## Nancy F Hansen

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/2849886/publications.pdf

Version: 2024-02-01

687363 794594 6,664 22 13 19 citations h-index g-index papers 25 25 25 9736 docs citations times ranked citing authors all docs

| #  | Article  | IF   | CITATIONS |
|----|--|------|-----------|
| 1  | KLF3 and PAX6 are candidate driver genes in late-stage, MSI-hypermutated endometrioid endometrial carcinomas. PLoS ONE, 2022, 17, e0251286.  | 2.5  | 2         |
| 2  | A complete reference genome improves analysis of human genetic variation. Science, 2022, 376, eabl3533.  | 12.6 | 144       |
| 3  | The complete sequence of a human genome. Science, 2022, 376, 44-53.  | 12.6 | 1,222     |
| 4  | Long-read mapping to repetitive reference sequences using Winnowmap2. Nature Methods, 2022, 19, 705-710.   | 19.0 | 80        |
| 5  | Comparative clinical and genomic analysis of neurofibromatosis type 2-associated cranial and spinal meningiomas. Scientific Reports, 2020, 10, 12563.  | 3.3  | 16        |
| 6  | Telomere-to-telomere assembly of a complete human X chromosome. Nature, 2020, 585, 79-84.  | 27.8 | 549       |
| 7  | A robust benchmark for detection of germline large deletions and insertions. Nature Biotechnology, 2020, 38, 1347-1355.  | 17.5 | 233       |
| 8  | Admixture mapping identifies genetic regions associated with blood pressure phenotypes in African Americans. PLoS ONE, 2020, 15, e0232048.   | 2.5  | 12        |
| 9  | Title is missing!. , 2020, 15, e0232048.   |      | O         |
| 10 | Title is missing!. , 2020, 15, e0232048.   |      | 0         |
| 11 | Title is missing!. , 2020, 15, e0232048.   |      | O         |
| 12 | Title is missing!. , 2020, 15, e0232048.   |      | 0         |
| 13 | Low mutation burden and frequent loss of CDKN2A/B and SMARCA2, but not PRC2, define premalignant neurofibromatosis type 1–associated atypical neurofibromas. Neuro-Oncology, 2019, 21, 981-992.  | 1.2  | 69        |
| 14 | The <i>FOXA2</i> transcription factor is frequently somatically mutated in uterine carcinosarcomas and carcinomas. Cancer, 2018, 124, 65-73.   | 4.1  | 27        |
| 15 | The Sleep Inbred Panel, a Collection of Inbred <i>Drosophila melanogaster</i> with Extreme Long and Short Sleep Duration. G3: Genes, Genomes, Genetics, 2018, 8, 2865-2873.  | 1.8  | 16        |
| 16 | Somatic mutation profiles of clear cell endometrial tumors revealed by whole exome and targeted gene sequencing. Cancer, 2017, 123, 3261-3268.   | 4.1  | 72        |
| 17 | First insight into the somatic mutation burden of neurofibromatosis type 2-associated grade I and grade II meningiomas: a case report comprehensive genomic study of two cranial meningiomas with vastly different clinical presentation. BMC Cancer, 2017, 17, 127. | 2.6  | 13        |
| 18 | Selection for long and short sleep duration in Drosophila melanogaster reveals the complex genetic network underlying natural variation in sleep. PLoS Genetics, 2017, 13, e1007098.   | 3.5  | 43        |

| #  | Article  | IF   | CITATIONS |
|----|--|------|-----------|
| 19 | The transcription factors Ets1 and Sox10 interact during murine melanocyte development. Developmental Biology, 2015, 407, 300-312.   | 2.0  | 14        |
| 20 | Shimmer: detection of genetic alterations in tumors using next-generation sequence data. Bioinformatics, 2013, 29, 1498-1503.  | 4.1  | 59        |
| 21 | Exome sequencing of serous endometrial tumors identifies recurrent somatic mutations in chromatin-remodeling and ubiquitin ligase complex genes. Nature Genetics, 2012, 44, 1310-1315. | 21.4 | 365       |
| 22 | A Draft Sequence of the Neandertal Genome. Science, 2010, 328, 710-722.  | 12.6 | 3,588     |