Tuya Pal

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/1441517/publications.pdf

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69	2,758	361413	182427
papers	citations	20 h-index	51 g-index

69	69	69	4418
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Diversity in cancer genomics research is a matter of equity and scientific discovery. Genetics in Medicine, 2022, 24, 549-551.	2.4	2
2	An overview of genetic services delivery for hereditary breast cancer. Breast Cancer Research and Treatment, 2022, 191, 491-500.	2.5	12
3	Oncotype DX Risk Recurrence Score and Total Mortality for Early-Stage Breast Cancer by Race/Ethnicity. Cancer Epidemiology Biomarkers and Prevention, 2022, 31, 821-830.	2.5	10
4	A pooled case-only analysis of obesity and breast cancer subtype among Black women in the southeastern United States. Cancer Causes and Control, 2022, 33, 515-524.	1.8	3
5	Socioeconomic disparities in psychosocial service recommendation and receipt among young Black breast cancer survivors. Supportive Care in Cancer, 2022, , 1.	2.2	1
6	Abstract P3-14-11: Mammaprint and Blueprint identify genomic differences in HR+ HER2- breast cancers from young Black and White women. Cancer Research, 2022, 82, P3-14-11-P3-14-11.	0.9	0
7	Evaluating breast cancer predisposition genes in women of African ancestry. Genetics in Medicine, 2022, 24, 1468-1475.	2.4	2
8	Anxiety and depression among Black breast cancer survivors: Examining the role of patient-provider communication and cultural values. Patient Education and Counseling, 2022, 105, 2391-2396.	2.2	7
9	Bilateral Oophorectomy and the Risk of Breast Cancer in BRCA1 Mutation Carriers: A Reappraisal. Cancer Epidemiology Biomarkers and Prevention, 2022, 31, 1351-1358.	2.5	3
10	Whole transcriptomic analysis of HR+ breast cancer in Black women classified as basal-type by BluePrint.. Journal of Clinical Oncology, 2022, 40, 517-517.	1.6	1
11	Oncologist participation in pilot testing a crowdsourcing platform to build a survivorship care risk model.. Journal of Clinical Oncology, 2022, 40, e13568-e13568.	1.6	0
12	Family communication of genetic test results among women with inherited breast cancer genes. Journal of Genetic Counseling, 2021, 30, 701-709.	1.6	11
13	Radiotherapy after breastâ€conserving surgery for elderly patients with earlyâ€stage breast cancer: A national registryâ€based study. International Journal of Cancer, 2021, 148, 857-867.	5.1	2
14	Qualitative Methods for Refining a Web-Based Educational Tool for Patients Focused on Inherited Cancer Predisposition. Journal of Cancer Education, 2021, , 1.	1.3	4
15	Abstract PD11-05: Diabetes decreases overall survival in women with breast cancer in the southern community cohort study. , 2021, , .		1
16	Sharing genetic test results with family members of BRCA, PALB2, CHEK2, and ATM carriers. Patient Education and Counseling, 2021, 104, 720-725.	2.2	21
17	Breast Cancer Disparities Through the Lens of the COVID-19 Pandemic. Current Breast Cancer Reports, 2021, 13, 110-112.	1.0	8
18	Framework for Implementing and Tracking a Molecular Tumor Board at a National Cancer Instituteâ€"Designated Comprehensive Cancer Center. Oncologist, 2021, 26, e1962-e1970.	3.7	11

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19	IMProving care After inherited Cancer Testing (IMPACT) study: protocol of a randomized trial evaluating the efficacy of two interventions designed to improve cancer risk management and family communication of genetic test results. BMC Cancer, 2021, 21, 1099.	2.6	5
20	Follow-up Interactive Long-Term Expert Ranking (FILTER): a crowdsourcing platform to adjudicate risk for survivorship care. JAMIA Open, 2021, 4, ooab090.	2.0	0
21	Impact of Genetic Testing on Risk-Management Behavior of Black Breast Cancer Survivors: A Longitudinal, Observational Study. Annals of Surgical Oncology, 2020, 27, 1659-1670.	1.5	6
22	Cancer Risks Associated With Germline PALB2 Pathogenic Variants: An International Study of 524 Families. Journal of Clinical Oncology, 2020, 38, 674-685.	1.6	270
23	Sex Disparity Observed for Oncotype DX Breast Recurrence Score in Predicting Mortality Among Patients with Early Stage ER-Positive Breast Cancer. Clinical Cancer Research, 2020, 26, 101-109.	7.0	14
24	Points to consider: is there evidence to support BRCA1/2 and other inherited breast cancer genetic testing for all breast cancer patients? A statement of the American College of Medical Genetics and Genomics (ACMG). Genetics in Medicine, 2020, 22, 681-685.	2.4	20
25	Update on multiâ€gene panel testing and communication of genetic test results. Breast Journal, 2020, 26, 1513-1519.	1.0	9
26	Strategies to enhance identification of hereditary breast cancer gene carriers. Expert Review of Molecular Diagnostics, 2020, 20, 861-865.	3.1	1
27	Contribution of Germline Predisposition Gene Mutations to Breast Cancer Risk in African American Women. Journal of the National Cancer Institute, 2020, 112, 1213-1221.	6.3	51
28	Disparities in Genetic Testing and Care Among Black Women with Hereditary Breast Cancer. Current Breast Cancer Reports, 2020, 12, 125-131.	1.0	29
29	Disparities in BRCA counseling across providers in a diverse population of young breast cancer survivors. Genetics in Medicine, 2020, 22, 1088-1093.	2.4	10
30	Breast cancer screening implications of risk modeling among female relatives of ATM and CHEK2 carriers. Cancer, 2020, 126, 1651-1655.	4.1	4
31	The big reveal: Family disclosure patterns of BRCA genetic test results among young Black women with invasive breast cancer. Journal of Genetic Counseling, 2020, 29, 410-422.	1.6	24
32	Patterns and covariates of benefit finding in young Black breast cancer survivors: A longitudinal, observational study. Psycho-Oncology, 2020, 29, 1115-1122.	2.3	7
33	Acceptability and outcomes of multigene panel testing among young Black breast cancer survivors. Breast Journal, 2020, 26, 2112-2114.	1.0	1
34	Cancer risk management among female BRCA1/2, PALB2, CHEK2, and ATM carriers. Breast Cancer Research and Treatment, 2020, 182, 421-428.	2.5	23
35	A Web-Based Tool to Automate Portions of Pretest Genetic Counseling for Inherited Cancer. Journal of the National Comprehensive Cancer Network: JNCCN, 2020, 18, 841-847.	4.9	18
36	Abstract P6-08-20: Cancer risk management and family sharing of genetic test results among women with inherited breast cancer genes. , 2020, , .		1

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37	Overall Mortality After Diagnosis of Breast Cancer in Men vs Women. JAMA Oncology, 2019, 5, 1589.	7.1	103
38	Genetic Testing Across Young Hispanic and Non-Hispanic White Breast Cancer Survivors: Facilitators, Barriers, and Awareness of the Genetic Information Nondiscrimination Act. Genetic Testing and Molecular Biomarkers, 2019, 23, 75-83.	0.7	27
39	A randomized controlled intervention to promote readiness to genetic counseling for breast cancer survivors. Psycho-Oncology, 2019, 28, 980-988.	2.3	8
40	Health beliefs associated with readiness for genetic counseling among high risk breast cancer survivors. Breast Journal, 2019, 25, 117-123.	1.0	7
41	Preferences for multigene panel testing for hereditary breast cancer risk among ethnically diverse BRCA-uninformative families. Journal of Community Genetics, 2018, 9, 81-92.	1.2	11
42	Considerations in Testing for Inherited Breast Cancer Predisposition in the Era of Personalized Medicine. Surgical Oncology Clinics of North America, 2018, 27, 1-22.	1.5	12
43	The Inherited Cancer Registry (ICARE) Initiative: An Academic-Community Partnership for Patients and Providers. Oncology Issues, 2018, 33, 54-63.	0.1	7
44	Psychosocial impact of BRCA testing in young Black breast cancer survivors. Psycho-Oncology, 2018, 27, 2778-2785.	2.3	7
45	Care delivery considerations for widespread and equitable implementation of inherited cancer predisposition testing. Expert Review of Molecular Diagnostics, 2017, 17, 57-70.	3.1	30
46	Correlation between germline mutations in MMR genes and microsatellite instability in ovarian cancer specimens. Familial Cancer, 2017, 16, 351-355.	1.9	18
47	Racial disparities in BRCA testing and cancer risk management across a populationâ€based sample of young breast cancer survivors. Cancer, 2017, 123, 2497-2505.	4.1	192
48	NCCN Guidelines Insights: Genetic/Familial High-Risk Assessment: Breast and Ovarian, Version 2.2017. Journal of the National Comprehensive Cancer Network: JNCCN, 2017, 15, 9-20.	4.9	408
49	Comment on â€œCan Breast Surgeons Provide Breast Cancer Genetic Testing? An American Society of Breast Surgeons Surveyâ€. Annals of Surgical Oncology, 2017, 24, 588-589.	1.5	2
50	Bilateral Oophorectomy and Breast Cancer Risk inBRCA1andBRCA2Mutation Carriers. Journal of the National Cancer Institute, 2017, 109, .	6.3	160
51	Hereditary Cancer: Example of a Public Health Approach to Ensure Population Health Benefits of Genetic Medicine. Healthcare (Switzerland), 2016, 4, 6.	2.0	10
52	Recruitment of a Populationâ€Based Sample of Young Black Women with Breast Cancer through a State Cancer Registry. Breast Journal, 2016, 22, 166-172.	1.0	12
53	Evolution of Hereditary Breast Cancer Genetic Services: Are Changes Reflected in the Knowledge and Clinical Practices of Florida Providers?. Genetic Testing and Molecular Biomarkers, 2016, 20, 569-578.	0.7	14
54	A high frequency of BRCA mutations in young black women with breast cancer residing in Florida. Cancer, 2015, 121, 4173-4180.	4.1	91

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55	Differences in BRCA counseling and testing practices based on ordering provider type. Genetics in Medicine, 2015, 17, 51-57.	2.4	47
56	Factors associated with genetic counseling and BRCA testing in a population-based sample of young Black women with breast cancer. Breast Cancer Research and Treatment, 2015, 151, 169-176.	2.5	78
57	Clinical Considerations in the Conduct of Cancer Next-Generation Sequencing Testing and Genetic Counseling. , 2015, , 81-101.		4
58	Factors Which Impact the Delivery of Genetic Risk Assessment Services Focused on Inherited Cancer Genomics: Expanding the Role and Reach of Certified Genetics Professionals. Journal of Genetic Counseling, 2014, 23, 522-530.	1.6	38
59	Educational Needs and Preferred Methods of Learning Among Florida Practitioners Who Order Genetic Testing for Hereditary Breast and Ovarian Cancer. Journal of Cancer Education, 2013, 28, 690-697.	1.3	17
60	Early Onset Breast Cancer in a Registry-based Sample of African-American Women:BRCAMutation Prevalence, and Other Personal and System-level Clinical Characteristics. Breast Journal, 2013, 19, 189-192.	1.0	32
61	Identification, Evaluation, and Treatment of Patients with Hereditary Cancer Risk within the United States. ISRN Oncology, 2013, 2013, 1-8.	2.1	8
62	Genetic Risk Assessments in Individuals at High Risk for Inherited Breast Cancer in the Breast Oncology Care Setting. Cancer Control, 2012, 19, 255-266.	1.8	37
63	Recruitment of Black Women for a Study of Inherited Breast Cancer Using a Cancer Registryâ€“Based Approach. Genetic Testing and Molecular Biomarkers, 2011, 15, 69-77.	0.7	23
64	Development of a Brochure for Increasing Awareness of Inherited Breast Cancer in Black Women. Genetic Testing and Molecular Biomarkers, 2011, 15, 59-67.	0.7	15
65	Updating and refining a study brochure for a cancer registry-based study of BRCA mutations among young African American breast cancer patients: lessons learned. Journal of Community Genetics, 2010, 1, 63-71.	1.2	10
66	Development of a culturally tailored genetic counseling booklet about hereditary breast and ovarian cancer for Black women. American Journal of Medical Genetics, Part A, 2010, 152A, 836-845.	1.2	20
67	Fertility in women with BRCA mutations: a case-control study. Fertility and Sterility, 2010, 93, 1805-1808.	1.0	69
68	BRCA1 and BRCA2 mutations account for a large proportion of ovarian carcinoma cases. Cancer, 2005, 104, 2807-2816.	4.1	622
69	BRCA1 and BRCA2 mutations in a study of African American breast cancer patients. Cancer Epidemiology Biomarkers and Prevention, 2004, 13, 1794-9.	2.5	27