

# Cognitive and clinical characteristics of patients with an carrying a C9orf72 repeat expansion: a population-based

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Citation Report

#	ARTICLE	IF	CITATIONS
1	Treatment and Management of Adult Motor Neuron Diseases. , 2011, , 169-178.		0
2	FTD/ALS families are no longer orphaned. <i>Neurology</i> , 2012, 79, 962-964.	1.5	4
3	Amyotrophic lateral sclerosis. <i>Current Opinion in Neurology</i> , 2012, 25, 530-535.	1.8	85
4	How do C9ORF72 repeat expansions cause amyotrophic lateral sclerosis and frontotemporal dementia. <i>Current Opinion in Neurology</i> , 2012, 25, 689-700.	1.8	169
5	Young-onset amyotrophic lateral sclerosis: historical and other observations. <i>Brain</i> , 2012, 135, 2883-2891.	3.7	65
6	Transmission of C9orf72 hexanucleotide repeat expansions in sporadic amyotrophic lateral sclerosis. <i>NeuroReport</i> , 2012, 23, 556-559.	0.6	16
7	Ask the Experts: Translating amyotrophic lateral sclerosis genetics to the clinic: implications for the patient. <i>Neurodegenerative Disease Management</i> , 2012, 2, 355-360.	1.2	0
8	The Use of Next-Generation Sequencing in Movement Disorders. <i>Frontiers in Genetics</i> , 2012, 3, 75.	1.1	21
9	Phenotype difference between ALS patients with expanded repeats in C9ORF72 and patients with mutations in other ALS-related genes. <i>Journal of Medical Genetics</i> , 2012, 49, 258-263.	1.5	157
10	Extensive genetics of ALS. <i>Neurology</i> , 2012, 79, 1983-1989.	1.5	145
11	Amyotrophic lateral sclerosis/frontotemporal dementia with predominant manifestations of obsessive-compulsive disorder associated to GGGGCC expansion of the c9orf72 gene. <i>Journal of Neurology</i> , 2012, 259, 2723-2725.	1.8	37
13	Cognitive and behavioral features of c9FTD/ALS. <i>Alzheimer's Research and Therapy</i> , 2012, 4, 29.	3.0	20
14	Expanding the genetics of amyotrophic lateral sclerosis and frontotemporal dementia. <i>Alzheimer's Research and Therapy</i> , 2012, 4, 30.	3.0	6
15	Neuropsychiatric features of C9orf72-associated behavioral variant frontotemporal dementia and frontotemporal dementia with motor neuron disease. <i>Alzheimer's Research and Therapy</i> , 2012, 4, 38.	3.0	30
16	Neuroimaging features of C9ORF72 expansion. <i>Alzheimer's Research and Therapy</i> , 2012, 4, 45.	3.0	29
17	The C9orf72 hexanucleotide repeat expansion in FTD and ALS. <i>Nature Reviews Neurology</i> , 2012, 8, 249-250.	4.9	14
18	A hexanucleotide repeat expansion in C9ORF72 causes familial and sporadic ALS in Taiwan. <i>Neurobiology of Aging</i> , 2012, 33, 2232.e11-2232.e18.	1.5	52
19	Analysis of C9orf72 repeat expansion in 563 Japanese patients with amyotrophic lateral sclerosis. <i>Neurobiology of Aging</i> , 2012, 33, 2527.e11-2527.e16.	1.5	98

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20	C9ORF72 repeat expansion in a large Italian ALS cohort: evidence of a founder effect. <i>Neurobiology of Aging</i> , 2012, 33, 2528.e7-2528.e14.	1.5	74
21	Investigation of C9orf72 in 4 Neurodegenerative Disorders. <i>Archives of Neurology</i> , 2012, 69, 1583.	4.9	89
22	The genetics and neuropathology of amyotrophic lateral sclerosis. <i>Acta Neuropathologica</i> , 2012, 124, 339-352.	3.9	346
23	Databases for neurogenetics: Introduction, overview, and challenges. <i>Human Mutation</i> , 2012, 33, 1311-1314.	1.1	3
24	Awaji Criteria for the Diagnosis of Amyotrophic Lateral Sclerosis. <i>Archives of Neurology</i> , 2012, 69, 1410.	4.9	211
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32	Frequency of the C9orf72 hexanucleotide repeat expansion in patients with amyotrophic lateral sclerosis and frontotemporal dementia: a cross-sectional study. <i>Lancet Neurology</i> , The, 2012, 11, 323-330.	4.9	1,039
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34	New therapy options for amyotrophic lateral sclerosis. <i>Expert Opinion on Pharmacotherapy</i> , 2013, 14, 1907-1917.	0.9	19
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41	Neurodegenerative Diseases: Integrative PPPM Approach as the Medicine of the Future. <i>Advances in Predictive, Preventive and Personalised Medicine</i> , 2013, , .	0.6	3
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43	<i>C9ORF72</i> Intermediate Repeat Copies Are a Significant Risk Factor for Parkinson Disease. <i>Annals of Human Genetics</i> , 2013, 77, 351-363.	0.3	69
44	Controversies and priorities in amyotrophic lateral sclerosis. <i>Lancet Neurology</i> , The, 2013, 12, 310-322.	4.9	454
45	Changes in cognition and behaviour in amyotrophic lateral sclerosis: nature of impairment and implications for assessment. <i>Lancet Neurology</i> , The, 2013, 12, 368-380.	4.9	363
46	UNC13A influences survival in Italian amyotrophic lateral sclerosis patients: a population-based study. <i>Neurobiology of Aging</i> , 2013, 34, 357.e1-357.e5.	1.5	59
47	<i>C9ORF72</i> Repeat Expansions in the Frontotemporal Dementias Spectrum of Diseases: A Flow-chart for Genetic Testing. <i>Journal of Alzheimer's Disease</i> , 2013, 34, 485-499.	1.2	93
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51	Autosomal Dominant Frontotemporal Lobar Degeneration Due to the <i>C9ORF72</i> Hexanucleotide Repeat Expansion: Late-Onset Psychotic Clinical Presentation. <i>Biological Psychiatry</i> , 2013, 74, 384-391.	0.7	105
52	Pathophysiological insights into ALS with <i>C9ORF72</i> expansions. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2013, 84, 931-935.	0.9	89
53	Neural substrates of episodic memory dysfunction in behavioural variant frontotemporal dementia with and without <i>C9ORF72</i> expansions. <i>NeuroImage: Clinical</i> , 2013, 2, 836-843.	1.4	35
54	The changing scene of amyotrophic lateral sclerosis. <i>Nature Reviews Neuroscience</i> , 2013, 14, 248-264.	4.9	860
55	Corticobasal and ataxia syndromes widen the spectrum of <i>C9ORF72</i> hexanucleotide expansion disease. <i>Clinical Genetics</i> , 2013, 83, 279-283.	1.0	128
56	Lack of <i>C9ORF72</i> coding mutations supports a gain of function for repeat expansions in amyotrophic lateral sclerosis. <i>Neurobiology of Aging</i> , 2013, 34, 2234.e13-2234.e19.	1.5	59

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62	Frontotemporal dementia with amyotrophic lateral sclerosis: A clinical comparison of patients with and without repeat expansions in <i>C9orf72</i> . <i>Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration</i> , 2013, 14, 172-176.	1.1	58
63	Grey matter correlates of clinical variables in amyotrophic lateral sclerosis (ALS): a neuroimaging study of ALS motor phenotype heterogeneity and cortical focality. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2013, 84, 766-773.	0.9	121
64	Genetics and Pathophysiology of Neurodegeneration with Brain Iron Accumulation (NBIA). <i>Current Neuropharmacology</i> , 2013, 11, 59-79.	1.4	100
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87	Neurodegenerative disorders. , 0, , 23-71.		0
88	Clinical Characteristics of C9ORF72-Linked Frontotemporal Lobar Degeneration. <i>Dementia and Geriatric Cognitive Disorders Extra</i> , 2013, 3, 251-262.	0.6	29
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92	Altered Intracellular Localization of SOD1 in Leukocytes from Patients with Sporadic Amyotrophic Lateral Sclerosis. <i>PLoS ONE</i> , 2013, 8, e75916.	1.1	38
143	Antisense Therapy in Neurology. <i>Journal of Personalized Medicine</i> , 2013, 3, 144-176.	1.1	53
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190	Genetics of ALS. , 2015, , 385-409.		0
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208	C9ORF72 hexanucleotide repeat expansion in ALS patients from the Central European Russia population. <i>Neurobiology of Aging</i> , 2015, 36, 2908.e5-2908.e9.	1.5	12
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210	Genetic analysis of amyotrophic lateral sclerosis in the Slovenian population. <i>Neurobiology of Aging</i> , 2015, 36, 1601.e17-1601.e20.	1.5	10
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