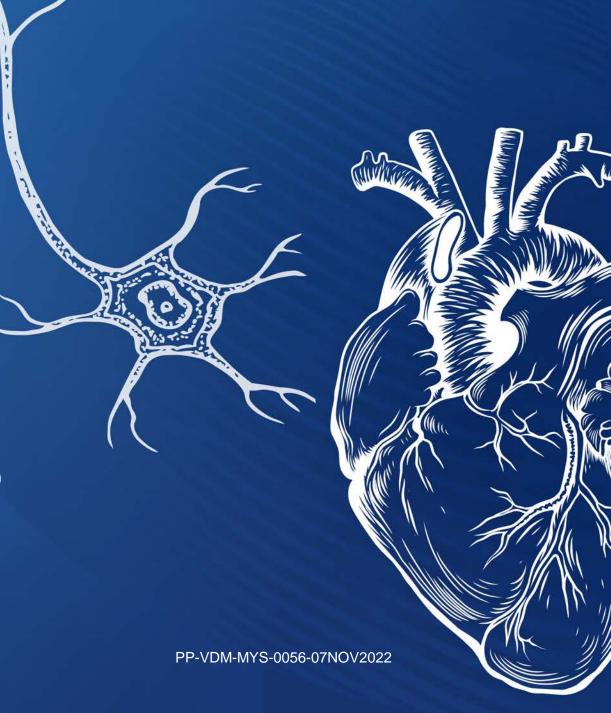


Ruling out Light-Chain (AL) amyloidosis in ATTR-CM

Transthyretin amyloid cardiomyopathy (ATTR-CM)

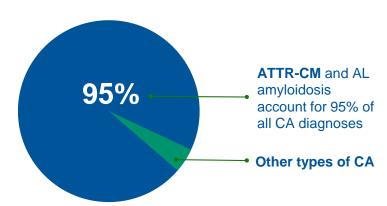




ATTR-CM and Light Chain (AL) amyloidosis have similar symptom presentation which may make diagnosis challenging

It is important to clinically differentiate between cardiac manifestations of ATTR and AL amyloidosis, as they have different clinical courses and treatment options, and AL amyloidosis is considered a hematologic urgency^{1,2}

Causes of Cardiac Amyloidosis (CA)³



ATTR-CM is under recognized, may present with symptoms similar to AL and may be misdiagnosed due to incomplete testing^{4,5}

Symptoms that may present in both ATTR-CM and **AL** amyloidosis

Cardiovascular^{3,6-8}

- Heart failure with potential intolerance to : Automatic neuropathy standard therapy
- Cardiac arrhythmia
- Aortic stenosis
- Low voltage relative to left ventricular (LV) mass
- Echocardiography showing increased LV wall thickness

Nervous System^{4,8-10}

- Orthostatic hypotension
- Peripheral sensory motor dysfunction
- Peripheral neuropathy
- Unexplained weight loss
- Sexual impotence

Ocular^{4,11,12}

- Vitreous opacity
- Glaucoma
- Periorbital purpura*

Musculoskeletal/Orthopedic⁴

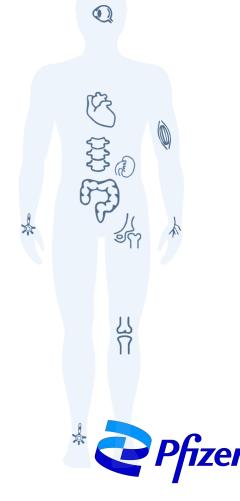
Carpal tunnel syndrome

Renal 10,13,14

- Renal impairment
- Nephrotic syndrome*
- Cardiorenal symptoms

Other 4,10,15

- Gastrointestinal complaints
- Macroglossia (large tongue)*
- Nail dystrophy*



^{*}More commonly seen in AL

Ruling out light chain amyloidosis (AL) is a key step in achieving a definitive ATTR-CM diagnosis¹⁶

SERUM AND URINE TESTS FOR RULING OUT AL AMYLOIDOSIS*

	Serum protein electrophoresis (SPEP) with immunofixation†	Urine protein electrophoresis (UPEP) with immunofixation [†]	Serum free light chain assay
What does it detect?	Clonal immunoglobulin and/or clonal light chain	Clonal immunoglobulin and/or clonal light chain	Free kappa chain Free lambda chain kappa: lambda ratio
Most sensitive test for:	Confirming clonal immunoglobulin production	Confirming clonal light chain production	Detecting low-level clonal light chain production; clonality assumed if ratio is far from 1:1
Normal range	No M-spike present	No M-spike present No proteinura ⁴	Kappa:lambda ratio=0.26- 1.65 [‡]

- AL amyloidosis is a hematological urgency and requires immediate treatment
- Survival of untreated patients with AL amyloidosis with cardiac involvement may be <6 months
- Cardiac localization of radiotracer by bone scintigraphy can occur in approximately 30% of patients with AL amyloidosis
- Specific disease-modifying therapies for AL amyloidosis are available

‡In patients with kidney disease, mild elevations in the kappa:lambda ratio are frequently encountered. In the setting of a normal SPEP/UPEP, a kappa:lambda ratio up to 2.5 can typically be considered normal.



^{*}If any of these tests are abnormal, bone scintigraphy should not be used to make the diagnosis of transthyretin amyloidosis, and a biopsy is recommended. Adapted from Witteles et al., 2019 Table 1, which uses SPIE (serum protein electrophoresis with immunofixation) and UPIE (urine protein electrophoresis with immunofixation) acronyms.

[†]SPEP and UPEP are more sensitive than protein electrophoresis without immunofixation and should be ordered as preferred test.

AL rule out is an important step in the diagnostic pathway. All three tests should be sent to evaluate for AL in order to maximize sensitivity and specificity¹⁷

Order Studies to Test for Abnormal Monoclonal Protein*

Serum Kappa/Lambda Free Light Chain Ratio

Abnormal result:

Ratio < 0.26 or > 1.65

Serum Protein Immunofixation
Abnormal result:

Monoclonal protein is detected

Urine Protein Immunofixation
Abnormal result:

Monoclonal protein is detected

Key Considerations:

- It is recommended that all three tests be sent to evaluate for AL amyloidosis. However, monoclonal protein presence does not rule out the possibility of ATTR-CM
- Amyloid subtyping is the only way to effectively differentiate between ATTR-CM and AL Amyloidosis

Adapted and reprinted with permission from Maurer MS, et al. Expert consensus recommendations for the suspicion and diagnosis of transthyretin cardiac amyloidosis. Circ Heart Fail. 2019;12:e006075. doi:10.1161/CIRCHEARTFAILURE.119.006075 © 2019 American Heart Association, Inc. All rights reserved.

*If any of these tests are abnormal, bone scintigraphy should not be used to make the diagnosis of transthyretin amyloidosis, and a biopsy is recommended.

Immunofixation of the serum and urine is a much more sensitive test and usually demonstrates a monoclonal band. Thus, immunofixation and serum-free light chains should always be measured when amyloidosis is suspected.¹⁷ Urine protein electrophoresis with immunofixation can be performed on spot or 24-h urine collection.



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