12 May 2023
EMA/88681/2023
Human Medicines Division

List of nationally authorised medicinal products

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

Procedure No. PSUSA/00000236/202210

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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
Riamet 20 mg/120 mg - Tabletten	SE/H/1778/001	1 - 24024	NOVARTIS PHARMA GMBH	AT
Riamet 20 mg/120 mg Tabletten	SE/H/1778/001	BE223562	NOVARTIS PHARMA N.V.	BE
Riamet 20 mg/120 mg comprimés	SE/H/1778/001	BE223562	NOVARTIS PHARMA N.V.	BE
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	50318.00.00	NOVARTIS PHARMA GMBH	DE
RIAMET 20 mg/120 mg, comprimé	SE/H/1778/001	34009 276 033 0 3	NOVARTIS PHARMA S.A.S.	FR
RIAMET® δηζθϊα 20/120 mg	SE/H/1778/001	250070104	NOVARTIS (HELLAS) S.A.C.I.	GR
Riamet 20 mg/120 mg, tabletten	SE/H/1778/001	RVG 25773	NOVARTIS PHARMA B.V.	NL
Riamet 20 mg/120 mg comprimidos	SE/H/1778/001	3466687	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Riamet 20 mg/120 mg tablett	SE/H/1778/001	16557	NOVARTIS SVERIGE AB	SE
Riamet® 20 mg/120 mg tablets	SE/H/1778/001	PL 23860/0024	NOVARTIS IRELAND LIMITED	XI