

| Client: Chicago-Eden                                |                              |                                |  |  |
|---|------------------------------|--------------------------------|--|--|
| Client Code: OUT                                    | Supervised By: Manager on du | Supervised By: Manager on duty |  |  |
| Address: 2358 Hassell Rd. Hoffman Estates, IL 60169 | Phone: 847-636-2667          | Fax:                           |  |  |

| Patient: Howe, John  |                 | Order ID: MK-1642732894316 |                                   |                                  |  |
|--|-----------------|----------------------------|-----------------------------------|----------------------------------|--|
| ID:  | DOB: 12/29/1956 | Gender: Male               | Collected:<br>01/20/2022 08:40 PM | Reported:<br>01/21/2022 19:19 PM |  |
| Unit:  |                 | Room:                      | 01/20/2022 00:40 1 14             | 01/21/2022 13:13 110             |  |
| Phone: (773) 472-1020                                      |                 | Specimen Type: NASAL SWAB  |                                   |                                  |  |
| Address: 940 w Gordon terrace , Chicago , Illinois , 60613 |                 | Physicians:                |                                   |                                  |  |

| Test                                  | Result   | Flag   | Unit | Ref Range |
|---------------------------------------|----------|--------|------|-----------|
| COVID- 19 SARS -COV- 2, RAPID ANTIGEN | NEGATIVE | NORMAL |      | NEGATIVE  |

This test is performed using COVID-19 Antigen Test kit by Access Bio for the Detection of COVID-19 (SARS-CoV-2) RNA. The test has been received Emergency Use Authorization (EUA) by US Food and Drug Administration. The test performance characteristics were determined by the M K LABS INC, Hoffman Estates, Illinois. The laboratory is certified under CLIA Wavier to perform testing on human clinical specimens.

## CLIA ID# 14D22 38829

This test was validated, and its performance determined by M K LABS INC. It has not been cleared or approved by FDA. Since FDA clearance is not required for clinical use of this laboratory developed test, this laboratory has established and validated the test's accuracy and precision, pursuant to the requirements of CLIA'88. Presumptive tests are indicated as (EUA). All other tests are confirmatory LCMS tests.

Lab Tech: SUPERVISOR Report Date: 01/21/2022 19:19 PM Lab Director: Samira Syed