

LZI Benzodiazepine Calibrators



IVD For In Vitro Diagnostic Use Only



Lin-Zhi International, Inc.

REF	Description	Quantity
0001	Negative Calibrator	1 x 5 mL
0132	Benzodiazepine 100 ng/mL Calibrator #2/Control 1	1 x 5 mL
0133	Benzodiazepine 200 ng/mL Calibrator #3/Cutoff A/Control 2	1 x 5 mL
0134	Benzodiazepine 300 ng/mL Calibrator #4/Cutoff B/Control 3	1 x 5 mL
0135	Benzodiazepine 1000 ng/mL Calibrator #5	1 x 5 mL

Intended Use

The Lin-Zhi International (LZI) Benzodiazepine Calibrators are for use as calibrators in the qualitative and semi-quantitative calibration of the LZI Benzodiazepine Enzyme Immunoassay (Ref# 0130/0131) on a number of automated clinical chemistry analyzers.

Description of the Calibrators:

The LZI Benzodiazepine Calibrators are human urine-based liquids, and ready to use. The 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 oxazepam into the Negative Calibrator. Calibrators are made from NIST traceable standards.

*Actual concentrations of these calibrators are within ± 10 % of the target value as determined 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 Benzodiazepine 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.
- 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 apply good laboratory practice to avoid any skin contact or ingestion.
- Do not use the calibrators beyond their expiration dates.
- For USA: Caution: Federal law restricts this device to sale by or on the order of a physician.

Preparation and Storage

The calibrators are provided 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 of the clinical analyzer should be discarded after use.

Procedure and Results

For qualitative calibration, use the 200 ng/mL or 300 ng/mL as your cutoff calibrator. For semi-quantitative calibration, use all five calibrators. Recalibration should be performed after reagent bottle change or there is a change in calibrators or reagent lot, and after instrument maintenance is performed. For interpretation of results, refer to the appropriate LZI Benzodiazepine Enzyme Immunoassay (Ref# 0130/0131) package insert (3).

Limitations

The LZI Benzodiazepine Calibrators are for use with the LZI Benzodiazepine Enzyme Immunoassay (Ref# 0130/0131) to detect benzodiazepines in human urine only.

Bibliography

1. Urine testing for Drug of Abuse. National Institute on Drug Abuse (NIDA) Research Monograph 73, 1986.
2. 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.
3. LZI Benzodiazepine Enzyme Immunoassay (Ref# 0130/0131) package insert.

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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