26 October 2017
EMA/726764/2017
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: amoxicillin / clavulanate

Procedure no.: PSUSA/00000188/201703

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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
Curam intravenös 500 mg/50 mg - Pulver zur Herstellung einer Injektions-/Infusionslösung	NL/H/0541/004	1-26283	SANDOZ GMBH	AT
Curam intravenös 500 mg/100 mg - Pulver zur Herstellung einer Injektions-/Infusionslösung	NL/H/0541/003	1-26284	SANDOZ GMBH	AT
Clavamox Duo - Trockensaft	939.404	1-22155	SANDOZ GMBH	AT
Clavamox intravenös 2,2 g - Trockensubstanz zur Infusionsbereitung	934.284	1-21170	SANDOZ GMBH	AT
Clavamox 625 mg - Filmtabletten	934.286	1-21175	SANDOZ GMBH	AT
Clavamox 1 g - Filmtabletten	936.596	1-21397	SANDOZ GMBH	AT
Clavamox intravenös 1,1 g - Trockensubstanz zur Infusionsbereitung	934.281	1-21171	SANDOZ GMBH	AT
Clavamox intravenös 550 mg - Trockenstechampulle	934.283	1-21173	SANDOZ GMBH	AT
Curam intravenös 1000 mg/200 mg - Pulver zur Herstellung einer Injektions-/Infusionslösung	NL/H/0541/001	1-26281	SANDOZ GMBH	AT
Curam intravenös 2000 mg/200 mg - Pulver zur Herstellung einer Infusionslösung	NL/H/0541/002	1-26282	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
Curam intravenös 500 mg/50 mg - Pulver zur Herstellung einer Injektions-/Infusionslösung	NL/H/0541/004	1-26283	SANDOZ GMBH	AT
Curam intravenös 500 mg/100 mg - Pulver zur Herstellung einer Injektions-/Infusionslösung	NL/H/0541/003	1-26284	SANDOZ GMBH	AT
Clavamox Duo – Trockensaft	939.404	1-22155	SANDOZ GMBH	AT
Clavamox intravenös 2,2 g - Trockensubstanz zur Infusionsbereitung	934.284	1-21170	SANDOZ GMBH	AT
Clavamox 625 mg – Filmtabletten	934.286	1-21175	SANDOZ GMBH	AT
Clavamox 1 g – Filmtabletten	936.596	1-21397	SANDOZ GMBH	AT
Clavamox intravenös 1,1 g - Trockensubstanz zur Infusionsbereitung	934.281	1-21171	SANDOZ GMBH	AT
Clavamox intravenös 550 mg - Trockenstechampulle	934.283	1-21173	SANDOZ GMBH	AT
Curam intravenös 1000 mg/200 mg - Pulver zur Herstellung einer Injektions-/Infusionslösung	NL/H/0541/001	1-26281	SANDOZ GMBH	AT
Curam intravenös 2000 mg/200 mg - Pulver zur Herstellung einer Infusionslösung	NL/H/0541/002	1-26282	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
Augmentin 400 mg/57 mg/5 ml Pulver zur Herstellung einer Suspension zum Einnehmen	UK/H/4737/001	1-22152	GLAXOSMITHKLINE PHARMA GMBH.	AT
Augmentin 2.000 mg/200 mg Pulver zur Herstellung einer Infusionslösung	DE/H/2809/004	1-18136	GLAXOSMITHKLINE PHARMA GMBH.	AT
Augmentin 1.000 mg/100 mg Pulver zur Herstellung einer Injektions- oder Infusionslösung	DE/H/2809/006	1-18135	GLAXOSMITHKLINE PHARMA GMBH.	AT
Augmentin 500 mg/50 mg Pulver zur Herstellung einer Injektions- oder Infusionslösung	DE/H/2809/005	1-18137	GLAXOSMITHKLINE PHARMA GMBH.	AT
Augmentin 875 mg/125 mg Filmtabletten	DE/H/2868/02	1-21396	GLAXOSMITHKLINE PHARMA GMBH.	AT
Clavucid Solutab 875/125, dispergeerbare tabletten	not available	BE202456	ASTELLAS PHARMA B.V., OFFICE BE	BE
CLAVUCID SOLUTAB 875mg/125mg, DISPERGIERBARE TABLETTEN	not available	BE202456	ASTELLAS PHARMA B.V., OFFICE BE	BE
Amoxiclav Sandoz 2000 mg/200 mg poeder voor oplossing voor infusie	NL/H/0541/002	BE271214	SANDOZ N.V.	BE
Amoxiclav Sandoz 1000 mg/200 mg poeder voor oplossing voor injectie / infusie	NL/H/0541/001	BE271196	SANDOZ N.V.	BE

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the Member State	Member State where product is authorised
Amoclane 875/125 mg poeder voor orale suspensie	not available	BE273856	EUROGENERICS SA	BE
Amoclane 875/125 mg poudre pour suspension buvable	not available	BE273856	EUROGENERICS N.V./S.A.	BE
Amoclane 875/125 mg Pulver zur Herstellung einer Suspension zum Einnehmen	not available	BE273856	EUROGENERICS N.V./S.A.	BE
Amoxiclav Sandoz 2000 mg/200 mg poeder voor oplossing voor infusie	NL/H/0541/002	BE271214	SANDOZ N.V.	BE
Amoxiclav Sandoz 1000 mg/200 mg poeder voor oplossing voor injectie / infusie	NL/H/0541/001	BE271196	SANDOZ N.V.	BE
Augmentin 875 mg/125 mg, comprimés pelliculés	DE/H/2868/02	BE439293	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Augmentin Retard 1.000 mg/62,5 mg Retardtabletten	BE/H/0168/001	BE240457	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Augmentin 1.000 mg/200 mg Pulver zur Herstellung einer Injektions- oder Infusionslösung	DE/H/2809/003	BE135213	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Augmentin P 1000 mg/100 mg poeder voor oplossing voor injectie of infusie	DE/H/2809/006	BE135204	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Augmentin 500 mg/125 mg filmomhulde tabletten	UK/H/4739/001	BE439284	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
Augmentin 500 mg/125 mg Pulver zur Herstellung einer Suspension zum Einnehmen in Beuteln	BE/H/0159/001	BE159345	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Augmentin 250 mg/62,5 mg/5 ml Pulver zur Herstellung einer Suspension zum Einnehmen	BE/H/209/02/MR	BE125413	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Augmentin 500 mg/125 mg, comprimés pelliculés	UK/H/4739/001	BE125334	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Augmentin 125 mg+31,25 mg+5 ml, poudre pour suspension buvable	BE/H/0209/001	BE125404	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Augmentin 2000 mg/200 mg, poudre pour solution pour perfusion	DE/H/2809/004	BE135511	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Augmentin P 500 mg+50 mg, poudre pour solution pour injection ou perfusion	DE/H/2809/005	BE135195	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Augmentin 875 mg/125 mg, comprimés pelliculés	DE/H/2868/02	BE200803	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Аугментин 250 mg/125 mg филмирани таблетки	UK/H/4735/001	20000315	GLAXOSMITHKLINE EOOD	BG
Аугментин 500 mg/125 mg филмирани таблетки	UK/H/4738/001	20000314	GLAXOSMITHKLINE EOOD	BG
Аугментин 125 mg/31,25 mg/5 ml прах за перорална суспензия	DE/H/2868/004	20000317	GLAXOSMITHKLINE EOOD	BG

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the Member State	Member State where product is authorised
Аугментин 250 mg/62,5 mg/5 ml прах за перорална суспензия	DE/H/2868/005	20000318	GLAXOSMITHKLINE EOOD	BG
Аугментин 400 mg/57 mg/5 ml прах за перорална суспензия	DE/H/2868/006	9900089	GLAXOSMITHKLINE EOOD	BG
Аугментин ES 600 mg/42,9 mg/5 ml прах за перорална суспензия	PT/H/0684/001	20050447	GLAXOSMITHKLINE EOOD	BG
Аугментин SR 1000 mg/62,5 mg таблетки с удължено освобождаване	BE/H/168/001	20060523	GLAXOSMITHKLINE EOOD	BG
Аугментин 875 mg/125 mg филмирани таблетки	DE/H/2868/02	20000316	GLAXOSMITHKLINE EOOD	BG
Forcid Solutab 875/125mg, δισκίο/διασπειρόμενο δισκίο	HU/H/0386/001	19680	ASTELLAS PHARMACEUTICALS A.E.B.E.	CY
Augmentin 400 mg/57 mg/5 ml κόνις για πόσιμο εναιώρημα (γεύση φράουλα)	UK/H/4737/01/MR	19702	SMITHKLINE BEECHAM LTD	CY
Augmentin 500 mg/125 mg επικαλυμμένα με λεπτό υμένιο δισκία	UK/H/4738/001	12656	SMITHKLINE BEECHAM LTD	CY
Augmentin ES 600 mg/42.9 mg/5 ml κόνις για πόσιμο εναιώρημα	PT/H/684/001	20148	SMITHKLINE BEECHAM LTD	CY
Augmentin 875 mg/125 mg επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/2868/02	19515	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
AUGMENTIN DUO, prášek pro perorální suspenzi	UK/H/4737/001	15/265/00-C	SMITHKLINE BEECHAM LTD	CZ
Augmentin 600 mg, prášek pro injekční nebo infuzní roztok	DE/H/2809/002	15/147/88-A/C	SMITHKLINE BEECHAM LTD	CZ
Augmentin 1,2 g, prášek pro injekční nebo infuzní roztok	DE/H/2809/003	15/147/88-B/C	SMITHKLINE BEECHAM LTD	CZ
Augmentin 625 mg, potahované tablety	UK/H/4738/001	15/141/84-B/C	SMITHKLINE BEECHAM LTD	CZ
Augmentin SR, 1000 mg/62,5 mg tablety s prodlouženým uvolňováním	BE/H/0168/001	15/200/03-C	SMITHKLINE BEECHAM LTD	CZ
Augmentin 1 g, potahované tablety	DE/H/2868/02	15/644/96-C	SMITHKLINE BEECHAM LTD	CZ
Augmentan 500 mg/50 mg Pulver zur Herstellung einer Injektions- oder Infusionslösung	DE/H/2809/005	86021.00.00	GLAXOSMITHKLINE GMBH & CO. KG	DE
Augmentan 100 mg/12,5 mg/ml Pulver zur Herstellung einer Suspension zum Einnehmen für Kinder	DE/H/2868/007-008/MR	86130.00.00	GLAXOSMITHKLINE GMBH & CO. KG	DE
Augmentan 100 mg/12,5 mg/ml Pulver zur Herstellung einer Suspension zum Einnehmen für Säuglinge	DE/H/2868/007-008/MR	86131.00.00	GLAXOSMITHKLINE GMBH & CO. KG	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
Augmentan 1000 mg/100 mg Pulver zur Herstellung einer Injektions- oder Infusionslösung	DE/H/2809/006/MR	86022.00.00	GLAXOSMITHKLINE GMBH & CO. KG	DE
Augmentan® i.v. 600 mg 500 mg/100 mg Pulver zur Herstellung einer Injektions- oder Infusionslösung	DE/H/2809/002	1927.02.04	GLAXOSMITHKLINE GMBH & CO. KG	DE
Augmentan® i.v. 1,2 g 1000 mg/200 mg Pulver zur Herstellung einer Injektions- oder Infusionslösung	DE/H/2809/003	1927.01.04	GLAXOSMITHKLINE GMBH & CO. KG	DE
Augmentan® Trockensaft 25 mg/6,25 mg pro ml 125 mg/31,25 mg/5 ml Pulver zur Herstellung einer Suspension zum Einnehmen	DE/H/2868/004	1927.00.03	GLAXOSMITHKLINE GMBH & CO. KG	DE
Augmentan® forte Trockensaft 50 mg/12,5 mg pro ml	DE/H/2868/05/MR	7351.00.00	GLAXOSMITHKLINE GMBH & CO. KG	DE
Augmentan® Kindersaft 400 mg/57 mg/5 ml Pulver zur Herstellung einer Suspension zum Einnehmen	DE/H/2868/06/MR	35944.00.00	GLAXOSMITHKLINE GMBH & CO. KG	DE
Augmentan® i.v. 2,2 g 2000 mg/200 mg Pulver zur Herstellung einer Infusionslösung	DE/H/2809/004/MR	1927.00.04	GLAXOSMITHKLINE GMBH & CO. KG	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
Augmentan® Filmtabletten 875/125 mg	DE/H/2868/02	34923.00.00	GLAXOSMITHKLINE GMBH & CO. KG	DE
Augmentan® Tropfen 50 mg/12,5 mg pro ml für Säuglinge 50 mg/12,5 mg/ml Pulver zur Herstellung einer Suspension zum Einnehmen	DE/H/2868/003	7351.01.00	GLAXOSMITHKLINE GMBH & CO. KG	DE
Augmentan® 500 mg/125 mg Filmtabletten	UK/H/4738/01/MR	92660.00.00	GLAXOSMITHKLINE GMBH & CO. KG	DE
Spektramox	not available	12255	MEDA AS	DK
Spektramox	not available	12257	MEDA AS	DK
Augmentin, 400 mg/57 mg/5 ml suukaudse suspensiooni pulber	UK/H/4737/001	213498	SMITHKLINE BEECHAM LTD	EE
Augmentin, 1000 mg/200 mg süste- või infusioonilahuse pulber	DE/H/2809/003	241698	SMITHKLINE BEECHAM LTD	EE
Augmentin, 500 mg+125 mg õhukese polümeerikattega tabletid	UK/H/4738/001	213298	SMITHKLINE BEECHAM LTD	EE
Augmentin, 875 mg/125 mg õhukese polümeerikattega tabletid	DE/H/2868/02	213398	SMITHKLINE BEECHAM LTD	EE
Augmentine 1.000 mg/200 mg polvo para solución inyectable y para perfusión	DE/H/2809/003	58.215	GLAXOSMITHKLINE S.A.	ES
Augmentine 500 mg/125 mg comprimidos recubiertos con película	UK/H/4739/001	56.685	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
Augmentine 100 mg/ml +12,5 mg/ml polvo para suspensión oral	DE/H/2868/008	59.051	GLAXOSMITHKLINE S.A.	ES
Augmentine 2.000 mg/200 mg polvo para solución para perfusión	DE/H/2809/004	58.216	GLAXOSMITHKLINE S.A.	ES
Augmentine 875 mg/125 mg polvo para suspensión oral en sobres	IT/H/0270/001	59.518	GLAXOSMITHKLINE, S.A.	ES
Augmentine 500 mg/125 mg polvo para suspensión oral en sobres	BE/H/0159/001	56.683	GLAXOSMITHKLINE S.A.	ES
Augmentine Plus 1.000 mg/62,5 mg comprimidos de liberación prolongada	BE/H/0168/001	65.459	GLAXOSMITHKLINE, S.A.	ES
Augmentine 875 mg/125 mg comprimidos recubiertos con película	DE/H/2868/02	59.515	GLAXOSMITHKLINE S.A.	ES
DUONASA 500 mg/125 mg comprimidos recubiertos con película.	not available	58988	LABORATORIOS NORMON, S.A.	ES
Augmentin 80/11,4 mg/ml jauhe oraalisuspensiota varten	UK/H/4737/001	12581	SMITHKLINE BEECHAM LTD	FI
Augmentin 875 mg/125 mg kalvopäällysteiset tabletit	DE/H/2868/02	12199	SMITHKLINE BEECHAM LTD	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
AMOXICILLINE/ACIDE CLAVULANIQUE BIOGARANÒ 100 mg/12,5 mg par ml, NOURRISSON, poudre pour suspension buvable (rapport amoxicilline/acide clavulanique : 8/1)	not available	34009 353 555 2 2	BIOGARAN	FR
AMOXICILLINE/ACIDE CLAVULANIQUE BIOGARAN® 1 g / 125 mg ADULTES, poudre pour suspension buvable en sachet-dose (rapport amoxicilline/acide clavulanique : 8/1)	not available	3400935335012	BIOGARAN	FR
AMOXICILLINE/ACIDE CLAVULANIQUE BIOGARAN® 1 g / 125 mg ADULTES, poudre pour suspension buvable en sachet-dose (rapport amoxicilline/acide clavulanique : 8/1)	not available	3400935334879	BIOGARAN	FR
AMOXICILLINE/ACIDE CLAVULANIQUE BIOGARAN® 100 mg/12,5 mg par ml, ENFANT, poudre pour suspension buvable (rapport amoxicilline/acide clavulanique : 8/1)	not available	34009 353 556 9 0	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/ACIDE CLAVULANIQUE BIOGARAN® 500 mg/62,5 mg ADULTES, comprimé pelliculé (rapport amoxicilline/acide clavulanique : 8/1)	not available	3400935335241	BIOGARAN	FR
AMOXICILLINE/ACIDE CLAVULANIQUE BIOGARAN® 500 mg/62,5 mg ADULTES, comprimé pelliculé (rapport amoxicilline/acide clavulanique : 8/1)	not available	3400935335470	BIOGARAN	FR
AUGMENTIN 100 mg/12,50 mg par ml NOURRISSONS, poudre pour suspension buvable (rapport amoxicilline/acide clavulanique : 8/1)	DE/H/2868/007-008/MR	NL16485	LABORATOIRE GLAXOSMITHKLINE	FR
AUGMENTIN 1 g/200 mg, poudre pour solution injectable / pour perfusion (IV)	DE/H/2809/003/MR	NL 13670-1	LABORATOIRE GLAXOSMITHKLINE	FR
AUGMENTIN 1 g/200 mg, poudre et solvant pour solution injectable / pour perfusion (IV).	DE/H/2809/003/MR	NL13670-2	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
AUGMENTIN 100 mg/12,50 mg par ml ENFANTS, poudre pour suspension buvable (rapport amoxicilline/acide clavulanique: 8/1)	DE/H/2868/007-008/MR	NL21290	LABORATOIRE GLAXOSMITHKLINE	FR
AUGMENTIN 2 g/200 mg ADULTES, poudre pour solution pour perfusion	DE/H/2809/004	NL13672	LABORATOIRE GLAXOSMITHKLINE	FR
DUAMENTIN 1 g/62,5 mg ADULTES, comprimé pelliculé (rapport amoxicilline/acide clavulanique: 16/1)	BE/H/168/001	NL26435	LABORATOIRE GLAXOSMITHKLINE	FR
AUGMENTIN 500 mg/50 mg, poudre pour solution injectable / pour perfusion (IV)	DE/H/2809/005/MR	NL13674-1	LABORATOIRE GLAXOSMITHKLINE	FR
Forcid Solutab 875/125mg, δισκίο/διασπειρόμενο δισκίο	HU/H/0386/001	53986/21-8-2008	ASTELLAS PHARMACEUTICALS A.E.B.E.	GR
Forcid Solutab 125/31,25 δισκίο/διασπειρόμενο δισκίο	NL/H/0224/001	53983/21-8-2008	ASTELLAS PHARMA EUROPE B.V.	GR
Forcid Solutab 500/125 δισκίο/διασπειρόμενο δισκίο	NL/H/0224/004	53985/21-8-2008	ASTELLAS PHARMA EUROPE B.V.	GR
Forcid Solutab 250/62,5 δισκίο/διασπειρόμενο δισκίο	NL/H/0224/002	53984/21-08-2008	ASTELLAS PHARMA EUROPE B.V.	GR
Augmentin 250 mg/62.5 mg/5 ml κόνις για πόσιμο εναιώρημα	BE/H/209/02/MR	39097/11-6-2007	GLAXOSMITHKLINE, S.A.	GR

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Augmentin 500 mg/125 mg επικαλυμμένα με λεπτό υμένιο δισκία	UK/H/4738/01/MR	1759904	GLAXOSMITHKLINE AEBE	GR
Augmentin 1000 mg/200 mg κόνις για ενέσιμο διάλυμα ή διάλυμα προς έγχυση	DE/H/2809/003/MR	1759908	GLAXOSMITHKLINE AEBE	GR
Augmentin 875 mg/125 mg επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/2868/02	1759911	GLAXOSMITHKLINE AEBE	GR
Augmentin 400 mg/57 mg/5 ml κόνις για πόσιμο εναιώρημα (γεύση φράουλα)	UK/H/4737/01/MR	15440/22-6-09	GLAXOSMITHKLINE AEBE	GR
Augmentin 875 mg + 125 mg filmom obložene tablete	not available	HR-H-170450439	GLAXOSMITHKLINE D.O.O.	HR
Augmentin 400 mg + 57 mg/5 ml prašak za oralnu suspenziju	not available	HR-H-709518864	GLAXOSMITHKLINE D.O.O.	HR
AUGMENTIN 1000 mg + 200 mg prašak za otopinu za injekciju ili infuziju	not available	UP/I-530-09/12-02/334	GLAXOSMITHKLINE D.O.O.	HR
Forcid Solutab 875 mg/125 mg diszpergálódó tabletta	HU/H/0386/001	OGYI-T- 9988/01	ASTELLAS PHARMA KFT	HU
Forcid Solutab 250 mg/62,5 mg diszpergálódó tabletta	not available	OGYI-T-9020/01	ASTELLAS PHARMA KFT	HU
Forcid Solutab 500 mg/125 mg diszpergálódó tabletta	not available	OGYI-T-9020/02	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
Augmentin 500 mg/100 mg por oldatos injekcióhoz vagy infúzióhoz	DE/H/2809/002	OGYI-T-1352/12	GLAXOSMITHKLINE KFT.	HU
Augmentin 250 mg/125 mg filmtabletta	UK/H/4735/001	OGYI-T-1352/03	GLAXOSMITHKLINE KFT.	HU
Augmentin DUO 400 mg/57 mg/5 ml por belsőleges szuszpenzióhoz	UK/H/4737/001	OGYI-T-1352/05	GLAXOSMITHKLINE KFT.	HU
Augmentin DUO 400 mg/57 mg/5 ml por belsőleges szuszpenzióhoz	UK/H/4737/001	OGYI-T-1352/06	GLAXOSMITHKLINE KFT.	HU
Augmentin 500 mg/100 mg por oldatos injekcióhoz vagy infúzióhoz	DE/H/2809/002	OGYI-T-1352/09	GLAXOSMITHKLINE KFT.	HU
Augmentin 1000 mg/200 mg por oldatos injekcióhoz vagy infúzióhoz	DE/H/2809/003/MR	OGYI-T-1352/13	GLAXOSMITHKLINE KFT.	HU
Augmentin 125 mg/31,25 mg/5 ml por belsőleges szuszpenzióhoz	DE/H/2868/004	OGYI-T-1352/01	GLAXOSMITHKLINE KFT.	HU
Augmentin 500 mg/125 mg filmtabletta	UK/H/4739/001	OGYI-T-1352/04	GLAXOSMITHKLINE KFT.	HU
Augmentin 250 mg/62,5 mg/5 ml por belsőleges szuszpenzióhoz	DE/H/2868/005	OGYI-T-1352/02	GLAXOSMITHKLINE KFT.	HU
Augmentin DUO 875 mg/125 mg filmtabletta	DE/H/2868/02	OGYI-T-1352/08	GLAXOSMITHKLINE KFT.	HU
Augmentin 250 mg/125 mg film-coated tablets	UK/H/4735/01/MR	PA 1077/93/7	GLAXOSMITHKLINE (IRELAND) LIMITED	IE

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the Member State	Member State where product is authorised
Augmentin DUO 400mg/57 mg/5ml powder for oral suspension	UK/H/4737/01/MR	PA 1077/19/2	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Augmentin Intravenous 500 mg/100 mg powder for solution for injection or infusion	DE/H/2809/002	PA 1077/93/1	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Augmentin Intravenous 1000 mg/200 mg powder for solution for injection or infusion	DE/H/2809/003	PA 1077/93/2	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Augmentin Paediatric 125 mg/31.25 mg per 5 ml powder for oral suspension	DE/H/2868/04	PA 1077/93/4	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Augmentin 500 mg/125 mg film-coated tablets	UK/H/4738/001	PA 1077/19/3	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Augmentin 875 mg/125 mg film-coated tablets	DE/H/2868/02	PA 1077/19/5	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Augmentin 875 mg/125 mg film-coated tablets	DE/H/2868/002	PA 678/12/6	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Augmentin 400 mg/57 mg/5 ml – (80 mg/11,4 mg/ml) mixtúruduft, dreifa (jarðarberjabragð)	UK/H/4737/01/MR	970163	GLAXOSMITHKLINE PHARMA A/S	IS
Augmentin 1000 mg/200 mg stungulyfs-/innrennslisstofn, lausn	DE/H/2809/003	980044	GLAXOSMITHKLINE PHARMA A/S	IS
Augmentin 500 mg/125 mg filmuhúðaðar töflur	UK/H/4738/01/MR	870191	GLAXOSMITHKLINE PHARMA A/S	IS
Augmentin 250 mg/62,5 mg/5 ml - (50 mg/12,5 mg/ml) mixtúruduft, dreifa	BE/H/209/02/MR	870194 (IS)	GLAXOSMITHKLINE PHARMA A/S	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
Augmentin 875 mg/125 mg filmuhúðaðar töflur	DE/H/2868/02	950055	GLAXOSMITHKLINE PHARMA A/S	IS
NEODUPLAMOX 875 mg/125 mg compresse rivestite con film	not available	026141147	VALEAS S.P.A.	IT
NEODUPLAMOX 875 mg/125 mg polvere per sospensione orale bustine	not available	026141198	VALEAS S.P.A.	IT
NEODUPLAMOX bambini 400 mg/57 mg/5 ml polvere per sospensione orale	not available	026141200	VALEAS S.P.A.	IT
NEODUPLAMOX bambini 400 mg/57 mg/5 ml polvere per sospensione orale	not available	026141212	VALEAS S.P.A.	IT
NEODUPLAMOX bambini 400 mg/57 mg/5 ml polvere per sospensione orale	not available	026141224	VALEAS S.P.A.	IT
NEODUPLAMOX bambini 400 mg/57 mg polvere per sospensione orale bustine	not available	026141236	VALEAS S.P.A.	IT
Augmentin 875 mg/125 mg compresse rivestite con film	DE/H/2868/02	026089211	GLAXOSMITHKLINE S.P.A.	IT
Augmentin 875 mg/125 mg compresse rivestite con film	DE/H/2868/02/MR	026089223	GLAXOSMITHKLINE S.P.A.	IT
Augmentin 875 mg/125 mg compresse rivestite con film	DE/H/2868/02	026089235	GLAXOSMITHKLINE S.P.A.	IT

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the Member State	Member State where product is authorised
Augmentin 875 mg/125 mg compresse rivestite con film	DE/H/2868/02	026089247	GLAXOSMITHKLINE S.P.A.	IT
Augmentin 875 mg/125 mg compresse rivestite con film	DE/H/2868/02	026089250	GLAXOSMITHKLINE S.P.A.	IT
Augmentin 875 mg/125 mg compresse rivestite con film	DE/H/2868/02	026089262	GLAXOSMITHKLINE S.P.A.	IT
Augmentin 875 mg/125 mg compresse rivestite con film	DE/H/2868/02	026089274	GLAXOSMITHKLINE S.P.A.	IT
Augmentin 875 mg/125 mg compresse rivestite con film	DE/H/2868/02	026089312	GLAXOSMITHKLINE S.P.A.	IT
Augmentin 875 mg/125 mg compresse rivestite con film	DE/H/2868/02	026089286	GLAXOSMITHKLINE S.P.A.	IT
Augmentin 875 mg/125 mg compresse rivestite con film	DE/H/2868/02	026089298	GLAXOSMITHKLINE S.P.A.	IT
Augmentin 875 mg/125 mg compresse rivestite con film	DE/H/2868/02	026089300	GLAXOSMITHKLINE S.P.A.	IT
Augmentin bambini 400 mg/57 mg polvere per sospensione orale in bustine	IT/H/0270/002	026089146	GLAXOSMITHKLINE S.P.A.	IT
Augmentin bambini 400 mg/57 mg/5 ml polvere per sospensione orale	UK/H/4737/001	026089110	GLAXOSMITHKLINE S.P.A.	IT
Augmentin bambini 400 mg/57 mg/5 ml polvere per sospensione orale	UK/H/4737/001	026089122	GLAXOSMITHKLINE S.P.A.	IT

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the Member State	Member State where product is authorised
Augmentin bambini 400 mg/57 mg/5 ml polvere per sospensione orale	UK/H/4737/001	026089134	GLAXOSMITHKLINE S.P.A.	IT
Augmentin 1000 mg/200 mg polvere per soluzione iniettabile o per infusione	DE/H/2809/003	026089072	GLAXOSMITHKLINE S.P.A.	IT
Augmentin 2000 mg/200 mg polvere per soluzione per infusione	DE/H/2809/004	026089084	GLAXOSMITHKLINE S.P.A.	IT
Augmentin 875 mg/125 mg polvere per sospensione orale in bustine	IT/H/0270/001	026089108	GLAXOSMITHKLINE S.P.A.	IT
Augmentin 875 mg/125 mg compresse rivestite con film	DE/H/2868/02	026089019	GLAXOSMITHKLINE S.P.A.	IT
Augmentin 2000 mg/200 mg polvere per soluzione per infusione.	DE/H/2809/004/MR	026089437	GLAXOSMITHKLINE S.P.A.	IT
Augmentin 2000 mg/200 mg polvere per soluzione per infusione.	DE/H/2809/004/MR	026089449	GLAXOSMITHKLINE S.P.A.	IT
Augmentin 2000 mg/200 mg polvere per soluzione per infusione.	DE/H/2809/004/MR	026089452	GLAXOSMITHKLINE S.P.A.	IT
Augmentin 1000 mg/200 mg polvere per soluzione iniettabile o per infusione	DE/H/2809/003	026089387	GLAXOSMITHKLINE S.P.A.	IT
Augmentin 1000 mg/200 mg polvere per soluzione iniettabile o per infusione	DE/H/2809/003	026089399	GLAXOSMITHKLINE S.P.A.	IT
Augmentin 1000 mg/200 mg polvere per soluzione iniettabile o per infusione	DE/H/2809/003	026089401	GLAXOSMITHKLINE S.P.A.	IT

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the Member State	Member State where product is authorised
Augmentin 1000 mg/200 mg polvere per soluzione iniettabile o per infusione.	DE/H/2809/003	026089413	GLAXOSMITHKLINE S.P.A.	IT
Augmentin 1000 mg/200 mg polvere per soluzione iniettabile o per infusione	DE/H/2809/003	026089425	GLAXOSMITHKLINE S.P.A.	IT
Augmentin 400 mg/57 mg/5 ml milteliai geriamajai suspensijai	DE/H/2868/006	LT/1/98/3675/001	SMITHKLINE BEECHAM LTD	LT
Augmentin 500 mg /125 mg plėvele dengtos tabletės	UK/H/4738/001	LT/1/98/3675/014	BEECHAM GROUP PLC	LT
Augmentin 500 mg /125 mg plėvele dengtos tabletės	UK/H/4738/01	LT/1/98/3675/015	BEECHAM GROUP PLC	LT
Augmentin 500 mg /125 mg plėvele dengtos tabletės	UK/H/4738/01	LT/1/98/3675/016	BEECHAM GROUP PLC	LT
Augmentin 500 mg /125 mg plėvele dengtos tabletės	UK/H/4738/01	LT/1/98/3675/017	BEECHAM GROUP PLC	LT
Augmentin 500 mg /125 mg plėvele dengtos tabletės	UK/H/4738/01	LT/1/98/3675/013	BEECHAM GROUP PLC	LT
Augmentin 500 mg /125 mg plėvele dengtos tabletės	UK/H/4738/01	LT/1/98/3675/018	BEECHAM GROUP PLC	LT
Augmentin 500 mg /125 mg plėvele dengtos tabletės	UK/H/4738/01	LT/1/98/3675/019	BEECHAM GROUP PLC	LT
Augmentin 500 mg /125 mg plėvele dengtos tabletės	UK/H/4738/01	LT/1/98/3675/020	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
Augmentin 500 mg /125 mg plėvele dengtos tabletės	UK/H/4738/01	LT/1/98/3675/021	BEECHAM GROUP PLC	LT
Augmentin 500 mg /125 mg plėvele dengtos tabletės	UK/H/4738/01	LT/1/98/3675/022	BEECHAM GROUP LTD	LT
Augmentin 500 mg /125 mg plevele dengtos tabletes	UK/H/4738/01	LT/1/98/3675/023	BEECHAM GROUP PLC	LT
Augmentin 875 mg /125 mg plėvele dengtos tabletės	DE/H/2868/02	LT/1/98/3675/002	BEECHAM GROUP PLC	LT
Augmentin 875 mg /125 mg plėvele dengtos tabletės	DE/H/2868/02	LT/1/98/3675/003	BEECHAM GROUP PLC	LT
Augmentin 875 mg /125 mg plėvele dengtos tabletės	DE/H/2868/02	LT/1/98/3675/004	BEECHAM GROUP PLC	LT
Augmentin 875 mg /125 mg plėvele dengtos tabletės	DE/H/2868/02	LT/1/98/3675/005	BEECHAM GROUP PLC	LT
Augmentin 875 mg /125 mg plėvele dengtos tabletės	DE/H/2868/02	LT/1/98/3675/006	BEECHAM GROUP PLC	LT
Augmentin 875 mg /125 mg plėvele dengtos tabletės	DE/H/2868/02	LT/1/98/3675/007	BEECHAM GROUP PLC	LT
Augmentin 875 mg /125 mg plėvele dengtos tabletės	DE/H/2868/02	LT/1/98/3675/008	BEECHAM GROUP PLC	LT
Augmentin 875 mg /125 mg plėvele dengtos tabletės	DE/H/2868/02	LT/1/98/3675/009	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
Augmentin 875 mg /125 mg plėvele dengtos tabletės	DE/H/2868/02	LT/1/98/3675/010	BEECHAM GROUP PLC	LT
Augmentin 875 mg /125 mg plėvele dengtos tabletės	DE/H/2868/02	LT/1/98/3675/011	BEECHAM GROUP PLC	LT
Augmentin 875 mg /125 mg plėvele dengtos tabletės	DE/H/2868/02	LT/1/98/3675/012	BEECHAM GROUP PLC	LT
Augmentin ES 600 mg/42,9 mg/5 ml milteliai geriamajai suspensijai	PT/H/0684/001	LT/1/98/1191/001	GLAXOSMITHKLINE LIETUVA UAB	LT
Augmentin ES 600 mg/42,9 mg/5 ml milteliai geriamajai suspensijai	PT/H/0684/001	LT/1/98/1191/002	GLAXOSMITHKLINE LIETUVA UAB	LT
Augmentin ES 600 mg/42,9 mg/5 ml milteliai geriamajai suspensijai	PT/H/0684/001	LT/1/98/1191/003	GLAXOSMITHKLINE LIETUVA UAB	LT
Augmentin ES 600 mg/42,9 mg/5 ml milteliai geriamajai suspensijai	PT/H/0684/001	LT/1/98/1191/004	GLAXOSMITHKLINE LIETUVA UAB	LT
Augmentin SR 1000 mg/62,5 mg pailginto atpalaidavimo tabletės	BE/H/0168/001	LT/1/98/1191/005	GLAXOSMITHKLINE LIETUVA UAB	LT
Augmentin 200 mg/28,5 mg/5 ml milteliai geriamajai suspensijai	UK/H/4737/002	LT/1/98/3992/001	SMITHKLINE BEECHAM LTD	LT
Clavucid Solutab 875/125, comprimés dispersibles	not available	1651/99060007	ASTELLAS PHARMA B.V., OFFICE BE	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
Amoclane 875/125 mg poudre pour suspension buvable	not available	0019/09010037	EUROGENERICS N.V./S.A.	LU
Augmentin 250 mg/62,5 mg/5 ml, poudre pour suspension buvable	BE/H/0209/002	260/01/09/6525	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Augmentin P 500 mg/50 mg poeder voor oplossing voor injectie of infusie	DE/H/2809/005	260/01/09/6519	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Augmentin P 1000 mg/100 mg poeder voor oplossing voor injectie of infusie	DE/H/2809/006	260/01/09/6520	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Augmentin 2.000 mg/200 mg Pulver zur Herstellung einer Infusionslösung	DE/H/2809/004	260/01/09/6522	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Augmentin 875 mg/125 mg Filmtabletten	DE/H/2868/02	2001/09 6529	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Augmentin Retard 1.000 mg/62,5 mg Retardtabletten	BE/H/168/001	2003 02 0023	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Augmentin 500 mg/125 mg poeder voor orale suspensie in zakjes	BE/H/0159/001	260/01/09/6526	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Augmentin 1000 mg+200 mg, poudre pour solution pour injection ou perfusion	DE/H/2809/003	260/01 09 6521	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Augmentin 500 mg/125 mg, comprimés pelliculés	UK/H/4739/001	2001/09 6523	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
Augmentin 125 mg+31,25 mg+5 ml, poudre pour suspension buvable	BE/H/0209/001	260/01/09/6524	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Augmentin 400 mg/57 mg/5 ml pulveris iekšķigi lietojamas suspensijas pagatavošanai	UK/H/4737/001	99-0039	GLAXOSMITHKLINE LATVIA SIA	LV
Augmentin 1000 mg/200 mg pulveris injekciju vai infūziju šķīduma pagatavošanai	DE/H/2809/003	99-0033	GLAXOSMITHKLINE LATVIA SIA	LV
Augmentin 875 mg/125 mg apvalkotās tabletes	DE/H/2868/02	99-0035	GLAXOSMITHKLINE LATVIA SIA	LV
Аугментин ES 600 mg/42,9 mg/5 ml прах за перорална суспензия	not available	AA 1051/00101	GLAXOSMITHKLINE EOOD	MT
Augmentin 250 mg/125 mg film-coated tablets	UK/H/4735/001	MA 192/01501	GLAXOSMITHKLINE (IRELAND) LIMITED	MT
Augmentin 500 mg/100 mg powder for solution for injection or infusion	DE/H/2809/002	MA 447/00204	BEECHAM GROUP LTD	MT
Augmentin 1000 mg/200 mg powder for solution for injection or infusion	DE/H/2809/003	MA 447/00203	BEECHAM GROUP LTD	MT
Augmentin 500 mg/125 mg film-coated tablets	UK/H/4738/001	MA 192/01503	GLAXOSMITHKLINE (IRELAND) LIMITED	MT
Augmentin 875 mg/125 mg film-coated tablets	DE/H/2868/02	MA 192/01502	GLAXOSMITHKLINE (IRELAND) LIMITED	MT
Augmentin 100 mg/12,5 mg/ml, poeder voor orale suspensie	DE/H/2868/007-008/MR	RVG 14740	GLAXOSMITHKLINE B.V.	NL
Augmentin 500 mg/125 mg filmdrasjerte tabletter	UK/H/4738/01	14-10121	GLAXOSMITHKLINE 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
Augmentin 250 mg + 125 mg tabletki powlekane	UK/H/4735/01/MR	R/0641	GLAXOSMITHKLINE EXPORT LTD	PL
Augmentin, (400 mg + 57 mg)/5 ml, proszek do sporządzania zawiesiny doustnej	UK/H/4737/001	4193	GLAXOSMITHKLINE EXPORT LTD	PL
Augmentin, 1000 mg + 200 mg, proszek do sporzadzania roztworu do wstrzykiwan lub infuzji	DE/H/2809/003	R/0797	GLAXOSMITHKLINE EXPORT LTD	PL
Augmentin, 500 mg + 125 mg, tabletki powlekane	UK/H/4739/001	R/3682	GLAXOSMITHKLINE EXPORT LTD	PL
Augmentin, 2000 mg + 200 mg, proszek do sporzadzania roztworu do infuzji	DE/H/2809/004	9644	GLAXOSMITHKLINE EXPORT LTD	PL
Augmentin ES, (600 mg + 42,9 mg)/5 ml, proszek do sporzadzania zawiesiny doustnej	PT/H/0684/001	12306	GLAXOSMITHKLINE EXPORT LTD	PL
Augmentin SR, 1000 mg + 62,5 mg, tabletki o przedłużonym uwalnianiu	BE/H/0168/001	12737	GLAXOSMITHKLINE EXPORT LTD	PL
Augmentin 875 mg + 125 mg tabletki powlekane	DE/H/2868/02	R/7175	GLAXOSMITHKLINE EXPORT LTD	PL
Noprilam, 125 mg/31,25 mg/5 ml, pó para suspensão oral	not available	4726899	BIAL - PORTELA & Cª , SA	PT
Noprilam, 250 mg/62,5 mg/5 ml, pó para suspensão oral	not available	4727491	BIAL - PORTELA & Cª , SA	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
Noprilam, 500 mg/125 mg, comprimidos revestidos por película	not available	4717690	BIAL - PORTELA & Cª, SA	PT
Noprilam DT, 875 mg/125 mg, comprimidos revestidos por película	not available	4715595	BIAL - PORTELA & Cª, SA	PT
Clavamox 125, 125 mg/31,25 mg/5 ml, pó para suspensão oral	not available	9596239	BIAL - PORTELA & Cª, SA	PT
Clavamox 250, 250 mg/62,5 mg/5 ml, pó para suspensão oral	not available	9596247	BIAL - PORTELA & Cª, SA	PT
Clavamox 500, 500 mg/125 mg, comprimidos revestidos por película	not available	4717591	BIAL - PORTELA & Cª, SA	PT
Clavamox DT 400, 400 mg/57 mg/5 ml, pó para suspensão oral	not available	2922698	BIAL - PORTELA & Cª, SA	PT
Clavamox DT, 875 mg/125 mg, comprimidos revestidos por película	not available	4716296	BIAL - PORTELA & Cª, SA	PT
Clavamox ES 600 mg/42,9 mg/ 5 ml pó para suspensão oral	not available	5324082	BIAL - PORTELA & Cª, SA	PT
Clavamox ES 600 mg/42,9 mg/ 5 ml pó para suspensão oral	not available	5324181	BIAL - PORTELA & Cª, SA	PT
Clavamox ES 600 mg/42,9 mg/ 5 ml pó para suspensão oral	not available	5324280	BIAL - PORTELA & Cª, SA	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
Clavamox ES 600 mg/42,9 mg/ 5 ml pó para suspensão oral	not available	5324389	BIAL - PORTELA & Cª, SA	PT
Clavamox 125, 125 mg/31,25 mg/5 ml, pó para suspensão oral	not available	4727897	BIAL - PORTELA & Cª, SA	PT
Clavamox 125, 125 mg/31,25 mg/5 ml, pó para suspensão oral	not available	9596213	BIAL - PORTELA & Cª, SA	PT
Clavamox 125, 125 mg/31,25 mg/5 ml, pó para suspensão oral	not available	4823191	BIAL - PORTELA & Cª, SA	PT
Noprilam, 125 mg/31,25 mg/5 ml, pó para suspensão oral	not available	2273993	BIAL - PORTELA & Cª, SA	PT
Noprilam, 125 mg/31,25 mg/5 ml, pó para suspensão oral	not available	4726998	BIAL - PORTELA & Cª, SA	PT
Clavamox 250, 250 mg/62,5 mg/5 ml, pó para suspensão oral	not available	4727699	BIAL - PORTELA & Cª, SA	PT
Clavamox 250, 250 mg/62,5 mg/5 ml, pó para suspensão oral	not available	9596254	BIAL - PORTELA & Cª, SA	PT
Clavamox 250, 250 mg/62,5 mg/5 ml, pó para suspensão oral	not available	4727798	BIAL - PORTELA & Cª, SA	PT
Noprilam, 250 mg/62,5 mg/5 ml, pó para suspensão oral	not available	2273894	BIAL - PORTELA & Cª, SA	PT
Noprilam, 250 mg/62,5 mg/5 ml, pó para suspensão oral	not available	4727590	BIAL - PORTELA & Cª, SA	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
Clavamox 500, 500 mg/125 mg, comprimidos revestidos por película	not available	9588822	BIAL - PORTELA & Cª, SA	PT
Noprilam, 500 mg/125 mg, comprimidos revestidos por película	not available	2273399	BIAL - PORTELA & Cª, SA	PT
Noprilam, 500 mg/125 mg, comprimidos revestidos por película	not available	4715496	BIAL - PORTELA & Cª, SA	PT
Clavamox DT 400, 400 mg/57 mg/5 ml, pó para suspensão oral	not available	2922789	BIAL - PORTELA & Cª, SA	PT
Clavamox DT, 875 mg/125 mg, comprimidos revestidos por película	not available	4716395	BIAL - PORTELA & Cª, SA	PT
Clavamox DT, 875 mg/125 mg, comprimidos revestidos por película	not available	9766105	BIAL - PORTELA & Cª, SA	PT
Clavamox DT, 875 mg/125 mg, comprimidos revestidos por película	not available	4716494	BIAL - PORTELA & Cª, SA	PT
Noprilam DT, 875 mg/125 mg, comprimidos revestidos por película	not available	4715694	BIAL - PORTELA & Cª, SA	PT
Noprilam DT, 875 mg/125 mg, comprimidos revestidos por película	not available	2517696	BIAL - PORTELA & Cª, SA	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
Noprilam DT, 875 mg/125 mg, comprimidos revestidos por película	not available	4715793	BIAL - PORTELA & Cª, SA	PT
Forcid Solutab 875/125, 875 mg + 125 mg, comprimidos dispersíveis	HU/H/0386/001	4346581	ASTELLAS FARMA LDA.	PT
Forcid Solutab 875/125, 875 mg + 125 mg, comprimidos dispersíveis	HU/H/0386/001	4279782	ASTELLAS FARMA LDA.	PT
Forcid Solutab 875/125, 875 mg + 125 mg, comprimidos dispersíveis	HU/H/0386/001	4279683	ASTELLAS FARMA LDA.	PT
Forcid Solutab 875/125, 875 mg + 125 mg, comprimidos dispersíveis	HU/H/0386/001	4279881	ASTELLAS FARMA LDA.	PT
Forcid Solutab 875/125, 875 mg + 125 mg, comprimidos dispersíveis	HU/H/0386/001	4280087	ASTELLAS FARMA LDA.	PT
Forcid Solutab 875/125, 875 mg + 125 mg, comprimidos dispersíveis	HU/H/0386/001	4279980	ASTELLAS FARMA LDA.	PT
Augmentin Duo 400 mg/57 mg/5 ml, pó para suspensão oral (sabor a morango)	UK/H/4737/01/MR	2922482	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Augmentin 125 mg/31,25 mg/5 ml, pó para suspensão oral	DE/H/2868/004	8605014	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Augmentin 500 mg/125 mg, comprimidos revestidos por película	UK/H/4738/01/MR	4717286	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Augmentin 500 mg/125 mg, comprimidos revestidos por película	UK/H/4738/01/MR	8604702	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, 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
Augmentin 500 mg/125 mg, comprimidos revestidos por película	UK/H/4738/01/MR	4714689	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Augmentin Forte 250 mg/62,5 mg/5 ml, pó para suspensão oral	DE/H/2868/05/MR	8605006	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Augmentin ES 600 mg/42,9 mg/ 5 ml pó para suspensão oral	PT/H/0684/001	5323688	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Augmentin ES 600 mg/42,9 mg/ 5 ml pó para suspensão oral	PT/H/0684/001	5323787	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Augmentin ES 600 mg/42,9 mg/ 5 ml pó para suspensão oral	PT/H/0684/001	5323886	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LIMITADA	PT
Augmentin ES 600 mg/42,9 mg/ 5 ml pó para suspensão oral	PT/H/0684/001	5323985	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Augmentin Duo 875 mg/125 mg, comprimidos revestidos por película	DE/H/2868/02	5751888	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Augmentin Duo 875 mg/125 mg, comprimidos revestidos por película	DE/H/2868/02	4714986	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Augmentin SR 1000 mg/62,5 mg comprimate cu eliberare prelungită	BE/H/0168/001	7554/2015/01	BEECHAM GROUP PLC	RO
Augmentin SR 1000 mg/62,5 mg comprimate cu eliberare prelungită	BE/H/0168/001	7554/2015/02	BEECHAM GROUP PLC	RO
Augmentin SR 1000 mg/62,5 mg comprimate cu eliberare prelungită	BE/H/0168/001	7554/2015/03	BEECHAM GROUP PLC	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
Augmentin SR 1000 mg/62,5 mg comprimate cu eliberare prelungita	BE/H/0168/001	7554/2015/05	BEECHAM GROUP PLC	RO
Augmentin 875 mg/125 mg comprimate filmate	DE/H/2868/02	8164/2015/01	BEECHAM GROUP PLC	RO
Augmentin 875 mg/125 mg comprimate filmate	DE/H/2868/02	8164/2015/02	BEECHAM GROUP PLC	RO
Augmentin 875 mg/125 mg comprimate filmate	DE/H/2868/02	8164/2015/03	BEECHAM GROUP PLC	RO
Augmentin 875 mg/125 mg comprimate filmate	DE/H/2868/02	8164/2015/04	BEECHAM GROUP PLC	RO
Augmentin 875 mg/125 mg comprimate filmate	DE/H/2868/02	8164/2015/05	BEECHAM GROUP PLC	RO
Augmentin 875 mg/125 mg comprimate filmate	DE/H/2868/02	8164/2015/06	BEECHAM GROUP PLC	RO
Augmentin 875 mg/125 mg comprimate filmate	DE/H/2868/02	8164/2015/07	BEECHAM GROUP PLC	RO
Augmentin 875 mg/125 mg comprimate filmate	DE/H/2868/02	8164/2015/10	BEECHAM GROUP PLC	RO
Augmentin 875 mg/125 mg comprimate filmate	DE/H/2868/02	8164/2015/11	BEECHAM GROUP PLC	RO
Augmentin 875 mg/125 mg comprimate filmate	DE/H/2868/02	8164/2015/12	BEECHAM GROUP PLC	RO
Augmentin 500 mg/125 mg comprimate filmate	UK/H/4738/01/MR	8395/2015/01	BEECHAM GROUP PLC	RO
Augmentin 500 mg/125 mg comprimate filmate	UK/H/4738/01/MR	8395/2015/02	BEECHAM GROUP PLC	RO
Augmentin 500 mg/125 mg comprimate filmate	UK/H/4738/01/MR	8395/2015/03	BEECHAM GROUP PLC	RO
Augmentin 500 mg/125 mg comprimate filmate	UK/H/4738/01/MR	8395/2015/04	BEECHAM GROUP PLC	RO
Augmentin 500 mg/125 mg comprimate filmate	UK/H/4738/01/MR	8395/2015/05	BEECHAM GROUP PLC	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
Augmentin 500 mg/125 mg comprimate filmate	UK/H/4738/01/MR	8395/2015/06	BEECHAM GROUP PLC	RO
Augmentin 500 mg/125 mg comprimate filmate	UK/H/4738/01/MR	8395/2015/07	BEECHAM GROUP PLC	RO
Augmentin 500 mg/125 mg comprimate filmate	UK/H/4738/01/MR	8395/2015/08	BEECHAM GROUP PLC	RO
Augmentin 500 mg/125 mg comprimate filmate	UK/H/4738/01/MR	8395/2015/09	BEECHAM GROUP PLC	RO
Augmentin 500 mg/125 mg comprimate filmate	UK/H/4738/01/MR	8395/2015/10	BEECHAM GROUP PLC	RO
Augmentin 500 mg/125 mg comprimate filmate	UK/H/4738/01/MR	8395/2015/11	BEECHAM GROUP PLC	RO
Augmentin 500 mg/125 mg comprimate filmate	UK/H/4738/01/MR	8395/2015/12	BEECHAM GROUP PLC	RO
Augmentin 500 mg/125 mg comprimate filmate	UK/H/4738/01/MR	8395/2015/13	BEECHAM GROUP PLC	RO
Augmentin 875 mg/125 mg comprimate filmate	DE/H/2868/02	8164/2015/08	BEECHAM GROUP PLC	RO
Augmentin intravenos 2000 mg/200 mg pulbere pentru soluţie perfuzabilă	DE/H/2809/004/MR	8740/2016/01	BEECHAM GROUP PLC	RO
AUGMENTIN ES 600 mg/42,9 mg/ 5 ml pulbere pentru suspensie orală	PT/H/0684/001	7850/2015/01	SMITHKLINE BEECHAM LTD	RO
AUGMENTIN ES 600 mg/42,9 mg/ 5 ml pulbere pentru suspensie orală	PT/H/0684/001	7850/2015/02	SMITHKLINE BEECHAM LTD	RO
AUGMENTIN ES 600 mg/42,9 mg/ 5 ml pulbere pentru suspensie orală	PT/H/0684/001	7850/2015/04	SMITHKLINE BEECHAM LTD	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
AUGMENTIN ES 600 mg/42,9 mg/ 5 ml pulbere pentru suspensie orală	PT/H/0684/001	7850/2015/03	SMITHKLINE BEECHAM LTD	RO
Augmentin BIS 400 mg/57 mg/5 ml pulbere pentru suspensie orală	UK/H/4737/001	9385/2016/02	SMITHKLINE BEECHAM LTD	RO
Augmentin BIS 400 mg/57 mg/5 ml pulbere pentru suspensie orală	UK/H/4737/001	9385/2016/03	SMITHKLINE BEECHAM LTD	RO
Augmentin BIS 400 mg/57 mg/5 ml pulbere pentru suspensie orală	UK/H/4737/001	9385/2016/01	SMITHKLINE BEECHAM LTD	RO
Augmentin BIS 400 mg/57 mg/5 ml pulbere pentru suspensie orală	UK/H/4737/001	9385/2016/04	SMITHKLINE BEECHAM LTD	RO
Augmentin BIS 400 mg/57 mg/5 ml pulbere pentru suspensie orală	UK/H/4737/001	9385/2016/05	SMITHKLINE BEECHAM LTD	RO
Augmentin BIS 400 mg/57 mg/5 ml pulbere pentru suspensie orală	UK/H/4737/001	9385/2016/06	SMITHKLINE BEECHAM LTD	RO
Spektramox 80 mg/ml+11,97 mg/ml pulver till oral suspension	not available	15980	MEDA AB	SE
Spektramox 50 mg/ml+12,5 mg/ml pulver till oral suspension	not available	10700	MEDA AB	SE
Spektramox 250 mg/125 mg filmdragerade tabletter	not available	12414	MEDA 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
Spektramox 500 mg/125 mg filmdragerade tabletter	not available	12415	MEDA AB	SE
Spektramox 875 mg/125 mg filmdragerade tabletter	not available	15979	MEDA AB	SE
Augmentin SR 1000 mg/62,5 mg tablete s podaljšanim sproščanjem	BE/H/0168/001	H/05/01951/001	GLAXOSMITHKLINE D.O.O.	SI
Augmentin SR 1000 mg/62,5 mg tablete s podaljšanim sproščanjem	BE/H/0168/001	H/05/01951/002	GLAXOSMITHKLINE D.O.O.	SI
Augmentin 400 mg/57 mg v 5 ml prašek za peroralno suspenzijo z okusom jagode	UK/H/4737/01/MR	H/99/00969/001	GLAXOSMITHKLINE D.O.O.	SI
Augmentin 400 mg/57 mg v 5 ml prašek za peroralno suspenzijo z okusom jagode	UK/H/4737/01/MR	H/99/00969/002	GLAXOSMITHKLINE D.O.O.	SI
Augmentin 400 mg/57 mg v 5 ml prašek za peroralno suspenzijo z okusom jagode	UK/H/4737/01/MR	H/99/00969/003	GLAXOSMITHKLINE D.O.O.	SI
Augmentin 400 mg/57 mg v 5 ml prašek za peroralno suspenzijo z okusom jagode	UK/H/4737/01/MR	H/99/00969/004	GLAXOSMITHKLINE D.O.O.	SI
Augmentin 400 mg/57 mg v 5 ml prašek za peroralno suspenzijo z okusom jagode	UK/H/4737/01/MR	H/99/00969/005	GLAXOSMITHKLINE D.O.O.	SI

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the Member State	Member State where product is authorised
Augmentin 400 mg/57 mg v 5 ml prašek za peroralno suspenzijo z okusom jagode	UK/H/4737/01/MR	H/99/00969/006	GLAXOSMITHKLINE D.O.O.	SI
Augmentin 400 mg/57 mg v 5 ml prašek za peroralno suspenzijo z okusom jagode	UK/H/4737/01/MR	H/99/00969/007	GLAXOSMITHKLINE D.O.O.	SI
Augmentin 400 mg/57 mg v 5 ml prašek za peroralno suspenzijo z okusom jagode	UK/H/4737/01/MR	H/99/00969/008	GLAXOSMITHKLINE D.O.O.	SI
Augmentin 400 mg/57 mg v 5 ml prašek za peroralno suspenzijo z okusom jagode	UK/H/4737/01/MR	H/99/00969/009	GLAXOSMITHKLINE D.O.O.	SI
Augmentin SR 1000 mg/62,5 mg tablete s podaljšanim sproščanjem	BE/H/0168/001	H/05/01951/003	GLAXOSMITHKLINE D.O.O.	SI
Augmentin SR 1000 mg/62,5 mg tablete s podaljšanim sproščanjem	BE/H/0168/001	H/05/01951/004	GLAXOSMITHKLINE D.O.O.	SI
Augmentin SR 1000 mg/62,5 mg tablete s podaljšanim sproščanjem	BE/H/0168/001	H/05/01951/005	GLAXOSMITHKLINE D.O.O.	SI
Augmentin SR 1000 mg/62,5 mg tablete s podaljšanim sproščanjem	BE/H/0168/001	H/05/01951/006	GLAXOSMITHKLINE D.O.O.	SI
Augmentin SR 1000 mg/62,5 mg tablete s podaljšanim sproščanjem	BE/H/0168/001	H/05/01951/007	GLAXOSMITHKLINE D.O.O.	SI
Augmentin SR 1000 mg/62,5 mg tablete s podaljšanim sproščanjem	BE/H/0168/001	H/05/01951/008	GLAXOSMITHKLINE D.O.O.	SI

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Augmentin SR 1000 mg/62,5 mg tablete s podaljšanim sproščanjem	BE/H/0168/001	H/05/01951/009	GLAXOSMITHKLINE D.O.O.	SI
MEGAMOX 625 Forte Filmom obalené tablety	not available	15/0109/99-S	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	SK
MEGAMOX 375 Filmom obalené tablety	not available	15/0108/99-S	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	SK
Amoksiklav 600 mg, prášok na injekčný/infúzny roztok	not available	15/0605/94-S	SANDOZ PHARMACEUTICALS D.D.	SK
Amoksiklav 1,2 g, prášok na injekčný/infúzny roztok	not available	15/0605/94-S	SANDOZ PHARMACEUTICALS D.D.	SK
Amoksiklav 1000 mg granulát na perorálnu suspenziu	IT/H/0245/001	15/0771/10-S	SANDOZ PHARMACEUTICALS D.D.	SK
Amoksiklav 600 mg, prášok na injekčný/infúzny roztok	not available	15/0605/94-S	SANDOZ PHARMACEUTICALS D.D.	SK
Amoksiklav 1,2 g, prášok na injekčný/infúzny roztok	not available	15/0605/94-S	SANDOZ PHARMACEUTICALS D.D.	SK
Amoksiklav 1000 mg granulát na perorálnu suspenziu	IT/H/0245/001	15/0771/10-S	SANDOZ PHARMACEUTICALS D.D.	SK
Augmentin 375 mg 250 mg/125 mg filmom obalené tablety	UK/H/4735/01/MR	15/0141/84-C/S	GLAXOSMITHKLINE SLOVAKIA S.R.O.	SK
Augmentin DUO 400 mg/57 mg/5 ml prášok na perorálnu suspenziu (jahodová príchut)	UK/H/4737/01/MR	15/0005/00-S	GLAXOSMITHKLINE SLOVAKIA S.R.O.	SK
Augmentin ES 600 mg/42,9 mg/5 ml prášok na perorálnu suspenziu	PT/H/684/001	15/0085/07-S	GLAXOSMITHKLINE SLOVAKIA S.R.O.	SK

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the Member State	Member State where product is authorised
Augmentin SR 1 000 mg/62,5 mg tablety s predĺženým uvoľňovaním	BE/H/168/001	15/0122/02-S	GLAXOSMITHKLINE SLOVAKIA S.R.O.	SK
Augmentin 1 g 875 mg/125 mg filmom obalené tablety	DE/H/2868/02	15/0682/96-S	GLAXOSMITHKLINE SLOVAKIA S.R.O.	SK
Co-amoxiclav 500mg/125mg Film-Coated Tablets	not available	PL 44041/0006	NOUMED LIFE SCIENCES	UK
Co-amoxiclav 500/125mg Tablets	not available	PL 04416/0556	SANDOZ LTD	UK
Co-amoxiclav 250/62.5 mg/5 ml Powder for Oral Suspension	not available	PL 04416/0515	SANDOZ LTD	UK
Co-amoxiclav 125/31.25 mg/5 ml Powder for Oral Suspension	not available	PL 04416/0514	SANDOZ LTD	UK
Co-Amoxiclav 250/125mg Tablets BP	not available	PL 04520/0054	SANDOZ GMBH	UK
Co-amoxiclav 500/125mg Tablets	not available	PL 04416/0556	SANDOZ LTD	UK
Co-amoxiclav 250/62.5 mg/5 ml Powder for Oral Suspension	not available	PL 04416/0515	SANDOZ LTD	UK
Co-amoxiclav 125/31.25 mg/5 ml Powder for Oral Suspension	not available	PL 04416/0514	SANDOZ LTD	UK
Co-Amoxiclav 250/125mg Tablets BP	not available	PL 04520/0054	SANDOZ GMBH	UK
Augmentin 375 mg Tablets	UK/H/4735/001	PL 00038/0270	BEECHAM GROUP PLC	UK
Augmentin DUO 400/57	UK/H/4737/001	PL 10592/0070	SMITHKLINE BEECHAM LTD	UK
Augmentin®-Duo 200/28	UK/H/4737/002	PL 10592/0072	SMITHKLINE BEECHAM LTD	UK
Augmentin Intravenous	DE/H/2809/002	PL 00038/0320	BEECHAM GROUP LTD	UK

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Augmentin 125/31 SF Suspension	DE/H/2868/004	PL 00038/0298	BEECHAM GROUP PLC	UK
Augmentin 1 g Tablets	DE/H/2868/02	PL 00038/0368	BEECHAM GROUP LTD	UK