09 June 2016
EMA/413583/2016
Procedure Management and Committees Support

List of nationally authorised medicinal products

Active substance: artemether / lumefantrin (dispersible tablet)

Procedure no.: PSUSA/00009060/201510

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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 Dispersible tablet 20/120 mg	not available	PL 00101/0957	NOVARTIS PHARMACEUTICALS UK LIMITED	UK