

LZI Norfentanyl Controls



IVD For In Vitro Diagnostic Use Only



Lin-Zhi International, Inc.

REF		Description	Quantity
0317		LZI Norfentanyl Level 1 Control (3.75 ng/mL)	1 x 5 mL
0318		LZI Norfentanyl Level 2 Control (6.25 ng/mL)	1 x 5 mL

Intended Use

The LZI Norfentanyl Controls are for use as assayed quality control materials to monitor the precision of the LZI Fentanyl and LZI Fentanyl II Enzyme Immunoassay (Ref# 0310/0311 and Ref# 0570/0571) on a number of automated clinical chemistry analyzers. (1, 2)

Description of the Controls

The LZI Norfentanyl Controls are human urine-based liquids and ready-to-use. The constituent is a processed drug-free human urine matrix containing buffers, stabilizers, and less than 0.1 % of sodium azide. The controls are prepared by spiking known concentrations of norfentanyl into the drug-free matrix.

*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 (3).

Precautions and Warning

- The LZI Norfentanyl Controls are for in vitro diagnostic use only. Harmful if swallowed.
- The controls 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 (4).
- The controls 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.
- Do not use the controls beyond their expiration dates.
- For USA: Cautions: Federal law restricts this device to sale by or on the order of a physician.

Preparation and Storage

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

Stability

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

Procedure and Results

Both levels of controls (3.75 ng/mL and 6.25 ng/mL) should be run daily to ensure proper assay performance. Additionally, run the controls with each new calibration and after specific maintenance or troubleshooting procedures are performed. For interpretation of results, refer to the appropriate LZI Fentanyl and LZI Fentanyl II Enzyme Immunoassay (Ref# 0310/0311 and Ref# 0570/0571) package insert. (1, 2)

Limitations

The LZI Norfentanyl Controls are for use with the LZI Fentanyl and LZI Fentanyl II Enzyme Immunoassay (Ref# 0310/0311 and Ref# 0570/0571) to detect norfentanyl in human urine only. Quality control materials should be used in accordance with local, state, and/or federal regulations or accreditation requirements. (1, 2)

Symbols Used

	Authorized Representative		In Vitro Diagnostic medical device		Safety Data Sheet
	Biological Risks		Lot Number		Temperature Limits
	CE Mark		Manufacturer	T.K.	Test Kit Number
	Consult Instructions for Use		Negative Control		Use-by Date
	Contents		Positive Control		
	Global Trade Item Number		Reference Number		

Bibliography

1. LZI Fentanyl Enzyme Immunoassay (Ref# 0310/0311) package insert.
2. LZI Fentanyl II Enzyme Immunoassay (Ref# 0570/0571) package insert.
3. Guidance for Industry and FDA Staff, Assayed and Unassayed Quality Control Material. U.S. Department of Health and Human Services. FDA, Document issued on June 7, 2007.
4. 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/npq/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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