Kunikazu Tanji

List of Publications by Year in descending order

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130	9,520	40	94
papers	citations	h-index	g-index
130	130	130	19304
all docs	docs citations	times ranked	citing authors

#	Article	IF	CITATIONS
1	Neuropathology of Multiple System Atrophy, a Glioneuronal Degenerative Disease. Cerebellum, 2024, 23, 2-12.	2.5	3
2	Inducible Systemic Gcn1 Deletion in Mice Leads to Transient Body Weight Loss upon Tamoxifen Treatment Associated with Decrease of Fat and Liver Glycogen Storage. International Journal of Molecular Sciences, 2022, 23, 3201.	4.1	2
3	Accumulation of Nonfibrillar TDP-43 in the Rough Endoplasmic Reticulum Is the Early-Stage Pathology in Amyotrophic Lateral Sclerosis. Journal of Neuropathology and Experimental Neurology, 2022, 81, 271-281.	1.7	5
4	Novel tankyrase inhibitors suppress TDP-43 aggregate formation. Biochemical and Biophysical Research Communications, 2021, 537, 85-92.	2.1	4
5	Role of VAPB and vesicular profiles in αâ€synuclein aggregates in multiple system atrophy. Brain Pathology, 2021, 31, e13001.	4.1	5
6	Effects of voluntary and forced exercises on motor function recovery in intracerebral hemorrhage rats. NeuroReport, 2020, 31, 189-196.	1.2	8
7	Nrp1 is Activated by Konjac Ceramide Binding-Induced Structural Rigidification of the a1a2 Domain. Cells, 2020, 9, 517.	4.1	2
8	Ribosome binding protein GCN1Âregulates the cell cycle and cell proliferation and is essential for the embryonic development of mice. PLoS Genetics, 2020, 16, e1008693.	3.5	20
9	Title is missing!. , 2020, 16, e1008693.		O
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	Title is missing!. , 2020, 16, e1008693. Phosphorylated TDP-43 aggregates in skeletal and cardiac muscle are a marker of myogenic degeneration in amyotrophic lateral sclerosis and various conditions. Acta Neuropathologica	5.2 1.3	0
13	Title is missing!., 2020, 16, e1008693. Phosphorylated TDP-43 aggregates in skeletal and cardiac muscle are a marker of myogenic degeneration in amyotrophic lateral sclerosis and various conditions. Acta Neuropathologica Communications, 2019, 7, 165. Konjac ceramide (kCer) regulates keratinocyte migration by Sema3A-like repulsion mechanism.		O 35
13	Title is missing!., 2020, 16, e1008693. Phosphorylated TDP-43 aggregates in skeletal and cardiac muscle are a marker of myogenic degeneration in amyotrophic lateral sclerosis and various conditions. Acta Neuropathologica Communications, 2019, 7, 165. Konjac ceramide (kCer) regulates keratinocyte migration by Sema3A-like repulsion mechanism. Biochemistry and Biophysics Reports, 2019, 17, 132-138. Autophagy Is a Common Degradation Pathway for Bunina Bodies and TDP-43 Inclusions in Amyotrophic	1.3	0 35 4
13 14 15	Title is missing!. , 2020, 16, e1008693. Phosphorylated TDP-43 aggregates in skeletal and cardiac muscle are a marker of myogenic degeneration in amyotrophic lateral sclerosis and various conditions. Acta Neuropathologica Communications, 2019, 7, 165. Konjac ceramide (kCer) regulates keratinocyte migration by Sema3A-like repulsion mechanism. Biochemistry and Biophysics Reports, 2019, 17, 132-138. Autophagy Is a Common Degradation Pathway for Bunina Bodies and TDP-43 Inclusions in Amyotrophic Lateral Sclerosis. Journal of Neuropathology and Experimental Neurology, 2019, 78, 910-921. Immunoreactivity of myelinâ€essociated oligodendrocytic basic protein in Lewy bodies. Neuropathology,	1.3	0 35 4

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19	Neurite Outgrowth and Morphological Changes Induced by 8-trans Unsaturation of Sphingadienine in kCer Molecular Species. International Journal of Molecular Sciences, 2019, 20, 2116.	4.1	4
20	An autopsy case of earlyâ€stage amyotrophic lateral sclerosis with TDPâ€43 immunoreactive neuronal, but not glial, inclusions. Neuropathology, 2019, 39, 224-230.	1.2	8
21	A mouse model of adult-onset multiple system atrophy. Neurobiology of Disease, 2019, 127, 339-349.	4.4	14
22	Atypical globular glial tauopathy with a combination of types I and II pathology. Neuropathology, 2019, 39, 127-134.	1.2	3
23	Role of the ISR-ATF4 pathway and its cross talk with Nrf2 in mitochondrial quality control. Journal of Clinical Biochemistry and Nutrition, 2019, 64, 1-12.	1.4	67
24	TRIM9 and TRIM67 Are New Targets in Paraneoplastic Cerebellar Degeneration. Cerebellum, 2019, 18, 245-254.	2.5	44
25	YOD1 attenuates neurogenic proteotoxicity through its deubiquitinating activity. Neurobiology of Disease, 2018, 112, 14-23.	4.4	23
26	AMBRA1, a novel αâ€synucleinâ€binding protein, is implicated in the pathogenesis of multiple system atrophy. Brain Pathology, 2018, 28, 28-42.	4.1	25
27	Alteration of autophagy-related proteins in peripheral blood mononuclear cells of patients with Parkinson's disease. Neurobiology of Aging, 2018, 63, 33-43.	3.1	54
28	Immunohistochemical localization of exoribonucleases (DIS3L2 and XRN1) in intranuclear inclusion body disease. Neuroscience Letters, 2018, 662, 389-394.	2.1	11
29	Gnetin C, a resveratrol dimer, reduces amyloid-β 1–42 (Aβ42) production and ameliorates Aβ42-lowered cell viability in cultured SH-SY5Y human neuroblastoma cells . Biomedical Research, 2018, 39, 105-115.	0.9	17
30	Colocalization of Bunina bodies and TDPâ€43 inclusions in a case of sporadic amyotrophic lateral sclerosis with Lewy bodyâ€like hyaline inclusions. Neuropathology, 2018, 38, 521-528.	1.2	11
31	Autophagy mediators (FOXO1, SESN3 and TSC2) in Lewy body disease and aging. Neuroscience Letters, 2018, 684, 35-41.	2.1	19
32	Interferon (IFN)-induced protein 35 (IFI35) negatively regulates IFN-Î ² -phosphorylated STAT1-RIG-I-CXCL10/CCL5 axis in U373MG astrocytoma cells treated with polyinosinic-polycytidylic acid. Brain Research, 2017, 1658, 60-67.	2,2	18
33	PLA2G6 accumulates in Lewy bodies in PARK14 and idiopathic Parkinson's disease. Neuroscience Letters, 2017, 645, 40-45.	2.1	34
34	Rebamipide reduces amyloid-l² 1–42 (Al²42) production and ameliorates Al²43-lowered cell viability in cultured SH-SY5Y human neuroblastoma cells. Neuroscience Research, 2017, 124, 40-50.	1.9	6
35	Alteration of mitochondrial protein PDHA1 in Lewy body disease and PARK14. Biochemical and Biophysical Research Communications, 2017, 489, 439-444.	2.1	16
36	Status epilepticus causing extensive microvacuolar change with astrocytosis and diffusion MRI abnormalities in the subcortical white matter. Journal of the Neurological Sciences, 2017, 382, 55-57.	0.6	2

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37	Alteration of Upstream Autophagyâ€Related Proteins (<scp>ULK1</scp> , <scp>ULK2</scp> ,) Tj ETQq1 1 0.78431 Pathology, 2016, 26, 359-370.	14 rgBT /O [.] 4.1	verlock 10 40
38	Accumulation of phosphorylated αâ€synuclein in subpial and periventricular astrocytes in multiple system atrophy of long duration. Neuropathology, 2016, 36, 157-167.	1.2	38
39	αâ€Synuclein pathology in the cranial and spinal nerves in Lewy body disease. Neuropathology, 2016, 36, 262-269.	1.2	17
40	Novel eosinophilic neuronal cytoplasmic inclusions in the external cuneate nucleus of humans. Neuropathology, 2016, 36, 441-447.	1.2	1
41	Guidelines for the use and interpretation of assays for monitoring autophagy (3rd edition). Autophagy, 2016, 12, 1-222.	9.1	4,701
42	The role of NUB1 in \hat{l}_{\pm} -synuclein degradation in Lewy body disease model mice. Biochemical and Biophysical Research Communications, 2016, 470, 635-642.	2.1	3
43	G proteinâ€coupled receptor 26 immunoreactivity in intranuclear inclusions associated with polyglutamine and intranuclear inclusion body diseases. Neuropathology, 2016, 36, 50-55.	1.2	11
44	Interferon-stimulated gene (ISG) 60, as well as ISG56 and ISG54, positively regulates TLR3/IFN-β/STAT1 axis in U373MG human astrocytoma cells. Neuroscience Research, 2016, 105, 35-41.	1.9	24
45	Isopentenyl diphosphate isomerase, a cholesterol synthesizing enzyme, is localized in <scp>L</scp> ewy bodies. Neuropathology, 2015, 35, 432-440.	1.2	31
46	An autopsy case of refractory epilepsy due to unilateral polymicrogyria in a 65â€yearâ€old man: Histogenesis of fourâ€layered polymicrogyric cortex. Neuropathology, 2015, 35, 569-574.	1.2	1
47	<scp>p</scp> 62 Deficiency Enhances αâ€Synuclein Pathology in Mice. Brain Pathology, 2015, 25, 552-564.	4.1	37
48	Localization of nuclear receptor subfamily 4, group A, member 3 (<scp>NR4A3</scp>) in <scp>L</scp> ewy body disease and multiple system atrophy. Neuropathology, 2015, 35, 503-509.	1.2	8
49	Emerging functional cross-talk between the Keap1-Nrf2 system and mitochondria. Journal of Clinical Biochemistry and Nutrition, 2015, 56, 91-97.	1.4	115
50	Carnosic acid attenuates apoptosis induced by amyloid- \hat{l}^2 $1\hat{a}$ \in "42 or $1\hat{a}$ \in "43 in SH-SY5Y human neuroblastoma cells. Neuroscience Research, 2015, 94, 1-9.	1.9	47
51	Desferrioxamine, an iron chelator, inhibits CXCL10 expression induced by polyinosinic–polycytidylic acid in U373MG human astrocytoma cells. Neuroscience Research, 2015, 94, 10-16.	1.9	9
52	Role of the <scp>K</scp> eap1/ <scp>N</scp> rf2 pathway in neurodegenerative diseases. Pathology International, 2015, 65, 210-219.	1.3	104
53	Filamentous aggregations of phosphorylated α-synuclein in Schwann cells (Schwann cell cytoplasmic) Tj ETQq1 1	0.784314 5.2	FrgBT /Ove
54	Retinoic acid-inducible gene-I-like receptor (RLR)-mediated antiviral innate immune responses in the lower respiratory tract: Roles of TRAF3 and TRAF5. Biochemical and Biophysical Research Communications, 2015, 467, 191-196.	2.1	4

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55	Trehalose intake induces chaperone molecules along with autophagy in a mouse model of Lewy body disease. Biochemical and Biophysical Research Communications, 2015, 465, 746-752.	2.1	70
56	Sortilin-related receptor CNS expressed 2 (SorCS2) is localized to Bunina bodies in amyotrophic lateral sclerosis. Neuroscience Letters, 2015, 608, 6-11.	2.1	8
57	Sigma-1 receptor is involved in degradation of intranuclear inclusions in a cellular model of Huntington's disease. Neurobiology of Disease, 2015, 74, 25-31.	4.4	54
58	The Role of Atg8 Homologue in Lewy Body Disease. , 2014, , 383-389.		0
59	An autopsy case of incipient <scp>P</scp> ick's disease: Immunohistochemical profile of earlyâ€stage <scp>P</scp> ick body formation. Neuropathology, 2014, 34, 386-391.	1.2	10
60	Accumulation of the sigmaâ€1 receptor is common to neuronal nuclear inclusions in various neurodegenerative diseases. Neuropathology, 2014, 34, 148-158.	1.2	52
61	<scp>ALS</scp> â€associated protein <scp>FIG4</scp> is localized in <scp>P</scp> ick and <scp>L</scp> ewy bodies, and also neuronal nuclear inclusions, in polyglutamine and intranuclear inclusion body diseases. Neuropathology, 2014, 34, 19-26.	1.2	27
62	Ubiquitinâ€negative, eosinophilic neuronal cytoplasmic inclusions associated with stress granules and autophagy: An immunohistochemical investigation of two cases. Neuropathology, 2014, 34, 140-147.	1.2	2
63	Phosphorylation of serine 349 of p62 in Alzheimer's disease brain. Acta Neuropathologica Communications, 2014, 2, 50.	5.2	43
64	Carnosic acid suppresses the production of amyloid- \hat{l}^2 1-42 and 1-43 by inducing an \hat{l}_\pm -secretase TACE/ADAM17 in U373MG human astrocytoma cells. Neuroscience Research, 2014, 79, 83-93.	1.9	49
65	ISG54 and ISG56 are induced by TLR3 signaling in U373MG human astrocytoma cells: Possible involvement in CXCL10 expression. Neuroscience Research, 2014, 84, 34-42.	1.9	24
66	Valosinâ€containing protein immunoreactivity in tauopathies, synucleinopathies, polyglutamine diseases and intranuclear inclusion body disease. Neuropathology, 2013, 33, 637-644.	1.2	20
67	Plasma matrix metalloproteinase-3 correlates with the clinical severity in men with multiple system atrophy. Neurology and Clinical Neuroscience, 2013, 1, 69-77.	0.4	3
68	Giant cell polymyositis and myocarditis associated with myasthenia gravis and thymoma. Neuropathology, 2013, 33, 281-287.	1.2	43
69	Alteration of autophagosomal proteins in the brain of multiple system atrophy. Neurobiology of Disease, 2013, 49, 190-198.	4.4	50
70	The Lewy Body in Parkinson's Disease and Related Neurodegenerative Disorders. Molecular Neurobiology, 2013, 47, 495-508.	4.0	323
71	Carnosic acid suppresses the production of amyloid- \hat{l}^2 $1\hat{a}$ \in "42 by inducing the metalloprotease gene TACE/ADAM17 in SH-SY5Y human neuroblastoma cells. Neuroscience Research, 2013, 75, 94-102.	1.9	45
72	Brain expression level and activity of HDAC6 protein in neurodegenerative dementia. Biochemical and Biophysical Research Communications, 2013, 430, 394-399.	2.1	34

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73	Keap1 Is Localized in Neuronal and Glial Cytoplasmic Inclusions in Various Neurodegenerative Diseases. Journal of Neuropathology and Experimental Neurology, 2013, 72, 18-28.	1.7	61
74	An autopsy case of preclinical multiple system atrophy (<scp>MSA</scp> â€ <scp>C</scp>). Neuropathology, 2013, 33, 667-672.	1.2	27
75	Endosomal sorting related protein CHMP2B is localized in Lewy bodies and glial cytoplasmic inclusions in α-synucleinopathy. Neuroscience Letters, 2012, 527, 16-21.	2.1	24
76	Autophagy-related proteins (p62, NBR1 and LC3) in intranuclear inclusions in neurodegenerative diseases. Neuroscience Letters, 2012, 522, 134-138.	2.1	35
77	p62/sequestosome 1 binds to TDPâ€43 in brains with frontotemporal lobar degeneration with TDPâ€43 inclusions. Journal of Neuroscience Research, 2012, 90, 2034-2042.	2.9	60
78	Optineurin immunoreactivity in neuronal nuclear inclusions of polyglutamine diseases (Huntington's,) Tj ETC	Qq Q,Q 0 rg	BT /Qverlock
79	Autophagic adapter protein NBR1 is localized in Lewy bodies and glial cytoplasmic inclusions and is involved in aggregate formation in α-synucleinopathy. Acta Neuropathologica, 2012, 124, 173-186.	7.7	92
80	Ubiquilin immunoreactivity in cytoplasmic and nuclear inclusions in synucleinopathies, polyglutamine diseases and intranuclear inclusion body disease. Acta Neuropathologica, 2012, 124, 149-151.	7.7	41
81	Abnormal tau deposition in neurons, but not in glial cells in the cerebral tissue surrounding arteriovenous malformation. Neuropathology, 2012, 32, 267-271.	1.2	4
82	Immunohistochemical analysis of Marinesco bodies, using antibodies against proteins implicated in the ubiquitinâ€proteasome system, autophagy and aggresome formation. Neuropathology, 2012, 32, 261-266.	1.2	30
83	Ubiquitinâ€related proteins in neuronal and glial intranuclear inclusions in intranuclear inclusion body disease. Pathology International, 2012, 62, 407-411.	1.3	17
84	Edaravone and carnosic acid synergistically enhance the expression of nerve growth factor in human astrocytes under hypoxia/reoxygenation. Neuroscience Research, 2011, 69, 291-298.	1.9	22
85	Synphilin-1-Binding Protein NUB1 is Colocalized With Nonfibrillar, Proteinase K-Resistant α-Synuclein in Presynapses in Lewy Body Disease. Journal of Neuropathology and Experimental Neurology, 2011, 70, 879-889.	1.7	15
86	Incipient intranuclear inclusion body disease in a 78-year-old woman. Neuropathology, 2011, 31, 188-193.	1.2	23
87	Enhancement of native and phosphorylated TDPâ€43 immunoreactivity by proteinase K treatment following autoclave heating. Neuropathology, 2011, 31, 401-404.	1.2	6
88	Accumulation of histone deacetylase 6, an aggresome-related protein, is specific to Lewy bodies and glial cytoplasmic inclusions. Neuropathology, 2011, 31, 561-568.	1.2	60
89	Alteration of autophagosomal proteins (LC3, GABARAP and GATE-16) in Lewy body disease. Neurobiology of Disease, 2011, 43, 690-697.	4.4	102
90	Immunohistochemical study of microscopic globular bodies of normal human brain. Biomedical Research, 2011, 32, 337-342.	0.9	1

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91	Proteinase K-resistant α-synuclein is deposited in presynapses in human Lewy body disease and A53T α-synuclein transgenic mice. Acta Neuropathologica, 2010, 120, 145-154.	7.7	87
92	Involvement of the peripheral nervous system in synucleinopathies, tauopathies and other neurodegenerative proteinopathies of the brain. Acta Neuropathologica, 2010, 120, 1-12.	7.7	131
93	Dynamic movements of Ro52 cytoplasmic bodies along microtubules. Histochemistry and Cell Biology, 2010, 133, 273-284.	1.7	20
94	TRIM9, a novel brain-specific E3 ubiquitin ligase, is repressed in the brain of Parkinson's disease and dementia with Lewy bodies. Neurobiology of Disease, 2010, 38, 210-218.	4.4	82
95	Widespread occurrence of eosinophilic neuronal cytoplasmic inclusions in an asymptomatic adult: A novel ubiquitin-negative filamentous inclusion. Neuropathology, 2010, 30, 648-653.	1.2	2
96	Retinoic acid-inducible gene-I is induced by double-stranded RNA and regulates the expression of CC chemokine ligand (CCL) 5 in human mesangial cells. Nephrology Dialysis Transplantation, 2010, 25, 3534-3539.	0.7	47
97	Edaravone improves the expression of nerve growth factor in human astrocytes subjected to hypoxia/reoxygenation. Neuroscience Research, 2010, 66, 284-289.	1.9	11
98	Accumulation of presynaptic proteinase K-resistant alpha-synuclein in Parkinson's disease. Neuroscience Research, 2010, 68, e192.	1.9	0
99	Alteration of biochemical and pathological properties of TDP-43 protein by a lipid mediator, 15-deoxy-Δ12,14-prostaglandin J2. Experimental Neurology, 2010, 222, 296-303.	4.1	15
100	Retinoic acid-inducible gene-l is constitutively expressed and involved in IFN-γ-stimulated CXCL9–11 production in intestinal epithelial cells. Immunology Letters, 2009, 123, 9-13.	2.5	21
101	Decreased Cystatin C Immunoreactivity in Spinal Motor Neurons and Astrocytes in Amyotrophic Lateral Sclerosis. Journal of Neuropathology and Experimental Neurology, 2009, 68, 1200-1206.	1.7	24
102	α-Synuclein pathology in the neostriatum in Parkinson's disease. Acta Neuropathologica, 2008, 115, 453-459.	7.7	52
103	Maturation process of TDP-43-positive neuronal cytoplasmic inclusions in amyotrophic lateral sclerosis with and without dementia. Acta Neuropathologica, 2008, 116, 193-203.	7.7	111
104	Ubiquitination of E3 ubiquitin ligase TRIM5α and its potential role. FEBS Journal, 2008, 275, 1540-1555.	4.7	97
105	Retinoic acid-inducible gene-l is induced by interferon- \hat{I}^3 and regulates CXCL11 expression in HeLa cells. Life Sciences, 2008, 82, 670-675.	4.3	16
106	Epitope mapping of 2E2-D3, a monoclonal antibody directed against human TDP-43. Neuroscience Letters, 2008, 434, 170-174.	2.1	35
107	Parkin is expressed in vascular endothelial cells. Neuroscience Letters, 2007, 419, 199-201.	2.1	4
108	Retinoic acid-inducible gene-I mediates RANTES/CCL5 expression in U373MG human astrocytoma cells stimulated with double-stranded RNA. Neuroscience Research, 2007, 58, 199-206.	1.9	39

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109	The Lewy body in Parkinson's disease: Molecules implicated in the formation and degradation of αâ€synuclein aggregates. Neuropathology, 2007, 27, 494-506.	1.2	415
110	Immunohistochemical localization of NUB1, a synphilin-1-binding protein, in neurodegenerative disorders. Acta Neuropathologica, 2007, 114, 365-371.	7.7	23
111	TDP-43-immunoreactive neuronal and glial inclusions in the neostriatum in amyotrophic lateral sclerosis with and without dementia. Acta Neuropathologica, 2007, 115, 115-122.	7.7	103
112	Interferon-Î ³ upregulates retinoic acid-inducible gene-l in human pericardial mesothelial cells. Acta Cardiologica, 2007, 62, 553-557.	0.9	7
113	NUB1 Suppresses the Formation of Lewy Body-Like Inclusions by Proteasomal Degradation of Synphilin-1. American Journal of Pathology, 2006, 169, 553-565.	3.8	56
114	Oncogenic protein UnpEL/Usp4 deubiquitinates Ro52 by its isopeptidase activity. Biochemical and Biophysical Research Communications, 2006, 339, 731-736.	2.1	30
115	Function and subcellular location of $Ro52\hat{l}^2$. Biochemical and Biophysical Research Communications, 2006, 340, 872-878.	2.1	8
116	Platelet-activating factor enhances the expression of nerve growth factor in normal human astrocytes under hypoxia. Molecular Brain Research, 2005, 133, 95-101.	2.3	17
117	Interaction of NUB1 with the proteasome subunit S5a. Biochemical and Biophysical Research Communications, 2005, 337, 116-120.	2.1	27
118	\hat{l}_{\pm} -Synuclein pathology affecting Bergmann glia of the cerebellum in patients with \hat{l}_{\pm} -synucleinopathies. Acta Neuropathologica, 2003, 105, 403-409.	7.7	54
119	15-Deoxy-Δ12,14-prostaglandin J2 inhibits the expression of granulocyte-macrophage colony-stimulating factor in endothelial cells stimulated with lipopolysaccharide. Prostaglandins and Other Lipid Mediators, 2003, 71, 293-299.	1.9	10
120	Demonstration of α-Synuclein Immunoreactivity in Neuronal and Glial Cytoplasm in Normal Human Brain Tissue Using Proteinase K and Formic Acid Pretreatment. Experimental Neurology, 2002, 176, 98-104.	4.1	128
121	Retinoic Acid-Inducible Gene-I Is Induced in Endothelial Cells by LPS and Regulates Expression of COX-2. Biochemical and Biophysical Research Communications, 2002, 292, 274-279.	2.1	116
122	Expression of $\hat{l}\pm$ -synuclein, the precursor of non-amyloid \hat{l}^2 component of Alzheimer's disease amyloid, in human cerebral blood vessels. Neuroscience Letters, 2002, 326, 5-8.	2.1	62
123	Immunohistochemical comparison of \hat{l}_{\pm} - and \hat{l}^2 -synuclein in adult rat central nervous system. Brain Research, 2002, 941, 118-126.	2.2	75
124	Platelet-activating factor enhances the expression of vascular endothelial growth factor in normal human astrocytes. Brain Research, 2002, 944, 65-72.	2.2	25
125	15â€Deoxyâ€ î" ^{12,14} â€prostaglandin J ₂ inhibitsCX3CL1/fractalkine expression human endothelial cells. Immunology and Cell Biology, 2002, 80, 531-536.	in 2.3	22
126	Soluble Interleukin-6 Receptor α Inhibits the Cytokine-Induced Fractalkine/CX3CL1 Expression in Human Vascular Endothelial Cells in Culture. Experimental Cell Research, 2001, 269, 35-41.	2.6	42

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127	Synergistic stimulation, by tumor necrosis factor-α and interferon-γ, of fractalkine expression in human astrocytes. Neuroscience Letters, 2001, 303, 132-136.	2.1	86
128	Desferrioxamine, an iron chelator, upregulates cyclooxygenase-2 expression and prostaglandin production in a human macrophage cell line. Biochimica Et Biophysica Acta - Molecular and Cell Biology of Lipids, 2001, 1530, 227-235.	2.4	47
129	Interleukin-1β stimulates galectin-9 expression in human astrocytes. NeuroReport, 2001, 12, 3755-3758.	1.2	59
130	Expression of ??-synuclein in a human glioma cell line and its up-regulation by interleukin-1??. NeuroReport, 2001, 12, 1909-1912.	1.2	63