

MALIGNANT HYPERTHERMIA TREATMENT WITH AGILUS

(dantrolene sodium hemiheptahydrate)¹

In combination with adequate support measures, AGILUS® is indicated for the treatment of malignant hyperthermia in adults and children of all ages.¹



AGILUS® should be administered rapidly by intravenous injection at an **initial dose of 2.5 mg/kg body weight** for both adult and paediatric patients.¹



1

Calculate

the number of vials of AGILUS® required (see table below). Each vial contains 120mg of dantrolene.



2

Reconstitute

each vial with 20 mL of water for injections and shake for approximately 1 minute, before inspecting for particulates. Further shaking may be necessary.



3

Administer

reconstituted AGILUS® without mixing with other solutions or giving via the same venous access.



4

Apply

other supportive measures alongside AGILUS®.

Dosing examples by body weight to achieve a loading dose of 2.5 mg/kg for both adults and children¹

Number of vials to be prepared ^a	Body weight range	Example dosing recommendation		
		Body weight	Dose to be administered	Volume to be administered ^a
1	Up to 48 kg	3 kg	7.5 mg	1.4 mL
		6 kg	15 mg	2.8 mL
		12 kg	30 mg	5.6 mL
		24 kg	60 mg	11.3 mL
		48 kg	120 mg	22.6 mL
2	From 49 kg to 96 kg	72 kg	180 mg	33.9 mL
		96 kg	240 mg	45.2 mL
3	From 97 kg	120 kg	300 mg	56.5 mL
		144 kg ^b	300 mg ^b	56.5 mL

a. Total volume of one reconstituted vial is 22.6mL
b. For all body weights, the initial dose and any repeat doses should not exceed 300 mg, equivalent to 2.5 vials



As long as the main clinical symptoms persist, bolus injection of 2.5 mg/kg should be repeated every **10 minutes** until physiological and metabolic abnormalities improve.¹ If a cumulative dose of 10 mg/kg or above is considered, the diagnosis of malignant hyperthermia should be re-examined.¹

Click or scan to view Prescribing information

Adverse events should be reported. Reporting forms and information can be found at www.mhra.gov.uk/yellowcard. Adverse events should also be reported to Norgine Pharmaceuticals Ltd on: Tel: +44 (0)1895 826 606 Email: medinfo@norgine.com



Note: AGILUS® is a skeletal muscle relaxant. The most commonly reported adverse event is skeletal muscle weakness.

1. AGILUS®. Summary of product characteristics

To request further copies of this material, please contact: ukisupport@norgine.com

Job code: UK-EC-AGI-2400039 Date of preparation: December 2024