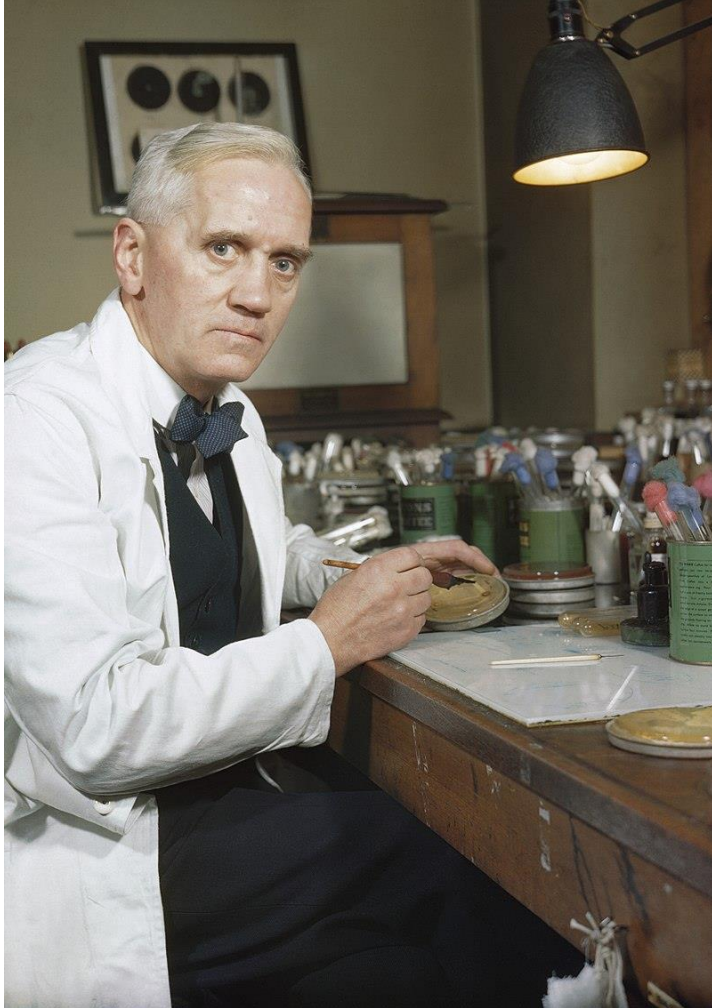


L'apport des solutions DATA & IT dans la prise en charge des patients : l'approche de bioMérieux

Stéphanie Le Page – PharmD, PhD
Global Medical Affairs

ANTIMICROBIAL RESISTANCE (AMR) IS A GLOBAL PUBLIC HEALTH CONCERN THAT WAS PREDICTED ...



“The time may come when penicillin can be bought by anyone in the shops. Then **there is the danger** that the ignorant man may easily under dose himself and, by exposing his microbes to non-lethal quantities of the drug, educate them to resist penicillin.”

*Sir Alexander Fleming
Nobel lecture, 1945*

THE CHALLENGES OF SEPSIS AND AMR

TODAY

1,270,000 deaths per year
as a result of AMR¹
1 death every 25 seconds

2050
10,000,000
global deaths annually¹

Illustration:

Sepsis, where accurate and rapid diagnostics are key

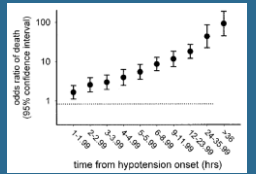
1
HOUR

*delay in time to effective
antibiotics*

=

7.6%

*Reduced
survival rate²*



48
HOUR

*delay in time to effective
antibiotics*

=

9.1

*Increase Odd Ratio
of Death³*

Each
DAY

*delay in time to effective
antibiotics*

=

*Increased 30 -Day
Mortality⁴*

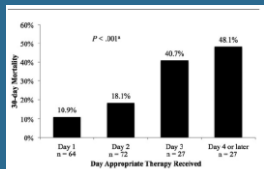


Figure 2. Relationship between day appropriate therapy was received and 30-day mortality. *P-value for χ^2 and χ^2 test for linear trend.

ANTIMICROBIAL RESISTANCE (AMR) IS A GLOBAL PUBLIC HEALTH CONCERN THAT WILL INCREASE IN MAGNITUDE IN THE FUTURE IF NOT BETTER HANDLED COLLECTIVELY

Increased length of stay



Increased cost of treatment



Precaution measures
are expensive



Constant monitoring



OUR PLEDGE TO CURBING ANTIMICROBIAL RESISTANCE

Providing **high medical value diagnostic solutions** to guide effective and rational use of antibiotics



Developing **innovative antibiotic susceptibility testing** and automated detection of resistance

Providing **actionable insights**, based on consolidated data to support medical decisions



Producing **education material and raising awareness** on the value of diagnosis and rational use of antibiotics.



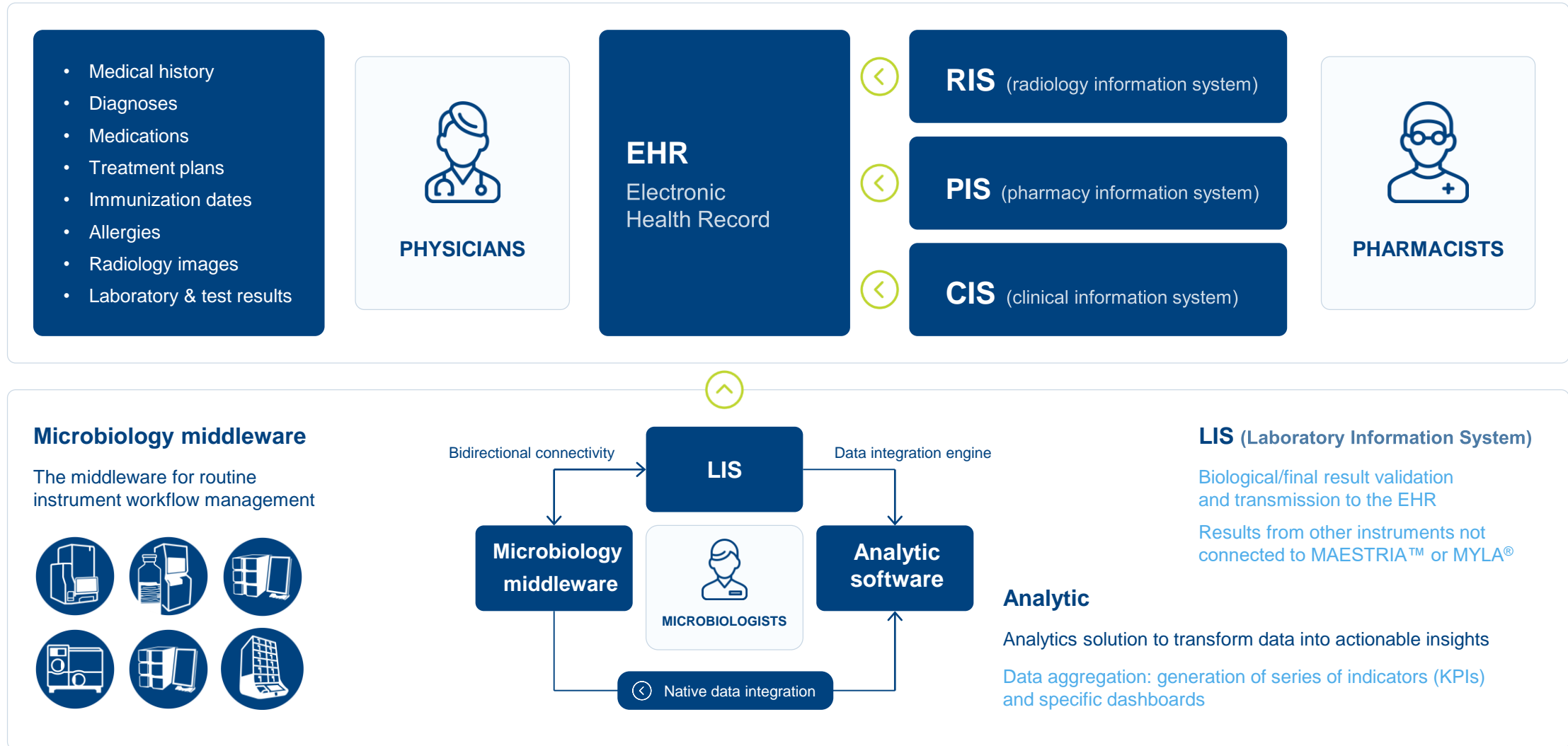
80%

OF OUR R&D BUDGET
IS DEDICATED TO
THE FIGHT AGAINST AMR

[DATA ENVIRONMENT IN HOSPITAL]

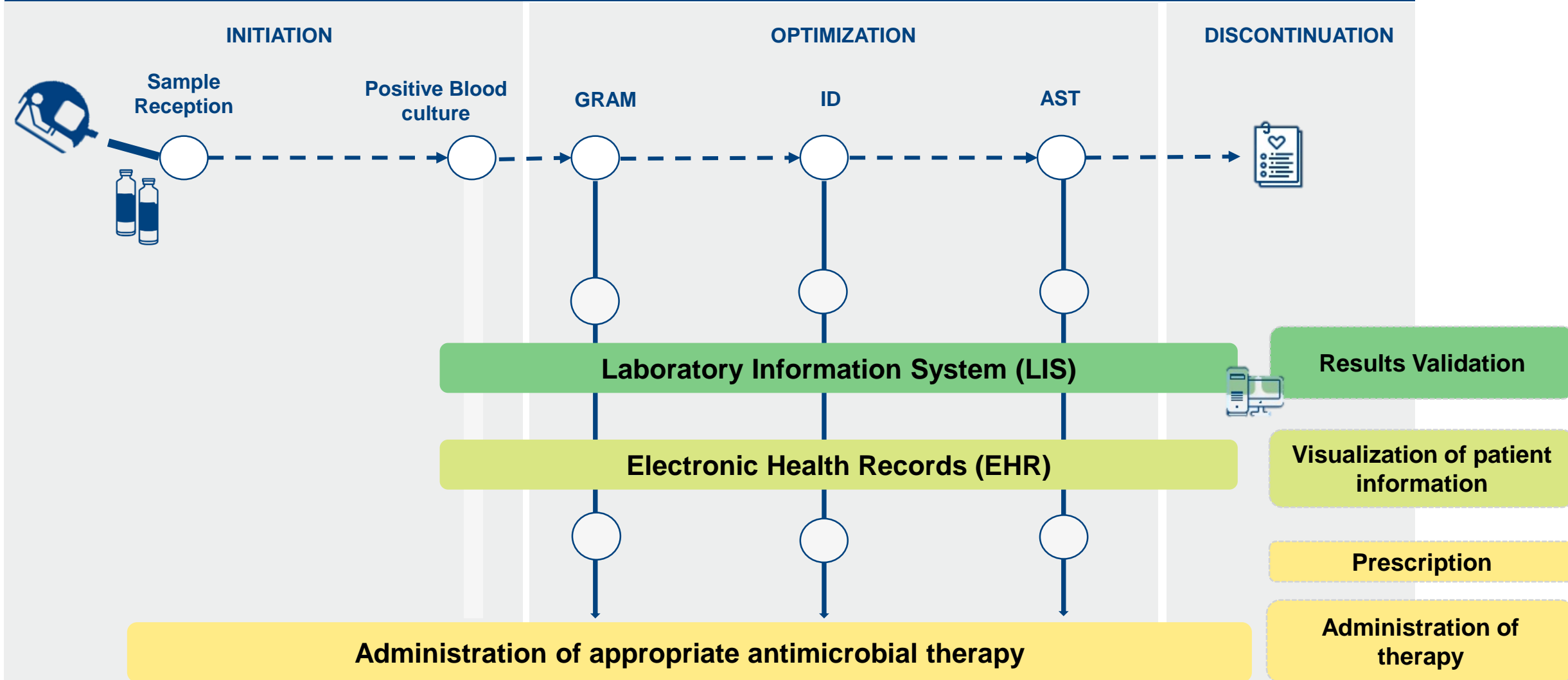


DATA ENVIRONMENT IN HOSPITALS

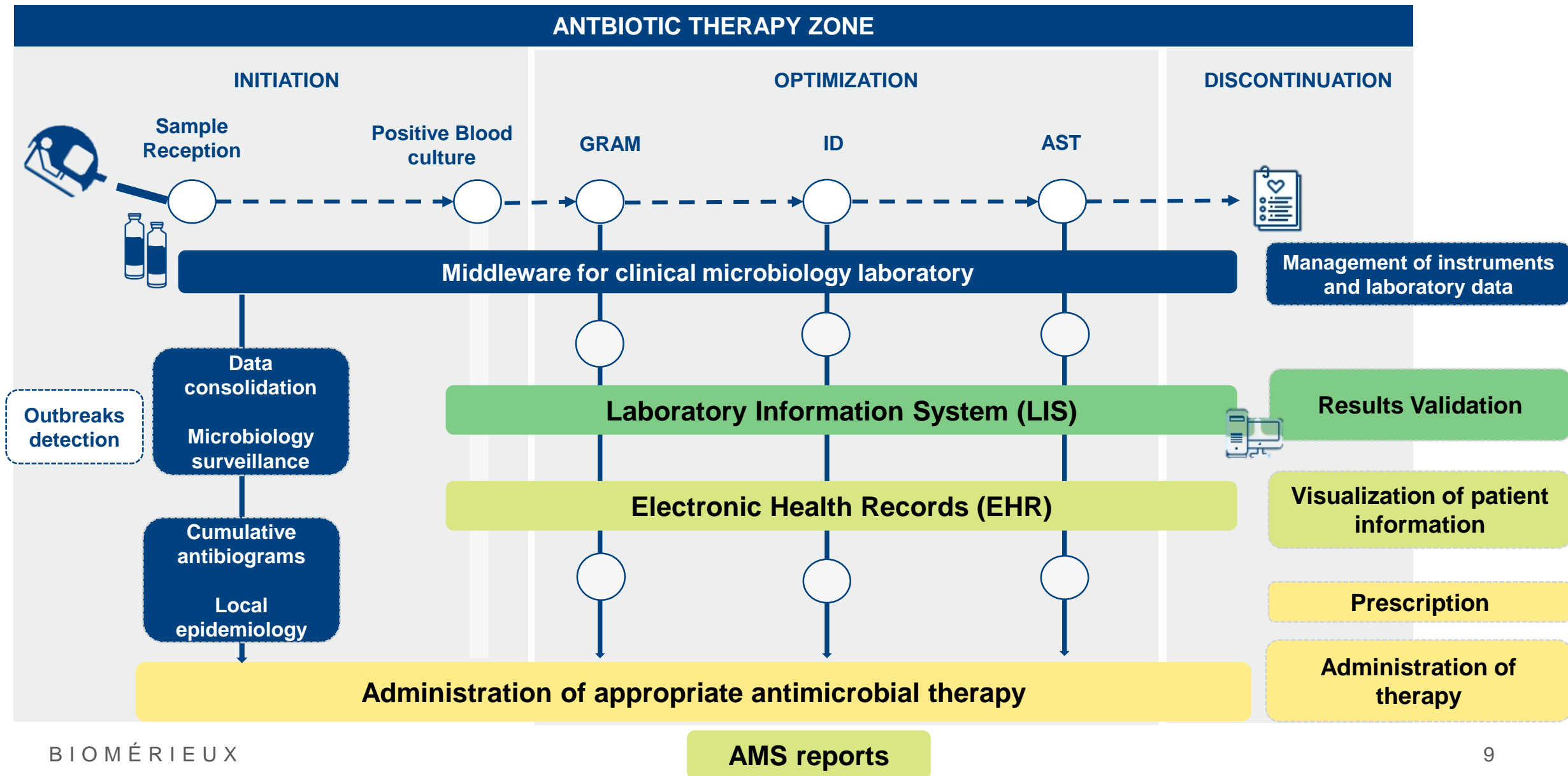


LABORATORY WORKFLOW: generation of data

ANTIBIOTIC THERAPY ZONE



LABORATORY WORKFLOW: generation of data



SEVERAL PRINCIPLES SHOULD BE APPLIED TO IMPROVE PATIENT OUTCOMES

IMPROVE LABORATORY MANAGEMENT EFFICIENCY

DECREASE THE TIME TO AN ACTIONABLE DIAGNOSTIC RESULT

**IMPROVE THE ACCURACY OF INITIAL EMPIRIC ANTIBIOTIC THERAPY
SHORTEN THE TIME TO ADJUSTMENT WHEN EMPIRIC THERAPY IS INCORRECT**

ANTICIPATE EMERGING THREATS THROUGH EFFECTIVE SURVEILLANCE

IMPROVING LABORATORY MANAGEMENT EFFICIENCY

FROM THE LAB

Microbiology middleware

ID patient / Nom du pat...ID échantillon



TABLEAU DE BORD DES ACTIVITÉS

MAESTRIA v5

Routine

Données non résolues

Suivi des hémocultures

LISTE DES PRÉLÈVEMENTS

LISTE DES FLACONS

 Flacon positif (5)
myVirtuO-R3

BLOOD CULTURE

Filter by Patient ID5

Bottles in progress10

Warning Volume0

Flacons orphelins0

Flacons anonymes0

Reloaded Bottles2

Negative bottles to unload6

Flacons en attente de chargement3

BC CONTAMINATION

Presumptive Contaminations0

Confirmed Contaminations1

SEPSIS CODE

Patient flag ICU40

Clinical study case27

MDRO

MDRO to Review6

Confirmed MDRO260

ESBL E. coli - Last 7 Days1

CRE E.coli - Last 7 Days1

ID MASS SPEC

BLOOD CS LISTING69

Demo Specimen7

ID/AST VITEK 2

Running AST0

Urine Specimens9

Need Help4

FROM THE LAB

Performance indicators generation



Lab Manager Dashboard



Institution

stephanie.lepage



Positivity rate

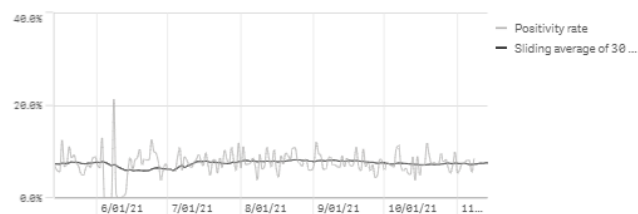


7.7%

Last 30 days

7.2%

vs. 30 days before



May 21

Nov 21



Contamination rate

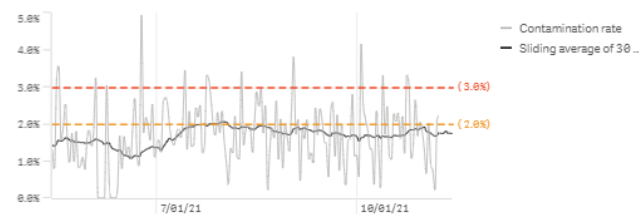


1.8%

Last 30 days

1.6%

vs. 30 days before



May 21

Nov 21



Bottle volume (mL)

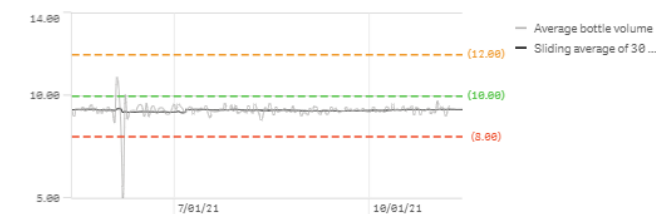


9.28

Last 30 days

9.27

vs. 30 days before



May 21

Nov 21



Mean load time (hh:mm)



3:00

Last 30 days

3:04

vs. 30 days before



May 21

Nov 21



BC instruments load rate

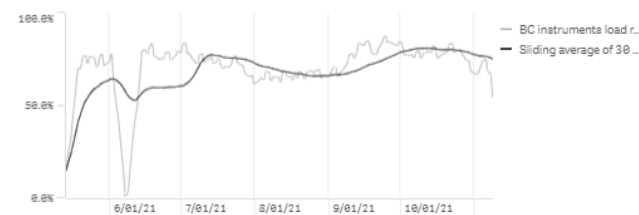


74.9%

Last 30 days

81.1%

vs. 30 days before



May 21

Nov 21



Number of identifications

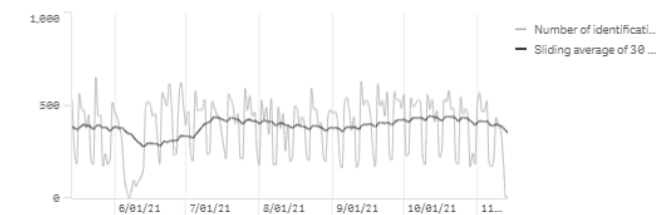


10,975

Last 30 days

13,133

vs. 30 days before



May 21

Nov 21

DECREASING THE TIME TO AN ACTIONABLE DIAGNOSTIC RESULT

THE CURRENT TYPICAL PATIENT DIAGNOSIS JOURNEY CAN BE REALLY LONG – THERE ARE OPPORTUNITIES TO DECREASE TIME TO RESULTS



*If customers use their own short incubation protocol for VITEK MS

**Gram Negative bacteria only

***Theoretical based on instrument run times

THE USE OF MULTIPLEX RAPID DIAGNOSTIC TESTS COMBINED WITH AMS CAN DECREASE TTR & IS ASSOCIATED WITH A SIGNIFICANT DECREASE IN MORTALITY FOR BLOOD STREAM INFECTIONS

Clinical Infectious Diseases

MAJOR ARTICLE



The Effect of Molecular Rapid Diagnostic Testing on Clinical Outcomes in Bloodstream Infections: A Systematic Review and Meta-analysis

Tristan T. Timbrook,^{1,4} Jacob B. Morton,^{1,4} Kevin W. McConeghy,² Aisling R. Caffrey,^{1,2,4} Eleftherios Mylonakis,³ and Kerry L. LaPlante^{1,2,4}

¹Rhode Island Infectious Diseases Research Program, Providence Veterans Affairs Medical Center, ²Center of Innovation in Long Term Services and Supports, Providence Veterans Affairs Medical Center, ³Infectious Diseases Division, Warren Alpert Medical School of Brown University, Providence, and ⁴College of Pharmacy, University of Rhode Island, Kingston

Background. Previous reports on molecular rapid diagnostic testing (mRDT) do not consistently demonstrate improved clinical outcomes in bloodstream infections (BSIs). This meta-analysis seeks to evaluate the impact of mRDT in improving clinical outcomes in BSIs.

Methods. We searched PubMed, CINAHL, Web of Science, and EMBASE through May 2016 for BSI studies comparing clinical outcomes between mRDT and conventional microbiology methods.

Results. Thirty-one studies were included with 5920 patients. The mortality risk was significantly lower with mRDT than with conventional microbiology methods (odds ratio [OR], 0.66; 95% confidence interval [CI], .54–.80), yielding a number needed to treat of 20. The mortality risk was slightly lower with mRDT in studies with antimicrobial stewardship programs (ASPs) (OR, 0.64; 95% CI, .51–.79), and non-ASP studies failed to demonstrate a significant decrease in mortality risk (0.72; .46–1.12). Significant decreases in mortality risk were observed with both gram-positive (OR, 0.73; 95% CI, .55–.97) and gram-negative organisms (0.51; .33–.78) but not yeast (0.90; .49–1.67). Time to effective therapy decreased by a weighted mean difference of –5.03 hours (95% CI, –8.60 to –1.45 hours), and length of stay decreased by –2.48 days (–3.90 to –1.06 days).

Conclusions. For BSIs, mRDT was associated with significant decreases in mortality risk in the presence of a ASP, but not in its absence. mRDT also decreased the time to effective therapy and the length of stay. mRDT should be considered as part of the standard of care in patients with BSIs.

Keywords. rapid diagnostic tests; bloodstream infections; meta-analysis; antimicrobial stewardship.

Improving Clinical Outcomes in Bloodstream Infections

Up to 1/3 reduction in mortality with molecular rapid diagnostics testing (mRDT) and antimicrobial stewardship programs (ASPs)¹

mRDT vs.
conventional microbiology



Mortality Odds Ratio, 0.72
95% CI, 0.46-1.12

mRDT with ASP vs.
conventional microbiology



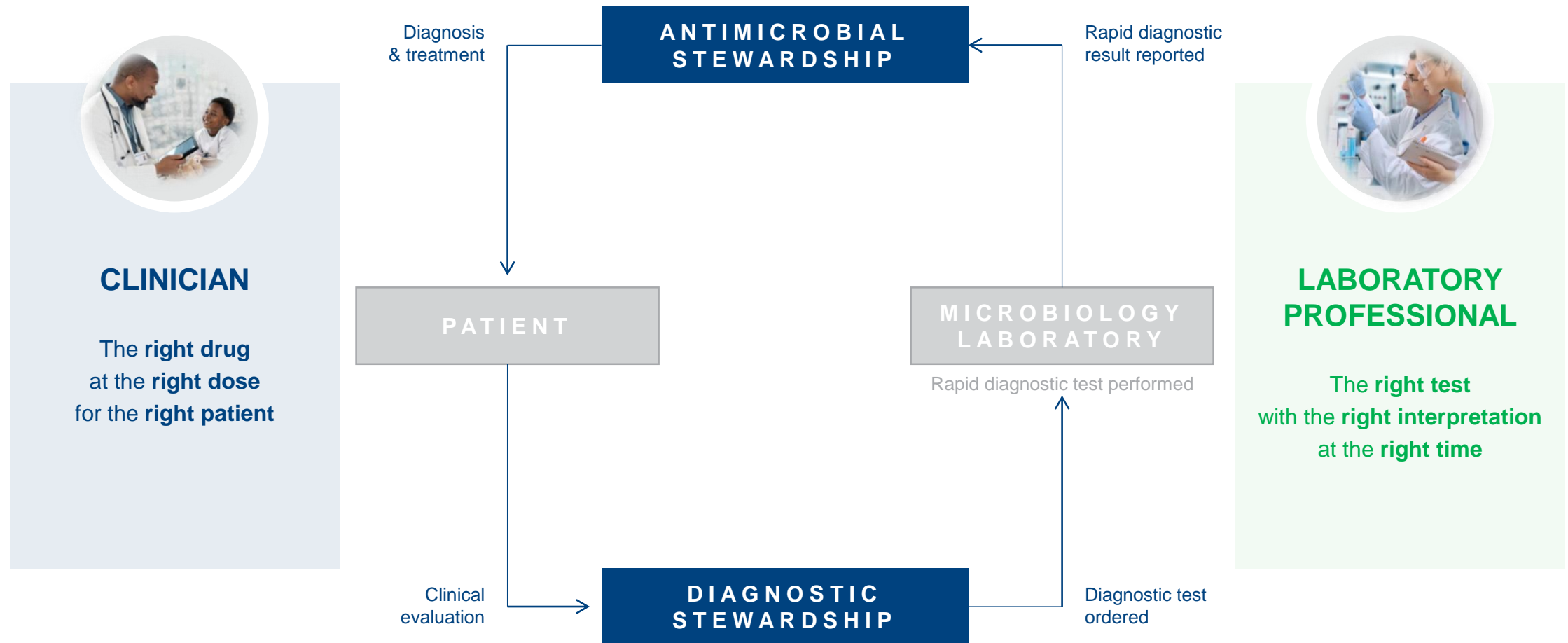
Mortality Odds Ratio, 0.64
95% CI, 0.51–.079

5920 patients
31 studies

BIOMÉRIEUX

“In BSI patients, mRDT should be considered as part of the standard care”

MICROBIOLOGY RESULTS ARE KEY TO MANAGE PATIENTS WITH INFECTION



ADVANCED ANALYTICS

TRANSFORM DIAGNOSTICS DATA INTO ACTIONABLE INSIGHTS

SUPPORT INFECTIOUS DISEASE MANAGEMENT



► Guiding decisions with Antibigram

► Identify sample quality issues
& ineffective collection practices



OPTIMIZE QUALITY RESULTS

DRIVE LAB PERFORMANCE



► Optimize microbiology lab workflow
and time-to-results



HOW DATA & IT CAN IMPROVE BSI



HOW TO IMPROVE BLOODSTREAM INFECTION DIAGNOSTICS?

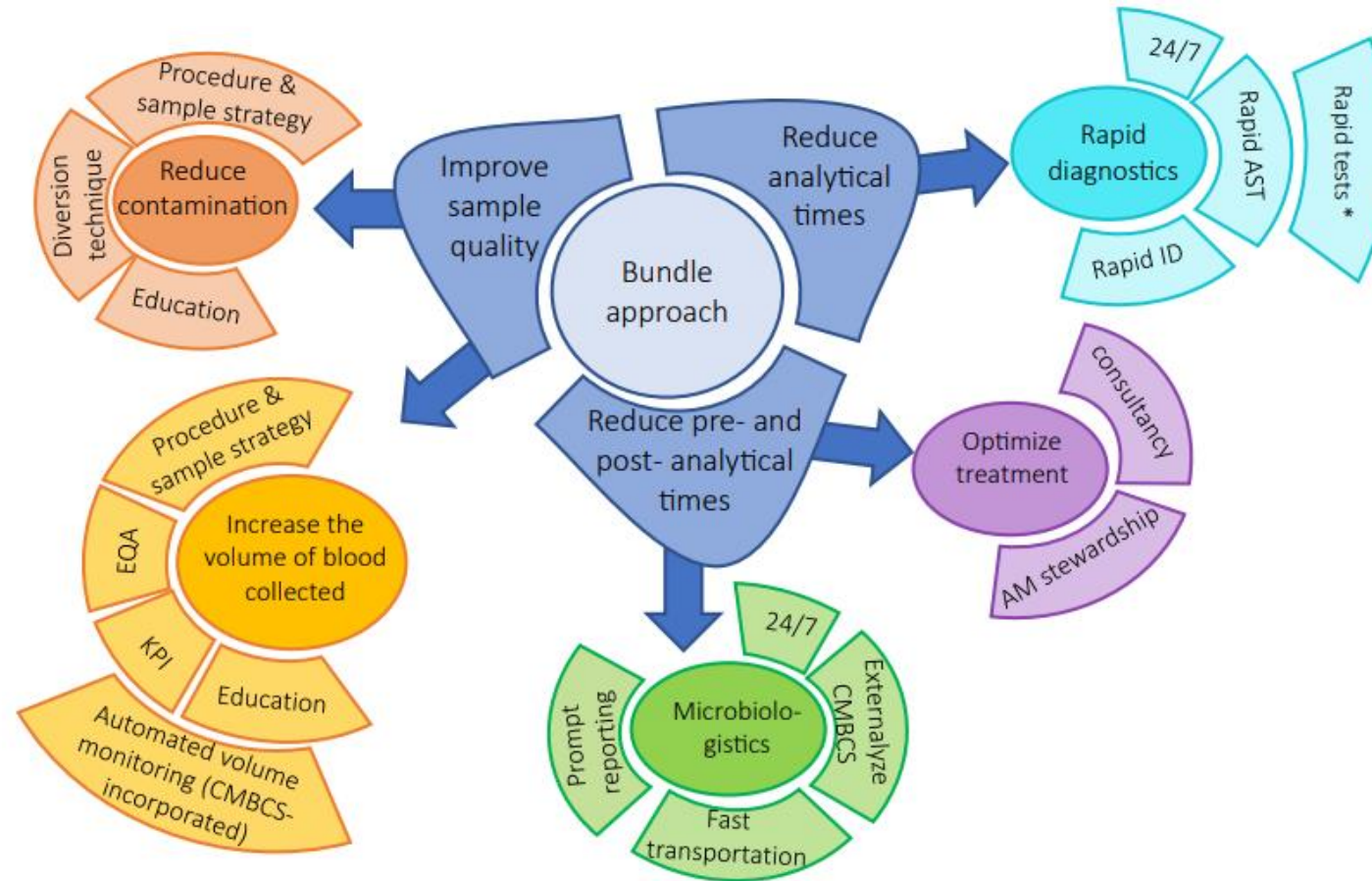


Fig. 1. Summary of all the actions to improve the bloodstream infection pathogen diagnostics. Types of actions belong to three complementary axes and actions aim to manage sample quality, times before and after analysis and analytical times. Each action per se is associated with a limited improvement but combination of several actions significantly improves diagnosis. Improvement is maximum when programme include actions on sampling quality, rapid diagnostics and logistics. KPI, key performance indicator; EQA, external quality assessment; CMBCS, continuous-monitoring blood culture system; AM stewardship, antimicrobial stewardship. *Rapid tests (e.g. *mecA* detection) may be needed in area of high level of resistance.

BLOOD CULTURE KPI



BLOOD SAMPLING

AMS KPI

Target:

**Contamination rate
< 3% (optimally < 1%)**

Contamination Rates



BOTTLE-FILLING

AMS KPI

Target:

Quantity 8-10 mL
Increasing the blood volume increases the chance of recovering microorganisms

Bottle Volume



SPECIMEN TRANSPORTATION

AMS KPI

Target:

< 4 hours

Mean Load Time



SPECIES IDENTIFICATION

AMS KPI

Target:

**Same day of Blood
Culture (BC) positivity**

TTR, Fast ID



AST

AMS KPI

Target:

**Preliminary or final
available on the same
day of next day**

TTR

CN

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MA

MAESTRIA™

BLOOD CULTURE KPI: CONTAMINATION RATE



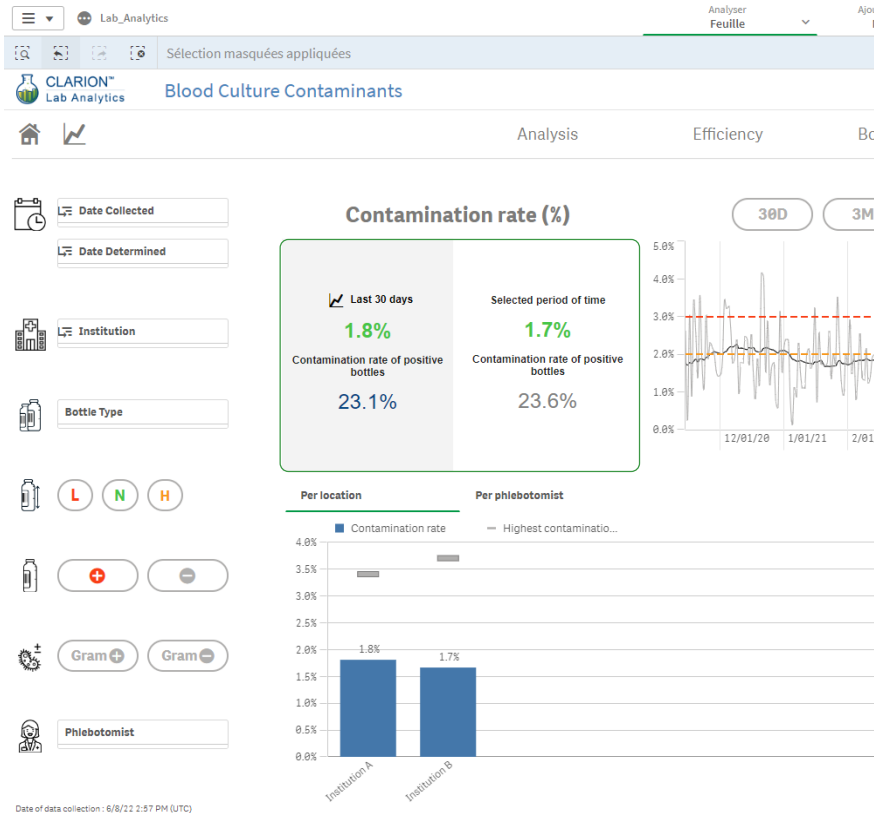
BLOOD SAMPLING

AMS KPI

Target:

Contamination rate
< 3% (optimally < 1%)

Contamination Rates



Impact on AMS:

- **Unnecessary treatment increase antibiotic exposure¹**
 - Unneeded intravenous antibiotics
 - Prolonged therapy
- **Potential adverse event including:**
 - Allergic reaction
 - Antibiotic resistance emergence
 - Disruption of the host microbiome
- **Cost associated:**
 - For the lab: increase unnecessary workflow
 - For the hospital: potential increase in length of hospital stay or follow up testing

Actions of improvement:

1. **Identify areas of high contamination** (e.g. specific ward)
2. **Targeted education** for specific institutions across network, specific units, and even specific individuals for proper blood culture collection technique
3. **Data can be utilized to improve resource utilization**



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BLOOD CULTURE KPI:

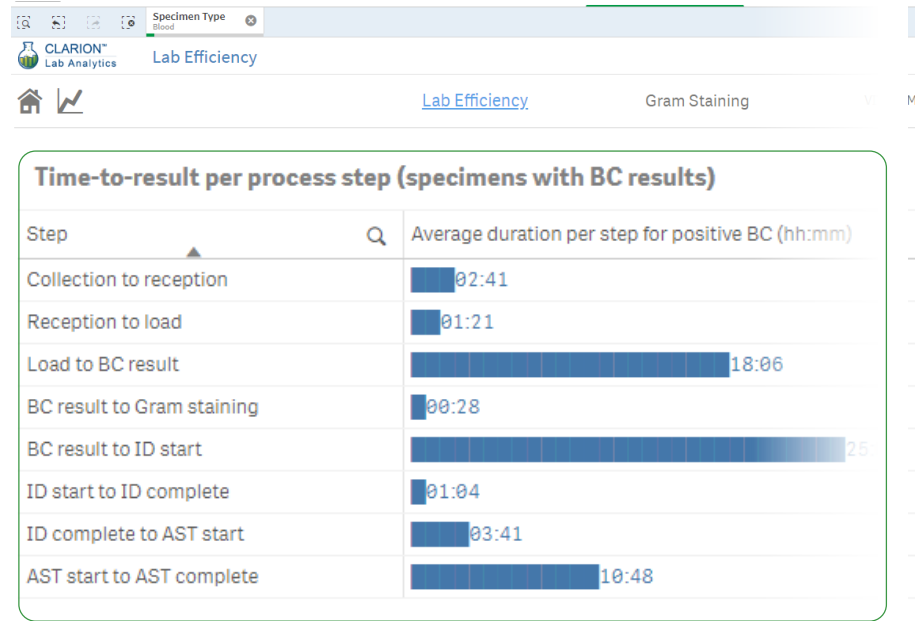


SPECIES IDENTIFICATION

AMS KPI

Target:
Same day of Blood Culture (BC) positivity

TTR, Fast ID



Impact on AMS:

- Better diagnose Bloodstream infection
- Appropriateness of empiric antibiotic therapy
- Impact on time to appropriate therapy
- Hospital cost¹
- Mortality and morbidity

Actions of improvement:

Rapid identification

- Rapid tests: syndromic panel
- FAST AST: VITEK® REVEAL
- 24/7: workflow modification
- Availability of the results: 24/7



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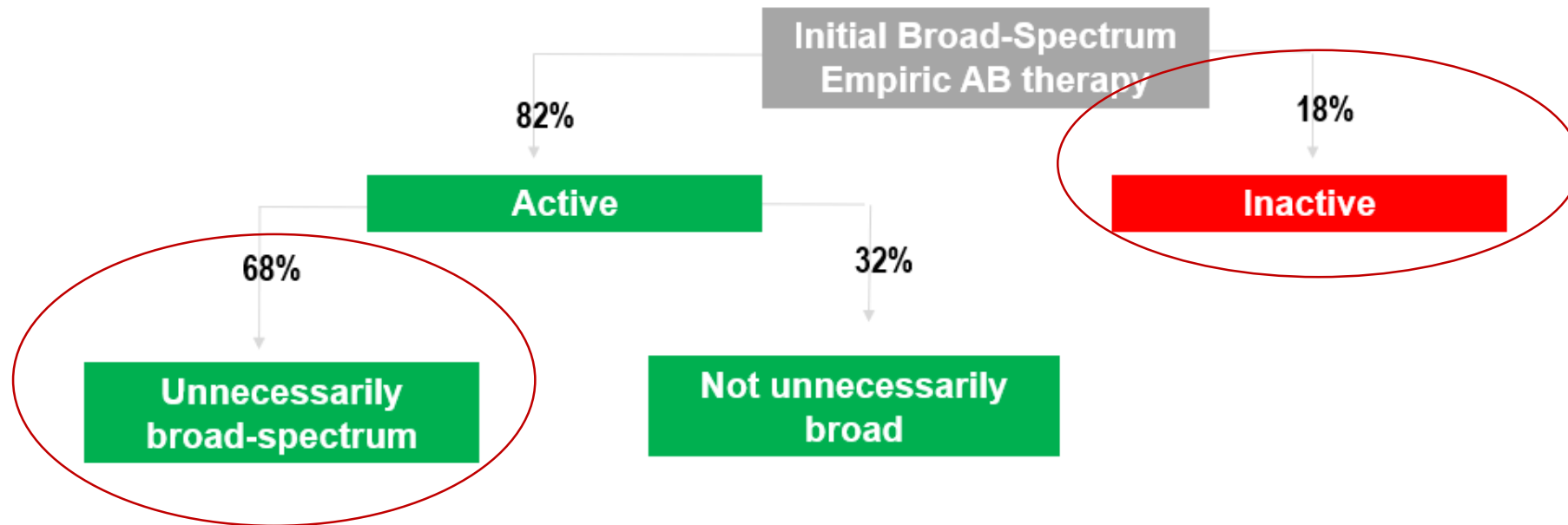


MAESTRIA™

IMPROVING THE ACCURACY OF INITIAL EMPIRIC ANTIBIOTIC THERAPY

EMPIRIC TREATMENTS CAN BE FREQUENTLY WRONG OR UNNECESSARILY BROAD

- 17 430 adults with culture positive sepsis admitted to 104 US hospitals
- Resistant G+ organisms isolated in 13.6% of patients & Resistant G- organisms in 13.2%



Rhee C et al. Prevalence of Antibiotic-Resistant Pathogens in Culture-Proven Sepsis and Outcomes Associated With Inadequate and Broad-Spectrum Empiric Antibiotic Use. *JAMA Network Open*. 2020;3(4):e202899.

WHO IS SUPPORTING THE USE OF SURVEILLANCE TOOLS TO IMPROVE DIAGNOSTIC STEWARDSHIP AND INFORM TREATMENT GUIDELINES



Diagnostic stewardship

A guide to implementation in
antimicrobial resistance
surveillance sites



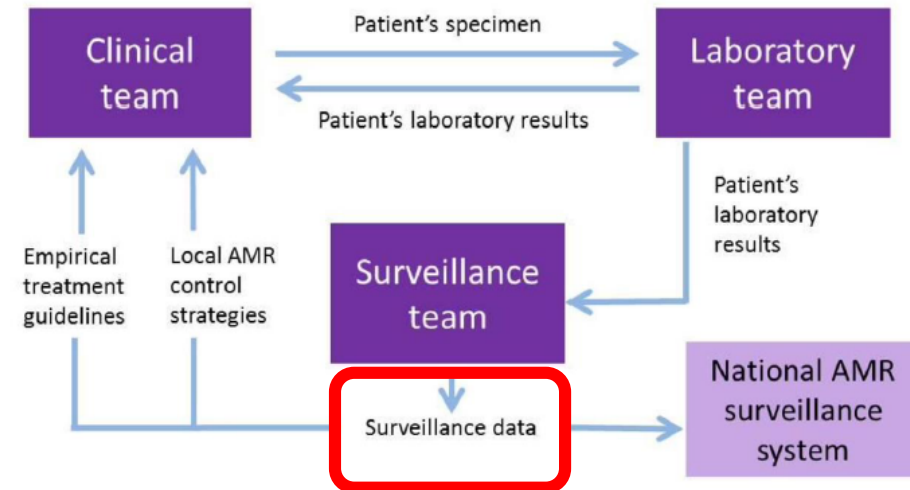
“coordinated guidance and interventions to improve appropriate use of microbiological diagnostics to guide therapeutic decisions. It should promote appropriate, timely diagnostic testing, including specimen collection, and pathogen identification and accurate, timely reporting of results to guide patient treatment.”

The main objective of microbiological diagnostic stewardship is to deliver:

- appropriate, timely diagnostic testing, including specimen collection, and pathogen identification and accurate, timely reporting of results
- patient management guided by timely microbiological data to deliver safer and more effective and efficient patient care

accurate and representative AMR surveillance data to inform treatment guidelines, and AMR control strategies.

Figure 1: Relationship between individual care and surveillance data



Good communication between the different professionals involved, namely the laboratory, clinical, and surveillance teams, plays a critical role in successful diagnostic stewardship.

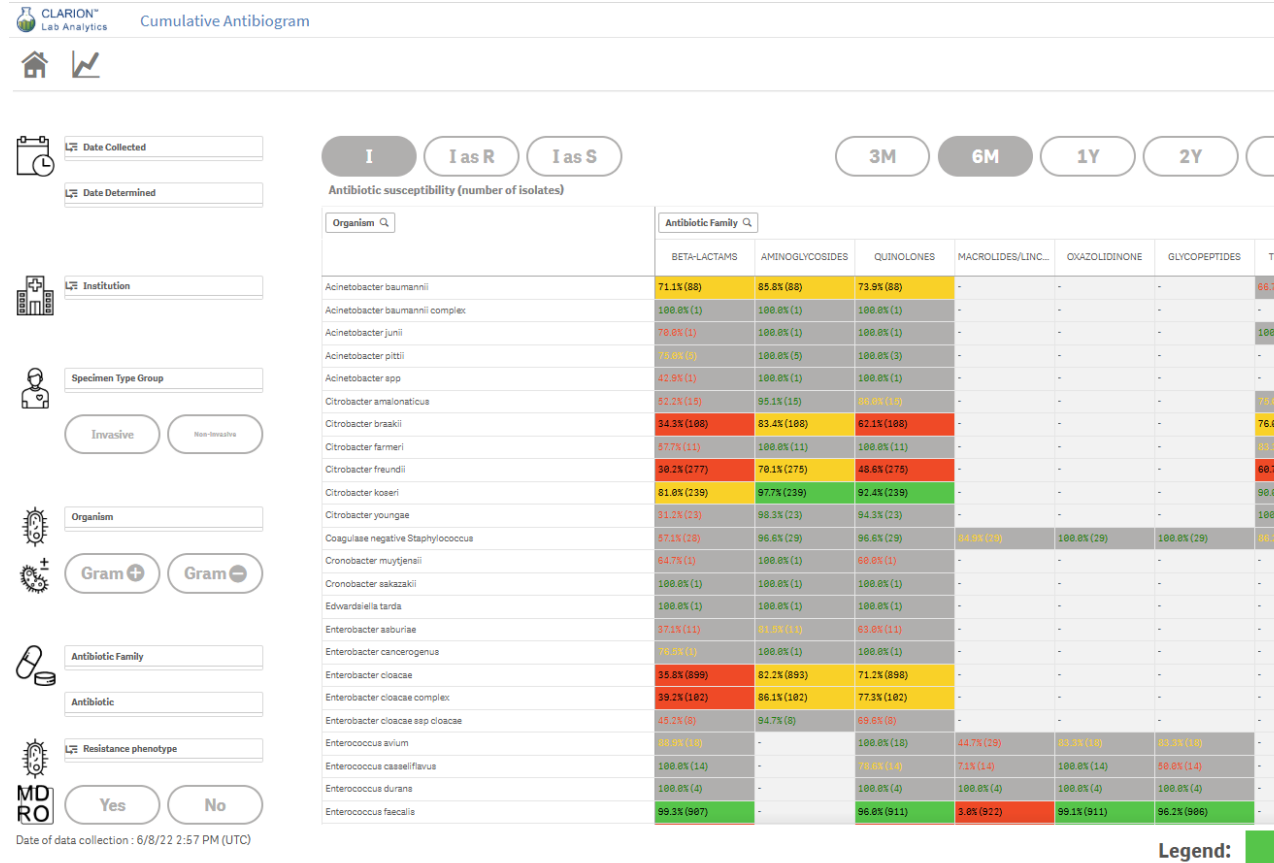
ROUTINE ANTIBIOGRAM



ROUTINE ANTIBIOGRAM

AMS KPI

- Deliver and analyse annually the routine antibiogram (min)
- Reflects % of first isolates (per patient) of a given species that is susceptible to each of the antimicrobial agents routinely tested (deduplication)
- Aggregate of all patients from all locations for the prior calendar year



Impact on AMS:

- Guiding clinicians in the selection of empirical antimicrobial therapy for initial infections before definitive susceptibility results become available or when definitive susceptibility results are not available^{1,2}
- ASP will update local empiric therapy guidelines (annually): antimicrobial formulary
- Monitoring changes resistance over time

Actions of improvement:

- Increase awareness of antibiotic resistance
- Education for the interpretation of real-time antibiograms



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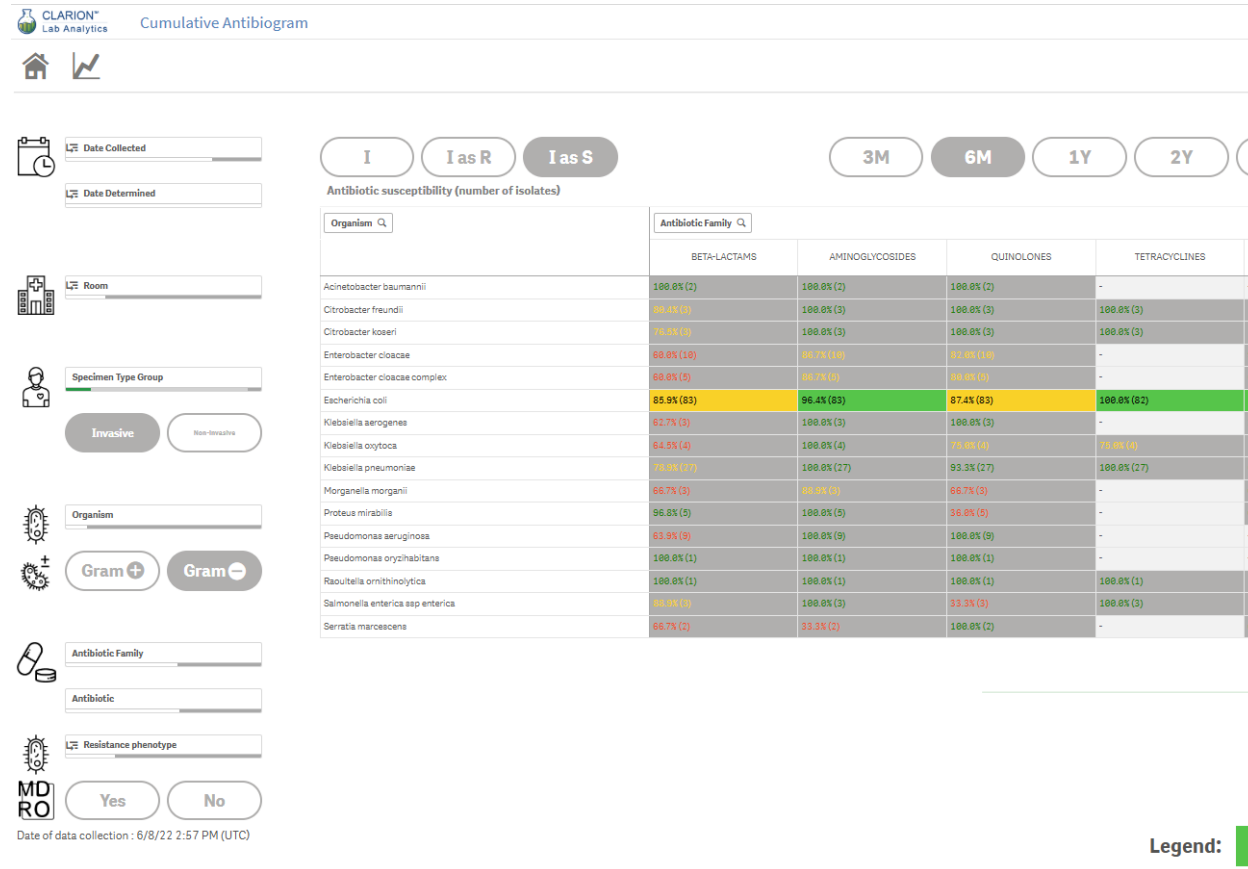
ADVANCED ANTIBIOGRAM



ROUTINE ANTIBIOGRAM

AMS KPI

- On demand by ASP (mainly ID pharmacist, ID physician)
- Advanced antibiogram allows ID and clinical microbiology experts to filter results to better match the particular patient in need.



Usage:

- If questions come up on susceptibility patterns on specific floor/unit
- Susceptibility patterns on rarely encountered organisms
- Specific patient condition

Impact on AMS:

- **Syndromic antibiogram** provides an increased likelihood of appropriate empiric antibiotic therapy for a specific infectious syndrome, considering the weighted incidence of pathogens causing the syndrome
- Assist with more targeted empirical therapy based on infection type, patient type, organism, isolate source, patient location

Actions of improvement:

- Increase awareness of antibiotic resistance
- Education for the interpretation of real-time antibiograms

CN

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MA

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[SHORTENING THE TIME TO ADJUSTMENT WHEN EMPIRIC THERAPY IS INCORRECT]

THE USE OF RDT COMBINED WITH AN ASP AND A CLINICAL DECISION SUPPORT SYSTEM CAN DECREASE THE TIME TO APPROPRIATE THERAPY FOR BLOOD STREAM INFECTIONS

Journal of
Antimicrobial
Chemotherapy

J Antimicrob Chemother 2024; 79 Suppl 1: i37–i43
<https://doi.org/10.1093/jac/dkac277>

Getting rapid diagnostic test data into the appropriate hands by leveraging pharmacy staff and a clinical surveillance platform: a case study from a US community hospital

Jeremy Frens^{1*}, Tyler Baumeister², Emily Sinclair¹, Dustin Zeigler¹, John Hurst³, Brandon Hill³, Sonya McElmeel⁴ and Stéphanie Le Page⁵

¹Department of Pharmacy, Cone Health, 1200 North Elm Street, Greensboro, NC, USA; ²Department of Pharmacy, Williamson Medical Center, Franklin, TN, USA; ³bioMérieux US Medical Affairs, bioMérieux, Durham, NC, USA; ⁴Department of Pharmacy, University of North Carolina Health, Chapel Hill, NC, USA; ⁵bioMérieux Global Medical Affairs Microbiology, bioMérieux, Marcy-l'Étoile, France

*Corresponding author. E-mail: Jeremy.frens@conehealth.com

Received 1 September 2023; accepted 17 June 2024

Objectives: To outline the procedural implementation and optimization of rapid diagnostic test (RDT) results for bloodstream infections (BSIs) and to evaluate the combination of RDTs with real-time antimicrobial stewardship team (AST) support plus clinical surveillance platform (CSP) software on time to appropriate therapy in BSIs at a single health system.

Methods: Blood culture reporting and communication were reported for four time periods: (i) a pre-BCID [BioFire® FilmArray® Blood Culture Identification (BCID) Panel] implementation period that consisted of literature review and blood culture notification procedure revision; (ii) a BCID implementation period that consisted of BCID implementation, real-time results notification via CSP, and creation of a treatment algorithm; (iii) a post-BCID implementation period; and (iv) a BCID2 implementation period. Time to appropriate therapy metrics was reported for the BCID2 time period.

Results: The mean time from BCID2 result to administration of effective antibiotics was 1.2 h (range 0–7.9 h) and time to optimal therapy was 7.6 h (range 0–113.8 h) during the BCID2 Panel implementation period. When comparing time to optimal antibiotic administration among patients growing ceftriaxone-resistant Enterobacterales, the BCID2 Panel group (mean 2.8 h) was significantly faster than the post-BCID Panel group (17.7 h; $P=0.0041$).

Conclusions: Challenges exist in communicating results to the appropriate personnel on the healthcare team who have the knowledge to act on these data and prescribe targeted therapy against the pathogen(s) identified. In this report, we outline the procedures for telephonic communication and CSP support that were implemented at our health system to distribute RDT data to individuals capable of assessing results, enabling timely optimization of antimicrobial therapy.

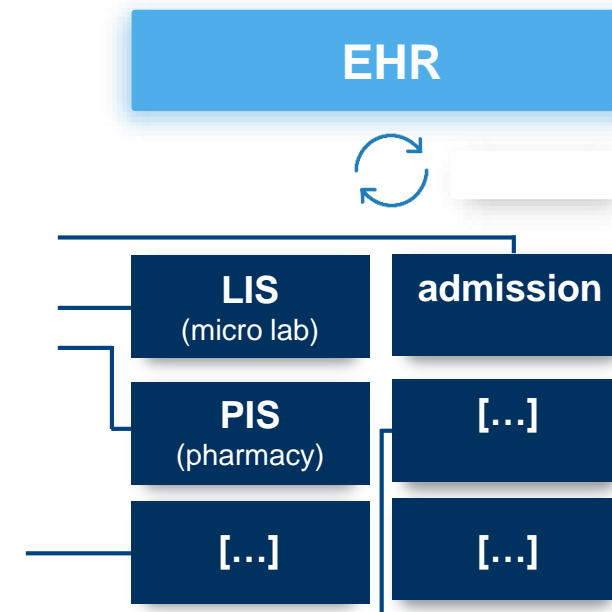
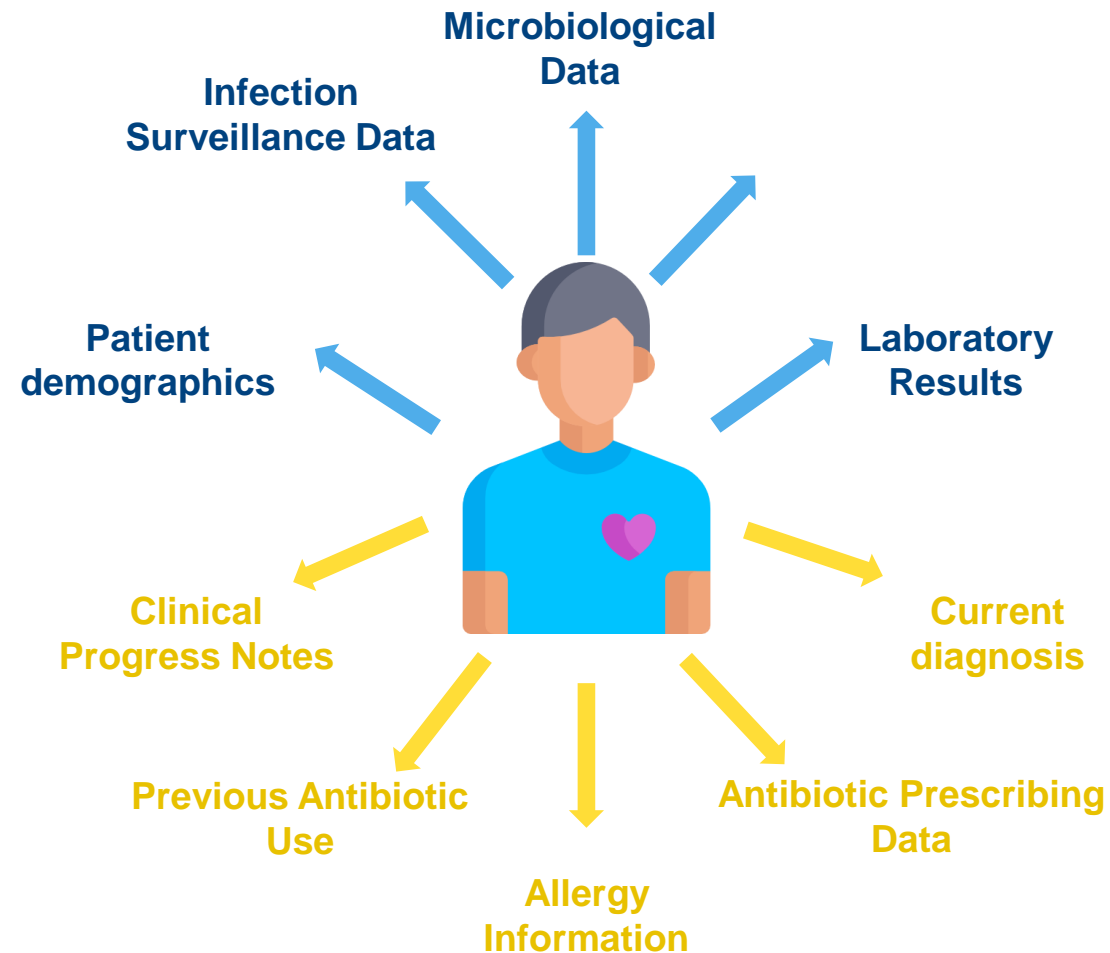
Time to effective antibiotic therapy for ESBL-producing organisms not empirically covered decreased from 17.7 to 2.8 hours after BCID2 implementation ($p = 0.0041$).

Table 2. Mean time to effective therapy from BCID2 result to administration of effective therapy for isolates not empirically covered

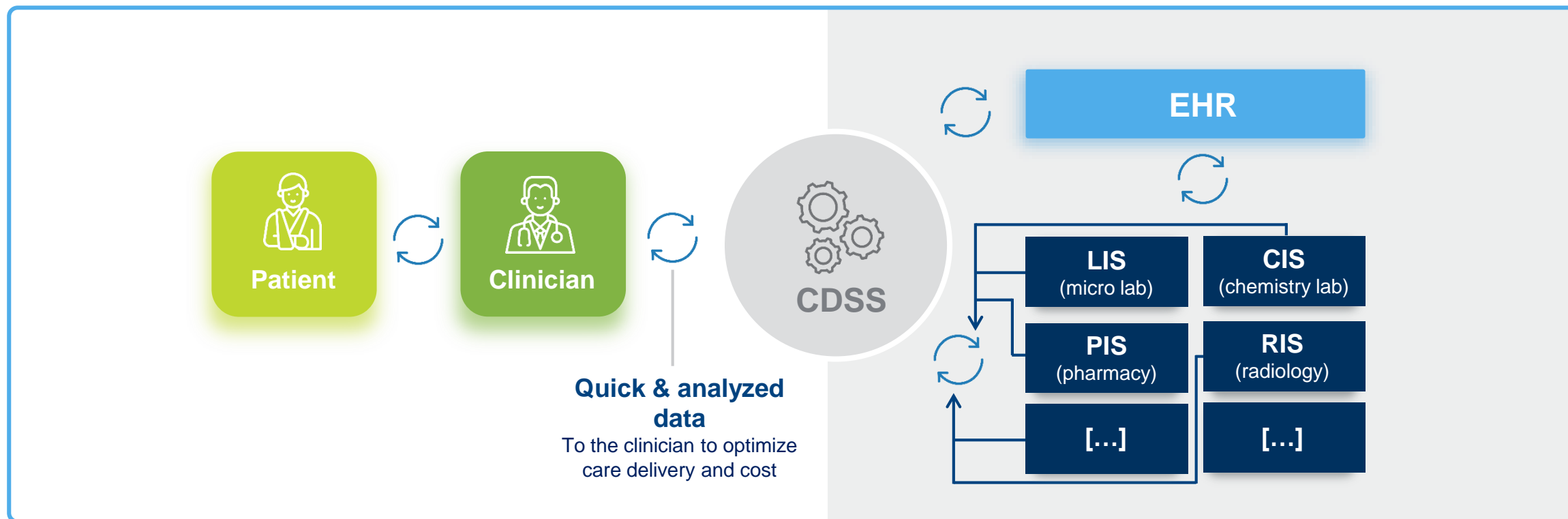
	Mean time to effective therapy from BCID2 result to administration of effective therapy for isolates not empirically covered, h (mean \pm SD)
Before BCID2— ceftriaxone-resistant isolates	17.7 \pm 22.0
After BCID2— CTX-M-positive isolates	2.8 \pm 2.1
P value	0.0041

“The combination of RDT, ASP support and CDSS can greatly aid in optimizing the timing of appropriate antibiotic therapy in patients with BSIs.”

TO TAKE RIGH DECISION, THERE IS A NEED TO ACCESS TO PATIENT DATA FROM SEVERAL DATA SOURCES



CLINICAL DECISION SUPPORT SYSTEMS (CDSS) INTEGRATED WITH EFFECTIVE DIAGNOSTICS
PLAY A KEY ROLE IN ANTIBIOTIC STEWARDSHIP BY GUIDING APPROPRIATE TREATMENT
DECISIONS, REDUCING UNNECESSARY ANTIBIOTIC USE, LOWERING ANTIBIOTIC CONSUMPTION,
ENHANCING GUIDELINE ADHERENCE, AND NARROWING THE SPECTRUM OF ANTIBIOTIC
PRESCRIBING.



LIS: Laboratory Information Systems CIS: Clinical Information Systems, RIS: Radiology Information Systems,
PIS: Pharmacy Information Systems, EHRs: Electronic Health Records, or EHRs,

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ALERTS EXPLAINING WHAT COULD BE DONE AND...



PROPOSING INTERVENTION TO SHORTEN THE TIME TO APPROPRIATE THERAPY

APSS+

Review

Cruz, Ross

Interventions

Meropenem

Person icon

Binoculars icon

Clipboard icon 10

Calendar icon

Back

Clipboard

Clipboard with red dot

Forward

Left arrow

Right arrow

Cruz, Ross (95952) ⓘ
Female - 86 years old (1938-11-08)
Admission date: 2025-04-22 07:04
8528-1 8528 (STM-Gynécologie, urologie et ORL)
ST-MATHIEU

Clipboard with green checkmark

More options

Summary ⓘ

Warning

Settings

VIP Comment

No documented VIP comment

Current admission

Reason of admission

Attending physician

Admission date 2025-04-22 07:04

Visit end

Anthropometric data

Height (cm) 157 cm ⓘ

Reminders

Calendar

Calendar with plus

Notes

Pin

Clipboard

Intervention

Posology

Date 2025-05-15
YYYY-MM-DD

Type Posology

Prescription assessment

Intervention

Interlocutor

Approval status Pending

Creation date
YYYY-MM-DD

Message

Working diagnosis:

Opinion:
The prescribed daily dose 2000 is lower than the recommended minimum daily dose 3000.
It is suggested to adjust the dose based on the upper end of the recommended dose, especially for the first doses, to fill the increased volume of distribution in morbidly obese patients. Meropenem shows time-dependent antibiotic killing, which means that its activity relates most to the percentage of time that serum drug concentrations remain above the minimum inhibitory concentration (MIC). You should consider a prolonged infusion(3-hr infusion, 1st dose over 30 minutes).

Suggestion:

Print

Document with green checkmark

Microscope with green checkmark

Dropdown arrow

Prescription with green checkmark

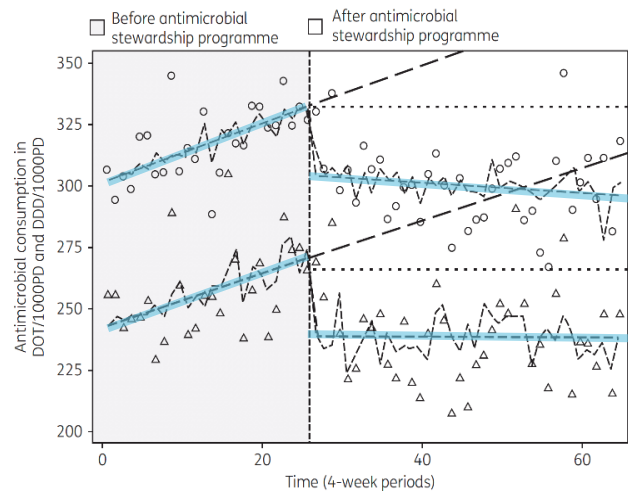
Settings with green checkmark

OK

Cancel

CLINICAL DECISION SUPPORT SYSTEMS ARE SUCCESSFUL WHEN INTEGRATED INTO WIDER STEWARDSHIP PROGRAMS

Consumption pre- and post-implementation



Antimicrobial consumption

~24%

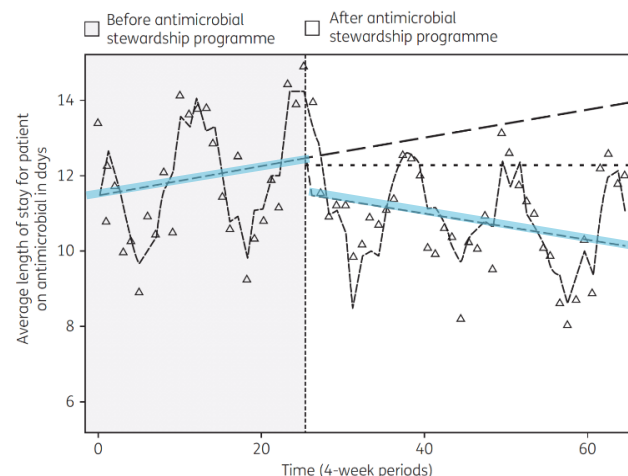
Antimicrobial Prescription Surveillance System for pharmacist-led prospective audit and feedback.

Interrupted time series analysis of intervention for **35,778 patients** receiving antimicrobials **over a 6-year period in CANADA.**

Surveillance system led to reductions in:

- Antimicrobial consumption
- Antimicrobial spending
- Patient length of stay
- Inappropriate prescriptions

Length of stay pre- and post-implementation



Length of stay

~2 days

○ Antimicrobial DOT/1000PD
△ Antimicrobial DDD/1000PD
----- Linear and model trend
— Linear projection of the trend prior to the intervention
..... Conservative mean projection

ANTICIPATING EMERGING THREATS THROUGH EFFECTIVE SURVEILLANCE



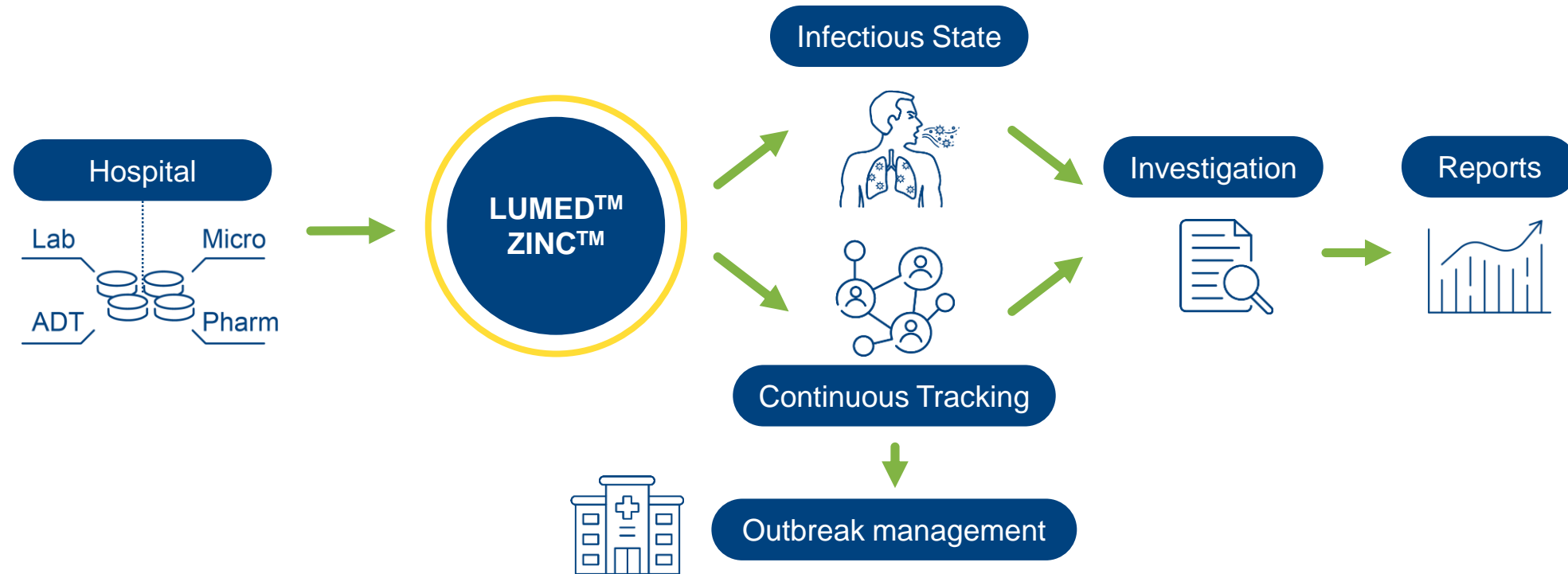
MDRO SURVEILLANCE

The dashboard displays the following components:

- Navigation & Filters:**
 - MDROs (Main Category)
 - AMS KPI (Antimicrobial Stewardship Key Performance Indicator)
 - Target: Monitoring of MDRO (report incidence and analyze trends)
 - MDRO dashboard (Section Header)
 - Filters: Date Collected, Date Determined, Institution, Specimen Type Group (Invasive, Non-Invasive), Organism (Gram+, Gram-), Antibiotic Family, Antibiotic, Resistance phenotype (Yes, No).
- Main Chart: Phenotypes occurrence (Top 15)**
 - Time Period: 3M, 6M, 1Y, 2Y, All (6M selected)
 - Legend: CRE ENTEROBACTER, VRE, ESBL E COLI, MRSA
 - Y-axis: Phenotypes occurrence (0 to 140)
 - X-axis: Time (May 2021 to Nov 2021)
 - Data points (approximate): May 2021 (39, 17, 8, 2), Jun 2021 (87, 24, 9, 2), Jul 2021 (97, 24, 18, 2), Aug 2021 (93, 21, 11, 2), Sep 2021 (92, 33, 18, 2), Oct 2021 (75, 24, 14, 8), Nov 2021 (13, 13, 7, 2).
- Phenotypes occurrence (Bar Chart)**
 - Y-axis: Phenotypes occurrence (0 to 500)
 - X-axis: MRSA, ESBL E COLI, VRE, CRE ENTEROBACTER
 - Data points: MRSA (487), ESBL E COLI (148), VRE (75), CRE ENTEROBACTER (18).
- Phenotypes occurrence per location (Bar Chart)**
 - Y-axis: Phenotypes occurrence per location (0 to 450)
 - X-axis: Two locations (unlabeled)
 - Data points: Location 1 (421), Location 2 (385).

LUMED ZINC FOR INFECTION PREVENTION AND CONTROL IS A REAL TIME TOOL TRACKING HOSPITAL ACQUIRED INFECTIONS AND OUTBREAKS (BOTH BACTERIAL & VIRAL) – PRIORITIZING & PROPOSING ACTIONS

Preventing the spread



**TRACK ALL PATIENTS
AT RISK IN REAL TIME**
From data to action:

ACTIVATE WORKFLOWS THAT PROMPT
IMMEDIATE ACTION BASED ON REAL-TIME
ALERTS TO MITIGATE THE RISK OF
SPREADING DANGEROUS PATHOGENS

KEY-MESSAGES

- **Improve laboratory management efficiency**
- **Decrease the time to an actionable diagnostic result**
- **Improve the accuracy of initial empiric antibiotic therapy**
- **Shorten the time to adjustment when empiric therapy is incorrect**
- **Anticipate emerging threats through effective surveillance**

QUIZZ MMM X BIOMÉRIEUX



- Quizz lancé en scannant le code QR
- Les résultats seront annoncés le 23 Mai à 13h30 au niveau du stand biomérieux x MMM
- Plusieurs cadeaux prévus pour les gagnants
- Bonne chance !





PIONEERING DIAGNOSTICS