

BD MAX™ StaphSR

## Simplify the standard of care for your presurgical screening

BD MAX™ StaphSR for detection of *Staphylococcus aureus* or methicillin-resistant *S. aureus* (MRSA) can be part of a comprehensive surgical site infection (SSI) prevention strategy.





## You may be missing opportunities to prevent SSIs

**Surgical site infections (SSIs)** affect up to **300,000 patients a year in the United States**.<sup>1</sup> SSIs are considered the most common and costly healthcare-acquired infection (HAI), accounting for **20% of all HAIs with an annual burden of \$3.5 to \$10 billion** for increased length of stay, visits to the emergency department, and readmissions.<sup>1</sup>

In one study, SSIs led to nearly twice the length of stay (10.56 days vs 5.64 days).<sup>2</sup> About 2–5% of patients undergoing inpatient surgery will have an SSI.<sup>1</sup>

It is important to test for both methicillin-resistant *S. aureus* (MRSA) and methicillin-sensitive *S. aureus* (MSSA). In one study, **50% of the *S. aureus*-related SSIs in this hospital population were caused by MSSA**, underscoring “the importance of screening pre-operatively for both MRSA and MSSA.”<sup>4</sup> Limited pre-surgical screening protocols that don’t cover both MRSA and MSSA could lead to missed opportunities to prevent SSIs.



**50%**

of *S. aureus*-related SSIs can come from MSSA<sup>4</sup>



**SSIs**

are the most common and costly healthcare-acquired infection (HAI)<sup>1</sup>



**3-6x**

higher risk of HAI in asymptomatic *S. aureus* carriers<sup>3</sup>

Pre-surgical screening with culture-only based methods may lead to less-sensitive detection of SSIs, slower results, and inconsistent timing of reports, which can inhibit the workflows of your surgical and antimicrobial stewardship teams.<sup>4</sup>

Rapid PCR screening and decolonization of nasal carriers of *S. aureus* upon admission can reduce the number of SSIs in a hospital.<sup>3</sup>

# BD MAX™ StaphSR offers accurate presurgical testing

## BD MAX™ StaphSR

- Identified more *S. aureus* (MRSA and MSSA) than direct culture.<sup>4</sup>
- Can provide results for 24 samples in just over 2 hours.<sup>5</sup>
- Can improve time-to-report consistency, thus streamlining workflows vs culture-based screening.<sup>4</sup>

### Performance of BD MAX™ StaphSR vs direct culture and broth-enriched culture<sup>4</sup>

Variable	Test	Sensitivity, % (n/total)	95% CI, %	Sensitivity, % (n/total)	95% CI, %
<i>S. aureus</i>	BD MAX™ StaphSR	98.2 (271/276)	95.8–99.2	93.1 (783/841)	91.2–94.6
	Direct culture	85.1 (235/276)	80.5–88.9	99.4 (836/841)	98.6–99.7
	Enriched culture	95.7 (264/276))	92.6–97.5	99.3 (835/841)	98.5–99.7
MRSA	BD MAX™ StaphSR	96.7 (29/30))	83.3–99.4	98.5 (1071/1087)	97.6–99.1
	Direct culture	76.7 (23/30)	59.1–88.2	99.4 (1080/1087)	98.7–99.7
	Enriched culture	96.7 (29/30))	83.3–99.4	99.4 (1081/1087)	98.8–99.7

Use of BD MAX™ StaphSR resulted in ~5-fold and 10-fold reductions in SSI rates per 100 surgeries vs screening for MRSA-only (direct culture) and no screening, respectively.<sup>4</sup>



# Real-world results you can count on

A recent study investigated whether **BD MAX™ StaphSR** could affect the occurrence of SSI<sup>1</sup>s caused by *S. aureus* in orthopedic, cardiac, neurosurgery, or any implantable device surgery patients in a multihospital setting compared to clinician-ordered MRSA-only testing using traditional culture (MRSA-SC) or no screening at all.<sup>4</sup>

\*Data involving surgeries, SSI<sup>1</sup>s and presurgical screening between February 2018 and February 2019 at Rhode Island Hospital.

<sup>#</sup>Two positive results during prior screening by BD MAX™ StaphSR consisted of one MSSA positive and one MRSA positive.

<sup>†</sup>SSI rate per 100 cases of respective surgery type.

<sup>‡</sup>5 MSSA infections and 1 MRSA infection.

<sup>§</sup>5 MSSA infections and 3 MRSA infections.

<sup>¶</sup>10 MSSA infections and 4 MRSA infections.

## Incidences of SSI<sup>1</sup>s for patients screened with BD MAX™ StaphSR vs MRSA-only (direct culture) testing using traditional culture (MRSA-CS) vs no testing<sup>\*4</sup>

Screening	Data
<b>Surgeries, N</b>	3388
<b>Prior screening, n (%)</b>	
BD MAX™ StaphSR	2050 (59.7)
MRSA-SC	724 (21.1)
No testing	614 (17.9)
<b>SSI<sup>1</sup>s, N</b>	28
<b>Pathogen, n</b>	
<i>S. aureus</i>	14
MRSA	14
<b>Prior screening, n</b>	
BD MAX™ StaphSR	6 (2 positive <sup>#</sup> ; 4 negative)
MRSA-SC	8 (1 positive; 7 negative)
No testing	14
<b>Rate of SSI<sup>1</sup> % (n/total)</b>	
BD MAX™ StaphSR	0.3 (6 <sup>#</sup> /2050)
MRSA-SC	1.10 (8 <sup>§</sup> /724)
None	2.28 (14 <sup>¶</sup> /614)

**This study shows that 50% of the *S. aureus*-related SSI<sup>1</sup>s were due to MSSA.<sup>4</sup>**

# Achieve infection prevention goals with BD MAX™ StaphSR

Minimizing the risk and number of SSIs acquired in the hospital can be achieved with **rapid PCR screening** and decolonization of nasal carriers of *S. aureus* upon admission.<sup>3</sup> **BD MAX™ StaphSR** can be a successful presurgical tool to facilitate patient management of SSI, including infection prevention measures and appropriate preoperative antibiotic prophylaxis.<sup>4</sup>



## Provides accurate pre-surgical testing for both MRSA and MSSA

An effective pre-surgical screening protocol should include testing for both MRSA and MSSA, as one study found that **50% of the *S. aureus*-related SSIs were due to MSSA.**<sup>4</sup>



## Enables presurgical prophylaxis

“The **BD MAX™ StaphSR** assay provided accurate detection of both *S. aureus* and MRSA nasal colonization in presurgical patients, allowing infection prevention measures, including **presurgical prophylaxis, to be implemented in a timely and consistent manner to avoid SSIs.**”<sup>4</sup>



## Improves infection control compliance to meet hospital performance goals

The **BD MAX™ StaphSR** assay enables better compliance with hospital-wide pre-operative guidelines for hospital performance measures for surgery, infectious disease and stewardship, and pharmacy departments.<sup>4</sup>



## Supports proper use of vancomycin and AMS

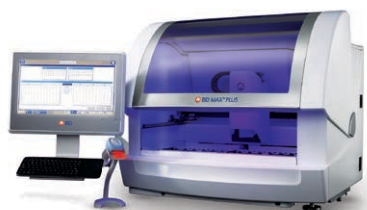
**BD MAX™ StaphSR** facilitates targeted, appropriate use of broad-spectrum vancomycin.<sup>4</sup> Vancomycin should not be administered to MRSA-negative patients as prophylaxis.<sup>1</sup> Reducing the use of inappropriate antibiotics in health care settings can help with antimicrobial stewardship (AMS) goals.<sup>6</sup>

# Streamline workflow and results reporting

“For the preoperative center, the benefit over the nasal culture screen was the consistent reporting time of the **BD MAX™ StaphSR** results to the pre-operative nursing service which allowed a streamlined workflow and reporting to the electronic medical record for the anesthesiologists and surgeons.”<sup>4</sup>

**BD MAX™ StaphSR can improve time-to-report consistency, thus streamlining workflows vs culture-based screening.**<sup>4</sup>

vs.



## BD MAX™ StaphSR orders

Specimens run daily on the BD MAX™ System (Monday-Friday).

**Results reported by morning next day.**



## Traditional culture orders

Specimens batched with the MRSA medical admissions.

**Results reported inconsistently to providers** because of arrival time in the microbiology lab and batch read times by two different shifts.







Up to 60% of SSIs are preventable and require a multi-disciplinary approach to solving.<sup>1</sup>

## Evidence-based recommendations on SSI:

Molecular testing with **BD MAX™ StaphSR** can help you to correctly identify and prophylactically treat patients colonized with MRSA and MSSA as well as limit the overutilization of vancomycin.<sup>4</sup> Consistent reporting time of the **BD MAX™ StaphSR results can streamline workflow and reporting to anesthesiologists and surgeons.**<sup>4</sup>

American College of Surgeons and Surgical Infection Society's guidelines on MRSA screening include the following recommendations:<sup>1</sup>

### Screening protocol

Implementation of global *S. aureus* screening and decolonization protocols should depend on institution's baseline SSI and MRSA rates.

### Joint replacement and cardiac procedures

"Clinical practice guidelines from the American Society of Health-System Pharmacists recommend screening and nasal mupirocin decolonization for *S. aureus*-colonized patients before total joint replacement and cardiac procedures."

### MRSA bundles

Screening, decolonization, contact precautions, and hand hygiene as part of a MRSA bundle are "highly effective if adhered to, otherwise there is no benefit."

### Decolonization protocol

"No standard decolonization protocol supported by literature; consider nasal mupirocin alone vs nasal mupirocin plus chlorhexidine gluconate bathing."  
"Decolonization protocols should be completed close to date of surgery to be effective."

### Prophylaxis

"Vancomycin should not be administered as prophylaxis to MRSA-negative patients." Inappropriate use of vancomycin alone in MRSA-negative patients was associated with increased risk of MSSA SSIs.

The Joint Commission's National Patient Safety Goals (2022):<sup>7</sup>

### Improve staff communication

"Get important test results to the right staff person on time."



## Streamlined integration into existing workflow with the BD MAX™ System family

- The BD MAX™ System family offers you a fully integrated, automated real-time PCR platform with a broad menu of molecular IVD and open-system tests.<sup>8</sup>
- The automated workflow and analytical performance reduces the need for manual tasks and achieves more rapid results.<sup>\*9,10</sup>  
\*When compared to culture or immunochromatographic antigen (IA).
- The compact and self-contained unitized reagent strips and the new reclosing septum cap simplify waste management and help reduce the risk of contamination.



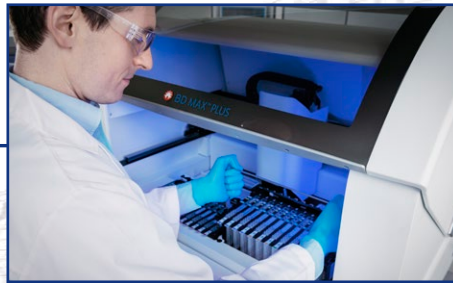
## Snap

Assemble unitized reagent strips with extraction and PCR reagents.



## Load

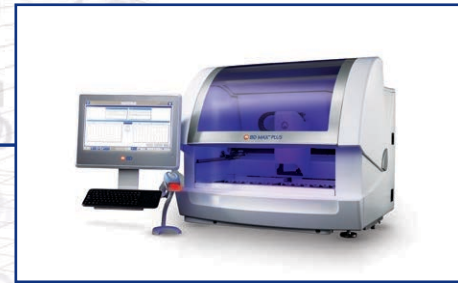
Load the Sample Buffer Tubes, PCR cartridges, and racks.



## Go

Come back in just over 2 to 3 hours for results.\*

\*Assay times may vary.



Less than **1.5 minutes** hands-on time per sample<sup>10,11</sup>



**24 patient** results in **just over 2 to 3 hours**<sup>11</sup>



**96 samples** per **8 hour shift**<sup>11</sup>

# Performance for presurgical screening

BD MAX™ StaphSR assay provides accurate and rapid detection of both *S. aureus* and MRSA nasal colonization in presurgical patients.<sup>4</sup>

Identified more *S. aureus* (MRSA and MSSA) than direct culture<sup>4</sup>

Resulted in ~5-fold reduction in SSI rates per 100 surgeries vs MRSA-only screening using traditional culture<sup>4</sup>



**Assay targets** : *Staphylococcus aureus* (SA) DNA and methicillin-resistant *Staphylococcus aureus* (MRSA) DNA

**mecA** : Yes

**mecC** : Yes

**MREJ Types** : 11 types detected

**Patient population** : Patients at risk for nasal colonization

**Specimen** : Nasal swab

**IVD part number** : 443419 (24 tests)

For more information about BD MAX™ Molecular Diagnostic System, please visit: [bd.com](https://www.bd.com)

HAI, healthcare-acquired infection; MRSA, methicillin-resistant *S. aureus*; MSSA, methicillin-sensitive *S. aureus*; MRSA-SC, methicillin-resistant *S. aureus* screening with traditional culture; PCR, polymerase chain reaction; SA, *S. aureus*; SSI, surgical site infection.

**References:** 1. Ban KA et al. *J Am Coll Surg.* 2017;224(1):59–74. 2. Shepard J et al. *JAMA Surg.* 2013;148(10):907–14. 3. Bode LG et al. *N Engl J Med.* 2010;362(1): 9–17. 4. Tansarli GS et al. *J Mol Diagn.* 2020;22(8):1063–9. 5. BD MAX™ Assay Run Times – Internal Document. 6 Joint Commission on Hospital Accreditation. *Jt Comm Perspect.* 2016;36(7):1–8. 7. The Joint Commission. Hospital: 2022 National Patient Safety Goals. Simplified version. Available at: [https://www.jointcommission.org/-/media/tjc/documents/standards/national-patient-safety-goals/2022/simple\\_2022-hap-npsg-goals-101921.pdf](https://www.jointcommission.org/-/media/tjc/documents/standards/national-patient-safety-goals/2022/simple_2022-hap-npsg-goals-101921.pdf). 8. BD MAX™ System User's Manual. Becton, Dickinson and Company: Sparks, MD. 9. Mortensen JE et al. *BMC Clin Pathol.* 2015;15:9. 10. Hirvonen JJ et al. *Eur J Clin Microbiol Infect Dis.* 2015;34(5):1005–9. 11. Felder RA, et al. *J Lab Autom.* 2014;19(5):468–73.

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