Elisabeth Syk Lundberg

List of Publications by Year
in descending order

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9 papers	220 citations	1306789 7 h-index	1473754 9 g-index
9 all docs	9 docs citations	9 times ranked	548 citing authors

#	Article	IF	Citations
1	Whole-Genome Sequencing of Cytogenetically Balanced Chromosome Translocations Identifies Potentially Pathological Gene Disruptions and Highlights the Importance of Microhomology in the Mechanism of Formation. Human Mutation, 2017, 38, 180-192.	1.1	58
2	Comprehensive structural variation genome map of individuals carrying complex chromosomal rearrangements. PLoS Genetics, 2019, 15, e1007858.	1.5	36
3	CTNND2â€"a candidate gene for reading problems and mild intellectual disability. Journal of Medical Genetics, 2015, 52, 111-122.	1.5	35
4	Endometrial stromal cells exhibit a distinct phenotypic and immunomodulatory profile. Stem Cell Research and Therapy, 2020, 11, 15.	2.4	32
5	Replicative and non-replicative mechanisms in the formation of clustered CNVs are indicated by whole genome characterization. PLoS Genetics, 2018, 14, e1007780.	1.5	28
6	Detailed gene dose analysis reveals recurrent focal gene deletions in pediatric B-cell precursor acute lymphoblastic leukemia. Leukemia and Lymphoma, 2016, 57, 2161-2170.	0.6	10
7	Meiotic segregation analyses of reciprocal translocations in spermatozoa and embryos: no support for predictive value regarding PGD outcome. Reproductive BioMedicine Online, 2017, 34, 645-652.	1.1	8
8	Ataxia in Patients With Bi-Allelic NFASC Mutations and Absence of Full-Length NF186. Frontiers in Genetics, 2019, 10, 896.	1.1	7
9	Flanking complex copy number variants in the same family formed through unequal crossing-over during meiosis. Mutation Research - Fundamental and Molecular Mechanisms of Mutagenesis, 2018, 812, 1-4.	0.4	6