

A Modern Approach to the Fundamentals of Sepsis Recognition, Management and Performance Improvement

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Chris Horvat, MD MHA

Improving Patients Safety and Quality in Latvia

Riga Stradins University

June 7, 2018

A Modern Approach to the Fundamentals of Sepsis Recognition, Management and Performance Improvement

Chris Horvat, MD MHA

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CRITICAL CARE MEDICINE

Conflict of Interest Disclosures

Children's Hospital of Pittsburgh Young Investigator Award



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Outline

American College of Critical Care Medicine Clinical Practice Parameters for Hemodynamic Support of Pediatric and Neonatal Septic Shock

Critical Care Medicine

June 2017 • Volume 45 • Number 6

ACCM Guidelines 2017



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Outline

Recognition Bundle (see AAP Trigger tool example Figure 2)

- Screen patient for septic shock using an institution trigger tool.
- Clinician assessment within 15 minutes for any patient who screens positive in the trigger tool.
- Initiate *Resuscitation Bundle* within 15 minutes for patient identified by the trigger tool whom the assessing clinician confirms suspicion of septic shock.

Resuscitation Bundle (see Algorithm Figure 3 and 4)

- Attain IV/IO access within 5 minutes.
- Appropriate fluid resuscitation begun within 30 minutes.
- Initiation of broad-spectrum empiric antibiotics within 60 minutes.
- Begin peripheral or central inotrope infusion therapy for fluid-refractory shock within 60 minutes.

Stabilization Bundle (see Algorithm Figure 3 and 4)

- Use multimodal monitoring to optimize fluid, hormonal, and cardiovascular therapies to attain hemodynamic goals.
- Confirm administration of appropriate antimicrobial therapy and source control.

Performance Bundle

- Measure adherence to Trigger, Resuscitation, and Stabilization Bundles.
- Perform root cause analysis to identify barriers to adherence.
- Provide an action plan to address identified barriers.

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Outline

- 1) **Historic perspective:** Foundation of resuscitation
- 2) **Recognition:** Augmenting clinical evaluation
- 3) **Resuscitation:** Guidelines for individualization
- 4) **Stabilization:** Bolstering clinical assessment
- 5) **Performance:** Collaboration and tracking



Dr. Peter Safar

Rules for Navigating Life

Rule No. 16

“When in doubt, THINK!”

Outline

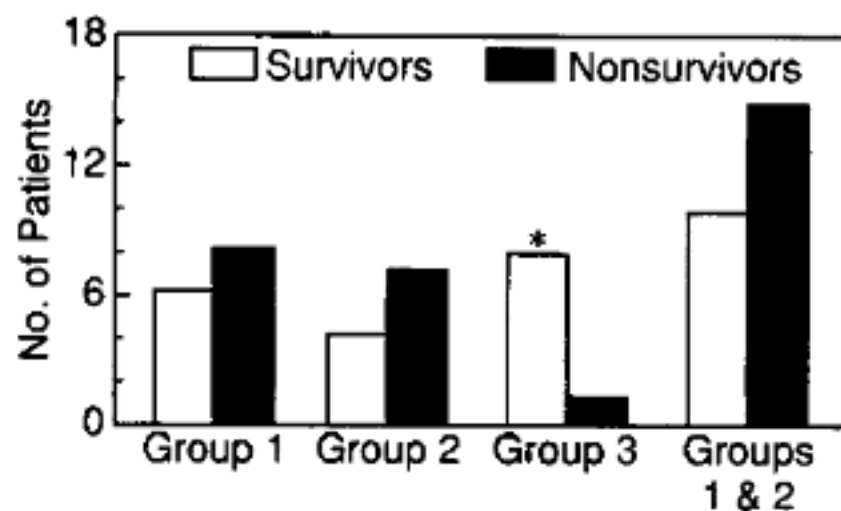
- 1) **Historic perspective: Foundation of resuscitation**
- 2) **Recognition: Augmenting clinical evaluation**
- 3) **Resuscitation: Guidelines for individualization**
- 4) **Stabilization: Bolstering clinical assessment**
- 5) **Performance: Collaboration and tracking**

Role of Early Fluid Resuscitation in Pediatric Septic Shock

Joseph A. Carcillo, MD; Alan L. Davis, MD; Arno Zaritsky, MD

JAMA, September 4, 1991 — Vol 266, No. 9

	1 h (mean \pm SD)	6 h (mean \pm SD)
Group 1 (n = 14; <20 mL/kg in 1 h)	11 \pm 6*	71 \pm 29†
Group 2 (n = 11; 20-40 mL/kg in 1 h)	32 \pm 5*	108 \pm 54
Group 3 (n = 9; >40 mL/kg in 1 h)	69 \pm 19*	117 \pm 29
All patients (n = 34)	33 \pm 26	95 \pm 42
Survivors (n = 18)	42 \pm 28†	97 \pm 49
Nonsurvivors (n = 16)	23 \pm 18	94 \pm 37



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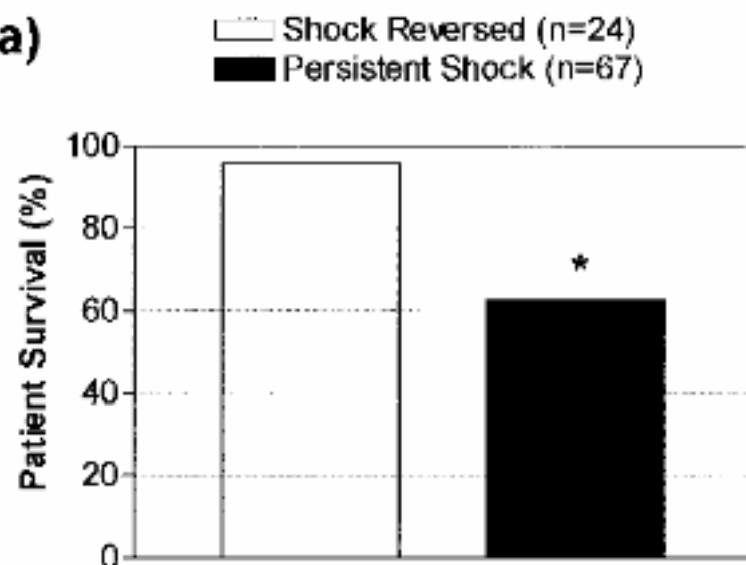
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Early Reversal of Pediatric-Neonatal Septic Shock by Community Physicians Is Associated With Improved Outcome

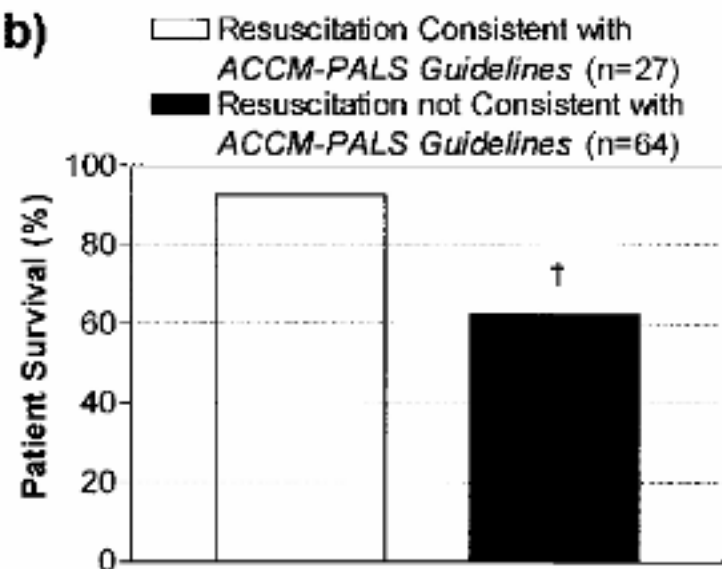
Yong Y. Han, MD*§; Joseph A. Carcillo, MD*‡§; Michelle A. Dragotta, RN§; Debra M. Bills, RN§; R. Scott Watson, MD, MPH*‡§; Mark E. Westerman, RT§; and Richard A. Orr, MD*‡§

PEDIATRICS Vol. 112 No. 4 October 2003 793

a)



b)



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Delayed Antimicrobial Therapy Increases Mortality and Organ Dysfunction Duration in Pediatric Sepsis*

Scott L. Weiss, MD¹; Julie C. Fitzgerald, MD, PhD¹; Fran Balamuth, MD, PhD²;
Elizabeth R. Alpern, MD, MSCE³; Jane Lavelle, MD²; Marianne Chilutti, MS⁴;
Robert Grundmeier, MD^{4,5}; Vinay M. Nadkarni, MD, MS¹; Neal J. Thomas, MD, MSc⁶

TABLE 5. PICU Mortality: Sepsis Recognition to Initial Antimicrobial Administration

Time to Initial Antibiotics (hr)	No. of Patients	% Mortality	% Difference	Unadjusted OR	95% CI
≤ 1	24	8	5	1.67	0.35–7.91
> 1	106	13			
≤ 2	55	7	10	2.43	0.74–7.99
> 2	75	17			
≤ 3	78	6	17	3.92	1.27–12.06
> 3	52	23			
≤ 4	91	8	15	3.60	1.23–10.52
> 4	39	23			

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The NEW ENGLAND JOURNAL *of* MEDICINE

ESTABLISHED IN 1812

JUNE 30, 2011

VOL. 364 NO. 26

Mortality after Fluid Bolus in African Children with Severe Infection

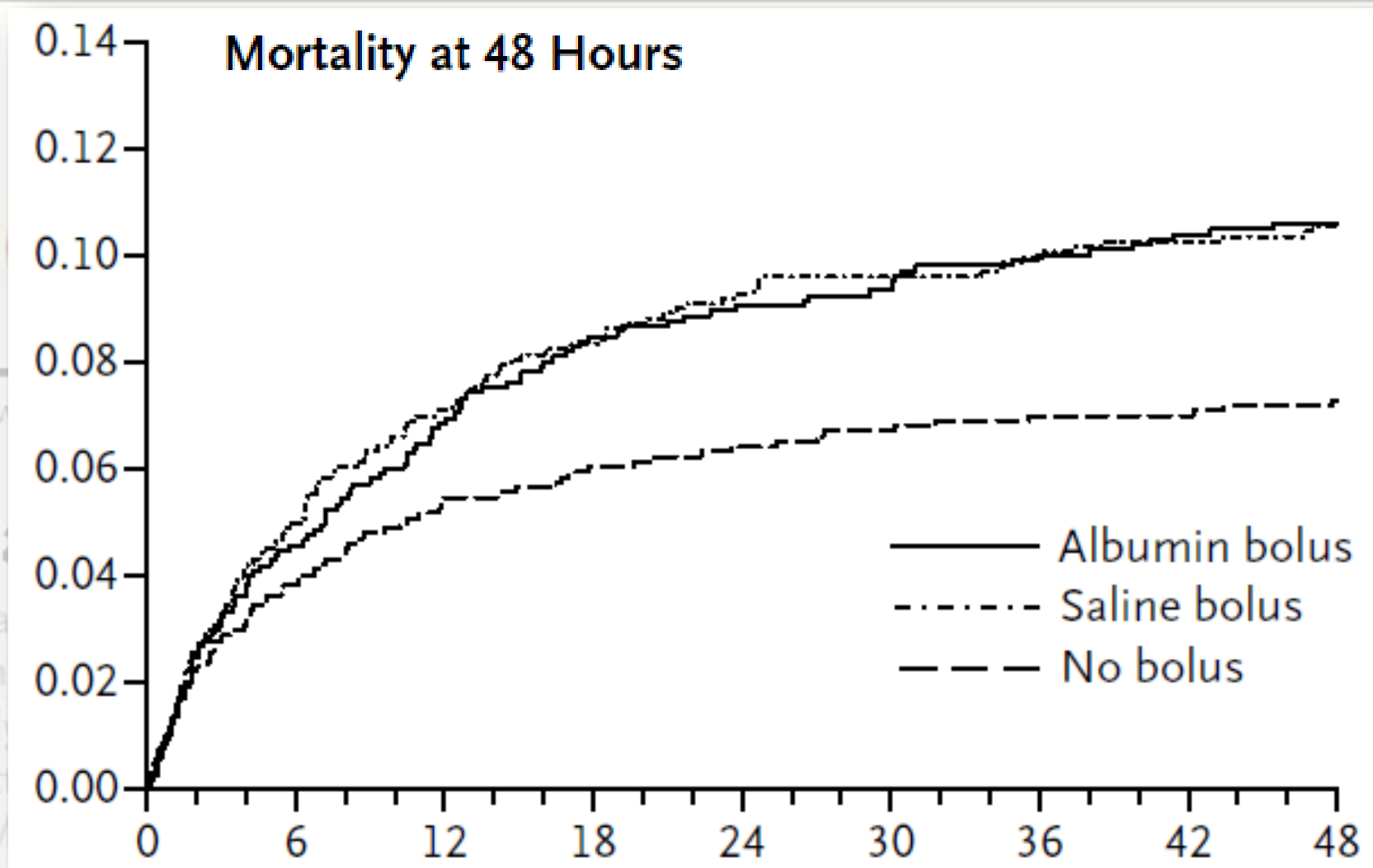
Kathryn Maitland, M.B., B.S., Ph.D., Sarah Kiguli, M.B., Ch.B., M.Med., Robert O. Opoka, M.B., Ch.B., M.Med., Charles Engoru, M.B., Ch.B., M.Med., Peter Olupot-Olupot, M.B., Ch.B., Samuel O. Akech, M.B., Ch.B., Richard Nyeko, M.B., Ch.B., M.Med., George Mtove, M.D., Hugh Reyburn, M.B., B.S., Trudie Lang, Ph.D., Bernadette Brent, M.B., B.S., Jennifer A. Evans, M.B., B.S., James K. Tibenderana, M.B., Ch.B., Ph.D., Jane Crawley, M.B., B.S., M.D., Elizabeth C. Russell, M.Sc., Michael Levin, F.Med.Sci., Ph.D., Abdel G. Babiker, Ph.D., and Diana M. Gibb, M.B., Ch.B., M.D., for the FEAST Trial Group*



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Table 1. (Continued.)

Variable	Albumin Bolus (N = 1050)	Saline Bolus (N = 1047)	No Bolus (N = 1044)	Total (N = 3141)
Laboratory assessments††				
Positive for malaria parasitemia — no./total no. (%)‡‡	590/1044 (57)	612/1042 (59)	591/1037 (57)	1793/3123 (57)
Hemoglobin — no./total no. (%)				
<5 g/dl	323/1024 (32)	332/1015 (33)	332/1015 (33)	987/3054 (32)
>10 g/dl	231/1024 (23)	230/1015 (23)	244/1015 (24)	705/3054 (23)

Charles Engoru, M.B., Ch.B., M.Med., Peter Olupot-Olupot, M.B., Ch.B., Samuel O. Akech, M.B., Ch.B.,
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SHOCK, Vol. 43, No. 1, pp. 68–73, 2015

**FLUID OVERLOAD IN PATIENTS WITH SEVERE SEPSIS AND
SEPTIC SHOCK TREATED WITH EARLY GOAL-DIRECTED THERAPY IS
ASSOCIATED WITH INCREASED ACUTE NEED FOR FLUID-RELATED
MEDICAL INTERVENTIONS AND HOSPITAL DEATH**

**Diana J. Kelm,^{*†} Jared T. Perrin,^{*} Rodrigo Cartin-Ceba,^{*†} Ognjen Gajic,^{*†}
Louis Schenck,[‡] and Cassie C. Kennedy^{*†}**

^{}Department of Internal Medicine and Divisions of [†]Pulmonary and Critical Care and [‡]Biomedical Statistics
and Informatics, Mayo Clinic, Rochester, Minnesota*

Received 28 Aug 2014; first review completed 15 Sep 2014; accepted in final form 17 Sep 2014

TABLE 3. Medical interventions and secondary outcomes in those with clinical evidence of fluid overload: univariate and multivariate analyses

Univariate analyses	Clinical evidence of fluid overload day 1 (n = 272)	Clinical evidence of persistent fluid overload (n = 182)
	Medical interventions, OR (95% CI)	
Thoracentesis ^a	3.38 (1.28–8.95)	3.10 (1.44–6.68)
Paracentesis [†]	0.58 (0.19–1.77)	0.57 (0.17–1.94)
Ultrafiltration [‡]	1.41 (0.66–3.02)	2.17 (1.06–4.43)
Diuretics [§]	1.15 (0.71–1.86)	1.77 (1.09–2.87)
	Secondary outcomes	
ICU LOS, mean difference, d	0.54 (–0.38 to 1.46)	0.39 (–0.51 to 1.30)
Hospital LOS, mean difference, d	0.99 (–2.25 to 4.22)	1.55 (–1.64 to 4.74)
30-d ICU readmission, OR (95% CI)	1.07 (0.62–1.84)	1.40 (0.839–2.34)
Hospital mortality, OR (95% CI)	2.33 (1.34–4.05)	1.89 (1.16–3.09)
Multivariate analyses	Medical interventions, OR (95% CI)	
Thoracentesis ^a	3.40 (1.37–10.3)	3.83 (1.74–9.15)
Ultrafiltration [‡]	—	1.90 (0.90–4.19)
Diuretics [§]	—	1.65 (1.00–2.72)
	Secondary outcomes	
30-d ICU readmission, OR (95% CI) [†]	—	1.61 (0.94–2.79)
Hospital mortality, OR (95% CI) [†]	2.27 (1.31–4.09)	1.92 (1.16–3.22)

*Departm



Early, Goal-Directed versus Usual Care

Rivers et al. EGTD NEJM 2001

ProCESS NEJM 2014

ARISE NEJM 2014

ProMISe NEJM 2015



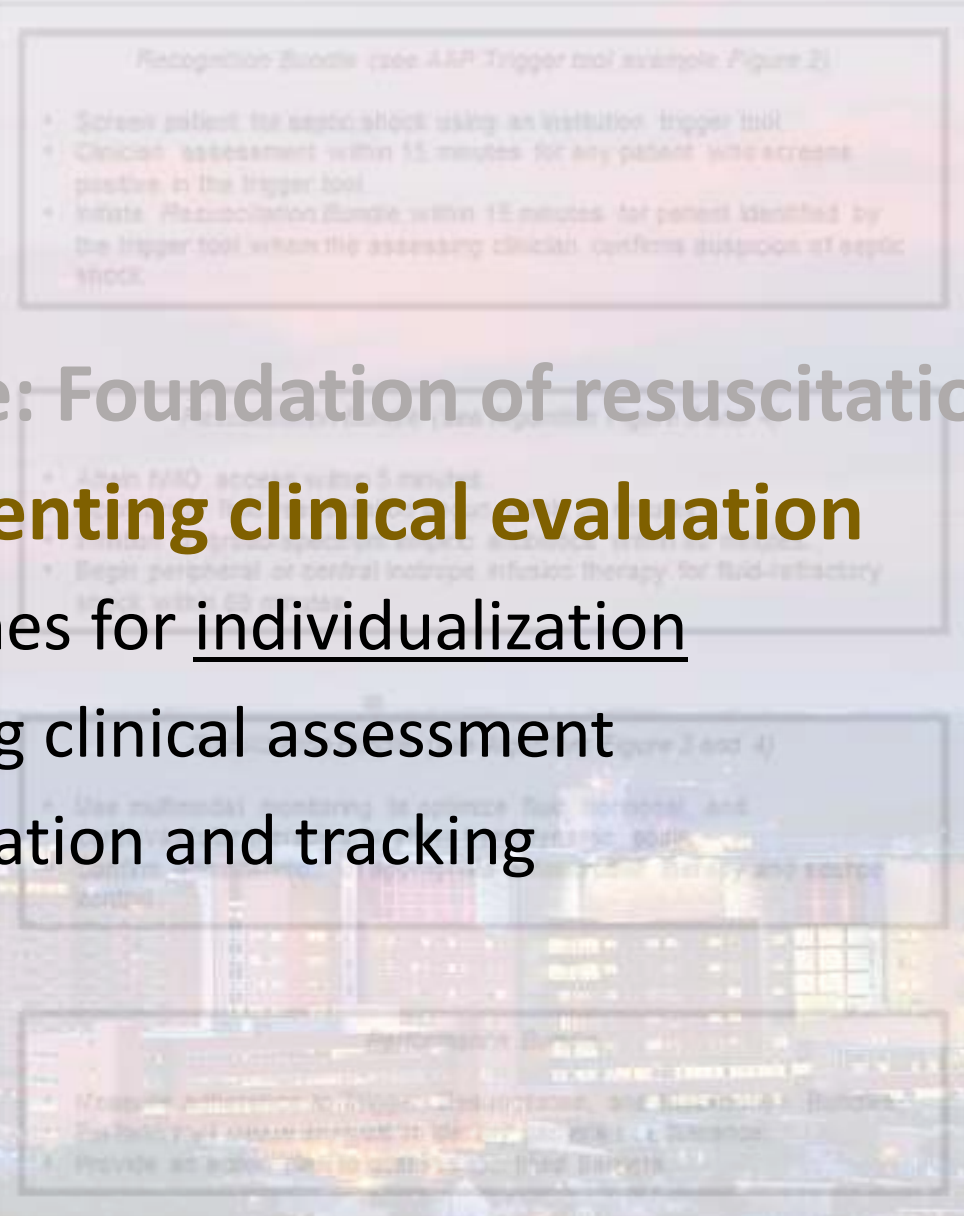
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Outline

- 1) Historic perspective: Foundation of resuscitation
- 2) Recognition: Augmenting clinical evaluation**
- 3) Resuscitation: Guidelines for individualization**
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- 5) Performance: Collaboration and tracking**



GET AHEAD

OF SEPSIS

KNOW THE RISKS. SPOT THE SIGNS. ACT FAST.



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CRITICAL CARE MEDICINE

Implementation of Goal-Directed Therapy for Children With Suspected Sepsis in the Emergency Department

abstract

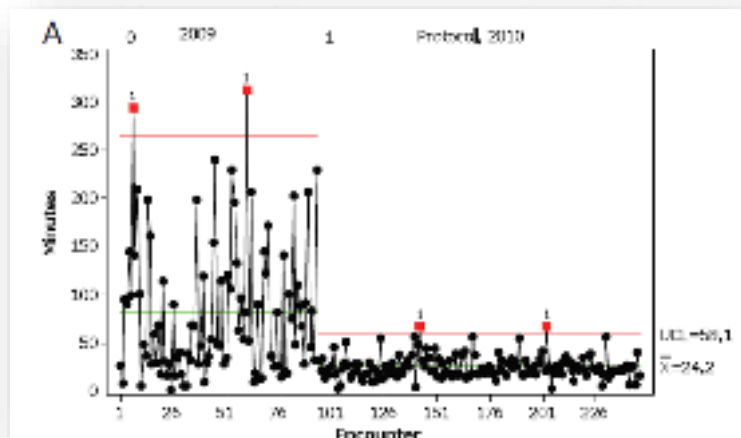
BACKGROUND: Suboptimal care for children with septic shock includes delayed recognition and inadequate fluid resuscitation.

OBJECTIVE: To describe the implementation of an emergency department (ED) protocol for the recognition of septic shock and facilitate

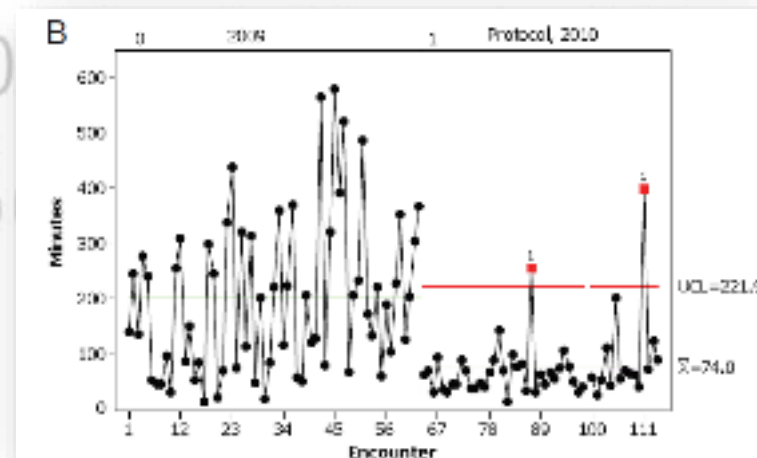
AUTHORS: Andrea T. Cruz, MD, MPH,^{a,b} Andrew M. Perry, MD,^a Eric A. Williams, MD, MS,^c Jeanine M. Graf, MD,^c Elizabeth R. Wuestner, MSN, RN,^d and Binita Patel, MD^a

Sections of ^aEmergency Medicine, ^bInfectious Diseases, and ^cCritical Care Medicine, Department of Pediatrics, Baylor College of Medicine, Houston, Texas; and ^dEmergency Department, Texas Children's Hospital, Houston, Texas

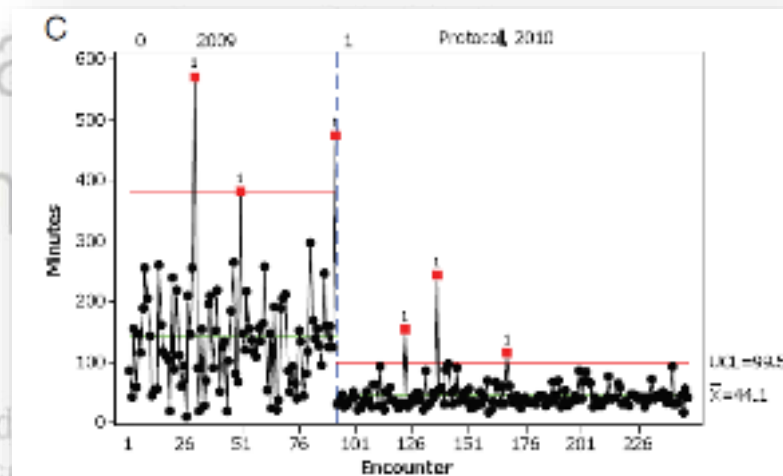
Time to 1st Bolus



Time to 3rd Bolus



Time to Antibiotic



BACKGROUND: Suboptimal care for children with septic shock includes delayed recognition and inadequate fluid resuscitation.

OBJECTIVE: To describe the implementation of an emergency department (ED) protocol for the recognition of septic shock and facilitate

MD,^a Eric A. Williams, MD, MD,^b Jeanine M. Graft, MD,^c Elizabeth R. Wuestner, MSN, RN,^d and Binita Patel, MD^a

Sections of ^aEmergency Medicine, ^bInfectious Diseases, and ^cCritical Care Medicine, Department of Pediatrics, Baylor College of Medicine, Houston, Texas; and ^dEmergency Department, Texas Children's Hospital, Houston, Texas

Improving Recognition of Pediatric Severe Sepsis in the Emergency Department: Contributions of a Vital Sign–Based Electronic Alert and Bedside Clinician Identification

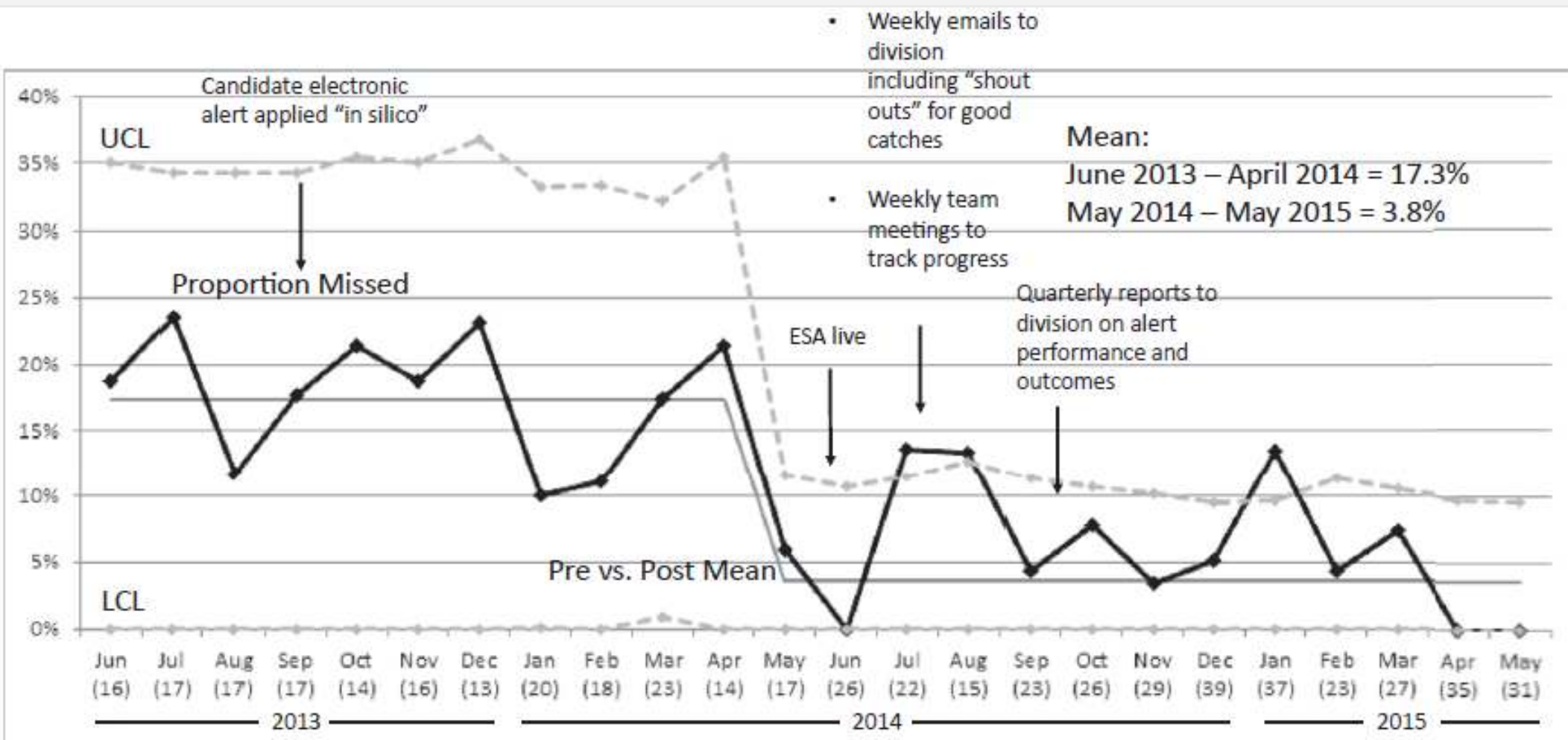


Fran Balamuth, MD, PhD*; Elizabeth R. Alpern, MD, MSCE; Mary Kate Abbadessa, MSN, RN; Katie Hayes, BS; Aileen Schast, PhD; Jane Lavelle, MD; Julie C. Fitzgerald, MD, PhD; Scott L. Weiss, MD, MSCE; Joseph J. Zorc, MD, MSCE

**Corresponding Author. E-mail: balamuthf@email.chop.edu.*

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Annals of Emergency Medicine 759

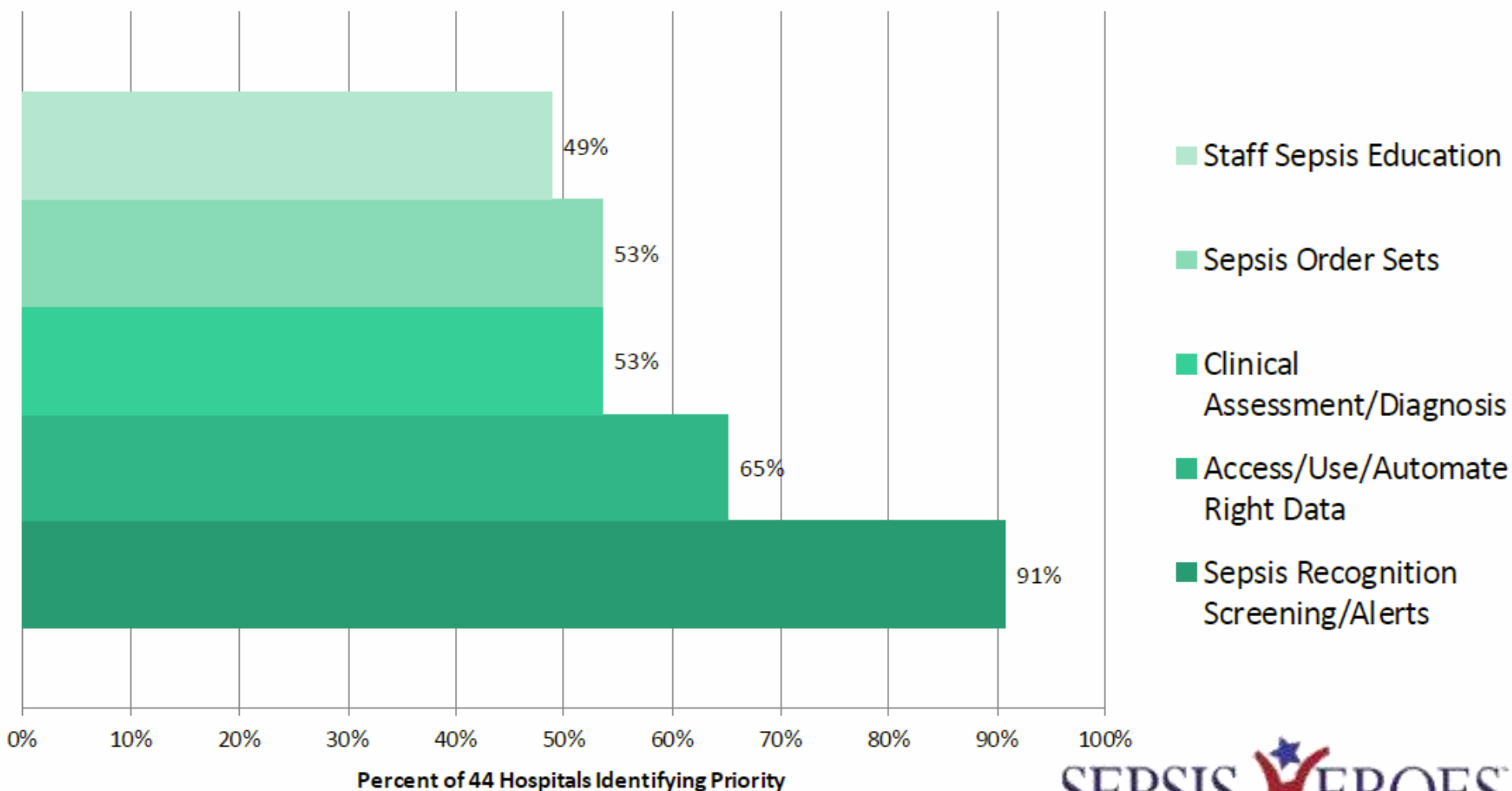


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Top Sepsis Improvement Priorities at IPSO Hospitals in Initial 12 Months of Participation



SEPSIS XEROES



Table 1. Criteria for positive sepsis screen. Positive screen = hypotension alone or ≥ 3 of the following 8 criteria:

1. Fever or hypothermia
2. Tachycardia
3. Tachypnea
4. Cap refill: ≥ 3 sec or < 1 sec
5. Pulse: decreased or bounding
6. Skin: mottled, flushed or petechiae/purpura
7. Mental status: depressed, highly irritable, confused
8. Presence of a high-risk medical condition:
 - Age < 1 mo
 - Severe developmental delay/intellectual disability
 - Central line (Mediport, Broviac, PICC)
 - Malignancy
 - Transplant
 - Asplenia (including Sickle Cell Disease)
 - Other immunocompromised

Age	Temp C	HR	RR	Systolic BP
0 d - 1 m	< 36.5 or > 38	> 205	> 60	< 60
≥ 1 m - 3 m	< 36 or > 38	> 205	> 60	< 70
≥ 3 m - 1 y	< 36 or > 38.5	> 190	> 60	< 70
1 y	< 36 or > 38.5	> 190	> 40	< 72
2 y	< 36 or > 38.5	> 140	> 40	< 74
3 y	< 36 or > 38.5	> 140	> 40	< 76
4 y	< 36 or > 38.5	> 140	> 34	< 78
5 y	< 36 or > 38.5	> 140	> 34	< 80
6 y	< 36 or > 38.5	> 140	> 30	< 82
7 y	< 36 or > 38.5	> 140	> 30	< 84
8 y	< 36 or > 38.5	> 140	> 30	< 86
9 y	< 36 or > 38.5	> 140	> 30	< 88
≥ 10 y - 13 y	< 36 or > 38.5	> 120	> 30	< 90
> 13	< 36 or > 38.5	> 110	> 20	< 90



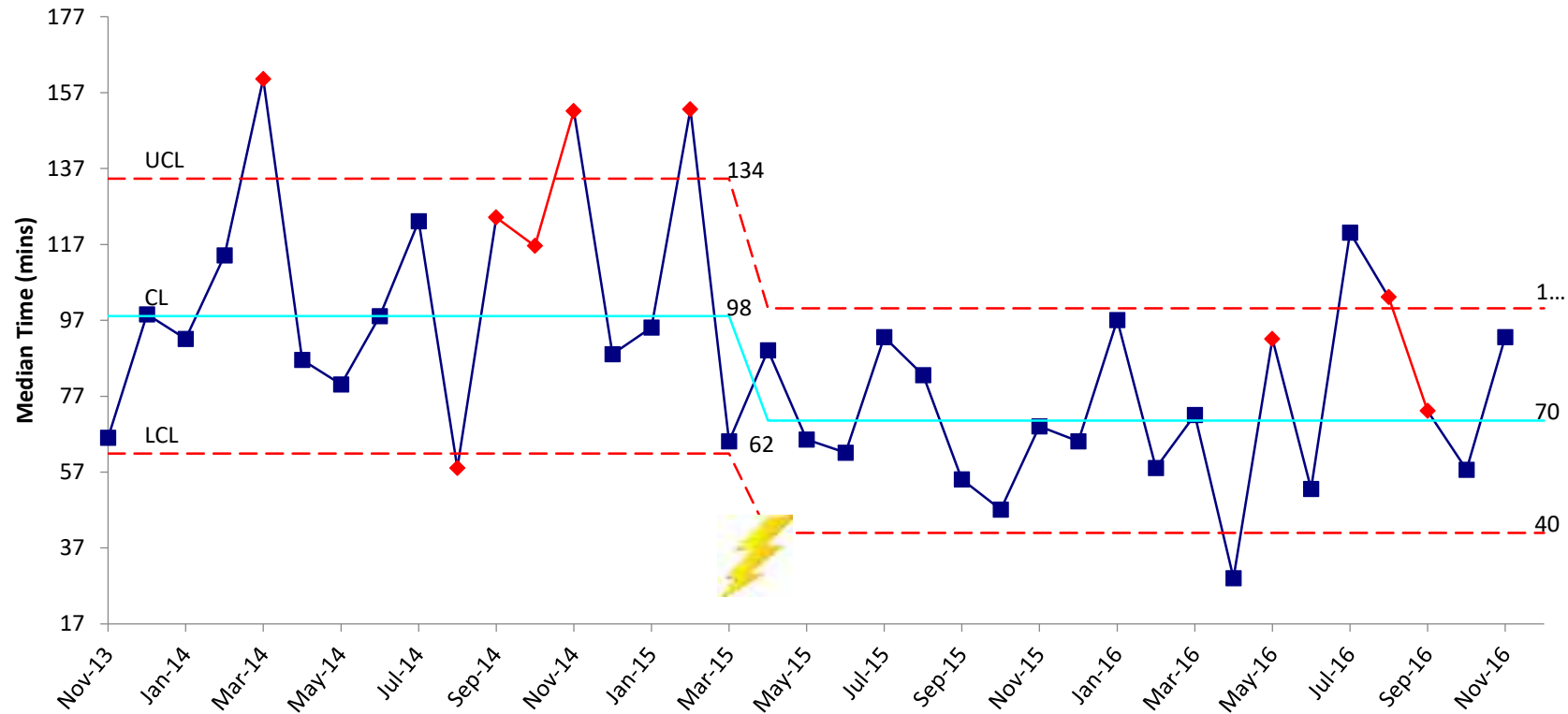
The Lightning Bolt Makes a Difference



Median Time to Antibiotics Control Chart

Goal is 60 minutes

Bolt started in March, 2015



Nov, 2013 – Dec, 2016

Outline

- 1) Historic perspective: Foundation of resuscitation
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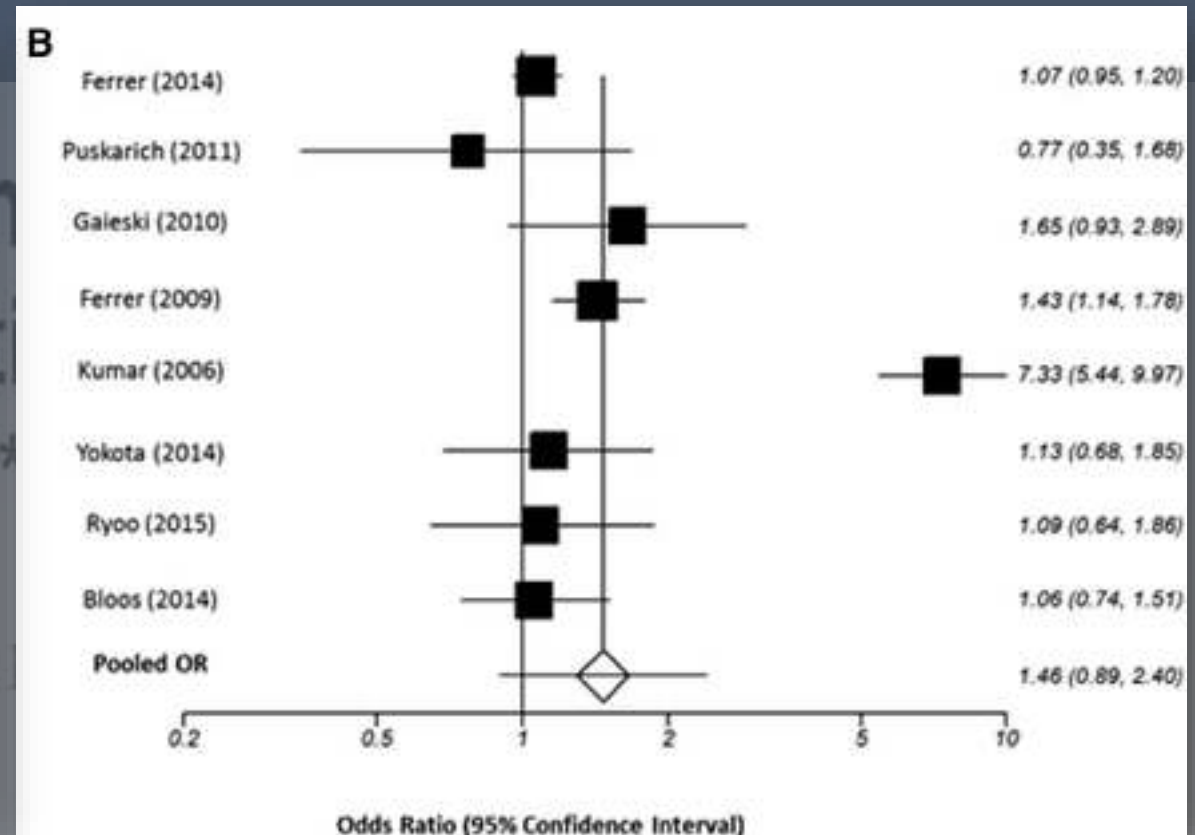
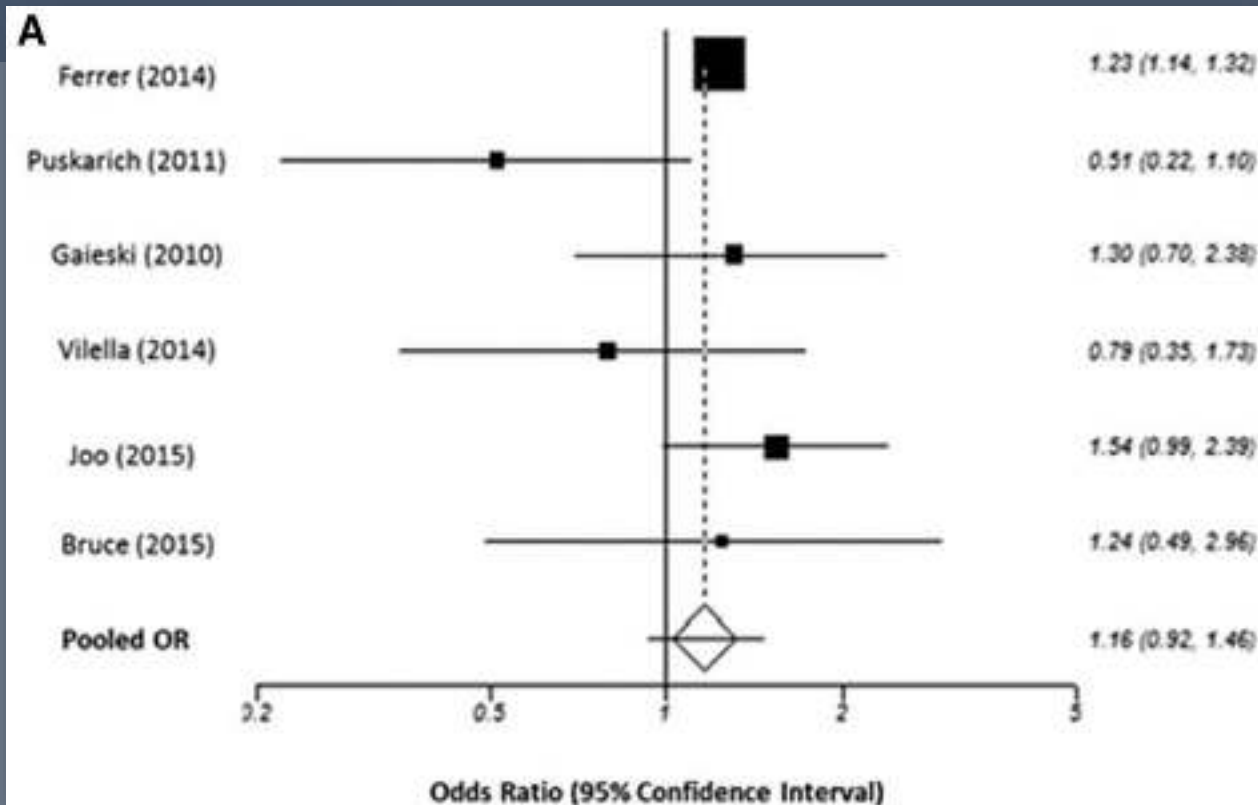
Time to Antibiotics

The Impact of Timing of Antibiotics on Outcomes in Severe Sepsis and Septic Shock: A Systematic Review and Meta-Analysis*

Sarah A. Sterling, MD; W. Ryan Miller, MD; Jason Pryor, MD; Michael A. Puskarich, MD;
Alan E. Jones, MD

Critical Care Medicine September 2015 • Volume 43 • Number 9

Time to Antibiotics



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Yes, but...



Video Credit: Daniel Izzo. *Bacteria Growth*. <https://www.youtube.com/watch?v=gEwzDydcIWc>.



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Guidelines - Time to Antibiotics

- 2012 ESIC/SCCM Guidelines - Administer antibiotics within 1 hour
- 2015 ESIC/SCCM Update - Administer antibiotics within 3 hours
- 2016 Update - Administer antibiotics within 1 hour

Time to Antibiotics in Major Clinical Trials



Rivers et al. EGTD NEJM
- Majority within 6 hours

Kumar et al. CCM
- Median of 6 [IQR 2-15] hrs

ProCESS NEJM
- Majority within 3 ± 1.75 hrs

ARISE NEJM
- Median of 70 [38-114] min

ProMISE NEJM
- All by median of 2.5 [1.8-3.5] hrs

Credit: Michael Alison, MD; Twitter 2015



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Time to Antibiotics in Major Clinical Trials

Rivers et al. – Majority within 6 hours

Kumar et al. – Median of 6 hours

ProCESS – Majority within 3 hours

ARISE – Median of 70 minutes

ProMISe – Median of 2.5 hours



Credit: Michael Alison, MD; Twitter 2015



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Factors Influencing Antibiotic Delivery

Patient and Organizational Factors Associated With Delays in Antimicrobial Therapy for Septic Shock*

Andre C. K. B. Amaral, MD¹; Robert A. Fowler, MDCM, MS(Epi), FRCPC¹; Ruxandra Pinto, PhD¹; Gordon D. Rubenfeld, MD, MSc¹; Paul Ellis, MD²; Brian Bookatz, MD³; John C. Marshall, MD, FRCSC, FACS⁴; Greg Martinka, MD⁵; Sean Keenan, MD⁶; Denny Laporta, MD⁷; Daniel Roberts, MD⁸; Anand Kumar, MD⁸; and the Cooperative Antimicrobial Therapy of Septic Shock Database Research Group

Critical Care Medicine

December 2016 • Volume 44 • Number 12



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Patient
Delays

ted With
Shock*

Andre C
Gordon I
Greg Ma
and the

Pinto, PhD¹;
D, FRCSC, FACS¹;
and Kumar, MD¹;

Variable	Difference in Minutes to a Standard Patient	% Change in Time to Antimicrobial	p
Year of entry in the cohort, per year	-9	-3.23 (-4.22 to -2.22)	<0.0001
Academic hospital ^a	+52	21.57 (3.13-43.34)	0.0224
Acute Physiology Score, per 5 points	+24	9.06 (6.63-11.53)	<0.0001
Temperature, per °C	-15	-5.68 (-7.62 to -3.89)	<0.0001
Age, per 10 yr	+16	5.94 (3.45-8.49)	<0.0001
Pre-shock length of stay ^b			
≤ 3 d	+50	25.34 (13.79-38.07)	<0.0001
Between 3 and 7 d	+121	61.55 (38.11-88.97)	<0.0001
> 7 d	+130	66.25 (44.88-90.79)	<0.0001
Immunosuppression	+18	6.92 (-1.595 to 16.16)	0.11407
Sex, male	-15	-5.41 (-11.90 to 1.56)	0.12517
Comorbidity	+85	14.15 (5.82-23.13)	0.00062
Community-acquired infection ^c	-53	-18.11 (-26.57 to -8.68)	0.00033
Hypertension	-33	-11.73 (-19.34 to -3.40)	0.00672
Transferred from ^d			
Emergency department	-47	-17.71 (-28.17 to -5.73)	0.00495
External hospital	+17	6.26 (-7.82 to 22.48)	0.40257
Medicine ward	+39	14.52 (1.32-29.45)	0.02999
Primary infection ^e			
Gastrointestinal	+18	7.64 (-6.87 to 24.42)	0.31884
Other	+35	14.66 (-0.11 to 31.66)	0.05189
Pneumonia	+45	18.69 (5.34-33.74)	0.00493
Need for source control	+19	7.425 (-3.34 to 19.39)	0.18361

Time to Antibiotics

1. Antibiotics ordered (written, verbal or EHR)
2. Order received by pharmacy
3. Dose and indication verification
4. Order prepared
5. Transported to nursing unit/bedside
6. Infusion pump prepared
7. Antibiotic initiated (infusion completed 30 minutes later)

Time to Antibiotics (Our Old Approach)

1. Meropenem ordered (written, verbal or EHR)
 - Resident/housestaff place order while senior staff resuscitate
2. Order received by pharmacy
3. Dose and indication verification
 - Phone call to verify indication given stewardship policies
4. Order prepared
5. Transported to nursing unit/bedside via tube station
 - Antibiotic waits in the station until a nurse is able to momentarily step away from the resuscitation
 - Eventually brought to bedside and set down in order to prepare vasopressors
6. Infusion pump prepared
7. Antibiotic initiated after verification (infusion completed 30 minutes later)

Time to Antibiotics (Current Approach)



Acute Care Pediatric Sepsis and Sepsis Shock

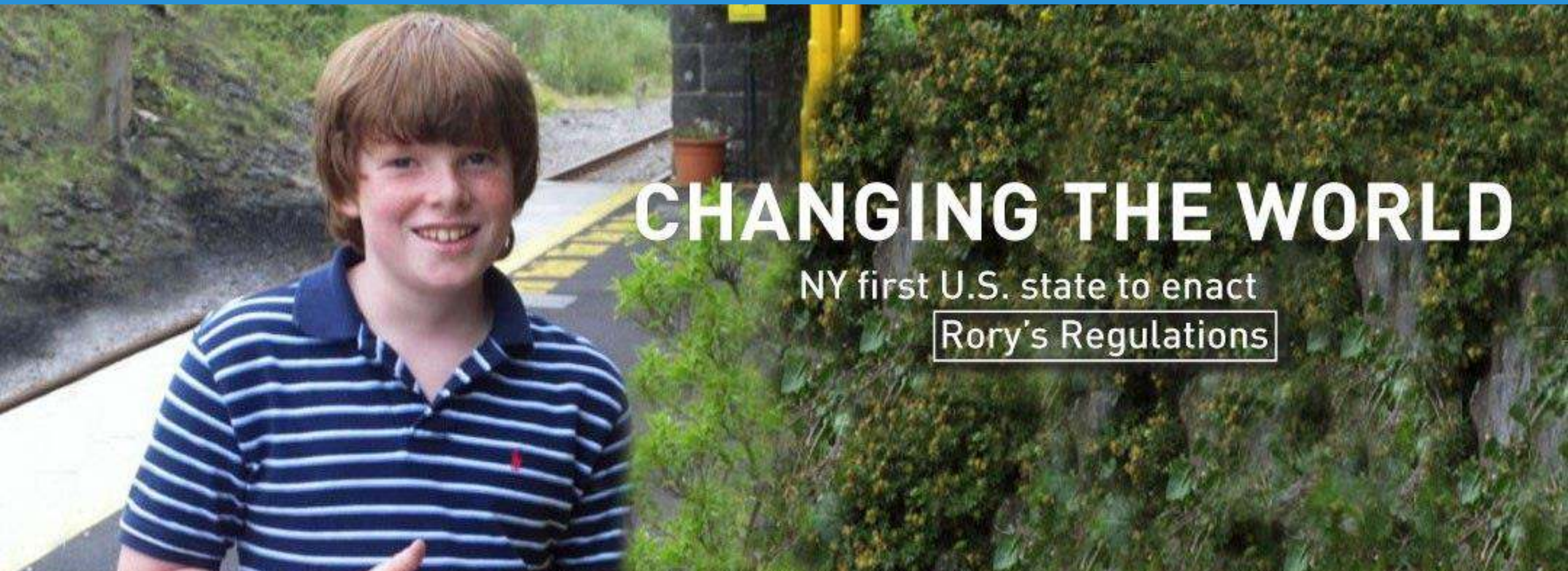
Clinical Effectiveness Guideline

PATIENT POPULATION	RECOMMENDED FIRST-DOSE ANTIBIOTICS
For patients > 60 days old and: - Fever without a source - Sickle cell disease - Neurosurgery patients - MSK (Ortho, Plastics, local MSK finding) - Renal, Urology, Renal Transplant, GU procedures - Meningitis/CNS disease (Use CNS dosing) - Pneumonia with/without effusion/empyema - Cardiac, CT Surg, Heart Transplant	Cefepime 50 mg/kg IV (max 2000 mg) Vancomycin 15 mg/kg IV (max 1500 mg) Consider anaerobic coverage if concern for aspiration
If allergy to cephalosporin/penicillin and not DMT patient (see below for allergy in DMT patient)	Meropenem 20 mg/kg (max 2000 mg) instead of cephalosporin/PCN Discuss alternatives with ID

- Sepsis recognition triggers bedside huddle
- One time dose of empiric, risk-factor based antibiotic ordered
- Phone call placed to pharmacy notifying patient has sepsis
- Antibiotic delivered to bedside and prioritized for initiation
- Reduction in time to abx to median of 1 hr (Still room to improve!)

THE RORY STAUNTON FOUNDATION

FOR SEPSIS PREVENTION



CHANGING THE WORLD

NY first U.S. state to enact

Rory's Regulations



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Time to Bundled Care

The NEW ENGLAND JOURNAL of MEDICINE

ORIGINAL ARTICLE

Time to Treatment and Mortality during Mandated Emergency Care for Sepsis

Christopher W. Seymour, M.D., Foster Gesten, M.D., Hallie C. Prescott, M.D.,
Marcus E. Friedrich, M.D., Theodore J. Iwashyna, M.D., Ph.D.,
Gary S. Phillips, M.A.S., Stanley Lemeshow, Ph.D., Tiffany Osborn, M.D., M.P.H.,
Kathleen M. Terry, Ph.D., and Mitchell M. Levy, M.D.



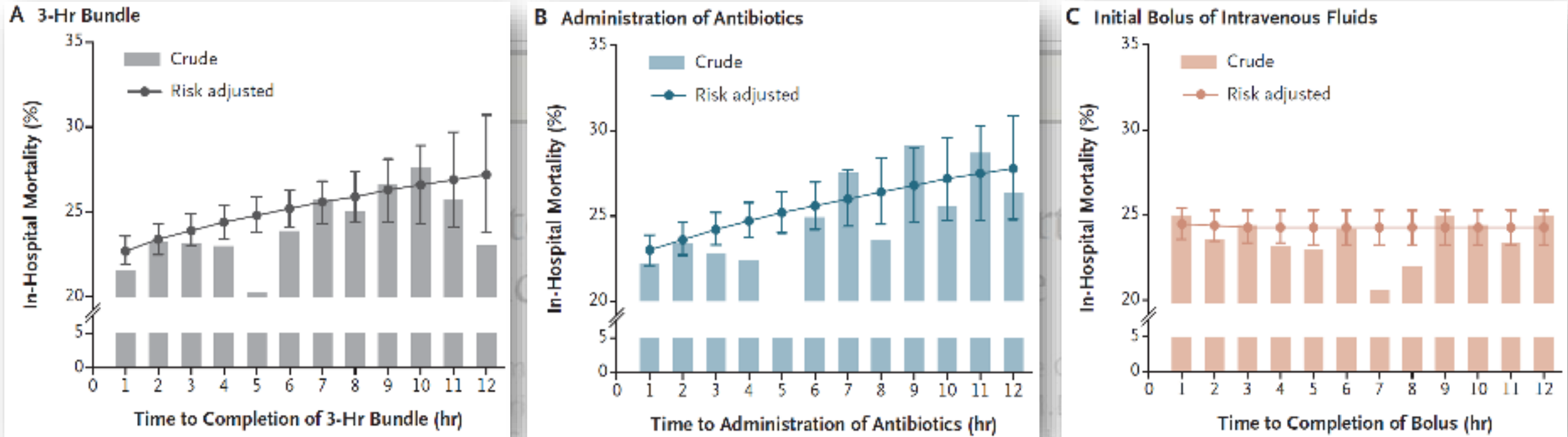
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Time to Bundled Care

The NEW ENGLAND JOURNAL of MEDICINE



Gary S. Phillips, M.A.S., Stanley Lemeshow, Ph.D., Hilary Osborn, M.D., M.P.H., Kathleen M. Terry, Ph.D., and Mitchell M. Levy, M.D.

THE RORY STAUNTON FOUNDATION

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Research Letter

October 2017

Epidemiology of Sepsis Among Adolescents in a Tertiary Care Community Hospital Emergency Department: Implications for Rory's Regulations

Idris V. R. Evans, MD, MSc^{1,2}; R. Scott Watson, MD, MPH³; Joseph Carcillo, MD^{1,2}; et al

» Author Affiliations | Article Information

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Table Characteristics of 899 Healthy Adolescents Presenting With 3 or More SIRS Criteria^a

Characteristic	Patients With Sepsis (n = 158)		Patients Without Sepsis (n = 741)
	Poor Course (n = 43)	No Poor Course (n = 115)	
Time until sepsis suspected, h			
≤24	35 (81.4)	112 (97.4)	0
>24	8 (18.6)	3 (2.6)	0
Age, mean (SD), y	16 (2)	16 (2)	16 (2)
Male sex	21 (48.8)	40 (34.8)	270 (36.4)
Race			
White	26 (60.5)	81 (70.4)	533 (71.8)
Black	10 (23.3)	22 (19.1)	134 (18.1)
Other	7 (16.3)	12 (10.4)	74 (10.0)
Positive blood culture results	4 (9.3)	2 (1.7)	0
All 4 SIRS criteria in 24 h	13 (30.2)	30 (26.1)	86 (11.6)
Maximum SOFA points in 24 h			
0	14 (32.6)	70 (60.9)	507 (68.4)
1	4 (9.3)	26 (22.6)	117 (15.8)
≥2	25 (58.1)	19 (16.5)	117 (15.8)
Hospital admission	26 (60.5)	30 (26.1)	180 (24.3)
Poor course			
Admission to intensive care	25 (58.1)	0	30 (4.0)
Transfer to acute care facility or hospital	27 (62.8)	0	83 (11.1)
In-hospital mortality	1 (2.3)	0	0
Serum lactate within 24 h	9 (20.9)	3 (2.6)	0
Serum lactate, median (IQR), mg/dL	16.2 (9.0-24.3)	9.0 (8.1-17.2)	0
Mechanical ventilation ^b	18 (41.9)	0	12 (1.6)
Vasopressor use ^b	4 (9.3)	0	0
Hospital length of stay, median (IQR), d	7 (1-15)	2 (1-3)	2 (1-2)



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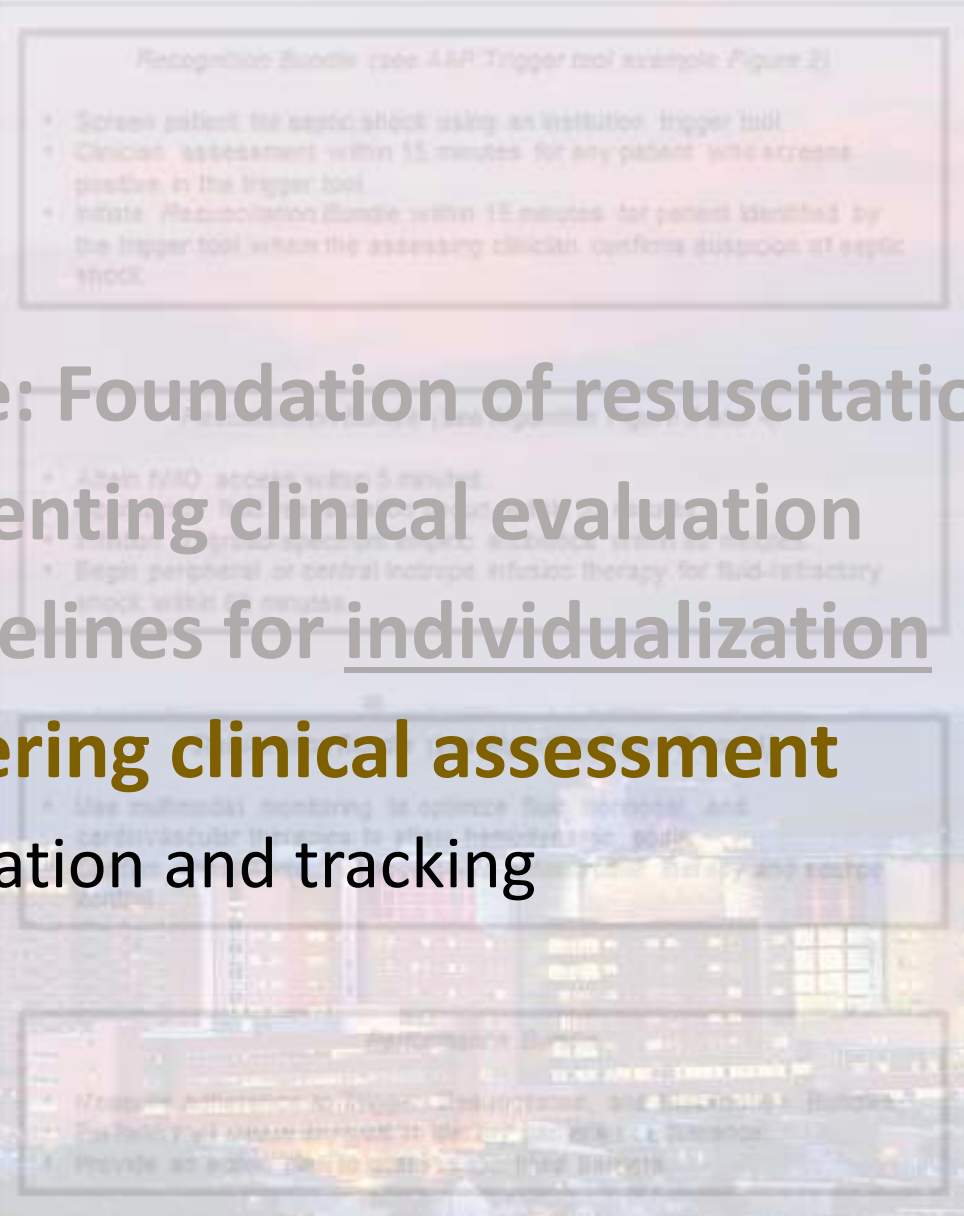
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Outline

- 1) Historic perspective: Foundation of resuscitation
- 2) Recognition: Augmenting clinical evaluation
- 3) Resuscitation: Guidelines for individualization
- 4) Stabilization: Bolstering clinical assessment**
- 5) Performance: Collaboration and tracking**



708 JAMA August 18, 2015 Volume 314, Number 7

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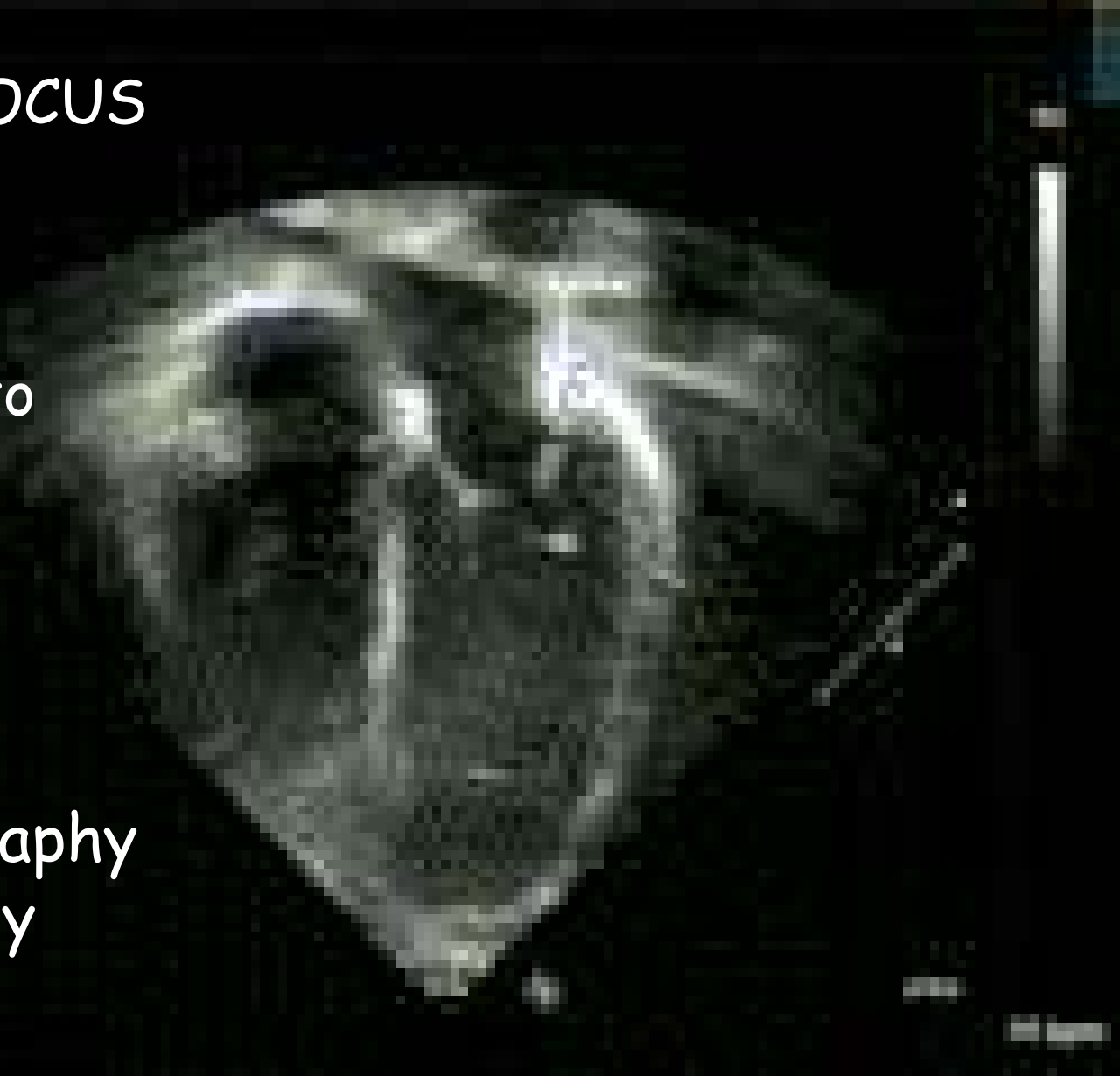
graph TD
    Start([Start]) --> DataCollection[Data Collection]
    DataCollection --> DataManagement[Data Management]
    DataManagement --> Decision1{Investigation Required?}
    Decision1 -- Yes --> DataCollection2[Data Collection]
    DataCollection2 --> DataManagement2[Data Management]
    DataManagement2 --> DataAnalysis[Data Analysis]
    DataAnalysis --> Reporting[Reporting]
    Reporting --> End([End])
    Decision1 -- No --> End
  
```

The flowchart illustrates the proposed decision support system for vehicle accident investigation. It begins with a 'Start' node, leading to 'Data Collection'. This step involves collecting data from various sources, including police reports, witness statements, and vehicle data. The collected data is then moved to 'Data Management', where it is organized and stored. A decision point follows: 'Investigation Required?'. If the answer is 'Yes', the process continues to 'Data Collection' (recollecting more data), then 'Data Management', and finally 'Data Analysis'. The analysis involves identifying causes, determining liability, and assessing damages. The results are then 'Reported' to the relevant authorities. If the answer to the decision point is 'No', the process proceeds directly to the 'End' node.



"In the right clinical context, [the FOCUS exam] can direct the clinician at the bedside in important next treatment interventions, optimize diagnostic efficiency, and assess the response to performed interventions"

-Labovitz et al. 2010
A Consensus Statement of the
American Society of Echocardiography
and American College of Emergency
Physicians

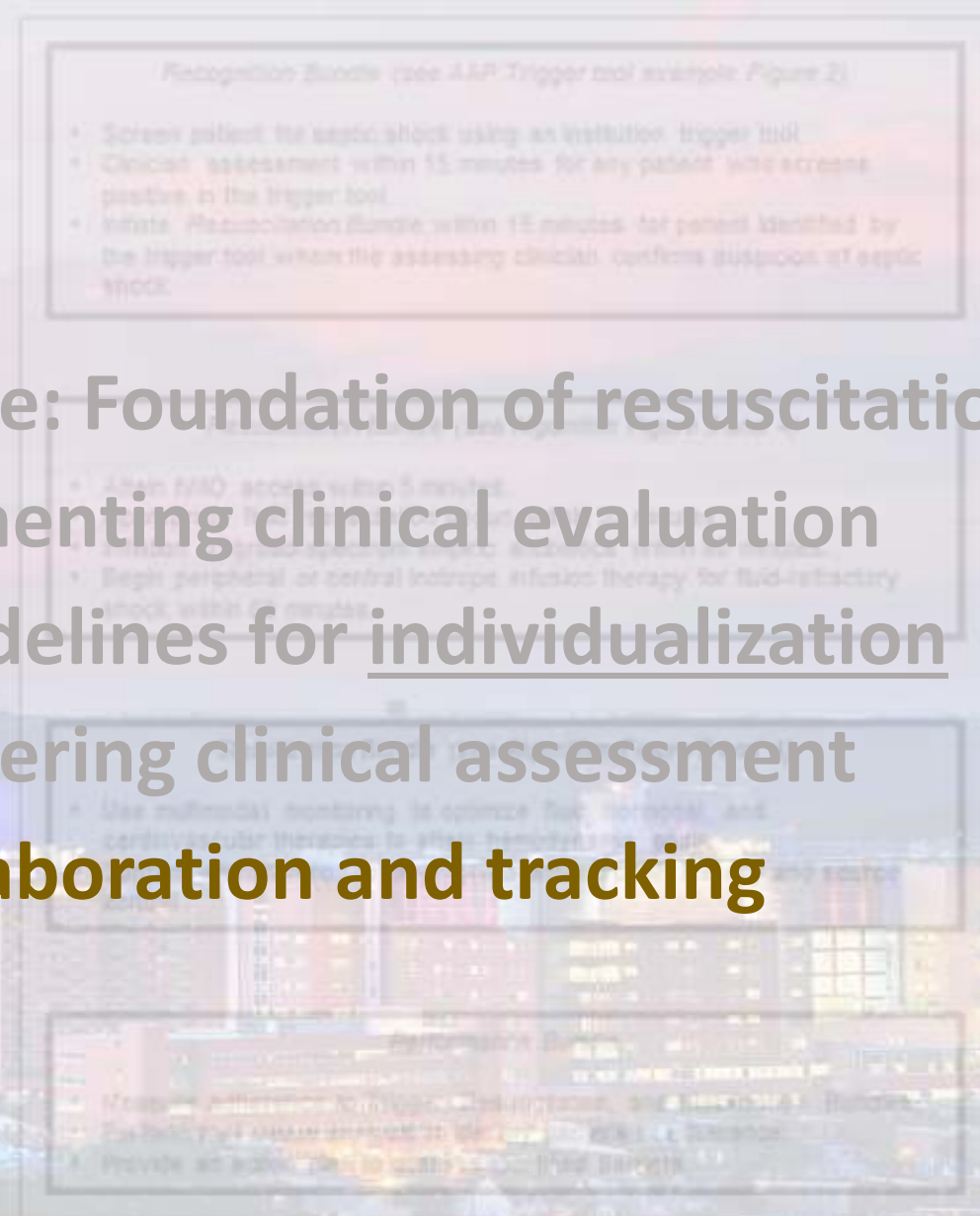


Adjuvant Approaches for Sepsis Subtypes

- Noninvasive hemodynamic assessment to help guide resuscitation
 - USCOM
 - Cardiotronic ICON
 - Bedside ultrasound
- Biomarkers of organ dysfunction and inflammatory response
 - Serum cortisol measured early
 - Adjuvant hydrocortisone infusion for persistent hypotension despite vasopressors
 - Markers of tissue perfusion and oxygen utilization (Lactate, SvO2)
 - Inflammatory cascade (CRP, Procalcitonin and Ferritin)
 - ADAMTS-13 assay
 - Plasma exchange for thrombocytopenia-associated MODs

Outline

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More Children Die From Sepsis Than Cancer

1 in 100 hospitalized children are diagnosed with severe sepsis/septic shock.

@ 1 in 10 will die

@ 4,500 deaths annually

@ 45% cases are hospital-onset.



Victoria

These are US stats. Global stats are worse.

SEPSIS  HEROES



CHILDREN'S
HOSPITAL
ASSOCIATION



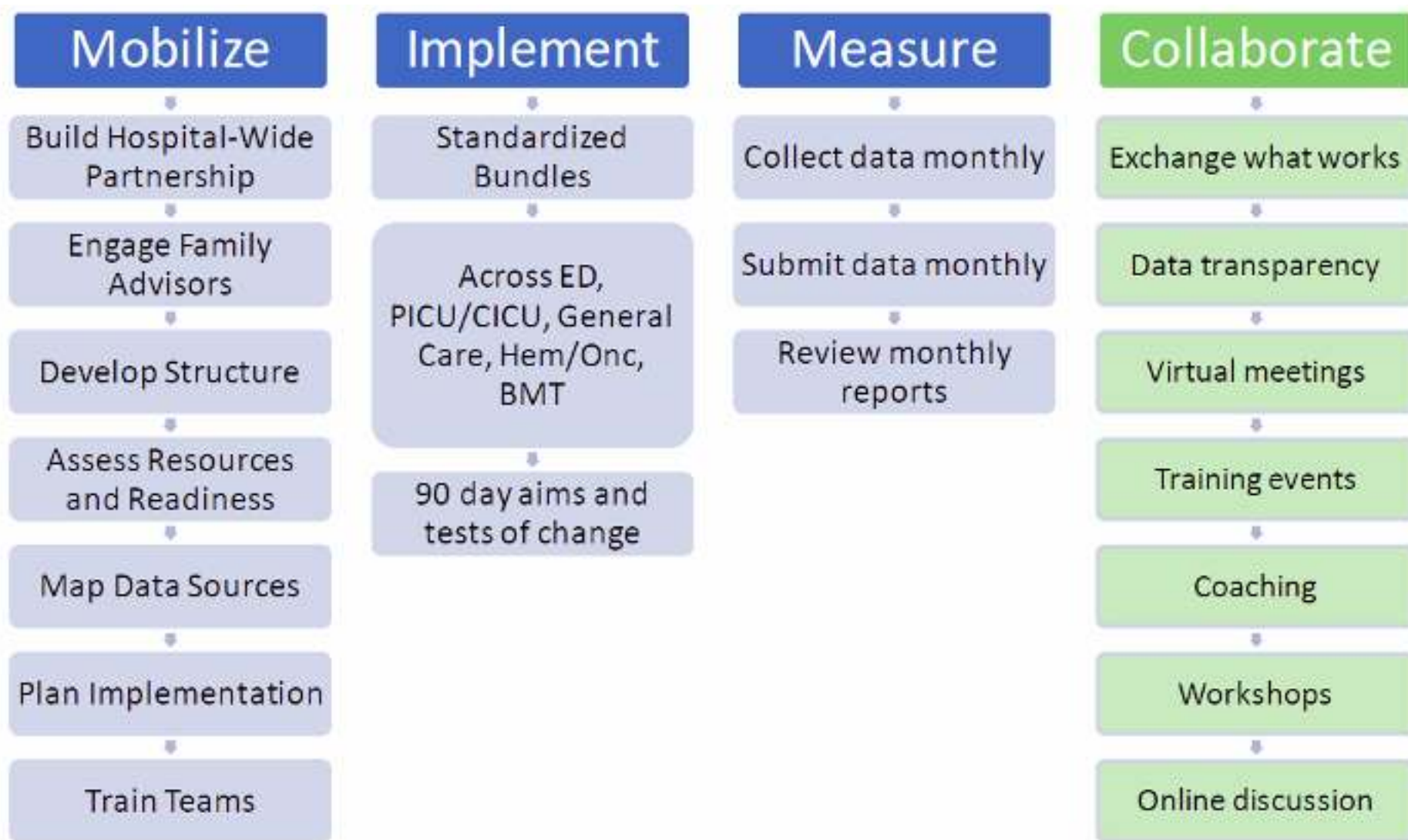
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SEPSIS XEROES



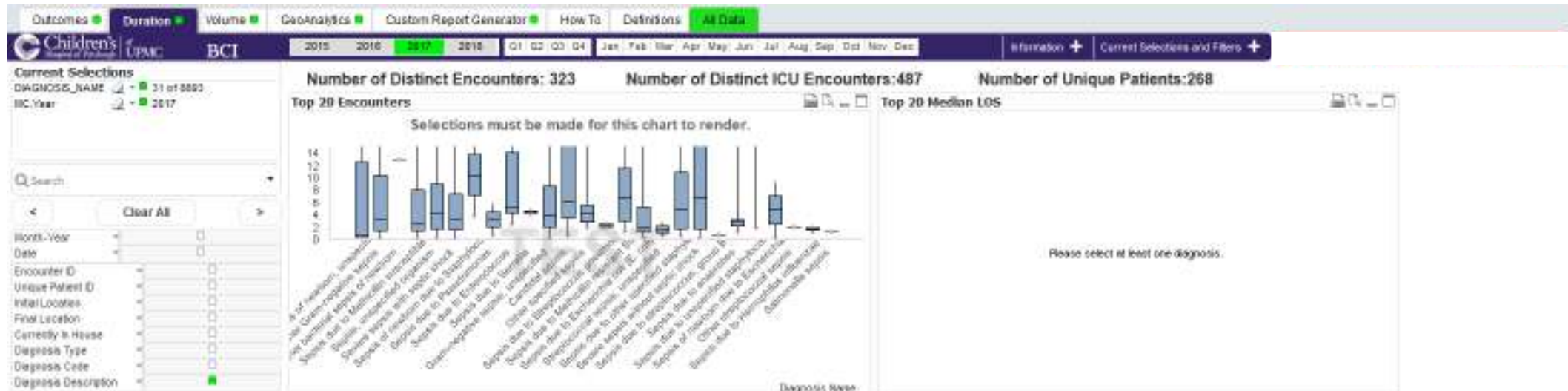
Sepsis and Septic Shock Prevalence and Mortality

	Sepsis or Septic Shock Prevalence (%)	Sepsis or Septic Shock Mortality (%)
Children's Hospital of Pittsburgh EHR definitions—all sepsis	23.3	2.2
Children's Hospital of Pittsburgh EHR definitions—septic shock	2.6	15.7
Watson et al. ¹	—	10.3
Hartman et al. ²	—	8.9
Ruth et al. ³ —both Angus criteria and ICD codes for sepsis	7.7	14.4
Ruth et al. ³ —Angus criteria	6.2	—
Ruth et al. ³ —ICD sepsis codes	3.1	—
Balamuth et al. ⁴ —Angus criteria	3.1	8.2
Balamuth et al. ⁴ —ICD sepsis codes	0.45	21.5
Weiss et al. ⁵ —SPROUT trial	8.2	25

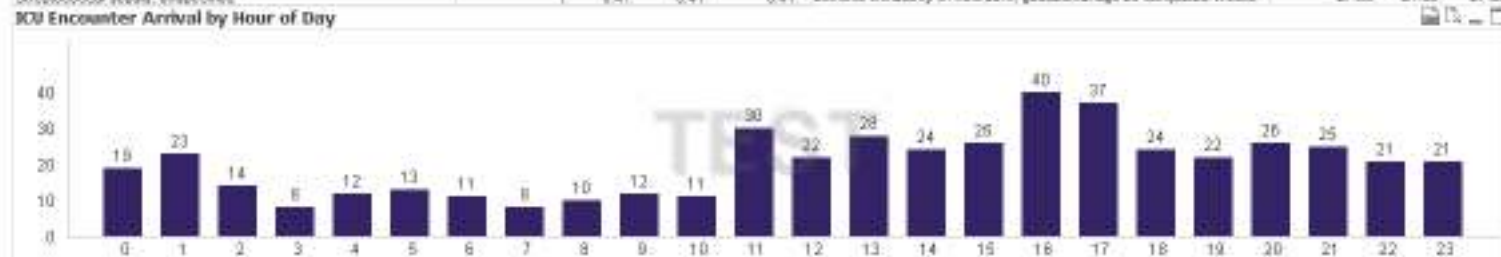
ICD, International Classification of Diseases.

Cohort Characteristics

	Sepsis and Septic Shock (N = 1,433)	Sepsis (n = 1,274)	Septic Shock (n = 159)
Age (mo), median [IQR]	42 [5–131]	41.7 [3.0–135.4]	64.5 [15.6–170.8]
Female, n (%)	649 (45.3)	577 (45.3)	72 (45.3)
Length of stay (d), median [IQR]	3.9 [2.0–9.1]	3.5 [1.9–7.4]	19.9 [7.2–43.7]
Length of stay (d), mean ± SD	10.6 ± 20.6	8.8 ± 18.6	35.7 ± 43.7
Hospital mortality, n (%)	32 (2.2)	7 (0.5)	25 (15.7)
30-d Mortality, n (%)	23 (1.6)	4 (0.3)	19 (11.9)
Extracorporeal support	11 (0.8)	0 (0)	11 (6.9)



Top 20 Encounters					Top 20 Median LOS				
Final Diagnosis	# Encount...	Min LOS	Max LOS	Median L...	Final Diagnosis	Median L...	Min LOS	Max LOS	
Sepsis, unspecified organism	52	8.40	84.16	6.89	Perforated intestinal perforation	128.73	128.73	128.73	
Sepsis due to Methicillin susceptible Staphylococcus aureus	7	8.02	33.88	3.88	Risk of larynx	98.01	98.01	98.01	
Sepsis due to Escherichia coli (E. coli)	5	8.71	7.94	1.57	Esophagus	78.18	78.18	78.18	
Sepsis due to Methicillin resistant Staphylococcus aureus	5	2.51	31.82	6.88	RESPIRATORY DISTRESS SYNDROME IN NEWBORN	72.29	72.29	72.29	
Other Gram-negative sepsis	3	3.87	19.57	17.95	Other pneumonia	61.18	61.18	61.18	
Other specified sepsis	3	1.97	4.48	3.81	ASPHYXIA AND STRANGULATION	54.95	54.95	54.95	
Gram-negative sepsis, unspecified	2	8.02	12.88	6.81	Respiratory distress syndrome of newborn	53.61	0.15	83.93	
Sepsis due to other specified staphylococcus	2	1.71	21.14	11.43	Traumatic subdural hemorrhage with loss of consciousness greater	48.92	40.92	48.92	
Sepsis due to Streptococcus pneumoniae	2	1.80	2.64	2.22	Malignant neoplasm of lower third of esophagus	48.34	40.34	48.34	
Sepsis of newborn due to Escherichia coli	2	8.10	9.23	4.88	DIAPHRAGMATIC HERNIA WITH OBSTRUCTION	39.88	39.88	39.88	
Bacterial sepsis of newborn, unspecified	1	1.45	1.45	1.45	Hepatorenal syndrome	33.00	33.00	33.00	
Other streptococcal sepsis	1	1.95	1.95	1.95	Single liveborn infant, delivered by cesarean	32.28	32.28	32.28	
Salmonella sepsis	1	1.20	1.28	1.28	Spastic diplegic cerebral palsy	31.99	31.99	31.99	
Sepsis due to anaerobes	1	15.74	15.74	15.74	Other inborn errors of metabolism of newborn	31.48	31.48	31.48	
Sepsis due to Haemophilus influenzae	1	2.01	2.81	2.81	OPEN FRACTURE OF VAULT OF SKULL WITH INTRACRANIAL INJURY	29.78	29.78	29.78	
Sepsis due to Pseudomonas	1	8.88	0.88	0.88	Chronic hepatic failure without coma	29.08	2.28	57.13	
Severe sepsis with septic shock	1	8.48	0.48	0.48	POSTINFLAMMATORY PULMONARY FIBROSIS	28.67	28.67	28.67	
Streptococcal sepsis, unspecified	1	8.41	0.41	0.41	Extreme immaturity of newborn, gestational age 26 completed weeks	27.83	27.83	27.83	



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Current Selections
 DIAGNOSIS_NAME 31 of 5093
 MCC Year 2017

Search

Clear All

Month-Year
 Date
 Encounter ID
 Unique Patient ID
 Initial Location
 Final Location
 Currently in House
 Diagnosis Type
 Diagnosis Code
 Diagnosis Description

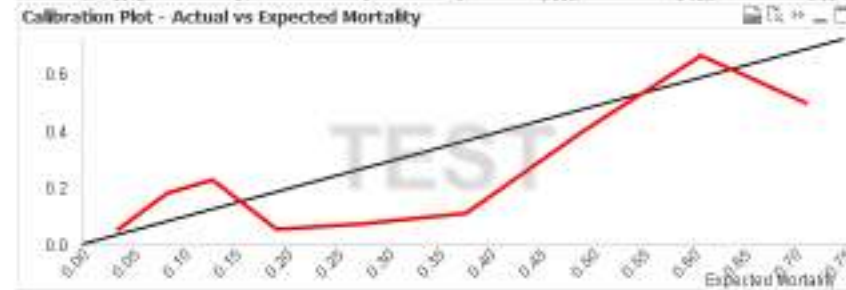
Number of Distinct Encounters: 323 Number of Distinct ICU Encounters: 487



Number of Unique Patients: 268



Month-Year	Admissions	Deaths Overall	ICU Deaths	Actual Mortal...	Expected Mortal...	Mortality Ratio
Jan-2017	33	25	17	6.72%	10.31%	0.65
Feb-2017	24	3	1	3.23%	7.00%	0.43
Mar-2017	21	3	2	9.52%	20.48%	0.46
Apr-2017	26	3	2	7.69%	10.00%	0.43
May-2017	28	4	4	13.79%	11.34%	1.22
Jun-2017	18	1	1	5.56%	14.93%	0.37
Jul-2017	26	4	2	7.69%	5.51%	1.40
Aug-2017	25	3	0	0.00%	6.63%	0.00
Sep-2017	32	3	2	6.25%	7.45%	0.84
Oct-2017	33	6	2	6.06%	10.29%	0.59

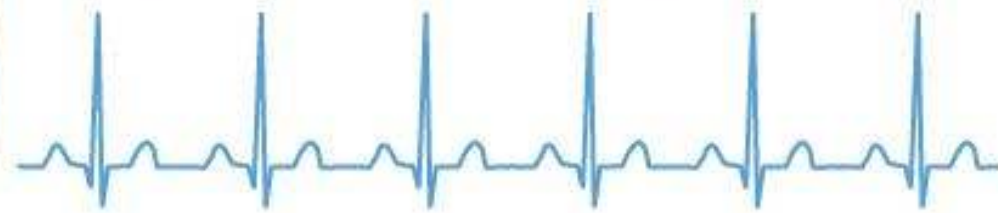


Summary

- 1) **Historic perspective:** Foundation of resuscitation
- 2) **Recognition:** Augmenting clinical evaluation
- 3) **Resuscitation:** Guidelines for individualization
- 4) **Stabilization:** Bolstering clinical assessment
- 5) **Performance:** Collaboration and tracking



Suspect **SEPSIS**



Save Lives

Image Credit: University of North Carolina at Chapel Hill School of Medicine



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Thank You!



Sajel
Kantawala



Gabriella
Butler



Chris Myers



Daniel
Rohm

Not Pictured:

Thomas Brown
Kelly Bricker
Janice Daugherty
Sue Park
Deneé Marasco
Kristi Russo



Bob Clark



Pat
Kochanek



Suresh
Srinivasan



Joe Carcillo

Thank You

Questions?

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