

Suggested Reading

Session 1

Brown EE, Lee YZJ, Halushka MK, et al. Genetic testing improves identification of transthyretin amyloid (ATTR) subtype in cardiac amyloidosis. <https://pubmed.ncbi.nlm.nih.gov/28494620/>

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Part 1: <https://www.ahajournals.org/doi/10.1161/HCI.0000000000000029>

Part 2: <https://www.ahajournals.org/doi/10.1161/HCI.0000000000000030>

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Witteles RM, Liedtke M. Avoiding catastrophe: understanding free light chain testing in the evaluation of ATTR amyloidosis. <https://pubmed.ncbi.nlm.nih.gov/33736459/>

Session 2

The American College of Medical Genetics and Genomics recommendations for reporting of incidental findings in clinical exome and genome sequencing. <https://www.ncbi.nlm.nih.gov/clinvar/docs/acmg/>

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Conceição I, Damy T, Romero M, et al. Early diagnosis of ATTR amyloidosis through targeted follow-up of identified carriers of TTR gene mutations. Amyloid. March 2019. <https://pubmed.ncbi.nlm.nih.gov/30793974/>

Dang D, Fournier P, Cariou E, et al. Gateway and journey of patients with cardiac amyloidosis. ESC Heart Fail. October 2020. <https://pubmed.ncbi.nlm.nih.gov/23474361/>

Davis MK, Fine NM, Small GR, et al. Establishing a cardiac amyloidosis clinic: a practical primer for cardiologists. Can J Cardiol. April 2021. <https://pubmed.ncbi.nlm.nih.gov/33485855/>

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