



EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

28 May 2020
EMA/372797/2020
Human Medicines Evaluation Division

List of nationally authorised medicinal products

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

Procedure no.: PSUSA/00000236/201910



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	SE/H/1778/001	BE223562	NOVARTIS PHARMA N.V.	BE
RIAMET 20 MG/120 MG, COMPRIMÉ	SE/H/1778/001	3400927603303	NOVARTIS PHARMA S.A.S.	FR
RIAMET® δηζθία 20/120 mg	SE/H/1778/001	250070102	NOVARTIS (HELLAS) S.A.C.I.	GR
RIAMET® δηζθία 20/120 mg	SE/H/1778/001	250070104	NOVARTIS (HELLAS) S.A.C.I.	GR
RIAMET® δηζθία 20/120 mg	SE/H/1778/001	250070105	NOVARTIS (HELLAS) S.A.C.I.	GR
Riamet 20 mg/120 mg comprimidos	SE/H/1778/001	63.764	NOVARTIS FARMACÉUTICA S.A.	ES
Riamet® 20 mg/120 mg Tabletten	SE/H/1778/001	50318.00.00	NOVARTIS PHARMA GMBH	DE
Riamet 20 mg/120 mg, tabletten	SE/H/1778/001	RVG 25773	NOVARTIS PHARMA B.V.	NL
RIAMET 20 mg/120 mg comprimés	SE/H/1778/001	BE223562	NOVARTIS PHARMA N.V.	BE
Riamet 20 mg/120 mg tablett	SE/H/1778/001	16557	NOVARTIS SVERIGE AB	SE
Riamet 20 mg/120 mg tabletten	SE/H/1778/001	BE223562	NOVARTIS PHARMA N.V.	BE
Riamet 20 mg/120 mg - Tabletten	SE/H/1778/001	1-24024	NOVARTIS PHARMA GMBH	AT
Riamet® 20 mg/120 mg tablets	SE/H/1778/001	PL 00101/0566	NOVARTIS PHARMACEUTICALS UK LIMITED	UK
RIAMET 20 mg/120 mg	SE/H/1778/001	3466687	NOVARTIS FARMA -	PT

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comprimidos			PRODUTOS FARMACÊUTICOS S.A.	
Riamet 20 mg/120 mg Tabletten	SE/H/1778/001	BE223562	NOVARTIS PHARMA N.V.	BE
RIAMET 20 MG/120 MG, COMPRIMÉ	SE/H/1778/001	3400927603303	NOVARTIS PHARMA S.A.S.	FR