Sanna-Maria Karppinen

List of Publications by Year in descending order

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18 papers	1,193 citations	623734 14 h-index	839539 18 g-index
19 all docs	19 docs citations	19 times ranked	2371 citing authors

#	Article	IF	Citations
1	Deletion of Col15a1 Modulates the Tumour Extracellular Matrix and Leads to Increased Tumour Growth in the MMTV-PyMT Mouse Mammary Carcinoma Model. International Journal of Molecular Sciences, 2021, 22, 9978.	4.1	8
2	Exploring the roles of MACIT and multiplexin collagens in stem cells and cancer. Seminars in Cancer Biology, 2020, 62, 134-148.	9.6	23
3	Collagen XV, a multifaceted multiplexin present across tissues and species. Matrix Biology Plus, 2020, 6-7, 100023.	3.5	29
4	Lack of collagen XVIII leads to lipodystrophy and perturbs hepatic glucose and lipid homeostasis. Journal of Physiology, 2020, 598, 3373-3393.	2.9	14
5	Toward understanding scarless skin wound healing and pathological scarring. F1000Research, 2019, 8, 787.	1.6	125
6	GATA4-dependent organ-specific endothelial differentiation controls liver development and embryonic hematopoiesis. Journal of Clinical Investigation, 2017, 127, 1099-1114.	8.2	102
7	Continuous Grading of Early Fibrosis in NAFLD Using Label-Free Imaging: A Proof-of-Concept Study. PLoS ONE, 2016, 11, e0147804.	2.5	34
8	Collagens XV and XVIII show different expression and localisation in cutaneous squamous cell carcinoma: type XV appears in tumor stroma, while XVIII becomes upregulated in tumor cells and lost from microvessels. Experimental Dermatology, 2016, 25, 348-354.	2.9	30
9	Germline alterations in the CLSPN gene in breast cancer families. Cancer Letters, 2008, 261, 93-97.	7.2	15
10	Germline alterations in the 53BP1 gene in breast and ovarian cancer families. Cancer Letters, 2007, 245, 337-340.	7.2	12
11	A recurrent mutation in PALB2 in Finnish cancer families. Nature, 2007, 446, 316-319.	27.8	402
12	Evaluation of the role of Finnish ataxia-telangiectasia mutations in hereditary predisposition to breast cancer. Carcinogenesis, 2006, 28, 1040-1045.	2.8	21
13	Screening for RAD51 and BRCA2 BRC repeat mutations in breast and ovarian cancer families. Cancer Letters, 2006, 236, 142-147.	7.2	15
14	Identification of a common polymorphism in the TopBP1 gene associated with hereditary susceptibility to breast and ovarian cancer. European Journal of Cancer, 2006, 42, 2647-2652.	2.8	37
15	Association of commonATM polymorphism with bilateral breast cancer. International Journal of Cancer, 2005, 116, 69-72.	5.1	42
16	RAD50 and NBS1 are breast cancer susceptibility genes associated with genomic instability. Carcinogenesis, 2005, 27, 1593-1599.	2.8	179
17	Mutation analysis of the ATRgene in breast and ovarian cancer families. Breast Cancer Research, 2005, 7, R495-501.	5.0	30
18	Analysis of 11q21-24 loss of heterozygosity candidate target genes in breast cancer: Indications of TSLC1 promoter hypermethylation. Genes Chromosomes and Cancer, 2002, 34, 384-389.	2.8	74