## Clindamycin

Newborn use only

Alert	In the Australian context, clindamycin is not used as first line therapy for infections in neonates. Infectious			
	Diseases consultation is recommended prior to commencement.			
	May be used for penicillin allergic patients or other patients for whom penicillin is inappropriate, provided			
	the target organism is also expected to be susceptible to clindamycin.			
	peopates <sup>(6)</sup>	yr alconol. Avoid exposure of > s	by mg/kg/day of benzyl alconol in	
Indication	Treatment of infections with susc	entible organisms where first-li	ne therapy is contraindicated or	
malcation	unavailable		the therapy is contraindicated of	
	Suitable infections may include in	traabdominal infections, skin a	nd soft tissue infections or bone a	nd ioint
	infections.			, <b>,</b>
Action	Binds to the 50S subunit of susce	ptible bacterial ribosomes and i	nhibits protein synthesis. <sup>(1)</sup>	
Drug type	Lincosamide antibiotic derived from	om lincomycin.	· · ·	
Trade name	Dalacin C, Clindamycin Mylan.			
Presentation	300 mg/2 mL, 600 mg/4 mL (150 mg/mL)			
Dose	IV <sup>(2)</sup> *	<b>.</b>		
	* In the Australian context, clind	amycin is not used as the first	line therapy for infections. Infect	ious
	Diseases consultation is recomm	ended.		_
	Corrected Gestational	Dose	Frequency	
	Age/Postmenstrual Age*			-
	≤32 weeks	5 mg/kg	8 <sup>th</sup> hourly	_
	33 <sup>+0</sup> -40 <sup>+6</sup> weeks	7 mg/kg	8 <sup>th</sup> hourly	-
	≥41 weeks	9 mg/kg	8 <sup>th</sup> hourly	
Dose adjustment	Therapeutic hypothermia – No in	formation.		
	ECMO – No information.			
	Renai Impairment – No dose adju	stment is necessary.	nont	
Maximum dasa	Repatic impairment – Ose with Ca	aution in severe nepatic impair	nent.	
Total cumulative				
dose				
Route	Intravenous			
Preparation	Draw up 0.5 mL (75 mg) of clindamycin and add 24.5 mL of sodium chloride 0.9% or glucose 5% to make a		make a	
•	final volume of 25 mL with a cond	centration of 3 mg/mL.	Ū.	
Administration	IV infusion over 1 hour			
Monitoring	Full blood count, hepatic and ren	al function during prolonged tre	eatment	
Contraindications	Serious allergic reaction to clindamycin or lincomycin or to any of the inactive ingredients.			
Precautions				
Drug interactions	CYP3A4 inhibitors may potentially	y increase the clindamycin conc	entrations and a risk of clindamyc	in
	toxicity.			
Adverse	Diarrhoea (mild-to-severe), nause	ea, vomiting, abdominal pain or	cramps, rash, itch.	
reactions				
Compatibility	Fluids: Glucose 5%, glucose in soc	dium chloride solutions, sodium	chloride 0.9%,	a diuma
	Y-Site <sup>(1)</sup> : Aciciovir, amikacin sulfat	e, aztreonam, cephamandole n	davmadatamidina, digavin, dana	soaium,
	enhedrine sulfate fentanyl furos	emide benarin sodium bydroc	ortisone sodium succinate gentai	micin
	morphine sulfate noradrenaline	(noreninenhrine) paracetamol	netilmicin sulfate nineracillin-taz	obactam
	(FDTA-free), potassium chloride.	remifentanil, sodium bicarbona	te, suxamethonium, tobramycin.	obactam
	vancomycin, zidovudine.	,		
Incompatibility	Azithromycin, calcium gluconate,	ceftriaxone, ciprofloxacin, cefa	lothin, ganciclovir, gentamicin, ka	namycin,
-	magnesium sulfate, penicillin or c	arbenicillin, pentamidine, phen	obarbital.	
Stability	Mylan: To reduce microbiological	hazard, use as soon as practica	ble after dilution. If storage is nec	essary,
	hold at 2 to 8°C for not more thar	n 24 hours. <sup>(8)</sup>		
Storage	Dalacin C: Store below 8°C. Do no	ot freeze.		
	Mylan brand: Store below 25°C.			

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Excipients	Dalacin C: Benzyl alcohol, disodium edetate, hydrochloric acid, sodium hydroxide, water for injections. Mylan brand: Disodium edetate, water for injections, hydrochloric acid and sodium hydroxide. Mylan brand does not contain benzyl alcohol.	
Special comments		
Evidence	<b>Background</b> Clindamycin is effective in vitro against many gram positive cocci, particularly Group A beta-haemolytic streptococci, <i>Streptococcus pneumoniae</i> , and methicillin-susceptibile and resistant <i>Staphylococcus aureus</i> , chough all of these may be resistant to clindamycin and susceptibility should be confirmed. It may also be effective against a wide range of gram positive anaerobic bacteria, including penicillin-resistant Bacteroides species. Aerobic gram negative bacteria are not usually susceptible to clindamycin. <sup>(3)</sup> It is used as the alternate to penicillin in streptococcal and staphylococcal infections and as a primary agent for nfections caused by penicillin resistant anaerobic bacilli. <sup>(4)</sup> It is approved for adults and children for systemic treatment of staphylococcal, streptococcal, and anaerobic bacterial infections and complicated ntraabdominal infections. <sup>(1, 5)</sup> Because of its profile and high oral bioavailability, it is also suggested as part of an oral multimodal alternative for prolonged parenteral antibiotic regimens e.g. to treat bone and joint or prosthesis-related infections. <sup>(1)</sup> <b>Efficacy</b> Gonzalez et al performed a prospective, multicentre clinical trial to determine pharmacokinetics (PK) and safety of intravenous clindamycin in preterm and term infants. <sup>(2)</sup> In this study, authors developed population based PK model using the combined PK data collected from 3 prospective clinical trials: Staph Trio, PTN POPS and CLINO1. From Staph Trio trial, authors enrolled 21 infants with median (range) number of clindamycin asmples per infant was 3 (2 to 4). They combined this data with additional PK samples collected from 41 preterm and term infants <121-day postnatal age in PTN POPS trial. The median (range) GA and PNA values from PTN POPS trial were 33 weeks (22-42 weeks) and 16 days (1 to 115) respectively. The median clindamycin dose was 5.1 mg/kg/dose (3.8 to 13.5) and 15 mg/kg/ay (7.6 to 40.6). The final population PK model developed by the authors using simulate	
Practice points		
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