



# Evaluation of polymerase chain reaction respiratory panel testing in antibiotic decision making

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## INTRODUCTION

- Diagnostic testing is widely used to limit the time from presentation to appropriate treatment.
- Polymerase chain reaction (PCR) respiratory panel is used in practice to determine a bacterial or viral etiology of a patient's respiratory illness.
- The BioFire FilmArray PCR rapidly detects 20 common viral and bacterial respiratory pathogens as an in-house lab which results in ~1 hour.
- The Diatherix PCR detects 31 viral and bacterial respiratory pathogens as a send-out lab which results in ~8 hours.
- The objective of this study was to evaluate the clinical utility in antibiotic (ABX) decision making based on the BioFire FilmArray Respiratory Panel compared to Diatherix PCR.

## METHODS

- A retrospective chart review was conducted using the electronic EHR at a large community hospital from January 1, 2019 to June 30, 2019.
- Patients greater 18 years old were included if they had a BioFire FilmArray PCR ordered and resulted during study period.
- Participants who were pregnant, from outside hospitals, had inappropriate sources or sample amounts, or died during admission were excluded
- Data collected included demographics, diagnosis, length of stay, 30 day readmission, physician specialty, collection source, and PCR results.
- Records were analyzed to determine if an BioFire results were utilized to change therapy
- Examples of an appropriate test utilization include:
  - PCR positive for viral pathogen prompts antibiotic discontinuation
  - PCR positive for atypical bacteria prompts initiation or continuation of a macrolide, fluoroquinolone or tetracycline.
  - Negative PCR prompts appropriate antibiotics continued for non-respiratory panel result reasons
- Results were compared to historical data collected on Diatherix TEM-PCR to evaluate if there were differences in type of test on action taken
- Chi-squared test was used, a *p*-value <0.05 was considered significant

## RESULTS

- A total of 274 BioFire and 195 Diatherix Respiratory Panels were ordered during pre-specified study periods
  - After exclusions were made, a total of 251 BioFire and 118 Diatherix Respiratory Panels were evaluated.
- With the implementation of BioFire, the percentage of appropriate source collection significantly improved (*p*=0.033).
- Collectively, ID (*p*=0.134) and non-ID providers (*p*=0.265) had a similar pattern of PCR utilization.
- After transitioning from Diatherix to BioFire, there was a decrease in the utilization of test results (44.1%, 34.7%, respectively). This result was not statistically significant.
- Of the panels evaluated (*n*=369), PCR results were not utilized in 230 (62.3%) cases.

## DISCUSSION

- We anticipated BioFire to improve antibiotic decision making and clinical outcomes as compared to Diatherix due to the oversight of in-house testing.
- The implementation of BioFire significantly improved appropriate source collection for PCR respiratory testing
- Based on the results of this study, the transition to BioFire testing did not improve utilization
- The overall utilization of respiratory PCR testing to drive antibiotic decision making at our facility was 37.7%. This low percentage of action based on PCR results was an unexpected finding of this evaluation.
- An evaluation of best practices with regard to PCR respiratory testing is needed to achieve better utilization among providers
- Additionally, a cost effectiveness analysis is warranted based on the results of this evaluation.

# Respiratory panel PCR results were used to guide antibiotic decision-making in one-third of patients.



## TABLE OF RESULTS

	Diatherix	BioFire	<i>p</i> -value
<b>Total Number Ordered</b>	n=195	n=274	
<b>Physician Specialty</b>			
Infectious Disease	40 (20.5%)	84 (30.7%)	
Nurse Practitioner	20 (10.3%)	72 (26.3%)	
Internal Medicine	7 (3.6%)	75 (27.4%)	
Hospitalist	61 (31.3%)	20 (7.3%)	
Resident Physician	35 (17.9%)	5 (1.6%)	
Pulmonology/Critical Care	17 (8.7%)	16 (5.8%)	
Other	15 (7.7%)	2 (0.7%)	
<b>Appropriate Source</b>	180 (92.3%)	265 (96.7%)	0.033
<b>Total Number Excluded</b>	77	23	
Outside hospital	39 (50.6%)	12 (52.2%)	
Inappropriate source/sample	15 (19.5%)	10 (43.5%)	
Death	23 (29.9%)	1 (4.3%)	
<b>Total Number Evaluated</b>	118	251	
Test results utilized	52 (44.1%)	87 (34.7%)	0.082
Test results not utilized	66 (55.9%)	164 (65.3%)	
<b>Number Evaluated by Specialty</b>			
<b>Infectious Disease</b>	27	78	
Test results utilized	13 (48.1%)	25 (32.1%)	0.134
Test results not utilized	14 (51.9%)	164 (65.3%)	
<b>Non-infectious disease</b>	91	173	
Test results utilized	39 (42.9%)	62 (35.8%)	0.265
Test results not utilized	52 (57.1%)	111 (64.2%)	
<b>Sig. Diff. by Test &amp; Provider Type</b>	<i>p</i> =0.627	<i>p</i> =0.559	

## DISCLOSURES

Authors of this presentation have nothing to disclose concerning possible financial or personal relationships with commercial entities that may have a direct or indirect interest in the subject matter of this presentation.

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