



EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

9 June 2017
EMA/372458/2017
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: artemether / lumefantrin (apart from the dispersible tablet)

Procedure no.: PSUSA/00000236/201610



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
Riamet® 20 mg/120 mg Tabletten	UK/H/0351/001	BE223562	NOVARTIS PHARMA N.V.	BE
RIAMET 20 MG/120 MG, COMPRIMÉ	UK/H/0351/001	3400927603303	NOVARTIS PHARMA S.A.S.	FR
RIAMET® δηζθία 20/120 mg	UK/H/0351/001	250070102	NOVARTIS (HELLAS) S.A.C.I.	GR
RIAMET® δηζθία 20/120 mg	UK/H/0351/001	250070104	NOVARTIS (HELLAS) S.A.C.I.	GR
RIAMET® δηζθία 20/120 mg	UK/H/0351/001	250070105	NOVARTIS (HELLAS) S.A.C.I.	GR
Riamet 20 mg/120 mg comprimidos	UK/H/0351/001	63.764	NOVARTIS FARMACÉUTICA S.A.	ES
РИАМЕТ 20 MG/120 MG ТАБЛЕТКИ	not available	20010737	NOVARTIS PHARMA GMBH	BG

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Riamet® 20 mg/120 mg Tabletten	UK/H/0351/001	50318.00.00	NOVARTIS PHARMA GMBH	DE
Riamet 20 mg/120 mg, tabletten	UK/H/0351/001	RVG 25773	NOVARTIS PHARMA B.V.	NL
RIAMET 20 mg/120 mg comprimés	UK/H/0351/001	BE223562	NOVARTIS PHARMA N.V.	BE
Riamet 20 mg/120 mg tablett	UK/H/0351/001	16557	NOVARTIS SVERIGE AB	SE
Riamet 20 mg/120 mg tabletten	UK/H/0351/001	BE223562	NOVARTIS PHARMA N.V.	BE
Riamet 20 mg/120 mg - Tabletten	UK/H/0351/001	1-24024	NOVARTIS PHARMA 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
Riamet® 20 mg/120 mg tablets	UK/H/0351/001	PL 00101/0566	NOVARTIS PHARMACEUTICALS UK LIMITED	UK
RIAMET 20 mg/120 mg comprimidos	UK/H/0351/001	3466687	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT