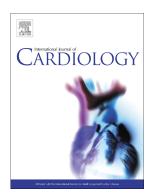
Left atrial strains in cardiac amyloidosis -does its subtype matter?



Attila Nemes

PII: S0167-5273(24)00698-3

DOI: https://doi.org/10.1016/j.ijcard.2024.132078

Reference: IJCA 132078

To appear in: International Journal of Cardiology

Received date: 14 March 2024

Revised date: 8 April 2024

Accepted date: 17 April 2024

Please cite this article as: A. Nemes, Left atrial strains in cardiac amyloidosis -does its subtype matter?, *International Journal of Cardiology* (2023), https://doi.org/10.1016/j.ijcard.2024.132078

This is a PDF file of an article that has undergone enhancements after acceptance, such as the addition of a cover page and metadata, and formatting for readability, but it is not yet the definitive version of record. This version will undergo additional copyediting, typesetting and review before it is published in its final form, but we are providing this version to give early visibility of the article. Please note that, during the production process, errors may be discovered which could affect the content, and all legal disclaimers that apply to the journal pertain.

© 2024 Published by Elsevier B.V.

Left atrial strains in cardiac amyloidosis -does its subtype matter?

Attila Nemes, MD, PhD, DSc

Department of Medicine, Albert Szent-Györgyi Medical School, University of Szeged, Szeged, Hungary

Address for correspondence: Attila Nemes, MD, PhD, DSc. Department of Medicine, Albert Szent-Györgyi Medical School, University of Szeged, H-6725 Szeged, Semmelweis street 8, Hungary, P.O. Box 427. Fax: +36-62-544568, Tel.: +36-62-545220. Email: nemes.attila@med.u-szeged.hu

I am reading the paper of Meucci et al. which aimed to perform a comparative analysis of left atrial (LA) remodelling between Fabry disease (FD) and transthyretin cardiac amyloidosis (ATTR-CA).[1] It could be concluded that ATTR-CA is characterized by a more advanced structural and functional LA remodeling as compared to FD despite the similar degree of left ventricular hypertrophy. I found the paper interesting, but have some important comments. In case of two-dimensional (2D) speckle-tracking echocardiography (STE), LA strains are measured on loops generated in a selected two-dimensional (2D) plane, while in case of threedimensional (3D) STE, digitally recorded 3D echocardiographic datasets help to create a virtual 3D cast, with which volumetric and functional (strain) analysis of a certain LA can be measured at the same time. While 2D-STE is an established procedure and its role is well supported by several studies in the field, 3D-STE is still subject of research, it needs of adequate training and dedicated equipment, suffers from many technical limitations and is not widespread enough.[2,3] In the study of Meucci et al., 2D-STE was used, and mean reservoir LA strain of ATTR-CA patients proved to be 6.9%.[1] In a recent 3D-STE study aiming to compare LA parameters between light-chain (AL) CA and hypertrophic cardiomyopathy, radial, circumferential (CS) and longitudinal (LS) LA reservoir strains for AL-CA proved to be -9.7%, 11.9% and 11.2%, respectively.[2] Although the comparison is difficult due to methodological differences, results could highlight differences in LA strains between different types of CA like AL-CA versus ATTR-CA suggesting more pronounced abnormalities in case of the latter one in correspondence with previous findings by Versteynlein et al. [4] They found that LA reservoir strain was markedly lower in ATTR-CA as compared to AL-CA (7.4 \pm 6.2% vs. 13.6 \pm 14.7, p = 0.017).[4] In accordance with this findings, lower reservoir LA-LS and LA-CS could be detected in ATTR-CA patients as compared to that of AL-CA patients in the study by Aimo et al.[5] These results could suggest further comparative studies between CA subtypes using more recent imaging techniques including 3D-STE, which allows

more detailed volumetric and strain analysis using specific unidimensional and complex LA strains. Moreover, further studies are warranted in patients with suspected isolated atrial amyloidosis.[6] To achieve the sample size, when creating different subgroups of patients, a multicenter would be needed. Moreover, even at the same time as above, same sort of evaluation of the right atrium in the description of atrial cardiomyopathy would also be an interesting topic as regards to atrial fibrillation, differentiation between CA and FD, etc.[7,8]

References

- Meucci MC, Lillo R, Mango F, et al. Left atrial structural and functional remodelling in Fabry disease and cardiac amyloidosis: A comparative analysis. Int J Cardiol (in press)
- Földeák D, Kormányos Á, Domsik P, et al. Left atrial dysfunction in lightchain cardiac amyloidosis and hypertrophic cardiomyopathy - A comparative threedimensional speckle-tracking echocardiographic analysis from the MAGYAR-Path Study. Rev Port Cardiol. 2017;36:905-913.
- 3. Nochioka K, Quarta CC, Claggett B, et al. Left atrial structure and function in cardiac amyloidosis. Eur Heart J Cardiovasc Imaging 2017;18:1128-1137.
- 4. Versteylen MO, Brons M, Teske AJ, Oerlemans MIFJ. Restrictive atrial dysfunction in cardiac amyloidosis: differences between immunoglobulin light chain and transthyretin cardiac amyloidosis patients. Biomedicines. 2022;10:1768.
- 5. Aimo A, Fabiani I, Giannoni A, et al. Multi-chamber speckle tracking imaging and diagnostic value of left atrial strain in cardiac amyloidosis. Eur Heart J Cardiovasc Imaging 2022;24:130-141.
- 6. Vergaro G, Aimo A, Rapezzi C, et al. Atrial amyloidosis: mechanisms and clinical manifestations. Eur J Heart Fail 2022;24:2019-2028.
- Tomaselli M, Badano LP, Cannone V, et al. Incremental Value of Right Atrial Strain Analysis to Predict Atrial Fibrillation Recurrence After Electrical Cardioversion J Am Soc Echocardiogr 2023;36:945-955.
- 8. Mattig I, Steudel T, Klingel K, et al. Right heart and left atrial strain to differentiate cardiac amyloidosis and Fabry disease Sci Rep 2024;14:2445.