

OTC



COVID-19 Saliva Antigen Test

- ✓ Viral Detection
- ✓ Saliva based, Easy Self-Operation
- ✓ 15 minutes Fast Testing
- ✓ At-Home / Non-labs sites use

Sensitivity: >90%

Specificity: >99%

- Data compared to RT-PCR confirmed cases results.
- Third Party University Clinical Reports.
- COVID-19 Designated Hospitals Reports.



Also available to: CE-IVD / Philippine-FDA
ANVISA / Chile ISP Listed.

WhatsApp: +86-571-87763175 | Web: reOpenTest.com

Contact: sales@reopentest.com



reOpenTest

Leading In-Vitro Diagnostic Testing.



COVID-19 Antibody Test

- ✓ IgM/IgG Antibody Test
- ✓ 15 minutes Fast Testing
- ✓ Individual Home-Testing

Sensitivity: >95%

Specificity: >99%

- Data compared to RT-PCR confirmed cases results.
- Third Party University Clinical Reports.
- COVID-19 Designated Hospitals Reports.



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reOpenTest vs EUA Approved

SARS-CoV-2 IgM/IgG Antibody Test

reOpenTest vs EUA Approved brand

Brand	Method	Sensitivity	Specificity	Data From
Cellex (EUA)	Lateral Flow Immunofluorescent	96.8%	97.8%	IFU
Roche (EUA)	Lateral Flow Immunofluorescent	76.9%	99.8%	IFU
Healgen (EUA)	Lateral Flow Immunofluorescent	96.7%	97.0%	IFU
reOpenTest *	Lateral Flow Immunofluorescent	94.5%	99.3%	IFU/Hospital Clinic Data / China(n=453)
		95.9%	97.8%	Hospital Clinic Data / Greece(n=120)
LUNGENE (Non-EUA)	Lateral Flow Immunofluorescent	87%	98.9%	IFU

* with ANVISA, CE-IVD, Chile Whitelist.

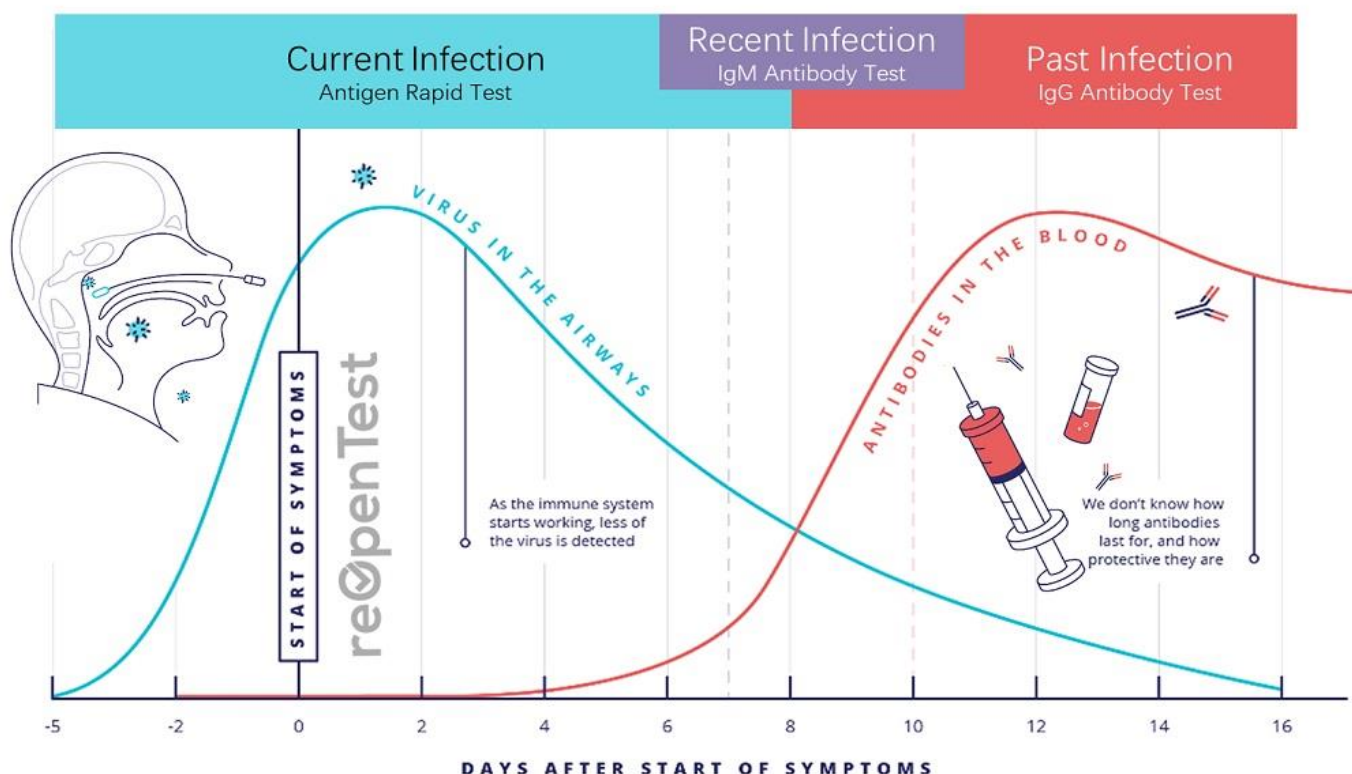
SARS-CoV-2 Antigen Rapid Test

reOpenTest vs EUA Approved brand

Brand	Method	Sensitivity	Specificity	Data From
Quidel Corporation	Lateral Flow Immunofluorescent	80%	100%	IFU
reOpenTest (Colloidal Gold)	Colloidal Gold	90%	98%	Korean University Medical Center Data(n=130)
Becton	Chromatographic Digital Immunoassay*	83%	100%	IFU
reOpenTest (FICA)	Fluorescence Immunochromatography*	88.6%	94.5%	3+ Hospitals Clinic Data (n=160)

* extra equipment required.

** FICA version shall view Test results under 365nm UV light.



All EUA Brands Data can be referred by FDA Reference:
<https://tinyurl.com/y8utovjj>
(Brands IFU included)

reOpenTest

What is the purpose of testing for COVID-19?

Viral Test Serology Test									
Group	reOpenTest			Person Quarantine Days					
	RT - PCR Antigen Test	Antibody Test		Early(1-3days)	Middle(3-14days)	Later(14+days)	Risk		
With Symptoms	Positive	Positive	Positive		✗ Active Infection		High +		
	Positive	Negative	Negative		✗ Active Infection		High +		
	Positive	Negative	Positive		✗ Active Infection		High +		
	Negative	Positive	Positive	Re - Test - Viral	✗ Active Infection	Re - Test - Viral	High -		
	Negative	Negative	Positive	N/A	Current Infection / Quarantine & Re - Test - Viral!		Middle		
	Negative	Negative	Negative		Quarantine & Re - Test - Viral!		Middle		
	Positive	Positive	Positive		✗ Active Infection		High +		
	Positive	Negative	Positive		✗ Active Infection		High +		
Without Symptoms, But At High Risk Of Exposure	Negative	Negative	Positive	Δ Past Infection / Quarantine to confirm		✓ Past Infection - Safe	Low +		
	Negative	Negative	Negative		Quarantine	Δ Safe	Middle		
	Negative	Positive	Positive	Re - Test - Viral	✗ Active Infection		High -		
	Negative	Negative	Positive	N/A	✓ Past Infection - Safe		Low		
Surveillance	Negative	Negative	Negative		No past infection		Low		

Certificates & Available Countries:

- ✓ Europe / CE-IVD
- ✓ Chile / ISP Listed
- ✓ France / <https://covid-19.sante.gouv.fr/tests>
- ✓ Bolivia / AGEMED

* Below is available to products not in this brand package.

- Brazil / ANVISA registered
- Philippine / FDA
- USA / FDA Listed

ATTACHED LIST

- Manufacturer Declaration – Antibody Test
- CE / NQA
- Clinical Report / For SARS-CoV-2 IgM/IgG Antibody Test
@ATTIKON University General Hospital, Athens, Greece.
- CE-IVD
- Clinical Report / For SARS-CoV-2 Antigen Rapid Test
@Yeungnam University Medical Center, Korea.



Carton Information

reOpenTest™

- Box: 260*120*65 mm
- Carton: 64*34*55 cm
- 1250 pcs/ctn
- GW:15kg