

Jose A Lopez-Escamez

List of Publications by Year in descending order

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Version: 2024-02-01

140
papers

5,537
citations

81743

39
h-index

102304

66
g-index

161
all docs

161
docs citations

161
times ranked

3332
citing authors

#	ARTICLE	IF	CITATIONS
1	Diagnostic criteria for Meniere's disease. <i>Journal of Vestibular Research: Equilibrium and Orientation</i> , 2015, 25, 1-7.	0.8	995
2	Tinnitus and tinnitus disorder: Theoretical and operational definitions (an international) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50,702 Td (m	0.9	150
3	Vestibular paroxysmia: Diagnostic criteria. <i>Journal of Vestibular Research: Equilibrium and Orientation</i> , 2017, 26, 409-415.	0.8	149
4	Utilizing Ethnic-Specific Differences in Minor Allele Frequency to Recategorize Reported Pathogenic Deafness Variants. <i>American Journal of Human Genetics</i> , 2014, 95, 445-453.	2.6	137
5	Long-term outcome and health-related quality of life in benign paroxysmal positional vertigo. <i>European Archives of Oto-Rhino-Laryngology</i> , 2005, 262, 507-511.	0.8	134
6	High Prevalence of Systemic Autoimmune Diseases in Patients with Meniere's Disease. <i>PLoS ONE</i> , 2011, 6, e26759.	1.1	125
7	Extended phenotype and clinical subgroups in unilateral Meniere disease: A cross-sectional study with cluster analysis. <i>Clinical Otolaryngology</i> , 2017, 42, 1172-1180.	0.6	120
8	Familial clustering and genetic heterogeneity in Meniere's disease. <i>Clinical Genetics</i> , 2014, 85, 245-252.	1.0	116
9	New Insights into Pathophysiology of Vestibular Migraine. <i>Frontiers in Neurology</i> , 2015, 6, 12.	1.1	99
10	Accompanying Symptoms Overlap during Attacks in Meniere's Disease and Vestibular Migraine. <i>Frontiers in Neurology</i> , 2014, 5, 265.	1.1	97
11	Clinical Subgroups in Bilateral Meniere Disease. <i>Frontiers in Neurology</i> , 2016, 7, 182.	1.1	96
12	Identification of two novel mutations in FAM136A and DTNA genes in autosomal-dominant familial Meniere's disease. <i>Human Molecular Genetics</i> , 2015, 24, 1119-1126.	1.4	95
13	Multiple positional nystagmus suggests multiple canal involvement in benign paroxysmal positional vertigo. <i>Acta Oto-Laryngologica</i> , 2005, 125, 954-961.	0.3	80
14	Genetic susceptibility to bilateral tinnitus in a Swedish twin cohort. <i>Genetics in Medicine</i> , 2017, 19, 1007-1012.	1.1	76
15	Meniere's disease. <i>Handbook of Clinical Neurology</i> / Edited By P J Vinken and G W Bruyn, 2016, 137, 257-277.	1.0	72
16	Innovations in Doctoral Training and Research on Tinnitus: The European School on Interdisciplinary Tinnitus Research (ESIT) Perspective. <i>Frontiers in Aging Neuroscience</i> , 2017, 9, 447.	1.7	72
17	Clinical Features, Familial History, and Migraine Precursors in Patients With Definite Vestibular Migraine: The VM Phenotypes Projects. <i>Headache</i> , 2018, 58, 534-544.	1.8	69
18	Proinflammatory cytokines and response to molds in mononuclear cells of patients with Meniere disease. <i>Scientific Reports</i> , 2018, 8, 5974.	1.6	67

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19	Impact of Treatment on Health-Related Quality of Life in Patients with Posterior Canal Benign Paroxysmal Positional Vertigo. <i>Otology and Neurotology</i> , 2003, 24, 637-641.	0.7	66
20	Genetics of vestibular disorders: pathophysiological insights. <i>Journal of Neurology</i> , 2016, 263, 45-53.	1.8	66
21	Time Course of Episodes of Definitive Vertigo in Ménière's Disease. <i>JAMA Otolaryngology</i> , 2008, 134, 1149.	1.5	65
22	Health-Related Quality of Life in Patients over Sixty Years Old with Benign Paroxysmal Positional Vertigo. <i>Gerontology</i> , 2004, 50, 82-86.	1.4	62
23	Variable expressivity and genetic heterogeneity involving DPT and SEMA3D genes in autosomal dominant familial Ménière's disease. <i>European Journal of Human Genetics</i> , 2017, 25, 200-207.	1.4	60
24	Allelic variants in TLR10 gene may influence bilateral affectation and clinical course of Ménière's disease. <i>Immunogenetics</i> , 2013, 65, 345-355.	1.2	59
25	A novel missense variant in <i>PRKCB</i> segregates low-frequency hearing loss in an autosomal dominant family with Ménière's disease. <i>Human Molecular Genetics</i> , 2016, 25, 3407-3415.	1.4	59
26	Excess of Rare Missense Variants in Hearing Loss Genes in Sporadic Meniere Disease. <i>Frontiers in Genetics</i> , 2019, 10, 76.	1.1	58
27	Differential Proinflammatory Signature in Vestibular Migraine and Meniere Disease. <i>Frontiers in Immunology</i> , 2019, 10, 1229.	2.2	57
28	Towards personalized medicine in Ménière's disease. <i>F1000Research</i> , 2018, 7, 1295.	0.8	57
29	Anterior semicircular canal benign paroxysmal positional vertigo and positional downbeating nystagmus. <i>American Journal of Otolaryngology - Head and Neck Medicine and Surgery</i> , 2006, 27, 173-178.	0.6	55
30	Systematic review of magnetic resonance imaging for diagnosis of Meniere disease. <i>Journal of Vestibular Research: Equilibrium and Orientation</i> , 2019, 29, 121-129.	0.8	54
31	Genetics of Tinnitus: An Emerging Area for Molecular Diagnosis and Drug Development. <i>Frontiers in Neuroscience</i> , 2016, 10, 377.	1.4	52
32	Association between Hyperacusis and Tinnitus. <i>Journal of Clinical Medicine</i> , 2020, 9, 2412.	1.0	51
33	Early Diagnosis and Management of Acute Vertigo from Vestibular Migraine and Ménière's Disease. <i>Neurologic Clinics</i> , 2015, 33, 619-628.	0.8	49
34	Position in bed is associated with left or right location in benign paroxysmal positional vertigo of the posterior semicircular canal. <i>American Journal of Otolaryngology - Head and Neck Medicine and Surgery</i> , 2002, 23, 263-266.	0.6	48
35	Regulation of Fn14 Receptor and NF- κ B Underlies Inflammation in Ménière's Disease. <i>Frontiers in Immunology</i> , 2017, 8, 1739.	2.2	48
36	Standardised profiling for tinnitus research: The European School for Interdisciplinary Tinnitus Research Screening Questionnaire (ESIT-SQ). <i>Hearing Research</i> , 2019, 377, 353-359.	0.9	48

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37	Peripheral vestibular disorders: an update. <i>Current Opinion in Neurology</i> , 2019, 32, 165-173.	1.8	48
38	Genetics of dizziness. <i>Current Opinion in Neurology</i> , 2014, 27, 98-104.	1.8	47
39	Burden of Rare Variants in the OTOG Gene in Familial Meniere's Disease. <i>Ear and Hearing</i> , 2020, 41, 1598-1605.	1.0	46
40	Motion sickness diagnostic criteria: Consensus Document of the Classification Committee of the Bárány Society. <i>Journal of Vestibular Research: Equilibrium and Orientation</i> , 2021, 31, 327-344.	0.8	46
41	Current Understanding and Clinical Management of Meniere's Disease: A Systematic Review. <i>Seminars in Neurology</i> , 2020, 40, 138-150.	0.5	44
42	HLA-DRB1*1101 Allele May Be Associated With Bilateral Ménière's Disease in Southern European Population. <i>Otology and Neurotology</i> , 2007, 28, 891-895.	0.7	43
43	Association of a functional polymorphism of <i>PTPN22</i> encoding a lymphoid protein phosphatase in bilateral Meniere's disease. <i>Laryngoscope</i> , 2010, 120, 103-107.	1.1	41
44	Genetics of Recurrent Vertigo and Vestibular Disorders. <i>Current Genomics</i> , 2011, 12, 443-450.	0.7	40
45	Impact of Bilaterality and Headache on Health-Related Quality of Life in Meniere's Disease. <i>Annals of Otolaryngology, Rhinology and Laryngology</i> , 2009, 118, 409-416.	0.6	38
46	Outlook for Tissue Engineering of the Tympanic Membrane. <i>Audiology Research</i> , 2015, 5, 117.	0.8	38
47	Genetic architecture of Meniere's disease. <i>Hearing Research</i> , 2020, 397, 107872.	0.9	38
48	Absence of COCH mutations in patients with Meniere disease. <i>European Journal of Human Genetics</i> , 2004, 12, 75-78.	1.4	37
49	Intronic Variants in the NFKB1 Gene May Influence Hearing Forecast in Patients with Unilateral Sensorineural Hearing Loss in Meniere's Disease. <i>PLoS ONE</i> , 2014, 9, e112171.	1.1	37
50	MICA-STR A.4 Is Associated With Slower Hearing Loss Progression in Patients With Ménière's Disease. <i>Otology and Neurotology</i> , 2012, 33, 223-229.	0.7	36
51	Hearing and vestibular disorders in patients with systemic lupus erythematosus. <i>Lupus</i> , 2013, 22, 437-442.	0.8	34
52	A Systematic Review of Extreme Phenotype Strategies to Search for Rare Variants in Genetic Studies of Complex Disorders. <i>Genes</i> , 2020, 11, 987.	1.0	34
53	CSVS, a crowdsourcing database of the Spanish population genetic variability. <i>Nucleic Acids Research</i> , 2021, 49, D1130-D1137.	6.5	34
54	Towards a unification of treatments and interventions for tinnitus patients: The EU research and innovation action UNITI. <i>Progress in Brain Research</i> , 2021, 260, 441-451.	0.9	31

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55	Genetic contribution to vestibular diseases. <i>Journal of Neurology</i> , 2018, 265, 29-34.	1.8	27
56	Recommendations on Collecting and Storing Samples for Genetic Studies in Hearing and Tinnitus Research. <i>Ear and Hearing</i> , 2019, 40, 219-226.	1.0	27
57	Systematic Review of Prevalence Studies and Familial Aggregation in Vestibular Migraine. <i>Frontiers in Genetics</i> , 2020, 11, 954.	1.1	26
58	Guía de Práctica Clínica Para el Diagnóstico y Tratamiento del Vértigo Posicional Paroxístico Benigno. Documento de Consenso de la Comisión de Otorrinología Sociedad Española de Otorrinolaringología y Cirugía de Cabeza y Cuello. <i>Acta Otorrinolaringológica Española</i> , 2018, 69, 345-366.	0.2	25
59	Burden of rare variants in synaptic genes in patients with severe tinnitus: An exome based extreme phenotype study. <i>EBioMedicine</i> , 2021, 66, 103309.	2.7	25
60	Cytokines and Inflammation in Meniere Disease. <i>Clinical and Experimental Otorhinolaryngology</i> , 2022, 15, 49-59.	1.1	25
61	Polymorphisms of CD16A and CD32 Fcγ receptors and circulating immune complexes in Ménière's disease: a case-control study. <i>BMC Medical Genetics</i> , 2011, 12, 2.	2.1	24
62	Relationship between headaches and tinnitus in a Swedish study. <i>Scientific Reports</i> , 2020, 10, 8494.	1.6	24
63	Genetics of Tinnitus: Time to Biobank Phantom Sounds. <i>Frontiers in Genetics</i> , 2017, 8, 110.	1.1	22
64	Genetics of vestibular syndromes. <i>Current Opinion in Neurology</i> , 2018, 31, 105-110.	1.8	22
65	Enrichment of damaging missense variants in genes related with axonal guidance signalling in sporadic Ménière's disease. <i>Journal of Medical Genetics</i> , 2020, 57, 82-88.	1.5	21
66	A pipeline combining multiple strategies for prioritizing heterozygous variants for the identification of candidate genes in exome datasets. <i>Human Genomics</i> , 2017, 11, 11.	1.4	20
67	Clinical and Functional Characterization of a Missense ELF2 Variant in a CANVAS Family. <i>Frontiers in Genetics</i> , 2018, 9, 85.	1.1	19
68	Expression of A, B, C and DR antigens in definite Ménière's disease in a Spanish population. <i>European Archives of Oto-Rhino-Laryngology</i> , 2002, 259, 347-350.	0.8	18
69	Functional variants of MIF, INFC and TFNA genes are not associated with disease susceptibility or hearing loss progression in patients with Ménière's disease. <i>European Archives of Oto-Rhino-Laryngology</i> , 2013, 270, 1521-1529.	0.8	18
70	Sex-Dependent Aggregation of Tinnitus in Swedish Families. <i>Journal of Clinical Medicine</i> , 2020, 9, 3812.	1.0	18
71	Identification of Potential Ménière's Disease Targets in the Adult Stria Vascularis. <i>Frontiers in Neurology</i> , 2021, 12, 630561.	1.1	18
72	Rare coding variants involving MYO7A and other genes encoding stereocilia link proteins in familial meniere disease. <i>Hearing Research</i> , 2021, 409, 108329.	0.9	18

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73	Monitoring of Caloric Response and Outcome in Patients With Benign Paroxysmal Positional Vertigo. <i>Otology and Neurotology</i> , 2007, 28, 798-802.	0.7	17
74	Short tone bursts are better than clicks for cervical vestibular-evoked myogenic potentials in clinical practice. <i>European Archives of Oto-Rhino-Laryngology</i> , 2012, 269, 1857-1863.	0.8	17
75	The pharmacological management of vertigo in Meniere disease. <i>Expert Opinion on Pharmacotherapy</i> , 2020, 21, 1753-1763.	0.9	17
76	Recurrent Vestibular Symptoms Not Otherwise Specified: Clinical Characteristics Compared With Vestibular Migraine and Meniere's Disease. <i>Frontiers in Neurology</i> , 2021, 12, 674092.	1.1	17
77	Clinical and Cytokine Profile in Patients with Early and Late Onset Meniere Disease. <i>Journal of Clinical Medicine</i> , 2021, 10, 4052.	1.0	16
78	Surface expression and distribution of Fc receptor III (CD 16 molecule) on human natural killer cells and polymorphonuclear neutrophils. <i>Microscopy Research and Technique</i> , 1994, 28, 277-285.	1.2	15
79	Poly(ADP-ribose) polymerase-1 (PARP-1) longer alleles spanning the promoter region may confer protection to bilateral Meniere's disease. <i>Acta Oto-Laryngologica</i> , 2009, 129, 1222-1225.	0.3	15
80	Large expert-curated database for benchmarking document similarity detection in biomedical literature search. <i>Database: the Journal of Biological Databases and Curation</i> , 2019, 2019, .	1.4	15
81	Systematic Review of Sequencing Studies and Gene Expression Profiling in Familial Meniere Disease. <i>Genes</i> , 2020, 11, 1414.	1.0	15
82	American Society of Anesthesiology Classification May Predict Severe Post-Tonsillectomy Haemorrhage in Children. <i>The Journal of Otolaryngology</i> , 2003, 32, 302.	0.6	15
83	Clinical Features of Headache in Patients With Diagnosis of Definite Vestibular Migraine: The VM-Phenotypes Projects. <i>Frontiers in Neurology</i> , 2018, 9, 395.	1.1	14
84	Heritability and Genetics Contribution to Tinnitus. <i>Otolaryngologic Clinics of North America</i> , 2020, 53, 501-513.	0.5	14
85	Functional Variants in <i>NOS1</i> and <i>NOS2A</i> Are Not Associated with Progressive Hearing Loss in Meniere's Disease in a European Caucasian Population. <i>DNA and Cell Biology</i> , 2011, 30, 699-708.	0.9	13
86	Dynamics of canal response to head-shaking test in benign paroxysmal positional vertigo. <i>Acta Oto-Laryngologica</i> , 2007, 127, 1246-1254.	0.3	12
87	Vestibular Evoked Myogenic Potentials and Health-Related Quality of Life in Patients With Vestibular Neuritis. <i>Otology and Neurotology</i> , 2010, 31, 954-958.	0.7	12
88	Dynamic visual acuity during head-thrust test in canal planes in healthy subjects and patients with vestibular neuritis. <i>Acta Oto-Laryngologica</i> , 2010, 130, 1260-1266.	0.3	12
89	Meniere's Disease: Genetics and the Immune System. <i>Current Otorhinolaryngology Reports</i> , 2018, 6, 24-31.	0.2	12
90	Practice Guidelines for the Diagnosis and Management of Benign Paroxysmal Positional Vertigo. Otolaryngology Committee of Spanish Otorhinolaryngology and Head and Neck Surgery Consensus Document. <i>Acta Otorrinolaringologica (English Edition)</i> , 2018, 69, 345-366.	0.1	12

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91	Unification of Treatments and Interventions for Tinnitus Patients (UNITI): a study protocol for a multi-center randomized clinical trial. <i>Trials</i> , 2021, 22, 875.	0.7	12
92	A Pilot Study Using Intratympanic Methylprednisolone for Treatment of Persistent Posterior Canal Benign Paroxysmal Positional Vertigo. <i>Journal of International Advanced Otology</i> , 2016, 12, 321-325.	1.0	11
93	Defective Î±â€œectorin may involve tectorial membrane in familial Meniere disease. <i>Clinical and Translational Medicine</i> , 2022, 12, .	1.7	11
94	Electron probe X-ray microanalysis of Ca and K distributions in the otolithic membrane. <i>Micron and Microscopica Acta</i> , 1992, 23, 349-350.	0.2	10
95	Standards for quantification of elements in the otolithic membrane by electron probe Xâ€œray microanalysis: Calibration curves and electron beam sensitivity. <i>Journal of Microscopy</i> , 1993, 171, 215-222.	0.8	10
96	Changes in elemental concentrations in K562 target cells after conjugation with human lymphocytes studied by X-ray microanalysis.. <i>Cell Biology International</i> , 1994, 18, 915-916.	1.4	10
97	Gentamicin Ototoxicity in Otoconia: Quantitative Electron Probe X-ray Microanalysis. <i>Acta Oto-Laryngologica</i> , 1994, 114, 18-23.	0.3	10
98	Cervical Metastasis of Occult Papillary Thyroid Carcinoma Associated with Epidermoid Carcinoma of the Larynx. <i>Orl</i> , 1999, 61, 224-226.	0.6	10
99	DNA Methylation Signature in Mononuclear Cells and Proinflammatory Cytokines May Define Molecular Subtypes in Sporadic Meniere Disease. <i>Biomedicines</i> , 2021, 9, 1530.	1.4	10
100	Electron probe microanalysis of gentamicin-induced changes on ionic composition of the vestibular gelatinous membrane. <i>Hearing Research</i> , 1994, 76, 60-66.	0.9	9
101	Abordaje prÃ¡ctico del vÃ©rtigo posicional paroxÃ©stico benigno recurrente. <i>Acta OtorrinolaringolÃ³gica EspaÃ±ola</i> , 2008, 59, 413-419.	0.2	8
102	Genetic and clinical heterogeneity in Meniere's disease. <i>Autoimmunity Reviews</i> , 2012, 11, 925-926.	2.5	8
103	Article Commentary: Role of vestibular testing in diagnosis of benign paroxysmal positional vertigo. <i>Otolaryngology - Head and Neck Surgery</i> , 2009, 141, 7-9.	1.1	7
104	MigraÃ±a vestibular: un diagnÃ³stico emergente. <i>Acta OtorrinolaringolÃ³gica EspaÃ±ola</i> , 2013, 64, 387-388.	0.2	7
105	Perspectivas para el tratamiento de la hipoacusia neurosensorial mediante regeneraciÃ³n celular del oÃ±do interno. <i>Acta OtorrinolaringolÃ³gica EspaÃ±ola</i> , 2015, 66, 286-295.	0.2	7
106	Generation and characterization of the human iPSC line PBMC1-iPS4F1 from adult peripheral blood mononuclear cells. <i>Stem Cell Research</i> , 2015, 15, 614-617.	0.3	7
107	AnÃ¡lisis de la producciÃ³n cientÃ­fica en otorrinolaringologÃ­a en EspaÃ±a durante el periodo 2011-2015. <i>Acta OtorrinolaringolÃ³gica EspaÃ±ola</i> , 2018, 69, 275-282.	0.2	7
108	A tinnitus symphony in 100 patients with Meniere's disease. <i>Clinical Otolaryngology</i> , 2019, 44, 1176-1180.	0.6	7

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109	A Systematic Review on the Association of Acquired Human Cytomegalovirus Infection with Hearing Loss. <i>Journal of Clinical Medicine</i> , 2020, 9, 4011.	1.0	7
110	X-ray Microanalytical Determination of P, S and K Concentrations in the Gelatinous Membrane of the Utricle. <i>Acta Oto-Laryngologica</i> , 1993, 113, 176-180.	0.3	6
111	Migraine, sudden sensorineural hearing loss and autoimmune ear disease. <i>Cephalalgia</i> , 2013, 33, 1206-1207.	1.8	6
112	High-frequency sensorineural hearing loss associated with vestibular episodic syndrome. <i>Clinical Otolaryngology</i> , 2017, 42, 856-859.	0.6	6
113	Genetic Inheritance and Its Contribution to Tinnitus. <i>Current Topics in Behavioral Neurosciences</i> , 2020, 51, 29-47.	0.8	6
114	Standardized Clinical Profiling in Spanish Patients with Chronic Tinnitus. <i>Journal of Clinical Medicine</i> , 2022, 11, 978.	1.0	6
115	Dependence between saccule and utricle in Ca and K concentrations determined by electron probe X-ray microanalysis. <i>Micron and Microscopica Acta</i> , 1992, 23, 367-368.	0.2	5
116	Bioinformatic Integration of Molecular Networks and Major Pathways Involved in Mice Cochlear and Vestibular Supporting Cells. <i>Frontiers in Molecular Neuroscience</i> , 2018, 11, 108.	1.4	5
117	A Predictive Model of Bilateral Sensorineural Hearing Loss in Meniere Disease Using Clinical Data. <i>Ear and Hearing</i> , 2022, 43, 1079-1085.	1.0	5
118	Do we need to reconsider the classification of vestibular migraine?. <i>Expert Review of Neurotherapeutics</i> , 2021, 21, 503-516.	1.4	4
119	Impact of treatment on benign positional vertigo-related quality of life. <i>International Congress Series</i> , 2003, 1240, 1329-1332.	0.2	3
120	Mechanically Induced Calcium Increases in Isolated Vestibular Hair Cells of the Guinea Pig. <i>Acta Oto-Laryngologica</i> , 1995, 115, 759-764.	0.3	2
121	A pilot study of sexual health in patients with Ménière's disease. <i>Acta Otorrinolaringologica (English Edition)</i> 2010, 78, 431-432.	1.0784314	2
122	Perspectives for the Treatment of Sensorineural Hearing Loss by Cellular Regeneration of the Inner Ear. <i>Acta Otorrinolaringologica (English Edition)</i> , 2015, 66, 286-295.	0.1	2
123	Generation of human iPSC line GRX-MCiPS4F-A2 from adult peripheral blood mononuclear cells (PBMCs) with Spanish genetic background. <i>Stem Cell Research</i> , 2015, 15, 337-340.	0.3	2
124	Video Head-Impulse Testing vs Clinical Diagnosis of Vestibular Disorders. <i>JAMA Otolaryngology - Head and Neck Surgery</i> , 2019, 145, 561.	1.2	2
125	La depresión de la investigación en otorrinolaringología en España tiene solución. <i>Acta Otorrinolaringológica Española</i> , 2020, 71, 1-2.	0.2	2
126	Systematic review of sound stimulation to elicit tinnitus residual inhibition. <i>Progress in Brain Research</i> , 2021, 262, 1-21.	0.9	2

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127	Emerging Topics in the Behavioral Neuroscience of Tinnitus. <i>Current Topics in Behavioral Neurosciences</i> , 2021, 51, 461-483.	0.8	2
128	Vestibular Contributions to Health and Disease. <i>Frontiers Research Topics</i> , 0, , .	0.2	2
129	Role of Genomic Medicine in Middle and Inner Ear Diseases. <i>Acta Otorrinolaringologica (English)</i> Tj ETQq1 1 0.784314 rgBT /Overlock 0.1	0.1	1
130	Magnetic resonance imaging of endolymphatic hydrops: Controversies and common ground. <i>Journal of Vestibular Research: Equilibrium and Orientation</i> , 2019, , 1-4.	0.8	1
131	Editorial: Epidemiology and Genetics of Vestibular Disorders. <i>Frontiers in Neurology</i> , 2021, 12, 743379.	1.1	1
132	MÃ©niÃ©re's Syndrome and Migraine. , 2014, , 129-141.		1
133	Role of inheritance in tinnitus: it is time to search the genome. <i>Actualidad MÃ©dica</i> , 2017, 102, 88-92.	0.1	1
134	Vestibular Myogenic Evoked Potentials in Patients With Benign Paroxysmal Positional Vertigo. <i>Acta Otorrinolaringologica (English Edition)</i> , 2007, 58, 495.	0.1	0
135	Potenciales evocadores miogÃ©nicos vestibulares en pacientes con vÃ©rtigo posicional paroxÃ©stico benigno. <i>Acta OtorrinolaringolÃ³gica EspaÃ±ola</i> , 2007, 58, 495.	0.2	0
136	Practical Approach to Recurrent Benign Paroxysmal Positional Vertigo. <i>Acta Otorrinolaringologica (English Edition)</i> , 2008, 59, 413-419.	0.1	0
137	Analysis of Scientific Production in Otolaryngology in Spain in the Period 2011â€“2015. <i>Acta Otorrinolaringologica (English Edition)</i> , 2018, 69, 275-282.	0.1	0
138	Burden of Rare Variants in Synaptic Genes in Patients with Severe Tinnitus: An Exome Based Extreme Phenotype Study. <i>SSRN Electronic Journal</i> , 0, , .	0.4	0
139	Balance Disorders Including Meniere Disease. , 2021, , .		0
140	Clinical and molecular genetics of Meniere disease. <i>Medizinische Genetik</i> , 2020, 32, 141-148.	0.1	0