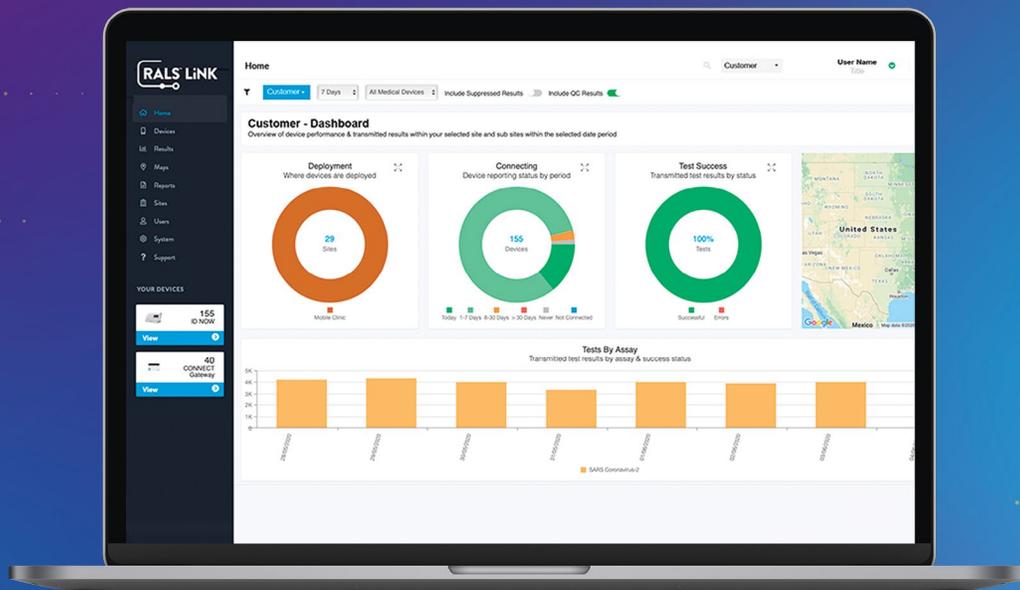




RALS™ LiNK FOR ID NOW™ CONNECTIVITY SOLUTION



The RALS™ LiNK dashboard can be personalized to show actionable data and analytics.



CONNECT Universal Gateway connects to ID NOW™.

Simple and Secure Results Reporting

Track, consolidate and report ID NOW™ COVID-19, Influenza A & B 2, Strep A 2 and RSV test results from multiple decentralized instruments and locations using RALS™ LiNK.

For COVID-19 testing, RALS LiNK automates many of the steps needed to report data as required by the [Coronavirus Aid, Relief, and Economic Security \(CARES\) Act, Section 18115,*](#) without the infrastructure burden.



RALS™ LiNK FOR ID NOW™ CONNECTIVITY SOLUTION

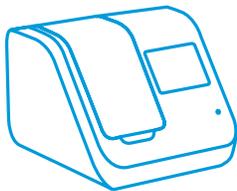
OUR RELIABLE DATA MANAGEMENT SYSTEM ALLOWS YOU TO:

- Consolidate results from multiple ID NOW™ instruments, streamlining the process for timely, customizable laboratory data reporting.
- Capture near real-time patient test results.
- Ensure tests are appropriately captured for processing and billing.
- Remotely monitor instrument location, status and performance.

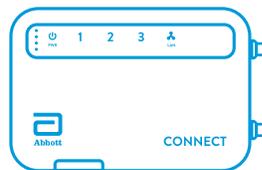


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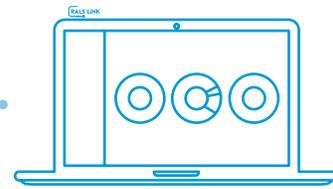
HOW RALS LiNK CONNECTIVITY WORKS WITH ID NOW:



A test is performed on an ID NOW™ instrument in a remote location.



ID NOW connects to a CONNECT Universal Gateway using an Ethernet® cable.



The CONNECT Universal Gateway encrypts and transmits data via a secure cellular connection from ID NOW to an online dashboard and/or existing EMR/LIS, if applicable.

RALS LiNK RECEIVES THESE DATA POINTS FROM ID NOW:

- Cartridge type
- Sample ID (order ID)
- Device serial number
- Result date
- Result summary
- Software version
- Operator ID
- Results type (patient/QC)

ID NOW ASSAYS SUPPORTED BY RALS LiNK:

- COVID-19*
- Influenza A & B 2
- Strep A 2
- RSV

FOR MORE INFORMATION, CONTACT YOUR LOCAL ABBOTT REPRESENTATIVE OR VISIT [ABBOTT.COM/POCT](https://www.abbott.com/poctralslink) OR [RALS.COM/RALSliNK](https://www.rals.com/ralslink)

*DATA CAPTURED FOR COVID-19 TESTS HELP FACILITATE COMPLIANCE WITH CARES ACT REPORTING REQUIREMENTS. FOR CARES ACT REPORTING REQUIREMENTS, VISIT [HHS.GOV/CORONAVIRUS](https://www.hhs.gov/coronavirus)

The ID NOW™ COVID-19 EUA has not been FDA cleared or approved. It has been authorized by the FDA under an emergency use authorization for use by authorized laboratories and patient care settings. The test has been authorized only for the detection of nucleic acid from SARS-CoV-2, not for any other viruses or pathogens, and is only authorized for the duration of the declaration that circumstances exist justifying the authorization of 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 sooner.

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