

Vaishali Sanchorawala

List of Publications by Year in descending order

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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	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.	1.4	22
2	Predictors and outcomes of acute kidney injury during autologous stem cell transplantation in AL amyloidosis. Nephrology Dialysis Transplantation, 2022, 37, 1281-1288.	0.4	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.	3.3	29
4	Predictive factors of outcomes in patients with AL amyloidosis treated with daratumumab. American Journal of Hematology, 2022, 97, 79-89.	2.0	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.	1.4	42
6	Beyond Survival in AL amyloidosis: Identifying and Satisfying Patients' Needs. Hemato, 2022, 3, 38-46.	0.2	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.	1.2	10
8	Update on the Contemporary Treatment of Light Chain Amyloidosis Including Stem Cell Transplantation. American Journal of Medicine, 2022, 135, S30-S37.	0.6	2
9	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
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.	1.4	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, , .	0.2	1
12	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
13	Myocardial Composition in Light-Chain Cardiac Amyloidosis More Than 1 Year After Successful Therapy. JACC: Cardiovascular Imaging, 2022, 15, 594-603.	2.3	6
14	Prevalence of plasma cell and lymphoproliferative disorders among blood relatives of patients with light chain amyloidosis. British Journal of Haematology, 2022, , .	1.2	0
15	Daratumumab in AL amyloidosis. Blood, 2022, 140, 2317-2322.	0.6	8
16	A novel substitution of proline (P32L) destabilises Î2-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
17	Predictors of hematologic response and survival with stem cell transplantation in AL amyloidosis: A 25-year longitudinal study. American Journal of Hematology, 2022, 97, 1189-1199.	2.0	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.	0.8	6

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19	Differences in the cytogenetic underpinnings of AL amyloidosis among African Americans and Caucasian Americans. <i>Blood Cancer Journal</i> , 2022, 12, .	2.8	0
20	Organ responses after highdose melphalan and stemcell transplantation in AL amyloidosis. <i>Leukemia</i> , 2021, 35, 916-919.	3.3	18
21	A pharmacist's review of the treatment of systemic light chain amyloidosis. <i>Journal of Oncology Pharmacy Practice</i> , 2021, 27, 187-198.	0.5	4
22	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
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.. <i>Journal of Clinical Oncology</i> , 2021, 39, 8003-8003.	0.8	15
24	Safety, Tolerability, and Efficacy of Selinexor in a Patient With Relapsed Light Chain (AL) Amyloidosis. <i>Clinical Lymphoma, Myeloma and Leukemia</i> , 2021, 21, e460-e463.	0.2	2
25	Detection of minimal residual disease by next generation sequencing in AL amyloidosis. <i>Blood Cancer Journal</i> , 2021, 11, 117.	2.8	6
26	Clinical Characteristics, Treatment Regimens, and Survival in Elderly Patients with AL Amyloidosis. <i>Clinical Lymphoma, Myeloma and Leukemia</i> , 2021, 21, 425-426.	0.2	4
27	Daratumumab-Based Treatment for Immunoglobulin Light-Chain Amyloidosis. <i>New England Journal of Medicine</i> , 2021, 385, 46-58.	13.9	268
28	Marked progress in AL amyloidosis survival: a 40-year longitudinal natural history study. <i>Blood Cancer Journal</i> , 2021, 11, 139.	2.8	45
29	Venetoclax induces deep hematologic remissions in t(11;14) relapsed/refractory AL amyloidosis. <i>Blood Cancer Journal</i> , 2021, 11, 10.	2.8	53
30	Predictive Factors of Overall Survival in Patients with Relapsed AL Amyloidosis Treated with Single Agent Daratumumab. <i>Blood</i> , 2021, 138, 2734-2734.	0.6	0
31	Early serum free light chain response after high-dose melphalan and stem cell transplantation predicts hematologic response in AL amyloidosis. <i>Bone Marrow Transplantation</i> , 2021, , .	1.3	0
32	The utility of repeat kidney biopsy in systemic immunoglobulin light chain amyloidosis. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2020, 27, 17-24.	1.4	8
33	Establishment of brain natriuretic peptide ϵ -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
34	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
35	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
36	The Amyloidosis Forum: a public private partnership to advance drug development in AL amyloidosis. <i>Orphanet Journal of Rare Diseases</i> , 2020, 15, 268.	1.2	9

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37	Race/ethnicity in systemic AL amyloidosis: perspectives on disease and outcome disparities. <i>Blood Cancer Journal</i> , 2020, 10, 118.	2.8	21
38	Systemic Amyloidosis Caused by Monoclonal Immunoglobulins. <i>Hematology/Oncology Clinics of North America</i> , 2020, 34, 1099-1113.	0.9	7
39	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
40	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
41	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
42	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
43	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
44	AL Amyloidosis in Myeloma: Red Flag Symptoms. <i>Clinical Lymphoma, Myeloma and Leukemia</i> , 2020, 20, 777-778.	0.2	6
45	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
46	The Role of Kidney Transplantation in Monoclonal Ig Deposition Disease. <i>Kidney International Reports</i> , 2020, 5, 485-493.	0.4	11
47	High-Dose Melphalan and Autologous Peripheral Blood Stem Cell Transplantation in AL Amyloidosis. <i>Acta Haematologica</i> , 2020, 143, 381-387.	0.7	19
48	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
49	Left Atrial Mechanics Associates With Paroxysmal Atrial Fibrillation in Light-Chain Amyloidosis Following Stem Cell Transplantation. <i>JACC: CardioOncology</i> , 2020, 2, 721-731.	1.7	11
50	Improved Quantification of Cardiac Amyloid Burden in Systemic Light Chain Amyloidosis. <i>JACC: Cardiovascular Imaging</i> , 2020, 13, 1325-1336.	2.3	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. <i>Blood</i> , 2020, 136, 48-50.	0.6	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. <i>Blood</i> , 2020, 136, 37-40.	0.6	5
53	Daratumumab plus CyBorD for patients with newly diagnosed AL amyloidosis: safety run-in results of ANDROMEDA. <i>Blood</i> , 2020, 136, 71-80.	0.6	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. <i>Journal of Clinical Oncology</i> , 2020, 38, 8546-8546.	0.8	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.	0.8	0
56	Racial and Ethnic Disparities in Systemic AL Amyloidosis: Examining Differences in Clinical Presentation and Outcomes. Blood, 2020, 136, 51-51.	0.6	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.	0.6	0
58	Incidence of Skin Hyperpigmentation in Black Patients Receiving Treatment with Immunomodulatory Medications. Blood, 2020, 136, 23-24.	0.6	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.	0.6	0
60	Prevalence and prognostic value of D-dimer elevation in patients with AL amyloidosis. American Journal of Hematology, 2019, 94, 1098-1103.	2.0	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.	1.4	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.	1.4	2
63	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
64	<p>Treatment Options For Relapsed/refractory Systemic Light-Chain (AL) Amyloidosis: Current Perspectives</p>. Journal of Blood Medicine, 2019, Volume 10, 373-380.	0.7	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.	1.3	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.	2.6	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.	2.0	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.	2.8	54
69	Delay treatment of AL amyloidosis at relapse until symptomatic: devil is in the details. Blood Advances, 2019, 3, 216-218.	2.5	25
70	Bortezomib ocular toxicities: Outcomes with ketotifen. American Journal of Hematology, 2019, 94, E80-E82.	2.0	11
71	Long-term outcome of kidney transplantation in AL amyloidosis. Kidney International, 2019, 95, 405-411.	2.6	57
72	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

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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. <i>Biology of Blood and Marrow Transplantation</i> , 2019, 25, e169-e173.	2.0	14
74	Development and validation of a survival staging system incorporating BNP in patients with light chain amyloidosis. <i>Blood</i> , 2019, 133, 215-223.	0.6	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). <i>Blood</i> , 2019, 134, 139-139.	0.6	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. <i>Blood</i> , 2019, 134, 3166-3166.	0.6	27
77	Successful Transition from Bortezomib Subcutaneous (SubQ) to Generic Intravenous (IV) Bortezomib: Cost Savings Initiative with Global Economic Impact. <i>Blood</i> , 2019, 134, 4758-4758.	0.6	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. <i>Blood</i> , 2019, 134, 4578-4578.	0.6	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. <i>Blood</i> , 2019, 134, 4353-4353.	0.6	2
80	Once AL amyloidosis: not always AL amyloidosis. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2018, 25, 139-140.	1.4	10
81	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
82	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
83	Neuralgic amyotrophy following high-dose melphalan and autologous peripheral blood stem cell transplantation for AL amyloidosis. <i>Bone Marrow Transplantation</i> , 2018, 53, 371-373.	1.3	4
84	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
85	Systemic immunoglobulin light chain amyloidosis. <i>Nature Reviews Disease Primers</i> , 2018, 4, 38.	18.1	350
86	Evaluation of a new continuous mononuclear cell collection procedure in a single transplant center cohort enriched for AL amyloidosis patients. <i>Transfusion and Apheresis Science</i> , 2018, 57, 411-415.	0.5	1
87	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
88	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
89	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
90	Echocardiography and Survival in Light Chain Cardiac Amyloidosis. <i>Circulation: Cardiovascular Imaging</i> , 2018, 11, e007826.	1.3	5

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91	Heparin-induced thrombocytopenia and thrombosis during high dose melphalan and autologous stem cell transplantation. <i>Blood</i> , 2018, 132, 755-757.	0.6	4
92	High-dose melphalan and stem cell transplantation in AL amyloidosis with elevated cardiac biomarkers. <i>Bone Marrow Transplantation</i> , 2018, 53, 1593-1595.	1.3	2
93	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
94	Safety, Tolerability and Response Rates of Daratumumab in Patients with Relapsed Light Chain (AL) Amyloidosis: Results of a Phase II Study. <i>Blood</i> , 2018, 132, 2005-2005.	0.6	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. <i>Blood</i> , 2018, 132, 4616-4616.	0.6	0
96	The Changing Face of Amyloidosis Referrals at a Tertiary Center over the Past 3 Decades. <i>Blood</i> , 2018, 132, 5536-5536.	0.6	0
97	A Woman in Her 40s With Headache and New-Onset Seizures. <i>JAMA Neurology</i> , 2017, 74, 476.	4.5	0
98	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
99	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
100	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
101	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
102	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
103	Hematologic relapse in AL amyloidosis after high-dose melphalan and stem cell transplantation. <i>Blood</i> , 2017, 130, 1383-1386.	0.6	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. <i>Bone Marrow Transplantation</i> , 2017, 52, 1349-1351.	1.3	13
105	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
106	Penile ulcers complicating systemic AL amyloidosis: a case report. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2016, 23, 203-204.	1.4	3
107	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
108	David C Seldin, MD, PhD: scientist, clinician, teacher, gentleman, 1957-2015. <i>Bone Marrow Transplantation</i> , 2016, 51, 323-323.	1.3	0

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109	Risk factors for venous thromboembolism in immunoglobulin light chain amyloidosis. <i>Haematologica</i> , 2016, 101, 86-90.	1.7	19
110	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
111	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
112	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
113	Immunoglobulin heavy light chain test quantifies clonal disease in patients with AL amyloidosis and normal serum free light chain ratio. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2016, 23, 214-220.	1.4	8
114	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
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. <i>Biology of Blood and Marrow Transplantation</i> , 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. <i>Hematology/ Oncology and Stem Cell Therapy</i> , 2016, 9, 86-88.	0.6	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. <i>Blood</i> , 2016, 128, 4523-4523.	0.6	1
118	Safety and Efficacy of Carfilzomib (CFZ) in Previously-Treated Systemic Light-Chain (AL) Amyloidosis. <i>Blood</i> , 2016, 128, 645-645.	0.6	46
119	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
120	Serum free light chain trends between orthotopic heart transplantation and auto-SCT in patients with AL amyloidosis. <i>Bone Marrow Transplantation</i> , 2015, 50, 868-869.	1.3	0
121	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
122	Vertebral compression fractures as the initial presentation of AL amyloidosis: case series and review of literature. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2015, 22, 156-162.	1.4	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. <i>Biology of Blood and Marrow Transplantation</i> , 2015, 21, 1445-1451.	2.0	55
124	Nonoperative Management of Spontaneous Splenic Rupture in a Patient with Light-Chain Amyloidosis: A Case Report. <i>Journal of Vascular and Interventional Radiology</i> , 2015, 26, 1578-1580.	0.2	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. <i>Blood</i> , 2015, 126, 5490-5490.	0.6	1
126	Heavy/Light Chain Quantification Identifies Clonal Plasma Cell Disease in Patients with AL Amyloidosis and Normal Serum Free Light Chain Ratio. <i>Blood</i> , 2015, 126, 2956-2956.	0.6	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. <i>Blood</i> , 2015, 126, 5499-5499.	0.6	0
128	Symptoms of Depression and Anxiety Assessed By the SF-36 Questionnaire in Patients with AL Amyloidosis. <i>Blood</i> , 2015, 126, 3299-3299.	0.6	2
129	A Retrospective Review of Engraftment Data for Tbo-Filgrastim Vs. Filgrastim in Patients Undergoing High Dose Chemotherapy and Autologous Stem Cell Transplantation. <i>Blood</i> , 2015, 126, 5484-5484.	0.6	0
130	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
131	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
132	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
133	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
134	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
135	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
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