11 June 2026
EMADOC-1700519818-3242706
Human Medicines Division

List of nationally authorised medicinal products

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

EURD list No. PSUSA/00000236/202510

Official address Domenico Scarlattilaan 6 ● 1083 HS Amsterdam ● The Netherlands
Address for visits and deliveries Refer to www.ema.europa.eu/how-to-find-us
Send us a question Go to www.ema.europa.eu/contact **Telephone** +31 (0)88 781 6000 An agency of the European Union

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.	AT
Riamet 20 mg/120 mg comprimés	SE/H/1778/001	34009 276 033 0 3	NOVARTIS PHARMA S.A.S.	BE
Riamet 20 mg/120 mg comprimidos	SE/H/1778/001	50318.00.00	NOVARTIS PHARMA GMBH	BE
Riamet 20 mg/120 mg tablett	SE/H/1778/001	RVG 25773	NOVARTIS PHARMA B.V.	BE
Riamet 20 mg/120 mg Tabletten	SE/H/1778/001	BE223562	NOVARTIS PHARMA N.V.	DE
Riamet 20 mg/120 mg tabletten	SE/H/1778/001	16557	NOVARTIS SVERIGE AB	FR
RIAMET 20 mg/120 mg, comprimé	SE/H/1778/001	BE223562	NOVARTIS PHARMA N.V.	NL
Riamet 20 mg/120 mg, tabletten	SE/H/1778/001	1 - 24024	NOVARTIS PHARMA GMBH	PT
Riamet® 20 mg/120 mg Tabletten	SE/H/1778/001	3466687	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	SE