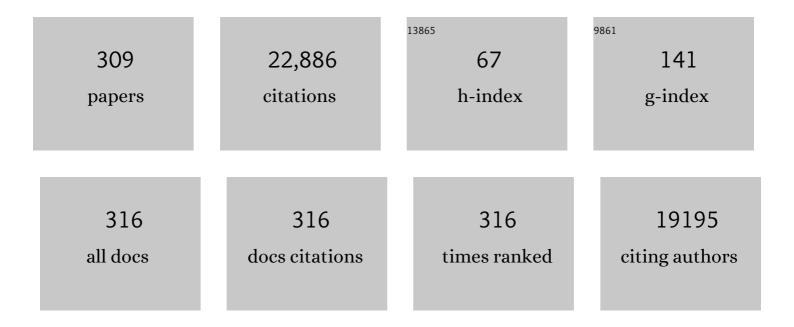
## Mathew S Maurer

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Extrapulmonary manifestations of COVID-19. Nature Medicine, 2020, 26, 1017-1032.	30.7	2,300
2	Tafamidis Treatment for Patients with Transthyretin Amyloid Cardiomyopathy. New England Journal of Medicine, 2018, 379, 1007-1016.	27.0	1,558
3	Nonbiopsy Diagnosis of Cardiac Transthyretin Amyloidosis. Circulation, 2016, 133, 2404-2412.	1.6	1,335
4	Frailty Assessment in the Cardiovascular Care of Older Adults. Journal of the American College of Cardiology, 2014, 63, 747-762.	2.8	850
5	Transthyretin Amyloid Cardiomyopathy. Journal of the American College of Cardiology, 2019, 73, 2872-2891.	2.8	573
6	<sup>99m</sup> Tc-Pyrophosphate Scintigraphy for Differentiating Light-Chain Cardiac Amyloidosis From the Transthyretin-Related Familial and Senile Cardiac Amyloidoses. Circulation: Cardiovascular Imaging, 2013, 6, 195-201.	2.6	499
7	Unveiling transthyretin cardiac amyloidosis and its predictors among elderly patients with severe aortic stenosis undergoing transcatheter aortic valve replacement. European Heart Journal, 2017, 38, 2879-2887.	2.2	489
8	Mavacamten for treatment of symptomatic obstructive hypertrophic cardiomyopathy (EXPLORER-HCM): a randomised, double-blind, placebo-controlled, phase 3 trial. Lancet, The, 2020, 396, 759-769.	13.7	481
9	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
10	The Impact of Frailty Status on Survival After Transcatheter Aortic Valve Replacement in Older Adults With Severe Aortic Stenosis. JACC: Cardiovascular Interventions, 2012, 5, 974-981.	2.9	411
11	Genotype and Phenotype of Transthyretin Cardiac Amyloidosis. Journal of the American College of Cardiology, 2016, 68, 161-172.	2.8	338
12	Cardiac Amyloidosis: Evolving Diagnosis and Management: A Scientific Statement From the American Heart Association. Circulation, 2020, 142, e7-e22.	1.6	338
13	Addressing Common Questions Encountered in the Diagnosis and Management of Cardiac Amyloidosis. Circulation, 2017, 135, 1357-1377.	1.6	319
14	Effects of Patisiran, an RNA Interference Therapeutic, on Cardiac Parameters in Patients With Hereditary Transthyretin-Mediated Amyloidosis. Circulation, 2019, 139, 431-443.	1.6	319
15	Expert Consensus Recommendations for the Suspicion and Diagnosis of Transthyretin Cardiac Amyloidosis. Circulation: Heart Failure, 2019, 12, e006075.	3.9	312
16	Multicenter Study of Planar Technetium 99m Pyrophosphate Cardiac Imaging. JAMA Cardiology, 2016, 1, 880.	6.1	304
17	THAOS – The Transthyretin Amyloidosis Outcomes Survey: initial report on clinical manifestations in patients with hereditary and wild-type transthyretin amyloidosis. Current Medical Research and Opinion, 2013, 29, 63-76.	1.9	246
18	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 Nuclear Cardiology, 2019, 26, 2065-2123.	2.1	230

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19	Heart Failure With a Normal Ejection Fraction. Circulation, 2003, 107, 656-658.	1.6	226
20	Single-beat estimation of end-diastolic pressure-volume relationship: a novel method with potential for noninvasive application. American Journal of Physiology - Heart and Circulatory Physiology, 2006, 291, H403-H412.	3.2	223
21	Multimorbidity in Older Adults With Cardiovascular Disease. Journal of the American College of Cardiology, 2018, 71, 2149-2161.	2.8	216
22	Review and Analysis of Existing Mobile Phone Apps to Support Heart Failure Symptom Monitoring and Self-Care Management Using the Mobile Application Rating Scale (MARS). JMIR MHealth and UHealth, 2016, 4, e74.	3.7	212
23	Prospective evaluation of the morbidity and mortality of wild-type and V122I mutant transthyretin amyloid cardiomyopathy: The Transthyretin Amyloidosis Cardiac Study (TRACS). American Heart Journal, 2012, 164, 222-228.e1.	2.7	209
24	Left ventricular mass predicts heart failure not related to previous myocardial infarction: the Cardiovascular Health Study. European Heart Journal, 2008, 29, 741-747.	2.2	203
25	Cardiac Care for Older Adults. Journal of the American College of Cardiology, 2011, 57, 1801-1810.	2.8	187
26	Diastolic dysfunction. Journal of the American College of Cardiology, 2004, 44, 1543-1549.	2.8	185
27	Left Heart Failure With a Normal Ejection Fraction: Identification of Different Pathophysiologic Mechanisms. Journal of Cardiac Failure, 2005, 11, 177-187.	1.7	184
28	Natural history and therapy of TTR-cardiac amyloidosis: emerging disease-modifying therapies from organ transplantation to stabilizer and silencer drugs. Heart Failure Reviews, 2015, 20, 163-178.	3.9	178
29	Transthyretin Stabilization by AG10 in Symptomatic Transthyretin AmyloidÂCardiomyopathy. Journal of the American College of Cardiology, 2019, 74, 285-295.	2.8	170
30	Ventricular Structure and Function in Hypertensive Participants With Heart Failure and a Normal Ejection Fraction. Journal of the American College of Cardiology, 2007, 49, 972-981.	2.8	166
31	Secondary Prevention of Atherosclerotic Cardiovascular Disease in Older Adults. Circulation, 2013, 128, 2422-2446.	1.6	166
32	Reduced Handgrip Strength as a Marker of Frailty Predicts Clinical Outcomes in Patients With Heart Failure Undergoing Ventricular Assist Device Placement. Journal of Cardiac Failure, 2014, 20, 310-315.	1.7	155
33	Prevalence and Prognostic Significance of Low QRS Voltage Among the Three Main Types of Cardiac Amyloidosis. American Journal of Cardiology, 2014, 114, 1089-1093.	1.6	154
34	Diagnosis and treatment of cardiac amyloidosis. A position statement of the European Society of Cardiology <scp>W</scp> orking <scp>G</scp> roup on <scp>M</scp> yocardial and <scp>P</scp> ericardial <scp>D</scp> iseases. European Journal of Heart Failure, 2021, 23, 512-526.	7.1	153
35	International External Validation Study of the 2014 European Society of Cardiology Guidelines on Sudden Cardiac Death Prevention in Hypertrophic Cardiomyopathy (EVIDENCE-HCM). Circulation, 2018, 137, 1015-1023.	1.6	149
36	Cardiac amyloidosis: the great pretender. Heart Failure Reviews, 2015, 20, 117-124.	3.9	147

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37	Cost-Effectiveness of Tafamidis Therapy for Transthyretin Amyloid Cardiomyopathy. Circulation, 2020, 141, 1214-1224.	1.6	147
38	Daratumumab plus CyBorD for patients with newly diagnosed AL amyloidosis: safety run-in results of ANDROMEDA. Blood, 2020, 136, 71-80.	1.4	146
39	Rationale and design of the Transcatheter Aortic Valve Replacement to UNload the Left ventricle in patients with ADvanced heart failure (TAVR UNLOAD) trial. American Heart Journal, 2016, 182, 80-88.	2.7	142
40	Knowledge Gaps in Cardiovascular Care of the Older Adult Population. Circulation, 2016, 133, 2103-2122.	1.6	139
41	Arterial Stiffness in Mild Primary Hyperparathyroidism. Journal of Clinical Endocrinology and Metabolism, 2005, 90, 3326-3330.	3.6	132
42	Cardiac Scintigraphy With Technetium-99m-Labeled Bone-Seeking Tracers for Suspected Amyloidosis. Journal of the American College of Cardiology, 2020, 75, 2851-2862.	2.8	131
43	Deprescribing in Older Adults With Cardiovascular Disease. Journal of the American College of Cardiology, 2019, 73, 2584-2595.	2.8	126
44	Development of Heart Failure in Chronic Hypertensive Dahl Rats. Hypertension, 2006, 47, 901-911.	2.7	125
45	Diflunisal for ATTR Cardiac Amyloidosis. Congestive Heart Failure, 2012, 18, 315-319.	2.0	124
46	Mechanism of Action and Clinical Application of Tafamidis in Hereditary Transthyretin Amyloidosis. Neurology and Therapy, 2016, 5, 1-25.	3.2	124
47	Effects of an Interatrial Shunt on Rest and Exercise Hemodynamics: Results of a Computer Simulation in Heart Failure. Journal of Cardiac Failure, 2014, 20, 212-221.	1.7	111
48	Tafamidis in Transthyretin Amyloid Cardiomyopathy. Circulation: Heart Failure, 2015, 8, 519-526.	3.9	110
49	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, 2019, 25, e1-e39.	1.7	107
50	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
51	What to Expect From the Evolving Field ofÂGeriatric Cardiology. Journal of the American College of Cardiology, 2015, 66, 1286-1299.	2.8	102
52	Polypharmacy in Older Adults Hospitalized for Heart Failure. Circulation: Heart Failure, 2020, 13, e006977.	3.9	102
53	ASNC/AHA/ASE/EANM/HFSA/ISA/SCMR/SNMMI expert consensus recommendations for multimodality imaging in cardiac amyloidosis: Part 2 of 2—Diagnostic criteria and appropriate utilization. Journal of Nuclear Cardiology, 2020, 27, 659-673.	2.1	97
54	Usefulness of Two-Dimensional Echocardiographic Parameters of the Left Side of the Heart to Predict Right Ventricular Failure After Left Ventricular Assist Device Implantation. American Journal of Cardiology, 2012, 109, 246-251.	1.6	96

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55	Perioperative Outcome and Longâ€Term Mortality for Heart Failure Patients Undergoing Intermediate― and Highâ€Risk Noncardiac Surgery: Impact of Left Ventricular Ejection Fraction. Congestive Heart Failure, 2010, 16, 45-49.	2.0	95
56	Nuclear imaging modalities for cardiac amyloidosis. Journal of Nuclear Cardiology, 2014, 21, 175-184.	2.1	93
57	Characterization of the <scp>inflammatoryâ€metabolic</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
58	Heart Failure with Preserved Ejection Fraction: Persistent Diagnosis, Therapeutic Enigma. Current Cardiovascular Risk Reports, 2011, 5, 440-449.	2.0	89
59	Metaâ€Analysis Global Group in Chronic (MAGGIC) Heart Failure Risk Score: Validation of a Simple Tool for the Prediction of Morbidity and Mortality in Heart Failure With Preserved Ejection Fraction. Journal of the American Heart Association, 2018, 7, e009594.	3.7	87
60	Pressure-Volume Relationships in Patients With Transthyretin (ATTR) Cardiac Amyloidosis Secondary to V122I Mutations and Wild-Type Transthyretin. Circulation: Heart Failure, 2011, 4, 121-128.	3.9	84
61	Serial scanning with technetium pyrophosphate (99mTc-PYP) in advanced ATTR cardiac amyloidosis. Journal of Nuclear Cardiology, 2016, 23, 1355-1363.	2.1	83
62	Cardiac Amyloidosis: Overlooked, Underappreciated, and Treatable. Annual Review of Medicine, 2020, 71, 203-219.	12.2	82
63	Home-Delivered Meals Postdischarge From Heart Failure Hospitalization. Circulation: Heart Failure, 2018, 11, e004886.	3.9	81
64	TTR (Transthyretin) Stabilizers Are Associated With Improved Survival in Patients With TTR Cardiac Amyloidosis. Circulation: Heart Failure, 2018, 11, e004769.	3.9	78
65	Association of Carpal Tunnel Syndrome With Amyloidosis, HeartÂFailure, and Adverse Cardiovascular Outcomes. Journal of the American College of Cardiology, 2019, 74, 15-23.	2.8	77
66	Myocardial contraction fraction: a volumetric index of myocardial shortening by freehand three-dimensional echocardiography. Journal of the American College of Cardiology, 2002, 40, 325-329.	2.8	73
67	Cardiac Transplantation Using Extended-Donor Criteria Organs for Systemic Amyloidosis Complicated by Heart Failure. Transplantation, 2007, 83, 539-545.	1.0	73
68	The feasibility of measuring frailty to predict disability and mortality in older medical intensive care unit survivors. Journal of Critical Care, 2014, 29, 401-408.	2.2	73
69	Unveiling outcomes in coexisting severe aortic stenosis and transthyretin cardiac amyloidosis. European Journal of Heart Failure, 2021, 23, 250-258.	7.1	71
70	Relation Between Six-Minute Walk Test Performance and Outcomes After Transcatheter Aortic Valve Implantation (from the PARTNER Trial). American Journal of Cardiology, 2013, 112, 700-706.	1.6	70
71	ASNC/AHA/ASE/EANM/HFSA/ISA/SCMR/SNMMI Expert Consensus Recommendations for Multimodality Imaging in Cardiac Amyloidosis: Part 2 of 2—Diagnostic Criteria and Appropriate Utilization. Journal of Cardiac Failure, 2019, 25, 854-865.	1.7	70
72	Rationale and Design of the Left Atrial Pressure Monitoring to Optimize Heart Failure Therapy Study (LAPTOP-HF). Journal of Cardiac Failure, 2015, 21, 479-488.	1.7	69

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73	The myocardial contraction fraction is superior to ejection fraction in predicting survival in patients with AL cardiac amyloidosis. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2015, 22, 61-66.	3.0	69
74	Comparison of Ventricular Structure and Function in Chinese Patients With Heart Failure and Ejection Fractions >55% Versus 40% to 55% Versus <40%. American Journal of Cardiology, 2009, 103, 845-851.	1.6	67
75	Left Atrial Decompression Pump forÂSevere Heart Failure With PreservedÂEjection Fraction. JACC: Heart Failure, 2015, 3, 275-282.	4.1	67
76	Comparison of Blood Volume Characteristics in Anemic Patients With Low Versus Preserved Left Ventricular Ejection Fractions. American Journal of Cardiology, 2008, 102, 1069-1072.	1.6	66
77	Gerotechnology for Older Adults With Cardiovascular Diseases. Journal of the American College of Cardiology, 2020, 76, 2650-2670.	2.8	66
78	Transthyretin Cardiac Amyloidoses in Older North Americans. Journal of the American Geriatrics Society, 2012, 60, 765-774.	2.6	64
79	Mechanisms Underlying Improvements in Ejection Fraction With Carvedilol in Heart Failure. Circulation: Heart Failure, 2009, 2, 189-196.	3.9	63
80	Clinical, ECG and echocardiographic clues to the diagnosis of TTR-related cardiomyopathy. Open Heart, 2016, 3, e000289.	2.3	62
81	Joint Commission Requirements for Discharge Instructions in Patients With Heart Failure: Is Understanding Important for Preventing Readmissions?. Journal of Cardiac Failure, 2014, 20, 641-649.	1.7	61
82	Gait Speed and Dependence in Activities of Daily Living in Older Adults With Severe Aortic Stenosis. Clinical Cardiology, 2012, 35, 307-314.	1.8	60
83	Avoiding misdiagnosis: expert consensus recommendations for the suspicion and diagnosis of transthyretin amyloidosis for the general practitioner. BMC Family Practice, 2020, 21, 198.	2.9	60
84	Design and Rationale of the Phase 3 ATTR-ACT Clinical Trial (Tafamidis in Transthyretin Cardiomyopathy) Tj ETQ	q0 Q Q rgB	T /Qyerlock 10
85	Prevalence and prognostic significance of exercise-induced supraventricular tachycardia in apparently healthy volunteers. American Journal of Cardiology, 1995, 75, 788-792.	1.6	58
86	Noninvasive Identification of ATTRwt Cardiac Amyloid: The Re-emergence of Nuclear Cardiology. American Journal of Medicine, 2015, 128, 1275-1280.	1.5	58
87	Can a Left Ventricular Assist Device in Individuals with Advanced Systolic Heart Failure Improve or Reverse Frailty?. Journal of the American Geriatrics Society, 2017, 65, 2383-2390.	2.6	58
88	Comparison of cardiac amyloidosis due to wild-type and V122I transthyretin in older adults referred to an academic medical center. Aging Health, 2013, 9, 229-235.	0.3	57
89	Stabilization of Cardiac Function With Diflunisal in Transthyretin (ATTR) Cardiac Amyloidosis. Journal of Cardiac Failure, 2020, 26, 753-759.	1.7	57
90	Cardiac Transplantation in Patients With Hypertrophic Cardiomyopathy. American Journal of Cardiology, 2012, 110, 568-574.	1.6	56

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91	A Prognostic Model for 6-Month Mortality in Elderly Survivors of Critical Illness. Chest, 2013, 143, 910-919.	0.8	56
92	Knowledge Gaps in Cardiovascular Care of Older Adults: A Scientific Statement from the American Heart Association, American College of Cardiology, and American Geriatrics Society: Executive Summary. Journal of the American Geriatrics Society, 2016, 64, 2185-2192.	2.6	56
93	Hip and knee arthroplasty are common among patients with transthyretin cardiac amyloidosis, occurring years before cardiac amyloid diagnosis: can we identify affected patients earlier?. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis. 2017. 24. 224-228.	3.0	56
94	ATTR Amyloidosis: Current and Emerging Management Strategies. JACC: CardioOncology, 2021, 3, 488-505.	4.0	56
95	Upright Posture and Postprandial Hypotension in Elderly Persons. Annals of Internal Medicine, 2000, 133, 533.	3.9	55
96	Interim 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, 2017, 24, 58-59.	3.0	55
97	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
98	Transthyretin Cardiac Amyloidosis in Black Americans. Circulation: Heart Failure, 2016, 9, e002558.	3.9	54
99	Transthyretin Cardiac Amyloidosis in Older Americans. Journal of Cardiac Failure, 2016, 22, 996-1003.	1.7	53
100	Amyloidosis cardiomyopathy. Current Opinion in Cardiology, 2018, 33, 571-579.	1.8	53
101	Diagnosing Transthyretin Cardiac Amyloidosis by Technetium Tc 99m Pyrophosphate. JACC: Cardiovascular Imaging, 2021, 14, 1221-1231.	5.3	52
102	Efficacy of Tafamidis in Patients With Hereditary and Wild-Type Transthyretin Amyloid Cardiomyopathy. JACC: Heart Failure, 2021, 9, 115-123.	4.1	52
103	Pathophysiology and Therapeutic Approaches to Cardiac Amyloidosis. Circulation Research, 2021, 128, 1554-1575.	4.5	52
104	Transcatheter Interatrial Shunt Device for the Treatment of Heart Failure. Circulation: Heart Failure, 2016, 9, .	3.9	51
105	Myocardial Contraction Fraction by M-Mode Echocardiography Is Superior to Ejection Fraction in Predicting Mortality in Transthyretin Amyloidosis. Journal of Cardiac Failure, 2018, 24, 504-511.	1.7	51
106	Independent Prognostic Value of Stroke Volume Index in Patients With Immunoglobulin Light Chain Amyloidosis. Circulation: Cardiovascular Imaging, 2018, 11, e006588.	2.6	51
107	Mechanisms of heart failure with well preserved ejection fraction in dogs following limited coronary microembolization. Cardiovascular Research, 2004, 64, 72-83.	3.8	50
108	Self-Reported Lack of Energy (Anergia) Among Elders in a Multiethnic Community. Journals of Gerontology - Series A Biological Sciences and Medical Sciences, 2008, 63, 707-714.	3.6	50

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109	The Transthyretin Amyloidosis Outcomes Survey (THAOS) registry: design and methodology. Current Medical Research and Opinion, 2013, 29, 77-84.	1.9	50
110	Impact of genotype and phenotype on cardiac biomarkers in patients with transthyretin amyloidosis – Report from the Transthyretin Amyloidosis Outcome Survey (THAOS). PLoS ONE, 2017, 12, e0173086.	2.5	50
111	Effects of age on outcome of tilt-table testing. American Journal of Cardiology, 1999, 83, 1055-1058.	1.6	48
112	Circulating Activated and Effector Memory T Cells Are Associated with Calcification and Clonal Expansions in Bicuspid and Tricuspid Valves of Calcific Aortic Stenosis. Journal of Immunology, 2011, 187, 1006-1014.	0.8	48
113	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. Circulation: Cardiovascular Imaging, 2021, 14, e000029.	2.6	48
114	Phase 1a/b study of monoclonal antibody CAEL-101 (11-1F4) in patients with AL amyloidosis. Blood, 2021, 138, 2632-2641.	1.4	48
115	Treating Anemia in Older Adults With Heart Failure With a Preserved Ejection Fraction With Epoetin Alfa. Circulation: Heart Failure, 2013, 6, 254-263.	3.9	47
116	Standardization of 99mTechnetium pyrophosphate imaging methodology to diagnose TTR cardiac amyloidosis. Journal of Nuclear Cardiology, 2018, 25, 181-190.	2.1	47
117	18Fluorine sodium fluoride positron emission tomography, a potential biomarker of transthyretin cardiac amyloidosis. Journal of Nuclear Cardiology, 2018, 25, 1559-1567.	2.1	46
118	Peptide probes detect misfolded transthyretin oligomers in plasma of hereditary amyloidosis patients. Science Translational Medicine, 2017, 9, .	12.4	44
119	The Lymphocytic Infiltration in Calcific Aortic Stenosis Predominantly Consists of Clonally Expanded T Cells. Journal of Immunology, 2007, 178, 5329-5339.	0.8	37
120	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
121	The Effect of Body Mass Index on Complications from Cardiac Surgery in the Oldest Old. Journal of the American Geriatrics Society, 2002, 50, 988-994.	2.6	36
122	Sex―and Raceâ€Related Differences in Characteristics and Outcomes of Hospitalizations for Heart Failure With Preserved Ejection Fraction. Journal of the American Heart Association, 2017, 6, .	3.7	36
123	Critical Comparison of Documents FromÂScientific Societies on CardiacÂAmyloidosis. Journal of the American College of Cardiology, 2022, 79, 1288-1303.	2.8	35
124	Personalized medicine approach for optimizing the dose of tafamidis to potentially ameliorate wild-type transthyretin amyloidosis (cardiomyopathy). Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2015, 22, 175-180.	3.0	34
125	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 Nuclear Cardiology, 2021, 28, 1769-1774.	2.1	34
126	The Columbia Cooperative Aging Program: An Interdisciplinary and Interdepartmental Approach to Geriatric Education for Medical Interns. Journal of the American Geriatrics Society, 2006, 54, 520-526.	2.6	33

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127	Unveiling wild-type transthyretin cardiac amyloidosis as a significant and potentially modifiable cause of heart failure with preserved ejection fraction. European Heart Journal, 2015, 36, 2595-2597.	2.2	33
128	Tafamidis—A Pricey Therapy for a Not-So-Rare Condition. JAMA Cardiology, 2020, 5, 247.	6.1	33
129	The Frailty Phenotype and Palliative Care Needs of Older Survivors of Critical Illness. Journal of the American Geriatrics Society, 2017, 65, 1168-1175.	2.6	31
130	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
131	Screening for ATTR amyloidosis in the clinic: overlapping disorders, misdiagnosis, and multiorgan awareness. Heart Failure Reviews, 2022, 27, 785-793.	3.9	31
132	The Prevalence and Impact of Anergia (Lack of Energy) in Subjects With Heart Failure and its Associations With Actigraphy. Journal of Cardiac Failure, 2009, 15, 145-151.	1.7	30
133	Gerontechnologies for Older Patients with Heart Failure: What is the Role of Smartphones, Tablets, and Remote Monitoring Devices in Improving Symptom Monitoring and Self-Care Management?. Current Cardiovascular Risk Reports, 2016, 10, 1.	2.0	30
134	How Should Physicians Assess Myocardial Contraction?. JACC: Cardiovascular Imaging, 2020, 13, 873-878.	5.3	30
135	Ventricular Pump Function in Heart Failure with Normal Ejection Fraction: Insights from Pressure-Volume Measurements. Progress in Cardiovascular Diseases, 2006, 49, 182-195.	3.1	28
136	Identifying Predictors of Taxane-Induced Peripheral Neuropathy Using Mass Spectrometry-Based Proteomics Technology. PLoS ONE, 2015, 10, e0145816.	2.5	28
137	Cardiac dysfunction in β-carotene-15,15′-dioxygenase-deficient mice is associated with altered retinoid and lipid metabolism. American Journal of Physiology - Heart and Circulatory Physiology, 2014, 307, H1675-H1684.	3.2	27
138	Noncompressibility of myocardium during systole with freehand three-dimensional echocardiography. Journal of the American Society of Echocardiography, 2002, 15, 1503-1506.	2.8	26
139	Wild-Type Transthyretin Cardiac Amyloidosis: Novel Insights From Advanced Imaging. Canadian Journal of Cardiology, 2016, 32, 1166.e1-1166.e10.	1.7	26
140	Comprehensive Proteomics Profiling Reveals Circulating Biomarkers of Hypertrophic Cardiomyopathy. Circulation: Heart Failure, 2021, 14, e007849.	3.9	26
141	Sex-Related Risk of Cardiac Involvement in Hereditary Transthyretin Amyloidosis. JACC: Heart Failure, 2021, 9, 736-746.	4.1	26
142	Clinical Outcomes After Left Ventricular Assist Device Implantation in Older Adults. JACC: Heart Failure, 2019, 7, 1069-1078.	4.1	25
143	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
144	Association Between Functional Impairment and Medication Burden in Adults with Heart Failure. Journal of the American Geriatrics Society, 2019, 67, 284-291.	2.6	24

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145	Systemic embolism in amyloid transthyretin cardiomyopathy. European Journal of Heart Failure, 2022, 24, 1387-1396.	7.1	23
146	Ventricular Volume and Length in Hypertensive Diastolic Heart Failure. Journal of the American Society of Echocardiography, 2005, 18, 1051-1057.	2.8	22
147	Mechanical Circulatory Support Device Utilization and Heart Transplant Waitlist Outcomes in Patients With Restrictive and Hypertrophic Cardiomyopathy. Circulation: Heart Failure, 2018, 11, e004665.	3.9	22
148	Remote Cardiac Monitoring in Patients With Heart Failure. JAMA Cardiology, 2022, 7, 556.	6.1	22
149	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
150	Myocardial Contraction Fraction. Journal of the American College of Cardiology, 2018, 71, 255-256.	2.8	21
151	Tafamidis for Transthyretin Amyloid Cardiomyopathy. New England Journal of Medicine, 2019, 380, 196-197.	27.0	21
152	Prescribing Patterns of HeartÂFailure-Exacerbating Medications Following a Heart Failure Hospitalization. JACC: Heart Failure, 2020, 8, 25-34.	4.1	21
153	Peripheral neuropathy symptoms in wild type transthyretin amyloidosis. Journal of the Peripheral Nervous System, 2020, 25, 265-272.	3.1	21
154	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
155	Improved Cardiovascular Disease Outcomes in Older Adults. F1000Research, 2016, 5, 112.	1.6	21
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