# Lucia Susani

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/8793120/publications.pdf

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|---|---|---|---|
| 19 papers | 940 citations | 759055 12 h-index | 839398 18 g-index |
| 19 all docs | 19 docs citations | 19 times ranked | 1207 citing authors |

| # | Article | IF | Citations |
|---|---|---|---|
| 1 | Chromosome Transplantation: A Possible Approach to Treat Human X-linked Disorders. Molecular Therapy - Methods and Clinical Development, 2020, 17, 369-377. | 1.8 | 10 |
| 2 | Chromosome Transplantation: Correction of the Chronic Granulomatous Disease Defect in Mouse Induced Pluripotent Stem Cells. Stem Cells, 2019, 37, 876-887. | 1.4 | 3 |
| 3 | Synonymous Mutations Add a Layer of Complexity in the Diagnosis of Human Osteopetrosis. Journal of Bone and Mineral Research, 2017, 32, 99-105. | 3.1 | 11 |
| 4 | A pre-screening FISH-based method to detect CRISPR/Cas9 off-targets in mouse embryonic stem cells. Scientific Reports, 2015, 5, 12327. | 1.6 | 20 |
| 5 | Chromosome transplantation as a novel approach for correcting complex genomic disorders. Oncotarget, 2015, 6, 35218-35230. | 0.8 | 10 |
| 6 | As Little as Needed: The Extraordinary Case of a Mild Recessive Osteopetrosis Owing to a Novel Splicing Hypomorphic Mutation in the TCIRG1 Gene. Journal of Bone and Mineral Research, 2014, 29, 1646-1650. | 3.1 | 22 |
| 7 | A Homozygous Contiguous Gene Deletion in Chromosome 16p13.3 Leads to Autosomal Recessive Osteopetrosis in a Jordanian Patient. Calcified Tissue International, 2012, 91, 250-254. | 1.5 | 7 |
| 8 | Characterization of a Novel Alu-Alu Recombination-Mediated Genomic Deletion in the TCIRG1 Gene in Five Osteopetrotic Patients. Journal of Bone and Mineral Research, 2009, 24, 162-167. | 3.1 | 11 |
| 9 | Cornelia de Lange syndrome mutations in SMC1A or SMC3 affect binding to DNA. Human Molecular Genetics, 2009, 18, 418-427. | 1.4 | 92 |
| 10 | Cell fusion is a physiological process in mouse liver. Hepatology, 2008, 48, 1655-1664. | 3.6 | 29 |
| 11 | Osteoclast-poor human osteopetrosis due to mutations in the gene encoding RANKL. Nature Genetics, 2007, 39, 960-962. | 9.4 | 346 |
| 12 | TCIRG1-dependent recessive osteopetrosis: Mutation analysis, functional identification of the splicing defects, andin vitro rescue by U1 snRNA. Human Mutation, 2004, 24, 225-235. | 1.1 | 90 |
| 13 | Chloride Channel ClCN7 Mutations Are Responsible for Severe Recessive, Dominant, and Intermediate Osteopetrosis. Journal of Bone and Mineral Research, 2003, 18, 1740-1747. | 3.1 | 202 |
| 14 | Transcription Map of Xq27: Candidates for Several X-Linked Diseases. Genomics, 1999, 57, 209-218. | 1.3 | 18 |
| 15 | YAC/STS Map across 12 Mb of Xq27 at 25-kb Resolution, Merging Xq26â€“qter. Genomics, 1996, 34, 42-54. | 1.3 | 26 |
| 16 | The ZNF75 Zinc Finger Gene Subfamily: Isolation and Mapping of the Four Members in Humans and Great Apes. Genomics, 1996, 35, 312-320. | 1.3 | 12 |
| 17 | ZNF75: Isolation of a cDNA Clone of the KRAB Zinc Finger Gene Subfamily Mapped in YACs 1 Mb Telomeric of HPRT. Genomics, 1993, 18, 223-229. | 1.3 | 19 |
| 18 | Isolation of a zinc finger motif (ZNF75) mapping on chromosome Xq26. Genomics, 1992, 13, 1231-1236. | 1.3 | 12 |

| # | Article | IF | Citations |
|---|---|---|---|
| 19 | Fidelity of a YAC clone in the region of human MCF-2 gene. Biochemical and Biophysical Research Communications, 1991, 181, 877-883. | 1.0 | 0 |