Jose A Lopez-Escamez

List of Publications by Year in Descending Order

134 papers	3,433 citations	33 h-index	53 g-index
161 ext. papers	4,584 ext. citations	3.4 avg, IF	5.54 L-index

#	Paper	IF	Citations
134	Cytokines and Inflammation in Meniere Disease.. *Clinical and Experimental Otorhinolaryngology*, **2022**,	3.4	3
133	Standardized Clinical Profiling in Spanish Patients with Chronic Tinnitus.. *Journal of Clinical Medicine* , **2022**, 11,	5.1	1
132	Defective ▯ectorin may involve tectorial membrane in familial Meniere disease. *Clinical and Translational Medicine*, **2022**, 12,	5.7	0
131	A Predictive Model of Bilateral Sensorineural Hearing Loss in Meniere Disease Using Clinical Data. *Ear and Hearing*, **2021**,	3.4	1
130	Unification of Treatments and Interventions for Tinnitus Patients (UNITI): a study protocol for a multi-center randomized clinical trial. *Trials*, **2021**, 22, 875	2.8	1
129	Genetic Inheritance and Its Contribution to Tinnitus. *Current Topics in Behavioral Neurosciences*, **2021**, 51, 29-47	3.4	3
128	Burden of rare variants in synaptic genes in patients with severe tinnitus: An exome based extreme phenotype study. *EBioMedicine*, **2021**, 66, 103309	8.8	10
127	Do we need to reconsider the classification of vestibular migraine?. *Expert Review of Neurotherapeutics*, **2021**, 21, 503-516	4.3	2
126	Recurrent Vestibular Symptoms Not Otherwise Specified: Clinical Characteristics Compared With Vestibular Migraine and Meni▯e's Disease. *Frontiers in Neurology*, **2021**, 12, 674092	4.1	4
125	CSVS, a crowdsourcing database of the Spanish population genetic variability. *Nucleic Acids Research*, **2021**, 49, D1130-D1137	20.1	8
124	Systematic review of sound stimulation to elicit tinnitus residual inhibition. *Progress in Brain Research*, **2021**, 262, 1-21	2.9	0
123	Emerging Topics in the Behavioral Neuroscience of Tinnitus. *Current Topics in Behavioral Neurosciences*, **2021**, 51, 461-483	3.4	1
122	Towards a unification of treatments and interventions for tinnitus patients: The EU research and innovation action UNITI. *Progress in Brain Research*, **2021**, 260, 441-451	2.9	12
121	Identification of Potential Meniere's Disease Targets in the Adult Stria Vascularis. *Frontiers in Neurology*, **2021**, 12, 630561	4.1	7
120	Motion sickness diagnostic criteria: Consensus Document of the Classification Committee of the B▯▯y Society. *Journal of Vestibular Research: Equilibrium and Orientation*, **2021**, 31, 327-344	2.5	16
119	Rare coding variants involving MYO7A and other genes encoding stereocilia link proteins in familial meniere disease. *Hearing Research*, **2021**, 409, 108329	3.9	4
118	Clinical and Cytokine Profile in Patients with Early and Late Onset Meniere Disease. *Journal of Clinical Medicine*, **2021**, 10,	5.1	3

117	Tinnitus and tinnitus disorder: Theoretical and operational definitions (an international multidisciplinary proposal). *Progress in Brain Research*, **2021**, 260, 1-25	2.9	35
116	Enrichment of damaging missense variants in genes related with axonal guidance signalling in sporadic Meniere's disease. *Journal of Medical Genetics*, **2020**, 57, 82-88	5.8	11
115	The pharmacological management of vertigo in Meniere disease. *Expert Opinion on Pharmacotherapy*, **2020**, 21, 1753-1763	4	6
114	Heritability and Genetics Contribution to Tinnitus. *Otolaryngologic Clinics of North America*, **2020**, 53, 501-513	2	9
113	Clinical and molecular genetics of Meniere disease. *Medizinische Genetik*, **2020**, 32, 141-148	0.5	
112	Relationship between headaches and tinnitus in a Swedish study. *Scientific Reports*, **2020**, 10, 8494	4.9	14
111	Current Understanding and Clinical Management of Meniere's Disease: A Systematic Review. *Seminars in Neurology*, **2020**, 40, 138-150	3.2	17
110	Genetic architecture of Meniere's disease. *Hearing Research*, **2020**, 397, 107872	3.9	16
109	Systematic Review of Prevalence Studies and Familial Aggregation in Vestibular Migraine. *Frontiers in Genetics*, **2020**, 11, 954	4.5	9
108	Systematic Review of Sequencing Studies and Gene Expression Profiling in Familial Meniere Disease. *Genes*, **2020**, 11,	4.2	5
107	Sex-Dependent Aggregation of Tinnitus in Swedish Families. *Journal of Clinical Medicine*, **2020**, 9,	5.1	7
106	Association between Hyperacusis and Tinnitus. *Journal of Clinical Medicine*, **2020**, 9,	5.1	19
105	Burden of Rare Variants in the OTOG Gene in Familial Meniere's Disease. *Ear and Hearing*, **2020**, 41, 1598-1605	3.4	20
104	A Systematic Review of Extreme Phenotype Strategies to Search for Rare Variants in Genetic Studies of Complex Disorders. *Genes*, **2020**, 11,	4.2	9
103	A Systematic Review on the Association of Acquired Human Cytomegalovirus Infection with Hearing Loss. *Journal of Clinical Medicine*, **2020**, 9,	5.1	3
102	Differential Proinflammatory Signature in Vestibular Migraine and Meniere Disease. *Frontiers in Immunology*, **2019**, 10, 1229	8.4	32
101	Magnetic resonance imaging of endolymphatic hydrops: Controversies and common ground. *Journal of Vestibular Research: Equilibrium and Orientation*, **2019**,	2.5	1
100	Video Head-Impulse Testing vs Clinical Diagnosis of Vestibular Disorders. *JAMA Otolaryngology - Head and Neck Surgery*, **2019**, 145, 561-562	3.9	1

99	Recommendations on Collecting and Storing Samples for Genetic Studies in Hearing and Tinnitus Research. *Ear and Hearing*, **2019**, 40, 219-226	3.4	14
98	Standardised profiling for tinnitus research: The European School for Interdisciplinary Tinnitus Research Screening Questionnaire (ESIT-SQ). *Hearing Research*, **2019**, 377, 353-359	3.9	21
97	Excess of Rare Missense Variants in Hearing Loss Genes in Sporadic Meniere Disease. *Frontiers in Genetics*, **2019**, 10, 76	4.5	20
96	A tinnitus symphony in 100 patients with Meniere's disease. *Clinical Otolaryngology*, **2019**, 44, 1176-1180	1.8	5
95	Large expert-curated database for benchmarking document similarity detection in biomedical literature search. *Database: the Journal of Biological Databases and Curation*, **2019**, 2019,	5	4
94	Systematic review of magnetic resonance imaging for diagnosis of Meniere disease. *Journal of Vestibular Research: Equilibrium and Orientation*, **2019**, 29, 121-129	2.5	27
93	Peripheral vestibular disorders: an update. *Current Opinion in Neurology*, **2019**, 32, 165-173	7.1	26
92	Proinflammatory cytokines and response to molds in mononuclear cells of patients with Meniere disease. *Scientific Reports*, **2018**, 8, 5974	4.9	33
91	Meniere□ Disease: Genetics and the Immune System. *Current Otorhinolaryngology Reports*, **2018**, 6, 24-31	0.5	6
90	Analysis of scientific production in otolaryngology in Spain in the period 2011-2015. *Acta Otorrinolaringol□gica Espa□ola*, **2018**, 69, 275-282	0.9	5
89	Genetic contribution to vestibular diseases. *Journal of Neurology*, **2018**, 265, 29-34	5.5	19
88	Practice Guidelines for the Diagnosis and Management of Benign Paroxysmal Positional Vertigo Otoneurology Committee of Spanish Otorhinolaryngology and Head and Neck Surgery Consensus Document. *Acta Otorrinolaringol□gica Espa□ola*, **2018**, 69, 345-366	0.9	13
87	Clinical and Functional Characterization of a Missense Variant in a CANVAS Family. *Frontiers in Genetics*, **2018**, 9, 85	4.5	11
86	Bioinformatic Integration of Molecular Networks and Major Pathways Involved in Mice Cochlear and Vestibular Supporting Cells. *Frontiers in Molecular Neuroscience*, **2018**, 11, 108	6.1	5
85	Clinical Features of Headache in Patients With Diagnosis of Definite Vestibular Migraine: The VM-Phenotypes Projects. *Frontiers in Neurology*, **2018**, 9, 395	4.1	9
84	Towards personalized medicine in M□i□e's disease. *F1000Research*, **2018**, 7,	3.6	17
83	Genetics of vestibular syndromes. *Current Opinion in Neurology*, **2018**, 31, 105-110	7.1	10
82	Clinical Features, Familial History, and Migraine Precursors in Patients With Definite Vestibular Migraine: The VM-Phenotypes Projects. *Headache*, **2018**, 58, 534-544	4.2	38

81	Practice Guidelines for the Diagnosis and Management of Benign Paroxysmal Positional Vertigo Otoneurology Committee of Spanish Otorhinolaryngology and Head and Neck Surgery Consensus Document. *Acta Otorrinolaringologica (English Edition)*, **2018**, 69, 345-366	0.1	4
80	Analysis of Scientific Production in Otolaryngology in Spain in the Period 2011–2015. *Acta Otorrinolaringologica (English Edition)*, **2018**, 69, 275-282	0.1	
79	Extended phenotype and clinical subgroups in unilateral Meniere disease: A cross-sectional study with cluster analysis. *Clinical Otolaryngology*, **2017**, 42, 1172-1180	1.8	58
78	Genetic susceptibility to bilateral tinnitus in a Swedish twin cohort. *Genetics in Medicine*, **2017**, 19, 1007-1012	8.1	47
77	Innovations in Doctoral Training and Research on Tinnitus: The European School on Interdisciplinary Tinnitus Research (ESIT) Perspective. *Frontiers in Aging Neuroscience*, **2017**, 9, 447	5.3	50
76	A pipeline combining multiple strategies for prioritizing heterozygous variants for the identification of candidate genes in exome datasets. *Human Genomics*, **2017**, 11, 11	6.8	10
75	High-frequency sensorineural hearing loss associated with vestibular episodic syndrome. *Clinical Otolaryngology*, **2017**, 42, 856-859	1.8	2
74	Variable expressivity and genetic heterogeneity involving DPT and SEMA3D genes in autosomal dominant familial Meniere's disease. *European Journal of Human Genetics*, **2017**, 25, 200-207	5.3	43
73	Genetics of Tinnitus: Time to Biobank Phantom Sounds. *Frontiers in Genetics*, **2017**, 8, 110	4.5	15
72	Regulation of Fn14 Receptor and NF-B Underlies Inflammation in Meniere's Disease. *Frontiers in Immunology*, **2017**, 8, 1739	8.4	35
71	Vestibular paroxysmia: Diagnostic criteria. *Journal of Vestibular Research: Equilibrium and Orientation*, **2016**, 26, 409-415	2.5	91
70	A novel missense variant in PRKCB segregates low-frequency hearing loss in an autosomal dominant family with Meniere's disease. *Human Molecular Genetics*, **2016**, 25, 3407-3415	5.6	43
69	A Pilot Study Using Intratympanic Methylprednisolone for Treatment of Persistent Posterior Canal Benign Paroxysmal Positional Vertigo. *Journal of International Advanced Otology*, **2016**, 12, 321-325	1.1	9
68	Clinical Subgroups in Bilateral Meniere Disease. *Frontiers in Neurology*, **2016**, 7, 182	4.1	59
67	Genetics of Tinnitus: An Emerging Area for Molecular Diagnosis and Drug Development. *Frontiers in Neuroscience*, **2016**, 10, 377	5.1	41
66	Genetics of vestibular disorders: pathophysiological insights. *Journal of Neurology*, **2016**, 263 Suppl 1, S45-53	5.5	48
65	Meniere's disease. *Handbook of Clinical Neurology / Edited By P J Vinken and G W Bruyn*, **2016**, 137, 257-77	3	46
64	Early Diagnosis and Management of Acute Vertigo from Vestibular Migraine and Meniere's Disease. *Neurologic Clinics*, **2015**, 33, 619-28, ix	4.5	35

63	Perspectives for the treatment of sensorineural hearing loss by cellular regeneration of the inner ear. *Acta Otorrinolaringológica Española*, **2015**, 66, 286-95	0.9	7
62	Generation and characterization of the human iPSC line PBMC1-iPS4F1 from adult peripheral blood mononuclear cells. *Stem Cell Research*, **2015**, 15, 614-7	1.6	6
61	Generation of human iPSC line GRX-MCiPS4F-A2 from adult peripheral blood mononuclear cells (PBMCs) with Spanish genetic background. *Stem Cell Research*, **2015**, 15, 337-40	1.6	1
60	Perspectives for the Treatment of Sensorineural Hearing Loss by Cellular Regeneration of the Inner Ear. *Acta Otorrinolaringologica (English Edition)*, **2015**, 66, 286-295	0.1	1
59	New insights into pathophysiology of vestibular migraine. *Frontiers in Neurology*, **2015**, 6, 12	4.1	63
58	Outlook for Tissue Engineering of the Tympanic Membrane. *Audiology Research*, **2015**, 5, 117	1.5	27
57	Diagnostic criteria for Meniere's disease. *Journal of Vestibular Research: Equilibrium and Orientation*, **2015**, 25, 1-7	2.5	632
56	Identification of two novel mutations in FAM136A and DTNA genes in autosomal-dominant familial Meniere's disease. *Human Molecular Genetics*, **2015**, 24, 1119-26	5.6	70
55	Utilizing ethnic-specific differences in minor allele frequency to recategorize reported pathogenic deafness variants. *American Journal of Human Genetics*, **2014**, 95, 445-53	11	118
54	Intronic variants in the NFKB1 gene may influence hearing forecast in patients with unilateral sensorineural hearing loss in Meniere's disease. *PLoS ONE*, **2014**, 9, e112171	3.7	25
53	Familial clustering and genetic heterogeneity in Meniere's disease. *Clinical Genetics*, **2014**, 85, 245-52	4	76
52	Accompanying Symptoms Overlap during Attacks in Meniere's Disease and Vestibular Migraine. *Frontiers in Neurology*, **2014**, 5, 265	4.1	64
51	Genetics of dizziness: cerebellar and vestibular disorders. *Current Opinion in Neurology*, **2014**, 27, 98-104	7.1	30
50	Meniere's Syndrome and Migraine **2014**, 129-141		1
49	Functional variants of MIF, INFG and TFNA genes are not associated with disease susceptibility or hearing loss progression in patients with Meniere's disease. *European Archives of Oto-Rhino-Laryngology*, **2013**, 270, 1521-9	3.5	14
48	Allelic variants in TLR10 gene may influence bilateral affectation and clinical course of Meniere's disease. *Immunogenetics*, **2013**, 65, 345-55	3.2	44
47	Hearing and vestibular disorders in patients with systemic lupus erythematosus. *Lupus*, **2013**, 22, 437-42	2.6	28
46	Migraine, sudden sensorineural hearing loss and autoimmune ear disease. *Cephalalgia*, **2013**, 33, 1206-7	6.1	6

45	Genetic and clinical heterogeneity in Meniere's disease. *Autoimmunity Reviews*, **2012**, 11, 925-6	13.6	7
44	Role of Genomic Medicine in Middle and Inner Ear Diseases. *Acta Otorrinolaringologica (English Edition)*, **2012**, 63, 470-479	0.1	
43	Short tone bursts are better than clicks for cervical vestibular-evoked myogenic potentials in clinical practice. *European Archives of Oto-Rhino-Laryngology*, **2012**, 269, 1857-63	3.5	12
42	MICA-STR A.4 is associated with slower hearing loss progression in patients with Ménière's disease. *Otology and Neurotology*, **2012**, 33, 223-9	2.6	26
41	A pilot study of sexual health in patients with Ménière's disease. *Acta Otorrinolaringologica (English Edition)*, **2011**, 62, 119-125	0.1	
40	Genetics of recurrent vertigo and vestibular disorders. *Current Genomics*, **2011**, 12, 443-50	2.6	28
39	Polymorphisms of CD16A and CD32 Fc receptors and circulating immune complexes in Ménière's disease: a case-control study. *BMC Medical Genetics*, **2011**, 12, 2	2.1	22
38	Functional variants in NOS1 and NOS2A are not associated with progressive hearing loss in Ménière's disease in a European Caucasian population. *DNA and Cell Biology*, **2011**, 30, 699-708	3.6	8
37	High prevalence of systemic autoimmune diseases in patients with Ménière's disease. *PLoS ONE*, **2011**, 6, e26759	3.7	99
36	Dynamic visual acuity during head-thrust test in canal planes in healthy subjects and patients with vestibular neuritis. *Acta Oto-Laryngologica*, **2010**, 130, 1260-6	1.6	8
35	Vestibular evoked myogenic potentials and health-related quality of life in patients with vestibular neuritis. *Otology and Neurotology*, **2010**, 31, 954-8	2.6	11
34	Association of a functional polymorphism of PTPN22 encoding a lymphoid protein phosphatase in bilateral Meniere's disease. *Laryngoscope*, **2010**, 120, 103-7	3.6	30
33	Poly(ADP-ribose) polymerase-1 (PARP-1) longer alleles spanning the promoter region may confer protection to bilateral Meniere's disease. *Acta Oto-Laryngologica*, **2009**, 129, 1222-5	1.6	10
32	Role of vestibular testing in diagnosis of benign paroxysmal positional vertigo. *Otolaryngology - Head and Neck Surgery*, **2009**, 141, 7-9; author reply 10-1	5.5	3
31	Impact of bilaterality and headache on health-related quality of life in Meniere's disease. *Annals of Otology, Rhinology and Laryngology*, **2009**, 118, 409-16	2.1	27
30	Abordaje práctico del vértigo posicional paroxístico benigno recurrente. *Acta Otorrinolaringológica Española*, **2008**, 59, 413-419	0.9	8
29	Practical Approach to Recurrent Benign Paroxysmal Positional Vertigo. *Acta Otorrinolaringologica (English Edition)*, **2008**, 59, 413-419	0.1	
28	Time course of episodes of definitive vertigo in Meniere's disease. *JAMA Otolaryngology*, **2008**, 134, 1149-54		51

27	Vestibular Myogenic Evoked Potentials in Patients With Benign Paroxysmal Positional Vertigo. *Acta Otorrinolaringologica (English Edition)*, **2007**, 58, 495	0.1	
26	HLA-DRB1*1101 allele may be associated with bilateral Ménière's disease in southern European population. *Otology and Neurotology*, **2007**, 28, 891-5	2.6	35
25	Potenciales evocadores miogénicos vestibulares en pacientes con vértigo posicional paroxístico benigno. *Acta Otorrinolaringológica Española*, **2007**, 58, 495	0.9	
24	Dynamics of canal response to head-shaking test in benign paroxysmal positional vertigo. *Acta Oto-Laryngologica*, **2007**, 127, 1246-54	1.6	9
23	Monitoring of caloric response and outcome in patients with benign paroxysmal positional vertigo. *Otology and Neurotology*, **2007**, 28, 798-800	2.6	14
22	Anterior semicircular canal benign paroxysmal positional vertigo and positional downbeating nystagmus. *American Journal of Otolaryngology - Head and Neck Medicine and Surgery*, **2006**, 27, 173-8	2.8	47
21	Long-term outcome and health-related quality of life in benign paroxysmal positional vertigo. *European Archives of Oto-Rhino-Laryngology*, **2005**, 262, 507-11	3.5	99
20	Multiple positional nystagmus suggests multiple canal involvement in benign paroxysmal vertigo. *Acta Oto-Laryngologica*, **2005**, 125, 954-61	1.6	62
19	Health-related quality of life in patients over sixty years old with benign paroxysmal positional vertigo. *Gerontology*, **2004**, 50, 82-6	5.5	46
18	Absence of COCH mutations in patients with Meniere disease. *European Journal of Human Genetics*, **2004**, 12, 75-8	5.3	33
17	Impact of treatment on health-related quality of life in patients with posterior canal benign paroxysmal positional vertigo. *Otology and Neurotology*, **2003**, 24, 637-41	2.6	56
16	Impact of treatment on benign positional vertigo-related quality of life. *International Congress Series*, **2003**, 1240, 1329-1332		3
15	American Society of Anesthesiology classification may predict severe post-tonsillectomy haemorrhage in children. *The Journal of Otolaryngology*, **2003**, 32, 302-7		13
14	Expression of A, B, C and DR antigens in definite Meniere's disease in a Spanish population. *European Archives of Oto-Rhino-Laryngology*, **2002**, 259, 347-50	3.5	15
13	Position in bed is associated with left or right location in benign paroxysmal positional vertigo of the posterior semicircular canal. *American Journal of Otolaryngology - Head and Neck Medicine and Surgery*, **2002**, 23, 263-6	2.8	39
12	Cervical metastasis of occult papillary thyroid carcinoma associated with epidermoid carcinoma of the larynx. *Orl*, **1999**, 61, 224-6	2	9
11	Mechanically induced calcium increases in isolated vestibular hair cells of the guinea pig. *Acta Oto-Laryngologica*, **1995**, 115, 759-64	1.6	2
10	Surface expression and distribution of Fc receptor III (CD16 molecule) on human natural killer cells and polymorphonuclear neutrophils. *Microscopy Research and Technique*, **1994**, 28, 277-85	2.8	13

9	Changes in elemental concentrations in K562 target cells after conjugation with human lymphocytes studied by X-ray microanalysis. *Cell Biology International*, **1994**, 18, 915-6	4.5	9
8	Electron probe microanalysis of gentamicin-induced changes on ionic composition of the vestibular gelatinous membrane. *Hearing Research*, **1994**, 76, 60-6	3.9	6
7	Gentamicin ototoxicity in otoconia: quantitative electron probe X-ray microanalysis. *Acta Oto-Laryngologica*, **1994**, 114, 18-23	1.6	7
6	Standards for quantification of elements in the otolithic membrane by electron probe X-ray microanalysis: calibration curves and electron beam sensitivity. *Journal of Microscopy*, **1993**, 171, 215-22	1.9	8
5	X-ray microanalytical determination of P, S and K concentrations in the gelatinous membrane of the utricle. *Acta Oto-Laryngologica*, **1993**, 113, 176-80	1.6	5
4	Electron probe X-ray microanalysis of Ca and K distributions in the otolithic membrane. *Micron and Microscopica Acta*, **1992**, 23, 349-350		9
3	Dependence between saccule and utricle in Ca and K concentrations determined by electron probe X-ray microanalysis. *Micron and Microscopica Acta*, **1992**, 23, 367-368		5
2	Rare Variants in theOTOGGene Are a Frequent Cause of Familial Meniere Disease		2
1	Excess of rare missense variants in hearing loss genes in sporadic Meniere disease		1