

# Nãria Morral

## List of Publications by Year in descending order

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53  
papers

4,046  
citations

147801

31  
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182427

51  
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54  
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54  
docs citations

54  
times ranked

2964  
citing authors

#	ARTICLE	IF	CITATIONS
1	Genomic DNA transfer with a high-capacity adenovirus vector results in improved in vivo gene expression and decreased toxicity. <i>Nature Genetics</i> , 1998, 18, 180-183.	21.4	641
2	Administration of helper-dependent adenoviral vectors and sequential delivery of different vector serotype for long-term liver-directed gene transfer in baboons. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 1999, 96, 12816-12821.	7.1	412
3	The origin of the major cystic fibrosis mutation ( $\Delta$ F508) in European populations. <i>Nature Genetics</i> , 1994, 7, 169-175.	21.4	323
4	High Doses of a Helper-Dependent Adenoviral Vector Yield Supraphysiological Levels of $\Delta$ 1-Antitrypsin with Negligible Toxicity. <i>Human Gene Therapy</i> , 1998, 9, 2709-2716.	2.7	249
5	Immune Responses to Reporter Proteins and High Viral Dose Limit Duration of Expression with Adenoviral Vectors: Comparison of E2a Wild Type and E2a Deleted Vectors. <i>Human Gene Therapy</i> , 1997, 8, 1275-1286.	2.7	175
6	Use of a Liver-Specific Promoter Reduces Immune Response to the Transgene in Adenoviral Vectors. <i>Human Gene Therapy</i> , 1999, 10, 1773-1781.	2.7	174
7	Lethal Toxicity, Severe Endothelial Injury, and a Threshold Effect with High Doses of an Adenoviral Vector in Baboons. <i>Human Gene Therapy</i> , 2002, 13, 143-154.	2.7	160
8	CAGT Microsatellite alleles within the cystic fibrosis transmembrane conductance regulator (CFTR) gene are not generated by unequal crossingover. <i>Genomics</i> , 1991, 10, 692-698.	2.9	129
9	Toxicological Comparison of E2a-Deleted and First-Generation Adenoviral Vectors Expressing $\Delta$ 1-Antitrypsin after Systemic Delivery. <i>Human Gene Therapy</i> , 1998, 9, 1587-1598.	2.7	118
10	The search for South European cystic fibrosis mutations: Identification of two new mutations, four variants, and intronic sequences. <i>Genomics</i> , 1991, 10, 193-200.	2.9	117
11	Sterol Regulatory Element-binding Protein-1 (SREBP-1) Is Required to Regulate Glycogen Synthesis and Gluconeogenic Gene Expression in Mouse Liver. <i>Journal of Biological Chemistry</i> , 2014, 289, 5510-5517.	3.4	102
12	Microsatellite haplotypes for cystic fibrosis: mutation frameworks and evolutionary tracers. <i>Human Molecular Genetics</i> , 1993, 2, 1015-1022.	2.9	97
13	Development of a complementing cell line and a system for construction of adenovirus vectors with E1 and E2a deleted. <i>Journal of Virology</i> , 1996, 70, 7030-7038.	3.4	89
14	Toxicity Associated with Repeated Administration of First-Generation Adenovirus Vectors Does Not Occur with a Helper-Dependent Vector. <i>Molecular Medicine</i> , 2000, 6, 179-195.	4.4	79
15	Multiplex PCR amplification of three microsatellites within the CFTR gene. <i>Genomics</i> , 1992, 13, 1362-1364.	2.9	73
16	Gene therapy for inherited retinal and optic nerve degenerations. <i>Expert Opinion on Biological Therapy</i> , 2018, 18, 37-49.	3.1	72
17	Basal Insulin Gene Expression Significantly Improves Conventional Insulin Therapy in Type 1 Diabetic Rats. <i>Diabetes</i> , 2002, 51, 130-138.	0.6	71
18	Analysis of the CFTR gene confirms the high genetic heterogeneity of the Spanish population: 43 mutations account for only 78% of CF chromosomes. <i>Human Genetics</i> , 1994, 93, 447-51.	3.8	65

#	ARTICLE	IF	CITATIONS
19	Novel targets and therapeutic strategies for type 2 diabetes. Trends in Endocrinology and Metabolism, 2003, 14, 169-175.	7.1	58
20	Lack of liver glycogen causes hepatic insulin resistance and steatosis in mice. Journal of Biological Chemistry, 2017, 292, 10455-10464.	3.4	58
21	ΔF508 GENE DELETION IN CYSTIC FIBROSIS IN SOUTHERN EUROPE. Lancet, The, 1989, 334, 1404.	13.7	56
22	Gene therapy for age-related macular degeneration. Expert Opinion on Biological Therapy, 2017, 17, 1235-1244.	3.1	53
23	Transcription releases protein VII from adenovirus chromatin. Virology, 2007, 369, 411-422.	2.4	50
24	Hepatic insulin expression improves glycemic control in type 1 diabetic rats. Diabetes Research and Clinical Practice, 2001, 52, 153-163.	2.8	44
25	Effects of glucose metabolism on the regulation of genes of fatty acid synthesis and triglyceride secretion in the liver. Journal of Lipid Research, 2007, 48, 1499-1510.	4.2	43
26	Haplotype analysis of 94 cystic fibrosis mutations with seven polymorphicCFTR DNA markers. , 1996, 8, 149-159.		42
27	A tetranucleotide repeat polymorphism in the cystic fibrosis gene. Human Genetics, 1991, 86, 625.	3.8	35
28	Cystic fibrosis in Spain: high frequency of mutation G542X in the Mediterranean coastal area. Human Genetics, 1993, 91, 66-70.	3.8	35
29	Complete detection of mutations in cystic fibrosis patients of Native American origin. Human Genetics, 1994, 94, 629-32.	3.8	35
30	Cystic fibrosis in a low-incidence population: two major mutations in Finland. Human Genetics, 1994, 93, 162-166.	3.8	35
31	Adenovirus-Mediated Expression of Glucokinase in the Liver as an Adjuvant Treatment for Type 1 Diabetes. Human Gene Therapy, 2002, 13, 1561-1570.	2.7	33
32	Helper-dependent Adenovirus-mediated Short Hairpin RNA Expression in the Liver Activates the Interferon Response. Journal of Biological Chemistry, 2008, 283, 2120-2128.	3.4	33
33	Uniparental inheritance of microsatellite alleles of the cystic fibrosis gene (CFTR): identification of a 50 kilobase deletion. Human Molecular Genetics, 1993, 2, 677-681.	2.9	32
34	CFTR haplotypic variability for normal and mutant genes in cystic fibrosis families from southern France. Human Genetics, 1996, 98, 336-344.	3.8	29
35	Reply to "Age of the ΔF508 cystic fibrosis mutation. Nature Genetics, 1994, 8, 216-218.	21.4	25
36	Comparative nucleic acid transfection efficacy in primary hepatocytes for gene silencing and functional studies. BMC Research Notes, 2011, 4, 8.	1.4	22

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37	Dinucleotide (CA/GT) repeat polymorphism in intron 17B of the cystic fibrosis transmembrane conductance regulator (CFTR) gene. <i>Human Genetics</i> , 1992, 88, 356.	3.8	21
38	Robust Hepatic Gene Silencing for Functional Studies Using Helper-Dependent Adenoviral Vectors. <i>Human Gene Therapy</i> , 2009, 20, 87-94.	2.7	21
39	Accurate single-day titration of adenovirus vectors based on equivalence of protein VII nuclear dots and infectious particles. <i>Journal of Virological Methods</i> , 2009, 159, 251-258.	2.1	19
40	Analysis of microsatellites by direct blotting electrophoresis and chemiluminescence detection. <i>Electrophoresis</i> , 1995, 16, 1886-1888.	2.4	18
41	Identical Intragenic Microsatellite Haplotype Found in Cystic Fibrosis Chromosomes Bearing Mutation G551D in Irish, English, Scottish, Breton and Czech Patients. <i>Human Heredity</i> , 1995, 45, 6-12.	0.8	18
42	Prenatal diagnosis of cystic fibrosis by multiplex PCR of mutation and microsatellite alleles. <i>Lancet</i> , The, 1991, 338, 458.	13.7	12
43	Gene targets of mouse miR-709: regulation of distinct pools. <i>Scientific Reports</i> , 2016, 6, 18958.	3.3	12
44	Constitutive Expression of Short Hairpin RNA in Vivo Triggers Buildup of Mature Hairpin Molecules. <i>Human Gene Therapy</i> , 2011, 22, 1483-1497.	2.7	11
45	Impact of silencing hepatic SREBP-1 on insulin signaling. <i>PLoS ONE</i> , 2018, 13, e0196704.	2.5	9
46	Vector and Helper Genome Rearrangements Occur During Production of Helper-Dependent Adenoviral Vectors. <i>Human Gene Therapy Methods</i> , 2013, 24, 1-10.	2.1	7
47	shRNA-Induced Interferon-Stimulated Gene Analysis. <i>Methods in Molecular Biology</i> , 2012, 820, 163-177.	0.9	6
48	Challenges for Gene Therapy of Type 1 Diabetes. <i>Current Gene Therapy</i> , 2002, 2, 403-414.	2.0	6
49	Role of non-coding RNAs on liver metabolism and NAFLD pathogenesis. <i>Human Molecular Genetics</i> , 2022, 31, R4-R21.	2.9	6
50	Enhancing hepatic mitochondrial fatty acid oxidation stimulates eating in food-deprived mice. <i>American Journal of Physiology - Regulatory Integrative and Comparative Physiology</i> , 2015, 308, R131-R137.	1.8	5
51	Aberrant gene expression induced by a high fat diet is linked to H3K9 acetylation in the promoter-proximal region. <i>Biochimica Et Biophysica Acta - Gene Regulatory Mechanisms</i> , 2021, 1864, 194691.	1.9	5
52	Haplotype analysis of 94 cystic fibrosis mutations with seven polymorphic CFTR DNA markers. <i>Human Mutation</i> , 1996, 8, 149-159.	2.5	5
53	Insights from a high-fat diet fed mouse model with a humanized liver. <i>PLoS ONE</i> , 2022, 17, e0268260.	2.5	1