

Clinical Bulletin: DxWound *ermA* and *ermB* Tests

Description

Antibiotic resistance is considered by the CDC to be a major public health threat in the United States.¹ Patients with antibiotic resistant infections have higher morbidity and mortality, including longer hospital stays with increased costs to the healthcare system.¹ To help mitigate the consequences of antibiotic resistance, the CDC has called for improved antibiotic stewardship and the development of new diagnostic tests.¹

Antibiotic resistance genes are genes present in bacteria that mediate resistance to a specific antibiotic or class of antibiotics. DxWound tests for several antibiotic resistance genes associated with bacteria that typically cause skin and soft tissue infections (SSTIs), including the erythromycin-resistance rRNA methylase (*erm*) genes *ermA* and *ermB*. The *ermA* and *ermB* genes have been found in *Staphylococcus*, *Streptococcus*, *Enterococcus*, and a broad range of other bacterial genera.^{2,3} They mediate resistance to macrolides (e.g. azithromycin) and clindamycin.^{2,4}

Indications for Ordering

The DxWound *ermA* and *ermB* tests are ordered based on the clinician's determination of medical necessity. Tests for *ermA* and *ermB* can be useful for any patient suspected of having a skin and soft tissue infection (SSTI) caused by *Staphylococcus*, *Streptococcus*, or *Enterococcus* and for which macrolides or clindamycin may be considered. The presence of *ermA* or *ermB* may indicate resistance to macrolides or clindamycin.^{2,4} Testing for *ermA* and *ermB* genes may help clinicians rapidly target antibiotic therapy.

Methodology

DxWound tests analyze DNA using real-time polymerase chain reaction (PCR) technology. The tests are performed on a swab sample taken directly from the site of infection without need for bacterial growth. The process is simple. First, cleanse and debride the wound as necessary. The Levine technique is then suggested for collection of the sample. The swab sample is collected in a solution that kills microorganisms at the same time as protecting microbial DNA, essentially preserving the wound microbiome in time at the point of specimen collection. DNA is detected directly from the patient specimen without culture enrichment. All DxWound tests have ≥95% accuracy and ≥99% reproducibility. Accuracy and reproducibility were established by testing reference specimens with characterized organisms or antibiotic resistance genes obtained from multiple sources including the FDA-CDC antimicrobial resistance isolate bank, commercial vendors, and other laboratories. For more information, please refer to the Technical Specifications bulletin.

Clinical Use

Macrolides and clindamycin may be used to treat infections caused by various *Staphylococcus* or *Streptococcus* species. However, resistance to these antibiotics has been growing. For example, erythromycin-resistant group A *Streptococcus* and clindamycin-resistant group B *Streptococcus* have been identified as antibiotic resistance threats in the United States by the CDC.¹

The *erm* enzymes are the most common and best studied resistance mechanism against macrolides and clindamycin.⁴ Macrolide antibiotics, such as azithromycin, as well as the structurally unrelated clindamycin, bind to the 50S ribosomal subunit of the bacterial rRNA complex, thereby inhibiting protein synthesis.³ The *erm* genes encode adenine-N6 methyltransferases, which add one or two methyl groups to a single adenine in 23S rRNA of the 50S ribosomal subunit. This modification reduces the binding of macrolides and clindamycin to the 50S ribosomal subunit resulting in resistance.³

Test Interpretation

These genetic tests identify if *ermA* and/or *ermB* genes are present. The *ermA* and *ermB* genes have a broad host range and have been associated with multiple bacterial genera.^{2,3}

Although these genes have been reported in a wide range of diverse organisms, most data on their prevalence and clinical impact have only involved *Staphylococcus*, *Streptococcus*, and *Enterococcus*. Pilot study data of DxWound samples indicate a strong association between the *ermA* and *ermB* genes with aerobic Gram-positive cocci, including *Enterococcus faecalis*, *Enterococcus faecium*, *Staphylococcus aureus*, *Staphylococcus epidermidis*, *Staphylococcus lugdunensis*, *Streptococcus agalactiae* (group B), and *Streptococcus pyogenes* (group A).⁵

The association of *ermA* and *Staphylococcus aureus* is highly likely when they are co-detected. If samples are positive for *ermA*, *S. aureus*, and coagulase-negative *Staphylococcus* species (CoNS), the probability of *S. aureus* being a carrier of *ermA* is 81% vs. 32% for CoNS.⁵ In the absence of *S. aureus*, an association of *ermA* with a CoNS or a β -hemolytic *Streptococcus* (group A *Streptococcus pyogenes* or group B *Streptococcus agalactiae*) is likely. In case of co-detection of *ermA*, a CoNS, and a β -hemolytic *Streptococcus*, the CoNS has more than 50% chance to be a carrier.⁵ The *ermA* gene is not often identified in *Enterococcus*. See Table 1 for additional information for *ermA* test interpretation.

Table 1. DxWound *ermA* Test Interpretation^{6,7}

DxWound Test Results	Potential Organism Present*	Common Antibiotic Coverage	
		Oral	IV
<i>ermA</i> + <i>Streptococcus pyogenes</i> or <i>Streptococcus agalactiae</i>	Macrolide/Clindamycin-Resistant <i>Streptococcus</i> **	Amoxicillin, Amox-Clav, Cephalexin, Linezolid [#]	Cefazolin, Ceftriaxone, Penicillin G, Vancomycin [#]
<i>ermA</i> + <i>Staphylococcus</i> species	Macrolide/Clindamycin-Resistant <i>Staphylococcus</i> species	Cephalexin, Dicloxacillin, Trimethoprim-Sulfamethoxazole	Cefazolin, Nafcillin, Oxacillin, Vancomycin [#]
<i>ermA</i> + <i>mecA</i> + <i>Staphylococcus</i> species	Macrolide/Clindamycin-Resistant Methicillin-Resistant <i>Staphylococcus</i> species	Doxycycline, Linezolid, Minocycline, TMP-SMX	Daptomycin, Linezolid, Vancomycin [#]

**Erythromycin-resistant group A *Streptococcus* and clindamycin-resistant group B *Streptococcus* are identified as antibiotic resistance threats in the United States by the CDC.

[#]Possible alternative if the patient has a true penicillin allergy

The association of *ermB* and *Enterococcus* is highly likely and consistent with the high prevalence of *ermB* in *Enterococcus* species reported in the literature. There are other potential carriers of *ermB* but the probability of these species carrying *ermB* in pilot study data was small. See Table 2 for additional information for *ermB* test interpretation.

Table 2. DxWound *ermB* Test Interpretation^{6,7}

DxWound Test Results ^{&}	Potential Organism Present*	Common Antibiotic Coverage	
		Oral	IV
<i>ermB</i> + <i>Streptococcus pyogenes</i> or <i>agalactiae</i>	Macrolide/Clindamycin-Resistant** <i>Streptococcus</i>	Amoxicillin, Amox-Clav, Cephalexin, Linezolid [#]	Cefazolin, Ceftriaxone, Penicillin G, Vancomycin [#]
<i>ermB</i> + <i>Staphylococcus</i> species	Macrolide/Clindamycin-Resistant <i>Staphylococcus</i>	Cephalexin, Dicloxacillin, Trimethoprim-Sulfamethoxazole	Cefazolin, Nafcillin, Oxacillin, Vancomycin [#]
<i>ermB</i> + <i>mecA</i> + <i>Staphylococcus</i> species	Macrolide/Clindamycin-Resistant Methicillin-Resistant <i>Staphylococcus</i> species	Doxycycline, Linezolid, Minocycline, TMP-SMX	Daptomycin, Linezolid, Vancomycin [#]

**Erythromycin-resistant group A *Streptococcus* and clindamycin-resistant group B *Streptococcus* are identified as antibiotic resistance threats in the United States by the CDC.

[&] Interpret with caution as *ermB* is commonly identified in *Enterococcus* species

[#]Possible alternative if patient has a true penicillin allergy

Limitations

- These tests detect the presence of genes and do not detect whether the genes are expressed
- These tests do not detect all known antibiotic resistance mechanisms, nor do they identify which species is associated with *ermA* or *ermB*
- It is possible that *ermA* or *ermB* is associated with an organism that is not included in these tests
- The DxWound Genetic Analysis Report does not make recommendations for treatment. All test results should be evaluated in the context of the patient's individual clinical presentation

Technical Assistance

For technical assistance with interpretation or to speak with one of our clinical support specialists, scientists or clinical pharmacists, please call Client Services at (877) 866-0603, Monday–Friday 5:00am to 5:00 pm (Pacific Time).

Continued on next page.

Reporting

All test results are reported, generally, within 1 business day of specimen receipt. These results are available in the online portal as a DxWound Report.

How to Order

Using a CogenDx test requisition (paper or electronic), medically necessary tests are ordered by individual test based on patient-specific elements identified during the clinical assessment and documented in the patient's medical record by the provider. Submit swab specimen in the collection device provided according to the directions in the Specimen Collection Manual. A completed Documentation for DxWound Testing form is required with each order.

References

1. Centers for Disease Control and Prevention, Antibiotic Resistance Threats in the United States, 2013.
2. Roberts MC. Update on macrolide-lincosamide-streptogramin, ketolide, and oxazolidinone resistance genes. *FEMS Microbiol Lett.* 2008; 282(2): 147-59.
3. Roberts MC, Sutcliffe J, Courvalin P, Jensen LB, Rood J, Seppala H. Nomenclature for macrolide and macrolide-lincosamide-streptogramin B resistance determinants. *Antimicrob Agents Chemother.* 1999; 43(12): 2823-30.
4. Fyfe C, Grossman TH, Kerstein K, Sutcliffe J. Resistance to Macrolide Antibiotics in Public Health Pathogens. *Cold Spring Harb Perspect Med.* 2016; 6(10).
5. Data on file, Millennium Health, LLC. 12-18-17.
6. Stevens DL, Bisno AL, Chambers HF, et al. Practice guidelines for the diagnosis and management of skin and soft tissue infections: 2014 update by the Infectious Diseases Society of America. *Clin Infect Dis.* 2014; 59(2): e10-52
7. Lipsky BA, Berendt AR, Cornia PB, et al. 2012 Infectious Diseases Society of America clinical practice guideline for the diagnosis and treatment of diabetic foot infections. *Clin Infect Dis.* 2012; 54(12): e132-73.



Contact CogenDx to learn more.

Client Services: (877) 866-0603

cogendx.com

Millennium Health, LLC tests were developed and their performance characteristics determined by Millennium Health. These tests have not been cleared or approved by the US Food and Drug Administration. FDA clearance or approval is not required for clinical use. Millennium Health is regulated under CLIA as qualified to perform high-complexity testing, and is accredited by the College of American Pathologists (CAP). These tests are used for clinical purposes and should not be regarded as investigational or for research. The test results should be used with other clinical and diagnostic findings for patient case management.

CogenDx is a brand of Millennium Health, LLC. The CogenDx logo is a service mark or registered service mark of Millennium Health, LLC or its subsidiaries in the United States and other countries. All other trademarks used herein are the property of their respective owners.