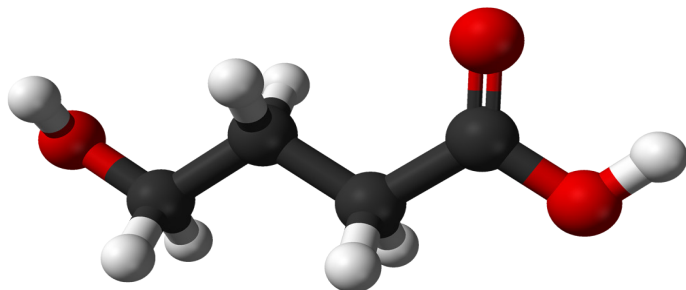


A Solid-Phase Method For Gamma-Hydroxybutyrate (GHB) In Urine Without Conversion To Gamma-Butyrolactone (GBL)



UCT Part Numbers

ZSGHB020
Clean Screen® GHB
200 mg, 10 mL

SBSTFA-1-1
Selectra-Sil BSTFA
with 1% TMCS 1g vial

Procedure:

1. Prepare Sample

- To 200 μ L of urine add internal standard and 100 μ L of 100 mM phosphate buffer (pH 6.0). Mix/vortex

2. Condition Clean Screen® GHB 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. Load Sample

- Place test tubes into vacuum manifold for collection
- The sample loading and wash are both collected
- Decant sample onto column. Aspirate at ~1 inch Hg

4. Wash Column

- Add 1 mL of CH_3OH / NH_4OH (99:1) to original sample test tube; Vortex
- Decant wash onto column

Note: Aspirate at ~1 inch of Hg

5. Concentrate

- Evaporate to dryness at 60°C using a stream of air or N_2

6. Sample Clean Up

- Add 200 μ L of dimethylformamide
- Add 1 mL of hexane saturated with dimethylformamide
- Mix by inversion for 5 minutes
- Centrifuge at 3000 rpm for 5 minutes
- Transfer lower dimethylformamide layer to a clean test tube

7. Concentrate

- Evaporate to dryness at < 50°C using a stream of air or N_2

8. Derivatize

- Add 100 μ L ethyl acetate and 100 μ L BSTFA with 1% TMCS.
- Mix/vortex.

9. Quantitate

- Inject 1 to 2 μ L onto gas chromatograph.



MSD Ions:

Analyte	Primary Ion	Secondary	Tertiary	Cerilliant #
GHB-D ₆ -di-TMS	239	240	241	G-006
GHB-di-TMS	233	234	235	G-001

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