

VERIGENE[®]

The VERIGENE[®] System | Enabling Better Care. Today.

VERIGENE[®] Gram-Positive Blood Culture Nucleic Acid Test (BC-GP)



VERIGENE[®] BC-GP is a qualitative, multiplexed, *in vitro*, diagnostic test performed on the VERIGENE System, that rapidly identifies genus, species, and genetic resistance markers for a broad panel of gram-positive bacteria directly from positive blood culture bottles.

While conventional microbiological methods may require 2 to 4 days to produce bacterial identification and susceptibility results, VERIGENE BC-GP provides results within 2.5 hours of blood culture positivity. Features include:

- Automation, with a sample to result system
- An on-demand and scalable workflow
- Fast time to results, with <5 minutes hands-on time and <2.5 hours run time

“Implementation of the BC-GP assay contributed to a reduction in time to appropriate antimicrobial therapy, regardless of patient population, and a decrease in LOS and overall hospital costs among patients without other significant comorbidities.”

Felsenstein S, Mender JM, Sposto R, et al.¹

VERIGENE[®] BC-GP

Species

Staphylococcus aureus

Staphylococcus epidermidis

Staphylococcus lugdunensis

Streptococcus anginosus Group

Streptococcus agalactiae

Streptococcus pneumoniae

Streptococcus pyogenes

Enterococcus faecalis

Enterococcus faecium

Genus

Staphylococcus spp.

Streptococcus spp.

Listeria spp.

Resistance

mecA (methicillin)*

vanA (vancomycin)**

vanB (vancomycin)**

*The assay detects the presence of the *mecA* gene in a sample, but does not determine which *Staphylococcus* species (*S. aureus* and or *S. epidermidis*) produced the gene.

**The assay detects the presence of the *vanA* or *vanB* gene in a sample, but does not determine which *Enterococcus* species (*E. faecalis* and/or *E. faecium*) produced the gene.

Performance

VERIGENE® BC-GP Performance vs. Reference Methods²

| Target | Positive Agreement (%) | Negative Agreement (%) |
|---|------------------------|------------------------|
| Species | | |
| <i>Staphylococcus aureus</i> | 99.1 | 100 |
| <i>Staphylococcus epidermidis</i> | 93.1 | 98.9 |
| <i>Staphylococcus lugdunensis</i> | 95.0 | 100 |
| <i>Streptococcus agalactiae</i> | 98.6 | 100 |
| <i>Streptococcus anginosus</i> Group | 100 | 99.8 |
| <i>Streptococcus pneumoniae</i> | 100 | 99.6 |
| <i>Streptococcus pyogenes</i> | 95.8 | 100 |
| <i>Enterococcus faecalis</i> | 96.9 | 99.9 |
| <i>Enterococcus faecium</i> | 97.1 | 100 |
| Genus | | |
| <i>Staphylococcus</i> spp. | 98.0 | 99.4 |
| <i>Streptococcus</i> spp. | 93.6 | 99.6 |
| <i>Listeria</i> spp. | 100 | 100 |
| Resistance | | |
| <i>mecA</i> — <i>S. aureus</i> * (methicillin) | 97.5 | 98.8 |
| <i>mecA</i> — <i>S. epidermidis</i> * (methicillin) | 92.0 | 81.5 |
| <i>vanA</i> ** (vancomycin) | 94.2 | 99.8 |
| <i>vanB</i> ** (vancomycin) | 100 | 100 |

*The assay detects the presence of the *mecA* gene in a sample, but does not determine which *Staphylococcus* species (*S. aureus* and or *S. epidermidis*) produced the gene.

**The assay detects the presence of the *vanA* or *vanB* gene in a sample, but does not determine which *Enterococcus* species (*E. faecalis* and/or *E. faecium*) produced the gene.

Usage

VERIGENE® BC-GP is indicated for use in conjunction with other clinical and laboratory findings to aid in the diagnosis of bacterial bloodstream infections; however, it is not to be used to monitor these infections. See the Package Insert for more information.

Ordering Information

| Product Name | Part Number |
|---|-------------|
| VERIGENE® Gram-Positive Blood Culture (BC-GP) Nucleic Acid Test Kit Includes: 20 BC-GP Test Cartridges 20 Extraction Trays | 20-005-018 |
| VERIGENE® Gram-Positive Blood Culture (BC-GP) Nucleic Acid Utility Kit Includes: 20 BC-GP Utility Trays | 20-012-018 |

References

1. Felsenstein S, Mender JM, Sposto R, et al. Impact of a rapid blood culture assay for gram-positive identification and detection of resistance markers in a pediatric hospital. Arch Pathol Lab Med 2016;140:267-75.
2. VERIGENE Gram-Positive Blood Culture Nucleic Acid Test (BC-GP) Package Insert (027-00030-01). Combined results obtained testing prospective fresh and/or frozen blood culture specimens.

Luminex
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For In Vitro Diagnostic Use. Products are region specific and may not be approved in some countries/regions. Please contact Luminex at support@luminexcorp.com to obtain the appropriate product information for your country of residence.

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