

### Success stories

### Metacognition and clinical reasoning

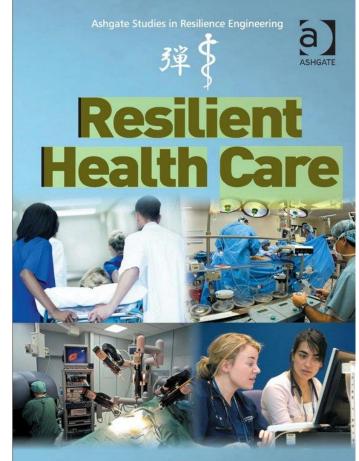
### Dr Stephen Gourley

### Deputy Chair, National Rural Health Alliance and

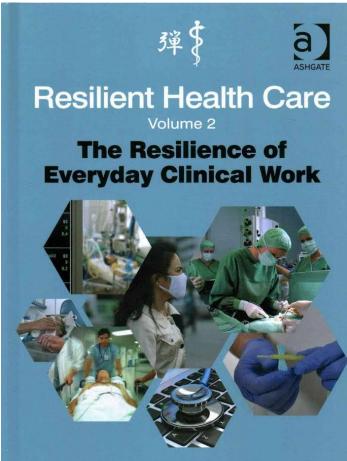
Director Emergency Medicine, Alice Springs Hospital, Northern Territory

Lean thinking Six-Sigma **Root cause analysis** Swiss cheese theory Airline safety models Checklists

Lean thinking Six-Sigma Root cause analysis Swiss cheese theory Airline safety models Checklists



ERIK HOLLNAGEL, JEFFREY BRAITHWAITE AND ROBERT L. WEARS



Robert L. Wears, Erik Hollnagel and Jeffrey Braithwaite

### **Resilient Health Care**

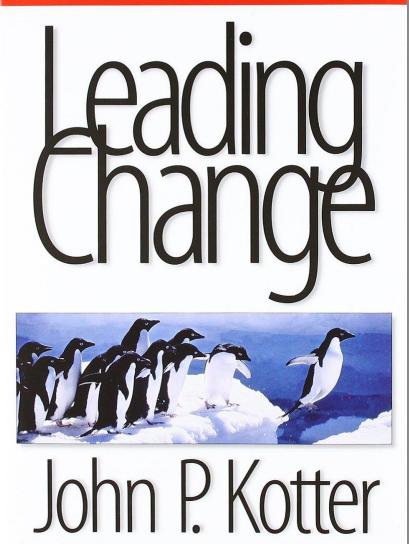
V O L U M E 3 Reconciling Work-as-Imagined and Work-as-Done



EDITED BY Jeffrey Braithwaite • Robert L. Wears • Erik Hollnagel



#### AN ACTION PLAN FROM THE WORLD'S FOREMOST EXPERT ON BUSINESS LEADERSHIP



HARVARD BUSINESS REVIEW PRESS

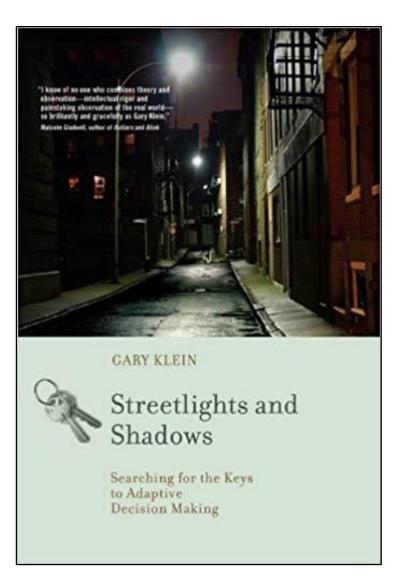
Artistry, Choice, & Leadership

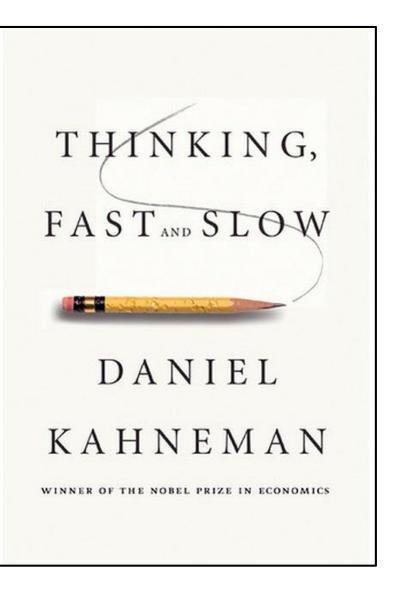
# REFRAMING Edition ORGANIZATIONS

LEE G. BOLMAN TERRENCE E. DEAL

Bestselling authors of Leading with Soul

II JOSSEY-BASS A Way Band





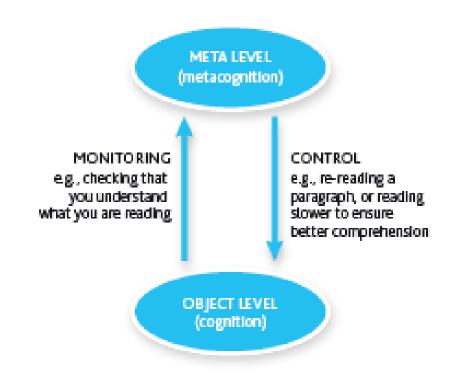
# METACOGNITION



Thinking about thinking.... Learning about learning.... Knowing about knowing .... Awareness of awareness...

# METACOGNITION

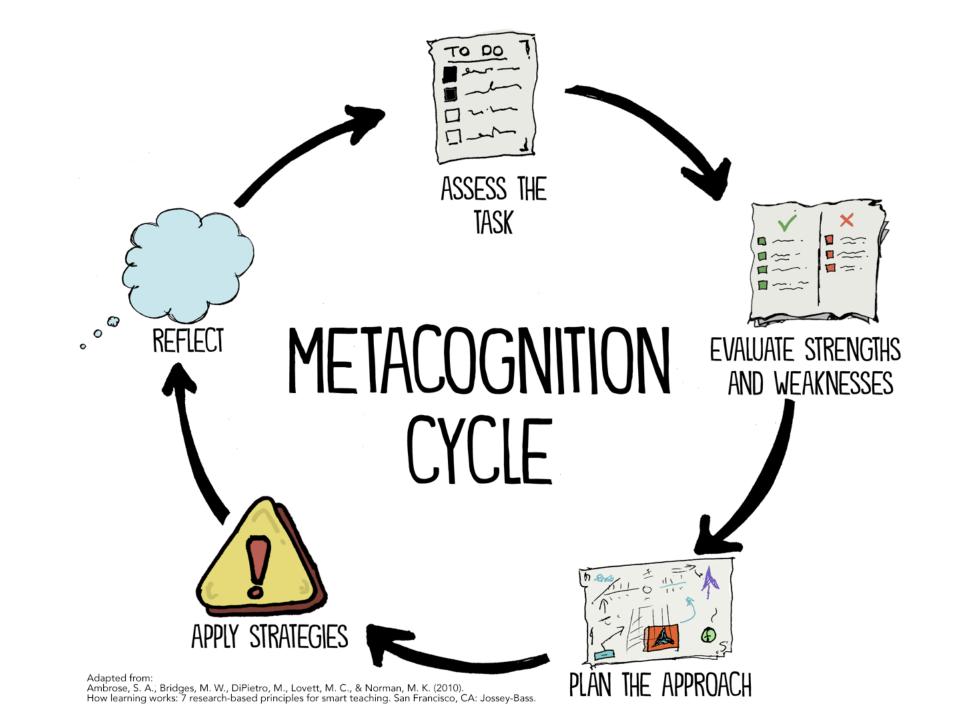


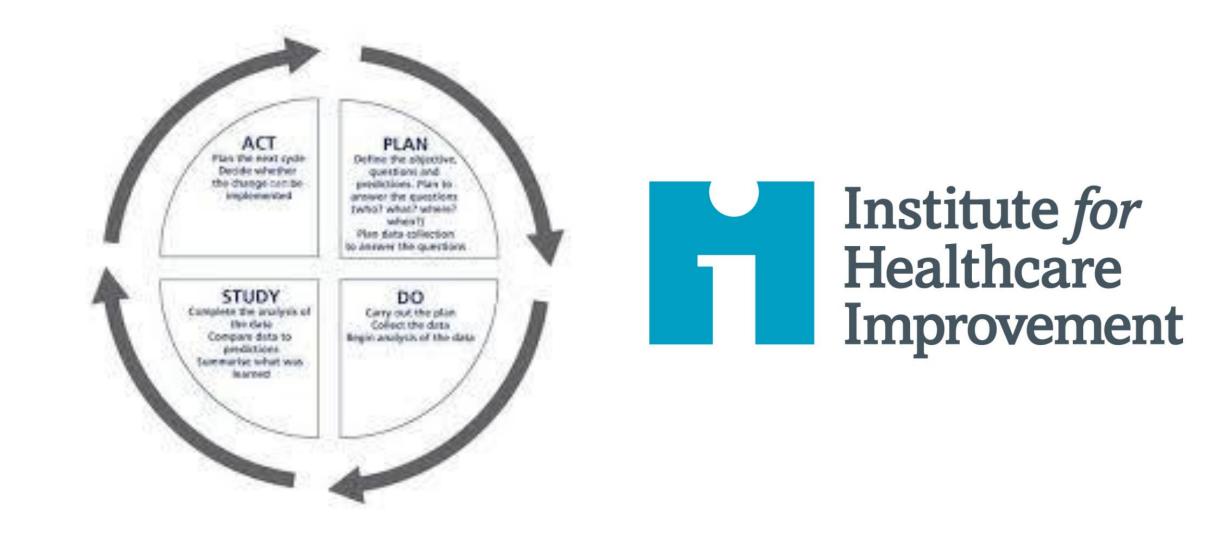


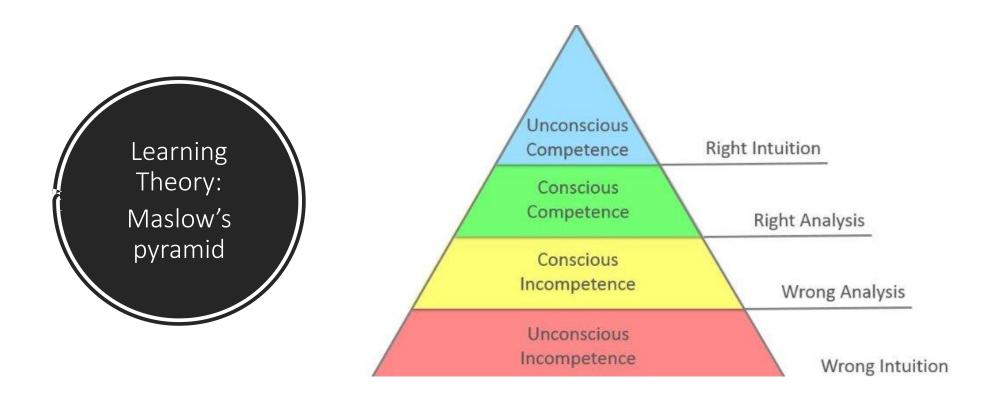
Regure 1. Nelson and Narens' (1990) Model of Metacognition

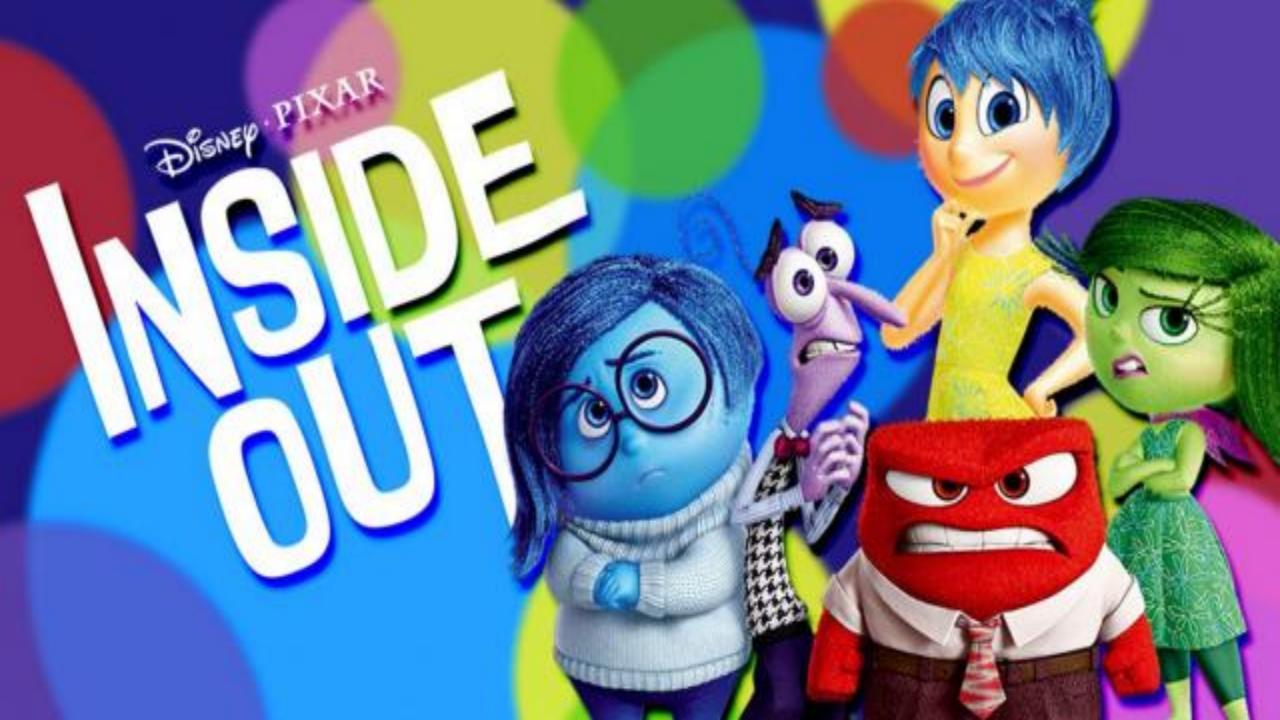
# METACOGNITION

- Content knowledge understanding your own capabilities often inaccurate and greater confidence in performing well is associated with a less accurate metacognitive judgment of performance
- Procedural knowledge about doing things. This type of knowledge is displayed as heuristics and strategies. A high degree of procedural knowledge can allow tasks to be done more automatically (like driving a car).
- Strategic knowledge your capability for using strategies to learn information
- Conditional knowledge knowing when and why to use content and procedural knowledge









# **Type 1 Thinking**



# **Type 1 Thinking**

- Fast
- Automatic
- FrequentEmotional
- Stereotypical
- Unconscious



### **TYPE 2 Thinking**

# SADNESS

## **TYPE 2 Thinking**

- Slow
- Effortful
- Infrequent
- Logical
- Calculating
- Conscious



# "H" ANGER

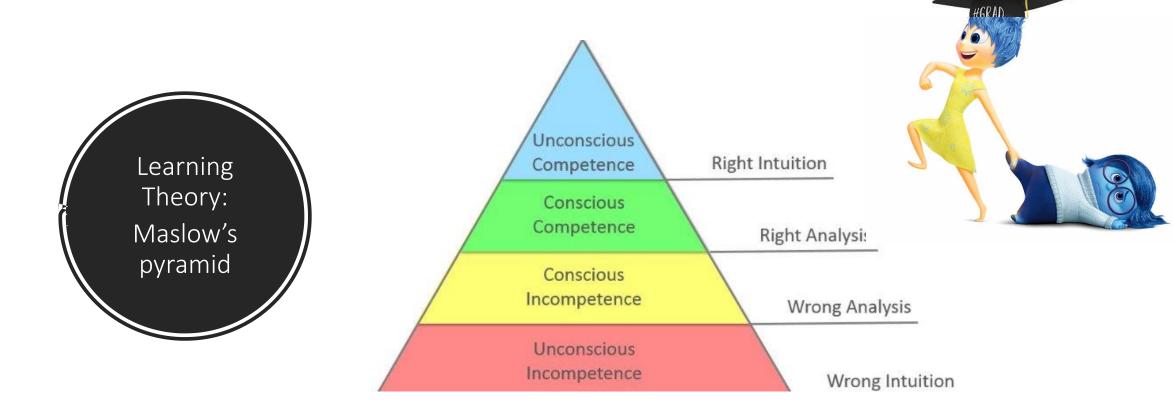
**#INSIDEOUT** 

6





### Type I thinking in training.....



# Cognitive Biases

Many described and seem to keep expanding Important to know they are there But more important to listen to Type 2 when there is a concern

Examples:

- Anchoring
- Availability
- Substitution
- Optimism and loss aversion

- Framing
- Sunk cost
- Overconfidence

....Biases help to explain why we are terrible at statistics and numbers

### Possible solutions What's new in safety?



## Possible solutions

- Resilience theory
  - Work in a complex adaptive system
  - Linear models to look at error are not fit for purpose
  - Linear solutions are unlikely to succeed (and haven't)
- Safety II model
  - Focus on what goes right, rather than what goes wrong
- Clinical leadership
  - Positive, supportive, collaborative
- Strong organizational culture
  - "The way we do things around here"
  - Focus on work-as-done, rather than work as imagined





# Thank you



Australasian College for Emergency Medicine



### Success stories

Supervising for safety

### Dr Amanda Stafford Clinical Lead Royal Perth Hospital Homeless Team Emergency Department Consultant Royal Perth Hospital, Western Australia



Clinical Supervision & Patient Safety

Dr Amanda Stafford Royal Perth Hospital

### **Clinical Supervision**

• A formal process of professional support and learning which enables individual clinicians to develop knowledge and competence and assume responsibility for their own practice.

• Cutcliffe and Butterworth, 2001

• "The current flaw is that many work practices depend on the judgement of junior doctors to recognise when they don't know or are out of their depth."

• Clinical Supervision at the point of care Clinical Excellence Commission 2012 NSW • Trainees are junior, mobile, inexperienced and may be "unconsciously incompetent" in their task and need to be actively supported and supervised rather than passively overseen."

• Clinical Supervision at the point of care Clinical Excellence Commission 2012 NSW

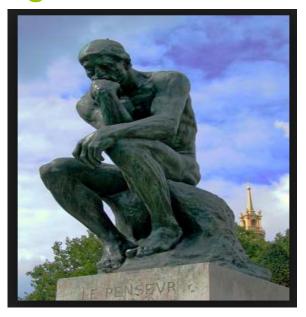
### Training as an Apprenticeship

• "The apprenticeship model persists as the major model of training, especially in procedural specialities."

• Clinical Supervision at the point of care Clinical Excellence Commission 2012 NSW

### Main EM Skills

### **Diagnostic Skills**



#### **Technical Skills**





Clinical Supervision at the point of care Clinical Excellence Commission 2012 NSW

• 54 RCAs 2008 and 2009 in NSW

• Clinical Supervision theme

- clinical reasoning
- decision making
- care provided

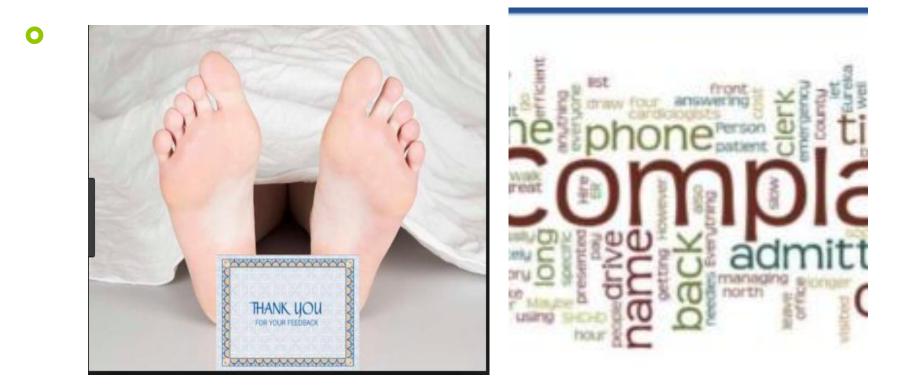
## Major Issues

<ul> <li>Lack of recognition of deterioration 31/54</li> </ul>	
<ul> <li>Inadequate treatment</li> </ul>	23/54
<ul> <li>Missed diagnosis</li> </ul>	14/54
<ul> <li>Delayed diagnosis</li> </ul>	4/54
<ul> <li>Wrong treatment</li> </ul>	4/54
<ul> <li>Delayed treatment</li> </ul>	3/54

# Major Contributors

- Availability of Senior Clinician
- Workload
- Rostering/Skill Mix
- Conflicting Priorities
- Lack of Senior Clinician Review or Input
- - escalation of care
- - appropriate level of care
- - culture re involving Senior Clinicians
- - effectiveness of supervision

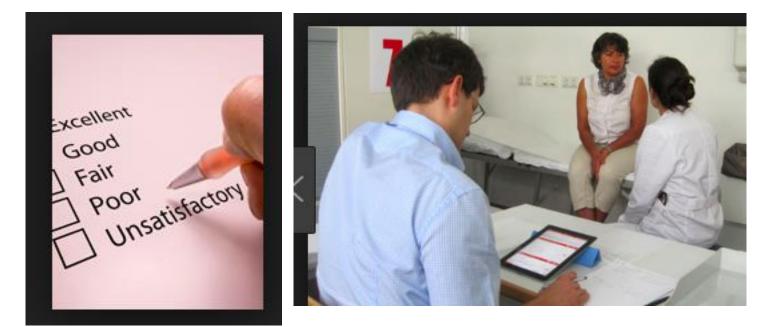
### Measuring the Effectiveness of Clinical Supervision



### More positive.....

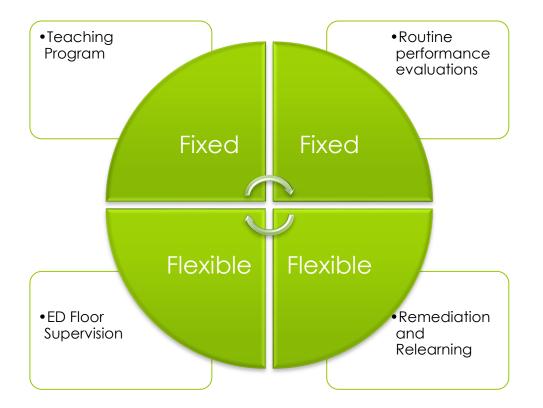
### • Term report & WBA

### • FACEM exam

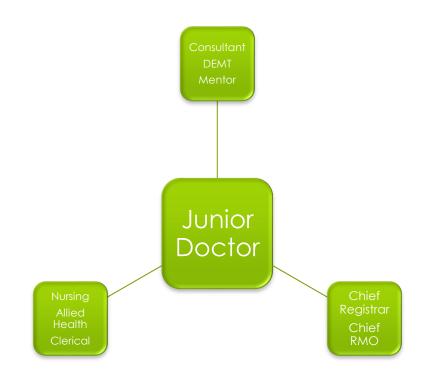




### Good Clinical Supervision Need a framework and system



## Multi-disciplinary Team



### Poorly performing doctors



### Thank You





Australasian College for Emergency Medicine



### Success stories

### Clinical practice improvement ED sepsis project

### Dr Anh Tran

Emergency Physician, Emergency Department Werribee Mercy Hospital, Victoria

# Patient story

2030 59 yo female (Type 2 DM) presented at post fall
Triage: Pulse 112 RR 20 BP 138/53 Sao2 95% room air GCS 15
Category 4
90 mins : Temp 40.1 pulse 109 RR 21 BP 113/86 Sao2 94% room air
Bloods and blood culture sent

### 150 minutes: "A/W r/v by EDMO"

**7.5 hours: Temp 39.5 pulse 121 RR 20 BP 95/50 Sao2 95% room air GCS 14/15** T/F resus, IV antibiotics, Nsaline 500mls

?lactate

9.5 Hours: stable, improved GCS referred to med team

# Sepsis care -audit

### VAED code: "Severe Sepsis" "Septic shock" N=18

Recognition at triage	67%
Started on a sepsis pathway	0%
IV antibiotics administered within 1 hour	22%
IV fluids administered within 1 hour	50%
Lactate measured	94%
2x blood cultures taken	67%
Adult Retrieval Victoria consulted (UCC or regional ED only)	22%
Transfer to other hospital (UCC only)	44%

# What to do?

- a. Keep auditing
- b. Communicate the audit to clinical staff in the hope they improve performance
- c. Schedule face to face education sessions to educate clinicians on the importance of good sepsis care
- d. Communicate the importance of timely interventions in sepsis
- e. Clinical practice improvement project



30th JANUARY 2019

### Implementing a sepsis bundle of care in the WMH emergency department End of project report February 2019



# WERRIBEE MERCY HOSPITAL

WMH ED has a 3 bed Resuscitation area (x1 paediatric), 9 monitored cubicles and 6 unmonitored cubicles, a designated fast track area and a 10 bed Short Stay Unit.

### Medical EFT :

FACEM EFT 7 JMO 25 Nursing 70

The emergency department sees 40000 patients per year.

This project coincided with the opening of the ICU at Werribee Mercy Hospital on August 1. (increased presentations)

Hospital inpatient services include: general medicine (including HDU/ICU), general surgery, obstetrics and gynaecology, special care nursery and psychiatry.

Limited subspecialty services





# Safer Care Victoria

Safer Care Victoria (SCV) is the state's healthcare quality and safety improvement agency. Works with patients, families and carers, clinicians and health services to monitor and improve the quality and safety of care delivered across Victorian public health system.

**ECCN- Emergency Care Clinical Network.** 

This network brings together clinicians who deliver emergency care

Within Urgent Care Centres, Emergency Departments and through Ambulance Victoria to improve the quality of care and patient experience in Victorian emergency departments.





# What are we trying to accomplish?

# Aims

#### Improve <u>early recognition</u> of sepsis

Improve early intervention

**Antibiotics within 1 hour** 

Improve <u>early escalation</u> of care

Measure 1: Proportion of patients identified at triage

Measure 2: Proportion of patients initiated on a sepsis pathway in ED

Measure 3: Proportion of patients administered IV fluids within one hour of presentation

Measure 4: Proportion of patients administered IV antibiotics within one hour of presentation

Measure 5: Proportion of patients with lactate measured

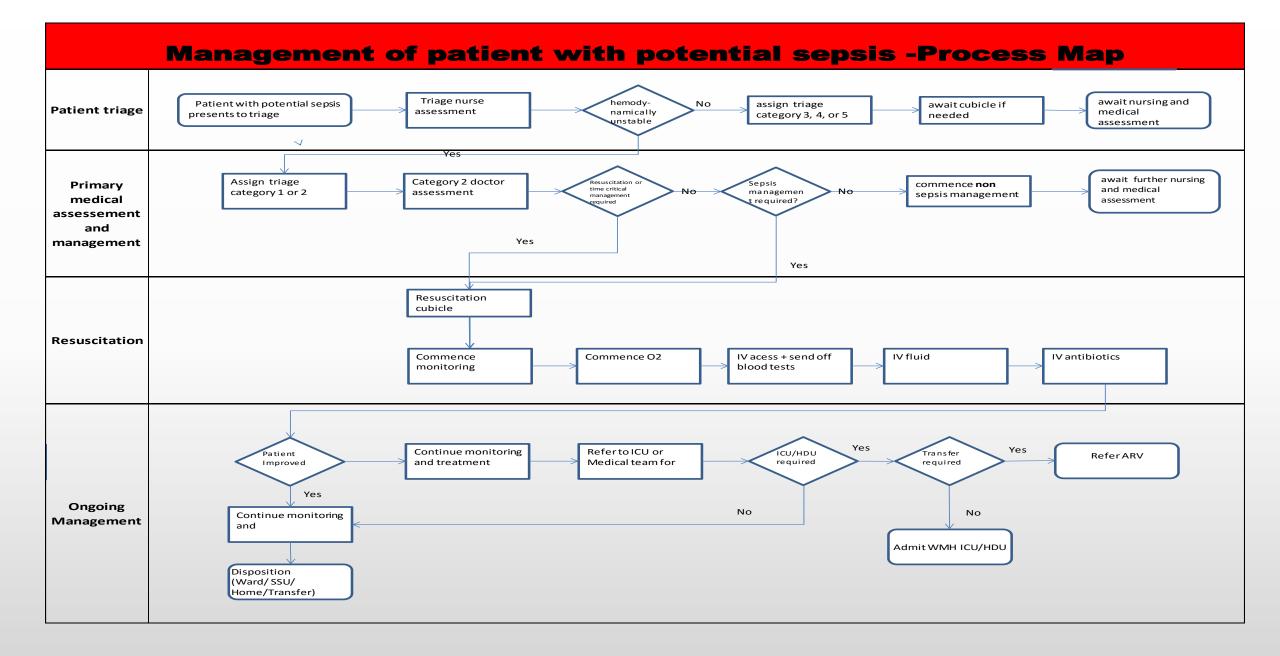
Measure 6: Proportion of patients with 2 sets of blood cultures taken within 6 hours of presentation

Measure 7: Proportion of patients where Adult Retrieval Victoria was consulted

Measure 8: Proportion of patients transferred to higher level of care (inter-hospital transfer or transfer to ICU/HDU)



# What change will lead to an improvement?





#### WMH Sepsis project affinity diagram

Why do patients presenting to the emergency department with suspected sepsis not receive optimum care?

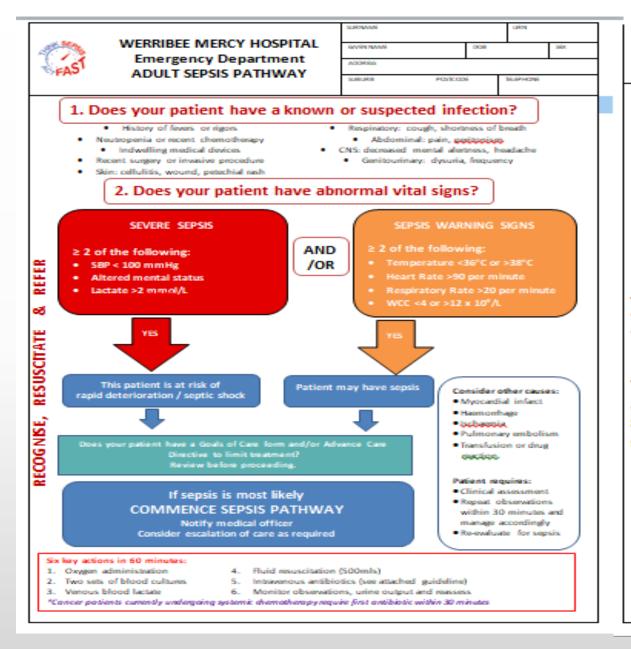
Recognition	Process	Clinician Knowledge	Task	Equipment	
Some patient groups not being assessed and managed appropriately eg	observations for suspected sepsis not standardised ie some not doing BP/Temp at	uncertainty re which bloods to take lactate: ?venous/ arterial blood culture: ?how many	other competing demands on Cat 2 doctor hence delayed charting of IV abs	planned calibration of lactate prevents lactate level being obtained hence delays	
current sepsis criteria not bieng followed	deteriorating patient in waiting room not reassessed	sets/ which site/ ?take before antibiotics	delay in cannulation due to bed availability, staff availability	not enough beds availab to commence interventio	
		when analyser is calibrating there is uncertainty about		no resus space available	
high risk patients are not	patients not reviewed post treatment in a timely manner	alternative actions resulting	delay in cannulation due to unsuccessful attempts and	when busy	
identified	ie there is not standardised review process		one operator attempting		
		uncertainty re what			
patients not assigned correct triage category	en a patient is septic the frequency of observations and process of review is not	interventions	delay in antibiotics due to getting the medication charted due to other competing		
recognition delayed due to	patients not reviewed post	uncertainty re which antibiotics to give and when to give ie benchmark time	demands, doctors need to look up what to give, doses		
increased workload	treatment in a timely manner ie there is not standardised review process	uncertainty re when to escalate ie trigger, time	delay in antibiotic due to increased preparation time		
	patients not reviewed post treatment in a timely manner ie there is not standardised	frame for referral , acceptable physiological parameters			
	review process				
		understanding of how much to give			
	patients not reviewed post treatment in a timely manner				
	ie there is not standardised review process				
		understanding re when to give intervention and the			
	patients not reviewed post treatment in a timely manner ie there is not standardised review process	best time frame			

### **ECCN Sepsis Project- Driver diagram**

Aim		Primary Drivers		Secondary Drivers	Change Ideas			
				recognition criteria and high risk groups are defined,	update sepsis criteria and apply to all patients. make criteria clear and easy to use			
			ı /	clincians are aware and use recognition criteria and early warnings signs to identify severe sepsis and commence necessary treatment	Make criteria visible at triage and clinical areas.			
	/	all sepsis is recognised and action is taken early	*	clincians are able to take early action when sepsis is recognised	standardise response when sepsis identified ie assign category 2 at triage immediately commence on a pathway, overhead call out, simplify initial response			
improved care of				evidence based interventions are defined and are standardised	make initiation of treatment not cubicle dependent			
patients presenting to the	timely and effective	A	evidence based interventions are embedded into process of care which is standardised	develop an escalation process for when resources a re not available eg resus beds, cubicles, allocated nursing staff, blood gas				
ED with suspected sepsis		provided to all patients with sepsis frontline staff supported to efficiently care for septic patients		clinicians follow standardised pathway for all patients identified with sepsis	analyser revise ECCN pathway (based on evidence)			
			frontline_staff	frontline staff	frontline staff		when usual management options are not possible eg resus cubicle occupied, alternative strategies are defined and known to clinicians	to make it more relevant locally ensure steps in pathway are clearly defined, understandable and relevant locally
				staff are supported and able to obtain assistance when unable to provide timely interventions eg IV access information is readily available to inform and	develop a escalation process to support clinicians when unable to take bloods or cannulate			
				all staff have the skills and knowledge required to look	improve cannulation and venepuncture skills in all clinical staff			
		clinicians have knowledge and skills required to provide good sepsis care		knowledge and skills required to provide	all clilnicans have an understanding of recognition criteria and importance early recognition and timely intervention	develop a sustainable education package to assist with improving awareness, recognition and understanding the the importance of timely treatment and steps required to improve the care of sepsis		
				increase knowledge of latest evidence based care	online pacakge to allow staff to learn in their own time			
				improve awareness of when and how to escalate care	feedback process for missed cases or "wins" to assist with case based learning			

# What changes will lead to improvement?

- Improving recognition
- Standardising care to assist with early, effective intervention (in line with evidence based practice)
- Improving clinician awareness and understanding



Eme	BEE MERCY HOSPITAL rgency Department LT SEPSIS PATHWAY
Activation of pathway	Date:
Signs/ Symptoms	1. Does your patient have two or more SRS criteria, and/or severe sepsis?       2. Does, your patients also have any of the following risk factors, signs or symptoms of infection?         Temperature <36°C or >38°C       History of fever or rigor         Heart Rate >00 bpm       Neutropenia or recent chemotherapy         &gag Rate >20/min       Indwelling medical divice         WCC <4 or > 12 x 10°/L       Recent surgery/invisive procedure         Systolic BP <100mmHg       Skin: cellulitis, wound, petechial rish         Altered mental state       Respiratory: cough, shortness of breath         Decessed/usioo?       CNS: decreased mental alertness, headache         Cool peripheries (hands and feet)       Genitourinary: dysuria, frequency
Medical review	Name: Time: CONTINUE PATHWAY: Y / N
1. Oxygen administration	Aim SpO, 92-96% (or 88-92% for COPD& chronic typeII respiratory failure) TIME: Initials
administration Ensure IV access	Large bore peripheral cannula inserted/available for fluid bolus, OR If central venous access device already available: type (if applicable)
2. Blood cultures	Two sets of blood cultures         TIME:         Initials           (2 peripheral; or 1 from all lumens of device or port if accessible, plus 1 peripheral)         *DO NOT DELAY ANTIBIOTIC ADMINISTRATION IF BLOOD CULTURES DIFFICULT TO OBTAIN
3. Lactabe	Venous blood lactate TIME: Initials Record lactate levelgppol/L
Pathology	Collect FBC, UEC, CRP, LFTs, coags and blood glucose level     Consider cross match if patient at risk of anaemia or known recent surgery     DO NOT WAIT for test results. Commence fluid resuscitation and antibiotics ASAP.
4. Fluid Resuscitate If hypotensive (SBP<100mmHg) or lactate >2mmol/L	Fluids must have medical officer authorisation and be prescribed on the IV Therapy Chat
If blood ;	pressure does not improve after fluid boluses EXCALATE care and consider inotropes

*							Empirie ontihintie o	والمتحد
		SURVERSE		LIKN			Double-click to show w	/hit
Emerge	ency Department	Service reacts	DOB		Silox		<ul> <li>Empirical regimens are interested.</li> </ul>	ende
-	SEPSIS PATHWAY	ADDRES	I		-		information is available	
ADULT SEPSIS PAIRWAT							<ul> <li>Ensure the patient's clinic</li> </ul>	
							uncertain, use the recomm     The following guidelines I	
<b>Clinically examine</b>	the patient for a focus of infection,	e.g. chest, urinary tract i	infection				please refer here for more	
5. Antibiotics	Check the patient's ALLERGY STA no penicillin allergy non-life-threatening penicilli Record antibiotic allergy and rea	icillin allengy (e.g. rash) in allengy (e.g. anaphylaxis	u		ials.		All doses recommended in may be required for patien Risk factors for high risk o organism, e.g. ESBL, Pseud No allergy to penicillin	nts wi f mul
i soooda uuda a faan Shee							UNKNOWN SOURCE OF INFECTION cefbratone 1g IV 24-hourly PLUS flucloxacillin 2g IV 4-hourly	
	For SUSPECTED, KNOWN or UNI Refer to empiric antibiotic guid		presumed site)	hnit	úals.		Add vancomycin IV if MRSA     Use meropenem 1g IV 8-ho	
	Antibiotics must be pr	rescribed on a medicat	ion chart by a i	medical of	ficer.		PEBRILE NEUTROPENIA piperacillin/tazobactam 4.5g IV 6-ho	urly
	ADMINISTER ANTIBIOTICS AS SC "Cancer patients currently under antibiotic within 30 minutes Time prescribed:;	going assemic chemother		Ini	sals.		<ul> <li>Consider adding stat gentari</li> <li>Add vancomycin IV if MRSA</li> <li>Use meropenem 1g IV 8-ho unconscious) as piperacillin</li> <li>Consider adding metonidar infection possible</li> </ul>	or li urly i tarol
Steroids	Consider hydrocortisone if patie	ant taking conticosteroids	ar known/Suspect	ted steroid d	eficien	cy	<ul> <li>Seek specialist advice if fung</li> </ul>	
	NOT improving - ESCALATE care, e.g.	ICU referral					INTRAVASCULAR DEVICE SOURCE (r	
Name of contact:	Tim	e:;					piperacillin/tacobactam 4.5g IV 8-he PLUS vancomycin IV (see dosing tab	
6. Monitoring	Monitor vital signs & fluid balan frequently as needed. Keep oxygen saturation 92-96% Assess for deterioration which Increasing respiratory rate (in	(88-92% if at risk of CO <sub>2</sub> n may include one or more	stantion) of the following:		more		Consider piperacillin/turoba     Use meropenem 1g IV 8-ho     Consider adding antifungal     colonisation with Candida	ctam urfy i
	S8P <100 mmHg						RESPIRATORY TRACT SOURCE	
	Decreased or no improvement Urine output < 0.5 ml/kg/hos If factate elevated repeat in 2	af .	DON'L ESCALATE O	are, e.g. ICU	referral		celluizzene 1g IV 24-hourly PLUS authromycin 500mg IV 24-hourly Consider oral oseltamivir 75 Consider celluizzone 1g IV 1	
	Initiate investigations as directed	Sputum	for MCS				<ul> <li>Replace celtriaxone with pip known respiratory colonisat</li> <li>Consider adding fluctoracilli</li> </ul>	ion v
Investigation	Urine MSU (or CSU) for MCS     Throat swab for respiratory m		swab for MCS or C. df/ficile testin	ng (il dianho	a pres	ent)	cases (e.g. rapid clinical dete	riora
Investigation Source control		ultiplex PCR 🔲 Stool fo		ng (if diantho	aa pres	ent)		riora

### e-click to show white space s Diseases consult (if available)

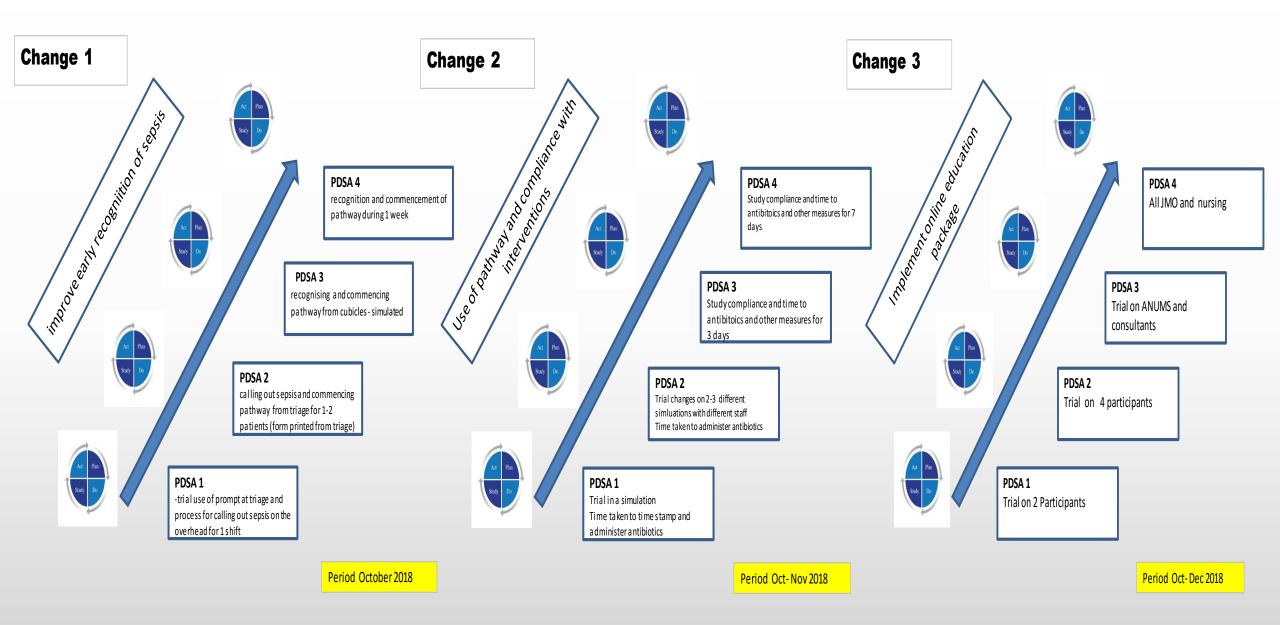
<ul> <li>Empirical regimens are intended for initial therapy ONLY (up to 48 hours) - modify as soon as additional information is available</li> <li>Ensure the patient's clinical findings and investigations are concordant with the presumed site of infection; if uncertain, use the recommendations for unknown site of infection</li> <li>The following guidelines have been adapted from Therapeutic Guidelines (TG): Antibiotic (version 15, 2014), please refer here for more detailed information if required or seek expert advice</li> <li>All doses recommended in this guideline are for normal renal function with CGD=S0ml/min, dose reductions may be required for patients with nenal impairment - see Appendix 2.6 (Table 2.33) TG for advice</li> <li>Risk factors for high risk of multidrug-resistant organisms: known colonisation with multidrug-resistant organism, e.g. ES8L, <i>Pseudomonos</i>, high risk travel (Indian subcontinent, Asia, Southem/Fastern Europe)</li> </ul>							
No allergy to penicillin	Non-life threatening penicillin allergy	Life threatening penicillin allergy					
UNKNOWN SOURCE OF INFECTION							
celtriaxone 1g IV 24-hourly PLUS	ceftriaxone 1g IV 24-hourly PLUS	ciprofloxacin 400mg IV 12-hourly PLUS					
fluclosacillin 2g IV 4-hourly	cefazolin 2g IV 6-hourly	vancomycin IV (see dosing table)					
<ul> <li>Add vancomycin IV if MRSA is su</li> <li>Use meropenem 1g IV 8-hourly i</li> </ul>	spected f high risk of multidrug-resistant organism						
FEBRILE NEUTROPENIA							
piperacillin/tarobactam 4.5g IV 6-hourly	celepime 2g IV 8-hourly	ciprofloxacin 400mg IV 12-hourly PLUS vancomycin IV (see dosing table)					
unconscious) as piperacillin-tarol	00mg IV 12-hourly (to celepime and cipro						
INTRAVASCULAR DEVICE SOURCE (remov	ve device)						
piperacillin/tazobactam 4.5g IV 8-hourly	cefacidime 2g IV 8 hourly PLUS	ciprofloxacin 400mg IV 12-hourly PLUS					
PLUS vancomycin IV (see dosing table)	wancomycin IV (see dosing table)	vancomycin IV (see dosing table)					
<ul> <li>Consider piperacillin/tazobactam</li> </ul>	4.5g IV 6-hourly in critically ill patients wi	th severe sepsis or septic shock					
<ul> <li>Use meropenem 1g IV 8-hourly i</li> </ul>	f high risk of multidrug-resistant organism						
<ul> <li>Consider adding antifungal cover colonisation with Candida</li> </ul>	if severe sepsis, high risk (e.g. prolonged i	ntravenous access) or known					
RESPIRATORY TRACT SOURCE	RESPIRATORY TRACT SOURCE						
celtriaxone 1g IV 24-hourly PLUS	celtriaxone 1g IV 24-hourly PLUS	moxifloxacin 400mg IV 24-hourly					
azithromycin 500mg IV 24-hourly	arithromycin 500mg IV 24-hourly						
<ul> <li>Consider oral oseltamivir 75mg 1</li> </ul>							
÷.	urly in critically ill patients with severe sep-	-					
	Ilin-tazobactam 4.5g IV 6-hourly OR mere						
	rith resistant organism, e.g. Pseudomonas,						
<ul> <li>Consider adding fluctoxacillin 2g IV 6-hourly and vancomycin if strongly suspect Staphylcoccus aurrus in severe cases (e.g. rapid clinical deterioration or cavitating pneumonia)</li> </ul>							
URINARY TRACT SOURCE							
celtriaxone 1g IV 24-hourly PLUS	ceftriazone 1g IV 24-hourly	ciprofloxacin 400mg IV 12-hourly					
amoxicillin 2g IV 6-hourly	and the second sec	and the second se					
Consider which are to be the bound of a side of the theory of the second side of the theory of theory of the theory of the theory of theory of theory of theory of th							

- Consider celtriaxone 1g IV 12-hourly in critically ill patients with severe sepsis or septic shock
- Use meropenem 1g IV 8-hourly if severe AND high risk of multidrug-resistant organism

# Online sepsis education package

Power point presentation 10 question quiz All clinical staff voluntary

# **SurveyMonkey**®

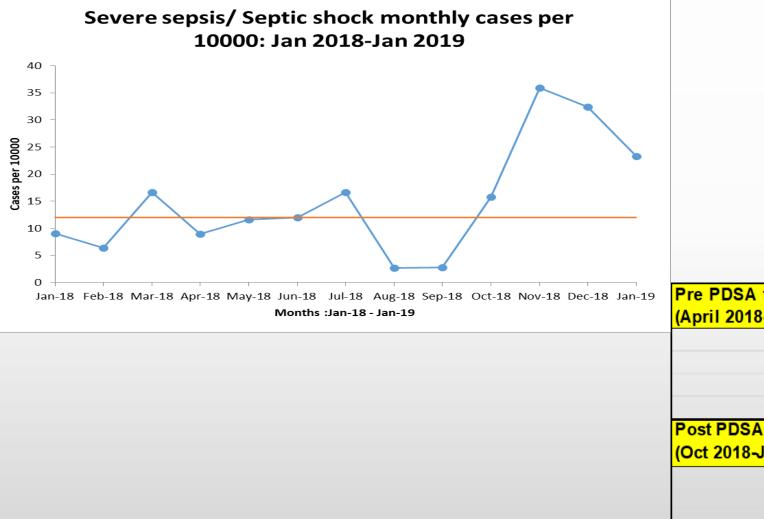




# How do we know if there has been an — improvement?

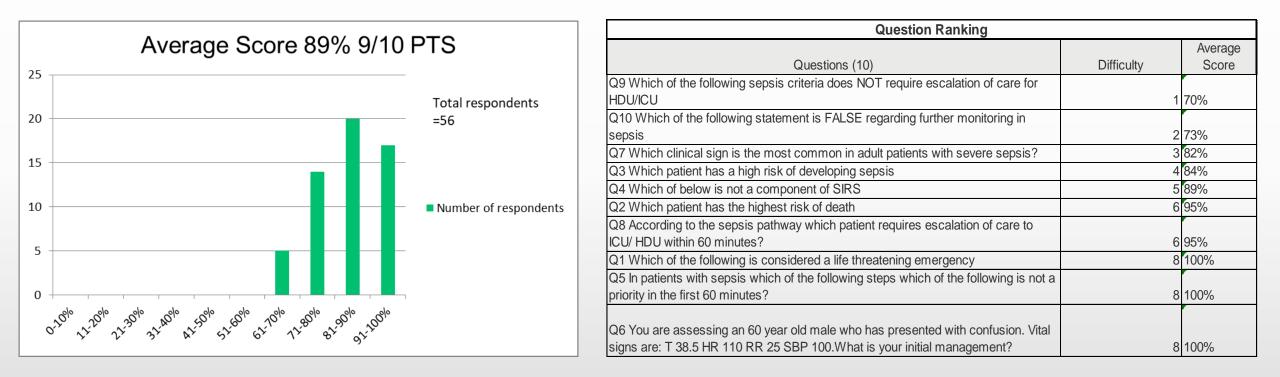


# Evaluation - Recognition



Pre PDSA testing/implementation		
(April 2018-Sept 2018)	Triage Category	Count
	Cat 1	0
	Cat 2	14
	Cat 3	3
	Cat 4	2
Post PDSA testing/implementation		
(Oct 2018-Jan 2019)	Triage Category	Count
	Cat 1	1
	Cat 2	28
	Cat 3	7

# Evaluation- Sepsis Education



Did this education package improve your understanding of sepsis												
	was not hel	pful	somewh	at helpful	neu	tral	help	oful	very h	elpful	Total	Weighted Average
star	1.85%	1	5.56%	3	3.70%	2	42.59%	23	46.30%	25	54	4.26
											Skipped	2

# Measures

Measure	Pre data (1 Jan - 30/06/2018)	Post Data (30/10/2018- 29/01/2019)
Recognition at triage	66%	96%
Started on sepsis pathway	0%	80%
IV antibiotics within 1 hour	22%	80%
IV fluids within 1 hour	50%	76%
Lactate measured	94%	100%
Blood cultures 2 sets within 6 hours	63%	83%



#### PATIENT STORY

31 yo male presented at 1617 with fevers, headache and throat pain and cough, vomiting

Met sepsis criteria with high fever, tachycardia and hypotension (Sa02 >95% GCS 15)

Assigned Category 2 and sepsis pathway commenced immediately

Bloods (Blood cultures x2 and lactate) taken at	17 minutes
IV fluids commenced at	18 minutes
IV antibiotics at	18 minutes
Persistently hypotensive despite initial treatment	
Escalation of care at (resuscitation cubicle)	90 minutes
Medical registrar review	135 minutes
ICU registrar review	165 minutes
Transferred to HDU	180 minutes
Outcome	
ICU length of stay	2 days

This patient presented on a day whereby the ED recorded its maximum number of presentations.

# Benefits

- Process for change following identification of patient safety problem/incident
- Systematic methodical process to bring about change
- Proactive vs reactive approach
- Frontline clinician engagement
- Builds collaboration within and outside the ED
- Improvements can be identified and communicated
- Areas for further improvement could be identified



# Challenges

Takes time

Improvement science knowledge not widespread

Service improvement team support

Availability

Clinical perspective

Trust in the approach and understanding change

Measurement

Support and resourcing of project teams

Engagement and support by leadership

Training

Building time to work on projects

Supporting clinical practice improvement in the emergency department to improve patient safety

- Increase knowledge and experience in improvement work
- Engage and support projects
- Advocate for resourcing of improvement projects
- Look for improvement opportunities both within and external to ED
- Shift focus from measurement to improvement work

### Project Team

- Project Lead :Margaret Daly (ANUM)
- Medical Project Lead: Anh Tran (Emergency Physician)
- Rachel Vorlander (Antimicrobial stewardship pharmacist)
- Prema Madaiah (RN)
- Stacey Paterson (RN)
- Abby Bean (Quality coordinator)



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