26 October 2017
EMA/726434/2017
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: amoxicillin

Procedure no.: PSUSA/00000187/201703

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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
Ospamox 250 mg – lösbare Tabletten	8331825	137432	SANDOZ GMBH	AT
Ospamox 250 mg – lösbare Tabletten	8331825	137432	SANDOZ GMBH	AT
Ospamox 500 mg – Filmtabletten	922.421	17.659	SANDOZ GMBH	AT
Ospamox 750 mg – Filmtabletten	922.420	17.657	SANDOZ GMBH	AT
Ospamox 1000 mg – Filmtabletten	922.419	17.653	SANDOZ GMBH	AT
Ospamox 125 mg/5 ml - Pulver für orale Suspension	AT/H/0116/003	17.661	SANDOZ GMBH	AT
Ospamox 500 mg/5 ml - Pulver für orale Suspension	AT/H/0116/005	1-24881	SANDOZ GMBH	AT
Ospamox 500 mg - lösbare Tabletten	NL/H/0455/001	1-25540	SANDOZ GMBH	AT
Ospamox 750 mg - lösbare Tabletten	NL/H/0455/002	1-25541	SANDOZ GMBH	AT
Amoxicillin Sandoz 750 mg – Filmtabletten	AT/H/0187/002	1-28901	SANDOZ GMBH	AT
Amoxicillin Sandoz 1000 mg – Filmtabletten	AT/H/0187/003	1-28902	SANDOZ GMBH	AT
Amoxicillin Hexal 500 mg – Filmtabletten	AT/H/0188/001	1-28778	HEXAL PHARMA GMBH	AT
Amoxicillin Hexal 750 mg – Filmtabletten	AT/H/0188/002	1-28779	HEXAL PHARMA GMBH	AT
Amoxicillin Hexal 1000 mg – Filmtabletten	AT/H/0188/003	1-28780	HEXAL PHARMA GMBH	AT
Amoxicillin 1A Pharma 500 mg – Filmtabletten	AT/H/0189/001	1-28793	1A PHARMA GMBH	AT
Amoxicillin Sandoz 500 mg – Filmtabletten	AT/H/0187/001	1-28900	SANDOZ GMBH	AT

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the Member State	Member State where product is authorised
Amoxicillin 1A Pharma 750 mg - Filmtabletten	AT/H/0189/002	1-28794	1A PHARMA GMBH	AT
Amoxicillin 1A Pharma 1000 mg – Filmtabletten	AT/H/0189/003	1-28795	1A PHARMA GMBH	AT
Clamoxyl I.V./I.M. 1 g poeder en oplosmiddel voor oplossing voor injectie	not available	BE109417	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
CLAMOXYL I.V./I.M. 1 g Pulver und Lösungsmittel zur Herstellung einer Injektionslösung	not available	BE109417	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Clamoxyl 250 mg/5 ml, poeder voor orale suspensie	FR/H/600/03/MR	BE080577	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Clamoxyl 250 mg/5 ml, Pulver zur Herstellung einer Suspension zum Einnehmen	FR/H/600/03/MR	BE080577	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Clamoxyl 125 mg/5 ml, poeder voor orale suspensie	FR/H/600/02/MR	BE080595	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Clamoxyl 125 mg/5 ml, Pulver zur Herstellung einer Suspension zum Einnehmen	FR/H/600/02/MR	BE080595	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Clamoxyl 1 g, dispergeerbare tabletten	FR/H/600/01/MR	BE164306	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Clamoxyl 1 g, Tabletten zur Herstellung einer Suspension zum Einnehmen	FR/H/0600/001	BE164306	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Clamoxyl 500 mg, harde capsules	UK/H/6220/02	BE191807	GLAXOSMITHKLINE PHARMACEUTICALS SA	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
CLAMOXYL 500 mg Kapseln	UK/H/6220/02	BE191807	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Clamoxyl 250 mg/5 ml, poudre pour suspension buvable	FR/H/600/03/MR	BE080577	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Clamoxyl I.V./I.M. 1 g, poudre pour solution injectable/pour perfusion	BE/H/0259/001	BE109417	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Clamoxyl 125 mg/5 ml, poudre pour suspension buvable	FR/H/600/02/MR	BE080595	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Clamoxyl 500 mg, gélules	UK/H/6220/002	BE191807	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Clamoxyl 1 g, comprimés dispersibles	FR/H/600/01/MR	BE164306	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Flemoxin 250 mg/5 ml Pulver zur Herstellung einer Suspension zum Einnehmen	not available	BE126987	ASTELLAS PHARMA B.V., OFFICE BE	BE
Flemoxin Solutab 500 mg lösliche Tabletten	not available	BE127005	ASTELLAS PHARMA B.V., OFFICE BE	BE
Flemoxin Solutab 1 g lösliche Tabletten	not available	BE171805	ASTELLAS PHARMA B.V., OFFICE BE	BE
Flemoxin 250 mg/5 ml poeder voor orale suspensie	not available	BE126987	ASTELLAS PHARMA B.V., OFFICE BE	BE
Flemoxin Solutab 500 mg oplosbare tabletten	not available	BE127005	ASTELLAS PHARMA B.V., OFFICE BE	BE
Flemoxin Solutab 1 g oplosbare tabletten	not available	BE171805	ASTELLAS PHARMA B.V., OFFICE BE	BE
Flemoxin Solutab, comprimés solubles à 1 g	not available	BE171805	ASTELLAS PHARMA B.V., OFFICE BE	BE
Flemoxin Solutab, comprimés solubles à 500 mg	not available	BE127005	ASTELLAS PHARMA B.V., OFFICE BE	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
Flemoxin 250 mg/5 ml poudre pour suspension buvable	not available	BE126987	ASTELLAS PHARMA B.V., OFFICE BE	BE
Clamoxyl I.V./I.M. 1 g poeder en oplosmiddel voor oplossing voor injectie	not available	BE109417	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Clamoxyl 250 mg/5 ml, poeder voor orale suspensie	FR/H/600/03/MR	BE080577	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Clamoxyl 125 mg/5 ml, poeder voor orale suspensie	FR/H/600/02/MR	BE080595	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Clamoxyl 1 g, dispergeerbare tabletten	FR/H/600/01/MR	BE164306	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Clamoxyl 500 mg, harde capsules	UK/H/6220/02	BE191807	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
CLAMOXYL 500 mg Kapseln	UK/H/6220/02	BE191807	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Clamoxyl I.V./I.M. 1 g, poudre pour solution injectable/pour perfusion	BE/H/0259/001	BE109417	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Clamoxyl 500 mg, gélules	UK/H/6220/002	BE191807	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Clamoxyl 1 g, comprimés dispersibles	FR/H/600/01/MR	BE164306	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Amoxil 500 mg καψάκια, σκληρά	UK/H/6220/02	19966	SMITHKLINE BEECHAM LTD	CY
Amoxil Forte 250 mg/5 ml κόνις για πόσιμο εναιώρημα	FR/H/600/003	3617	SMITHKLINE BEECHAM LTD	CY
Amoxil 500 mg καψάκια, σκληρά	UK/H/6220/02	19966	SMITHKLINE BEECHAM LTD	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
Amoxil Forte 250 mg/5 ml κόνις για πόσιμο εναιώρημα	FR/H/600/003	3617	SMITHKLINE BEECHAM LTD	CY
Amoxicillin Micro Labs 250 mg Filmtabletten	DE/H/4385/001	94792.00.00	MICRO LABS GMBH	DE
Amoxicillin Micro Labs 250 mg Filmtabletten	DE/H/4385/001	94792.00.00	MICRO LABS GMBH	DE
Amoxicillin Micro Labs 250 mg Filmtabletten	DE/H/4385/001	94792.00.00	MICRO LABS GMBH	DE
Amoxicillin Micro Labs 250 mg Filmtabletten	DE/H/4385/001	94792.00.00	MICRO LABS GMBH	DE
Amoxicillin Micro Labs 250 mg Filmtabletten	DE/H/4385/001	94792.00.00	MICRO LABS GMBH	DE
Amoxicillin Micro Labs 250 mg Filmtabletten	DE/H/4385/001	94792.00.00	MICRO LABS GMBH	DE
Amoxicillin Micro Labs 250 mg Filmtabletten	DE/H/4385/001	94792.00.00	MICRO LABS GMBH	DE
Amoxicillin Micro Labs 250 mg Filmtabletten	DE/H/4385/001	94792.00.00	MICRO LABS GMBH	DE
Amoxicillin Micro Labs 250 mg Filmtabletten	DE/H/4385/001	94792.00.00	MICRO LABS GMBH	DE
Amoxicillin Micro Labs 250 mg Filmtabletten	DE/H/4385/001	94792.00.00	MICRO LABS GMBH	DE
Amoxicillin Micro Labs 250 mg Filmtabletten	DE/H/4385/001	94792.00.00	MICRO LABS GMBH	DE
Amoxicillin Micro Labs 250 mg Filmtabletten	DE/H/4385/001	94792.00.00	MICRO LABS GMBH	DE
Amoxicillin Micro Labs 250 mg Filmtabletten	DE/H/4385/001	94792.00.00	MICRO LABS GMBH	DE
Amoxicillin Micro Labs 250 mg Filmtabletten	DE/H/4385/001	94792.00.00	MICRO LABS GMBH	DE
Amoxicillin Micro Labs 250 mg Filmtabletten	DE/H/4385/001	94792.00.00	MICRO LABS GMBH	DE

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the Member State	Member State where product is authorised
Amoxicillin Micro Labs 250 mg Filmtabletten	DE/H/4385/001	94792.00.00	MICRO LABS GMBH	DE
Amoxicillin Micro Labs 250 mg Filmtabletten	DE/H/4385/001	94792.00.00	MICRO LABS GMBH	DE
Amoxicillin Micro Labs 250 mg Filmtabletten	DE/H/4385/001	94792.00.00	MICRO LABS GMBH	DE
Amoxicillin Micro Labs 250 mg Filmtabletten	DE/H/4385/001	94792.00.00	MICRO LABS GMBH	DE
Amoxicillin Micro Labs 500 mg Filmtabletten	DE/H/4385/002	94793.00.00	MICRO LABS GMBH	DE
Amoxicillin Micro Labs 500 mg Filmtabletten	DE/H/4385/002	94793.00.00	MICRO LABS GMBH	DE
Amoxicillin Micro Labs 500 mg Filmtabletten	DE/H/4385/002	94793.00.00	MICRO LABS GMBH	DE
Amoxicillin Micro Labs 500 mg Filmtabletten	DE/H/4385/002	94793.00.00	MICRO LABS GMBH	DE
Amoxicillin Micro Labs 500 mg Filmtabletten	DE/H/4385/002	94793.00.00	MICRO LABS GMBH	DE
Amoxicillin Micro Labs 500 mg Filmtabletten	DE/H/4385/002	94793.00.00	MICRO LABS GMBH	DE
Amoxicillin Micro Labs 500 mg Filmtabletten	DE/H/4385/002	94793.00.00	MICRO LABS GMBH	DE
Amoxicillin Micro Labs 500 mg Filmtabletten	DE/H/4385/002	94793.00.00	MICRO LABS GMBH	DE
Amoxicillin Micro Labs 500 mg Filmtabletten	DE/H/4385/002	94793.00.00	MICRO LABS GMBH	DE
Amoxicillin Micro Labs 500 mg Filmtabletten	DE/H/4385/002	94793.00.00	MICRO LABS GMBH	DE
Amoxicillin Micro Labs 500 mg Filmtabletten	DE/H/4385/002	94793.00.00	MICRO LABS GMBH	DE
Amoxicillin Micro Labs 500 mg Filmtabletten	DE/H/4385/002	94793.00.00	MICRO LABS GMBH	DE
Amoxicillin Micro Labs 500 mg Filmtabletten	DE/H/4385/002	94793.00.00	MICRO LABS GMBH	DE

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the Member State	Member State where product is authorised
Amoxicillin Micro Labs 500 mg Filmtabletten	DE/H/4385/002	94793.00.00	MICRO LABS GMBH	DE
Amoxicillin Micro Labs 500 mg Filmtabletten	DE/H/4385/002	94793.00.00	MICRO LABS GMBH	DE
Amoxicillin Micro Labs 500 mg Filmtabletten	DE/H/4385/002	94793.00.00	MICRO LABS GMBH	DE
Amoxicillin Micro Labs 500 mg Filmtabletten	DE/H/4385/002	94793.00.00	MICRO LABS GMBH	DE
Amoxicillin Micro Labs 500 mg Filmtabletten	DE/H/4385/002	94793.00.00	MICRO LABS GMBH	DE
Amoxicillin Micro Labs 500 mg Filmtabletten	DE/H/4385/002	94793.00.00	MICRO LABS GMBH	DE
Amoxicillin Micro Labs 750 mg Filmtabletten	DE/H/4385/003	94794.00.00	MICRO LABS GMBH	DE
Amoxicillin Micro Labs 750 mg Filmtabletten	DE/H/4385/003	94794.00.00	MICRO LABS GMBH	DE
Amoxicillin Micro Labs 750 mg Filmtabletten	DE/H/4385/003	94794.00.00	MICRO LABS GMBH	DE
Amoxicillin Micro Labs 750 mg Filmtabletten	DE/H/4385/003	94794.00.00	MICRO LABS GMBH	DE
Amoxicillin Micro Labs 750 mg Filmtabletten	DE/H/4385/003	94794.00.00	MICRO LABS GMBH	DE
Amoxicillin Micro Labs 750 mg Filmtabletten	DE/H/4385/003	94794.00.00	MICRO LABS GMBH	DE
Amoxicillin Micro Labs 750 mg Filmtabletten	DE/H/4385/003	94794.00.00	MICRO LABS GMBH	DE
Amoxicillin Micro Labs 750 mg Filmtabletten	DE/H/4385/003	94794.00.00	MICRO LABS GMBH	DE
Amoxicillin Micro Labs 750 mg Filmtabletten	DE/H/4385/003	94794.00.00	MICRO LABS GMBH	DE
Amoxicillin Micro Labs 750 mg Filmtabletten	DE/H/4385/003	94794.00.00	MICRO LABS GMBH	DE
Amoxicillin Micro Labs 750 mg Filmtabletten	DE/H/4385/003	94794.00.00	MICRO LABS GMBH	DE

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the Member State	Member State where product is authorised
Amoxicillin Micro Labs 750 mg Filmtabletten	DE/H/4385/003	94794.00.00	MICRO LABS GMBH	DE
Amoxicillin Micro Labs 750 mg Filmtabletten	DE/H/4385/003	94794.00.00	MICRO LABS GMBH	DE
Amoxicillin Micro Labs 750 mg Filmtabletten	DE/H/4385/003	94794.00.00	MICRO LABS GMBH	DE
Amoxicillin Micro Labs 750 mg Filmtabletten	DE/H/4385/003	94794.00.00	MICRO LABS GMBH	DE
Amoxicillin Micro Labs 750 mg Filmtabletten	DE/H/4385/003	94794.00.00	MICRO LABS GMBH	DE
Amoxicillin Micro Labs 750 mg Filmtabletten	DE/H/4385/003	94794.00.00	MICRO LABS GMBH	DE
Amoxicillin Micro Labs 750 mg Filmtabletten	DE/H/4385/003	94794.00.00	MICRO LABS GMBH	DE
Amoxicillin Micro Labs 750 mg Filmtabletten	DE/H/4385/003	94794.00.00	MICRO LABS GMBH	DE
Helicomp® Sandoz®, 20 mg/1000 mg/500 mg magensaftresistente Hartkapseln/Filmtablette n	not available	63912.00.00	HEXAL AG	DE
OMEP Plus Amoxicillin + Clarithromycin	not available	65160.00.00	HEXAL AG	DE
Imacillin, granulat til oral suspension	not available	11372	MEDA AS	DK
Imacillin, opløselige tabletter	not available	15288	MEDA AS	DK
Imacillin, opløselige tabletter	not available	15289	MEDA AS	DK
Imacillin, opløselige tabletter	not available	15290	MEDA AS	DK
Imacillin, opløselige tabletter	not available	15291	MEDA AS	DK
Imacillin, opløselige tabletter	not available	16713	MEDA AS	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
AMITRÓN® 250 mg/5 ml Suspensión extemporánea	not available	54.625	LDP LABORATORIOS TORLAN, S.A.	ES
Clamoxyl 500 mg cápsulas	not available	50.239	GLAXOSMITHKLINE S.A.	ES
CLAMOXYL 1 g intramuscular	not available	55522	GLAXOSMITHKLINE, S.A.	ES
Clamoxyl 250 mg polvo para suspensión oral en sobre	not available	50.944	GLAXOSMITHKLINE S.A.	ES
Clamoxyl 1 g polvo para suspensión oral en sobre	not available	59.132	GLAXOSMITHKLINE S.A.	ES
Clamoxyl 250 mg/5 ml polvo para suspensión oral en frasco	not available	52.015	GLAXOSMITHKLINE S.A.	ES
Clamoxyl 1 g comprimidos	FR/H/600/01	59.133	GLAXOSMITHKLINE S.A.	ES
Clamoxyl 750 mg comprimidos	not available	51.426	GLAXOSMITHKLINE S.A.	ES
Clamoxyl 500 mg polvo para suspensión oral en sobre	not available	58.840	GLAXOSMITHKLINE S.A.	ES
AMITRÓN® 500 mg Inyectable	not available	56.265	LDP LABORATORIOS TORLAN, S.A.	ES
AMITRÓN® 1g Inyectable	not available	56.266	LDP LABORATORIOS TORLAN, S.A.	ES
AMITRÓN® 250 mg Inyectable	not available	56.264	LDP LABORATORIOS TORLAN, S.A.	ES
AMITRÓN® 500 mg Cápsulas	not available	54.626	LDP LABORATORIOS TORLAN, S.A.	ES
Clamoxyl 500 mg cápsulas	not available	50.239	GLAXOSMITHKLINE S.A.	ES
CLAMOXYL 1 g intramuscular	not available	55522	GLAXOSMITHKLINE, 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
Clamoxyl 250 mg polvo para suspensión oral en sobre	not available	50.944	GLAXOSMITHKLINE S.A.	ES
Clamoxyl 1 g polvo para suspensión oral en sobre	not available	59.132	GLAXOSMITHKLINE S.A.	ES
Clamoxyl 1 g comprimidos	FR/H/600/01	59.133	GLAXOSMITHKLINE S.A.	ES
Clamoxyl 750 mg comprimidos	not available	51.426	GLAXOSMITHKLINE S.A.	ES
AMOXICILLINE SANDOZ 500 mg, gélule	NL 21558	34009 367 960 1 0	SANDOZ	FR
HICONCIL 500 mg, gélule	not available	316 122-9	BRISTOL-MYERS SQUIBB	FR
HICONCIL 500 mg, gélule	not available	316 121-2	BRISTOL-MYERS SQUIBB	FR
HICONCIL 500 mg, gélule	not available	316 123-5	BRISTOL-MYERS SQUIBB	FR
HICONCIL 500 mg, gélule	not available	553 445-6	BRISTOL-MYERS SQUIBB	FR
HICONCIL 500 mg, gélule	not available	554 346-1	BRISTOL-MYERS SQUIBB	FR
HICONCIL 500 mg, gélule	not available	342 243-4	BRISTOL-MYERS SQUIBB	FR
HICONCIL 125 mg /5 ml, pour poudre suspension buvable	not available	3400931703464	BRISTOL-MYERS SQUIBB	FR
HICONCIL 250 mg /5 ml, pour poudre suspension buvable	not available	3400931761594	BRISTOL-MYERS SQUIBB	FR
HICONCIL 500 mg /5 ml, pour poudre suspension buvable	not available	3400932026616	BRISTOL-MYERS SQUIBB	FR
CLAMOXYL 1 g, poudre et solvant pour solution injectable (IM)	not available	VNL11851	LABORATOIRE GLAXOSMITHKLINE	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
CLAMOXYL 2 g, poudre pour solution injectable (IV)	not available	VNL11492	LABORATOIRE GLAXOSMITHKLINE	FR
CLAMOXYL 500 mg, poudre pour solution injectable (IM, IV)	not available	VNL11494-1	LABORATOIRE GLAXOSMITHKLINE	FR
CLAMOXYL 1 g, poudre pour solution injectable (IM - IV)	not available	VNL11491-1	LABORATOIRE GLAXOSMITHKLINE	FR
CLAMOXYL 125 mg/5 ml, poudre pour suspension buvable	not available	VNL11074	LABORATOIRE GLAXOSMITHKLINE	FR
CLAMOXYL 250 mg/5 ml, poudre pour suspension buvable	not available	VNL11075	LABORATOIRE GLAXOSMITHKLINE	FR
CLAMOXYL 1 g, comprimé dispersible	not available	NL15119	LABORATOIRE GLAXOSMITHKLINE	FR
CLAMOXYL 1 g, poudre pour suspension buvable en sachet-dose	not available	NL15559	LABORATOIRE GLAXOSMITHKLINE	FR
CLAMOXYL 500 mg/5 ml, poudre pour suspension buvable	not available	VNL11666	LABORATOIRE GLAXOSMITHKLINE	FR
CLAMOXYL 500 mg, gélule	not available	VNL9721	LABORATOIRE GLAXOSMITHKLINE	FR
CLAMOXYL 1 g, poudre et solvant pour solution injectable (IM)	not available	VNL11851	LABORATOIRE GLAXOSMITHKLINE	FR
CLAMOXYL 2 g, poudre pour solution injectable (IV)	not available	VNL11492	LABORATOIRE GLAXOSMITHKLINE	FR
CLAMOXYL 500 mg, poudre pour solution injectable (IM, IV)	not available	VNL11494-1	LABORATOIRE GLAXOSMITHKLINE	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
CLAMOXYL 1 g, poudre pour solution injectable (IM - IV)	not available	VNL11491-1	LABORATOIRE GLAXOSMITHKLINE	FR
CLAMOXYL 125 mg/5 ml, poudre pour suspension buvable	not available	VNL11074	LABORATOIRE GLAXOSMITHKLINE	FR
CLAMOXYL 250 mg/5 ml, poudre pour suspension buvable	not available	VNL11075	LABORATOIRE GLAXOSMITHKLINE	FR
CLAMOXYL 1 g, comprimé dispersible	not available	NL15119	LABORATOIRE GLAXOSMITHKLINE	FR
CLAMOXYL 1 g, poudre pour suspension buvable en sachet-dose	not available	NL15559	LABORATOIRE GLAXOSMITHKLINE	FR
CLAMOXYL 500 mg/5 ml, poudre pour suspension buvable	not available	VNL11666	LABORATOIRE GLAXOSMITHKLINE	FR
CLAMOXYL 500 mg, gélule	not available	VNL9721	LABORATOIRE GLAXOSMITHKLINE	FR
AMOXICILLINE BIOGARAN 500 mg, gélule	not available	3400933856731	BIOGARAN	FR
AMOXICILLINE BIOGARAN 500 mg, gélule	not available	3400933857042	BIOGARAN	FR
AMOXICILLINE BIOGARAN 500 mg, gélule	not available	3400933856960	BIOGARAN	FR
AMOXICILLINE BIOGARAN 500 mg, gélule	not available	3400933857103	BIOGARAN	FR
AMOXICILLINE BIOGARAN 500 mg, gélule	not available	3400933857271	BIOGARAN	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
AMOXICILLINE SANDOZ 125 mg/5 ml, poudre pour suspension buvable	NL 21556	368 829-6	SANDOZ	FR
AMOXICILLINE SANDOZ 250 mg/5 ml, poudre pour suspension buvable	NL 21557	365 824-3	SANDOZ	FR
AMOXICILLINE SANDOZ 500 mg, gélule	NL 21558	34009 367 960 1 0	SANDOZ	FR
AMOXICILLINE RPG 250 mg/5 ml, poudre pour suspension buvable en flacon	not available	NL 21 383	RANBAXY PHARMACIE GENERIQUES	FR
AMOXICILLINE RPG 500 mg/5 ml, poudre pour suspension buvable en flacon	not available	NL 21 384	RANBAXY PHARMACIE GENERIQUES	FR
Amoxil 500 mg/5 ml κόνις για πόσιμο εναιώρημα	FR/H/600/04/MR	0933603	GLAXOSMITHKLINE AEBE	GR
AMOXIL	FR/H/0600/001	85885/11/16-02-2012	GLAXOSMITHKLINE AEBE	GR
AMOXIL	UK/H/6220/02	49566/23-06-2014	GLAXOSMITHKLINE AEBE	GR
Amoxil 250 mg/5 ml κόνις για πόσιμο εναιώρημα	FR/H/600/03/MR	0933602	GLAXOSMITHKLINE AEBE	GR
Amoxil 1 g κόνις για ενέσιμο διάλυμα ή διάλυμα προς έγχυση	BE/H/0259/001	0933607	GLAXOSMITHKLINE AEBE	GR
AMOXIL	FR/H/0600/001	85885/11/16-02-2012	GLAXOSMITHKLINE AEBE	GR
AMOXIL	UK/H/6220/02	49566/23-06-2014	GLAXOSMITHKLINE AEBE	GR
Amoxil 250 mg/5 ml κόνις για πόσιμο εναιώρημα	FR/H/600/03/MR	0933602	GLAXOSMITHKLINE AEBE	GR
Duomox 1000 mg tabletta	not available	OGYI-T-5460/05	ASTELLAS PHARMA KFT	HU
Duomox 250 mg tabletta	not available	OGYI-T-5460/01	ASTELLAS 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
Duomox 500 mg tabletta	not available	OGYI-T-5460/03	ASTELLAS PHARMA KFT	HU
Duomox 750 mg tabletta	not available	OGYI-T-5460/04	ASTELLAS PHARMA KFT	HU
Oramox (Amoxicillin Oral Suspension BP) 125mg / 5ml	not available	PA0298/019/001	ATHLONE LABORATORIES LIMITED	IE
Oramox (Amoxicillin Oral Suspension BP) 250mg/5ml	not available	PA0298/019/002	ATHLONE LABORATORIES LIMITED	IE
Oramox 250mg Hard Capsules	not available	PA0298/019/003	ATHLONE LABORATORIES LIMITED	IE
Oramox 500mg Hard Capsules	not available	PA0298/019/004	ATHLONE LABORATORIES LIMITED	IE
Amoxil 3g Powder for Oral Suspension Sachets	UK/H/6220/07	PA 1077/33/6	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Amoxil Vials 500mg, powder for solution for injection or infusion	UK/H/6220/05	PA 1077/33/4	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Amoxil Paediatric 125mg/1.25ml Powder for Oral Suspension	UK/H/6220/03	PA 1077/33/5	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Amoxil 3g Powder for Oral Suspension Sachets	UK/H/6220/07	PA 1077/33/6	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Amoxil Vials 500mg, powder for solution for injection or infusion	UK/H/6220/05	PA 1077/33/4	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Amoxil Paediatric 125mg/1.25ml Powder for Oral Suspension	UK/H/6220/03	PA 1077/33/5	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Amoxicillin 250 mg/5 ml powder for oral suspension	not available	PA0126/282/004	CLONMEL HEALTHCARE LTD.	IE
Amoxicillin 250 mg hard capsules	not available	PA0126/282/001	CLONMEL HEALTHCARE 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
Amoxicillin 125mg / 5ml powder for oral suspension	not available	PA0126/282/003	CLONMEL HEALTHCARE LTD.	IE
Amoxicillin 125mg / 5ml powder for oral suspension	not available	PA0126/282/003	CLONMEL HEALTHCARE LTD.	IE
Amoxicillin 250 mg/5 ml powder for oral suspension	not available	PA0126/282/004	CLONMEL HEALTHCARE LTD.	IE
Amoxicillin 500 mg hard capsules	not available	PA0126/282/002	CLONMEL HEALTHCARE LTD.	IE
Amoxicillin 250 mg hard capsules	not available	PA0126/282/001	CLONMEL HEALTHCARE LTD.	IE
Amoxicillin 500 mg hard capsules	not available	PA0126/282/002	CLONMEL HEALTHCARE LTD.	IE
Amoxicillin Sandoz 100 mg/ml mixtúruduft, dreifa	AT/H/0116/005	IS/1/15/062/01	SANDOZ A/S	IS
SINTOPEN " 250 MG/5 ML GRANULATO PER SOSPENSIONE ORALE"	not available	023053135	MAGIS FARMACEUTICI SRL	IT
SINTOPEN " 1 G COMPRESSE"	not available	023053123	MAGIS FARMACEUTICI SRL	IT
ZIMOX 1 g compresse solubili e masticabili	not available	023086174	PFIZER ITALIA S.R.L.	IT
ZIMOX 500 mg capsule rigide	not available	023086061	PFIZER ITALIA S.R.L.	IT
ZIMOX 500 mg compresse solubili e masticabili	not available	023086162	PFIZER ITALIA S.R.L.	IT
ZIMOX 1 g compresse	not available	023086150	PFIZER ITALIA S.R.L.	IT
ZIMOX 250 mg/5 ml polvere per sospensione orale	not available	023086097	PFIZER 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
ZIMOX 100 mg/ml gocce orali, sospensione	not available	023086186	PFIZER ITALIA S.R.L.	IT
Amoxina 1 g compresse dispersibili	not available	023966094	AESCULAPIUS FARMACEUTICI S.R.L.	IT
Amoxina 1 g compresse dispersibili	not available	023966118	AESCULAPIUS FARMACEUTICI S.R.L.	IT
Amoxina 250mg/5ml granulato per sospensione orale	not available	023966106	AESCULAPIUS FARMACEUTICI S.R.L.	IT
Amoxina 250mg/5ml granulato per sospensione orale	not available	023966082	AESCULAPIUS FARMACEUTICI S.R.L.	IT
VELAMOX 1 g compresse dispersibili	not available	023097102	MEDIOLANUM FARMACEUTICI SPA	IT
VELAMOX 250 mg/7 ml polvere per sospensione orale	not available	023097037	MEDIOLANUM FARMACEUTICI SPA	IT
VELAMOX 500 mg capsule rigide	not available	023097013	MEDIOLANUM FARMACEUTICI SPA	IT
Amoxil 1 g disperguojamosios tabletės	FR/H/0600/001	LT/1/04/0031/006	BEECHAM GROUP PLC	LT
Amoxil 250 mg/5 ml milteliai geriamajai suspensijai	FR/H/600/003	LT/1/04/0031/001	BEECHAM GROUP PLC	LT
Amoxil 1 g disperguojamosios tabletės	FR/H/0600/001	LT/1/04/0031/004	BEECHAM GROUP PLC	LT
Amoxil 1 g disperguojamosios tabletės	FR/H/0600/001	LT/1/04/0031/005	BEECHAM GROUP PLC	LT
Amoxil 500 mg kietos kapsulės	UK/H/6220/02	LT/1/04/0031/002	BEECHAM GROUP PLC	LT
Amoxil 500 mg kietos kapsules	UK/H/6220/02	LT/1/04/0031/003	BEECHAM GROUP PLC	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
Amoxil 1 g disperguojamosios tabletės	FR/H/0600/001	LT/1/04/0031/006	BEECHAM GROUP PLC	LT
Amoxil 250 mg/5 ml milteliai geriamajai suspensijai	FR/H/600/003	LT/1/04/0031/001	BEECHAM GROUP PLC	LT
Amoxil 1 g disperguojamosios tabletės	FR/H/0600/001	LT/1/04/0031/004	BEECHAM GROUP PLC	LT
Amoxil 1 g disperguojamosios tabletės	FR/H/0600/001	LT/1/04/0031/005	BEECHAM GROUP PLC	LT
Amoxil 500 mg kietos kapsulės	UK/H/6220/02	LT/1/04/0031/002	BEECHAM GROUP PLC	LT
Amoxil 500 mg kietos kapsules	UK/H/6220/02	LT/1/04/0031/003	BEECHAM GROUP PLC	LT
Clamoxyl I.V./I.M. 1 g poeder en oplosmiddel voor oplossing voor injectie	not available	260/08 10 0038	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
CLAMOXYL I.V./I.M. 1 g Pulver und Lösungsmittel zur Herstellung einer Injektionslösung	not available	260/08 10 0038	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Clamoxyl 1 g, dispergeerbare tabletten	FR/H/600/01	2008 10 0030	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Clamoxyl 1 g, Tabletten zur Herstellung einer Suspension zum Einnehmen	FR/H/0600/001	2008 10 0030	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Clamoxyl 500 mg capsules	not available	260/08 10 0031	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
CLAMOXYL 500 mg Kapseln	not available	260/08 10 0031	GLAXOSMITHKLINE PHARMACEUTICALS SA	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
Clamoxyl 125 mg/5 ml, Pulver zur Herstellung einer Suspension zum Einnehmen	FR/H/600/02/MR	0260/08100032	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Clamoxyl 250 mg/5 ml, Pulver zur Herstellung einer Suspension zum Einnehmen	FR/H/600/03/MR	0260/08100033	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Clamoxyl 125 mg/5 ml, poudre pour suspension buvable	FR/H/600/02/MR	0260/08100032	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Clamoxyl 250 mg/5 ml, poudre pour suspension buvable	FR/H/600/03/MR	0260/08100033	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Clamoxyl I.V./I.M. 1 g, poudre pour solution injectable/pour perfusion	BE/H/0259/001	260/08 10 0038	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Clamoxyl 500 mg, gélules	not available	260/08 10 0031	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Clamoxyl 1 g, comprimés dispersibles	FR/H/600/01/MR	2008 10 0030	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
CLAMOXYL I.V./I.M. 1 g Pulver und Lösungsmittel zur Herstellung einer Injektionslösung	not available	260/08 10 0038	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Clamoxyl 500 mg capsules	not available	260/08 10 0031	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
CLAMOXYL 500 mg Kapseln	not available	260/08 10 0031	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Clamoxyl 125 mg/5 ml, Pulver zur Herstellung einer Suspension zum Einnehmen	FR/H/600/02/MR	0260/08100032	GLAXOSMITHKLINE PHARMACEUTICALS SA	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
Clamoxyl 250 mg/5 ml, poudre pour suspension buvable	FR/H/600/03/MR	0260/08100033	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Clamoxyl I.V./I.M. 1 g, poudre pour solution injectable/pour perfusion	BE/H/0259/001	260/08 10 0038	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Clamoxyl 1 g, comprimés dispersibles	FR/H/600/01/MR	2008 10 0030	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Flemoxin Solutab, comprimés solubles à 500 mg	not available	BE127005	ASTELLAS PHARMA B.V., OFFICE BE	LU
Flemoxin Solutab, comprimés solubles à 1 g	not available	1996060293	ASTELLAS PHARMA B.V., OFFICE BE	LU
Amoxil 500 mg cietās kapsulas	UK/H/6220/02	04-0196	GLAXOSMITHKLINE LATVIA SIA	LV
Ospamox 1000 mg apvalkotās tabletes	AT/H/0187/003	10-0081	SANDOZ PHARMACEUTICALS D.D.	LV
Amoxil 250 mg/5 ml powder for oral suspension	FR/H/600/03/MR	MA447/00105	BEECHAM GROUP PLC	MT
Amoxil and associated names 500 mg capsules, hard	UK/H/6220/02	MA447/00103	BEECHAM GROUP PLC	MT
Amoxil 250 mg/5 ml powder for oral suspension	FR/H/600/03/MR	MA447/00105	BEECHAM GROUP PLC	MT
Amoxil and associated names 500 mg capsules, hard	UK/H/6220/02	MA447/00103	BEECHAM GROUP PLC	MT
IMACILLIN®	not available	94-1502	MEDA AS	NO
IMACILLIN®	not available	8005	MEDA AS	NO
IMACILLIN®	not available	6187	MEDA AS	NO
IMACILLIN®	not available	7886	MEDA AS	NO
IMACILLIN®	not available	7887	MEDA AS	NO

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the Member State	Member State where product is authorised
IMACILLIN®	not available	7888	MEDA AS	NO
IMACILLIN®	not available	7889	MEDA AS	NO
IMACILLIN®	not available	7890	MEDA AS	NO
Ospamox 500 mg, tabletki powlekane	AT/H/0188/001	16801	SANDOZ GMBH	PL
Ospamox 750 mg, tabletki powlekane	AT/H/0188/002	16802	SANDOZ GMBH	PL
OSPAMOX 1000 MG, 1000 MG, TABLETKI POWLEKANE	AT/H/0188/003	16687	SANDOZ GMBH	PL
Clamoxyl 500 mg cápsulas	UK/H/6220/02	4667481	BEECHAM PORTUGUESA, PRODUTOS FARMACÊUTICOS E QUÍMICOS, LDA	PT
Clamoxyl 500 mg cápsulas	UK/H/6220/02	8352112	BEECHAM PORTUGUESA, PRODUTOS FARMACÊUTICOS E QUÍMICOS, LDA	PT
Clamoxyl 250 mg/5 ml pó para suspensão oral	FR/H/0600/003	8352203	SMITH KLINE & FRENCH PORTUGUESA-PRODUTOS FARMACEUTICOS LDA	PT
Clamoxyl 1 g comprimidos dispersíveis	FR/H/600/01/MR	8436816	SMITH KLINE & FRENCH PORTUGUESA-PRODUTOS FARMACEUTICOS LDA	PT
Clamoxyl 500 mg/5 ml pó para suspensão oral	FR/H/0600/004	8352211	SMITH KLINE & FRENCH PORTUGUESA-PRODUTOS FARMACEUTICOS LDA	PT
Clamoxyl 1 g comprimidos dispersíveis	FR/H/0600/001	4667580	SMITH KLINE & FRENCH PORTUGUESA-PRODUTOS FARMACEUTICOS LDA	PT
Flemoxin Solutab®	not available	2234284	ASTELLAS FARMA LDA.	PT
Flemoxin Solutab®	not available	4695383	ASTELLAS FARMA LDA.	PT
Flemoxin Solutab®	not available	2234482	ASTELLAS FARMA LDA.	PT
Flemoxin Solutab®	not available	4695581	ASTELLAS FARMA LDA.	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
Clamoxyl 500 mg cápsulas	UK/H/6220/02	4667481	BEECHAM PORTUGUESA, PRODUTOS FARMACÊUTICOS E QUÍMICOS, LDA	PT
Clamoxyl 500 mg cápsulas	UK/H/6220/02	8352112	BEECHAM PORTUGUESA, PRODUTOS FARMACÊUTICOS E QUÍMICOS, LDA	PT
Clamoxyl 250 mg/5 ml pó para suspensão oral	FR/H/0600/003	8352203	SMITH KLINE & FRENCH PORTUGUESA-PRODUTOS FARMACEUTICOS LDA	PT
Clamoxyl 1 g comprimidos dispersíveis	FR/H/600/01/MR	8436816	SMITH KLINE & FRENCH PORTUGUESA-PRODUTOS FARMACEUTICOS LDA	PT
Clamoxyl 500 mg/5 ml pó para suspensão oral	FR/H/0600/004	8352211	SMITH KLINE & FRENCH PORTUGUESA-PRODUTOS FARMACEUTICOS LDA	PT
Clamoxyl 1 g comprimidos dispersíveis	FR/H/0600/001	4667580	SMITH KLINE & FRENCH PORTUGUESA-PRODUTOS FARMACEUTICOS LDA	PT
OSPAMOX 125 MG/5 ML	AT/H/0116/003	5022082	SANDOZ FARMACÊUTICA LDA.	PT
AMOXICILINA SANDOZ 1000 mg comprimate filmate	AT/H/0187/003	5475/2013/08	S.C. SANDOZ S.R.L.	RO
AMOXICILINA SANDOZ 500 mg comprimate filmate	AT/H/0187/001	5474/2013/01	S.C. SANDOZ S.R.L.	RO
AMOXICILINA SANDOZ 500 mg comprimate filmate	AT/H/0187/001	5474/2013/02	S.C. SANDOZ S.R.L.	RO
AMOXICILINA SANDOZ 500 mg comprimate filmate	AT/H/0187/001	5474/2013/04	S.C. SANDOZ 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
AMOXICILINA SANDOZ 500 mg comprimate filmate	AT/H/0187/001	5474/2013/03	S.C. SANDOZ S.R.L.	RO
AMOXICILINA SANDOZ 500 mg comprimate filmate	AT/H/0187/001	5474/2013/05	S.C. SANDOZ S.R.L.	RO
AMOXICILINA SANDOZ 500 mg comprimate filmate	AT/H/0187/001	5474/2013/06	S.C. SANDOZ S.R.L.	RO
AMOXICILINA SANDOZ 500 mg comprimate filmate	AT/H/0187/001	5474/2013/07	S.C. SANDOZ S.R.L.	RO
AMOXICILINA SANDOZ 500 mg comprimate filmate	AT/H/0187/001	5474/2013/09	S.C. SANDOZ S.R.L.	RO
AMOXICILINA SANDOZ 500 mg comprimate filmate	AT/H/0187/001	5474/2013/08	S.C. SANDOZ S.R.L.	RO
AMOXICILINA SANDOZ 500 mg comprimate filmate	AT/H/0187/001	5474/2013/10	S.C. SANDOZ S.R.L.	RO
AMOXICILINA SANDOZ 1000 mg comprimate filmate	AT/H/0187/003	5475/2013/01	S.C. SANDOZ S.R.L.	RO
AMOXICILINA SANDOZ 1000 mg comprimate filmate	AT/H/0187/003	5475/2013/02	S.C. SANDOZ S.R.L.	RO
AMOXICILINA SANDOZ 1000 mg comprimate filmate	AT/H/0187/003	5475/2013/04	S.C. SANDOZ S.R.L.	RO
AMOXICILINA SANDOZ 1000 mg comprimate filmate	AT/H/0187/003	5475/2013/03	S.C. SANDOZ 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
AMOXICILINA SANDOZ 1000 mg comprimate filmate	AT/H/0187/003	5475/2013/05	S.C. SANDOZ S.R.L.	RO
AMOXICILINA SANDOZ 1000 mg comprimate filmate	AT/H/0187/003	5475/2013/06	S.C. SANDOZ S.R.L.	RO
AMOXICILINA SANDOZ 1000 mg comprimate filmate	AT/H/0187/003	5475/2013/07	S.C. SANDOZ S.R.L.	RO
AMOXICILINA SANDOZ 1000 mg comprimate filmate	AT/H/0187/003	5475/2013/10	S.C. SANDOZ S.R.L.	RO
AMOXICILINA SANDOZ 1000 mg comprimate filmate	AT/H/0187/003	5475/2013/09	S.C. SANDOZ S.R.L.	RO
Amimox 375 mg filmdragerade tabletter	not available	11538	MEDA AB	SE
Amimox 500 mg filmdragerade tabletter	not available	11647	MEDA AB	SE
Imacillin 750 mg tabletter	not available	11801	MEDA AB	SE
Amimox 750 mg filmdragerade tabletter	not available	11539	MEDA AB	SE
Amimox 50 mg/ml granulat till oral suspension	not available	11541	MEDA AB	SE
Imacillin 1 g tabletter	not available	11802	MEDA AB	SE
Amimox 100 mg/ml granulat till oral suspension	not available	11837	MEDA AB	SE
Amimox 125 mg granulat till oral suspension, dospåse	not available	11540	MEDA AB	SE
Duomox 1 000	not available	15/0383/95-S	ASTELLAS PHARMA S.R.O.	SK
Duomox 125	not available	15/0383/95-S	ASTELLAS PHARMA S.R.O.	SK

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the Member State	Member State where product is authorised
Duomox 250	not available	15/0383/95-S	ASTELLAS PHARMA S.R.O.	SK
Duomox 375	not available	15/0383/95-S	ASTELLAS PHARMA S.R.O.	SK
Duomox 750	not available	15/0383/95-S	ASTELLAS PHARMA S.R.O.	SK
Duomox 500	not available	15/0383/95-S	ASTELLAS PHARMA S.R.O.	SK
Amoxicillin 125mg/5ml Oral Suspension Sugar Free	not available	PL 30464/0088	ATHLONE PHARMACEUTICALS LIMITED	UK
Amoxicillin 250mg/5ml Oral Suspension Sugar Free	not available	PL 30464/0089	ATHLONE PHARMACEUTICALS LIMITED	UK
Amoxicillin 125mg/5ml Oral Suspension Sugar Free	not available	PL 30464/0090	ATHLONE PHARMACEUTICALS LIMITED	UK
Amoxicillin 250mg/5ml Oral Suspension Sugar Free	not available	PL 30464/0091	ATHLONE PHARMACEUTICALS LIMITED	UK
Amoxicillin 500 mg Capsules	not available	PL 30464/0106	ATHLONE PHARMACEUTICALS LIMITED	UK
Amoxicillin Oral Suspension BP 250 mg/5 ml	not available	PL 30464/0010	ATHLONE PHARMACEUTICALS LIMITED	UK
Amoxicillin 250 mg Capsules	not available	PL 30464/0105	ATHLONE PHARMACEUTICALS LIMITED	UK
Amoxicillin Sugar Free Suspension BP 125mg/5ml	not available	PL 21880/0123	MEDREICH PLC	UK
Amoxicillin Sugar Free Suspension BP 250mg/5ml	not available	PL 21880/0124	MEDREICH PLC	UK
Amoxil Capsules 250 mg	UK/H/6220/01	PL 00038/0103	BEECHAM GROUP PLC	UK
Amoxil Capsules 500 mg	UK/H/6220/02	PL 00038/0105	BEECHAM GROUP PLC	UK
Amoxil Vials for Injection 1 g	UK/H/6220/06	PL 00038/0225	BEECHAM GROUP PLC	UK

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the Member State	Member State where product is authorised
Amoxil Sachets 3 G Sucrose-Free	UK/H/6220/07	PL 00038/0334	BEECHAM GROUP PLC	UK
Amoxil Paediatric Suspension	UK/H/6220/03	PL 00038/0107	BEECHAM GROUP PLC	UK
Amoxil Vials for Injection 250 mg	UK/H/6220/04	0038/0221	BEECHAM GROUP PLC	UK
Amoxil Vials for Injection 500 mg	UK/H/6220/05	PL 00038/0222	BEECHAM GROUP PLC	UK
Amoxil Capsules 250 mg	UK/H/6220/01	PL 00038/0103	BEECHAM GROUP PLC	UK
Amoxil Capsules 500 mg	UK/H/6220/02	PL 00038/0105	BEECHAM GROUP PLC	UK
Amoxil Vials for Injection 1 g	UK/H/6220/06	PL 00038/0225	BEECHAM GROUP PLC	UK
Amoxil Sachets 3 G Sucrose-Free	UK/H/6220/07	PL 00038/0334	BEECHAM GROUP PLC	UK
Amoxil Paediatric Suspension	UK/H/6220/03	PL 00038/0107	BEECHAM GROUP PLC	UK
Amoxil Vials for Injection 250 mg	UK/H/6220/04	0038/0221	BEECHAM GROUP PLC	UK
Amoxil Vials for Injection 500 mg	UK/H/6220/05	PL 00038/0222	BEECHAM GROUP PLC	UK
Amoxicillin 250mg Capsules	not available	PL 36722/0019	SPECIAL CONCEPT DEVELOPMENT (UK) LTD	UK
Amoxicillin 500mg Capsules	not available	PL 36722/0020	SPECIAL CONCEPT DEVELOPMENT (UK) LTD	UK
Amoxicillin 250 mg/5 ml powder for oral suspension	AT/H/0116/004	PL 04416/0485	SANDOZ LTD	UK
Amoxicillin 500 mg hard capsules	AT/H/0116/002	PL 04416/0577	SANDOZ LTD	UK
Amoxicillin 250 mg hard capsules	AT/H/0116/001	PL 04416/0576	SANDOZ LTD	UK
Amoxicillin 125mg/5ml Suspension	AT/H/0116/003	PL 04416/0484	SANDOZ LTD	UK

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the Member State	Member State where product is authorised
Amoxicillin Mixture BP 125mg/5ml	not available	PL 06464/1420	WAYMADE PLC	UK
Amoxicillin Mixture BP 250mg/5ml	not available	PL 06464/1421	WAYMADE PLC	UK
Amoxicillin 250mg Capsules	not available	PL 06464/1418	WAYMADE PLC	UK
Amoxicillin 500mg Capsules	not available	PL 06464/1419	WAYMADE PLC	UK
Amoxicillin 500 mg Capsules	not available	PL 44041/0002	NOUMED LIFE SCIENCES	UK
Amoxicillin 250 mg Capsules	not available	PL 40496/0041	BRILL PHARMA LIMITED	UK
Amoxicillin 500 mg Capsules	not available	PL 40496/0042	BRILL PHARMA LIMITED	UK