10 November 2022
EMA/759338/2022
Human Medicines Division

List of nationally authorised medicinal products

Active substance(s): amoxicillin/clavulanate

Procedure No. PSUSA/00000188/202203

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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
Clavamox Duo - Trockensaft	not available	1-22155	SANDOZ GMBH	AT
Augmentin 400 mg/57 mg/5 ml Pulver zur Herstellung einer Suspension zum Einnehmen Multifruchtgeschmack	DE/H/2868/006	138128	GLAXOSMITHKLINE PHARMA GMBH.	AT
Curam intravenös 500 mg/100 mg - Pulver zur Herstellung einer Injektions-/Infusionslösung	NL/H/0541/003	1-26284	SANDOZ GMBH	AT
Curam intravenös 1000 mg/200 mg - Pulver zur Herstellung einer Injektions-/Infusionslösung	NL/H/0541/001	1-26281	SANDOZ GMBH	AT
Augmentin 875 mg/125 mg Filmtabletten	DE/H/2868/002	1-21396	GLAXOSMITHKLINE PHARMA GMBH.	AT
Augmentin 500 mg/125 mg Filmtabletten	DE/H/6032/001	1-17839	GLAXOSMITHKLINE PHARMA GMBH.	AT
Curam intravenös 500 mg/50 mg - Pulver zur Herstellung einer Injektions-/Infusionslösung	NL/H/0541/004	1-26283	SANDOZ GMBH	AT
Curam intravenös 2000 mg/200 mg - Pulver zur Herstellung einer Infusionslösung	NL/H/0541/002	1-26282	SANDOZ GMBH	AT
Augmentin P 500 mg/50 mg poeder voor oplossing voor injectie of infusie	DE/H/2809/005	BE135195	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Augmentin P 500 mg/50 mg Pulver zur Herstellung einer Injektions-/Infusionslösung	DE/H/2809/005	BE135195	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Augmentin 1.000 mg/200 mg Pulver zur Herstellung	DE/H/2809/003	BE135213	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
einer Injektions- oder Infusionslösung				
Augmentin 2.000 mg/200 mg Pulver zur Herstellung einer Infusionslösung	DE/H/2809/004	BE135511	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Augmentin 1000 mg/200 mg poeder voor oplossing voor injectie of infusie	DE/H/2809/003	BE135213	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Augmentin 2000 mg/200 mg poeder voor oplossing voor infusie	DE/H/2809/004	BE135511	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Augmentin 1000 mg/200 mg, poudre pour solution injectable/pour perfusion	DE/H/2809/003	BE135213	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 injectable/pour perfusion	DE/H/2809/005	BE135195	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Augmentin P 1.000 mg/100 mg Pulver zur Herstellung einer Injektions-/Infusionslösung	DE/H/2809/006	BE135204	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 P 1000 mg/100 mg, poudre pour solution injectable/pour perfusion	DE/H/2809/006	BE135204	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Augmentin 500 mg/125 mg Pulver zur Herstellung einer Suspension zum Einnehmen in Beuteln	BE/H/0159/01	BE159345	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Augmentin 500 mg/125 mg poeder voor orale suspensie in zakjes	BE/H/0159/01	BE159345	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Augmentin 500 mg/125	BE/H/0159/001	BE159345	GLAXOSMITHKLINE	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
mg, poudre pour suspension buvable en sachet			PHARMACEUTICALS SA	
Amoxiclav Sandoz 1000 mg/200 mg poeder voor oplossing voor injectie / infusie	NL/H/0541/001	BE508720	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 875 mg/125 mg Filmtabletten	DE/H/2868/02	BE200803	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Augmentin 875 mg/125 mg Filmtabletten	DE/H/2868/02	BE439293	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Augmentin 875/125 mg filmomhulde tabletten	DE/H/2868/02	BE200803	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Augmentin 875/125 mg filmomhulde tabletten	DE/H/2868/02	BE439293	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Augmentin 875 mg/125 mg, comprimés pelliculés	DE/H/2868/02	BE200803	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Augmentin 500 mg/125 mg, comprimés pelliculés	DE/H/6032/001	BE439284	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Augmentin 500 mg/125 mg Filmtabletten	DE/H/6032/001	BE125334	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Augmentin 500 mg/125 mg Filmtabletten	DE/H/6032/001	BE439284	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Augmentin 500 mg/125 mg filmomhulde tabletten	DE/H/6032/001	BE125334	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Augmentin 500 mg/125 mg filmomhulde tabletten	DE/H/6032/001	BE439284	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Augmentin 500 mg/125 mg, comprimés pelliculés	DE/H/6032/001	BE125334	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Amoxiclav Sandoz 2000 mg/200 mg poeder voor oplossing voor infusie	NL/H/0541/002	BE508737	SANDOZ N.V.	BE

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Amoxiclav Sandoz 2000 mg/200 mg poeder voor oplossing voor infusie	NL/H/0541/002	BE271214	SANDOZ N.V.	BE
Augmentin 250 mg/62,5 mg/5 ml, poudre pour suspension buvable	BE/H/0209/02	BE125413	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Augmentin 125 mg/31,25 mg/5 ml Pulver zur Herstellung einer Suspension zum Einnehmen	BE/H/0209/001	BE125404	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Augmentin 125 mg/31,25 mg/5 ml poeder voor orale suspensie	BE/H/0209/001	BE125404	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Augmentin 250 mg/62,5 mg/5 ml, poudre pour suspension buvable	BE/H/0209/002	BE125413	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Augmentin 250 mg/62,5 mg/5 ml poeder voor orale suspensie	BE/H/0209/02	BE125413	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Augmentin 125 mg/31,25 mg/5 ml, poudre pour suspension buvable	BE/H/0209/01	BE125404	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Amoclane 875/125 mg poeder voor orale suspensie	not available	BE273856	EUROGENERICS N.V./S.A.	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
Аугментин ES 600 mg/42,9 mg/5 ml прах за перорална суспензия	PT/H/0684/001	20050447	GLAXOSMITHKLINE TRADING SERVICES LIMITED	BG
Аугментин 125 mg/31,25 mg/5 ml прах за	DE/H/2868/04	20000317	GLAXOSMITHKLINE TRADING SERVICES LIMITED	BG

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перорална суспензия				
Аугментин 250 mg/62,5 mg/5 ml прах за перорална суспензия	DE/H/2868/005	20000318	GLAXOSMITHKLINE TRADING SERVICES LIMITED	BG
Аугментин 400 mg/57 mg/5 ml прах за перорална суспензия	DE/H/2868/06	9900089	GLAXOSMITHKLINE TRADING SERVICES LIMITED	BG
Аугментин 875 mg/125 mg филмирани таблетки	DE/H/2868/02	20000316	GLAXOSMITHKLINE TRADING SERVICES LIMITED	BG
Аугментин 500 mg/125 mg филмирани таблетки	DE/H/6032/001	20000314	GLAXOSMITHKLINE TRADING SERVICES LIMITED	BG
Augmentin ES 600 mg/42.9 mg/5 ml κόνις για πόσιμο εναιώρημα	PT/H/0684/001	20148	GLAXOSMITHKLINE (IRELAND) LIMITED	CY
Augmentin Mixed Fruit 400 mg/57 mg/5 ml κόνις για πόσιμο εναιώρημα (βελτιωτικό γεύσης ανάμικτα φρούτα)	DE/H/2868/006	022824	GLAXOSMITHKLINE (IRELAND) LIMITED	CY
Augmentin 875 mg/125 mg ep??a??μμ??a με ?ept? ?μ???? d?s??a	DE/H/2868/002	19515	GLAXOSMITHKLINE (IRELAND) LIMITED	CY
Augmentin 400 mg/57 mg/5 ml κόνις για πόσιμο εναιώρημα (γεύση φράουλα)	PT/H/2363/001	19702	GLAXOSMITHKLINE (IRELAND) LIMITED	CY
Augmentin 500 mg/125 mg επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/6032/001	12656	GLAXOSMITHKLINE (IRELAND) LIMITED	CY
Augmentin 400 mg/5 ml + 57 mg/5 ml prášek pro perorální suspenzi	DE/H/2868/06	15/685/16-C	GLAXOSMITHKLINE (IRELAND) LIMITED	CZ
Augmentin 1 g potahované tablety	DE/H/2868/002	15/644/96-C	GLAXOSMITHKLINE (IRELAND) LIMITED	CZ
Augmentin 625 mg potahované tablety	DE/H/6032/001	15/141/84-B/C	GLAXOSMITHKLINE (IRELAND) LIMITED	CZ
Augmentan 1000 mg/100 mg Pulver zur Herstellung	DE/H/2809/006	86022.00.00	GLAXOSMITHKLINE GMBH & CO. KG	DE

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einer Injektions-/Infusionslösung				
Augmentan 100 mg/12,5 mg/ml Pulver zur Herstellung einer Suspension zum Einnehmen für Kinder	DE/H/2868/007	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/2868/008	86131.00.00	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/2868/04	1927.00.03	GLAXOSMITHKLINE GMBH & CO. KG	DE
Augmentan forte Trockensaft 50 mg/12,5 mg pro ml 250 mg/62,5 mg/5 ml Pulver zur Herstellung einer Suspension zum Einnehmen	DE/H/2868/005	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/2868/006	35944.00.00	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 Filmtabletten	DE/H/2868/002	34923.00.00	GLAXOSMITHKLINE 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
875/125 mg 875 mg/125 mg Filmtabletten			CO. KG	
Augmentan 500 mg/125 mg Filmtabletten	DE/H/6032/001	92660.00.00	GLAXOSMITHKLINE GMBH & CO. KG	DE
Augmentan 40 mg/ml + 5,7 mg/ml Pulver zur Herstellung einer Suspension zum Einnehmen	DE/H/6223/001	2202820.00.00	GLAXOSMITHKLINE GMBH & CO. KG	DE
Spektramox	not available	12255	VIATRIS APS	DK
Spektramox	not available	12257	VIATRIS APS	DK
Augmentin Fruit, 400 mg/57 mg/5 ml suukaudse suspensiooni pulber	DE/H/2868/006	947217	GLAXOSMITHKLINE TRADING SERVICES LIMITED	EE
Augmentin, 875 mg/125 mg õhukese polümeerikattega tabletid	DE/H/2868/002	213398	GLAXOSMITHKLINE (IRELAND) LIMITED	EE
Augmentin, 400 mg/57 mg/5 ml suukaudse suspensiooni pulber	PT/H/2363/001	213498	GLAXOSMITHKLINE (IRELAND) LIMITED	EE
Augmentin, 500 mg/125 mg õhukese polümeerikattega tabletid	DE/H/6032/001	213298	GLAXOSMITHKLINE (IRELAND) LIMITED	EE
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 100 mg/ml +12,5 mg/ml polvo para suspensión oral	DE/H/2868/008	59.051	GLAXOSMITHKLINE, S.A.	ES
Augmentine 875 mg/125 mg comprimidos recubiertos con película	DE/H/2868/002	59.515	GLAXOSMITHKLINE S.A.	ES
Augmentine 500 mg/125 mg comprimidos	DE/H/6032/001	81.482	GLAXOSMITHKLINE S.A.	ES

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recubiertos con película				
AUGMENTIN 500 mg/62,5 mg, comprimé pelliculé (rapport amoxicilline/acide clavulanique : 8/1)	not available	NL25012	LABORATOIRE GLAXOSMITHKLINE	FR
AUGMENTIN 1 g/200 mg, poudre pour solution injectable / pour perfusion (IV)	DE/H/2809/003	NL13670-1	LABORATOIRE GLAXOSMITHKLINE	FR
AUGMENTIN 1 g/200 mg, poudre et solvant pour solution injectable / pour perfusion (IV)	DE/H/2809/003	NL13670-2	LABORATOIRE GLAXOSMITHKLINE	FR
AUGMENTIN 2 g/200 mg ADULTES, poudre pour solution pour perfusion	DE/H/2809/004	NL13672	LABORATOIRE GLAXOSMITHKLINE	FR
AUGMENTIN 1 g/125 mg, poudre pour suspension buvable en sachet-dose (rapport amoxicilline/acide clavulanique : 8/1)	not available	NL22398	LABORATOIRE GLAXOSMITHKLINE	FR
AUGMENTIN 100 mg/12,50 mg par ml NOURRISSONS, poudre pour suspension buvable en flacon (rapport amoxicilline/acide clavulanique : 8/1) Amoxicilline/acide clavulanique	DE/H/2868/008	NL16485	GLAXOSMITHKLINE TRADING SERVICES LIMITED	FR
AUGMENTIN 100 mg/12,50 mg par ml ENFANTS, poudre pour suspension buvable en flacon (rapport amoxicilline/acide clavulanique : 8/1) Amoxicilline/acide clavulanique	DE/H/2868/007	NL21290	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 80 mg/ 11,4 mg/ mL, poudre pour suspension buvable (rapport amoxicilline/ acide clavulanique : 7/1)	DE/H/2868/06	NL47194	LABORATOIRE GLAXOSMITHKLINE	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	NL25190	LABORATOIRE GLAXOSMITHKLINE	FR
AMOXICILLINE/ACIDE CLAVULANIQUE EG 100 mg/12,5 mg par ml NOURRISSONS, poudre pour suspension buvable en flacon (rapport amoxicilline/acide clavulanique: 8/1)	not available	NL29905	EG LABO LABORATOIRES EUROGENERICS - DO NOT USE	FR
AMOXICILLINE/ACIDE CLAVULANIQUE EG LABO - LABORATOIRES EUROGENERICS 100 mg/12,5 mg par ml ENFANTS, poudre pour suspension buvable en flacon (rapport amoxicilline/acide clavulanique: 8/1)	not available	NL29910	EG LABO LABORATOIRES EUROGENERICS	FR
AMOXICILLINE / ACIDE CLAVULANIQUE BIOGARAN 500 mg/62,5 mg, comprimé pelliculé (rapport amoxicilline/acide clavulanique : 8/1)	not available	NL25189	LABORATOIRE GLAXOSMITHKLINE	FR
AUGMENTIN 250 mg/31,25 mg NOURRISSONS, poudre	not available	NL52586	LABORATOIRE GLAXOSMITHKLINE	FR

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pour suspension buvable en sachet (rapport amoxicilline/acide clavulanique : 8/1)				
AUGMENTIN 500 mg/62,5 mg ENFANTS, poudre pour suspension buvable en sachet (rapport amoxicilline/acide clavulanique : 8/1)	not available	NL52587	LABORATOIRE GLAXOSMITHKLINE	FR
AMOXICILLINE/ACIDE CLAVULANIQUE ZENTIVA LAB 500 mg/62,50 mg ADULTES, comprimé pelliculé (rapport amoxicilline/acide clavulanique : 8/1)	not available	NL25155	LABORATOIRE GLAXOSMITHKLINE	FR
AMOXICILLINE/ACIDE CLAVULANIQUE GLAXOSMITHKLINE 1 g/125 mg ADULTES, poudre pour suspension buvable en sachet-dose (Rapport amoxicilline/acide clavulanique : 8/1)	not available	NL37616	LABORATOIRE GLAXOSMITHKLINE	FR
AMOXICILLINE/ACIDE CLAVULANIQUE BIOGARAN 100 mg/12,5 mg par ml ENFANTS, poudre pour suspension buvable (Rapport Amoxicilline/Acide clavulanique: 8/1)	not available	NL25244	LABORATOIRE GLAXOSMITHKLINE	FR
AMOXICILLINE/ACIDE CLAVULANIQUE BIOGARAN 100 mg/12,5 mg par ml NOURRISSONS, poudre pour suspension buvable (Rapport Amoxicilline/Acide	not available	NL25243	LABORATOIRE GLAXOSMITHKLINE	FR

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clavulanique: 8/1)				
AMOXICILLINE/ACIDE CLAVULANIQUE GLAXOSMITHKLINE 100 mg/12,50 mg par ml NOURRISSONS, poudre pour suspension buvable en flacon (Rapport amoxicilline/acide clavulanique : 8/1)	not available	NL37619	LABORATOIRE GLAXOSMITHKLINE	FR
AMOXICILLINE/ACIDE CLAVULANIQUE GLAXOSMITHKLINE 100 mg/12,50 mg par ml ENFANTS, poudre pour suspension buvable en flacon (Rapport amoxicilline/acide clavulanique : 8/1)	not available	NL37617	LABORATOIRE GLAXOSMITHKLINE	FR
AMOXICILLINE/ACIDE CLAVULANIQUE GLAXOSMITHKLINE 500 mg/62,5 mg ADULTES, comprimé pelliculé (rapport amoxicilline/acide clavulanique : 8/1)	not available	NL37615	LABORATOIRE GLAXOSMITHKLINE	FR
AUGMENTIN 40 mg/5,7 mg/mL poudre pour suspension buvable (rapport amoxicilline/acide clavulanique 7/1)	DE/H/6223/001	NL49594	LABORATOIRE GLAXOSMITHKLINE	FR
AMOXICILLINE/ACIDE CLAVULANIQUE ZENTIVA LAB 1 g/125 mg ADULTES, poudre pour suspension buvable en sachet-dose (rapport amoxicilline/acide clavulanique: 8/1)	not available	NL25156	LABORATOIRE GLAXOSMITHKLINE	FR

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AMOXICILLINE/ACIDE CLAVULANIQUE ZENTIVA LAB 100 mg/12,50 mg par ml NOURRISSONS, poudre pour suspension buvable en flacon (rapport amoxicilline/acide clavulanique : 8/1)	not available	NL16931	LABORATOIRE GLAXOSMITHKLINE	FR
AMOXICILLINE/ACIDE CLAVULANIQUE ZENTIVA LAB 100 mg/12,50 mg par ml ENFANTS, poudre pour suspension buvable en flacon (rapport amoxicilline/acide clavulanique : 8/1)	not available	NL25205	LABORATOIRE GLAXOSMITHKLINE	FR
Augmentin 400 mg/57 mg/5 ml ????? ??a p?s?µ? e?a????µa (?e?s? f???t??)	DE/H/2868/006	1759915	GLAXOSMITHKLINE SINGLE MEMBER A.E.B.E.	GR
Augmentin 875 mg/125 mg ep??a??µµ??a µe ?ept? ?µ???? d?s??a	DE/H/2868/002	1759911	GLAXOSMITHKLINE SINGLE MEMBER A.E.B.E.	GR
Augmentin 400 mg/57 mg/5 ml κόνις για πόσιμο εναιώρημα (γεύση φράουλα)	PT/H/2363/001	1759912	GLAXOSMITHKLINE SINGLE MEMBER A.E.B.E.	GR
Augmentin 500 mg/125 mg επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/6032/001	1759904	GLAXOSMITHKLINE SINGLE MEMBER A.E.B.E.	GR
Augmentin 250 mg/62.5 mg/5 ml κόνις για πόσιμο εναιώρημα	BE/H/0209/002	1759909	GLAXOSMITHKLINE SINGLE MEMBER A.E.B.E.	GR
Augmentin 875 mg + 125 mg filmom obložene tablete	not available	HR-H-170450439	GLAXOSMITHKLINE TRADING SERVICES LIMITED	HR
Augmentin 400 mg + 57 mg/5 ml prašak za oralnu suspenziju s okusom voća	DE/H/2868/006	HR-H-377920132	GLAXOSMITHKLINE TRADING SERVICES LIMITED	HR

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Augmentin 250 mg/62,5 mg/5 ml por belsoleges szuszpenzióhoz	DE/H/2868/005	OGYI-T-1352/02	GLAXOSMITHKLINE TRADING SERVICES LIMITED	HU
Augmentin DUO 400 mg/57 mg/5 ml vegyes gyümölcsízu por belsoleges szuszpenzióhoz	DE/H/2868/006	OGYI-T-1352/14	GLAXOSMITHKLINE TRADING SERVICES LIMITED	HU
Augmentin DUO 400 mg/57 mg/5 ml vegyes gyümölcsízu por belsoleges szuszpenzióhoz	DE/H/2868/006	OGYI-T-1352/15	GLAXOSMITHKLINE TRADING SERVICES LIMITED	HU
Augmentin DUO 875 mg/125 mg filmtabletta	DE/H/2868/02	OGYI-T-1352/08	GLAXOSMITHKLINE TRADING SERVICES LIMITED	HU
Augmentin 250 mg/125 mg filmtabletta	IE/H/0715/001	OGYI-T-1352/03	GLAXOSMITHKLINE TRADING SERVICES LIMITED	HU
Augmentin 500 mg/125 mg filmtabletta	DE/H/6032/001	OGYI-T-1352/04	GLAXOSMITHKLINE TRADING SERVICES LIMITED	HU
Germentin 125 mg/31.25 mg per 5 ml Powder for Oral Suspension	not available	PA 0577/034/002	MCDERMOTT LABORATORIES LTD	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 Duo Mixed Fruit 400 mg/57 mg/5 ml Powder for Oral Suspension	DE/H/2868/006	PA 1077/19/6	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Germentin 250 mg/125 mg Film-coated Tablets	not available	PA0577/034/003	MCDERMOTT LABORATORIES LTD	IE
Augmentin 875 mg/125 mg film-coated tablets	DE/H/2868/002	PA 1077/19/5	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Augmentin 250 mg/125 mg film-coated tablets	IE/H/0715/001	PA 1077/93/7	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Augmentin 500 mg/125 mg film-coated tablets	DE/H/6032/001	PA 1077/19/3	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
CLAVULIN bambini 400 mg/57 mg/5 ml polvere per sospensione orale	not available	026138204	GLAXOSMITHKLINE S.P.A.	IT
CLAVULIN bambini 400	not available	026138216	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
mg/57 mg/5 ml polvere per sospensione orale				
CLAVULIN bambini 400 mg/57 mg/5 ml polvere per sospensione orale	not available	026138228	GLAXOSMITHKLINE S.P.A.	IT
Augmentin 875 mg/125 mg polvere per sospensione orale in bustine	IT/H/0270/001	026089324	GLAXOSMITHKLINE S.P.A.	IT
Augmentin 875 mg/125 mg polvere per sospensione orale in bustine	IT/H/0270/001	026089351	GLAXOSMITHKLINE S.P.A.	IT
Augmentin 875 mg/125 mg polvere per sospensione orale in bustine	IT/H/0270/001	026089336	GLAXOSMITHKLINE S.P.A.	IT
Augmentin 875 mg/125 mg polvere per sospensione orale in bustine	IT/H/0270/001	026089363	GLAXOSMITHKLINE S.P.A.	IT
Augmentin 875 mg/125 mg polvere per sospensione orale in bustine	IT/H/0270/001	026089348	GLAXOSMITHKLINE S.P.A.	IT
Augmentin bambini 400 mg/57 mg polvere per sospensione orale in bustine	IT/H/0270/002	026089375	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 875 mg/125 mg polvere per sospensione orale in bustine	IT/H/0270/001	026089108	GLAXOSMITHKLINE S.P.A.	IT
Augmentin 400 mg/57 mg/5 ml polvere per sospensione orale (aroma frutti misti)	DE/H/2868/006	026089514	GLAXOSMITHKLINE S.P.A.	IT
Augmentin 400 mg/57 mg/5 ml polvere per sospensione orale (aroma	DE/H/2868/006	026089488	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
frutti misti)				
Augmentin 400 mg/57 mg/5 ml polvere per sospensione orale (aroma frutti misti)	DE/H/2868/006	026089464	GLAXOSMITHKLINE S.P.A.	IT
Augmentin 400 mg/57 mg/5 ml polvere per sospensione orale (aroma frutti misti)	DE/H/2868/006	026089490	GLAXOSMITHKLINE S.P.A.	IT
Augmentin 400 mg/57 mg/5 ml polvere per sospensione orale (aroma frutti misti)	DE/H/2868/006	026089502	GLAXOSMITHKLINE S.P.A.	IT
Augmentin 400 mg/57 mg/5 ml polvere per sospensione orale (aroma frutti misti)	DE/H/2868/006	026089476	GLAXOSMITHKLINE S.P.A.	IT
Neoduplamox bambini 400 mg/57 mg/5 ml polvere per sospensione orale (aroma frutti misti)	not available	026141 200	VALEAS S.P.A.	IT
Neoduplamox bambini 400 mg/57 mg/5 ml polvere per sospensione orale (aroma frutti misti)	not available	026141 212	VALEAS S.P.A.	IT
Neoduplamox bambini 400 mg/57 mg/5 ml polvere per sospensione orale (aroma frutti misti)	not available	026141 224	VALEAS S.P.A.	IT
Neoduplamox bambini 400 mg/57 mg polvere per sospensione orale bustine	not available	026141 236	VALEAS S.P.A.	IT
Augmentin 875 mg/125 mg compresse rivestite con film	DE/H/2868/002	026089211	GLAXOSMITHKLINE S.P.A.	IT
Augmentin 875 mg/125 mg compresse rivestite con film	DE/H/2868/002	026089223	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/2868/002	026089235	GLAXOSMITHKLINE S.P.A.	IT
Augmentin 875 mg/125 mg compresse rivestite con film	DE/H/2868/002	026089247	GLAXOSMITHKLINE S.P.A.	IT
Augmentin 875 mg/125 mg compresse rivestite con film	DE/H/2868/002	026089250	GLAXOSMITHKLINE S.P.A.	IT
Augmentin 875 mg/125 mg compresse rivestite con film	DE/H/2868/002	026089262	GLAXOSMITHKLINE S.P.A.	IT
Augmentin 875 mg/125 mg compresse rivestite con film	DE/H/2868/002	026089274	GLAXOSMITHKLINE S.P.A.	IT
Augmentin 875 mg/125 mg compresse rivestite con film	DE/H/2868/002	026089312	GLAXOSMITHKLINE S.P.A.	IT
Augmentin 875 mg/125 mg compresse rivestite con film	DE/H/2868/002	026089286	GLAXOSMITHKLINE S.P.A.	IT
Augmentin 875 mg/125 mg compresse rivestite con film	DE/H/2868/002	026089298	GLAXOSMITHKLINE S.P.A.	IT
Augmentin 875 mg/125 mg compresse rivestite con film	DE/H/2868/002	026089300	GLAXOSMITHKLINE S.P.A.	IT
Augmentin 875 mg/125 mg compresse rivestite con film	DE/H/2868/002	026089019	GLAXOSMITHKLINE S.P.A.	IT
Clavulin 875 mg/125 mg compresse rivestite con film	not available	026138139	GLAXOSMITHKLINE S.P.A.	IT
Augmentin bambini 400 mg/57 mg/5 ml polvere per sospensione orale	PT/H/2363/001	026089110	GLAXOSMITHKLINE S.P.A.	IT
Augmentin bambini 400 mg/57 mg/5 ml polvere	PT/H/2363/001	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
per sospensione orale				
Augmentin bambini 400 mg/57 mg/5 ml polvere per sospensione orale	PT/H/2363/001	026089134	GLAXOSMITHKLINE S.P.A.	IT
Neoduplamox 875 mg /125 mg compresse rivestite con film	not available	026141 147	VALEAS S.P.A.	IT
CLAVULIN 875 mg/125 mg polvere per sospensione orale in bustine	not available	026138192	GLAXOSMITHKLINE S.P.A.	IT
CLAVULIN bambini 400 mg/57 mg polvere per sospensione orale in bustine	not available	026138230	GLAXOSMITHKLINE S.P.A.	IT
Neoduplamox 875 mg/125 mg polvere per sospensione orale bustine	not available	026141 198	VALEAS S.P.A.	IT
Augmentin 400 mg/57 mg/5 ml milteliai geriamajai suspensijai	DE/H/2868/006	LT/1/98/3675/001	GLAXOSMITHKLINE (IRELAND) LIMITED	LT
Augmentin 875 mg /125 mg plėvele dengtos tabletės	DE/H/2868/002	LT/1/98/3675/002	GLAXOSMITHKLINE (IRELAND) LIMITED	LT
Augmentin 875 mg /125 mg plėvele dengtos tabletės	DE/H/2868/002	LT/1/98/3675/003	GLAXOSMITHKLINE (IRELAND) LIMITED	LT
Augmentin 875 mg /125 mg plėvele dengtos tabletės	DE/H/2868/02	LT/1/98/3675/004	GLAXOSMITHKLINE (IRELAND) LIMITED	LT
Augmentin 875 mg /125 mg plevele dengtos tabletes	DE/H/2868/02	LT/1/98/3675/005	GLAXOSMITHKLINE (IRELAND) LIMITED	LT
Augmentin 875 mg /125 mg plevele dengtos tabletes	DE/H/2868/002	LT/1/98/3675/006	GLAXOSMITHKLINE (IRELAND) LIMITED	LT
Augmentin 875 mg /125 mg plevele dengtos tabletes	DE/H/2868/002	LT/1/98/3675/007	GLAXOSMITHKLINE (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
Augmentin 875 mg /125 mg plėvele dengtos tabletės	DE/H/2868/002	LT/1/98/3675/008	GLAXOSMITHKLINE (IRELAND) LIMITED	LT
Augmentin 875 mg /125 mg plevele dengtos tabletes	DE/H/2868/002	LT/1/98/3675/009	GLAXOSMITHKLINE (IRELAND) LIMITED	LT
Augmentin 875 mg /125 mg plevele dengtos tabletes	DE/H/2868/002	LT/1/98/3675/010	GLAXOSMITHKLINE (IRELAND) LIMITED	LT
Augmentin 875 mg /125 mg plevele dengtos tabletes	DE/H/2868/002	LT/1/98/3675/011	GLAXOSMITHKLINE (IRELAND) LIMITED	LT
Augmentin 875 mg /125 mg plevele dengtos tabletes	DE/H/2868/002	LT/1/98/3675/012	GLAXOSMITHKLINE (IRELAND) LIMITED	LT
Augmentin 500 mg/125 mg plevele dengtos tabletes	DE/H/6032/001	LT/1/98/3675/014	GLAXOSMITHKLINE (IRELAND) LIMITED	LT
Augmentin 500 mg/125 mg plėvele dengtos tabletės	DE/H/6032/001	LT/1/98/3675/015	GLAXOSMITHKLINE (IRELAND) LIMITED	LT
Augmentin 500 mg/125 mg plėvele dengtos tabletės	DE/H/6032/001	LT/1/98/3675/016	GLAXOSMITHKLINE (IRELAND) LIMITED	LT
Augmentin 500 mg/125 mg plevele dengtos tabletes	DE/H/6032/001	LT/1/98/3675/017	GLAXOSMITHKLINE (IRELAND) LIMITED	LT
Augmentin 500 mg/125 mg plevele dengtos tabletes	DE/H/6032/001	LT/1/98/3675/013	GLAXOSMITHKLINE (IRELAND) LIMITED	LT
Augmentin 500 mg/125 mg plėvele dengtos tabletės	DE/H/6032/001	LT/1/98/3675/018	GLAXOSMITHKLINE (IRELAND) LIMITED	LT
Augmentin 500 mg/125 mg plėvele dengtos tabletės	DE/H/6032/001	LT/1/98/3675/019	GLAXOSMITHKLINE (IRELAND) LIMITED	LT
Augmentin 500 mg/125 mg plėvele dengtos tabletės	DE/H/6032/001	LT/1/98/3675/020	GLAXOSMITHKLINE (IRELAND) LIMITED	LT
Augmentin 500 mg/125 mg plėvele dengtos tabletės	DE/H/6032/001	LT/1/98/3675/021	GLAXOSMITHKLINE (IRELAND) LIMITED	LT
Augmentin 500 mg/125 mg plėvele dengtos tabletės	DE/H/6032/001	LT/1/98/3675/022	GLAXOSMITHKLINE (IRELAND) LIMITED	LT
Augmentin 500 mg/125 mg plėvele dengtos tabletės	DE/H/6032/001	LT/1/98/3675/023	GLAXOSMITHKLINE (IRELAND) LIMITED	LT
Augmentin 200 mg/28,5	DE/H/6223/001	LT/1/98/3992/001	GLAXOSMITHKLINE (IRELAND)	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
mg/5 ml milteliai geriamajai suspensijai			LIMITED	
Augmentin P 500 mg/50 mg Pulver zur Herstellung einer Injektions-/Infusionslösung	DE/H/2809/005	2001 09 6519	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Augmentin 1.000 mg/200 mg Pulver zur Herstellung einer Injektions-/Infusionslösung	DE/H/2809/003	2001 09 6521	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Augmentin 2.000 mg/200 mg Pulver zur Herstellung einer Infusionslösung	DE/H/2809/004	2001 09 6522	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Augmentin 1000 mg/200 mg, poudre pour solution injectable/pour perfusion	DE/H/2809/003	2001 09 6521	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Augmentin 2000 mg/200 mg, poudre pour solution pour perfusion	DE/H/2809/004	2001 09 6522	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Augmentin P 500 mg/50 mg, poudre pour solution injectable/pour perfusion	DE/H/2809/005	2001 09 6519	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Augmentin P 1.000 mg/100 mg Pulver zur Herstellung einer Injektions-/Infusionslösung	DE/H/2809/006	2001 09 6520	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Augmentin P 1000 mg/100 mg, poudre pour solution injectable/pour perfusion	DE/H/2809/006	2001 09 6520	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Augmentin 500 mg/125 mg, poudre pour suspension buvable en sachet	BE/H/0159/01	2001096526	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Augmentin 500 mg/125 mg Pulver zur Herstellung einer Suspension zum Einnehmen in Beuteln	BE/H/0159/01	2001096526	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Augmentin 875 mg/125 mg	DE/H/2868/02	2001 09 6529	GLAXOSMITHKLINE	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
Filmtabletten			PHARMACEUTICALS SA	
Augmentin 875 mg/125 mg, comprimés pelliculés	DE/H/2868/02	2001 09 6529	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Augmentin 500 mg/125 mg Filmtabletten	DE/H/6032/001	2001 09 6523	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Augmentin 500 mg/125 mg, comprimés pelliculés	DE/H/6032/001	2001 09 6523	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Augmentin 250 mg/62,5 mg/5 ml, poudre pour suspension buvable	BE/H/0209/02	2001096525	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Augmentin 125 mg/31,25 mg/5 ml Pulver zur Herstellung einer Suspension zum Einnehmen	BE/H/0209/01	2001096524	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Augmentin 250 mg/62,5 mg/5 ml Pulver zur Herstellung einer Suspension zum Einnehmen	BE/H/0209/02	2001096525	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Augmentin 125 mg/31,25 mg/5 ml, poudre pour suspension buvable	BE/H/0209/01	2001096524	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Amoclane 875/125 mg poudre pour suspension buvable	not available	2009010037	EUROGENERICS N.V./S.A.	LU
Augmentin Fruit 400 mg/57 mg/5 ml pulveris iekšķīgi lietojamas suspensijas pagatavošanai (vairāku augļu garša)	DE/H/2868/006	17-0184	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LV
Augmentin 875 mg/125 mg apvalkotas tabletes	DE/H/2868/02	99-0035	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LV
Augmentin 500 mg/125 mg apvalkotās tabletes	DE/H/6032/001	99-0034	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LV
Augmentin Duo 400 mg/57 mg/5 ml powder for oral	DE/H/2868/006	MA192/01505	GLAXOSMITHKLINE (IRELAND) LIMITED	MT

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
suspension (mixed fruit flavour)				
Augmentin 875 mg/125 mg film-coated tablets	DE/H/2868/02	MA 192/01502	GLAXOSMITHKLINE (IRELAND) LIMITED	MT
Augmentin 250 mg/125 mg film-coated tablets	IE/H/0715/001	MA 192/01501	GLAXOSMITHKLINE (IRELAND) LIMITED	MT
Augmentin DUO 400 mg/57 mg/5 ml powder for oral suspension	PT/H/2363/001	MA 192/01504	GLAXOSMITHKLINE (IRELAND) LIMITED	MT
Augmentin 500 mg/125 mg film-coated tablets	UK/H/4738/01	MA192/01503	GLAXOSMITHKLINE (IRELAND) LIMITED	MT
Augmentin 100 mg/12,5 mg/ml, poeder voor orale suspensie	DE/H/2868/007	RVG 14740	GLAXOSMITHKLINE B.V.	NL
Augmentin 500 mg/125 mg, filmomhulde tabletten	DE/H/6032/001	RVG 09840	GLAXOSMITHKLINE B.V.	NL
Augmentin 400 mg/57 mg/5 ml pulver til mikstur, suspensjon (blandet fruktsmak)	DE/H/2868/006	16-11188	GLAXOSMITHKLINE AS	NO
Augmentin 500 mg/125 mg filmdrasjerte tabletter	DE/H/6032/001	14-10121	GLAXOSMITHKLINE AS	NO
Augmentin ES, (600 mg + 42,9 mg)/5 ml, proszek do sporzadzania zawiesiny doustnej	PT/H/0684/001	12306	GLAXOSMITHKLINE (IRELAND) LIMITED	PL
Augmentin MFF, (400 mg + 57 mg)/5 ml, proszek do sporządzania zawiesiny doustnej	DE/H/2868/006	24335	GLAXOSMITHKLINE (IRELAND) LIMITED	PL
Augmentin, 875 mg + 125, mg tabletki powlekane	DE/H/2868/02	R/7175	GLAXOSMITHKLINE (IRELAND) LIMITED	PL
Augmentin, 250 mg + 125 mg, tabletki powlekane	IE/H/0715/001	R/0641	GLAXOSMITHKLINE (IRELAND) LIMITED	PL
Augmentin, 500 mg + 125 mg, tabletki powlekane	DE/H/6032/001	R/3682	GLAXOSMITHKLINE (IRELAND) LIMITED	PL
Noprilam, 500 mg/125 mg, comprimidos revestidos por	not available	4717690	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
película				
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 250, 250 mg/62,5 mg/5 ml, pó para suspensão oral	not available	9596247	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
Clavamox 500, 500 mg/125 mg, comprimidos revestidos por película	not available	4717591	BIAL - PORTELA & Cª , SA	PT
Clavamox 500, 500 mg/125 mg, comprimidos revestidos por película	not available	9588822	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
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
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	PT/H/0684/001	5323787	GLAXOSMITHKLINE -	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
mg/42,9 mg/ 5 ml pó para suspensão oral			PRODUTOS FARMACEUTICOS, LDA	
Augmentin ES 600 mg/42,9 mg/ 5 ml pó para suspensão oral	PT/H/0684/001	5323886	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	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 125 mg/31,25 mg/5 ml, pó para suspensão oral	DE/H/2868/04	8605014	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Augmentin Forte 250 mg/62,5 mg/5 ml, pó para suspensão oral	DE/H/2868/005	8605006	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Augmentin Duo 400 mg/5 ml + 57 mg/5 ml pó para suspensão oral (sabor misto de frutas)	DE/H/2868/006	5738711	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Augmentin Duo 400 mg/5 ml + 57 mg/5 ml pó para suspensão oral (sabor misto de frutas)	DE/H/2868/006	5738422	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Augmentin Duo 400 mg/5 ml + 57 mg/5 ml pó para suspensão oral (sabor misto de frutas)	DE/H/2868/006	5738414	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Augmentin Duo 400 mg/5 ml + 57 mg/5 ml pó para suspensão oral (sabor misto de frutas)	DE/H/2868/006	5738729	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Clavamox 125, 125 mg/31,25 mg/5 ml, pó para suspensão oral	not available	9596239	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	not available	9596213	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
para suspensão oral				
Clavamox 125, 125 mg/31,25 mg/5 ml, pó para suspensão oral	not available	4823191	BIAL - PORTELA & Cª , SA	PT
Augmentin Duo 875 mg/125 mg, comprimidos revestidos por película	DE/H/2868/002	5751888	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Augmentin Duo 875 mg/125 mg, comprimidos revestidos por película	DE/H/2868/002	4714986	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Augmentin Duo 400 mg/57 mg/5 ml, pó para suspensão oral (sabor a morango)	PT/H/2363/001	2922482	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Clavamox DT 400, 400 mg/57 mg/5 ml, pó para suspensão oral	not available	2922698	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
Augmentin 500 mg/125 mg, comprimidos revestidos por película	DE/H/6032/001	4717286	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Augmentin 500 mg/125 mg, comprimidos revestidos por película	DE/H/6032/001	8604702	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Augmentin 500 mg/125 mg, comprimidos revestidos por película	DE/H/6032/001	4714689	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Noprilam DT, 875 mg/125 mg, comprimidos revestidos por película	not available	4715595	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
Noprilam, 125 mg/31,25 mg/5 ml, pó para suspensão oral	not available	4726899	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 DT, 875 mg/125 mg, comprimidos revestidos por película	not available	4716296	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
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
Clavamox ES 600 mg/42,9 mg/ 5 ml pó para suspensão oral	not available	5324389	BIAL - PORTELA & Cª , SA	PT
AUGMENTIN ES 600 mg/42,9 mg/ 5 ml pulbere	PT/H/0684/001	7850/2015/01	GLAXOSMITHKLINE (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
pentru suspensie orala				
AUGMENTIN ES 600 mg/42,9 mg/ 5 ml pulbere pentru suspensie orala	PT/H/0684/001	7850/2015/02	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
AUGMENTIN ES 600 mg/42,9 mg/ 5 ml pulbere pentru suspensie orală	PT/H/0684/001	7850/2015/04	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
AUGMENTIN ES 600 mg/42,9 mg/ 5 ml pulbere pentru suspensie orală	PT/H/0684/001	7850/2015/03	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Augmentin FP 400 mg/57 mg/5 ml pulbere pentru suspensie orală (aromă de fructe de pădure)	DE/H/2868/006	11562/2019/01	GLAXOSMITHKLINE TRADING SERVICES LIMITED	RO
Augmentin FP 400 mg/57 mg/5 ml pulbere pentru suspensie orală (aromă de fructe de pădure)	DE/H/2868/006	11562/2019/02	GLAXOSMITHKLINE TRADING SERVICES LIMITED	RO
Augmentin FP 400 mg/57 mg/5 ml pulbere pentru suspensie orală (aromă de fructe de pădure)	DE/H/2868/006	11562/2019/03	GLAXOSMITHKLINE TRADING SERVICES LIMITED	RO
Augmentin FP 400 mg/57 mg/5 ml pulbere pentru suspensie orală (aromă de fructe de pădure)	DE/H/2868/006	11562/2019/04	GLAXOSMITHKLINE TRADING SERVICES LIMITED	RO
Augmentin FP 400 mg/57 mg/5 ml pulbere pentru suspensie orală (aromă de fructe de pădure)	DE/H/2868/006	11562/2019/08	GLAXOSMITHKLINE TRADING SERVICES LIMITED	RO
Augmentin FP 400 mg/57 mg/5 ml pulbere pentru suspensie orală (aromă de fructe de pădure)	DE/H/2868/006	11562/2019/07	GLAXOSMITHKLINE TRADING SERVICES LIMITED	RO
Augmentin FP 400 mg/57 mg/5 ml pulbere pentru suspensie orală (aromă de fructe de pădure)	DE/H/2868/006	11562/2019/06	GLAXOSMITHKLINE TRADING SERVICES 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
Augmentin FP 400 mg/57 mg/5 ml pulbere pentru suspensie orală (aromă de fructe de pădure)	DE/H/2868/006	11562/2019/05	GLAXOSMITHKLINE TRADING SERVICES LIMITED	RO
Augmentin 875 mg/125 mg comprimate filmate	DE/H/2868/002	8164/2015/01	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Augmentin 875 mg/125 mg comprimate filmate	DE/H/2868/002	8164/2015/02	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Augmentin 875 mg/125 mg comprimate filmate	DE/H/2868/002	8164/2015/03	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Augmentin 875 mg/125 mg comprimate filmate	DE/H/2868/002	8164/2015/04	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Augmentin 875 mg/125 mg comprimate filmate	DE/H/2868/002	8164/2015/05	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Augmentin 875 mg/125 mg comprimate filmate	DE/H/2868/002	8164/2015/06	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Augmentin 875 mg/125 mg comprimate filmate	DE/H/2868/02	8164/2015/07	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Augmentin 875 mg/125 mg comprimate filmate	DE/H/2868/02	8164/2015/10	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Augmentin 875 mg/125 mg comprimate filmate	DE/H/2868/02	8164/2015/11	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Augmentin 875 mg/125 mg comprimate filmate	DE/H/2868/02	8164/2015/12	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Augmentin 875 mg/125 mg comprimate filmate	DE/H/2868/02	8164/2015/08	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Augmentin 875 mg/125 mg comprimate filmate	DE/H/2868/02	8164/2015/09	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Augmentin 500 mg/125 mg comprimate filmate	DE/H/6032/001	8395/2015/01	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Augmentin 500 mg/125 mg comprimate filmate	DE/H/6032/001	8395/2015/02	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Augmentin 500 mg/125 mg comprimate filmate	DE/H/6032/001	8395/2015/03	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Augmentin 500 mg/125 mg comprimate filmate	DE/H/6032/001	8395/2015/04	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Augmentin 500 mg/125 mg comprimate filmate	DE/H/6032/001	8395/2015/05	GLAXOSMITHKLINE (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
Augmentin 500 mg/125 mg comprimate filmate	DE/H/6032/001	8395/2015/06	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Augmentin 500 mg/125 mg comprimate filmate	DE/H/6032/001	8395/2015/07	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Augmentin 500 mg/125 mg comprimate filmate	DE/H/6032/001	8395/2015/08	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Augmentin 500 mg/125 mg comprimate filmate	DE/H/6032/001	8395/2015/09	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Augmentin 500 mg/125 mg comprimate filmate	DE/H/6032/001	8395/2015/10	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Augmentin 500 mg/125 mg comprimate filmate	DE/H/6032/001	8395/2015/11	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Augmentin 500 mg/125 mg comprimate filmate	DE/H/6032/001	8395/2015/12	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Augmentin 500 mg/125 mg comprimate filmate	DE/H/6032/001	8395/2015/13	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Spektramox 500 mg/125 mg filmdragerade tabletter	not available	12415	VIATRIS AB	SE
Spektramox 875 mg/125 mg filmdragerade tabletter	not available	15979	VIATRIS AB	SE
Spektramox 50 mg/ml+12,5 mg/ml pulver till oral suspension	not available	10700	MEDA AB	SE
Spektramox 80 mg/ml+12 mg/ml pulver till oral suspension	not available	15980	MEDA AB	SE
Augmentin 875 mg/125 mg filmsko obložene tablete	DE/H/2868/002	H/99/00239/001	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Augmentin 875 mg/125 mg filmsko obložene tablete	DE/H/2868/002	H/99/00239/002	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Augmentin 875 mg/125 mg filmsko obložene tablete	DE/H/2868/002	H/99/00239/003	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Augmentin 875 mg/125 mg filmsko obložene tablete	DE/H/2868/002	H/99/00239/004	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Augmentin 875 mg/125 mg filmsko obložene tablete	DE/H/2868/002	H/99/00239/005	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Augmentin 875 mg/125 mg filmsko obložene tablete	DE/H/2868/002	H/99/00239/006	GLAXOSMITHKLINE TRADING SERVICES LIMITED	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 875 mg/125 mg filmsko obložene tablete	DE/H/2868/002	H/99/00239/007	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Augmentin 875 mg/125 mg filmsko obložene tablete	DE/H/2868/002	H/99/00239/008	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Augmentin 875 mg/125 mg filmsko obložene tablete	DE/H/2868/002	H/99/00239/009	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Augmentin 875 mg/125 mg filmsko obložene tablete	DE/H/2868/002	H/99/00239/010	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Augmentin 875 mg/125 mg filmsko obložene tablete	DE/H/2868/002	H/99/00239/011	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Augmentin 875 mg/125 mg filmsko obložene tablete	DE/H/2868/002	H/99/00239/012	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
MEGAMOX 625 Forte Filmom obalené tablety	not available	15/0109/99-S	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	SK
Amoksiklav 600 mg prášok na injekcný/infúzny roztok	not available	15/0063/20-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
MEGAMOX 375 Filmom obalené tablety	not available	15/0108/99-S	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	SK
MEGAMOX 375 Filmom obalené tablety	not available	15/0108/99-S	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	SK
Augmentin ES 600 mg/42,9 mg/5 ml prášok na perorálnu suspenziu	PT/H/0684/001	15/0085/07-S	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SK
Augmentin DUO s príchutou miešaného ovocia 400 mg/5 ml + 57 mg/5 ml prášok na perorálnu suspenziu	DE/H/2868/006	15/0375/17-S	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SK
Augmentin 1 g 875 mg/125 mg filmom obalené tablety	DE/H/2868/002	15/0682/96-S	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SK
Augmentin 375 mg 250 mg/125 mg filmom obalené tablety	IE/H/0715/001	15/0141/84-C/S	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SK
Augmentin 625 mg 500 mg/125 mg filmom	DE/H/6032/001	15/0126/13-S	GLAXOSMITHKLINE TRADING SERVICES 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
obalené tablety				
Amoksiklav 1000 mg granulát na perorálnu suspenziu	IT/H/0245/001	15/0771/10-S	SANDOZ PHARMACEUTICALS D.D.	SK
Augmentin 125/31 Suspension	DE/H/2868/04	PL 00038/0298	BEECHAM GROUP PLC	XI
Augmentin 250/62 Suspension	DE/H/2868/005	PL 00038/0337	BEECHAM GROUP PLC	XI
Augmentin Duo 400/57 powder for oral suspension	DE/H/2868/006	PL 10592/0303	SMITHKLINE BEECHAM LTD	XI
Augmentin 1 g Tablets	DE/H/2868/002	PL 00038/0368	BEECHAM GROUP LTD	XI
Augmentin 375 mg Tablets	IE/H/0715/001	PL 00038/0270	BEECHAM GROUP PLC	XI
Augmentin 400/57.	PT/H/2363/001	PL 10592/0070	SMITHKLINE BEECHAM LTD	XI
Augmentin 625 mg Tablets	DE/H/6032/001	PL 00038/0362	BEECHAM GROUP LTD	XI
Augmentin Duo 200/28	DE/H/6223/001	PL 10592/0072	SMITHKLINE BEECHAM LTD	XI