

# Yoshihide Asano

## List of Publications by Year in descending order

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Version: 2024-02-01

218  
papers

6,149  
citations

71061

41  
h-index

98753

67  
g-index

228  
all docs

228  
docs citations

228  
times ranked

5670  
citing authors

#	ARTICLE	IF	CITATIONS
1	Tocilizumab in systemic sclerosis: a randomised, double-blind, placebo-controlled, phase 3 trial. <i>Lancet Respiratory Medicine</i> , 2020, 8, 963-974.	5.2	348
2	Clinical Correlations With Dermatomyositis-Specific Autoantibodies in Adult Japanese Patients With Dermatomyositis. <i>Archives of Dermatology</i> , 2011, 147, 391.	1.7	293
3	Increased Expression of Integrin $\alpha 3 \beta 1$ Contributes to the Establishment of Autocrine TGF- $\beta 2$ Signaling in Scleroderma Fibroblasts. <i>Journal of Immunology</i> , 2005, 175, 7708-7718.	0.4	207
4	Impaired Smad7-Smurf1-mediated negative regulation of TGF- $\beta 2$ signaling in scleroderma fibroblasts. <i>Journal of Clinical Investigation</i> , 2004, 113, 253-264.	3.9	181
5	Endothelial Fli1 Deficiency Impairs Vascular Homeostasis. <i>American Journal of Pathology</i> , 2010, 176, 1983-1998.	1.9	178
6	Increased Expression of Integrin $\alpha 5 \beta 1$ Induces the Myofibroblastic Differentiation of Dermal Fibroblasts. <i>American Journal of Pathology</i> , 2006, 168, 499-510.	1.9	159
7	Vasculopathy in scleroderma. <i>Seminars in Immunopathology</i> , 2015, 37, 489-500.	2.8	143
8	Simultaneous downregulation of KLF5 and Fli1 is a key feature underlying systemic sclerosis. <i>Nature Communications</i> , 2014, 5, 5797.	5.8	120
9	Naturally Occurring Antibodies in Humans Can Neutralize a Variety of Influenza Virus Strains, Including H3, H1, H2, and H5. <i>Journal of Virology</i> , 2011, 85, 11048-11057.	1.5	102
10	Future treatments in systemic sclerosis. <i>Journal of Dermatology</i> , 2010, 37, 54-70.	0.6	98
11	Clinical significance of surfactant protein D as a serum marker for evaluating pulmonary fibrosis in patients with systemic sclerosis. <i>Arthritis and Rheumatism</i> , 2001, 44, 1363-1369.	6.7	92
12	Systemic sclerosis. <i>Journal of Dermatology</i> , 2018, 45, 128-138.	0.6	92
13	Transforming Growth Factor- $\beta 2$ Regulates DNA Binding Activity of Transcription Factor Fli1 by p300/CREB-binding Protein-associated Factor-dependent Acetylation. <i>Journal of Biological Chemistry</i> , 2007, 282, 34672-34683.	1.6	87
14	Phosphatidylinositol 3-Kinase Is Involved in $\alpha 2(I)$ Collagen Gene Expression in Normal and Scleroderma Fibroblasts. <i>Journal of Immunology</i> , 2004, 172, 7123-7135.	0.4	86
15	Immunization with DNA topoisomerase I and Freund's complete adjuvant induces skin and lung fibrosis and autoimmunity via interleukin-6 signaling. <i>Arthritis and Rheumatism</i> , 2011, 63, 3575-3585.	6.7	81
16	Involvement of $\alpha 5 \beta 1$ Integrin in the Establishment of Autocrine TGF- $\beta 2$ Signaling in Dermal Fibroblasts Derived from Localized Scleroderma. <i>Journal of Investigative Dermatology</i> , 2006, 126, 1761-1769.	0.3	80
17	Constitutive Thrombospondin-1 Overexpression Contributes to Autocrine Transforming Growth Factor- $\beta 2$ Signaling in Cultured Scleroderma Fibroblasts. <i>American Journal of Pathology</i> , 2005, 166, 1451-1463.	1.9	79
18	TLR4, rather than TLR2, regulates wound healing through TGF- $\beta 2$ and CCL5 expression. <i>Journal of Dermatological Science</i> , 2014, 73, 117-124.	1.0	75

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19	Fibrosis, Vascular Activation, and Immune Abnormalities Resembling Systemic Sclerosis in Bleomycin-Treated Fli1-Haploinsufficient Mice. <i>Arthritis and Rheumatology</i> , 2015, 67, 517-526.	2.9	75
20	Increased Expression Levels of Integrin $\alpha 2 \beta 5$ on Scleroderma Fibroblasts. <i>American Journal of Pathology</i> , 2004, 164, 1275-1292.	1.9	73
21	Transcription Factor Fli1 Regulates Collagen Fibrillogenesis in Mouse Skin. <i>Molecular and Cellular Biology</i> , 2009, 29, 425-434.	1.1	69
22	Epithelial Fli1 deficiency drives systemic autoimmunity and fibrosis: Possible roles in scleroderma. <i>Journal of Experimental Medicine</i> , 2017, 214, 1129-1151.	4.2	69
23	The impact of Fli1 deficiency on the pathogenesis of systemic sclerosis. <i>Journal of Dermatological Science</i> , 2010, 59, 153-162.	1.0	68
24	Adiponectin is an endogenous anti-fibrotic mediator and therapeutic target. <i>Scientific Reports</i> , 2017, 7, 4397.	1.6	64
25	Review: Frontiers of Antifibrotic Therapy in Systemic Sclerosis. <i>Arthritis and Rheumatology</i> , 2017, 69, 257-267.	2.9	62
26	Phosphorylation of Fli1 at Threonine 312 by Protein Kinase C $\delta$ Promotes Its Interaction with p300/CREB-Binding Protein-Associated Factor and Subsequent Acetylation in Response to Transforming Growth Factor $\beta 2$ . <i>Molecular and Cellular Biology</i> , 2009, 29, 1882-1894.	1.1	61
27	Amelioration of Tissue Fibrosis by Toll-like Receptor 4 Knockout in Murine Models of Systemic Sclerosis. <i>Arthritis and Rheumatology</i> , 2015, 67, 254-265.	2.9	57
28	The Pathogenesis of Systemic Sclerosis: An Understanding Based on a Common Pathologic Cascade across Multiple Organs and Additional Organ-Specific Pathologies. <i>Journal of Clinical Medicine</i> , 2020, 9, 2687.	1.0	57
29	Serum Levels of Galectin-3: Possible Association with Fibrosis, Aberrant Angiogenesis, and Immune Activation in Patients with Systemic Sclerosis. <i>Journal of Rheumatology</i> , 2012, 39, 539-544.	1.0	54
30	Serum adiponectin levels inversely correlate with the activity of progressive skin sclerosis in patients with diffuse cutaneous systemic sclerosis. <i>Journal of the European Academy of Dermatology and Venereology</i> , 2012, 26, 354-360.	1.3	53
31	Growth of clonogenic myeloblastic leukemic cells in the presence of human recombinant erythropoietin in addition to various human recombinant hematopoietic growth factors. <i>Blood</i> , 1988, 72, 1682-1686.	0.6	52
32	The Role of IL-32 in Cutaneous T-Cell Lymphoma. <i>Journal of Investigative Dermatology</i> , 2014, 134, 1428-1435.	0.3	52
33	Effect of (-)-epigallocatechin gallate on leukemic blast cells from patients with acute myeloblastic leukemia. <i>Life Sciences</i> , 1996, 60, 135-142.	2.0	51
34	Diagnostic criteria, severity classification and guidelines of localized scleroderma. <i>Journal of Dermatology</i> , 2018, 45, 755-780.	0.6	51
35	Diagnostic criteria, severity classification and guidelines of eosinophilic fasciitis. <i>Journal of Dermatology</i> , 2018, 45, 881-890.	0.6	50
36	Constitutively phosphorylated Smad3 interacts with Sp1 and p300 in scleroderma fibroblasts. <i>Rheumatology</i> , 2006, 45, 157-165.	0.9	48

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37	The wound/burn guidelines "6: Guidelines for the management of burns. <i>Journal of Dermatology</i> , 2016, 43, 989-1010.	0.6	48
38	Skin Barrier Dysfunction and Low Antimicrobial Peptide Expression in Cutaneous T-cell Lymphoma. <i>Clinical Cancer Research</i> , 2014, 20, 4339-4348.	3.2	47
39	Multifaceted contribution of the TLR4-activated IRF5 transcription factor in systemic sclerosis. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2015, 112, 15136-15141.	3.3	47
40	CXCL17 Attenuates Imiquimod-Induced Psoriasis-like Skin Inflammation by Recruiting Myeloid-Derived Suppressor Cells and Regulatory T Cells. <i>Journal of Immunology</i> , 2017, 198, 3897-3908.	0.4	47
41	Rituximab therapy is more effective than cyclophosphamide therapy for Japanese patients with anti- $\alpha$ -topoisomerase $\beta$ -positive systemic sclerosis-associated interstitial lung disease. <i>Journal of Dermatology</i> , 2019, 46, 1006-1013.	0.6	47
42	Fli1 deficiency contributes to the suppression of endothelial CXCL5 expression in systemic sclerosis. <i>Archives of Dermatological Research</i> , 2014, 306, 331-338.	1.1	45
43	Serum levels of tissue inhibitor of metalloproteinase-1 and 2 in patients with eosinophilic fasciitis. <i>British Journal of Dermatology</i> , 2004, 151, 407-412.	1.4	43
44	TBX4 is involved in the super-enhancer-driven transcriptional programs underlying features specific to lung fibroblasts. <i>American Journal of Physiology - Lung Cellular and Molecular Physiology</i> , 2018, 314, L177-L191.	1.3	42
45	Decreased cathepsin V expression due to Fli1 deficiency contributes to the development of dermal fibrosis and proliferative vasculopathy in systemic sclerosis. <i>Rheumatology</i> , 2013, 52, 790-799.	0.9	41
46	CXCL13 produced by macrophages due to Fli1 deficiency may contribute to the development of tissue fibrosis, vasculopathy and immune activation in systemic sclerosis. <i>Experimental Dermatology</i> , 2018, 27, 1030-1037.	1.4	41
47	Increased expression of chemerin in endothelial cells due to Fli1 deficiency may contribute to the development of digital ulcers in systemic sclerosis. <i>Rheumatology</i> , 2015, 54, 1308-1316.	0.9	40
48	An orally-active adiponectin receptor agonist mitigates cutaneous fibrosis, inflammation and microvascular pathology in a murine model of systemic sclerosis. <i>Scientific Reports</i> , 2018, 8, 11843.	1.6	39
49	A possible contribution of endothelial $\alpha$ 1 downregulation due to Fli1 deficiency to the development of digital ulcers in systemic sclerosis. <i>Experimental Dermatology</i> , 2015, 24, 127-132.	1.4	38
50	Serum Adhesion Molecule Levels as Prognostic Markers in Patients with Early Systemic Sclerosis: A Multicentre, Prospective, Observational Study. <i>PLoS ONE</i> , 2014, 9, e88150.	1.1	38
51	Interleukin-31 promotes fibrosis and T helper 2 polarization in systemic sclerosis. <i>Nature Communications</i> , 2021, 12, 5947.	5.8	38
52	Clinical significance of serum growth differentiation factor-15 levels in systemic sclerosis: association with disease severity. <i>Modern Rheumatology</i> , 2012, 22, 668-675.	0.9	37
53	A Possible Contribution of Altered Cathepsin B Expression to the Development of Skin Sclerosis and Vasculopathy in Systemic Sclerosis. <i>PLoS ONE</i> , 2012, 7, e32272.	1.1	36
54	Diagnostic criteria, severity classification and guidelines of systemic sclerosis. <i>Journal of Dermatology</i> , 2018, 45, 633-691.	0.6	35

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55	Association of anti-RNA polymerase III antibody and malignancy in Japanese patients with systemic sclerosis. <i>Journal of Dermatology</i> , 2015, 42, 524-527.	0.6	34
56	A potential contribution of antimicrobial peptide LL-37 to tissue fibrosis and vasculopathy in systemic sclerosis. <i>British Journal of Dermatology</i> , 2016, 175, 1195-1203.	1.4	33
57	Bosentan reverses the pro-fibrotic phenotype of systemic sclerosis dermal fibroblasts via increasing DNA binding ability of transcription factor Fli1. <i>Arthritis Research and Therapy</i> , 2014, 16, R86.	1.6	31
58	A possible contribution of lipocalin-2 to the development of dermal fibrosis, pulmonary vascular involvement and renal dysfunction in systemic sclerosis. <i>British Journal of Dermatology</i> , 2015, 173, 681-689.	1.4	31
59	Systemic Sclerosis Dermal Fibroblasts Suppress Th1 Cytokine Production via Galectin-9 Overproduction due to Fli1 Deficiency. <i>Journal of Investigative Dermatology</i> , 2017, 137, 1850-1859.	0.3	31
60	The Prevalence and Clinical Significance of Anti-U1 RNA Antibodies in Patients with Systemic Sclerosis. <i>Journal of Investigative Dermatology</i> , 2003, 120, 204-210.	0.3	30
61	Low Herpesvirus Entry Mediator (HVEM) Expression on Dermal Fibroblasts Contributes to a Th2-Dominant Microenvironment in Advanced Cutaneous T-Cell Lymphoma. <i>Journal of Investigative Dermatology</i> , 2012, 132, 1280-1289.	0.3	30
62	Histological features of localized scleroderma <i>en coup de sabre</i> : a study of 16 cases. <i>Journal of the European Academy of Dermatology and Venereology</i> , 2014, 28, 1805-1810.	1.3	30
63	Epigenetic suppression of Fli1, a potential predisposing factor in the pathogenesis of systemic sclerosis. <i>International Journal of Biochemistry and Cell Biology</i> , 2015, 67, 86-91.	1.2	30
64	Fli1 deficiency contributes to the downregulation of endothelial protein C receptor in systemic sclerosis: a possible role in prothrombotic conditions. <i>British Journal of Dermatology</i> , 2016, 174, 338-347.	1.4	29
65	Skin thickness score as a surrogate marker of organ involvements in systemic sclerosis: a retrospective observational study. <i>Arthritis Research and Therapy</i> , 2019, 21, 129.	1.6	29
66	Serum chemokine levels as prognostic markers in patients with early systemic sclerosis: a multicenter, prospective, observational study. <i>Modern Rheumatology</i> , 2013, 23, 1076-1084.	0.9	28
67	Progranulin Overproduction Due to Fli1 Deficiency Contributes to the Resistance of Dermal Fibroblasts to Tumor Necrosis Factor in Systemic Sclerosis. <i>Arthritis and Rheumatology</i> , 2015, 67, 3245-3255.	2.9	28
68	A possible contribution of visfatin to the resolution of skin sclerosis in patients with diffuse cutaneous systemic sclerosis via a direct anti-fibrotic effect on dermal fibroblasts and Th1 polarization of the immune response. <i>Rheumatology</i> , 2013, 52, 1239-1244.	0.9	27
69	Clinical correlation of brachial artery flow-mediated dilation in patients with systemic sclerosis. <i>Modern Rheumatology</i> , 2014, 24, 106-111.	0.9	27
70	Plasma plasmin-alpha2-plasmin inhibitor complex levels are increased in systemic sclerosis patients with pulmonary hypertension. <i>British Journal of Rheumatology</i> , 2003, 42, 240-243.	2.5	26
71	High-dose intravenous immunoglobulin infusion in polyarteritis nodosa. <i>Clinical Rheumatology</i> , 2006, 25, 396-398.	1.0	26
72	Endothelin Receptor Blockade Ameliorates Vascular Fragility in Endothelial Cell-Specific Fli1 Knockout Mice by Increasing Fli1 DNA Binding Ability. <i>Arthritis and Rheumatology</i> , 2015, 67, 1335-1344.	2.9	26

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73	Effect of human recombinant granulocyte/macrophage colony-stimulating factor and native granulocyte colony-stimulating factor on clonogenic leukemic blast cells. <i>Cancer Research</i> , 1987, 47, 5647-8.	0.4	26
74	Serum apelin levels: clinical association with vascular involvements in patients with systemic sclerosis. <i>Journal of the European Academy of Dermatology and Venereology</i> , 2013, 27, 37-42.	1.3	25
75	High-dose intravenous immunoglobulin infusion as treatment for diffuse scleroderma. <i>British Journal of Dermatology</i> , 2007, 156, 1058-1060.	1.4	24
76	Increased serum soluble CD147 levels in patients with systemic sclerosis: association with scleroderma renal crisis. <i>Clinical Rheumatology</i> , 2012, 31, 835-839.	1.0	24
77	Serum Autotaxin Levels Correlate with Pruritus in Patients with Atopic Dermatitis. <i>Journal of Investigative Dermatology</i> , 2014, 134, 1745-1747.	0.3	23
78	Successful experience of rituximab therapy for systemic sclerosis-associated interstitial lung disease with concomitant systemic lupus erythematosus. <i>Journal of Dermatology</i> , 2014, 41, 418-420.	0.6	23
79	The impact of transcription factor Fli1 deficiency on the regulation of angiogenesis. <i>Experimental Dermatology</i> , 2017, 26, 912-918.	1.4	23
80	Effects of bosentan on nondigital ulcers in patients with systemic sclerosis. <i>British Journal of Dermatology</i> , 2012, 166, 417-421.	1.4	22
81	Serum resistin levels: a possible correlation with pulmonary vascular involvement in patients with systemic sclerosis. <i>Rheumatology International</i> , 2014, 34, 1165-1170.	1.5	22
82	Prediction of therapeutic response before and during i.v. cyclophosphamide pulse therapy for interstitial lung disease in systemic sclerosis: A longitudinal observational study. <i>Journal of Dermatology</i> , 2018, 45, 1425-1433.	0.6	22
83	Elevated serum galectin-9 levels in patients with atopic dermatitis. <i>Journal of Dermatology</i> , 2015, 42, 723-726.	0.6	21
84	Tamibarotene Ameliorates Bleomycin-Induced Dermal Fibrosis by Modulating Phenotypes of Fibroblasts, Endothelial Cells, and Immune Cells. <i>Journal of Investigative Dermatology</i> , 2016, 136, 387-398.	0.3	21
85	Safety and tolerability of bosentan for digital ulcers in Japanese patients with systemic sclerosis: Prospective, multicenter, open-label study. <i>Journal of Dermatology</i> , 2017, 44, 13-17.	0.6	21
86	Increased production of soluble inducible costimulator in patients with diffuse cutaneous systemic sclerosis. <i>Archives of Dermatological Research</i> , 2013, 305, 17-23.	1.1	20
87	Clinical significance of monitoring serum adiponectin levels during intravenous pulse cyclophosphamide therapy in interstitial lung disease associated with systemic sclerosis. <i>Modern Rheumatology</i> , 2013, 23, 323-329.	0.9	20
88	A potential contribution of altered cathepsin L expression to the development of dermal fibrosis and vasculopathy in systemic sclerosis. <i>Experimental Dermatology</i> , 2016, 25, 287-292.	1.4	20
89	Rationally-based therapeutic disease modification in systemic sclerosis: Novel strategies. <i>Seminars in Cell and Developmental Biology</i> , 2020, 101, 146-160.	2.3	20
90	Analysis of two distinct B cell activation pathways mediated by a monoclonal T helper cell. II. T helper cell secretion of interleukin 4 selectively inhibits antigen-specific B cell activation by cognate, but not noncognate, interactions with T cells. <i>Journal of Immunology</i> , 1988, 140, 419-26.	0.4	20

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91	Fli1 Deficiency Induces CXCL6 Expression in Dermal Fibroblasts and Endothelial Cells, Contributing to the Development of Fibrosis and Vasculopathy in Systemic Sclerosis. <i>Journal of Rheumatology</i> , 2017, 44, 1198-1205.	1.0	19
92	Interleukin-25 is involved in cutaneous T-cell lymphoma progression by establishing a T helper 2-dominant microenvironment. <i>British Journal of Dermatology</i> , 2018, 178, 1373-1382.	1.4	19
93	Fli1-haploinsufficient dermal fibroblasts promote skin-localized transdifferentiation of Th2-like regulatory T cells. <i>Arthritis Research and Therapy</i> , 2018, 20, 23.	1.6	19
94	Age-Related Degeneracy of T Cell Repertoire: Influence of the Aged Environment on T Cell Allorecognition. <i>Gerontology</i> , 1990, 36, 3-9.	1.4	18
95	Clinical significance of serum levels of matrix metalloproteinase-13 in patients with systemic sclerosis. <i>Rheumatology</i> , 2006, 45, 303-307.	0.9	18
96	A potential contribution of decreased galectin-7 expression in stratified epithelia to the development of cutaneous and oesophageal manifestations in systemic sclerosis. <i>Experimental Dermatology</i> , 2019, 28, 536-542.	1.4	18
97	Efficacy of low-dose imatinib mesylate for cutaneous involvement in systemic sclerosis: a preliminary report of three cases. <i>Modern Rheumatology</i> , 2012, 22, 94-99.	0.9	17
98	Clinical significance of serum retinol binding protein-4 levels in patients with systemic sclerosis. <i>Journal of the European Academy of Dermatology and Venereology</i> , 2013, 27, 337-344.	1.3	17
99	Dynamics of serum angiopoietin-2 levels correlate with efficacy of intravenous pulse cyclophosphamide therapy for interstitial lung disease associated with systemic sclerosis. <i>Modern Rheumatology</i> , 2013, 23, 884-890.	0.9	17
100	The wound/burn guidelines "4": Guidelines for the management of skin ulcers associated with connective tissue disease/vasculitis. <i>Journal of Dermatology</i> , 2016, 43, 729-757.	0.6	17
101	A case of taxane-induced scleroderma: a different expression profile of Fli1 proteins in dermal fibroblasts and microvascular endothelial cells compared with systemic sclerosis. <i>British Journal of Dermatology</i> , 2011, 164, 1393-1395.	1.4	16
102	Systemic sclerosis complicated with localized scleroderma-like lesions induced by Krbner phenomenon. <i>Journal of Dermatological Science</i> , 2018, 89, 282-289.	1.0	16
103	Serum interleukin-34 levels in patients with systemic sclerosis: Clinical association with interstitial lung disease. <i>Journal of Dermatology</i> , 2018, 45, 1216-1220.	0.6	16
104	Epitopes associated with MHC restriction site of T cells. III. I-J epitope on MHC-restricted T helper cells. <i>Journal of Experimental Medicine</i> , 1987, 166, 1613-1626.	4.2	15
105	Clinical features of scleroderma patients with contracture of phalanges. <i>Clinical Rheumatology</i> , 2007, 26, 1275-1277.	1.0	15
106	Serum omentin levels: A possible contribution to vascular involvement in patients with systemic sclerosis. <i>Journal of Dermatology</i> , 2015, 42, 461-466.	0.6	15
107	Circulating galectin-1 concentrations in systemic sclerosis: potential contribution to digital vasculopathy. <i>International Journal of Rheumatic Diseases</i> , 2016, 19, 622-627.	0.9	15
108	A case of peplomycin-induced scleroderma. <i>British Journal of Dermatology</i> , 2004, 150, 1213-1214.	1.4	14

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109	Linear Connective Tissue Nevus. <i>Pediatric Dermatology</i> , 2007, 24, 439-441.	0.5	14
110	The wound/burn guidelines â€“ 3: Guidelines for the diagnosis and treatment for diabetic ulcer/gangrene. <i>Journal of Dermatology</i> , 2016, 43, 591-619.	0.6	14
111	Critical contribution of the interleukinâ€6/signal transducer and activator of transcription 3 axis to vasculopathy associated with systemic sclerosis. <i>Journal of Dermatology</i> , 2017, 44, 967-971.	0.6	14
112	Clinical significance of monitoring serum adiponectin levels during intravenous pulse cyclophosphamide therapy in interstitial lung disease associated with systemic sclerosis. <i>Modern Rheumatology</i> , 2013, 23, 323-329.	0.9	14
113	Effect of Interleukin 10 on the Hematopoietic Progenitor Cells from Patients with Aplastic Anemia. <i>Stem Cells</i> , 1999, 17, 147-151.	1.4	13
114	Significant attenuation of macrovascular involvement by bosentan in a patient with diffuse cutaneous systemic sclerosis with multiple digital ulcers and gangrene. <i>Modern Rheumatology</i> , 2011, 21, 548-552.	0.9	13
115	Effect of the chimeric soluble granulocyte colony-stimulating factor receptor on the proliferation of leukemic blast cells from patients with acute myeloblastic leukemia. <i>Cancer Research</i> , 1997, 57, 3395-7.	0.4	13
116	Anti-U1RNP antibodies in patients with localized scleroderma. <i>Archives of Dermatological Research</i> , 2001, 293, 455-459.	1.1	12
117	Serum levels of interleukinâ€18â€binding protein isoform a: Clinical association with inflammation and pulmonary hypertension in systemic sclerosis. <i>Journal of Dermatology</i> , 2016, 43, 912-918.	0.6	12
118	Nucleosome in patients with systemic sclerosis: possible association with immunological abnormalities via abnormal activation of T and B cells. <i>Annals of the Rheumatic Diseases</i> , 2016, 75, 1858-1865.	0.5	12
119	Altered Properties of Endothelial Cells and Mesenchymal Stem Cells Underlying the Development of Sclerodermaâ€like Vasculopathy in KLF5<sup>+/â~</sup>;Fliâ€1<sup>+/â~</sup> Mice. <i>Arthritis and Rheumatology</i> , 2020, 72, 2136-2146.	2.9	12
120	Interleukin-10 inhibits the autocrine growth of leukemic blast cells from patients with acute myeloblastic leukemia. <i>International Journal of Hematology</i> , 1997, 66, 445.	0.7	12
121	Circulating soluble CD40 ligand in patients with eosinophilic fasciitis. <i>Annals of the Rheumatic Diseases</i> , 2003, 62, 190-191.	0.5	11
122	ICAM-1 Deficiency Exacerbates Sarcoid-Like Granulomatosis Induced by Propionibacterium acnes through Impaired IL-10 Production by Regulatory T Cells. <i>American Journal of Pathology</i> , 2013, 183, 1731-1739.	1.9	11
123	Serum vaspin levels: A possible correlation with digital ulcers in patients with systemic sclerosis. <i>Journal of Dermatology</i> , 2015, 42, 528-531.	0.6	11
124	Gastroesophageal Reflux Disease-Related Disorders of Systemic Sclerosis Based on the Analysis of 66 Patients. <i>Digestion</i> , 2018, 98, 201-208.	1.2	11
125	Increased expression of aquaporin-1 in dermal fibroblasts and dermal microvascular endothelial cells possibly contributes to skin fibrosis and edema in patients with systemic sclerosis. <i>Journal of Dermatological Science</i> , 2019, 93, 24-32.	1.0	11
126	Altered dynamics of transforming growth factor Æ (TGF-Æ) receptors in scleroderma fibroblasts. <i>Annals of the Rheumatic Diseases</i> , 2011, 70, 384-387.	0.5	10



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127	Improvement of endothelial function in parallel with the amelioration of dry cough and dyspnea due to interstitial pneumonia by intravenous cyclophosphamide pulse therapy in patients with systemic sclerosis: a preliminary report of two cases. <i>Modern Rheumatology</i> , 2012, 22, 598-601.	0.9	10
128	Serum levels of matrix metalloproteinase-13 in patients with eosinophilic fasciitis. <i>Journal of Dermatology</i> , 2014, 41, 746-748.	0.6	10
129	A potential contribution of psoriasin to vascular and epithelial abnormalities and inflammation in systemic sclerosis. <i>Journal of the European Academy of Dermatology and Venereology</i> , 2018, 32, 291-297.	1.3	10
130	Systemic sclerosis: Is the epithelium a missing piece of the pathogenic puzzle?. <i>Journal of Dermatological Science</i> , 2019, 94, 259-265.	1.0	10
131	Facile fabrication of PEG-coated PLGA microspheres via SPG membrane emulsification for the treatment of scleroderma by ECM degrading enzymes. <i>Colloids and Surfaces B: Biointerfaces</i> , 2019, 179, 453-461.	2.5	10
132	Serum S100A12 levels: Possible association with skin sclerosis and interstitial lung disease in systemic sclerosis. <i>Experimental Dermatology</i> , 2021, 30, 409-415.	1.4	10
133	Generation of T cell repertoire. Two distinct mechanisms for generation of T suppressor cells, T helper cells, and T augmenting cells. <i>Journal of Immunology</i> , 1989, 142, 365-73.	0.4	10
134	Unprecedented success of rituximab therapy for prednisolone- and immunosuppressant-resistant systemic sclerosis-associated interstitial lung disease. <i>Scandinavian Journal of Rheumatology</i> , 2017, 46, 247-252.	0.6	9
135	A possible implication of reduced levels of LIF, LIFR, and gp130 in vasculopathy related to systemic sclerosis. <i>Archives of Dermatological Research</i> , 2017, 309, 833-842.	1.1	9
136	Association of serum CCL20 levels with pulmonary vascular involvement and primary biliary cholangitis in patients with systemic sclerosis. <i>International Journal of Rheumatic Diseases</i> , 2021, 24, 711-718.	0.9	9
137	Serum H $\alpha$ -ficolin levels: Clinical association with interstitial lung disease in patients with systemic sclerosis. <i>Journal of Dermatology</i> , 2017, 44, 1168-1171.	0.6	8
138	Recent advances in the treatment of skin involvement in systemic sclerosis. <i>Inflammation and Regeneration</i> , 2017, 37, 12.	1.5	8
139	Rapid alteration of serum interleukin-6 levels may predict the reactivity of i.v. cyclophosphamide pulse therapy in systemic sclerosis-associated interstitial lung disease. <i>Journal of Dermatology</i> , 2018, 45, 1221-1224.	0.6	8
140	Fli1 deficiency induces endothelial adipsin expression, contributing to the onset of pulmonary arterial hypertension in systemic sclerosis. <i>Rheumatology</i> , 2020, 59, 2005-2015.	0.9	8
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