

Lysergic Acid Diethylamide (LSD) In Urine For GC or GC/MS Confirmations Using: 200 mg Clean Screen® Extraction Column



UCT Part Numbers

ZSDAU020		ZCAU020
Clean Screen® DAU	Or	Clean Screen® DAU
10 mL, 200 mg sorbent		10 mL, 200 mg sorbent
Without Tips		With CLEAN-THRU® Tips

Procedure:

1. Prepare Sample

- To 2 mL 100 mM phosphate buffer (pH= 6.0) add internal standard. Add 5 mL of urine.
- Mix/vortex.
- Sample pH should be 6.0 ± 0.5 .
- Adjust pH accordingly with 100 mM monobasic or dibasic sodium phosphate.
- Centrifuge as appropriate.

2. Condition Clean Screen® Extraction Column

- 1 x 3 mL CH_3OH .
- 1 x 3 mL D.I. H_2O .
- 1 x 1 mL 100 mM phosphate buffer (pH = 6.0).

Note: Aspirate at < 3 inches Hg to prevent sorbent drying.

3. Apply Sample

- Load at 1 mL/minute.

4. Wash Column

- 1 x 3 mL D.I. H_2O .
- 1 x 1 mL 100 mM acetic acid.
- 1 x 3 mL CH_3OH .
- Dry column (5 minutes at > 10 inches Hg).

5. Elute LSD

- 1 x 3 mL $\text{CH}_2\text{Cl}_2/\text{IPA}/\text{NH}_4\text{OH}$ (78:20:2);
 - Collect eluate at 1 to 2 mL/minute.
- Note:** Prepare elution solvent daily. Add IPA/ NH_4OH , mix, then add CH_2Cl_2 (pH 11-12).

6. Dry Eluate

- Evaporate to dryness at < 40°C.

7. Derivatize

- Add 20 μL ethyl acetate and 20 μL BSTFA (with 1% TMCS)***.
- Overlay with N_2 and cap. Mix/vortex.
- React 20 minutes at 70°C.

Remove from heat source to cool.

Note: Do not evaporate BSTFA solution.

8. Quantitate

- Inject 1 to 2 μL onto gas chromatograph.
- For MSD monitor the following ions:

Analyte	Primary Ion***	Secondary	Tertiary	Cerilliant #
LSD-D ₃ -TMS*	298	296	271	L-006
LSD-TMS	395	293	268	L-005

* Suggested internal standard for GC/MS: D₃-LSD

*** Part # SBSTFA-1-1,10,25,100

**** Quantitation ion



UCT, LLC • 2731 Bartram Road • Bristol, PA 19007 800.385.3153 • 215.781.9255

www.unitedchem.com Email: methods@unitedchem.com

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