ARCHITECT
Total T_3 Calibrators

REF 7K64-02

en

Total T3
7K64
G8-0469 / R02
S7K6V0

INTENDED USE

The ARCHITECT Total T_3 Calibrators are for the calibration of the ARCHITECT iSystem when used for the quantitative determination of total triiodothyronine (Total T_3) in human serum and plasma. Refer to the ARCHITECT assay-specific reagent package insert for additional information.

CONTENTS

6 Bottles (4 mL each) of ARCHITECT Total T_3 Calibrators prepared in human serum. Preservative: Sodium azide.

The calibrators yield the following concentrations:

	Total T_3 Concentration	
Calibrator	ng/mL	nmol/L
CAL A	0.2	0.31
CAL B	0.5	0.77
CAL C	0.75	1.15
CAL D	1.5	2.30
CAL E	3.5	5.38
CAL F	6.0	9.22

STANDARDIZATION

The calibrators are manufactured using Certified Reference Material (CRM), Triiodo-L-Thyronine (T_3), and signal matched to Abbott internal reference standards, which are traceable to the CRM Triiodo-L-Thyronine (T_3) at each concentration level. The concentration of the CRM Triiodo-L-Thyronine (T_3) used to make the internal reference standards is determined by HPLC.

PRECAUTIONS

- IVD
- For *In Vitro* Diagnostic Use
- Rx ONLY
- **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 these reagents and human specimens be handled in accordance with the OSHA Standard on Bloodborne Pathogens. Biosafety Level 2 or other appropriate biosafety practices should be used for materials that contain or are suspected of containing infectious agents.[1-4]
- The human serum used in the ARCHITECT Total T_3 Calibrators is nonreactive for HBsAg, HIV-1 RNA or HIV-1 Ag, anti-HCV, and anti-HIV-1/HIV-2.

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

Safety Data Sheets are available at www.abbottdiagnostics.com or contact your local representative.

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

STORAGE

- Calibrators are stable until the expiration date when stored and handled as directed.
- Do not use past expiration date.
- 2°C – 8°C

PREPARATION FOR USE

Calibrators may be used immediately after removal from 2-8°C storage.

Prior to each use, mix by gentle inversion.

After each use, tightly close the caps and return the calibrators to 2-8°C storage.

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

	Caution
	Consult instructions for use
	Manufacturer
	Temperature limitation
	Use by/Expiration date
CAL A	Calibrator (A,B,C,D,E or F)
CONTAINS AZIDE	Contains Sodium Azide. Contact with acids liberates very toxic gas.
DISTRIBUTED IN THE USA BY	Distributed in the USA by
INFORMATION FOR USA ONLY	Information needed for United States of America only
IVD	*In Vitro* Diagnostic Medical Device
LOT	Lot Number
PRODUCT OF IRELAND	Product of Ireland
REF	List Number
Rx ONLY	For use by or on the order of a physician only (applicable to USA classification only).

ARCHITECT is a trademark of Abbott Laboratories in various jurisdictions.

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Created October 2016.

Abbott

Architect
Total T_4 Controls

REF 7K66-10

en
Total T4
7K66
G6-2781 / R04
C7K660

Read Highlighted Changes: Revised December 2015.

INTENDED USE

The ARCHITECT Total T_4 Controls are for the verification of the accuracy and precision of the ARCHITECT iSystem when used for the quantitative determination of thyroxine (Total T_4) in human serum and plasma. Refer to the ARCHITECT assay-specific reagent package insert for additional information.

CONTENTS

3 Bottles (8 mL each) of ARCHITECT Total T_4 Controls. The Low Control, Medium Control, and High Control contain T_4 prepared in human serum. Preservative: Sodium Azide.

The following concentration ranges may be used for individual replicate control specifications on the ARCHITECT iSystem:

	Target Concentration / Range			
Control	µg/dL		nmol/L	
CONTROL L	4.2	2.73 - 5.43	54.1	35.14 - 69.88
CONTROL M	7.4	5.44 - 9.92	95.2	70.01 - 127.67
CONTROL H	14.6	9.93 - 19.71	187.9	127.8 - 253.67

Each laboratory should establish its own concentration ranges for new control lots at each control level. This can be accomplished by assaying a minimum of 20 replicates over several (3-5) days. Sources of variation that can be expected should be included in this study in order to be representative of future system performance. These may include:

- Multiple stored calibrations
- Multiple reagent lots
- Multiple calibrator lots
- Multiple processing modules
- Data points collected at different times of the day

These results should be applied to your laboratory's quality control practices.

PRECAUTIONS

- IVD
- For *In Vitro* Diagnostic Use
- **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 these reagents and human specimens be handled in accordance with the OSHA Standard on Bloodborne Pathogens. Biosafety Level 2 or other appropriate biosafety practices should be used for materials that contain or are suspected of containing infectious agents.
- The human serum used in the ARCHITECT Total T_4 Controls is nonreactive for HBsAg, HIV-1 RNA or HIV-1 Ag, anti-HCV, and anti-HIV-1/HIV-2.

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

Safety Data Sheets are available at www.abbottdiagnostics.com or contact your local representative.

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

STORAGE

- Controls are stable until the expiration date when stored and handled as directed.
- Do not use past expiration date.
- 8°C / 2°C

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

	Caution
	Consult instructions for use
	Manufacturer
	Temperature limitation
	Use by/Expiration date
CONC	Concentration
CONTAINS AZIDE	Contains Sodium Azide. Contact with acids liberates very toxic gas.
CONTROL L	Control Low, Medium, High (L,M,H)
DISTRIBUTED IN THE USA BY	Distributed in the USA by
INFORMATION FOR USA ONLY	Information needed for United States of America only
IVD	*In Vitro* Diagnostic Medical Device
LOT	Lot Number
PRODUCT OF IRELAND	Product of Ireland
RANGE	Range
REF	List Number

ARCHITECT is a trademark of Abbott Laboratories in various jurisdictions.

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Revised December 2015.

Abbott

ARCHITECT Total T_4

REF 7K66-27
REF 7K66-37
REF 7K66-32

en
Total T4
7K66
G65409R02
B7K6T0

Read Highlighted Changes: Revised May 2017.

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.

NAME

ARCHITECT Total T_4

INTENDED USE

The ARCHITECT Total T_4 (TT_4) assay is a Chemiluminescent Microparticle Immunoassay (CMIA) for the quantitative determination of thyroxine (Total T_4) in human serum and plasma on the ARCHITECT iSystem.

SUMMARY AND EXPLANATION OF THE TEST

Thyroxine (T_4) is an iodine-containing hormone which has a molecular weight of approximately 777 daltons and is secreted by the thyroid gland. T_4 and its associate thyroid hormone T_3 are responsible for regulating diverse biochemical processes throughout the body which are essential for normal metabolic and neural activity.[1]

Although T_3 has greater biologic potency[2], T_4 is normally present in human serum in approximately 50-fold excess of circulating T_3 and accounts for more than 90% of the circulating protein-bound iodine. T_4 is 99.9% bound to serum thyroxine binding proteins (TBP). The hormone is transported bound primarily to thyroxine binding globulin (TBG) and secondarily by thyroxine binding prealbumin (TBPA) and albumin.[3] Less than 0.05% of the total circulating T_4 is unbound and therefore biologically active.[4, 5] Clinically, T_4 measurements have long been recognized as an aid in the assessment and diagnosis of thyroid status. Elevated T_4 values are characteristically seen in patients with overt hyperthyroidism, while T_4 levels are generally depressed in patients with overt hypothyroidism. Normal T_4 levels accompanied by high T_3 values are seen in patients with T_3-thyrotoxicosis.[6] T_4 levels are altered by physiological or pathological changes in TBP capacity.[3, 4] Thyroxine binding globulin (TBG) capacity has a pronounced effect on the concentration of thyroid hormones. Consequently, T_4 levels may be elevated with increased concentrations of TBG, such as in pregnancy, administration of oral contraceptives or estrogen, infectious and chronic active hepatitis, biliary cirrhosis or congenital increase in TBG levels.[7-9] Conversely, when TBG levels are decreased, such as in nephrotic syndrome, androgen therapy, glucocorticoid therapy, major systemic illness or congenital decrease of TBG, T_4 may be reduced.

Drugs which compete for protein binding sites, such as phenylbutazone, diphenylhydantoin or salicylates, can result in a depressed T_4 measurement.[7-9] Serum T_4 levels in neonates and infants are higher than values in the normal adult, due to the increased concentration of TBG in neonate serum.[10]

While in many cases T_4 values give good indications of thyroid status, T_4 values should be normalized for individual variations in thyroxine binding protein (TBP) capacity. The Free Thyroxine Index (FTI) is conventionally used to achieve this measurement.[11, 12]

To ensure maximum diagnostic accuracy, the final definition of thyroid status should be determined in conjunction with other thyroid function tests such as TSH, Free T_4, Total T_3, FTI and clinical evaluation by the physician.

The ARCHITECT Total T_4 assay is to be used as an aid in the assessment of thyroid status.

BIOLOGICAL PRINCIPLES OF THE PROCEDURE

The ARCHITECT Total T_4 assay is a two-step immunoassay to determine the presence of thyroxine (Total T_4) in human serum and plasma using CMIA technology with flexible assay protocols, referred to as Chemiflex.

1. Sample and anti-T_4 coated paramagnetic microparticles are combined. Bound T_4 is removed from the binding sites on thyroxine binding globulin, prealbumin and albumin. The T_4 present in the sample binds to the anti-T_4 coated microparticles.
2. After washing, T_3 acridinium-labeled conjugate is added to create a reaction mixture.
3. Following another wash cycle, Pre-Trigger and Trigger Solutions are added to the reaction mixture.
4. The resulting chemiluminescent reaction is measured as relative light units (RLUs). There is an inverse relationship between the amount of Total T_4 in the sample and the RLUs detected by the ARCHITECT iSystem optics.

For additional information on system and assay technology, refer to the ARCHITECT System Operations Manual, Section 3.

REAGENTS

Kit Contents

ARCHITECT Total T_4 7K66

NOTE: Some kit sizes are not available in all countries or for use on all ARCHITECT iSystems. Please contact your local distributor.

REF	7K66-27	7K66-37	7K66-32
Σ	100	500	2000
MICROPARTICLES	1 x 6.6 mL	1 x 27.0 mL	4 x 27.0 mL
CONJUGATE	1 x 5.9 mL	1 x 26.3 mL	4 x 26.3 mL

MICROPARTICLES Anti-T_4 (sheep) coated Microparticles in TRIS buffer with sheep IgG stabilizers. Minimum concentration: 0.05% solids. Preservative: Sodium Azide.

CONJUGATE T_3 acridinium-labeled Conjugate in MES buffer with NaCl and Triton X-100 stabilizers. Minimum concentration: 0.2 ng/mL. Preservative: ProClin.

Other Reagents

PRE-TRIGGER SOLUTION ARCHITECT Pre-Trigger Solution containing 1.32% (w/v) hydrogen peroxide.

TRIGGER SOLUTION ARCHITECT Trigger Solution containing 0.35 N sodium hydroxide.

WASH BUFFER ARCHITECT Wash Buffer containing phosphate buffered saline solution. Preservatives: antimicrobial agents.

NOTE: Bottle and volume varies based on order.

Warnings and Precautions

- IVD
- For *In Vitro* Diagnostic Use

Safety Precautions

CAUTION: This product requires the handling of human specimens. It is recommended that all human-sourced materials be considered potentially infectious and handled in accordance with the OSHA Standard on Bloodborne Pathogens. Biosafety Level 2 or other appropriate biosafety practices should be used for materials that contain or are suspected of containing infectious agents.[13-16]

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

The following warnings and precautions apply to: CONJUGATE	
	
WARNING	Contains methylisothiazolones.
H317	May cause an allergic skin reaction.
Prevention	
P261	Avoid breathing mist / vapors / spray.
P272	Contaminated work clothing should not be allowed out of the workplace.
P280	Wear protective gloves / protective clothing / eye protection.
Response	
P302+P352	IF ON SKIN: Wash with plenty of water.
P333+P313	If skin irritation or rash occurs: Get medical advice / attention.
P362+P364	Take off contaminated clothing and wash it before reuse.
Disposal	
P501	Dispose of contents / container in accordance with local regulations.

Safety Data Sheets are available at www.abbottdiagnostics.com or contact your local representative.

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

Reagent Handling

- Do not use reagent kits beyond the expiration date.
- **Do not pool reagents within a kit or between kits.**
- Before loading the reagent kit on the system for the first time, the microparticle bottle requires mixing to resuspend microparticles that may have settled during shipment. For microparticle mixing instructions, refer to the **PROCEDURE, Assay Procedure** section of this package insert.
- **Septums MUST be used to prevent reagent evaporation and contamination and to ensure reagent integrity. Reliability of assay results cannot be guaranteed if septums are not used according to the instructions in this package insert.**
  - To avoid contamination, wear clean gloves when placing a septum on an uncapped reagent bottle.
  - Once a septum has been placed on an open reagent bottle, **do not invert the bottle** as this will result in reagent leakage and may compromise assay results.
  - Over time, residual liquids may dry on the septum surface. These are typically dried salts and have no effect on assay efficacy.

For a detailed discussion of handling precautions during system operation, refer to the ARCHITECT System Operations Manual, Section 7.

Reagent Storage

When stored and handled as directed, reagents are stable until the expiration date.

	Storage Temperature	Maximum Storage Time	Additional Storage Instructions
Unopened/ Opened*	2-8°C	Until expiration date	May be used immediately after removal from 2-8°C storage. Store in upright position.
On board	System temperature	30 days	Discard after 30 days. For information on tracking onboard time, refer to the ARCHITECT System Operations Manual, Section 5.

* Reagents may be stored on or off the ARCHITECT iSystem. If reagents are removed from the system, store them at 2-8°C (with septums and replacement caps) in an upright position. For reagents stored off the system, it is recommended that they be stored in their original trays and boxes to ensure they remain upright. **If the microparticle bottle does not remain upright (with a septum installed) while in refrigerated storage off the system, the reagent kit must be discarded.** For information on unloading reagents, refer to the ARCHITECT System Operations Manual, Section 5.

Indications of Reagent Deterioration

When a control value is out of the specified range, it may indicate deterioration of the reagents or errors in technique. Associated test results are invalid, and samples must be retested. Assay recalibration may be necessary. For troubleshooting information, refer to the ARCHITECT System Operations Manual, Section 10.

INSTRUMENT PROCEDURE

The ARCHITECT Total T_4 assay file must be installed on the ARCHITECT iSystem from an ARCHITECT iSystem Assay CD-ROM prior to performing the assay.

For detailed information on assay file installation and viewing and editing assay parameters, refer to the ARCHITECT System Operations Manual, Section 2.

For information on printing assay parameters, refer to the ARCHITECT System Operations Manual, Section 5.

For a detailed description of system procedures, refer to the ARCHITECT System Operations Manual.

Alternate Result Units

Edit assay parameter "Result concentration units" to select an alternate unit.

Conversion formula:

(Concentration in Default result unit) x (Conversion factor) = (Concentration in Alternate result unit)

Default result unit	Conversion factor	Alternate result unit
µg/dL	12.87	nmol/L

SPECIMEN COLLECTION AND PREPARATION FOR ANALYSIS

Specimen Types

Validated specimen types to be used with this assay:

Specimen Types	Collection Tubes
Human serum	Serum
	Serum separator tubes
Human plasma	Sodium heparin
	Lithium heparin
	Lithium heparin plasma separator tubes
	Potassium EDTA

- Other specimen collection tube types have not been validated with this assay.
- The instrument does not provide the capability to verify specimen type. It is the responsibility of the operator to verify that the correct specimen types are used in the assay.
- When serial specimens are being evaluated, the same type of specimen should be used throughout the study.

Specimen Conditions

- Do not use specimens with the following conditions:
  - heat-inactivated
- To prevent cross contamination, use of disposable pipettes or pipette tips is recommended.
- **For optimal results, serum and plasma specimens should be free of fibrin, red blood cells or other particulate matter.**
- **Ensure that complete clot formation in serum specimens has taken place prior to centrifugation. Some specimens, especially those from patients receiving anticoagulant or thrombolytic therapy may exhibit increased clotting time. If the specimen is centrifuged before a complete clot forms, the presence of fibrin may cause erroneous results.**

Preparation for Analysis

- Follow the tube manufacturer's processing instructions for specimen collection tubes.
- Mix thawed specimens thoroughly by low speed vortexing or by inverting 10 times. Visually inspect the specimens. If layering or stratification is observed, continue mixing until specimens are visibly homogeneous.
- To ensure consistency in results, centrifuge specimens before testing if
  - they contain fibrin, red blood cells, or other particulate matter or
  - they were frozen and thawed.
- Inspect all specimens for bubbles. Remove bubbles with an applicator stick before analysis. Use a new applicator stick for each specimen to prevent cross contamination.

Specimen Storage

Specimen Type	Storage Temperature	Maximum Storage Time
Serum/Plasma	2-8°C	≤ 6 days

If testing will be delayed more than 24 hours, remove serum or plasma from the clot, serum separator or red blood cells.

If testing will be delayed more than 6 days, specimens should be frozen at -10°C or colder.

Specimens stored frozen at -10°C or colder for 6 days showed no performance difference.

Avoid multiple freeze/thaw cycles.

Specimen Shipping

- Package and label specimens in compliance with applicable state, federal, and international regulations covering the transport of clinical specimens and infectious substances.
- Do not exceed the storage limitations listed above.

PROCEDURE

Materials Provided

7K66 ARCHITECT Total T_4 Reagent Kit

Materials Required but not Provided

- ARCHITECT Total T_4 Assay file obtained from the ARCHITECT iSystem e-Assay CD-ROM found on www.abbottdiagnostics.com.
- 7K66-02 ARCHITECT Total T_4 Calibrators
- 7K66-12 ARCHITECT Total T_4 Controls
- ARCHITECT Pre-Trigger Solution
- ARCHITECT Trigger Solution
- ARCHITECT Wash Buffer
- ARCHITECT Reaction Vessels
- ARCHITECT Sample Cups
- ARCHITECT Septum
- ARCHITECT Replacement Caps
- Pipettes or pipette tips (optional) to deliver the volumes specified on the patient or control order screen.

For information on materials required for maintenance procedures, refer to the ARCHITECT System Operations Manual, Section 9.

Assay Procedure

- Before loading the reagent kit on the system for the first time, the microparticle bottle requires mixing to resuspend microparticles that may have settled during shipment. After the first time the microparticles have been loaded, no further mixing is required.
  - **Invert the microparticle bottle 30 times.**
  - Visually inspect the bottle to ensure microparticles are resuspended. If microparticles are still adhered to the bottle, continue to invert the bottle until the microparticles have been completely resuspended.
  - **If the microparticles do not resuspend, DO NOT USE. Contact your local Abbott representative.**
  - Once the microparticles have been resuspended, place a septum on the bottle. For instructions about placing septums on bottles, refer to the **Reagent Handling** section of this package insert.
- Load the reagent kit on the ARCHITECT iSystem.
  - Verify that all necessary reagents are present.
  - Ensure that septums are present on all reagent bottles.
- Order tests.
  - For information on ordering patient specimens and controls and for general operating procedures, refer to the ARCHITECT System Operations Manual, Section 5.
- Minimum sample cup volume is calculated by the system and printed on the Orderlist report. To minimize the effects of evaporation, verify adequate sample cup volume is present prior to running the test.

  Maximum number of replicates sampled from the same sample cup: 10
  - Priority:

    Sample volume for first test: 74 µL

    Sample volume for each additional test from same sample cup: 24 µL
  - ≤ 3 hours on board:

    Sample volume for first test: 150 µL

    Sample volume for each additional test from same sample cup: 24 µL
  - > 3 hours on board: Additional sample volume is required. Refer to the ARCHITECT System Operations Manual, Section 5 for information on sample evaporation and volumes.
  - If using primary or aliquot tubes, use the sample gauge to ensure sufficient patient specimen is present.

- Order calibration, if necessary.
  - For information on ordering calibrations, refer to the ARCHITECT System Operations Manual, Section 6.
- Prepare ARCHITECT Total T_4 Calibrators and Controls.
  - Mix calibrator(s) and controls by gentle inversion before use.
  - Hold bottles **vertically** and dispense recommended volumes into each respective sample cup.
  - Recommended volumes:

    for each calibrator: 4 drops

    for each control: 4 drops
- Load samples.
  - For information on loading samples, refer to the ARCHITECT System Operations Manual, Section 5.
- Press RUN.
- For optimal performance, it is important to perform routine maintenance as described in the ARCHITECT System Operations Manual, Section 9. Perform maintenance more frequently when required by laboratory procedures.

Specimen Dilution Procedures

Specimens with a Total T_4 value exceeding 24.00 µg/dL are flagged with the code ">24.00" and may be diluted using the Manual Dilution Procedure.

Manual Dilution Procedure

Suggested dilution: 1:2

It is recommended that dilutions not exceed 1:2.

1. Add 75 µL of the patient specimen to 75 µL of ARCHITECT Total T_4 Calibrator A.

   To avoid contamination of Calibrator A, dispense several drops of Calibrator A into a clean test tube prior to pipetting.
2. The operator must enter the dilution factor (2) in the patient or control order screen. The system will use this dilution factor to automatically calculate the concentration of the sample before dilution. This will be the reported result. The dilution should be performed so that the reported result reads greater than 6.0 µg/dL.

   If the operator does not enter the dilution factor, the reported result will be that of the diluted sample. This result (before dilution factor is applied) should be greater than 3.00 µg/dL.

For detailed information on ordering dilutions, refer to the ARCHITECT System Operations Manual, Section 5.

Calibration

- Test Calibrators A - F in duplicate. The calibrators should be priority loaded.

  A single replicate of each control level must be tested to evaluate the assay calibration. Ensure that assay control values are within the ranges specified in the respective control package insert.
- Calibration Range: 0.0 - 24.0 µg/dL.
- Once an ARCHITECT Total T_4 calibration is accepted and stored, all subsequent samples may be tested without further calibration unless:
  - A reagent kit with a new lot number is used or
  - Controls are out of range.
- For detailed information on how to perform an assay calibration, refer to the ARCHITECT System Operations Manual, Section 6.

Quality Control Procedures

The recommended control requirement for the ARCHITECT Total T_4 assay is that a single replicate of each control level be tested once every 24 hours each day of use. If the quality control procedures in your laboratory require more frequent use of controls to verify test results, follow your laboratory-specific procedures.

Ensure that assay control values are within the concentration ranges specified in the control package insert.

Verification of Assay Claims

For protocols to verify package insert claims, refer to the ARCHITECT System Operations Manual, Appendix B.

The ARCHITECT Total T_4 assay belongs to method group 1.

RESULTS

Calculation

The ARCHITECT Total T_4 assay utilizes a 4 Parameter Logistic Curve fit data reduction method (4PLC, Y-weighted) to generate a calibration curve.

For information on alternate result units, refer to the **INSTRUMENT PROCEDURE, Alternate Result Units** section of this package insert.

Measuring Interval

Measuring interval is defined as the range of values in µg/dL which meets the limits of acceptable performance for both imprecision and linearity. The measuring interval for the ARCHITECT Total T_4 assay is 3.00 µg/dL (Limit of Quantitation - LoQ) to 24.00 µg/dL.

Flags

Some results may contain information in the Flags field. For a description of the flags that may appear in this field, refer to the ARCHITECT System Operations Manual, Section 5.

LIMITATIONS OF THE PROCEDURE

- For diagnostic purposes, results should be used in conjunction with other data; e.g., symptoms, results of other thyroid tests, clinical impressions, etc.
- If the Total T_4 results are inconsistent with clinical evidence, additional testing is suggested to confirm the result.
- Performance of this test has not been established with neonatal specimens.

EXPECTED VALUES

A normal range of 4.87 µg/dL to 11.72 µg/dL (central 95% interval) was obtained by testing serum specimens from 437 individuals determined as normal by AxSYM Ultrasensitive hTSH II and AxSYM Free T_4 assays. It is recommended that each laboratory establish its own normal range, which may be unique to the population it serves depending upon geographical, patient, dietary, or environmental factors.

SPECIFIC PERFORMANCE CHARACTERISTICS

Precision

The ARCHITECT Total T_4 assay is designed to have a precision of ≤ 10% (total CV) for concentrations in the range of the low control (4.2 µg/dL), medium control (7.4 µg/dL), and high control (14.6 µg/dL). A study based on guidance from Clinical and Laboratory Standards Institute (CLSI, formerly NCCLS) document EP5-A[17] was performed for the ARCHITECT Total T_4 assay. A three member processed human serum based panel was assayed, using two lots of reagents, in replicates of two at two separate times per day for 20 testing days. Data from this study are shown in the following table.*

Panel Member	Reagent Lot	Instrument	n	Mean Conc. Value (µg/dL)	Within Run SD	Within Run %CV	Total SD	Total %CV
1	1	1	80	4.20	0.155	3.7	0.188	4.5
1	2	1	80	4.32	0.176	4.1	0.196	4.5
1	1	2	80	4.45	0.133	3.0	0.167	3.8
1	2	2	80	4.30	0.136	3.2	0.151	3.5
2	1	1	80	7.46	0.251	3.4	0.300	4.0
2	2	1	80	7.35	0.222	3.0	0.268	3.6
2	1	2	80	7.90	0.234	3.0	0.301	3.8
2	2	2	80	7.32	0.210	2.9	0.224	3.1
3	1	1	80	14.87	0.896	6.0	1.084	7.3
3	2	1	80	14.94	0.521	3.5	0.655	4.4
3	1	2	80	16.22	0.616	3.8	0.738	4.5
3	2	2	80	15.13	0.595	3.9	0.703	4.6

* Representative data; results in individual laboratories may vary from these data.

Recovery

The ARCHITECT Total T_4 assay is designed to have a mean recovery of 100 ± 10% when analyzing samples spiked with known amounts of T_4.

T_4 was added to five normal human serum samples. The concentration of Total T_4 was determined using the ARCHITECT Total T_4 assay and the resulting percent recovery was calculated.*

Sample	Endogenous T_4 Concentration (µg/dL)	T_4 Added (µg/dL)	Observed Total T_4 Concentration (µg/dL)	%Recovery*
1	7.472	1.20	8.825	112.7
2	7.301	1.20	8.487	98.9
3	7.574	1.20	8.631	88.2
4	6.760	1.20	7.911	95.9
5	8.547	1.20	10.015	122.3

Mean Recovery 103.6%

$$\text{* \% Recovery} = \frac{\text{Observed Total } T_4 \text{ Conc. (µg/dL)} - \text{Endogenous Total } T_4 \text{ Conc. (µg/dL)}}{T_4 \text{ Added (µg/dL)}} \times 100$$

Representative data; results in individual laboratories may vary from these data.

Sensitivity

The ARCHITECT Total T_4 assay is designed to have a Limit of Quantitation (LoQ) of ≤ 3.0 µg/dL. The LoQ is defined as the lowest concentration at which analyte in a sample can be accurately quantitated with precision of ≤ 10% CV.

A study was performed based on guidance from the CLSI document EP17-A2[18] with four zero-level samples and 8 samples with T_4 concentrations ranging from 2.0 to 3.5 µg/dL. The samples were tested in at least 5 separate runs over a minimum of 3 days using 2 reagent lots and 6 instruments. In this study, the Limit of Blank (LoB) was 0.70 µg/dL, Limit of Detection (LoD) was 0.91 µg/dL and LoQ was 2.0 µg/dL.*

* Representative data; results in individual laboratories may vary from these data.

Analytical Specificity

The ARCHITECT Total T_4 assay is designed to have a mean analytical specificity of ≤ 3.2% cross reactivity with triiodothyronine (T_3) at a concentration of 100 µg/dL in a sample containing approximately 3 µg/dL of Total T_4 as confirmed by a study based on guidance from CLSI document EP7-A.[19]

Interference

The ARCHITECT Total T_4 assay is designed to have a mean potential interference from hemoglobin, bilirubin, triglycerides, and protein of < 10% at the levels indicated below as confirmed by a study based on guidance from CLSI document EP7-A.[19]

Hemoglobin	≤ 500 mg/dL
Bilirubin	≤ 20 mg/dL
Triglycerides	≤ 3000 mg/dL
Protein	≥ 4.5 and ≤ 12 g/dL

Accuracy by Correlation

The ARCHITECT Total T_4 assay is designed to have a slope of 1.00 ± 0.20 and a correlation coefficient (r) of ≥ 0.90 when compared to the AxSYM Total T_4 assay.

A study was performed where specimens were tested using ARCHITECT Total T_4 assay and AxSYM Total T_4 assay. Data from this study were analyzed using least squares and Passing Bablok[20] regression methods and are summarized in the following table.*

Abbott ARCHITECT Total T_4 vs. Abbott AxSYM Total T_4

Method	Number of Specimens	Intercept	Slope	Correlation Coefficient
Least Squares Linear Regression	656	+0.26	0.96	0.97
Passing-Bablok Linear Regression**	656	-0.20	0.94	0.97

In this evaluation, serum specimens tested ranged from 1.03 to 20.55 µg/dL with the ARCHITECT Total T_4 assay and from 1.12 to 22.46 µg/dL with the AxSYM Total T_4 assay.

* Representative data: variables such as differences in sampling size and sample population may impact the correlation of the assay; therefore, results in individual laboratories may vary from these data.

** A linear regression method with no special assumptions regarding the distribution of the samples and the measurement errors.[20]

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Key to Symbols

	Consult instructions for use
	Manufacturer
	Sufficient for
	Temperature limitation
	Use by/Expiration date
CONJUGATE	Conjugate
CONTAINS: AZIDE	Contains Sodium Azide. Contact with acids liberates very toxic gas.
CONTROL NO.	Control Number
DISTRIBUTED IN THE USA BY	Distributed in the USA by
INFORMATION FOR USA ONLY	Information needed for United States of America only
IVD	*In Vitro* Diagnostic Medical Device
LOT	Lot Number
MICROPARTICLES	Microparticles
PRE-TRIGGER SOLUTION	Pre-Trigger Solution
PRODUCT OF IRELAND	Product of Ireland
REACTION VESSELS	Reaction Vessels
REAGENT LOT	Reagent Lot
REF	List Number
REPLACEMENT CAPS	Replacement Caps
SAMPLE CUPS	Sample Cups
SEPTUM	Septum
SN	Serial number
TRIGGER SOLUTION	Trigger Solution
WARNING: SENSITIZER	Warning: May cause an allergic reaction.
WASH BUFFER	Wash Buffer

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Revised May 2017.

Abbott

ARCHITECT
Total T_4 Calibrators

REF 7K66-02

en
Total T4
7K66
G6-7577 / R02
S7K6T0

Revised May 2016

INTENDED USE

The ARCHITECT Total T_4 Calibrators are for the calibration of the ARCHITECT iSystem when used for the quantitative determination of thyroxine (Total T_4) in human serum and plasma. Refer to the ARCHITECT assay-specific reagent package insert for additional information.

CONTENTS

6 Bottles (4 mL each) of ARCHITECT Total T_4 Calibrators. Calibrators A - F prepared in human serum. Preservative: Sodium Azide.

The calibrators yield the following concentrations:

Calibrators	Total T_4 Concentration (µg/dL)	Total T_4 Concentration (nmol/L)
CAL A	0.0	0.0
CAL B	3.0	38.6
CAL C	6.0	77.2
CAL D	12.0	154.4
CAL E	18.0	231.7
CAL F	24.0	308.9

STANDARDIZATION

The calibrators are matched to an Abbott internal reference standard. This internal reference standard is manufactured by gravimetric methods using USP reference L-Thyroxine.

PRECAUTIONS

- Rx ONLY
- IVD
- For *In Vitro* Diagnostic Use
- **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 these reagents and human specimens be handled in accordance with the OSHA Standard on Bloodborne Pathogens. Biosafety Level 2 or other appropriate biosafety practices should be used for materials that contain or are suspected of containing infectious agents.[1-4]
- The human serum used in the ARCHITECT Total T_4 Calibrators is nonreactive for HBsAg, HIV-1 RNA or HIV-1 Ag, anti-HCV, and anti-HIV-1/HIV-2.

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

Safety Data Sheets are available at www.abbottdiagnostics.com or contact your local representative.

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

STORAGE

- Calibrators are stable until the expiration date when stored and handled as directed.
- Do not use past expiration date.
- 2°C – 8°C

PREPARATION FOR USE

- The calibrators may be used immediately after removal from 2-8°C storage. Prior to each use mix by gentle inversion. After each use, tightly close the caps and return the calibrators to 2-8°C storage.

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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

	Caution
	Consult instructions for use
	Manufacturer
	Temperature limitation
	Use by/Expiration date
CAL A	Calibrator (A,B,C,D,E or F)
CONTAINS AZIDE	Contains Sodium Azide. Contact with acids liberates very toxic gas.
DISTRIBUTED IN THE USA BY	Distributed in the USA by
INFORMATION FOR USA ONLY	Information needed for United States of America only
IVD	*In Vitro* Diagnostic Medical Device
LOT	Lot Number
PRODUCT OF IRELAND	Product of Ireland
REF	List Number
Rx ONLY	For use by or on the order of a physician only (applicable to USA classification only).

ARCHITECT is a trademark of Abbott Laboratories in various jurisdictions.

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Revised May 2016.

Abbott

ARCHITECT Anti-TPO

REF 2K47-20
REF 2K47-25

en
Anti-TPO
2K47
G1-0392/R04
B2K470

Read Highlighted Changes: Revised February 2015.

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.

NAME

ARCHITECT Anti-TPO

INTENDED USE

ARCHITECT Anti-TPO is a Chemiluminescent Microparticle Immunoassay (CMIA) for the quantitative determination of the IgG class of thyroid peroxidase autoantibodies (anti-TPO) in human serum and plasma on the ARCHITECT iSystem. The ARCHITECT Anti-TPO assay is intended for use as an aid in the diagnosis of thyroid disease.

SUMMARY AND EXPLANATION OF THE TEST

It was first demonstrated by Trotter et al. in 1957[1] and subsequently by Roitt and Doniach in 1958[2] that many patients with Hashimoto's thyroiditis had detectable autoantibodies in their blood directed at a thyroid antigen distinct from thyroglobulin. This antigen was termed thyroid microsomal and it has since been demonstrated that most if not all anti-thyroid microsomal autoantibodies recognize thyroid peroxidase (TPO).[3]

TPO is a membrane-bound glycoprotein enzyme with an approximate mass of 107kD. The *in vivo* function is the iodination of tyrosine in the synthesis of T_3 and T_4.[4] Autoimmune reactivity to TPO is believed to be polyclonal and heterogeneous in nature with a minimum of six antigenic determinants being recognized, comprising both conformational and linear epitopes.[5, 6] In addition, the proportion of each immunoglobulin class (G or M) or subclass (G1 – G4) as well as their affinity varies widely from patient to patient.[7, 8] Unlike autoantibodies to thyroglobulin (anti-Tg), autoantibodies to TPO fix complement,[9] are potentially deleterious and may have a pathogenic role in (destructive) autoimmune thyroid disease.[10, 11] Anti-TPO antibodies are found often in conjunction with anti-Tg in the majority of cases of Hashimoto's thyroiditis, Primary Myxedema, and Graves' disease. The relationship of autoimmune thyroid disease to pregnancy has been the subject of considerable interest with the recognition of the postpartum thyroid disease syndromes.[12] Anti-TPO antibodies are demonstrable in most cases of postpartum thyroiditis and it has been found that the presence of autoantibody in early pregnancy was associated with a high risk of asymptomatic postpartum hypothyroidism.[13-17]

It is common to find anti-TPO antibodies in the absence of autoantibodies to thyroglobulin, particularly in patients with small goitres and up to 64% of cases of autoimmune hypothyroidism have been reported to be associated with anti-TPO antibodies alone.[18] In addition, anti-TPO antibodies are frequently found in patients with other autoimmune diseases such as Rheumatoid Arthritis, Addison's Disease and Type I Diabetes.[19-21] They are also detectable at low levels in up to 20% of asymptomatic individuals,[22] particularly the elderly[23] and more often in women than in men, although the clinical significance of these autoantibodies is unclear.

BIOLOGICAL PRINCIPLES OF THE PROCEDURE

The ARCHITECT Anti-TPO assay is a two-step immunoassay for the quantitative determination of anti-TPO in human serum and plasma using CMIA technology with flexible assay protocols, referred to as Chemiflex.

1. Sample, assay diluent, and TPO coated paramagnetic microparticles are combined and incubated. The anti-TPO present in the sample binds to the TPO coated microparticles.
2. After washing, anti-human IgG acridinium-labeled conjugate is added to create a reaction mixture.
3. Following another incubation and wash, Pre-Trigger and Trigger Solutions are added to the reaction mixture.
4. The resulting chemiluminescent reaction is measured as relative light units (RLUs). There is a direct relationship between the amount of anti-TPO in the sample and the RLUs detected by the ARCHITECT iSystem optics.

For additional information on system and assay technology, refer to the ARCHITECT System Operations Manual, Section 3.

REAGENTS

Kit Contents

ARCHITECT Anti-TPO 2K47

NOTE: Some kit sizes are not available in all countries or for use on all ARCHITECT iSystems. Please contact your local distributor.

REF	2K47-25	2K47-20
Σ	100	400
MICROPARTICLES	1 x 6.6 mL	4 x 6.6 mL
CONJUGATE	1 x 5.9 mL	4 x 5.9 mL
ASSAY DILUENT	1 x 10.0 mL	4 x 10.0 mL

MICROPARTICLES Thyroid peroxidase (recombinant) coated microparticles in MES buffer with protein (bovine) stabilizer. Minimum concentration: 0.10 % solids. Preservative: antimicrobial agents.

CONJUGATE Anti-human IgG (mouse, monoclonal) acridinium-labeled conjugate in MES buffer with protein (bovine) stabilizer. Minimum concentration: 80.0 ng/mL. Preservative: antimicrobial agents.

ASSAY DILUENT Assay Diluent in MES buffer. Preservative: antimicrobial agents.

Other Reagents

PRE-TRIGGER SOLUTION ARCHITECT Pre-Trigger Solution containing 1.32% (w/v) hydrogen peroxide.

TRIGGER SOLUTION ARCHITECT Trigger Solution containing 0.35 N sodium hydroxide.

WASH BUFFER ARCHITECT Wash Buffer containing phosphate buffered saline solution. Preservatives: antimicrobial agents.

Warnings and Precautions

- IVD
- For *In Vitro* Diagnostic Use

Safety Precautions

CAUTION: This product requires the handling of human specimens. It is recommended that all human-sourced materials be considered potentially infectious and handled in accordance with the OSHA Standard on Bloodborne Pathogens. Biosafety Level 2 or other appropriate biosafety practices should be used for materials that contain or are suspected of containing infectious agents.[24-27]

The following warnings and precautions apply to: MICROPARTICLES	
	
WARNING	Contains potassium ferricyanide.
H361	Suspected of damaging fertility or the unborn child.
Prevention	
P201	Obtain special instructions before use.
P280	Wear protective gloves / protective clothing / eye protection.
Response	
P308+P313	IF exposed or concerned: Get medical advice / attention.
Disposal	
P501	Dispose of contents / container in accordance with local regulations.

The following warnings and precautions apply to: ASSAY DILUENT	
	
WARNING	
H319	Causes serious eye irritation.
Prevention	
P264	Wash hands thoroughly after handling.
P280	Wear protective gloves / protective clothing / eye protection.
Response	
P305+P351+P338	IF IN EYES: Rinse cautiously with water for several minutes. Remove contact lenses, if present and easy to do. Continue rinsing.
P337+P313	If eye irritation persists: Get medical advice / attention.

Safety Data Sheets are available at www.abbottdiagnostics.com or contact your local representative.

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

Reagent Handling

- Do not use reagent kits beyond the expiration date.
- **Do not pool reagents within a kit or between kits.**
- Before loading the reagent kit on the system for the first time, the microparticle bottle requires mixing to resuspend microparticles that may have settled during shipment. For microparticle mixing instructions, refer to the **PROCEDURE, Assay Procedure** section of this package insert.
- **Septums MUST be used to prevent reagent evaporation and contamination and to ensure reagent integrity. Reliability of assay results cannot be guaranteed if septums are not used according to the instructions in this package insert.**
  - **To avoid contamination, wear clean gloves when placing a septum on an uncapped reagent bottle.**
  - Once a septum has been placed on an open reagent bottle, **do not invert the bottle** as this will result in reagent leakage and may compromise assay results.
  - Over time, residual liquids may dry on the septum surface. These are typically dried salts and have no effect on assay efficacy.

For a detailed discussion of handling precautions during system operation, refer to the ARCHITECT System Operations Manual, Section 7.

Reagent Storage

When stored and handled as directed, reagents are stable until the expiration date.

	Storage Temperature	Maximum Storage Time	Additional Storage Instructions
Unopened/ Opened*	2-8°C	Until expiration date	May be used immediately after removal from 2-8°C storage. Store in upright position.
On board	System temperature	30 days	Discard after 30 days. For information on tracking onboard time, refer to the ARCHITECT System Operations Manual, Section 5.

* Reagents may be stored on or off the ARCHITECT iSystem. If reagents are removed from the system, store them at 2-8°C (with septums and replacement caps) in an upright position. For reagents stored off the system, it is recommended that they be stored in their original trays and boxes to ensure they remain upright. **If the microparticle bottle does not remain upright (with a septum installed) while in refrigerated storage off the system, the reagent kit must be discarded.** For information on unloading reagents, refer to the ARCHITECT System Operations Manual, Section 5.

After reagents are removed from the system, one must initiate a scan to update the onboard stability timer.

Indications of Reagent Deterioration

When a control value is out of the specified range, it may indicate deterioration of the reagents or errors in technique. Associated test results are invalid, and samples must be retested. Assay recalibration may be necessary. For troubleshooting information, refer to the ARCHITECT System Operations Manual, Section 10.

INSTRUMENT PROCEDURE

The ARCHITECT Anti-TPO assay file must be installed on the ARCHITECT iSystem prior to performing the assay. For detailed information on assay file installation and viewing and editing assay parameters, refer to the ARCHITECT System Operations Manual, Section 2.

For information on printing assay parameters, refer to the ARCHITECT System Operations Manual, Section 5.

For a detailed description of system procedures, refer to the ARCHITECT System Operations Manual.

The default result unit for the ARCHITECT Anti-TPO assay is IU/mL.

SPECIMEN COLLECTION AND PREPARATION FOR ANALYSIS

Specimen Types

Verified specimen types to be used with this assay:

	Collection Tubes	
Specimen Types	**Glass**	**Plastic**
Serum	No additive (uncoated)	Serum separator tubes
Plasma	Lithium heparin Plasma separator tubes with lithium heparin EDTA	Lithium heparin Plasma separator tubes with lithium heparin Sodium heparin EDTA

- Other specimen collection tube types have not been tested with this assay.
- When serial specimens are being evaluated, the same type of specimen should be used throughout the study.
- Performance has not been established for the use of cadaveric specimens or the use of body fluids other than human serum or plasma.

- The instrument does not provide the capability to verify specimen type. It is the responsibility of the operator to verify that the correct specimen types are used in the assay.

Specimen Conditions

- Do not use specimens with the following conditions:
  - heat-inactivated
  - obvious microbial contamination
- For accurate results, serum and plasma specimens should be free of fibrin, red blood cells, and other particulate matter. Serum specimens from patients receiving anticoagulant or thrombolytic therapy may contain fibrin due to incomplete clot formation.
- Ensure that complete clot formation in serum specimens has taken place prior to centrifugation. Some specimens, especially those from patients receiving anticoagulant or thrombolytic therapy, may exhibit increased clotting time. If the specimen is centrifuged before a complete clot forms, the presence of fibrin may cause erroneous results.
- To prevent cross contamination, use of disposable pipettes or pipette tips is recommended.

Preparation for Analysis

- Follow the tube manufacturer's processing instructions for specimen collection tubes.
- Patient specimens with a cloudy or turbid appearance must be centrifuged prior to testing. Following centrifugation, avoid the lipid layer (if present) when pipetting the specimen into a sample cup or secondary tube.
- Specimens must be mixed THOROUGHLY after thawing by LOW speed vortexing or by gently inverting, and centrifuged prior to use to remove red blood cells or particulate matter to ensure consistency in the results. Multiple freeze-thaw cycles of specimens should be avoided.
- All samples (patient specimens, controls, and calibrators) should be tested within 3 hours of being placed on board the ARCHITECT System. Refer to the ARCHITECT System Operations Manual, Section 5, for a more detailed discussion of onboard sample storage constraints.
- Inspect all specimens for bubbles. Remove bubbles with an applicator stick before analysis. Use a new applicator stick for each specimen to prevent cross contamination.

Specimen Storage

Specimen Type	Storage Temperature	Maximum Storage Time
Serum/Plasma	Room temperature	≤ 8 hours
	2-8°C	≤ 72 hours
	-10°C or colder	≤ 30 days

If testing will be delayed for more than 8 hours, remove serum or plasma from the serum or plasma separator, red blood cells or clot. Specimens removed from the separator gel, cells or clot may be stored up to 72 hours at 2-8°C.

Specimens can be stored up to 30 days frozen at -10°C or colder.

Specimen Shipping

- Package and label specimens in compliance with applicable state, federal, and international regulations covering the transport of clinical specimens and infectious substances.
- It is recommended that specimens be removed from the clot, red blood cells, or separator gel.
- Specimens may be shipped ambient for up to 8 hours after collection or on wet or dry ice for up to 72 hours after collection.
- Do not exceed the storage time limitations listed above.

PROCEDURE

Materials Provided

2K47 ARCHITECT Anti-TPO Reagent Kit

Materials Required but not Provided

- ARCHITECT Anti-TPO Assay file obtained from the ARCHITECT iSystem e-Assay CD-ROM found on www.abbottdiagnostics.com.
- 2K47-01 ARCHITECT Anti-TPO Calibrators
- 2K47-10 ARCHITECT Anti-TPO Controls
- ARCHITECT Pre-Trigger Solution
- ARCHITECT Trigger Solution
- ARCHITECT Wash Buffer
- ARCHITECT Reaction Vessels
- ARCHITECT Sample Cups
- ARCHITECT Septum
- ARCHITECT Replacement Caps
- Pipettes or pipette tips (optional) to deliver the volumes specified on the patient or control order screen.

For information on materials required for maintenance procedures, refer to the ARCHITECT System Operations Manual, Section 9.

Assay Procedure

- Before loading the reagent kit on the system for the first time, the microparticle bottle requires mixing to resuspend microparticles that may have settled during shipment. After the first time the microparticles have been loaded, no further mixing is required.
  - **Invert the microparticle bottle 30 times.**
  - Visually inspect the bottle to ensure microparticles are resuspended. If microparticles are still adhered to the bottle, continue to invert the bottle until the microparticles have been completely resuspended.
  - **If the microparticles do not resuspend, DO NOT USE. Contact your local Abbott representative.**
  - Once the microparticles have been resuspended, place a septum on the bottle. For instructions about placing septums on bottles, refer to the **Reagent Handling** section of this package insert.
- Load the reagent kit on the ARCHITECT iSystem.
  - Verify that all necessary reagents are present.
  - Ensure that septums are present on all reagent bottles.
- Order calibration, if necessary.
  - For information on ordering calibrations, refer to the ARCHITECT System Operations Manual, Section 6.
- Order tests.
  - For information on ordering patient specimens and controls and for general operating procedures, refer to the ARCHITECT System Operations Manual, Section 5.
- Minimum sample cup volume is calculated by the system and printed on the Orderlist report. Verify adequate sample cup volume is present prior to running the test.

  Maximum number of replicates sampled from the same sample cup: 10
  - Priority:

    Sample volume for first test: 75 µL

    Sample volume for each additional test from same sample cup: 25 µL
  - ≤ 3 hours on board:

    Sample volume for first test: 150 µL

    Sample volume for each additional test from same sample cup: 25 µL
  - To minimize the effects of evaporation, all samples (patient specimens, calibrators and controls) must be tested within 3 hours of being placed on board the ARCHITECT iSystem.
  - If using primary or aliquot tubes, use the sample gauge to ensure sufficient patient specimen is present.
- Prepare ARCHITECT Anti-TPO Calibrators and Controls

- ARCHITECT Anti-TPO Calibrators and Controls should be prepared according to their respective package inserts.
- Hold bottles **vertically** and dispense recommended volumes into each respective sample cup.
- Recommended volumes:
  for each calibrator: 5 drops
  for each control: 5 drops

- Load samples.
  - For information on loading samples, refer to the ARCHITECT System Operations Manual, Section 5.
- Press RUN. The system performs the following functions:
  - Moves the sample to the aspiration point.
  - Loads a reaction vessel (RV) into the process path.
  - Aspirates and transfers sample into the RV.
  - Advances the RV one position and transfers assay diluent and microparticles into the RV.
  - Mixes, incubates, and washes the reaction mixture.
  - Adds conjugate to the RV.
  - Mixes, incubates, and washes the reaction mixture.
  - Adds pre-trigger and trigger solutions.
  - Measures chemiluminescent emission to determine the quantity of anti-TPO in the sample.
  - Aspirates contents of RV to liquid waste and unloads RV to solid waste.
  - Calculates the result.
- For additional information on principles of operation, refer to the ARCHITECT System Operations Manual, Section 3.
- For optimal performance, it is important to perform routine maintenance as described in the ARCHITECT System Operations Manual, Section 9. Perform maintenance more frequently when required by laboratory procedures.

Specimen Dilution Procedures

Specimens with an anti-TPO value exceeding 1000.00 IU/mL are flagged with the code "> 1000.00" and may be diluted using the Automated Dilution Protocol or the Manual Dilution Procedure.

Automated Dilution Protocol

The system performs a 1:2 dilution of the specimen and automatically calculates the concentration of the specimen before dilution and reports the result.

Specimens with an anti-TPO value exceeding 2000.00 IU/mL are flagged with the code "> 2000.00" and may be diluted using the Manual Dilution Procedure.

Manual Dilution Procedure

Suggested dilution: 1:20

1. Prior to diluting the specimen, dispense approximately 10 drops of ARCHITECT Anti-TPO Calibrator A into a clean test tube for use in the next step.
2. Transfer 190 µL of ARCHITECT Anti-TPO Calibrator A from the test tube prepared in the prior step into another clean test tube and add 10 µL of the patient specimen.
3. The operator must enter the dilution factor in the Patient or Control order screen. The system will use this dilution factor to automatically calculate the concentration of the sample before dilution and report the result. The result should be > 5.61 IU/mL before the dilution factor is applied.

For detailed information on ordering dilutions, refer to the ARCHITECT System Operations Manual, Section 5.

Calibration

- Test Calibrators A-F in duplicate. The calibrators should be priority loaded.
  A single sample of each control level must be tested to evaluate the assay calibration. Ensure that assay control values are within the ranges specified in the respective control package insert.
- Calibration Range: 0.00 - 1000.00 IU/mL.
- Once an ARCHITECT Anti-TPO calibration is accepted and stored, all subsequent samples may be tested without further calibration unless:
  - A reagent kit with a new lot number is used or
  - Controls are out of range.
- For detailed information on how to perform an assay calibration, refer to the ARCHITECT System Operations Manual, Section 6.

Quality Control Procedures

The recommended control requirement for the ARCHITECT Anti-TPO assay is that a single sample of each control level be tested once every 24 hours each day of use. If the quality control procedures in your laboratory require more frequent use of controls to verify test results, follow your laboratory-specific procedures.

The ARCHITECT Anti-TPO Control values must be within the acceptable ranges specified in the control package insert. If a control is out of its specified range, the associated test results are invalid and must be retested. Recalibration may be indicated.

Verification of Assay Claims

For protocols to verify package insert claims, refer to the ARCHITECT System Operations Manual, Appendix B.

The ARCHITECT Anti-TPO assay belongs to method group 1.

RESULTS

Calculation

The ARCHITECT Anti-TPO assay utilizes a 4 Parameter Logistic Curve fit data reduction method (4PLC, Y-weighted) to generate a calibration curve.

Flags

Some results may contain information in the Flags field. For a description of the flags that may appear in this field, refer to the ARCHITECT System Operations Manual, Section 5.

LIMITATIONS OF THE PROCEDURE

- Antibody measurement represents one parameter in a multi-criteria diagnostic process. When making a diagnosis of thyroid disease, a combination of test methods should be used in conjunction with clinical symptoms.
- About 20% of asymptomatic specimens may present with anti-TPO autoantibodies reflecting the prevalence in apparently healthy populations. The prevalence of anti-TPO may also depend on age, gender, and geographic region of the selected population.
- Some specimens may not dilute linearly because of the heterogeneity of the autoantibodies with respect to physiochemical properties.
- Specimens from patients who have received preparations of mouse monoclonal antibodies for diagnosis or therapy may contain human anti-mouse antibodies (HAMA). Such specimens may show either falsely elevated or depressed values when tested with assay kits that employ mouse monoclonal antibodies. Assay results that are not consistent with other clinical observations may require additional information for diagnosis.[26-29]
- Heterophilic antibodies in human serum can react with reagent immunoglobulins, interfering with *in vitro* immunoassays. The presence of heterophilic antibodies in a patient specimen may cause anomalous values to be observed. Additional information may be required for diagnosis.[30]

EXPECTED VALUES

In a study, human serum specimens were collected from a population of 236 apparently healthy individuals. All specimens delivered TSH values within the normal reference range. Of this study population, 9 specimens delivered positive results on a commercially available anti-TPO assay device and were excluded from further normal range analysis. The 97.5 percentile concentration of the remaining population was 5.61 IU/mL. In this study population, the normal range

is < 5.61 IU/mL. A total of 97.8% (222/227) of the population gave values within this normal range.* This normal range is suggested as a guideline and each laboratory should establish a normal range appropriate to their patient populations, giving due consideration to age, gender, geographical location and their clinical practice.

* Representative data; results in individual laboratories may vary from these data.

SPECIFIC PERFORMANCE CHARACTERISTICS

Precision

The ARCHITECT Anti-TPO assay is designed to have an assay precision of ≤ 10% total CV for samples ≥ 5.61 IU/mL.

A study was performed for the ARCHITECT Anti-TPO assay with guidance from the National Committee for Clinical Laboratory Standards (NCCLS) Protocol EP5-A.[31] ARCHITECT Anti-TPO Positive Control and three human panels were assayed using three lots of reagents in replicates of two at two separate times per day for 20 days on three instruments. Each reagent lot used a single calibration curve throughout the study. Data from this study are summarized in the following table.*

Sample	Instrument	Reagent Lot	n	Mean Conc. Value (IU/mL)	Within Run SD	Within Run %CV	Total SD	Total %CV
Positive Control	1	A	80	74.85	2.11	2.8	2.16	2.9
		B	80	74.20	1.89	2.6	2.04	2.7
		C	80	74.63	2.01	2.7	2.17	2.9
	2	A	80	77.42	2.11	2.7	3.25	4.2
		B	80	75.32	1.92	2.5	2.54	3.4
		C	80	74.59	1.73	2.3	2.57	3.4
	3	A	80	75.41	2.13	2.8	2.52	3.3
		B	80	75.48	1.90	2.5	2.13	2.8
		C	80	76.66	2.42	3.2	2.87	3.7
Panel 1	1	A	80	1.57	0.08	4.8	0.10	6.5
		B	80	1.46	0.06	3.8	0.09	5.8
		C	80	1.64	0.09	5.6	0.10	6.1
	2	A	80	1.60	0.09	5.3	0.12	7.6
		B	80	1.53	0.06	3.9	0.11	7.2
		C	80	1.52	0.10	6.7	0.12	7.7
	3	A	80	1.47	0.08	5.3	0.11	7.8
		B	80	1.47	0.07	4.7	0.13	8.5
		C	80	1.52	0.14	9.5	0.15	9.8
Panel 2	1	A	80	20.98	0.65	3.1	0.76	3.6
		B	80	21.14	0.61	2.9	0.66	3.1
		C	80	21.51	0.71	3.3	0.75	3.5
	2	A	80	21.27	0.61	2.9	0.98	4.6
		B	80	21.62	0.66	3.0	0.90	4.2
		C	80	20.82	0.67	3.2	0.85	4.1
	3	A	80	21.00	0.73	3.5	0.86	4.1
		B	80	21.77	0.60	2.7	0.84	3.8
		C	80	21.24	0.70	3.3	0.89	4.2
Panel 3	1	A	80	214.78	5.14	2.4	6.48	3.0
		B	80	221.79	4.73	2.1	5.82	2.6
		C	80	216.71	5.36	2.5	6.36	2.9
	2	A	80	219.32	4.41	2.0	8.61	3.9
		B	80	224.54	4.04	1.8	13.37	6.0
		C	80	218.73	5.76	2.6	13.18	6.0
	3	A	80	212.91	6.11	2.9	6.84	3.2
		B	80	225.46	5.15	2.3	5.67	2.5
		C	80	228.17	5.80	2.5	7.09	3.1

* Representative data; results in individual laboratories may vary from these data.

Functional Sensitivity

In a study, human panels ranging in concentration from 0.16-1.20 IU/mL were tested in replicates of 2 over 10 days on one instrument using two reagent lots and three calibrations for a total of 40 replicates per panel. The total %CVs (combining variance components for replicate, run, day and reagent lot) were calculated and plotted against the mean concentration. A reciprocal curve was fitted through the data and the functional sensitivity value was calculated as the concentration corresponding to the 20% CV level of the fitted curve. The lowest ARCHITECT Anti-TPO assay value exhibiting a 20% CV is 0.50 IU/mL.*

* Representative data; results in individual laboratories may vary from these data.

Analytical Sensitivity

The ARCHITECT Anti-TPO assay is designed to have an analytical sensitivity of ≤ 1.0 IU/mL. The analytical sensitivity of the ARCHITECT Anti-TPO assay, defined as the concentration at two standard deviations above the ARCHITECT Anti-TPO Calibrator A (0.0 IU/mL) was calculated to be 0.16 IU/mL* at the 95% level of confidence (based upon one study with n=48 runs, 10 replicates of Calibrator A and 4 replicates of Calibrator B per run).

* Representative data; results in individual laboratories may vary from these data.

Linearity

The ARCHITECT Anti-TPO assay is linear between 3.0 and 1000.0 IU/mL based on a study performed with guidance from NCCLS protocol EP6-A.[32]

Autodilution Verification

The ARCHITECT Anti-TPO automated dilution protocol is designed to recover within 15% of manually diluted specimens. In a study, the automated dilution protocol (1:2) was compared to a manual 1:2 dilution procedure using 9 human specimens with anti-TPO levels that were greater than Calibrator E (250 IU/mL). The manual dilution was performed with ARCHITECT Anti-TPO Calibrator A. The observed percent recovery results are summarized in the following table.*

Sample ID	Automated Dilution (IU/mL)	Manual Dilution (IU/mL)	% Recovery**
1	861.25	859.27	100.2
2	684.49	703.64	97.3
3	844.36	847.62	99.6
4	724.55	757.09	95.7
5	709.46	688.49	103.1
6	1105.65	1106.18	100.0
7	948.43	931.80	101.8
8	840.77	851.72	98.7
9	966.48	998.45	96.8

* Representative data; results in individual laboratories may vary from these data.

$$** \% \text{ Recovery} = \frac{\text{Automated Dilution (IU/mL)}}{\text{Manual Dilution (IU/mL)}} \times 100$$

Interference

Interference from elevated levels of bilirubin, hemoglobin, triglycerides, and total protein in the ARCHITECT Anti-TPO assay is designed to be ≤ 15% at the levels indicated.

A study based on guidance from the NCCLS Protocol EP7-A[33] was performed for the ARCHITECT Anti-TPO assay. Specimens with anti-TPO levels between 45.07 and 361.64 IU/mL were supplemented with the following potentially interfering compounds. The average amount of interference observed during the study ranged from -3.6% to +3.7%.*

Potentially Interfering Substance	Potentially Interfering Substance Concentration
Bilirubin	20 mg/dL
Hemoglobin	1000 mg/dL
Total Protein (Low)	4 g/dL
Total Protein (High)	10 g/dL
Triglycerides	1000 mg/dL

* Representative data; results in individual laboratories may vary from these data.

Evaluation of Autoimmune Disease Specimens and High Titer IgG Samples

Potential interference from autoimmune disease specimens and high titer IgG samples in the ARCHITECT Anti-TPO assay is designed to be ≤ 15%. In a study, the ARCHITECT Anti-TPO assay was evaluated by testing specimens with known autoimmune diseases and elevated IgG. Specimens were evaluated with anti-TPO levels spiked between 131.44 and 568.78 IU/mL. Mean absolute % Interference is summarized in the following table.*

Clinical Condition	Mean Absolute % Interference
Anti-Nuclear Antibody (ANA)	1.6
Rheumatoid Arthritis (RA)	1.6
Systemic Lupus Erythematosus (SLE)	1.1
Insulin Dependent Diabetes Mellitus (IDDM)	1.0
Crohn's Disease	2.4
Multiple Sclerosis	1.7
Ulcerative Colitis	1.5
Hyperglobulinemia (high IgG)	0.9

* Representative data; results in individual laboratories may vary from these data.

Evaluation of Other Potential Interferents

Potential interference from HAMA and rheumatoid factor (RF) in the ARCHITECT Anti-TPO assay is designed to be ≤ 15%. In a study, the ARCHITECT Anti-TPO assay was evaluated by testing specimens with HAMA and RF to further assess the clinical specificity. Specimens positive for HAMA and specimens positive for RF were evaluated for % interference with anti-TPO levels spiked between 163.0 and 184.3 IU/mL. Mean absolute % interference is summarized in the following table.*

Other Potential Interferents	Number of Specimens	Mean Absolute % Interference
HAMA Positive	10	2.1
RF Positive	10	1.6

* Representative data; results in individual laboratories may vary from these data.

Clinical Sensitivity

In two studies, clinical sensitivity was evaluated by testing 139 clinically defined Hashimoto's thyroiditis specimens and 125 Graves' disease specimens. The clinical diagnosis was based on the criteria of the respective laboratory. The presence of autoantibodies against thyroglobulin and/or TPO was not necessarily a diagnostic criterion of these Graves' disease and Hashimoto's thyroiditis specimens. Data from these studies are summarized in the following table.*

	Hashimoto's Thyroiditis		Graves' Disease	
	n	% Positive	n	% Positive
Study 1	89	64.0	75	92.0
Study 2	50	74.0	50	100.0

* Representative data; results in individual laboratories may vary from these data.

Concordance

The performance of the ARCHITECT Anti-TPO was compared to a commercially available immunoassay for the determination of anti-TPO. A total of 500 specimens were evaluated in a study, encompassing a population of apparently healthy individuals and patients with autoimmune thyroid disease (Graves' disease and Hashimoto's thyroiditis). Specimens were tested in replicates of one using the ARCHITECT Anti-TPO assay with three reagent lots on three instruments and compared with a commercially available immunoassay (Comparison Assay). Data from this study are summarized in the following table.*

	Comparison Assay	
ARCHITECT Anti-TPO	Negative	Positive
Negative	242	32
Positive	5	221
Concordance = 92.6 %		

* Representative data; results in individual laboratories may vary from these data.

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Key to Symbols

	Consult instructions for use
	Manufacturer
	Sufficient for
	Temperature limitation
	Use by/Expiration date
ASSAY DILUENT	Assay Diluent
CONJUGATE	Conjugate
CONTROL NO.	Control Number
EC REP	Authorized Representative in the European Community
IVD	*In Vitro* Diagnostic Medical Device
LOT	Lot Number
MICROPARTICLES	Microparticles
PRE-TRIGGER SOLUTION	Pre-Trigger Solution
PRODUCED FOR ABBOTT BY	Produced for Abbott by
PRODUCT OF USA	Product of USA
REACTION VESSELS	Reaction Vessels
REAGENT LOT	Reagent Lot
REF	List Number
REPLACEMENT CAPS	Replacement Caps
SAMPLE CUPS	Sample Cups
SEPTUM	Septum
SN	Serial number
TRIGGER SOLUTION	Trigger Solution
WARNING: EYE IRRITANT	Warning: Causes serious eye irritation.
WARNING: REPRODUCTIVE HAZARD	Warning: Reproductive Hazard
WASH BUFFER	Wash Buffer


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Revised February 2015.

Abbott

ARCHITECT
Anti-TPO Calibrators

REF 2K47-01

en
Anti-TPO
2K47
G1-0435 / R03
S2K470

Revised April 2015.

INTENDED USE

The ARCHITECT Anti-TPO Calibrators are for the calibration of the ARCHITECT iSystem when used for the quantitative determination of the IgG class of thyroid peroxidase autoantibodies (anti-TPO) in human serum and plasma. Refer to the ARCHITECT Anti-TPO reagent package insert for additional information.

CONTENTS

6 Bottles (4.0 mL each) of ARCHITECT Anti-TPO Calibrators. Calibrator A contains phosphate buffer with protein (bovine) stabilizer. Calibrators B-F contain human plasma in phosphate buffer with protein (bovine) stabilizers. Preservative: antimicrobial agents.

The calibrators yield the following concentrations:

Calibrator	Anti-TPO Concentration (IU/mL)
CAL A	0.0
CAL B	5.0
CAL C	20.0
CAL D	62.5
CAL E	250.0
CAL F	1000.0

STANDARDIZATION

The ARCHITECT Anti-TPO Calibrators are standardized against International Reference Preparations MRC 66/387.

PRECAUTIONS

- IVD
- For *In Vitro* Diagnostic Use
- **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 these reagents and human specimens be handled in accordance with the OSHA Standard on Bloodborne Pathogens, Biosafety Level 2 or other appropriate biosafety practices should be used for materials that contain or are suspected of containing infectious agents.[1-4]
- Calibrators B-F contain human plasma, donor units of which have been tested and found to be reactive for anti-TPO and nonreactive for HBsAg, HIV-1 RNA or HIV-1 Ag, anti-HIV-1/HIV-2 and anti-HCV.

Safety Data Sheets are available at www.abbottdiagnostics.com or contact your local representative.

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

STORAGE

- Calibrators are stable until the expiration date when stored and handled as directed.
- Do not use past expiration date.
- -10°C

PREPARATION FOR ANALYSIS

- Self-defrosting freezers are not suitable for storage.
- Thaw completely at room temperature (15-30°C) for 45-60 minutes.
- Prior to use, mix THOROUGHLY by inversion 5-10 times.
- After each use, immediately return the thawed calibrators to refrigerated storage (2-8°C) for up to 30 days after thaw.

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Key to Symbols

	Caution
	Consult instructions for use
	Manufacturer
	Temperature limitation
	Use by/Expiration date
AFTER THAW	After thaw store at
CAL A	Calibrator (A,B,C,D,E or F)
EC REP	Authorized Representative in the European Community
INFORMATION FOR USA ONLY	Information needed for United States of America only
IVD	*In Vitro* Diagnostic Medical Device
LOT	Lot Number
PRODUCED FOR ABBOTT BY	Produced for Abbott by
PRODUCT OF USA	Product of USA
REF	List Number
UNTIL FIRST USE	Until first use store at

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Revised April 2015.

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ARCHITECT
Anti-TPO Controls

REF 2K47-10

en
Anti-TPO
2K47
JL840832 / R04
C2K470

Read Highlighted Changes: Revised April 2015.

INTENDED USE

The ARCHITECT Anti-TPO Controls are for the estimation of test precision and the detection of systematic analytical deviations of the ARCHITECT iSystem (reagents, calibrators and instrument) when used for the quantitative determination of the IgG class of thyroid peroxidase autoantibodies (anti-TPO) in human serum and plasma. Refer to the ARCHITECT Anti-TPO reagent package insert for additional information.

CONTENTS

2 Bottles (4.0 mL each) of ARCHITECT Anti-TPO Controls. The negative control and positive control contain human plasma in phosphate buffer with protein (bovine) stabilizers. Preservative: antimicrobial agents.

The following concentration ranges may be used for individual replicate control specifications on the ARCHITECT iSystem:

Control	Target Concentration (IU/mL)	Range (IU/mL)
CONTROL −	0.5	≤ 2.15
CONTROL +	75	45 - 105

Each laboratory should establish its own concentration ranges for new control lots at each control level. This can be accomplished by assaying a minimum of 20 replicates over several (3-5) days. Sources of variation that can be expected should be included in this study in order to be representative of future system performance. These may include:

- Multiple stored calibrations
- Multiple reagent lots
- Multiple calibrator lots
- Multiple processing modules
- Data points collected at different times of the day

These results should be applied to your laboratory's quality control practices.

ARCHITECT ANTI-TPO NEGATIVE CONTROL PROCEDURE

- Order the ARCHITECT Anti-TPO Negative Control as a patient sample. The negative control cannot be ordered as a control. This information pertains only to the ARCHITECT Anti-TPO Negative Control (2K47L).
- For the ARCHITECT Anti-TPO Negative Control, do not configure or use the Westgard or Levey-Jennings software screens for quality control analysis. The use of these screens for the negative control will result in data that is not statistically valid.
- Manual verification of the validity of the ARCHITECT Anti-TPO Negative Control is required any time the negative control is run. For control values that fall outside of the validity range listed in the ARCHITECT Anti-TPO Control Package Insert refer to the ARCHITECT System Operation Manual, Section 10 for troubleshooting information.
- Because the ARCHITECT Anti-TPO Negative Control is run as a patient sample, patient values will not be flagged by the ARCHITECT iSystem if a negative control is outside of its control range. Only release patient results if a valid negative control value is obtained.

PRECAUTIONS

- IVD
- For *In Vitro* Diagnostic Use
- **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 these reagents and human specimens be handled in accordance with the OSHA Standard on Bloodborne Pathogens. Biosafety Level 2 or other appropriate biosafety practices should be used for materials that contain or are suspected of containing infectious agents.[1-4]
- Controls contain human plasma, donor units of which have been tested and found to be reactive for anti-TPO and nonreactive for HBsAg, HIV-1 RNA or HIV-1 Ag, anti-HIV-1/HIV-2 and anti-HCV.

Safety Data Sheets are available at www.abbottdiagnostics.com or contact your local representative.

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

STORAGE

- Controls are stable until the expiration date when stored and handled as directed.
- Do not use past expiration date.
- -10°C

PREPARATION FOR ANALYSIS

- Self-defrosting freezers are not suitable for storage.
- Thaw completely at room temperature (15-30°C) for 45-60 minutes.
- Prior to use, mix THOROUGHLY by inversion 5-10 times.
- After each use, immediately return the thawed controls to refrigerated storage (2-8°C) for up to 30 days after thaw.

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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–Third Edition*. CLSI Document M29-A3. Wayne, PA: CLSI; 2005.

Key to Symbols

	Caution
	Consult instructions for use
	Manufacturer
	Temperature limitation
	Use by/Expiration date
AFTER THAW	After thaw store at
CONC	Concentration
CONTROL -	Negative Control
CONTROL +	Positive Control
EC REP	Authorized Representative in the European Community
INFORMATION FOR USA ONLY	Information needed for United States of America only
IVD	*In Vitro* Diagnostic Medical Device
LOT	Lot Number
PRODUCED FOR ABBOTT BY	Produced for Abbott by
PRODUCT OF USA	Product of USA
RANGE	Range
REF	List Number
UNTIL FIRST USE	Until first use store at

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