

Description of Repeat Carbapenem-resistant Enterobacterales (CRE) Submitted to a State Laboratory from the Same Individuals — December 2018 – March 2022

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Background

- CRE and carbapenemase-producing CRE (CP-CRE) can colonize the GI tract
- CRE and CP-CRE have been persistently detected in long-term care facility residents
- Re-identification of CRE and CP-CRE in other healthcare settings is less well described

Objective

- To describe repeated submission of CRE to a state laboratory, including persistence of detection and progression from colonization to infection

Methods

- Review of CRE isolates submitted to Kentucky Division of Laboratory Services
- Specimens collected December 2018 through March 2022
- Tested for carbapenemase production using modified carbapenem inactivation method (mCIM)
- Positive mCIM followed by Cepheid Xpert Carba-R assay for detection of specific carbapenemases
- Descriptive statistics derived and summarized

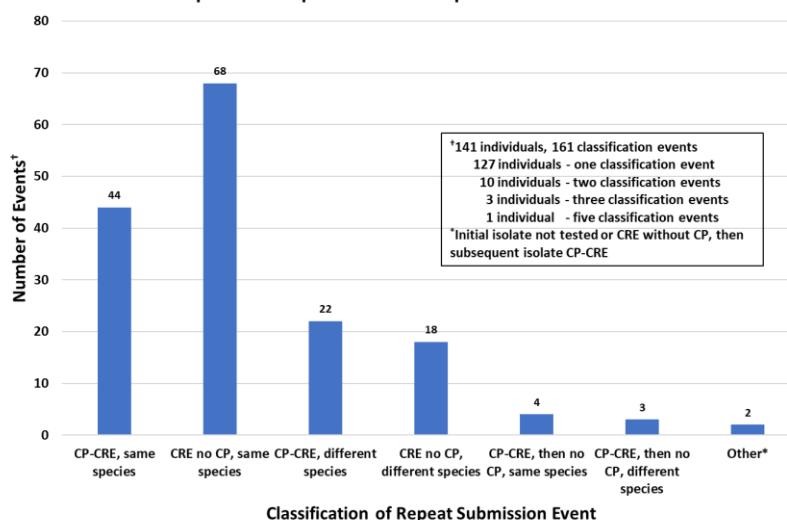
Results

- 1,822 isolates submitted to DLS from 1,498 distinct individuals
- 324 submissions were repeat submissions; from 214 individuals
- 141 individuals were tested >14 days after collection of initial specimen
- Same CP-CRE identified from 42 individuals (median 71 days, IQR 40-151 days, max 857 days)
- Same CRE without CP identified from 66 individuals (median 111.5 days, IQR 55-237 days, max 676 days)
- Urine was most common specimen source among resubmissions (67%)
- Of 47 individuals with a discrepant resubmission result, only four had the same species identified initially as CP-CRE and then subsequently as non-CP-CRE
- Ten individuals had screening and then clinical specimens with the same carbapenemase gene or species

Conclusions

- Persistence can occur with CP-CRE and with CRE without CP
- Detection can occur after more than 500 days
- The in vivo loss of CP by CRE appears to occur infrequently
- Clinicians and infection preventionists should remain vigilant for persistent CRE colonization during clinical and infection control decision making

Classification of Repeat Versus Initial Submission by Comparison of Species and Carbapenemase Production



1,498 individuals; 1,822 isolates

Repeat submissions

- 214 individuals (14.3%), 324 isolates
- 141 individuals, testing >14 days apart

Same CP-CRE (species & mechanism)

- 42 individuals (29.8%)
- Median 71 days apart (IQR 40-151), max 857 days

Same non-CP-CRE species

- 66 individuals (46.8%)
- Median 111.5 days apart (IQR 55-237), max 676 days

CP-CRE, then non-CP-CRE, same species

- KPC *Klebsiella aerogenes* → non-CP CR-*Klebsiella aerogenes*
- VIM *Enterobacter cloacae* → non-CP CR-*Enterobacter cloacae*
- OXA-48 *Klebsiella pneumoniae* → non-CP CR-*Klebsiella pneumoniae* (x2)

CP-CRE, then non-CP-CRE, different species

- KPC *Klebsiella aerogenes* → non-CP CR-*Klebsiella ozaenae*
- KPC *Klebsiella oxytoca* → non-CP CR-*Klebsiella aerogenes*
- KPC *Klebsiella oxytoca* → non-CP CR-*Enterobacter cloacae*

Screening, then clinical specimen (clinical specimen source)

- Non-CP CR-*Klebsiella pneumoniae* – 53 days (urine)
- Non-CP CR-*Enterobacter cloacae* – 60 days (wound)
- KPC *Klebsiella pneumoniae* – 70 days (urine)
- KPC *Escherichia coli* – 73 days (blood)
- KPC *Klebsiella pneumoniae* – 347 days (urine)
- KPC *Enterobacter cloacae* → KPC *Citrobacter koseri* – 21 days (sputum)
- KPC *Citrobacter freundii* → KPC *Enterobacter cloacae* – 33 days (blood)
- KPC *Klebsiella pneumoniae* → KPC *Enterobacter cloacae* – 38 days (blood)
- KPC *Citrobacter freundii* → KPC *Proteus mirabilis* – 139 days (urine)
- KPC & VIM *Enterobacter cloacae* → non-CP CR-*Enterobacter cloacae* – 750 days (blood)

