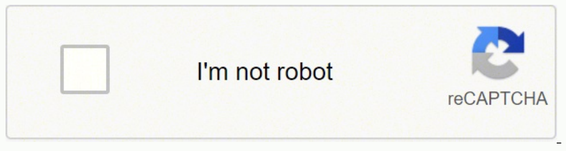
Abbott alinity specifications



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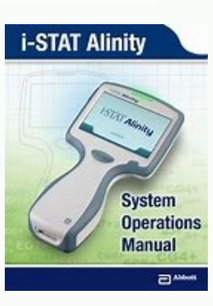
Abbott alinity specifications













Abbott alinity m specifications.

Profile: Researcher Patient Clinician Lab Professional Default Biomedical researchers, academics, and others engaged in medical research Patients and caregivers Doctors, nurses or any other medical professionals directly involved in the diagnosis and treatment of patients Laboratorians or any other individuals directly involved in the medical laboratory setting Please select your persona Forgot password Edit Password Your session is about to expire. Click Ok to extend or Cancel to log out. Your session has expired. Please log in again. Published since 1987, America’s Civil War strives to deliver to our readers the best articles on the most formativ... The American experience comes alive through thoroughly researched stories, outstanding photography and artwork. Aviation History magazine is an authoritative, in-depth history of world aviation from its origins to the... Biographies, battles, eyewitness accounts & period photos of America’s greatest internal conflict. Civil War T... MHQ: The Quarterly Journal of Military History takes you on an exciting journey to the world's greatest b... Military History magazine presents land, naval & air warfare from ancient times to the late 20th cent... Vietnam magazine presents the full & true stories from America’s most controversial & divisive wa... Wild West magazine presents the great American frontier from its beginnings to today. America’s western f... World War II magazine covers every aspect of history's greatest modern conflict with vivid, revealing, an... © 2021 World History Group, LLC. All Rights Reserved. Abbott has recalled two of its Alinity COVID-19 test kits after finding they can yield false positive results. Abbott is recalling its Alinity m SARS-CoV-2 AMP kit and Alinity m Resp-4-Plex AMP kit, which use swab samples taken from the nose, mouth and throat to test for COVID-19, according to an Oct. 14 news release. The tests are administered in healthcare settings and must be sent to a lab for results. The FDA warned COVID-19 test labs last month that the test kits may be faulty. The tests require lab equipment that uses software to automate the mixing of the chemicals. Abbott said an error in the software may cause a test sample in one well of the machine's tray to overflow into another test sample, which could cause a false positive result. Abbott said it is working on fixing the software issue at each lab site where the tests are processed. The FDA has classified the recall as Class I, the most serious type of recall, meaning use of the device could cause serious injury or death. Abbott said it has not received any reports of adverse events related to the test. Read the full news release here. The SARS-CoV-2 IgG assay is a chemiluminescent microparticle immunoassay (CMIA) intended for the qualitative detection of IgG antibodies to SARS-CoV-2 in human serum, serum separator tube and plasma (ACD, CPD, CPDA-1, dipotassium EDTA, tripotassium EDTA, lithium heparin, lithium heparin separator tube, sodium citrate, sodium heparin). The SARS-CoV-2 IgG assay is intended for use as an aid in identifying individuals with an adaptive immune response to SARS-CoV-2, indicating recent or prior infection. At this time, it is unknown for how long antibodies persist following infection and if the presence of antibodies confers protective immunity. The SARS-CoV-2 IgG assay should not be used to diagnose acute SARS-CoV-2 infection.9,10 Download SARS-CoV-2 IgG EUA Resources Fact Sheet for Recipients: Understanding the Abbott SARS-CoV-2 IgG EUA Assay Results Fact Sheet for Healthcare Providers: Interpreting Abbott SARS-CoV-2 IgG EUA Assay Results CLINICAL PERFORMANCE 9,10 The positive agreement between the ARCHITECT i2000SR and the Alinity i was 100% and the negative agreement was 99.00%. a Five specimens from 1 immunocompromised patient were excluded from the study. Refer to the LIMITATIONS OF THE PROCEDURE section of the package insert for further information. When the results from these specimens were included, the PPA at ≥ 14 days post-symptom onset was 96.77% (95% CI: 90.86, 99.33). Healthcare challenges such as increasing consolidation, rising costs, reimbursement cuts and staff shortages are having a direct impact on labs. At the same time, molecular labs want to enhance their operational productivity to further solidify their positions as trusted healthcare partners. TO ACHIEVE THESE GOALS, LABS NEED TO DRIVE IMPROVEMENTS IN KEY AREAS: Lab schedule efficiencies are often impacted by batch instrument limitations (need for batching and sorting samples). Four to six platforms are often needed for routine PCR testing, causing inefficiencies and requiring more staff training and broader knowledge. Highly skilled staff spend hours each day on tedious tasks. Data is not easily or effectively used to optimize performance. Alinity m is a fully integrated and automated molecular diagnostics analyzer that uses innovative technology to deliver the next level of flexibility and efficiency to your lab. We surveyed health systems and labs around the world to develop the Alinity m to meet your needs, now and into the future. SPECIFICATIONS transforming your lab to meet tomorrow's needs: Providing true random access and STAT prioritization PREDICTABLE OPERATIONAL PERFORMANCE AND EFFICIENCY Ability to manage the majority of routine PCR tests on a single platform DRIVES CONSOLIDATION ON A SINGLE PLATFORM Driving standardization through simple, universal design elements STREAMLINED OPERATIONS AND SIMPLIFIED WORKFLOW Delivering on the Abbott legacy of high-quality product designs and assays INCREASED LAB CONFIDENCE ReadiFlex enables you to achieve true random access by balancing flexibility, turnaround time, and throughput. Perform any test, any time, including STAT samples Rapid turnaround time for all assays Consistent throughput regardless of assay mix ALINITY m ASSAYS Abbott has been on the forefront of innovation for more than 130 years. Alinity m benefits from this legacy and commitment to high quality assay development. - Innovative technology underpinned by robust design to provide accurate, consistent results - Multiplex technology enables you to improve your workflow and increase productivity - Comprehensive assay menu includes Resp-4-Plex, SARS-CoV-2, CMV, EBV, HBV, HCV, HPV, STI and HIV-1 (including Dried Blood Spot sample type). ASSAY SPECIFICATIONS Assay availability varies by country, please contact your local sales representative for more information. DRIVE LAB EFFICIENCY THROUGH: Operational efficiencies Reduces complexity and streamlines workflow STAFF EFFICIENCIES Improve resource flexibility and ease of training COST EFFICIENCIES Drives maintenance and service cost optimization ENVISION THE LAB OF THE FUTURE TRANSFORM CHALLENGES INTO OPPORTUNITIES. GOALS INTO REALITIES. COMPROMISES INTO SUCCESSES. DO IT ALL with A SINGLE SOLUTION THAT CAN TRANSFORM YOUR LAB. discover more Have an Abbott Molecular Laboratory representative contact you about our lab solutions. Let's discuss your lab Receive exclusive access to resources with the Abbott Customer Portal (Diagnostics Portal). Register Now Representatives are ready to offer support, gather feedback, and answer questions. Contact Tech Support Surging ahead. Staying informed. Learn more now

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