

Cora H Nijboer

List of Publications by Year in descending order

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Version: 2024-02-01

37
papers

2,142
citations

218677

26
h-index

345221

36
g-index

37
all docs

37
docs citations

37
times ranked

3236
citing authors

#	ARTICLE	IF	CITATIONS
1	Cognitive performance during adulthood in a rat model of neonatal diffuse white matter injury. <i>Psychopharmacology</i> , 2022, 239, 745.	3.1	0
2	Nutritional Supplementation Reduces Lesion Size and Neuroinflammation in a Sex-Dependent Manner in a Mouse Model of Perinatal Hypoxic-Ischemic Brain Injury. <i>Nutrients</i> , 2022, 14, 176.	4.1	7
3	Feasibility and safety of intranasally administered mesenchymal stromal cells after perinatal arterial ischaemic stroke in the Netherlands (PASSION): a first-in-human, open-label intervention study. <i>Lancet Neurology</i> , The, 2022, 21, 528-536.	10.2	50
4	Intranasal mesenchymal stem cell therapy to boost myelination after encephalopathy of prematurity. <i>Glia</i> , 2021, 69, 655-680.	4.9	18
5	Nasal administration of mesenchymal stem cells reverses chemotherapy-induced peripheral neuropathy in mice. <i>Brain, Behavior, and Immunity</i> , 2021, 93, 43-54.	4.1	23
6	Neuroprotection offered by mesenchymal stem cells in perinatal brain injury: Role of mitochondria, inflammation, and reactive oxygen species. <i>Journal of Neurochemistry</i> , 2021, 158, 59-73.	3.9	38
7	The impact of trophic and immunomodulatory factors on oligodendrocyte maturation: Potential treatments for encephalopathy of prematurity. <i>Glia</i> , 2021, 69, 1311-1340.	4.9	10
8	Chronic social stress lessens the metabolic effects induced by a high-fat diet. <i>Journal of Endocrinology</i> , 2021, 249, 19-30.	2.6	4
9	Regenerative Therapies to Restore Interneuron Disturbances in Experimental Models of Encephalopathy of Prematurity. <i>International Journal of Molecular Sciences</i> , 2021, 22, 211.	4.1	8
10	Postnatal Nutrition to Improve Brain Development in the Preterm Infant: A Systematic Review From Bench to Bedside. <i>Frontiers in Physiology</i> , 2019, 10, 961.	2.8	31
11	Intranasal Stem Cell Treatment as a Novel Therapy for Subarachnoid Hemorrhage. <i>Stem Cells and Development</i> , 2018, 27, 313-325.	2.1	45
12	Origin and dynamics of oligodendrocytes in the developing brain: Implications for perinatal white matter injury. <i>Glia</i> , 2018, 66, 221-238.	4.9	188
13	SOX4 inhibits oligodendrocyte differentiation of embryonic neural stem cells in vitro by inducing Hes5 expression. <i>Stem Cell Research</i> , 2018, 33, 110-119.	0.7	29
14	Forkhead box protein P1, a key player in neuronal development?. <i>Neural Regeneration Research</i> , 2018, 13, 801.	3.0	3
15	Repair of neonatal brain injury: bringing stem cell-based therapy into clinical practice. <i>Developmental Medicine and Child Neurology</i> , 2017, 59, 997-1003.	2.1	35
16	A quantitative method for microstructural analysis of myelinated axons in the injured rodent brain. <i>Scientific Reports</i> , 2017, 7, 16492.	3.3	34
17	FOXP1 Promotes Embryonic Neural Stem Cell Differentiation by Repressing Jagged1 Expression. <i>Stem Cell Reports</i> , 2017, 9, 1530-1545.	4.8	56
18	Delayed administration of neural stem cells after hypoxia-ischemia reduces sensorimotor deficits, cerebral lesion size, and neuroinflammation in neonatal mice. <i>Pediatric Research</i> , 2017, 81, 127-135.	2.3	28

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19	Impaired oligodendrocyte maturation in preterm infants: Potential therapeutic targets. <i>Progress in Neurobiology</i> , 2016, 136, 28-49.	5.7	110
20	Prevention of chemotherapy-induced peripheral neuropathy by the small-molecule inhibitor pifithrin- α . <i>Pain</i> , 2015, 156, 2184-2192.	4.2	60
21	The Neonatal Brain Is Not Protected by Osteopontin Peptide Treatment after Hypoxia-Ischemia. <i>Developmental Neuroscience</i> , 2015, 37, 142-152.	2.0	14
22	Development of Cerebral Gray and White Matter Injury and Cerebral Inflammation over Time after Inflammatory Perinatal Asphyxia. <i>Developmental Neuroscience</i> , 2015, 37, 78-94.	2.0	34
23	Assessment of long-term safety and efficacy of intranasal mesenchymal stem cell treatment for neonatal brain injury in the mouse. <i>Pediatric Research</i> , 2015, 78, 520-526.	2.3	74
24	The rodent endovascular puncture model of subarachnoid hemorrhage: mechanisms of brain damage and therapeutic strategies. <i>Journal of Neuroinflammation</i> , 2014, 11, 2.	7.2	77
25	Intranasally administered mesenchymal stem cells promote a regenerative niche for repair of neonatal ischemic brain injury. <i>Experimental Neurology</i> , 2014, 261, 53-64.	4.1	132
26	Long-Term Functional Consequences and Ongoing Cerebral Inflammation after Subarachnoid Hemorrhage in the Rat. <i>PLoS ONE</i> , 2014, 9, e90584.	2.5	70
27	Intranasal Administration of Human MSC for Ischemic Brain Injury in the Mouse: In Vitro and In Vivo Neuroregenerative Functions. <i>PLoS ONE</i> , 2014, 9, e112339.	2.5	76
28	Mitochondrial JNK phosphorylation as a novel therapeutic target to inhibit neuroinflammation and apoptosis after neonatal ischemic brain damage. <i>Neurobiology of Disease</i> , 2013, 54, 432-444.	4.4	67
29	Astrocyte GRK2 as a novel regulator of glutamate transport and brain damage. <i>Neurobiology of Disease</i> , 2013, 54, 206-215.	4.4	17
30	Intranasal Mesenchymal Stem Cell Treatment for Neonatal Brain Damage: Long-Term Cognitive and Sensorimotor Improvement. <i>PLoS ONE</i> , 2013, 8, e51253.	2.5	143
31	Extent of Bilateral Neuronal Network Reorganization and Functional Recovery in Relation to Stroke Severity. <i>Journal of Neuroscience</i> , 2012, 32, 4495-4507.	3.6	208
32	Targeting the p53 pathway to protect the neonatal ischemic brain. <i>Annals of Neurology</i> , 2011, 70, 255-264.	5.3	88
33	Inhibition of the JNK/AP-1 pathway reduces neuronal death and improves behavioral outcome after neonatal hypoxic-ischemic brain injury. <i>Brain, Behavior, and Immunity</i> , 2010, 24, 812-821.	4.1	80
34	Cell-specific roles of GRK2 in onset and severity of hypoxic-ischemic brain damage in neonatal mice. <i>Brain, Behavior, and Immunity</i> , 2010, 24, 420-426.	4.1	31
35	GRK2: A Novel Cell-Specific Regulator of Severity and Duration of Inflammatory Pain. <i>Journal of Neuroscience</i> , 2010, 30, 2138-2149.	3.6	103
36	Alternate Pathways Preserve Tumor Necrosis Factor- α Production After Nuclear Factor- κ B Inhibition in Neonatal Cerebral Hypoxia-Ischemia. <i>Stroke</i> , 2009, 40, 3362-3368.	2.0	50

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37	A Dual Role of the NF- κ B Pathway in Neonatal Hypoxic-Ischemic Brain Damage. <i>Stroke</i> , 2008, 39, 2578-2586.	2.0	101