



07404085001V1.0

# TPLA2 Calibrator Set

Mediace TPLA Gen.2 Calibrator Set

REF 07404085 190

5 × 2 mL Calibrator

SEKISUI

## English

### System information

For use on Roche/Hitachi **cobas c** analyzers the calibrator codes are 770-774.

Please note: For technical reasons it is necessary to specify dummy units (mg/dL) on Roche/Hitachi **cobas c** analyzers. The values can be converted to the correct units via the host computer.

### Intended use

Mediace TPLA Gen.2 Calibrator Set contains 5 calibrators for use in the calibration of the Sekisui "Mediace TPLA Gen.2" immunoturbidimetric assay (Cat. No. 07404182190).

### Summary

The Mediace TPLA Gen.2 Calibrator Set contains 5 calibrators based on human serum and bovine serum albumin (BSA).

The concentrations of the calibrator components have been adjusted to ensure optimal calibration of the Sekisui "Mediace TPLA Gen.2" assay on clinical chemistry analyzers.

### Reagents – working solutions

#### Reactive components:

Human serum and bovine serum albumin (BSA) with chemical additives and material of biological origin as specified.

The origin of the biological additive is as follows:

Analyte	Origin
Anti-Treponema pallidum antibodies	human

#### Non-reactive components:

BSA; gelatin; stabilizers and preservatives.

The antibody levels are lot-specific. The exact calibrator values are given on the label of each vial and in the electronically available or enclosed value sheets.

The values are also encoded in electronic files sent via the **cobas** link to the analyzers.

### Calibrator values<sup>1</sup>

The calibrator values were determined using the Sekisui Mediace TPLA Gen.2 immunoturbidimetric method.

Traceability information is given in the relevant Method Sheets for the system reagents.

### Precautions and warnings

For in vitro diagnostic use.

Exercise the normal precautions required for handling all laboratory reagents.

Calibrators contain < 0.1 % sodium azide as a preservative. Sodium azide may react with lead and copper plumbing to form potentially explosive metal azide buildup. Flush with copious amounts of water when discarding material.

Disposal of all waste material should be in accordance with local guidelines. Safety data sheet available for professional user on request.

All human material should be considered potentially infectious. All products derived from human blood are prepared exclusively from the blood of donors tested individually and shown to be free from HBsAg and antibodies to HCV and HIV.

The testing methods applied were FDA-approved or cleared in compliance with the European Directive 98/79/EC, Annex II, List A.

However, as no testing method can rule out the potential risk of infection with absolute certainty, the material should be handled with the same level of care as a patient specimen. In the event of exposure, the directives of the responsible health authorities should be followed.<sup>2,3</sup>

### Handling

Calibrator bottles 1-5 are ready for use. Mix carefully before use. Avoid the formation of foam.

The enclosed barcoded labels are intended exclusively for Roche/Hitachi **cobas c** systems to identify the calibrator. Attach the barcoded labels to the tubes carrying the sample cups containing the calibrator material.

### Storage and stability<sup>1</sup>

Storage:

Store at 2-8 °C.

Do not freeze.

Criterion for the stability data stated:  
Recovery within ± 15 % of initial value.

Stability:

Unopened: up to the stated expiration date at 2-8 °C

After opening: 8 weeks at 2-8 °C

Store calibrator tightly capped when not in use.

### Materials provided

- See "Reagents – working solutions" section
- Barcoded labels

### Materials required (but not provided)

- System reagents and clinical chemistry analyzers
- General laboratory equipment

### Assay

Use Mediace TPLA Gen.2 Calibrator Set as specified in the relevant Method Sheet for the system reagents.

### References

- Data on file at Sekisui Medical Co., Ltd.
- Occupational Safety and Health Standards: bloodborne pathogens. (29 CFR Part 1910.1030). Fed. Register.
- Directive 2000/54/EC of the European Parliament and Council of 18 September 2000 on the protection of workers from risks related to exposure to biological agents at work.

A point (period/stop) is always used in this Method Sheet as the decimal separator to mark the border between the integral and the fractional parts of a decimal numeral. Separators for thousands are not used.

### Symbols

Roche Diagnostics uses the following symbols and signs in addition to those listed in the ISO 15223-1 standard.

CONTENT	Contents of kit
CALIBRATOR	Calibrator
→	Volume after reconstitution or mixing
GTIN	Global Trade Item Number

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Additions, deletions or changes are indicated by a change bar in the margin.

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