

Diether Lambrechts

List of Publications by Citations

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

477
papers

36,016
citations

90
h-index

178
g-index

596
ext. papers

43,702
ext. citations

10.6
avg, IF

6.43
L-index

#	Paper	IF	Citations
477	Effects of KRAS, BRAF, NRAS, and PIK3CA mutations on the efficacy of cetuximab plus chemotherapy in chemotherapy-refractory metastatic colorectal cancer: a retrospective consortium analysis. <i>Lancet Oncology, The</i> , 2010 , 11, 753-62	21.7	1653
476	Plasma HDL cholesterol and risk of myocardial infarction: a mendelian randomisation study. <i>Lancet, The</i> , 2012 , 380, 572-80	40	1523
475	Large-scale association analysis identifies 13 new susceptibility loci for coronary artery disease. <i>Nature Genetics</i> , 2011 , 43, 333-8	36.3	1394
474	Large-scale genotyping identifies 41 new loci associated with breast cancer risk. <i>Nature Genetics</i> , 2013 , 45, 353-61, 361e1-2	36.3	813
473	Addition of cetuximab to oxaliplatin-based first-line combination chemotherapy for treatment of advanced colorectal cancer: results of the randomised phase 3 MRC COIN trial. <i>Lancet, The</i> , 2011 , 377, 2103-14	40	762
472	VEGF is a modifier of amyotrophic lateral sclerosis in mice and humans and protects motoneurons against ischemic death. <i>Nature Genetics</i> , 2003 , 34, 383-94	36.3	712
471	Gene prioritization through genomic data fusion. <i>Nature Biotechnology</i> , 2006 , 24, 537-44	44.5	685
470	Association analysis identifies 65 new breast cancer risk loci. <i>Nature</i> , 2017 , 551, 92-94	50.4	643
469	Role of PlGF in the intra- and intermolecular cross talk between the VEGF receptors Flt1 and Flk1. <i>Nature Medicine</i> , 2003 , 9, 936-43	50.5	631
468	Phenotype molding of stromal cells in the lung tumor microenvironment. <i>Nature Medicine</i> , 2018 , 24, 1277-1289	50.5	607
467	Associations of breast cancer risk factors with tumor subtypes: a pooled analysis from the Breast Cancer Association Consortium studies. <i>Journal of the National Cancer Institute</i> , 2011 , 103, 250-63	9.7	513
466	Treatment of motoneuron degeneration by intracerebroventricular delivery of VEGF in a rat model of ALS. <i>Nature Neuroscience</i> , 2005 , 8, 85-92	25.5	429
465	Association between BRCA1 and BRCA2 mutations and survival in women with invasive epithelial ovarian cancer. <i>JAMA - Journal of the American Medical Association</i> , 2012 , 307, 382-90	27.4	427
464	VEGF: once regarded as a specific angiogenic factor, now implicated in neuroprotection. <i>BioEssays</i> , 2004 , 26, 943-54	4.1	423
463	Multiple independent variants at the TERT locus are associated with telomere length and risks of breast and ovarian cancer. <i>Nature Genetics</i> , 2013 , 45, 371-84, 384e1-2	36.3	422
462	Thrombomodulin mutations in atypical hemolytic-uremic syndrome. <i>New England Journal of Medicine</i> , 2009 , 361, 345-57	59.2	418
461	Genome-wide association analysis of more than 120,000 individuals identifies 15 new susceptibility loci for breast cancer. <i>Nature Genetics</i> , 2015 , 47, 373-80	36.3	406

460	Parent-of-origin-specific allelic associations among 106 genomic loci for age at menarche. <i>Nature</i> , 2014 , 514, 92-97	50.4	401
459	Deficiency or inhibition of oxygen sensor Phd1 induces hypoxia tolerance by reprogramming basal metabolism. <i>Nature Genetics</i> , 2008 , 40, 170-80	36.3	383
458	Polygenic Risk Scores for Prediction of Breast Cancer and Breast Cancer Subtypes. <i>American Journal of Human Genetics</i> , 2019 , 104, 21-34	11	363
457	Tumour hypoxia causes DNA hypermethylation by reducing TET activity. <i>Nature</i> , 2016 , 537, 63-68	50.4	354
456	Gain of function of mutant p53 by coaggregation with multiple tumor suppressors. <i>Nature Chemical Biology</i> , 2011 , 7, 285-95	11.7	345
455	Genome-wide association studies identify four ER negative-specific breast cancer risk loci. <i>Nature Genetics</i> , 2013 , 45, 392-8, 398e1-2	36.3	327
454	Inhibition of the Glycolytic Activator PFKFB3 in Endothelium Induces Tumor Vessel Normalization, Impairs Metastasis, and Improves Chemotherapy. <i>Cancer Cell</i> , 2016 , 30, 968-985	24.3	325
453	Prediction of breast cancer risk based on profiling with common genetic variants. <i>Journal of the National Cancer Institute</i> , 2015 , 107,	9.7	324
452	Progranulin functions as a neurotrophic factor to regulate neurite outgrowth and enhance neuronal survival. <i>Journal of Cell Biology</i> , 2008 , 181, 37-41	7.3	322
451	Role and therapeutic potential of VEGF in the nervous system. <i>Physiological Reviews</i> , 2009 , 89, 607-48	47.9	319
450	Vitamin D deficiency is highly prevalent in COPD and correlates with variants in the vitamin D-binding gene. <i>Thorax</i> , 2010 , 65, 215-20	7.3	308
449	Underestimated and under-recognized: the late consequences of acute coronary syndrome (GRACE UK-Belgian Study). <i>European Heart Journal</i> , 2010 , 31, 2755-64	9.5	306
448	A genome-wide association study identifies susceptibility loci for ovarian cancer at 2q31 and 8q24. <i>Nature Genetics</i> , 2010 , 42, 874-9	36.3	277
447	Neurovascular signalling defects in neurodegeneration. <i>Nature Reviews Neuroscience</i> , 2008 , 9, 169-81	13.5	277
446	A locus on 19p13 modifies risk of breast cancer in BRCA1 mutation carriers and is associated with hormone receptor-negative breast cancer in the general population. <i>Nature Genetics</i> , 2010 , 42, 885-92	36.3	276
445	PIK3CA mutations are not a major determinant of resistance to the epidermal growth factor receptor inhibitor cetuximab in metastatic colorectal cancer. <i>Clinical Cancer Research</i> , 2009 , 15, 3184-8	12.9	276
444	Genome dynamics of the human embryonic kidney 293 lineage in response to cell biology manipulations. <i>Nature Communications</i> , 2014 , 5, 4767	17.4	275
443	Markers of response for the antiangiogenic agent bevacizumab. <i>Journal of Clinical Oncology</i> , 2013 , 31, 1219-30	2.2	272

442	p53 induces formation of NEAT1 lncRNA-containing paraspeckles that modulate replication stress response and chemosensitivity. <i>Nature Medicine</i> , 2016 , 22, 861-8	50.5	271
441	GWAS meta-analysis and replication identifies three new susceptibility loci for ovarian cancer. <i>Nature Genetics</i> , 2013 , 45, 362-70, 370e1-2	36.3	267
440	Genomic analyses identify hundreds of variants associated with age at menarche and support a role for puberty timing in cancer risk. <i>Nature Genetics</i> , 2017 , 49, 834-841	36.3	257
439	VEGF: a modifier of the del22q11 (DiGeorge) syndrome?. <i>Nature Medicine</i> , 2003 , 9, 173-82	50.5	256
438	A common variant at the TERT-CLPTM1L locus is associated with estrogen receptor-negative breast cancer. <i>Nature Genetics</i> , 2011 , 43, 1210-4	36.3	253
437	Genome-wide association analysis identifies three new breast cancer susceptibility loci. <i>Nature Genetics</i> , 2012 , 44, 312-8	36.3	237
436	Large-scale genomic analyses link reproductive aging to hypothalamic signaling, breast cancer susceptibility and BRCA1-mediated DNA repair. <i>Nature Genetics</i> , 2015 , 47, 1294-1303	36.3	226
435	Common variants at 19p13 are associated with susceptibility to ovarian cancer. <i>Nature Genetics</i> , 2010 , 42, 880-4	36.3	210
434	Self-Maintaining Gut Macrophages Are Essential for Intestinal Homeostasis. <i>Cell</i> , 2018 , 175, 400-415.e13	56.2	201
433	A genetic <i>Xenopus laevis</i> tadpole model to study lymphangiogenesis. <i>Nature Medicine</i> , 2005 , 11, 998-1004	54.5	191
432	Identification of 12 new susceptibility loci for different histotypes of epithelial ovarian cancer. <i>Nature Genetics</i> , 2017 , 49, 680-691	36.3	190
431	Identification of ten variants associated with risk of estrogen-receptor-negative breast cancer. <i>Nature Genetics</i> , 2017 , 49, 1767-1778	36.3	186
430	Centrosome Amplification Is Sufficient to Promote Spontaneous Tumorigenesis in Mammals. <i>Developmental Cell</i> , 2017 , 40, 313-322.e5	10.2	181
429	Plasma circulating tumor DNA as an alternative to metastatic biopsies for mutational analysis in breast cancer. <i>Annals of Oncology</i> , 2014 , 25, 1959-1965	10.3	181
428	Identification of six new susceptibility loci for invasive epithelial ovarian cancer. <i>Nature Genetics</i> , 2015 , 47, 164-71	36.3	177
427	Genomic Characterization of Primary Invasive Lobular Breast Cancer. <i>Journal of Clinical Oncology</i> , 2016 , 34, 1872-81	2.2	175
426	Large-scale gene-centric analysis identifies novel variants for coronary artery disease. <i>PLoS Genetics</i> , 2011 , 7, e1002260	6	175
425	Prognostic significance of POLE proofreading mutations in endometrial cancer. <i>Journal of the National Cancer Institute</i> , 2015 , 107, 402	9.7	169

424	The role of fatty acid oxidation in lymphangiogenesis. <i>Nature</i> , 2017 , 542, 49-54	50.4	167
423	Functional variants at the 11q13 risk locus for breast cancer regulate cyclin D1 expression through long-range enhancers. <i>American Journal of Human Genetics</i> , 2013 , 92, 489-503	11	167
422	Optimized filtering reduces the error rate in detecting genomic variants by short-read sequencing. <i>Nature Biotechnology</i> , 2011 , 30, 61-8	44.5	163
421	25(OH)D2 half-life is shorter than 25(OH)D3 half-life and is influenced by DBP concentration and genotype. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2014 , 99, 3373-81	5.6	154
420	VEGF pathway genetic variants as biomarkers of treatment outcome with bevacizumab: an analysis of data from the AVITA and AVOREN randomised trials. <i>Lancet Oncology</i> , 2012 , 13, 724-33	21.7	154
419	Patient-derived organoids from endometrial disease capture clinical heterogeneity and are amenable to drug screening. <i>Nature Cell Biology</i> , 2019 , 21, 1041-1051	23.4	146
418	Low penetrance breast cancer susceptibility loci are associated with specific breast tumor subtypes: findings from the Breast Cancer Association Consortium. <i>Human Molecular Genetics</i> , 2011 , 20, 3289-303	5.6	140
417	CPS1 maintains pyrimidine pools and DNA synthesis in KRAS/LKB1-mutant lung cancer cells. <i>Nature</i> , 2017 , 546, 168-172	50.4	136
416	The BRCA1- Δ 11q Alternative Splice Isoform Bypasses Germline Mutations and Promotes Therapeutic Resistance to PARP Inhibition and Cisplatin. <i>Cancer Research</i> , 2016 , 76, 2778-90	10.1	136
415	CHEK2*1100delC heterozygosity in women with breast cancer associated with early death, breast cancer-specific death, and increased risk of a second breast cancer. <i>Journal of Clinical Oncology</i> , 2012 , 30, 4308-16	2.2	134
414	Lessons from the adjuvant bevacizumab trial on colon cancer: what next?. <i>Journal of Clinical Oncology</i> , 2011 , 29, 1-4	2.2	133
413	CYP2D6 genotype and adjuvant tamoxifen: meta-analysis of heterogeneous study populations. <i>Clinical Pharmacology and Therapeutics</i> , 2014 , 95, 216-27	6.1	131
412	Semiautomated isolation and molecular characterisation of single or highly purified tumour cells from CellSearch enriched blood samples using dielectrophoretic cell sorting. <i>British Journal of Cancer</i> , 2013 , 108, 1358-67	8.7	131
411	Genomic landscape of carcinogen-induced and genetically induced mouse skin squamous cell carcinoma. <i>Nature Medicine</i> , 2015 , 21, 946-54	50.5	127
410	Epigenetic analysis leads to identification of HNF1B as a subtype-specific susceptibility gene for ovarian cancer. <i>Nature Communications</i> , 2013 , 4, 1628	17.4	124
409	Genome-wide association study identifies a common variant associated with risk of endometrial cancer. <i>Nature Genetics</i> , 2011 , 43, 451-4	36.3	121
408	PALB2, CHEK2 and ATM rare variants and cancer risk: data from COGS. <i>Journal of Medical Genetics</i> , 2016 , 53, 800-811	5.8	121
407	Sequencing an Ashkenazi reference panel supports population-targeted personal genomics and illuminates Jewish and European origins. <i>Nature Communications</i> , 2014 , 5, 4835	17.4	115

406	Relief of hypoxia by angiogenesis promotes neural stem cell differentiation by targeting glycolysis. <i>EMBO Journal</i> , 2016 , 35, 924-41	13	114
405	Evidence of gene-environment interactions between common breast cancer susceptibility loci and established environmental risk factors. <i>PLoS Genetics</i> , 2013 , 9, e1003284	6	112
404	Somatic mutation profiling and associations with prognosis and trastuzumab benefit in early breast cancer. <i>Journal of the National Cancer Institute</i> , 2013 , 105, 960-7	9.7	112
403	Inhibition of tumor angiogenesis and growth by a small-molecule multi-FGF receptor blocker with allosteric properties. <i>Cancer Cell</i> , 2013 , 23, 477-88	24.3	110
402	A pan-cancer blueprint of the heterogeneous tumor microenvironment revealed by single-cell profiling. <i>Cell Research</i> , 2020 , 30, 745-762	24.7	108
401	Germline mutation in BRCA1 or BRCA2 and ten-year survival for women diagnosed with epithelial ovarian cancer. <i>Clinical Cancer Research</i> , 2015 , 21, 652-7	12.9	107
400	Genome-Wide Meta-Analyses of Breast, Ovarian, and Prostate Cancer Association Studies Identify Multiple New Susceptibility Loci Shared by at Least Two Cancer Types. <i>Cancer Discovery</i> , 2016 , 6, 1052-67	24.4	104
399	Transient PLK4 overexpression accelerates tumorigenesis in p53-deficient epidermis. <i>Nature Cell Biology</i> , 2016 , 18, 100-10	23.4	104
398	Novel role for vascular endothelial growth factor (VEGF) receptor-1 and its ligand VEGF-B in motor neuron degeneration. <i>Journal of Neuroscience</i> , 2008 , 28, 10451-9	6.6	104
397	The P450 oxidoreductase *28 SNP is associated with low initial tacrolimus exposure and increased dose requirements in CYP3A5-expressing renal recipients. <i>Pharmacogenomics</i> , 2011 , 12, 1281-91	2.6	102
396	Discriminating mild from critical COVID-19 by innate and adaptive immune single-cell profiling of bronchoalveolar lavages. <i>Cell Research</i> , 2021 , 31, 272-290	24.7	102
395	A transcriptome-wide association study of 229,000 women identifies new candidate susceptibility genes for breast cancer. <i>Nature Genetics</i> , 2018 , 50, 968-978	36.3	101
394	Common breast cancer susceptibility loci are associated with triple-negative breast cancer. <i>Cancer Research</i> , 2011 , 71, 6240-9	10.1	100
393	Quiescent Endothelial Cells Upregulate Fatty Acid Oxidation for Vasculoprotection via Redox Homeostasis. <i>Cell Metabolism</i> , 2018 , 28, 881-894.e13	24.6	99
392	Lineage-dependent gene expression programs influence the immune landscape of colorectal cancer. <i>Nature Genetics</i> , 2020 , 52, 594-603	36.3	96
391	An Integrated Gene Expression Landscape Profiling Approach to Identify Lung Tumor Endothelial Cell Heterogeneity and Angiogenic Candidates. <i>Cancer Cell</i> , 2020 , 37, 21-36.e13	24.3	93
390	Breast cancer risk variants at 6q25 display different phenotype associations and regulate ESR1, RMND1 and CCDC170. <i>Nature Genetics</i> , 2016 , 48, 374-86	36.3	93
389	19p13.1 is a triple-negative-specific breast cancer susceptibility locus. <i>Cancer Research</i> , 2012 , 72, 1795-803	30.1	93

388	HIF-1 β Promotes Glutamine-Mediated Redox Homeostasis and Glycogen-Dependent Bioenergetics to Support Postimplantation Bone Cell Survival. <i>Cell Metabolism</i> , 2016 , 23, 265-79	24.6	92
387	Identification of a BRCA2-specific modifier locus at 6p24 related to breast cancer risk. <i>PLoS Genetics</i> , 2013 , 9, e1003173	6	90
386	Evidence that breast cancer risk at the 2q35 locus is mediated through IGFBP5 regulation. <i>Nature Communications</i> , 2014 , 4, 4999	17.4	87
385	Identification and molecular characterization of a new ovarian cancer susceptibility locus at 17q21.31. <i>Nature Communications</i> , 2013 , 4, 1627	17.4	85
384	Genetic predisposition for beta cell fragility underlies type 1 and type 2 diabetes. <i>Nature Genetics</i> , 2016 , 48, 519-27	36.3	83
383	No evidence that protein truncating variants in BRIP1 are associated with breast cancer risk: implications for gene panel testing. <i>Journal of Medical Genetics</i> , 2016 , 53, 298-309	5.8	83
382	Obesity and survival among women with ovarian cancer: results from the Ovarian Cancer Association Consortium. <i>British Journal of Cancer</i> , 2015 , 113, 817-26	8.7	80
381	Fine-scale mapping of the FGFR2 breast cancer risk locus: putative functional variants differentially bind FOXA1 and E2F1. <i>American Journal of Human Genetics</i> , 2013 , 93, 1046-60	11	80
380	Vitamin D status at breast cancer diagnosis: correlation with tumor characteristics, disease outcome, and genetic determinants of vitamin D insufficiency. <i>Carcinogenesis</i> , 2012 , 33, 1319-26	4.6	80
379	VEGF at the neurovascular interface: therapeutic implications for motor neuron disease. <i>Biochimica Et Biophysica Acta - Molecular Basis of Disease</i> , 2006 , 1762, 1109-21	6.9	80
378	Phylogenetic analysis of metastatic progression in breast cancer using somatic mutations and copy number aberrations. <i>Nature Communications</i> , 2017 , 8, 14944	17.4	79
377	Expression profiling of budding cells in colorectal cancer reveals an EMT-like phenotype and molecular subtype switching. <i>British Journal of Cancer</i> , 2017 , 116, 58-65	8.7	79
376	ABCA transporter gene expression and poor outcome in epithelial ovarian cancer. <i>Journal of the National Cancer Institute</i> , 2014 , 106,	9.7	79
375	LBA81 Keeping exhausted T-cells in check in COVID-19. <i>Annals of Oncology</i> , 2020 , 31, S1208	10.3	78
374	(Epi)genetic variation in ageing of metabolic fitness. <i>Archives of Public Health</i> , 2015 , 73,	2.6	78
373	The 15q24/25 susceptibility variant for lung cancer and chronic obstructive pulmonary disease is associated with emphysema. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2010 , 181, 486-93 ^{10.2}		77
372	Association of vitamin D levels and risk of ovarian cancer: a Mendelian randomization study. <i>International Journal of Epidemiology</i> , 2016 , 45, 1619-1630	7.8	77
371	Genome-wide association study identifies 32 novel breast cancer susceptibility loci from overall and subtype-specific analyses. <i>Nature Genetics</i> , 2020 , 52, 572-581	36.3	76

370	The role of genetic breast cancer susceptibility variants as prognostic factors. <i>Human Molecular Genetics</i> , 2012 , 21, 3926-39	5.6	75
369	Height and Breast Cancer Risk: Evidence From Prospective Studies and Mendelian Randomization. <i>Journal of the National Cancer Institute</i> , 2015 , 107,	9.7	74
368	Assessing interactions between the associations of common genetic susceptibility variants, reproductive history and body mass index with breast cancer risk in the breast cancer association consortium: a combined case-control study. <i>Breast Cancer Research</i> , 2010 , 12, R110	8.3	74
367	Role of Gas6 in erythropoiesis and anemia in mice. <i>Journal of Clinical Investigation</i> , 2008 , 118, 583-96	15.9	74
366	Single-nucleotide polymorphisms associated with outcome in metastatic renal cell carcinoma treated with sunitinib. <i>British Journal of Cancer</i> , 2013 , 108, 887-900	8.7	73
365	Somatic profiling of the epidermal growth factor receptor pathway in tumors from patients with advanced colorectal cancer treated with chemotherapy ± cetuximab. <i>Clinical Cancer Research</i> , 2013 , 19, 4104-13	12.9	72
364	Endovascular treatment of the descending thoracic aorta. <i>European Journal of Vascular and Endovascular Surgery</i> , 2003 , 26, 437-44	2.3	71
363	Identification of nine new susceptibility loci for endometrial cancer. <i>Nature Communications</i> , 2018 , 9, 3166	17.4	70
362	Matrix-binding vascular endothelial growth factor (VEGF) isoforms guide granule cell migration in the cerebellum via VEGF receptor Flk1. <i>Journal of Neuroscience</i> , 2010 , 30, 15052-66	6.6	68
361	Associations of common variants at 1p11.2 and 14q24.1 (RAD51L1) with breast cancer risk and heterogeneity by tumor subtype: findings from the Breast Cancer Association Consortium. <i>Human Molecular Genetics</i> , 2011 , 20, 4693-706	5.6	66
360	BRCA2 Polymorphic Stop Codon K3326X and the Risk of Breast, Prostate, and Ovarian Cancers. <i>Journal of the National Cancer Institute</i> , 2016 , 108,	9.7	65
359	The association of the 4q25 susceptibility variant for atrial fibrillation with stroke is limited to stroke of cardioembolic etiology. <i>Stroke</i> , 2010 , 41, 1850-7	6.7	65
358	Identification of four novel susceptibility loci for oestrogen receptor negative breast cancer. <i>Nature Communications</i> , 2016 , 7, 11375	17.4	64
357	Lineage-specific functions of TET1 in the postimplantation mouse embryo. <i>Nature Genetics</i> , 2017 , 49, 1061-1072	36.3	63
356	Meta-analysis of vascular endothelial growth factor variations in amyotrophic lateral sclerosis: increased susceptibility in male carriers of the -2578AA genotype. <i>Journal of Medical Genetics</i> , 2009 , 46, 840-6	5.8	63
355	Genome-wide CRISPR screening identifies TMEM106B as a proviral host factor for SARS-CoV-2. <i>Nature Genetics</i> , 2021 , 53, 435-444	36.3	62
354	Uncovering the genomic heterogeneity of multifocal breast cancer. <i>Journal of Pathology</i> , 2015 , 236, 457-66	9.6	61
353	Genome-wide significant risk associations for mucinous ovarian carcinoma. <i>Nature Genetics</i> , 2015 , 47, 888-97	36.3	60

352	High-throughput interrogation of PIK3CA, PTEN, KRAS, FBXW7 and TP53 mutations in primary endometrial carcinoma. <i>Gynecologic Oncology</i> , 2013 , 128, 327-34	4.9	60
351	Fine-scale mapping of the 5q11.2 breast cancer locus reveals at least three independent risk variants regulating MAP3K1. <i>American Journal of Human Genetics</i> , 2015 , 96, 5-20	11	59
350	Single-cell profiling of myeloid cells in glioblastoma across species and disease stage reveals macrophage competition and specialization. <i>Nature Neuroscience</i> , 2021 , 24, 595-610	25.5	59
349	A single-cell map of intratumoral changes during anti-PD1 treatment of patients with breast cancer. <i>Nature Medicine</i> , 2021 , 27, 820-832	50.5	57
348	Fine-mapping of 150 breast cancer risk regions identifies 191 likely target genes. <i>Nature Genetics</i> , 2020 , 52, 56-73	36.3	56
347	Five endometrial cancer risk loci identified through genome-wide association analysis. <i>Nature Genetics</i> , 2016 , 48, 667-674	36.3	56
346	Platinum sensitivity-related germline polymorphism discovered via a cell-based approach and analysis of its association with outcome in ovarian cancer patients. <i>Clinical Cancer Research</i> , 2011 , 17, 5490-500	12.9	55
345	Microglial upregulation of progranulin as a marker of motor neuron degeneration. <i>Journal of Neuropathology and Experimental Neurology</i> , 2010 , 69, 1191-200	3.1	55
344	The Cancer Cell Oxygen Sensor PHD2 Promotes Metastasis via Activation of Cancer-Associated Fibroblasts. <i>Cell Reports</i> , 2015 , 12, 992-1005	10.6	54
343	Mismatch repair deficiency endows tumors with a unique mutation signature and sensitivity to DNA double-strand breaks. <i>ELife</i> , 2014 , 3, e02725	8.9	54
342	Lipid availability determines fate of skeletal progenitor cells via SOX9. <i>Nature</i> , 2020 , 579, 111-117	50.4	53
341	Joint associations of a polygenic risk score and environmental risk factors for breast cancer in the Breast Cancer Association Consortium. <i>International Journal of Epidemiology</i> , 2018 , 47, 526-536	7.8	53
340	Functional mechanisms underlying pleiotropic risk alleles at the 19p13.1 breast-ovarian cancer susceptibility locus. <i>Nature Communications</i> , 2016 , 7, 12675	17.4	53
339	Low expression VEGF haplotype increases the risk for tetralogy of Fallot: a family based association study. <i>Journal of Medical Genetics</i> , 2005 , 42, 519-22	5.8	53
338	Monocyte-driven atypical cytokine storm and aberrant neutrophil activation as key mediators of COVID-19 disease severity. <i>Nature Communications</i> , 2021 , 12, 4117	17.4	53
337	Implementing liquid biopsies into clinical decision making for cancer immunotherapy. <i>Oncotarget</i> , 2017 , 8, 48507-48520	3.3	52
336	Chromosomal Instability in Cell-Free DNA as a Highly Specific Biomarker for Detection of Ovarian Cancer in Women with Adnexal Masses. <i>Clinical Cancer Research</i> , 2017 , 23, 2223-2231	12.9	52
335	Associations of obesity and circulating insulin and glucose with breast cancer risk: a Mendelian randomization analysis. <i>International Journal of Epidemiology</i> , 2019 , 48, 795-806	7.8	52

334	Genetic modifiers of CHEK2*1100delC-associated breast cancer risk. <i>Genetics in Medicine</i> , 2017 , 19, 599-603	5.1	51
333	Hypomorphic Missense Variants Confer Moderate Risks of Breast Cancer. <i>Cancer Research</i> , 2017 , 77, 2789-2799	10.1	49
332	Vascular endothelial growth factor counteracts the loss of phospho-Akt preceding motor neurone degeneration in amyotrophic lateral sclerosis. <i>Neuropathology and Applied Neurobiology</i> , 2007 , 33, 499-509	5.3	49
331	Comparison of 6q25 breast cancer hits from Asian and European Genome Wide Association Studies in the Breast Cancer Association Consortium (BCAC). <i>PLoS ONE</i> , 2012 , 7, e42380	3.7	49
330	Shared genetics underlying epidemiological association between endometriosis and ovarian cancer. <i>Human Molecular Genetics</i> , 2015 , 24, 5955-64	5.6	48
329	Biology of breast cancer during pregnancy using genomic profiling. <i>Endocrine-Related Cancer</i> , 2014 , 21, 545-54	5.7	48
328	Common non-synonymous SNPs associated with breast cancer susceptibility: findings from the Breast Cancer Association Consortium. <i>Human Molecular Genetics</i> , 2014 , 23, 6096-111	5.6	48
327	Genome-wide association and transcriptome studies identify target genes and risk loci for breast cancer. <i>Nature Communications</i> , 2019 , 10, 1741	17.4	47
326	Genetic variability in the multidrug resistance associated protein-1 (ABCC1/MRP1) predicts hematological toxicity in breast cancer patients receiving (neo-)adjuvant chemotherapy with 5-fluorouracil, epirubicin and cyclophosphamide (FEC). <i>Annals of Oncology</i> , 2013 , 24, 1513-25	10.3	47
325	Fine-mapping of the HNF1B multicancer locus identifies candidate variants that mediate endometrial cancer risk. <i>Human Molecular Genetics</i> , 2015 , 24, 1478-92	5.6	46
324	A variant at chromosome 9p21 is associated with recurrent myocardial infarction and cardiac death after acute coronary syndrome: the GRACE Genetics Study. <i>European Heart Journal</i> , 2010 , 31, 1132-41	9.5	46
323	The molecular genetic basis of ovarian cancer and its roadmap towards a better treatment. <i>Gynecologic Oncology</i> , 2010 , 117, 358-65	4.9	46
322	Shared heritability and functional enrichment across six solid cancers. <i>Nature Communications</i> , 2019 , 10, 431	17.4	45
321	The role of KRAS rs61764370 in invasive epithelial ovarian cancer: implications for clinical testing. <i>Clinical Cancer Research</i> , 2011 , 17, 3742-50	12.9	45
320	Adult body mass index and risk of ovarian cancer by subtype: a Mendelian randomization study. <i>International Journal of Epidemiology</i> , 2016 , 45, 884-95	7.8	45
319	Developing Organoids from Ovarian Cancer as Experimental and Preclinical Models. <i>Stem Cell Reports</i> , 2020 , 14, 717-729	8	44
318	Tamoxifen Metabolism and Efficacy in Breast Cancer: A Prospective Multicenter Trial. <i>Clinical Cancer Research</i> , 2018 , 24, 2312-2318	12.9	44
317	Differences in MWCNT- and SWCNT-induced DNA methylation alterations in association with the nuclear deposition. <i>Particle and Fibre Toxicology</i> , 2018 , 15, 11	8.4	44

316	Conservation of copy number profiles during engraftment and passaging of patient-derived cancer xenografts. <i>Nature Genetics</i> , 2021 , 53, 86-99	36.3	44
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42	Fine-mapping of 150 breast cancer risk regions identifies 178 high confidence target genes		2
41	Genome-wide association study identifies 32 novel breast cancer susceptibility loci from overall and subtype-specific analyses		2
40	Conservation of copy number profiles during engraftment and passaging of patient-derived cancer xenografts		2
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25	Polygenic Risk Modelling for Prediction of Epithelial Ovarian Cancer Risk		1
24	Clinical practices underlie COVID-19 patient respiratory microbiome composition and its interactions with the host		1
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22	A variant in FTO gene shows association with histological ulceration in cutaneous melanoma. <i>Journal of Cutaneous Pathology</i> , 2020 , 47, 98-101	1.7	1
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