

Vaishali Sanchorawala

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/5414261/publications.pdf>

Version: 2024-02-01

253
papers

9,725
citations

50170

46
h-index

42291

92
g-index

256
all docs

256
docs citations

256
times ranked

4451
citing authors

#	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.	2.0	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.	2.0	539
3	Systemic immunoglobulin light chain amyloidosis. Nature Reviews Disease Primers, 2018, 4, 38.	18.1	350
4	Consensus guidelines for the conduct and reporting of clinical trials in systemic light-chain amyloidosis. Leukemia, 2012, 26, 2317-2325.	3.3	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.	0.6	323
6	Daratumumab-Based Treatment for Immunoglobulin Light-Chain Amyloidosis. New England Journal of Medicine, 2021, 385, 46-58.	13.9	268
7	Lenalidomide and dexamethasone in the treatment of AL amyloidosis: results of a phase 2 trial. Blood, 2007, 109, 492-496.	0.6	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.	0.6	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.	0.6	200
10	Light-Chain (AL) Amyloidosis: Diagnosis and Treatment. Clinical Journal of the American Society of Nephrology: CJASN, 2006, 1, 1331-1341.	2.2	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.	0.6	161
12	Amyloidosis of the gastrointestinal tract: a 13-year, single-center, referral experience. Haematologica, 2013, 98, 141-146.	1.7	155
13	Long-term outcome of patients with AL amyloidosis treated with high-dose melphalan and stem-cell transplantation. Blood, 2007, 110, 3561-3563.	0.6	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.	0.6	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.	1.3	149
16	Daratumumab plus CyBORd for patients with newly diagnosed AL amyloidosis: safety run-in results of ANDROMEDA. Blood, 2020, 136, 71-80.	0.6	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.	0.6	118

#	ARTICLE	IF	CITATIONS
19	Pomalidomide and dexamethasone in the treatment of AL amyloidosis: results of a phase 1 and 2 trial. <i>Blood</i> , 2016, 128, 1059-1062.	0.6	117
20	Safety, tolerability, and response rates of daratumumab in relapsed AL amyloidosis: results of a phase 2 study. <i>Blood</i> , 2020, 135, 1541-1547.	0.6	111
21	Effect of Dose-Intensive Intravenous Melphalan and Autologous Blood Stem-Cell Transplantation on AL Amyloidosis-Associated Renal Disease. <i>Annals of Internal Medicine</i> , 2001, 134, 746.	2.0	111
22	Improvement in quality of life of patients with AL amyloidosis treated with high-dose melphalan and autologous stem cell transplantation. <i>Blood</i> , 2004, 104, 1888-1893.	0.6	109
23	Long-term outcome of patients with AL amyloidosis treated with high-dose melphalan and stem cell transplantation: 20-year experience. <i>Blood</i> , 2015, 126, 2345-2347.	0.6	109
24	A phase 1/2 study of the oral proteasome inhibitor ixazomib in relapsed or refractory AL amyloidosis. <i>Blood</i> , 2017, 130, 597-605.	0.6	108
25	Monoclonal gammopathy of undetermined significance in systemic transthyretin amyloidosis (ATTR). <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2018, 25, 62-67.	1.4	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. <i>Bone Marrow Transplantation</i> , 2004, 33, 381-388.	1.3	107
27	Persistent Pleural Effusions in Primary Systemic Amyloidosis. <i>Chest</i> , 2003, 124, 969-977.	0.4	106
28	Cardiac Transplantation Followed by Dose-Intensive Melphalan and Autologous Stem-Cell Transplantation for Light Chain Amyloidosis and Heart Failure. <i>Transplantation</i> , 2010, 90, 905-911.	0.5	103
29	Kidney dysfunction during lenalidomide treatment for AL amyloidosis. <i>Nephrology Dialysis Transplantation</i> , 2011, 26, 881-886.	0.4	99
30	Update on treatment of light chain amyloidosis. <i>Haematologica</i> , 2014, 99, 209-221.	1.7	93
31	Serum free light-chain responses after high-dose intravenous melphalan and autologous stem cell transplantation for AL (primary) amyloidosis. <i>Bone Marrow Transplantation</i> , 2005, 36, 597-600.	1.3	92
32	Serum Free Light Chain Responses after High-Dose Intravenous Melphalan and Autologous Stem Cell Transplantation for AL (Primary) Amyloidosis. <i>Blood</i> , 2004, 104, 942-942.	0.6	82
33	Rationale, application and clinical qualification for NT-proBNP as a surrogate end point in pivotal clinical trials in patients with AL amyloidosis. <i>Leukemia</i> , 2016, 30, 1979-1986.	3.3	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. <i>British Journal of Haematology</i> , 1999, 104, 553-559.	1.2	68
35	Incidence and outcome of acute renal failure complicating autologous stem cell transplantation for AL amyloidosis. <i>Kidney International</i> , 2003, 63, 1868-1873.	2.6	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. <i>Bone Marrow Transplantation</i> , 2006, 38, 339-343.	1.3	62

#	ARTICLE	IF	CITATIONS
37	Long-term follow-up from a phase 1/2 study of single-agent bortezomib in relapsed systemic AL amyloidosis. <i>Blood</i> , 2014, 124, 2498-2506.	0.6	62
38	Longitudinal systolic strain, cardiac function improvement, and survival following treatment of light-chain (AL) cardiac amyloidosis. <i>European Heart Journal Cardiovascular Imaging</i> , 2017, 18, 1057-1064.	0.5	60
39	High-dose intravenous melphalan with autologous stem cell transplantation in AL amyloidosis-associated end-stage renal disease. <i>Kidney International</i> , 2003, 63, 1051-1057.	2.6	59
40	Increases in B-type natriuretic peptide (BNP) during treatment with lenalidomide in AL amyloidosis. <i>Blood</i> , 2010, 116, 5071-5072.	0.6	59
41	Long-term outcome of kidney transplantation in AL amyloidosis. <i>Kidney International</i> , 2019, 95, 405-411.	2.6	57
42	Spontaneous rupture of the spleen in AL amyloidosis. <i>American Journal of Hematology</i> , 2003, 74, 131-135.	2.0	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. <i>Biology of Blood and Marrow Transplantation</i> , 2015, 21, 1445-1451.	2.0	55
44	Early Detection of Multiorgan Light-Chain Amyloidosis by Whole-Body ¹⁸ F-Fluorbetapir PET/CT. <i>Journal of Nuclear Medicine</i> , 2019, 60, 1234-1239.	2.8	54
45	Venetoclax induces deep hematologic remissions in t(11;14) relapsed/refractory AL amyloidosis. <i>Blood Cancer Journal</i> , 2021, 11, 10.	2.8	53
46	Melphalan, lenalidomide and dexamethasone for the treatment of immunoglobulin light chain amyloidosis: results of a phase II trial. <i>Haematologica</i> , 2013, 98, 789-792.	1.7	50
47	Clarification on the definition of complete haematologic response in light-chain (AL) amyloidosis. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2021, 28, 1-2.	1.4	49
48	AL amyloidosis associated with B-cell lymphoproliferative disorders: Frequency and treatment outcomes. <i>American Journal of Hematology</i> , 2006, 81, 692-695.	2.0	47
49	Localized amyloidosis of the breast: a case series. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2011, 18, 72-75.	1.4	47
50	Low-dose continuous oral melphalan for the treatment of primary systemic (AL) amyloidosis. <i>British Journal of Haematology</i> , 2002, 117, 886-889.	1.2	46
51	Safety and Efficacy of Carfilzomib (CFZ) in Previously-Treated Systemic Light-Chain (AL) Amyloidosis. <i>Blood</i> , 2016, 128, 645-645.	0.6	46
52	Marked progress in AL amyloidosis survival: a 40-year longitudinal natural history study. <i>Blood Cancer Journal</i> , 2021, 11, 139.	2.8	45
53	Quantitative serum free light chain assay in the diagnostic evaluation of AL amyloidosis. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2005, 12, 210-215.	1.4	44
54	Myocardial infarction with acute coronary arteries caused by amyloid light-chain AL amyloidosis: a case report and literature review. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2011, 18, 160-164.	1.4	42

#	ARTICLE	IF	CITATIONS
55	Guidelines for high dose chemotherapy and stem cell transplantation for systemic AL amyloidosis: EHA-ISA working group guidelines. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2022, 29, 1-7.	1.4	42
56	Safety and efficacy of high-dose melphalan and auto-SCT in patients with AL amyloidosis and cardiac involvement. <i>Bone Marrow Transplantation</i> , 2014, 49, 434-439.	1.3	41
57	Improved Quantification of Cardiac Amyloid Burden in Systemic Light Chain Amyloidosis. <i>JACC: Cardiovascular Imaging</i> , 2020, 13, 1325-1336.	2.3	41
58	Bortezomib in a phase 1 trial for patients with relapsed AL amyloidosis: cardiac responses and overall effects. <i>QJM - Monthly Journal of the Association of Physicians</i> , 2011, 104, 957-970.	0.2	40
59	Assessment of minimal residual disease using multiparametric flow cytometry in patients with AL amyloidosis. <i>Blood Advances</i> , 2020, 4, 880-884.	2.5	40
60	Association of acquired von Willebrand syndrome with AL amyloidosis. <i>American Journal of Hematology</i> , 2007, 82, 363-367.	2.0	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. <i>Blood</i> , 2012, 120, 4445-4446.	0.6	38
62	Bortezomib and high-dose melphalan conditioning for stem cell transplantation for AL amyloidosis: a pilot study. <i>Haematologica</i> , 2011, 96, 1890-1892.	1.7	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). <i>Blood</i> , 2019, 134, 139-139.	0.6	34
64	Predictive factors for hematopoietic engraftment after autologous peripheral blood stem cell transplantation for AL amyloidosis. <i>Bone Marrow Transplantation</i> , 2005, 35, 567-575.	1.3	33
65	Successful treatment of AL amyloidosis with high-dose melphalan and autologous stem cell transplantation in patients over age 65. <i>Blood</i> , 2006, 108, 3945-3947.	0.6	33
66	Tandem cycles of high-dose melphalan and autologous stem cell transplantation increases the response rate in AL amyloidosis. <i>Bone Marrow Transplantation</i> , 2007, 40, 557-562.	1.3	33
67	Durable hematologic complete responses can be achieved with lenalidomide in AL amyloidosis. <i>Blood</i> , 2010, 116, 1990-1991.	0.6	33
68	Bendamustine With Dexamethasone in Relapsed/Refractory Systemic Light-Chain Amyloidosis: Results of a Phase II Study. <i>Journal of Clinical Oncology</i> , 2020, 38, 1455-1462.	0.8	31
69	High-Dose Melphalan and Stem Cell Transplantation in Patients on Dialysis Due to Immunoglobulin Light-Chain Amyloidosis and Monoclonal Immunoglobulin Deposition Disease. <i>Biology of Blood and Marrow Transplantation</i> , 2018, 24, 127-132.	2.0	31
70	Hematologic relapse in AL amyloidosis after high-dose melphalan and stem cell transplantation. <i>Blood</i> , 2017, 130, 1383-1386.	0.6	30
71	An overview of high-dose melphalan and stem cell transplantation in the treatment of AL amyloidosis. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2007, 14, 261-269.	1.4	29
72	A randomized phase 3 study of ixazomib+dexamethasone versus physician's choice in relapsed or refractory AL amyloidosis. <i>Leukemia</i> , 2022, 36, 225-235.	3.3	29

#	ARTICLE	IF	CITATIONS
73	Azotemia associated with use of lenalidomide in plasma cell dyscrasias. <i>Leukemia and Lymphoma</i> , 2008, 49, 1108-1115.	0.6	28
74	Regression of cardiac wall thickness following chemotherapy and stem cell transplantation for light chain (AL) amyloidosis. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2011, 18, 130-131.	1.4	27
75	A longitudinal evaluation of health-related quality of life in patients with <scp>AL</scp> amyloidosis: associations with health outcomes over time. <i>British Journal of Haematology</i> , 2017, 179, 461-470.	1.2	27
76	Presence of t(11;14) in AL amyloidosis as a marker of response when treated with a bortezomib-based regimen. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2020, 27, 244-249.	1.4	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. <i>Blood</i> , 2019, 134, 3166-3166.	0.6	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. <i>Journal of Clinical Oncology</i> , 2013, 31, 2749-2750.	0.8	26
79	Patient outcomes in light chain (AL) amyloidosis: The clock is ticking from symptoms to diagnosis. <i>European Journal of Haematology</i> , 2020, 105, 495-501.	1.1	26
80	Hepatic response after high-dose melphalan and stem cell transplantation in patients with AL amyloidosis associated liver disease. <i>Haematologica</i> , 2009, 94, 1029-1032.	1.7	25
81	Delay treatment of AL amyloidosis at relapse until symptomatic: devil is in the details. <i>Blood Advances</i> , 2019, 3, 216-218.	2.5	25
82	Establishment of brain natriuretic peptide -based criteria for evaluating cardiac response to treatment in light chain (AL) amyloidosis. <i>British Journal of Haematology</i> , 2020, 188, 424-427.	1.2	25
83	Lymphadenopathy as a manifestation of amyloidosis: a case series. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2014, 21, 256-260.	1.4	24
84	Guidelines for non-transplant chemotherapy for treatment of systemic AL amyloidosis: EHA-ISA working group. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2023, 30, 3-17.	1.4	22
85	Race/ethnicity in systemic AL amyloidosis: perspectives on disease and outcome disparities. <i>Blood Cancer Journal</i> , 2020, 10, 118.	2.8	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). <i>Blood</i> , 2014, 124, 3450-3450.	0.6	21
87	Cardiac Amyloidosis: Evolving Approach to Diagnosis and Management. <i>Current Treatment Options in Cardiovascular Medicine</i> , 2011, 13, 528-542.	0.4	20
88	Modified high-dose melphalan and autologous SCT for AL amyloidosis or high-risk myeloma: analysis of SWOG trial S0115. <i>Bone Marrow Transplantation</i> , 2013, 48, 1537-1542.	1.3	20
89	Macroglossia "not always AL amyloidosis. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2011, 18, 83-86.	1.4	19
90	Clinical presentation and treatment responses in IgM-related AL amyloidosis. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2015, 22, 229-235.	1.4	19

#	ARTICLE	IF	CITATIONS
91	Risk factors for venous thromboembolism in immunoglobulin light chain amyloidosis. <i>Haematologica</i> , 2016, 101, 86-90.	1.7	19
92	High-Dose Melphalan and Autologous Peripheral Blood Stem Cell Transplantation in AL Amyloidosis. <i>Acta Haematologica</i> , 2020, 143, 381-387.	0.7	19
93	A Case of Atypical Light Chain Deposition Disease—Diagnosis and Treatment. <i>Clinical Journal of the American Society of Nephrology: CJASN</i> , 2007, 2, 858-867.	2.2	18
94	Oral Cyclic Melphalan and Dexamethasone for Patients With AL Amyloidosis. <i>Clinical Lymphoma, Myeloma and Leukemia</i> , 2010, 10, 469-472.	0.2	18
95	Depression and anxiety in patients with AL amyloidosis as assessed by the SF-36 questionnaire: experience in 1226 patients. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2016, 23, 188-193.	1.4	18
96	Treatment patterns and health care resource utilization among patients with relapsed/refractory systemic light chain amyloidosis. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2018, 25, 1-7.	1.4	18
97	Organ responses after highdose melphalan and stemcell transplantation in AL amyloidosis. <i>Leukemia</i> , 2021, 35, 916-919.	3.3	18
98	Immunologic recovery after autologous blood stem cell transplantation in patients with AL-amyloidosis. <i>Bone Marrow Transplantation</i> , 2001, 28, 1105-1109.	1.3	17
99	Amyloidotic Cardiomyopathy: Multidisciplinary Approach to Diagnosis and Treatment. <i>Heart Failure Clinics</i> , 2011, 7, 385-393.	1.0	17
100	Challenges in the management of patients with systemic light chain (AL) amyloidosis during the COVID-19 pandemic. <i>British Journal of Haematology</i> , 2020, 190, 346-357.	1.2	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. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2009, 16, 103-107.	1.4	16
102	High Dose Melphalan and Autologous Peripheral Blood Stem Cell Transplantation in AL Amyloidosis. <i>Hematology/Oncology Clinics of North America</i> , 2014, 28, 1131-1144.	0.9	16
103	Validation of new renal staging system in AL amyloidosis treated with high dose melphalan and stem cell transplantation. <i>American Journal of Hematology</i> , 2016, 91, E458-60.	2.0	16
104	Psychometric validation of the SF-36 Health Survey in light chain amyloidosis: results from community-based and clinic-based samples. <i>Patient Related Outcome Measures</i> , 2017, Volume 8, 157-167.	0.7	16
105	Transbronchial biopsies safely diagnose amyloid lung disease. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2017, 24, 37-41.	1.4	15
106	Orthotopic heart transplant rejection in association with immunomodulatory therapy for AL amyloidosis: A case series and review of the literature. <i>American Journal of Transplantation</i> , 2019, 19, 3185-3190.	2.6	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.. <i>Journal of Clinical Oncology</i> , 2021, 39, 8003-8003.	0.8	15
108	A Phase I Dose-Escalation Study of Carfilzomib in Patients with Previously-Treated Systemic Light-Chain (AL) Amyloidosis. <i>Blood</i> , 2014, 124, 4741-4741.	0.6	15

#	ARTICLE	IF	CITATIONS
109	Short and long-term outcome of treatment with high-dose melphalan and stem cell transplantation for multiple myeloma-associated AL amyloidosis. <i>Annals of Hematology</i> , 2010, 89, 579-584.	0.8	14
110	A second course of high-dose melphalan and auto-SCT for the treatment of relapsed AL amyloidosis. <i>Bone Marrow Transplantation</i> , 2011, 46, 976-980.	1.3	14
111	A new era of amyloidosis: the trends at a major US referral centre. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2019, 26, 192-196.	1.4	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. <i>Biology of Blood and Marrow Transplantation</i> , 2019, 25, e169-e173.	2.0	14
113	Quantitative [18F]florbetapir PET/CT may identify lung involvement in patients with systemic AL amyloidosis. <i>European Journal of Nuclear Medicine and Molecular Imaging</i> , 2020, 47, 1998-2009.	3.3	14
114	Comparing measures of hematologic response after high-dose melphalan and stem cell transplantation in AL amyloidosis. <i>Blood Cancer Journal</i> , 2020, 10, 88.	2.8	14
115	Amyloid Deposits in the Bone Marrow of Patients with Immunoglobulin Light Chain Amyloidosis Do Not Impact Stem Cell Mobilization or Engraftment. <i>Biology of Blood and Marrow Transplantation</i> , 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. <i>Bone Marrow Transplantation</i> , 2017, 52, 1349-1351.	1.3	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. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2018, 25, 148-155.	1.4	13
118	Use of melphalan (M)/dexamethasone (D)/bortezomib in AL amyloidosis. <i>Journal of Clinical Oncology</i> , 2010, 28, 8024-8024.	0.8	13
119	Long-term outcome of patients with monoclonal Ig deposition disease treated with high-dose melphalan and stem cell transplantation. <i>Bone Marrow Transplantation</i> , 2011, 46, 161-162.	1.3	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. <i>Biology of Blood and Marrow Transplantation</i> , 2016, 22, 1729-1732.	2.0	12
121	The six-minute walk test in patients with AL amyloidosis: a single centre case series. <i>British Journal of Haematology</i> , 2017, 177, 388-394.	1.2	12
122	Modified High-Dose Melphalan and Autologous Stem Cell Transplantation for Immunoglobulin Light Chain Amyloidosis. <i>Biology of Blood and Marrow Transplantation</i> , 2018, 24, 1823-1827.	2.0	12
123	Outcomes of patients with AL amyloidosis and low serum free light chain levels at diagnosis. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2018, 25, 156-159.	1.4	12
124	Prevalence and prognostic value of D-dimer elevation in patients with AL amyloidosis. <i>American Journal of Hematology</i> , 2019, 94, 1098-1103.	2.0	12
125	Predictors of hematologic response and survival with stem cell transplantation in AL amyloidosis: A 25-year longitudinal study. <i>American Journal of Hematology</i> , 2022, 97, 1189-1199.	2.0	12
126	Plerixafor-augmented peripheral blood stem cell mobilization in AL amyloidosis with cardiac involvement: a case series. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2014, 21, 149-153.	1.4	11

#	ARTICLE	IF	CITATIONS
127	Bortezomib ocular toxicities: Outcomes with ketotifen. American Journal of Hematology, 2019, 94, E80-E82.	2.0	11
128	The Role of Kidney Transplantation in Monoclonal Ig Deposition Disease. Kidney International Reports, 2020, 5, 485-493.	0.4	11
129	Left Atrial Mechanics Associates With Paroxysmal Atrial Fibrillation in Light-Chain Amyloidosis Following Stem Cell Transplantation. JACC: CardioOncology, 2020, 2, 721-731.	1.7	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.	0.6	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.	1.4	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.	1.4	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.	0.6	10
134	Predictive factors of outcomes in patients with AL amyloidosis treated with daratumumab. American Journal of Hematology, 2022, 97, 79-89.	2.0	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.	1.2	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.	1.4	9
137	The Amyloidosis Forum: a public private partnership to advance drug development in AL amyloidosis. Orphanet Journal of Rare Diseases, 2020, 15, 268.	1.2	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.	1.4	8
139	Treatment Options For Relapsed/refractory Systemic Light-Chain (AL) Amyloidosis: Current Perspectives. Journal of Blood Medicine, 2019, Volume 10, 373-380.	0.7	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.	1.4	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.	0.6	8
142	Phase I/II study of bortezomib (B) in patients with systemic AL-amyloidosis (AL). Journal of Clinical Oncology, 2007, 25, 8050-8050.	0.8	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.	1.4	8
144	Daratumumab in AL amyloidosis. Blood, 2022, 140, 2317-2322.	0.6	8

#	ARTICLE	IF	CITATIONS
145	Systemic AL amyloidosis with an undetectable plasma cell dyscrasia: A zebra without stripes. <i>American Journal of Hematology</i> , 2020, 95, E45-E48.	2.0	7
146	Systemic Amyloidosis Caused by Monoclonal Immunoglobulins. <i>Hematology/Oncology Clinics of North America</i> , 2020, 34, 1099-1113.	0.9	7
147	Predictors and outcomes of acute kidney injury during autologous stem cell transplantation in AL amyloidosis. <i>Nephrology Dialysis Transplantation</i> , 2022, 37, 1281-1288.	0.4	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. <i>Journal of Clinical Oncology</i> , 2020, 38, 8546-8546.	0.8	7
149	Successful Treatment of AL Amyloidosis Patients over Age 65 with High-Dose Melphalan and Autologous Stem Cell Transplantation (HDM/SCT). <i>Blood</i> , 2004, 104, 923-923.	0.6	7
150	High-dose melphalan and autologous stem cell transplantation for AL amyloidosis: recent trends in treatment-related mortality and 1-year survival at a single institution. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2011, 18, 127-129.	1.4	6
151	A solitary mediastinal mass due to localized AL amyloidosis: case report and review of the literature. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2013, 20, 127-130.	1.4	6
152	Hospital admissions following outpatient administration of high-dose melphalan and autologous SCT for AL amyloidosis. <i>Bone Marrow Transplantation</i> , 2014, 49, 1345-1346.	1.3	6
153	Simultaneous presentation of kappa-restricted chronic lymphocytic leukemia and lambda light chain AL amyloidosis. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2014, 21, 124-127.	1.4	6
154	Effect of severe hypoalbuminemia on toxicity of high-dose melphalan and autologous stem cell transplantation in patients with AL amyloidosis. <i>Bone Marrow Transplantation</i> , 2016, 51, 1318-1322.	1.3	6
155	Predictive value of the new renal response criteria in AL amyloidosis treated with high dose melphalan and stem cell transplantation. <i>American Journal of Hematology</i> , 2018, 93, E129-E132.	2.0	6
156	AL Amyloidosis in Myeloma: Red Flag Symptoms. <i>Clinical Lymphoma, Myeloma and Leukemia</i> , 2020, 20, 777-778.	0.2	6
157	Detection of minimal residual disease by next generation sequencing in AL amyloidosis. <i>Blood Cancer Journal</i> , 2021, 11, 117.	2.8	6
158	Role of high-dose melphalan and autologous peripheral blood stem cell transplantation in AL amyloidosis. <i>American Journal of Blood Research</i> , 2012, 2, 9-17.	0.6	6
159	Myocardial Composition in Light-Chain Cardiac Amyloidosis More Than 1 Year After Successful Therapy. <i>JACC: Cardiovascular Imaging</i> , 2022, 15, 594-603.	2.3	6
160	Birtamimab in patients with Mayo stage IV AL amyloidosis: Rationale for confirmatory affirm-AL phase 3 study. <i>Journal of Clinical Oncology</i> , 2022, 40, TPS8076-TPS8076.	0.8	6
161	Risk of second primary malignancy in patients with AL amyloidosis treated with lenalidomide. <i>American Journal of Hematology</i> , 2013, 88, 719-719.	2.0	5
162	Echocardiography and Survival in Light Chain Cardiac Amyloidosis. <i>Circulation: Cardiovascular Imaging</i> , 2018, 11, e007826.	1.3	5

#	ARTICLE	IF	CITATIONS
163	Long term outcome of patients treated on clinical trials of immunomodulatory agents for the treatment of Immunoglobulin light chain (AL) amyloidosis: A pooled analysis. American Journal of Hematology, 2019, 94, E194-E196.	2.0	5
164	Health-Related Quality of Life in Patients with AL Amyloidosis Treated with Daratumumab, Bortezomib, Cyclophosphamide, and Dexamethasone: Results from the Phase 3 Andromeda Study. Blood, 2020, 136, 37-40.	0.6	5
165	Phase 2 Study of Bendamustine in Combination with Dexamethasone (Ben/Dex) in Patients with Previously-Treated Systemic Light Chain (AL) Amyloidosis. Blood, 2014, 124, 3480-3480.	0.6	5
166	Amyloidomics comes of age. Blood, 2012, 119, 1795-1796.	0.6	4
167	Microbiologically documented infections in patients undergoing high-dose melphalan and autologous stem cell transplantation for the treatment of light chain amyloidosis. Transplant Infectious Disease, 2013, 15, 187-194.	0.7	4
168	Single agent lenalidomide three times a week induces hematologic responses in AL amyloidosis patients on dialysis. American Journal of Hematology, 2014, 89, 706-708.	2.0	4
169	Neuralgic amyotrophy following high-dose melphalan and autologous peripheral blood stem cell transplantation for AL amyloidosis. Bone Marrow Transplantation, 2018, 53, 371-373.	1.3	4
170	Heparin-induced thrombocytopenia and thrombosis during high dose melphalan and autologous stem cell transplantation. Blood, 2018, 132, 755-757.	0.6	4
171	High-dose melphalan and autologous peripheral blood stem cell transplantation in patients with AL amyloidosis and cardiac defibrillators. Bone Marrow Transplantation, 2019, 54, 1304-1309.	1.3	4
172	Cardiac biomarkers and health-related quality of life in patients with light chain (<sc>AL</sc>) amyloidosis. British Journal of Haematology, 2019, 185, 998-1001.	1.2	4
173	A pharmacist's review of the treatment of systemic light chain amyloidosis. Journal of Oncology Pharmacy Practice, 2021, 27, 187-198.	0.5	4
174	Clinical Characteristics, Treatment Regimens, and Survival in Elderly Patients with AL Amyloidosis. Clinical Lymphoma, Myeloma and Leukemia, 2021, 21, 425-426.	0.2	4
175	Feasibility of Second Autologous Peripheral Blood Stem Cell (PBSC) Collection Followed by a Second Cycle of High Dose Melphalan (HDM) in Patients Relapsing after an Initial Course of HDM for the Treatment of AL Amyloidosis.. Blood, 2004, 104, 5226-5226.	0.6	4
176	Treatment of AL Amyloidosis with Two Cycles of Induction Therapy with Bortezomib and Dexamethasone Followed by Bortezomib-High Dose Melphalan Conditioning and Autologous Stem Cell Transplantation. Blood, 2012, 120, 2019-2019.	0.6	4
177	Correlation Between 24-Hour Urine Protein and Random Urine Protein-Creatinine Ratio in Amyloid Light-Chain Amyloidosis. Kidney Medicine, 2022, 4, 100427.	1.0	4
178	Penile ulcers complicating systemic AL amyloidosis: a case report. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2016, 23, 203-204.	1.4	3
179	Successful Transition from Bortezomib Subcutaneous (SubQ) to Generic Intravenous (IV) Bortezomib: Cost Savings Initiative with Global Economic Impact. Blood, 2019, 134, 4758-4758.	0.6	3
180	Increases In B-Type Natriuretic Peptide (BNP) During Treatment with Lenalidomide In AL Amyloidosis. Blood, 2010, 116, 3021-3021.	0.6	3

#	ARTICLE	IF	CITATIONS
181	Health-related quality of life in patients with light chain amyloidosis treated with bortezomib, cyclophosphamide, and dexamethasone ± daratumumab: Results from the ANDROMEDA study. American Journal of Hematology, 2022, 97, 719-730.	2.0	3
182	Febrile reactions occurring with second cycle of high-dose melphalan and SCT in patients with AL amyloidosis: a melphalan recall reaction. Bone Marrow Transplantation, 2010, 45, 21-24.	1.3	2
183	REDUCTION IN VENTRICULAR WALL THICKNESS FOLLOWING HIGH-DOSE CHEMOTHERAPY AND STEM-CELL TRANSPLANTATION FOR AL CARDIAC AMYLOIDOSIS. Journal of the American College of Cardiology, 2010, 55, A36.E351.	1.2	2
184	Vertebral compression fractures as the initial presentation of AL amyloidosis: case series and review of literature. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2015, 22, 156-162.	1.4	2
185	Nonoperative Management of Spontaneous Splenic Rupture in a Patient with Light-Chain Amyloidosis: A Case Report. Journal of Vascular and Interventional Radiology, 2015, 26, 1578-1580.	0.2	2
186	High-dose melphalan and stem cell transplantation in AL amyloidosis with elevated cardiac biomarkers. Bone Marrow Transplantation, 2018, 53, 1593-1595.	1.3	2
187	Updated analysis of phase 2 study of bendamustine and dexamethasone in patients with relapsed/refractory systemic light chain (AL) amyloidosis. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2019, 26, 113-114.	1.4	2
188	Safety, Tolerability, and Efficacy of Selinexor in a Patient With Relapsed Light Chain (AL) Amyloidosis. Clinical Lymphoma, Myeloma and Leukemia, 2021, 21, e460-e463.	0.2	2
189	The Use of Next Generation Gene Sequencing to Measure Minimal Residual Disease in Patients with AL Amyloidosis and Low Plasma Cell Burden: A Feasibility Study. Blood, 2019, 134, 4353-4353.	0.6	2
190	Durable Responses to Lenalidomide (Revlimid®) in Patients with AL Amyloidosis: Follow Up of a Phase II Trial. Blood, 2007, 110, 192-192.	0.6	2
191	A Phase I Trial of Pomalidomide, Bortezomib (Velcade), and Dexamethasone (PVD) As Initial Treatment of AL Amyloidosis and Light Chain Deposition Disease. Blood, 2014, 124, 4767-4767.	0.6	2
192	Symptoms of Depression and Anxiety Assessed By the SF-36 Questionnaire in Patients with AL Amyloidosis. Blood, 2015, 126, 3299-3299.	0.6	2
193	Update on the Contemporary Treatment of Light Chain Amyloidosis Including Stem Cell Transplantation. American Journal of Medicine, 2022, 135, S30-S37.	0.6	2
194	A novel substitution of proline (P32L) destabilises β 2-microglobulin inducing hereditary systemic amyloidosis. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2022, , 1-8.	1.4	2
195	Optimal dosing of high-dose melphalan prior to autologous hematopoietic stem cell transplantation in a patient with AL amyloidosis and a solitary kidney. Hematology/ Oncology and Stem Cell Therapy, 2016, 9, 86-88.	0.6	1
196	Evaluation of a new continuous mononuclear cell collection procedure in a single transplant center cohort enriched for AL amyloidosis patients. Transfusion and Apheresis Science, 2018, 57, 411-415.	0.5	1
197	Safety of autologous stem cell transplantation in patients with known Human T-cell Lymphotropic Viruses Type 1 and 2 infection: A case series of four patients. American Journal of Hematology, 2019, 94, E317-E319.	2.0	1
198	Safety and Efficacy of Propylene Glycol-Free Melphalan (Evomela) in Patients with AL Amyloidosis Undergoing Autologous Stem Cell Transplantation: Preliminary Results of a Phase II Study. Blood, 2019, 134, 4578-4578.	0.6	1

#	ARTICLE	IF	CITATIONS
199	Bortezomib and High Dose Melphalan Followed by Autologous Stem Cell Transplantation (BortHDM/SCT) for the Treatment of AL Amyloidosis: Results of a Feasibility Study.. Blood, 2009, 114, 4353-4353.	0.6	1
200	High-Dose Melphalan and Autologous Stem Cell Transplantation In AL Amyloidosis and Monoclonal Immunoglobulin Deposition Disease Associated End-Stage Renal Disease Requiring Dialysis. Blood, 2010, 116, 3553-3553.	0.6	1
201	Macroglossia “ Not Always AL Amyloidosis. Blood, 2010, 116, 5007-5007.	0.6	1
202	Long-Term Outcome Of Patients With AL Amyloidosis Treated With High-Dose Melphalan and Stem Cell Transplantation: 19 Year Experience At a Single Center. Blood, 2013, 122, 3328-3328.	0.6	1
203	The Incidence of Atrial Fibrillation Among Patients with AL Amyloidosis Undergoing High Dose Melphalan and Stem Cell Transplantation (HDM/SCT): Experience at a Single Institution. Blood, 2015, 126, 5490-5490.	0.6	1
204	Final Results of a Phase 2 Study of Bendamustine in Combination with Dexamethasone in Patients with Previously Treated Systemic Light-Chain (AL) Amyloidosis. Blood, 2016, 128, 4523-4523.	0.6	1
205	A phase II trial of lenalidomide for patients with AL amyloidosis. Journal of Clinical Oncology, 2006, 24, 7524-7524.	0.8	1
206	Treatment of AL Amyloidosis with Tandem Cycles of High Dose Melphalan and Autologous Stem Cell Transplantation. , 2004, , 124-126.		1
207	AL (Immunoglobulin Light-Chain) Amyloidosis. , 2008, , 551-569.		1
208	Outcome of Patients with AL Amyloidosis Who Do Not Achieve Hematologic Complete Response After Treatment with High Dose Melphalan and Autologous Transplantation: Results In a Series of 421 Patients. Blood, 2010, 116, 2394-2394.	0.6	1
209	Infectious Complications In Patients with AL (Immunoglobulin Light Chain) Amyloidosis Undergoing Treatment with High-Dose Melphalan and Autologous Stem-Cell Transplantation (HDM/SCT). Blood, 2011, 118, 2041-2041.	0.6	1
210	Modified High-Dose Melphalan and Autologous Stem Cell Transplantation (mHDM/SCT) In the Treatment of AL Amyloidosis (AL) and/or High-Risk Myeloma (hM): Analysis of SWOG Trial S0115. Blood, 2011, 118, 2004-2004.	0.6	1
211	Clinical Presentation and Treatment Responses In IgM-Related AL Amyloidosis. Blood, 2013, 122, 1991-1991.	0.6	1
212	Standard 30-minute Monitoring Time and Less Intensive Pre-medications is Safe in Patients Treated With Subcutaneous Daratumumab for Multiple Myeloma and Light Chain Amyloidosis. Clinical Lymphoma, Myeloma and Leukemia, 2022, , .	0.2	1
213	Serum free light chain trends between orthotopic heart transplantation and auto-SCT in patients with AL amyloidosis. Bone Marrow Transplantation, 2015, 50, 868-869.	1.3	0
214	David C Seldin, MD, PhD: scientist, clinician, teacher, gentleman, 1957“2015. Bone Marrow Transplantation, 2016, 51, 323-323.	1.3	0
215	A Woman in Her 40s With Headache and New-Onset Seizures. JAMA Neurology, 2017, 74, 476.	4.5	0
216	Treatment of AL Amyloidosis with Tandem Cycles of High Dose Melphalan and Autologous Stem Cell Transplantation. , 2004, , 124-126.		0

#	ARTICLE	IF	CITATIONS
217	Quantitative Serum Free Light-Chain Assay in the Diagnostic Evaluation of AL Amyloidosis. , 2004, , 90-92.		0
218	Pulsed Low Dose Intravenous Melphalan in Patients with AL Amyloidosis, Ineligible for Aggressive Treatment with High-Dose Melphalan and Stem Cell Transplantation.. Blood, 2004, 104, 2393-2393.	0.6	0
219	Association of Acquired Von Willebrand Syndrome with Primary (AL) Amyloidosis.. Blood, 2005, 106, 4078-4078.	0.6	0
220	High-Dose Melphalan and Autologous Stem Cell Transplantation in Unusual Non-Amyloid Light Chain Deposition Disorders.. Blood, 2005, 106, 5476-5476.	0.6	0
221	Early Serum Free Light Chain Responses Following High-Dose Melphalan and Stem Cell Transplantation for AL Amyloidosis Predict Treatment Outcomes.. Blood, 2005, 106, 1160-1160.	0.6	0
222	Treatment of AL Amyloidosis with Tandem Cycles of High Dose Melphalan and Autologous Stem Cell Transplantation: Final Analysis of a Prospective Trial.. Blood, 2006, 108, 612-612.	0.6	0
223	Hepatic Response after High-Dose Melphalan and Stem Cell Transplantation for AL Amyloidosis Associated Liver Disease.. Blood, 2007, 110, 2873-2873.	0.6	0
224	Localized AL Amyloidosis of the Breast: A Case Series.. Blood, 2009, 114, 4906-4906.	0.6	0
225	Second Autologous Peripheral Blood Stem Cell Transplantation with High Dose Melphalan (HDM/SCT) in Patients Relapsing After An Initial Course of HDM/SCT for the Treatment of AL Amyloidosis.. Blood, 2009, 114, 2318-2318.	0.6	0
226	Early Serum Free Light Chain Responses Following High-Dose Melphalan and Stem Cell Transplantation for AL Amyloidosis.. Blood, 2009, 114, 4352-4352.	0.6	0
227	Oral Cyclic Melphalan and Dexamethasone in the Treatment of Patients with AL Amyloidosis; Ineligible for High-Dose Melphalan and Stem Cell Transplantation.. Blood, 2009, 114, 1883-1883.	0.6	0
228	Long Term Results of High-Dose Melphalan and Autologous Stem Cell Transplantation in Non-Amyloid Monoclonal Immunoglobulin Deposition Disorders.. Blood, 2009, 114, 4356-4356.	0.6	0
229	Lenalidomide Treatment In Patients with AL Amyloidosis Associated End-Stage Renal Disease and Dialysis. Blood, 2010, 116, 3022-3022.	0.6	0
230	High-Dose Melphalan and Stem Cell Transplantation for Patients with AL Amyloidosis and Cardiac Involvement. Blood, 2011, 118, 2043-2043.	0.6	0
231	Melphalan, Lenalidomide and Dexamethasone Combination Therapy In Patients with AL Amyloidosis. Blood, 2011, 118, 2924-2924.	0.6	0
232	Risk of Second Primary Malignancies in Patients with AL Amyloidosis Treated with Lenalidomide. Blood, 2012, 120, 1873-1873.	0.6	0
233	Lenalidomide and Dexamethasone in the Treatment of AL Amyloidosis: Final Results of A Phase II Trial. Blood, 2012, 120, 4084-4084.	0.6	0
234	Results after long-term follow-up from the CAN2007 phase I/II study of weekly or twice-weekly bortezomib in patients (pts) with relapsed systemic light-chain (AL) amyloidosis.. Journal of Clinical Oncology, 2013, 31, 8545-8545.	0.8	0

#	ARTICLE	IF	CITATIONS
235	Proteasome Inhibitor Based Protocol For Antibody Mediated Rejection In Kidney Transplantation. Blood, 2013, 122, 4728-4728.	0.6	0
236	Pomalidomide and Dexamethasone in Patients with Relapsed AL (Light Chain) Amyloidosis: Results of a Phase 1 Study. Blood, 2014, 124, 3463-3463.	0.6	0
237	Clinical Presentation and Treatment Responses in IgM AL Amyloidosis, a Series of 106 Patients. Blood, 2014, 124, 4750-4750.	0.6	0
238	Heavy/Light Chain Quantification Identifies Clonal Plasma Cell Disease in Patients with AL Amyloidosis and Normal Serum Free Light Chain Ratio. Blood, 2015, 126, 2956-2956.	0.6	0
239	Effect of Severe Hypoalbuminemia on Myelosuppression and Other Toxicities of High Dose Melphalan and Autologous Stem Cell Transplantation in AL Amyloidosis Patients. Blood, 2015, 126, 5499-5499.	0.6	0
240	A Retrospective Review of Engraftment Data for Tbo-Filgrastim Vs. Filgrastim in Patients Undergoing High Dose Chemotherapy and Autologous Stem Cell Transplantation. Blood, 2015, 126, 5484-5484.	0.6	0
241	Induction Therapy with Bortezomib and Dexamethasone and Conditioning with High-Dose Melphalan and Bortezomib Followed By Autologous Stem Cell Transplantation for AL Amyloidosis: Long Term Follow-up Analysis. Blood, 2018, 132, 4616-4616.	0.6	0
242	The Changing Face of Amyloidosis Referrals at a Tertiary Center over the Past 3 Decades. Blood, 2018, 132, 5536-5536.	0.6	0
243	Successful transition from bortezomib subcutaneous to generic intravenous bortezomib: Cost savings initiative with global economic impact.. Journal of Clinical Oncology, 2020, 38, e19375-e19375.	0.8	0
244	Predictive Factors of Overall Survival in Patients with Relapsed AL Amyloidosis Treated with Single Agent Daratumumab. Blood, 2021, 138, 2734-2734.	0.6	0
245	Early serum free light chain response after high-dose melphalan and stem cell transplantation predicts hematologic response in AL amyloidosis. Bone Marrow Transplantation, 2021, , .	1.3	0
246	Beyond Survival in AL amyloidosis: Identifying and Satisfying Patientsâ€™ Needs. Hemato, 2022, 3, 38-46.	0.2	0
247	Racial and Ethnic Disparities in Systemic AL Amyloidosis: Examining Differences in Clinical Presentation and Outcomes. Blood, 2020, 136, 51-51.	0.6	0
248	Amyloidosis Appointment Companion: A Virtual Healthcare Tool to Optimize Shared Decision Making and Improve Patient Experience and Provider Satisfaction for Telehealth and in-Person Appointments. Blood, 2020, 136, 38-39.	0.6	0
249	Incidence of Skin Hyperpigmentation in Black Patients Receiving Treatment with Immunomodulatory Medications. Blood, 2020, 136, 23-24.	0.6	0
250	Modified High Dose Versus High Dose Melphalan Conditioning in Older Patients Undergoing Autologous Stem Cell Transplantation for Immunoglobulin Light Chain Amyloidosis. Blood, 2020, 136, 4-5.	0.6	0
251	AL amyloidosis: who, what, when, why, and where. Oncology, 2012, 26, 164, 166, 169.	0.4	0
252	Prevalence of plasma cell and lymphoproliferative disorders among blood relatives of patients with light chain amyloidosis. British Journal of Haematology, 2022, , .	1.2	0

#	ARTICLE	IF	CITATIONS
253	Differences in the cytogenetic underpinnings of AL amyloidosis among African Americans and Caucasian Americans. Blood Cancer Journal, 2022, 12, .	2.8	0