

# Anthony E Kline

## List of Publications by Year in descending order

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68  
papers

3,137  
citations

126708

33  
h-index

155451

55  
g-index

70  
all docs

70  
docs citations

70  
times ranked

1846  
citing authors

#	ARTICLE	IF	CITATIONS
1	Persistent cognitive dysfunction after traumatic brain injury: A dopamine hypothesis. <i>Neuroscience and Biobehavioral Reviews</i> , 2009, 33, 981-1003.	2.9	221
2	Attenuation of Working Memory and Spatial Acquisition Deficits after a Delayed and Chronic Bromocriptine Treatment Regimen in Rats Subjected to Traumatic Brain Injury by Controlled Cortical Impact. <i>Journal of Neurotrauma</i> , 2002, 19, 415-425.	1.7	142
3	Chronic methylphenidate treatment enhances water maze performance following traumatic brain injury in rats. <i>Neuroscience Letters</i> , 2000, 280, 163-166.	1.0	112
4	Administration of haloperidol and risperidone after neurobehavioral testing hinders the recovery of traumatic brain injury-induced deficits. <i>Life Sciences</i> , 2008, 83, 602-607.	2.0	100
5	Empirical Comparison of Typical and Atypical Environmental Enrichment Paradigms on Functional and Histological Outcome after Experimental Traumatic Brain Injury. <i>Journal of Neurotrauma</i> , 2010, 27, 1047-1057.	1.7	100
6	Emerging Therapies in Traumatic Brain Injury. <i>Seminars in Neurology</i> , 2015, 35, 083-100.	0.5	100
7	Acute treatment with the 5-HT1A receptor agonist 8-OH-DPAT and chronic environmental enrichment confer neurobehavioral benefit after experimental brain trauma. <i>Behavioural Brain Research</i> , 2007, 177, 186-194.	1.2	99
8	Chronic administration of antipsychotics impede behavioral recovery after experimental traumatic brain injury. <i>Neuroscience Letters</i> , 2008, 448, 263-267.	1.0	85
9	Bromocriptine Reduces Lipid Peroxidation and Enhances Spatial Learning and Hippocampal Neuron Survival in a Rodent Model of Focal Brain Trauma. <i>Journal of Neurotrauma</i> , 2004, 21, 1712-1722.	1.7	84
10	Gender associations with chronic methylphenidate treatment and behavioral performance following experimental traumatic brain injury. <i>Behavioural Brain Research</i> , 2007, 181, 200-209.	1.2	84
11	Environmental Enrichment as a Viable Neurorehabilitation Strategy for Experimental Traumatic Brain Injury. <i>Journal of Neurotrauma</i> , 2014, 31, 873-888.	1.7	82
12	Amantadine improves water maze performance without affecting motor behavior following traumatic brain injury in rats. <i>Restorative Neurology and Neuroscience</i> , 1999, 14, 285-294.	0.4	79
13	Found in translation: Understanding the biology and behavior of experimental traumatic brain injury. <i>Neuroscience and Biobehavioral Reviews</i> , 2015, 58, 123-146.	2.9	75
14	Combination therapies for neurobehavioral and cognitive recovery after experimental traumatic brain injury: Is more better?. <i>Progress in Neurobiology</i> , 2016, 142, 45-67.	2.8	75
15	Evaluation of a Combined Therapeutic Regimen of 8-OH-DPAT and Environmental Enrichment after Experimental Traumatic Brain Injury. <i>Journal of Neurotrauma</i> , 2010, 27, 2021-2032.	1.7	73
16	Environmental enrichment promotes robust functional and histological benefits in female rats after controlled cortical impact injury. <i>Experimental Neurology</i> , 2013, 247, 410-418.	2.0	68
17	Environmental enrichment-mediated functional improvement after experimental traumatic brain injury is contingent on task-specific neurobehavioral experience. <i>Neuroscience Letters</i> , 2008, 431, 226-230.	1.0	67
18	Acute systemic administration of interleukin-10 suppresses the beneficial effects of moderate hypothermia following traumatic brain injury in rats. <i>Brain Research</i> , 2002, 937, 22-31.	1.1	66

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19	Temporal Effects of Environmental Enrichment Mediated Functional Improvement After Experimental Traumatic Brain Injury in Rats. <i>Neurorehabilitation and Neural Repair</i> , 2011, 25, 558-564.	1.4	66
20	Time dependent alterations in dopamine tissue levels and metabolism after experimental traumatic brain injury in rats. <i>Neuroscience Letters</i> , 2004, 372, 127-131.	1.0	64
21	Divergent Long-Term Consequences of Chronic Treatment with Haloperidol, Risperidone, and Bromocriptine on Traumatic Brain Injury Induced Cognitive Deficits. <i>Journal of Neurotrauma</i> , 2015, 32, 590-597.	1.7	64
22	Methylphenidate treatment following ablation-induced hemiplegia in rat: Experience during drug action alters effects on recovery of function. <i>Pharmacology Biochemistry and Behavior</i> , 1994, 48, 773-779.	1.3	62
23	Abbreviated Environmental Enrichment Enhances Neurobehavioral Recovery Comparably to Continuous Exposure After Traumatic Brain Injury. <i>Neurorehabilitation and Neural Repair</i> , 2011, 25, 343-350.	1.4	61
24	Differential effects of single versus multiple administrations of haloperidol and risperidone on functional outcome after experimental brain trauma. <i>Critical Care Medicine</i> , 2007, 35, 919-924.	0.4	59
25	A Relatively Brief Exposure to Environmental Enrichment after Experimental Traumatic Brain Injury Confers Long-Term Cognitive Benefits. <i>Journal of Neurotrauma</i> , 2012, 29, 2684-2688.	1.7	58
26	The Therapeutic Efficacy Conferred by the 5-HT <sub>1A</sub> Receptor Agonist 8-Hydroxy-2-(di-n-propylamino)tetralin (8-OH-DPAT) after Experimental Traumatic Brain Injury Is Not Mediated by Concomitant Hypothermia. <i>Journal of Neurotrauma</i> , 2004, 21, 175-185.	1.7	55
27	Old Dog, New Tricks: The Attentional Set-Shifting Test as a Novel Cognitive Behavioral Task after Controlled Cortical Impact Injury. <i>Journal of Neurotrauma</i> , 2014, 31, 926-937.	1.7	54
28	A delayed and chronic treatment regimen with the 5-HT <sub>1A</sub> receptor agonist 8-OH-DPAT after cortical impact injury facilitates motor recovery and acquisition of spatial learning. <i>Behavioural Brain Research</i> , 2008, 194, 79-85.	1.2	51
29	Deciphering of Mitochondrial Cardiolipin Oxidative Signaling in Cerebral Ischemia-Reperfusion. <i>Journal of Cerebral Blood Flow and Metabolism</i> , 2015, 35, 319-328.	2.4	51
30	Protective effects of the 5-HT <sub>1A</sub> receptor agonist 8-hydroxy-2-(di-n-propylamino)tetralin against traumatic brain injury-induced cognitive deficits and neuropathology in adult male rats. <i>Neuroscience Letters</i> , 2002, 333, 179-182.	1.0	50
31	Traumatic Brain Injury-Induced Cognitive and Histological Deficits Are Attenuated by Delayed and Chronic Treatment with the 5-HT <sub>1A</sub> -Receptor Agonist Buspirone. <i>Journal of Neurotrauma</i> , 2012, 29, 1898-1907.	1.7	47
32	Evaluation of a Combined Treatment Paradigm Consisting of Environmental Enrichment and the 5-HT <sub>1A</sub> Receptor Agonist Buspirone after Experimental Traumatic Brain Injury. <i>Journal of Neurotrauma</i> , 2012, 29, 1960-1969.	1.7	46
33	A Combined Therapeutic Regimen of Buspirone and Environmental Enrichment Is More Efficacious than Either Alone in Enhancing Spatial Learning in Brain-Injured Pediatric Rats. <i>Journal of Neurotrauma</i> , 2014, 31, 1934-1941.	1.7	37
34	Donepezil Is Ineffective in Promoting Motor and Cognitive Benefits after Controlled Cortical Impact Injury in Male Rats. <i>Journal of Neurotrauma</i> , 2013, 30, 557-564.	1.7	33
35	Traumatic injury compromises nucleocytoplasmic transport and leads to TDP-43 pathology. <i>ELife</i> , 2021, 10, .	2.8	33
36	Paths to Successful Translation of New Therapies for Severe Traumatic Brain Injury in the Golden Age of Traumatic Brain Injury Research: A Pittsburgh Vision. <i>Journal of Neurotrauma</i> , 2020, 37, 2353-2371.	1.7	31

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37	Combining the Antipsychotic Drug Haloperidol and Environmental Enrichment after Traumatic Brain Injury Is a Double-Edged Sword. <i>Journal of Neurotrauma</i> , 2017, 34, 451-458.	1.7	30
38	The neurobehavioral benefit conferred by a single systemic administration of 8-OH-DPAT after brain trauma is confined to a narrow therapeutic window. <i>Neuroscience Letters</i> , 2007, 416, 165-168.	1.0	28
39	Biologic and Plastic Effects of Experimental Traumatic Brain Injury Treatment Paradigms and Their Relevance to Clinical Rehabilitation. <i>PM and R</i> , 2011, 3, S18-27.	0.9	28
40	5-hydroxytryptamine 1A (5-HT 1A ) receptor agonists: A decade of empirical evidence supports their use as an efficacious therapeutic strategy for brain trauma. <i>Brain Research</i> , 2016, 1640, 5-14.	1.1	28
41	Environmental enrichment, alone or in combination with various pharmacotherapies, confers marked benefits after traumatic brain injury. <i>Neuropharmacology</i> , 2019, 145, 13-24.	2.0	28
42	Abbreviated environmental enrichment confers neurobehavioral, cognitive, and histological benefits in brain-injured female rats. <i>Experimental Neurology</i> , 2016, 286, 61-68.	2.0	27
43	The Therapeutic Efficacy of Environmental Enrichment and Methylphenidate Alone and in Combination after Controlled Cortical Impact Injury. <i>Journal of Neurotrauma</i> , 2017, 34, 444-450.	1.7	26
44	Elucidating opportunities and pitfalls in the treatment of experimental traumatic brain injury to optimize and facilitate clinical translation. <i>Neuroscience and Biobehavioral Reviews</i> , 2018, 85, 160-175.	2.9	26
45	Refining environmental enrichment to advance rehabilitation based research after experimental traumatic brain injury. <i>Experimental Neurology</i> , 2017, 294, 12-18.	2.0	23
46	Chronic treatment with galantamine rescues reversal learning in an attentional set-shifting test after experimental brain trauma. <i>Experimental Neurology</i> , 2019, 315, 32-41.	2.0	22
47	Galantamine and Environmental Enrichment Enhance Cognitive Recovery after Experimental Traumatic Brain Injury But Do Not Confer Additional Benefits When Combined. <i>Journal of Neurotrauma</i> , 2017, 34, 1610-1622.	1.7	21
48	Elucidating the role of 5-HT1A and 5-HT7 receptors on 8-OH-DPAT-induced behavioral recovery after experimental traumatic brain injury. <i>Neuroscience Letters</i> , 2012, 515, 153-156.	1.0	20
49	Delayed and Abbreviated Environmental Enrichment after Brain Trauma Promotes Motor and Cognitive Recovery That Is Not Contingent on Increased Neurogenesis. <i>Journal of Neurotrauma</i> , 2019, 36, 756-767.	1.7	20
50	Early life stress increases vulnerability to the sequelae of pediatric mild traumatic brain injury. <i>Experimental Neurology</i> , 2020, 329, 113318.	2.0	20
51	Comparable impediment of cognitive function in female and male rats subsequent to daily administration of haloperidol after traumatic brain injury. <i>Experimental Neurology</i> , 2017, 296, 62-68.	2.0	19
52	Preclinical Models of Traumatic Brain Injury: Emerging Role of Glutamate in the Pathophysiology of Depression. <i>Frontiers in Pharmacology</i> , 2018, 9, 579.	1.6	17
53	Early Life Stress Preceding Mild Pediatric Traumatic Brain Injury Increases Neuroinflammation but Does Not Exacerbate Impairment of Cognitive Flexibility during Adolescence. <i>Journal of Neurotrauma</i> , 2021, 38, 411-421.	1.7	17
54	Environmental enrichment and amantadine confer individual but nonadditive enhancements in motor and spatial learning after controlled cortical impact injury. <i>Brain Research</i> , 2019, 1714, 227-233.	1.1	15

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55	Dose-dependent neurorestorative effects of amantadine after cortical impact injury. <i>Neuroscience Letters</i> , 2019, 694, 69-73.	1.0	13
56	Rehabilitative Success After Brain Trauma by Augmenting a Subtherapeutic Dose of Environmental Enrichment With Galantamine. <i>Neurorehabilitation and Neural Repair</i> , 2017, 31, 977-985.	1.4	12
57	Chronic unpredictable stress during adolescence protects against adult traumatic brain injury-induced affective and cognitive deficits. <i>Brain Research</i> , 2021, 1767, 147544.	1.1	11
58	Systemic administration of donepezil attenuates the efficacy of environmental enrichment on neurobehavioral outcome after experimental traumatic brain injury. <i>Restorative Neurology and Neuroscience</i> , 2018, 36, 45-57.	0.4	9
59	Intermittent treatment with haloperidol or quetiapine does not disrupt motor and cognitive recovery after experimental brain trauma. <i>Behavioural Brain Research</i> , 2018, 340, 159-164.	1.2	9
60	Disruption of basal forebrain cholinergic neurons after traumatic brain injury does not compromise environmental enrichment-mediated cognitive benefits. <i>Brain Research</i> , 2021, 1751, 147175.	1.1	7
61	Aripiprazole and environmental enrichment independently improve functional outcome after cortical impact injury in adult male rats, but their combination does not yield additional benefits. <i>Experimental Neurology</i> , 2019, 314, 67-73.	2.0	6
62	Intranasally Administered L-Myc-Immortalized Human Neural Stem Cells Migrate to Primary and Distal Sites of Damage after Cortical Impact and Enhance Spatial Learning. <i>Stem Cells International</i> , 2021, 2021, 1-11.	1.2	5
63	Intermittent Administration of Haloperidol after Cortical Impact Injury Neither Impedes Spontaneous Recovery Nor Attenuates the Efficacy of Environmental Enrichment. <i>Journal of Neurotrauma</i> , 2019, 36, 1606-1614.	1.7	4
64	Albeit nocturnal, rats subjected to traumatic brain injury do not differ in neurobehavioral performance whether tested during the day or night. <i>Neuroscience Letters</i> , 2018, 665, 212-216.	1.0	3
65	Spontaneous recovery of traumatic brain injury-induced functional deficits is not hindered by daily administration of lorazepam. <i>Behavioural Brain Research</i> , 2018, 339, 215-221.	1.2	3
66	Preclinical neurorehabilitation with environmental enrichment confers cognitive and histological benefits in a model of pediatric asphyxial cardiac arrest. <i>Experimental Neurology</i> , 2021, 335, 113522.	2.0	3
67	Spontaneous recovery after controlled cortical impact injury is not impeded by intermittent administration of the antipsychotic drug risperidone. <i>Neuroscience Letters</i> , 2018, 682, 69-73.	1.0	2
68	Brain injury and recovery. <i>Brain Research</i> , 2016, 1640, 1-4.	1.1	0