## Torben A Kruse

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/3573719/publications.pdf

Version: 2024-02-01

86 papers

4,078 citations

29 h-index

172457

60 g-index

90 all docs 90 docs citations

90 times ranked 8662 citing authors

| #  | Article   | IF          | Citations |
|----|---|-------------|-----------|
| 1  | Multiple independent variants at the TERT locus are associated with telomere length and risks of breast and ovarian cancer. Nature Genetics, 2013, 45, 371-384.   | 21.4        | 493       |
| 2  | Association of Type and Location of <i>BRCA1 </i> 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       |
| 3  | Identification of 12 new susceptibility loci for different histotypes of epithelial ovarian cancer.<br>Nature Genetics, 2017, 49, 680-691.  | 21.4        | 356       |
| 4  | Identification of ten variants associated with risk of estrogen-receptor-negative breast cancer. Nature Genetics, 2017, 49, 1767-1778.  | 21.4        | 289       |
| 5  | Mutational spectrum in a worldwide study of 29,700 families with <i>BRCA1</i> or <i>BRCA2</i> mutations. Human Mutation, 2018, 39, 593-620.   | 2.5         | 224       |
| 6  | Identification of six new susceptibility loci for invasive epithelial ovarian cancer. Nature Genetics, 2015, 47, 164-171.   | 21.4        | 221       |
| 7  | Prediction of Breast and Prostate Cancer Risks in Male <i>BRCA1 </i> and <ibrca2 <="" i=""> Mutation Carriers Using Polygenic Risk Scores. Journal of Clinical Oncology, 2017, 35, 2240-2250.</ibrca2>  | 1.6         | 152       |
| 8  | Evaluation of Nine Somatic Variant Callers for Detection of Somatic Mutations in Exome and Targeted Deep Sequencing Data. PLoS ONE, 2016, 11, e0151664.   | <b>2.</b> 5 | 144       |
| 9  | Large scale multifactorial likelihood quantitative analysis of <i>BRCA1</i> and <i>BRCA2</i> variants:<br>An ENIGMA resource to support clinical variant classification. Human Mutation, 2019, 40, 1557-1578.   | 2.5         | 102       |
| 10 | Classifications within Molecular Subtypes Enables Identification of BRCA1/BRCA2 Mutation Carriers by RNA Tumor Profiling. PLoS ONE, 2013, 8, e64268.  | <b>2.</b> 5 | 89        |
| 11 | 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        |
| 12 | Familial Isolated Hyperparathyroidism as a Variant of Multiple Endocrine Neoplasia Type $1$ in a Large Danish Pedigree1. Journal of Clinical Endocrinology and Metabolism, 2000, 85, 165-167.   | 3.6         | 74        |
| 13 | Hereditary Breast Cancer: Clinical, Pathological and Molecular Characteristics. Breast Cancer: Basic and Clinical Research, 2014, 8, BCBCR.S18715.  | 1.1         | 71        |
| 14 | Ruxolitinib and interferon-α2 combination therapy for patients with polycythemia vera or myelofibrosis: a phase II study. Haematologica, 2020, 105, 2262-2272.  | 3.5         | 67        |
| 15 | Molecular signature of different lesion types in the brain white matter of patients with progressive multiple sclerosis. Acta Neuropathologica Communications, 2019, 7, 205.  | <b>5.</b> 2 | 61        |
| 16 | Allergic rhinitis $\hat{a}\in$ a total genome-scan for susceptibility genes suggests a locus on chromosome 4q24-q27. European Journal of Human Genetics, 2001, 9, 945-952.  | 2.8         | 59        |
| 17 | Whole Blood Transcriptional Profiling Reveals Deregulation of Oxidative and Antioxidative Defence Genes in Myelofibrosis and Related Neoplasms. Potential Implications of Downregulation of Nrf2 for Genomic Instability and Disease Progression. PLoS ONE, 2014, 9, e112786. | 2.5         | 59        |
| 18 | A new locus for Seckel syndrome on chromosome 18p11.31-q11.2. European Journal of Human Genetics, 2001, 9, 753-757.   | 2.8         | 54        |

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|----|--|-----|-----------|
| 19 | Mathematical modelling as a proof of concept for MPNs as a human inflammation model for cancer development. PLoS ONE, 2017, 12, e0183620.  | 2.5 | 51        |
| 20 | The <i>BRCA1</i> c. 5096G>A p.Arg1699Gln (R1699Q) intermediate risk variant: breast and ovarian cancer risk estimation and recommendations for clinical management from the ENIGMA consortium. Journal of Medical Genetics, 2018, 55, 15-20. | 3.2 | 50        |
| 21 | Long non-coding RNA expression profiles predict metastasis in lymph node-negative breast cancer independently of traditional prognostic markers. Breast Cancer Research, 2015, 17, 55.   | 5.0 | 49        |
| 22 | 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        |
| 23 | Molecular Concordance Between Primary Breast Cancer and Matched Metastases. Breast Journal, 2016, 22, 420-430.   | 1.0 | 44        |
| 24 | Clonal expansion and linear genome evolution through breast cancer progression from pre-invasive stages to asynchronous metastasis. Oncotarget, 2015, 6, 5634-5649.  | 1.8 | 42        |
| 25 | 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        |
| 26 | Differential Dynamics of CALR Mutant Allele Burden in Myeloproliferative Neoplasms during Interferon Alfa Treatment. PLoS ONE, 2016, 11, e0165336.   | 2.5 | 38        |
| 27 | Safety and efficacy of combination therapy of interferonâ€Î±2 and ruxolitinib in polycythemia vera and myelofibrosis. Cancer Medicine, 2018, 7, 3571-3581.   | 2.8 | 38        |
| 28 | Search for a shared segment on chromosome 10q26 in patients with bipolar affective disorder or schizophrenia from the Faroe Islands. American Journal of Medical Genetics Part A, 2002, 114, 196-204.  | 2.4 | 34        |
| 29 | Assessing Associations between the AURKA-HMMR-TPX2-TUBG1 Functional Module and Breast Cancer Risk in BRCA1/2 Mutation Carriers. PLoS ONE, 2015, 10, e0120020.  | 2.5 | 34        |
| 30 | Human longevity and variation in DNA damage response and repair: study of the contribution of sub-processes using competitive gene-set analysis. European Journal of Human Genetics, 2014, 22, 1131-1136.                                    | 2.8 | 31        |
| 31 | Myelinâ€specific <scp>T</scp> cells induce interleukinâ€1 beta expression in lesionâ€reactive microglialâ€like cells in zones of axonal degeneration. Glia, 2016, 64, 407-424.   | 4.9 | 28        |
| 32 | The microRNA-132/212 family fine-tunes multiple targets in Angiotensin II signalling in cardiac fibroblasts. JRAAS - Journal of the Renin-Angiotensin-Aldosterone System, 2015, 16, 1288-1297.   | 1.7 | 27        |
| 33 | 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        |
| 34 | Genomic profiling of a randomized trial of interferon- $\hat{l}\pm$ vs hydroxyurea in MPN reveals mutation-specific responses. Blood Advances, 2022, 6, 2107-2119.   | 5.2 | 26        |
| 35 | The subclonal structure and genomic evolution of oral squamous cell carcinoma revealed by ultra-deep sequencing. Oncotarget, 2017, 8, 16571-16580.   | 1.8 | 25        |
| 36 | Genomic Analyses of Breast Cancer Progression Reveal Distinct Routes of Metastasis Emergence.<br>Scientific Reports, 2017, 7, 43813.   | 3.3 | 24        |

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|----|--|-----|-----------|
| 37 | Identification of metastasis driver genes by massive parallel sequencing of successive steps of breast cancer progression. PLoS ONE, 2018, 13, e0189887.   | 2.5 | 24        |
| 38 | Association of miR-548c-5p, miR-7-5p, miR-210-3p, miR-128-3p with recurrence in systemically untreated breast cancer. Oncotarget, 2018, 9, 9030-9042.  | 1.8 | 22        |
| 39 | CFP suppresses breast cancer cell growth by TES-mediated upregulation of the transcription factor DDIT3. Oncogene, 2019, 38, 4560-4573.  | 5.9 | 22        |
| 40 | Subtypes in BRCA-mutated breast cancer. Human Pathology, 2019, 84, 192-201.  | 2.0 | 22        |
| 41 | Microarray-Based RNA Profiling of Breast Cancer: Batch Effect Removal Improves Cross-Platform Consistency. BioMed Research International, 2014, 2014, 1-11.  | 1.9 | 21        |
| 42 | Use of next generation sequencing in head and neck squamous cell carcinomas: A review. Oral Oncology, 2014, 50, 1035-1040.   | 1.5 | 21        |
| 43 | Effect of thrombopoietin receptor agonists on markers of coagulation and P-selectin in patients with immune thrombocytopenia. Platelets, 2019, 30, 206-212.  | 2.3 | 21        |
| 44 | 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        |
| 45 | RNA profiling reveals familial aggregation of molecular subtypes in non-BRCA1/2 breast cancer families. BMC Medical Genomics, 2014, 7, 9.  | 1.5 | 18        |
| 46 | The impact of interferon-alpha2 on HLA genes in patients with polycythemia vera and related neoplasms. Leukemia and Lymphoma, 2017, 58, 1914-1921.   | 1.3 | 17        |
| 47 | Investigating a case of possible field cancerization in oral squamous cell carcinoma by the use of next-generation sequencing. Oral Oncology, 2017, 68, 74-80.   | 1.5 | 15        |
| 48 | Sorted peripheral blood cells identify <i>CALR</i> mutations in B- and T-lymphocytes. Leukemia and Lymphoma, 2018, 59, 973-977.  | 1.3 | 15        |
| 49 | Tumor-specific genetic aberrations in cell-free DNA of gastroesophageal cancer patients. Journal of Gastroenterology, 2019, 54, 108-121.   | 5.1 | 14        |
| 50 | Long-Term Efficacy and Safety of Recombinant Interferon Alpha-2 Vs. Hydroxyurea in Polycythemia Vera: Preliminary Results from the Three-Year Analysis of the Daliah Trial - a Randomized Controlled Phase III Clinical Trial. Blood, 2018, 132, 580-580.  | 1.4 | 14        |
| 51 | Safety and Efficacy of Combination Therapy of Interferon-Alpha2 + JAK1-2 Inhibitor in the Philadelphia-Negative Chronic Myeloproliferative Neoplasms. Preliminary Results from the Danish Combi-Trial - an Open Label, Single Arm, Non-Randomized Multicenter Phase II Study. Blood, 2015, 126, 824-824. | 1.4 | 14        |
| 52 | Transcriptional Profiling of Whole Blood Identifies a Unique 5-Gene Signature for Myelofibrosis and Imminent Myelofibrosis Transformation. PLoS ONE, 2014, 9, e85567.  | 2.5 | 13        |
| 53 | Epithelial ovarian cancer and the use of circulating tumor DNA: A systematic review. Gynecologic Oncology, 2021, 161, 884-895.   | 1.4 | 12        |
| 54 | Spotting and validation of a genome wide oligonucleotide chip with duplicate measurement of each gene. Biochemical and Biophysical Research Communications, 2006, 344, 1111-1120.  | 2.1 | 11        |

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|----|--|-------------|-----------|
| 55 | Tumour-infiltrating CD4-, CD8- and FOXP3-positive immune cells as predictive markers of mortality in BRCA1- and BRCA2-associated breast cancer. British Journal of Cancer, 2021, 125, 1388-1398.   | 6.4         | 11        |
| 56 | Whole Blood Gene Expression Profiling in patients undergoing colon cancer surgery identifies differential expression of genes involved in immune surveillance, inflammation and carcinogenesis. Surgical Oncology, 2018, 27, 208-215.  | 1.6         | 10        |
| 57 | High Expression of Carcinoembryonic Antigen-Related Cell Adhesion Molecule(CEACAM) 6 In Primary Myelofibrosis. Blood, 2010, 116, 4116-4116.  | 1.4         | 10        |
| 58 | Myeloproliferative Neoplasms in Danish Twins. Acta Haematologica, 2018, 139, 195-198.  | 1.4         | 8         |
| 59 | Heterogeneity and tumor evolution reflected in liquid biopsy in metastatic breast cancer patients: a review. Cancer and Metastasis Reviews, 2022, 41, 433-446.   | <b>5.</b> 9 | 8         |
| 60 | Feature Selection for Predicting Tumor Metastases in Microarray Experiments using Paired Design. Cancer Informatics, 2007, 3, 117693510700300.   | 1.9         | 7         |
| 61 | The gene expression and immunohistochemical timeâ€course of diphenylcyclopropenoneâ€induced contact allergy in healthy humans following repeated epicutaneous challenges. Experimental Dermatology, 2017, 26, 926-933.   | 2.9         | 7         |
| 62 | Molecular characterization of sorted malignant B cells from patients clinically identified with mantle cell lymphoma. Experimental Hematology, 2020, 84, 7-18.e12.   | 0.4         | 7         |
| 63 | Significantly Upregulated Thrombo-Inflammatory Genes Are Normoregulated or Significantly<br>Downregulated during Treatment with Interferon-Alpha2 in Patients with Philadelphia-Negative<br>Chronic Myeloproliferative Neoplasms. Blood, 2019, 134, 2978-2978.   | 1.4         | 6         |
| 64 | A 7-Gene Signature Depicts the Biochemical Profile of Early Prefibrotic Myelofibrosis. PLoS ONE, 2016, 11, e0161570.   | 2.5         | 6         |
| 65 | Increased oxidative stress with substantial dysregulation of genes related to oxidative stress and DNA repair after laparoscopic colon cancer surgery. Surgical Oncology, 2020, 35, 71-78.   | 1.6         | 5         |
| 66 | Increased Expression of Proteasome-Related Genes In Patients with Primary Myelofibrosis. Blood, 2010, 116, 4117-4117.  | 1.4         | 5         |
| 67 | Interferon-alfa2 Treatment of Patients with Polycythemia Vera and Related Neoplasms Impacts Deregulation of Oxidative Stress Genes and Antioxidative Defence Mechanisms. Potential Implications of IFN-Alfa Induced Changes in TP53, NRF2 and CXCR4 for Genomic Instability and CD34+ Mobilisation. Blood. 2018. 132. 4326-4326. | 1.4         | 3         |
| 68 | Interferon-alfa2 Treatment of Patients with Polycythemia Vera and Related Neoplasms Influences<br>Deregulated Inflammation and Immune Genes in Polycythemia Vera and Allied Neoplasms. Blood, 2018,<br>132, 5490-5490.   | 1.4         | 3         |
| 69 | The Optimal Sequencing Depth of Tumor Biopsies for Identifying Clonal Cell Populations. Journal of Molecular Diagnostics, 2019, 21, 790-795.   | 2.8         | 2         |
| 70 | Extracellular Matrix-Related Genes Are Deregulated in Peripheral Blood from Patients with Myelofibrosis and Related Neoplasms. Blood, 2018, 132, 5491-5491.  | 1.4         | 2         |
| 71 | Search for a shared segment on chromosome 10q26 in patients with bipolar affective disorder or schizophrenia from the Faroe Islands. American Journal of Medical Genetics Part A, 2002, 114, 196-204.  | 2.4         | 1         |
| 72 | The Impact of Interferon on Interferon-Related Genes in Polycythemia Vera and Allied Neoplasms. Blood, 2018, 132, 4328-4328.   | 1.4         | 1         |

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|----|---|-----|-----------|
| 73 | Genomic Profiling of a Phase III Clinical Trial of Interferon Versus Hydroxyurea in MPN Patients<br>Reveals Mutation-Specific and Treatment-Specific Patterns of Response. Blood, 2019, 134, 4202-4202.             | 1.4 | 1         |
| 74 | Increased Gene Expression of Histone Deacetylases In Patients with Philadelphia-Negative Chronic Myeloproliferative Neoplasms. Blood, 2010, 116, 4119-4119.   | 1.4 | 1         |
| 75 | Whole Blood Transcriptional Profiling Reveals Highly Deregulated Atherosclerosis Genes in Myeloproliferative Cancer. Blood, 2018, 132, 3071-3071.   | 1.4 | 1         |
| 76 | Highly Deregulated Fibulins in Patients with Philadelphia-Negative Chronic Myeloproliferative Neoplasms. Blood, 2019, 134, 5396-5396.   | 1.4 | 1         |
| 77 | Latent growth curve modeling of incomplete timecourse data in microarray gene expression studies. , 2012, , .   |     | 0         |
| 78 | OTEH-4. Deeper insight into intratumoral heterogeneity by MRI and PET-guided stereotactic biopsies from glioblastoma patients. Neuro-Oncology Advances, 2021, 3, ii11-ii11.   | 0.7 | 0         |
| 79 | Comparison of the Metastasis Predictive Potential of mRNA and Long Non-Coding RNA Profiling in Systemically Untreated Breast Cancer. Cancers, 2021, 13, 4907.   | 3.7 | 0         |
| 80 | Enhanced Gene Expression of EZH2 In Patients with Primary Myelofibrosis. Blood, 2010, 116, 4118-4118.   | 1.4 | 0         |
| 81 | Gene Expression Profiling with Principal Component Analysis Depicts the Biological Continuum From Essential Thrombocythemia Over Polycythemia Vera to Myelofibrosis. Blood, 2010, 116, 4115-4115.                   | 1.4 | О         |
| 82 | The Impact of Interferon-alpha2 on HLA-Genes in Patients with Polycythemia Vera and Related Neoplasms. Blood, 2015, 126, 4097-4097.   | 1.4 | 0         |
| 83 | Genetic Evidence for Involvement of Human Endogenous Retrovirus Herv-Fc1 in the Pathogenesis of MPNs. Blood, 2018, 132, 5488-5488.  | 1.4 | 0         |
| 84 | The Impact of the Mutational Landscape upon the Molecular Responses to Interferon-Alfa2 in Calr-Mutated MPN Patients. Blood, 2018, 132, 4327-4327.  | 1.4 | 0         |
| 85 | The Impact of Somatic Mutations upon the Response to Combination Therapy with Ruxolitinib and Interferon in MPN Patients. Blood, 2021, 138, 3589-3589.  | 1.4 | 0         |
| 86 | Abstract P5-07-07: Mapping clonal evolution and tumor heterogeneity by whole exome sequencing of tissue and plasma circulating tumor DNA in metastatic breast cancer. Cancer Research, 2022, 82, P5-07-07-P5-07-07. | 0.9 | 0         |