

# Nãria Morral

## List of Publications by Year in descending order

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53  
papers

4,046  
citations

147801

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docs citations

54  
times ranked

2964  
citing authors

#	ARTICLE	IF	CITATIONS
1	Role of non-coding RNAs on liver metabolism and NAFLD pathogenesis. <i>Human Molecular Genetics</i> , 2022, 31, R4-R21.	2.9	6
2	Insights from a high-fat diet fed mouse model with a humanized liver. <i>PLoS ONE</i> , 2022, 17, e0268260.	2.5	1
3	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
4	Gene therapy for inherited retinal and optic nerve degenerations. <i>Expert Opinion on Biological Therapy</i> , 2018, 18, 37-49.	3.1	72
5	Impact of silencing hepatic SREBP-1 on insulin signaling. <i>PLoS ONE</i> , 2018, 13, e0196704.	2.5	9
6	Lack of liver glycogen causes hepatic insulin resistance and steatosis in mice. <i>Journal of Biological Chemistry</i> , 2017, 292, 10455-10464.	3.4	58
7	Gene therapy for age-related macular degeneration. <i>Expert Opinion on Biological Therapy</i> , 2017, 17, 1235-1244.	3.1	53
8	Gene targets of mouse miR-709: regulation of distinct pools. <i>Scientific Reports</i> , 2016, 6, 18958.	3.3	12
9	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
10	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
11	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
12	shRNA-Induced Interferon-Stimulated Gene Analysis. <i>Methods in Molecular Biology</i> , 2012, 820, 163-177.	0.9	6
13	Comparative nucleic acid transfection efficacy in primary hepatocytes for gene silencing and functional studies. <i>BMC Research Notes</i> , 2011, 4, 8.	1.4	22
14	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
15	Robust Hepatic Gene Silencing for Functional Studies Using Helper-Dependent Adenoviral Vectors. <i>Human Gene Therapy</i> , 2009, 20, 87-94.	2.7	21
16	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
17	Helper-dependent Adenovirus-mediated Short Hairpin RNA Expression in the Liver Activates the Interferon Response. <i>Journal of Biological Chemistry</i> , 2008, 283, 2120-2128.	3.4	33
18	Effects of glucose metabolism on the regulation of genes of fatty acid synthesis and triglyceride secretion in the liver. <i>Journal of Lipid Research</i> , 2007, 48, 1499-1510.	4.2	43

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19	Transcription releases protein VII from adenovirus chromatin. <i>Virology</i> , 2007, 369, 411-422.	2.4	50
20	Novel targets and therapeutic strategies for type 2 diabetes. <i>Trends in Endocrinology and Metabolism</i> , 2003, 14, 169-175.	7.1	58
21	Adenovirus-Mediated Expression of Glucokinase in the Liver as an Adjuvant Treatment for Type 1 Diabetes. <i>Human Gene Therapy</i> , 2002, 13, 1561-1570.	2.7	33
22	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
23	Basal Insulin Gene Expression Significantly Improves Conventional Insulin Therapy in Type 1 Diabetic Rats. <i>Diabetes</i> , 2002, 51, 130-138.	0.6	71
24	Challenges for Gene Therapy of Type 1 Diabetes. <i>Current Gene Therapy</i> , 2002, 2, 403-414.	2.0	6
25	Hepatic insulin expression improves glycemic control in type 1 diabetic rats. <i>Diabetes Research and Clinical Practice</i> , 2001, 52, 153-163.	2.8	44
26	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
27	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
28	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
29	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
30	Toxicological Comparison of E2a-Deleted and First-Generation Adenoviral Vectors Expressing $\alpha$ -Antitrypsin after Systemic Delivery. <i>Human Gene Therapy</i> , 1998, 9, 1587-1598.	2.7	118
31	High Doses of a Helper-Dependent Adenoviral Vector Yield Supraphysiological Levels of $\alpha$ -Antitrypsin with Negligible Toxicity. <i>Human Gene Therapy</i> , 1998, 9, 2709-2716.	2.7	249
32	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
33	CFTR haplotypic variability for normal and mutant genes in cystic fibrosis families from southern France. <i>Human Genetics</i> , 1996, 98, 336-344.	3.8	29
34	Haplotype analysis of 94 cystic fibrosis mutations with seven polymorphic CFTR DNA markers. , 1996, 8, 149-159.		42
35	Haplotype analysis of 94 cystic fibrosis mutations with seven polymorphic CFTR DNA markers. <i>Human Mutation</i> , 1996, 8, 149-159.	2.5	5
36	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

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37	Analysis of microsatellites by direct blotting electrophoresis and chemiluminescence detection. Electrophoresis, 1995, 16, 1886-1888.	2.4	18
38	Identical Intragenic Microsatellite Haplotype Found in Cystic Fibrosis Chromosomes Bearing Mutation G551D in Irish, English, Scottish, Breton and Czech Patients. Human Heredity, 1995, 45, 6-12.	0.8	18
39	Analysis of the CFTR gene confirms the high genetic heterogeneity of the Spanish population: 43 mutations account for only 78% of CF chromosomes. Human Genetics, 1994, 93, 447-51.	3.8	65
40	Complete detection of mutations in cystic fibrosis patients of Native American origin. Human Genetics, 1994, 94, 629-32.	3.8	35
41	Cystic fibrosis in a low-incidence population: two major mutations in Finland. Human Genetics, 1994, 93, 162-166.	3.8	35
42	The origin of the major cystic fibrosis mutation ( $\Delta$ F508) in European populations. Nature Genetics, 1994, 7, 169-175.	21.4	323
43	Reply to "Age of the $\Delta$ F508 cystic fibrosis mutation. Nature Genetics, 1994, 8, 216-218.	21.4	25
44	Cystic fibrosis in Spain: high frequency of mutation G542X in the Mediterranean coastal area. Human Genetics, 1993, 91, 66-70.	3.8	35
45	Microsatellite haplotypes for cystic fibrosis: mutation frameworks and evolutionary tracers. Human Molecular Genetics, 1993, 2, 1015-1022.	2.9	97
46	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
47	Multiplex PCR amplification of three microsatellites within the CFTR gene. Genomics, 1992, 13, 1362-1364.	2.9	73
48	Dinucleotide (CA/GT) repeat polymorphism in intron 17B of the cystic fibrosis transmembrane conductance regulator (CFTR) gene. Human Genetics, 1992, 88, 356.	3.8	21
49	CA/GT Microsatellite alleles within the cystic fibrosis transmembrane conductance regulator (CFTR) gene are not generated by unequal crossingover. Genomics, 1991, 10, 692-698.	2.9	129
50	The search for South European cystic fibrosis mutations: Identification of two new mutations, four variants, and intronic sequences. Genomics, 1991, 10, 193-200.	2.9	117
51	Prenatal diagnosis of cystic fibrosis by multiplex PCR of mutation and microsatellite alleles. Lancet, The, 1991, 338, 458.	13.7	12
52	A tetranucleotide repeat polymorphism in the cystic fibrosis gene. Human Genetics, 1991, 86, 625.	3.8	35
53	$\Delta$ F508 GENE DELETION IN CYSTIC FIBROSIS IN SOUTHERN EUROPE. Lancet, The, 1989, 334, 1404.	13.7	56