

## Overview

### Useful For

Confirming carbapenemase production from pure isolates of Enterobacterales or *Pseudomonas aeruginosa*

### Additional Tests

Test Id	Reporting Name	Available Separately	Always Performed
CARNB	Carbapenemase-Carba NP Test	No	Yes

### Special Instructions

- [Infectious Specimen Shipping Guidelines](#)

### Method Name

Colorimetric Detection of Carbapenem Hydrolysis

### NY State Available

Yes

## Specimen

### Specimen Type

Varies

### Shipping Instructions

1. For shipping information see [Infectious Specimen Shipping Guidelines](#).
2. Place specimen in a large infectious container (T146) and label as an etiologic agent/infectious substance.

### Necessary Information

**Specimen source and organism identification are required.**

### Specimen Required

#### Preferred:

**Specimen Type:** Pure culture of Enterobacterales or *Pseudomonas aeruginosa* from source cultured

**Supplies:** Infectious Container, Large (T146)

**Container/Tube:** Agar slant or other appropriate media

#### Collection Instructions:

1. Perform isolation of bacterial isolate.
2. Submit Enterobacterales or *Pseudomonas aeruginosa* isolate in pure culture actively growing. **Do not submit mixed**

cultures.

3. Place the slant into the secondary infectious container for shipment.
4. Each isolate must be submitted under a separate order.

**Acceptable:**

**Specimen Type:** Enterobacterales or *Pseudomonas aeruginosa* isolate swab

**Supplies:**

- E-Swab (T853)
- Infectious Container, Large (T146)

**Container/Tube:** E-Swab collection and transport system

**Collection Instructions:**

1. Perform isolation of bacterial isolate.
2. Utilize the E-swab to obtain an adequate sample of Enterobacterales or *Pseudomonas aeruginosa* isolate in pure culture. **Do not submit mixed cultures.**
3. Place swab into the transport system containing 1-mL liquid Amies transport medium.
4. If needed, break off end of swab and close the transport tube.
5. Place the transport system into the secondary infectious container for shipment.
6. Each isolate must be submitted under a separate order.

**Forms**

If not ordering electronically, complete, print, and send a [Microbiology Test Request](#) (T244) with the specimen.

**Specimen Minimum Volume**

See Specimen Required

**Reject Due To**

Agar plate	Reject
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**Specimen Stability Information**

Specimen Type	Temperature	Time	Special Container
Varies	Ambient (preferred)		
	Refrigerated		

**Clinical & Interpretive**

**Clinical Information**

Gram-negative bacilli (GNB) with acquired carbapenemases have disseminated worldwide, rendering them a global threat. The therapeutic armamentarium for infections caused by carbapenem-resistant Enterobacterales (CRE) is limited, and CRE infections have been associated with significant mortality. Enterobacterales harboring *Klebsiella pneumoniae* carbapenemase (KPC) are endemic in some regions of the United States, and although still sporadic, GNB harboring New Delhi metallo-beta-lactamase (NDM) have been reported from several states. Timely detection of these

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carbapenemases (along with emerging carbapenemases such as OXA-48 and VIM) is important. Detection is challenging since isolates may have only borderline reductions in susceptibility to carbapenems, and carbapenem resistance may be mediated by mechanisms other than carbapenemases (eg, *AmpC* or extended-spectrum beta-lactamase with decreased membrane permeability). While molecular methods are confirmatory, testing may not be immediately available and may be limited by the number of targets assayed. The Carba NP test is preferred over the *mCIM* (modified carbapenem inactivation method) test due to faster turnaround time.

If an isolate is suspected to possess KPC or NDM carbapenemase (eg, due to local epidemiology), CARBI / Carbapenem Resistance Genes, Molecular Detection, PCR, Varies may be preferred over this Carba NP test.

**Reference Values**

Negative

**Interpretation**

A positive result indicates production of a carbapenemase by the isolate submitted for testing. A negative result indicates lack of production of a carbapenemase by the isolate submitted for testing.

**Cautions**

Results of the Carba NP test should be interpreted along with antimicrobial susceptibility testing results. Phenotypic resistance to carbapenems may be due to traits other than carbapenemase production (eg, *AmpC* or extended-spectrum beta-lactamase production with decreased membrane permeability). Additionally, a positive test is only indicative of carbapenemase production in general; the assay does not determine the type of carbapenemase present (eg, NDM-1, KPC, OXA-48-like). If an isolate is suspected to possess KPC or NDM carbapenemase (eg, due to local epidemiology), CARBI / Carbapenem Resistance Genes, Molecular Detection, PCR, Varies may be preferred.

False-negative results may occur due to plasmid loss in isolates submitted for testing, the presence of a nonexpressed carbapenemase gene, or low-level carbapenemase expression.

**Supportive Data**

We evaluated 271 gram -negative bacilli (of which 131 were carbapenemase producers and of which 201 were Enterobacterales) using the Carba NP test and the modified Hodge test. Sensitivity for detection of carbapenemase production was comparable (Carba NP, 100 versus modified Hodge test, 98%,  $p=0.08$ ), but the Carba NP test was more specific (100 versus 80%,  $p<0.0001$ ) and faster.(1)

**Clinical Reference**

1. Vasoo S, Cunningham SA, Kohner PC, et al. Comparison of a novel, rapid chromogenic biochemical assay, the Carba NP test, with the modified Hodge test for detection of carbapenemase-producing Gram-negative bacilli. *J Clin Microbiol*. 2013;51(9):3097-3101
2. Nordmann P, Poirel L, Dortet L. Rapid detection of carbapenemase-producing *Enterobacteriaceae*. *Emerg Infect Dis*. 2012;18(9):1503-1507
3. Clinical and Laboratory Standards Institute (CLSI): Performance Standards for Antimicrobial Susceptibility Testing. 35th ed. CLSI Supplement M100, CLSI; 2025

## Performance

### Method Description

A pure bacterial isolate is emulsified into cell lysis buffer in 2 tubes: one contains the base indicator solution (phenol red with zinc salts) alone and the other contains the base indicator solution plus imipenem (6 mg/mL). The tubes are incubated at 37 degrees C for 2 hours. A positive reaction is indicated by a color change from red to yellow as a result of hydrolysis of the beta-lactam ring of imipenem. (Vasoo S, Cunningham SA, Kohner PC, et al. Comparison of a novel, rapid chromogenic biochemical assay, the Carba NP test, with the modified Hodge test for detection of carbapenemase-producing Gram-negative bacilli. J Clin Microbiol. 2013;51[9]:3097-3101)

### PDF Report

No

### Day(s) Performed

Monday through Friday

### Report Available

2 to 4 days

### Performing Laboratory Location

Mayo Clinic Laboratories - Rochester Main Campus

## Fees & Codes

### Fees

- Authorized users can sign in to [Test Prices](#) for detailed fee information.
- Clients without access to Test Prices can contact [Customer Service](#) 24 hours a day, seven days a week.
- Prospective clients should contact their account representative. For assistance, contact [Customer Service](#).

### Test Classification

This test was developed and its performance characteristics determined by Mayo Clinic in a manner consistent with CLIA requirements. It has not been cleared or approved by the US Food and Drug Administration.

### CPT Code Information

87182

### LOINC® Information

Test ID	Test Order Name	Order LOINC® Value
CARNP	Carbapenemase-Carba NP Test	74676-8

Result ID	Test Result Name	Result LOINC® Value
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## Test Definition: CARNP

Carbapenemase Detection-Carba NP Test,  
Varies

CARNP	Carbapenemase-Carba NP Test	74676-8
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