

## Overview

### Useful For

Definitive identification of amyloid proteins

### Reflex Tests

Test Id	Reporting Name	Available Separately	Always Performed
MLCPC	Microdissection, Laser Capture	No, (Bill Only)	No
MSPTC	Mass Spectrometry	No, (Bill Only)	No

### Testing Algorithm

[Testing begins with a Congo red stain. If the Congo red stain is positive, testing proceeds with microdissection at an additional charge to obtain tissue of interest and subsequently perform mass spectrometry at an additional charge to identify protein involved.](#)

In some instances, per pathologist discretion, a different initial Congo red stain may be performed using SS2PC / Special Stain, Group II, Other (Bill Only).

-If the stain is negative for amyloid, then this test will not be performed, and only the SS2PC will be charged.

-If the stain is positive for amyloid, this test will be performed, and the SS2PC billing charge will be credited.

A pathology consultation is typically not required. If the amyloid subtyping results do not fit the clinical findings, PATHC / Pathology Consultation may be added if appropriate, upon client approval.

For more information see [Amyloidosis: Laboratory Approach to Diagnosis](#).

### Special Instructions

- [Amyloidosis: Laboratory Approach to Diagnosis](#)
- [Pathology Consultation Ordering Algorithm](#)

### Method Name

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

### NY State Available

Yes

## Specimen

### Specimen Type

AMYLOID

**Ordering Guidance**

[This test \*\*should only\*\* be ordered on patients for whom a primary diagnosis has already been established. If a patient does not have a primary diagnosis, order PATHC / Pathology Consultation or refer to the Pathology Consultation Ordering Algorithm.](#)

If a pathology consultation is desired in addition to this test, order PATHC / Pathology Consultation alone and send the required paperwork with specimen. Indicate that amyloid protein identification is desired. If needed, this test will be added by the reviewing pathologist and will be reported with the consultation. For more information see PATHC / Pathology Consultation.

**Shipping Instructions**

Attach the green pathology address label included in the kit to the outside of the transport container.

**Necessary Information**

1. Preliminary pathology report and history are required.
2. Include performed Congo red slide
3. A brief explanatory note or consultative letter is also recommended.

**Specimen Required**

**Specimen Type:** Formalin-fixed or B5-fixed paraffin-embedded tissue block

**Collection Instructions:**

1. Do not send fixed tissue slides for testing. Testing can only be done on paraffin-embedded tissue blocks.
2. [If Congo red stain has already been performed, send Congo red](#) stained slide along with the tissue block.

**Forms**

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

- [Cardiovascular Test Request](#) (T724)
- [Hematopathology/Cytogenetics Test Request](#) (T726)
- [Renal Diagnostics Test Request](#) (T830)

**Reject Due To**

Fixed tissue slides Wet/frozen tissue Cytological smears Nonformalin fixed tissue Nonparaffin embedded tissue	Reject
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**Specimen Stability Information**

Specimen Type	Temperature	Time	Special Container
AMYLOID	Ambient (preferred)		
	Refrigerated		

**Clinical & Interpretive****Clinical Information**

Amyloidosis is a group of hereditary and acquired diseases unified by extracellular tissue deposition of misfolded proteins resulting in end organ damage. Amyloidosis can be a systemic or localized disease. Although many cases of amyloidosis are hereditary, most are acquired as the result of an underlying monoclonal B-cell/plasma cell malignancy, as a phenomenon of aging, or as the result of long-standing chronic inflammation. Specific amyloid-related diseases are therefore associated with specific amyloid proteins. These include kappa or lambda immunoglobulin light chains (AL amyloid), transthyretin (ATTR amyloid), serum amyloid A (SAA amyloid), and other uncommon subtypes. Because treatment of patients with amyloidosis differs radically for the different amyloid subtypes, it is critically important to accurately identify the proteins that constitute the amyloid deposits.

The basic diagnosis of amyloidosis is typically achieved by Congo red staining of paraffin-embedded tissue biopsy specimens obtained from diverse anatomic sites and demonstrating Congo red-positive, apple-green birefringent, amyloid deposits in the tissues. The next step is to definitively subtype the amyloid deposits. This test fulfills that need. It relies on laser microdissection of Congo red-positive amyloid deposits followed by analysis by liquid chromatography tandem mass spectrometry to accurately determine the identity of the proteins that constitute the amyloid.

**Reference Values**

An interpretive report will be provided.

**Interpretation**

An interpretation will be provided.

**Cautions**

In rare instances amyloid deposits may show a false-negative result by Congo red staining. Because this test depends on the presence of Congo red-positive amyloid, these cases may not be identified as amyloid by this testing algorithm. Correlation with clinical and pathologic features and other laboratory test results is recommended to definitively exclude a diagnosis of amyloidosis.

**Clinical Reference**

1. Theis JD, Dasari S, Vrana JA, Kurtin PJ, Dogan A. Shotgun-proteomics-based clinical testing for diagnosis and classification of amyloidosis. *J Mass Spectrom*. 2013;48(10):1067-1077
2. Said SM, Sethi S, Valeri AM, et al. Renal amyloidosis: origin and clinicopathologic correlations of 474 recent cases. *Clin J Am Soc Nephrol*. 2013;8(9):1515-1523
3. Dasari S, Theis JD, Vrana JA, et al. Amyloid typing by mass spectrometry in clinical practice: a comprehensive review of

16,175 samples. Mayo Clin Proc. 2020;95(9):1852-1864. doi:10.1016/j.mayocp.2020.06.029

4. Klein CJ, Vrana JA, Theis JD, et al. Mass spectrometric-based proteomic analysis of amyloid neuropathy type in nerve tissue. Arch Neurol. 2011;68(2):195-199

5. Vrana JA, Gamez JD, Madden BJ, Theis JD, Bergen HR III, Dogan A. Classification of amyloidosis by laser microdissection and mass spectrometry-based proteomic analysis in clinical biopsy specimens. Blood. 2009;114(24):4957-4959

## Performance

### Method Description

Affected areas are removed from paraffin-embedded tissues by laser microdissection. Protein digestion is performed, followed by liquid chromatography tandem mass spectrometry.(Unpublished Mayo method)

### PDF Report

No

### Day(s) Performed

Monday through Friday

### Report Available

7 to 15 days

### Specimen Retention Time

Submitted block: Not retained; Congo red-stained slides performed at Mayo Clinic: Indefinitely

### Performing Laboratory Location

Rochester

## Fees & 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 account representative. 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. It has not been cleared or approved by the US Food and Drug Administration.

### CPT Code Information

88313

82542 (if appropriate)

88380 (if appropriate)

**LOINC® Information**

Test ID	Test Order Name	Order LOINC® Value
AMPIP	Amyloid Protein ID, Par, LC MS/MS	101405-9

Result ID	Test Result Name	Result LOINC® Value
71185	Interpretation	50595-8
71186	Participated in the Interpretation	No LOINC Needed
71187	Report electronically signed by	19139-5
71189	Material Received	81178-6
71592	Disclaimer	62364-5
72109	Case Number	80398-1