

Hepatitis C testing

3-step procedure

Easy

- Simple 3-step procedure with point-of-care results
- Approved for fingerstick and venipuncture collection

Flexible

- Appropriate for testing in public health settings, physician offices, community health clinics, emergency rooms and outreach settings
- Ideal for batch testing

Accurate

- > 98% accurate
- Comparable to lab-accurate results
- First and only FDA-approved, CLIA-waived, rapid HCV test

Easy, flexible procedure and point-of-care results that help change testing for HCV







Feature	Benefit
5 μL specimen collection loops provided	Small sample size required
Built-in procedural control	Demonstrates assay validity
External controls available	Ensures proper performance of the test
Simple 3-step procedure	Minimal training required
20-minute test-read window	Conducive to batch testing and walk-away procedure

Time to results	20 minutes	
Test-read window	20-40 minutes	
Sample type	Fingerstick whole blood and venipuncture whole blood	
CLIA complexity	Waived	
External controls	Positive and negative (purchased separately)	
Internal control	IgG specific	
Storage requirements	2°C-30°C (36°F-86°F)	
Operating requirements	15°C-37°C (59°F-99°F)	
Method	Lateral flow	
Device shelf life	18 months from date of manufacture	
Kit size	25 count (Cat. No. 1001-0181) 100 count (Cat. No. 1001-0180)	
CPT code	86803 (CPT code)/G0472 (HCPCS)	
Additional items available	Visual Reference Panel (Cat. No. 1001-0343) Test Kit Controls (Cat. No. 1001-0182)	

Performance	Positive percent agreement	Negative percent agreement
Fingerstick whole blood	97.9%	98.5%
Venipuncture whole blood	99.5%	99.0%

Refer to the package insert for complete information on the OraQuick® HCV Rapid Antibody Test.



Clinical data has not been collected to demonstrate the performance of the OraQuick® HCV Rapid Antibody Test in individuals under 15 years of age or for pregnant women.

OraQuick

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