Nadia Stefanova

List of Publications by Year in descending order

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		116194	104191
117	5,480 citations	36	69
papers	citations	h-index	g-index
123	123	123	5770
all docs	docs citations	times ranked	citing authors

#	Article	IF	CITATIONS
1	Host oligodendrogliopathy and α-synuclein strains dictate disease severity in multiple system atrophy. Brain, 2023, 146, 237-251.	3.7	10
2	The Compound ATH434 Prevents Alpha-Synuclein Toxicity in a Murine Model of Multiple System Atrophy. Journal of Parkinson's Disease, 2022, 12, 105-115.	1.5	9
3	Spreading of Aggregated α-Synuclein in Sagittal Organotypic Mouse Brain Slices. Biomolecules, 2022, 12, 163.	1.8	4
4	Disease-Modifying Therapies for Multiple System Atrophy: Where Are We in 2022?. Journal of Parkinson's Disease, 2022, 12, 1369-1387.	1.5	10
5	Microglia in Parkinson's Disease. Journal of Parkinson's Disease, 2022, 12, S105-S112.	1.5	18
6	A multiplex pedigree with pathologically confirmed multiple system atrophy and Parkinson's disease with dementia. Brain Communications, 2022, 4, .	1.5	3
7	Shared Genetics of Multiple System Atrophy and Inflammatory Bowel Disease. Movement Disorders, 2021, 36, 449-459.	2.2	16
8	Signs of early cellular dysfunction in multiple system atrophy. Neuropathology and Applied Neurobiology, 2021, 47, 268-282.	1.8	16
9	Multiple System Atrophy. , 2021, , 1-29.		O
10	Neuropathology of multiple system atrophy: Kurt Jellinger's legacy. Journal of Neural Transmission, 2021, 128, 1481-1494.	1.4	6
10	Neuropathology of multiple system atrophy: Kurt Jellinger`s legacy. Journal of Neural Transmission, 2021, 128, 1481-1494. <scp>ATH434</scp> Reduces αâ€Synucleinâ€Related Neurodegeneration in a Murine Model of Multiple System Atrophy. Movement Disorders, 2021, 36, 2605-2614.	1.4 2.2	6
	2021, 128, 1481-1494. <scp>ATH434</scp> Reduces αâ€Synucleinâ€Related Neurodegeneration in a Murine Model of Multiple		
11	2021, 128, 1481-1494. <scp>ATH434</scp> Reduces αâ€Synucleinâ€Related Neurodegeneration in a Murine Model of Multiple System Atrophy. Movement Disorders, 2021, 36, 2605-2614. Current experimental disease-modifying therapeutics for multiple system atrophy. Journal of Neural	2.2	11
11 12	2021, 128, 1481-1494. ⟨scp⟩ATH434⟨/scp⟩ Reduces αâ€Synucleinâ€Related Neurodegeneration in a Murine Model of Multiple System Atrophy. Movement Disorders, 2021, 36, 2605-2614. Current experimental disease-modifying therapeutics for multiple system atrophy. Journal of Neural Transmission, 2021, 128, 1529-1543. Is Multiple System Atrophy a Prion-like Disorder?. International Journal of Molecular Sciences, 2021,	2.2	11
11 12 13	2021, 128, 1481-1494.	2.2 1.4 1.8	11 11 12
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11 12 13 14	 2021, 128, 1481-1494. ⟨scp>ATH434⟨ scp>⟩ Reduces αâ€6ynucleinâ€Related Neurodegeneration in a Murine Model of Multiple System Atrophy. Movement Disorders, 2021, 36, 2605-2614. Current experimental disease-modifying therapeutics for multiple system atrophy. Journal of Neural Transmission, 2021, 128, 1529-1543. Is Multiple System Atrophy a Prion-like Disorder?. International Journal of Molecular Sciences, 2021, 22, 10093. Toll-like receptor 4 deficiency facilitates α-synuclein propagation and neurodegeneration in a mouse model of prodromal Parkinson's disease. Parkinsonism and Related Disorders, 2021, 91, 59-65. Therapeutic potential of iron modulating drugs in a mouse model of multiple system atrophy. Neurobiology of Disease, 2021, 159, 105509. The Concept of α-Synuclein Strains and How Different Conformations May Explain Distinct 	2.2 1.4 1.8 1.1	11 11 12 12 8

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19	Targeting \hat{l}_{\pm} -synuclein by PD03 AFFITOPE® and Anle138b rescues neurodegenerative pathology in a model of multiple system atrophy: clinical relevance. Translational Neurodegeneration, 2020, 9, 38.	3.6	22
20	Signs of Chronic Hypoxia Suggest a Novel Pathophysiological Event in <scp>αâ€Synucleinopathies</scp> . Movement Disorders, 2020, 35, 2333-2338.	2.2	8
21	Transcriptional profiling of multiple system atrophy cerebellar tissue highlights differences between the parkinsonian and cerebellar sub-types of the disease. Acta Neuropathologica Communications, 2020, 8, 76.	2.4	20
22	Assessment of the Retina of Plp- $\hat{l}\pm$ -Syn Mice as a Model for Studying Synuclein-Dependent Diseases. , 2020, 61, 12.		5
23	Genes to treat excitotoxicity ameliorate the symptoms of the disease in mice models of multiple system atrophy. Journal of Neural Transmission, 2020, 127, 205-212.	1.4	6
24	MSA: From basic mechanisms to experimental therapeutics. Parkinsonism and Related Disorders, 2020, 73, 94-104.	1.1	13
25	High-salt diet does not boost neuroinflammation and neurodegeneration in a model of \hat{l}_{\pm} -synucleinopathy. Journal of Neuroinflammation, 2020, 17, 35.	3.1	11
26	Shock waves promote spinal cord repair via TLR3. JCI Insight, 2020, 5, .	2.3	15
27	The molecular tweezer CLR01 reduces aggregated, pathologic, and seeding-competent α-synuclein in experimental multiple system atrophy. Biochimica Et Biophysica Acta - Molecular Basis of Disease, 2019, 1865, 165513.	1.8	25
28	Neuroinflammation and Glial Phenotypic Changes in Alpha-Synucleinopathies. Frontiers in Cellular Neuroscience, 2019, 13, 263.	1.8	54
29	Increased anxiety-like behavior following circuit-specific catecholamine denervation in mice. Neurobiology of Disease, 2019, 125, 55-66.	2.1	25
30	Induced pluripotent stem cells in multiple system atrophy: recent developments and scientific challenges. Clinical Autonomic Research, 2019, 29, 385-395.	1.4	2
31	L-dopa response pattern in a rat model of mild striatonigral degeneration. PLoS ONE, 2019, 14, e0218130.	1.1	O
32	Anle138b modulates αâ€synuclein oligomerization and prevents motor decline and neurodegeneration in a mouse model of multiple system atrophy. Movement Disorders, 2019, 34, 255-263.	2.2	72
33	The Relevance of Iron in the Pathogenesis of Multiple System Atrophy: A Viewpoint. Journal of Alzheimer's Disease, 2018, 61, 1253-1273.	1.2	47
34	Progressive striatonigral degenerationÂin a transgenic mouse model of multiple system atrophy: translational implications for interventional therapies. Acta Neuropathologica Communications, 2018, 6, 2.	2.4	50
35	Multiple system atrophy: experimental models and reality. Acta Neuropathologica, 2018, 135, 33-47.	3.9	20
36	Translational therapies for multiple system atrophy: Bottlenecks and future directions. Autonomic Neuroscience: Basic and Clinical, 2018, 211, 7-14.	1.4	11

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37	Recommendations of the Global Multiple System Atrophy Research Roadmap Meeting. Neurology, 2018, 90, 74-82.	1.5	23
38	Region-Specific Effects of Immunotherapy With Antibodies Targeting \hat{l}_{\pm} -synuclein in a Transgenic Model of Synucleinopathy. Frontiers in Neuroscience, 2018, 12, 452.	1.4	31
39	Limited effects of dysfunctional macroautophagy on the accumulation of extracellularly derived α-synuclein in oligodendroglia: implications for MSA pathogenesis. BMC Neuroscience, 2018, 19, 32.	0.8	11
40	Very lateâ€onset pure autonomic failure. Movement Disorders, 2017, 32, 1106-1108.	2.2	4
41	Lower Affinity of Isradipine for L-Type Ca ²⁺ Channels during Substantia Nigra Dopamine Neuron-Like Activity: Implications for Neuroprotection in Parkinson's Disease. Journal of Neuroscience, 2017, 37, 6761-6777.	1.7	72
42	Distinct Parameters in the EEG of the PLP α-SYN Mouse Model for Multiple System Atrophy Reinforce Face Validity. Frontiers in Behavioral Neuroscience, 2017, 10, 252.	1.0	14
43	Toll-like receptor 4 stimulation with monophosphoryl lipid A ameliorates motor deficits and nigral neurodegeneration triggered by extraneuronal α-synucleinopathy. Molecular Neurodegeneration, 2017, 12, 52.	4.4	73
44	Anle138b Partly Ameliorates Motor Deficits Despite Failure of Neuroprotection in a Model of Advanced Multiple System Atrophy. Frontiers in Neuroscience, 2016, 10, 99.	1.4	23
45	Changes in the miRNA-mRNA Regulatory Network Precede Motor Symptoms in a Mouse Model of Multiple System Atrophy: Clinical Implications. PLoS ONE, 2016, 11, e0150705.	1.1	26
46	Preface. Movement Disorders, 2016, 31, 151-151.	2.2	0
47	Toward disease modification in multiple system atrophy: Pitfalls, bottlenecks, and possible remedies. Movement Disorders, 2016, 31, 235-240.	2.2	9
48	Review: Multiple system atrophy: emerging targets for interventional therapies. Neuropathology and Applied Neurobiology, 2016, 42, 20-32.	1.8	50
49	Neuroprotection by Epigenetic Modulation in a Transgenic Model of Multiple System Atrophy. Neurotherapeutics, 2016, 13, 871-879.	2.1	17
50	Glia and alpha-synuclein in neurodegeneration: A complex interaction. Neurobiology of Disease, 2016, 85, 262-274.	2.1	156
51	Overexpression of \hat{l}_{\pm} -synuclein in oligodendrocytes does not increase susceptibility to focal striatal excitotoxicity. BMC Neuroscience, 2015, 16, 86.	0.8	5
52	Involvement of Peripheral Nerves in the Transgenic PLP-α-Syn Model of Multiple System Atrophy: Extending the Phenotype. PLoS ONE, 2015, 10, e0136575.	1.1	17
53	Animal models of multiple system atrophy. Clinical Autonomic Research, 2015, 25, 9-17.	1.4	55
54	Shock Wave Treatment Protects From Neuronal Degeneration via a Tollâ€Like Receptor 3 Dependent Mechanism: Implications of a Firstâ€Ever Causal Treatment for Ischemic Spinal Cord Injury. Journal of the American Heart Association, 2015, 4, e002440.	1.6	28

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55	Enteric nervous system α-synuclein immunoreactivity in idiopathic REM sleep behavior disorder. Neurology, 2015, 85, 1761-1768.	1.5	121
56	Alpha-synuclein immunoreactivity patterns in the enteric nervous system. Neuroscience Letters, 2015, 602, 145-149.	1.0	40
57	Failure of Neuroprotection Despite Microglial Suppression by Delayed-Start Myeloperoxidase Inhibition in a Model of Advanced Multiple System Atrophy: Clinical Implications. Neurotoxicity Research, 2015, 28, 185-194.	1.3	38
58	Animal Models of Multiple-System Atrophy. , 2015, , 887-904.		0
59	Multiple System Atrophy: Genetic or Epigenetic?. Experimental Neurobiology, 2014, 23, 277-291.	0.7	28
60	Towards translational therapies for multiple system atrophy. Progress in Neurobiology, 2014, 118, 19-35.	2.8	35
61	Multiple system atrophy as emerging template for accelerated drug discovery in α-synucleinopathies. Parkinsonism and Related Disorders, 2014, 20, 793-799.	1.1	18
62	The Role of Glia in Alpha-Synucleinopathies. Molecular Neurobiology, 2013, 47, 575-586.	1.9	61
63	Tollâ€like receptor 4 is required for αâ€synuclein dependent activation of microglia and astroglia. Glia, 2013, 61, 349-360.	2.5	542
64	Oligodendroglial alpha-synucleinopathy and MSA-like cardiovascular autonomic failure: Experimental evidence. Experimental Neurology, 2013, 247, 531-536.	2.0	46
65	Models of Multiple System Atrophy. Current Topics in Behavioral Neurosciences, 2013, 22, 369-393.	0.8	16
66	Bladder dysfunction in a transgenic mouse model of multiple system atrophy. Movement Disorders, 2013, 28, 347-355.	2.2	50
67	Intact Olfaction in a Mouse Model of Multiple System Atrophy. PLoS ONE, 2013, 8, e64625.	1.1	20
68	CaV1.2 Calcium Channel Expression in Reactive Astrocytes is associated with the Formation of Amyloid-Î ² Plaques in an Alzheimer's Disease Mouse Model. Journal of Alzheimer's Disease, 2013, 37, 439-451.	1.2	39
69	Cell Fate Analysis of Embryonic Ventral Mesencephalic Grafts in the 6-OHDA Model of Parkinson's Disease. PLoS ONE, 2012, 7, e50178.	1.1	1
70	Systemic proteasome inhibition triggers neurodegeneration in a transgenic mouse model expressing human \hat{l}_{\pm} -synuclein under oligodendrocyte promoter: implications for multiple system atrophy. Acta Neuropathologica, 2012, 124, 51-65.	3.9	73
71	Myeloperoxidase Inhibition Ameliorates Multiple System Atrophy-Like Degeneration in a Transgenic Mouse Model. Neurotoxicity Research, 2012, 21, 393-404.	1.3	96
72	Kappa opioid receptor activation blocks progressive neurodegeneration after kainic acid injection. Hippocampus, 2011, 21, 1010-1020.	0.9	20

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73	Toll-Like Receptor 4 Promotes α-Synuclein Clearance and Survival of Nigral Dopaminergic Neurons. American Journal of Pathology, 2011, 179, 954-963.	1.9	230
74	Glial dysfunction in the pathogenesis of \hat{l}_{\pm} -synucleinopathies: emerging concepts. Acta Neuropathologica, 2011, 121, 675-693.	3.9	164
75	Nogo-B is associated with cytoskeletal structures in human monocyte-derived macrophages. BMC Research Notes, 2011, 4, 6.	0.6	20
76	Erythropoietin is neuroprotective in a transgenic mouse model of multiple system atrophy. Movement Disorders, 2011, 26, 507-515.	2.2	17
77	Mesenchymal Stem Cells in a Transgenic Mouse Model of Multiple System Atrophy: Immunomodulation and Neuroprotection. PLoS ONE, 2011, 6, e19808.	1.1	77
78	Etiology, Pathology, and Pathogenesis. Blue Books of Neurology, 2010, 34, 321-339.	0.1	2
79	Targeted overexpression of human α-synuclein in oligodendroglia induces lesions linked to MSA -like progressive autonomic failure. Experimental Neurology, 2010, 224, 459-464.	2.0	65
80	Multiple System Atrophy: Animal Models. , 2010, , 229-232.		0
81	Multiple system atrophy: an update. Lancet Neurology, The, 2009, 8, 1172-1178.	4.9	406
82	Striatal transplantation in a rodent model of multiple system atrophy: Effects on Lâ€Dopa response. Journal of Neuroscience Research, 2009, 87, 1679-1685.	1.3	23
83	Mitochondrial inhibitor 3â€nitroproprionic acid enhances oxidative modification of alphaâ€synuclein in a transgenic mouse model of multiple system atrophy. Journal of Neuroscience Research, 2009, 87, 2728-2739.	1.3	78
84	Recent developments in multiple system atrophy. Journal of Neurology, 2009, 256, 1791-1808.	1.8	86
85	Striatal transplantation for multiple system atrophy â€" Are grafts affected by α-synucleinopathy?. Experimental Neurology, 2009, 219, 368-371.	2.0	28
86	Multiple system atrophy: A primary oligodendrogliopathy. Annals of Neurology, 2008, 64, 239-246.	2.8	279
87	Rasagiline is neuroprotective in a transgenic model of multiple system atrophy. Experimental Neurology, 2008, 210, 421-427.	2.0	79
88	Axl and Growth Arrest–Specific Gene 6 Are Frequently Overexpressed in Human Gliomas and Predict		
	Poor Prognosis in Patients with Glioblastoma Multiforme. Clinical Cancer Research, 2008, 14, 130-138.	3.2	240
89		2.2	13

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91	High dose levodopa therapy is not toxic in multiple system atrophy: Experimental evidence. Movement Disorders, 2007, 22, 969-973.	2.2	6
92	Animal models of multiple system atrophy. Trends in Neurosciences, 2005, 28, 501-506.	4.2	77
93	In vitro models of multiple system atrophy. Movement Disorders, 2005, 20, S53-S56.	2.2	22
94	Failure of caspase inhibition in the double-lesion rat model of striatonigral degeneration (multiple) Tj ETQq0 0 C	rgBT/Ove	erlock 10 Tf 50
95	Riluzole improves motor deficits and attenuates loss of striatal neurons in a sequential double lesion rat model of striatonigral degeneration (parkinson variant of multiple system atrophy). Journal of Neural Transmission, 2005, 112, 1025-1033.	1.4	34
96	Evaluation of [1231]IBZM pinhole SPECT for the detection of striatal dopamine D2 receptor availability in rats. NeuroImage, 2005, 24, 822-831.	2.1	24
97	Evidence for dopaminergic re-innervation by embryonic allografts in an optimized rat model of the Parkinsonian variant of multiple system atrophy. Brain Research Bulletin, 2005, 68, 54-58.	1.4	11
98	Oxidative Stress in Transgenic Mice with Oligodendroglial $\hat{l}\pm$ -Synuclein Overexpression Replicates the Characteristic Neuropathology of Multiple System Atrophy. American Journal of Pathology, 2005, 166, 869-876.	1.9	191
99	Double-Lesion Animal Models of Multiple System Atrophy. , 2005, , 571-583.		O
100	Neuroprotective agents for clinical trials in Parkinson's disease: A systematic assessment. Neurology, 2004, 62, 158-159.	1.5	18
101	Failure of neuronal protection by inhibition of glial activation in a rat model of striatonigral degeneration. Journal of Neuroscience Research, 2004, 78, 87-91.	1.3	28
102	Effects of pulsatile l-DOPA treatment in the double lesion rat model of striatonigral degeneration (multiple system atrophy). Neurobiology of Disease, 2004, 15, 630-639.	2.1	28
103	Neuropathological and behavioral changes induced by various treatment paradigms with MPTP and 3-nitropropionic acid in mice: towards a model of striatonigral degeneration (multiple system) Tj ETQq $1\ 1\ 0.784$	431 4. øgBT	/Overlock 10
104	Tumor necrosis factor-?-induced cell death in U373 cells overexpressing ?-synuclein. Journal of Neuroscience Research, 2003, 73, 334-340.	1.3	37
105	Dystonia is predictive of subsequent altered dopaminergic responsiveness in a chronic 1-methyl-4-phenyl-1,2,3,6-tetrahydropyridine+3-nitropropionic acid model of striatonigral degeneration in monkeys. Neuroscience Letters, 2002, 335, 34-38.	1.0	33
106	Subacute systemic 3-nitropropionic acid intoxication induces a distinct motor disorder in adult C57Bl/6 mice: behavioural and histopathological characterisation. Neuroscience, 2002, 114, 1005-1017.	1.1	147
107	Ultrastructure of \hat{l} ±-synuclein-positive aggregations in U373 astrocytoma and rat primary glial cells. Neuroscience Letters, 2002, 323, 37-40.	1.0	19
108	Freezing of gait in postmortem-confirmed atypical parkinsonism. Movement Disorders, 2002, 17, 1041-1045.	2.2	46

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109	Glial cell death induced by overexpression of ?-synuclein. Journal of Neuroscience Research, 2001, 65, 432-438.	1.3	87
110	Overstimulation of the α1B-adrenergic receptor causes a "seizure plus―syndrome. Nature Medicine, 2001, 7, 132-132.	15.2	5
111	Multiple System Atrophy. Seminars in Neurology, 2001, 21, 033-040.	0.5	17
112	Failure of Neuroprotection by Embryonic Striatal Grafts in a Double Lesion Rat Model of Striatonigral Degeneration (Multiple System Atrophy). Experimental Neurology, 2000, 164, 166-175.	2.0	16
113	Sexual Dimorphism of the Bed Nucleus of the Stria Terminalis and the Amygdala. Advances in Anatomy, Embryology and Cell Biology, 2000, 158, III-X, 1-78.	1.0	22
114	Depression in alpha-synucleinopathies: prevalence, pathophysiology and treatment., 2000,, 335-343.		22
115	γ-Aminobutyric acid-immunoreactive neurons in the amygdala of the rat – sex differences and effect of early postnatal castration. Neuroscience Letters, 1998, 255, 175-177.	1.0	27
116	Sex and age differences of neurons expressing gaba-immunoreactivity in the rat bed nucleus of the stria terminalis. International Journal of Developmental Neuroscience, 1998, 16, 443-448.	0.7	12
117	Distribution of GABA-immunoreactive nerve cells in the bed nucleus of the stria terminalis in male and female rats. European Journal of Histochemistry, 1997, 41, 23-8.	0.6	17