Laboratory Results
AZOVA Testing Network Partner

Patient	Cale Shapera
DOB	05/30/1988
Gender	Male
Passport/Citizen ID	HK181590
Ordering Provider	AZOVA Test Observation Proctor
Sample Type	Nasal swab
Collected	04/06/2022 06:21 PM (GMT -07:00)
Final report date	04/06/2022 06:53 PM (GMT -07:00)

AZOVA Patient Services
144 S Main St
Alpine, Utah, 84004
United States

Results

Test Type	Result Unit	Result Range
BTNX Rapid Antigen Test	Negative	[Positive, Negative, Invalid]

The Rapid ResponseTM COVID-19 Antigen Rapid Test Device is an in vitro immunochromatographic assay for the direct and qualitative detection of SARS-CoV-2 viral nucleoprotein antigens from nasal and nasopharyngeal secretions from individuals suspected of COVID-19 within 6 days of symptom onset and and from individuals without symptoms or other epidemiological reasons to suspect COVID-19 infection, when tested twice over two (or three) days with at least 24 hours (and no more than 36 hours) between tests. This test is authorized for use at the Point of Care i.e., in patient care settings

Interpretation: Only one colored band appears, in the control region (C). No apparent colored band appears in the test region (T).

Patient fact sheet: https://www.btnx.com/files/COV-19C25/1110036511-COV-19C25%20(NP-AN)-IFU-V1.2-CAN.pdf

Ordering Provider:
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This test was performed by an authorized AZOVA Testing Network partner.