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Retrospective Review of the utility of MRSA NAAT screening to predict MRSA Infections

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Retrospective review of the utility of Methicillin-Resistant Staphylococcus aureus (MRSA) Nucleic Acid Amplification Test (NAAT) screening to predict MRSA infections



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Background

• Appropriate and timely antibiotic de-escalation is a cornerstone of antimicrobial stewardship. Cultures and sensitivities may not be known for up to 96 hours, exposing patients to broad-spectrum antibiotics.¹

•The MRSA NAAT is a screening tool that is used to detect nasal colonization of MRSA and has been shown to have a strong negative predictive value (NPV) for MRSA pneumonia in multiple studies.²⁻⁵

• In a retrospective study of patients with suspected MRSA pneumonia, utilization of MRSA NAAT was associated with

| • Difference in prevale • Patients may be • Lower prevalen | | | | | | | |
|-----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|--------------------------------------------------------|----------------------|------------------------------------|-------------------------|-------------------------|-------------------------------------------------------|------------------------------------|
| Predictive Value for A | | | | Prevalence | | | |
| •NPV of 98.2% is simi •As expected, specific •Consistent with othe | alence | Prev | Number Positive for MRSA 491 | | Number 4296 | TestMRSA NAAT w/ corresponding clinical culture | |
| <u>Predictive Value for D</u> •The majority of cultu pulmonary, urinary, | L.4% | 11 | | | | | |
| NPV was 94-99% For wound cultures, to lower prevalence Strong predictor that | 9% •NPV was 94-9 •For wound cu to lower preva | | 209 | | 4296 | Clinical culture | |
| For abdominal cultu Small sample size fo strength of NPV | | | | | | | |
| •For CNS, there were | | / | V, and NP | pecificity, PP | ensitivity, S | S | |
| Unable to calculate s Limits strength of NF | Accuracy (95% CI) | NPV, (95% CI) | PPV (95% CI) | Specificity (95% CI) | Sensitivity (95% CI) | Number | Culture Source |
| For unknown culture of these cultures is r | 90.3% (89.3-91.1) | 98.2% (97.7-98.6) | 28.7% (24.8-32.9) | 91.4% (90.5-92.3) | 67.5% (60.7-73.8) | 4296 | All sources |
| WebIntelligence Only able to pull dat Limits sample size | 89% (87.4-90.4) | 99.1% (98.5-99.5) | 13.8% (9.4-19.3) | 89.5% (87.9-90.9) | 68.3% (51.9-81.9) | 1702 | Blood |
| Unable to identify ty Certain culture type | 94.6% (92.8-96) | 98.9% (97.8-99.5) | 55% (43.5-66.2) | 95.2% (93.5-96.6) | 84.6% (71.9-93.1) | 809 | Pulmonary |
| •Limits ability to inter | 90.2% (88-92.2) | 99.9% (99.2-100) | 7.3% (2.7-15.2) | 90.3% (88-92.3) | 85.7% (42.1-99.6) | 789 | Urine |
| •Confirm culture type samples | 88.6% (86-90.8) | 94.1% (91.9-95.8) | 54.1% (43.7-64.2) | 92.8% (90.4-94.7) | 59.6% (48.6-69.8) | 710 | Wound |
| Repeat calculations types Share results with in impact on practice | 86.9% (80.5-91.8) | 94.7% (89.4-97.8) | 38.1% (18.1-61.6) | 90.6% (84.4-94.9) | 53.3% (26.6-78.7) | 153 | Unknown |
| | 94.5% (86.6-98.5) | 95.8% (88.3-99.1) | 0% (0-97.5) | 98.6% (92.3-100) | 0% (0-70.8) | 73 | Intra- abdominal |
| Parente DM, Cunha CB, Myloresistant Staphylococcus aur diagnostic meta-analysis wit 67:1–7. Dangerfield B, Chung A, Web Staphylococcus aureus (MRS Agents Chemother. 2014;58) Smith MN, Brotherton AL, Lu Clinical Utility of Methicillin- MRSA Pneumonia. Ann Phar | 93.6% (82.5-98.7) | 100% (92-100) | 0% (0-70.8) | 93.6% (82.5-98.7) | NA | 47 | Central nervous system (CNS) |
| | 92.3% (64-99.8) | 100 (69.2-100) | 66.7% (9.4-99.2) | 90.9% (58.7-99.8) | 100% (15.8-100) | 13 | Miscellaneous |

Discussion

Results

nce between MRSA NAAT and culture asymptomatic carriers ce helps strengthen NPV

Sources

ar to other retrospective studies ity and NPV are high studies

approximately 2 days less of empiric antibiotic therapy.⁶

•There are fewer retrospective studies on the utility and NPV for other infection types, such as skin and soft tissue infections and intra-abdominal infections. •A small prospective cohort study looking at MRSA NAATs for patients in the emergency department with skin and soft tissue infections found that the MRSA NAAT was a better predictor of MRSA infection than risk factors for MRSA. In this study, the NPV was found to be 72.8%.⁷

•Conversely, a large retrospective cohort study of 200,000 patients found that MRSA NAAT had a NPV greater than 90% for most types of infections, skin and soft tissue infections.⁵

• Discordance is likely due to the differences in prevalence of MRSA in different regions. NPV is driven by prevalence, and MRSA NAAT screening is considered to have a stronger NPV in areas with a low incidence of MRSA.³

•There are less data to understand predictive value of MRSA NAAT screening in the Oregon region given the lack of studies in this area. The purpose of this study is to identify the prevalence of MRSA in the Oregon region as well as the NPV of MRSA NAAT screening, eliciting its utility in predicting MRSA infection and optimizing antimicrobial therapy.

ifferent Culture Sources re sources were blood, followed by and wound NPV may be stronger in this region due infection is not MRSA

es, NPV was still high at 95% cultures of abdominal source limits

no positive MRSA cultures ensitivity

s, NPV was 94.7%, but the exact source ot yet known

Limitations

for previous 15 months

be of wound culture based on report

Objectives

Primary Outcome

• Evaluate the sensitivity, specificity, positive predictive value (PPV), and NPV of MRSA NAAT to predict MRSA infections within a regional health system in Oregon

Secondary Outcome

•Evaluate the sensitivity, specificity, PPV, and NPV of MRSA NAAT to predict MRSA infections for specific culture sources within a regional health system in Oregon

Methods

<u>Study Design</u> •Retrospective study

Inclusion Criteria

- •Age 18 years or older
- •Admitted to a regional health system in Oregon between
- 6/30/2021 and 10/24/2022
- MRSA NAAT collected
- •Corresponding culture collected within 7 days of MRSA NAAT collection

s have small sample sizes pret NPV

Next Steps

- in unknown and miscellaneous
- o finalize NPV for different culture

fectious disease clinicians to determine

References

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Exclusion Criteria

•Rejected or inconclusive MRSA NAAT •Rejected or inconclusive culture

Data Collection

•Retrospective report using WebIntelligence

<u>Data Analysis</u>

•Sensitivity, specificity, positive predictive value (PPV) and NPV were calculated using a script written in R •Exact binomial method was used to calculate 95% confidence intervals

resistant Staphylococcus aureus nasal polymerase chain reaction assay in critically ill patients with nosocomial pneumonia. J Crit Care 2017; 38:168–71. 5. Mergenhagen KA, Starr KE, Wattengel BA, Lesse AJ, Sumon Z, Sellick JA. Determining the Utility of Methicillin-Resistant Staphylococcus aureus Nares Screening in Antimicrobial Stewardship. Clin Infect Dis. 2020;71(5):1142-1148. doi:10.1093/cid/ciz974 6. Baby N, Faust AC, Smith T, Sheperd LA, Knoll L, Goodman EL. Nasal methicillin-resistant Staphylococcus aureus (MRSA) PCR testing reduces the duration of MRSA-targeted therapy in patients with suspected MRSA pneumonia. Antimicrob Agents Chemother 2017; 61:1-8.

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