



*Minnesota Hospital Association*

# MHA/OHA HIIN Antibiotic Stewardship/MDRO Collaborative

July 11, 2017



# Reminders



- For best sound quality, dial in at **1-800-791-2345** and enter code **11076**
- Please use the chat box to ask questions!

*Please note – this webinar is being recorded.*

# Housekeeping

- Education Credit
  - Nursing Education Credit – 1 hour
  - Pharmacy Education Credit – 0.1
    - Pharmacists, please list your license number on the sign-in sheet to receive credit

# Agenda

- Welcome
- Presentation:
  - Antimicrobial Stewardship: The Practical Aspects of Leveraging Technology by Kimberly Boeser, PharmD, MPH, BCPS AQ-ID
- Questions/discussion
- ASP 101 reminders
- Wrap-up

# ANTIMICROBIAL STEWARDSHIP: THE PRACTICAL ASPECTS OF LEVERAGING TECHNOLOGY

***KIMBERLY BOESER, PHARM.D., MPH, BCPS AQ-ID***

INFECTIOUS DISEASES CLINICAL PHARMACIST

ANTIMICROBIAL STEWARDSHIP COORDINATOR

DIRECTOR, PGY2 INFECTIOUS DISEASE RESIDENCY PROGRAM



# THE CALL FOR STEWARDSHIP

**Stewardship:** the conducting, supervising, or managing of something; *especially*: the careful and responsible management of something entrusted to one's care

**Antimicrobial Stewardship:** coordinated interventions designed to improve and measure the appropriate selection, dosing, route and duration of antimicrobial therapy

- Primary Goal: optimize clinical outcomes, while minimizing unintended consequences of antimicrobial use
  - Toxicity
  - Selection of pathogenic organisms (MRSA, VRE, ESBL gram negative bacteria)
  - Emergence of RESISTANCE
- Secondary Goal: reduce health care costs w/out adversely impacting quality of care

# ANTIMICROBIAL STEWARDSHIP

**Growing body of evidence demonstrates that ASPs dedicated to improving antibiotic use,**

- Improve the quality of patient care and patient safety
  - Increase infection cure rates
  - Reduce treatment failures
  - Reduce adverse events associated with antimicrobial therapy
- Decrease antibiotic resistance
  - significantly reduce hospital rates of (*Clostridium difficile* infections) (CDI)
    - Fairview quality initiative 2017
- Provide hospitals with opportunity for cost savings

**2014 CDC recommended that all acute care hospitals implement Antibiotic Stewardship Programs**

**ASPs can be implemented effectively in a wide variety of hospitals**

- SUCCESS is dependent on defined leadership and a coordinated multidisciplinary approach

# Implementing an Antibiotic Stewardship Program: Guidelines by the Infectious Diseases Society of America and the Society for Healthcare Epidemiology of America

Tamar F. Barlam,<sup>1,a</sup> Sara E. Cosgrove,<sup>2,a</sup> Lilian M. Abbo,<sup>3</sup> Conan MacDougall,<sup>4</sup> Audrey N. Schuetz,<sup>5</sup> Edward J. Septimus,<sup>6</sup> Arjun Srinivasan,<sup>7</sup> Timothy H. Dellit,<sup>8</sup> Yngve T. Falck-Ytter,<sup>9</sup> Neil O. Fishman,<sup>10</sup> Cindy W. Hamilton,<sup>11</sup> Timothy C. Jenkins,<sup>12</sup> Pamela A. Lipsett,<sup>13</sup> Preeti N. Malani,<sup>14</sup> Larissa S. May,<sup>15</sup> Gregory J. Moran,<sup>16</sup> Melinda M. Neuhauser,<sup>17</sup> Jason G. Newland,<sup>18</sup> Christopher A. Oehl,<sup>19</sup> Matthew H. Samore,<sup>20</sup> Susan K. Seo,<sup>21</sup> and Kavita K. Trivedi<sup>22</sup>

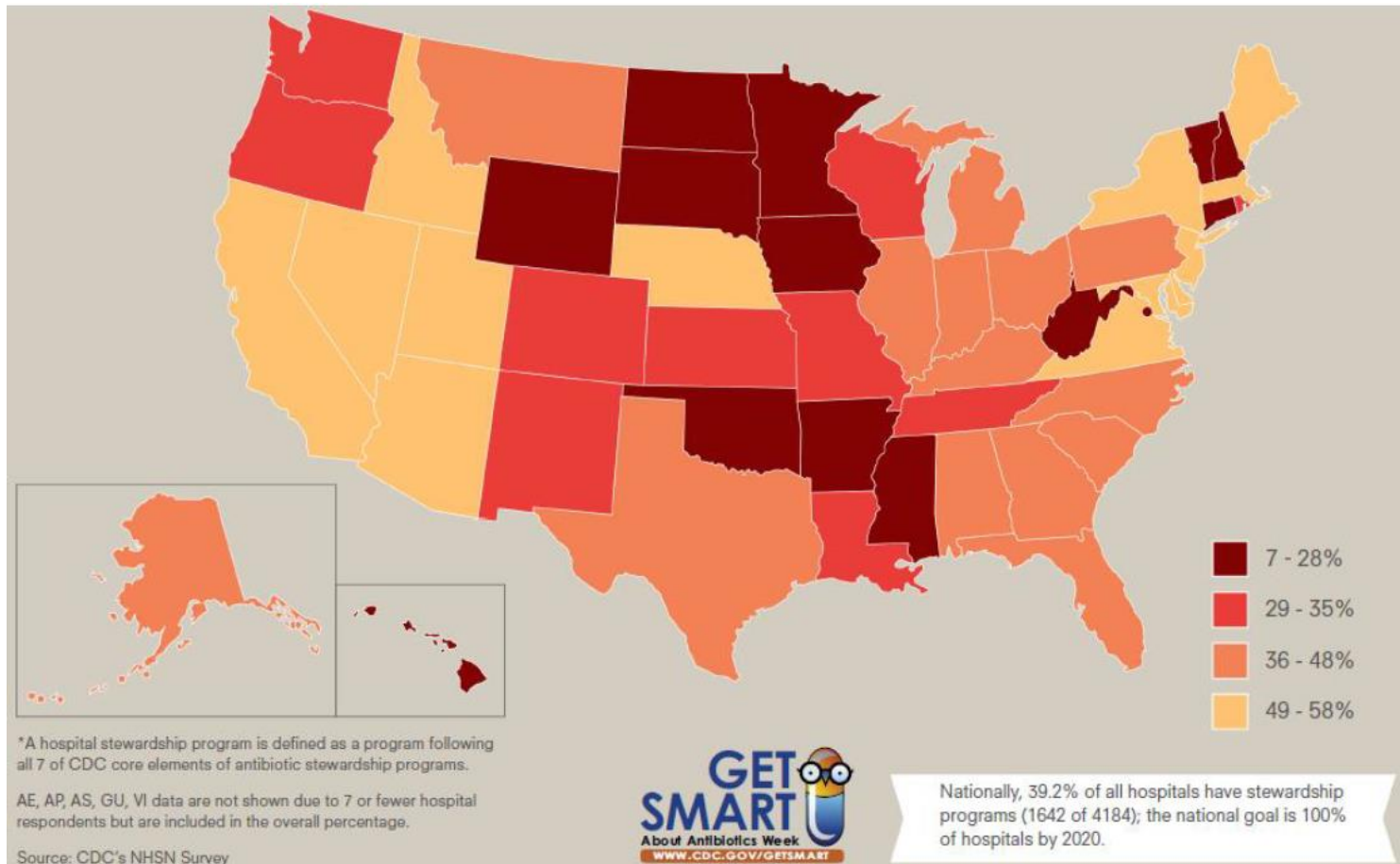
<sup>1</sup>Section of Infectious Diseases, Boston University School of Medicine, Boston, Massachusetts; <sup>2</sup>Division of Infectious Diseases, Johns Hopkins University School of Medicine, Baltimore, Maryland; <sup>3</sup>Division of Infectious Diseases, University of Miami Miller School of Medicine, Miami, Florida; <sup>4</sup>Department of Clinical Pharmacy, School of Pharmacy, University of California, San Francisco; <sup>5</sup>Department of Medicine, Weill Cornell Medical Center/New York–Presbyterian Hospital, New York, New York; <sup>6</sup>Department of Internal Medicine, Texas A&M Health Science Center College of Medicine, Houston; <sup>7</sup>Division of Healthcare Quality Promotion, Centers for Disease Control and Prevention, Atlanta, Georgia; <sup>8</sup>Division of Allergy and Infectious Diseases, University of Washington School of Medicine, Seattle; <sup>9</sup>Department of Medicine, Case Western Reserve University and Veterans Affairs Medical Center, Cleveland, Ohio; <sup>10</sup>Department of Medicine, University of Pennsylvania Health System, Philadelphia; <sup>11</sup>Hamilton House, Virginia Beach, Virginia; <sup>12</sup>Division of Infectious Diseases, Denver Health, Denver, Colorado; <sup>13</sup>Department of Anesthesiology and Critical Care Medicine, Johns Hopkins University Schools of Medicine and Nursing, Baltimore, Maryland; <sup>14</sup>Division of Infectious Diseases, University of Michigan Health System, Ann Arbor; <sup>15</sup>Department of Emergency Medicine, University of California, Davis; <sup>16</sup>Department of Emergency Medicine, David Geffen School of Medicine, University of California, Los Angeles Medical Center, Sylmar; <sup>17</sup>Department of Veterans Affairs, Hines, Illinois; <sup>18</sup>Department of Pediatrics, Washington University School of Medicine in St. Louis, Missouri; <sup>19</sup>Section on Infectious Diseases, Wake Forest University School of Medicine, Winston-Salem, North Carolina; <sup>20</sup>Department of Veterans Affairs and University of Utah, Salt Lake City; <sup>21</sup>Infectious Diseases, Memorial Sloan Kettering Cancer Center, New York, New York; and <sup>22</sup>Trivedi Consults, LLC, Berkeley, California

Evidence-based guidelines for implementation and measurement of antibiotic stewardship interventions in inpatient populations including long-term care were prepared by a multidisciplinary expert panel of the Infectious Diseases Society of America and the Society for Healthcare Epidemiology of America. The panel included clinicians and investigators representing internal medicine, emergency medicine, microbiology, critical care, surgery, epidemiology, pharmacy, and adult and pediatric infectious diseases specialties. These recommendations address the best approaches for antibiotic stewardship programs to influence the optimal use of antibiotics.

**Keywords.** antibiotic stewardship; antibiotic stewardship programs; antibiotics; implementation.



Downloaded from <http://ajph.org/> on November 10, 2015



# **NATIONAL EFFORTS FOR ANTIMICROBIAL STEWARDSHIP**

# APPROVED: New Antimicrobial Stewardship Standard

The Joint Commission recently announced a new Medication Management (MM) standard for **hospitals, critical access hospitals, and nursing care centers**. Standard MM.09.01.01 addresses antimicrobial stewardship and becomes **effective January 1, 2017**.

Current scientific literature emphasizes the need to reduce the use of inappropriate antimicrobials in all health care settings due to antimicrobial resistance.

k:  
dship

1.C.9 The hospital has written policies and procedures whose purpose is to improve antibiotic use (antibiotic stewardship).

☐ Yes

☐ No

1.C.10 The hospital has designated a leader (e.g., physician, pharmacist, etc.) responsible for program outcomes of antibiotic stewardship activities at the hospital.

☐ Yes

☐ No

1.C.11 The hospital's antibiotic stewardship policy and procedures requires practitioners to document in the medical record or during order entry an indication for all antibiotics, in addition to other required elements such as dose and duration.

☐ Yes

☐ No

1.C.12 The hospital has a formal procedure for all practitioners to review the appropriateness of any antibiotics prescribed after 48 hours from the initial orders (e.g., antibiotic time out).

☐ Yes

☐ No

1.C.13 The hospital monitors antibiotic use (consumption) at the unit and/or hospital level.

☐ Yes

☐ No



No citation risk for 1.C.9 through 1.C.13; for information only.

# NATIONAL STRATEGY GOALS

1

Slow the development of resistant bacteria and prevent the spread of resistant infections

2

Strengthen national one-health surveillance efforts to combat resistance

3

Advance development and use of rapid and innovative diagnostic tests for identification and characterization of resistance bacteria

4

Accelerate basic and applied research and development for new antibiotics, other therapeutics and vaccines

5

Improve international collaboration and capacities for antibiotic resistance prevention, surveillance, control and antibiotic research and development

# CDC CORE ELEMENTS

<http://www.cdc.gov/getsmart/healthcare/implementation/core-elements.html>

## Leadership Commitment

- Formal Statement
- Designated resources: human, financial, IT

## Accountability

- Single leader responsible for program outcomes

## Drug Expertise

- Single pharmacist leader

## Action

- Implement improvement

## Tracking

- Regular reporting on antibiotic prescribing and resistance

## Reporting

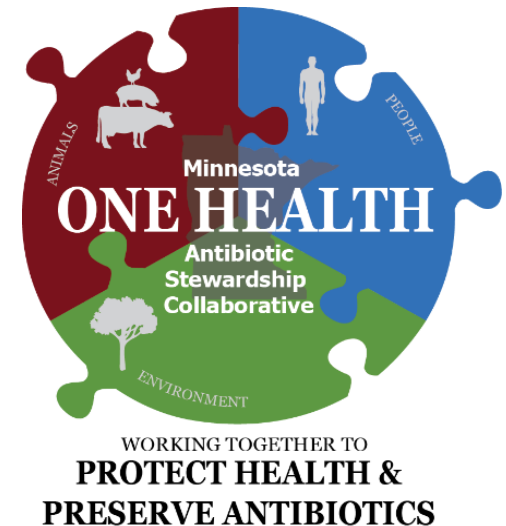
- Regular reporting on antibiotic use and resistance to doctors, nurses and relevant staff

## Education

- Educating clinicians about resistance and optimal prescribing

# STATE-WIDE COLLABORATIVE: MN ONE HEALTH ANTIMICROBIAL STEWARDSHIP

- **Multi-partner initiative to address antibiotic use**
- **Inter-agency approach by government**
  - Minnesota Department Health
  - Department of Agriculture
  - Pollution Control Agency
  - Board of Animal Health
- **Stakeholders from academia, clinical practice, health and agriculture advocacy groups**
- **Mission**
  - *Provide a collaborative environment to promote judicious antibiotic use and reduce the impact of antibiotic resistant pathogens*
- **Vision**
  - *Minnesota leaders in human, animal, and environment health will work together to raise awareness and change behaviors to preserve antibiotics and treat infections effectively*



## One Health Antibiotic Stewardship

One Health Antibiotic Stewardship Home

About Us

News and Events

State Plan and Data

Footprint Model

Resource Library

For Everyone  
for Human Healthcare  
for Animal and  
Agriculture  
for Environmental

Contact Us



# WORKING TOGETHER TO PROTECT HEALTH & PRESERVE ANTIBIOTICS



## Minnesota One Health Antibiotic Stewardship Collaborative

### What is One Health Antibiotic Stewardship?

Antibiotics are powerful tools for fighting and preventing infections. However, widespread use of antibiotics has resulted in an alarming increase in antibiotic-resistant infections. Antibiotic stewardship consists of coordinated interventions that promote judicious antibiotic use and reduce the impact of antibiotic resistant pathogens. A One Health approach recognizes that human, agricultural and companion animal, and environmental health are interconnected, and issues such as antibiotic stewardship require a collaborative effort across multiple disciplines. We believe that a One Health approach will create an informed public and professionals that can communicate, and practice a more holistic approach to antibiotic use.

### Workgroup member affiliations

Abbott Northwestern Hospital  
Allina Health  
Association for Professionals in Infection Control and Epidemiology – Minnesota  
Blue Cross Blue Shield  
Children's Hospitals and Clinics of Minnesota  
Emergency Physicians Professional Association- Minnesota  
HealthEast  
HealthPartners  
Hennepin County Medical Center  
Land O'Lakes  
Leading Age Minnesota  
M Health  
Mayo Clinic  
Merck Research Labs  
Metropolitan Council

### About One Health Antibiotic Stewardship

Learn what antibiotic resistance is, its effect on humans, animals, and the environment, and why One Health Antibiotic Stewardship is important.

### Current State Plan and Data

Review the state plan, its progress. Find Minnesota antibiotic (antimicrobial susceptibility) data.

News and Events

Footprint Model

### Resource Library



RESOURCE LIBRARY FOR  
Everyone



RESOURCE LIBRARY FOR  
Human  
Healthcare



RESOURCE LIBRARY FOR  
Animal Healthcare  
and Agriculture



RESOURCE LIBRARY FOR  
Environmental  
Sciences



### Stay Informed!

- ▶ [Subscribe to Minnesota One Health Antibiotic Stewardship Collaborative Updates](#)
- ▶ [Contact Us](#)

# ANTIMICROBIAL STEWARDSHIP AT FAIRVIEW-UMMC

## **January 2007-UMMC implemented first Antimicrobial Stewardship Program**

- Co-leadership with Dr. Susan Kline and Dr. Kimberly Boeser
  - 0.5 FTE ID staff physician support
  - 1.0 FTE Infectious Diseases Clinical Pharmacist
- Developed restricted antimicrobial guidelines and disease state guidelines
  - 2014 approved as system guidelines
- Expanded training
  - 2012 PGY2 ID resident
  - 2013 ID fellows training
  - 2015 added a second PGY2 ID resident
  - 2 U of MN College of Pharmacy students per block
- Maintained data collection from 2007-2014
  - Interventions and acceptance rates
  - ABX \$/pt day
  - Tracking of MDR-pathogens and *C. difficile* (Infection Prevention)
  - Morbidity and Mortality (hospital-wide)

## **January 2014-Fairview Southdale and Ridges Hospitals implemented ASPs**

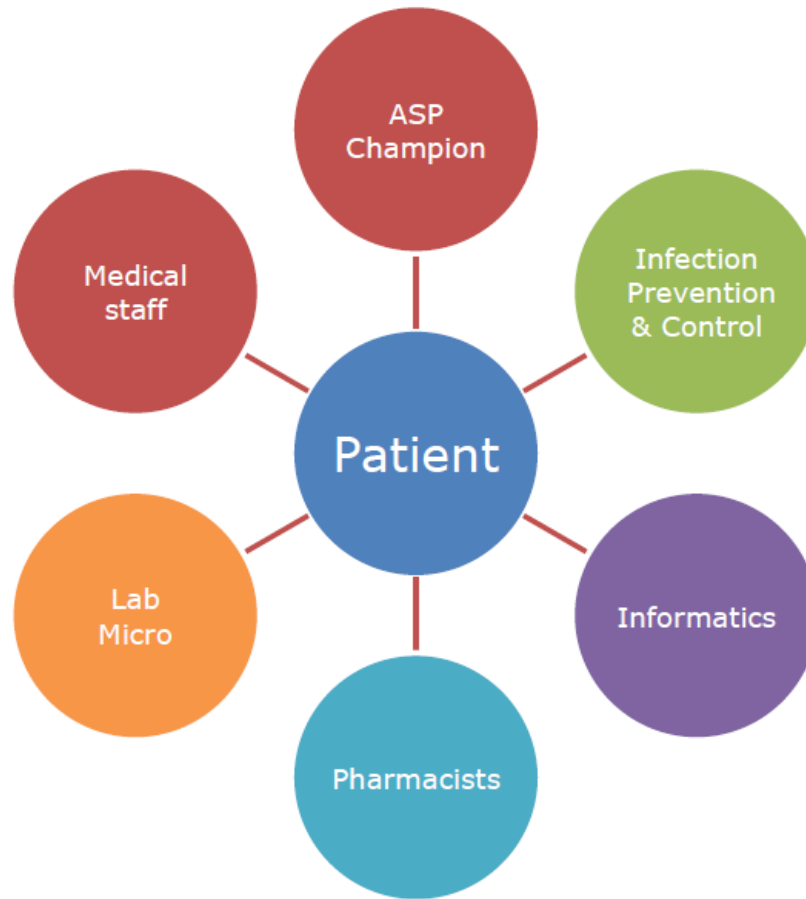
- Co-leadership with Dr. Steve Dittes and Dr. Michelle Borchart, Dr. Emily Medcraft and Dr. Ron Greenberg
- Round 2-3 days a week with ID staff on patients on restricted antimicrobials



# STRUCTURE OF OUR ASP TEAM

- ID PharmD + ID staff (+/- Pharmacy resident(s)-PGY1 or PGY2 ID, ID fellow, pharmacy students)
- Patients are flagged by order for restricted antimicrobial (34) and focus on 12 standard interventions
- Pharmacy team reviews all patients and determine which patients to round on with ID staff
- Round daily with ID staff
- Review all pertinent labs, imaging, drug profile, and cultures and make determination for ongoing antimicrobial approvals
- Place a progress note in the electronic medical record (EMR)
- Communicate with primary teams
- Follow-up on recommendations and monitor for adverse effects and duration of therapy of antimicrobials

# MULTIDISCIPLINARY APPROACH



# CURRENT EXPANSION EFFORTS

- Formal Antimicrobial Stewardship Programs at 3 out of 7 system hospitals
  - Joint efforts between Infectious Disease Providers, Pharmacists, and Infection Prevention
  - Active intervention with real-time feedback
- Current efforts to expand to all 7 hospitals
- Site Gap Analysis were completed 12/2016, updated 7/2017
- System-wide Antimicrobial Stewardship Steering Committee was developed & Project Plan Created
  - Kick-off meeting in May->Project time-line completion by 12/2017
- System Coordinator Role to take effect in May 2017
- Primary goal is to ensure regulatory compliance at all sites
- IT investments for Antimicrobial Stewardship and Infection Prevention (Epic ICON 500)
- Tracking and monitoring-Antibiotic Use and Antibiotic Resistance
  - CDC NHSN AUR module will be the gold standard for reporting and tracking
- Long-term goals->expansion to long term care/rehab facilities and ambulatory/clinic settings

# **LEVERING TECHNOLOGY AND THE EMR FOR AS PRACTICE**

## Leadership Commitment

- Formal Statement
- Designated resources: human, financial, IT

## Accountability

- Single leader responsible for program outcomes

## Drug Expertise

- Single pharmacist leader

## Action

- Implement improvement

## Tracking

- Regular reporting on antibiotic prescribing and resistance

## Reporting

- Regular reporting on antibiotic use and resistance to doctors, nurses and relevant staff

## Education

- Educating clinicians about resistance and optimal prescribing

# ACTION

- **Patient Identification and Assessment**
  - Report generated daily for patient started on a restricted antimicrobial
  - AS clinical tool built in Epic to gain efficiency in clinical review
  - Documentation in the EMR
  - Restricted Antimicrobial Guidelines and Disease State Guidelines
- **Verigene® Gram negative bacteremia AS Response Team**
  - AMT on-call 7days a week 8a-10p
  - Microlab pages AMT on-call for all positive blood cultures with gram negative pathogens
  - AMT/ID pharmacist makes assessment of patient, rounds on patient with ID staff or discusses over the phone
  - Treatment algorithm designed based on validation and antibiogram and order sets built for treating highly, multi-drug resistant bacteremias
  - Standard documentation via an AMT note is placed in the chart, primary medical team is paged
- **Cdiff Risk Assessment Tool**
  - Join effort with IP and EVS
  - 3 pronged approach to early intervention: pharmacist medication stewardship, provider, staff, patient and family education by IP, enhanced bleach cleaning by EVS
  - Physician champions-Dr. Susan Kline and Dr. Alison Galdys
- **Medication Use Evaluations**
  - ~6-8 completed annually, system-wide
  - Examples for 2017: carbapenems, fluoroquinolones, micafungin, vancomycin
  - Findings presented at our System Antibiotic Subcommittee with actions/recommendations for improvement of utilization
- **PCN Allergy Assessment/PAST**
  - Collaboration with ID Division
  - Physician champion- Dr. Pia Franco
  - Training received from ALK

# EPIC PATIENT LISTS AND IVENTS

Restricted Antimicrobials - UMMC East 106 Patients										
Refreshed just now  Search All My Lists										
Unit	AMT Follow-Up	Patient Name	MRN	Admit Date	Service	RX ID Consulted	antimicrob	AMT DOT - Non-restricted	AMT DOT - Restricted	RX Antimicrobial Stewardship Notes
UU U4E		<div></div>		06/22/2...	Cardiol...	✗	✓	12	19	+ ID consult  VF arrest s/p ECMO and PNA vs pulm edema Klebsiella, Enterobacter, MSSA, Acinetobacter, Stenotrophomonas in sputum  6/28 ID recs -Start meropenem
UU U4E				06/26/2...	Cardiot... Surgery	✗	✓	—	16	Post-op infection vs HCAP. PAST candidate? d1= 7/4-7/12 Merrem
UU U5A				07/06/2...	Hospita...	✗	✓	4	4	7/3 sputum cx no orgs AMT to review 7/10 Possible PAST candidate HAP Aztreonam (7/8- Vanc (7/8- Lvq (7/8-
UU U5A				07/09/2...	Pulmon...	✗	✓	3	3	CF exacerbation (h/o steno, ESBL E. Coli) Mero (7/9- ); nafcillin - hives LVQ (7/9- TMP/SMX (7/9- Tobi nebs (7/9-  7/10 rec - consider ertapenem

**Antimicrobial Stewardship Notes** [Comment](#)

AMT to review 7/10  
Possible PAST candidate  
HAP  
Aztreonam (7/8-  
Vanc (7/8-  
Lvq (7/8-

Last edited by Kobic, Emir, RPH on 07/10/17 at 0929

**General Information**

Type:

Subtype:

Status:

Significance:

Value:

ent:  minutes

Outcomes:

Ant:

Re:

**Associated Orders**

**Documentation**

Title	Number
AMT - appropriate therapy	1021
AMT - follow-up in 24-48 hours	1022
AMT - ID Consult	1032
AMT - will review TODAY	1023

**Associated Users**

User	Role
------	------

# AS MONITORING TOOL

## Gram stain [351709450]

Order Status: Completed

Specimen Description  
Special Requests  
Gram Stain

Specimen: Sputum  
Sputum  
Screen

--  
<10 Squamous epithelial cells/low power field  
<25 PMNs/low power field  
Moderate Mixed gram positive and gram negative bacteria present.  
FINAL 07/10/2017

Collected: 07/10/17 1945

Updated: 07/10/17 2214

## Sputum Culture Aerobic Bacterial [351709449]

Order Status: Completed

Specimen Description  
Special Requests  
Culture Micro  
Micro Report Status

Specimen: Sputum

Specimen: Sputum  
Sputum  
Screen  
Moderate growth Normal flora  
FINAL 07/10/2017

Collected: 07/10/17 1945

Updated: 07/10/17 2117

## Sputum Culture Aerobic Bacterial [351452341]

Order Status: Completed

Specimen Description  
Special Requests  
Culture Micro  
Micro Report Status

Specimen: Sputum  
Sputum  
Screen  
Moderate growth Normal flora  
FINAL 07/10/2017

Collected: 07/08/17 1140

Updated: 07/10/17 1302

## Strep pneumo Agn Ur greater or equal to 13yrs or CSF any age [351485336]

Order Status: Completed

Specimen Description  
S Pneumoniae Antigen

Specimen: Urine  
Unspecified Urine

--  
Negative, no Streptococcus pneumoniae antigen detected by immunochromatographic membrane assay. A negative Streptococcus pneumoniae antigen result does not rule out infection with Streptococcus pneumoniae.  
FINAL 07/09/2017

Collected: 07/08/17 1745

Updated: 07/09/17 0006

## Gram stain [351479222]

Order Status: Completed

Specimen Description  
Special Requests  
Gram Stain

Specimen: Sputum  
Sputum  
Screen

--  
<10 Squamous epithelial cells/low power field  
<25 PMNs/low power field  
Moderate Mixed gram positive and gram negative bacteria present.  
FINAL 07/08/2017

Collected: 07/08/17 1140

Updated: 07/08/17 1543

## Radiology (Last 504 hours)

07/08 1139

CT Chest Pulmonary Embolism w Contrast

[Images](#)

07/07 1841

XR Chest Port 1 View

[Images](#)

07/07 0711

CT Abdomen Pelvis w Contrast

[Images](#)

05/29 0000

UIS IMAGING - HIM SCAN



# EMR DOCUMENTATION

University of Minnesota Medical Center, Fairview  
Antimicrobial Management Team (AMT) Note

**Antimicrobial Stewardship Program**-- *A joint venture between Fairview  
Pharmacy Services and UM Physicians to optimize antibiotic management*

*NOT a formal Consult-Restricted Antibiotic Review*

To:  
Unit:  
Allergies:  
Infection History:

Brief Summary:

HPI:

Interval History:

Assessment:

Recommendation/Interventions:

- 1).
- 2).
- 3).

Intervention Examples:

1. Change to more appropriate antibiotic based on lab data
2. Change to alternative unrestricted anti-infective
3. Discontinue one or more antibiotics (PO or IV)
4. Change from IV to PO antibiotics
5. Better empiric antibiotic therapy
6. Antibiotic dosage change
7. Consult recommended (eg. Infectious Disease, Pulmonary/Critical Care, Renal, Urology, etc.)
8. Additional/Further diagnostic testing recommended
9. Simplify antibiotic regimen (eg. Inpatients on redundant or excessively broad spectrum antibiotics)
10. Recommend change in post-op antibiotic duration
11. Other (Duration of Therapy)
12. Agree with management

Discussed w/ ID Staff-Dr. XX

Current ~~Antibiotic~~ therapy:

Previous Antibiotic therapy:

Vital Signs and other clinical features:

Temperature

Imaging:

Culture Results:

Date	Culture Site	Organism

## Meropenem (Merrem®) Use Guidelines:

**Reasons for restriction:** To prevent unnecessary use and preserve the efficacy for severe infections. Meropenem will be reserved for infections that are polymicrobial and/or contain resistant gram negative bacteria such as *Pseudomonas*, *Enterobacter*, *Serratia* or an ESBL producing *E. coli* or *Klebsiella*. It is also restricted due to costs. For example *Pseudomonas* susceptible to piperacillin/tazobactam should not be treated with meropenem unless the patient has an IgE mediated allergy noted (see page 2).

### FDA Approved Indications:

- Skin or soft tissue infection
- Intra-abdominal infection
- Bacterial Meningitis
- Pediatric-bacterial meningitis

### Fairview Indications:

- Multidrug resistant gram negative pathogens, e.g. *Pseudomonas aeruginosa*, *Burkholderia cepacia*, or Extended Spectrum Beta Lactamase (ESBL) gram negative bacteria which are only sensitive to meropenem
  - Ertapenem is a formulary carbapenem that has excellent susceptibility to ESBL producing organisms and may be considered an alternative over meropenem if Pseudomonal coverage is not necessary based on site of infection or documented cultures.
- A dose of 500mg IV Q6H is the approved dosing for all indication except Cystic Fibrosis patients and CNS infections.
  - Higher dose meropenem may be used if a documented pathogen has a higher MIC reported of  $\geq 2$ .
  - 500 mg IV Q6H is our standard dosing as many of our gram negative pathogens maintain low susceptibility MICs to meropenem and utilizing lower dose but closer frequency (500 mg IV Q6H vs. 1 gm IV Q8H) optimizing time dependent killing

### Dosing:

#### Adult:

Standard dose: 500 mg IV Q6H  
Standard dosing for CF patients: 1 gm IV Q8 hours  
Bacterial meningitis: 2 gm IV Q8H

#### Pediatric:

Bacterial meningitis  $\geq 3$  months = 40 mg/kg IV Q8 hours  
Skin and/or SQ tissue = 10 mg/kg IV Q8 hours  
Complicated abdominal infections = 20 mg/kg IV Q8 hours

### Cost:

Antibiotic	Cost/dose	Cost/day	Alternative Agent	Cost/dose	Cost/day
Meropenem 500 mg vial	\$6.40	\$25.60			
Meropenem 1 gm vial	\$12.81	\$38.43			
			Primaxin (Imipenem/Cil) 500 mg vial	\$7.50-8.50	\$30.00-34.00
			Primaxin	\$4.50	\$18.00

# VERIGENE®

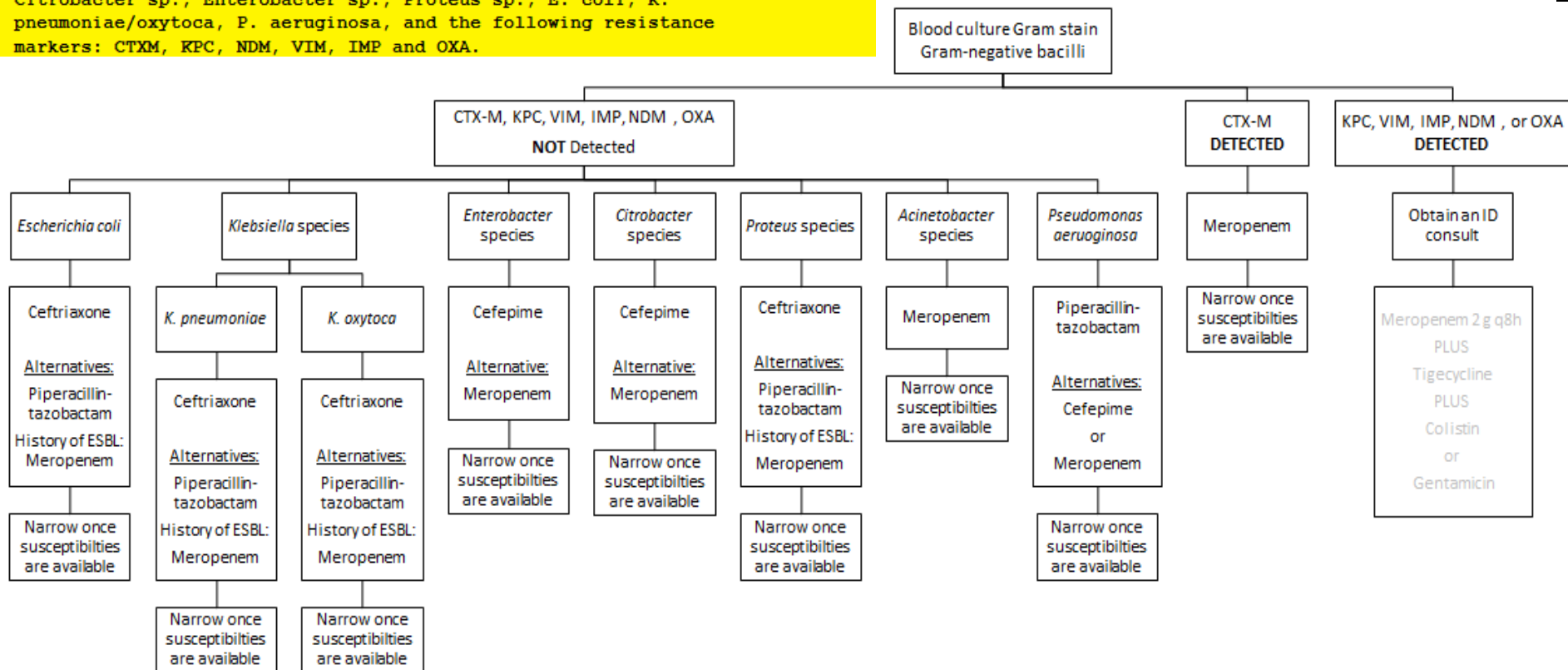
## GRAM-NEGATIVE BLOOD CULTURE ASSAY (BC-GN)

Cultured on the 1st day of incubation: *Klebsiella pneumoniae*  
Critical Value/Significant Value, preliminary result only, called to and read back by MARK O. RN @2312 4/1/17. CT

(Note)

POSITIVE for KLEBSIELLA PNEUMONIAE by Verigene multiplex nucleic acid test. Final identification and antimicrobial susceptibility testing will be verified by standard methods.

Specimen tested with Verigene multiplex, gram-negative blood culture nucleic acid test for the following targets: *Acinetobacter* sp., *Citrobacter* sp., *Enterobacter* sp., *Proteus* sp., *E. coli*, *K. pneumoniae/oxytoca*, *P. aeruginosa*, and the following resistance markers: CTXM, KPC, NDM, VIM, IMP and OXA.



# CDIFF RISK ASSESSMENT TOOL

CDI Pilot Units (153 Patients)					Last Refreshed: 1528	Search All My Lists
UNIT	Patient Name/Age/Sex	C-DIFF RISK SCORING SYSTEM Score Column	C Diff Follow-Up	C-Diff score change	C-DIFF RISK SCORING SYSTEM Time Since Reviewed Column	Admission Date
UUU6C		22		=	26 hrs 18 mins	2/25/16
UU5CBM		21		=	16 hrs 58 mins	12/30/16
UU5CBM		19		=	477 hrs 56 mins	11/3/16
UUU4E		19		=	17 hrs 6 mins	9/16/16
UUU4E		19		19		12/30/16
UU5CBM		18		18		12/28/16
UU5CBM		18		18		12/30/16

[Rx Snapshot](#)
[Antimicrobial Stewardship](#)
[Rx Med Rec at Admn](#)
[MAR ADMINISTRATIONS](#)
[Rx FV TPN Monitoring](#)
[c-diff risk \(testing\)](#)
 Report:

C-DIFF Risk : 22 [Last reviewed: Mason, Jocelyn, RPH at 01/02/17 1310] [Comment](#)

Pharmacy CDI Review Complete on 12/12/2016 11:42 AM by Jocelyn Mason  
 Intervention: discontinue PPI  
 Estimate Time Spent: 10 min

IP CDI Review Complete on 12/12/2016 11:48 AM by Jocelyn Mason  
 EVS Notified: Yes  
 Sign Placed: Yes  
 Education: RN/Pt/Family/Provider  
 EVS Cleaned: Yes  
 Estimate Time Spent: 15 min

Last edited by Mason, Jocelyn, RPH on 01/02/17 at 1310

- [PPI active in last 60 days: 2 points - \[Last updated: 01/03/17 1529\]](#) [Comment](#)
- [Fluoroquinolone ordered in last 90 days: 1 points - \[Last updated: 01/03/17 1529\]](#) [Comment](#)
- [Age 50-80 years: 1 points - \[Last updated: 01/03/17 1529\]](#) [Comment](#)
- [At least 1 day in ICU \(1 point\): 1 points - \[Last updated: 01/03/17 1529\]](#) [Comment](#)
- [Readmitted in last 30 days: 2 points - \[Last updated: 01/03/17 1529\]](#) [Comment](#)
- [Most recent albumin < 3.5 : 1 points - \[Last updated: 01/03/17 1529\]](#) [Comment](#)
- [Currently on antineoplastic medication: 3 points - \[Last updated: 01/03/17 1529\]](#) [Comment](#)
- [History of C-diff diagnosis or positive result in last year : 3 points - \[Last updated: 01/03/17 1529\]](#) [Comment](#)
- [Antibiotic ordered in last 90 days: 1 points - \[Last updated: 01/03/17 1529\]](#) [Comment](#)
- [Beta Lactam AND glycopeptide ordered in last 90 days: 2 points - \[Last updated: 01/03/17 1529\]](#) [Comment](#)
- [Beta-lactam AND Fluoroquinolone ordered in last 90 days: 2 points - \[Last updated: 01/03/17 1529\]](#) [Comment](#)
- [Transplant Patient \(or on wait list\): 3 points - \[Last updated: 01/03/17 1511\]](#) [Comment](#)

# PCN ALLERGY ASSESSMENT PROGRAM

Type:

Service:

Date of Service:

07:49 AM

☐ Cosign Required

Have you ever taken drugs similar to penicillin?

Cephalosporins such as Cephalexin (Keflex), Cefepime (Maxipime), Ceftriaxone (Rocephin) Carbapenems such as Imipenem (Primaxin), Meropenem (Merrem), Etrapanem (Invanz)

\*\*\*

Procedure was reviewed and discussed with the patient and or family member: {YES NO:124710::"Yes"}

The patient met University of Minnesota Medical Center Penicillin Skin Test Protocol for Penicillin Allergy Skin Testing: {YES NO:124710::"Yes"}

## Step 1: Puncture Test Results:

The test was read after \*\*\* minutes of placement.

\*\*\*Insert picture

Histamine control response (+): {YES NO:124710::"Yes"} (\*\*mm)

Normal saline control response (-): {YES NO:124710::"Yes"} (\*\*mm)

PRE-PEN response (PRP): {YES NO:124710::"Yes"} (\*\*mm)

Penicillin G response (PG): {YES NO:124710::"Yes"} (\*\*mm)

## Assessment:

The histamine response was {Positive / Negative:124514}, determining that the patient \*\*\*does or does not have the ability to mount an allergic reaction.

The normal saline response was {Positive / Negative:124514}, determining that the patient \*\*\*does or does not have skin too sensitive for testing.

The penicillin and PRE-PEN sites \*\*\*did or did not produce larger wheals than the controls or were {Positive / Negative:124514} for any reaction and therefore, the patient \*\*\*does or does not appear initially allergic and the testing \*\*\*can or cannot continue to the second portion of testing with intradermal testing.

## Step 2: Intradermal Test Results:

The test was read after \*\*\*minutes of placement.

\*\*\* Insert picture here

## Leadership Commitment

- Formal Statement
- Designated resources: human, financial, IT

## Accountability

- Single leader responsible for program outcomes

## Drug Expertise

- Single pharmacist leader

## Action

- Implement improvement

## Tracking

- Regular reporting on antibiotic prescribing and resistance

## Reporting

- Regular reporting on antibiotic use and resistance to doctors, nurses and relevant staff

## Education

- Educating clinicians about resistance and optimal prescribing

# EPIC ICON MODULE IMPLEMENTATION

- **Required the upgraded version of Epic in order to build the reporting structure**
- **Significant financial investment from Fairview and MHealth for IT infrastructure upgrades**
  - Epic upgrade
  - Currently undergoing Sunquest LIS (MicroLab) upgrade
- **Currently in the build and testing the functionality of the module components**
  - Goal is to utilize the module (Ivents) for total assessment, intervention and documentation->allows easy method for tracking and reporting
  - Leveraging our technology to gain efficiencies with clinical activities
- **Creating a AS Dashboard**
- **Hurdles: Technology upgrades take time, time intensive to build and test, many different groups involved and everyone has their own priorities**
- **Go-Live: October 2<sup>nd</sup>, 2017**

# NATIONAL HEALTHCARE SAFETY NETWORK (NHSN) AU(R) MODULE

- AUR=Antibiotic Use and Resistance module
- Launched in 2012
- Goal=provide a mechanism for facilities to report and analyze antimicrobial use and/or resistance as part of local or regional efforts to reduce antimicrobial resistance
- Must coordinate with their laboratory and/or pharmacy information software providers to configure their system to enable the generation of standard formatted file(s) to be imported into NHSN
- Data can be system wide, hospital specific, and/or unit specific
- Ideal state=aggregate information of antibiotic use at a regional or national level->create antibiotic benchmarks



# STANDARDIZE ANTIMICROBIAL ADMINISTRATION RATIO (SAAR)

- **Measurement for tracking and reporting continues to be difficult and facilities risk not being compliant with this core element**
- **SAAR could be used for benchmarking for antibiotic use**
  - Same concept as Standard Infection Ration (SIR) that Infection Prevention uses
  - Gives the observed antimicrobial use over predicted
  - May allow for risk adjusted comparisons
- **CDC has been collaborating with organization to make this module doable and helpful**
  - Grant was available from the CDC for Departments of Health
- **SAAR may be more helpful if reported by specific patient populations or grouped by antibiotics**
  - Agents mainly for healthcare associated pathogens
  - Agents mainly for community pathogens
  - Agents active against MRSA
  - Agents frequently use for surgical prophylaxis
  - All agents

# NHSN AUR MODULE

## National Healthcare Safety Network

### Rate Table - All Submitted AU Data - Antimicrobial Utilization Rates by Location

#### Rate per 1,000 Days Present

As of: February 3, 2012 at 3:52 PM

Date Range: All AU\_RATELOCATION

Org ID=10846 CDC Location=IN:ACUTE:CC:M Location=INMEDCC

Summary Yr/Mon	Antimicrobial Category	Antimicrobial Class	Antimicrobial Days	Days Present	Rate per 1000 Days Present
2011M01	Antibacterial	-- All --	90165	10000	9,016.500
2011M01	Antibacterial	Aminoglycosides	438	10000	43.800
2011M01	Antibacterial	Carbapenems	12	10000	1.200
2011M01	Antibacterial	Cephalosporins	57	10000	5.700
2011M01	Antibacterial	Fluoroquinolones	12	10000	1.200
2011M01	Antibacterial	Folate pathway inhibitors	6	10000	0.600

# **NATIONAL STRATEGY OBJECTIVES**

- **95% of eligible hospitals report antibiotic use data to National Healthcare Safety Network Antibiotic Use & Resistance by 2020**
- **Reduce inappropriate inpatient antibiotic use by 20% from 2014 levels**
- **Reduce inappropriate outpatient antibiotic use by 50% from 2010 levels**



A collaboration between University of Minnesota Physicians and University of Minnesota Medical Center

# ONGOING IMPACT OF AN ANTIMICROBIAL STEWARDSHIP PROGRAM AT A LARGE ACADEMIC MEDICAL CENTER, 9 YEARS OF EXPERIENCE

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## ABSTRACT

**Background:** The University of Minnesota Medical Center (UMMC), Fairview is a 300 bed tertiary care facility. UMMC has had a long-standing, comprehensive antimicrobial stewardship program (ASP).

**Methods:** The stewardship team is comprised of a full-time PharmD, and part-time ID staff members who rotate through the service. The team allows providers to order restricted antimicrobials, per hospital guidelines and policies, without upfront approval, followed by a chart review. Recommendations are placed in the electronic medical record as a progress note. Verbal recommendations may also be made. The number of patient on restricted antimicrobial, total number of interventions and acceptance rates, antimicrobial cost per pt day, antimicrobial utilization, and patient outcomes are evaluated annually.

**Results:** There was a downward trend in Hospital Acquired (HA) C. difficile diarrhea from 2007 to 2014 from 1.2 to 0.3/1000 patient days (pt day). Rates appear stable from 2014-2016 with adjustment for change to NHSN lab based surveillance only. From 2009-2014 a decrease was seen in HA VRE infections from 0.33 to 0.22/1000 pt days and in HA MRSA infections from 0.2 to 0.08/1000 pt days. VRE and MRSA HAI rates increased in 2015 - first quarter 2016. Newly acquired HA ESB infections increased from 2009-2016 at 0.09 to 0.20/1000 pt days. CRE is an emerging problem during the ASP history.

Cost savings, after adjusting for inflation, continued from year to year. The greatest cost savings was from 2006-08 in which antimicrobial costs/pt day declined by 7%, antibiotics costs declined by \$7.40/pt day. In 2012, we observed our lowest antibiotic cost/pt day at \$36.36 which is a difference of \$19.03 before implementation of the program. From 2013-2015, we have observed a sustained average antibiotic cost per patient day of \$42.84.

**Conclusion:** The ASP has continued to cost justify the program. Our antibiotic costs/pt day have leveled off in the last 3 years and remained low despite rising antibiotic costs due to market inflation and drug shortages. We began to observe a decrease in HA VRE and C. difficile infections after 3 years of operation, and MRSA after 5 years. The effects of the program and the Infection Prevention Department appear to be synergistic. Future areas for focus include preventing rising multi-drug resistant organism HAIs, focus on non-restricted antibiotics that are overused and contribute to C. difficile diarrhea and use of the procalcitonin test to help optimize antibiotic use.

## INTRODUCTION

**Objective:** To track and measure the impact of the antimicrobial stewardship program (ASP) on key outcome measures over time since implementation in 2007, compared to pre-intervention period of 2006.

- Interventions made by the ASP team (~1,900-2,400/year)
- Acceptance Rate (~80%/year)
- Morbidity, mortality, length of stay (LOS)
- Antibiotic costs
- Antimicrobial resistance trends
- Antibiotic usage (since institution of EPIC in 2011 downward trend)

## CLINICAL OUTCOMES

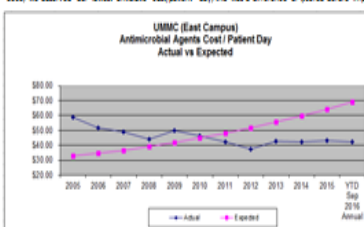
### University HealthSystem Consortium Data

Discharge Year	Cases	Mean LOS Observed	Mean LOS Expected	LOS Index	% Deaths Observed	% Deaths Expected	Mortality Index	UHC Risk Model
2006	28,071	5.21	5.33	0.98	1.66	1.98	0.84	Pre-2012 model
2007	29,194	5.19	5.36	0.97	1.71	1.93	0.89	"
2008	31,194	5.12	5.31	0.97	1.41	1.67	0.85	"
2009	30,312	5.28	5.58	0.95	1.53	1.83	0.84	"
2010	28,709	5.39	5.78	0.93	1.58	2.07	0.76	"
2011	28,805	5.62	5.80	0.97	1.86	2.22	0.84	"
2012	28,216	5.71	5.99	0.95	1.76	2.05	0.86	2012 model
2013	28,302	5.98	5.80	1.03	1.97	1.85	1.07	2015 model, 2012 model mortality index = 0.81
2014	27,738	6.02	5.93	1.02	2.06	2.01	1.03	2015 model
2015	26,701	6.10	6.34	0.96	2.22	2.32	0.96	"
2016 (Jan-Aug)	18,584	6.30	6.76	0.93	2.07	2.43	0.85	"

## ANTIMICROBIAL COSTS TABLE

Antimicrobial Agents Activity	2006	2007	2008	2009	2010	2011	2012	2013	2014	2015	YTD Sep 2016 Annual
Antimicrobial Agents Total Cost	\$ 5,712,500	\$ 4,254,176	\$ 4,041,570	\$ 4,398,065	\$ 5,061,640	\$ 5,706,207	\$ 5,321,557	\$ 5,077,654	\$ 4,089,295	\$ 4,280,557	\$ 4,254,155
Total Patient Days	76,731	95,709	95,937	95,902	101,940	97,479	95,345	95,404	95,404	95,671	103,780
Costs / Patient Day	\$69.02	\$44.57	\$42.05	\$45.97	\$49.84	\$56.40	\$55.82	\$53.23	\$42.85	\$44.74	\$41.23

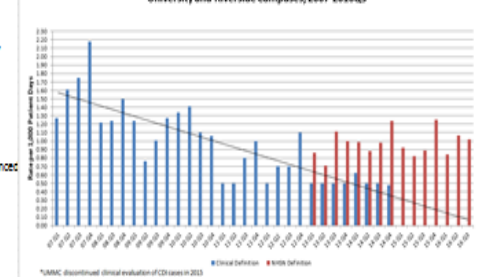
From 2006 to 2009 there was a \$2.47 increase in antimicrobial cost/patient day; primary expenditure increase was due to antifungal utilization and vancomycin resistant Enterococcus (VRE) treatment. Antibiotic cost was adjusted for percent inflation for the market. Inflation increased steadily from 2005-2011 (4.9%, 5.2%, 5.1%, 7.5%, 6.4%, 7.5% and 7.5%) while the University HAI (patient day) decreased. In 2011, the HAI (patient day) was below the expected market cost. In 2012, we observed our lowest antibiotic cost/patient day, this was a difference of \$19.03 before implementation of the stewardship program.



AMT Interventions & Acceptance Rates - Adult	2007	2008	2009	2010	2011	2012	2013
Total Interventions	1,991	2,165	2,665	2,340	2,098	2,264	2,417
Total Accepts	1,153	1,235	1,626	1,381	1,267	1,376	1,531
Total Declines	468	518	541	439	453	458	423
Total Agree w/Management	370	412	498	520	378	430	463

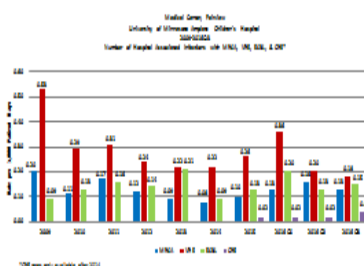
## CDI RATE TREND 2007-2016

Differences in Clinical and NHSN Surveillance Definitions of Hospital Acquired CDI, University and Riverside Campuses, 2007-2016<sup>1,2</sup>



\*HAI Definitions: Beginning in 2009 all positive VRE, MRSA and ESB infections, which are identified on a patient for the first time, from all hospital units are evaluated to determine if the culture represents a hospital acquired infection (HAI) as defined by NHSN diagnostic criteria. HAI includes: BSI, UTI, SSTI, SSI, pneumonias, bone and joint infections, cardiovascular, CNS, GI, EENT, URS, reproductive and systemic infections.  
\*Active surveillance cultures for VRE stool carriage are done in BMT patients  
\*and active surveillance for MRSA nasal carriage is done in ICUs.  
\*C. difficile HAI cases are also identified using NHSN criteria  
Additional infection control interventions were taken for C. difficile with enhanced environmental cleaning in those rooms with patients with CDI

## MULTIDRUG RESISTANT HAI RATES



## SUMMARY

- HAI trends, MRSA and VRE rates have decreased during the intervention period but ESB and CRE rates have increased.
- C. difficile diarrhea rate has decreased during the intervention period.
- Morbidity, mortality and LOS have remained stable since implementation of ASP.
- Antibiotic costs decline over time of ASP. Cost of ASP is justified.
- Effects of ASP and infection prevention appear to be synergistic.

# WHAT DO WE DO NOW?

- Keep fighting!
- Leadership and Engagement
- Education and Training
- Protect our patients
  - Quick identification and treatment
  - Active Surveillance
- Support local and national efforts including legislation
- Work with industry
- Be Stewards...everyone...including you!

Resistance anywhere is resistance everywhere.  
Antibiotic overuse increases the development  
of drug-resistant germs and  
limits treatment options for infections.  
You have a role in preventing antibiotic resistance.

# QUESTIONS & DISCUSSION

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# ASP 101 Reminders

## *MHA/OHA Acute Care ASP 101* Implementation Timeline

### Phase 1: CDC Core Elements 1-3

June 2017

#### Events

- ☐ **June 20** - ASP 101 Kick off webinar - overview of ASP initiative across the continuum of care

#### Homework

- ☐ Review Kansas DOH ASP Toolkit for Rural and Critical Access Hospitals pg. 1-14

July 2017

#### Action Items

- ☐ **July 11** - MHA/OHA collaborative webinar “Leveraging the EMR to Promote ASP activities” (Register [online](#))
- ☐ Develop an ASP team
- ☐ Draft a leadership ASP statement of support (example provided)

#### Due

- ☐ ASP team in place
- ☐ Leadership ASP statement of support for your facility

### Phase 2: CDC Core Element 4

August 2017

#### Events

- ☐ **Aug. 8** - ASP 101 Sharing call and presentation, “TJC ASP Lessons Learned” (Register [online](#))

#### Homework

- ☐ Review Kansas DOH ASP Toolkit for Rural and Critical Access Hospitals Pg. 15-24

September 2017

#### Action Items

- ☐ **Sept. 12** - MHA/OHA Collaborative Webinar (Register [online](#))
- ☐ Based on the facility ASP statement of support draft an ASP policy that supports optimal antibiotic use (example provided)

#### Due

- ☐ Facility specific ASP policy and procedure



# MHA/OHA HIIN Contacts

## ■ OHA

- James Guliano, Vice President Quality Programs
- Rosalie Weakland, Senior Director Quality Programs
- Subcontractor – HSAG
  - Christine Bailey, Director, Quality Improvement and Patient Safety

## ■ MHA

- Tania Daniels, Vice President, Quality and Patient Safety
- Lali Silva, Senior Director Quality and Process Improvement
- Susan Klammer, Quality & Process Improvement Specialist

# Thank you for joining us!

## **Next Webinar:**

**“TJC ASP Lessons Learned”**

**Tuesday, August 8 at 11:30am CST/ 12:30pm EST**

**Join online: <https://zoom.us/j/537497272>**