



# Test Definition: FNMEN

Neisseria Meningitidis IgG Vaccine Response

## Overview

### Method Name

Multi-Analyte Immunodetection (MAID)

### NY State Available

Yes

## Specimen

### Specimen Type

Serum

### Specimen Required

#### Container/Tube:

**Preferred:** Red top tube

**Acceptable:** Serum gel tube

**Specimen Volume:** 0.5 mL

**Collection Instructions:** Draw blood in a plain red-top tube(s), serum gel tube(s) is acceptable. Spin down and send 0.5 mL of serum refrigerated in a plastic vial.

**Note:** Serum gel tube is acceptable, but must pour off into a plastic vial.

### Specimen Minimum Volume

0.3 mL

### Reject Due To

### Specimen Stability Information

Specimen Type	Temperature	Time	Special Container
Serum	Refrigerated (preferred)	14 days	
	Ambient	7 days	
	Frozen	30 days	

## Clinical & Interpretive

### Reference Values

Reference Ranges (pre-vaccination):

Serogroup A	<4.0 ug/mL
Serogroup C	<5.0 ug/mL
Serogroup Y	<4.0 ug/mL
Serogroup W-135	<3.0 ug/mL

This assay measures serum IgG antibodies recognizing polysaccharide antigens from the four Neisseria meningitidis serogroups included in the licensed meningococcal vaccine. The meningococcal vaccine response is best evaluated by testing pre-vaccination and post-vaccination samples in parallel. A two-fold or greater increase for at least two sero-groups is expected when comparing post-vaccination to pre-vaccination results. N. meningitidis IgG levels peak approximately one month post-vaccination, but decline markedly by two years.

## Performance

### PDF Report

No

### Day(s) Performed

Tuesday

### Report Available

3 to 11 days

### Performing Laboratory Location

Quest Diagnostics

## 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 analytical performance characteristics have been determined by Quest Diagnostics. It has not been cleared or approved by FDA. This assay has been validated pursuant to the CLIA regulations and is use for clinical purposes.

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**CPT Code Information**

86317/x4

**LOINC® Information**

Test ID	Test Order Name	Order LOINC® Value
FNMEN	N. meningitidis IgG Vacc Response	Not Provided

Result ID	Test Result Name	Result LOINC® Value
Z0726	Serogroup A	42986-0
Z0532	Serogroup C	42985-2
Z0533	Serogroup Y	39618-4
Z0534	Serogroup W-135	39610-1