## Enrico Moro

## List of Publications by Year in descending order

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59 3,514 32 58
papers citations h-index g-index

62 62 62 4650 all docs docs citations times ranked citing authors

#	Article	IF	CITATIONS
1	Y Chromosome Microdeletions and Alterations of Spermatogenesis*. Endocrine Reviews, 2001, 22, 226-239.	8.9	347
2	Deletion and expression analysis of AZFa genes on the human Y chromosome revealed a major role for DBY in male infertility. Human Molecular Genetics, 2000, 9, 1161-1169.	1.4	227
3	In vivo Wnt signaling tracing through a transgenic biosensor fish reveals novel activity domains. Developmental Biology, 2012, 366, 327-340.	0.9	227
4	Laboratory guidelines for molecular diagnosis of Y-chromosomal microdeletions. Journal of Developmental and Physical Disabilities, 1999, 22, 292-299.	3 <b>.</b> 6	218
5	Long-range gene regulation links genomic type 2 diabetes and obesity risk regions to <i>HHEX</i> , <i>SOX4</i> , and <i>IRX3</i> . Proceedings of the National Academy of Sciences of the United States of America, 2010, 107, 775-780.	3.3	189
6	High frequency of well-defined Y-chromosome deletions in idiopathic Sertoli cell-only syndrome. Human Reproduction, 1998, 13, 302-307.	0.4	186
7	Wnt/ $\hat{l}^2$ -Catenin Signaling Defines Organizing Centers that Orchestrate Growth and Differentiation of the Regenerating Zebrafish Caudal Fin. Cell Reports, 2014, 6, 467-481.	2.9	163
8	Human male infertility and Y chromosome deletions: role of the AZF-candidate genes DAZ, RBM and DFFRY. Human Reproduction, 1999, 14, 1710-1716.	0.4	138
9	Wnt Signaling Regulates Postembryonic Hypothalamic Progenitor Differentiation. Developmental Cell, 2012, 23, 624-636.	3.1	90
10	Wnt activation promotes neuronal differentiation of Glioblastoma. Cell Death and Disease, 2013, 4, e500-e500.	2.7	89
11	Pituitary tumor transforming gene-null male mice exhibit impaired pancreatic beta cell proliferation and diabetes. Proceedings of the National Academy of Sciences of the United States of America, 2003, 100, 3428-3432.	3.3	87
12	Zebrafish pancreas development. Molecular and Cellular Endocrinology, 2009, 312, 24-30.	1.6	79
13	Y Chromosome Microdeletions in Cryptorchidism and Idiopathic Infertility*. Journal of Clinical Endocrinology and Metabolism, 1999, 84, 3660-3665.	1.8	74
14	Diverse Chemical Scaffolds Support Direct Inhibition of the Membrane-bound O-Acyltransferase Porcupine. Journal of Biological Chemistry, 2012, 287, 23246-23254.	1.6	72
15	Intracardiac flow dynamics regulate atrioventricular valve morphogenesis. Cardiovascular Research, 2014, 104, 49-60.	1.8	67
16	Y Chromosome Microdeletions in Cryptorchidism and Idiopathic Infertility. Journal of Clinical Endocrinology and Metabolism, 1999, 84, 3660-3665.	1.8	67
17	Generation and application of signaling pathway reporter lines in zebrafish. Molecular Genetics and Genomics, 2013, 288, 231-242.	1.0	66
18	Lef1-dependent Wnt $\hat{l}^2$ -catenin signalling drives the proliferative engine that maintains tissue homeostasis during lateral line development. Development (Cambridge), 2011, 138, 3931-3941.	1.2	65

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19	RAB8B Is Required for Activity and Caveolar Endocytosis of LRP6. Cell Reports, 2013, 4, 1224-1234.	2.9	65
20	The pathogenesis of lysosomal storage disorders: beyond the engorgement of lysosomes to abnormal development and neuroinflammation. Human Molecular Genetics, 2018, 27, R119-R129.	1.4	59
21	Impaired Mitochondrial ATP Production Downregulates Wnt Signaling via ER Stress Induction. Cell Reports, 2019, 28, 1949-1960.e6.	2.9	56
22	A Smad3 transgenic reporter reveals TGF-beta control of zebrafish spinal cord development. Developmental Biology, 2014, 396, 81-93.	0.9	52
23	Using transgenic reporters to visualize bone and cartilage signaling during development in vivo. Frontiers in Endocrinology, 2012, 3, 91.	1.5	47
24	Glucocerebrosidase deficiency in zebrafish affects primary bone ossification through increased oxidative stress and reduced Wnt/l²-catenin signaling. Human Molecular Genetics, 2015, 24, 1280-1294.	1.4	46
25	Novel insulin-like 3 (INSL3) gene mutation associated with human cryptorchidism. American Journal of Medical Genetics Part A, 2001, 103, 348-349.	2.4	43
26	Emilin3 is required for notochord sheath integrity and interacts with Scube2 to regulate notochord-derived Hedgehog signals. Development (Cambridge), 2013, 140, 4594-4601.	1.2	38
27	Male Infertility Caused by a de Novo Partial Deletion of the DAZ Cluster on the Y Chromosome 1. Journal of Clinical Endocrinology and Metabolism, 2000, 85, 4069-4073.	1.8	37
28	A novel functional role of iduronate-2-sulfatase in zebrafish early development. Matrix Biology, 2010, 29, 43-50.	1.5	37
29	Zebrafish reporter lines reveal in vivo signaling pathway activities involved in pancreatic cancer. DMM Disease Models and Mechanisms, 2014, 7, 883-94.	1.2	37
30	Simplet/Fam53b is required for Wnt signal transduction by regulating $\hat{l}^2$ -catenin nuclear localization. Development (Cambridge), 2014, 141, 3529-3539.	1.2	35
31	Role of the AZFa candidate genes in male infertility. Journal of Endocrinological Investigation, 2000, 23, 646-651.	1.8	34
32	A living biosensor model to dynamically trace glucocorticoid transcriptional activity during development and adult life in zebrafish. Molecular and Cellular Endocrinology, 2014, 392, 60-72.	1.6	34
33	Perturbations in cell signaling elicit early cardiac defects in mucopolysaccharidosis type II. Human Molecular Genetics, 2017, 26, 1643-1655.	1.4	34
34	Y chromosome microdeletions in infertile men with varicocele. Molecular and Cellular Endocrinology, 2000, 161, 67-71.	1.6	33
35	Different insulin-like 3 (INSL3) gene mutations not associated with human cryptorchidism. Journal of Endocrinological Investigation, 2001, 24, RC13-RC15.	1.8	33
36	Developmental defects and neuromuscular alterations due to mitofusin 2 gene (MFN2) silencing in zebrafish: a new model for Charcot-Marie-Tooth type 2A neuropathy. Neuromuscular Disorders, 2011, 21, 58-67.	0.3	33

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37	Male Infertility Caused by a de Novo Partial Deletion of the DAZ Cluster on the Y Chromosome. Journal of Clinical Endocrinology and Metabolism, 2000, 85, 4069-4073.	1.8	33
38	Y-chromosomal DNA haplotypes in infertile European males carrying Y-microdeletions. Journal of Endocrinological Investigation, 2000, 23, 671-676.	1.8	32
39	Absence of testicular DAZ gene expression in idiopathic severe testiculopathies. Human Reproduction, 1999, 14, 2286-2292.	0.4	27
40	FGF signaling deregulation is associated with early developmental skeletal defects in animal models for mucopolysaccharidosis type II (MPSII). Human Molecular Genetics, 2018, 27, 2262-2275.	1.4	27
41	Analysis of beta cell proliferation dynamics in zebrafish. Developmental Biology, 2009, 332, 299-308.	0.9	24
42	Sertoli Cell Function in Infertile Patients with and without Microdeletions of theAzoospermia Factorson the Y Chromosome Long Arm1. Journal of Clinical Endocrinology and Metabolism, 2001, 86, 2414-2419.	1.8	20
43	Synthesis and biological evaluation of imidazo[1,2-a]pyrimidines and imidazo[1,2-a]pyridines as new inhibitors of the Wnt/ $\hat{l}^2$ -catenin signaling. European Journal of Medicinal Chemistry, 2014, 83, 45-56.	2.6	16
44	"Janus―efficacy of CX-5011: CK2 inhibition and methuosis induction by independent mechanisms. Biochimica Et Biophysica Acta - Molecular Cell Research, 2020, 1867, 118807.	1.9	14
45	Temporal control of Wnt signaling is required for habenular neuron diversity and brain asymmetry. Development (Cambridge), 2020, 147, .	1.2	14
46	Sertoli Cell Function in Infertile Patients with and without Microdeletions of the Azoospermia Factors on the Y Chromosome Long Arm. Journal of Clinical Endocrinology and Metabolism, 2001, 86, 2414-2419.	1.8	14
47	Y chromosome. Lancet, The, 2000, 355, 234-235.	6.3	13
48	A novel approach for the analysis of DAZ gene copy number in severely idiopathic infertile men. Journal of Endocrinological Investigation, 2002, 25, RC1-RC3.	1.8	13
49	The Golgi  casein kinase' Fam20C is a genuine  phosvitin kinase' and phosphorylates polyserine stretches devoid of the canonical consensus. FEBS Journal, 2018, 285, 4674-4683.	2.2	10
50	Inhibin B plasma concentrations in infertile patients with DAZ gene deletions treated with FSH. European Journal of Endocrinology, 2002, 146, 801-806.	1.9	9
51	Lysosomal Function and Axon Guidance: Is There a Meaningful Liaison?. Biomolecules, 2021, 11, 191.	1.8	8
52	CDY 1 analysis in infertile patients with DAZ deletions. Journal of Endocrinological Investigation, 2001, 24, RC4-RC6.	1.8	7
53	Zebrafish spata2 is expressed at early developmental stages. International Journal of Developmental Biology, 2007, 51, 241-246.	0.3	7
54	Response to local dihydrotestosterone treatment in a patient with partial androgen-insensitivity syndrome due to a novel mutation in the androgen receptor gene. American Journal of Medical Genetics Part A, 2002, 107, 259-260.	2.4	6

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55	A transcriptional and post-transcriptional dysregulation of Dishevelled 1 and 2 underlies the Wnt signaling impairment in type I Gaucher disease experimental models. Human Molecular Genetics, 2020, 29, 274-285.	1.4	4
56	Lysosomal Storage Disorders: Molecular Basis and Therapeutic Approaches. Biomolecules, 2021, 11, 964.	1.8	3
57	Novel insulinâ€ike 3 (INSL3) gene mutation associated with human cryptorchidism. American Journal of Medical Genetics Part A, 2001, 103, 348-349.	2.4	1
58	af9 Regulates gata2 Expression During Early Hemangioblast Specification and Vascular Pattern Formation In Zebrafish Blood, 2010, 116, 2600-2600.	0.6	1
59	Simplet/Fam53b is required for Wnt signal transduction by regulating $\hat{I}^2$ -catenin nuclear localization. Journal of Cell Science, 2014, 127, e1-e1.	1.2	0