

Test Definition: PERAM

Perampanel, Serum

Overview

Useful For

Monitoring serum concentration of perampanel, in specific clinical conditions (ie, severe kidney impairment, mild to moderate hepatic impairment, and end-stage kidney disease)

Assessing compliance

Assessing potential toxicity

Method Name

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

NY State Available

Yes

Specimen

Specimen Type

Serum Red

Specimen Required

Collection Container/Tube: Red top (serum gel/SST is not acceptable)

Submission Container/Tube: Plastic vial

Specimen Volume: 1 mL **Collection Instructions:**

- 1. Draw blood immediately before next scheduled dose.
- 2. For sustained-release formulations ONLY, draw blood a minimum of 12 hours after last dose.
- 3. Centrifuge and aliquot serum into a plastic vial within 2 hours of collection.

Forms

If not ordering electronically, complete, print, and send a Therapeutics Test Request (T831) with the specimen.

Specimen Minimum Volume

0.5 mL

Reject Due To

Gross	OK
hemolysis	
Gross lipemia	ОК
Gross icterus	ОК



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Specimen Stability Information

Specimen Type	Temperature	Time	Special Container
Serum Red	Refrigerated (preferred)	28 days	
	Ambient	28 days	
	Frozen	28 days	

Clinical & Interpretive

Clinical Information

Perampanel (Fycompa) is approved for adjunctive therapy to treat primary generalized tonic-clonic seizures in patients aged 12 years and older as well as the treatment of partial-onset seizures with or without secondarily generalized seizures in patients with epilepsy 4 years of age and older.

Reference Values

180-980 ng/mL

Interpretation

The serum concentration should be interpreted in the context of the patient's clinical response and may provide useful information in patients showing poor response or adverse effects, particularly when perampanel is coadministered with other anticonvulsant drugs.

Most individuals display optimal response to perampanel with serum levels 180 to 980 ng/mL. Some individuals may respond well outside of this range or may display toxicity within the therapeutic range; thus, interpretation should include clinical evaluation. Toxic levels have not been well established. Therapeutic ranges are based on specimen collected at trough (ie, immediately before the next dose).

Cautions

This test cannot be performed on whole blood.

Clinical Reference

- 1. Reimers A, Berg JA, Burns ML, Brodtkorb E, Johannessen SI, Johannessen Landmark C: Reference ranges for antiepileptic drugs revisited: a practical approach to establish national guidelines. Drug Des Devel Ther. 2018 Feb 8;12:271-280. doi: 10.2147/DDDT.S154388
- 2. Hiemke C, Bergemann N, Clement HW, et al: Consensus guidelines for therapeutic drug monitoring in neuropsychopharmacology: Update 2017. Pharmacopsychiatry. 2018 Jan;51(1-02):9-62. doi: 10.1055/s-0043-116492
- 3. Rifai N, Horvath AR, Wittwer CT, eds: Tietz Textbook of Clinical Chemistry and Molecular Diagnostics. 6th ed. Elsevier; 2018

Performance



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Method Description

The serum sample is diluted in acetonitrile internal standard. The protein precipitate is centrifuged, and a portion of the supernatant is diluted with mobile phase for detection by a tandem mass spectrometer. (Unpublished Mayo method)

PDF Report

No

Day(s) Performed

Tuesday, Friday

Report Available

2 to 5 days

Specimen Retention Time

14 days

Performing Laboratory Location

Rochester

Fees & Codes

Fees

- Authorized users can sign in to <u>Test Prices</u> for detailed fee information.
- Clients without access to Test Prices can contact <u>Customer Service</u> 24 hours a day, seven days a week.
- Prospective clients should contact their account representative. For assistance, contact <u>Customer Service</u>.

Test Classification

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

CPT Code Information

80299

LOINC® Information

Decult ID	Took Dooult Name	Decult LOING® Value
PERAM	Perampanel, S	88895-8
Test ID	Test Order Name	Order LOINC® Value

Result ID	Test Result Name	Result LOINC® Value
609438	Perampanel, S	88895-8