<pre></pre>	Bigs of divide deterioration (e.g. altered level of consciousness or total Q-ADDS score of 24)         If you suspect neutropaenic sepsis, refer to local guidelines if available, otherwise continue screening on this pathway         Screening initiated:       D/ MM / YY         Are ANY of the following risk factors present? (lock all that apply)         Are ANY of the following risk factors present? (lock all that apply)         Are ANY of the following risk factors present? (lock all that apply)         Are ANY of the following risk factors present? (lock all that apply)         Are ANY of the following risk factors present? (lock all that apply)         Are ANY reason to suspect an infection?         Manourised / Applenia / Neutropaenia         Bit bere ANY reason to suspect an infection?         Abdomen / GIT         Stare alwy reason to suspect an infection?         Abdomen / GIT         Skin / Joint / Prosthesis / Device         Yes, but source is unclear at present         Bit we oxygen requipement heave ANY moderate risk criteria?         Bit we oxygen requipement heave ANY moderate risk criteria?         Bit we oxygen requipement heave ANY moderate risk criteria?         Bit we oxygen requipement heave ANY moderate risk criteria?         Bit we oxygen requipement heave ANY moderate risk criteria?         Bit we oxygen requipement heave ANY moderate risk criteria?         Bit we oxygen requipement heave A
Absence of risk factors does not exclude sepsises as a cause of deterioration         Bit expresentation within 48 hours         Bit expresentation with 48 hours         Bit expresentation with 48 hours Bit expresention with 48 hours Bit expresentation with 48 hours Bit expresenta	Absence of risk factors does not exclude sepsis as a cause of deterinduon         Absence of risk factors does not exclude sepsis as a cause of deterinduon         Balance of risk factors does not exclude sepsis as a cause of deterinduon         Balance of risk factors does not exclude sepsis as a cause of deterinduon         Balance of risk factors does not exclude sepsis as a cause of deterinduon         Balance of risk factors does not exclude sepsis as a cause of deterinduon         Balance of risk factors does not exclude sepsis as a cause of deterinduon         Balance of risk factors does not exclude sepsis as a cause of deterinduon         Balance of risk factors does not exclude sepsis as a cause of deterinduon         Balance of risk factors does not exclude sepsis as a cause of deterinduon         Balance of risk factors does not exclude sepsis as a cause of deterinduon         Balance of risk factors does not exclude sepsis as a cause of deterinduon         Balance of risk factors does not exclude sepsis as a cause of deterinduon         Balance of risk factors does not exclude sepsis as a cause of deterinduon         Balance of risk factors does not exclude sepsis as a cause of deterinduon         Balance of risk factors does not exclude sepsis as a cause of deterinduon         Balance of risk factors does not exclude sepsis as a cause of deterinduon         Balance of risk factors does not exclude risk for exclude sepsis deterinduon         Balance of risk factors does not exclude risk for exclude risk for e
<ul> <li>Until proven otherwise</li> <li>Obtain immediate senior medical review</li> <li>Consider transfer to resuscitation area</li> <li>Commence resuscitation</li> <li>Senior medical review attended: DD / MM / YY</li> <li>M: MM (24hr)</li> <li>Does the senior medical reviewer think sepsis or septic shock is likely?</li> <li>Sepsis / septic shock likely</li> <li>YES</li> <li>Commence resuscitation and treatment for sepsis NOW (See page 2)</li> </ul>	Senior medical review attended: DD / M/ / YY RA: M (24hr) Does the senior medical reviewer think sepsis or septic shock is likely? Sepsis / septic shock likely Sepsis / septic shock unlikely ✓ YES
	Signature Log Every person documenting in this clinical pathway must supply a sample of their initials and signature below

Queensland		(Affix identification label here)			
Government		URN:			
Emergency Department		Family name:			
Non-pregnant Adult Sepsis Pathway		Given name(s):			
For tertiary and secondary facilities		Address:			
		Date of birth: Sex:	M F I		
	Notify nursing team leader and SMO the patient has potential sepsis or septic shock (tick when notified)				
	ACTIONS 1–4 to be commenced for: • Neutropaenic or meningococcal sepsis within 30 minutes of recognition • Septic Shock within 1 hour of recognition of shock (mortality 20–23%) • Sepsis within 3 hours of triage (mortality 10–12%) (Document variance in comments section if key tasks not commenced)				
	1. Measure (or remeasure) lactate	C.	Lactate collected		
щ	<ul> <li>2. Take blood cultures x 2 sets</li> <li>Collect prior to antibiotics unless this would delay treatment for &gt;1 hour</li> <li>If patient has a central line collect an additional (third) set of blood cultures via the line</li> <li>Collect FBC, UEC, BGL, LFT, lipase and VBG</li> <li>For septic shock add coagulation studies</li> <li>Collect other relevant cultures but do not delay antibiotics</li> </ul>				
RESUSCITATE	<ul> <li>3. Commence appropriate IV antibiotics</li> <li>Identify likely source of infection (including relevant imaging findings)</li> <li>Prescribe antibiotics according to guidelines. Modify for allergies or prior microbiological sensitivities</li> <li>Notify nursing staff of urgent need to administer antibiotics and ensure completed</li> <li>Recommend consulting microbiologist or infectious diseases physician (particularly fir septic shock, recent overseas travel, risk factors for multi-resistant organisms, IV drug use, morbid obesity or dialysis patient)</li> </ul>				
	<ul> <li>4. Commence IV or intraosseous fluids if clinically indicated</li> <li>Consider volume of fluid based on patient's weight, cardiac function, contorbidities, current volume status and haemodynamics</li> <li>If bolus indicated, rapidly infuse 250mL–500mL IV or intraosseous 0.9% NaCl or Hartmann's over 5 minutes</li> <li>Assess response to fluid and consider repeating bolus if clinically indicated &gt; do NOT exceed 30mL/kg without SMO input</li> </ul>				
	5. Consider vasopressors/inotropes for hypo (e.g. Noradrenaline: usual commencing do	tension during or after fluid resuscitation se 5mcg/min)	Vasopressors/ inotropes considered (or not indicated)		
	6. Facilitate rapid source control - if this requ notification of appropriate surgical or inter	ires operative intervention ensure early ventional team	Source control facilitated (or not required)		
	<ul> <li>7. Reassess and monitor response to resuscitation - aim for:</li> <li>Oxygen saturation &gt;94% (88–92% if COPD)</li> <li>Systolic BP &gt;100mm Hg</li> <li>Urine output &gt;0.5 to 1.0mL/kg/hr - consider IDC with hourly monitoring</li> <li>Lactate &lt;2mmol/L</li> <li>If haemodynamic status not improving or if vasopressors/inotropes commenced refer to ICU</li> </ul>				
REVIEW	<ul> <li>8. Early referral to relevant inpatient team with clinical hand-over, and document:</li> <li>Appropriate criteria to ensure escalation of signs of deterioration</li> <li>Requirement to review antibiotics as soon as possible</li> <li>Need for infectious diseases, microbiologist or AMS team review, particularly in septic shock</li> </ul>		Referral completed and documented		
	Handover risk of deterioration to receiving nurse when patient transferred out of ED       Date and time complete:         An emergency call can be initiated at any time if you are clinically concerned.       DD / MM       HH : MM (24hr)       initials				
	ED staff name:	Ward staff name:			
Comn	nents / Variance from Actions				

DO NOT WRITE IN THIS BINDING MARGIN