List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	New Criteria for Response to Treatment in Immunoglobulin Light Chain Amyloidosis Based on Free Light Chain Measurement and Cardiac Biomarkers: Impact on Survival Outcomes. Journal of Clinical Oncology, 2012, 30, 4541-4549.	1.6	735
2	A European collaborative study of treatment outcomes in 346 patients with cardiac stage III AL amyloidosis. Blood, 2013, 121, 3420-3427.	1.4	385
3	A staging system for renal outcome and early markers of renal response to chemotherapy in AL amyloidosis. Blood, 2014, 124, 2325-2332.	1.4	366
4	Systemic immunoglobulin light chain amyloidosis. Nature Reviews Disease Primers, 2018, 4, 38.	30.5	350
5	Daratumumab-Based Treatment for Immunoglobulin Light-Chain Amyloidosis. New England Journal of Medicine, 2021, 385, 46-58.	27.0	268
6	Longitudinal Left Ventricular Function for Prediction of Survival in Systemic Light-Chain Amyloidosis. Journal of the American College of Cardiology, 2012, 60, 1067-1076.	2.8	253
7	Prophylactic implantation of cardioverter-defibrillator in patients with severe cardiac amyloidosis and high risk for sudden cardiac death. Heart Rhythm, 2008, 5, 235-240.	0.7	214
8	Immunohistochemistry in the classification of systemic forms of amyloidosis: a systematic investigation of 117 patients. Blood, 2012, 119, 488-493.	1.4	200
9	Premature telomeric loss in rheumatoid arthritis is genetically determined and involves both myeloid and lymphoid cell lineages. Proceedings of the National Academy of Sciences of the United States of America, 2003, 100, 13471-13476.	7.1	185
10	Translocation t(11;14) Is Associated With Adverse Outcome in Patients With Newly Diagnosed AL Amyloidosis When Treated With Bortezomib-Based Regimens. Journal of Clinical Oncology, 2015, 33, 1371-1378.	1.6	185
11	First-in-Human Phase I/II Study of NEOD001 in Patients With Light Chain Amyloidosis and Persistent Organ Dysfunction. Journal of Clinical Oncology, 2016, 34, 1097-1103.	1.6	176
12	Autologous/reduced-intensity allogeneic stem cell transplantation vs autologous transplantation in multiple myeloma: long-term results of the EBMT-NMAM2000 study. Blood, 2013, 121, 5055-5063.	1.4	171
13	Homeostatic control of T-cell generation in neonates. Blood, 2003, 102, 1428-1434.	1.4	158
14	Cryo-EM structure of a light chain-derived amyloid fibril from a patient with systemic AL amyloidosis. Nature Communications, 2019, 10, 1103.	12.8	120
15	Hereditary Apolipoprotein Al-Associated Amyloidosis in Surgical Pathology Specimens. Journal of Molecular Diagnostics, 2009, 11, 257-262.	2.8	116
16	In vivo detection of nerve injury in familial amyloid polyneuropathy by magnetic resonance neurography. Brain, 2015, 138, 549-562.	7.6	112
17	Non-invasive predictors of survival in cardiac amyloidosis. European Journal of Heart Failure, 2007, 9, 617-624.	7.1	109
18	Assessment of disease severity and outcome in patients with systemic light-chain amyloidosis by the high-sensitivity troponin T assay. Blood, 2010, 116, 2455-2461.	1.4	109

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19	Polymorphism of Amyloid Fibrils In Vivo. Angewandte Chemie - International Edition, 2016, 55, 4822-4825.	13.8	109
20	A phase 1/2 study of the oral proteasome inhibitor ixazomib in relapsed or refractory AL amyloidosis. Blood, 2017, 130, 597-605.	1.4	108
21	Defining the pathogenic role of telomerase mutations in myelodysplastic syndrome and acute myeloid leukemia. Human Mutation, 2009, 30, 1567-1573.	2.5	107
22	AL amyloidosis patients with low amyloidogenic free light chain levels at first diagnosis have an excellent prognosis. Blood, 2017, 130, 632-642.	1.4	104
23	Evaluation of the cytogenetic aberration pattern in amyloid light chain amyloidosis as compared with monoclonal gammopathy of undetermined significance reveals common pathways of karyotypic instability. Blood, 2008, 111, 4700-4705.	1.4	103
24	Bortezomib, Melphalan, and Dexamethasone for Light-Chain Amyloidosis. Journal of Clinical Oncology, 2020, 38, 3252-3260.	1.6	102
25	Amyloid in biopsies of the gastrointestinal tract—a retrospective observational study on 542 patients. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2016, 468, 569-577.	2.8	93
26	Carpal tunnel syndrome and spinal canal stenosis: harbingers of transthyretin amyloid cardiomyopathy?. Clinical Research in Cardiology, 2019, 108, 1324-1330.	3.3	93
27	Treatment with intravenous melphalan and dexamethasone is not able to overcome the poor prognosis of patients with newly diagnosed systemic light chain amyloidosis and severe cardiac involvement. Blood, 2010, 116, 522-528.	1.4	84
28	Gain of chromosome 1q21 is an independent adverse prognostic factor in light chain amyloidosis patients treated with melphalan/dexamethasone. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2014, 21, 9-17.	3.0	84
29	Postallograft lenalidomide induces strong NK cell–mediated antimyeloma activity and risk for T cell–mediated GvHD: Results from a phase I/II dose-finding study. Experimental Hematology, 2013, 41, 134-142.e3.	0.4	83
30	Melphalan 140 mg/m ² or 200 mg/m ² for autologous transplantation in myeloma: results from the Collaboration to Collect Autologous Transplant Outcomes in Lymphoma and Myeloma (CALM) study. A report by the EBMT Chronic Malignancies Working Party. Haematologica, 2018, 103, 514-521.	3.5	70
31	Cryo-EM reveals structural breaks in a patient-derived amyloid fibril from systemic AL amyloidosis. Nature Communications, 2021, 12, 875.	12.8	70
32	Prognostic impact of cytogenetic aberrations in AL amyloidosis patients after high-dose melphalan: a long-term follow-up study. Blood, 2016, 128, 594-602.	1.4	67
33	Daratumumab for systemic AL amyloidosis: prognostic factors and adverse outcome with nephrotic-range albuminuria. Blood, 2020, 135, 1517-1530.	1.4	67
34	Improved outcomes after heart transplantation for cardiac amyloidosis in the modern era. Journal of Heart and Lung Transplantation, 2018, 37, 611-618.	0.6	66
35	High-dose melphalan with autologous stem cell transplantation after VAD induction chemotherapy for treatment of amyloid light chain amyloidosis: a single centre prospective phase II study. British Journal of Haematology, 2004, 127, 543-551.	2.5	62
36	Allogeneic and syngeneic hematopoietic cell transplantation in patients with amyloid light-chain amyloidosis: a report from the European Group for Blood and Marrow Transplantation. Blood, 2006, 107, 2578-2584.	1.4	62

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37	Evaluation of the serum-free light chain test in untreated patients with AL amyloidosis. Haematologica, 2008, 93, 459-462.	3.5	62
38	Sjögren's syndrome and localized nodular cutaneous amyloidosis: Coincidence or a distinct clinical entity?. Arthritis and Rheumatism, 2008, 58, 1992-1999.	6.7	61
39	Hyperdiploidy is less frequent in AL amyloidosis compared with monoclonal gammopathy of undetermined significance and inversely associated with translocation t(11;14). Blood, 2011, 117, 3809-3815.	1.4	60
40	Common Fibril Structures Imply Systemically Conserved Protein Misfolding Pathways Inâ€Vivo. Angewandte Chemie - International Edition, 2017, 56, 7510-7514.	13.8	59
41	Venetoclax induces deep hematologic remissions in t(11;14) relapsed/refractory AL amyloidosis. Blood Cancer Journal, 2021, 11, 10.	6.2	53
42	Rapid Progression of Left Ventricular Wall Thickness Predicts Mortality in Cardiac Light-chain Amyloidosis. Journal of Heart and Lung Transplantation, 2007, 26, 1313-1319.	0.6	52
43	Tandem Autologous Stem Cell Transplantation Improves Outcomes in Newly Diagnosed Multiple Myeloma with Extramedullary Disease and High-Risk Cytogenetics: A Study from the Chronic Malignancies Working Party of the European Society for Blood and Marrow Transplantation. Biology of Blood and Marrow Transplantation, 2019, 25, 2134-2142.	2.0	52
44	Amyloid in endomyocardial biopsies. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2010, 456, 523-532.	2.8	50
45	Clarification on the definition of complete haematologic response in light-chain (AL) amyloidosis. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2021, 28, 1-2.	3.0	49
46	Sural nerve injury in familial amyloid polyneuropathy. Neurology, 2017, 89, 475-484.	1.1	48
47	Staged heart transplantation and chemotherapy as a treatment option in patients with severe cardiac lightâ€chain amyloidosis. European Journal of Heart Failure, 2009, 11, 1014-1020.	7.1	45
48	Extramedullary relapses after allogeneic non-myeloablative stem cell transplantation in multiple myeloma patients do not negatively affect treatment outcome. Bone Marrow Transplantation, 2008, 41, 779-784.	2.4	44
49	Rapid assessment of longitudinal systolic left ventricular function using speckle tracking of the mitral annulus. Clinical Research in Cardiology, 2012, 101, 273-280.	3.3	43
50	Prevalence of Germline Mutations in the TTR Gene in a Consecutive Series of Surgical Pathology Specimens With ATTR Amyloid. American Journal of Surgical Pathology, 2009, 33, 58-65.	3.7	39
51	Lenalidomide/melphalan/dexamethasone in newly diagnosed patients with immunoglobulin light chain amyloidosis: results of a prospective phase 2 study with long-term follow up. Haematologica, 2017, 102, 1424-1431.	3.5	39
52	Current status of hematopoietic cell transplantation in the treatment of systemic amyloid light-chain amyloidosis. Bone Marrow Transplantation, 2012, 47, 895-905.	2.4	37
53	Skeletal scintigraphy indicates disease severity of cardiac involvement in patients with senile systemic amyloidosis. International Journal of Cardiology, 2013, 164, 179-184.	1.7	37
54	Aggregation of Full-length Immunoglobulin Light Chains from Systemic Light Chain Amyloidosis (AL) Patients Is Remodeled by Epigallocatechin-3-gallate. Journal of Biological Chemistry, 2017, 292, 2328-2344.	3.4	37

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55	A Randomized Phase III Trial of Melphalan and Dexamethasone (MDex) Versus Bortezomib, Melphalan and Dexamethasone (BMDex) for Untreated Patients with AL Amyloidosis. Blood, 2016, 128, 646-646.	1.4	37
56	Role of mutations and post-translational modifications in systemic AL amyloidosis studied by cryo-EM. Nature Communications, 2021, 12, 6434.	12.8	36
57	The impact of allogeneic stem cell transplantation on the natural course of poor-risk chronic lymphocytic leukemia as defined by the EBMT consensus criteria: a retrospective donor versus no donor comparison. Annals of Oncology, 2014, 25, 200-206.	1.2	35
58	Peripheral blood or bone marrow cells in reduced-intensity or myeloablative conditioning allogeneic HLA identical sibling donor transplantation for multiple myeloma. Haematologica, 2007, 92, 1513-1518.	3.5	34
59	Prognosis and Staging of AL Amyloidosis. Acta Haematologica, 2020, 143, 388-400.	1.4	34
60	Primary Results from the Phase 3 Tourmaline-AL1 Trial of Ixazomib-Dexamethasone Versus Physician's Choice of Therapy in Patients (Pts) with Relapsed/Refractory Primary Systemic AL Amyloidosis (RRAL). Blood, 2019, 134, 139-139.	1.4	34
61	Allogeneic hematopoietic stem cell transplantation for poor-risk CLL: dissecting immune-modulating strategies for disease eradication and treatment of relapse. Bone Marrow Transplantation, 2015, 50, 1279-1285.	2.4	33
62	Cytogenetic intraclonal heterogeneity of plasma cell dyscrasia in AL amyloidosis as compared with multiple myeloma. Blood Advances, 2018, 2, 2607-2618.	5.2	33
63	Fatal amyloid formation in a patient's antibody light chain is caused by a single point mutation. ELife, 2020, 9, .	6.0	33
64	Mutational landscape reflects the biological continuum of plasma cell dyscrasias. Blood Cancer Journal, 2017, 7, e537-e537.	6.2	32
65	Myocardial contraction fraction derived from cardiovascular magnetic resonance cine images—reference values and performance in patients with heart failure and left ventricular hypertrophy. European Heart Journal Cardiovascular Imaging, 2017, 18, 1414-1422.	1.2	32
66	Genome-wide association study of immunoglobulin light chain amyloidosis in three patient cohorts: comparison with myeloma. Leukemia, 2017, 31, 1735-1742.	7.2	32
67	CD38 as Immunotherapeutic Target in Light Chain Amyloidosis and Multiple Myeloma—Association With Molecular Entities, Risk, Survival, and Mechanisms of Upfront Resistance. Frontiers in Immunology, 2018, 9, 1676.	4.8	32
68	The impact of stem cell transplantation on the natural course of peripheral T-cell lymphoma: a real-world experience. Annals of Hematology, 2018, 97, 1241-1250.	1.8	31
69	Local vs. systemic pulmonary amyloidosis—impact on diagnostics and clinical management. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2018, 473, 627-637.	2.8	31
70	Long-term survival of 1338 MM patients treated with tandem autologous vs. autologous-allogeneic transplantation. Bone Marrow Transplantation, 2020, 55, 1810-1816.	2.4	31
71	First report of ibrutinib in IgM-related amyloidosis: few responses, poor tolerability, and short survival. Blood, 2018, 131, 368-371.	1.4	30
72	Treatment of AL amyloidosis with bendamustine: a study of 122 patients. Blood, 2018, 132, 1988-1991.	1.4	30

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73	Performance analysis of AL amyloidosis cardiac biomarker staging systems with special focus on renal failure and atrial arrhythmia. Haematologica, 2019, 104, 1451-1459.	3.5	29
74	Letermovir prophylaxis is effective in preventing cytomegalovirus reactivation after allogeneic hematopoietic cell transplantation: single-center real-world data. Annals of Hematology, 2021, 100, 2087-2093.	1.8	29
75	A randomized phase 3 study of ixazomib–dexamethasone versus physician's choice in relapsed or refractory AL amyloidosis. Leukemia, 2022, 36, 225-235.	7.2	29
76	Deferred autologous stem cell transplantation in systemic AL amyloidosis. Blood Cancer Journal, 2018, 8, 101.	6.2	28
77	Bortezomib-based induction followed by stem cell transplantation in light chain amyloidosis: results of the multicenter HOVON 104 trial. Haematologica, 2019, 104, 2274-2282.	3.5	27
78	Results of the Phase 3 VITAL Study of NEOD001 (Birtamimab) Plus Standard of Care in Patients with Light Chain (AL) Amyloidosis Suggest Survival Benefit for Mayo Stage IV Patients. Blood, 2019, 134, 3166-3166.	1.4	27
79	Centre characteristics and procedureâ€related factors have an impact on outcomes of allogeneic transplantation for patients with <scp>CLL</scp> : a retrospective analysis from the European Society for Blood and Marrow Transplantation (<scp>EBMT</scp>). British Journal of Haematology, 2017, 178, 521-533.	2.5	26
80	Flow cytometryâ€based characterization of underlying clonal B and plasma cells in patients with light chain amyloidosis. Cancer Medicine, 2016, 5, 1464-1472.	2.8	25
81	Prognostic value of novel imaging parameters derived from standard cardiovascular magnetic resonance in high risk patients with systemic light chain amyloidosis. Journal of Cardiovascular Magnetic Resonance, 2019, 21, 53.	3.3	25
82	Localized immunoglobulin light chain amyloidosis: Novel insights including prognostic factors for local progression. American Journal of Hematology, 2020, 95, 1158-1169.	4.1	25
83	Protease resistance of <i>ex vivo</i> amyloid fibrils implies the proteolytic selection of disease-associated fibril morphologies. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2021, 28, 243-251.	3.0	25
84	Risk factors for AA amyloidosis in Germany. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2015, 22, 1-7.	3.0	24
85	Obesity is a significant susceptibility factor for idiopathic AA amyloidosis. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2018, 25, 37-45.	3.0	24
86	MR neurography biomarkers to characterize peripheral neuropathy in AL amyloidosis. Neurology, 2018, 91, e625-e634.	1.1	24
87	Tissue biopsy for the diagnosis of amyloidosis: experience from some centres. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2022, 29, 8-13.	3.0	24
88	Risk Stratification in Cardiac Amyloidosis: Novel Approaches. Transplantation, 2005, 80, S151-S155.	1.0	23
89	Familial Mediterranean fever in Germany: clinical presentation and amyloidosis risk. Scandinavian Journal of Rheumatology, 2013, 42, 52-58.	1.1	23
90	Cerebral amyloid angiopathy – an underdiagnosed entity in younger adults with lobar intracerebral hemorrhage?. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2013, 20, 45-47.	3.0	22

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91	Incremental value of cardiac magnetic resonance for the evaluation of cardiac tumors in adults: experience of a high volume tertiary cardiology centre. International Journal of Cardiovascular Imaging, 2017, 33, 879-888.	1.5	22
92	Guidelines for non-transplant chemotherapy for treatment of systemic AL amyloidosis: EHA-ISA working group. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2023, 30, 3-17.	3.0	22
93	Three German fibrinogen Aα-chain amyloidosis patients with the p.Glu526Val mutation. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2008, 453, 25-31.	2.8	21
94	Modulation of B Cells and Homing Marker on NK Cells Through Extracorporeal Photopheresis in Patients With Steroid-Refractory/Resistant Graft-VsHost Disease Without Hampering Anti-viral/Anti-leukemic Effects. Frontiers in Immunology, 2018, 9, 2207.	4.8	21
95	Outcome of a Salvage Third Autologous Stem Cell Transplantation in Multiple Myeloma. Biology of Blood and Marrow Transplantation, 2018, 24, 1372-1378.	2.0	20
96	Outcome of Patients With Newly Diagnosed Systemic Light-Chain Amyloidosis Associated With Deletion of 17p. Clinical Lymphoma, Myeloma and Leukemia, 2018, 18, e493-e499.	0.4	20
97	Pomalidomide and dexamethasone grant rapid haematologic responses in patients with relapsed and refractory AL amyloidosis: a European retrospective series of 153 patients. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis. 2020. 27. 231-236.	3.0	20
98	Magnetization transfer ratio quantifies polyneuropathy in hereditary transthyretin amyloidosis. Annals of Clinical and Translational Neurology, 2020, 7, 799-807.	3.7	20
99	Validation of the Criteria of Response to Treatment In AL Amyloidosis Blood, 2010, 116, 1364-1364.	1.4	19
100	Indications for High-Dose Chemotherapy with Autologous Stem Cell Support in Patients with Systemic Amyloid Light Chain Amyloidosis. Transplantation, 2005, 80, S160-S163.	1.0	18
101	Amyloid in bone marrow smears in systemic light-chain amyloidosis. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2017, 24, 52-59.	3.0	18
102	Prognostic significance of tumor burden assessed by whole-body magnetic resonance imaging in multiple myeloma patients treated with allogeneic stem cell transplantation. Haematologica, 2018, 103, 336-343.	3.5	18
103	Modified body mass index and time interval between diagnosis and operation affect survival after liver transplantation for hereditary amyloidosis: a singleâ€center analysis. Clinical Transplantation, 2013, 27, 40-48.	1.6	17
104	Allogeneic Transplantation in Multiple Myeloma—Does It Still Have a Place?. Journal of Clinical Medicine, 2020, 9, 2180.	2.4	17
105	Challenges in the management of patients with systemic light chain (AL) amyloidosis during the COVIDâ€19 pandemic. British Journal of Haematology, 2020, 190, 346-357.	2.5	17
106	Indications for Liver Transplantation in Patients with Amyloidosis: A Single-Center Experience with 11 Cases. Transplantation, 2005, 80, S156-S159.	1.0	16
107	Impact of CR before and after allogeneic and autologous transplantation in multiple myeloma: results from the EBMT NMAM2000 prospective trial. Bone Marrow Transplantation, 2015, 50, 505-510.	2.4	16
108	Shaping of CD56bri Natural Killer Cells in Patients With Steroid-Refractory/Resistant Acute Graft-vsHost Disease via Extracorporeal Photopheresis. Frontiers in Immunology, 2019, 10, 547.	4.8	16

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109	Osteopontin: a novel predictor of survival in patients with systemic light-chain amyloidosis. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2014, 21, 202-210.	3.0	15
110	Prognostic significance of semiautomatic quantification of left ventricular long axis shortening in systemic light-chain amyloidosis. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2015, 22, 45-53.	3.0	15
111	Novel recurrent chromosomal aberrations detected in clonal plasma cells of light chain amyloidosis patients show potential adverse prognostic effect: first results from a genome-wide copy number array analysis. Haematologica, 2017, 102, 1281-1290.	3.5	15
112	AL amyloidosis with a localized B cell neoplasia. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2019, 474, 353-363.	2.8	15
113	Seeded fibrils of the germline variant of human λ-III immunoglobulin light chain FOR005 have a similar core as patient fibrils with reduced stability. Journal of Biological Chemistry, 2020, 295, 18474-18484.	3.4	15
114	Bone involvement in patients with systemic AL amyloidosis mimics lytic myeloma bone disease. Haematologica, 2008, 93, 955-956.	3.5	14
115	Association of Antigen-Specific T-cell Responses with Antigen Expression and Immunoparalysis in Multiple Myeloma. Clinical Cancer Research, 2015, 21, 1712-1721.	7.0	14
116	Response Assessment in Myeloma: Practical Manual on Consistent Reporting in an Era of Dramatic Therapeutic Advances. Biology of Blood and Marrow Transplantation, 2017, 23, 1193-1202.	2.0	14
117	Efficacy and tolerability of the histone deacetylase inhibitor panobinostat in clinical practice. Hematological Oncology, 2018, 36, 210-216.	1.7	14
118	Potential therapeutic targets in plasma cell disorders: A flow cytometry study. Cytometry Part B - Clinical Cytometry, 2017, 92, 145-152.	1.5	13
119	Quantification of number of CD38 sites on bone marrow plasma cells in patients with light chain amyloidosis and smoldering multiple myeloma. Cytometry Part B - Clinical Cytometry, 2018, 94, 767-776.	1.5	13
120	A novel risk score to predict survival in advanced heart failure due to cardiac amyloidosis. Clinical Research in Cardiology, 2020, 109, 700-713.	3.3	13
121	Eight novel loci implicate shared genetic etiology in multiple myeloma, AL amyloidosis, and monoclonal gammopathy of unknown significance. Leukemia, 2020, 34, 1187-1191.	7.2	13
122	<scp>Daratumumab, lenalidomide, and dexamethasone</scp> in systemic <scp>lightâ€chain</scp> amyloidosis: High efficacy, relevant toxicity and main adverse effect of gain 1q21. American Journal of Hematology, 2021, 96, E253-E257.	4.1	13
123	Impact of time to diagnosis on Mayo stages, treatment outcome, and survival in patients with AL amyloidosis and cardiac involvement. European Journal of Haematology, 2021, 107, 449-457.	2.2	13
124	Immunoglobulin light-chain amyloidosis shares genetic susceptibility with multiple myeloma. Leukemia, 2014, 28, 2254-2256.	7.2	12
125	Regional differences in prognostic value of cardiac valve plane displacement in systemic light-chain amyloidosis. Journal of Cardiovascular Magnetic Resonance, 2017, 19, 87.	3.3	12
126	First Climpse on Real-World Efficacy Outcomes for 2000 Patients with Systemic Light Chain Amyloidosis in Europe: A Retrospective Observational Multicenter Study By the European Myeloma Network. Blood, 2020, 136, 50-51.	1.4	12

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127	MLN9708, a Novel, Investigational Oral Proteasome Inhibitor, in Patients with Relapsed or Refractory Light-Chain Amyloidosis (AL): Results of a Phase 1 Study. Blood, 2012, 120, 731-731.	1.4	12
128	Anterior Aortic Plane Systolic Excursion: A Novel Indicator of Transplant-Free Survival in Systemic Light-Chain Amyloidosis. Journal of the American Society of Echocardiography, 2016, 29, 1188-1196.	2.8	11
129	Incidence of Second Primary Malignancies after Autologous Transplantation for Multiple Myeloma in the Era of Novel Agents. Biology of Blood and Marrow Transplantation, 2018, 24, 930-936.	2.0	11
130	Lenalidomide and dexamethasone in relapsed/refractory immunoglobulin light chain (AL) amyloidosis: results from aÂlarge cohort of patients with long followâ€up. British Journal of Haematology, 2021, 195, 230-243.	2.5	11
131	Incidence of Acute Graft-Versus-Host Disease and Survival after Allogeneic Hematopoietic Cell Transplantation over Time: A Study from the Transplant Complications and Chronic Malignancies Working Party of the EBMT. Blood, 2018, 132, 2120-2120.	1.4	11
132	Reduction in Absolute Involved Free Light Chain and Difference between Involved and Uninvolved Free Light Chain Is Associated with Prolonged Major Organ Deterioration Progression-Free Survival in Patients with Newly Diagnosed AL Amyloidosis Receiving Bortezomib, Cyclophosphamide, and Dexamethasone with or without Daratumumab: Results from Andromeda. Blood, 2020, 136, 48-50.	1.4	11
133	A Randomized Phase III Trial of Melphalan and Dexamethasone (MDex) Versus Bortezomib, Melphalan and Dexamethasone (BMDex) for Untreated Patients with AL Amyloidosis. Blood, 2014, 124, 35-35.	1.4	11
134	Common Fibril Structures Imply Systemically Conserved Protein Misfolding Pathways In Vivo. Angewandte Chemie, 2017, 129, 7618-7622.	2.0	10
135	Treosulfan conditioning for allogeneic transplantation in multiple myeloma – improved overall survival in first line haematopoietic stem cell transplantation – a large retrospective study by the Chronic Malignancies Working Party of the EBMT. British Journal of Haematology, 2020, 189, e213-e217.	2.5	10
136	Analysis of the complete lambda light chain germline usage in patients with AL amyloidosis and dominant heart or kidney involvement. PLoS ONE, 2022, 17, e0264407.	2.5	10
137	Response to bendamustine is associated with a survival advantage in a heavily pretreated patients with AL amyloidosis. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2017, 24, 56-57.	3.0	9
138	Comparison of Different Stem Cell Mobilization Regimens in AL Amyloidosis Patients. Biology of Blood and Marrow Transplantation, 2017, 23, 1870-1878.	2.0	9
139	CT features in amyloidosis of the respiratory system – Comprehensive analysis in a tertiary referral center cohort. European Journal of Radiology, 2020, 129, 109123.	2.6	9
140	Outcome of Third Salvage Autologous Stem Cell Transplantation in Multiple Myeloma. Blood, 2016, 128, 993-993.	1.4	9
141	Humoral Responses and Chronic GVHD Exacerbation after COVID-19 Vaccination Post Allogeneic Stem Cell Transplantation. Vaccines, 2022, 10, 330.	4.4	9
142	Solid state NMR assignments of a human λ-III immunoglobulin light chain amyloid fibril. Biomolecular NMR Assignments, 2021, 15, 9-16.	0.8	8
143	Real-world outcomes in non-endemic hereditary transthyretin amyloidosis with polyneuropathy: a 20-year German single-referral centre experience. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2021, 28, 91-99.	3.0	8
144	CD4+CD25highCD127low Regulatory T Cells in Peripheral Blood Are Not an Independent Factor for Chronic Graft-versus-Host Disease after Allogeneic Stem Cell Transplantation. Scientific World Journal, The, 2012, 2012, 1-10.	2.1	7

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145	Stem cell transplantation in multiple myeloma and other plasma cell disorders (report from an EBMT) Tj ETQq 11	0.784314 1.3	rgBT /Overlo
146	Genome-wide association study of clinical parameters in immunoglobulin light chain amyloidosis in three patient cohorts. Haematologica, 2017, 102, e411-e414.	3.5	7
147	Reduced intensity conditioning regimens including alkylating chemotherapy do not alter survival outcomes after allogeneic hematopoietic cell transplantation in chronic lymphocytic leukemia compared to low-intensity non-myeloablative conditioning. Journal of Cancer Research and Clinical Oncology. 2019. 145. 2823-2834.	2.5	7
148	Conditioningâ€based outcomes after allogeneic transplantation for myeloma following a prior autologous transplant (1991â€2012) on behalf of EBMT CMWP. European Journal of Haematology, 2020, 104, 181-189.	2.2	7
149	Stem Cell Mobilization and Autologous Transplant for Immunoglobulin Light-Chain Amyloidosis. Hematology/Oncology Clinics of North America, 2020, 34, 1133-1144.	2.2	7
150	Upfront stem cell transplantation for newly diagnosed multiple myeloma with del(17p) and t(4;14): a study from the CMWP-EBMT. Bone Marrow Transplantation, 2021, 56, 210-217.	2.4	7
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