Sample Information

Patient ID: [redacted]
Lab Sample ID: [redacted]
Specimen Type: NP Swab
Collected: 07/22/2020
Received: 07/23/2020
Reported: 07/27/2020

Test(s) Requested

05700 SARS-CoV-2, qPCR

Tested For Result Reference Range
SARS-CoV-2, qPCR NEGATIVE N/A
SARS-CoV-2

Sample Comments

This test was developed and its performance characteristics determined by Aegis Sciences Corporation. This test has not been FDA cleared or approved. This test has been authorized by FDA under an Emergency Use Authorization for use by authorized laboratories. This test has been validated by Aegis Sciences Corporation in accordance with the FDA Guidance Document (Policy for Diagnostic Tests for Coronavirus Disease-2019 during the Public Health Emergency, FDA-2020-D-0987) issued on March 16, 2020. FDA independent review of this validation is pending. This test is authorized only for the detection of nucleic acid from SARS-CoV-2, not for any other viruses or pathogens. This test is only authorized for the duration of the declaration that circumstances exist justifying the authorization of the emergency use of in vitro diagnostic tests for detection and/or diagnosis of COVID-19 under section 584(b)(1) of the Act, 21 U.S.C. 360bbb-3(b)(1), unless the authorization is terminated or revoked.
TEST ORDERED: 94500-6~SARS coronavirus 2 RNA~139900~SARS-CoV-2, NAA

<table>
<thead>
<tr>
<th>RESULT</th>
<th>VALUE</th>
<th>UNITS</th>
<th>REFERENCE RANGES</th>
<th>ABNORMAL</th>
<th>RESULT STATUS</th>
</tr>
</thead>
<tbody>
<tr>
<td>139901 ~ SARS-CoV-2, NAA</td>
<td>LDTDET ~ Detected</td>
<td>Not Detect</td>
<td>Abnormal</td>
<td>Final</td>
<td></td>
</tr>
<tr>
<td>94500-6 ~ SARS coronavirus 2 RNA</td>
<td>260373001 ~ Detected</td>
<td>Not Detect</td>
<td>Abnormal</td>
<td>Final</td>
<td></td>
</tr>
</tbody>
</table>

This test was developed and its performance characteristics determined by LabCorp Laboratories. This test has not been FDA cleared or approved. This test has been authorized by FDA under an Emergency Use Authorization (EUA). This test has been validated in accordance with the FDA's Guidance Document (Policy for Diagnostics Testing in Laboratories Certified to Perform High Complexity Testing under CLIA prior to Emergency Use Authorization for Coronavirus Disease-2019 during the Public Health Emergency) issued on February 29th, 2020. FDA Independent review of this validation is pending. This test is only authorized for the duration of time the declaration that circumstances exist justifying the authorization of the emergency use of in vitro diagnostic tests for detection of SARS-CoV-2 virus and/or diagnosis of COVID-19 infection under section 564(b)(1) of the Act, 21 U.S.C. 360bbb-3(b)(1), unless the authorization is terminated or revoked sooner.
**Test Name**
SARS CoV 2 RNA (COVID 19), QUALITATIVE NAAT
SARS CoV 2 RNA

**Result:** NOT DETECTED

A Not Detected (negative) test result for this test means that SARS-CoV-2 RNA was not present in the specimen above the limit of detection. A negative result does not rule out the possibility of COVID-19 and should not be used as the sole basis for treatment or patient management decisions. If COVID-19 is still suspected, based on exposure history together with other clinical findings, re-testing should be considered in consultation with public health authorities. Laboratory test results should always be considered in the context of clinical observations and epidemiological data in making a final diagnosis and patient management decisions.

**REFERENCE RANGE:** NOT DETECTED

Please review the "Fact Sheets" and FDA authorized labeling available for health care providers and patients using the following websites:

This test has been authorized by the FDA under an Emergency Use Authorization (EUA) for use by Authorized laboratories.

Due to the current public health emergency, Quest Diagnostics is receiving a high volume of samples from a wide variety of swabs and media for COVID-19 testing. In order to serve patients during this public health crisis, samples from appropriate clinical sources are being tested. Negative test results derived from specimens received in non-commercially manufactured viral collection and transport media, or in media and sample collection kits not yet authorized by FDA for COVID-19 testing should be cautiously evaluated and the patient potentially subjected to extra precautions such as additional clinical monitoring, including collection of an additional specimen.

**Methodology:** Nucleic Acid Amplification Test (NAAT) includes PCR or TMA

Additional information about COVID-19 can be found at the Quest Diagnostics website.
Aegis Sciences Corporation
501 Great Circle Road
Nashville, TN 37228
(615) 255-2400
Lab Director: Matthew T. Hardison, PhD
CLIA#: 44D2062333

Laboratory Report

Client Information

Client: Contemporary Family Practice-
Report To: Kenneth Zeledorf
1560 W Lacey Blvd #101
Hanford, CA 93230

Sample Information

Patient ID: [Redacted]
Lab Sample ID: [Redacted]
Specimen Type: NP Swab
Collected: 09/10/2020
Received: 09/10/2020
Reported: 09/11/2020

Test(s) Requested

05700  SARS-CoV-2, qPCR

<table>
<thead>
<tr>
<th>Tested For</th>
<th>Result</th>
<th>Reference Range</th>
</tr>
</thead>
<tbody>
<tr>
<td>SARS-CoV-2, qPCR</td>
<td>NEGATIVE</td>
<td>N/A</td>
</tr>
<tr>
<td>SARS-CoV-2</td>
<td></td>
<td></td>
</tr>
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</table>

Sample Comments

This test was developed and its performance characteristics determined by Aegis Sciences Corporation. This test has not been FDA cleared or approved. This test has been authorized by FDA under an Emergency Use Authorization for use by authorized laboratories. This test has been validated by Aegis Sciences Corporation in accordance with the FDA Guidance Document (Policy for Diagnostic Tests for Coronavirus Disease-2019 during the Public Health Emergency, FDA-2020-D-0887) issued on March 16, 2020. This test is authorized only for the detection of nucleic acid from SARS-CoV-2, not for any other viruses or pathogens. This test is only authorized for the duration of the declaration that circumstances exist justifying the authorization of the emergency use of in vitro diagnostic tests for detection and/or diagnosis of COVID-19 under section 564(b)(1) of the Act, 21 U.S.C. 360bbb-3(b)(1), unless the authorization is terminated or revoked.
September 28, 2020

SUBJECT:

To Whom It May Concern:

This letter is to inform you of the rapid SARS-CoV-2 test that took place at Contemporary Family Practice on 9/28/2020. Your test results are Negative.

You may return to work. No quarantine is required at this time.

Should you need further assistance, please feel free to contact this office.

Best regards,

Hannah Alvarado
CFP | Medical Assistant
VA Chemistry/Hematology

Personal Health Record of [Redacted]

Information last updated in My HealtheVet on [Redacted]

Nasopharyngeal structure (body structure) Specimen
COVID-19 (VAPAHCS-ABBOTT) Test Details

Collected on 28 Sep 2020 @ 1930 at FRESNO VA MEDICAL CENTER

Test results slightly outside the reference range are not unusual. Your provider has reviewed your test results and will contact you for any important issues. If you have further questions, please do not hesitate to contact your primary care provider. View Comments to see the history of an amended test result.

Test Name: COVID-19 (VAPAHCS-ABBOTT)
Result: Not Detected
Units: 
Reference Range: Not Detected
Lab Test: Human Coronavirus RNA Qi PCR-VAMC LAB 1

Ordering Provider: FELIX, CYNTHIA
Ordering Location: FRESNO VA MEDICAL CENTER
Performing Location: PALO ALTO VA MEDICAL CENTER 3851 MIRANDA AVE, PALO ALTO, CA 94304-1207
Status: Final

Interpretation:

Comments: Results were obtained using Abbott RealTime SARS-CoV-2 EUA Assay. This assay is for in vitro diagnostic use only. The test has been validated at VAPA HCS but FDA independent review is pending. Optimal performance of this test requires appropriate specimen collection, storage, and transport to the test site as per CDC guidelines. Detection of SARS-CoV-2 RNA may be affected by sample collection methods, patient factors (e.g., presence of symptoms), and/or stage of infection. False-negative results may arise from degradation of the viral RNA during shipping/storage. The impacts of vaccines, antiviral therapeutics, antibiotics, chemotherapeutic or immunosuppressant drugs have not been evaluated. As with any molecular test, mutations within the target regions of Abbott RealTime SARS-CoV-2 assay could affect primer and/or probe binding, resulting in failure to detect the presence of virus. Performance has only been established with the specimen types listed in the Intended Use. Other specimen types have not been evaluated. Results should be interpreted by a trained professional in consultation with the patient's history and clinical signs and symptoms, and epidemiological risk factors. Laboratories are required to report all positive results to the appropriate public health authorities. Negative results do not preclude infection with the SARS-CoV-2 virus and should not be the sole basis of a patient treatment management or public health decision. Follow up testing should be performed according to the current CDC recommendations. This test was performed at Veterans Affairs Palo Alto Health Care System (CLIA# 05D05686226), 3801 Miranda Avenue, Palo Alto, CA 94304 Medical Director: Kristen Jensen, MD

The reference ranges may be different from what you have seen in the past. This is because of better technology and changes in testing standards. Ask your health care provider if you have questions.

VA laboratory results are available to you 36 hours after the laboratory analysis is finalized. Your VA provider may need additional time to review the results. NOTE: COVID-19 results are an exception to the hold period. COVID-19 results are available immediately after receipt by VA.

This information is a copy of your VA medical record, and may not reflect the most recent changes. If you think the information is not accurate, please contact your VA facility directly.
01 Oct 2020 @ 1338 ET

EDWARDS, ANAMARIA (Chu, P - Fresno VA)

General: General Inquiry

Department of Veterans Affairs
2615 E. Clinton Ave
Fresno, CA 93703
Telephone Advice: 559-228-6933
Toll Free 1-888-826-2838

VA Central California Health Care System
2615 E. Clinton Ave, Fresno CA 93703
Phone: (559) 225-6100

OCT 01, 2020

To whom it may concern:

[Redacted] is a patient at the VA Central California HCS, Fresno.

Patient tested Negative for COVID-19,
Test Date: September 28, 2020

[Redacted] may return to work on [Redacted]

Thank you,
Cynthia Felix NP-C
VA Primary Care
VACCHS

Ana-Maria Edwards, LVN
LVN, Primary Care

This message may not be from the person you initially contacted.
It may have been reassigned in order to efficiently address your question.
Urgent Care of Hanford
1028 N. Douty St., Suite 1
Hanford, CA 93230
Phone: (559) 530-2526
Fax: (559) 410-8215

Patient Name: [redacted]
Birth Date: [redacted]
Sex: [redacted]

Rapid Test: SARS2COVID-19
Lot: [redacted]
Exp: 2/14/21

Results:
☑ Negative

Symptoms: (Choose all that apply)
☑ Cough  ☑ Fever/Chills  ☑ Shortness of breath  ☑ Nausea/Vomiting
☑ Sore Throat  ☑ Headache  ☑ Fatigue  ☑ Muscle or Body aches
☑ Diarrhea  ☑ Congestion or Runny nose  ☑ Loss of Taste or Smell
☑ Unknown  ☑ Positive Exposure  ☑ Traveled

Remarks: ____________________________

Provider’s Name (Print): [redacted]
Provider’s Signature: [redacted]

HCP Office Stamp: [image]
COVID RESULTS | WORK NOTE

10/23/2020 11:02:02 PM AEGIS SCIENCES CORPORATION

Laboratory Report

Client Information
Client: Contemporary Family Practice
Report To: Kenneth Zelisko
1560 W Lacey Blvd #101
Harford, CA 93230

Sample Information
Patient ID:
Lab Sample ID:
Specimen Type: NP Swab
Collected: 10/21/2020
Received: 10/22/2020
Reported: 10/23/2020

Test(s) Requested
05700 SARS-CoV-2, qPCR

<table>
<thead>
<tr>
<th>Tested For</th>
<th>Result</th>
<th>Reference Range</th>
</tr>
</thead>
<tbody>
<tr>
<td>SARS-CoV-2, qPCR</td>
<td>NEGATIVE</td>
<td>N/A</td>
</tr>
</tbody>
</table>

Sample Comments
The Thermo Fisher Scientific, Inc. TaPath COVID-19 Omni Kit used to perform this test has been authorized by FDA under an Emergency Use Authorization for use by authorized laboratories. This test has not been cleared or approved by the FDA. This test is authorized only for the detection of nucleic acid from SARS-CoV-2, not for any other viruses or pathogens. This test is only authorized for the duration of the declaration that circumstances exist justifying the authorization of emergency use of the test. It is not intended to be used as a diagnostic test for detection and/or diagnosis of COVID-19 as defined in section 581 of the Act, 21 U.S.C. 360bb-360bb-360bb, unless the authorization is terminated or revised.
Your COVID-19 test result

NEGATIVE

A negative result for this test means that SARS-CoV-2 RNA (the cause of COVID-19) was not detected in the collection sample.

What does it mean to have a negative test result?

A negative test result does not completely rule out being infected with COVID-19.

If you test negative for COVID-19, this means the virus was not detected at the time your specimen was collected. It is still possible that you were very early in your infection at the time of your specimen collection and that you could test positive later.

Also, you could be exposed later and still develop the illness. For all these reasons, it is important to follow CDC guidance, including but not limited to frequent hand washing, social distancing, wearing a face covering, covering coughs and sneezes, monitoring symptoms, and cleaning and disinfectant of frequently touched surfaces - even after a negative test result.

Test information

Patient's name

Patient's date of birth

Test type
SARS-COV-2 RNA, QL, RT PCR (COVID-19)

Provider
MERCADO INNA

Collection date
November 4, 2020 at 3:10 PM ET

Collection location
1405 HERNDON AVENUE, CLOVIS, CA 93611

MinuteClinic contact information

Customer Service: (886) 389-2727
COVID | RAPID TEST RESULTS

Today’s Date: 11/2/2020
Patient Name: [Redacted]
DOB: [Redacted]

To Whom It May Concern:

This letter serves to inform of the above patient's rapid test results for SARS-CoV-2 (COVID-19). The test was conducted at Contemporary Family Practice. The results are the following:

☐ POSITIVE  ☒ NEGATIVE

If POSITIVE, please see attached the CDC quarantine guidelines. Also, the Kings County Health Department will be in contact with further instructions.

If NEGATIVE, please be advised that the patient above has been seen by the medical provider of CFP. The patient has been assessed and appears to be clear of COVID-19. The patient is released to return to work | school.

Should you need further assistance, please feel free to contact this office.

Best regards,

CFP | Administration
Sample Information

Patient ID: [redacted]
Lab Sample ID: [redacted]
Specimen Type: Nasal Swab
Collected: 11/04/2020
Received: 11/05/2020
Reported: 11/05/2020

Test(s) Requested

05700 SARS-CoV-2, qPCR

Tested For

<table>
<thead>
<tr>
<th>SARS-CoV-2, qPCR</th>
<th>Result</th>
<th>Reference Range</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>NEGATIVE</td>
<td>N/A</td>
</tr>
</tbody>
</table>

CERTIFICATE FOR RETURN TO SCHOOL OR WORK

Patient Name: [redacted]

Has been under my care 11/04/20 and is able to return

Limitations/Remarks: does not need to quarantine

Date: 11/09/20

Provider Signature: [redacted]

MD/PA/NP
Visalia Medical Clinic

5400 W. Hillsdale Ave
Visalia, CA 93291
(559) 738-7555
Fax: (559) 738-7527

Exam Date: 11/14/2020 1:27:00PM
Referring Phys: Castellano, Juanita
Ordering Phys: Castellano, Juanita

Result Report

<table>
<thead>
<tr>
<th>Test</th>
<th>Result</th>
<th>Units</th>
<th>Flag</th>
<th>Reference Range</th>
</tr>
</thead>
</table>

Patient Information Sheet

Covid Comments
Positive results indicate the presence of viral antigens, but clinical correlation with patient history and other diagnostic information is necessary to determine infection status.

Covid Comments
Positive results do not rule out bacterial infection, or co-infection, with other viruses.

Covid Comments
Negative results do not rule out COVID-19 and should not be used as the sole basis for treatment, or patient management decisions, including infection control decisions.

Covid Comments
Negative results should be considered in the context of a patient's recent exposures, history and the presence of clinical signs and symptoms consistent with COVID-19.
Date: 11/19/20

Patient Name: [Redacted]
DOB: [Redacted]

☐ Patient is currently being tested for COVID-19. While results are pending, patient is to be Quarantined for 14 days.

☒ Patient has tested negative for COVID-19 and is able to return to work.

Provider signature

Date: 11/19/20
# 2019 Novel Coronavirus (COVID-19). NAA 139900

<table>
<thead>
<tr>
<th>NAME</th>
<th>VALUE</th>
<th>REFERENCE RANGE</th>
<th>LAB</th>
</tr>
</thead>
<tbody>
<tr>
<td>SARS-CoV-2, NAA</td>
<td>Not Detected</td>
<td>Not Detected</td>
<td>SEQCO</td>
</tr>
</tbody>
</table>

- This nucleic acid amplification test was developed and its performance
- characteristics determined by LabCorp Laboratories. Nucleic acid
- amplification tests include PCR and TMA. This test has not been FDA
- cleared or approved. This test has been authorized by FDA under an
- Emergency Use Authorization (EUA). This test is only authorized for
- the duration of time the declaration that circumstances exist

- justifying the authorization of the emergency use of in vitro
- diagnostic tests for detection of SARS-CoV-2 virus and/or diagnosis
- of COVID-19 infection under section 564(b)(1) of the Act, 21 U.S.C.
- 350bbb-3(b) (1), unless the authorization is terminated or revoked
- sooner.

- When diagnostic testing is negative, the possibility of a false
- negative result should be considered in the context of a patient’s
- recent exposures and the presence of clinical signs and symptoms
- consistent with COVID-19. An individual without symptoms of COVID-19
- and who is not shedding SARS-CoV-2 virus would expect to have a
- negative (not detected) result in this assay.

**PERFORMING LAB: Integrated Genetics Sequenom, 3595 John Hopkins Court, San Diego, CA 921211121, Phone - 8778217266, Director - MDPhD Cacheris**
To: [Redacted]
DOB: [Redacted]

The result of your COVID-19 Nasal RT-PCR test that was administered on 11/19/2020 is Positive.

Your Public Health Department has been told of this result as is required by law.

**What is Self-Isolation?**
When you have COVID-19, it is important to self-isolate (or stay at home and separate yourself from others). You should stay in a separate room away from other people in your household to keep them safe and prevent them from getting ill.

**How can you care for yourself at home?**
There is no specific treatment for the virus that causes COVID-19. Most people with COVID-19 will have mild illness and can get better with proper home care without the need to see a medical provider. Here are steps that you can take to help you get better:
- Rest & drink plenty of fluids
- Take acetaminophen (Tylenol®) as directed to reduce fever and pain (talk to your child’s medical provider before giving children younger than 2 years old over-the-counter cold medications)

Remember: over-the-counter medicines do not cure the illness and do not stop you from spreading germs.

**What are the symptoms of COVID-19?**
Common symptoms of COVID-19 include: fever or chills, cough, shortness of breath or difficulty breathing, fatigue, muscle or body pain, headache, congestion or runny nose, sore throat, nausea, vomiting, diarrhea, or new loss of taste or smell. Please note that this list does not include all possible symptoms.

**When should you get medical care?**
Write down when your symptoms started and continue to monitor your health. Get medical care if your symptoms get worse, especially if you are at a higher risk of serious illness. People who are 65 years and older or have a health problem such as a chronic disease (e.g., diabetes) or a weak immune system are considered high risk. Call ahead before visiting your medical provider, you may be able to get advice by phone.

**Call 911 if you start to have emergency warning signs**
If you have an emergency warning sign such as: severe difficulty breathing and cannot catch your breath, persistent pain or pressure in the chest, new confusion, and/or bluish lips or face, or the inability to wake or stay awake, seek emergency medical care immediately.

If you call 911, tell the dispatch personnel that you have COVID-19. If you need help finding healthcare, call your county’s Information Line or 2-1-1. You can call 24 hours a day, 7 days a week. For more information, visit http://www.211.org/.

**What steps can you take to protect others in your home and community?**
1. Stay home except to get medical care
To: [Redacted]
DOB: [Redacted]

The result of your COVID-19 Nasal RT-PCR test that was administered on 11/25/2020 is Negative.

Your Public Health Department has been notified of this result as required by law.

If you feel well and do not have a fever or cough, you may go about your normal activity abiding by your state and local health department recommendation in regards to ‘Stay at Home’ or ‘Shelter in Place’ orders.

Regardless of your result, it is still important for you to take the following actions for care at home:

- Wash hands thoroughly and frequently with soap and water for at least 20 seconds. If soap and water are not readily available, use an alcohol-based hand sanitizer that contains at least 60% alcohol. Avoid touching eyes, nose, and mouth with unwashed hands.
- To cough or sneeze, cover mouth and nose with a tissue and immediately dispose of it. If no tissue is available, cough/sneeze into the inside of elbow, not hands. Wash hands thoroughly with soap and water immediately following.
- Avoid sharing eating utensils, towels, linens, clothes or other items. Wash items thoroughly with soap and water.
- Practice physical distancing. This includes avoiding crowded public places where close contact with others may occur. Maintain distance of 6 feet (2 meters) from others, when possible. Avoid contact with people who are sick.
- Use cloth face coverings when you are:
  - Inside of, or in line to enter, any indoor public place
  - Using public transportation, a taxi, private car service, or ride-sharing vehicle
  - Outside and unable to physical distance 6 feet (2 meters) away from others
- Use household detergent and water to clean frequently touched surfaces such as tabletops, light switches, handles, phones, keyboards, toilets, faucets, and doorknobs. Dirty surfaces should be cleaned, then disinfected using common household disinfectant.
- Open windows and use a fan (if possible) in shared spaces for good airflow.
- Be alert for symptoms. Watch for fever, cough, shortness of breath, or other symptoms of COVID-19. Take your temperature if symptoms develop. Follow CDC guidance if symptoms develop.

If you were recently exposed to someone with COVID-19, you should still stay at home and self-quarantine (separate yourself from others) for 14 days, starting from the most recent day that you were possibly exposed to COVID-19. You should continue to monitor for symptoms for up to 14 days after your last possible exposure. If you have additional questions please contact your primary provider. It has been our great privilege in assisting you in your healthcare.
**Laboratory Report**

**Client Information**

Client: Contemporary Family Practice-
Report To: Kenneth Zeisdorf
1560 W Lacey Blvd #101
Hanford, CA 93230

**Sample Information**

Patient ID: [Redacted]
Lab Sample ID: [Redacted]
Specimen Type: NP Swab
Collected: 11/20/2020
Received: 11/21/2020
Reported: 11/22/2020

**Test(s) Requested**

<table>
<thead>
<tr>
<th>Test Code</th>
<th>Description</th>
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</thead>
<tbody>
<tr>
<td>05700</td>
<td>SARS-CoV-2, qPCR</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Tested For</th>
<th>Result</th>
<th>Reference Range</th>
</tr>
</thead>
<tbody>
<tr>
<td>SARS-CoV-2, qPCR</td>
<td>NEGFATIVE</td>
<td>N/A</td>
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</tbody>
</table>

**Sample Comments**

The Thermo Fisher Scientific, Inc. TaqPath COVID-19 Combo Kit used to perform this test has been authorized by FDA under an Emergency Use Authorization for use by authorized laboratories. The test has not been otherwise FDA cleared or approved. This test is authorized only for the detection of nucleic acid from SARS-CoV-2, not for any other viruses or pathogens. This test is only authorized for the duration of the declaration that circumstances exist justifying the authorization of the emergency use of in vitro diagnostic tests for detection and/or diagnosis of COVID-19 under section 564(b)(1) of the Act, 21 U.S.C. 360bbb-3(b)(1), unless the authorization is terminated or revoked.

Visit [https://www.aegislabs.com/our-services/covid19-testing](https://www.aegislabs.com/our-services/covid19-testing) for COVID-19 testing resources.
NEGATIVE for SARS-CoV-2.

This means that SARS-CoV-2 (the virus that causes COVID-19) was not detected in the patient's sample collected on Nov 22, 2020.

- A negative result does not rule out a SARS-CoV-2 infection, irrespective of the sample type. If clinical suspicions exist, no clinical action and management should be based solely on a negative test result. Clinical symptoms, patient history, and epidemiological information must be considered when interpreting a negative result.
- Variability in collection technique can reduce the sensitivity of the test, and SARS-CoV-2 may not be detected in early stages of infection.
- False negatives are possible.
- If this patient is experiencing symptoms, consider collecting a new sample for COVID-19 testing or testing for other respiratory viruses.
- Collection of multiple specimens may be necessary to detect the SARS-CoV-2 virus.

DETAILS

<table>
<thead>
<tr>
<th>Test</th>
<th>Results</th>
</tr>
</thead>
<tbody>
<tr>
<td>SARS-CoV-2 amplification test</td>
<td>NEGATIVE (Not Detected)</td>
</tr>
<tr>
<td>SARS-CoV-2 Target ORF1ab gene</td>
<td>Cycle Threshold (Ct) Value¹</td>
</tr>
<tr>
<td>N gene</td>
<td>0</td>
</tr>
</tbody>
</table>

To learn more about the technical details of this test, please see the test methodology and limitations section.

HELPFUL INFORMATION FOR THE PATIENT

Because it is possible for this test to give a false negative in some people with COVID-19, if you have symptoms of illness (such as fever, cough, and/or shortness of breath), you should discuss your symptoms and your test results with your doctor, who can decide how to care for you.
Point of Care Testing

<table>
<thead>
<tr>
<th>Procedure</th>
<th>Units</th>
<th>Reference Range</th>
</tr>
</thead>
<tbody>
<tr>
<td>POC COVID-19 Quidel</td>
<td>negative</td>
<td>[negative]</td>
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</tbody>
</table>

Interpretive Data

**POC COVID-19 Quidel**
Positive results indicate the presence of viral antigens, but clinical correlation with patient history and other diagnostic information is necessary to determine infection status. Positive results do not rule out bacterial infection, or co-infection, with other viruses. Negative results do not rule out COVID-19 and should not be used as the sole basis for treatment, or patient management decisions, including infection control decisions. Negative results should be considered in the context of a patient’s recent exposures, history and the presence of clinical signs and symptoms consistent with COVID-19. Negative results from patients with symptom onset beyond five days should be considered for confirmation testing with a molecular assay.

For emergency use Facts Sheet, please copy the links below into an external browser.

Healthcare Provider Fact Sheet:


Patient Fact Sheet:


LEGEND: c=Corrected, @=Abnormal, C=Critical, L=Low, H=High, f=Result Comment, i=Interp Data, *=Performing Lab
COVID | RAPID TEST RESULTS

Today's Date: 12/01/2020
Patient Name: [redacted]
DOB: [redacted]

To Whom It May Concern:

This letter serves to inform of the above patient's rapid test results for SARS-CoV-2 (COVID-19). The test was conducted at Contemporary Family Practice (CFP). The results are the following:

☑ POSITIVE ☐ NEGATIVE

If POSITIVE, please read the CDC quarantine guidelines by visiting their website at https://www.cdc.gov/coronavirus/2019-ncov/if-you-are-sick/quarantine.html. Also, the Kings County Health Department will be in contact with further instructions.

If NEGATIVE, please be advised that the patient above has been seen by the medical provider of CFP. The patient has been assessed and appears to be clear of COVID-19. The patient is released to return to work | school.

Should you need further assistance, please feel free to contact this office.

Best regards,
CONTEMPORARY FAMILY PRACTICE
KENNETH E. ZELSDORF, FNP-C
1500 W. LACEY BLVD., SUITE 101
HANFORD, CA 93230
(559) 772-8285

CFP | Administration
To: [Redacted]
DOB: [Redacted]

The result of your COVID-19 Nasal RT-PCR test that was administered on 12/01/2020 is Negative.

Your Public Health Department has been notified of this result as required by law.

If you feel well and do not have a fever or cough, you may go about your normal activity abiding by your state and local health department recommendation in regards to 'Stay at Home' or 'Shelter in Place' orders.

Regardless of your result, it is still important for you to take the following actions for care at home:

- Wash hands thoroughly and frequently with soap and water for at least 20 seconds. If soap and water are not readily available, use an alcohol based hand sanitizer that contains at least 60% alcohol. Avoid touching eyes, nose, and mouth with unwashed hands.
- To cough or sneeze, cover mouth and nose with a tissue and immediately dispose of it. If no tissue is available, cough/sneeze into the inside of elbow, not hands. Wash hands thoroughly with soap and water immediately following.
- Avoid sharing eating utensils, towels, linens, clothes or other items. Wash items thoroughly with soap and water.
- Practice physical distancing. This includes avoiding crowded public places where close contact with others may occur. Maintain distance of 6 feet (2 meters) from others, when possible. Avoid contact with people who are sick.
- Use cloth face coverings when you are:
  - Inside of, or in line to enter, any indoor public place
  - Using public transportation, a taxi, private car service, or ride-sharing vehicle
  - Outside and unable to physical distance 6 feet (2 meters) away from others
- Use household detergent and water to clean frequently touched surfaces such as tabletops, light switches, handles, phones, keyboards, toilets, faucets, and doorknobs. Dirty surfaces should be cleaned, then disinfected using common household disinfectant.
- Open windows and use a fan (if possible) in shared spaces for good airflow.
- Be alert for symptoms. Watch for fever, cough, shortness of breath, or other symptoms of COVID-19. Take your temperature if symptoms develop. Follow CDC guidance if symptoms develop.

If you were recently exposed to someone with COVID-19, you should still stay at home and self-quarantine (separate yourself from others) for 14 days, starting from the most recent day that you were possibly exposed to COVID-19. You should continue to monitor for symptoms for up to 14 days after your last possible exposure. If you have additional questions please contact your primary provider. It has been our great privilege in assisting you in your healthcare.
COVID | RAPID TEST RESULTS

Today's Date: 10/16/20
Patient Name: [Redacted]
DOB: [Redacted]

To Whom It May Concern:

This letter serves to inform of the above patient's rapid test results for SARS-CoV-2 (COVID-19). The test was conducted at Contemporary Family Practice. The results are the following:

☐ POSITIVE ☒ NEGATIVE

If POSITIVE, please see attached the CDC quarantine guidelines. Also, the Kings County Health Department will be in contact with further instructions.

If NEGATIVE, you may return to work. No quarantine is required at this time.

Should you need further assistance, please feel free to contact this office.

Best regards,

CONTEMPORARY FAMILY PRACTICE
KENNETH E. ZELSDORF, FNP-C
1560 W. LACEY BLVD., SUITE 101
HANFORD, CA 93230
(559) 772-8285

CFP | Administration
<table>
<thead>
<tr>
<th>Laboratory Flowsheet</th>
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<td>14:47 PDT</td>
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**Immunology Results**

SARS-CoV-2 by PCR * NOT DETECTED
SARS-CoV-2 by PCR: DETECTED

Date / Time: October 28 2020 03:50 PDT

SARS-CoV-2 by PCR: DETECTED

Contributor System: QUEST_RLN

Status: Auth (Verified)

1.) (Medium Importance) Result Comment by Contributor_system, QUEST_RLN on October 31 2020 19:41 PDT

A Detected result is considered a positive test result for COVID-19. This indicates that RNA from SARS-CoV-2 (formerly 2019-nCoV) was detected, and the patient is infected with the virus and presumed to be contagious. If requested by public health authority, specimen will be sent for additional testing.

REFERENCE RANGE: NOT DETECTED

Please review the "Fact Sheets" and FDA authorized labeling available for health care providers and patients using the following websites:

This test has been authorized by the FDA under an Emergency Use Authorization (EUA) for use by authorized laboratories.

Due to the current public health emergency, Quest Diagnostics is receiving a high volume of samples from a wide variety of swabs and media for COVID-19 testing. In order to serve patients during this public health crisis, samples from appropriate clinical sources are being tested. Negative test results derived from specimens received in non-commercially manufactured viral collection and transport media, or in media and sample collection kits not yet authorized by FDA for COVID-19 testing should be cautiously evaluated and the patient potentially subjected to extra precautions such as additional clinical monitoring, including collection of an additional specimen.

Methodology: Nucleic Acid Amplification Test (NAAT) includes RT-PCR or TMA

Additional information about COVID-19 can be