

Texas Department of State Health Services

VIM-CRPA in West Texas

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Objectives

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- Describe the cases of VIM-CRPA in West Texas
- Summarize the containment strategy used to slow the spread of VIM-CRPA

9/30/2019



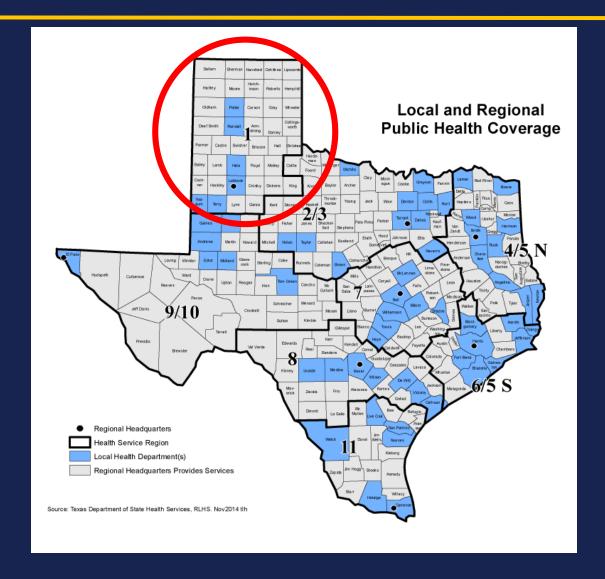
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Background

Prior to the Epi-Aid



Texas Public Health



- 254 counties
- 8 public health regions
- 64 local health departments

Texas Notifiable Multidrug-Resistant Organisms

- Carbapenem-resistant Enterobacteriaceae (CRE)
- Multi-drug resistant Acinetobacter (MDRA)
- Vancomycin Intermediate Staphylococcus aureus (VISA)
- Vancomycin Resistant Staphylococcus aureus (VRSA)



Source: Texas Notifiable Conditions List

Pseudomonas aeruginosa

- Gram-negative found in most environments
 - Soil, water, plants, animals
- Seen in community and hospitalacquired infections (HAIs)
 - Much more common in HAIs
 - 5th most common pathogen in all HAIs
 - 2nd most common in ventilator-associated pneumonias (VAPs)VAPs
- Admission to ICU, burns, neutropenia, or cystic fibrosis are significant risk factors



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MDRO P. aeruginosa

- Many resistance mechanisms exist
 - High rate of multidrug resistance compared to other organisms
- Resistance rates higher in LTACHs than ICUs
- Second most common MDRO found on healthcare workers during routine patient care (17.4%)
- Recovered in 22% of ICU rooms
- Environmental reservoirs contribute to spread
 - Water or humidity related
 - Showers, sinks, artificial nails, ultrasound gel, soap dispensers
 - Forms biofilms prevents penetration of cleaning agents; difficult to eradicate

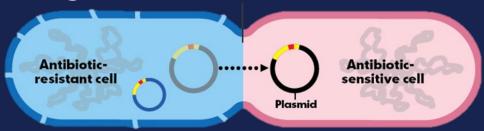


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Carbapenemases

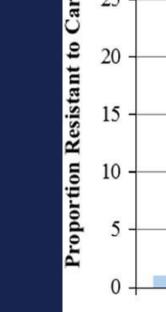
- 1. Multiple mechanisms lead to carbapenem resistance
- 2. Carbapenemases are enzymes that degrade carbapenem antibiotics
 - a. Often on mobile genes that can transfer to Gramnegative organisms



- 3. Common carbapenemases: KPC, NDM, VIM, OXA-48-type, and IMP
- 4. Found in Enterobacteriaceae, Pseudomonas aeruginosa, and Acinetobacter baumannii

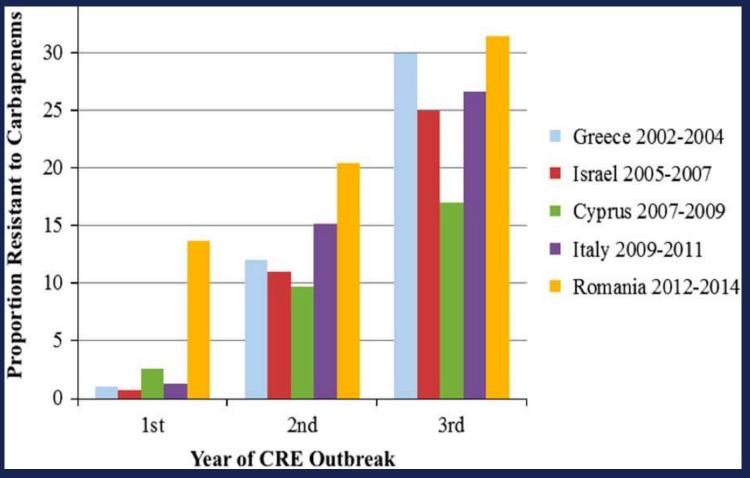


Carbapenemases Can Spread Rapidly



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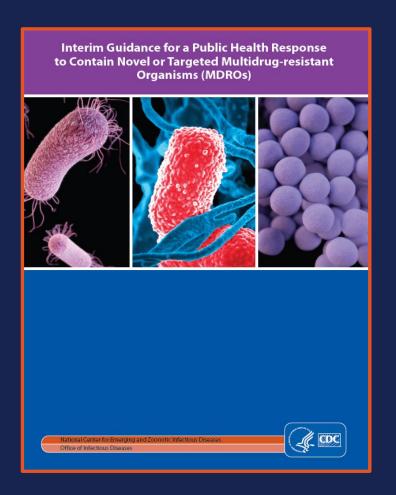
CDC Response Guidance



Health Services

Goal:

Slow spread of novel or rare multidrug-resistant organisms or mechanisms

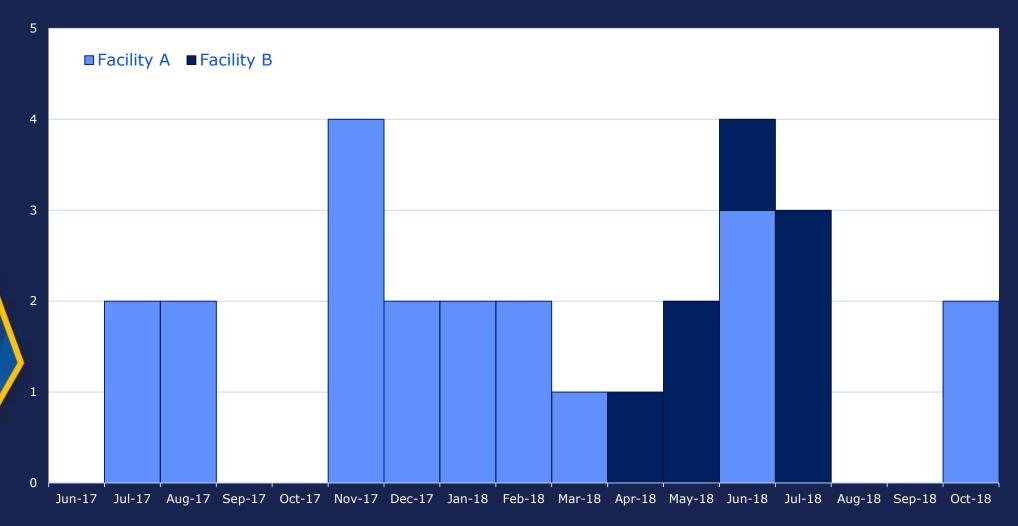


Initial 4 cases identified Summer 2017





Epidemic Curve of VIM-CRPA June 2017-October 2018





Characteristics of West Texas VIM-CRPA

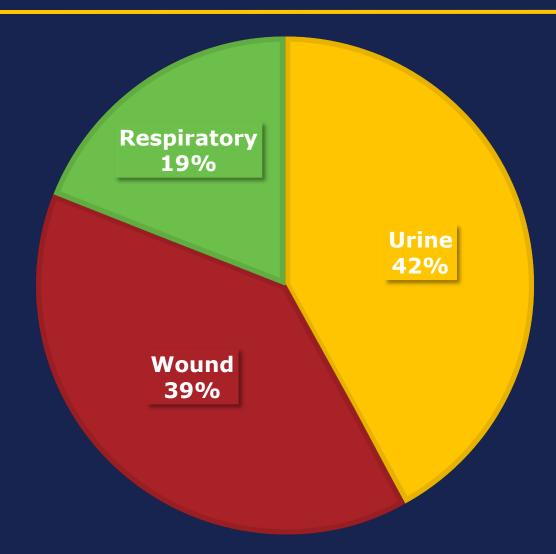
June 2017-October 2018

- 27 patients
 - 1 resident of New Mexico
- 62% Male
- Median age: 63 years old
- 81% on antibiotics before culture
- 96% had an invasive procedure in the last year



Specimen Sources



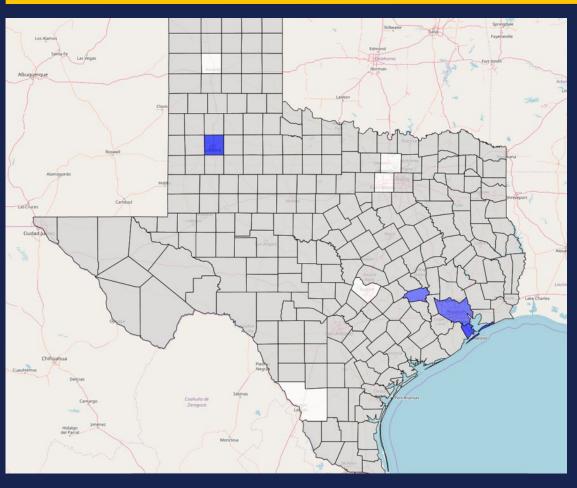


Resistance Patterns



Resistance Pattern	Number of Specimens (%)
Susceptible to colistin only	7 (27%)
Susceptible to colistin and one other class	18 (69%)
Susceptible to colistin and two other classes	1 (4%)

High Prevalence



 Total cases of VIM CRPA in Texas

> Region 1 with 27 Region 6/5S with 5



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During the Epi-Aid

3 Week Visit from the CDC

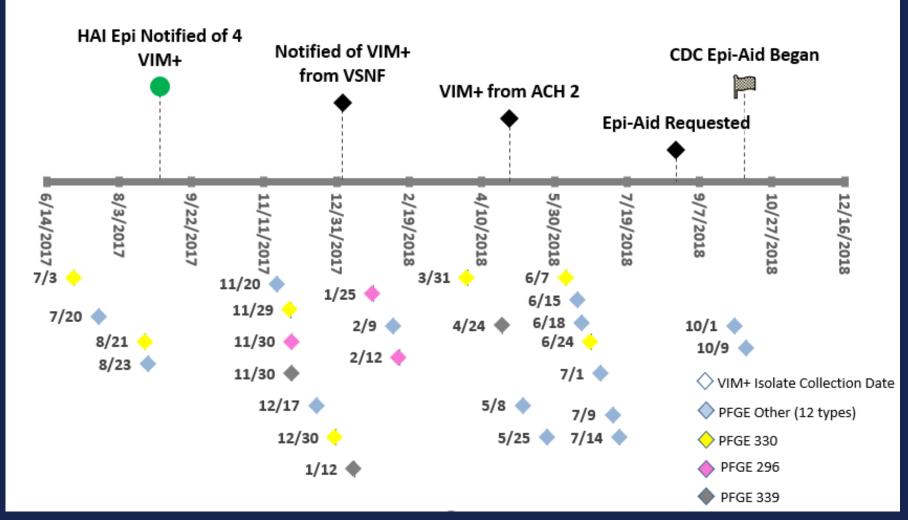
West Texas VIM CRPA Timeline

July 2017 – October 2018



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Epidemic Stages

- 0 No cases reported
- 1 Sporadic occurrence
 - Single cases not epidemiologically related
- 2 Single facility outbreaks
 - ≥2 epi-linked cases in one facility
- 3 Regional spread
 - >1 facility cluster within one referral network
- 4 Interregional spread
 - Multiple clusters occurring within different referral networks
- 5 Endemic
 - Most facilities are repeatedly seeing cases admitted from unrelated sources



The Goal of the Epi-Aid

To Develop & Implement Regional Containment Strategy

- Elements of a successful regional strategy:
 - Led by a central public health authority
 - Participation of most or all of facilities in the region
 - Surveillance cultures/screening
 - Inter-facility communication
- Goal: Decrease spread of MDROs in the regional network of facilities



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The Containment Strategy

Systematic public health response to slow the spread of emerging AR





Preparing to Implement a Regional Prevention Strategy

- 1. Define the region through referral networks
- 2. Increase regional awareness of issue
- 3. Facilitate detection
- 4. Assess baseline regional prevalence
- 5. Assess baseline infection control at high risk facilities



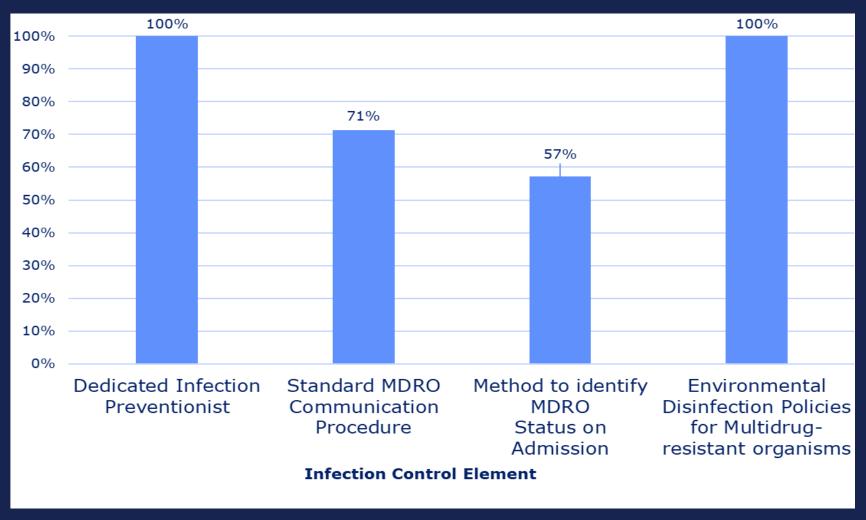
Onsite Assessments

- Conducted at 11 healthcare facilities in West Texas
 - 3 short stay acute care hospitals (ACH)
 - 1 long term acute care hospital (LTACH)
 - 1 inpatient rehabilitation facility (IHR)
 - 4 skilled nursing facilities (SNFs)
 - 1 ventilator skilled nursing facility (vSNF)
 - 1 outpatient wound care center



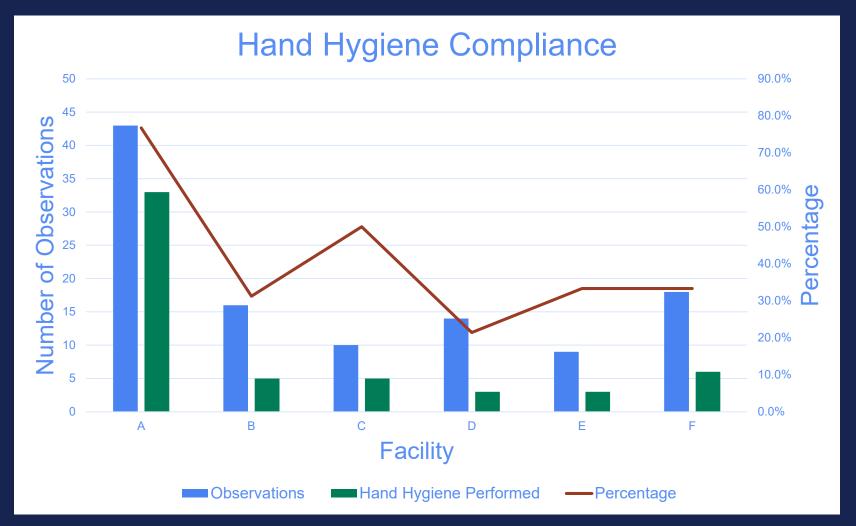
Onsite Assessment Results





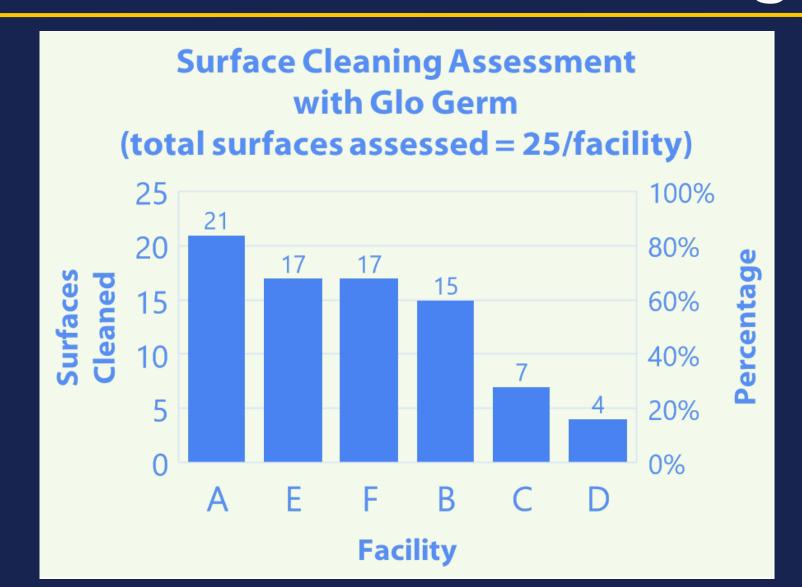
Hand Hygiene





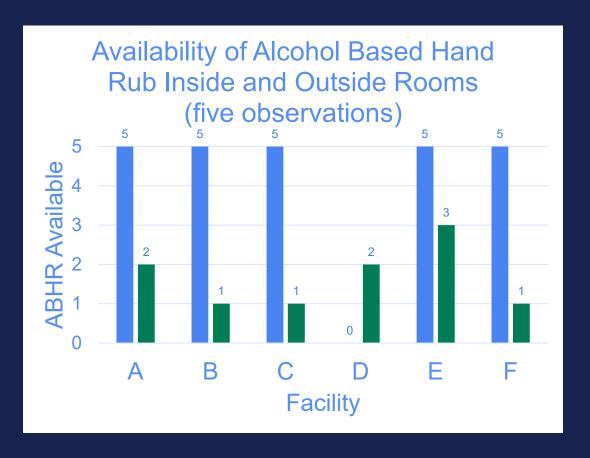
Environmental Cleaning

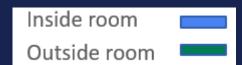




Alcohol Based Hand Rub

- Few easily accessible ABHR
 - Facilitiesstated issueswith fire code
 - Many unclear what local regulation is







Point Prevalence Survey (PPS)

Evaluate the presence of CP-MDROs

- Conducted at 6 different facilities
- 261 colonization swabs collected
 - No additional VIM CRPA positives identified
 - 2 Previously unknown KPC +
- 1 facility screened directly after the epi-aid
 - 68 colonization swabs
 - No additional positives





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PPS Results



Facility	Total	Tested	0/0	Findings
	Attempted			
A - ACH	50	33	66	0
B – ACH	16	7	44	0
C – LTACH	30	23	77	0
D - vSNF	80	39	49	2 KPC *
E - SNF	55	44	80	0
F - SNF	80	47	59	0
G – SNF	79	68	86	0
Total – 7	390	261	67	2 KPC

Surveillance Cultures

Skin/Wound and Sputum Samples

- 1 acute care facility collected clinical specimens along with the PPS rectal swabs
- From two ICUs
- 34 patients (100%) agreed to skin/wound
- 9 patients (26%) agreed to sputum
- Results 1 CRPA, 1 CRAB



Environmental Sampling

Collected due to high P. aeruginosa rates at ACH1

- 45 samples collected
- Sites:
 - sinks
 - drains
 - toilets
 - showers
 - water samples
 - patient areas in the burn unit
 - medical ICU
 - emergency department



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Environmental Sampling Results

- 27% showed *P. aeruginosa* growth (VIM was detected but not isolated in 3)
- 4 KPC+ CRE
- 2 OXA+ CRAB
- 1 VIM+ Pseudomonas monteilli
- 2 First Catch water samples were over the EPA guideline 1 grew P. aeruginosa



Whole Genome Sequencing

Illumina MiSeq Platform

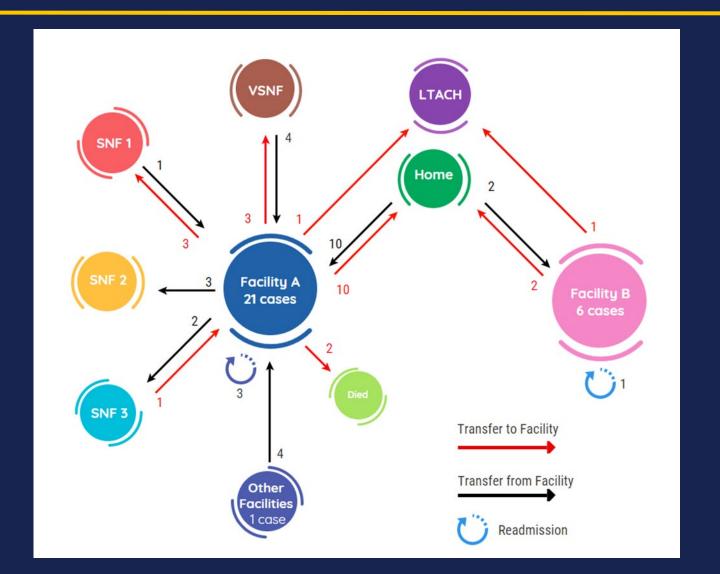
- 26 investigation related isolates tested
- All were sequence type ST308
- All carried blavim-2 gene
- Ranging between 0 88 SNPs but majority were very closely related
- The isolates were compared to 5 from Texas and 19 from other states
 - West Texas samples showed to be unique



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Connections



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Educational Resources

Carbapenem-resistant Acinetobacter baumannii (CRAB)



What is Acinetobacter baumannii?

- A. QRUTREDQU is a bacteria found in soil and is a common contaminant of medical
 equipment, surfaces and skin. It is a cause of healthcare-associated infections. These
 infections are common and can lead to severe infection or death.
- A. QAWINGOU can become resistant to some of our strongest antibiotics called carbapenems. Examples include imigenem or meropenem. This is called Carbapenem-resistant A. QAWINGOU CRAB.
- This organism can carry any of the 5 plasmid-encoded enzymes of primary public health concern that degrade carbapenems: OXA (oxacillinase) (most common in CRAB), KPC, NDM, IMP, and VIM. Enzymes are also known as resistance mechanisms. CRAB with multiple resistance mechanisms can lead to epidemic sureard.
- These organisms can be carried by individuals without causing illness (called asymptomatic carriage). These individuals can spread CRAB to others.
- High-risk patients include those who require medical devices like ventilators, urinary catheters, intravenous catheters, and/or those who are taking long-term courses of antibiotics.
- CRAB can be transmitted from person to person or through shared equipment or healthcare personnel.
- Testing individuals helps prevent spread in a facility and helps their doctor treat them should they become ill.

Prevention in Healthcare Settings

The Texas Department of State Health Services (DSHS) serves as the Antibiotic Resistance Laboratory Network (ARLN) regional lab for the Mountain Region. In order by understand the occurrence of this organism in your community and prevent further spread of the bacteria, epidemiologists may perform infection control consultations or coordinate the collection of patient samples at healthcare facilities where care was received by a disents with CRAB or other resistance mechanisms.





What can you do?

To prevent the spread, healthcare personnel should follow infection control precautions including:

- · Adhering to hand hygiene recommendations
- Carefully cleaning and disinfecting rooms and medical equipment
- Wearing a gown and gloves when performing care of patients/residents that may lead to contamination of healthcare personal hands or clothes (bathing, assisting with toilleting) in addition to using standard precautions.
- Adhering to guidelines for use of personal protective equipment (PPE) in patients who
 require transmission-based precautions
- . When possible, catgrting individuals and dedicating equipment and staff
- · Only prescribing antibiotics when necessary
- Daily cleaning and disinfection of surfaces close to the individual (bed rails, tray table) and other frequently touched surfaces
- Participating in public health initiatives to prevent CRAB from spreading

For more information, please contact the Texas Department of State Health Services (1-888-983-7111) or at the Contact of Disease Control and Reporting at hims bean only probablished SE Austrana, 518 of the

Carbapenem-resistant Enterobacteriaceae (CRE)



What is Enterobacteriaceae?

- A large family of gram-negative rods including Enterobacter, Klebsiella, and E. coil found in normal gut flora; they are opportunistic pathogens.
- They are the most commonly encountered bacteria in clinical microbiology labs. Infections can lead to severe infection or death. They are difficult to treat because they have high levels of resistance to antibiotics.
- Enterobacteriaceae can become resistant to carbapenem antibiotics (such as imipenem or meropenem) – some of our strongest antibiotics. They can carry any of the 5 plasmidencoded enzymes of primary public health concern that degrade carbapenems: KPC (most common in US), NDM, VIM, IMP, and OXX. Enzymes are also known as resistance mechanisms. CRE with multiple resistance mechanisms can lead to epidemic spread.
- High-risk patients include those who require medical devices like ventilators (breathing machines), urinary catheters, intravenous catheters, and/or are taking long courses of
- They can be carried on individuals without causing illness (caused asymptomatic carriage). These individuals can spread CRE to others or become ill from it.
- Can be transmitted person to person or through shared equipment or healthcare personnel.

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- Daily cleaning and disinfection of areas close to the individual (bed rails, tray table) and other frequently touched surfaces
- · Participating in public health initiatives to prevent CRE from spreading

Carbapenem-resistant Pseudomonas aeruginosa (CRPA)



What is Pseudomonas aeruginosa?

- P. aeruginosa is bacteria that thrives in moist places like water and soil. It is a leading cause of healthcare-associated infections (CAUTI, CLABSI, VAE). These infections are common and can lead to severe infection or death.
- P. aeruginosa can become resistant to some of our strongest antibiotics called carbapenems. Examples include imigenem or meropenem. This is called carbapenemresistant P. aeruginosa or CRPA.
- This organism can carry any of the 5 plasmid-encoded enzymes of primary public health concern that degrade carbapenems: VIM (Verona iptograp-encoded metallo-beta-lactamase) (most common in CRPA), KPC, NDM, IMP, and OXA. Enzymes are also known as resistance mechanisms. CRPA with multiple resistance mechanisms can lead to epidemic spread.
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For more information, please contact the Texas Department of State Health Services (1-888-983-7111) or yield the Centers of Disease Control and Prevention at https://www.odc.gov/na/pdfs/cot/CRE-gadance-503.cd

For more information, please contact the Texas Department of State Health Services (1-888-963-7111) or visit the Centers of Disease Central and Provention at https://www.cdc.com/ha/ddfs/trefCRE-cuidence-508.pdf

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CRPA from spreak

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Inter-Facility Infection Prevention Transfer Form



Inter-Facility Infection Prevention Transfer Form

This form must be filled out for transfer to accepting facility with information communicated prior to or with transfer.

Please attach copies of latest culture reports with susceptibilities if available.

Sending Healthcare Facility:

Patient/Resident Last Name	me First Name Date of Birth		Medical Record Number

Name of Sending Facility		Phone Numl	ber	Address	
Sending Facility Contacts	NAME		PHONE		EMAIL
Case Manager/Admin/SW					
Infection Prevention					
Personal Protective Fouriement for Safe Patient Contact and Infection Prevention					

Personal Protective Equipment for Safe Patient Contact and Infection Prevention Please check what is needed:

Standard Precautions

☐ Standard









☐ Surgical (Droplet Mask)

Fit-Teste
N95

□ Cough/uncontrolled

respiratory secretions

Incontinent of urine

Acute diarrhea or incontinent of stool

Does patient currently have an infection, colonization OR a history (in the last 12 months) of a positive culture of a multidrug-resistant organism (MDRO) or other organism of epidemiological significance?	(Last 12 months) Check if YES	Curren. Check if YES
Methicillin-resistant Staphylococcus aureus (MRSA)		
Vancomycin-resistant Enterococcus (VRE)		
Clostridium difficile		
Acinetobacter, multidrug-resistant		
E. coli, Klebsiella, Proteus, etc. w/ Extended Spectrum β-Lactamase (ESBL)		
Carbapenem-resistant Enterobacteriaceae (CRE)		
Carbapanem-resistant Pseudomonas aeruginosa (CRPA)		
Other:		
Cultures pending:		
SVA(PTOMS: Check any that currently apply:		

Draining wounds

fluid/drainage Concerning rash (e.g.

Other uncontained body

1	vesicum)	
De	erson completing form:	

None of the symptoms listed present



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Next Steps

Post 3 week Epi-Aid Visit

Follow up

- 1. Provide written feedback to facilities
- 2. Regional Webinar on Containment Strategies
- 3. Host conference calls check-ups with facilities
- 4. Regional Containment Kickoff Meeting
- 5. Educational webinars to address gaps in facilities infection control



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West Texas gives VIIVI the B.O.O.T.



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- Kickoff meeting for the implementation of the regional containment strategy
- Be prompt (investigate new cases and perform contact screening)
- Obtain isolates (submit clinical isolates to AR Laboratory Network, conduct active surveillance)
- Optimize Infection Prevention
- Transfer using the regional interfacility notification form – every time!





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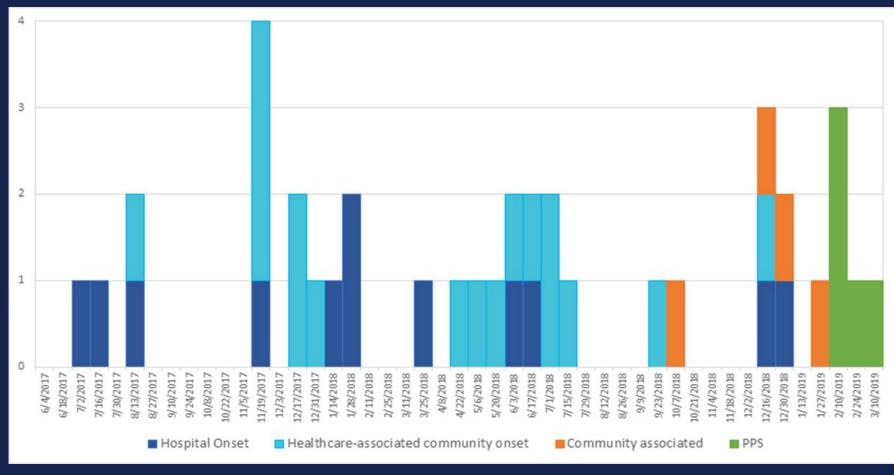
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Regional Prevention Strategy

- 1. Detection
 - Continue to recruit submission to ARLN
- 2. Infection control
 - Return site visits every 6 months
- 3. Inter-facility notification
 - Implement MDRO Transfer Form
- 4. Targeted screening in response to cases
- 5. Active surveillance at high-risk facilities
 - Every other month PPS at facilities involved
 - Begin admission screening at ACH1 and ACH2

Hospital or Community Associated?

Collection dates 6/4/17 - 3/10/19







PPS Summary

- Total swabs collected: 751
- Previously unknown VIM+ CRPA: 5
- Previously unknown KPC+ CRE: 2

Date (2019)	PPS#	# Screened	Positives
28-Jan	1	37	2 initial cases
11-Feb	2	80	3
25-Feb	3	75	1
12-Mar	4	78	1
25-Mar	5	77	0
8-Apr	6	78	0
Total	6	425	7



Looking into the City Water

- Water samples are going to be collected for testing in May
- Testing will focus
 on free residual
 chlorine, total
 chlorine, pH and
 hardness

Facility	Water age range
ACH2	2-5 days
LTACH	2-5 days
SNF	2-5 days
vSNF	2-5 days
SNF	2-5 days
ACH1	5-7 days*
SNF	1-2 days
SSLC	5-7 days
Wound Care Center 1	2-5 days
Wound Care Center 2	2-5 days

Thank you!



- CDC Epi-Aid Team
- Local LRN
- Texas Antibiotic Resistance Lab Network (ARLN)
- Support from State Health Departments
- Participating Healthcare Facilities



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Questions or Comments?

Contact Information
TexasARLN@DSHS.Texas.gov
MDROTexas@DSHS.Texas.gov