## Reinhard Dengler

List of Publications by Year in descending order

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236925 197818 2,560 61 25 49 citations h-index g-index papers 62 62 62 3308 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	Electrodiagnostic criteria for diagnosis of ALS. Clinical Neurophysiology, 2008, 119, 497-503.	1.5	927
2	Early treatment with aspirin plus extended-release dipyridamole for transient ischaemic attack or ischaemic stroke within 24 h of symptom onset (EARLY trial): a randomised, open-label, blinded-endpoint trial. Lancet Neurology, The, 2010, 9, 159-166.	10.2	111
3	4-Aminopyridine Induced Activity Rescues Hypoexcitable Motor Neurons from Amyotrophic Lateral Sclerosis Patient-Derived Induced Pluripotent Stem Cells. Stem Cells, 2016, 34, 1563-1575.	3.2	109
4	Basal ganglia pathology in ALS is associated with neuropsychological deficits. Neurology, 2015, 85, 1301-1309.	1.1	96
5	Alterations in the hypothalamic melanocortin pathway in amyotrophic lateral sclerosis. Brain, 2016, 139, 1106-1122.	7.6	80
6	Human brain connectivity: Clinical applications for clinical neurophysiology. Clinical Neurophysiology, 2020, 131, 1621-1651.	1.5	68
7	Hippocampal degeneration in patients with amyotrophic lateral sclerosis. Neurobiology of Aging, 2014, 35, 2639-2645.	3.1	62
8	Quantitative Susceptibility MRI to Detect Brain Iron in Amyotrophic Lateral Sclerosis. Radiology, 2018, 289, 195-203.	7.3	61
9	Peripheral nerve ultrasound in amyotrophic lateral sclerosis phenotypes. Muscle and Nerve, 2015, 51, 669-675.	2.2	55
10	Structural and diffusion imaging versus clinical assessment to monitor amyotrophic lateral sclerosis. NeuroImage: Clinical, 2016, 11, 408-414.	2.7	51
11	Interaction of physical function, quality of life and depression in Amyotrophic lateral sclerosis: characterization of a large patient cohort. BMC Neurology, 2015, 15, 84.	1.8	49
12	Lower motor neuron involvement in ALS assessed by motor unit number index (MUNIX): Long-term changes and reproducibility. Clinical Neurophysiology, 2016, 127, 1984-1988.	1.5	45
13	Significance of CSF NfL and tau in ALS. Journal of Neurology, 2018, 265, 2633-2645.	3.6	45
14	Supraglottal Injection of Botulinum Toxin Type A in Adductor Type Spasmodic Dysphonia With Both Intrinsic and Extrinsic Hyperfunction. Laryngoscope, 1998, 108, 55-63.	2.0	40
15	Quantifying disease progression in amyotrophic lateral sclerosis using peripheral nerve sonography. Muscle and Nerve, 2016, 54, 391-397.	2.2	40
16	Enhanced audio–visual interactions in the auditory cortex of elderly cochlear-implant users. Hearing Research, 2015, 328, 133-147.	2.0	37
17	Neural mechanisms underlying cognitive inflexibility in Parkinson's disease. Neuropsychologia, 2016, 93, 142-150.	1.6	37
18	Prior probabilities modulate cortical surprise responses: A study of event-related potentials. Brain and Cognition, 2016, 106, 78-89.	1.8	35

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19	Decomposing card-sorting performance: Effects of working memory load and age-related changes Neuropsychology, 2016, 30, 579-590.	1.3	34
20	Impaired set-shifting in amyotrophic lateral sclerosis: An event-related potential study of executive function Neuropsychology, 2016, 30, 120-134.	1.3	33
21	Dual routes to cortical orienting responses: Novelty detection and uncertainty reduction. Biological Psychology, 2015, 105, 66-71.	2.2	32
22	Meta-analytical and electrophysiological evidence for executive dysfunction in primary dystonia. Cortex, 2016, 82, 133-146.	2.4	32
23	Auditory and audio–visual processing in patients with cochlear, auditory brainstem, and auditory midbrain implants: An <scp>EEG</scp> study. Human Brain Mapping, 2017, 38, 2206-2225.	3.6	32
24	Residual Neural Processing of Musical Sound Features in Adult Cochlear Implant Users. Frontiers in Human Neuroscience, 2014, 8, 181.	2.0	31
25	The Axon Guidance Protein Semaphorin 3A Is Increased in the Motor Cortex of Patients With Amyotrophic Lateral Sclerosis. Journal of Neuropathology and Experimental Neurology, 2016, 75, 326-333.	1.7	29
26	Amyotrophic lateral sclerosis affects cortical and subcortical activity underlying motor inhibition and action monitoring. Human Brain Mapping, 2015, 36, 2878-2889.	3.6	27
27	Cognitive Flexibility in Primary Dystonia. Journal of the International Neuropsychological Society, 2016, 22, 662-670.	1.8	26
28	Succinylcholine Metabolite Succinic Acid Alters Steady State Activation in Muscle Sodium Channels. Anesthesiology, 2000, 92, 1385-1391.	2.5	23
29	Circulating Insulin-like Growth Factor-1 and Insulin-like Growth Factor Binding Protein-3 predict Three-months Outcome after Ischemic Stroke. Experimental and Clinical Endocrinology and Diabetes, 2017, 125, 485-491.	1.2	22
30	Dopaminergic modulation of performance monitoring in Parkinson's disease: An event-related potential study. Scientific Reports, 2017, 7, 41222.	3.3	21
31	Toward <i>in vivo</i> determination of peripheral nervous system immune activity in amyotrophic lateral sclerosis. Muscle and Nerve, 2019, 59, 567-576.	2.2	21
32	Attention shifting in Parkinson's disease: An analysis of behavioral and cortical responses Neuropsychology, 2014, 28, 929-944.	1.3	20
33	The anesthetic propofol modulates gating in paramyotonia congenita mutant muscle sodium channels. Muscle and Nerve, 2001, 24, 736-743.	2.2	19
34	Intraspinal administration of human spinal cord-derived neural progenitor cells in the <i>G93A-SOD1 </i> mouse model of ALS delays symptom progression, prolongs survival and increases expression of endogenous neurotrophic factors. Journal of Tissue Engineering and Regenerative Medicine, 2017, 11, 751-764.	2.7	19
35	Peripheral nerve atrophy together with higher cerebrospinal fluid progranulin indicate axonal damage in amyotrophic lateral sclerosis. Muscle and Nerve, 2018, 57, 273-278.	2.2	17
36	Neural correlates of cognitive set shifting in amyotrophic lateral sclerosis. Clinical Neurophysiology, 2016, 127, 3537-3545.	1.5	16

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37	Diagnostic support for selected neuromuscular diseases using answer-pattern recognition and data mining techniques: a proof of concept multicenter prospective trial. BMC Medical Informatics and Decision Making, 2016, $16, 31$ .	3.0	15
38	Human centromedian-parafascicular complex signals sensory cues for goal-oriented behavior selection. NeuroImage, 2017, 152, 390-399.	4.2	15
39	Dyspnea in Amyotrophic Lateral Sclerosis: Rasch-Based Development and Validation of a Patient-Reported Outcome (DALS-15). Journal of Pain and Symptom Management, 2018, 56, 736-745.e2.	1.2	14
40	Interhemispheric connectivity in amyotrophic lateral sclerosis: A near-infrared spectroscopy and diffusion tensor imaging study. Neurolmage: Clinical, 2016, 12, 666-672.	2.7	11
41	Sonographic and 3T-MRI-based evaluation of the tongue in ALS. NeuroImage: Clinical, 2020, 26, 102233.	2.7	11
42	Valence-specific conflict moderation in the dorso-medial PFC and the caudate head in emotional speech. Social Cognitive and Affective Neuroscience, 2015, 10, 165-171.	3.0	10
43	Differential involvement of forearm muscles in ALS does not relate to sonographic structural nerve alterations. Clinical Neurophysiology, 2018, 129, 1438-1443.	1.5	9
44	AANEM – IFCN glossary of terms in neuromuscular electrodiagnostic medicine and ultrasound. Clinical Neurophysiology, 2020, 131, 1662-1663.	1.5	8
45	Dyspnea as a Fatigue-Promoting Factor in ALS and the Role of Objective Indicators of Respiratory Impairment. Journal of Pain and Symptom Management, 2020, 60, 430-438.e1.	1.2	8
46	CLIPPERS Syndrome: An Entity to be Faced in Neurosurgery. World Neurosurgery, 2015, 84, 2077.e1-2077.e3.	1.3	7
47	Attenuated error-related potentials in amyotrophic lateral sclerosis with executive dysfunctions. Clinical Neurophysiology, 2017, 128, 1496-1503.	1.5	7
48	The upper cervical spinal cord in ALS assessed by cross-sectional and longitudinal 3T MRI. Scientific Reports, 2020, 10, 1783.	3.3	7
49	Dopaminergic modulation of novelty repetition in Parkinson's disease: A study of P3 event-related brain potentials. Clinical Neurophysiology, 2020, 131, 2841-2850.	1.5	6
50	Characteristics of pain and the burden it causes in patients with amyotrophic lateral sclerosis – a longitudinal study. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2021, , 1-8.	1.7	5
51	The Dyspnea-ALS-Scale (DALS-15) optimizes individual treatment in patients with amyotrophic lateral sclerosis (ALS) suffering from dyspnea. Health and Quality of Life Outcomes, 2019, 17, 95.	2.4	4
52	A Multi-Center Cohort Study on Characteristics of Pain, Its Impact and Pharmacotherapeutic Management in Patients with ALS. Journal of Clinical Medicine, 2021, 10, 4552.	2.4	4
53	Ultrasound assessment of nerve and nerve root thickness. Do they contribute to the diagnosis of ALS?. Clinical Neurophysiology, 2014, 125, 1719-1720.	1.5	3
54	Intraspinal cavernous bleeding during early pregnancy. Journal of Neurology, 2016, 263, 2127-2129.	3.6	3

#	Article	lF	CITATION
55	Lack of an association between attention-deficit/hyperactivity disorder (ADHD) and amyotrophic lateral sclerosis (ALS). Journal of the Neurological Sciences, 2018, 385, 7-11.	0.6	2
56	Letters to the editor. Muscle and Nerve, 1993, 16, 876-885.	2.2	1
57	Upper motor neuron involvement in amyotrophic lateral sclerosis. Do we have a new diagnostic tool?. Clinical Neurophysiology, 2021, 132, 618-619.	1.5	1
58	Pain-Related Coping Behavior in ALS: The Interplay between Maladaptive Coping, the Patient's Affective State and Pain. Journal of Clinical Medicine, 2022, 11, 944.	2.4	1
59	Guidelines for low intensity transcranial electrical stimulation – An overdue step in a fairly uncontrolled field. Clinical Neurophysiology, 2017, 128, 1770-1771.	1.5	0
60	Association between attention-deficit/hyperactivity disorder (ADHD) and amyotrophic lateral sclerosis (ALS). Journal of the Neurological Sciences, 2018, 391, 152.	0.6	0
61	Diagnostic criteria of ALS. Are the Gold Coast Criteria the ultimate solution?. Clinical Neurophysiology, 2021, 132, 3177-3178.	1.5	0