# Rob B Van Der Luijt

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/2403114/publications.pdf

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|---|---|---|---|
| 24<br>papers | 3,360<br>citations | 430442<br>18<br>h-index | 610482<br>24<br>g-index |
| 24<br>all docs | 24<br>docs citations | 24<br>times ranked | 7444<br>citing authors |

| # | Article | IF | Citations |
|---|---|---|---|
| 1 | Association analysis identifies 65 new breast cancer risk loci. Nature, 2017, 551, 92-94. | 13.7 | 1,099 |
| 2 | Genome-wide association analysis of more than 120,000 individuals identifies 15 new susceptibility loci for breast cancer. Nature Genetics, 2015, 47, 373-380. | 9.4 | 513 |
| 3 | 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. | 9.4 | 493 |
| 4 | Identification of 12 new susceptibility loci for different histotypes of epithelial ovarian cancer. Nature Genetics, 2017, 49, 680-691. | 9.4 | 356 |
| 5 | Targeted sequencing by proximity ligation for comprehensive variant detection and local haplotyping. Nature Biotechnology, 2014, 32, 1019-1025. | 9.4 | 231 |
| 6 | Genetically Predicted Body Mass Index and Breast Cancer Risk: Mendelian Randomization Analyses of Data from 145,000 Women of European Descent. PLoS Medicine, 2016, 13, e1002105. | 3.9 | 118 |
| 7 | MEN1 redefined, a clinical comparison of mutation-positive and mutation-negative patients. BMC Medicine, 2016, 14, 182. | 2.3 | 95 |
| 8 | The BRCA1 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. | 1.5 | 50 |
| 9 | MicroRNA Related Polymorphisms and Breast Cancer Risk. PLoS ONE, 2014, 9, e109973. | 1.1 | 49 |
| 10 | DNA Glycosylases Involved in Base Excision Repair May Be Associated with Cancer Risk in BRCA1 and BRCA2 Mutation Carriers. PLoS Genetics, 2014, 10, e1004256. | 1.5 | 47 |
| 11 | Association of Genomic Domains in BRCA1 and BRCA2 with Prostate Cancer Risk and Aggressiveness. Cancer Research, 2020, 80, 624-638. | 0.4 | 39 |
| 12 | Identification and characterization of novel associations in the CASP8/ALS2CR12 region on chromosome 2 with breast cancer risk. Human Molecular Genetics, 2015, 24, 285-298. | 1.4 | 38 |
| 13 | Assessing Associations between the AURKA-HMMR-TPX2-TUBG1 Functional Module and Breast Cancer Risk in BRCA1/2 Mutation Carriers. PLoS ONE, 2015, 10, e0120020. | 1.1 | 34 |
| 14 | BRCA1 Circos: a visualisation resource for functional analysis of missense variants. Journal of Medical Genetics, 2015, 52, 224-230. | 1.5 | 32 |
| 15 | Does rapid genetic counseling and testing in newly diagnosed breast cancer patients cause additional psychosocial distress? results from a randomized clinical trial. Genetics in Medicine, 2016, 18, 137-144. | 1.1 | 28 |
| 16 | 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. | 2.2 | 26 |
| 17 | Behavioral and psychosocial effects of rapid genetic counseling and testing in newly diagnosed breast cancer patients: Design of a multicenter randomized clinical trial. BMC Cancer, 2011, 11, 6. | 1.1 | 22 |
| 18 | Association of genetic susceptibility variants for type 2 diabetes with breast cancer risk in women of European ancestry. Cancer Causes and Control, 2016, 27, 679-693. | 0.8 | 21 |

| # | Article | IF | Citations |
|---|---|---|---|
| 19 | 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. | 1.1 | 18 |
| 20 | Uptake of prenatal diagnostic testing for retinoblastoma compared to other hereditary cancer syndromes in the Netherlands. Familial Cancer, 2017, 16, 271-277. | 0.9 | 16 |
| 21 | Germline and somatic mosaicism in a family with multiple endocrine neoplasia type 1 (MEN1) syndrome. European Journal of Endocrinology, 2019, 180, K15-K19. | 1.9 | 16 |
| 22 | Difference in CXCR4 expression between sporadic and VHL-related hemangioblastoma. Familial Cancer, 2016, 15, 607-616. | 0.9 | 11 |
| 23 | The association between cancer family history and ovarian cancer risk in BRCA1/2 mutation carriers: can it be explained by the mutation position?. European Journal of Human Genetics, 2018, 26, 848-857. | 1.4 | 5 |
| 24 | Recontacting non-BRCA1/2 breast cancer patients for germline CHEK2 c.1100del pathogenic variant testing: uptake and patient experiences. Hereditary Cancer in Clinical Practice, 2021, 19, 9. | 0.6 | 3 |