/001	BE294366	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium Duo 2 mg/125 mg	DK/H/3047/001	BE294341	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE

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Imodium Duo 2 mg/125 mg	DK/H/3047/001	BE294366	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium Duo 2 mg/125 mg	DK/H/3047/001	BE294366	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium Duo 2 mg/125 mg	DK/H/3047/001	BE294366	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium Duo 2 mg/125 mg	DK/H/3047/001	BE294341	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium Duo 2 mg/125 mg	DK/H/3047/001	BE294366	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium Duo 2 mg/125 mg	DK/H/3047/001	BE294341	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium Duo 2 mg/125 mg	DK/H/3047/001	BE294366	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium Duo 2 mg/125 mg	DK/H/3047/001	BE294341	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium Duo 2 mg/125 mg	DK/H/3047/001	BE294341	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium Duo 2 mg/125 mg	DK/H/3047/001	BE294341	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium Duo 2 mg/125 mg comprimés	DK/H/3047/001	BE294341	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE

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Imodium Duo 2 mg/125 mg comprimés	DK/H/3047/001	BE294366	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium Duo 2 mg/125 mg comprimés	DK/H/3047/001	BE294366	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium Duo 2 mg/125 mg comprimés	DK/H/3047/001	BE294341	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium Duo 2 mg/125 mg comprimés	DK/H/3047/001	BE294366	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium Duo 2 mg/125 mg comprimés	DK/H/3047/001	BE294341	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium Duo 2 mg/125 mg comprimés	DK/H/3047/001	BE294366	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium Duo 2 mg/125 mg comprimés	DK/H/3047/001	BE294366	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium Duo 2 mg/125 mg comprimés	DK/H/3047/001	BE294366	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium Duo 2 mg/125 mg comprimés	DK/H/3047/001	BE294341	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium Duo 2 mg/125 mg comprimés	DK/H/3047/001	BE294341	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium Duo 2 mg/125 mg comprimés	DK/H/3047/001	BE294341	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE

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Imodium Duo 2 mg/125 mg comprimés	DK/H/3047/001	BE294341	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium Duo 2 mg/125 mg comprimés	DK/H/3047/001	BE294341	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium Duo 2 mg/125 mg comprimés	DK/H/3047/001	BE294366	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium Duo 2 mg/125 mg comprimés	DK/H/3047/001	BE294366	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium Instant 2 mg comprimés orodispersibles	not available	BE 181422	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium Instant 2 mg comprimés orodispersibles	not available	BE 181422	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium Instant 2 mg orodispergeerbare tabletten	not available	BE 181422	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium Instant 2 mg orodispergeerbare tabletten	not available	BE 181422	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium Instant 2 mg Schmelztabletten	not available	BE 181422	JOHNSON & JOHNSON CONSUMER N.V./S.A.	BE
Imodium Instant 2 mg Schmelztabletten	not available	BE 181422	JOHNSON & JOHNSON CONSUMER N.V./S.A.	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
ИМОДИУМ 2 mg капсули	not available	20010160	MCNEIL HEALTHCARE (IRELAND) LIMITED	BG
ИМОДИУМ 2 mg капсули	not available	20010160	MCNEIL HEALTHCARE (IRELAND) LIMITED	BG
Имодиум Плюс 2 mg/125 mg таблетки	DK/H/3047/001/MR	20180231	MCNEIL HEALTHCARE (IRELAND) LIMITED	BG
Имодиум Плюс 2 mg/125 mg таблетки	DK/H/3047/001/MR	20180231	MCNEIL HEALTHCARE (IRELAND) LIMITED	BG
Имодиум Плюс 2 mg/125 mg таблетки	DK/H/3047/001/MR	20180231	MCNEIL HEALTHCARE (IRELAND) LIMITED	BG
Имодиум Плюс 2 mg/125 mg таблетки	DK/H/3047/001/MR	20180231	MCNEIL HEALTHCARE (IRELAND) LIMITED	BG
Имодиум Плюс 2 mg/125 mg таблетки	DK/H/3047/001/MR	20180231	MCNEIL HEALTHCARE (IRELAND) LIMITED	BG
Имодиум Плюс 2 mg/125 mg таблетки	DK/H/3047/001/MR	20180231	MCNEIL HEALTHCARE (IRELAND) LIMITED	BG
Имодиум Плюс 2 mg/125 mg таблетки	DK/H/3047/001/MR	20180231	MCNEIL HEALTHCARE (IRELAND) LIMITED	BG
Имодиум Плюс 2 mg/125 mg таблетки	DK/H/3047/001/MR	20180231	MCNEIL HEALTHCARE (IRELAND) LIMITED	BG
Имодиум Плюс 2 mg/125 mg таблетки	DK/H/3047/001/MR	20180231	MCNEIL HEALTHCARE (IRELAND) LIMITED	BG

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Имодиум Плюс 2 mg/125 mg таблетки	DK/H/3047/001/MR	20180231	MCNEIL HEALTHCARE (IRELAND) LIMITED	BG
Имодиум Плюс 2 mg/125 mg таблетки	DK/H/3047/001/MR	20180231	MCNEIL HEALTHCARE (IRELAND) LIMITED	BG
Имодиум Плюс 2 mg/125 mg таблетки	DK/H/3047/001/MR	20180231	MCNEIL HEALTHCARE (IRELAND) LIMITED	BG
Имодиум Плюс 2 mg/125 mg таблетки	DK/H/3047/001/MR	20180231	MCNEIL HEALTHCARE (IRELAND) LIMITED	BG
Имодиум Плюс 2 mg/125 mg таблетки	DK/H/3047/001/MR	20180231	MCNEIL HEALTHCARE (IRELAND) LIMITED	BG
Имодиум Плюс 2 mg/125 mg таблетки	DK/H/3047/001/MR	20180231	MCNEIL HEALTHCARE (IRELAND) LIMITED	BG
Имодиум Плюс 2 mg/125 mg таблетки	DK/H/3047/001/MR	20180231	MCNEIL HEALTHCARE (IRELAND) LIMITED	BG
Imodium Plus	DK/H/3047/001/MR	022813	JOHNSON & JOHNSON HELLAS CONSUMER AE	CY
Imodium Plus	DK/H/3047/001/MR	022813	JOHNSON & JOHNSON HELLAS CONSUMER AE	CY
Imodium Plus	DK/H/3047/001/MR	022813	JOHNSON & JOHNSON HELLAS CONSUMER AE	CY
Imodium Plus	DK/H/3047/001/MR	022813	JOHNSON & JOHNSON HELLAS CONSUMER AE	CY

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Imodium Plus	DK/H/3047/001/MR	022813	JOHNSON & JOHNSON HELLAS CONSUMER AE	CY
Imodium Plus	DK/H/3047/001/MR	022813	JOHNSON & JOHNSON HELLAS CONSUMER AE	CY
Imodium Plus	DK/H/3047/001/MR	022813	JOHNSON & JOHNSON HELLAS CONSUMER AE	CY
Imodium Plus	DK/H/3047/001/MR	022813	JOHNSON & JOHNSON HELLAS CONSUMER AE	CY
Imodium Plus	DK/H/3047/001/MR	022813	JOHNSON & JOHNSON HELLAS CONSUMER AE	CY
Imodium Plus	DK/H/3047/001/MR	022813	JOHNSON & JOHNSON HELLAS CONSUMER AE	CY
Imodium Plus	DK/H/3047/001/MR	022813	JOHNSON & JOHNSON HELLAS CONSUMER AE	CY
Imodium Plus	DK/H/3047/001/MR	022813	JOHNSON & JOHNSON HELLAS CONSUMER AE	CY
Imodium Plus	DK/H/3047/001/MR	022813	JOHNSON & JOHNSON HELLAS CONSUMER AE	CY
Imodium Plus	DK/H/3047/001/MR	022813	JOHNSON & JOHNSON HELLAS CONSUMER AE	CY
Imodium Plus	DK/H/3047/001/MR	022813	JOHNSON & JOHNSON HELLAS CONSUMER AE	CY

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Imodium Plus	DK/H/3047/001/MR	022813	JOHNSON & JOHNSON HELLAS CONSUMER AE	CY
Imodium™ Capsules	not available	6139	JANSSEN-CILAG INTERNATIONAL NV	CY
Imodium™ Capsules	not available	6139	JANSSEN-CILAG INTERNATIONAL NV	CY
Imodium™ Capsules	not available	6139	JANSSEN-CILAG INTERNATIONAL NV	CY
IMODIUM 2 mg tvrdé tobolky	not available	49/071/92-S/C	MCNEIL HEALTHCARE (IRELAND) LIMITED	CZ
IMODIUM 2 mg tvrdé tobolky	not available	49/071/92-S/C	MCNEIL HEALTHCARE (IRELAND) LIMITED	CZ
Imodium Plus 2 mg/125 mg tablety	DK/H/3047/001/MR	49/432/17-C	MCNEIL HEALTHCARE (IRELAND) LIMITED	CZ
Imodium Plus 2 mg/125 mg tablety	DK/H/3047/001/MR	49/432/17-C	MCNEIL HEALTHCARE (IRELAND) LIMITED	CZ
Imodium Plus 2 mg/125 mg tablety	DK/H/3047/001/MR	49/432/17-C	MCNEIL HEALTHCARE (IRELAND) LIMITED	CZ
Imodium Plus 2 mg/125 mg tablety	DK/H/3047/001/MR	49/432/17-C	MCNEIL HEALTHCARE (IRELAND) LIMITED	CZ
Imodium Plus 2 mg/125 mg tablety	DK/H/3047/001/MR	49/432/17-C	MCNEIL HEALTHCARE (IRELAND) LIMITED	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
Imodium Plus 2 mg/125 mg tablet	DK/H/3047/001/MR	49/432/17-C	MCNEIL HEALTHCARE (IRELAND) LIMITED	CZ
Imodium Plus 2 mg/125 mg tablet	DK/H/3047/001/MR	49/432/17-C	MCNEIL HEALTHCARE (IRELAND) LIMITED	CZ
Imodium Plus 2 mg/125 mg tablet	DK/H/3047/001/MR	49/432/17-C	MCNEIL HEALTHCARE (IRELAND) LIMITED	CZ
Imodium Plus 2 mg/125 mg tablet	DK/H/3047/001/MR	49/432/17-C	MCNEIL HEALTHCARE (IRELAND) LIMITED	CZ
Imodium Plus 2 mg/125 mg tablet	DK/H/3047/001	49/432/17-C	MCNEIL HEALTHCARE (IRELAND) LIMITED	CZ
Imodium Plus 2 mg/125 mg tablet	DK/H/3047/001/MR	49/432/17-C	MCNEIL HEALTHCARE (IRELAND) LIMITED	CZ
Imodium Plus 2 mg/125 mg tablet	DK/H/3047/001/MR	49/432/17-C	MCNEIL HEALTHCARE (IRELAND) LIMITED	CZ
Imodium Plus 2 mg/125 mg tablet	DK/H/3047/001/MR	49/432/17-C	MCNEIL HEALTHCARE (IRELAND) LIMITED	CZ
Imodium Plus 2 mg/125 mg tablet	DK/H/3047/001/MR	49/432/17-C	MCNEIL HEALTHCARE (IRELAND) LIMITED	CZ
Imodium Plus 2 mg/125 mg tablet	DK/H/3047/001/MR	49/432/17-C	MCNEIL HEALTHCARE (IRELAND) LIMITED	CZ
Imodium Plus 2 mg/125 mg tablet	DK/H/3047/001/MR	49/432/17-C	MCNEIL HEALTHCARE (IRELAND) LIMITED	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
IMODIUM RAPID 2 MG tablety dispergovatelne v ústech	not available	49/112/14-C	MCNEIL HEALTHCARE (IRELAND) LIMITED	CZ
IMODIUM RAPID 2 MG tablety dispergovatelne v ústech	not available	49/112/14-C	MCNEIL HEALTHCARE (IRELAND) LIMITED	CZ
IMODIUM RAPID 2 MG tablety dispergovatelne v ústech	not available	49/112/14-C	MCNEIL HEALTHCARE (IRELAND) LIMITED	CZ
IMODIUM RAPID 2 MG tablety dispergovatelne v ústech	not available	49/112/14-C	MCNEIL HEALTHCARE (IRELAND) LIMITED	CZ
IMODIUM 2 mg Hartkapseln	not available	6761963.00.01	JANSSEN-CILAG GMBH	DE
IMODIUM 2 mg Hartkapseln	not available	6761963.00.01	JANSSEN-CILAG GMBH	DE
IMODIUM LINGUAL 2 mg Plättchen (Lyophilisat zum Einnehmen)	not available	34563.00.00	JANSSEN-CILAG GMBH	DE
IMODIUM N 0,2 mg/ml Lösung zum Einnehmen	not available	6761963.00.00	JANSSEN-CILAG GMBH	DE
IMODIUM N 0,2 mg/ml Lösung zum Einnehmen	not available	6761963.00.00	JANSSEN-CILAG GMBH	DE
Imodium® akut 2 mg Hartkapseln	not available	14373.00.00	JOHNSON & JOHNSON GMBH	DE

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Imodium® akut 2 mg Hartkapseln	not available	14373.00.00	JOHNSON & JOHNSON GMBH	DE
Imodium® akut Duo 2 mg/125 mg Tabletten	DK/H/3047/001/MR	65706.00.00	JOHNSON & JOHNSON GMBH	DE
Imodium® akut Duo 2 mg/125 mg Tabletten	DK/H/3047/001/MR	65706.00.00	JOHNSON & JOHNSON GMBH	DE
Imodium® akut Duo 2 mg/125 mg Tabletten	DK/H/3047/001/MR	65706.00.00	JOHNSON & JOHNSON GMBH	DE
Imodium® akut Duo 2 mg/125 mg Tabletten	DK/H/3047/001/MR	65706.00.00	JOHNSON & JOHNSON GMBH	DE
Imodium® akut lingual 2 mg Schmelztabletten	not available	34564.00.00	JOHNSON & JOHNSON GMBH	DE
Imodium® akut lingual 2 mg Schmelztabletten	not available	34564.00.00	JOHNSON & JOHNSON GMBH	DE
Imodium® akut lingual 2 mg Schmelztabletten	not available	34564.00.00	JOHNSON & JOHNSON GMBH	DE
Imodium Plus, tabletter	DK/H/3047/001	39258	MCNEIL DENMARK APS	DK
Imodium Plus, tabletter	DK/H/3047/001	39258	MCNEIL DENMARK APS	DK
Imodium Plus, tabletter	DK/H/3047/001	39258	MCNEIL DENMARK APS	DK

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Imodium Plus, tabletter	DK/H/3047/001	39258	MCNEIL DENMARK APS	DK
Imodium Plus, tabletter	DK/H/3047/001	39258	MCNEIL DENMARK APS	DK
Imodium Plus, tabletter	DK/H/3047/001	39258	MCNEIL DENMARK APS	DK
Imodium Plus, tabletter	DK/H/3047/001	39258	MCNEIL DENMARK APS	DK
Imodium Plus, tabletter	DK/H/3047/001	39258	MCNEIL DENMARK APS	DK
Imodium Plus, tabletter	DK/H/3047/001	39258	MCNEIL DENMARK APS	DK
Imodium Plus, tabletter	DK/H/3047/001	39258	MCNEIL DENMARK APS	DK
Imodium Plus, tabletter	DK/H/3047/001	39258	MCNEIL DENMARK APS	DK
Imodium Plus, tabletter	DK/H/3047/001	39258	MCNEIL DENMARK APS	DK
Imodium Plus, tabletter	DK/H/3047/001	39258	MCNEIL DENMARK APS	DK
Imodium Plus, tabletter	DK/H/3047/001	39258	MCNEIL DENMARK APS	DK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Imodium Plus, tabletter	DK/H/3047/001	39258	MCNEIL DENMARK APS	DK
Imodium Plus, tabletter	DK/H/3047/001	39258	MCNEIL DENMARK APS	DK
Imodium Smelt, frysetørrede tabletter	not available	48356	MCNEIL DENMARK APS	DK
Imodium Smelt, frysetørrede tabletter	not available	48356	MCNEIL DENMARK APS	DK
Imodium Smelt, frysetørrede tabletter	not available	48356	MCNEIL DENMARK APS	DK
Imodium Smelt, frysetørrede tabletter	not available	48356	MCNEIL DENMARK APS	DK
Imodium Smelt, frysetørrede tabletter	not available	48356	MCNEIL DENMARK APS	DK
Imodium, tabletter	not available	13970	MCNEIL DENMARK APS	DK
Imodium, tabletter	not available	13970	MCNEIL DENMARK APS	DK
Imodium, tabletter	not available	13970	MCNEIL DENMARK APS	DK
Imodium instant, 2 mg suus dispergeeruvad tabletid	not available	831013	MCNEIL HEALTHCARE (IRELAND) LIMITED	EE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Imodium instant, 2 mg suus dispergeeruvad tabletid	not available	831013	MCNEIL HEALTHCARE (IRELAND) LIMITED	EE
IMODIUM, 2 mg kõvakapslid	not available	120195	MCNEIL HEALTHCARE (IRELAND) LIMITED	EE
Loperamid-ratiopharm 2 mg, õhukese polümeerikattega tabletid	not available	063894	RATIOPHARM GMBH	EE
Elissan 2 mg comprimidos	not available	55157	LABORATORIOS SERRA PAMIES S.A.	ES
Fortasec 2 mg cápsulas duras	not available	56523	JOHNSON & JOHNSON S.A.	ES
Fortasec 2 mg cápsulas duras	not available	56523	JOHNSON & JOHNSON S.A.	ES
FORTASEC Flas 2 mg liofilizado oral	not available	65349	JOHNSON & JOHNSON S.A.	ES
FORTASEC Flas 2 mg liofilizado oral	not available	65349	JOHNSON & JOHNSON S.A.	ES
IMODIUM DUO 2 mg/ 125 mg comprimidos	DK/H/3047/001/MR	69.021	JOHNSON & JOHNSON S.A.	ES
IMODIUM DUO 2 mg/ 125 mg comprimidos	DK/H/3047/001/MR	69.021	JOHNSON & JOHNSON 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
IMODIUM DUO 2 mg/ 125 mg comprimidos	DK/H/3047/001/MR	69.021	JOHNSON & JOHNSON S.A.	ES
IMODIUM DUO 2 mg/ 125 mg comprimidos	DK/H/3047/001/MR	69.021	JOHNSON & JOHNSON S.A.	ES
IMODIUM DUO 2 mg/ 125 mg comprimidos	DK/H/3047/001/MR	69.021	JOHNSON & JOHNSON S.A.	ES
IMODIUM DUO 2 mg/ 125 mg comprimidos	DK/H/3047/001/MR	69.021	JOHNSON & JOHNSON S.A.	ES
IMODIUM DUO 2 mg/ 125 mg comprimidos	DK/H/3047/001/MR	69.021	JOHNSON & JOHNSON S.A.	ES
IMODIUM DUO 2 mg/ 125 mg comprimidos	DK/H/3047/001/MR	69.021	JOHNSON & JOHNSON S.A.	ES
IMODIUM DUO 2 mg/ 125 mg comprimidos	DK/H/3047/001/MR	69.021	JOHNSON & JOHNSON S.A.	ES
IMODIUM DUO 2 mg/ 125 mg comprimidos	DK/H/3047/001/MR	69.021	JOHNSON & JOHNSON S.A.	ES
IMODIUM DUO 2 mg/ 125 mg comprimidos	DK/H/3047/001/MR	69.021	JOHNSON & JOHNSON S.A.	ES
IMODIUM DUO 2 mg/ 125 mg comprimidos	DK/H/3047/001/MR	69.021	JOHNSON & JOHNSON S.A.	ES
IMODIUM DUO 2 mg/ 125 mg comprimidos	DK/H/3047/001/MR	69.021	JOHNSON & JOHNSON 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
IMODIUM DUO 2 mg/ 125 mg comprimidos	DK/H/3047/001/MR	69.021	JOHNSON & JOHNSON S.A.	ES
IMODIUM DUO 2 mg/ 125 mg comprimidos	DK/H/3047/001/MR	69.021	JOHNSON & JOHNSON S.A.	ES
IMODIUM DUO 2 mg/ 125 mg comprimidos	DK/H/3047/001/MR	69.021	JOHNSON & JOHNSON S.A.	ES
Loperan 2 mg cápsulas duras.	not available	55.317	CHIESI ESPAÑA S.A.U.	ES
Salvacolina 0,2 mg/ml solución oral	not available	36.747	LABORATORIOS SALVAT, S.A.	ES
Salvacolina 2 mg comprimidos	not available	32.202	LABORATORIOS SALVAT, S.A.	ES
Salvacolina Flas 2 mg comprimidos bucodispersables	not available	82401	LABORATORIOS SALVAT, S.A.	ES
Sindiar 2 mg cápsulas duras	not available	57915	TEVA PHARMA S.L.U.,	ES
Imodium 2 mg kapsel, hård	not available	8087	MCNEIL, A DIVISION OF JANSSEN-CILAG OY	FI
Imodium 2 mg kapsel, mjuk	not available	28359	MCNEIL, A DIVISION OF JANSSEN-CILAG OY	FI
Imodium 2 mg kapsel, mjuk	not available	28359	MCNEIL, A DIVISION OF JANSSEN-CILAG OY	FI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Imodium 2 mg kapsel, mjuk	not available	28359	MCNEIL, A DIVISION OF JANSSEN-CILAG OY	FI
Imodium 2 mg kapseli, kova	not available	8087	MCNEIL, A DIVISION OF JANSSEN-CILAG OY	FI
Imodium 2 mg kapseli, pehmeä	not available	28359	MCNEIL, A DIVISION OF JANSSEN-CILAG OY	FI
Imodium 2 mg kapseli, pehmeä	not available	28359	MCNEIL, A DIVISION OF JANSSEN-CILAG OY	FI
Imodium 2 mg kapseli, pehmeä	not available	28359	MCNEIL, A DIVISION OF JANSSEN-CILAG OY	FI
Imodium 2 mg tablett, filmdragerad	not available	11114	MCNEIL, A DIVISION OF JANSSEN-CILAG OY	FI
Imodium 2 mg tablett, filmdragerad	not available	11114	MCNEIL, A DIVISION OF JANSSEN-CILAG OY	FI
Imodium 2 mg tabletti, kalvopäällysteinen	not available	11114	MCNEIL, A DIVISION OF JANSSEN-CILAG OY	FI
Imodium 2 mg tabletti, kalvopäällysteinen	not available	11114	MCNEIL, A DIVISION OF JANSSEN-CILAG OY	FI
Imodium Plus 2 mg/125 mg tabletter	DK/H/3047/001	22042	MCNEIL, A DIVISION OF JANSSEN-CILAG OY	FI
Imodium Plus 2 mg/125 mg tabletter	DK/H/3047/001	22042	MCNEIL, A DIVISION OF JANSSEN-CILAG OY	FI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Imodium Plus 2 mg/125 mg tabletter	DK/H/3047/001	22042	MCNEIL, A DIVISION OF JANSSEN-CILAG OY	FI
Imodium Plus 2 mg/125 mg tabletter	DK/H/3047/001	22042	MCNEIL, A DIVISION OF JANSSEN-CILAG OY	FI
Imodium Plus 2 mg/125 mg tabletter	DK/H/3047/001	22042	MCNEIL, A DIVISION OF JANSSEN-CILAG OY	FI
Imodium Plus 2 mg/125 mg tabletter	DK/H/3047/001	22042	MCNEIL, A DIVISION OF JANSSEN-CILAG OY	FI
Imodium Plus 2 mg/125 mg tabletter	DK/H/3047/001	22042	MCNEIL, A DIVISION OF JANSSEN-CILAG OY	FI
Imodium Plus 2 mg/125 mg tabletter	DK/H/3047/001	22042	MCNEIL, A DIVISION OF JANSSEN-CILAG OY	FI
Imodium Plus 2 mg/125 mg tabletter	DK/H/3047/001	22042	MCNEIL, A DIVISION OF JANSSEN-CILAG OY	FI
Imodium Plus 2 mg/125 mg tabletter	DK/H/3047/001	22042	MCNEIL, A DIVISION OF JANSSEN-CILAG OY	FI
Imodium Plus 2 mg/125 mg tabletter	DK/H/3047/001	22042	MCNEIL, A DIVISION OF JANSSEN-CILAG OY	FI
Imodium Plus 2 mg/125 mg tabletter	DK/H/3047/001	22042	MCNEIL, A DIVISION OF JANSSEN-CILAG OY	FI
Imodium Plus 2 mg/125 mg tabletter	DK/H/3047/001	22042	MCNEIL, A DIVISION OF JANSSEN-CILAG OY	FI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Imodium Plus 2 mg/125 mg tabletter	DK/H/3047/001	22042	MCNEIL, A DIVISION OF JANSSEN-CILAG OY	FI
Imodium Plus 2 mg/125 mg tabletter	DK/H/3047/001	22042	MCNEIL, A DIVISION OF JANSSEN-CILAG OY	FI
Imodium Plus 2 mg/125 mg tabletter	DK/H/3047/001	22042	MCNEIL, A DIVISION OF JANSSEN-CILAG OY	FI
Imodium Plus 2 mg/125 mg tabletti	DK/H/3047/001	22042	MCNEIL, A DIVISION OF JANSSEN-CILAG OY	FI
Imodium Plus 2 mg/125 mg tabletti	DK/H/3047/001	22042	MCNEIL, A DIVISION OF JANSSEN-CILAG OY	FI
Imodium Plus 2 mg/125 mg tabletti	DK/H/3047/001	22042	MCNEIL, A DIVISION OF JANSSEN-CILAG OY	FI
Imodium Plus 2 mg/125 mg tabletti	DK/H/3047/001	22042	MCNEIL, A DIVISION OF JANSSEN-CILAG OY	FI
Imodium Plus 2 mg/125 mg tabletti	DK/H/3047/001	22042	MCNEIL, A DIVISION OF JANSSEN-CILAG OY	FI
Imodium Plus 2 mg/125 mg tabletti	DK/H/3047/001	22042	MCNEIL, A DIVISION OF JANSSEN-CILAG OY	FI
Imodium Plus 2 mg/125 mg tabletti	DK/H/3047/001	22042	MCNEIL, A DIVISION OF JANSSEN-CILAG OY	FI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Imodium Plus 2 mg/125 mg tabletti	DK/H/3047/001	22042	MCNEIL, A DIVISION OF JANSSEN-CILAG OY	FI
Imodium Plus 2 mg/125 mg tabletti	DK/H/3047/001	22042	MCNEIL, A DIVISION OF JANSSEN-CILAG OY	FI
Imodium Plus 2 mg/125 mg tabletti	DK/H/3047/001	22042	MCNEIL, A DIVISION OF JANSSEN-CILAG OY	FI
Imodium Plus 2 mg/125 mg tabletti	DK/H/3047/001	22042	MCNEIL, A DIVISION OF JANSSEN-CILAG OY	FI
Imodium Plus 2 mg/125 mg tabletti	DK/H/3047/001	22042	MCNEIL, A DIVISION OF JANSSEN-CILAG OY	FI
Imodium Plus 2 mg/125 mg tabletti	DK/H/3047/001	22042	MCNEIL, A DIVISION OF JANSSEN-CILAG OY	FI
Imodium Plus 2 mg/125 mg tabletti	DK/H/3047/001	22042	MCNEIL, A DIVISION OF JANSSEN-CILAG OY	FI
Imodium Plus 2 mg/125 mg tabletti	DK/H/3047/001	22042	MCNEIL, A DIVISION OF JANSSEN-CILAG OY	FI
DIARETYL 2 mg, gélule	not available	34009 358 421 4 5	COOPERATION PHARMACEUTIQUE FRANCAISE	FR
DIARETYL 2 mg, gélule	not available	34009 341 154 8 6	COOPERATION PHARMACEUTIQUE FRANCAISE	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
DIASTROLIB 2 mg, lyophilisat oral	not available	NL 31999	TEVA SANTÉ	FR
IMODIUM 0,2 mg/ml ENFANTS, solution buvable	not available	34009 300 798 4 3	JANSSEN-CILAG	FR
IMODIUM 0,2 mg/ml ENFANTS, solution buvable	not available	34009 318 865 9 4	JANSSEN-CILAG	FR
IMODIUM 2 mg, gélule	not available	34009 362 197 8 6	JANSSEN-CILAG	FR
IMODIUM 2 mg, gélule	not available	34009 318 860 7 5	JANSSEN-CILAG	FR
IMODIUM 2 mg, gélule	not available	34009 339 930 4 7	JANSSEN-CILAG	FR
IMODIUM 2 mg, gélule	not available	34009 362 197 8 6	JANSSEN-CILAG	FR
IMODIUM 2 mg, gélule	not available	34009 318 860 7 5	JANSSEN-CILAG	FR
IMODIUMCAPS 2 mg, gélule	not available	34009 385 718 4 4	JOHNSON & JOHNSON SANTÉ BEAUTÉ FRANCE	FR
IMODIUMCAPS 2 mg, gélule	not available	34009 385 719 0 5	JOHNSON & JOHNSON SANTÉ BEAUTÉ 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
IMODIUMDUO, comprimé	DK/H/3047/001/MR	34009 378 819 3 7	JOHNSON & JOHNSON SANTÉ BEAUTÉ FRANCE	FR
IMODIUMDUO, comprimé	DK/H/3047/001/MR	34009 378 822 4 8	JOHNSON & JOHNSON SANTÉ BEAUTÉ FRANCE	FR
IMODIUMDUO, comprimé	DK/H/3047/001/MR	34009 378 823 0 9	JOHNSON & JOHNSON SANTÉ BEAUTÉ FRANCE	FR
IMODIUMDUO, comprimé	DK/H/3047/001/MR	34009 378 819 3 7	JOHNSON & JOHNSON SANTÉ BEAUTÉ FRANCE	FR
IMODIUMDUO, comprimé	DK/H/3047/001/MR	34009 378 821 8 7	JOHNSON & JOHNSON SANTÉ BEAUTÉ FRANCE	FR
IMODIUMDUO, comprimé	DK/H/3047/001/MR	34009 378 822 4 8	JOHNSON & JOHNSON SANTÉ BEAUTÉ FRANCE	FR
IMODIUMDUO, comprimé	DK/H/3047/001/MR	34009 378 825 3 8	JOHNSON & JOHNSON SANTÉ BEAUTÉ FRANCE	FR
IMODIUMDUO, comprimé	DK/H/3047/001/MR	34009 378 823 0 9	JOHNSON & JOHNSON SANTÉ BEAUTÉ FRANCE	FR
IMODIUMDUO, comprimé	DK/H/3047/001/MR	34009 378 821 8 7	JOHNSON & JOHNSON SANTÉ BEAUTÉ FRANCE	FR
IMODIUMDUO, comprimé	DK/H/3047/001/MR	34009 378 824 7 7	JOHNSON & JOHNSON SANTÉ BEAUTÉ FRANCE	FR
IMODIUMDUO, comprimé	DK/H/3047/001/MR	34009 378 825 3 8	JOHNSON & JOHNSON SANTÉ BEAUTÉ 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
IMODIUMDUO, comprimé	DK/H/3047/001/MR	34009 378 820 1 9	JOHNSON & JOHNSON SANTÉ BEAUTÉ FRANCE	FR
IMODIUMDUO, comprimé	DK/H/3047/001/MR	34009 378 824 7 7	JOHNSON & JOHNSON SANTÉ BEAUTÉ FRANCE	FR
IMODIUMDUO, comprimé	DK/H/3047/001/MR	34009 378 827 6 7	JOHNSON & JOHNSON SANTÉ BEAUTÉ FRANCE	FR
IMODIUMDUO, comprimé	DK/H/3047/001/MR	34009 378 820 1 9	JOHNSON & JOHNSON SANTÉ BEAUTÉ FRANCE	FR
IMODIUMDUO, comprimé	DK/H/3047/001/MR	34009 378 827 6 7	JOHNSON & JOHNSON SANTÉ BEAUTÉ FRANCE	FR
IMODIUMLIQUICAPS 2 mg, capsule molle	not available	34009 218 008 7 6	JOHNSON & JOHNSON SANTÉ BEAUTÉ FRANCE	FR
IMODIUMLIQUICAPS 2 mg, capsule molle	not available	34009 218 010 1 9	JOHNSON & JOHNSON SANTÉ BEAUTÉ FRANCE	FR
IMODIUMLIQUICAPS 2 mg, capsule molle	not available	34009 218 009 3 7	JOHNSON & JOHNSON SANTÉ BEAUTÉ FRANCE	FR
LOPERAMIDE LYOC 2 mg, lyophilisat oral	not available	NL22126	TEVA SANTÉ	FR
Lopéramide Mylan 2 mg, gélule	not available	NL 21216	MYLAN S.A.S	FR
Imodium Plus	DK/H/3047/001	92817/17/12-02-2019	JOHNSON & JOHNSON HELLAS CONSUMER AE	GR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Imodium Plus	DK/H/3047/001	92817/17/12-02-2019	JOHNSON & JOHNSON HELLAS CONSUMER AE	GR
Imodium Plus	DK/H/3047/001	92817/17/12-02-2019	JOHNSON & JOHNSON HELLAS CONSUMER AE	GR
Imodium Plus	DK/H/3047/001	92817/17/12-02-2019	JOHNSON & JOHNSON HELLAS CONSUMER AE	GR
Imodium Plus	DK/H/3047/001	92817/17/12-02-2019	JOHNSON & JOHNSON HELLAS CONSUMER AE	GR
Imodium Plus	DK/H/3047/001	92817/17/12-02-2019	JOHNSON & JOHNSON HELLAS CONSUMER AE	GR
Imodium Plus	DK/H/3047/001	92817/17/12-02-2019	JOHNSON & JOHNSON HELLAS CONSUMER AE	GR
Imodium Plus	DK/H/3047/001	92817/17/12-02-2019	JOHNSON & JOHNSON HELLAS CONSUMER AE	GR
Imodium Plus	DK/H/3047/001	92817/17/12-02-2019	JOHNSON & JOHNSON HELLAS CONSUMER AE	GR
Imodium Plus	DK/H/3047/001	92817/17/12-02-2019	JOHNSON & JOHNSON HELLAS CONSUMER AE	GR
Imodium Plus	DK/H/3047/001	92817/17/12-02-2019	JOHNSON & JOHNSON HELLAS CONSUMER AE	GR
Imodium Plus	DK/H/3047/001	92817/17/12-02-2019	JOHNSON & JOHNSON HELLAS CONSUMER AE	GR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Imodium Plus	DK/H/3047/001	92817/17/12-02-2019	JOHNSON & JOHNSON HELLAS CONSUMER AE	GR
Imodium Plus	DK/H/3047/001	92817/17/12-02-2019	JOHNSON & JOHNSON HELLAS CONSUMER AE	GR
Imodium Plus	DK/H/3047/001	92817/17/12-02-2019	JOHNSON & JOHNSON HELLAS CONSUMER AE	GR
Imodium Plus	DK/H/3047/001	92817/17/12-02-2019	JOHNSON & JOHNSON HELLAS CONSUMER AE	GR
IMODIUM® INSTANT	not available	70410/18/18-12-2019	JOHNSON & JOHNSON HELLAS CONSUMER AE	GR
IMODIUM® ORIGINAL	not available	72014/17/25-1-2018	JOHNSON & JOHNSON HELLAS CONSUMER AE	GR
Imodium 2 mg raspadljive tablete za usta	not available	HR-H-493726499	JOHNSON & JOHNSON S.E.D.O.O.	HR
Imodium COMBO 2 mg/125 mg tablete	not available	HR-H-849757099	JOHNSON & JOHNSON S.E.D.O.O.	HR
Imodium 2 mg kemény kapszula	not available	OGYI-T-2221/01	JOHNSON & JOHNSON KFT.	HU
Imodium 2 mg kemény kapszula	not available	OGYI-T-2221/02	JOHNSON & JOHNSON KFT.	HU
Imodium Instant 2 mg szájban diszpergálódó tabletta	not available	OGYI-T-02221/03-04-05-06	JOHNSON & JOHNSON KFT.	HU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Imodium Instant 2 mg szájban diszpergálódó tabletta	not available	OGYI-T-02221/03-04-05-06	JOHNSON & JOHNSON KFT.	HU
Imodium Instant 2 mg szájban diszpergálódó tabletta	not available	OGYI-T-02221/03-04-05-06	JOHNSON & JOHNSON KFT.	HU
Imodium Instant 2 mg szájban diszpergálódó tabletta	not available	OGYI-T-02221/03-04-05-06	JOHNSON & JOHNSON KFT.	HU
Imodium Plus 2 mg/125 mg tabletta	DK/H/3047/001/MR	OGYI-T-23360/01-16	JOHNSON & JOHNSON KFT.	HU
Imodium Plus 2 mg/125 mg tabletta	DK/H/3047/001/MR	OGYI-T-23360/01-16	JOHNSON & JOHNSON KFT.	HU
Imodium Plus 2 mg/125 mg tabletta	DK/H/3047/001/MR	OGYI-T-23360/01-16	JOHNSON & JOHNSON KFT.	HU
Imodium Plus 2 mg/125 mg tabletta	DK/H/3047/001/MR	OGYI-T-23360/01-16	JOHNSON & JOHNSON KFT.	HU
Imodium Plus 2 mg/125 mg tabletta	DK/H/3047/001/MR	OGYI-T-23360/01-16	JOHNSON & JOHNSON KFT.	HU
Imodium Plus 2 mg/125 mg tabletta	DK/H/3047/001/MR	OGYI-T-23360/01-16	JOHNSON & JOHNSON KFT.	HU
Imodium Plus 2 mg/125 mg tabletta	DK/H/3047/001/MR	OGYI-T-23360/01-16	JOHNSON & JOHNSON KFT.	HU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Imodium Plus 2 mg/125 mg tabletta	DK/H/3047/001/MR	OGYI-T-23360/01-16	JOHNSON & JOHNSON KFT.	HU
Imodium Plus 2 mg/125 mg tabletta	DK/H/3047/001/MR	OGYI-T-23360/01-16	JOHNSON & JOHNSON KFT.	HU
Imodium Plus 2 mg/125 mg tabletta	DK/H/3047/001/MR	OGYI-T-23360/01-16	JOHNSON & JOHNSON KFT.	HU
Imodium Plus 2 mg/125 mg tabletta	DK/H/3047/001/MR	OGYI-T-23360/01-16	JOHNSON & JOHNSON KFT.	HU
Imodium Plus 2 mg/125 mg tabletta	DK/H/3047/001/MR	OGYI-T-23360/01-16	JOHNSON & JOHNSON KFT.	HU
Imodium Plus 2 mg/125 mg tabletta	DK/H/3047/001/MR	OGYI-T-23360/01-16	JOHNSON & JOHNSON KFT.	HU
Imodium Plus 2 mg/125 mg tabletta	DK/H/3047/001/MR	OGYI-T-23360/01-16	JOHNSON & JOHNSON KFT.	HU
Imodium Plus 2 mg/125 mg tabletta	DK/H/3047/001/MR	OGYI-T-23360/01-16	JOHNSON & JOHNSON KFT.	HU
Imodium Plus 2 mg/125 mg tabletta	DK/H/3047/001/MR	OGYI-T-23360/01-16	JOHNSON & JOHNSON KFT.	HU
Imodium Plus 2 mg/125 mg tabletta	DK/H/3047/001/MR	OGYI-T-23360/01-16	JOHNSON & JOHNSON KFT.	HU
Imorevin 2 mg/125 mg tabletta	NL/H/2663/001	OGYI-T-22515/01	SAGER PHARMA KFT.	HU
Imorevin 2 mg/125 mg tabletta	NL/H/2663/001	OGYI-T-22515/02	SAGER PHARMA KFT.	HU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Imorevin 2 mg/125 mg tabletta	NL/H/2663/001	OGYI-T-22515/03	SAGER PHARMA KFT.	HU
Imorevin 2 mg/125 mg tabletta	NL/H/2663/001	OGYI-T-22515/04	SAGER PHARMA KFT.	HU
Imorevin 2 mg/125 mg tabletta	NL/H/2663/001	OGYI-T-22515/05	SAGER PHARMA KFT.	HU
Imorevin 2 mg/125 mg tabletta	NL/H/2663/001	OGYI-T-22515/06	SAGER PHARMA KFT.	HU
Imorevin 2 mg/125 mg tabletta	NL/H/2663/001	OGYI-T-22515/07	SAGER PHARMA KFT.	HU
Imorevin 2 mg/125 mg tabletta	NL/H/2663/001	OGYI-T-22515/08	SAGER PHARMA KFT.	HU
Imorevin 2 mg/125 mg tabletta	NL/H/2663/001	OGYI-T-22515/09	SAGER PHARMA KFT.	HU
Imorevin 2 mg/125 mg tabletta	NL/H/2663/001	OGYI-T-22515/10	SAGER PHARMA KFT.	HU
Imorevin 2 mg/125 mg tabletta	NL/H/2663/001	OGYI-T-22515/11	SAGER PHARMA KFT.	HU
Imorevin 2 mg/125 mg tabletta	NL/H/2663/001	OGYI-T-22515/12	SAGER PHARMA KFT.	HU
Imorevin 2 mg/125 mg tabletta	NL/H/2663/001	OGYI-T-22515/13	SAGER PHARMA KFT.	HU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Imorevin 2 mg/125 mg tabletta	NL/H/2663/001	OGYI-T-22515/14	SAGER PHARMA KFT.	HU
Imorevin 2 mg/125 mg tabletta	NL/H/2663/001	OGYI-T-22515/15	SAGER PHARMA KFT.	HU
Imorevin 2 mg/125 mg tabletta	NL/H/2663/001	OGYI-T-22515/16	SAGER PHARMA KFT.	HU
Imorevin 2 mg/125 mg tabletta	NL/H/2663/001	OGYI-T-22515/17	SAGER PHARMA KFT.	HU
Imorevin 2 mg/125 mg tabletta	NL/H/2663/001	OGYI-T-22515/18	SAGER PHARMA KFT.	HU
Arret 2mg Hard Capsules	not available	PA 330/42/1	JOHNSON & JOHNSON (IRELAND) LTD.	IE
Arret 2mg Hard Capsules	not available	PA 330/42/1	JOHNSON & JOHNSON (IRELAND) LTD.	IE
Arret 2mg Hard Capsules	not available	PA 330/42/1	JOHNSON & JOHNSON (IRELAND) LTD.	IE
Arret 2mg Hard Capsules	not available	PA 330/42/1	JOHNSON & JOHNSON (IRELAND) LTD.	IE
Boots Pharmaceuticals Diarrhoea Relief 2 mg Hard Capsules	not available	PA 23081/002/001	TAW PHARMA (IRELAND) LTD	IE
Imodium 2 mg Capsules	not available	PA 330/45/2	JOHNSON & JOHNSON (IRELAND) LTD.	IE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Imodium Instants 2 mg Orodispersible Tablets	not available	PA 330/45/1	JOHNSON & JOHNSON (IRELAND) LTD.	IE
Imodium Instants 2 mg Orodispersible Tablets	not available	PA 330/45/1	JOHNSON & JOHNSON (IRELAND) LTD.	IE
Imodium Instants 2 mg Orodispersible Tablets	not available	PA 330/45/1	JOHNSON & JOHNSON (IRELAND) LTD.	IE
Imodium Instants 2 mg Orodispersible Tablets	not available	PA 330/45/1	JOHNSON & JOHNSON (IRELAND) LTD.	IE
Imodium Plus 2mg/125mg Tablets	DK/H/3047/001/MR	PA 330/50/1	JOHNSON & JOHNSON (IRELAND) LTD.	IE
Imodium Plus 2mg/125mg Tablets	DK/H/3047/001/MR	PA 330/50/1	JOHNSON & JOHNSON (IRELAND) LTD.	IE
Imodium Plus 2mg/125mg Tablets	DK/H/3047/001/MR	PA 330/50/1	JOHNSON & JOHNSON (IRELAND) LTD.	IE
Imodium Plus 2mg/125mg Tablets	DK/H/3047/001/MR	PA 330/50/1	JOHNSON & JOHNSON (IRELAND) LTD.	IE
Imodium Plus 2mg/125mg Tablets	DK/H/3047/001/MR	PA 330/50/1	JOHNSON & JOHNSON (IRELAND) LTD.	IE
Imodium Plus 2mg/125mg Tablets	DK/H/3047/001/MR	PA 330/50/1	JOHNSON & JOHNSON (IRELAND) LTD.	IE
Imodium Plus 2mg/125mg Tablets	DK/H/3047/001/MR	PA 330/50/1	JOHNSON & JOHNSON (IRELAND) LTD.	IE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Imodium Plus 2mg/125mg Tablets	DK/H/3047/001/MR	PA 330/50/1	JOHNSON & JOHNSON (IRELAND) LTD.	IE
Imodium Plus 2mg/125mg Tablets	DK/H/3047/001/MR	PA 330/50/1	JOHNSON & JOHNSON (IRELAND) LTD.	IE
Imodium Plus 2mg/125mg Tablets	DK/H/3047/001/MR	PA 330/50/1	JOHNSON & JOHNSON (IRELAND) LTD.	IE
Imodium Plus 2mg/125mg Tablets	DK/H/3047/001/MR	PA 330/50/1	JOHNSON & JOHNSON (IRELAND) LTD.	IE
Imodium Plus 2mg/125mg Tablets	DK/H/3047/001/MR	PA 330/50/1	JOHNSON & JOHNSON (IRELAND) LTD.	IE
Imodium Plus 2mg/125mg Tablets	DK/H/3047/001/MR	PA 330/50/1	JOHNSON & JOHNSON (IRELAND) LTD.	IE
Imodium Plus 2mg/125mg Tablets	DK/H/3047/001/MR	PA 330/50/1	JOHNSON & JOHNSON (IRELAND) LTD.	IE
Imodium Plus 2mg/125mg Tablets	DK/H/3047/001/MR	PA 330/50/1	JOHNSON & JOHNSON (IRELAND) LTD.	IE
Imodium Plus 2mg/125mg Tablets	DK/H/3047/001/MR	PA 330/50/1	JOHNSON & JOHNSON (IRELAND) LTD.	IE
Imodium® LiquiRelief 2mg Soft Capsules	not available	PA 330/45/3	JOHNSON & JOHNSON (IRELAND) LTD.	IE
Imodium 2 mg töflur	not available	MTNR 900168 (IS)	MCNEIL SWEDEN AB	IS

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
Imodium 2 mg töflur	not available	MTNR 900168 (IS)	MCNEIL SWEDEN AB	IS
Imodium 2 mg töflur	not available	MTNR 900168 (IS)	MCNEIL SWEDEN AB	IS
Imodium 2 mg töflur	not available	MTNR 900168 (IS)	MCNEIL SWEDEN AB	IS
DISSENTEN 2 mg compresse	not available	023694021	SPA – SOCIETÀ PRODOTTI ANTIBIOTICI S.P.A.	IT
DISSENTEN 2 mg compresse	not available	023694058	SPA – SOCIETÀ PRODOTTI ANTIBIOTICI S.P.A.	IT
DISSENTEN ANTIDIARREA 2 mg compresse	not available	043905013	SPA – SOCIETÀ PRODOTTI ANTIBIOTICI S.P.A.	IT
ENTEROG ANTIDIARROICO	not available	026112033	SANOFI S.R.L.	IT
IMODIUM 2 mg capsule molli	not available	023673104	JOHNSON & JOHNSON S.P.A.	IT
IMODIUM 2 mg capsule rigide	not available	023673066	JOHNSON & JOHNSON S.P.A.	IT
Imodium 2 mg compresse orosolubili	not available	023673092	JOHNSON & JOHNSON S.P.A.	IT
Lopemid 2mg capsule rigide	not available	023691013	VISUFARMA SPA	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
Losipaco 2 mg/125 mg compresse	NL/H/2666/001	042141034	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Losipaco 2 mg/125 mg compresse	NL/H/2666/001	042141046	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Losipaco 2 mg/125 mg compresse	NL/H/2666/001	042141010	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Losipaco 2 mg/125 mg compresse	NL/H/2666/001	042141022	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Losipaco 2 mg/125 mg compresse	NL/H/2666/001	042141059	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Losipaco 2 mg/125 mg compresse	NL/H/2666/001	042141061	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	IT
Aloper 2 mg/125 mg tabletės	not available	LT/1/16/3919/001	SIA INGEN PHARMA	LT
Aloper 2 mg/125 mg tabletės	not available	LT/1/16/3919/002	SIA INGEN PHARMA	LT
Imodium 2 mg kietosios kapsulės	not available	LT/1/93/0292/001	MCNEIL HEALTHCARE (IRELAND) LIMITED	LT
Imodium instant 2 mg burnoje disperguojamos tabletės	not available	LT/1/01/1997/001-002	MCNEIL HEALTHCARE (IRELAND) LIMITED	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
Imodium instant 2 mg burnoje disperguojamos tabletės	not available	LT/1/01/1997/001-002	MCNEIL HEALTHCARE (IRELAND) LIMITED	LT
Imodium 0,2 mg/ml solution buvable	not available	2008069794	JOHNSON & JOHNSON CONSUMER N.V./S.A.	LU
Imodium 2 mg gélules	not available	2008069795	JOHNSON & JOHNSON CONSUMER N.V./S.A.	LU
Imodium 2 mg gélules	not available	2008069795	JOHNSON & JOHNSON CONSUMER N.V./S.A.	LU
Imodium 2 mg gélules	not available	2008069795	JOHNSON & JOHNSON CONSUMER N.V./S.A.	LU
Imodium Duo 2 mg/125 mg comprimés	DK/H/3047/001/MR	2008010024	JOHNSON & JOHNSON CONSUMER N.V./S.A.	LU
Imodium Duo 2 mg/125 mg comprimés	DK/H/3047/001/MR	2008010024	JOHNSON & JOHNSON CONSUMER N.V./S.A.	LU
Imodium Duo 2 mg/125 mg comprimés	DK/H/3047/001/MR	2008010024	JOHNSON & JOHNSON CONSUMER N.V./S.A.	LU
Imodium Duo 2 mg/125 mg comprimés	DK/H/3047/001/MR	2008010024	JOHNSON & JOHNSON CONSUMER N.V./S.A.	LU
Imodium Duo 2 mg/125 mg comprimés	DK/H/3047/001/MR	2008010024	JOHNSON & JOHNSON CONSUMER N.V./S.A.	LU
Imodium Duo 2 mg/125 mg comprimés	DK/H/3047/001/MR	2008010024	JOHNSON & JOHNSON CONSUMER N.V./S.A.	LU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Imodium Duo 2 mg/125 mg comprimés	DK/H/3047/001/MR	2008010024	JOHNSON & JOHNSON CONSUMER N.V./S.A.	LU
Imodium Duo 2 mg/125 mg comprimés	DK/H/3047/001/MR	2008010024	JOHNSON & JOHNSON CONSUMER N.V./S.A.	LU
Imodium Duo 2 mg/125 mg comprimés	DK/H/3047/001/MR	2008010024	JOHNSON & JOHNSON CONSUMER N.V./S.A.	LU
Imodium Duo 2 mg/125 mg comprimés	DK/H/3047/001/MR	2008010024	JOHNSON & JOHNSON CONSUMER N.V./S.A.	LU
Imodium Duo 2 mg/125 mg comprimés	DK/H/3047/001/MR	2008010024	JOHNSON & JOHNSON CONSUMER N.V./S.A.	LU
Imodium Duo 2 mg/125 mg comprimés	DK/H/3047/001/MR	2008010024	JOHNSON & JOHNSON CONSUMER N.V./S.A.	LU
Imodium Duo 2 mg/125 mg comprimés	DK/H/3047/001/MR	2008010024	JOHNSON & JOHNSON CONSUMER N.V./S.A.	LU
Imodium Duo 2 mg/125 mg comprimés	DK/H/3047/001/MR	2008010024	JOHNSON & JOHNSON CONSUMER N.V./S.A.	LU
Imodium Duo 2 mg/125 mg comprimés	DK/H/3047/001/MR	2008010024	JOHNSON & JOHNSON CONSUMER N.V./S.A.	LU
Imodium Duo 2 mg/125 mg comprimés	DK/H/3047/001/MR	2008010024	JOHNSON & JOHNSON CONSUMER N.V./S.A.	LU
Imodium Duo 2 mg/125 mg comprimés	DK/H/3047/001/MR	2008010024	JOHNSON & JOHNSON CONSUMER N.V./S.A.	LU
Imodium Duo 2 mg/125 mg comprimés	DK/H/3047/001/MR	2008010024	JOHNSON & JOHNSON CONSUMER N.V./S.A.	LU
Imodium Duo 2 mg/125 mg comprimés	DK/H/3047/001/MR	2008010024	JOHNSON & JOHNSON CONSUMER N.V./S.A.	LU
Imodium Instant 2 mg comprimés orodispersibles	not available	2008069796	JOHNSON & JOHNSON CONSUMER N.V./S.A.	LU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Imodium Instant 2 mg comprimés orodispersibles	not available	2008069796	JOHNSON & JOHNSON CONSUMER N.V./S.A.	LU
IMODIUM 2 mg cietās kapsulas N6	not available	98-0799	MCNEIL HEALTHCARE (IRELAND) LIMITED	LV
IMODIUM INSTANT 2 mg mutē disperģējamās tabletes	not available	02-0131	MCNEIL HEALTHCARE (IRELAND) LIMITED	LV
IMODIUM INSTANT 2 mg mutē disperģējamās tabletes	not available	02-0131	MCNEIL HEALTHCARE (IRELAND) LIMITED	LV
Imodium™ Capsules	not available	018/01401	JANSSEN-CILAG INTERNATIONAL NV	MT
Imodium™ Capsules	not available	018/01401	JANSSEN-CILAG INTERNATIONAL NV	MT
Imodium 0,2 mg/ml drank	not available	RVG 08423	JOHNSON & JOHNSON CONSUMER B.V.	NL
Imodium 0,2 mg/ml drank	not available	RVG 08423	JOHNSON & JOHNSON CONSUMER B.V.	NL
Imodium 2 mg harde capsules	not available	RVG 06945	JOHNSON & JOHNSON CONSUMER B.V.	NL
Imodium 2 mg harde capsules	not available	RVG 06945	JOHNSON & JOHNSON CONSUMER B.V.	NL

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Imodium 2 mg harde capsules	not available	RVG 06945	JOHNSON & JOHNSON CONSUMER B.V.	NL
Imodium 2 mg harde capsules	not available	RVG 06945	JOHNSON & JOHNSON CONSUMER B.V.	NL
Imodium Duo 2 mg/125 mg, tabletten	DK/H/3047/001/MR	RVG 33869	JOHNSON & JOHNSON CONSUMER B.V.	NL
Imodium Duo 2 mg/125 mg, tabletten	DK/H/3047/001/MR	RVG 33869	JOHNSON & JOHNSON CONSUMER B.V.	NL
Imodium Duo 2 mg/125 mg, tabletten	DK/H/3047/001/MR	RVG 33869	JOHNSON & JOHNSON CONSUMER B.V.	NL
Imodium Duo 2 mg/125 mg, tabletten	DK/H/3047/001/MR	RVG 33869	JOHNSON & JOHNSON CONSUMER B.V.	NL
Imodium Duo 2 mg/125 mg, tabletten	DK/H/3047/001/MR	RVG 33869	JOHNSON & JOHNSON CONSUMER B.V.	NL
Imodium Duo 2 mg/125 mg, tabletten	DK/H/3047/001/MR	RVG 33869	JOHNSON & JOHNSON CONSUMER B.V.	NL
Imodium Duo 2 mg/125 mg, tabletten	DK/H/3047/001/MR	RVG 33869	JOHNSON & JOHNSON CONSUMER B.V.	NL
Imodium Duo 2 mg/125 mg, tabletten	DK/H/3047/001/MR	RVG 33869	JOHNSON & JOHNSON CONSUMER B.V.	NL
Imodium Duo 2 mg/125 mg, tabletten	DK/H/3047/001/MR	RVG 33869	JOHNSON & JOHNSON CONSUMER B.V.	NL

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Imodium Duo 2 mg/125 mg, tabletten	DK/H/3047/001/MR	RVG 33869	JOHNSON & JOHNSON CONSUMER B.V.	NL
Imodium Duo 2 mg/125 mg, tabletten	DK/H/3047/001/MR	RVG 33869	JOHNSON & JOHNSON CONSUMER B.V.	NL
Imodium Duo 2 mg/125 mg, tabletten	DK/H/3047/001/MR	RVG 33869	JOHNSON & JOHNSON CONSUMER B.V.	NL
Imodium Duo 2 mg/125 mg, tabletten	DK/H/3047/001/MR	RVG 33869	JOHNSON & JOHNSON CONSUMER B.V.	NL
Imodium Duo 2 mg/125 mg, tabletten	DK/H/3047/001/MR	RVG 33869	JOHNSON & JOHNSON CONSUMER B.V.	NL
Imodium Duo 2 mg/125 mg, tabletten	DK/H/3047/001/MR	RVG 33869	JOHNSON & JOHNSON CONSUMER B.V.	NL
Imodium Duo 2 mg/125 mg, tabletten	DK/H/3047/001/MR	RVG 33869	JOHNSON & JOHNSON CONSUMER B.V.	NL
Imodium Instant smelttablet 2 mg orodispergeerbare tabletten	not available	RVG 33724	JOHNSON & JOHNSON CONSUMER B.V.	NL
Imodium Instant smelttablet 2 mg orodispergeerbare tabletten	not available	RVG 33724	JOHNSON & JOHNSON CONSUMER B.V.	NL
Kruidvat diarreeremmer loperamide HCl 2 mg	not available	RVG 125673	MAREL B.V.	NL

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
smelttablet, orodispergeerbare tabletten				
Loperamide HCl Disphar 2 mg orodispergeerbare tabletten	not available	RVG 119415	DISPHAR INTERNATIONAL B.V.	NL
Loperamide HCl RXT 2 mg diarreeremmende capsules, capsules	not available	RVG 56839	RXT	NL
Losibere 2 mg / 125 mg, tabletten	NL/H/2730/001	RVG 111976	DISPHAR INTERNATIONAL B.V.	NL
Losidaru 2 mg / 125 mg, tabletten	NL/H/2907/001	RVG 113191	DISPHAR INTERNATIONAL B.V.	NL
LOSIMED DUO 2 mg / 125 mg, tabletten	NL/H/2665/001	RVG 111984	DAVANTIS HEALTH B.V.	NL
Losimido 2 mg / 125 mg tabletten	NL/H/2663/001	RVG 111982	DISPHAR INTERNATIONAL B.V.	NL
Losipaco 2 mg/125 mg, tabletten	NL/H/2666/001	RVG 111985	RECORDATI INDUSTRIA CHIMICA E FARMACEUTICA S.P.A.	NL
Imodium	not available	7633	MCNEIL SWEDEN AB	NO
Imodium	not available	7633	MCNEIL SWEDEN AB	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
Imodium	not available	7633	MCNEIL SWEDEN AB	NO
Imodium	not available	7253	MCNEIL SWEDEN AB	NO
Imodium Comp 2 mg / 125 mg tabletter	DK/H/3047/001/MR	06-4059	MCNEIL SWEDEN AB	NO
Imodium Comp 2 mg / 125 mg tabletter	DK/H/3047/001	06-4059	MCNEIL SWEDEN AB	NO
Imodium Comp 2 mg / 125 mg tabletter	DK/H/3047/001	06-4059	MCNEIL SWEDEN AB	NO
Imodium Comp 2 mg / 125 mg tabletter	DK/H/3047/001	06-4059	MCNEIL SWEDEN AB	NO
Imodium Comp 2 mg / 125 mg tabletter	DK/H/3047/001	06-4059	MCNEIL SWEDEN AB	NO
Imodium Comp 2 mg / 125 mg tabletter	DK/H/3047/001	06-4059	MCNEIL SWEDEN AB	NO
Imodium Comp 2 mg / 125 mg tabletter	DK/H/3047/001	06-4059	MCNEIL SWEDEN AB	NO
Imodium Comp 2 mg / 125 mg tabletter	DK/H/3047/001	06-4059	MCNEIL SWEDEN AB	NO
Imodium Comp 2 mg / 125 mg tabletter	DK/H/3047/001	06-4059	MCNEIL SWEDEN AB	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
Imodium Comp 2 mg / 125 mg tabletter	DK/H/3047/001	06-4059	MCNEIL SWEDEN AB	NO
Imodium Comp 2 mg / 125 mg tabletter	DK/H/3047/001	06-4059	MCNEIL SWEDEN AB	NO
Imodium Comp 2 mg / 125 mg tabletter	DK/H/3047/001	06-4059	MCNEIL SWEDEN AB	NO
Imodium Comp 2 mg / 125 mg tabletter	DK/H/3047/001	06-4059	MCNEIL SWEDEN AB	NO
Imodium Comp 2 mg / 125 mg tabletter	DK/H/3047/001	06-4059	MCNEIL SWEDEN AB	NO
Imodium Comp 2 mg / 125 mg tabletter	DK/H/3047/001	06-4059	MCNEIL SWEDEN AB	NO
Imodium Comp 2 mg / 125 mg tabletter	DK/H/3047/001	06-4059	MCNEIL SWEDEN AB	NO
Dissenten, 2 mg, tabletki	not available	R/0207	SPA – SOCIETÀ PRODOTTI ANTIBIOTICI S.P.A.	PL
Imodium Instant, 2 mg, tabletka ulegająca rozpadowi w jamie ustnej	not available	9091	MCNEIL HEALTHCARE (IRELAND) LIMITED	PL
Imodium Instant, 2 mg, tabletka ulegająca rozpadowi w jamie ustnej	not available	9091	MCNEIL HEALTHCARE (IRELAND) LIMITED	PL

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Stoperan S, 2 mg + 125 mg, tabletki	PL/H/0483/001	21502	US PHARMACIA SP. Z O.O.	PL
IMODIUM 2 mg cápsulas	not available	9385104	JOHNSON & JOHNSON LDA	PT
IMODIUM 2 mg cápsulas	not available	9385112	JOHNSON & JOHNSON LDA	PT
Imodium Plus 2 mg /125 mg comprimidos	DK/H/3047/001	5045877, 5047261, 5037346, 5037353	JOHNSON & JOHNSON, LDA.	PT
Imodium Plus 2 mg /125 mg comprimidos	DK/H/3047/001	5045877, 5047261, 5037346, 5037353	JOHNSON & JOHNSON, LDA.	PT
Imodium Plus 2 mg /125 mg comprimidos	DK/H/3047/001	5045877, 5047261, 5037346, 5037353	JOHNSON & JOHNSON, LDA.	PT
Imodium Plus 2 mg /125 mg comprimidos	DK/H/3047/001	5045877, 5047261, 5037346, 5037353	JOHNSON & JOHNSON, LDA.	PT
Imodium Rapid 2 mg comprimido orodispersível	not available	2438489	JOHNSON & JOHNSON LDA	PT
Imodium, 0.2 mg/ml, solução oral	not available	9385211	JOHNSON & JOHNSON LDA	PT
Enterium 2 mg capsule	not available	6467/2014/01	SANIENCE S.R.L.	RO
IMODIUM 2 mg capsule	not available	11594/2019/01-02	MCNEIL HEALTHCARE (IRELAND) LIMITED	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
IMODIUM 2 mg capsule	not available	11594/2019/01-02	MCNEIL HEALTHCARE (IRELAND) LIMITED	RO
Imodium 2 mg comprimate orodispersabile	not available	12758/2019/01-02-03-04	MCNEIL HEALTHCARE (IRELAND) LIMITED	RO
Imodium 2 mg comprimate orodispersabile	not available	12758/2019/01-02-03-04	MCNEIL HEALTHCARE (IRELAND) LIMITED	RO
Imodium 2 mg comprimate orodispersabile	not available	12758/2019/01-02-03-04	MCNEIL HEALTHCARE (IRELAND) LIMITED	RO
Imodium 2 mg comprimate orodispersabile	not available	12758/2019/01-02-03-04	MCNEIL HEALTHCARE (IRELAND) LIMITED	RO
Imodium Plus 2 mg/125 mg comprimate	DK/H/3047/001/MR	10809/2018/01	MCNEIL HEALTHCARE (IRELAND) LIMITED	RO
Imodium Plus 2 mg/125 mg comprimate	DK/H/3047/001/MR	10809/2018/03	MCNEIL HEALTHCARE (IRELAND) LIMITED	RO
Imodium Plus 2 mg/125 mg comprimate	DK/H/3047/001/MR	10809/2018/02	MCNEIL HEALTHCARE (IRELAND) LIMITED	RO
Imodium Plus 2 mg/125 mg comprimate	DK/H/3047/001/MR	10809/2018/04	MCNEIL HEALTHCARE (IRELAND) LIMITED	RO
Loperamid Solacium 2 mg capsule	not available	12577/2019/01	S.C. SOLACIUM PHARMA S.R.L.	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
Loperamid Solacium 2 mg capsule	not available	12577/2019/01	S.C. SOLACIUM PHARMA S.R.L.	RO
Loperamid Terapia 2 mg capsule	not available	12294/2019/01	TERAPIA S.A.	RO
Loprex 2 mg capsule	not available	13558/2020/01	LAROPHARM SRL	RO
Dimor Comp 2 mg/125 mg tabletter	NL/H/2730/001	48046	NORDIC DRUGS AB	SE
Imodium 2 mg munsönderfallande tablett	not available	23069	MCNEIL SWEDEN AB	SE
Imodium 2 mg munsönderfallande tablett	not available	23069	MCNEIL SWEDEN AB	SE
Imodium 2 mg munsönderfallande tablett	not available	23069	MCNEIL SWEDEN AB	SE
Imodium 2 mg munsönderfallande tablett	not available	23069	MCNEIL SWEDEN AB	SE
Imodium 2 mg munsönderfallande tablett	not available	23069	MCNEIL SWEDEN AB	SE
Imodium 2 mg munsönderfallande tablett	not available	23069	MCNEIL SWEDEN AB	SE
Imodium 2 mg munsönderfallande tablett	not available	23069	MCNEIL SWEDEN AB	SE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Imodium 2 mg tablett	not available	11165	MCNEIL SWEDEN AB	SE
Imodium 2 mg tablett	not available	11165	MCNEIL SWEDEN AB	SE
Imodium 2 mg tablett	not available	11165	MCNEIL SWEDEN AB	SE
Imodium 2 mg tablett	not available	11165	MCNEIL SWEDEN AB	SE
Imodium Plus 2 mg/125 mg tabletter	DK/H/3047/001/MR	23601	MCNEIL SWEDEN AB	SE
Imodium Plus 2 mg/125 mg tabletter	DK/H/3047/001/MR	23601	MCNEIL SWEDEN AB	SE
Imodium Plus 2 mg/125 mg tabletter	DK/H/3047/001/MR	23601	MCNEIL SWEDEN AB	SE
Imodium Plus 2 mg/125 mg tabletter	DK/H/3047/001/MR	23601	MCNEIL SWEDEN AB	SE
Imodium Plus 2 mg/125 mg tabletter	DK/H/3047/001/MR	23601	MCNEIL SWEDEN AB	SE
Imodium Plus 2 mg/125 mg tabletter	DK/H/3047/001/MR	23601	MCNEIL SWEDEN AB	SE
Imodium Plus 2 mg/125 mg tabletter	DK/H/3047/001/MR	23601	MCNEIL SWEDEN AB	SE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Imodium Plus 2 mg/125 mg tabletter	DK/H/3047/001/MR	23601	MCNEIL SWEDEN AB	SE
Imodium Plus 2 mg/125 mg tabletter	DK/H/3047/001/MR	23601	MCNEIL SWEDEN AB	SE
Imodium Plus 2 mg/125 mg tabletter	DK/H/3047/001/MR	23601	MCNEIL SWEDEN AB	SE
Imodium Plus 2 mg/125 mg tabletter	DK/H/3047/001/MR	23601	MCNEIL SWEDEN AB	SE
Imodium Plus 2 mg/125 mg tabletter	DK/H/3047/001/MR	23601	MCNEIL SWEDEN AB	SE
Imodium Plus 2 mg/125 mg tabletter	DK/H/3047/001/MR	23601	MCNEIL SWEDEN AB	SE
Imodium Plus 2 mg/125 mg tabletter	DK/H/3047/001/MR	23601	MCNEIL SWEDEN AB	SE
Imodium Plus 2 mg/125 mg tabletter	DK/H/3047/001/MR	23601	MCNEIL SWEDEN AB	SE
Imolopesim 2 mg/125 mg tabletter	NL/H/2907/001	48971	ORIFARM GENERICS A/S	SE
IMODIUM 2 mg tvrdé kapsuly	not available	49/0071/92-S	MCNEIL HEALTHCARE (IRELAND) LIMITED	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
IMODIUM 2 mg tvrdé kapsuly	not available	49/0071/92-S	MCNEIL HEALTHCARE (IRELAND) LIMITED	SK
Imodium Instant 2 mg orodispergovateľné tablety	not available	49/0401/12-S	MCNEIL HEALTHCARE (IRELAND) LIMITED	SK
Imodium Instant 2 mg orodispergovateľné tablety	not available	49/0401/12-S	MCNEIL HEALTHCARE (IRELAND) LIMITED	SK
Imodium Instant 2 mg orodispergovateľné tablety	not available	49/0401/12-S	MCNEIL HEALTHCARE (IRELAND) LIMITED	SK
Imodium Instant 2 mg orodispergovateľné tablety	not available	49/0401/12-S	MCNEIL HEALTHCARE (IRELAND) LIMITED	SK
Imodium Plus 2 mg/125 mg tablety	DK/H/3047/001/MR	49/0283/18-S	MCNEIL HEALTHCARE (IRELAND) LIMITED	SK
Imodium Plus 2 mg/125 mg tablety	DK/H/3047/001/MR	49/0283/18-S	MCNEIL HEALTHCARE (IRELAND) LIMITED	SK
Imodium Plus 2 mg/125 mg tablety	DK/H/3047/001/MR	49/0283/18-S	MCNEIL HEALTHCARE (IRELAND) LIMITED	SK
Imodium Plus 2 mg/125 mg tablety	DK/H/3047/001/MR	49/0283/18-S	MCNEIL HEALTHCARE (IRELAND) LIMITED	SK
Imodium Plus 2 mg/125 mg tablety	DK/H/3047/001/MR	49/0283/18-S	MCNEIL HEALTHCARE (IRELAND) LIMITED	SK
Imodium Plus 2 mg/125 mg tablety	DK/H/3047/001/MR	49/0283/18-S	MCNEIL HEALTHCARE (IRELAND) LIMITED	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
Imodium Plus 2 mg/125 mg tablet	DK/H/3047/001/MR	49/0283/18-S	MCNEIL HEALTHCARE (IRELAND) LIMITED	SK
Imodium Plus 2 mg/125 mg tablet	DK/H/3047/001/MR	49/0283/18-S	MCNEIL HEALTHCARE (IRELAND) LIMITED	SK
Imodium Plus 2 mg/125 mg tablet	DK/H/3047/001/MR	49/0283/18-S	MCNEIL HEALTHCARE (IRELAND) LIMITED	SK
Imodium Plus 2 mg/125 mg tablet	DK/H/3047/001/MR	49/0283/18-S	MCNEIL HEALTHCARE (IRELAND) LIMITED	SK
Imodium Plus 2 mg/125 mg tablet	DK/H/3047/001/MR	49/0283/18-S	MCNEIL HEALTHCARE (IRELAND) LIMITED	SK
Imodium Plus 2 mg/125 mg tablet	DK/H/3047/001/MR	49/0283/18-S	MCNEIL HEALTHCARE (IRELAND) LIMITED	SK
Imodium Plus 2 mg/125 mg tablet	DK/H/3047/001/MR	49/0283/18-S	MCNEIL HEALTHCARE (IRELAND) LIMITED	SK
Imodium Plus 2 mg/125 mg tablet	DK/H/3047/001/MR	49/0283/18-S	MCNEIL HEALTHCARE (IRELAND) LIMITED	SK
Imodium Plus 2 mg/125 mg tablet	DK/H/3047/001/MR	49/0283/18-S	MCNEIL HEALTHCARE (IRELAND) LIMITED	SK
Imodium Plus 2 mg/125 mg tablet	DK/H/3047/001/MR	49/0283/18-S	MCNEIL HEALTHCARE (IRELAND) LIMITED	SK
Boots Diarrhoea Relief 2mg Capsules	not available	PL 00014/0611	THE BOOTS COMPANY PLC	XI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Imodium 1 mg/5 ml oral solution	not available	PL 00242/0040	JANSSEN-CILAG LIMITED	XI
Imodium Dual Action Relief Tablets	DK/H/3047/001/MR	PL 15513/0342	MCNEIL PRODUCTS LIMITED (HIGH WYCOMBE)	XI
Imodium Dual Action Relief Tablets	DK/H/3047/001/MR	PL 15513/0342	MCNEIL PRODUCTS LIMITED (HIGH WYCOMBE)	XI
Imodium Dual Action Relief Tablets	DK/H/3047/001/MR	PL 15513/0342	MCNEIL PRODUCTS LIMITED (HIGH WYCOMBE)	XI
Imodium Dual Action Relief Tablets	DK/H/3047/001/MR	PL 15513/0342	MCNEIL PRODUCTS LIMITED (HIGH WYCOMBE)	XI
Imodium Dual Action Relief Tablets	DK/H/3047/001/MR	PL 15513/0342	MCNEIL PRODUCTS LIMITED (HIGH WYCOMBE)	XI
Imodium Dual Action Relief Tablets	DK/H/3047/001/MR	PL 15513/0342	MCNEIL PRODUCTS LIMITED (HIGH WYCOMBE)	XI
Imodium Dual Action Relief Tablets	DK/H/3047/001/MR	PL 15513/0342	MCNEIL PRODUCTS LIMITED (HIGH WYCOMBE)	XI
Imodium Dual Action Relief Tablets	DK/H/3047/001/MR	PL 15513/0342	MCNEIL PRODUCTS LIMITED (HIGH WYCOMBE)	XI
Imodium Dual Action Relief Tablets	DK/H/3047/001/MR	PL 15513/0342	MCNEIL PRODUCTS LIMITED (HIGH WYCOMBE)	XI
Imodium Dual Action Relief Tablets	DK/H/3047/001/MR	PL 15513/0342	MCNEIL PRODUCTS LIMITED (HIGH WYCOMBE)	XI
Imodium Dual Action Relief Tablets	DK/H/3047/001/MR	PL 15513/0342	MCNEIL PRODUCTS LIMITED (HIGH WYCOMBE)	XI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Imodium Dual Action Relief Tablets	DK/H/3047/001/MR	PL 15513/0342	MCNEIL PRODUCTS LIMITED (HIGH WYCOMBE)	XI
Imodium Dual Action Relief Tablets	DK/H/3047/001/MR	PL 15513/0342	MCNEIL PRODUCTS LIMITED (HIGH WYCOMBE)	XI
Imodium Dual Action Relief Tablets	DK/H/3047/001/MR	PL 15513/0342	MCNEIL PRODUCTS LIMITED (HIGH WYCOMBE)	XI
Imodium Dual Action Relief Tablets	DK/H/3047/001	PL 15513/0342	MCNEIL PRODUCTS LIMITED (HIGH WYCOMBE)	XI
Imodium Dual Action Relief Tablets	DK/H/3047/001/MR	PL 15513/0342	MCNEIL PRODUCTS LIMITED (HIGH WYCOMBE)	XI
Imodium Dual Action Relief Tablets	DK/H/3047/001/MR	PL 15513/0342	MCNEIL PRODUCTS LIMITED (HIGH WYCOMBE)	XI
Loperamide 2 mg Capsules	not available	PL 00289/1980	TEVA UK LIMITED	XI
Loperamide 2 mg Capsules	not available	PL 00289/1979	TEVA UK LIMITED	XI
Loperamide 2 mg Tablets	not available	PL 36687/0311	TORRENT PHARMA (UK) LTD.	XI
Loperamide 2 mg Tablets	not available	PL 36687/0312	TORRENT PHARMA (UK) LTD.	XI
Loperamide 2 mg Tablets	not available	PL 36687/0313	TORRENT PHARMA (UK) LTD.	XI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Loperamide 2mg Tablets	not available	PL 20117/0304	MORNINGSIDE HEALTHCARE LTD	XI
Loperamide 2mg Tablets	not available	PL 20117/0315	MORNINGSIDE HEALTHCARE LTD	XI
Loperamide Hydrochloride 2mg Capsules	not available	PL 11311/0516	TILLOMED LABORATORIES LTD	XI
Teva Diarrhoea Relief 2 mg Capsules	not available	PL 00289/1979	TEVA UK LIMITED	XI
Tillomed Diarrhoea Relief (Loperamide Hydrochloride) 2mg Capsules	not available	PL 11311/0515	TILLOMED LABORATORIES LTD	XI
Value Health Diarrhoea Relief 2mg Capsules	not available	PL 00014/0611	THE BOOTS COMPANY PLC	XI
Waitrose Acute Diarrhoea Relief 2mg Capsules	not available	PL 31308/0001	BRUNEL HEALTHCARE MANUFACTURING LIMITED	XI