

Sepsis QAPI 2022

| POOR | Meeting Benchmark | Exceeding Benchmark |
|------|-------------------|---------------------|
|------|-------------------|---------------------|

| Jan | Feb | Mar | Apr | May | Jun | Jul | Aug | Sep | Oct | Nov | Dec |
|---------------|-----------------------|-----------------------|-----------------------|-----------------------|---------------|-----|-----|-----|-----|-----|-----|
| 56.25% | 100% | 91.67% | 72.23% | 81.82% | 88.89% | | | | | | |
| 9/16 | 7/7 | 11/12 | 8/11 | 9/11 | 8/9 | | | | | | |
| 6 excluded | 11 excluded | 13 excluded | 10 excluded | 10 excluded | 6 excluded | | | | | | |

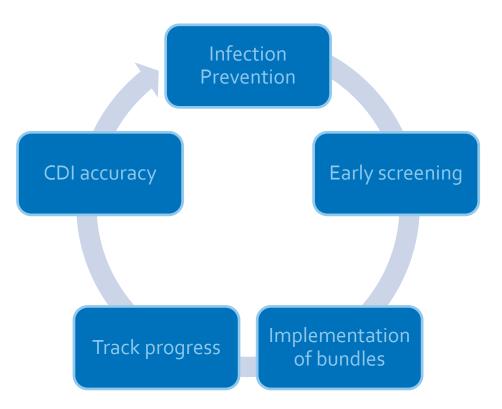
KRMC Sepsis Bundle Compliance



CY 2019=48.89% CY 2020=60.42% CY 2021=65.49% CY 2022=78.79%

Where we are at:

- Continue:
 - PI program with multi-disciplinary collaboration
 - Monthly meetings
 - Gap analysis with NHA (done)
 - Sepsis surveillance and screening
 - Data dashboard
 - Manual abstraction and tracking
 - Timely feedback for outliers
 - ER provider OPPE (concurrent)
 - EMR optimization
 - Education
- Goals: (70% overall compliance goal)
 - Communication to help drive bundle compliance
 - T2 Biosystems analysis (hold)
 - Track ER specific data and readmissions
 - Door/Order to antibiotic time



Outliers by Quarter:

| | Initial Lactate | Blood Cultures | Late Antibiotic | Repeat Lactate | Fluid bolus 3oml/kg | Persistent hypotension | Vasopressor Initiated | Focused Exam (within 6 hrs of fluid bolus) |
|----|--------------------|-------------------|--------------------|-------------------|------------------------|---------------------------|--------------------------|---|
| Qı | 2 | 3 | 3 | 2 | 4 | 1 | | 2 |
| Q2 | 1 | 1 | 1 | 1 | 1 | | 1 | 2 |

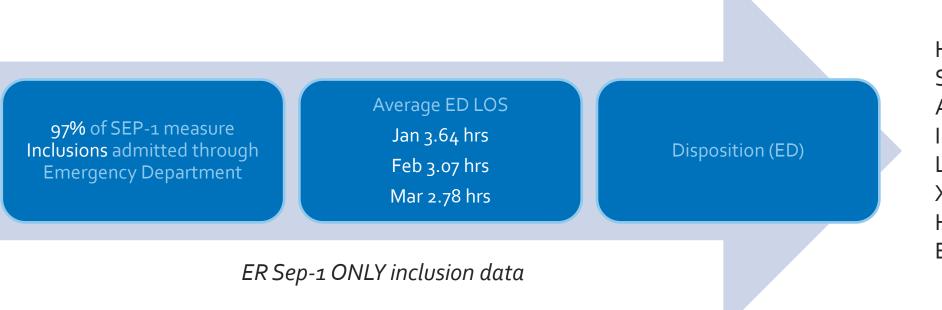
Outlier opportunities:

- AKI baseline if available within 6 hrs OR any reason for AKI besides infection
- Sepsis workup on known infectious patients with non-infectious visits
- Provider documentation of 1) target fluid volume 2) reason WHY giving <30ml/kg
- Tissue perfusion exam: Attest or review 5/8 parameters



This does not match # of fallouts, as looking at all bundle requirements for each case.

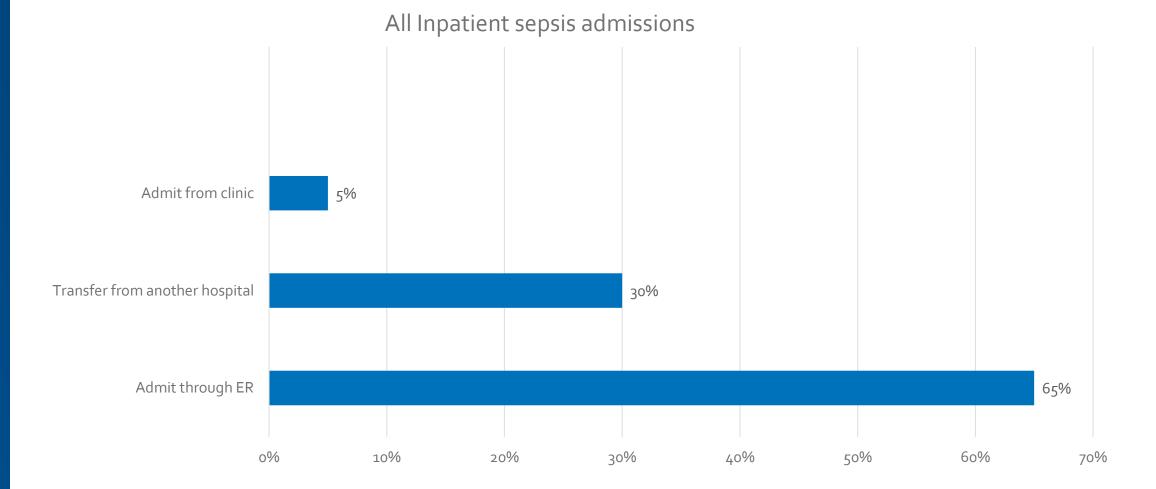
ER Specific data: Q1



Home x16 SNF x8 ALF x1 Inpt rehab x2 LTAC x1 Xfer acute care x2 Hospice x1 Expire x4

- ED outliers:
 - -----(late antibiotics)
 - -----(nursing error fluids)
 - -----(no blood culture, lactic, abx documentation)
 - -----(fluid ordered, then discontinued by pharmacy-no explanation)

Hospital-wide data: Q1



Average LOS: (includes all inpt sepsis diagnoses)

| DRG | LOS | Qı | Q2 | Q3 | Q4 |
|---|-------------------|----|----|----|----|
| DRG 907 Other OR Procedures for injuries with MCC | Ave (9.6 days) | 1 | | | |
| DRG 896 Alcohol, drug abuse or dependence without rehab | Ave (7.0 days) | 1 | | | |
| DRG 870 Septicemia or severe sepsis with MV >96 hrs | Ave (14.5 days) | 1 | | | |
| DRG 871 Septicemia or severe sepsis without MV >96 hrs with MCC | Ave (6.2 days) | 39 | | | |
| DRG 872 Septicemia or severe sepsis without MV >96 hrs without MCC | Ave (4.2 days) | 4 | | | |
| DRG 853 Infectious and parasitic disease with OR procedure with MCC | Ave (12.5 days) | 5 | | | |
| DRG 854 Infectious and parasitic disease with OR procedure with CC | Ave (6.6 days) | 1 | | | |
| DRG 720 Septicemia and disseminated infections | Based on severity | 2 | | | |
| DRG 710 Infectious and parasitic disease including HIV W OR Procedure | Based on severity | 2 | | | |
| DRG 441 Disorders of liver except malignancy, cirrhosis, or alcoholic hepatitis | Ave (6.4 days) | 1 | | | |
| DRG 521 Hip Replacement with Principal Diagnosis of Hip Fx with MCC | Ave (7.2 days) | 1 | | | |
| DRG 314 Other Circulatory System Diagnosis with MCC | Ave (6.6 days) | 3 | | | |
| DRG 070 Non-specific Cerebrovascular Disorders with MCC | Ave (6.2 days) | 1 | | | |

| Jan | Feb | Mar | | | | |
|-----------|-----------|-----------|--|--|--|--|
| 7.04 days | 7.72 days | 7.05 days | | | | |

MCC=Major complication or comorbidity

Readmissions: January

| Initial Account Number | Patient Name | Financial Class Name | Initial Attending Provider | Initial DRG ID | Initial DRG | LOS | Initial Discharge Disposition | Initial Admit Date | Initial Discharge Date | Readm Account Number | Readm DRG ID | Readmission Summary | Readm LOS | | Readm DC Disposition |
|------------------------------|-----------------|-------------------------|----------------------------------|----------------------|-------------|-----|-------------------------------------|-----------------------|------------------------------|----------------------------|-----------------|---------------------|--------------|--|-------------------------|
|------------------------------|-----------------|-------------------------|----------------------------------|----------------------|-------------|-----|-------------------------------------|-----------------------|------------------------------|----------------------------|-----------------|---------------------|--------------|--|-------------------------|

patient information removed



V5.11 Fluid Updates: (OLD to NEW)

- That administration of 30ml/kg of crystalloid fluid would be detrimental or harmful for the patient despite having hypotension, lactate >4, or septic shock documentation
- AND the patient must have one of the following conditions within that note:
 - <u>Advanced or end-stage heart failure (must include NYHA class III</u>, IV or symptoms with minimal exertion)
 - <u>Advanced or end-stage chronic renal disease</u> (must include stage IV, V, or ESRD)
- AND the VOLUME in place of the 30ml/kg that the patient should receive with ORDER
- ✓ OR that a portion of the crystalloid volume was administered as colloids

The amount of volume to bolus

This can be a specific volume (ex: 1500ml) OR weight based (ex: 20ml/kg)
Still must be a balanced crystalloid fluid, but once target reached could change to bicarb, etc.

- The reason for the lesser volume (not limited to):
 - Heart Failure
 - Renal Failure
 - Fluid Overload
 - Blood Pressure responded to lesser volume
 - Portion given as colloids
- Evaluate tissue perfusion/volume reassessment (within 6hrs of crystalloid start time)
 - I have evaluated the patient post fluid bolus
 - Patient evaluated after 1L NS, plan to give 500ml additional
 - Sepsis re-examination complete

IBW always an option if the patient BMI >30. MUST document "IBW fluid orders due to obesity" or use specific IBW order

Canned Text

oSepsisFluid-HeartFailure 1SepsisFluid-Hypotension 2SepsisFluid-Overload 3SepsisFluid-RenalFailure 4SepsisFluid-Colloids 5SepsisFluid-Reassessment 6SepsisFluid-IBW

| Search Canned Text | Support Text | | |
|---------------------------|---------------------------|-----------------------|---------------------------|
| Favorites | | | |
| JSepsisFluid-HeartFailure | 1SepsisFluid-Hypotension | 2SepsisFluid-Overload | 3SepsisFluid-RenalFailure |
| 4SepsisFluid-Colloids | 5SepsisFluid-Reassessment | 6SepsisFluid-IBW | |
| | | | |

Opportunity: Providers please add date/time with sepsis reassessment

| ✓ Course |
|--|
| Hospital Course A B i U D C Crystalloid fluid bolus of ml administered for sepsis target fluid volume due to heart failure. Crystalloid fluid bolus of ml administered for sepsis target volume due to fluid overload. Crystalloid fluid bolus of ml administered for sepsis target volume due to fluid overload. Crystalloid fluid bolus of ml administered for sepsis target volume due to fluid overload. Crystalloid fluid bolus of ml administered for sepsis target fluid volume due to renal failure. Crystalloid fluid bolus of ml administered for sepsis target fluid volume due to renal failure. Crystalloid fluid bolus of ml, along withml of albumin for sepsis target fluid volume. Sepsis re-examination complete. Using Ideal Body Weight (IBW) to determine target fluid resuscitation volume d/t patient obesity; pharmacy to calculate IBW and add volume to MAR. |
| Reevaluation(s) |

Fluid research

- Sepsis Alliance
 - SMART and SALT-ED Trials with 29k patients comparing saline to balanced crystalloid
 - Saline led to a higher incidence of AKI (acute kidney injury)
 - Issues:
 - » High sodium and chloride content
 - » Hyperchloremia may lead to renal vasoconstriction, reduced renal blood flow, & kidney injury
 - » Acidosis
 - Recommendations=
 - Balanced crystalloids instead of NS (LR and Normosol)
 - Albumin in patients who receive large volumes of crystalloids over just using crystalloids alone

Surviving Sepsis Campaign: International Guidelines for the Management of Sepsis and Septic Shock 2021. Evans, Rhodes, Alhazzani et al. *Crit Care Med 2021*;49(11):e1063-e1143 Neyra JA et al *Crit Care Med*. 2017:43(9): 1938-1944. Yunos et al. (2012)

Self et al NEJM (2018); 378:9. Semler et al NEJM (2018); 378:9

Repeat volume status and tissue perfusion assessment

- Specified time frame (start of fluid bolus to 6-hrs post septic shock presentation)
- May consist of any one of the following three:
 - 1. <u>Provider documentation attesting to performing exam</u>:
 - Example: Sepsis focused exam performed (yes) or "review of systems complete"
 - Does the heading "exam" or "review of systems" count?
 - 2. <u>Provider documentation indicating they completed a review of at least 5 of the following 8</u> <u>parameters:</u>
 - O2 saturation, Capillary refill, Cardiopulmonary assessment, Peripheral pulses, Shock index, Skin color or condition, UOP, VS
 - 3. <u>Documentation that one of the following was completed:</u>
 - CVP
 - Sv02
 - ECHO

CMS v5.12 Update

- New documentation allowed for <u>blood culture</u> (Can select YES if there is an attempt to collect the specimen, but the attempt resulted in failure to collect (dehydrated etc); there must be a time directly associated with documentation indicating that the blood culture was collected during the specified time.
 - ✓ Blood culture attempted
 - ✓ Blood culture x 3 attempts
 - ✓ Unable to collect blood cultures
- <u>Colloids</u> are now included for crystalloid fluid requirement if given at greater than 125ml/hr
 - ✓ Must have start/stop in designated time period of at least 126ml/hr
 - Example: Target goal 1100ml with 1L NS + 100ml albumin
- New criteria for pregnant 20 weeks through day 3 post-delivery
 - \checkmark Account for physiologic changes during pregnancy
 - 1. SBP <85 (non-pregnant <90)
 - 2. If lactate >2, was obtained during active delivery DO NOT USE; active delivery defined as documentation of uterine contractions resulting in cervical change (dilation or effacement)
 - 3. SIRS differences: Temperature <a>100.4 or <96.8F (non-pregnant >100.9); Heart rate >110 (non-pregnant >90); RR >24 (non-pregnant >20); WBC >15,000 (non-pregnant >12,000)
 - 4. Organ dysfunction differences: Cr >1.2 (non-pregnant 2.0)

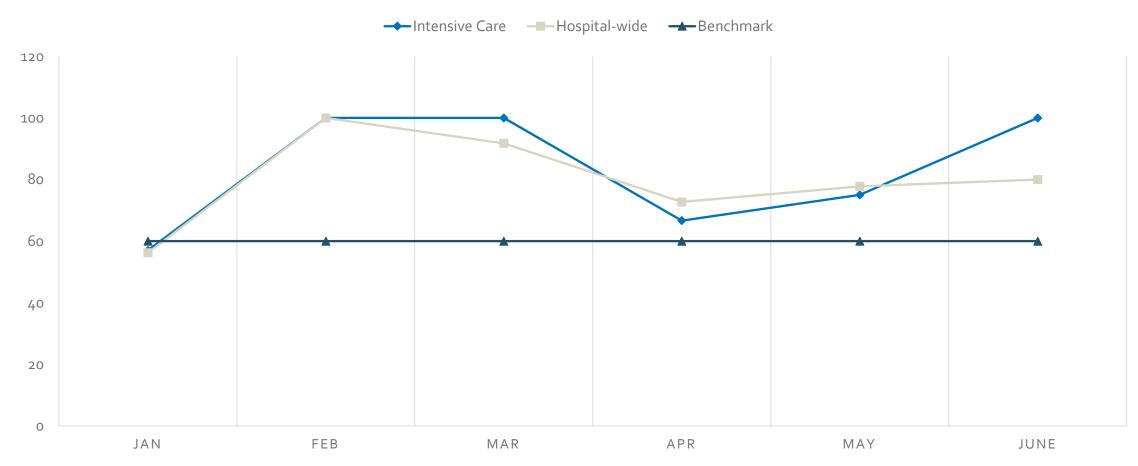
Maternal sepsis has increased over the last decade due to older maternal age, multi-fetal gestation, obesity, diabetes, and assisted reproductive technologies



ICU Sepsis Bundle Performance

| Sep-1 Hospital-wide | 76.67% |
|--------------------------|--------|
| Sep-1 Intensive Care | 79.17% |
| Sep-1 National Benchmark | 60.00% |

ICU SEP-1 BUNDLE COMPLIANCE 2022



KRMC Sep-1 Outcomes (Nebraska Hospital Association) submission

- Sep-1 measure compliance: Up 30 percentage points from 2019 (49% to 79%)
- Mortality (2019) 16.7%, (2020) 16.5%, (2021) 12.9%, (2022) 13.9%
 - Note: (2021) 15.5% of Sep-1 patients met septic shock criteria, CY 2022 32% YTD.
- Length of stay for Sep-1 MEETS (2021) 4.61 days vs 7.27 days for those who did not pass the measure (2022 thru Q2)

5.79 days vs 7.29 days.

- Patients discharging to home from severe sepsis/septic shock has increased 10% from 2019, at a current rate of 55%.
- Sep-1 Readmission rates have increased from 2019/2020-this is an active agenda item. Our current rate is 13.6%.

