Vaishali Sanchorawala

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	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. American Journal of Hematology, 2005, 79, 319-328.	4.1	1,179
2	High-Dose Melphalan and Autologous Stem-Cell Transplantation in Patients with AL Amyloidosis: An 8-Year Study. Annals of Internal Medicine, 2004, 140, 85.	3.9	539
3	Systemic immunoglobulin light chain amyloidosis. Nature Reviews Disease Primers, 2018, 4, 38.	30.5	350
4	Consensus guidelines for the conduct and reporting of clinical trials in systemic light-chain amyloidosis. Leukemia, 2012, 26, 2317-2325.	7.2	332
5	Dose-Intensive Melphalan With Blood Stem-Cell Support for the Treatment of AL (Amyloid Light-Chain) Amyloidosis: Survival and Responses in 25 Patients. Blood, 1998, 91, 3662-3670.	1.4	323
6	Daratumumab-Based Treatment for Immunoglobulin Light-Chain Amyloidosis. New England Journal of Medicine, 2021, 385, 46-58.	27.0	268
7	Lenalidomide and dexamethasone in the treatment of AL amyloidosis: results of a phase 2 trial. Blood, 2007, 109, 492-496.	1.4	262
8	Outcome of AL amyloidosis after high-dose melphalan and autologous stem cell transplantation: long-term results in a series of 421 patients. Blood, 2011, 118, 4346-4352.	1.4	259
9	Acquired factor X deficiency in patients with amyloid light-chain amyloidosis: incidence, bleeding manifestations, and response to high-dose chemotherapy. Blood, 2001, 97, 1885-1887.	1.4	200
10	Light-Chain (AL) Amyloidosis: Diagnosis and Treatment. Clinical Journal of the American Society of Nephrology: CJASN, 2006, 1, 1331-1341.	4.5	191
11	Efficacy and safety of once-weekly and twice-weekly bortezomib in patients with relapsed systemic AL amyloidosis: results of a phase 1/2 study. Blood, 2011, 118, 865-873.	1.4	161
12	Amyloidosis of the gastrointestinal tract: a 13-year, single-center, referral experience. Haematologica, 2013, 98, 141-146.	3.5	155
13	Long-term outcome of patients with AL amyloidosis treated with high-dose melphalan and stem-cell transplantation. Blood, 2007, 110, 3561-3563.	1.4	154
14	Weekly and twice-weekly bortezomib in patients with systemic AL amyloidosis: results of a phase 1 dose-escalation study. Blood, 2009, 114, 1489-1497.	1.4	153
15	An overview of the use of high-dose melphalan with autologous stem cell transplantation for the treatment of AL amyloidosis. Bone Marrow Transplantation, 2001, 28, 637-642.	2.4	149
16	Daratumumab plus CyBorD for patients with newly diagnosed AL amyloidosis: safety run-in results of ANDROMEDA. Blood, 2020, 136, 71-80.	1.4	146
17	Tolerability and Efficacy of Thalidomide for the Treatment of Patients with Light Chain–Associated (AL) Amyloidosis. Clinical Lymphoma and Myeloma, 2003, 3, 241-246.	2.1	137
18	Development and validation of a survival staging system incorporating BNP in patients with light chain amyloidosis. Blood, 2019, 133, 215-223.	1.4	118

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19	Pomalidomide and dexamethasone in the treatment of AL amyloidosis: results of a phase 1 and 2 trial. Blood, 2016, 128, 1059-1062.	1.4	117
20	Safety, tolerability, and response rates of daratumumab in relapsed AL amyloidosis: results of a phase 2 study. Blood, 2020, 135, 1541-1547.	1.4	111
21	Effect of Dose-Intensive Intravenous Melphalan and Autologous Blood Stem-Cell Transplantation on AL Amyloidosis–Associated Renal Disease. Annals of Internal Medicine, 2001, 134, 746.	3.9	111
22	Improvement in quality of life of patients with AL amyloidosis treated with high-dose melphalan and autologous stem cell transplantation. Blood, 2004, 104, 1888-1893.	1.4	109
23	Long-term outcome of patients with AL amyloidosis treated with high-dose melphalan and stem cell transplantation: 20-year experience. Blood, 2015, 126, 2345-2347.	1.4	109
24	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
25	Monoclonal gammopathy of undetermined significance in systemic transthyretin amyloidosis (ATTR). Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2018, 25, 62-67.	3.0	108
26	High-dose intravenous melphalan and autologous stem cell transplantation as initial therapy or following two cycles of oral chemotherapy for the treatment of AL amyloidosis: results of a prospective randomized trial. Bone Marrow Transplantation, 2004, 33, 381-388.	2.4	107
27	Persistent Pleural Effusions in Primary Systemic Amyloidosis. Chest, 2003, 124, 969-977.	0.8	106
28	Cardiac Transplantation Followed by Dose-Intensive Melphalan and Autologous Stem-Cell Transplantation for Light Chain Amyloidosis and Heart Failure. Transplantation, 2010, 90, 905-911.	1.0	103
29	Kidney dysfunction during lenalidomide treatment for AL amyloidosis. Nephrology Dialysis Transplantation, 2011, 26, 881-886.	0.7	99
30	Update on treatment of light chain amyloidosis. Haematologica, 2014, 99, 209-221.	3.5	93
31	Serum free light-chain responses after high-dose intravenous melphalan and autologous stem cell transplantation for AL (primary) amyloidosis. Bone Marrow Transplantation, 2005, 36, 597-600.	2.4	92
32	Serum Free Light Chain Responses after High-Dose Intravenous Melphalan and Autologous Stem Cell Transplantation for AL (Primary) Amyloidosis Blood, 2004, 104, 942-942.	1.4	82
33	Rationale, application and clinical qualification for NT-proBNP as a surrogate end point in pivotal clinical trials in patients with AL amyloidosis. Leukemia, 2016, 30, 1979-1986.	7.2	73
34	Intermediate-dose intravenous melphalan and blood stem cells mobilized with sequential GM+G-CSF or G-CSF alone to treat AL (amyloid light chain) amyloidosis. British Journal of Haematology, 1999, 104, 553-559.	2.5	68
35	Incidence and outcome of acute renal failure complicating autologous stem cell transplantation for AL amyloidosis. Kidney International, 2003, 63, 1868-1873.	5.2	63
36	Clinical and molecular characteristics of patients with non-amyloid light chain deposition disorders, and outcome following treatment with high-dose melphalan and autologous stem cell transplantation. Bone Marrow Transplantation, 2006, 38, 339-343.	2.4	62

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37	Long-term follow-up from a phase 1/2 study of single-agent bortezomib in relapsed systemic AL amyloidosis. Blood, 2014, 124, 2498-2506.	1.4	62
38	Longitudinal systolic strain, cardiac function improvement, and survival following treatment of light-chain (AL) cardiac amyloidosis. European Heart Journal Cardiovascular Imaging, 2017, 18, 1057-1064.	1.2	60
39	High-dose intravenous melphalan with autologous stem cell transplantation in AL amyloidosis-associated end-stage renal disease. Kidney International, 2003, 63, 1051-1057.	5.2	59
40	Increases in B-type natriuretic peptide (BNP) during treatment with lenalidomide in AL amyloidosis. Blood, 2010, 116, 5071-5072.	1.4	59
41	Long-term outcome of kidney transplantation in AL amyloidosis. Kidney International, 2019, 95, 405-411.	5.2	57
42	Spontaneous rupture of the spleen in AL amyloidosis. American Journal of Hematology, 2003, 74, 131-135.	4.1	56
43	Induction Therapy with Bortezomib Followed by Bortezomib-High Dose Melphalan and Stem Cell Transplantation for Light Chain Amyloidosis: Results of a Prospective Clinical Trial. Biology of Blood and Marrow Transplantation, 2015, 21, 1445-1451.	2.0	55
44	Early Detection of Multiorgan Light-Chain Amyloidosis by Whole-Body ¹⁸ F-Florbetapir PET/CT. Journal of Nuclear Medicine, 2019, 60, 1234-1239.	5.0	54
45	Venetoclax induces deep hematologic remissions in t(11;14) relapsed/refractory AL amyloidosis. Blood Cancer Journal, 2021, 11, 10.	6.2	53
46	Melphalan, lenalidomide and dexamethasone for the treatment of immunoglobulin light chain amyloidosis: results of a phase II trial. Haematologica, 2013, 98, 789-792.	3.5	50
47	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
48	AL amyloidosis associated with B-cell lymphoproliferative disorders: Frequency and treatment outcomes. American Journal of Hematology, 2006, 81, 692-695.	4.1	47
49	Localized amyloidosis of the breast: a case series. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2011, 18, 72-75.	3.0	47
50	Low-dose continuous oral melphalan for the treatment of primary systemic (AL) amyloidosis. British Journal of Haematology, 2002, 117, 886-889.	2.5	46
51	Safety and Efficacy of Carfilzomib (CFZ) in Previously-Treated Systemic Light-Chain (AL) Amyloidosis. Blood, 2016, 128, 645-645.	1.4	46
52	Marked progress in AL amyloidosis survival: a 40-year longitudinal natural history study. Blood Cancer Journal, 2021, 11, 139.	6.2	45
53	Quantitative serum free light chain assay in the diagnostic evaluation of AL amyloidosis. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2005, 12, 210-215.	3.0	44
54	Myocardial infarction with "clean coronaries―caused by amyloid light-chain AL amyloidosis: a case report and literature review. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2011, 18, 160-164.	3.0	42

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55	Guidelines for high dose chemotherapy and stem cell transplantation for systemic AL amyloidosis: EHA-ISA working group guidelines. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2022, 29, 1-7.	3.0	42
56	Safety and efficacy of high-dose melphalan and auto-SCT in patients with AL amyloidosis and cardiac involvement. Bone Marrow Transplantation, 2014, 49, 434-439.	2.4	41
57	Improved Quantification of CardiacÂAmyloid Burden in SystemicÂLight ChainÂAmyloidosis. JACC: Cardiovascular Imaging, 2020, 13, 1325-1336.	5.3	41
58	Bortezomib in a phase 1 trial for patients with relapsed AL amyloidosis: cardiac responses and overall effects. QJM - Monthly Journal of the Association of Physicians, 2011, 104, 957-970.	0.5	40
59	Assessment of minimal residual disease using multiparametric flow cytometry in patients with AL amyloidosis. Blood Advances, 2020, 4, 880-884.	5.2	40
60	Association of acquired von Willebrand syndrome with AL amyloidosis. American Journal of Hematology, 2007, 82, 363-367.	4.1	39
61	High-dose melphalan and stem cell transplantation for patients with AL amyloidosis: trends in treatment-related mortality over the past 17 years at a single referral center. Blood, 2012, 120, 4445-4446.	1.4	38
62	Bortezomib and high-dose melphalan conditioning for stem cell transplantation for AL amyloidosis: a pilot study. Haematologica, 2011, 96, 1890-1892.	3.5	34
63	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
64	Predictive factors for hematopoietic engraftment after autologous peripheral blood stem cell transplantation for AL amyloidosis. Bone Marrow Transplantation, 2005, 35, 567-575.	2.4	33
65	Successful treatment of AL amyloidosis with high-dose melphalan and autologous stem cell transplantation in patients over age 65. Blood, 2006, 108, 3945-3947.	1.4	33
66	Tandem cycles of high-dose melphalan and autologous stem cell transplantation increases the response rate in AL amyloidosis. Bone Marrow Transplantation, 2007, 40, 557-562.	2.4	33
67	Durable hematologic complete responses can be achieved with lenalidomide in AL amyloidosis. Blood, 2010, 116, 1990-1991.	1.4	33
68	Bendamustine With Dexamethasone in Relapsed/Refractory Systemic Light-Chain Amyloidosis: Results of a Phase II Study. Journal of Clinical Oncology, 2020, 38, 1455-1462.	1.6	31
69	High-Dose Melphalan and Stem Cell Transplantation in Patients on Dialysis Due to Immunoglobulin Light-Chain Amyloidosis and Monoclonal Immunoglobulin Deposition Disease. Biology of Blood and Marrow Transplantation, 2018, 24, 127-132.	2.0	31
70	Hematologic relapse in AL amyloidosis after high-dose melphalan and stem cell transplantation. Blood, 2017, 130, 1383-1386.	1.4	30
71	An overview of high-dose melphalan and stem cell transplantation in the treatment of AL amyloidosis. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2007, 14, 261-269.	3.0	29
72	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

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73	Azotemia associated with use of lenalidomide in plasma cell dyscrasias. Leukemia and Lymphoma, 2008, 49, 1108-1115.	1.3	28
74	Regression of cardiac wall thickness following chemotherapy and stem cell transplantation for light chain (AL) amyloidosis. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2011, 18, 130-131.	3.0	27
75	A longitudinal evaluation of healthâ€related quality of life in patients with <scp>AL</scp> amyloidosis: associations with health outcomes over time. British Journal of Haematology, 2017, 179, 461-470.	2.5	27
76	Presence of t(11;14) in AL amyloidosis as a marker of response when treated with a bortezomib-based regimen. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2020, 27, 244-249.	3.0	27
77	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
78	New Hematologic Response Criteria Predict Survival in Patients With Immunoglobulin Light Chain Amyloidosis Treated With High-Dose Melphalan and Autologous Stem-Cell Transplantation. Journal of Clinical Oncology, 2013, 31, 2749-2750.	1.6	26
79	Patient outcomes in light chain (AL) amyloidosis: The clock is ticking from symptoms to diagnosis. European Journal of Haematology, 2020, 105, 495-501.	2.2	26
80	Hepatic response after high-dose melphalan and stem cell transplantation in patients with AL amyloidosis associated liver disease. Haematologica, 2009, 94, 1029-1032.	3.5	25
81	Delay treatment of AL amyloidosis at relapse until symptomatic: devil is in the details. Blood Advances, 2019, 3, 216-218.	5.2	25
82	Establishment of brain natriuretic peptide ―based criteria for evaluating cardiac response to treatment in light chain (AL) amyloidosis. British Journal of Haematology, 2020, 188, 424-427.	2.5	25
83	Lymphadenopathy as a manifestation of amyloidosis: a case series. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2014, 21, 256-260.	3.0	24
84	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
85	Race/ethnicity in systemic AL amyloidosis: perspectives on disease and outcome disparities. Blood Cancer Journal, 2020, 10, 118.	6.2	21
86	Long-Term Outcome of a Phase 1 Study of the Investigational Oral Proteasome Inhibitor (PI) Ixazomib at the Recommended Phase 3 Dose (RP3D) in Patients (Pts) with Relapsed or Refractory Systemic Light-Chain (AL) Amyloidosis (RRAL). Blood, 2014, 124, 3450-3450.	1.4	21
87	Cardiac Amyloidosis: Evolving Approach to Diagnosis and Management. Current Treatment Options in Cardiovascular Medicine, 2011, 13, 528-542.	0.9	20
88	Modified high-dose melphalan and autologous SCT for AL amyloidosis or high-risk myeloma: analysis of SWOG trial S0115. Bone Marrow Transplantation, 2013, 48, 1537-1542.	2.4	20
89	Macroglossia – not always AL amyloidosis. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2011, 18, 83-86.	3.0	19
90	Clinical presentation and treatment responses in IgM-related AL amyloidosis. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2015, 22, 229-235.	3.0	19

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91	Risk factors for venous thromboembolism in immunoglobulin light chain amyloidosis. Haematologica, 2016, 101, 86-90.	3.5	19
92	High-Dose Melphalan and Autologous Peripheral Blood Stem Cell Transplantation in AL Amyloidosis. Acta Haematologica, 2020, 143, 381-387.	1.4	19
93	A Case of Atypical Light Chain Deposition Disease—Diagnosis and Treatment. Clinical Journal of the American Society of Nephrology: CJASN, 2007, 2, 858-867.	4.5	18
94	Oral Cyclic Melphalan and Dexamethasone for Patients With AL Amyloidosis. Clinical Lymphoma, Myeloma and Leukemia, 2010, 10, 469-472.	0.4	18
95	Depression and anxiety in patients with AL amyloidosis as assessed by the SF-36 questionnaire: experience in 1226 patients. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2016, 23, 188-193.	3.0	18
96	Treatment patterns and health care resource utilization among patients with relapsed/refractory systemic light chain amyloidosis. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2018, 25, 1-7.	3.0	18
97	Organ responses after highdose melphalan and stemcell transplantation in AL amyloidosis. Leukemia, 2021, 35, 916-919.	7.2	18
98	Immunologic recovery after autologous blood stem cell transplantation in patients with AL-amyloidosis. Bone Marrow Transplantation, 2001, 28, 1105-1109.	2.4	17
99	Amyloidotic Cardiomyopathy: Multidisciplinary Approach to Diagnosis and Treatment. Heart Failure Clinics, 2011, 7, 385-393.	2.1	17
100	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
101	Spontaneous rupture of the liver in a patient with systemic AL amyloidosis undergoing treatment with high-dose melphalan and autologous stem cell transplantation: A case report with literature review. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2009, 16, 103-107.	3.0	16
102	High Dose Melphalan and Autologous Peripheral Blood Stem Cell Transplantation in AL Amyloidosis. Hematology/Oncology Clinics of North America, 2014, 28, 1131-1144.	2.2	16
103	Validation of new renal staging system in AL amyloidosis treated with high dose melphalan and stem cell transplantation. American Journal of Hematology, 2016, 91, E458-60.	4.1	16
104	Psychometric validation of the SF-36 Health Survey in light chain amyloidosis: results from community-based and clinic-based samples. Patient Related Outcome Measures, 2017, Volume 8, 157-167.	1.2	16
105	Transbronchial biopsies safely diagnose amyloid lung disease. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2017, 24, 37-41.	3.0	15
106	Orthotopic heart transplant rejection in association with immunomodulatory therapy for AL amyloidosis: A case series and review of the literature. American Journal of Transplantation, 2019, 19, 3185-3190.	4.7	15
107	Subcutaneous daratumumab + bortezomib, cyclophosphamide, and dexamethasone (VCd) in patients with newly diagnosed light chain (AL) amyloidosis: Updated results from the phase 3 ANDROMEDA study Journal of Clinical Oncology, 2021, 39, 8003-8003.	1.6	15
108	A Phase I Dose-Escalation Study of Carfilzomib in Patients with Previously-Treated Systemic Light-Chain (AL) Amyloidosis. Blood, 2014, 124, 4741-4741.	1.4	15

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109	Short and long-term outcome of treatment with high-dose melphalan and stem cell transplantation for multiple myeloma-associated AL amyloidosis. Annals of Hematology, 2010, 89, 579-584.	1.8	14
110	A second course of high-dose melphalan and auto-SCT for the treatment of relapsed AL amyloidosis. Bone Marrow Transplantation, 2011, 46, 976-980.	2.4	14
111	A new era of amyloidosis: the trends at a major US referral centre. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2019, 26, 192-196.	3.0	14
112	Induction Therapy with Bortezomib and Dexamethasone and Conditioning with High-Dose Melphalan and Bortezomib Followed by Autologous Stem Cell Transplantation for Immunoglobulin Light Chain Amyloidosis: Long-Term Follow-Up Analysis. Biology of Blood and Marrow Transplantation, 2019, 25, e169-e173.	2.0	14
113	Quantitative [18F]florbetapir PET/CT may identify lung involvement in patients with systemic AL amyloidosis. European Journal of Nuclear Medicine and Molecular Imaging, 2020, 47, 1998-2009.	6.4	14
114	Comparing measures of hematologic response after high-dose melphalan and stem cell transplantation in AL amyloidosis. Blood Cancer Journal, 2020, 10, 88.	6.2	14
115	Amyloid Deposits in the Bone Marrow of Patients with Immunoglobulin Light Chain Amyloidosis Do Not Impact Stem Cell Mobilization or Engraftment. Biology of Blood and Marrow Transplantation, 2012, 18, 1935-1938.	2.0	13
116	The incidence of atrial fibrillation among patients with AL amyloidosis undergoing high-dose melphalan and stem cell transplantation: experience at a single institution. Bone Marrow Transplantation, 2017, 52, 1349-1351.	2.4	13
117	A library of ATTR amyloidosis patient-specific induced pluripotent stem cells for disease modelling and <i>in vitro</i> testing of novel therapeutics. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2018, 25, 148-155.	3.0	13
118	Use of melphalan (M)/dexamethasone (D)/bortezomib in AL amyloidosis Journal of Clinical Oncology, 2010, 28, 8024-8024.	1.6	13
119	Long-term outcome of patients with monoclonal Ig deposition disease treated with high-dose melphalan and stem cell transplantation. Bone Marrow Transplantation, 2011, 46, 161-162.	2.4	12
120	The Effect of Bone Marrow Plasma Cell Burden on Survival in Patients with Light Chain Amyloidosis Undergoing High-Dose Melphalan and Autologous Stem Cell Transplantation. Biology of Blood and Marrow Transplantation, 2016, 22, 1729-1732.	2.0	12
121	The six-minute walk test in patients with AL amyloidosis: a single centre case series. British Journal of Haematology, 2017, 177, 388-394.	2.5	12
122	Modified High-Dose Melphalan and Autologous Stem Cell Transplantation for Immunoglobulin Light Chain Amyloidosis. Biology of Blood and Marrow Transplantation, 2018, 24, 1823-1827.	2.0	12
123	Outcomes of patients with AL amyloidosis and low serum free light chain levels at diagnosis. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2018, 25, 156-159.	3.0	12
124	Prevalence and prognostic value of Dâ€dimer elevation in patients with AL amyloidosis. American Journal of Hematology, 2019, 94, 1098-1103.	4.1	12
125	Predictors of hematologic response and survival with stem cell transplantation in <scp>AL</scp> amyloidosis: A 25â€year longitudinal study. American Journal of Hematology, 2022, 97, 1189-1199.	4.1	12
126	Plerixafor-augmented peripheral blood stem cell mobilization in AL amyloidosis with cardiac involvement: a case series. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2014, 21, 149-153.	3.0	11

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127	Bortezomib ocular toxicities: Outcomes with ketotifen. American Journal of Hematology, 2019, 94, E80-E82.	4.1	11
128	The Role of Kidney Transplantation in Monoclonal Ig Deposition Disease. Kidney International Reports, 2020, 5, 485-493.	0.8	11
129	Left Atrial Mechanics Associates With Paroxysmal Atrial Fibrillation in Light-Chain Amyloidosis Following StemÂCell Transplantation. JACC: CardioOncology, 2020, 2, 721-731.	4.0	11
130	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
131	Multiple arterial and venous thromboembolic complications in AL amyloidosis and cardiac involvement: a case report and literature review. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2012, 19, 156-160.	3.0	10
132	Once AL amyloidosis: not always AL amyloidosis. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2018, 25, 139-140.	3.0	10
133	Relapse Rate and Long-Term Survival of AL Amyloidosis Patients Treated with High-Dose Melphalan and Autologous Stem Cell Transplantation (HDM/SCT) Blood, 2006, 108, 3094-3094.	1.4	10
134	Predictive factors of outcomes in patients with <scp>AL</scp> amyloidosis treated with daratumumab. American Journal of Hematology, 2022, 97, 79-89.	4.1	10
135	Summary of the EHA-ISA Working Group Guidelines for High-dose Chemotherapy and Stem Cell Transplantation for Systemic AL Amyloidosis. HemaSphere, 2022, 6, e681.	2.7	10
136	Diaphragm paralysis in primary systemic amyloidosis. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2005, 12, 193-196.	3.0	9
137	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
138	Immunoglobulin heavy light chain test quantifies clonal disease in patients with AL amyloidosis and normal serum free light chain ratio. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2016, 23, 214-220.	3.0	8
139	<p>Treatment Options For Relapsed/refractory Systemic Light-Chain (AL) Amyloidosis: Current Perspectives</p> . Journal of Blood Medicine, 2019, Volume 10, 373-380.	1.7	8
140	The utility of repeat kidney biopsy in systemic immunoglobulin light chain amyloidosis. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2020, 27, 17-24.	3.0	8
141	Safety, Tolerability and Response Rates of Daratumumab in Patients with Relapsed Light Chain (AL) Amyloidosis: Results of a Phase II Study. Blood, 2018, 132, 2005-2005.	1.4	8
142	Phase I/II study of bortezomib (B) in patients with systemic AL-amyloidosis (AL). Journal of Clinical Oncology, 2007, 25, 8050-8050.	1.6	8
143	Neurological manifestations of hereditary transthyretin amyloidosis: a focus on diagnostic delays. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2022, 29, 184-189.	3.0	8
144	Daratumumab in AL amyloidosis. Blood, 2022, 140, 2317-2322.	1.4	8

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145	Systemic AL amyloidosis with an undetectable plasma cell dyscrasia: A zebra without stripes. American Journal of Hematology, 2020, 95, E45-E48.	4.1	7
146	Systemic Amyloidosis Caused by Monoclonal Immunoglobulins. Hematology/Oncology Clinics of North America, 2020, 34, 1099-1113.	2.2	7
147	Predictors and outcomes of acute kidney injury during autologous stem cell transplantation in AL amyloidosis. Nephrology Dialysis Transplantation, 2022, 37, 1281-1288.	0.7	7
148	Ixazomib-dexamethasone (Ixa-Dex) vs physician's choice (PC) in relapsed/refractory (RR) primary systemic AL amyloidosis (AL) patients (pts) by prior proteasome inhibitor (PI) exposure in the phase III TOURMALINE-AL1 trial Journal of Clinical Oncology, 2020, 38, 8546-8546.	1.6	7
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