Saundra S Buys

List of Publications by Year in descending order

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101543 9,053 81 36 citations h-index papers

g-index 85 85 85 12152 docs citations times ranked citing authors all docs

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82

#	Article	IF	CITATIONS
1	Risks of Breast, Ovarian, and Contralateral Breast Cancer for <i>BRCA1</i> and <i>BRCA2</i> Mutation Carriers. JAMA - Journal of the American Medical Association, 2017, 317, 2402.	7.4	1,898
2	Effect of Screening on Ovarian Cancer Mortality. JAMA - Journal of the American Medical Association, 2011, 305, 2295.	7.4	1,080
3	Breast-Cancer Risk in Families with Mutations in <i>PALB2</i> . New England Journal of Medicine, 2014, 371, 497-506.	27.0	745
4	Genetic/Familial High-Risk Assessment: Breast, Ovarian, and Pancreatic, Version 2.2021, NCCN Clinical Practice Guidelines in Oncology. Journal of the National Comprehensive Cancer Network: JNCCN, 2021, 19, 77-102.	4.9	498
5	Association of Type and Location of <i>BRCA1</i> li>and <i>BRCA2</i> Mutations With Risk of Breast and Ovarian Cancer. JAMA - Journal of the American Medical Association, 2015, 313, 1347.	7.4	390
6	Identification of 12 new susceptibility loci for different histotypes of epithelial ovarian cancer. Nature Genetics, 2017 , 49 , 680 - 691 .	21.4	356
7	A study of over 35,000 women with breast cancer tested with a 25â€gene panel of hereditary cancer genes. Cancer, 2017, 123, 1721-1730.	4.1	303
8	Identification of ten variants associated with risk of estrogen-receptor-negative breast cancer. Nature Genetics, 2017, 49, 1767-1778.	21.4	289
9	Genome-wide association study identifies 32 novel breast cancer susceptibility loci from overall and subtype-specific analyses. Nature Genetics, 2020, 52, 572-581.	21.4	265
10	The Breast Cancer Family Registry: an infrastructure for cooperative multinational, interdisciplinary and translational studies of the genetic epidemiology of breast cancer. Breast Cancer Research, 2004, 6, R375-89.	5.0	255
11	Mutational spectrum in a worldwide study of 29,700 families with <i>BRCA1 </i>) or <i>BRCA2 </i>) i > mutations. Human Mutation, 2018, 39, 593-620.	2.5	224
12	Identification of six new susceptibility loci for invasive epithelial ovarian cancer. Nature Genetics, 2015, 47, 164-171.	21.4	221
13	Breast cancer risk variants at 6q25 display different phenotype associations and regulate ESR1, RMND1 and CCDC170. Nature Genetics, 2016, 48, 374-386.	21.4	125
14	Combating subclonal evolution of resistant cancer phenotypes. Nature Communications, 2017, 8, 1231.	12.8	124
15	Fine-mapping of 150 breast cancer risk regions identifies 191 likely target genes. Nature Genetics, 2020, 52, 56-73.	21.4	120
16	10-year performance of four models of breast cancer risk: a validation study. Lancet Oncology, The, 2019, 20, 504-517.	10.7	116
17	Extended mortality results for ovarian cancer screening in the PLCO trial with median 15 years follow-up. Gynecologic Oncology, 2016, 143, 270-275.	1.4	111
18	Expanding Access to BRCA1/2 Genetic Counseling with Telephone Delivery: A Cluster Randomized Trial. Journal of the National Cancer Institute, 2014, 106, dju328-dju328.	6.3	105

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19	Coffee, tea, caffeine intake, and the risk of cancer in the PLCO cohort. British Journal of Cancer, 2015, 113, 809-816.	6.4	99
20	Early Detection of Ovarian Cancer using the Risk of Ovarian Cancer Algorithm with Frequent CA125 Testing in Women at Increased Familial Risk – Combined Results from Two Screening Trials. Clinical Cancer Research, 2017, 23, 3628-3637.	7.0	99
21	Identification of four novel susceptibility loci for oestrogen receptor negative breast cancer. Nature Communications, 2016, 7, 11375.	12.8	93
22	The spectrum of apoptotic defects and clinical manifestations, including systemic lupus erythematosus, in humans with CD95 (Fas/APO-1) mutations. Arthritis and Rheumatism, 1999, 42, 1833-1842.	6.7	90
23	Cancer Risks Associated With <i>BRCA1</i> and <i>BRCA2</i> Pathogenic Variants. Journal of Clinical Oncology, 2022, 40, 1529-1541.	1.6	90
24	Male breast cancer in BRCA1 and BRCA2 mutation carriers: pathology data from the Consortium of Investigators of Modifiers of BRCA1/2. Breast Cancer Research, 2016, 18, 15.	5.0	88
25	Polygenic risk scores and breast and epithelial ovarian cancer risks for carriers of BRCA1 and BRCA2 pathogenic variants. Genetics in Medicine, 2020, 22, 1653-1666.	2.4	82
26	Functional mechanisms underlying pleiotropic risk alleles at the 19p13.1 breast–ovarian cancer susceptibility locus. Nature Communications, 2016, 7, 12675.	12.8	78
27	Breast cancer risk prediction using a polygenic risk score in the familial setting: a prospective study from the Breast Cancer Family Registry and kConFab. Genetics in Medicine, 2017, 19, 30-35.	2.4	53
28	Age-specific breast cancer risk by body mass index and familial risk: prospective family study cohort (ProF-SC). Breast Cancer Research, 2018, 20, 132.	5.0	51
29	Cohort Profile: The Breast Cancer Prospective Family Study Cohort (ProF-SC). International Journal of Epidemiology, 2016, 45, 683-692.	1.9	48
30	Characterization of the Cancer Spectrum in Men With Germline <i>BRCA1</i> Aland <i>BRCA2</i> Aland <i>BRCA2</i> Aland <i>BRCA2</i> Aland <i>BRCA2</i> Aland <i>BRCA2</i> AlandA	7.1	48
31	DNA Glycosylases Involved in Base Excision Repair May Be Associated with Cancer Risk in BRCA1 and BRCA2 Mutation Carriers. PLoS Genetics, 2014, 10, e1004256.	3.5	47
32	Rare Mutations in <i>RINT1</i> Predispose Carriers to Breast and Lynch Syndrome–Spectrum Cancers. Cancer Discovery, 2014, 4, 804-815.	9.4	44
33	Regular use of aspirin and other non-steroidal anti-inflammatory drugs and breast cancer risk for women at familial or genetic risk: a cohort study. Breast Cancer Research, 2019, 21, 52.	5.0	44
34	Inheritance of deleterious mutations at both BRCA1 and BRCA2 in an international sample of $32,295$ women. Breast Cancer Research, $2016,18,112.$	5.0	42
35	Risk-reducing salpingo-oophorectomy, natural menopause, and breast cancer risk: an international prospective cohort of BRCA1 and BRCA2 mutation carriers. Breast Cancer Research, 2020, 22, 8.	5.0	41
36	Association of Genomic Domains in <i>BRCA1</i> and <i>BRCA2</i> with Prostate Cancer Risk and Aggressiveness. Cancer Research, 2020, 80, 624-638.	0.9	39

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37	Recreational Physical Activity Is Associated with Reduced Breast Cancer Risk in Adult Women at High Risk for Breast Cancer: A Cohort Study of Women Selected for Familial and Genetic Risk. Cancer Research, 2020, 80, 116-125.	0.9	37
38	Comparison of Clinical, Maternal, and Self Pubertal Assessments: Implications for Health Studies. Pediatrics, $2016,138,\ldots$	2.1	36
39	Oral Contraceptive Use and Breast Cancer Risk: Retrospective and Prospective Analyses From a BRCA1 and BRCA2 Mutation Carrier Cohort Study. JNCI Cancer Spectrum, 2018, 2, pky023.	2.9	33
40	Identification of independent association signals and putative functional variants for breast cancer risk through fine-scale mapping of the 12p11 locus. Breast Cancer Research, 2016, 18, 64.	5.0	31
41	Height and Body Mass Index as Modifiers of Breast Cancer Risk in <i>BRCA1</i> / <i>2</i> Mutation Carriers: A Mendelian Randomization Study. Journal of the National Cancer Institute, 2019, 111, 350-364.	6.3	30
42	An original phylogenetic approach identified mitochondrial haplogroup T1a1 as inversely associated with breast cancer risk in BRCA2 mutation carriers. Breast Cancer Research, 2015, 17, 61.	5.0	26
43	An international survey of surveillance schemes for unaffected BRCA1 and BRCA2 mutation carriers. Breast Cancer Research and Treatment, 2016, 157, 319-327.	2.5	26
44	Alcohol Consumption, Cigarette Smoking, and Risk of Breast Cancer for <i>BRCA1</i> and <i>BRCA2</i> Mutation Carriers: Results from The BRCA1 and BRCA2 Cohort Consortium. Cancer Epidemiology Biomarkers and Prevention, 2020, 29, 368-378.	2.5	24
45	Polygenic risk modeling for prediction of epithelial ovarian cancer risk. European Journal of Human Genetics, 2022, 30, 349-362.	2.8	23
46	Candidate Genetic Modifiers for Breast and Ovarian Cancer Risk in <i>BRCA1</i> and <i>BRCA2</i> Mutation Carriers. Cancer Epidemiology Biomarkers and Prevention, 2015, 24, 308-316.	2.5	22
47	Breast Cancer Risk Perceptions among Relatives of Women with Uninformative Negative <i>BRCA1/2</i> Test Results: The Moderating Effect of the Amount of Shared Information. Journal of Genetic Counseling, 2016, 25, 258-269.	1.6	21
48	A Phase 1 dose-escalation study of disulfiram and copper gluconate in patients with advanced solid tumors involving the liver using S-glutathionylation as a biomarker. BMC Cancer, 2021, 21, 510.	2.6	21
49	Mendelian randomisation study of height and body mass index as modifiers of ovarian cancer risk in 22,588 BRCA1 and BRCA2 mutation carriers. British Journal of Cancer, 2019, 121, 180-192.	6.4	19
50	Breast and Prostate Cancer Risks for Male <i>BRCA1</i> and <i>BRCA2</i> Pathogenic Variant Carriers Using Polygenic Risk Scores. Journal of the National Cancer Institute, 2022, 114, 109-122.	6.3	19
51	No clinical utility of KRAS variant rs61764370 for ovarian or breast cancer. Gynecologic Oncology, 2016, 141, 386-401.	1.4	18
52	Pubertal development in girls by breast cancer family history: the LEGACY girls cohort. Breast Cancer Research, 2017, 19, 69.	5.0	18
53	Association of breast cancer risk in BRCA1 and BRCA2 mutation carriers with genetic variants showing differential allelic expression: identification of a modifier of breast cancer risk at locus 11q22.3. Breast Cancer Research and Treatment, 2017, 161, 117-134.	2.5	18
54	Do cancer survivors develop healthier lifestyle behaviors than the cancer-free population in the PLCO study?. Journal of Cancer Survivorship, 2017, 11, 233-245.	2.9	18

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55	Multigene and Drug Interaction Approach for Tamoxifen Metabolite Patterns Reveals Possible Involvement of CYP2C9, CYP2C19, and <i>ABCB1</i> . Journal of Clinical Pharmacology, 2016, 56, 1570-1581.	2.0	17
56	Fiber intake and the risk of head and neck cancer in the prostate, lung, colorectal and ovarian (PLCO) cohort. International Journal of Cancer, 2019, 145, 2342-2348.	5.1	17
57	Analysis of Serial Ovarian Volume Measurements and Incidence of Ovarian Cancer: Implications for Pathogenesis. Journal of the National Cancer Institute, 2014, 106, .	6.3	16
58	The impact of folate intake on the risk of head and neck cancer in the prostate, lung, colorectal, and ovarian cancer screening trial (PLCO) cohort. British Journal of Cancer, 2018, 118, 299-306.	6.4	16
59	Common variants in breast cancer risk loci predispose to distinct tumor subtypes. Breast Cancer Research, 2022, 24, 2.	5.0	15
60	Integrative analyses reveal signaling pathways underlying familial breast cancer susceptibility. Molecular Systems Biology, 2016, 12, 860.	7.2	14
61	Non-invasive optical spectroscopic monitoring of breast development during puberty. Breast Cancer Research, 2017, 19, 12.	5.0	14
62	Comparing 5-Year and Lifetime Risks of Breast CancerÂusing the Prospective Family Study Cohort. Journal of the National Cancer Institute, 2021, 113, 785-791.	6.3	13
63	Patient Interactions With an Automated Conversational Agent Delivering Pretest Genetics Education: Descriptive Study. Journal of Medical Internet Research, 2021, 23, e29447.	4.3	11
64	Breast cancer family history and allele-specific DNA methylation in the legacy girls study. Epigenetics, 2018, 13, 240-250.	2.7	10
65	Comparison of methods to assess onset of breast development in the LEGACY Girls Study: methodological considerations for studies of breast cancer. Breast Cancer Research, 2018, 20, 33.	5.0	9
66	A Phase I Study of Neoadjuvant Chemotherapy With Nab-Paclitaxel, Doxorubicin, and Cyclophosphamide in Patients With Stage II to III Breast Cancer. Clinical Breast Cancer, 2017, 17, 503-509.	2.4	7
67	Population-based relative risks for specific family history constellations of breast cancer. Cancer Causes and Control, 2019, 30, 581-590.	1.8	7
68	Association of Prepubertal and Adolescent Androgen Concentrations With Timing of Breast Development and Family History of Breast Cancer. JAMA Network Open, 2019, 2, e190083.	5.9	7
69	Association of germline genetic variants with breast cancer-specific survival in patient subgroups defined by clinic-pathological variables related to tumor biology and type of systemic treatment. Breast Cancer Research, 2021, 23, 86.	5.0	7
70	Effect of volume and pH on surface receptor number in macrophages. Journal of Cellular Physiology, 1989, 140, 371-378.	4.1	4
71	21-gene recurrence score testing utilization among older women from different races: A population-based study. Journal of Geriatric Oncology, 2021, 12, 206-211.	1.0	4
72	Weight is More Informative than Body Mass Index for Predicting Postmenopausal Breast Cancer Risk: Prospective Family Study Cohort (ProF-SC). Cancer Prevention Research, 2022, 15, 185-191.	1.5	4

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73	Body Mass Index and Mammographic Density in a Multiracial and Multiethnic Population-Based Study. Cancer Epidemiology Biomarkers and Prevention, 2022, 31, 1313-1323.	2.5	3
74	Response: Table 1 Journal of the National Cancer Institute, 2016, 108, djw173.	6.3	2
75	A competing risks model with binary time varying covariates for estimation of breast cancer risks in <i>BRCA1</i> families. Statistical Methods in Medical Research, 2021, 30, 2165-2183.	1.5	2
76	Evaluation of the relevance and access of EHR-based variables to support personalized medicine in breast cancer. Cogent Medicine, 2016, 3, 1234661.	0.7	1
77	Considerations When Using Breast Cancer Risk Models for Women with Negative BRCA1/BRCA2 Mutation Results. Journal of the National Cancer Institute, 2020, 112, 418-422.	6.3	1
78	Recreational Physical Activity and Outcomes After Breast Cancer in Women at High Familial Risk. JNCI Cancer Spectrum, 2021, 5, pkab090.	2.9	1
79	Maternal and prenatal factors and age at thelarche in the LEGACY Girls Study cohort: implications for breast cancer risk. International Journal of Epidemiology, 2023, 52, 272-283.	1.9	1
80	31: Impact of Multiple CYP Variants on Tamoxifen Metabolite Concentrations. American Journal of Clinical Pathology, 2015, 143, A015-A015.	0.7	0
81	OUP accepted manuscript. International Journal of Epidemiology, 2022, , .	1.9	O