

Mathew S Maurer

List of Publications by Year in descending order

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309
papers

22,886
citations

13865

67
h-index

9860

141
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316
all docs

316
docs citations

316
times ranked

19195
citing authors

#	ARTICLE	IF	CITATIONS
1	The Wiggers Diagram: Hemodynamic Changes in Cardiac Amyloidosis. Journal of Cardiac Failure, 2023, 29, 217-219.	1.7	2
2	Transthyretin cardiac amyloidosis in continental Western Europe: an insight through the Transthyretin Amyloidosis Outcomes Survey (THAOS). European Heart Journal, 2022, 43, 391-400.	2.2	105
3	The importance of SPECT cardiac reconstruction for accurate 99mTc-pyrophosphate interpretation in TTR amyloidosis. Journal of Nuclear Cardiology, 2022, 29, 1478-1480.	2.1	2
4	Screening for ATTR amyloidosis in the clinic: overlapping disorders, misdiagnosis, and multiorgan awareness. Heart Failure Reviews, 2022, 27, 785-793.	3.9	31
5	Addendum 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. Journal of Cardiac Failure, 2022, 28, e1-e4.	1.7	8
6	Surveillance for disease progression of transthyretin amyloidosis after heart transplantation in the era of novel disease modifying therapies. Journal of Heart and Lung Transplantation, 2022, 41, 199-207.	0.6	9
7	Prescriptions for Potentially Inappropriate Medications from the Beers Criteria Among Older Adults Hospitalized for Heart Failure. Journal of Cardiac Failure, 2022, 28, 906-915.	1.7	7
8	Too Stiff But Still Got Rhythm. JACC: Cardiovascular Imaging, 2022, 15, 30-32.	5.3	4
9	Clinico-histopathologic and single-nuclei RNA-sequencing insights into cardiac injury and microthrombi in critical COVID-19. JCI Insight, 2022, 7, .	5.0	14
10	Amyloid Cardiomyopathy in Older Adults. Current Geriatrics Reports, 2022, 11, 1.	1.1	0
11	Proposed Cardiac End Points for Clinical Trials in Immunoglobulin Light Chain Amyloidosis: Report From the Amyloidosis Forum Cardiac Working Group. Circulation: Heart Failure, 2022, 15, CIRCHEARTFAILURE121009038.	3.9	6
12	Critical Comparison of Documents From Scientific Societies on Cardiac Amyloidosis. Journal of the American College of Cardiology, 2022, 79, 1288-1303.	2.8	35
13	Impact of bone marrow minimal residual disease status on quality of organ response in systemic AL amyloidosis. American Journal of Hematology, 2022, 97, .	4.1	1
14	Association of Transthyretin Val122Ile Variant With Incident Heart Failure Among Black Individuals. JAMA - Journal of the American Medical Association, 2022, 327, 1368.	7.4	19
15	Remote Cardiac Monitoring in Patients With Heart Failure. JAMA Cardiology, 2022, 7, 556.	6.1	22
16	Racial Differences in Val122Ile-Associated Transthyretin Cardiac Amyloidosis. Journal of Cardiac Failure, 2022, 28, 950-959.	1.7	8
17	Proteomics profiling reveals a distinct high-risk molecular subtype of hypertrophic cardiomyopathy. Heart, 2022, 108, 1807-1814.	2.9	4
18	Impact of light chain isotype on clinical features and outcomes in systemic AL amyloidosis. Leukemia and Lymphoma, 2022, , 1-5.	1.3	0

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19	Characteristics of patients with autonomic dysfunction in the Transthyretin Amyloidosis Outcomes Survey (THAOS). Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2022, 29, 175-183.	3.0	7
20	Sex Differences in Wild-Type Transthyretin Amyloidosis: An Analysis from the Transthyretin Amyloidosis Outcomes Survey (THAOS). Cardiology and Therapy, 2022, 11, 393-405.	2.6	7
21	Systemic embolism in amyloid transthyretin cardiomyopathy. European Journal of Heart Failure, 2022, 24, 1387-1396.	7.1	23
22	Clinical and genetic profile of patients enrolled in the Transthyretin Amyloidosis Outcomes Survey (THAOS): 14-year update. Orphanet Journal of Rare Diseases, 2022, 17, .	2.7	22
23	Relation of Body Mass Index to Transthyretin Cardiac Amyloidosis Particularly in Black and Hispanic Patients (from the SCAN-MP Study). American Journal of Cardiology, 2022, 177, 116-120.	1.6	2
24	Radiopharmaceutical supply disruptions and the use of 99mTc-hydroxymethylene diphosphonate as an alternative to 99mTc-pyrophosphate for the diagnosis of transthyretin cardiac amyloidosis: An ASNC Information Statement. Journal of Nuclear Cardiology, 2022, 29, 2748-2760.	2.1	4
25	The diagnostic challenges of cardiac amyloidosis: A practical approach to the two main types. Blood Reviews, 2021, 45, 100720.	5.7	15
26	Difference in Metabolomic Response to Exercise between Patients with and without Hypertrophic Cardiomyopathy. Journal of Cardiovascular Translational Research, 2021, 14, 246-255.	2.4	16
27	Unveiling outcomes in coexisting severe aortic stenosis and transthyretin cardiac amyloidosis. European Journal of Heart Failure, 2021, 23, 250-258.	7.1	71
28	Association of Midlife Cardiovascular Risk Factors With the Risk of Heart Failure Subtypes Later in Life. Journal of Cardiac Failure, 2021, 27, 435-444.	1.7	6
29	Predicting the development of adverse cardiac events in patients with hypertrophic cardiomyopathy using machine learning. International Journal of Cardiology, 2021, 327, 117-124.	1.7	12
30	Impact of Tafamidis on Health-Related Quality of Life in Patients With Transthyretin Amyloid Cardiomyopathy (from the Tafamidis in Transthyretin Cardiomyopathy Clinical Trial). American Journal of Cardiology, 2021, 141, 98-105.	1.6	21
31	Diagnosing Transthyretin Cardiac Amyloidosis by Technetium Tc 99m Pyrophosphate. JACC: Cardiovascular Imaging, 2021, 14, 1221-1231.	5.3	52
32	Frailty subtypes and recovery in older survivors of acute respiratory failure: a pilot study. Thorax, 2021, 76, 350-359.	5.6	6
33	Inclusion of Performance Parameters and Patient Context in the Clinical Practice Guidelines for Heart Failure. Journal of Cardiac Failure, 2021, 27, 190-197.	1.7	2
34	Efficacy of Tafamidis in Patients With Hereditary and Wild-Type Transthyretin Amyloid Cardiomyopathy. JACC: Heart Failure, 2021, 9, 115-123.	4.1	52
35	Sex Differences in the Phenotype of Transthyretin Cardiac Amyloidosis Due to Val122Ile Mutation: Insights from Noninvasive Pressure-Volume Analysis. Journal of Cardiac Failure, 2021, 27, 67-74.	1.7	20
36	Anticoagulation with warfarin compared to novel oral anticoagulants for atrial fibrillation in adults with transthyretin cardiac amyloidosis: comparison of thromboembolic events and major bleeding. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2021, 28, 30-34.	3.0	31

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37	Transthyretin cardiac amyloidosis: A treatable form of heart failure with a preserved ejection fraction. Trends in Cardiovascular Medicine, 2021, 31, 59-66.	4.9	14
38	Editorial for “Reference Ranges, Diagnostic and Prognostic Utility of Native T1 Mapping and Extracellular Volume for Cardiac Amyloidosis: A Meta-analysis”. Journal of Magnetic Resonance Imaging, 2021, 53, 1469-1470.	3.4	1
39	Response by Unlu et al to Letter Regarding Article, “Polypharmacy in Older Adults Hospitalized for Heart Failure”. Circulation: Heart Failure, 2021, 14, e008305.	3.9	1
40	Diagnosis and treatment of cardiac amyloidosis: a position statement of the ESC Working Group on Myocardial and Pericardial Diseases. European Heart Journal, 2021, 42, 1554-1568.	2.2	434
41	Diagnosis and treatment of cardiac amyloidosis. A position statement of the European Society of Cardiology Working Group on Myocardial and Pericardial Diseases. European Journal of Heart Failure, 2021, 23, 512-526.	7.1	153
42	The association of prior carpal tunnel syndrome surgery with adverse cardiovascular outcomes and long-term mortality after aortic valve replacement. IJC Heart and Vasculature, 2021, 33, 100741.	1.1	0
43	ATTR amyloidosis during the COVID-19 pandemic: insights from a global medical roundtable. Orphanet Journal of Rare Diseases, 2021, 16, 204.	2.7	11
44	False Positive 99mTc-Pyrophosphate Scanning Leading to Inappropriate Tafamidis Prescriptions. JACC: Cardiovascular Imaging, 2021, 14, 2042-2044.	5.3	13
45	Cognition predicts days-alive-out-of-hospital after LVAD implantation. International Journal of Artificial Organs, 2021, 44, 952-955.	1.4	5
46	Pathophysiology and Therapeutic Approaches to Cardiac Amyloidosis. Circulation Research, 2021, 128, 1554-1575.	4.5	52
47	Characteristics of Patients with Late- vs. Early-Onset Val30Met Transthyretin Amyloidosis from the Transthyretin Amyloidosis Outcomes Survey (THAOS). Neurology and Therapy, 2021, 10, 753-766.	3.2	14
48	Prefrailty, impairment in physical function, and risk of incident heart failure among older adults. Journal of the American Geriatrics Society, 2021, 69, 2486-2497.	2.6	14
49	Prediction of Genotype Positivity in Patients With Hypertrophic Cardiomyopathy Using Machine Learning. Circulation Genomic and Precision Medicine, 2021, 14, e003259.	3.6	8
50	Phenotypic Differences of Glu89Gln Genotype in ATTR Amyloidosis From Endemic Loci: Update From THAOS. Cardiology and Therapy, 2021, 10, 481-490.	2.6	8
51	ASNC/AHA/ASE/EANM/HFSA/ISA/SCMR/SNMMI Expert Consensus Recommendations for Multimodality Imaging in Cardiac Amyloidosis: Part 1 of “Evidence Base and Standardized Methods of Imaging. Circulation: Cardiovascular Imaging, 2021, 14, e000029.	2.6	48
52	ASNC/AHA/ASE/EANM/HFSA/ISA/SCMR/SNMMI Expert Consensus Recommendations for Multimodality Imaging in Cardiac Amyloidosis: Part 2 of “Diagnostic Criteria and Appropriate Utilization. Circulation: Cardiovascular Imaging, 2021, 14, e000030.	2.6	16
53	Comprehensive Proteomics Profiling Reveals Circulating Biomarkers of Hypertrophic Cardiomyopathy. Circulation: Heart Failure, 2021, 14, e007849.	3.9	26
54	Addendum to ASNC/AHA/ASE/EANM/HFSA/ISA/SCMR/SNMMI expert consensus recommendations for multimodality imaging in cardiac amyloidosis: Part 1 of “evidence base and standardized methods of imaging. Journal of Nuclear Cardiology, 2021, 28, 1769-1774.	2.1	34

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55	Relation between Modified Body Mass Index and Adverse Outcomes after Aortic Valve Implantation. American Journal of Cardiology, 2021, 153, 94-100.	1.6	2
56	Gene Editing â€” A Cure for Transthyretin Amyloidosis?. New England Journal of Medicine, 2021, 385, 558-559.	27.0	8
57	Deep Learning Analysis of Echocardiographic Images to Predict Positive Genotype in Patients With Hypertrophic Cardiomyopathy. Frontiers in Cardiovascular Medicine, 2021, 8, 669860.	2.4	10
58	Factors associated with changes in serum transthyretin after treatment with tafamidis and outcomes in transthyretin cardiac amyloidosis. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2021, , 1-2.	3.0	4
59	Cardiopulmonary exercise testing in patients with Cardiac Amyloidosis. Clinical Lymphoma, Myeloma and Leukemia, 2021, 21, 545-548.	0.4	6
60	Two Decades of Cardiac Amyloidosis. JACC: CardioOncology, 2021, 3, 522-533.	4.0	20
61	Recognition and Implications of Undiagnosed Cardiac Amyloid Patients in HFpEF Trials. JACC: Heart Failure, 2021, 9, 803-806.	4.1	3
62	Cardiac Scintigraphy and Screening for Transthyretin Cardiac Amyloidosis. Circulation, 2021, 144, 1005-1007.	1.6	8
63	Effects of Septal Reduction Therapy on Acute Cardiovascular Events and All-Cause Mortality in Patients with Hypertrophic Cardiomyopathy. International Heart Journal, 2021, 62, 1035-1041.	1.0	2
64	Phase 1a/b study of monoclonal antibody CAEL-101 (11-1F4) in patients with AL amyloidosis. Blood, 2021, 138, 2632-2641.	1.4	48
65	The Potential Role of EHR data in optimizing eligibility criteria definition for cardiovascular outcome trials. International Journal of Medical Informatics, 2021, 156, 104587.	3.3	0
66	Sex-Related Risk of Cardiac Involvement in Hereditary Transthyretin Amyloidosis. JACC: Heart Failure, 2021, 9, 736-746.	4.1	26
67	Temporal Trends of Wild-Type Transthyretin Amyloid Cardiomyopathy in the Transthyretin Amyloidosis Outcomes Survey. JACC: CardioOncology, 2021, 3, 537-546.	4.0	21
68	Cardiovascular Diseases That Have Emerged From the Darkness. Journal of the American Heart Association, 2021, 10, e021095.	3.7	5
69	ATTR Amyloidosis: Current and Emerging Management Strategies. JACC: CardioOncology, 2021, 3, 488-505.	4.0	56
70	Lack of Association Between Neurohormonal Blockade and Survival in Transthyretin Cardiac Amyloidosis. Journal of the American Heart Association, 2021, 10, e022859.	3.7	19
71	Can lightning strike twice? Wild-type transthyretin cardiac amyloidosis associated with rare liver disease. Oxford Medical Case Reports, 2021, 2021, omab113.	0.4	0
72	Impact of Light Chain Isotype on Clinical Features and Outcomes in Systemic AL Amyloidosis. Blood, 2021, 138, 4726-4726.	1.4	0

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73	A Proposal to Accelerate Widespread Implementation of Geriatric Cardiology. Trends in Cardiovascular Medicine, 2021, , .	4.9	1
74	158â€fRacial differences in val122lle associated transthyretin cardiac amyloidosis. European Heart Journal Supplements, 2021, 23, .	0.1	0
75	Abstract 9735: Cost Effectiveness of Screening for Valvular Heart Disease Using the Valvenet Deep Learning Electrocardiogram Model. Circulation, 2021, 144, .	1.6	0
76	Estimating cancer risk from 99mTc pyrophosphate imaging for transthyretin cardiac amyloidosis. Journal of Nuclear Cardiology, 2020, 27, 215-224.	2.1	7
77	Cardiac Amyloidosis. , 2020, , , 301-310.e3.		1
78	Eosinophils, Lymphocytes, and Myocytes, Oh My: HIV-Associated Myocarditis. American Journal of Medicine, 2020, 133, 52-55.	1.5	1
79	ASNC/AHA/ASE/EANM/HFSA/ISA/SCMR/SNMMI expert consensus recommendations for multimodality imaging in cardiac amyloidosis: Part 2 of 2â€Diagnostic criteria and appropriate utilization. Journal of Nuclear Cardiology, 2020, 27, 659-673.	2.1	97
80	99mTechnetium pyrophosphate scintigraphy with cadmium zinc telluride cameras is a highly sensitive and specific imaging modality to diagnose transthyretin cardiac amyloidosis. Journal of Nuclear Cardiology, 2020, 27, 371-380.	2.1	14
81	Prescribing Patterns of HeartâFailure-Exacerbating Medications Following a Heart Failure Hospitalization. JACC: Heart Failure, 2020, 8, 25-34.	4.1	21
82	Tafamidisâ€A Pricey Therapy for a Not-So-Rare Condition. JAMA Cardiology, 2020, 5, 247.	6.1	33
83	Stabilization of Cardiac Function With Diflunisal in Transthyretin (ATTR) Cardiac Amyloidosis. Journal of Cardiac Failure, 2020, 26, 753-759.	1.7	57
84	Markers of nutritional status and inflammation in transthyretin cardiac amyloidosis: association with outcomes and the clinical phenotype. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2020, 27, 73-80.	3.0	17
85	The quintessential form of diastolic heart failure in older adults: Wild type transthyretin cardiac amyloidosis. Clinical Cardiology, 2020, 43, 171-178.	1.8	5
86	Comparing outcomes for infiltrative and restrictive cardiomyopathies under the new heart transplant allocation system. Clinical Transplantation, 2020, 34, e14109.	1.6	14
87	Diuretic Dose and NYHA Functional Class Are Independent Predictors of Mortality in Patients With Transthyretin Cardiac Amyloidosis. JACC: CardioOncology, 2020, 2, 414-424.	4.0	37
88	The Amyloidosis Forum: a public private partnership to advance drug development in AL amyloidosis. Orphanet Journal of Rare Diseases, 2020, 15, 268.	2.7	9
89	Response by Kazi et al to Letter Regarding Article, â€Cost-Effectiveness of Tafamidis Therapy for Transthyretin Amyloid Cardiomyopathyâ€ Circulation, 2020, 142, e212-e213.	1.6	1
90	Cardiac Amyloidosis is Underdiagnosed in Patients Undergoing Transcatheter Aortic Valve Replacement. Structural Heart, 2020, 4, 512-514.	0.6	1

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91	Polypharmacy in Older Adults Hospitalized for Heart Failure. <i>Circulation: Heart Failure</i> , 2020, 13, e006977.	3.9	102
92	Outcomes after heart transplantation for AL compared to ATTR cardiac amyloidosis. <i>Clinical Transplantation</i> , 2020, 34, e14028.	1.6	15
93	Gerontechnology for Older Adults With Cardiovascular Diseases. <i>Journal of the American College of Cardiology</i> , 2020, 76, 2650-2670.	2.8	66
94	Systemic Amyloidosis due to Monoclonal Immunoglobulins. <i>Hematology/Oncology Clinics of North America</i> , 2020, 34, 1055-1068.	2.2	5
95	Avoiding misdiagnosis: expert consensus recommendations for the suspicion and diagnosis of transthyretin amyloidosis for the general practitioner. <i>BMC Family Practice</i> , 2020, 21, 198.	2.9	60
96	Mavacamten for treatment of symptomatic obstructive hypertrophic cardiomyopathy (EXPLORER-HCM): a randomised, double-blind, placebo-controlled, phase 3 trial. <i>Lancet</i> , 2020, 396, 759-769.	13.7	481
97	A Peak into the Pace of Cardiac Amyloidosis. <i>JACC: Clinical Electrophysiology</i> , 2020, 6, 1155-1157.	3.2	1
98	Transition of a Large Tertiary Heart Failure Program in Response to the COVID-19 Pandemic. <i>Circulation: Heart Failure</i> , 2020, 13, e007516.	3.9	17
99	Early-Onset of Transthyretin Amyloidosis in a Young Afro-Caribbean Woman With Thr60Ala Mutation. <i>JACC: Case Reports</i> , 2020, 2, 2063-2067.	0.6	2
100	Cardiac Scintigraphy With Technetium-99m-Labeled Bone-Seeking Tracers for Suspected Amyloidosis. <i>Journal of the American College of Cardiology</i> , 2020, 75, 2851-2862.	2.8	131
101	Cardiac Amyloidosis: Evolving Diagnosis and Management: A Scientific Statement From the American Heart Association. <i>Circulation</i> , 2020, 142, e7-e22.	1.6	338
102	DISCOVERY: prevalence of transthyretin (TTR) mutations in a US-centric patient population suspected of having cardiac amyloidosis. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2020, 27, 223-230.	3.0	17
103	Initial Monthly Cost of Tafamidis—the Real Price for Patients? Reply. <i>JAMA Cardiology</i> , 2020, 5, 848.	6.1	3
104	Norma: Participant 1,001 and the Value of Clinical Research. <i>Journal of the American Geriatrics Society</i> , 2020, 68, 2407-2408.	2.6	0
105	Untangling the physiology of transthyretin cardiac amyloidosis by leveraging echocardiographically derived pressure–volume indices. <i>European Heart Journal</i> , 2020, 41, 1448-1450.	2.2	9
106	Extrapulmonary manifestations of COVID-19. <i>Nature Medicine</i> , 2020, 26, 1017-1032.	30.7	2,300
107	Peripheral neuropathy symptoms in wild type transthyretin amyloidosis. <i>Journal of the Peripheral Nervous System</i> , 2020, 25, 265-272.	3.1	21
108	Cost-Effectiveness of Tafamidis Therapy for Transthyretin Amyloid Cardiomyopathy. <i>Circulation</i> , 2020, 141, 1214-1224.	1.6	147

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109	Phase 3 Multicenter Study of Revusiran in Patients with Hereditary Transthyretin-Mediated (hATTR) Amyloidosis with Cardiomyopathy (ENDEAVOUR). Cardiovascular Drugs and Therapy, 2020, 34, 357-370.	2.6	55
110	How Should Physicians Assess Myocardial Contraction?. JACC: Cardiovascular Imaging, 2020, 13, 873-878.	5.3	30
111	Impaired systemic venous capacitance: the neglected mechanism in patients with heart failure and a preserved ejection fraction?. European Journal of Heart Failure, 2020, 22, 173-176.	7.1	9
112	Cardiac Amyloidosis: Overlooked, Underappreciated, and Treatable. Annual Review of Medicine, 2020, 71, 203-219.	12.2	82
113	Salt Taste Sensitivity and Heart Failure Outcomes Following Heart Failure Hospitalization. American Journal of Cardiology, 2020, 127, 58-63.	1.6	2
114	Cardiac amyloidosis in severe aortic stenosis: we can find it but what should we do?. European Journal of Heart Failure, 2020, 22, 1863-1865.	7.1	2
115	Disparities, Uncertainties, and Societal Cost: Precision Medicine and Transthyretin Amyloidosis. American Journal of Medicine, 2020, 133, 892-894.	1.5	2
116	Older Adults Can Successfully Monitor Symptoms Using an Inclusively Designed Mobile Application. Journal of the American Geriatrics Society, 2020, 68, 1313-1318.	2.6	20
117	Characterization of the <scp>inflammatoryâ€œmetabolic</scp> phenotype of heart failure with a preserved ejection fraction: a hypothesis to explain influence of sex on the evolution and potential treatment of the disease. European Journal of Heart Failure, 2020, 22, 1551-1567.	7.1	93
118	Durable Mechanical Circulatory Support in Patients With Amyloid Cardiomyopathy. Circulation: Heart Failure, 2020, 13, e007931.	3.9	15
119	Outcomes By Cardiac Stage in Newly Diagnosed AL Amyloidosis: Results from Andromeda. Blood, 2020, 136, 44-45.	1.4	5
120	Daratumumab plus CyBorD for patients with newly diagnosed AL amyloidosis: safety run-in results of ANDROMEDA. Blood, 2020, 136, 71-80.	1.4	146
121	Abstract 15821: Patients With Amyloid Cardiomyopathy Who Receive Durable Mechanical Circulatory Support Are at Increased Risk for Mortality and Early Adverse Events. Circulation, 2020, 142, .	1.6	0
122	Sars-Cov-2 Infection and Systemic Light Chain Amyloidosis: The International Society of Amyloidosis' Survey. Blood, 2020, 136, 11-11.	1.4	0
123	Abstract 15242: Increased Mortality Among African American Patients With Heart Failure Caused by Hereditary Transthyretin Amyloid Cardiomyopathy. Circulation, 2020, 142, .	1.6	0
124	Abstract 14398: African Americans With Transthyretin Cardiac Amyloidosis Have a Lower Prevalence of Atrial Fibrillation but Increased Thromboembolic Events Than Caucasians. Circulation, 2020, 142, .	1.6	1
125	Left Ventricular Assist Device Therapy in Older Adults: Addressing Common Clinical Questions. Journal of the American Geriatrics Society, 2019, 67, 2410-2419.	2.6	13
126	Comparison of Effectiveness of Alcohol Septal Ablation Versus Ventricular Septal Myectomy on Acute Care Use for Cardiovascular Disease in Patients With Hypertrophic Cardiomyopathy. American Journal of Cardiology, 2019, 124, 1272-1278.	1.6	1

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127	Transthyretin Stabilization by AG10 in Symptomatic Transthyretin Amyloid Cardiomyopathy. Journal of the American College of Cardiology, 2019, 74, 285-295.	2.8	170
128	The Truth Is Unfolding About Transthyretin Cardiac Amyloidosis. Circulation, 2019, 140, 27-30.	1.6	9
129	Association between reduced myocardial contraction fraction and cardiovascular disease outcomes: The Multi-Ethnic Study of Atherosclerosis. International Journal of Cardiology, 2019, 293, 10-16.	1.7	16
130	Days alive and outside of hospital from diagnosis of transthyretin vs. light chain cardiac amyloidosis. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2019, 26, 4-5.	3.0	1
131	Chronotropic incompetence and autonomic dysfunction as mechanisms of dyspnoea in patients with late stage cardiac amyloidosis. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2019, 26, 134-135.	3.0	3
132	One year follow up analysis of the phase 1a/b study of chimeric fibril-reactive monoclonal antibody 11-1F4 in patients with AL amyloidosis. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2019, 26, 115-116.	3.0	24
133	Cost-effectiveness of technetium pyrophosphate scintigraphy versus heart biopsy for the diagnosis of transthyretin amyloidosis. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2019, 26, 71-72.	3.0	3
134	Association of Carpal Tunnel Syndrome With Amyloidosis, Heart Failure, and Adverse Cardiovascular Outcomes. Journal of the American College of Cardiology, 2019, 74, 15-23.	2.8	77
135	Application of Proteomics Profiling for Biomarker Discovery in Hypertrophic Cardiomyopathy. Journal of Cardiovascular Translational Research, 2019, 12, 569-579.	2.4	17
136	Beyond the Valve and into the Muscle: A Review of Coexisting Aortic Stenosis and Transthyretin Cardiac Amyloidosis. Structural Heart, 2019, 3, 462-468.	0.6	2
137	Perspectives on Implementing a Multidomain Approach to Caring for Older Adults With Heart Failure. Journal of the American Geriatrics Society, 2019, 67, 2593-2599.	2.6	12
138	ASNC/AHA/ASE/EANM/HFSA/ISA/SCMR/SNMMI expert consensus recommendations for multimodality imaging in cardiac amyloidosis: Part 1 of "evidence base and standardized methods of imaging. Journal of Nuclear Cardiology, 2019, 26, 2065-2123.	2.1	230
139	Expert Consensus Recommendations for the Suspicion and Diagnosis of Transthyretin Cardiac Amyloidosis. Circulation: Heart Failure, 2019, 12, e006075.	3.9	312
140	ASNC/AHA/ASE/EANM/HFSA/ISA/SCMR/SNMMI Expert Consensus Recommendations for Multimodality Imaging in Cardiac Amyloidosis: Part 2 of "Diagnostic Criteria and Appropriate Utilization. Journal of Cardiac Failure, 2019, 25, 854-865.	1.7	70
141	ASNC/AHA/ASE/EANM/HFSA/ISA/SCMR/SNMMI Expert Consensus Recommendations for Multimodality Imaging in Cardiac Amyloidosis: Part 1 of "Evidence Base and Standardized Methods of Imaging. Journal of Cardiac Failure, 2019, 25, e1-e39.	1.7	107
142	Anticoagulation with Warfarin versus Novel Oral Anticoagulants in Atrial Fibrillation in Amyloid Transthyretin Amyloidosis Cardiomyopathy: A Retrospective Cohort Study. Journal of Cardiac Failure, 2019, 25, S82.	1.7	3
143	Amyloidosis of the Brain and Heart. JACC: Heart Failure, 2019, 7, 129-131.	4.1	5
144	Facilitated Peer Mentorship to Support Aging Research: A REACH Evaluation of the CoMPAdRE Program. Journal of the American Geriatrics Society, 2019, 67, 804-810.	2.6	8

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145	Transthyretin Amyloid Cardiomyopathy. Journal of the American College of Cardiology, 2019, 73, 2872-2891.	2.8	573
146	Trends and causes of hospitalizations in patients with amyloidosis. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2019, 26, 177-178.	3.0	5
147	Cardiac Amyloidosis A Rare Disease in Older Adults Hospitalized for Heart Failure?. Circulation: Heart Failure, 2019, 12, e006169.	3.9	13
148	Deprescribing in Older Adults With Cardiovascular Disease. Journal of the American College of Cardiology, 2019, 73, 2584-2595.	2.8	126
149	Indications for Î²-Blocker Prescriptions in Heart Failure with Preserved Ejection Fraction. Journal of the American Geriatrics Society, 2019, 67, 1461-1466.	2.6	14
150	Myocardial Contraction Fraction Predicts Cardiovascular Events in Patients With Hypertrophic Cardiomyopathy and Normal Ejection Fraction. Journal of Cardiac Failure, 2019, 25, 450-456.	1.7	19
151	Top Ten Tips Palliative Care Clinicians Should Know About Caring for Patients with Left Ventricular Assist Devices. Journal of Palliative Medicine, 2019, 22, 437-441.	1.1	11
152	Demyelinating Neuropathy in a Patient Treated With Revusiran for Transthyretin (Thr60Ala) Amyloidosis. Journal of Clinical Neuromuscular Disease, 2019, 20, 120-128.	0.7	8
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