LZI Oxycodone Calibrators

300 ng/mL Cutoff





IVD For In Vitro Diagnostic Use Only

Lin-Zhi International, Inc.

REF	Description	Quantity
0001	LZI Universal Negative Calibrator	1 x 5 mL
0622	LZI Oxycodone 300 Low Calibrator (150 ng/mL)	1 x 5 mL
0623	LZI Oxycodone 300 Cutoff Calibrator (300 ng/mL)	1 x 5 mL
0624	LZI Oxycodone 300 Intermediate Calibrator (500 ng/mL)	1 x 5 mL
0625	LZI Oxycodone 300 High Calibrator (800 ng/mL)	1 x 5 mL

Intended Use

The LZI Oxycodone 300 Calibrators are for use as calibrators in the qualitative and semi-quantitative calibration of the LZI Oxycodone III Enzyme Immunoassay (Ref# 0610/0611) at the cutoff value of 300 ng/mL on automated clinical chemistry analyzers (1).

Description of the Calibrators:

The LZI Oxycodone 300 Calibrators are human urine-based liquids and ready-to-use. The LZI Universal Negative Calibrator is a processed drug-free human urine matrix containing buffers, stabilizers, and less than 0.1 % of sodium azide. The calibrators are prepared by spiking known concentrations of oxycodone into the negative calibrator.

*Actual concentrations of these calibrators are confirmed by GC/MS or LC/MS. Values are provided only as guidelines and laboratories should determine the ranges based on their own test system and tolerance (2).

Precautions and Warning

- The LZI Oxycodone 300 Calibrators are for in vitro diagnostic use only. Harmful if swallowed.
- The calibrators contain sodium azide, which may react with lead or copper plumbing to form potentially explosive metal azide. When disposing such liquids always flush with a large volume of water to prevent azide build-up (3).
- The calibrators are prepared from non-sterile human urine. They are not tested by licensed reagents for the presence of antibodies to human immunodeficiency viruses, the hepatitis antigens, and/or anti- hepatitis antibodies. They should be handled as potentially infectious. Always use good laboratory practice to avoid any skin contact or ingestion.
- <u>Do not use the calibrators beyond their expiration dates.</u>
- For USA: Caution: Federal law restricts this device to sale by or on the order of a physician.

Preparation and Storage

The calibrators are ready-to-use. No reconstitution is required. Label the cap before removal to identify it with the original bottle. The calibrators should be stored refrigerated at 2-8°C when not in use.

Stability

When stored refrigerated at 2-8°C, the calibrators are stable either opened-recapped or unopened, until the expiration date printed on the vial label. Store calibrators tightly capped when not in use. Calibrator solution dispensed in the sample cups and left on board a clinical analyzer should be discarded after use.

Procedure and Results

For qualitative calibration, use the 300 ng/mL as the cutoff calibrator. For semi-quantitative calibration, use all five calibrators. Recalibration should be performed after reagent bottle change, a change in calibrators or reagent lot, and after instrument maintenance is performed. For interpretation of results, refer to the appropriate LZI Oxycodone III Enzyme Immunoassay (Ref# 0610/0611) package insert (1).

Limitations

The LZI Oxycodone 300 Calibrators are for use with the LZI Oxycodone III Enzyme Immunoassay (Ref# 0610/0611) to detect oxycodone in human urine only.

Symbols Used

EC REP	Authorized Representative	CONTENTS	Contents	REF	Reference Number
₩	Biological Risks	GTIN	Global Trade Item Number	SDS	Safety Data Sheet
CALIBRATOR	Calibrator	IVD	In Vitro Diagnostic medical device	2°C	Temperature Limits
CE	CE Mark	LOT	Lot Number	T.K.	Test Kit Number
Πi	Consult Instructions for Use	**	Manufacturer	\subseteq	Use-by Date

Bibliography

- LZI Oxycodone III Enzyme Immunoassay (Ref# 0610/0611) package insert.
- Guidance for Industry, Abbreviated 510(k) Submissions for In Vitro Diagnostic Calibrators. U.S. Department of Health and Human Services. FDA, Document issued on February 22, 1999.
- Sodium Azide. National Institute for Occupational Safety (NIOSH). Pocket Guide to Chemical Hazards. Third Printing, September 2007. Available online at: https://www.cdc.gov/niosh/npg/default.html

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

Notice: Adulteration of reagents, use of instruments without appropriate capabilities, or other failure to follow instructions as set forth in this labeling can affect performance characteristics, and stated or implied label claims.



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EC REP

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