## Philip N Hawkins

# List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

10,300 40 101 101 h-index g-index citations papers 5.85 12,408 111 7.4 avg, IF L-index ext. citations ext. papers

#	Paper	IF	Citations
101	Cardiac Magnetic Resonance-Derived Extracellular Volume Mapping for the Quantification of Hepatic and Splenic Amyloid. <i>Circulation: Cardiovascular Imaging</i> , <b>2021</b> , CIRCIMAGING121012506	3.9	7
100	Search for AL amyloidosis risk factors using Mendelian randomization. <i>Blood Advances</i> , <b>2021</b> , 5, 2725-27	<b>′3⁄1</b> 8	3
99	99mTc-DPD scintigraphy in immunoglobulin light chain (AL) cardiac amyloidosis. <i>European Heart Journal Cardiovascular Imaging</i> , <b>2021</b> , 22, 1304-1311	4.1	6
98	Quantitative cardiovascular magnetic resonance myocardial perfusion mapping to assess hyperaemic response to adenosine stress. <i>European Heart Journal Cardiovascular Imaging</i> , <b>2021</b> , 22, 273	3- <del>2</del> 1	3
97	Longitudinal strain is an independent predictor of survival and response to therapy in patients with systemic AL amyloidosis. <i>European Heart Journal</i> , <b>2021</b> ,	9.5	9
96	Two types of systemic amyloidosis in a single patient. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , <b>2020</b> , 27, 275-276	2.7	3
95	Diagnostic imaging of cardiac amyloidosis. <i>Nature Reviews Cardiology</i> , <b>2020</b> , 17, 413-426	14.8	39
94	Echocardiographic phenotype and prognosis in transthyretin cardiac amyloidosis. <i>European Heart Journal</i> , <b>2020</b> , 41, 1439-1447	9.5	50
93	The Impact of Longitudinal Strain on Haematological and Cardiac Response and Survival in Patients with Systemic AL Amyloidosis. <i>Blood</i> , <b>2020</b> , 136, 40-40	2.2	1
92	TRAP1 chaperone protein mutations and autoinflammation. Life Science Alliance, 2020, 3,	5.8	7
91	Assessment of Multivessel Coronary Artery Disease Using Cardiovascular Magnetic Resonance Pixelwise Quantitative Perfusion Mapping. <i>JACC: Cardiovascular Imaging</i> , <b>2020</b> , 13, 2546-2557	8.4	7
90	Eight novel loci implicate shared genetic etiology in multiple myeloma, AL amyloidosis, and monoclonal gammopathy of unknown significance. <i>Leukemia</i> , <b>2020</b> , 34, 1187-1191	10.7	12
89	Noninvasive Mapping of the Electrophysiological Substrate in Cardiac Amyloidosis and Its Relationship to Structural Abnormalities. <i>Journal of the American Heart Association</i> , <b>2019</b> , 8, e012097	6	14
88	A prospective observational study of 915 patients with systemic AL amyloidosis treated with upfront bortezomib. <i>Blood</i> , <b>2019</b> , 134, 2271-2280	2.2	77
87	The UK National Amyloidosis Centre. <i>European Heart Journal</i> , <b>2019</b> , 40, 1661-1664	9.5	1
86	Acute changes in cardiac structural and tissue characterisation parameters following haemodialysis measured using cardiovascular magnetic resonance. <i>Scientific Reports</i> , <b>2019</b> , 9, 1388	4.9	15
85	Recognising and understanding cryopyrin-associated periodic syndrome in adults. <i>British Journal of Nursing</i> , <b>2019</b> , 28, 1180-1186	0.7	3

### (2016-2019)

84	Coronary Physiology. <i>JACC: Cardiovascular Imaging</i> , <b>2019</b> , 12, 1958-1969	8.4	66
83	Native T1 and Extracellular Volume in Transthyretin Amyloidosis. <i>JACC: Cardiovascular Imaging</i> , <b>2019</b> , 12, 810-819	8.4	89
82	Cardiac Structural and Functional Consequences of Amyloid Deposition by Cardiac Magnetic Resonance and Echocardiography and Their Prognostic Roles. <i>JACC: Cardiovascular Imaging</i> , <b>2019</b> , 12, 823-833	8.4	55
81	Role of Tc-DPD scintigraphy in imaging extra-cardiac light chain (AL) amyloidosis. <i>British Journal of Haematology</i> , <b>2018</b> , 183, 506-509	4.5	3
8o	Systemic immunoglobulin light chain amyloidosis. <i>Nature Reviews Disease Primers</i> , <b>2018</b> , 4, 38	51.1	207
79	Autosomal dominant familial Mediterranean fever in Northern European Caucasians associated with deletion of p.M694 residue-a case series and genetic exploration. <i>Rheumatology</i> , <b>2017</b> , 56, 209-213	3.9	38
78	High prevalence of recurrent nocturnal desaturations in systemic AL amyloidosis: a cross-sectional pilot study. <i>Sleep Medicine</i> , <b>2017</b> , 32, 191-197	4.6	3
77	Six-minute walk test (6MWT) in AL amyloidosis - baseline and 12-month follow-up after chemotherapy. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis,</i> <b>2017</b> , 24, 62-63	2.7	3
76	A good clonal response to chemotherapy in AL amyloidosis is associated with improved quality of life and function at 1 year. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , <b>2017</b> , 24, 72-73	2.7	1
75	Hyperimmunoglobulin D syndrome in an Indian family undiagnosed for 11 years. <i>International Journal of Rheumatic Diseases</i> , <b>2017</b> , 20, 2236-2237	2.3	1
74	International multi-centre study of pregnancy outcomes with interleukin-1 inhibitors. <i>Rheumatology</i> , <b>2017</b> , 56, 2102-2108	3.9	54
73	Magnetic Resonance in Transthyretin Cardiac Amyloidosis. <i>Journal of the American College of Cardiology</i> , <b>2017</b> , 70, 466-477	15.1	176
72	001 Multiparametric mapping to understand pathophysiology in cardiac amyloidosis. <i>Heart</i> , <b>2017</b> , 103, A1-A2	5.1	11
71	Genome-wide association study of clinical parameters in immunoglobulin light chain amyloidosis in three patient cohorts. <i>Haematologica</i> , <b>2017</b> , 102, e411-e414	6.6	7
70	European Collaborative Study Defining Clinical Profile Outcomes and Novel Prognostic Criteria in Monoclonal Immunoglobulin M-Related Light Chain Amyloidosis. <i>Journal of Clinical Oncology</i> , <b>2016</b> , 34, 2037-45	2.2	46
69	EULAR recommendations for the management of familial Mediterranean fever. <i>Annals of the Rheumatic Diseases</i> , <b>2016</b> , 75, 644-51	2.4	241
68	Systemic amyloidosis. <i>Lancet, The</i> , <b>2016</b> , 387, 2641-2654	40	475
67	Comparison of Free Light Chain Assays: Freelite and N Latex in Diagnosis, Monitoring, and Predicting Survival in Light Chain Amyloidosis. <i>American Journal of Clinical Pathology</i> , <b>2016</b> , 146, 78-85	1.9	7

66	Nonbiopsy Diagnosis of Cardiac Transthyretin Amyloidosis. <i>Circulation</i> , <b>2016</b> , 133, 2404-12	16.7	792
65	Therapeutic Clearance of Amyloid by Antibodies to Serum Amyloid P Component. <i>New England Journal of Medicine</i> , <b>2015</b> , 373, 1106-14	59.2	250
64	Natural history and outcomes in localised immunoglobulin light-chain amyloidosis: a long-term observational study. <i>Lancet Haematology,the</i> , <b>2015</b> , 2, e241-50	14.6	76
63	Prognostic Value of Late Gadolinium Enhancement Cardiovascular Magnetic Resonance in Cardiac Amyloidosis. <i>Circulation</i> , <b>2015</b> , 132, 1570-9	16.7	320
62	Emerging treatments for amyloidosis. <i>Kidney International</i> , <b>2015</b> , 87, 516-26	9.9	20
61	A European collaborative study of cyclophosphamide, bortezomib, and dexamethasone in upfront treatment of systemic AL amyloidosis. <i>Blood</i> , <b>2015</b> , 126, 612-5	2.2	232
60	T1 mapping and survival in systemic light-chain amyloidosis. European Heart Journal, 2015, 36, 244-51	9.5	247
59	CMR-based differentiation of AL and ATTR cardiac amyloidosis. <i>JACC: Cardiovascular Imaging</i> , <b>2014</b> , 7, 133-42	8.4	182
58	Stringent patient selection improves outcomes in systemic light-chain amyloidosis after autologous stem cell transplantation in the upfront and relapsed setting. <i>Haematologica</i> , <b>2014</b> , 99, e260-3	6.6	11
57	Two types of amyloid in a single heart. <i>Blood</i> , <b>2014</b> , 124, 3025-7	2.2	20
56	Efficacy of bortezomib, cyclophosphamide and dexamethasone in treatment-nalle patients with high-risk cardiac AL amyloidosis (Mayo Clinic stage III). <i>Haematologica</i> , <b>2014</b> , 99, 1479-85	6.6	83
55	Utility and limitations of 3,3-diphosphono-1,2-propanodicarboxylic acid scintigraphy in systemic amyloidosis. <i>European Heart Journal Cardiovascular Imaging</i> , <b>2014</b> , 15, 1289-98	4.1	131
54	Online registry for mutations in hereditary amyloidosis including nomenclature recommendations. <i>Human Mutation</i> , <b>2014</b> , 35, E2403-12	4.7	162
53	Native T1 mapping in transthyretin amyloidosis. <i>JACC: Cardiovascular Imaging</i> , <b>2014</b> , 7, 157-65	8.4	265
52	The Prognostic Significance of Phenotypically NormallPlasma Cells in Chemotherapy Treated AL Patients with Underlying MGUS and Multiple Myeloma. <i>Blood</i> , <b>2014</b> , 124, 2073-2073	2.2	1
51	Acute hepatitis in a child heterozygous for the I259V MEFV gene variant. <i>Prague Medical Report</i> , <b>2014</b> , 115, 128-33	0.7	3
50	A European Collaborative Study of 230 Patients to Assess the Role of Cyclophosphamide, Bortezomib and Dexamethasone in Upfront Treatment of Patients with Systemic AL Amyloidosis. <i>Blood</i> , <b>2014</b> , 124, 305-305	2.2	
49	Longterm Outcomes and Improved Renal Function with Autologous Stem Cell Transplantation (ASCT) in Light Chain Deposition Disease (LCDD). <i>Blood</i> , <b>2014</b> , 124, 1198-1198	2.2	

### (2009-2013)

48	Measurement of Tissue interstitial volume in healthy patients and those with amyloidosis with equilibrium contrast-enhanced MR imaging. <i>Radiology</i> , <b>2013</b> , 268, 858-64	20.5	40
47	A European collaborative study of treatment outcomes in 346 patients with cardiac stage III AL amyloidosis. <i>Blood</i> , <b>2013</b> , 121, 3420-7	2.2	267
46	New criteria for response to treatment in immunoglobulin light chain amyloidosis based on free light chain measurement and cardiac biomarkers: impact on survival outcomes. <i>Journal of Clinical Oncology</i> , <b>2012</b> , 30, 4541-9	2.2	553
45	A Matched Comparison of Cyclophosphamide, Bortezomib and Dexamethasone (CVD) Versus Cyclophosphamide, Thalidomide and Dexamethasone (CTD) in the Treatment of Mayo Cardiac Stage III Patients with AL Amyloidosis <i>Blood</i> , <b>2012</b> , 120, 2966-2966	2.2	1
44	Continuous Therapy with Lenalidomide Correlates with Improved Progression Free Survival in Heavily Pre-Treated Patients with AL Amyloidosis <i>Blood</i> , <b>2012</b> , 120, 2978-2978	2.2	
43	Treatment and Outcome of 267 Patients with IgM-Related AL Amyloidosis. <i>Blood</i> , <b>2012</b> , 120, 4074-4074	2.2	1
42	A case series and systematic literature review of anakinra and immunosuppression in idiopathic recurrent pericarditis. <i>Journal of Cardiology Cases</i> , <b>2011</b> , 4, e93-e97	0.6	24
41	ALchemy - A Large Prospective <b>R</b> eal World' Study of Chemotherapy in AL Amyloidosis. <i>Blood</i> , <b>2011</b> , 118, 992-992	2.2	4
40	European Collaborative Study of Treatment Outcomes in 347 Patients with Systemic AL Amyloidosis with Mayo Stage III Disease. <i>Blood</i> , <b>2011</b> , 118, 995-995	2.2	3
39	Complete and Very Good Partial Responses Are Attainable Endpoints in Elderly Patients (>75 years) with AL Amyloidosis and Are Associated with Improved Overall Survival,. <i>Blood</i> , <b>2011</b> , 118, 3975-3975	2.2	1
38	Cyclophosphamide, Bortezomib and Dexamethasone (CVD) Therapy in AL Amyloidosis Is Associated with High Clonal Response Rates and Prolonged Progression Free Survival,. <i>Blood</i> , <b>2011</b> , 118, 3978-3978	2.2	О
37	Validation of the Criteria of Response to Treatment In AL Amyloidosis <i>Blood</i> , <b>2010</b> , 116, 1364-1364	2.2	15
36	A European Collaborative Study of Treatment Outcomes In 428 Patients with Systemic AL Amyloidosis. <i>Blood</i> , <b>2010</b> , 116, 988-988	2.2	4
35	Remarkable Efficacy of IL-1 Receptor Antagonist In Schnitzler's Syndrome: a Series of 6 Cases. <i>Blood</i> , <b>2010</b> , 116, 3958-3958	2.2	
34	Transient Post Chemotherapy Rise in NT Pro-BNP in AL Amyloidosis : Implications for Organ Response Assessment <i>Blood</i> , <b>2009</b> , 114, 1791-1791	2.2	3
33	A New Staging System for AL Amyloidosis Incorporating Serum Free Light Chains, cardiac Troponin-T and NT-ProBNP <i>Blood</i> , <b>2009</b> , 114, 2796-2796	2.2	6
32	Cyclophosphamide, Thalidomide and Dexamethasone (CTD) Versus Melphalan Plus Dexamethasone (MD) for Newly-Diagnosed Systemic AL Amyloidosis lResults From the UK Amyloidosis Treatment Trial <i>Blood</i> , <b>2009</b> , 114, 2869-2869	2.2	10
31	In AL Amyloidosis, Both Oral Melphalan and Dexamethasone (Mel-Dex) and Risk-Adapted Cyclophosphamide, Thalidomide and Dexamethasone (CTD) Have Similar Efficacy as Upfront Treatment <i>Blood</i> , <b>2009</b> , 114, 745-745	2.2	6

30	Early Detection of Cardiac Systolic Functional Impairment and Correlation with NT-ProBNP Change in AL Amyloidosis by Cardiac Lateral Wall Tissue Doppler S Wave <i>Blood</i> , <b>2009</b> , 114, 2814-2814	2.2	
29	Is There a Role for Thalidomide Maintenance in the Treatment of AL Amyloidosis? <i>Blood</i> , <b>2009</b> , 114, 1863-1863	2.2	
28	Role of NT-ProBNP to Assess the Adequacy of Treatment Response in AL Amyloidosis <i>Blood</i> , <b>2008</b> , 112, 1689-1689	2.2	3
27	Risk-Adapted Cyclophosphamide, Thalidomide and Dexamethasone (CTD) for the Treatment of Systemic AL Amyloidosis: Long Term Outcomes among 202 Patients <i>Blood</i> , <b>2008</b> , 112, 1733-1733	2.2	1
26	Significant Activity of Bortezomib-Based Therapy in Patients with Primary Systemic (AL) Amyloidosis. <i>Blood</i> , <b>2008</b> , 112, 869-869	2.2	4
25	Diagnostic performance of 123I-labeled serum amyloid P component scintigraphy in patients with amyloidosis. <i>American Journal of Medicine</i> , <b>2006</b> , 119, 355.e15-24	2.4	101
24	Systemic amyloidosis. <i>Current Opinion in Pharmacology</i> , <b>2006</b> , 6, 214-20	5.1	73
23	Outcome of autologous stem cell transplantation for AL amyloidosis in the UK. <i>British Journal of Haematology</i> , <b>2006</b> , 134, 417-25	4.5	72
22	Efficacy and Safety of Bortezomib in Systemic AL Amyloidosis - A Preliminary Report <i>Blood</i> , <b>2006</b> , 108, 129-129	2.2	8
21	Superior mesenteric artery syndrome in a young girl following spinal surgery for scoliosis. <i>British Journal of Hospital Medicine (London, England: 2005)</i> , <b>2005</b> , 66, 700-1	0.8	
20	Definition of organ involvement and treatment response in immunoglobulin light chain amyloidosis (AL): a consensus opinion from the 10th International Symposium on Amyloid and Amyloidosis, Tours, France, 18-22 April 2004. <i>American Journal of Hematology</i> , <b>2005</b> , 79, 319-28	7.1	971
19	Standard Oral Melphalan Chemotherapy for AL Amyloidosis Revisited Using the Serum Free Light Chain Assay <i>Blood</i> , <b>2005</b> , 106, 3495-3495	2.2	2
18	Spectrum of clinical features in Muckle-Wells syndrome and response to anakinra. <i>Arthritis and Rheumatism</i> , <b>2004</b> , 50, 607-12		650
17	Intermediate Dose Intravenous Melphalan and Dexamethasone Treatment in 144 Patients with Systemic AL Amyloidosis <i>Blood</i> , <b>2004</b> , 104, 755-755	2.2	5
16	Impact of Chromosomal Abnormalities Revealed by Interphase FISH on Survival in Primary Light Chain Amyloidosis <i>Blood</i> , <b>2004</b> , 104, 4875-4875	2.2	
15	Hereditary systemic amyloidosis with renal involvement. <i>Journal of Nephrology</i> , <b>2003</b> , 16, 443-8	4.8	29
14	Misdiagnosis of hereditary amyloidosis as AL (primary) amyloidosis. <i>New England Journal of Medicine</i> , <b>2002</b> , 346, 1786-91	59.2	516
13	Serum amyloid P component scintigraphy for diagnosis and monitoring amyloidosis. <i>Current Opinion in Nephrology and Hypertension</i> , <b>2002</b> , 11, 649-55	3.5	108

#### LIST OF PUBLICATIONS

12	to distal chromosome 1q. <i>Arthritis and Rheumatism</i> , <b>2000</b> , 43, 2034-40		37
11	Bleeding symptoms and coagulation abnormalities in 337 patients with AL-amyloidosis. <i>British Journal of Haematology</i> , <b>2000</b> , 110, 454-60	4.5	155
10	Serum amyloid P component scintigraphy in familial amyloid polyneuropathy: regression of visceral amyloid following liver transplantation. <i>European Journal of Nuclear Medicine and Molecular Imaging</i> , <b>1998</b> , 25, 709-13	8.8	57
9	Allogeneic bone marrow transplantation for systemic AL amyloidosis. <i>British Journal of Haematology</i> , <b>1998</b> , 100, 226-8	4.5	45
8	SAA1 alleles as risk factors in reactive systemic AA amyloidosis. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , <b>1998</b> , 5, 262-5	2.7	84
7	Instability, unfolding and aggregation of human lysozyme variants underlying amyloid fibrillogenesis. <i>Nature</i> , <b>1997</b> , 385, 787-93	50.4	980
6	Amyloidosis: a review of recent diagnostic and therapeutic developments. <i>British Journal of Haematology</i> , <b>1997</b> , 99, 245-56	4.5	143
5	Apolipoprotein E4 genotype is not a risk factor for systemic AA amyloidosis or familial amyloid polyneuropathy. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , <b>1995</b> , 2, 163-166	2.7	7
4	Studies with radiolabelled serum amyloid P component provide evidence for turnover and regression of amyloid deposits in vivo. <i>Clinical Science</i> , <b>1994</b> , 87, 289-95	6.5	70
3	Biochemical effect of liver transplantation in two Swedish patients with familial amyloidotic polyneuropathy (FAP-met30). <i>Clinical Genetics</i> , <b>1991</b> , 40, 242-6	4	278
2	Evaluation of systemic amyloidosis by scintigraphy with 123I-labeled serum amyloid P component. <i>New England Journal of Medicine</i> , <b>1990</b> , 323, 508-13	59.2	422
1	Systemic Amyloidosis886-892		