

Test Definition: PVLE

Paraneoplastic Vision Loss Evaluation, Serum

Reporting Title: Paraneoplastic Vision Loss Eval, S

Performing Location: Rochester

Ordering Guidance:

Multiple neurological phenotype-specific autoimmune/paraneoplastic evaluations are available. For more information as well as phenotype-specific testing options, refer to <u>Autoimmune Neurology Test Ordering Guide</u>.

For a list of antibodies performed with each evaluation, see Autoimmune Neurology Antibody Matrix.

Necessary Information:

Provide the following information:

- -Relevant clinical information
- -Ordering provider name, phone number, mailing address, and e-mail address

Specimen Requirements:

Patient Preparation:

- 1. For optimal antibody detection, specimen collection is recommended prior to initiation of immunosuppressant medication.
- 2. This test should not be requested for patients who have recently received radioisotopes, therapeutically or diagnostically, because of potential assay interference. The specific waiting period before specimen collection will depend on the isotope administered, the dose given, and the clearance rate in the individual patient. Specimens will be screened for radioactivity prior to analysis. Radioactive specimens received in the laboratory will be held 1 week and assayed if sufficiently decayed or canceled if radioactivity remains.

Supplies: Sarstedt Aliquot Tube, 5 mL (T914)

Collection Container/Tube:

Preferred: Red top **Acceptable:** Serum gel

Submission Container/Tube: Plastic vial

Specimen Volume: 4 mL

Collection Instructions: Centrifuge and aliquot serum into a plastic vial.

Forms:

If not ordering electronically, complete, print, and send a <u>Neurology Specialty Testing Client Test Request</u> (T732) with the specimen.

| Specimen Type | Temperature | Time | Special Container |
|---------------|--------------------------|----------|-------------------|
| Serum | Refrigerated (preferred) | 28 days | |
| | Frozen | 28 days | |
| | Ambient | 72 hours | |

Result Codes:

| Result ID | Reporting Name | Туре | Unit | LOINC® | |
|-----------|-------------------------|--------------|------|---------|--|
| 83077 | CRMP-5-IgG, S | Alphanumeric | | 72504-4 | |
| 610009 | Recoverin Immunoblot, S | Alphanumeric | | 83003-4 | |



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| 607411 | Paraneoplas Vision Loss Interp, S | Alphanumeric | In Process |
|--------|-----------------------------------|--------------|------------|
| 618908 | IFA Notes | Alphanumeric | 48767-8 |

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

Supplemental Report:

No

Components:

| Test Id | Reporting Name | CPT Units | CPT Code | Always Performed | Available Separately |
|---------|-----------------------------------|-----------|----------|------------------|----------------------|
| PVLEI | Paraneoplas Vision Loss Interp, S | | | Yes | No |
| CRMS | CRMP-5-IgG, S | 1 | 86255 | Yes | No |
| RCVBS | Recoverin Immunoblot, S | 1 | 84182 | Yes | Yes |

CPT Code Information:

86255 x1

84182 x1

84182 CRMWS (if appropriate)

86256 CRMTS (if appropriate)

Reflex Tests:

| Test Id | Reporting Name | CPT Units | CPT Code | Always Performed | Available Separately |
|---------|----------------------------|-----------|----------|------------------|----------------------|
| CRMWS | CRMP-5-IgG Western Blot, S | 1 | 84182 | No | Yes |
| CRMTS | CRMP-5-IgG Titer, S | 1 | 86256 | No | No |

Result Codes for Reflex Tests:

| Test ID | Result ID | Reporting Name | Туре | Unit | LOINC® |
|---------|-----------|----------------------------|--------------|-------|---------|
| CRMWS | 83107 | CRMP-5-IgG Western Blot, S | Alphanumeric | | 47401-5 |
| CRMTS | 43436 | CRMP-5-IgG Titer, S | Alphanumeric | titer | 94815-8 |

Reference Values:

COLLAPSIN RESPONSE-MEDIATOR PROTEIN-5-IgGNegative

RECOVERIN IMMUNOBLOT

Negative

COLLAPSIN RESPONSE-MEDIATOR PROTEIN-5 TITER

<1:240

COLLAPSIN RESPONSE-MEDIATOR PROTEIN-5 WESTERN BLOT



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Negative

Titers lower than 1:240 are detectable by recombinant CRMP-5 Western blot analysis. CRMP-5 Western blot analysis will be done on request on stored serum (held 4 weeks). This supplemental testing is recommended in cases of chorea, vision loss, cranial neuropathy, and myelopathy. Call 1-800-533-1710 to request CRMP-5 Western blot.

Neuron-restricted patterns of IgG staining that do not fulfill criteria for CRMP-5-IgG may be reported as "unclassified antineuronal IgG." Complex patterns that include non-neuronal elements may be reported as "uninterpretable."