

Test Definition: FSERT

Sertraline (Zoloft) and Desmethylsertraline

Reporting Title: Sertraline (Zoloft) **Performing Location:** Medtox Laboratories, Inc.

Specimen Requirements: Submit only 1 of the following specimens:

Plasma

Specimen Type: Plasma Container/Tube: Green Top Specimen Volume: 2 mL Collection Instructions: Draw blood in a green-top (sodium heparin) tube, **plasma gel tube is not acceptable.** Spin down and send 2 mL of sodium heparin plasma refrigerated in a plastic vial.

Serum

Specimen Type: Serum Container/Tube: Red Specimen Volume: 2 mL

Collection Instructions: Draw blood in a plain, red-top tube, **serum gel tube is not acceptable.** Spin down and send 2 mL of serum refrigerated in a plastic vial.

Specimen Type	Temperature	Time	Special Container
Varies	Refrigerated (preferred)	7 days	
	Frozen	180 days	
	Ambient	72 hours	

Result Codes:

Result ID	Reporting Name	Туре	Unit	LOINC®
Z2298	Sertraline	Alphanumeric		6906-2
Z2299	Desmethylsertraline	Alphanumeric		6897-3

LOINC® and CPT codes are provided by the performing laboratory.

Supplemental Report:

No

CPT Code Information:

80332

Reference Values:

Sertraline:



Test Definition: FSERT

Sertraline (Zoloft) and Desmethylsertraline

Reference Range: 30 - 200 ng/mL Report Limit 10 ng/mL

Desmethylsertraline: ng/mL No reference range provided

The stated reference range is the range of observed steady-state concentrations in individuals receiving therapeutic dosage regimens of sertraline. This is not a defined therapeutic range. Report Limit 10 ng/mL