

List of records with final decision on inclusion or exclusion in the systematic review and meta-analysis

1: Youssef ME, Yahya G, Popoviciu MS, Cavalu S, Abd-Eldayem MA, Saber S. Unlocking the Full Potential of SGLT2 Inhibitors: Expanding Applications beyond Glycemic Control. *Int J Mol Sci.* 2023 Mar 23;24(7):6039. doi: 10.3390/ijms24076039.

Excluded (reason: outside the field of interest)

2: Navarro-Saez MDC, Feijoo-Massó C, Bravo Ferrer ZDC, Oliva Morera JC, Balado González AM, Palau-Domínguez A, Guillamon Toran L, Comet Monte R, Fernández-Codina A. Trends in diagnosis of cardiac transthyretin amyloidosis: 3-year analysis of scintigraphic studies: Prevalence of myocardial uptake and its predictor factors. *Int J Cardiovasc Imaging.* 2023 Apr 1. doi: 10.1007/s10554-023-02840-y.

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3: Cersosimo A, Bonelli A, Lombardi CM, Moreo A, Pagnesi M, Tomasoni D, Arabia G, Vizzardi E, Adamo M, Farina D, Metra M, Inciardi RM. Multimodality imaging in the diagnostic management of concomitant aortic stenosis and transthyretin-related wild-type cardiac amyloidosis. *Front Cardiovasc Med.* 2023 Mar 14;10:1108696. doi: 10.3389/fcvm.2023.1108696.

Excluded (reason: outside the field of interest)

4: Ruiz-Hueso R, Salamanca-Bautista P, Quesada-Simón MA, Yun S, Conde-Martel A, Morales-Rull JL, Suárez-Gil R, García-García JÁ, Llàcer P, Fonseca-Aizpuru EM, Amores-Arriaga B, Martínez-González Á, Armengou-Arxe A, Peña-Somovilla JL, López-Reboiro ML, Aramburu-Bodas Ó; PREVAMIC Investigators Group. Estimating the Prevalence of Cardiac Amyloidosis in Old Patients with Heart Failure-Barriers and Opportunities for Improvement: The PREVAMIC Study. *J Clin Med.* 2023 Mar 15;12(6):2273. doi: 10.3390/jcm12062273.

Excluded (reason: outside the field of interest)

5: Rauf MU, Hawkins PN, Cappelli F, Perfetto F, Zampieri M, Argiro A, Petrie A, Law S, Porcari A, Razvi Y, Bomsztyk J, Ravichandran S, Ioannou A, Patel R, Starr N, Hutt DF, Mahmood S, Wisniewski B, Martinez-Naharro A, Venneri L, Whelan C, Roczenio D, Gilbertson J, Lachmann HJ, Wechalekar AD, Rapezzi C, Serenelli M, Massa P, Caponetti AG, Ponziani A, Accietto A, Giovannetti A, Satri G, Sguazzotti M, Gagliardi C, Biagini E, Longhi S, Fontana M, Gillmore JD. Tc-99m labelled bone scintigraphy in suspected cardiac amyloidosis. *Eur Heart J.* 2023 Mar 22;ehad139. doi: 10.1093/eurheartj/ehad139.

Excluded (reason: outside the field of interest)

6: Tsushima T, Terao T, Narita K, Fukumoto A, Ikeda D, Kamura Y, Kuzume A, Tabata R, Miura D, Takeuchi M, Matsue K. Clinical Characteristics and Outcomes of Cyclin D1-Positive AL Amyloidosis. *Am J Clin Pathol.* 2023 Mar 20;aqad013. doi: 10.1093/ajcp/aqad013.

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7: Raval M, Siddiq S, Sharma K, Sanghvi L, Jain A, Patel S, Trivedi J, Uttam Chandani K, Patel D, Desai R. A review of recent advances in the diagnosis of cardiac amyloidosis, treatment of its cardiac complications, and disease-modifying therapies. *F1000Res.* 2023 Feb 20;12:192. doi: 10.12688/f1000research.130285.1.

Excluded (reason: outside the field of interest)

8: Nuzzi V, Porcari A, Gigli M, Zaja F, Dore F, Bussani R, Sinagra G, Merlo M. A case report of isolated cardiac light chain amyloidosis without clinically overt heart failure: an under-recognized presentation. *Eur Heart J Case Rep.* 2023 Feb 13;7(3):ytad072. doi: 10.1093/ehjcr/ytad072. PMID: 36909839; PMCID: PMC9998033.

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9: Naiki H, Yamaguchi A, Sekijima Y, Ueda M, Ohashi K, Hatakeyama K, Ikeda Y, Hoshii Y, Shintani-Domoto Y, Miyagawa-Hayashino A, Tsujikawa H, Endo J, Arai T, Ando Y. Steep increase in the number of transthyretin-positive cardiac biopsy cases in Japan: evidence obtained by the nation-wide pathology consultation for the typing diagnosis of amyloidosis. *Amyloid.* 2023 Feb 16:1-6. doi: 10.1080/13506129.2023.2180334.

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10: Le Bras P, Plard L, Le Gouill C, Piriou N, Touchefeu Y. Transthyretin Cardiac Amyloidosis Mimicking Immune Checkpoint-Induced Myocarditis in a Patient Treated with Atezolizumab and Bevacizumab for Advanced Hepatocellular Carcinoma: A Case Report. *Case Rep Oncol.* 2022 Nov 8;15(3):967-973. doi: 10.1159/000526534.

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11: Testa S, Kumar J, Goodell AJ, Zehnder JL, Alexander KM, Sidana S, Arai S, Witteles RM, Liedtke M. Prevalence, mutational spectrum and clinical implications of clonal hematopoiesis of indeterminate potential in plasma cell dyscrasias. *Semin Oncol.* 2022 Dec;49(6):465-475. doi: 10.1053/j.seminoncol.2022.11.001.

Excluded (reason: outside the field of interest)

12: Schneider SM, Kochar K, Ruge M, Marek-Iannucci S, Datta T, Hajduczuk A, Ullah W, Rajapreyar I, Brailovsky Y. Cardiogenic Shock Due to Atrial Arrhythmia as the Initial Presentation of Transthyretin Cardiac Amyloidosis. *JACC Case Rep.* 2022 Nov 16;4(22):1490-1495. doi: 10.1016/j.jaccas.2022.07.041.

Excluded (reason: outside the field of interest)

13: Yadav P, Lee YH, Panday H, Kant S, Bajwa N, Parashar R, Jha SK, Jha NK, Nand P, Lee SS, Jha AK. Implications of Microorganisms in Alzheimer's Disease. *Curr Issues Mol Biol.* 2022 Sep 30;44(10):4584-4615. doi: 10.3390/cimb44100314.

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14: Porcari A, Hutt DF, Grigore SF, Quigley AM, Rowczenio D, Gilbertson J, Patel R, Razvi Y, Ioannou A, Rauf MU, Martinez-Naharro A, Venneri L, Hawkins PN, Fontana M, Gillmore JD. Comparison of different technetium-99m-labelled bone tracers for imaging cardiac amyloidosis. *Eur J Prev Cardiol.* 2023 Feb 14;30(3):e4-e6. doi: 10.1093/eurjpc/zwac237.

Excluded (reason: outside the field of interest)

15: Jain A, Zahra F. Transthyretin Amyloid Cardiomyopathy (ATTR-CM). 2022 Sep 26. In: StatPearls [Internet]. Treasure Island (FL): StatPearls Publishing; 2023 Jan–.

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16: Yang H, Li R, Ma F, Wei Y, Liu Y, Sun Y, He X, Zeng H, Yan J, Wang DW, Wang H. An echo score raises the suspicion of cardiac amyloidosis in Chinese with heart failure with preserved ejection fraction. *ESC Heart Fail.* 2022 Dec;9(6):4280-4290. doi: 10.1002/ehf2.14164.

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17: Wang J, Yang S, Liao P, Zeng L, Ling W, Wan L, Weng J, Zhong L. Incidence and effect of secondary cardiac amyloidosis on outcomes of patients with t(11;14) multiple myeloma. *Front Cardiovasc Med.* 2022 Sep 2;9:994384. doi: 10.3389/fcvm.2022.994384.

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18: Ge Y, Pandya A, Cuddy SAM, Singh A, Singh A, Dorbala S. Modeling the Cost and Health Impacts of Diagnostic Strategies in Patients with Suspected Transthyretin Cardiac Amyloidosis. *J Am Heart Assoc.* 2022 Sep 20;11(18):e026308. doi: 10.1161/JAHA.122.026308. Epub 2022 Sep 14.

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19: Tzagournissakis M, Foukarakis E, Samonakis D, Tsilimbaris M, Michaelidou K, Mathioudakis L, Marinis A, Giannakoudakis E, Spanaki C, Skoula I, Erimaki S, Amoiridis G, Koutsis G, Koukouraki S, Stylianou K, Plaitakis A, Mitsias PD, Zaganas I. High Hereditary Transthyretin-Related Amyloidosis Prevalence in Crete: Genetic Heterogeneity and Distinct Phenotypes. *Neurol Genet.* 2022 Sep 9;8(5):e200013. doi: 10.1212/NXG.0000000000200013.

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20: Davies DR, Redfield MM, Scott CG, Minamisawa M, Grogan M, Dispenzieri A, Chareonthaitawee P, Shah AM, Shah SJ, Wehbe RM, Solomon SD, Reddy YNV, Borlaug BA, AbouEzzeddine OF. A Simple Score to Identify Increased Risk of Transthyretin Amyloid Cardiomyopathy in Heart Failure With Preserved Ejection Fraction. *JAMA Cardiol.* 2022 Oct 1;7(10):1036-1044. doi: 10.1001/jamacardio.2022.1781.

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21: Westin O, Fosbøl EL, Maurer MS, Leicht BP, Hasbak P, Mylin AK, Rørvig S, Lindkær TH, Johannesen HH, Gustafsson F. Screening for Cardiac Amyloidosis 5 to 15 Years After Surgery for Bilateral Carpal Tunnel Syndrome. *J Am Coll Cardiol.* 2022 Sep 6;80(10):967-977. doi: 10.1016/j.jacc.2022.06.026.

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22: Donà C, Nitsche C, Koschutnik M, Heitzinger G, Mascherbauer K, Kammerlander AA, Dannenberg V, Halavina K, Rettl R, Duca F, Traub-Weidinger T, Puchinger J, Gunacker PC, Lamm G, Vock P, Lileg B, Philipp V, Staudenherz A, Calabretta R, Hacker M, Agis H, Bartko P, Hengstenberg C, Fontana M, Goliash G, Mascherbauer J. Unveiling Cardiac Amyloidosis, its Characteristics, and Outcomes Among Patients With MR Undergoing Transcatheter Edge-to-Edge MV Repair. *JACC Cardiovasc Interv.* 2022 Sep 12;15(17):1748-1758. doi: 10.1016/j.jcin.2022.06.009.

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23: Porcari A, Fontana M, Gillmore JD. Transthyretin cardiac amyloidosis. *Cardiovasc Res.* 2023 Feb 3;118(18):3517-3535. doi: 10.1093/cvr/cvac119.

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24: Ramsell S, Arias Bermudez C, Takem Baiyee CAM, Rodgers B, Parikh S, Almaani S, Sharma N, LoRusso S, Freimer M, Redder E, Bumma N, Vallkati A, Efebera Y, Kahwash R, Campbell CM. Beta-Adrenergic Antagonist Tolerance in Amyloid Cardiomyopathy. *Front Cardiovasc Med.* 2022 Jul 11;9:907597. doi: 10.3389/fcvm.2022.907597.

Excluded (reason: outside the field of interest)

25: Rossi M, Varrà GG, Porcari A, Saro R, Pagura L, Lalario A, Dore F, Bussani R, Sinagra G, Merlo M. Re-Definition of the Epidemiology of Cardiac Amyloidosis. *Biomedicines.* 2022 Jun 30;10(7):1566. doi: 10.3390/biomedicines10071566. PMID: 35884871; PMCID: PMC9313045.

Excluded (reason: review)

26: Gao YJ, Shen KN, Chang L, Feng J, Mao YY, Zhang L, Cao XX, Zhou DB, Li J. Prevalence, clinical characteristics and treatment outcome of factor X deficiency in a consecutive cohort of primary light-chain amyloidosis. *Leuk Res.* 2022 Sep;120:106917. doi: 10.1016/j.leukres.2022.106917.

Excluded (reason: outside the field of interest)

27: Nitsche C, Mascherbauer K, Calabretta R, Koschutnik M, Dona C, Dannenberg V, Hofer F, Halavina K, Kammerlander AA, Traub-Weidinger T, Goliash G, Hengstenberg C, Hacker M, Mascherbauer J. Prevalence and Outcomes of Cardiac Amyloidosis in All-Comer Referrals for Bone Scintigraphy. *J Nucl Med.* 2022 Dec;63(12):1906-1911. doi: 10.2967/jnumed.122.264041.

Included in the systematic review and meta-analysis

28: Lizarazo Ortega DA, Valderrama BP, González-Robledo G, Trujillo PB. A Case of Wild-type Cardiac Transthyretin Amyloidosis Diagnosed by Non-invasive Methods. *Curr Med Imaging.* 2023;19(4):402-406. doi: 10.2174/1573405618666220610091446.

Excluded (reason: outside the field of interest)

29: Porcari A, Rossi M, Dore F, Imazio M, Fontana M, Merlo M, Sinagra G. I dieci punti che il cardiologo deve conoscere su scintigrafia miocardica con traccianti ossei, amiloidosi e cuore [Ten questions for the cardiologist about

cardiac scintigraphy with bone tracers, amyloidosis and the heart]. G Ital Cardiol (Rome). 2022 Jun;23(6):424-432. Italian. doi: 10.1714/3810.37938.

Excluded (reason: outside the field of interest)

30: Wang M, Pan W, Xu Y, Zhang J, Wan J, Jiang H. Microglia-Mediated Neuroinflammation: A Potential Target for the Treatment of Cardiovascular Diseases. J Inflamm Res. 2022 May 25;15:3083-3094. doi: 10.2147/JIR.S350109.

Excluded (reason: outside the field of interest)

31: Hussain M, Krywanczyk A, Donnellan E, Martyn T, Hassan OA, Alkharabsheh S, Watson C, Tang WH, Kwon D, Cremer P, Cheng F, Kanj M, Griffin B, Tan C, Rodriguez ER, Hanna M, Jaber W, Collier P. Association Between Atrial Uptake on Cardiac Scintigraphy With Technetium-99m-Pyrophosphate Labeled Bone-Seeking Tracers and Atrial Fibrillation. Circ Cardiovasc Imaging. 2022 May;15(5):e013829. doi: 10.1161/CIRCIMAGING.121.013829.

Excluded (reason: outside the field of interest)

32: Halme HL, Ihalainen T, Suomalainen O, Loimaala A, Mätzke S, Uusitalo V, Sipilä O, Hippeläinen E. Convolutional neural networks for detection of transthyretin amyloidosis in 2D scintigraphy images. EJNMMI Res. 2022 May 7;12(1):27. doi: 10.1186/s13550-022-00897-9.

Included in the systematic review but not in the meta-analysis (for possible patient data overlap with another study)

33: Aimo A, Merlo M, Porcari A, Georgiopoulos G, Pagura L, Vergaro G, Sinagra G, Emdin M, Rapezzi C. Redefining the epidemiology of cardiac amyloidosis. A systematic review and meta-analysis of screening studies. Eur J Heart Fail. 2022 Dec;24(12):2342-2351. doi: 10.1002/ehf.2532.

Excluded (reason: review)

34: Salvalaggio A, Cipriani A, Righetto S, Artioli P, Sinigiani G, De Michieli L, Cason M, Pilichou K, Cecchin D, Briani C. Incidental cardiac uptake of ^{99m}Tc-diphosphonates is predictive of poor outcome: data from 9616 bone scintigraphies. J Nucl Cardiol. 2022 Dec;29(6):3419-3425. doi: 10.1007/s12350-022-02961-2.

Included in the systematic review and in the meta-analysis

35: Gill SS, Fellin E, Stampke L, Zhao Y, Masri A. Clinical Clues and Diagnostic Workup of Cardiac Amyloidosis. Methodist Debakey Cardiovasc J. 2022 Mar 14;18(2):36-46. doi: 10.14797/mdcvj.1061.

Excluded (reason: outside the field of interest)

36: Xu J, Qiu Z, Yan M, Wang B, Song Z, Liu H, Wang M, Cen X. Prognostic Factors of AL-PCMM and AL-MM: A Single-Center Retrospective Study. Int J Med Sci. 2022 Mar 14;19(3):588-595. doi: 10.7150/ijms.61712.

Excluded (reason: outside the field of interest)

37: Rapezzi C, Aimo A, Serenelli M, Barison A, Vergaro G, Passino C, Panichella G, Sinagra G, Merlo M, Fontana M, Gillmore J, Quarta CC, Maurer MS, Kittleson MM, Garcia-Pavia P, Emdin M. Critical Comparison of Documents From Scientific Societies on Cardiac Amyloidosis: JACC State-of-the-Art Review. J Am Coll Cardiol. 2022 Apr 5;79(13):1288-1303.

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38: Barge-Caballero G, Barge-Caballero E, López-Pérez M, Bilbao-Quesada R, González-Babarro E, Gómez-Otero I, López-López A, Gutiérrez-Feijoo M, Varela-Román A, González-Juanatey C, Díaz-Castro Ó, Crespo-Leiro MG. Syncope in patients with transthyretin amyloid cardiomyopathy: clinical features and outcomes. Postgrad Med. 2022 May;134(4):420-428. doi: 10.1080/00325481.2022.2054174.

Excluded (reason: outside the field of interest)

39: Stan C, Mititelu R, Adam RD, Jurcuț R. Awareness of Nuclear Medicine Physicians in Romania Regarding the Diagnostic of Cardiac Amyloidosis-A Survey- Based Study. Diagnostics (Basel). 2022 Feb 21;12(2):556. doi: 10.3390/diagnostics12020556.

Excluded (reason: outside the field of interest)

40: Chiu A, Dasari S, Kurtin PJ, Theis JD, Vrana JA, Dispenzieri A, Rech KL, Dao LN, Howard MT, Grogan M, McPhail ED. Bone marrow amyloid: a comprehensive analysis of 1,469 samples, including amyloid type, clinical features, and morphologic distribution. *Amyloid*. 2022 Sep;29(3):156-164. doi: 10.1080/13506129.2022.2031963.

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41: Takashio S, Tokitsu T, Matsukawa M, Sakaino N, Fujimoto K, Sakamoto T, Noda K, Tsunoda R, Misumi I, Hirai N, Doi H, Koide S, Mizuno Y, Hirose T, Kurokawa H, Kajiwarra I, Ohba K, Miyamoto S, Araki S, Yamamoto E, Matsushita K, Ueda M, Tsujita K; Kumamoto Cardiac Amyloid Survey Investigators. Incidence, clinical characteristics, and diagnostic approach in transthyretin amyloid cardiomyopathy: The Kumamoto Cardiac Amyloidosis Survey. *J Cardiol*. 2022 Jul;80(1):49-55. doi: 10.1016/j.jjcc.2022.01.002.

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42: Uusitalo V, Suomalainen O, Loimaala A, Mätzke S, Heliö T. Prognostic Value of ^{99m}Tc-HMDP Scintigraphy in Elderly Patients With Chronic Heart Failure. *Heart Lung Circ*. 2022 May;31(5):629-637. doi: 10.1016/j.hlc.2021.11.018.

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43: Bourque JM, Schepart A, Bhambri R, Castaño A, O'Brien A, Chen Y, Prasad S, Roy A, Grodin JL. Temporal Trends in Diagnostic Testing Patterns for Wild-Type Transthyretin Amyloid Cardiomyopathy in the Medicare Fee-for-Service Population. *Am J Cardiol*. 2022 Mar 15;167:98-103. doi: 10.1016/j.amjcard.2021.11.048.

Excluded (reason: outside the field of interest)

44: Abdelazeem B, Manasrah N, Yousaf A, Gjeka R, Kunadi A. Light Chain (AL) Cardiac Amyloidosis: A Diagnostic Dilemma. *Cureus*. 2021 Nov 5;13(11):e19278. doi: 10.7759/cureus.19278.

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45: Singal AK, Bansal R, Singh A, Dorbala S, Sharma G, Gupta K, Saxena A, Bhargava B, Karthikeyan G, Ramakrishnan S, Bisoi AK, Hote MP, Rajashekar P, Chowdhury UK, Devagourou V, Patel C, Ray R, Arawa SK, Mishra S. Concomitant Transthyretin Amyloidosis and Severe Aortic Stenosis in Elderly Indian Population: A Pilot Study. *JACC CardioOncol*. 2021 Oct 19;3(4):565-576. doi: 10.1016/j.jacc.2021.08.008.

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46: Korosoglou G, Giusca S, André F, Aus dem Siepen F, Nunninger P, Kristen AV, Frey N. Diagnostic Work-Up of Cardiac Amyloidosis Using Cardiovascular Imaging: Current Standards and Practical Algorithms. *Vasc Health Risk Manag*. 2021 Oct 23;17:661-673. doi: 10.2147/VHRM.S295376.

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47: Bäcker HC, Galle SE, Lentzsch S, Freibott CE, Shoap S, Strauch RJ, Rosenwasser MP. Flexor tenosynovectomy in carpal tunnel syndrome as a screening tool for early diagnosis of amyloidosis. *Ir J Med Sci*. 2022 Oct;191(5):2427-2430. doi: 10.1007/s11845-021-02832-8.

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48: Shimon S, Zikri M, Haberman D, Livschitz S, Tshori S, Fabricant Y, Meledin V, Gandelman G, Goland S, George J. Transthyretin cardiac amyloidosis in patients after TAVR: clinical and echocardiographic findings and long term survival. *ESC Heart Fail*. 2021 Dec;8(6):4549-4561. doi: 10.1002/ehf2.13667.

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49: Fabbri G, Serenelli M, Cantone A, Sanguetoli F, Rapezzi C. Transthyretin amyloidosis in aortic stenosis: clinical and therapeutic implications. *Eur Heart J Suppl*. 2021 Oct 8;23(Suppl E):E128-E132. doi: 10.1093/eurheartj/suab107.

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50: Nguyen FD, Rodriguez M, Krittanawong C, Witteles R, Lenihan DJ. Misconceptions and Facts About Cardiac Amyloidosis. *Am J Cardiol*. 2021 Dec 1;160:99-105. doi: 10.1016/j.amjcard.2021.08.045.

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51: Roshankar G, White GC, Cadet S, Fine NM, Chan D, White JA, Jimenez-Zepeda V, Slomka PJ, Miller RJH. Quantitative technetium pyrophosphate and cardiovascular magnetic resonance in patients with suspected cardiac amyloidosis. *J Nucl Cardiol*. 2022 Oct;29(5):2679-2690. doi: 10.1007/s12350-021-02806-4.

Excluded (reason: outside the field of interest)

52: Bonelli A, Paris S, Nardi M, Henein MY, Agricola E, Troise G, Faggiano P. Aortic Valve Stenosis and Cardiac Amyloidosis: A Misleading Association. *J Clin Med*. 2021 Sep 18;10(18):4234. doi: 10.3390/jcm10184234.

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53: Patel KP, Scully PR, Nitsche C, Kammerlander AA, Joy G, Thornton G, Hughes R, Williams S, Tillin T, Captur G, Chacko L, Kelion A, Sabharwal N, Newton JD, Kennon S, Ozkor M, Mullen M, Hawkins PN, Gillmore JD, Menezes L, Pugliese F, Hughes AD, Fontana M, Lloyd G, Treibel TA, Mascherbauer J, Moon JC. Impact of afterload and infiltration on coexisting aortic stenosis and transthyretin amyloidosis. *Heart*. 2022 Jan;108(1):67-72. doi: 10.1136/heartjnl-2021-319922.

Excluded (reason: outside the field of interest)

54: AbouEzzeddine OF, Davies DR, Scott CG, Fayyaz AU, Askew JW, McKie PM, Noseworthy PA, Johnson GB, Dunlay SM, Borlaug BA, Chareonthaitawee P, Roger VL, Dispenzieri A, Grogan M, Redfield MM. Prevalence of Transthyretin Amyloid Cardiomyopathy in Heart Failure With Preserved Ejection Fraction. *JAMA Cardiol*. 2021 Nov 1;6(11):1267-1274. doi: 10.1001/jamacardio.2021.3070.

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55: Dayco J, Weaver M, Sumbal N, Theisen R, Raheem S. Ruptured Bullae: A Case of Transthyretin Cardiac Amyloidosis. *Cureus*. 2021 Jul 11;13(7):e16318. doi: 10.7759/cureus.16318.

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56: Tini G, Sessarego E, Benenati S, Vianello PF, Musumeci B, Autore C, Canepa M. Yield of bone scintigraphy screening for transthyretin-related cardiac amyloidosis in different conditions: Methodological issues and clinical implications. *Eur J Clin Invest*. 2021 Dec;51(12):e13665. doi: 10.1111/eci.13665.

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57: Oye M, Dhruva P, Kandah F, Oye M, Missov E. Cardiac amyloid presenting as cardiogenic shock: case series. *Eur Heart J Case Rep*. 2021 Jul 26;5(7):ytab252. doi: 10.1093/ehjcr/ytab252.

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58: Melero Polo J, Roteta Unceta-Barrenechea A, Revilla Martí P, Pérez-Palacios R, Gracia Gutiérrez A, Bueno Juana E, Andrés Gracia A, Atienza Ayala S, Aibar Arregui MÁ. Echocardiographic markers of cardiac amyloidosis in patients with heart failure and left ventricular hypertrophy. *Cardiol J*. 2023;30(2):266-275. doi: 10.5603/CJ.a2021.0085.

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59: Pan Y, Wang Y, Zhou Y, Xi Y, Zhang Y. ^{99m}Tc-Pyrophosphate Retention in Atelectatic Pulmonary Tissues of a Patient With Transthyretin Cardiac Amyloidosis. *Clin Nucl Med*. 2021 Oct 1;46(10):820-821. doi: 10.1097/RLU.0000000000003816.

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60: Godara A, Riesenburger RI, Zhang DX, Varga C, Fogaren T, Siddiqui NS, Yu A, Wang A, Mastroianni M, Dowd R, Nail TJ, McPhail ED, Kurtin PJ, Theis JD, Toskic D, Arkun K, Pilichowska M, Kryzanski J, Patel AR, Comenzo R.

Association between spinal stenosis and wild-type ATTR amyloidosis. *Amyloid*. 2021 Dec;28(4):226-233. doi: 10.1080/13506129.2021.1950681.

Excluded (reason: outside the field of interest)

61: Goland S, Volodarsky I, Fabricant Y, Livschitz S, Tshori S, Cuciuc V, Zilberman L, Fugenfirov I, Meledin V, Shimoni S, Josfberg S, George J. Wild-type TTR amyloidosis among patients with unexplained heart failure and systolic LV dysfunction. *PLoS One*. 2021 Jul 9;16(7):e0254104. doi: 10.1371/journal.pone.0254104.

Excluded (reason: outside the field of interest)

62: Inomata T, Tahara N, Nakamura K, Endo J, Ueda M, Ishii T, Kitano Y, Koyama J. Diagnosis of wild-type transthyretin amyloid cardiomyopathy in Japan: red-flag symptom clusters and diagnostic algorithm. *ESC Heart Fail*. 2021 Aug;8(4):2647-2659. doi: 10.1002/ehf2.13473.

Excluded (reason: outside the field of interest)

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