Vaishali Sanchorawala

List of Publications by Year in descending order

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253 papers 9,725 citations

50276 46 h-index 92 g-index

256 all docs

256 docs citations

256 times ranked

4451 citing authors

#	Article	IF	CITATIONS
1	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
2	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
3	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
4	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
5	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
6	Beyond Survival in AL amyloidosis: Identifying and Satisfying Patients' Needs. Hemato, 2022, 3, 38-46.	0.6	0
7	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
8	Update on the Contemporary Treatment of Light Chain Amyloidosis Including Stem Cell Transplantation. American Journal of Medicine, 2022, 135, S30-S37.	1.5	2
9	Correlation Between 24-Hour Urine Protein and Random Urine Protein-Creatinine Ratio in Amyloid Light-Chain Amyloidosis. Kidney Medicine, 2022, 4, 100427.	2.0	4
10	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
11	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, , .	0.4	1
12	Healthâ€related quality of life in patients with <scp>light chain</scp> amyloidosis treated with bortezomib, cyclophosphamide, and dexamethasone ± daratumumab: Results from the <scp>ANDROMEDA</scp> study. American Journal of Hematology, 2022, 97, 719-730.	4.1	3
13	Myocardial Composition in Light-Chain Cardiac Amyloidosis More Than 1 Year After Successful Therapy. JACC: Cardiovascular Imaging, 2022, 15, 594-603.	5.3	6
14	Prevalence of plasma cell and lymphoproliferative disorders among blood relatives of patients with light chain amyloidosis. British Journal of Haematology, 2022, , .	2.5	0
15	Daratumumab in AL amyloidosis. Blood, 2022, 140, 2317-2322.	1.4	8
16	A novel substitution of proline (P32L) destabilises \hat{I}^2 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.	3.0	2
17	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
18	Birtamimab in patients with Mayo stage IV AL amyloidosis: Rationale for confirmatory affirm-AL phase 3 study Journal of Clinical Oncology, 2022, 40, TPS8076-TPS8076.	1.6	6

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19	Differences in the cytogenetic underpinnings of AL amyloidosis among African Americans and Caucasian Americans. Blood Cancer Journal, 2022, 12, .	6.2	O
20	Organ responses after highdose melphalan and stemcell transplantation in AL amyloidosis. Leukemia, 2021, 35, 916-919.	7.2	18
21	A pharmacist's review of the treatment of systemic light chain amyloidosis. Journal of Oncology Pharmacy Practice, 2021, 27, 187-198.	0.9	4
22	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
23	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
24	Safety, Tolerability, and Efficacy of Selinexor in a Patient With Relapsed Light Chain (AL) Amyloidosis. Clinical Lymphoma, Myeloma and Leukemia, 2021, 21, e460-e463.	0.4	2
25	Detection of minimal residual disease by next generation sequencing in AL amyloidosis. Blood Cancer Journal, 2021, 11, 117.	6.2	6
26	Clinical Characteristics, Treatment Regimens, and Survival in Elderly Patients with AL Amyloidosis. Clinical Lymphoma, Myeloma and Leukemia, 2021, 21, 425-426.	0.4	4
27	Daratumumab-Based Treatment for Immunoglobulin Light-Chain Amyloidosis. New England Journal of Medicine, 2021, 385, 46-58.	27.0	268
28	Marked progress in AL amyloidosis survival: a 40-year longitudinal natural history study. Blood Cancer Journal, 2021, 11, 139.	6.2	45
29	Venetoclax induces deep hematologic remissions in $t(11;14)$ relapsed/refractory AL amyloidosis. Blood Cancer Journal, 2021, 11, 10.	6.2	53
30	Predictive Factors of Overall Survival in Patients with Relapsed AL Amyloidosis Treated with Single Agent Daratumumab. Blood, 2021, 138, 2734-2734.	1.4	0
31	Early serum free light chain response after high-dose melphalan and stem cell transplantation predicts hematologic response in AL amyloidosis. Bone Marrow Transplantation, 2021, , .	2.4	O
32	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
33	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
34	Systemic AL amyloidosis with an undetectable plasma cell dyscrasia: A zebra without stripes. American Journal of Hematology, 2020, 95, E45-E48.	4.1	7
35	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
36	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

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37	Race/ethnicity in systemic AL amyloidosis: perspectives on disease and outcome disparities. Blood Cancer Journal, 2020, 10, 118.	6.2	21
38	Systemic Amyloidosis Caused by Monoclonal Immunoglobulins. Hematology/Oncology Clinics of North America, 2020, 34, 1099-1113.	2.2	7
39	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
40	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
41	Challenges in the management of patients with systemic light chain (AL) amyloidosis during the COVIDâ \in 19 pandemic. British Journal of Haematology, 2020, 190, 346-357.	2.5	17
42	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 the International Society of Amyloidosis, 2020, 27, 244-249.	3.0	27
43	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
44	AL Amyloidosis in Myeloma: Red Flag Symptoms. Clinical Lymphoma, Myeloma and Leukemia, 2020, 20, 777-778.	0.4	6
45	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
46	The Role of Kidney Transplantation in Monoclonal Ig Deposition Disease. Kidney International Reports, 2020, 5, 485-493.	0.8	11
47	High-Dose Melphalan and Autologous Peripheral Blood Stem Cell Transplantation in AL Amyloidosis. Acta Haematologica, 2020, 143, 381-387.	1.4	19
48	Assessment of minimal residual disease using multiparametric flow cytometry in patients with AL amyloidosis. Blood Advances, 2020, 4, 880-884.	5.2	40
49	Left Atrial Mechanics Associates With Paroxysmal Atrial Fibrillation in Light-Chain Amyloidosis Following StemÂCell Transplantation. JACC: CardioOncology, 2020, 2, 721-731.	4.0	11
50	Improved Quantification of CardiacÂAmyloid Burden in SystemicÂLight ChainÂAmyloidosis. JACC: Cardiovascular Imaging, 2020, 13, 1325-1336.	5.3	41
51	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
52	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.	1.4	5
53	Daratumumab plus CyBorD for patients with newly diagnosed AL amyloidosis: safety run-in results of ANDROMEDA. Blood, 2020, 136, 71-80.	1.4	146
54	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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55	Successful transition from bortezomib subcutaneous to generic intravenous bortezomib: Cost savings initiative with global economic impact Journal of Clinical Oncology, 2020, 38, e19375-e19375.	1.6	О
56	Racial and Ethnic Disparities in Systemic AL Amyloidosis: Examining Differences in Clinical Presentation and Outcomes. Blood, 2020, 136, 51-51.	1.4	0
57	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.	1.4	О
58	Incidence of Skin Hyperpigmentation in Black Patients Receiving Treatment with Immunomodulatory Medications. Blood, 2020, 136, 23-24.	1.4	0
59	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.	1.4	О
60	Prevalence and prognostic value of Dâ€dimer elevation in patients with AL amyloidosis. American Journal of Hematology, 2019, 94, 1098-1103.	4.1	12
61	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
62	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.	3.0	2
63	Safety of autologous stem cell transplantation in patients with known Human Tâ€cell Lymphotropic Viruses Type 1 and 2 infection: A case series of four patients. American Journal of Hematology, 2019, 94, E317-E319.	4.1	1
64	<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
65	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.	2.4	4
66	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
67	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.	4.1	5
68	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
69	Delay treatment of AL amyloidosis at relapse until symptomatic: devil is in the details. Blood Advances, 2019, 3, 216-218.	5.2	25
70	Bortezomib ocular toxicities: Outcomes with ketotifen. American Journal of Hematology, 2019, 94, E80-E82.	4.1	11
71	Long-term outcome of kidney transplantation in AL amyloidosis. Kidney International, 2019, 95, 405-411.	5.2	57
72	Cardiac biomarkers and healthâ€related quality of life in patients with light chain (<scp>AL</scp>) amyloidosis. British Journal of Haematology, 2019, 185, 998-1001.	2.5	4

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73	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
74	Development and validation of a survival staging system incorporating BNP in patients with light chain amyloidosis. Blood, 2019, 133, 215-223.	1.4	118
75	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
76	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
77	Successful Transition from Bortezomib Subcutaneous (SubQ) to Generic Intravenous (IV) Bortezomib: Cost Savings Initiative with Global Economic Impact. Blood, 2019, 134, 4758-4758.	1.4	3
78	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.	1.4	1
79	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.	1.4	2
80	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
81	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
82	Predictive value of the new renal response criteria in AL amyloidosis treated with high dose melphalan and stem cell transplantation. American Journal of Hematology, 2018, 93, E129-E132.	4.1	6
83	Neuralgic amyotrophy following high-dose melphalan and autologous peripheral blood stem cell transplantation for AL amyloidosis. Bone Marrow Transplantation, 2018, 53, 371-373.	2.4	4
84	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
85	Systemic immunoglobulin light chain amyloidosis. Nature Reviews Disease Primers, 2018, 4, 38.	30.5	350
86	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.	1.0	1
87	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
88	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
89	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
90	Echocardiography and Survival in Light Chain Cardiac Amyloidosis. Circulation: Cardiovascular Imaging, 2018, 11, e007826.	2.6	5

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91	Heparin-induced thrombocytopenia and thrombosis during high dose melphalan and autologous stem cell transplantation. Blood, 2018, 132, 755-757.	1.4	4
92	High-dose melphalan and stem cell transplantation in AL amyloidosis with elevated cardiac biomarkers. Bone Marrow Transplantation, 2018, 53, 1593-1595.	2.4	2
93	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
94	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
95	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.	1.4	O
96	The Changing Face of Amyloidosis Referrals at a Tertiary Center over the Past 3 Decades. Blood, 2018, 132, 5536-5536.	1.4	0
97	A Woman in Her 40s With Headache and New-Onset Seizures. JAMA Neurology, 2017, 74, 476.	9.0	O
98	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
99	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
100	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
101	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
102	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
103	Hematologic relapse in AL amyloidosis after high-dose melphalan and stem cell transplantation. Blood, 2017, 130, 1383-1386.	1.4	30
104	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
105	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
106	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.	3.0	3
107	Effect of severe hypoalbuminemia on toxicity of high-dose melphalan and autologous stem cell transplantation in patients with AL amyloidosis. Bone Marrow Transplantation, 2016, 51, 1318-1322.	2.4	6
108	David C Seldin, MD, PhD: scientist, clinician, teacher, gentleman, 1957–2015. Bone Marrow Transplantation, 2016, 51, 323-323.	2.4	0

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109	Risk factors for venous thromboembolism in immunoglobulin light chain amyloidosis. Haematologica, 2016, 101, 86-90.	3.5	19
110	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
111	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
112	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
113	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
114	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
115	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
116	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.	0.9	1
117	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.	1.4	1
118	Safety and Efficacy of Carfilzomib (CFZ) in Previously-Treated Systemic Light-Chain (AL) Amyloidosis. Blood, 2016, 128, 645-645.	1.4	46
119	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
120	Serum free light chain trends between orthotopic heart transplantation and auto-SCT in patients with AL amyloidosis. Bone Marrow Transplantation, 2015, 50, 868-869.	2.4	0
121	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
122	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.	3.0	2
123	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
124	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.	0.5	2
125	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.	1.4	1
126	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.	1.4	0

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127	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.	1.4	O
128	Symptoms of Depression and Anxiety Assessed By the SF-36 Questionnaire in Patients with AL Amyloidosis. Blood, 2015, 126, 3299-3299.	1.4	2
129	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.	1.4	0
130	Hospital admissions following outpatient administration of high-dose melphalan and autologous SCT for AL amyloidosis. Bone Marrow Transplantation, 2014, 49, 1345-1346.	2.4	6
131	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
132	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
133	Simultaneous presentation of kappa-restricted chronic lymphocytic leukemia and lambda light chain AL amyloidosis. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2014, 21, 124-127.	3.0	6
134	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
135	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
136	Update on treatment of light chain amyloidosis. Haematologica, 2014, 99, 209-221.	3.5	93
137	Single agent lenalidomide three times a week induces hematologic responses in AL amyloidosis patients on dialysis. American Journal of Hematology, 2014, 89, 706-708.	4.1	4
138	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
139	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
140	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.	1.4	5
141	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
142	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.	1.4	2
143	Pomalidomide and Dexamethasone in Patients with Relapsed AL (Light Chain) Amyloidosis: Results of a Phase 1 Study. Blood, 2014, 124, 3463-3463.	1.4	0
144	Clinical Presentation and Treatment Responses in IgM AL Amyloidosis, a Series of 106 Patients. Blood, 2014, 124, 4750-4750.	1.4	0

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