

# Patrick R Benusiglio

## List of Publications by Year in descending order

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Version: 2024-02-01

42  
papers

2,383  
citations

361413

20  
h-index

265206

42  
g-index

44  
all docs

44  
docs citations

44  
times ranked

3906  
citing authors

#	ARTICLE	IF	CITATIONS
1	Revisiting Li-Fraumeni Syndrome From <i>TP53</i> Mutation Carriers. <i>Journal of Clinical Oncology</i> , 2015, 33, 2345-2352.	1.6	525
2	Hereditary diffuse gastric cancer: updated clinical guidelines with an emphasis on germline <i>CDH1</i> mutation carriers. <i>Journal of Medical Genetics</i> , 2015, 52, 361-374.	3.2	479
3	Hereditary diffuse gastric cancer: updated clinical practice guidelines. <i>Lancet Oncology</i> , The, 2020, 21, e386-e397.	10.7	237
4	<i>CDH1</i> germline mutations and the hereditary diffuse gastric and lobular breast cancer syndrome: a multicentre study. <i>Journal of Medical Genetics</i> , 2013, 50, 486-489.	3.2	131
5	Lobular breast cancer: incidence and genetic and non-genetic risk factors. <i>Breast Cancer Research</i> , 2015, 17, 37.	5.0	126
6	Reassessing the clinical spectrum associated with hereditary leiomyomatosis and renal cell carcinoma syndrome in French <i>scp&gt;FH&lt;/scp&gt;</i> mutation carriers. <i>Clinical Genetics</i> , 2017, 92, 606-615.	2.0	103
7	Common variants in the ATM, BRCA1, BRCA2, CHEK2 and TP53 cancer susceptibility genes are unlikely to increase breast cancer risk. <i>Breast Cancer Research</i> , 2007, 9, R27.	5.0	94
8	Renal cell tumour characteristics in patients with the Birt-Hogg-DubÃ© cancer susceptibility syndrome: a retrospective, multicentre study. <i>Orphanet Journal of Rare Diseases</i> , 2014, 9, 163.	2.7	78
9	Association of single-nucleotide polymorphisms in the cell cycle genes with breast cancer in the British population. <i>Carcinogenesis</i> , 2008, 29, 333-341.	2.8	68
10	Pattern multiplicity and fumarate hydratase (FH)/S-(2-succino)-cysteine (2SC) staining but not eosinophilic nucleoli with perinucleolar halos differentiate hereditary leiomyomatosis and renal cell carcinoma-associated renal cell carcinomas from kidney tumors without FH gene alteration. <i>Modern Pathology</i> , 2018, 31, 974-983.	5.5	65
11	Common ERBB2 polymorphisms and risk of breast cancer in a white British population: a caseâ€“control study. <i>Breast Cancer Research</i> , 2005, 7, R204-9.	5.0	55
12	Lung cancer is also a hereditary disease. <i>European Respiratory Review</i> , 2021, 30, 210045.	7.1	39
13	A germline mutation in <i>PBRM1</i> predisposes to renal cell carcinoma. <i>Journal of Medical Genetics</i> , 2015, 52, 426-430.	3.2	38
14	HapMap-based study of the 17q21 ERBB2 amplicon in susceptibility to breast cancer. <i>British Journal of Cancer</i> , 2006, 95, 1689-1695.	6.4	35
15	Genetic Evidence of a Precisely Tuned Dysregulation in the Hypoxia Signaling Pathway during Oncogenesis. <i>Cancer Research</i> , 2014, 74, 6554-6564.	0.9	32
16	Cleft lip, cleft palate, hereditary diffuse gastric cancer and germline mutations in <i>CDH1</i> . <i>International Journal of Cancer</i> , 2013, 132, 2470-2470.	5.1	29
17	Clinical implications of CTNNA1 germline mutations in asymptomatic carriers. <i>Gastric Cancer</i> , 2019, 22, 899-903.	5.3	27
18	Lung Adenocarcinoma as Part of the Li-Fraumeni Syndrome Spectrum. <i>JAMA Oncology</i> , 2017, 3, 1736.	7.1	24

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19	Hereditary diffuse gastric cancer syndrome: improved performances of the 2015 testing criteria for the identification of probands with a <i>CDH1</i> germline mutation. <i>Journal of Medical Genetics</i> , 2015, 52, 563-565.	3.2	22
20	Cancer predisposition and germline CTNNA1 variants. <i>European Journal of Medical Genetics</i> , 2021, 64, 104316.	1.3	21
21	Germline <i>CDKN2A</i> / <i>P16INK4A</i> mutations contribute to genetic determinism of sarcoma. <i>Journal of Medical Genetics</i> , 2017, 54, 607-612.	3.2	19
22	Response to systemic therapy in fumarate hydratase-deficient renal cell carcinoma. <i>European Journal of Cancer</i> , 2021, 151, 106-114.	2.8	18
23	Hereditary breast and ovarian cancer: successful systematic implementation of a group approach to genetic counselling. <i>Familial Cancer</i> , 2017, 16, 51-56.	1.9	15
24	Common variation in EMSY and risk of breast and ovarian cancer: a case-control study using HapMap tagging SNPs. <i>BMC Cancer</i> , 2005, 5, 81.	2.6	14
25	Classification of 101 BRCA1 and BRCA2 variants of uncertain significance by cosegregation study: A powerful approach. <i>American Journal of Human Genetics</i> , 2021, 108, 1907-1923.	6.2	14
26	Germline ATG2B/GSKIP-containing 14q32 duplication predisposes to early clonal hematopoiesis leading to myeloid neoplasms. <i>Leukemia</i> , 2022, 36, 126-137.	7.2	10
27	Low risk of invasive lobular carcinoma of the breast in carriers of <i>BRCA1</i> (hereditary breast) Tj ETQq1 1 0.784314 rgBT /Overl... 25, 16-19.	1.0	7
28	Utility of a mainstreamed genetic testing pathway in breast and ovarian cancer patients during the COVID-19 pandemic. <i>European Journal of Medical Genetics</i> , 2020, 63, 104098.	1.3	7
29	Case Report: Expanding the tumour spectrum associated with the Birt-Hogg-Dubois cancer susceptibility syndrome. <i>F1000Research</i> , 2014, 3, 159.	1.6	7
30	International Delphi consensus guidelines for follow-up after prophylactic total gastrectomy: the Life after Prophylactic Total Gastrectomy (LAP-TG) study. <i>Gastric Cancer</i> , 2022, 25, 1094-1104.	5.3	7
31	The Birt-Hogg-Dugois cancer predisposition syndrome: Current challenges. <i>Intractable and Rare Diseases Research</i> , 2015, 4, 162-163.	0.9	6
32	CDH1 germline mutations: different syndromes, same management?. <i>Genetics in Medicine</i> , 2017, 19, 965-966.	2.4	6
33	Whole-Body MRI Screening in Children With Li-Fraumeni and Other Cancer Predisposition Syndromes.. <i>American Journal of Roentgenology</i> , 2016, 206, W52-W52.	2.2	5
34	Focus on <i>ERBB2</i> . <i>Pharmacogenomics</i> , 2008, 9, 825-828.	1.3	4
35	Gastrointestinal relapse of multiple myeloma and sustained response to lenalidomide: a case report. <i>Journal of Medical Case Reports</i> , 2011, 5, 110.	0.8	3
36	Deficit of Risk-Reducing Salpingo-Oophorectomies in BRCA1/2 Mutation Carriers After Telephone Genetic Counseling. <i>Journal of Clinical Oncology</i> , 2017, 35, 1255-1255.	1.6	3

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37	Overcoming the challenges associated with universal screening for Lynch syndrome in colorectal and endometrial cancer. <i>Genetics in Medicine</i> , 2020, 22, 1422-1423.	2.4	3
38	Reply: a bias in genotyping of the ERBB2 (HER2) Ile655Val variant. <i>Carcinogenesis</i> , 2005, 26, 2213-2213.	2.8	1
39	Germline Variants in Highly Selected Patients With Prostate Cancer. <i>JAMA Oncology</i> , 2019, 5, 1367.	7.1	1
40	Preimplantation genetic testing in patients with genetic susceptibility to cancer. <i>Familial Cancer</i> , 2022, , 1.	1.9	1
41	Do Female <i>BRCA2</i> Pathogenic Variant Carriers Have an Increased Risk of Lung Cancer?. <i>Journal of Clinical Oncology</i> , 2022, , JCO2200488.	1.6	1
42	Serous ovarian carcinoma in patients with Lynch syndrome: Caution is warranted. <i>Gynecologic Oncology Reports</i> , 2018, 26, 69-70.	0.6	0