Ahmed Bouhouche

List of Publications by Year
in descending order

37 papers	1,797 citations	516710 16 h-index	330143 37 g-index
38 all docs	38 docs citations	38 times ranked	3271 citing authors

#	Article	IF	Citations
1	Novel ITGB2 Mutation Is Responsible for a Severe Form of Leucocyte Adhesion Deficiency Type 1. BioMed Research International, 2022, 2022, 1-8.	1.9	2
2	Gene Panel Sequencing Identifies Novel Pathogenic Mutations in Moroccan Patients with Familial Parkinson Disease. Journal of Molecular Neuroscience, 2021, 71, 142-152.	2.3	4
3	Identification of the novel SDR42E1 gene that affects steroid biosynthesis associated with the oculocutaneous genital syndrome. Experimental Eye Research, 2021, 209, 108671.	2.6	3
4	Characterization of Recessive Parkinson Disease in a Large Multicenter Study. Annals of Neurology, 2020, 88, 843-850.	5.3	40
5	Novel pathogenic VPS13A mutation in Moroccan family with Choreoacanthocytosis: a case report. BMC Medical Genetics, 2020, 21, 47.	2.1	1
6	A Specific Diplotype H1j/H2 of the MAPT Gene Could Be Responsible for Parkinsonâ€™s Disease with Dementia. Case Reports in Genetics, 2020, 2020, 1-5.	0.2	2
7	Association of vitamin D status with multiple sclerosis in a case-control study from Morocco. Revue Neurologique, 2018, 174, 150-156.	1.5	14
8	Genetic Analysis of Undiagnosed Juvenile GM1-Gangliosidosis by Microarray and Exome Sequencing. Case Reports in Genetics, 2018, 2018, 1-8.	0.2	3
9	Non-Motor Symptoms of Parkinsonâ€™s Disease and Their Impact on Quality of Life in a Cohort of Moroccan Patients. Frontiers in Neurology, 2018, 9, 170.	2.4	78
10	Deep Brain Stimulation in Moroccan Patients With Parkinson's Disease: The Experience of Neurology Department of Rabat. Frontiers in Neurology, 2018, 9, 532.	2.4	10
11	Mutation Analysis of Consanguineous Moroccan Patients with Parkinsonâ€™s Disease Combining Microarray and Gene Panel. Frontiers in Neurology, 2017, 8, 567.	2.4	19
12	LRRK2 G2019S Mutation: Prevalence and Clinical Features in Moroccans with Parkinsonâ€™s Disease. Parkinson's Disease, 2017, 2017, 1-7.	1.1	36
13	Evidence for prehistoric origins of the G2019S mutation in the North African Berber population. PLoS ONE, 2017, 12, e0181335.	2.5	13
14	Clinical and genetic data of Huntington disease in Moroccan patients. African Health Sciences, 2016, 15, 1232.	0.7	6
15	A Novel Homozygous p.L539F Mutation Identified inPINK1Gene in a Moroccan Patient with Parkinsonism. BioMed Research International, 2016, 2016, 1-5.	1.9	10
16	Mutations in CAPN1 Cause Autosomal-Recessive Hereditary Spastic Paraplegia. American Journal of Human Genetics, 2016, 98, 1038-1046.	6.2	96
17	KIF1Cmutations in two families with hereditary spastic paraparesis and cerebellar dysfunction. Journal of Medical Genetics, 2014, 51, 137-142.	3.2	67
18	Exome Sequencing Links Corticospinal Motor Neuron Disease to Common Neurodegenerative Disorders. Science, 2014, 343, 506-511.	12.6	466

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19	Clinical and Genetic Study of Friedreichâ€™s Ataxia and Ataxia with Vitamin E Deficiency in 44 Moroccan Families. World Journal of Neuroscience, 2014, 04, 299-305.	0.1	0
20	CLN6 p.I154del Mutation Causing Late Infantile Neuronal Ceroid Lipofuscinosis in a Large Consanguineous Moroccan Family. Indian Journal of Pediatrics, 2013, 80, 694-696.	0.8	3
21	Inhabitual autosomal recessive form of dentin dysplasia type I in a large consanguineous Moroccan family. European Journal of Medical Genetics, 2013, 56, 442-444.	1.3	11
22	An autosomal recessive leucoencephalopathy with ischemic stroke, dysmorphic syndrome and retinitis pigmentosa maps to chromosome 17q24.2-25.3. BMC Medical Genetics, 2012, 13, 18.	2.1	1
23	Vitamin E in ataxia and neurodegenerative diseases: A review. World Journal of Neuroscience, 2012, 02, 217-222.	0.1	2
24	Alteration of Fatty-Acid-Metabolizing Enzymes Affects Mitochondrial Form and Function in Hereditary Spastic Paraplegia. American Journal of Human Genetics, 2012, 91, 1051-1064.	6.2	179
25	A locus for bilateral occipital polymicrogyria maps to chromosome 6q16â€“q22. Neurogenetics, 2009, 10, 35-42.	1.4	10
26	A Novel GDAP1 Mutation P78L Responsible for CMT4A Disease in Three Moroccan Families. Canadian Journal of Neurological Sciences, 2007, 34, 421-426.	0.5	10
27	Spastic paraplegia 5: Locus refinement, candidate gene analysis and clinical description. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2007, 144B, 854-861.	1.7	13
28	A novel locus for autosomal recessive spastic ataxia on chromosome 17p. Human Genetics, 2007, 121, 413-420.	3.8	26
29	Autosomal recessive mutilating sensory neuropathy with spastic paraplegia maps to chromosome 5p15.31â€“14.1. European Journal of Human Genetics, 2006, 14, 249-252.	2.8	31
30	Autosomal recessive axonal Charcot-Marie-Tooth disease (ARCMT2): phenotype-genotype correlations in 13 Moroccan families. Brain, 2006, 130, 1062-1075.	7.6	30
31	Mapping of a new form of pure autosomal recessive spastic paraplegia (SPG28). Annals of Neurology, 2005, 57, 567-571.	5.3	60
32	Mutation in the epsilon subunit of the cytosolic chaperonin-containing t-complex peptide-1 (Cct5) gene causes autosomal recessive mutilating sensory neuropathy with spastic paraplegia. Journal of Medical Genetics, 2005, 43, 441-443.	3.2	111
33	High incidence of SMN1 gene deletion in Moroccan adult-onset spinal muscular atrophy patients. Journal of Neurology, 2003, 250, 1209-1213.	3.6	9
34	Mutations in MTMR13, a New Pseudophosphatase Homologue of MTMR2 and Sbf1, in Two Families with an Autosomal Recessive Demyelinating Form of Charcot-Marie-Tooth Disease Associated with Early-Onset Glaucoma. American Journal of Human Genetics, 2003, 72, 1141-1153.	6.2	263
35	Clinical comparison between AVED patients with 744 del A mutation and Friedreich ataxia with GAA expansion in 15 Moroccan families. Journal of the Neurological Sciences, 2002, 198, 25-29.	0.6	54
36	Genetic, cytogenetic and physical refinement of the autosomal recessive CMT linked to 5q31â€“q33: exclusion of candidate genes including EGR1. European Journal of Human Genetics, 1999, 7, 849-859.	2.8	15

#	Article	IF	Citations
37	A Locus for an Axonal Form of Autosomal Recessive Charcot-Marie-Tooth Disease Maps to Chromosome 1q21.2-q21.3. American Journal of Human Genetics, 1999, 65, 722-727.	6.2	89