Queensland	(Affix identifica	tion label here)
Government	URN:	
	Family name:	
* PAEDIATRIC	Given name(s):	
Sepsis Pathway	Address:	
Facility:	Date of birth:	Sex: M F I
Clinical pathways never replace clinical judgement. U		an 16 years.
16–18 year olds may use the adult or paediatric sepsi		
Sepsis is infection with organ dysfunction. S	Sepsis is a MEDICAL EMERGENC	·Y. ■
SCREEN AND RECOGNISE		
Screening initiated: DD / MM / YY HH : MM (2	24hr)	
Could it be sepsis?		
☐ Signs of infection or history and evidence of fever of	or hypothermia	
PLUS ANY of the following		
	red behaviour or reduced level of cons	ciousness
	younger than 3 months sis admission within the last 30 days	
	riginal or Torres Strait Islander person	
*For Oncology patients refer to 'Management of Susp	pected Neutropenic Sepsis Pathway (S	W796)'
	▼ YES	
Document full set of observations in CEWT include	ding blood pressure and AVPU	
	↓ THEN	
Does the patient have ANY features of severe illne		
Severe respiratory distress, tachypnoea or apnoea		red AVPU
Severe tachycardia (CEWT heart rate score 3)☐ Hypotension (CEWT blood pressure score ≥2)		r skin perfusion or cold extremitie tate ≥2mmol/L (if known)
Other laboratory features of severe illness (if known		ate =21111101/E (II KIIOWII)
Low platelets	•	CRP
These laboratory tests are not mandatory		
YES		₩ NO
	Do you still sus	spect sepsis?
	▼ YES	↓ NO
Patient is highly likely to HAVE sepsis or	Patient MAY have sepsis	Patient UNLIKELY to
septic shock	Targeted history and examination	have sepsis now
Immediate senior medical review or call Retrieval Services Queensland (RSQ)	Obtain senior medical review or consider calling RSQ	Reassess and escalate as indicated
1300 799 127	Consider calling NOW	do maioatoa
Immediate monitoring in close observation area		
↓ THEN	▼ THEN	→ THEN
Senior medical review attended: HH: MM (24hr)		☐ Give Paediatric Sepsis
(ZTIII)		Checklist to parent or
Does the senior clinician think sepsis is likely?		carer (tear off back page)
Yes – sepsis with shock Yes – sepsis without	t shock OR Unlikely sepsis	



VES

Sepsis has been diagnosed by a senior medical doctor Start resuscitation and treatment for sepsis NOW (next page) Escalate to MET, PICU, ICU or RSQ 1300 799 127

Signatur	Signature Log Every person documenting in this clinical pathway must supply a sample of their initials and signature below						
Initials	Signature	Print name	Role				





(Affix identification lab	el here))		
URN:				
Family name:				
Given name(s):				
Address:				
Date of birth:	Sex:	M	F	

PAEDIATRIC	Given n	ame(s):		
Sepsis Pathway	Address	s:		
	Date of		Sex:	□M □F □I
ACUTE RESUSCITATION TREATMENT BUNDLI		5.1.4.1.		
	_	1		
Complete actions 1–6 within: 1 hour of recognition of shock or where there is high lik 3 hours to administer antimicrobials where there is less all relevant microbiological samples according to suspen	likelihoo	d of organ dysfunct	tion and sepsis. Prior	ritise timely collection of
1. Notify the Senior Medical Officer or RSQ for revie				☐ Consultant notified
 Refer to Consultant Paediatrician Notify Nursing Team Leader or Senior Nurse on call 	ı			
2. Monitor oxygen saturations and maintain 94% or		,		Oxygen saturations
	J			maintained
 3. IV or intraosseous access and blood culture Obtain intraosseous access after two failed attempts Take blood culture (2–6mL) prior to antibiotics Take lactate, VBG and blood glucose level Take FBC, CRP, Chem20, coagulation studies and second second			te cultures	☐ Blood cultures obtained ☐ Lactate taken
4. Commence appropriate IV or intraosseous antibio • Check allergies and presence of MRSA risk factors • Prescribe antibiotics according to the guidelines in T • Give intramuscular antibiotics if failed IV or intraosse	ac sultures	Antibiotic commenced		
Suspected source of infection: Sepsis where meningitis possible OR bacterial me Sepsis (source unknown, but bacterial meningitis of Febrile neutropenia (refer to 'Management of Suspon Neutropenic Sepsis Pathway [SW796]') Toxic Shock Syndrome	excluded)	Cellulitis, s	keletal or soft tissue nous access device	
5. Commence fluid resuscitation • Administer rapid isotonic fluid bolus IV or intraossed • Consider repeating up to 40–60mL/kg isotonic fluid • Observe for signs of fluid overload (hepatomegaly) • If hypoglycaemic, then give 2mL/kg glucose 10% • Consider second IV or intraosseous access		-	ponse	Fluid bolus commenced
6. Consider inotropic support and prepare early • Consider IV or intraosseous adrenaline infusion if no status after 40–60mL/kg of fluid • Prepare adrenaline (epinephrine) infusion by diluting chloride 0.9% or glucose 5%; commence infusion are infusion chart for equivalent mL/hr for child's weight, • Call PICU, ICU or RSQ 1300 799 127	g 1mg (1r t 0.1–0.5	mL of 1:1000) to 50	mL with sodium	☐ Inotrope considered
BEREAVEMENT				
Refer to CHQ Bereavement Service (1800 080 316) o Offer for family to spend time with child after death Ensure sepsis is documented on the death certificate		Inform Sepsis Car	e Coordinator of sep	
REASSESS				
Does the patient have ANY persistant signs of seps ☐ Tachypnoea (CEWT respiratory score ≥2) ☐ Tachycardia (CEWT heart rate score ≥2) ☐ Hypotension (CEWT blood pressure score ≥2) ☐ Lactate ≥2mmol/L	☐ Alte	ered AVPU	apillary refill ≥3 seco	bundle? nds or cold extremities
▼ YES			₩ NO	
Deteriorating or persistent signs of sepsis Escalate via local policy Notify Sepior Medical Officer and call PICL ICL or		Resolving signs • De-escalate as provided to review		tient for signs of

- Notify Senior Medic RSQ 1300 799 127
- Follow Sepsis Management Plan (next page)
- deterioration
- Follow Sepsis Management Plan (next page)





	(Affix identification I	abel her	e)		
URN:					
Family name:					
Given name(s):					
Address:					
Date of birth:		Sex:	M	F	

Sepsis Management Plan

DETERIORATING OR PERSISTENT SIGNS OF SEPSIS Level of care: Critical

RESOLVING SIGNS OF SEPSIS Level of care: Inpatient

COMMUNICATE

Discussions with family to include:

- Explanation of sepsis
- Parent and carer information sheet (tear off Information for Parents page at back)
- Family questions
- · Goals of care
- Social work, welfare support and other allied health services
- Indigenous Health Liaison Officers (IHLO)
- Interpreter supports

Discussions with family to include:

- Explanation of sepsis
- Parent and carer information sheet (tear off Information for Parents page at back)

Heart rate

- Family questions
- · Social work and welfare support
- Indigenous Health Liaison Officers (IHLO)
- Interpreter supports

MONITOR

Continuous: • SpO₂ Heart rate Respiratory rate Arterial blood pressure (if required) 15 minutes: AVPU · Non-invasive blood pressure · Capillary refill time 60 minutes:

Temperature (until resolved)

- Strict fluid balance
- Urine output

4 hourly:

- Lactate
- Blood sugar level
- Venous blood gas Temperature (once resolved)

Continuous:

- SpO₂
- Respiratory rate

60 minutes:

- Blood pressure
 - Temperature (until resolved) Urine output
- Strict fluid balance

4 hourly:

- AVPU
- Temperature (once resolved)

REASSESS

Patients may move between streams according to clinical response. Patients who are deteriorating or have persistent signs of sepsis require more frequent monitoring. Obtain senior medical officer advice on changing sepsis management plan stream.

Clinically reassess after interventions, monitored vital sign changes or every 60 minutes as a minimum:

- Tachypnoea (CEWT respiratory score ≥2)
- Tachycardia (CEWT heart rate score ≥2)
- Hypotension (CEWT blood pressure score ≥2)
- Altered AVPU
- Poor skin perfusion; capillary refill ≥3 seconds or cold extremities
- Urine output less than 1mL/kg/hr
- Lactate ≥2mmol/L (4 hourly)

If deteriorating or persistent signs of sepsis are still present:

- Escalate via local policy
- · Notify Senior Medical Officer and call PICU, ICU or RSQ 1300 799 127

Clinically reassess after interventions, monitored vital sign changes or every 60 minutes as a minimum:

- Tachypnoea (CEWT respiratory score ≤1)
- Tachycardia (CEWT heart rate score ≤1)
- Hypotension (CEWT blood pressure score ≤1)
- Improving AVPU
- Improved skin perfusion; capillary refill <3 seconds or warm extremities
- Urine output greater than or equal to 1mL/kg/hr

After 12 hours, if no intervention reassess every 4 hours

After 24 hours, if no intervention follow local de-escalation policy

INVESTIGATE

Collect relevant outst	anding microbiology samples:
Urine	☐ Blood cultures
☐ CSF (when stable)	Other relevant sources
Stool	(e.g. surgical specimens following
Respiratory	source control)

Collect relevant outstanding microbiology samples:					
Urine	☐ Blood cultures				
☐ CSF	Other relevant sources				
Stool	(e.g. surgical specimens following				
Respiratory	source control)				





(Affix identification lab	oel here)		
URN:				
Family name:				
Given name(s):				
Address:				
Date of birth:	Sex:	M	F	

Sepsis Management Plan (continued)

DETERIORATING OR PERSISTENT SIGNS OF SEPSIS

Level of care: Critical

RESOLVING SIGNS OF SEPSIS Level of care: Inpatient

ANTIMICROBIAL OPTIMISATION

- · Reconsider source and need for source control
- Review microbiology results in consultation with laboratory
- Review appropriateness of antimicrobial cover and consider additional risk factors
- Consider ID expert guidance as per local referral pathway.
 QCH oncall service available Ph: 07 3068 1111
- Ensure Therapeutic Drug Monitoring where appropriate
- Review microbiology results in consultation with laboratory
- Review appropriateness of antimicrobials and consider de-escalation, targeting or cessation

DOCUMENT

Antimicrobial Stewardship:

- Document confirmed or suspected source of infection in health record
- Document plan to continue, change or cease antimicrobials
- Consider longer-term central IV access if required
- Review antimicrobial allergy history if applicable and refer to ID or immunology for assessment

Other documentation:

- Document sepsis in health record
- Document when patient is seen by Sepsis Care Coordinator
- Document variations to assist future optimisation of the pathway

Antimicrobial Stewardship:

- Document confirmed or suspected source of infection in health record
- Document plan to continue, change or cease antimicrobials
- Review antimicrobial allergy history if applicable and refer to ID or immunology for assessment

Other documentation:

- Document sepsis in health record
- Document when patient is seen by Sepsis Care Coordinator
- Document variations to assist future optimisation of the pathway

HANDOVER AND DISCHARGE

Handover to ward:

- Document psychosocial support required in health record (e.g. social work, IHLO, interpreter)
- Document clinicians involved in handovers in the health record
- Involve parents and carers in handover and provide information
- Handover to also include provisional sepsis diagnosis, comorbidities, management plan for medicines and medical conditions

Discharge planning:

- · Give resources to family
- · Identify GP and document in health record
- · Discuss supports required with family and GP
- Consider nurse navigator, hospital in the home or other referral
- · Give local patient experience survey to family

RESOURCES

Clinical:

- Queensland Paediatric Sepsis Program clinical resources for health professionals
- Children's Resuscitation Emergency Drug Dosage Guide (CREDD). Consider using CREDD for weight adjusted dosing measurements
- · National Sepsis Clinical Care Standard, including discharge planning guide, GP letter template and other resources
- Surviving Sepsis Campaign Guidelines January 2020

Family:

- · Queensland Paediatric Sepsis Program family resources
- Find an Aboriginal Community Controlled Health Organisation (ACCHO) near you

Bereavement:

· Children's Health Queensland Bereavement Service

Table 1: PAEDIATRIC Empiric Prescribing Guidelines for Community Acquired Sepsis

• Where appropriate, screen patient for additional risk factors such as vaccination status, recent travel, multi-drug resistant organisms, immunocompromise, animal exposure, antenatal exposure or water-exposed soft tissue or skeletal infections. Contact paediatric ID specialist or microbiologist for advice

• Antimicrobial should be assessed with culture results and ID or microbiology at 24 to 48 hours of antimicrobial therapy

Suspected se infection	ource of	Initial, empirical antibiotic regimen	Immediate severe type hypersensitivity (e.g. anaphylaxis) to first line antimicrobial)**
Febrile neutropenia		Oncology patients: please refer to 'Management of Suspected Neutropenic Non-oncology patients: please manage as per the 'Paediatric Sepsis Pathw	Sepsis Paediatric Pathway' ay' (below)
Septic shock inotrop		ALL sources: ADD Gentamicin* PLUS Vancomycin to empirical regime where not already recommended EXCEPT in North Queensland if risk factors for melioidosis (wet season or flooding) REPLACE Cefotaxime with Meropenem and ADD Vancomycin where not already recommended	
	£812	Neonates and infants up to 2 months of age	
Sepsis where		cefOTAXIME IV PLUS Ampicillin (OR Amoxicillin) IV	cefOTAXIME IV
Meningitis possible OR		Infants and children older than 2 months of age • cefOTAXIME (OR cefTRIAXONE) IV	ciPROFLOXAcin IV PLUS Vancomycin IV
Bacterial	6	If Gram positive cocci in CSF	- CII NOI LOAACIITV / LOO VAIICOITIYCIIT IV
Meningitis		cefOTAXIME (OR cefTRIAXONE) IV PLUS Vancomycin IV	ciPROFLOXAcin IV PLUS Vancomycin IV
		All ages – if encephalitis suspected: ADD Aciclovir IV	,
		Neonates and infants up to 2 months of age	
		Ampicillin (OR Amoxicillin) IV PLUS Gentamicin* IV If at risk of nmMRSA	cefOTAXIME IV
Sepsis (source		Ampicillin (OR Amoxicillin) IV PLUS Gentamicin* IV PLUS linCOMYCIN (OR Clindamycin) IV	cefOTAXIME IV PLUS linCOMYCIN (OR Clindamycin) IV
unknown, but bacterial		Infants and children older than 2 months of age	
meningitis		cefOTAXIME (OR cefTRIAXONE) IV	ciPROFLOXAcin IV PLUS Vancomycin IV
excluded)		If at risk of nmMRSA • cefOTAXIME (OR cefTRIAXONE) IV PLUS linCOMYCIN	
		(OR Clindamycin) IV	• ciPROFLOXAcin IV PLUS Lincomycin (OR Clindamycin) IV
		If at risk of multi-resistant MRSA	
		cefOTAXIME (OR cefTRIAXONE) IV PLUS Vancomycin IV	ciPROFLOXAcin IV PLUS Vancomycin IV
		Neonates and infants up to 2 months of age • Ampicillin (OR Amoxicillin) IV PLUS Gentamicin* IV	cefOTAXIME IV
		Infants and children more than 2 months of age	- CEIOTAXIIVIE IV
		Benzylpenicillin IV	cefOTAXIME (OR cefTRIAXONE) IV
Bacterial	ALA	Severe pneumonia (requiring PICU admission)	,
Pneumonia	A STATE OF THE STA	All ages: cefOTAXIME IV	ciPROFLOXAcin IV PLUS Vancomycin IV
(Community acquired)		If empyema OR S. aureus (including nmMRSA) pneumonia suspected	PROFILOVA : IV RUIGI : : (OR OF death : 1) IV
, ,		cefOTAXIME (OR cefTRIAXONE) IV PLUS linCOMYCIN (OR Clindamycin) IV	ciPROFLOXAcin IV PLUS Lincomycin (OR Clindamycin) IV
		If life threatening pneumonia/empyema OR multi-resistant MRSA susp	ected
		cefOTAXIME (OR cefTRIAXONE) IV PLUS linCOMYCIN (OR Clindamycin) IV PLUS Vancomycin IV	ciPROFLOXAcin IV PLUS linCOMYCIN (OR Clindamycin) IV PLUS Vancomycin IV
lutus abdaminal		Ampicillin (OR Amoxicillin) IV PLUS Gentamicin* IV PLUS	cefOTAXIME (OR cefTRIAXONE) IV PLUS Metronidazole IV
Intra-abdominal source		Metronidazole IV	, ,
Urinary source	6	Ampicillin (OR Amoxicillin) IV PLUS Gentamicin* IV	Gentamicin* IV
		All ages and Hib immune, with skeletal infection, periorbital cellulitis w	
		• Flucloxacillin IV	• linCOMYCIN (OR Clindamycin) IV
		If younger than 5 years of age and NOT Hib immune, with skeletal infection of the ceforaxime IV	• ciPROFLOXAcin IV PLUS linCOMYCIN (OR Clindamycin) IV
		If at risk of nmMRSA	CIT NOT ECONOMITY TECO IIITOOVITOIN (ON OIIITAAITYCIII) IV
		ADD linCOMYCIN (OR Clindamycin) IV to appropriate therapy as above	• ciPROFLOXAcin IV PLUS linCOMYCIN (OR Clindamycin) IV
Severe cellulitis	1666	If at risk of multi-resistant MRSA	
or skeletal or soft tissue	12. 4 数	ADD Vancomycin IV to appropriate therapy as above	ciPROFLOXAcin IV PLUS Vancomycin IV
infection	1	Suspected necrotising fasciitis • cefOTAXIME IV PLUS Vancomycin IV PLUS linCOMYCIN	Meropenem IV PLUS Vancomycin IV PLUS linCOMYCIN
		(OR Clindamycin) IV PLUS consider IVIG 2g/kg	(OR Clindamycin) IV
		If external wound/inoculation associated with necrotising fasciitis	
		Meropenem IV PLUS Vancomycin IV PLUS linCOMYCIN (OR Clindamycin) IV	Meropenem IV PLUS Vancomycin IV PLUS linCOMYCIN (OR Clindamycin) IV
		Open fractures with severe tissue damage and contamination	(Ort CillidatifyCill) IV
		Piperacillin/Tazobactam IV	ciPROFLOXAcin IV PLUS linCOMYCIN (OR Clindamycin) IV
Central	Januarina	Consider removal of device	
venous access device source		Piperacillin/Tazobactam IV PLUS Vancomycin IV	cefTAZIDIME IV PLUS Vancomycin IV
Toxic shock s	•	ceFAZolin IV PLUS Lincomycin (OR Clindamycin) IV PLUS Vancomycin PLUS consider IVIG 2g/kg	Vancomycin IV PLUS Lincomycin (OR Clindamycin) IV
* If Pseudomonas ac	eruginosa is cult	ured, seek ID advice on appropriate directed therapy.	

If Pseudomonas aeruginosa is cultured, seek ID advice on appropriate directed therapy

^{**} The recommendations provided for immediate type hypersensitivity in this table are for an initial dose only in the emergency treatment of sepsis. Please contact a paediatric ID specialist for any subsequent dosing.

For more information, and **ongoing** prescribing information please refer to 'CHQ Paediatric Antibiocard: Empirical Antibiotic Guidelines' and the 'CHQ guideline: Empiric antibiotic guidelines for Paediatric Intensive care unit (PICU)'.

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Table 2: Antimicrobial Dose Recommendations for Sepsis by Age

- Term neonates >36 weeks post-menstrual age to adolescents.
- For premature neonates, refer to NeoMedQ, ANMF or Neofax; available via CKN or QCH Guidelines.

Antimicrobial	Dose recommend	dation by age (normal renal function)
Aciclovir IV	Birth to 3 months of age	• 20mg/kg IV 8 hourly
	Older than 3 months of age and less than 12 years of age	• 500mg/m² (maximum 1g) IV 8 hourly
	12 years of age and older	• 10mg/kg (maximum 1g) IV 8 hourly
Ampicillin (OR Amoxicillin) IV	Neonates	Week 1 of life: 50mg/kg IV 12 hourly Week 2–4 of life: 50mg/kg IV 8 hourly Meningitis: 100mg/kg/dose (on ID advice)
	Older than 1 month of age	• 50mg/kg (maximum 2g) IV 6 hourly
Benzylpenicillin IV	Neonates	Week 1 of life: 60mg/kg IV 12 hourly Week 2–4 of life: 60mg/kg IV 8 hourly
	Older than 1 month of age	60mg/kg (maximum 2.4g) IV 6 hourly
cefaZOLin IV	Neonates	Seek ID/specialist advice
	Older than 1 month of age	• 50mg/kg IV 8 hourly (maximum 2g)
cefOTAXIME IV or IM* for neonate	Neonates	Week 1 of life: 50mg/kg IV/IM 8 hourly Week 2–4 of life: 50mg/kg IV/IM 6 hourly
	Older than 1 month of age	• 50mg/kg (maximum 2g) IV/IM 6 hourly
cefTRIAXONE IV or IM*	Neonates	cefTRIAXONE contra-indicated (risk of kernicterus) – use cefOTAXIME
	Older than 1 month of age	• 50mg/kg (maximum 2g) IV/IM 12 hourly
cefTAZIDIME IV	Neonates	• 50mg/kg IV 12 hourly
	Older than 1 month of age	• 50mg/kg (maximum 2g) IV 8 hourly
ciPROFLOXAcin IV	Neonates	Seek ID/specialist advice
	Older than 1 month of age	• 10mg/kg (maximum 400mg) IV 8 hourly
Clindamycin IV	Neonates	• 7mg/kg IV 8 hourly
	Older than 1 month of age	• 10mg/kg (maximum 600mg) IV 6 hourly
Flucloxacillin IV	Neonates	Week 1 of life: 50mg/kg IV 12 hourly Week 2–3 of life: 50mg/kg IV 8 hourly Week 4 of life: 50mg/kg IV 6 hourly
	Older than 1 month of age	• 50mg/kg (maximum 2g) IV 6 hourly
Gentamicin IV	Neonates	Week 1–4 of life: 5mg/kg IV once daily
	Older than 1 month and less than 10 years of age	7.5mg/kg IV once daily (maximum 320mg)
	10 years of age and older	7mg/kg IV once daily (maximum 700mg)
	ALL ages: perform Therapeutic Drug Monitoring (TDM) – d check trough pre-2nd dose)	lose based on Adjusted body weight (neonates or renal impairment,
IinCOMYCIN IV	Neonates	No neonatal dosing recommendation for linCOMYCIN – use Clindamycin IV
	Older than 1 month of age	• 15mg/kg (maximum 1.2g) IV 8 hourly
Meropenem IV	All ages	40mg/kg (maximum 2g) IV 8 hourly
Metronidazole IV	Neonates	15mg/kg IV load, then 7.5mg/kg IV 8 hourly
	Older than 1 month of age	• 7.5mg/kg (maximum 500mg) IV 8 hourly
Piperacillin/ Tazobactam IV	Neonates	Week 1 of life: 100mg/kg IV 12 hourly Week 2–4 of life: 100mg/kg IV 8 hourly
(dose based on piperacillin component)	Older than 1 month of age	• 100mg/kg (maximum 4g) IV 6 hourly
Vancomycin IV	Neonates	Week 1 of life: 15mg/kg IV 12 hourly Week 2–4 of life: 15mg/kg IV 8 hourly
	Older than 1 month of age	15mg/kg (maximum 750mg) IV 6 hourly
	ALL ages: perform TDM – dose based on Actual body weig	ght
*Prioritise IV/IO access and a		ation in sepsis may result in subtherapeutic doses due to reduced muscular perfusion

- 1. Antibiotic Therapeutic Guidelines (Oct 2021). Therapeutic Guidelines Committee, North Melbourne, Victoria. Available on CKN
- AMH Children's Dosing Companion [Online]. Adelaide: Australian Medicines Handbook Pty Ltd; 2020. Last updated July 2022. Available on CKN.
 The Australasian Neonatal Medicines Formulary (ANMF) [Online]. Accessed 6 Oct 2022. Last updated 11/10/22. Available on CKN.
 Neofax 2022. Micromedex Healthcare solutions. Truven Health Analytics. US. Available on CKN.

- NeoMedQ Neonatal Medicines [Online]. Accessed 6 Oct 2022. Last updated Aug 2019. Available on CKN.
 BNF for Children 1/10/22. BMJ Group, London, UK. Available on CKN.

Table 3: PAEDIATRIC Antimicrobial Administration Guidelines for Community Acquired Sepsis

- · Commence IV antibiotics as soon as possible after blood cultures have been taken. Do not delay antibiotic administration while awaiting blood test results.
- If multiple IV antimicrobial orders are prescribed, administer in order of shortest to longest infusion times to ensure completed as quickly as possible. For example:
- » Septic shock requiring inotropes: inject IV cefotaxime over 3–5 minutes, followed by IV gentamicin over 30 minutes, followed by IV vancomycin over 60 minutes.
- Ensure adequate saline flush between incompatible agents.
- Where possible use separate dedicated lines for resuscitation fluid and for medications. If not possible, pause either the antibiotic or the resuscitation fluid to administer. You may administer via Y-site, but not concurrent delivery.
- Use CREDD where this is the locally recommended resource.

Antimicrobial (tradename/ brand)	Strength (powder volume) [volume]	Reconstitution	Final concentration PIV = Peripheral IV CVL = Central	Intravenous (IV) administration	Compatible IV fluids	Additional information
Aciclovir (DBL) Intravenous	25mg/mL [10mL; 20mL]	Reconstitution not required	PIV: Dilute to 5mg/mL CVL: 25mg/mL	Infuse over 60 minutes	Sodium Chloride 0.9%Glucose 5%Hartmann'sPlasma-Lyte via Y-site	Extravasation risk Ensure adequate hydration
Amoxicillin (Fisamox, Ibiamox, Amoxil) Intravenous	1g (0.8mL)	Water for injection Add 9.2mL to 1g vial (100mg/mL)	PIV or CVL: • Dilute to 50mg/mL or weaker	Infuse over 30 minutes	Sodium Chloride 0.9% Glucose 5%, 10% via Y-site Hartmann's	Flush well between aminoglycosides Rapid IV injection may cause seizures
AMPicillin (Austrapen, Auspen, Ibimicyn) Intravenous	500mg (0.3mL) 1g (0.7mL)	Water for injection Add 4.7mL to 500mg vial Add 9.3mL to 1g vial (100mg/mL)	PIV or CVL: • Undiluted; 100mg/mL • Dilute to 30mg/mL for infusion	50mg/kg UP TO ≤500mg: Inject undiluted over 3–5 minutes 100mg/kg <i>OR</i> >500mg: Infuse over 15–30 minutes	Sodium Chloride 0.9% Glucose 5%, 10% Ringer's via Y-site	Flush well between aminoglycosides Rapid IV injection may cause seizures
Benzylpenicillin (BenPen) Intravenous	600mg (0.4mL) 1.2g (0.8mL) 3g (2mL)	Water for injection Add 1.6mL to 600mg vial Add 3.2mL to 1.2g vial Add 8mL to 3g vial (300mg/mL)	PIV: Dilute to 60mg/mL CVL: Undiluted; 300mg/mL	Infuse over 30 minutes	Sodium Chloride 0.9% Glucose 5% Plasma-Lyte via Y-site	Flush well between aminoglycosides Rapid IV injection may cause electrolyte imbalance and seizures
CefaZOLin (AFT, Hospira, Sandoz, Alphapharm) Intravenous	1g (0.5mL)	Water for injection Add 9.5mL to 1g vial (100mg/mL)	PIV or CVL: • Undiluted; 100mg/mL • Dilute to 20mg/mL for infusion	Inject undiluted over 3–5 minutes; <i>OR</i> Infuse over 10–60 minutes	Sodium Chloride 0.9% Glucose 5%, 10% Hartmann's Plasma-Lyte via Y-site	Flush well between aminoglycosides
cefOTAXIME (Sandoz, DBL) Intravenous OR Intramuscular	1g (0.4mL) 2g (1mL)	Water for injection IV: Add 4.6mL to 1g vial Add 9mL to 2g vial (200mg/mL)	PIV or CVL: • Undiluted; 200mg/mL • Dilute to 60mg/mL for infusion	Inject undiluted over 3–5 minutes; <i>OR</i> Infuse over 15–30 minutes	Sodium Chloride 0.9% Glucose 5%, 10% Hartmann's	Flush well between aminoglycosides More rapid injection may cause cardiac arrhythmias
		IM: • Add 2.6mL to 1g vial • Add 5mL to 2g vial (330mg/mL)	IM: • Undiluted; 330mg/mL		C-01039 Medication admini. of solutions to be Injected	
cefTAZIDIME (Sandoz, AFT) Intravenous	1g (0.9mL) 2g (1.8mL)	Water for injection Add 5mL to 1g vial Add 10mL to 2g vial (170mg/mL)	PIV or CVL: • Undiluted; 170mg/mL • Dilute to 40mg/mL for infusion	Inject undiluted over 3–5 minutes; <i>OR</i> Infuse over 15–30 minutes	Sodium Chloride 0.9%Glucose: 5%, 10%Hartmann'sPlasma-Lyte via Y-site	Flush well between aminoglycosides
CefTRIAXone (AFT, Alphapharm, Hospira) Intravenous OR	1g (0.6mL)	Water for injection IV: Add 9.4mL to 1g vial (100mg/mL)	PIV or CVL: • Dilute to 40mg/mL	Dilute and inject over 5 minutes; <i>OR</i> Infuse over 30 minutes	Sodium Chloride 0.9% Glucose 5%, 10% Incompatible with Hartmann's & Ringer's	Flush well between aminoglycosides, or calcium containing solutions Not recommended for use in neonates
Intramuscular		IM: • Add 2.3mL to 1g vial (350mg/mL)	IM: • Undiluted; 350mg/mL		C-01039 Medication admini of solutions to be Injected	

Table 3 (continued)

Antimicrobial (tradename/ brand)	Strength (powder volume) [volume]	Reconstitution	Final concentration PIV = Peripheral IV CVL = Central	Intravenous (IV) administration	Compatible IV fluids	Additional information
Ciprofloxacin (Aspen, DBL) Intravenous	2mg/mL [100mL]	Reconstitution not required	PIV or CVL: • Undiluted; 2mg/mL • Dilute to 1mg/mL	Infuse over 60 minutes	Sodium Chloride 0.9%Glucose: 5%, 10%Hartmann'sPlasma-Lyte via Y-site	Extravasation risk Ensure adequate hydration
Clindamycin (Mylan, Dalacin C) Intravenous	150mg/mL [4mL]	Reconstitution not required	PIV or CVL: • Dilute to 18mg/mL or weaker	Infuse over 20–60 minutes Maximum infusion rate: 20mg/kg/hr or 30mg/minute	Sodium Chloride 0.9%Glucose: 5%, 10%Hartmann'sPlasma-Lyte via Y-site	Rapid IV injection may cause hypotension and cardiac arrest
Flucioxacillin (Flucil, Flubiclox, Hospira) Intravenous	500mg (0.4mL) 1g (0.7mL)	 Water for injection Add 9.6mL to 500mg vial Add 19.3mL to 1g vial (50mg/mL) 	PIV or CVL: • Undiluted; 50mg/mL or dilute to convenient volume	Infuse over at least 30 minutes May give over 3–5 minutes (phlebitis risk)	Sodium Chloride 0.9%Glucose 5%Hartmann'sPlasma-Lyte via Y-site	Extravasation risk Flush well between aminoglycosides
Gentamicin (Pfizer) Intravenous	40mg/mL [2mL]	Reconstitution not required	PIV or CVL: • Dilute to 10mg/mL or weaker	Infuse over 30 minutes	 Sodium Chloride 0.9% Glucose: 5%, 10% Hartmann's Plasma-Lyte via Y-site 	Therapeutic drug monitoring (TDM) required Rapid IV injection may cause ototoxicity Flush well between cephalosporins and penicillin
Lincomycin (Lincocin, SXP) Intravenous	300mg/mL [2mL]	Reconstitution not required	PIV or CVL: • Dilute to 10mg/mL or weaker	≤1g: Infuse over 60 minutes >1g: Maximum infusion rate 1g/hour	Sodium Chloride 0.9%Glucose 5%, 10%Hartmann'sPlasma-Lyte via Y-site	Rapid IV injection may cause hypotension and cardiac arrest
Meropenem (DBL, Kabi, Ranbaxy) Intravenous	500mg (0.4mL) 1g (0.9mL)	 Water for injection Add 9.6mL to 500mg vial Add 19.1mL to 1g vial (50mg/mL) 	PIV or CVL: • Undiluted; 50mg/mL or dilute to convenient volume	Inject undiluted over 3–5 minutes; <i>OR</i> Infuse over 15–30 minutes	Sodium Chloride 0.9%Glucose 5%, 10%Plasma-Lyte via Y-site	
Metronidazole (DBL, Claris, Sandoz) Intravenous	5mg/mL [100mL]	Reconstitution not required	PIV or CVL: • Undiluted; 5mg/mL or dilute to a convenient volume	• Infuse over 20–30 minutes	Sodium Chloride 0.9%Glucose 5%Hartmann's via Y-sitePlasma-Lyte via Y-site	
Piperacillin/ Tazobactam (DBL, AFT, Kabi, Tazocin EF) Intravenous	Piperacillin 4000mg Tazobactam 500mg; (3mL)	Water for injection Add 17mL to 4/0.5g vial (200mg/mL)	PIV or CVL: • Dilute to 90mg/mL or weaker	Infuse over 30 minutes	Sodium Chloride 0.9% Glucose 5% Hartmann's via Y-site (AFT, Tazocin EF only) Plasma-Lyte via Y-site	Flush well between aminoglycosides Concentrations expressed as piperacillin component
Vancomycin (DBL, AN, Vancocin CP, Alphapharm) Intravenous	500mg; 1g (powder volume negligible)	 Water for injection Add 10mL to 500mg vial Add 20mL to 1g vial (50mg/mL) 	PIV: Dilute to 5mg/mL or weaker CVL: Dilute to 10mg/mL or weaker	• Infuse over 60–120 minutes	 Sodium Chloride 0.9% Glucose 5%, 10% Hartmann's Plasma-Lyte via Y-site 	TDM required Extravasation risk If Red Man Syndrome occurs, slow infusion rate

References:

The Royal Children's Hospital Paediatric Injectable Guidelines, June 2020, Melbourne, Australia. Available on CKN.
 Burridge, N., Ed. (2022). The Australian Injectable Drugs Handbook 8th edition. Collingwood, The Society of Hospital Pharmacists of Australia. Available on CKN.



Information for parents, carers and families of children with sepsis

What is sepsis?

Sepsis happens when the body has an extreme response to an infection and starts to injure its own tissues and organs. Sepsis can be triggered by any infection (viral, fungal, bacterial) but most commonly occurs with bacterial infections of the brain, lungs, bladder, kidneys, abdomen, skin and soft tissues.

Care for your child in hospital

Your child's healthcare team team will provide urgent treatments including:

- Insertion of a cannula, collection of blood tests and administration of antibiotics.
- Give fluids and other medicines, via a cannula, to support your child's circulation.
- Monitor your child's response to treatment.
- Consult with a sepsis expert.
- Arrange for transfer to the most appropriate place for your child's care which may be a general ward or Paediatric Intensive Care Unit (PICU).

There will be many people in your child's healthcare team, which may include doctors, nurses and a social worker. You are your child's key support and advocate; let your healthcare team know about your child's condition, their progress and any changes that concern you.

Your healthcare team should talk to you about:

- What a diagnosis of sepsis means for your child in the short, medium and long term.
- Plans for your child's treatment, who will provide this care and their response to treatment.
- What to expect during your child's recovery.
- How to inform the healthcare team if you are concerned your child is getting worse.
- Support you can receive in hospital.



Ryan's Rule

You and your family will be informed about your child's treatment options and involved in decisions about their care. If you have concerns that your child's health condition is getting worse or not improving, discuss this initially with the healthcare team. You can also search 'Ryan's Rule' on the Children's Health Queensland website to learn about raising concerns.





Support for your family in hospital

Dealing with a complex health issue like sepsis and a hospital admission can be stressful and challenging for all family members. Speak to your child's healthcare team about ways to access additional support which may include:

- Social workers who can provide help to adjust and manage your child's health condition and admission.
- Welfare workers who can provide practical support with accommodation, finances, travel, and social needs.

Children and medical procedures

It is common for children to struggle with some medical procedures. Reassure your child of your support. It helps children to know what is going to



happen, why the procedure needs to happen and who will be involved. For more ideas, scan this QR code and read our blog on supporting your child through a procedure.

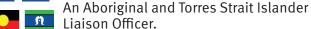
Cultural support

Let your healthcare team know if you need:





A translator or interpreter.



Sepsis resources

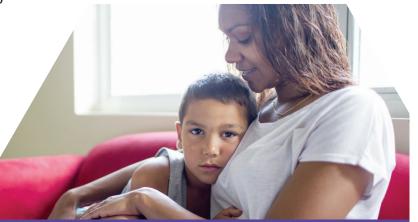
Sepsis on the Children's Health Queensland website has information for families including:

- 'Journeying through Sepsis' video series to support you through each stage of your child's sepsis journey.
- Paediatric Sepsis Family Support Network
- · Paediatric Sepsis Peer Mentor Program.

For more information visit Sepsis on the Children's Health Queensland website at www.childrens.health.qld.gov.au/sepsis or scan the QR code below.

Questions you could ask your child's healthcare team

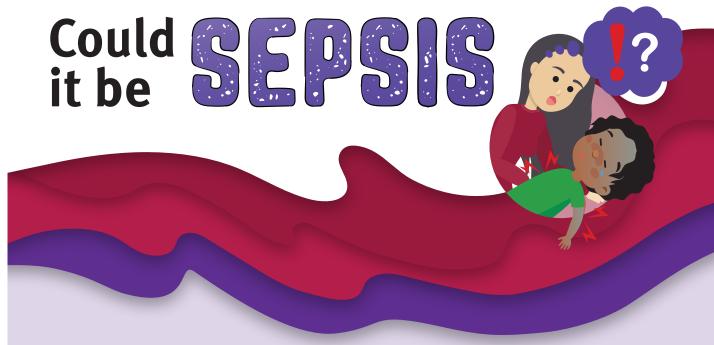
- What will my child's treatment be?
- Who will provide this treatment?
- How will my child be affected by sepsis and it's treatment?
- What complications of sepsis and the treatment should I be aware of?
- How did my child become unwell with sepsis?
- Who is my main contact person within the hospital for my child's care?
- What should I expect as my child recovers in hospital after the initial critical care for sepsis?
- How can I escalate my concerns if my child is getting worse?
- What supports are available to me, my child and my family in hospital?
- What should I expect with my child's recovery after discharge from hospital?
- What are the potential long-term impacts of my child's sepsis diagnosis?
- Is my child likely to come back to hospital?
- What are signs my child is getting unwell again, and when should we return to hospital or our GP?
- What supports are available to my child and our family following discharge from hospital?





Illnesses can change – trust your gut feeling. Even if your child has recently had sepsis, if you think they may have sepsis again come back to hospital and ask 'Could it be sepsis?'.

Visit www.childrens.health.qld.gov.au/sepsis



Sepsis is a **medical emergency** and needs immediate treatment.

Sepsis happens when the body has an extreme response to an infection and starts to injure its own tissues and organs. Sepsis can damage many parts of the body and can result in death. The best chance of getting better from sepsis is to treat it quickly.

Knowing if your child has sepsis can be difficult because many of the symptoms in the beginning are the same as mild infections. The difference is that your child's symptoms don't improve or may worsen.

Sepsis is rare, but any child can develop sepsis and we all need to know what to look out for.

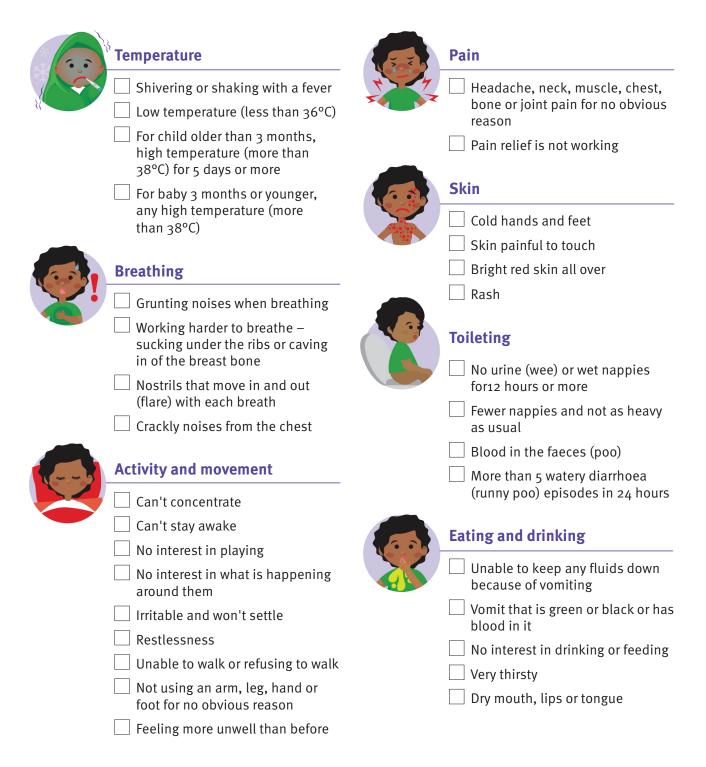
your child best, so trust your gut feeling. If your child is more unwell than ever before or this illness is different from other times – ask your doctor or nurse "Could it be sepsis?".





Paediatric Sepsis checklist

If you think your child is not getting better, or they are getting sicker, trust your gut feeling. Tick the boxes that apply to your child and ask your doctor or nurse "Could it be sepsis?".





Illnesses can change – trust your gut feeling. Even if your child has recently seen a doctor, if you think they may have sepsis, come back to hospital and ask "Could it be sepsis?".