

Artesunate

Artesunate dosing is	weight based; e	nsure accu	racy of docur	mented weigl	nt before admi	nistration
Form	Artesunate Artesun® 60mg powder for injection 1 ampoule of 1mL sodium bicarbonate 1 ampoule of 5mL sodium chloride solution for injection				Store at room temperature in outer box for light protection.	
Reconstitution	Determine the number of vials needed					
	Weight	<25 kg	26-50 kg	51-75 kg	76-100 kg	101- 125kg
	60mg vial	1	2	3	4	5
Compatibility &	 Draw up 1 mL of the supplied sodium bicarbonate solvent Add to the artesunate powder. Shake for several minutes until the powder is dissolved and the solution clear. For intravenous use after reconstitution draw up 5 mL of the supplied sodium chloride 0.9% solvent Add to the reconstituted artesunate solution, which yields a solution containing artesunate 10mg/mL (60mg in 6mL) Shake to mix well No further dilution required					
Stability	Reconstituted solution should be used immediately					
Administration	Inject the desired volume (0.24 mL/kg) slowly over 1-2 minutes.					
Monitoring	Monitor blood pressure, heart rate, respiratory rate, signs of hypersensitivity and haemoglobin levels. Monitor patients for 4 weeks after treatment for evidence of haemolytic anaemia.					
Additional Information	 Give 2.4mg/kg IV/IM at 0, 12, 24 hours, then every 24 hours until oral treatment can be substituted e.g. 168mg in a 70kg patient No dose adjustment required in renal or hepatic impairment This is an Unlicensed medication in Ireland- please contact pharmacy to ensure adequate stock available. Stock kept in ED and Pharmacy Discuss all patients with ID 					

Information provided relates to Artesun® Fosun Pharma