

Deborah J Marsh

List of Publications by Year in descending order

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90
papers

4,026
citations

172457

29
h-index

118850

62
g-index

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all docs

90
docs citations

90
times ranked

5501
citing authors

#	ARTICLE	IF	CITATIONS
1	Three-Dimensional Modelling of Ovarian Cancer: From Cell Lines to Organoids for Discovery and Personalized Medicine. <i>Frontiers in Bioengineering and Biotechnology</i> , 2022, 10, 836984.	4.1	22
2	The Anti-ROR1 Monoclonal Antibody Zilovertamab Inhibits the Proliferation of Ovarian and Endometrial Cancer Cells. <i>Pharmaceutics</i> , 2022, 14, 837.	4.5	6
3	Ubiquitin chromatin remodelling after DNA damage is associated with the expression of key cancer genes and pathways. <i>Cellular and Molecular Life Sciences</i> , 2021, 78, 1011-1027.	5.4	10
4	Studying the Oncosuppressive Functions of PTENP1 as a ceRNA. <i>Methods in Molecular Biology</i> , 2021, 2324, 165-185.	0.9	1
5	Abstract 1062: Inhibition of ovarian and endometrial cancer cell proliferation by an anti-ROR1 monoclonal antibody. , 2021, , .		0
6	PARP Inhibitors Display Differential Efficacy in Models of BRCA Mutant High-Grade Serous Ovarian Cancer. <i>International Journal of Molecular Sciences</i> , 2021, 22, 8506.	4.1	8
7	Amphiregulin increases migration and proliferation of epithelial ovarian cancer cells by inducing its own expression via PI3-kinase signaling. <i>Molecular and Cellular Endocrinology</i> , 2021, 533, 111338.	3.2	6
8	An organotypic model of high-grade serous ovarian cancer to test the anti-metastatic potential of ROR2 targeted Polyion complex nanoparticles. <i>Journal of Materials Chemistry B</i> , 2021, 9, 9123-9135.	5.8	11
9	Histone Monoubiquitination in Chromatin Remodelling: Focus on the Histone H2B Interactome and Cancer. <i>Cancers</i> , 2020, 12, 3462.	3.7	26
10	Toward Systems Pathology for PTEN Diagnostics. <i>Cold Spring Harbor Perspectives in Medicine</i> , 2020, 10, a037127.	6.2	4
11	The role of the free β -subunit of human chorionic gonadotropin in human malignancy. , 2020, , 269-281.		0
12	Writing Histone Monoubiquitination in Human Malignancyâ€™The Role of RING Finger E3 Ubiquitin Ligases. <i>Genes</i> , 2019, 10, 67.	2.4	35
13	Parafibromin-deficient (HPT-JT Type, CDC73 Mutated) Parathyroid Tumors Demonstrate Distinctive Morphologic Features. <i>American Journal of Surgical Pathology</i> , 2019, 43, 35-46.	3.7	74
14	Combining serum microRNA and CA-125 as prognostic indicators of preoperative surgical outcome in women with high-grade serous ovarian cancer. <i>Gynecologic Oncology</i> , 2018, 148, 181-188.	1.4	25
15	Histone H2B. , 2018, , 2384-2388.		0
16	CDC73. , 2018, , 991-995.		0
17	Abstract 3538: Targeting the E3 ubiquitin ligase RNF20 in ovarian cancer. , 2018, , .		0
18	Abstract A06: Cisplatin-induced DNA damage modifies the chromatin landscape of histone H2B monoubiquitination in a p53-dependent manner. , 2018, , .		0

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19	Comprehensive analyses of somatic TP53 mutation in tumors with variable mutant allele frequency. <i>Scientific Data</i> , 2017, 4, 170120.	5.3	9
20	Lessons learnt from outstanding mid-career women in endocrine cancer research. <i>Endocrine-Related Cancer</i> , 2016, 23, E5-E7.	3.1	0
21	Assessing mutant p53 in primary high-grade serous ovarian cancer using immunohistochemistry and massively parallel sequencing. <i>Scientific Reports</i> , 2016, 6, 26191.	3.3	162
22	The RING finger domain E3 ubiquitin ligases BRCA1 and the RNF20/RNF40 complex in global loss of the chromatin mark histone H2B monoubiquitination (H2Bub1) in cell line models and primary high-grade serous ovarian cancer. <i>Human Molecular Genetics</i> , 2016, 25, ddu362.	2.9	26
23	Host-Guest Complexes of Carboxylated Pillar[n]arenes With Drugs. <i>Journal of Pharmaceutical Sciences</i> , 2016, 105, 3615-3625.	3.3	40
24	Comparison of Methodologies to Detect Low Levels of Hemolysis in Serum for Accurate Assessment of Serum microRNAs. <i>PLoS ONE</i> , 2016, 11, e0153200.	2.5	160
25	Histone H2B. , 2016, , 1-5.		0
26	CDC73. , 2016, , 1-5.		0
27	Abstract B05: Assessment of TP53 mutation status in primary high-grade serous ovarian cancer and cell line models: Comparison between immunohistochemistry and next-generation sequencing.. , 2016, , .		0
28	Cowden Syndrome. , 2016, , 1218-1222.		0
29	Match that PhD. <i>Nature</i> , 2015, 523, 247-247.	27.8	2
30	Histone H2B monoubiquitination: roles to play in human malignancy. <i>Endocrine-Related Cancer</i> , 2015, 22, T19-T33.	3.1	108
31	Networks regulating ubiquitin and ubiquitin-like proteins promise new therapeutic targets. <i>Endocrine-Related Cancer</i> , 2015, 22, E1-E3.	3.1	1
32	A novel truncated form of S100P predicts disease-free survival in patients with lymph node positive breast cancer. <i>Cancer Letters</i> , 2015, 368, 64-70.	7.2	19
33	Genomic alterations as mediators of miRNA dysregulation in ovarian cancer. <i>Genes Chromosomes and Cancer</i> , 2015, 54, 1-19.	2.8	23
34	Abstract 1047: A role for the free beta subunit of human chorionic gonadotropin in sensitivity of epithelial ovarian cancer cells to platinum-based chemotherapeutics. , 2015, , .		0
35	Novel serum protein biomarker panel revealed by mass spectrometry and its prognostic value in breast cancer. <i>Breast Cancer Research</i> , 2014, 16, R63.	5.0	90
36	Factors that May Influence the Willingness of Cancer Patients to Consent for Biobanking. <i>Biopreservation and Biobanking</i> , 2014, 12, 409-414.	1.0	10

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37	Histones and Their Modifications in Ovarian Cancer – Drivers of Disease and Therapeutic Targets. <i>Frontiers in Oncology</i> , 2014, 4, 144.	2.8	46
38	Cowden Syndrome. , 2014, , 1-6.		0
39	Tissue biomarkers of breast cancer and their association with conventional pathologic features. <i>British Journal of Cancer</i> , 2013, 108, 351-360.	6.4	27
40	Abstract A29: Loss of histone H2B monoubiquitination in ovarian cancer – new therapeutic targeting opportunities based on chromatin relaxation. , 2013, , .		1
41	Abstract 330: Utilization of Sleeping Beauty mutagenesis for the identification of potential driver genes of ovarian cancer.. , 2013, , .		0
42	Abstract A10: A mutagenesis screen identifies tumor suppressors and kinases as potential driver genes of ovarian cancer. , 2013, , .		0
43	The tumor suppressor CDC73 interacts with the ring finger proteins RNF20 and RNF40 and is required for the maintenance of histone 2B monoubiquitination. <i>Human Molecular Genetics</i> , 2012, 21, 559-568.	2.9	85
44	Hypercalcaemia due to parathyroid carcinoma presenting in the third trimester of pregnancy. <i>Australian and New Zealand Journal of Obstetrics and Gynaecology</i> , 2012, 52, 204-207.	1.0	13
45	Insulin-like growth factor binding protein-3 inhibits migration of endometrial cancer cells. <i>Cancer Letters</i> , 2012, 317, 41-48.	7.2	11
46	Elevated levels of circulating microRNA-200 family members correlate with serous epithelial ovarian cancer. <i>BMC Cancer</i> , 2012, 12, 627.	2.6	163
47	Gonadotropin signalling in epithelial ovarian cancer. <i>Cancer Letters</i> , 2012, 324, 152-159.	7.2	50
48	Abstract 2167: The tumor suppressor CDC73/parafibromin is required for the maintenance of histone 2B monoubiquitination both in vitro and in vivo. , 2012, , .		0
49	Abstract 3150: miR-100 in ovarian cancer cell lines. , 2012, , .		0
50	Multiple Endocrine Neoplasia: Types 1 and 2. <i>Advances in Oto-Rhino-Laryngology</i> , 2011, 70, 84-90.	1.6	14
51	Mutant AKT1 in Proteus Syndrome. <i>New England Journal of Medicine</i> , 2011, 365, 2141-2142.	27.0	10
52	Metastatic parathyroid carcinoma initially misdiagnosed as parathyroid adenoma: the role of parafibromin in increasing diagnostic accuracy. <i>Internal Medicine Journal</i> , 2011, 41, 695-699.	0.8	6
53	Involvement of Insulin-like Growth Factor-binding Protein-3 in the Effects of Histone Deacetylase Inhibitor MS-275 in Hepatoma Cells. <i>Journal of Biological Chemistry</i> , 2011, 286, 29540-29547.	3.4	18
54	Abstract 4962: Assessing serum miRNA as putative biomarkers for serous epithelial ovarian cancer. , 2011, , .		0

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55	CDC73/HRPT2 CpG island hypermethylation and mutation of 5' untranslated sequence are uncommon mechanisms of silencing parafibromin in parathyroid tumors. <i>Endocrine-Related Cancer</i> , 2010, 17, 273-282.	3.1	37
56	The chemokine CXCL1 induces proliferation in epithelial ovarian cancer cells by transactivation of the epidermal growth factor receptor. <i>Endocrine-Related Cancer</i> , 2010, 17, 929-940.	3.1	98
57	Gonadotropin-induced ovarian cancer cell migration and proliferation require extracellular signal-regulated kinase 1/2 activation regulated by calcium and protein kinase C β . <i>Endocrine-Related Cancer</i> , 2010, 17, 335-349.	3.1	40
58	The Use of Denaturing High Performance Liquid Chromatography (DHPLC) for Mutation Scanning of Hereditary Cancer Genes. <i>Methods in Molecular Biology</i> , 2010, 653, 133-145.	0.9	10
59	Accuracy of Combined Protein Gene Product 9.5 and Parafibromin Markers for Immunohistochemical Diagnosis of Parathyroid Carcinoma. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2009, 94, 434-441.	3.6	120
60	The effect of disease-associated HRPT2 mutations on splicing. <i>Journal of Endocrinology</i> , 2009, 201, 387-396.	2.6	10
61	Denaturing High Performance Liquid Chromatography Detection of SDHB, SDHD, and VHL Germline Mutations in Pheochromocytoma. <i>Journal of Surgical Research</i> , 2009, 157, 55-62.	1.6	20
62	Rapamycin treatment for a child with germline PTEN mutation. <i>Nature Clinical Practice Oncology</i> , 2008, 5, 357-361.	4.3	114
63	Cowden Syndrome. , 2008, , 759-762.		0
64	Nucleolar localization of parafibromin is mediated by three nucleolar localization signals. <i>FEBS Letters</i> , 2007, 581, 5070-5074.	2.8	44
65	Molecular diagnosis of primary hyperparathyroidism in familial cancer syndromes. <i>Expert Opinion on Medical Diagnostics</i> , 2007, 1, 377-392.	1.6	21
66	Rapid Mutation Screening for HRPT2 and MEN1 Mutations Associated with Familial and Sporadic Primary Hyperparathyroidism. <i>Journal of Molecular Diagnostics</i> , 2006, 8, 559-566.	2.8	16
67	Loss of Nuclear Expression of Parafibromin Distinguishes Parathyroid Carcinomas and Hyperparathyroidism-Jaw Tumor (HPT-JT) Syndrome-related Adenomas From Sporadic Parathyroid Adenomas and Hyperplasias. <i>American Journal of Surgical Pathology</i> , 2006, 30, 1140-1149.	3.7	213
68	Genetic Testing in Pheochromocytoma- and Paraganglioma-Associated Syndromes. <i>Annals of the New York Academy of Sciences</i> , 2006, 1073, 104-111.	3.8	28
69	Clinical Presentation and Penetrance of Pheochromocytoma/Paraganglioma Syndromes. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2006, 91, 827-836.	3.6	560
70	Protein Chip Discovery of Secreted Proteins Regulated by the Phosphatidylinositol 3-Kinase Pathway in Ovarian Cancer Cell Lines. <i>Cancer Research</i> , 2006, 66, 1376-1383.	0.9	24
71	Identification of a functional bipartite nuclear localization signal in the tumor suppressor parafibromin. <i>Oncogene</i> , 2005, 24, 6241-6248.	5.9	65
72	A HIF1 α Regulatory Loop Links Hypoxia and Mitochondrial Signals in Pheochromocytomas. <i>PLoS Genetics</i> , 2005, 1, e8.	3.5	394

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73	Gene Expression of Parathyroid Tumors. <i>Cancer Research</i> , 2004, 64, 7405-7411.	0.9	96
74	Diagnosis of Proteus syndrome was correct. <i>American Journal of Medical Genetics Part A</i> , 2004, 130A, 214-215.	2.4	8
75	A molecular diagnosis of hyperparathyroidismâ€”jaw tumor syndrome in an adolescent with recurrent kidney stones. <i>Journal of Pediatrics</i> , 2004, 145, 567.	1.8	16
76	HRPT2 and parathyroid cancer. <i>Lancet Oncology</i> , The, 2004, 5, 78.	10.7	0
77	K40E: a novel succinate dehydrogenase (SDH)B mutation causing familial pheochromocytoma and paraganglioma. <i>Clinical Endocrinology</i> , 2004, 61, 510-514.	2.4	21
78	Von Hippelâ€”Lindau Disease. , 2004, , 1329-1333.		0
79	Cowden Syndrome. , 2004, , 301-304.		0
80	Novel succinate dehydrogenase subunit B (SDHB) mutations in familial pheochromocytomas and paragangliomas, but an absence of somatic SDHB mutations in sporadic pheochromocytomas. <i>Oncogene</i> , 2003, 22, 1358-1364.	5.9	108
81	Germline Inactivation of PTEN and Dysregulation of the Phosphoinositol-3-Kinase/Akt Pathway Cause Human Lhermitte-Duclos Disease in Adults. <i>American Journal of Human Genetics</i> , 2003, 73, 1191-1198.	6.2	213
82	Genome-Wide Copy Number Imbalances Identified in Familial and Sporadic Medullary Thyroid Carcinoma. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2003, 88, 1866-1872.	3.6	54
83	A Case Report in Favor of a Multistep Adrenocortical Tumorigenesis. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2003, 88, 998-1001.	3.6	54
84	Comparative Genomic Hybridization Analysis of Adrenocortical Tumors. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2002, 87, 3467-3474.	3.6	125
85	Genetics of pheochromocytoma and paraganglioma. <i>Current Opinion in Endocrinology, Diabetes and Obesity</i> , 2002, 9, 79-86.	0.6	9
86	Independent Genetic Events Associated with the Development of Multiple Parathyroid Tumors in Patients with Primary Hyperparathyroidism. <i>American Journal of Pathology</i> , 2002, 161, 1299-1306.	3.8	36
87	Transcriptional repression of the RET proto-oncogene by a mitogen activated protein kinase-dependent signalling pathway. <i>Gene</i> , 2002, 298, 9-19.	2.2	17
88	Genetic insights into familial cancers â€” update and recent discoveries. <i>Cancer Letters</i> , 2002, 181, 125-164.	7.2	75
89	Rapid Mutation Scanning of Genes Associated with Familial Cancer Syndromes Using Denaturing High-Performance Liquid Chromatography. <i>Neoplasia</i> , 2001, 3, 236-244.	5.3	31
90	Mutational Analysis and Genotype-Phenotype Correlation of the PHEX Gene in X-Linked Hypophosphatemic Rickets. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2001, 86, 3889-3899.	3.6	27