

## Overview

**Method Name**

Immunoassay

**NY State Available**

Yes

## Specimen

**Specimen Type**

Serum

**Specimen Required****Specimen Type:** Serum**Container/Tube:** Red**Specimen Volume:** 1 mL**Collection Instructions:** Draw blood in a plain, red-top tube, serum gel tube is acceptable. Spin down and send 1 mL of serum refrigerated in a plastic vial.**Specimen Minimum Volume**

0.5 mL

**Reject Due To**

Hemolysis	Mild OK; Gross reject
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**Specimen Stability Information**

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

## Clinical & Interpretive

**Clinical Information**

Discrimination between recent (primary) and past cytomegalovirus (CMV) infection can be an important tool in the clinical management of pregnant women. Although nearly all individuals with recent CMV infection are positive for CMV IgM, individuals with past CMV may also express CMV IgM due to long-term IgM persistence or viral reactivation; thus,

detection of CMV IgM is not a reliable indicator of recent CMV infection. Measurement of CMV IgG avidity can assist in discriminating recent from past CMV infection. A low avidity index is a reliable indicator of CMV infection within the previous 6 months, a high avidity index essentially excludes the possibility that infection occurred within the previous 4 months. Avidity index values should be considered within the context of other laboratory findings and clinical signs.

**Reference Values**

>0.70

**Interpretation**

<0.60 Low Avidity Index

0.60-0.70 Intermediate Avidity Index

>0.70 High Avidity Index

**Performance****PDF Report**

No

**Day(s) Performed**

Sunday, Tuesday through Friday

**Report Available**

4 to 10 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 performance characteristics have been determined by Quest Diagnostics Infectious Disease.](#) It has not been cleared or approved by FDA. This assay has been validated pursuant to the CLIA regulations and is used for clinical purposes.

**CPT Code Information**

86644

**LOINC® Information**

Test ID	Test Order Name	Order LOINC® Value
FCYTG	Cytomegalovirus IgG Avidity	52984-2

Result ID	Test Result Name	Result LOINC® Value
Z5329	CMV IgG Avidity Index	52984-2