



## PAROXETINE IN BLOOD, PLASMA/ SERUM AND URINE. LC-MSMS CONFIRMATIONS USING 200 mg CLEAN SCREEN<sup>®</sup> DAU EXTRACTION COLUMN

Part #:

ZSDAU020 – CLEAN SCREEN<sup>®</sup> DAU 200 mg, 10 mL Tube

### 1. PREPARE SAMPLE:

To 1 mL of 100 mM phosphate buffer (pH 6.0) add internal standards\*.  
Add 1 mL whole blood, Serum/Plasma or Urine. Add 2 mL of 100 mM phosphate buffer (pH 6.0).  
Vortex and centrifuge as appropriate.

### 2. CONDITION COLUMN:

1 x 3 mL CH<sub>3</sub>OH  
1 x 3 mL D.I. H<sub>2</sub>O  
1 x 3 mL 100 mM phosphate buffer (pH 6.0).  
**Note:** aspirate at < 3 inches Hg to prevent sorbent drying out.

### 3. APPLY SAMPLE:

Load sample at 1-2 mL / minute.

### 4. WASH COLUMN:

1 x 3 mL D.I. H<sub>2</sub>O  
1 x 3 mL 100 mM acetic acid  
1 x 3 mL CH<sub>3</sub>OH  
Dry column (5 minutes at > 10 inches Hg).

### 5. ELUTE PAROXETINE:

1 x 3 mL Ethyl Acetate: Acetonitrile: NH<sub>4</sub>OH (78:20:2)  
Collect eluate at 1-2 mL / minute.

### 6. EVAPORATION:

Evaporate eluates under a gentle stream of nitrogen < 40 °C  
Dissolve residue in 100 µL Methanol.

### INSTRUMENT CONDITIONS:

**Column:** 50 x 2.1 mm (3 µm) Selectra<sup>®</sup> Phenyl (UCT, LLC)

Mobile phase:	<u>Time</u>	<u>Acetonitrile</u>	<u>0.1% Formic Acid aq</u>
	0	10	90
	15	50	50
	16	10	90
	20	10	10

**Flow rate:** 0.35 mL/ minute

**Injection Volume:** 5 µL

**Column Temperature:** ambient

**Detector:** API 2000 MS/MS.

<u>Compound</u>	<u>MRM Transition</u>
Paroxetine	330.0 / 190.1
Paroxetine-D6	336.0 / 76.1

CHROMATOGRAM :

