Correction to

Dorbala, Sharmila; Ando, Yukio; Bokhari, Sabahat; Dispenzieri, Angela; Falk, Rodney H; Ferrari, Victor A; Fontana, Marianna; Gheysens, Olivier; Gillmore, Julian D; Glaudemans, Andor W J M

Published in:
Journal of Nuclear Cardiology

DOI:
10.1007/s12350-021-02711-w

Publication date:
2021

Correction to: ASNC/AHA/ASE/EANM/HFSA/ISA/SCMR/SNMMI expert consensus recommendations for multimodality imaging in cardiac amyloidosis: Part 1 of 2—evidence base and standardized methods of imaging

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The original article can be found online at https://doi.org/10.1007/s12350-019-01760-6.

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J Nucl Cardiol
1071-3581/$34.00

Published online: 14 July 2021

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doi:10.1007/s12350-021-02711-w

CORRECTION TO:

https://doi.org/10.1007/s12350-019-01760-6

- In the **Introduction** SCMR was listed incorrectly. SCMR is the Society for Cardiovascular Magnetic Resonance.
- **Figure 1** erroneously printed without Yen sign (¥) in 'Final Diagnosis.' Please see revised Figure 1.
- **Acknowledgments** erroneously printed without reviewers Richard Cheng, MD and Roy John, MD.

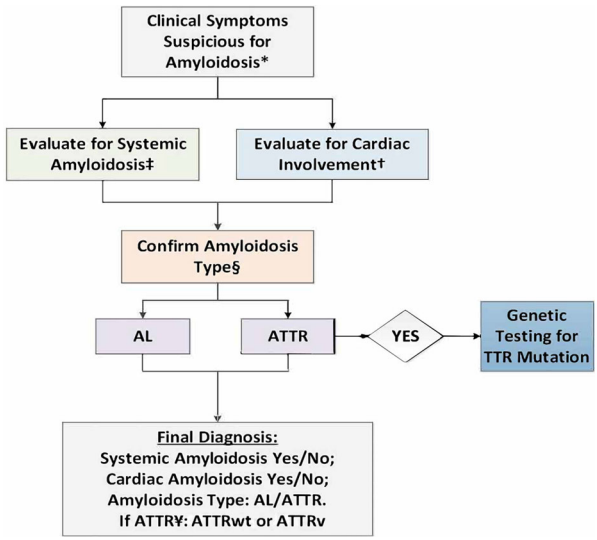

Figure 1. Systematic evaluation of cardiac amyloidosis. A comprehensive evaluation of cardiac amyloidosis includes consideration of clinical symptoms, evaluation of cardiac involvement (biomarkers and cardiac imaging), evaluation of systemic amyloidosis (serum, urine testing, and biopsy), followed by typing of amyloid deposits into AL or ATTR, and documentation of mutations in patients with ATTR amyloidosis. *__Clinical symptoms__: heart failure, peripheral/autonomic neuropathy, macroglossia, carpal tunnel syndrome, periorbital bruising, stroke, atrial fibrillation, postural hypotension, fatigue, weight loss, pedal edema, renal dysfunction, diarrhea, constipation. †__Evaluation for cardiac amyloidosis__: ECG, ECHO, CMR, EMB, ^{99m}Tc-PYP/DPD/HMDP/^{123}I-*m*IBG/PET, NT-proBNP, troponin T. ‡__Evaluation for systemic amyloidosis__: AL: detect plasma cell clone: serum and urine immunofixation, serum FLC assay and immunoglobulin analysis; AL: detect systemic organ involvement: 24-hour urine protein, alkaline phosphatase, eGFR, cardiac biomarkers (NT-proBNP, troponins); Tissue biopsy: EMB/Fatpad/Bone marrow/Other with Congo red staining. §__Confirm Amyloidosis Type__: ATTR: IHC and MS of Biopsy or ^{99m}Tc-PYP/DPD/HMDP Grade 2 or3 if a clonal process is excluded; AL: MS or IHC of Biopsy. ¥__Confirm TTR mutation in patients with ATTR amyloidosis__: genetic testing for TTR mutations. *AL*, amyloid light chain; *ATTR*, amyloid transthyretin; *CMR*, cardiac magnetic resonance imaging; *DPD*, -3,3-diphosphono-1,2-propanodicarboxylic acid; *ECG*, electrocardiogram; *EMB*, endomyocardial biopsy; *ECHO*, echocardiogram; *eGFR*, estimated glomerular filtration rate; *HMDP*, hydroxymethylenediphosphonate; *IHC*, immunohistochemistry; *mIBG*, meta-iodobenzylguanidine; *MS*, mass spectroscopy; *v*, hereditary; *PYP*, pyrophosphate; *Tc*, technetium; *wt*, wild-type.