SULFAMETHOXAZOLE AND TRIMETHOPRIM (IV)

(previously known as cotrimoxazole) This drug must be guardrailed

Trade Name	Sulfamethoxazole and Trimethoprim Concentrate Injection BP Bactrim® Bactrimel® (Section 29)			
Class	Antibiotic, sulphonamide derivative + folate antagonist			
Mechanism of Action	Sulfamethoxazole (SMX) interferes with bacterial folic acid synthesis and cell growth. Trimethoprim (TMP) inhibits enzymes in the folic acid pathway.			
Indications	Indication 1: Pneumocystis carinii prophylaxis Indication 2: Pneumocystis carinii treatment			
Contraindications	Jaundice - increases risk of kernicterus as sulfamethoxazole competes for protein binding sites usually available to bilirubin. G6PD deficiency - increased risk of haemolytic anaemia			
Supplied As	Sulfamethoxazole and Trimethoprim Concentrate Injection BP containing 400mg of sulfamethoxazole and 80mg of trimethoprim per 5mL ampoule (480mg/5mL)			
Dilution	Shake the ampoule well before use then dilute:			
	Drug	5% Glucose Added	Final Volume	Concentration
	96mg (1mL)	29mL	30mL	3.2mg/mL
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	Drug	5% Glucose Added	Final Volume	Concentration
	96mg (1mL)	14mL	15mL	6.4mg/mL
	This solution is not stable and any remaining portion should be discarded immediately after use.			
	In severe fluid restriction: Sulfamethoxazole/trimethoprim may be given undiluted via a central line (480mg/5mL = 96mg/mL) Ensure the dose volume is at least 0.5ml for accurate administration			
Dosage / Interval	All dose references in this profile relate to 'cotrimoxazole' but			
Must chart guardrail and use Alaris pump	must be charted as the combination of sulphamethoxazole and trimethoprim			
	Indicati	on 1: Prophylaxis		
	450mg/m ² up to a maximum dose of 960mg			
	$m^2 = (0.05 \times wt(kg)) + 0.05$ Indication 2: Treatment = 60mg/kg/dose			
	Indication 2: Treatment - 60mg/kg/dose			

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Guardrail	Concentration: Min – 3.2mg/mL Max – 96mg/mL			
	Soft Alert Min: 30mg/kg/hr Hard Alert Max: 60mg/kg/hr			
	Soft Alert Max: 40mg/kg/hr Default Setting: 40mg/kg/hr			
Interval	Indication 1: Twice a day for 3 days of the week			
	Indication 2: 12 hourly for 14 days			
Administration	Indication 1: Prophylaxis is usually given orally so use suspension and see oral sulfamethoxazole +trimethoprim, drug profile Indication 2: Treatment can be given orally (see oral sulfamethoxazole +trimethoprim, drug profile) or iv infusion over 60 minutes			
Compatible With	5% and 10% glucose, 0.45% and 0.9% sodium chloride, glucose/sodium chloride combinations and Hartman's Solution.			
Incompatible With	Do not mix with other medications.			
	(NICU tend to use relatively high concentrations and there's no data on compatibility with other medicines available)			
Monitoring	Nil			
Stability	Prepare solutions immediately before use and commence infusion within 30 minutes of preparation			
	Discard any remaining solution after 24 hours. (NB: concentrated solutions should be discarded immediately after use)			
Storage	Store below 30 C, protect from light, Do Not Refrigerate.			
	Do not use any solution that is cloudy or has visible precipitate			
Adverse Reactions	Skin rashes, stop at first sign of rash due to risk of Stevens Johnson Syndrome.			
	Blood dyscrasias, hepatitis, vomiting, cough, breathing difficulties			
Metabolism	Eliminated in the urine. Protein binding 68%.			
Comments	Intravenous administration is not to be used in treatment of newborn infants except on advice of a consultant for management of Pneumocystis carinii. pH of injection = approx 10, sodium content = 37mg/5mL			
References	 BNF for Children 2006 NZHPA Notes on Injectable Drugs 5th Edition 2004 Medicines for Children, RCPCH, 1999 Neofax, 1999 BNF for Children 2007 			

Updated By	P Schmidt, B Robertshawe February 2006 A Lynn, B Robertshawe Oct 2007 A Lynn, B Robertshawe, F Robertson May 2009 (new pumps) A Lynn, B Robertshawe September 2009 A Lynn, B Robertshawe July 2012 (re-order profile) M Wallenstein, A Lynn, B Robertshawe September 2020 (update)
	A Lynn, B Robertshawe February 2022