

**Overview**
**Useful For**

Assessing compliance

Monitoring for appropriate therapeutic level

Assessing toxicity

**Method Name**

Liquid Chromatography-Tandem Mass Spectrometry (LC-MS/MS)

**NY State Available**

Yes

**Specimen**
**Specimen Type**

Serum Red

**Specimen Required**

**Container/Tube:** Red top

**Specimen Volume:** 0.5 mL

**Forms**

If not ordering electronically, complete, print, and send 1 of the following forms with the specimen:

[-Neurology Specialty Testing Client Test Request \(T732\)](#)

[-Therapeutics Test Request \(T831\)](#)

**Specimen Minimum Volume**

0.3 mL

**Reject Due To**

Gross hemolysis	OK
Gross lipemia	OK
Gross icterus	OK

**Specimen Stability Information**

Specimen Type	Temperature	Time	Special Container
Serum Red	Refrigerated (preferred)	14 days	
	Ambient	14 days	
	Frozen	14 days	

## Clinical and Interpretive

### Clinical Information

Diazepam, a benzodiazepine derivative, is an anxiolytic agent that reduces neuronal depolarization resulting in decreased action potentials. It enhances the action of gamma-aminobutyric acid (GABA) by tightly binding to A-type GABA receptors, thus opening the membrane channels and allowing the entry of chloride ions. It is also used as a muscle relaxant, procedural sedation agent, and sedative-hypnotic agent to treat withdrawal states (ie, ethanol), along with other conditions (seizures).

Diazepam is metabolized to several metabolites in the liver including temazepam, oxazepam, and nordiazepam (desmethyldiazepam) and the clearance of the drug is reduced considerably in the elderly and in patients with hepatic disease.

Therapeutic assessment typically includes measurement of both the parent drug (diazepam) and the active metabolite (nordiazepam).

### Reference Values

[Therapeutic concentrations](#)

Diazepam and Nordiazepam: 200-2,500 ng/mL

### Interpretation

For seizures:

Serum concentrations are not usually monitored during early therapy because response to the drug can be monitored clinically as seizure control. If seizures resume despite adequate therapy, another anticonvulsant must be considered.

Toxicity is commonly seen when diazepam plus nordiazepam concentrations exceed 3,000 ng/mL. Adverse effects of benzodiazepines in therapeutic doses usually reflect the drug's pharmacology and include sedation, slurred speech, and ataxia. Respiratory depression/arrest may occur with large overdoses or following rapid IV injection with short-acting benzodiazepines.

### Cautions

No significant cautionary statements

### Clinical Reference

1. Langman LJ, Bechtel L, Holstege CP: Chapter 35: Clinical Toxicology, In Tietz Textbook of Clinical Chemistry and Molecular Diagnostics. Edited by CA Burtis, ER Ashwood, DE Bruns. WB Saunders Company, Philadelphia, PA, 2011, pp 1109-1188
2. Tietz Textbook of Clinical Chemistry and Molecular Diagnostics, Edited by CA Burtis, ER Ashwood, DE Bruns, WB Saunders Company, Philadelphia, PA, 2011, Table 60.2, pp 1109-1188

## Performance

### Method Description

The internal standard mixture containing chlordiazepoxide-d5, diazepam-d4, and nordiazepam-d5 is added to serum samples. The serum samples are treated with 0.1M phosphate buffer and extracted via liquid/liquid extraction with 50/50 (hexane/ethyl acetate). The organic layer from the extraction is dried under nitrogen, reconstituted in 95/5 (H2O/Methanol) and injected on a liquid chromatography-tandem mass spectrometer (LC-MS/MS). (Unpublished Mayo method)

**PDF Report**

No

**Day(s) and Time(s) Test Performed**

Friday, Varies

**Analytic Time**

Same day/1 day

**Maximum Laboratory Time**

7 days

**Specimen Retention Time**

14 days

**Performing Laboratory Location**

Rochester

**Fees and Codes****Fees**

- Authorized users can sign in to [Test Prices](#) for detailed fee information.
- Clients without access to Test Prices can contact [Customer Service](#) 24 hours a day, seven days a week.
- Prospective clients should contact their Regional Manager. For assistance, contact [Customer Service](#).

**Test Classification**

This test was developed and its performance characteristics determined by Mayo Clinic in a manner consistent with CLIA requirements. This test has not been cleared or approved by the U.S. Food and Drug Administration.

**CPT Code Information**

80346

G0480 (if appropriate)

**LOINC® Information**

Test ID	Test Order Name	Order LOINC Value
DIA	Diazepam and Nordiazepam, S	49044-1

Result ID	Test Result Name	Result LOINC Value
8629	Diazepam	3548-5
2475	Nordiazepam	3537-8



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Result ID	Test Result Name	Result LOINC Value
2459	Diazepam and Nordiazepam	16757-7