

Created September 2020.

For use under an Emergency Use Authorization (EUA) Only

Prescription Use only.

For *In Vitro* Diagnostic Use Only.

Package insert instructions must be carefully followed. Reliability of assay results cannot be guaranteed if there are any deviations from the instructions in this package insert.

For laboratory professional use only.

NAME

AdviseDx SARS-CoV-2 IgM Control Kit (also referred to as CoV-2 IgM CtrlS or SARS-CoV-2 IgM CtrlS)

INTENDED USE

The AdviseDx SARS-CoV-2 IgM Control Kit is for the estimation of test precision and the detection of systematic analytical deviations of the ARCHITECT i System when used for the qualitative detection of IgM antibodies to SARS-CoV-2 in human serum, serum separator tube, and plasma (dipotassium EDTA, tripotassium EDTA, lithium heparin, lithium heparin separator tube, sodium heparin).

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For additional information, refer to the AdviseDx SARS-CoV-2 IgM reagent package insert and the ARCHITECT System Operations Manual.

The AdviseDx SARS-CoV-2 IgM assay is only for use under the Food and Drug Administration's Emergency Use Authorization.

CONTENTS

The **CONTROL**- contains human plasma.

The **CONTROL**+ contains inactivated, cell-free, human blood-derived material, reactive for anti-SARS-CoV-2 IgM.

Preservatives: sodium azide and antimicrobial agents.

The controls are at the following ranges. The ranges may be used for individual replicate control specifications on the ARCHITECT i System.

Anti-SARS-CoV-2 IgM

Control	Quantity	^{RANGE} (Index [S/C])
CONTROL -	1 x 4.0 mL	≤ 0.52
CONTROL +	1 x 4.0 mL	1.39 - 6.67

NOTE: The insert ranges for the controls are not lot specific and represent the total range of values which may be generated throughout the life of the product. It is recommended that each laboratory establish its own means and acceptable ranges which should fall within the package insert ranges. Sources of variation that can be expected include:

- Calibration
- Control lot
- Reagent lot
- Calibrator lot
- Instrument


PRECAUTIONS

For Use Under An Emergency Use Authorization Only.

This assay is only for *in vitro* diagnostic use under the FDA Emergency Use Authorization.

- **IVD**
- For *In Vitro* Diagnostic Use
- **Rx ONLY**
- This product has not been FDA cleared or approved; this test has been authorized by FDA under an EUA for use by laboratories certified under CLIA, that meet requirements to perform moderate or high complexity tests.
- This product has been authorized only for the presence of IgM antibodies against SARS-CoV-2, not for any other viruses or pathogens;
- This product 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 Federal Food, Drug, and Cosmetic Act, 21 U.S.C. § 360bbb-3(b)(1), unless the authorization is terminated or revoked sooner.

Safety Precautions

-  **CAUTION:** This product contains human-sourced and/or potentially infectious components. Refer to the CONTENTS section of this package insert. No known test method can offer complete assurance that products derived from human sources or inactivated microorganisms will not transmit infection. Therefore, all human-sourced materials should be considered potentially infectious. It is recommended that this product, human specimens, and all consumables contaminated with potentially infectious materials be handled in accordance with the OSHA Standard on Bloodborne Pathogens. Biosafety Level 2 or other appropriate regional, national, and institutional biosafety practices should be used for materials that contain, are suspected of containing, or are contaminated with infectious agents. [1](#), [2](#), [3](#), [4](#)
- The human-sourced materials used in the **CONTROL +** have been tested and found to be reactive for anti-SARS-CoV-2 IgM and nonreactive for HBsAg, HIV-1 RNA or HIV-1 Ag, anti-HIV-1/HIV-2, and anti-HCV.
- The human-sourced material used in the **CONTROL -** has been tested and found to be nonreactive for anti-SARS-CoV-2 IgM, HBsAg, HIV-1 RNA or HIV-1 Ag, anti-HIV-1/HIV-2, and anti-HCV.

The following warnings and precautions apply to: CONTROL + and CONTROL -	
Contains sodium azide.	
EUH032	Contact with acids liberates very toxic gas.
P501	Dispose of contents / container in accordance with local regulations.

Follow local chemical disposal regulations based on your location along with recommendations and content in the Safety Data Sheet to determine the safe disposal of this product.

For the most current hazard information, see the product Safety Data Sheet.

Safety Data Sheets are available at www.corelaboratory.abbott or contact your local representative.

For a detailed discussion of safety precautions during system operation, refer to the ARCHITECT System Operations Manual, Section 8.

PREPARATION FOR USE

- Thaw completely at room temperature (15 to 30°C).

- Prior to each use, mix by gentle inversion.

STORAGE

- This product is shipped on dry ice.
- Protect from light.
- Do not use past expiration date.

	Storage Temperature	Maximum Storage Time	Additional Storage Instructions
Unopened	-20°C or colder	Until expiration date	
Opened	2 to 8°C	Up to 45 days after thaw, not to exceed expiration date printed on the bottle	Return to original carton to protect from light. Store tightly capped. Store in upright position.

INSTRUMENT PROCEDURE

- To obtain the recommended volume requirements for the controls, hold the bottle vertically, and dispense 4 drops of the negative control and 4 drops of the positive control into each sample cup in the assigned position.
- For information on configuring control data, refer to the ARCHITECT System Operations Manual, Section 2.
- For instructions on ordering and loading controls on the instrument, refer to the ARCHITECT System Operations Manual, Section 5.

QUALITY CONTROL PROCEDURES

The recommended control requirement for the AdviseDx SARS-CoV-2 IgM assay is that a single sample of each control level be tested once every 24 hours each day of use.

INDICATIONS OF INSTABILITY OR DETERIORATION

Instability or deterioration should be suspected if there are precipitates, visible signs of leakage, or if controls do not meet the appropriate package insert and/or ARCHITECT System Operations Manual criteria.

NOTE: The AdviseDx SARS-CoV-2 IgM controls may present a cloudy or turbid appearance following thaw. This is not necessarily a sign of deterioration if the controls meet the appropriate

criteria.

LIMITATIONS









Control values have not been established for assays other than the AdviseDx SARS-CoV-2 IgM assay. If the user wishes to use this control material with other assays, it is their responsibility to establish the appropriate ranges.

The controls are not calibrators and should not be used for assay calibration.

BIBLIOGRAPHY

1. US Department of Labor, Occupational Safety and Health Administration, 29 CFR Part 1910.1030, Bloodborne pathogens.
2. US Department of Health and Human Services. *Biosafety in Microbiological and Biomedical Laboratories*. 5th ed. Washington, DC: US Government Printing Office; December 2009.
3. World Health Organization. *Laboratory Biosafety Manual*. 3rd ed. Geneva: World Health Organization; 2004.
4. Clinical and Laboratory Standards Institute (CLSI). *Protection of Laboratory Workers From Occupationally Acquired Infections; Approved Guideline—Fourth Edition*. CLSI Document M29-A4. Wayne, PA: CLSI; 2014.

Key to Symbols

ISO 15223 Symbols	
	Caution
	Consult instructions for use
	Manufacturer
	Temperature limitation
	Upper limit of temperature
	Use by/Expiration date
	Negative Control
	Positive Control

ISO 15223 Symbols

IVD	<i>In Vitro</i> Diagnostic Medical Device
LOT	Lot Number
REF	List Number

Other Symbols

AFTER THAW	After thaw store at
CONTAINS: AZIDE	Contains Sodium Azide. Contact with acids liberates very toxic gas.
FOR USE WITH	Identifies products to be used together
GTIN	Global Trade Item Number
INFORMATION FOR USA ONLY	Information needed for United States of America only
PRODUCT OF IRELAND	Product of Ireland
PROTECT FROM LIGHT	Protect from light
RANGE	Range
Rx ONLY	For use by or on the order of a physician only (applicable to USA classification only).
UNTIL FIRST USE	Until first use store at

Note for number formatting:

- A space is used as thousands separator (example: 10 000 specimens).
- A period is used to separate the integer part from the fractional part of a number written in decimal form (example: 3.12%).

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Customer Service: Contact your local representative or find country-specific contact information on www.corelaboratory.abbott

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