

of the Indiana Hospital Association

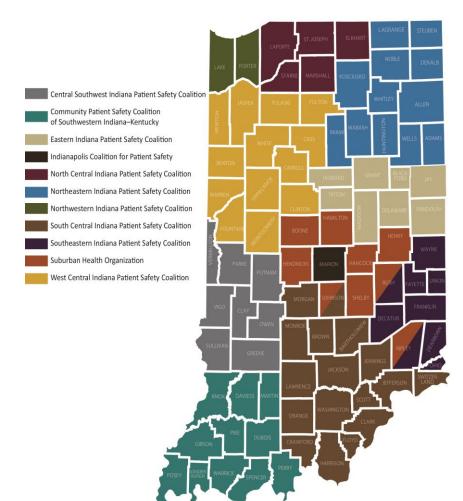
Sepsis Awareness Month: Back & to the Future: Maternal Sepsis

Brittany Waggoner, RN, MSN IHA Maternal and Infant Quality Improvement Advisor

Sept. 22, 2022

Our Mission





Advancing Health in Indiana

- Engage and inspire health care providers
- Create safe cultures
- Create reliable systems of care
- Prevent patient harm in Indiana

PREVENT PATIENT HARM

To create high reliability organizations who collaborate and engage in continuous improvement to achieve best in class outcomes

IMPROVE COMMUNITY HEALTH

To partner with communities and stakeholders to develop, plan, and coordinate initiatives that span the prevention and care continuum

INCREASE PATIENT AND FAMILY ENGAGEMENT

To engage patients and families in all aspects of their care and seek their input and inclusion in advisory capacities throughout organizations

LEAD A CULTURE OF SAFETY

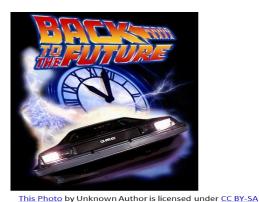
To create an environment of mutual trust, respect, and transparency among organizations, patients, and families

Sepsis: Back and to the Future



IHA 2022 Sepsis Awareness Month Webinars							
	1-Sept.	3 p.m. ET	Indiana Sepsis State of the State				
	8-Sept.	3 p.m. ET	Sepsis Pathophysiology & Bundle Compliance				
	15-Sept.	3 p.m. ET	Sepsis Diagnostic Advances				
	22-Sept.	3 p.m. ET	Maternal Sepsis				
	29-Sept.	3 p.m. ET	Sepsis Fluid Management Advances				
	6-Oct.	3 p.m. ET	Personal Hygiene and Sepsis Prevention				

Click on link to register for each webinar



Sepsis Webinar Details



2022 IHA Clinical Webinar Series - 3 - 4 p.m. ET

Sepsis: Back & to the Future (Click link to register)

Sept. 1: Indiana Sepsis 2022: Current State of the State and New Resources,

Rebecca Hancock PhD, RN, CNS, Patient Quality & Safety Advisor, IHA Chris Newkirk, BSN, RN, CCM, Clinical Quality Advisor, Columbus Regional Health

Sept. 8: Sepsis Back to Basics: Pathophysiology and Bundle Compliance,

Tom Ahrens, PhD, RN, FAAN

Sept. 15: Sepsis Future: Advances in Sepsis Diagnostics,

Dr. Sandy Estrada, Pharm.D., Clinical Consultant

Sept. 22: Sepsis Future: Focus on Maternal Sepsis,

Brittany Waggoner, Patient Safety & Quality Advisor, RN, MSN, CNS, IHA

Sept. 29: Sepsis Future: Fluid Management

Danielle Herr BSN, CCRN, Therapy Development Specialist

Vince Holly, MSN, RN, CCNS, ACNS-BC, CCRN, FCNS, Indiana University Health-Bloomington

Oct. 6: <u>Back to the Basics with Personal Hygiene for Infection Prevention</u>

Rebecca Hancock, Patient Quality & Safety Advisor, IHA Annette Handy, Clinical Director, Patient Safety Center, IHA

September is Sepsis Awareness Month—SET YOUR HOSPITAL GOALS!



- Updated 2022 Sepsis Toolkit coming August
 - ✓ Updated Social Media messages—connect with your marketing department & share IHA posts
 - ✓ Send photos of sepsis/COVID-19 infection prevention activities with caption to Casey Hutchens, <u>chutchens@ihaconnect.org</u>
 - ✓ Patient & Caregiver Education QR Codes on table tents, & posters
 - \checkmark Consider local mayoral proclamation for Sept 13, World Sepsis Day
 - √ Share "I am a Sepsis Champion" selfies on Sept 13 via social media
 - $\sqrt{}$ Updated data-based state sepsis goals

September Webinars, Thursdays 3-4pm *Back & to the Future with Sepsis*



www.survivesepsis.com



Sepsis Patient Discharge Education (Updated)





English



Spanish



Sepsis Patient and Family Education



Indiana Patient Safety Center of the Indiana Hospital Association

www.survivesepsis.com

SEE IT. STOP IT. SURVIVE IT.

I am a sepsis survivor, what now?

What can I do to improve my recovery?

Caregiver Instructions for Patients Recovering from Sepsis





ient-Safety

Adams Memorial Hospital





Ascension St. Vincent Indianapolis





Central Southwest Patient Safety Coalition





Columbus Regional Health





Harrison County Hospital





Indiana University Health



NICOLE

amber.

Partchara



St. Mary Medical Center





St. Mary's Medical Center







DECEMBER 2021 Thank you sepsis superheroes for acknowledgment of the sepsis BPA with notification to the provider for prompt attention.









1. Describe incidence of national and Indiana maternal sepsis and outcomes

2. Describe maternal sepsis identification & treatment recommendations

IHA Quality Reporting Table



The IHA table can be found at

Regulatory and Reporting (ihaconnect.org).



Current and Proposed CMS Quality Measures for Reporting in 2022 through 2028 *Revised 8/19/2022*

INPATIENT Current								
Measures Collected and Submitted by Hospital								
	HIQ	RP	VBP	HITECH				
MEASURE Bolded measures must be manually abstracted and submitted to HQR site quarterly.	Reporting effective date	Affects APU	Reporting effective date	Affects Reimburse ment	Promoting Interopera bility Program			
Structural Measure								
Maternal Morbidity	Oct 2021	FY 2023						
Hospital Commitment to Health Equity CY 2023 CY 2025								

Maternal Morbidity



Hospital Inpatient Quality Reporting (IQR) Program Maternal Morbidity Structural Measure Quick Reference Guide Fiscal Year (FY) 2023

Accessing and Completing the Maternal Morbidity Structural Measure in the Hospital Quality Reporting (HQR) Secure Portal	Helpful Tips
 In your Internet browser, navigate to <u>https://hqr.cms.gov</u>. The HQR home page will open. Enter your HARP user ID and Password. Click on Login. Select a device to verify your account. Click on Next. 	 CMS defines a statewide or national Perinatal Quality Improvement (QI) Collaborative as a statewide or multi-state network working to improve maternal and child health outcomes by addressing the quality and safety of perinatal care.
 Continue the two-factor authentication by entering your security code. Click on Continue. On the Terms & Conditions page, scroll down to the bottom of the Terms & Conditions. Click on Accept. 	 Examples include the Centers for Disease Control and Prevention's (CDC's) National Network of Perinatal Quality Collaboratives or Health Resources and Services Administration's (HRSA's) Alliance for Innovation on Maternal Health (AIM) program.
The HQR home page will open.Under the Dashboard, on the left-hand side of the screen, click on	 There are two parts to this measure's question. Both parts of the measure's question must be considered by hospitals when determining which final answer choice is appropriate.
 Data Submissions. Click on Structural Measures. Click on the Select a Response drop-down box and select your response: Yes No N/A (Our hospital does not provide inpatient labor/delivery care) Click on Save and Return. 	 For example, part one of the question assesses a hospital's participation in a statewide and/or national Perinatal QI Collaborative. Part two of the question assesses a hospital's implementation, through participation in the collaborative(s), of patient safety practices and/or bundles related to maternal morbidity. In order to select (A) Yes, a hospital must be able to answer "Yes" to both parts of the question if a hospital deems a "No" response is correct to either part of the question, then their attestation for the entire question must be (B) No.
 The structural measure data should display as "Submitted" on the Provider Participation Report. 	 If a facility does not provide labor/delivery care, the IPPS Measure Exception Form (used for the PC-01 measure) cannot be applied to the Maternal Morbidity Structural Measure. The facility will need to provide a response to the measure by selecting (C) N/A (Our hospital does not
Reporting and Submission Periods	provide inpatient labor/delivery care).
 For the CY 2021 reporting period/FY 2023 payment determination, the reporting period is a shortened period from October 1, 2021 through December 31, 2021. For CY 2021, if a facility participated in a perinatal quality improvement (QI) collaborative anytime during Q4 2021 (October 1 through December 31, 2021), it would satisfy the requirement. 	 Facilities should allow ample time before the deadline to review and, if necessary, correct their response. Facilities can update/correct their submitted response until the submission deadline. Immediately after that deadline, the HQR Secure Portal will be locked.
The submission period will be April 1 through May 16, 2022.	

2 parts for hospital with L & D:

- 1. Participation in a statewide and/or national Perinatal QI Collaborative
- 2. Implementation through participation in the collaborative(s), safety practices and/or bundles related to maternal morbidity

February 2022

Maternal MorbidityQRGFy2023 (qualityreportingcenter.com)

Guest Speaker





Brittany Waggoner, RN, MSN, XXX Maternal & Infant Quality Improvement Advisor Indiana Hospital Association





- Sepsis is an important cause of maternal morbidity and mortality.
- The Centers for Disease Control and Prevention notes that the proportion of U.S. maternal deaths from sepsis (12.7%) is similar to the proportion of deaths from obstetric hemorrhage (11.4%) and hypertensive disorders (7.4%).
- It is estimated that 63 to 73% of maternal deaths from sepsis are preventable.
- Furthermore, for each maternal death, there are 50 women who experience life threatening morbidity from sepsis.

National Center for Chronic Disease Prevention and Health Promotion





National Sepsis Statistics

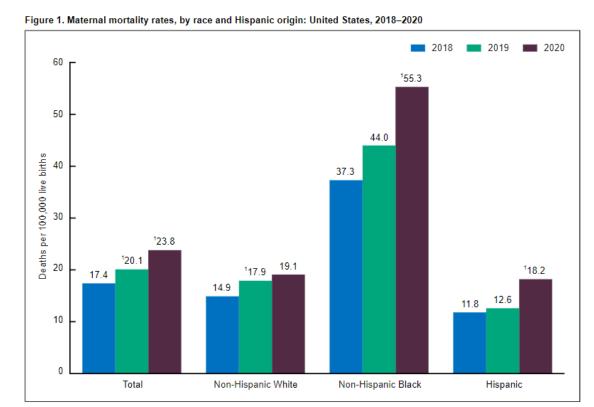
Maternal Mortality Rates (MMR) in the United States, 2020



	2018	2019	2020
Live Births	3,791,712	3,747,540	2,613,647
Maternal Deaths	658	754	861
Maternal Mortality Rate	17.4	20.1	23.8

Non-Hispanic Black MMR in the United States, 2020





¹Statistically significant increase in rate from previous year (p < 0.05).

NOTE: Race groups are single race.

SOURCE: National Center for Health Statistics, National Vital Statistics System, Mortality.

Non-Hispanic Black Maternal Mortality Rate 55.3

2.9 times the rate for non-Hispanic Whites (19.1)

The increases from 2019 to 2020 for non-Hispanic Black and Hispanic women were significant

The increases from 2019 to 2020 for non-Hispanic White women was not significant

Maternal Mortality due to Sepsis in the United States





Pregnancy-Related Deaths: Data from Maternal Mortality Review Committees in 36 US States, 2017-2019



Table 4. Underlying causes of pregnancy-related deaths*, overall and by race or ethnicity¹, data from Maternal Mortality Review Committees in 36 US states, 2017–2019¹

	Total							N	lon Hisp	anic				
			Hispanic		AIAN Asian		in	Black		NHOPI		White		
	N	%	n	%	n	%	n	%	n	%	n	%	n	%
Mental health conditions ²	224	22.7	34	24.1	2	-	1	3.1	21	7.0	0	-	159	34.8
Hemorrhage ³	135	13.7	30	21.3	2	-	10	31.3	33	10.9	1	-	53	11.6
Cardiac and coronary conditions ⁴	120	12.0	45	10.0			7	21.0	40	15.0	0		40	10.7
	126	12.8	15	10.6	1	-	7	21.9	48	15.9	0	-	49	10.7
Infection	91	9.2	15	10.6	1	-	0	0.0	23	7.6	0	-	49	10.7
Embolism- thrombotic	86	8.7	9	6.4	0	-	2	6.3	36	11.9	0	-	34	7.4
Cardiomyopathy	84	8.5	5	3.6	0	-	2	6.3	42	13.9	0	-	33	7.2
Hypertensive disorders of pregnancy	64	6.5	7	5.0	0	-	1	3.1	30	9.9	1	-	22	4.8
Amniotic fluid embolism	37	3.8	6	4.3	1	-	7	21.9	10	3.3	2	-	9	2.0
Injury ⁵	35	3.6	5	3.6	1	-	1	3.1	15	5.0	0	-	10	2.2
Cerebrovascular accident	25	2.5	2	1.4	0	-	0	0.0	10	3.3	0		13	2.8
Cancer	19	1.9	3	2.1	0	-	1	3.1	7	2.3	0	-	7	1.5
Metabolic/ endocrine conditions	12	1.2	2	1.4	0	-	0	0.0	6	2.0	0	-	3	0.7
					a states				4	1.3	1		5	1.1



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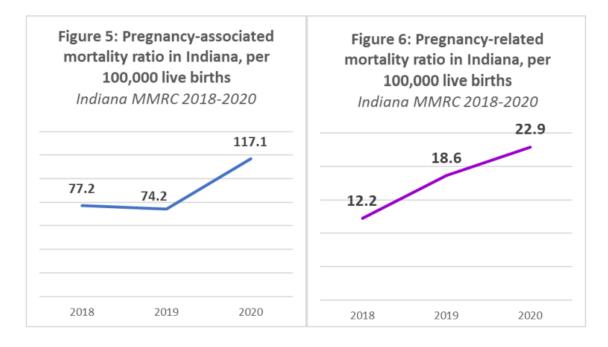


Indiana Maternal Mortality Review Committee





- 2022 Annual Report
 - 2020 Key Findings
 - 92 pregnancy-associated deaths occurred during pregnancy or within one year of the end of pregnancy.
 - 79% of reviewed pregnancy-associated deaths in 2020 were preventable.



Causes for all 2018-2020 Pregnancy Related Deaths



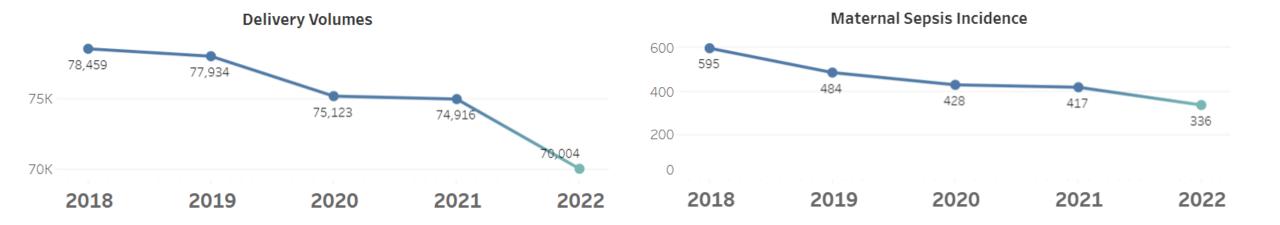
Indiana Maternal Mortality Review Committee 2022 Annual Report





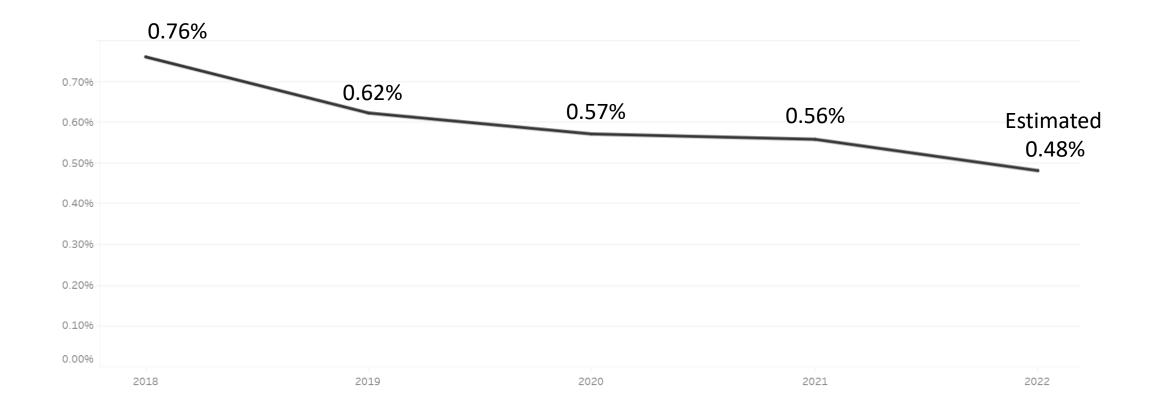






Indiana Maternal Sepsis Rate









Maternal Sepsis Readmission Rate

Maternal Sepsis Readmissions = 23 Delivery Volumes =74,916

Maternal Sepsis Readmission Rate = 3.07%

It's Not Just About Mortality....





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Pathophysiology of Sepsis



Antepartum	Intrapartum/ Immed. Postpartum	Post-discharge		
Septic abortion	Chorioamnionitis/ intraamniotic infection	Pneumonia/influenza		
Chorioamnionitis/ intraamniotic infection	Endometritis	Pyelonephritis		
Pneumonia/ influenza	Pneumonia/influenza	Wound Infection/ Necrotizing Fasciitis		
Pyelonephritis	Pyelonephritis	Mastitis		
Appendicitis	Wound Infection/ Necrotizing Fasciitis	Cholecystitis		

Pathophysiology

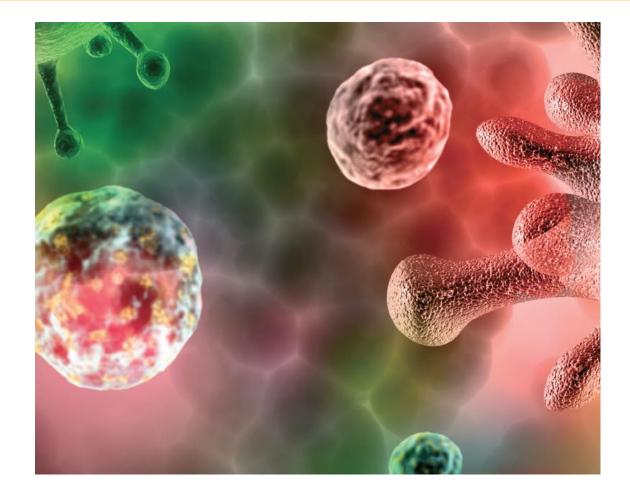


Pregnancy Sepsis Cardiovascular: Cardiovascular: ✤ Systemic vascular resistance (25–30%) ✤ Systemic vascular resistance Blood pressure ↑ Blood volume (40–45%) Blood pressure ↑ Heart rate ↑ Heart rate (10–20 bpm) Vasodilatation ↑ Cardiac output (40%) Myocardial depression Aorto-caval compression Respiratory: Respiratory: ↑ Pulmonary microvascular and plasma colloid pressure pressure and permeability ↓ Residual volume Acute lung injury ✤ Functional residual capacity ↑ Tidal volume Renal: ↑ Minute ventilation Ischaemia Compensated respiratory alkalosis Vasoconstriction Cytokine-mediated renal Renal: cell injury ↑ Renal plasma flow ↑ Glomerular filtration rate Renal collecting system dilatation Coagulation ↑ Procoagulant effects Coagulation ↑ Thrombin production ↑ Factors I, II, VII, VIII, IX, XII ↓ Activated Protein C ↑ (x5) plasminogen activator inhibitors (PAI) I & II Fibrinolysis (increased PAII) ←→ Anti-thrombin and Protein C

2019 Royal College of Obstetricians and Gynaecologists49Greer et al.

Pathogens





- Bacterial
- Fungal
- Viral
- No Causative Organism Identified

Non-OB



Tobacco use



Low socioeconomic status

Minorities

Presence of comorbidities



Transfusion

Pregnancy & Postpartum





Antepartum

- PROM
- Multiple Gestation
- Reproductive Technologies
- Preeclampsia
- Preterm Labor

Intrapartum

- PROM
- Multiple Vaginal Exams
- Prolonged 2nd Stage Labor

Postpartum

- Retained Products
- Hemorrhage
- Operative Vaginal or Cesarean Birth
- Multips



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Screening & Diagnoses

CMQCC's Two-Step Screening & Diagnosis Method

Comparing Normal Pregnancy Physiology and SIRS Criteria



Pregnancy Physiology	SIRS
↑ or ↓ Temperature	Temperature >38°C or <36°C (>100.4°F or <96.8°F)
HR ↑ 17%	HR >90 bpm
RR 🛧 in labor	RR >20 breaths/min
PaCO ₂ 28-32 mmHg	PaCO ₂ <32 mmHg
WBCs ↑ 8% to 5,000-12,000/mm ³ (up to 15,000/mm ³ seen) during pregnancy; during intrapartum may ↑ to 25,000-30,000/mm ³	WBC >12,000mm ³ or <4,000mm ³

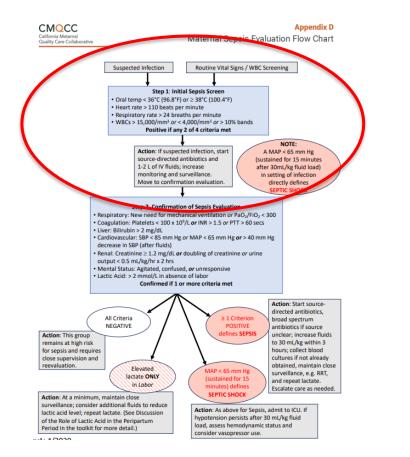
SEP-1 Criteria Measure Update

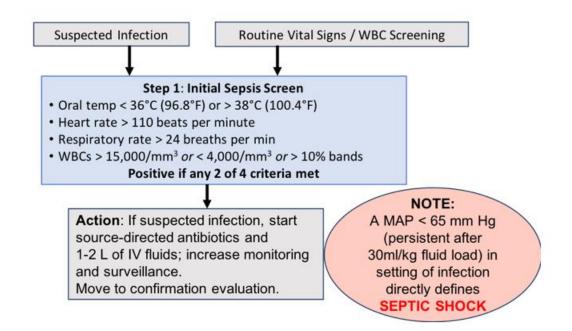


Non-Pregnant Criteria	Pregnant 20 weeks through Day 3 Post-delivery Criteria		
Temperature >38.3 C or <36.0 C (>100.9 F or <96.8 F)	Temperature ≥38 C or <36.0 C (≥100.4 or <96.8 F)		
<mark>Heart rate (pulse) >90</mark>	<mark>Heart rate (pulse) >110</mark>		
Respiration >20 per minute	Respiration >24 per minute		
White blood cell count >12,000 or 10% bands	White blood cell count >15,000 or 10% bands		

Screening & Diagnosing Step # 1







Confirmation of Sepsis Step #2



Tests to Evaluate End Organ Injury

Laboratory values

- CBC (including % immature neutrophils [bands], Platelets)
- Coagulation status (PT, INR, PTT)
- Comprehensive Metabolic Panel (specifically include bilirubin, creatinine)
- Venous Lactic Acid

Bedside assessment

- Urine output (place Foley catheter with urometer)
- Pulse oximetry
- Mental status assessment

Confirmation of Sepsis <u>Step #2</u>



Measure of End	Criteria			
Organ Injury	Positive if one (1) or more criteria are met			
Respiratory	Acute respiratory failure as evidenced by acute need for invasive or non-invasive			
function*	mechanical ventilation, OR			
	• PaO ₂ /FiO ₂ < 300			
Coagulation status	 Platelets < 100 x 10⁹/L, OR 			
	 International Normalized Ratio (INR) > 1.5, OR 			
	 Partial Thromboplastin Time (PTT) > 60 seconds 			
Liver function	Bilirubin > 2 mg/dL			
Cardiovascular	Persistent hypotension after fluid administration:			
function	 SBP < 85 mm Hg, OR 			
	○ MAP < 65 mm Hg, OR			
	○ > 40 mm Hg decrease in SBP			
Renal function	Creatinine > 1.2mg/dL, OR			
	 Doubling of serum creatinine, OR 			
	Urine output less 0.5 mL/kg/hour (for 2 hours)			
Mental status	Agitation, confusion, or unresponsiveness			
assessment				
Lactic acid	 > 2 mmol/L in absence of labor 			
	(Lactic acid not used for diagnosis in labor, but remains important for treatment.)			

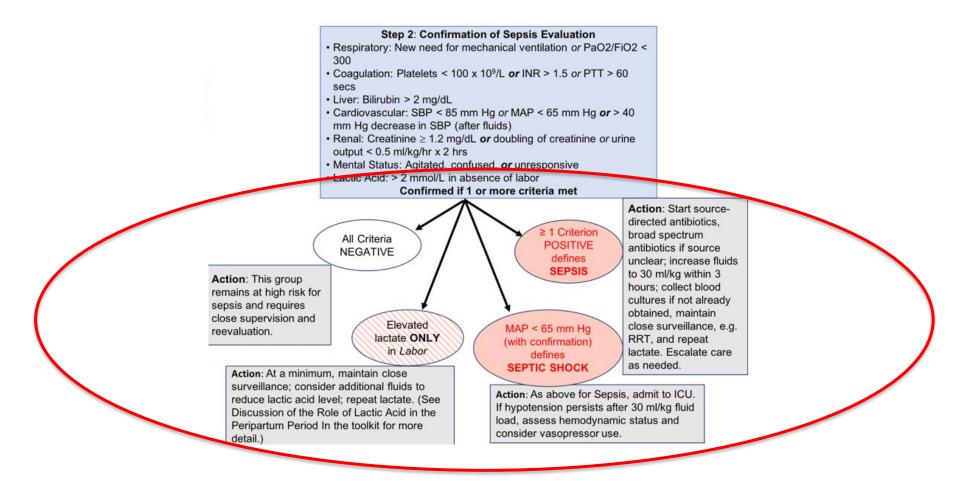


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Assessment & Treatment

Assessment & Treatment





Sepsis Treatment



- Narrow spectrum antibiotics if not already started
- If source unclear, give broad spectrum antibiotics
- Increase fluids to 30 mL/kg within 3 hours if not already done
- Repeat lactate
- Blood cultures, if not already drawn
- Call for RRT to escalate care, as needed

Fluid Management





30ml/kg Crystalloid IV Fluid Bolus
Begin within 3 hours
• Each hour increases mortality rate by 7.6%
Large-bore IV
Considerations
 Pulmonary edema ARDS Preeclampsia

Antibiotic Therapy



Broad spectrum

- Most cases are polymicrobial
- Ampicillin, Gentamycin, Clindamycin, Vancomycin, Metronidazole, Antivirals
- Reassess therapy daily
- Toxicity versus therapeutic

Antibiotic Regiment (CMQCC)



Antibiotic Regimen by Condition

(See full recommendations in Toolkit)

Condition	Antibiotic Choices	Duration	Notes
Chorioamnionitis	Ampicillin 2 g IV q6h	Generally limited	For post-cesarean
/ intraamniotic		to the peripartum	delivery: one additional
infection (Plante,	PLUS	period	dose of the chosen
et al 2019, ACOG)	Gentalizin 2 mg/kg IV load, then 5mg/kg		regimen is indicated. Add clindamycin 90 mg IV or
ACOG)	are (24b)	Duration of	Metronidazole 500 mg IV
		therapy is unclear,	for at least one additional
	Alternate regimens: (Based on local	but there are	dose.
	antibiotic resistance patterns)	some	
		recommendations	For post-vaginal delivery:
	Ampicillin-sulbactam 3g IV q6h OR	to continue until	No additional antibiotic
	Dinemacillin terreheaters 2.275 r IV rCh OD	afebrile for 24h	doses required, but if
	Piperacillin-tazobactam 3.375g IV q6h OR		additional doses of antibiotics are given,
	Cefoxitin 2g IV q6H OR		clindamycin is not
			indicated.

Antibiotic Regiment Cont.

-



Source infection	Recommended antibiotics		
Abdominal infections	Ceftriaxone, cefotaxime, ceftazidime, or cefepime plus metronidazole;		
	Complicated cases may require monotherapy with a carbapenem or piperacillin-tazobactam		
Chorioamnionitis	Ampicillin plus gentamicin. Add anaerobic coverage with clindamycin or metronidazole if cesarear delivery required		
Community- acquired pneumonia	Cefotaxime, ceftriaxone, ertapenem, or ampicillin plus azithromycin, clarithromycin, or erythromycin		
Endomyometritis	Ampicillin, gentamicin, and metronidazole (or clindamycin); Alternatively, may use cefotaxime or ceftriaxone plus metronidazole		
Hospital-acquired pneumonia	Low risk patients: Piperacillin-tazobactam, meropenem, imipenem, or cefepime		
	High mortality risk patients: double coverage for pseudomonas (beta lactam plus an aminoglycoside or a quinolone) and MRSA coverage with vancomycin or linezolid		
Skin and soft tissues (necrotizing)			
Urinary tract infections	Gentamicin with ampicillin; Alternatively, may use monotherapy with a carbapenem or piperacillin-tazobactam		

Septic Shock



- Admit to ICU
- Hemodynamic Monitoring
- Vasopressors



Lactic Acid



Not used for diagnosis in labor, but remains important for treatment Trends can be used to evaluate the effectiveness of treatment

IHAconnect.org/Quality-Patient-Safety

Anxiety Fatigue

- Sleep disturbances
- Post-traumatic stress syndrome



Individualized

Potential for

– Depression



Faces of Sepsis





Cortina Pride



Sarah S.



Samantha Bullock

Erza H.



Jennifer Inskip



Shay and Amelia B.



Kourtney N.



Steph Wasson

ty-Patient-Safety





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Thank you!





Brittany Waggoner, RN, MSN Maternal & Infant Quality Improvement Advisor Indiana Hospital Association



Nov 1-2 Nov 1-2 Nov 1-2 Nov 1-2 Nov Years Advancing Health Care Together

Nov. 1 – 2 The Westin Indianapolis

Learn more and register on our website:



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