## Benoit Marin

## List of Publications by Year in descending order

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		87723	40881
138	9,591	38	93
papers	citations	h-index	g-index
150	150	150	13230
all docs	docs citations	times ranked	citing authors

#	Article	IF	CITATIONS
1	Hypermetabolism is a reality in amyotrophic lateral sclerosis compared to healthy subjects. Journal of the Neurological Sciences, 2021, 420, 117257.	0.3	23
2	Incidence of motor neuron disease/amyotrophic lateral sclerosis in South Africa: a 4â€year prospective study. European Journal of Neurology, 2021, 28, 81-89.	1.7	18
3	Time-trend evolution and determinants of sex ratio in Amyotrophic Lateral Sclerosis: a dose–response meta-analysis. Journal of Neurology, 2021, 268, 2973-2984.	1.8	13
4	Lack of association between Toxocara canis and multiple sclerosis: A population-based case–control study. Multiple Sclerosis Journal, 2020, 26, 258-259.	1.4	3
5	Continuous hemoglobin and plethysmography variability index monitoring can modify blood transfusion practice and is associated with lower mortality. Journal of Clinical Monitoring and Computing, 2020, 34, 683-691.	0.7	7
6	Screening Questionnaires to Detect Neurological Disorders in Developing Countries: A Systematic Review. Neuroepidemiology, 2020, 54, 24-32.	1.1	2
7	COVID-19 studies registration worldwide for prospective studies with a specific focus on the fast-tracking of French ethic procedures. Anaesthesia, Critical Care & Delicine, 2020, 39, 481-482.	0.6	4
8	Impact of plate shape on the conservation of food praxis in institutionalised elderly adults with severe Alzheimer's disease or mixed dementia: Praxalim an observational before-after non-randomized study. International Journal of Nursing Studies Advances, 2020, 2, 100005.	0.9	3
9	Comorbidities of epilepsy in low and middle-income countries: systematic review and meta-analysis. Scientific Reports, 2020, 10, 9015.	1.6	22
10	Increased resting energy expenditure compared with predictive theoretical equations in amyotrophic lateral sclerosis. Nutrition, 2020, 77, 110805.	1.1	9
11	Cytomegalovirus (CMV) Shedding in French Day-Care Centers: A Nationwide Study of Epidemiology, Risk Factors, Centers' Practices, and Parents' Awareness of CMV. Journal of the Pediatric Infectious Diseases Society, 2020, 9, 686-694.	0.6	6
12	Predictive factors for gastrostomy at time of diagnosis and impact on survival in patients with amyotrophic lateral sclerosis. Clinical Nutrition, 2020, 39, 3112-3118.	2.3	14
13	Predictive formulas for estimation of height in sub-Saharan African older people: A new formula (EPIDEMCA study). Nutrition, 2020, 73, 110725.	1.1	1
14	Validity of death certificates in the identification of cases of amyotrophic lateral sclerosis (ALS) in the Limousin region, France. A population-based study. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2020, 21, 228-234.	1.1	5
15	Body mass index and peripheral arterial disease, a "U-shaped―relationship in elderly African population– the EPIDEMCA study. Vasa - European Journal of Vascular Medicine, 2020, 49, 50-56.	0.6	5
16	Peroral endoscopic pyloromyotomy is efficacious and safe for refractory gastroparesis: prospective trial with assessment of pyloric function. Endoscopy, 2019, 51, 40-49.	1.0	104
17	Cumulative incidence of restenosis in the endovascular treatment of extracranial carotid artery stenosis: a meta-analysis. Journal of NeuroInterventional Surgery, 2019, 11, 916-923.	2.0	13
18	Co-morbidities of mental disorders and chronic physical diseases in developing and emerging countries: a meta-analysis. BMC Public Health, 2019, 19, 304.	1.2	96

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19	Residential exposure to ultra high frequency electromagnetic fields emitted by Global System for Mobile (GSM) antennas and amyotrophic lateral sclerosis incidence: A geo-epidemiological population-based study. Environmental Research, 2019, 176, 108525.	3.7	9
20	Pain experienced by infants and toddlers at urine collection bag removal: A randomized, controlled, clinical trial. International Journal of Nursing Studies, 2019, 95, 1-6.	2.5	3
21	Global, regional, and national burden of neurological disorders, 1990–2016: a systematic analysis for the Global Burden of Disease Study 2016. Lancet Neurology, The, 2019, 18, 459-480.	4.9	2,625
22	Amyotrophic lateral sclerosis mortality rates among ethnic groups in a predominant admixed population in Latin America: a population-based study in Ecuador. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2019, 20, 404-412.	1.1	4
23	Associations of mental disorders and neurotropic parasitic diseases: a meta-analysis in developing and emerging countries. BMC Public Health, 2019, 19, 1645.	1.2	17
24	Resting energy expenditure equations in amyotrophic lateral sclerosis, creation of an ALS-specific equation. Clinical Nutrition, 2019, 38, 1657-1665.	2.3	13
25	Clinical features and prognosis of amyotrophic lateral sclerosis in Africa: the TROPALS study. Journal of Neurology, Neurosurgery and Psychiatry, 2019, 90, 20-29.	0.9	29
26	Pectoral I Block Does Not Improve Postoperative Analgesia After Breast Cancer Surgery. Regional Anesthesia and Pain Medicine, 2018, 43, 596-604.	1,1	64
27	Age-specific ALS incidence: a dose–response meta-analysis. European Journal of Epidemiology, 2018, 33, 621-634.	2.5	46
28	Trauma and amyotrophic lateral sclerosis: a european population-based case-control study from the EURALS consortium. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2018, 19, 118-125.	1,1	26
29	Hypermetabolism is a deleterious prognostic factor in patients with amyotrophic lateral sclerosis. European Journal of Neurology, 2018, 25, 97-104.	1.7	85
30	Ankle–Brachial Index: An Ubiquitous Marker of Cognitive Impairment—The EPIDEMCA Study. Angiology, 2018, 69, 497-506.	0.8	5
31	Prevalence of toxoplasmosis and its association with dementia in older adults in Central Africa: a result from the <scp>EPIDEMCA</scp> programme. Tropical Medicine and International Health, 2018, 23, 1304-1313.	1.0	10
32	Methodological Challenges of Neuroepidemiological Studies in Low- and Middle-Income Countries. , 2018, , 3-12.		2
33	Other Neurocognitive Disorders in Tropical Health (Amyotrophic Lateral Sclerosis and Parkinson's) Tj ETQq1 1	0.784314	ł ṛgBT /Over
34	Global, regional, and national burden of motor neuron diseases 1990–2016: a systematic analysis for the Global Burden of Disease Study 2016. Lancet Neurology, The, 2018, 17, 1083-1097.	4.9	163
35	Referral bias in ALS epidemiological studies. PLoS ONE, 2018, 13, e0195821.	1.1	22
36	Meta-analysis of perinatal factors associated with epilepsy in tropical countries. Epilepsy Research, 2018, 146, 54-62.	0.8	5

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37	Updated evidence of the association between toxocariasis and epilepsy: Systematic review and meta-analysis. PLoS Neglected Tropical Diseases, 2018, 12, e0006665.	1.3	51
38	Variation in worldwide incidence of amyotrophic lateral sclerosis: a meta-analysis. International Journal of Epidemiology, 2017, 46, dyw061.	0.9	202
39	Accuracy of Teledentistry for Diagnosing Dental Pathology Using Direct Examination as a Gold Standard: Results of the Tel-e-dent Study of Older Adults Living in Nursing Homes. Journal of the American Medical Directors Association, 2017, 18, 528-532.	1.2	55
40	Nationwide incidence of motor neuron disease using the French health insurance information system database. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2017, 18, 426-433.	1.1	14
41	Prevalence, awareness, treatment, and control of hypertension in older people in Central Africa: the EPIDEMCA study. Journal of the American Society of Hypertension, 2017, 11, 449-460.	2.3	14
42	Current issues in ALS epidemiology: Variation of ALS occurrence between populations and physical activity as a risk factor. Revue Neurologique, 2017, 173, 244-253.	0.6	23
43	ALS and frontotemporal dementia belong to a common disease spectrum. Revue Neurologique, 2017, 173, 273-279.	0.6	56
44	Validity of medico-administrative data related to amyotrophic lateral sclerosis in France: A population-based study. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2017, 18, 24-31.	1.1	9
45	Characteristics and Prognosis of Oldest Old Subjects with Amyotrophic Lateral Sclerosis. Neuroepidemiology, 2017, 49, 64-73.	1.1	13
46	Exploring the diagnosis delay and ALS functional impairment at diagnosis as relevant criteria for clinical trial enrolment*. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2017, 18, 519-527.	1.1	17
47	Systematic review and meta-analysis estimating association of cysticercosis and neurocysticercosis with epilepsy. PLoS Neglected Tropical Diseases, 2017, 11, e0005153.	1.3	65
48	Feasibility of a cardiovascular cohort in a Sub-Saharan Africa community: preliminary report of the pilot project TAHES (TanvÃ" Health Study) in Benin. Global Health Action, 2017, 10, 1270528.	0.7	10
49	A screening questionnaire for convulsive seizures: A three-stage field-validation in rural Bolivia. PLoS ONE, 2017, 12, e0173945.	1.1	18
50	Evaluation of the application of the European guidelines for the diagnosis and clinical care of amyotrophic lateral sclerosis ( <scp>ALS</scp> ) patients in six French <scp>ALS</scp> centres. European Journal of Neurology, 2016, 23, 787-795.	1.7	20
51	Population-Based Evidence that Survival in Amyotrophic Lateral Sclerosis Is Related to Weight Loss at Diagnosis. Neurodegenerative Diseases, 2016, 16, 225-234.	0.8	39
52	Randomized Trials in Developing Countries: Different Priorities and Study Design?. Frontiers of Neurology and Neuroscience, 2016, 39, 136-146.	3.0	2
53	Pseudobulbar affect (PBA) in an incident ALS cohort: results from the Apulia registry (SLAP). Journal of Neurology, 2016, 263, 316-321.	1.8	18
54	Longâ€term efficacy of rituximab in IgM antiâ€myelinâ€associated glycoprotein neuropathy: RIMAG followâ€up study. Journal of the Peripheral Nervous System, 2016, 21, 10-14.	1.4	25

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55	Current Issues in Randomized Clinical Trials of Neurodegenerative Disorders at Enrolment and Reporting: Diagnosis, Recruitment, Representativeness of Patients, Ethnicity, and Quality of Reporting. Frontiers of Neurology and Neuroscience, 2016, 39, 24-36.	3.0	4
56	Non small cell lung cancer (NSCLC) patients harboring BRAF mutation: Clinical characteristics and management in real world setting. Cohort BRAF EXPLORE GFPC 02-14. Annals of Oncology, 2016, 27, vi439.	0.6	1
57	A prospective hospital study of alcohol use disorders, comorbid psychiatric conditions and withdrawal prognosis. Annals of General Psychiatry, 2016, 15, 22.	1.2	8
58	Epidemiology of amyotrophic lateral sclerosis: A review of literature. Revue Neurologique, 2016, 172, 37-45.	0.6	90
59	Impact of Continuing First-Line EGFR Tyrosine Kinase Inhibitor Therapy Beyond RECIST Disease Progression in Patients with Advanced EGFR-Mutated Non-Small-Cell Lung Cancer (NSCLC): Retrospective GFPC 04-13 Study. Targeted Oncology, 2016, 11, 167-174.	1.7	16
60	Prevalence of Multiple Sclerosis in the City of Volta Redonda - Rio De Janeiro, Brazil Using the Capture-Recapture Method. Neuroepidemiology, 2016, 46, 88-95.	1.1	6
61	Stratification of ALS patients' survival: a population-based study. Journal of Neurology, 2016, 263, 100-111.	1.8	27
62	Non-self-sufficiency as a primary outcome measure in ALS trials. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2016, 17, 77-84.	1.1	9
63	Clinical and demographic factors and outcome of amyotrophic lateral sclerosis in relation to population ancestral origin. European Journal of Epidemiology, 2016, 31, 229-245.	2.5	87
64	Association between mild cognitive impairment and dementia and undernutrition among elderly people in Central Africa: some results from the EPIDEMCA (Epidemiology of Dementia in Central Africa) programme. British Journal of Nutrition, 2015, 114, 306-315.	1.2	26
65	Magnitude of Cardiovascular Risk Factors in Rural and Urban Areas in Benin: Findings from a Nationwide Steps Survey. PLoS ONE, 2015, 10, e0126441.	1.1	25
66	Extrapyramidal and cognitive signs in amyotrophic lateral sclerosis: A population based cross-sectional study. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2015, 16, 324-330.	1.1	26
67	Prevalence of peripheral artery disease in the elderly population in urban and rural areas of Central Africa: the EPIDEMCA study. European Journal of Preventive Cardiology, 2015, 22, 1462-1472.	0.8	30
68	Prevalence of multiple sclerosis in the city of Volta Redonda $\hat{a} \in$ Rio de Janeiro, Brazil- using the capture-recapture method. Journal of the Neurological Sciences, 2015, 357, e301.	0.3	0
69	Amyotrophic Lateral Sclerosis: An Aging-Related Disease. Current Geriatrics Reports, 2015, 4, 142-153.	1.1	17
70	Synthesis of a molecularly imprinted sorbent for selective solid-phase extraction of $\hat{l}^2$ -N-methylamino-l-alanine. Talanta, 2015, 144, 1021-1029.	2.9	12
71	Experimental pain sensitivity in subjects with major depression. International Journal of Psychiatry in Medicine, 2015, 50, 219-237.	0.8	6
72	Hepatocyte growth factor measurement in AL amyloidosis. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2015, 22, 112-116.	1.4	8

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73	Analgesic interaction between ondansetron and acetaminophen after tonsillectomy in children: The <scp>P</scp> aratron randomized, controlled trial. European Journal of Pain, 2015, 19, 661-668.	1.4	19
74	BMAALS: A French national project searching for a link between amyotrophic lateral sclerosis and the neurotoxic amino acid L-BMAA. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2014, 15, 155-156.	1.1	5
75	Searching for a link between the L-BMAA neurotoxin and amyotrophic lateral sclerosis: a study protocol of the French BMAALS programme. BMJ Open, 2014, 4, e005528-e005528.	0.8	25
76	Epilepsy: Asia versus Africa. Epilepsia, 2014, 55, 1317-1321.	2.6	14
77	Populationâ€based epidemiology of amyotrophic lateral sclerosis ( <scp>ALS</scp> ) in an ageing <scp>E</scp> urope – the <scp>F</scp> rench register of <scp>ALS</scp> in <scp>L</scp> imousin ( <scp>FRAL</scp> im register). European Journal of Neurology, 2014, 21, 1292.	1.7	55
78	Consensus Conference on a Composite Endpoint for Clinical Trials on Immunosuppressive Drugs in Lung Transplantation. Transplantation, 2014, 98, 1331-1338.	0.5	4
79	Human cytomegalovirus quantification in toddlers saliva from day care centers and emergency unit: A feasibility study. Journal of Clinical Virology, 2014, 61, 371-377.	1.6	26
80	Epidemiology, causes, and treatment of epilepsy in sub-Saharan Africa. Lancet Neurology, The, 2014, 13, 1029-1044.	4.9	212
81	First-ever population-based study on status epilepticus in French Island of La Reunion (France) – Incidence and fatality. Seizure: the Journal of the British Epilepsy Association, 2014, 23, 769-773.	0.9	26
82	Randomized open-label non-comparative multicenter phase II trial of sequential erlotinib and docetaxel versus docetaxel alone in patients with non-small-cell lung cancer after failure of first-line chemotherapy: GFPC 10.02 study. Lung Cancer, 2014, 85, 415-419.	0.9	18
83	Physical activity and amyotrophic lateral sclerosis: A European populationâ€based case–control study. Annals of Neurology, 2014, 75, 708-716.	2.8	79
84	Epidemiology of dementia in Central Africa (EPIDEMCA): protocol for a multicentre population-based study in rural and urban areas of the Central African Republic and the Republic of Congo. SpringerPlus, 2014, 3, 338.	1.2	47
85	Epidemiological evidence that physical activity is not a risk factor for ALS. European Journal of Epidemiology, 2014, 29, 459-475.	2.5	44
86	Development of an analytical procedure for quantifying the underivatized neurotoxin β-N-methylamino-l-alanine in brain tissues. Analytical and Bioanalytical Chemistry, 2014, 406, 4627-4636.	1.9	23
87	Transperineal versus transvaginal ultrasound cervical length measurement and preterm labor. Archives of Gynecology and Obstetrics, 2014, 290, 465-469.	0.8	14
88	Reliability of the measurement of the abdominal aortic diameter by novice operators using a pocket-sized ultrasound system. Archives of Cardiovascular Diseases, 2013, 106, 644-650.	0.7	34
89	Validation of the analytical procedure for the determination of the neurotoxin $\hat{l}^2$ -N-methylamino-l-alanine in complex environmental samples. Analytica Chimica Acta, 2013, 771, 42-49.	2.6	39
90	rTMS for pharmacoresistant major depression in the clinical setting of a psychiatric hospital: Effectiveness and effects of age. Journal of Affective Disorders, 2013, 150, 677-681.	2.0	22

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91	Newlyâ€diagnosed epileptic seizures in three populations: Geneva (EPIGEN), Martinique (EPIMART), and the Reunion Island (EPIREUN). Epileptic Disorders, 2013, 15, 243-254.	0.7	3
92	Placebo-controlled trial of rituximab in IgM anti-myelin–associated glycoprotein neuropathy. Neurology, 2013, 80, 2217-2225.	1.5	167
93	Measurement of the patellar tendon-tibial plateau angle and tuberosity advancement in dogs with cranial cruciate ligament rupture. Veterinary and Comparative Orthopaedics and Traumatology, 2013, 26, 469-478.	0.2	32
94	Dietary BMAA Exposure in an Amyotrophic Lateral Sclerosis Cluster from Southern France. PLoS ONE, 2013, 8, e83406.	1.1	116
95	Toxocariasis and Epilepsy: Systematic Review and Meta-Analysis. PLoS Neglected Tropical Diseases, 2012, 6, e1775.	1.3	120
96	Juvenile and adult-onset ALS/MND among Africans: incidence, phenotype, survival: A review. Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders, 2012, 13, 276-283.	2.3	29
97	Obesity and cervical ripening failure risk. Journal of Maternal-Fetal and Neonatal Medicine, 2012, 25, 304-307.	0.7	25
98	Measurement and Interpretation of the Ankle-Brachial Index. Circulation, 2012, 126, 2890-2909.	1.6	1,232
99	Assessment of left ventricular ejection fraction using an ultrasonic stethoscope in critically ill patients. Critical Care, 2012, 16, R29.	2.5	31
100	Intracameral cefuroxime injection at the end of cataract surgery to reduce the incidence of endophthalmitis: French study. Journal of Cataract and Refractive Surgery, 2012, 38, 1370-1375.	0.7	86
101	The dangers of inadequate understanding of epilepsy in Madagascar. Lancet Neurology, The, 2012, 11, 748-749.	4.9	3
102	Lanreotide autogel 90Âmg and lymphorrhea prevention after axillary node dissection in breast cancer: A phase III double blind, randomized, placebo-controlled trial. European Journal of Surgical Oncology, 2012, 38, 902-909.	0.5	11
103	First assessment at home of amyotrophic lateral sclerosis (ALS) patients by a nutrition network in the French region of Limousin. Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders, 2012, 13, 538-543.	2.3	8
104	FDG-PET/CT Metabolic Tumor Volume: A New Prognostic Marker in Hodgkin Lymphoma?. Blood, 2012, 120, 3632-3632.	0.6	0
105	Alteration of nutritional status at diagnosis is a prognostic factor for survival of amyotrophic lateral sclerosis patients. Journal of Neurology, Neurosurgery and Psychiatry, 2011, 82, 628-634.	0.9	212
106	Contribution of geolocalisation to neuroepidemiological studies: Incidence of ALS and environmental factors in Limousin, France. Journal of the Neurological Sciences, 2011, 309, 115-122.	0.3	29
107	Validation of a risk-assessment scale and a risk-adapted monitoring plan for academic clinical research studies — The Pre-Optimon study. Contemporary Clinical Trials, 2011, 32, 16-24.	0.8	34
108	Troubles nutritionnels lors de la sclérose latérale amyotrophique (SLA). Nutrition Clinique Et Metabolisme, 2011, 25, 205-216.	0.2	2

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109	The epidemiology and treatment of ALS: Focus on the heterogeneity of the disease and critical appraisal of therapeutic trials. Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders, 2011, 12, 1-10.	2.3	107
110	Basic critical care echocardiography: Validation of a curriculum dedicated to noncardiologist residents*. Critical Care Medicine, 2011, 39, 636-642.	0.4	158
111	Efficacy of L-asparaginase with methotrexate and dexamethasone (AspaMetDex regimen) in patients with refractory or relapsing extranodal NK/T-cell lymphoma, a phase 2 study. Blood, 2011, 117, 1834-1839.	0.6	346
112	Newly diagnosed epileptic seizures: Focus on an elderly population on the French island of Réunion in the Southern Indian Ocean. Epilepsia, 2011, 52, 2203-2208.	2.6	16
113	Can Mortality Data Be Used to Estimate Amyotrophic Lateral Sclerosis Incidence?. Neuroepidemiology, 2011, 36, 29-38.	1.1	61
114	Lymph node assessment with (18)F-FDG-PET and MRI in uterine cervical cancer. Anticancer Research, 2011, 31, 3865-71.	0.5	25
115	mRNA levels of enzymes and receptors implicated in arachidonic acid metabolism in gliomas. Clinical Biochemistry, 2010, 43, 827-835.	0.8	8
116	Onchocercose etÂépilepsie. Epilepsies, 2010, 22, 116-119.	0.0	3
117	Validation of the Use of Historical Events to Estimate the Age of Subjects Aged 65 Years and Over in Cotonou (Benin). Neuroepidemiology, 2010, 35, 12-16.	1.1	50
118	Prognostic Significance of BAD and AIF Apoptotic Pathways in Diffuse Large B-Cell Lymphoma. Clinical Lymphoma, Myeloma and Leukemia, 2010, 10, 118-124.	0.2	13
119	Fertility after embolization of the uterine arteries to treat obstetrical hemorrhage: a review of 53 cases. Fertility and Sterility, 2010, 94, 2574-2579.	0.5	70
120	Vital and functional outcomes of the first-ever hemispheric stroke, epidemiological comparative study between Kunming (China) and Limoges (France). Annals of Physical and Rehabilitation Medicine, 2010, 53, 547-558.	1.1	6
121	Cytogenetic Studies in Human Cells Exposed <i>In Vitro</i> to GSM-900ÂMHz Radiofrequency Radiation Using R-Banded Karyotyping. Radiation Research, 2010, 174, 712-718.	0.7	15
122	Increased CXCLâ€13 levels in human African trypanosomiasis meningoâ€encephalitis. Tropical Medicine and International Health, 2009, 14, 529-534.	1.0	23
123	Incidence of newly diagnosed epileptic seizures in a French South Indian Ocean Island, La Réunion (EPIREUN). Epilepsia, 2009, 50, 2207-2212.	2.6	13
124	L-Asparaginase-based treatment of 15 western patients with extranodal NK/T-cell lymphoma and leukemia and a review of the literature. Annals of Oncology, 2009, 20, 110-116.	0.6	111
125	Incidence of amyotrophic lateral sclerosis in the Limousin region of France, 1997–2007. Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders, 2009, 10, 216-220.	2.3	39
126	Prognosis of ALS: Comparing data from the Limousin referral centre, France, and a Uruguayan population. Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders, 2009, 10, 355-360.	2.3	17

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127	A Randomized Phase II Trial Assessing in Advanced Non-small Cell Lung Cancer Patients with Stable Disease after Two Courses of Cisplatin-Gemcitabine an Early Modification of Chemotherapy Doublet with Paclitaxel-Gemcitabine Versus Continuation of Cisplatin-Gemcitabine Chemotherapy (GFPC 03-01) Tj ETQq1	1 <sup>0</sup> .78431	<b>4</b> rgBT /O∨€
128	Non-AIDS-defining deaths and immunodeficiency in the era of combination antiretroviral therapy. Aids, 2009, 23, 1743-1753.	1.0	200
129	HGF Measurement in AL Amyloidosis Blood, 2009, 114, 1783-1783.	0.6	O
130	Perilesional brain oedema in calcific neurocysticercosis: a target to prevent seizure recurrence?. Lancet Neurology, The, 2008, 7, 1075-1076.	4.9	1
131	Phase angle is a prognostic factor for survival in amyotrophic lateral sclerosis. Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders, 2008, 9, 273-278.	2.3	60
132	Achievement a Negative 18-FDG/PET Status at the End of Procedure by Tailored Treatment According Pre-Transplantation Pet Status in Lymphomas Improve 5 Years-OS and EFS Blood, 2008, 112, 2193-2193.	0.6	0
133	Focused training for goal-oriented hand-held echocardiography performed by noncardiologist residents in the intensive care unit. Intensive Care Medicine, 2007, 33, 1795-1799.	3.9	765
134	Onchocerciasis-related epilepsy? Prospects at a time of uncertainty. Trends in Parasitology, 2006, 22, 17-20.	1.5	26
135	Logistic regression model of the clinical response to 5-fluorouracil based chemotherapy for metastatic colorectal cancer patients. Anticancer Research, 2006, 26, 3885-92.	0.5	O
136	Qu'est-ce qu'une régression logistique ?. Revue Des Maladies Respiratoires, 2005, 22, 159-162.	1.7	20
137	Onchocerciasis and epilepsy. Lancet, The, 1994, 343, 983.	6.3	11
138	Juvenile and adult-onset ALS/MND among Africans: incidence, phenotype, survival: A review. Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders, 0, , 1-8.	2.3	0