Newborn Use Only

Alert	Note: 1 unit = 1000 milliunits	
	When using for diabetes insipidus (DI), Paediatric Endocrine consultation should be obtained.	
	Management should be in intensive care where monitoring and expertise are readily available.	
Indication	1. Treatment of refractory hypotension.	
	2. Adjunctive treatment of pulmonary hypertension.	
	3. Acute antidiuretic hormone (ADH) replacement when diagnosis of diabetes insipidus established.	
	[The drug of choice for the treatment of diabetes insipidus is desmopressin (dDAVP). An	
	argipressin infusion should be considered in the initial management of post-surgical or post-	
	traumatic DI.]	
	4. Adjunct in acute massive haemorrhage of gastrointestinal tract or oesophageal varices (specialist use only) [Terlipressin or octreotide preferred].	
Action	Antidiuretic hormone secreted by the posterior pituitary. Its release is mediated either by high serum	
	osmolality or by a hypotension/low right atrial pressure baroreflex. Argipressin acts via V1A receptors in	
	blood vessels, causing vasoconstriction, and via V ₂ receptors in the renal tubules, causing anti-diuresis	
	Provokes vasodilatation in some vascular beds via its action on oxytocin receptors.	
Drug Type	Vasopressor.	
Trade Name	Pitressin.	
Presentation	20 unit/1 mL ampoule	
Dose	For hypotension:	
	10-50 milliunit/kg/hour infusion.	
	For a day of the sector of the	
	For pulmonary hypertension:	
	10-20 milliunit/kg/hour (can be commenced at 6 milliunit/kg/hour to a maximum 70 milliunit/kg/hour	
	For diabetes insipidus:	
	Starting dose: 0.5 milliunit/kg/hour	
	Dose range: 0.5 to 1 milliunit/kg/hour. May increase to 2 milliunit/kg/hour.	
	The final wean may be from 0.5 to 0.25 milliunit/kg/hour.	
	For acute massive gastrointestinal bleeding:	
	May not be best agent for this indication.	
	Commence argipressin 120milliunit/kg/hour. Increase (titrate) over 2 hours to maximal dose of 600	
	milliunit/kg/hour. Monitor carefully for side effects including fluid retention, electrolyte abnormalities	
	hypertension and cardiac arrhythmias. If bleeding not controlled at dose 600 milliunit/kg/hour	
	(10milliunit/kg/minute) then unlikely to be controlled at higher doses and other measures should be	
	used.	
Dose adjustment		
Maximum dose	For hypotension: 120 milliunit/kg/hour (2 milliunit/kg/minute). [Note up to 480 milliunit/kg/hour (8	
	milliunit/kg/minute) has been reported.]	
	For acute massive gastrointestinal bleeding: 600 milliunit/kg/hour (10 milliunit/kg/min).	
Total cumulative		
dose		
Route	Continuous IV infusion.	
Preparation	Note: 1 unit = 1000 milliunits.	
	HYPOTENSION/PULMONARY HYPERTENSION:	
	Single strength continuous IV infusion	
	Infusion strength Prescribed amount	
	1 mL/hour = 50 milliunit/kg/hour 2.5 unit/kg argipressin and make up to 50 mL	
	Draw up 0.125 mL/kg argin ressin (2.5 unit/kg) and add sodium chloride 0.0% or glucose 5% to make a	
	Draw up 0.125 mL/kg argipressin (2.5 unit/kg) and add sodium chloride 0.9% or glucose 5% to make a final volume of 50 mL with a final concentration of 50 milliunit/kg/mL	
	Draw up 0.125 mL/kg argipressin (2.5 unit/kg) and add sodium chloride 0.9% or glucose 5% to make a final volume of 50 mL with a final concentration of 50 milliunit/kg/mL. Infusing at a rate of 1 mL/hour = 50 milliunit/kg/hour.	

	DOUBLE STRENGTH continuous IV inf	usion
	Infusion strength	Prescribed amount
	1 mL/hour = 100	5 unit/kg argipressin and make up to 50 mL
	milliunit/kg/hour	
	Draw up 0.25 mL/kg argipressin (5 uni volume of 50 mL with a final concentr Infusing at a rate of 1 mL/hour = 100 u	
		the fractions
	QUADRUPLE STRENGTH continuous I	
	Infusion strength	Prescribed amount
	1 mL/hour = 200 milliunit/kg/hour	10 unit/kg argipressin and make up to 50 mL
	Draw up 0.5 mL/kg argipressin (10 uni volume of 50 mL with a final concentr Infusing at a rate of 1 mL/hour = 200 i	-
	DIABETES INSIPIDUS	
	Continuous IV infusion	Prescribed amount
	Infusion strength 1 mL/hour = 0.8 milliunit/kg/hour	40 milliunit/kg argipressin and make up to 50 mL.
	Add 0.1 mL (2 units) of argipressin (20 make a 4 milliunit/mL solution FURTHER DILUTE	unit/mL ampoule) to 500 mL bag of sodium chloride 0.9% to
	volume of 50 mL with a final concentr Infusing at a rate of 1 mL/hour = 0.8 n	-
	GASTROINTESTINAL BLEEDING	linfusion
	QUADRUPLE STRENGTH continuous I Infusion strength	Prescribed amount
	1 mL/hour = 200 milliunit/kg/hour	10 unit/kg argipressin and make up to 50 mL
	Draw up 0.5 mL/kg argipressin (10 uni volume of 50 mL with a final concentr Infusing at a rate of 1 mL/hour = 200 i	-
Administration		central line. Use with caution via a peripheral line.
Monitoring	Continuous heart rate, ECG and blood	· · ·
		Illy monitored and may require the weaning of other
	vasopressors.	
	Assess urine output and peripheral pe	rfusion frequently.
	Fluid balance and serum sodium.	nd outroupoption
	Observe IV site closely for blanching a For diabetes insipidus:	nu extravasation.
	-	range 0.5 to 1 milliunit/kg/hour aiming for:
	 urine output 2–4 mL/kg/hour, 	
	 neutral fluid balance, 	
	 maintain plasma sodium 145–150 	mmol/L
	-	of 20–30 minutes, so a change in infusion rate is reflected 1 hour
	later.	
Contraindications	Hypersensitivity to argipressin.	
		sin (Vasopressin) Page 2 of 6

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Argipressin (Vasopressin)

Newborn Use Only

Precautions	Use in hypotension:
Precautions	Argipressin causes water retention and hyponatraemia.
	May cause ischaemia related to infusion site.
	Acute ECG or biochemical evidence of myocardial ischaemia.
	Previously documented chronic and/or severe liver dysfunction (INR > 2, direct bilirubin > 50
	micromol/L) or clinical evidence of portal hypertension.
	Documented or high suspicion of mesenteric ischaemia.
	Use in diabetes insipidus:
	The mainstay of initial therapy is accurate fluid and electrolyte management. ADH administration
	should only be considered after a reasonable period of observation establishes that DI is persistent (at
	least 4–6 hours, but preferably longer in acute situations). Early or over vigorous ADH administration
	may provoke cerebral oedema,
	Prior to starting the infusion, it is advisable to allow the patient to drift into a slightly negative fluid
	balance. This can be easily achieved by not replacing all the previous hour(s) urine output. Once the
	argipressin infusion has commenced, continue the fluid regimen of replacement of previous hour's
	losses plus insensible losses.
	Use in gastrointestinal bleeding: There are few reports of argipressin use for gastrointestinal bleeding
	in newborns. The dose regimen is unclear and other agents may be more effective.
Drug Interactions	Noradrenaline (norepinephrine) and heparin—when used with argipressin may decrease the
0	antidiuretic effect of argipressin.
Adverse	Causes water retention and hyponatraemia. Early or over vigorous administration may provoke
Reactions	cerebral oedema,
	Cardiac complications include coronary ischaemia, myocardial infarction, ventricular arrhythmias
	(ventricular tachycardia and asystole) and severe hypertension. Other reported adverse effects include
	severe GI ischaemia leading to bowel necrosis, hyponatraemia, anaphylaxis, bronchospasm, urticaria,
	angioedema, rashes, venous thrombosis, local irritation at injection site and peripheral vasoconstrictio
	leading to cutaneous gangrene. ^{1,2}
Compatibility	Fluids: Glucose 5%, sodium chloride 0.9%
	Vicita (17.19). Civing other drugs via Vicita may change the infusion rate of arginressin. A dedicated lin
	Y-site (17,18): Giving other drugs via Y-site may change the infusion rate of argipressin. A dedicated line is preferred.
	Aciclovir, amikacin, amiodarone, aminophylline, amphotericin B liposome, atenolol, atropine,
	azithromycin, calcium chloride, calcium gluconate, cefazolin, cefepime, cefotaxime, ceftazidime,
	ceftriaxone, chloramphenicol, ciprofloxacin, dexamethasone, dexmedetomidine, digoxin, dobutamine,
	dopamine, epinephrine, epoetin alfa, fentanyl, fluconazole, folic acid, ganciclovir, gentamicin,
	glycopyrrolate, heparin, hydrocortisone, imipenem-cilastatin, lidocaine, linezolid, magnesium sulfate,
	meropenem, metronidazole, midazolam, milrinone, morphine sulfate, naloxone, nitroprusside,
	octreotide, pantoprazole, penicillin G, phenobarbital, piperacillin-tazobactam, potassium chloride,
	propranolol, protamine, pyridoxine, ranitidine, remifentanil, rocuronium, sodium acetate, sodium
	bicarbonate, theophylline, thiamine, ticarcillin-clavulanate, tobramycin, urokinase, vancomycin,
	vecuronium, zidovudine.
	Variable Y-site compatibility (use caution): Ampicillin, furosemide, insulin (regular), sulfamethoxazole-
	trimethoprim
Incompatibility	Fluids: No information.
	Y-site (17,18): Diazepam, diazoxide, indometacin, phenytoin.
Stability	Diluted solution: Discard remainder after use.
	Change infusion solution every 24 hours
Storage	Ampoule: Store below 25°C.
Excipients	Acetic acid and water for injection.
Special	Administration via a central line is preferred as extravasation may cause tissue necrosis.
Special	Autoristication via a central line is preferred as extravasation may cause lissue necrosis.

Newborn Use Only

Evidence	Efficacy:
LVIACIICC	Newborns with hypotension: A pilot trial in preterm infants born < 30 weeks gestation with refractory
	hypotension (n = 20) of argipressin (0.01 units/kg/h to maximum 0.04 units/kg/hour) versus dopamine
	(5 to 20 microg/kg/min) reported similar increases in blood pressure. Infants receiving argipressin
	received fewer doses of surfactant, had lower PaCO ₂ values and were less tachycardic. No difference in
	clinical outcome was reported. ³ The role of argipressin for hypotension in newborns is unclear (LOE II,
	GOR D).
	Children with refractory hypotension: A review ⁴ of argipressin/terlipressin as rescue therapy in
	children with catecholamine-resistant shock or cardio-circulatory arrest found 31 reports (428 patients)
	including a single RCT of argipressin in children with vasodilatory shock. Infants with vasodilatory shock
	were randomised to low-dose argipressin (0.0005 to 0.002 units/kg/minute = 0.03 to 0.12
	units/kg/hour) or placebo in addition to open-label vasoactive agents. There was no difference in time
	to vasoactive-free haemodynamic stability, a trend to increasing mortality and no benefit in respect to
	organ-failure-free days from use of argipressin. In observational studies, argipressin from 0.00002
	unit/kg/minute to 0.002 unit/kg/minute (0.0012 to 0.12 units/kg/hour) increased blood pressure, urine
	output, and decreased serum lactate. In most reports, argipressin and terlipressin led to a reduction of
	catecholamine dose. However, mortality remained high (188/428; 43.9%) despite the use of argipressin
	or terlipressin. ⁴ The role of argipressin for refractory hypotension in newborns is unclear. (LOE IV in
	newborns, GOR D) American College of Critical Care Medicine Clinical Practice Parameters for
	Hemodynamic Support of Pediatric and Neonatal Septic Shock included argipressin as an option for
	management of vasodilatory shock (hypotension with low vascular resistance) refractory to
	catecholamines including noradrenaline (norepinephrine). ⁵
	Use in pulmonary hypertension: In a case series of 10 newborn infants with severe persistent
	pulmonary hypertension of the newborn on nitric oxide, argipressin 0.0002 ± 0.0002 U/kg/minute
	(0.012 ± 0.012 units/kg/hour) was associated with an improvement in oxygenation index, peak effect 6
	hours after initiation, and a reduction in inhaled nitric oxide dose, improvement in blood pressure and
	urine output ($p < 0.05$), without drop in the serum sodium level or worsening in serum lactate level. ⁶
	The role of argipressin for pulmonary hypertension in newborns is unclear. (LOE IV, GOR D)
	Use in congenital diaphragmatic hernia : In a case series of 13 infants with CDH treated with argipressin
	for refractory hypotension, argipressin (range 0.0001–0.002 units/kg/min) increased mean arterial
	pressure and decreased pulmonary/systemic pressure ratio, heart rate, and fraction of inspired oxygen.
	In 6 of 13 patients, extracorporeal membrane oxygenation therapy was no longer indicated after
	treatment. The role of argipressin in newborns with CDH and refractory hypotension is unclear. (LOE IV,
	GOR D) ^{7,8}
	Infants with diabetes insipidus: the drug of choice for the treatment of diabetes insipidus is
	desmopressin (dDAVP), a synthetic analog of arginine vasopressin, but with a 2,000- to 3,000-fold lower
	vasopressor effect. An aqueous argipressin infusion should be considered in the initial management of
	post-surgical or post-traumatic DI. It has the advantage of having a relatively short half-life so that the
	dose can be titrated against the urine output. It has the disadvantage of requiring significant
	observation and adjustment of the infusion rate depending upon the hourly urine output. Management
	should be in PICU where monitoring and expertise are readily available. In children with DI secondary
	to brain injury, an initial infusion of aqueous argipressin 0.00025 to 0.001 units/kg/hour (0.25 to 1.0
	milliunits/kg/hour titrated to urine output 2–3 ml/kg/hour, urine specific gravity 1.010–1.020 and
	serum sodium 140–145 mEq/L, was effective. ⁸ Argipressin infusion can be used in infants with diabetes
	insipidus where dDAVP is not though appropriate. (LOE IV GOR C)
	Infants with gastrointestinal bleeding: There are only case reports of argipressin being used for
	gastrointestinal bleeding. ^{9,10,11} Argipressin use was reported in 15 children with severe oesophageal
	variceal bleeding and 2 with peptic ulcer bleeding with control from use of argipressin alone in 9 of 17
	episodes. Argipressin was commenced at 0.1 to 0.2 units/minute with titration over 2 hours to control
	bleeding. The maximum delivered dosage ranged from 0.004 to 0.04 units/kg/min (0.24 to 2.4
	units/kg/hour). Control of bleeding did not improve with high dose argipressin and there was a
	significantly greater incidence of complications in those patients receiving ≥ 0.01 units/kg/min (0.6
	units/kg/hour). Complications included electrolyte abnormalities (Na, K, Cl or Ca) in 10 infants, fluid
	overload (4 infants), hypertension (4 infants) and cardiac dysrhythmias (2 infants). ¹⁰

Argipressin (Vasopressin)

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	Meta-analysis of studies in adults with acute variceal bleeds found that although vasopressor agents
	reduced mortality and achieved haemostasis, trials of argipressin were not conclusive and argipressin was less effective for haemostasis compared to octreotide or somatostatin. ^{12, 13} Argipressin may be used in combination with nitroglycerin so as to balance its vasoconstrictive effect. Major side effects associated with the use of argipressin include myocardial ischaemia, life threatening arrythmias, mesenteric ischaemia and limb vasoconstriction or ischaemia. Other minor complications include water retention with sodium depletion, benign arrhythmia and acrocyanosis. Monitor cardiac rate and rhythm, and watch for peripheral ischaemia. Terlipressin may be preferred over argipressin as it has the convenience of bolus administration, decreased cardiotoxicity and its ability to control up to 79% of variceal hemorrhage. ^{14,15} (GOR D) Pharmacokinetics: The pharmacology of argipressin in newborns and children has not been sufficiently investigated and data on potential short and long-term adverse effects are still lacking. ^{14,16} Half-life approximately 30 minutes, clinical duration of action 2–3 hours. Safety:
	Safety data of argipressin in paediatric patients is limited. ^{14,16} Potent vasoconstrictor action may cause
	ischaemia. Complications are more common when argipressin is co-administered with moderate to
	high doses of noradrenaline (norepinephrine). Hyponatraemia occurs frequently during argipressin infusion requiring close monitoring of serum sodium and water intake. ^{1,2} For control of gastrointestinal
	haemorrhage, argipressin was associated with electrolyte abnormalities (Na, K, Cl or Ca) in 10 infants,
	fluid overload (4 infants), hypertension (4 infants) and cardiac dysrhythmias (2 infants), particularly at
Practice points	doses ≥ 0.01 units/kg/minute. ¹⁰
References	1. Baldasso E, Garcia PC, Piva JP, Branco RG, Tasker RC. Pilot safety study of low-dose vasopressin in
	 Inon-septic critically ill children. Intensive care medicine. 2009;35:355-9. Davalos MC, Barrett R, Seshadri S, Walters HL, 3rd, Delius RE, Zidan M, Mastropietro CW. Hyponatremia during arginine vasopressin therapy in children following cardiac surgery. Pediatric critical care medicine : a journal of the Society of Critical Care Medicine and the World Federation of Pediatric Intensive and Critical Care Societies. 2013;14:290-7. Rios DR, Kaiser JR. Vasopressin versus dopamine for treatment of hypotension in extremely low birth weight infants: a randomized, blinded pilot study. The Journal of pediatrics. 2015;166:850-5. Meyer S, McGuire W, Gottschling S, Mohammed Shamdeen G, Gortner L. The role of vasopressin and terlipressin in catecholamine-resistant shock and cardio-circulatory arrest in children: review of the literature. Wiener medizinische Wochenschrift. 2011;161:192-203. Davis AL, Carcillo JA, Aneja RK, Deymann AJ, Lin JC, Nguyen TC, Okhuysen-Cawley RS, Relvas MS, Rozenfeld RA, Skippen PW, Stojadinovic BJ, Williams EA, Yeh TS, Balamuth F, Brierley J, de Caen AR, Cheifetz IM, Choong K, Conway E, Jr., Cornell T, Doctor A, Dugas MA, Feldman JD, Fitzgerald JC, Flori HR, Fortenberry JD, Graciano AL, Greenwald BM, Hall MW, Han YY, Hernan LJ, Irazuzta JE, Iselin E, van der Jagt EW, Jeffries HE, Kache S, Katyal C, Kissoon NT, Kon AA, Kutko MC, MacLaren G, Maul T, Mehta R, Odetola F, Parbuoni K, Paul R, Peters MJ, Ranjit S, Reuter-Rice KE, Schnitzler EJ, Scott HF, Torres A, Jr., Weingarten-Abrams J, Weiss SL, Zimmerman JJ, Zuckerberg AL. American College of Critical Care Medicine. 2017;45:1061-93. Mohamed A, Nasef N, Shah V, McNamara PJ. Vasopressin as a rescue therapy for refractory pulmonary hypertension in neonates: case series. Pediatric Critical Care Medicine. 2014;15:148-54. Acker SN, Kinsella JP, Abman SH, Gien J. Vasopressin improves hemodynamic status in infants with congenital diaphragmatic hernia. Journal of Pediatrics. 2014;165:53-8

11. Liebman WM. Diagnosis and management of upper gastrointestinal hemorrhage in children. Pediatr Ann. 1976;5:690-9.
12. D'Amico G, Pagliaro L, Pietrosi G, Tarantino I. Emergency sclerotherapy versus vasoactive drugs for bleeding oesophageal varices in cirrhotic patients. The Cochrane database of systematic reviews. 2010:CD002233.
13. Wells M, Chande N, Adams P, Beaton M, Levstik M, Boyce E, Mrkobrada M. Meta-analysis: vasoactive medications for the management of acute variceal bleeds. Aliment Pharmacol Ther. 2012;35:1267-78.
14. Agrawal A, Singh VK, Varma A, Sharma R. Therapeutic applications of vasopressin in pediatric patients. Indian pediatrics. 2012;49:297-305.
15. Arora NK, Ganguly S, Mathur P, Ahuja A, Patwari A. Upper gastrointestinal bleeding: Etiology and management. Indian Journal of Pediatrics. 2002;69:155-68.
16. Biban P, Gaffuri M. Vasopressin and terlipressin in neonates and children with refractory septic shock. Current drug metabolism. 2013;14:186-92.
 Micromedex solutions. Accessed on 13 April 2021. Australia Ijectable drugs handbook. 8th edition. Accessed on 13 April 2021.

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