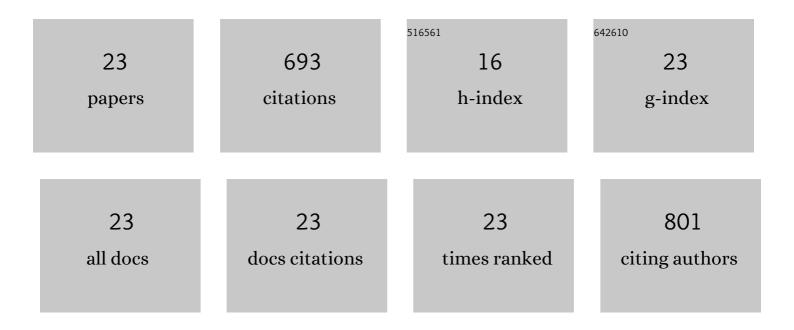
Semi Kim

List of Publications by Year in descending order

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SEMIKIM

#	Article	IF	CITATIONS
1	Tetraspanin TM4SF5 mediates loss of contact inhibition through epithelial-mesenchymal transition in human hepatocarcinoma. Journal of Clinical Investigation, 2008, 118, 1354-1366.	3.9	103
2	TMPRSS4 induces invasion and epithelial-mesenchymal transition through upregulation of integrin Â5 and its signaling pathways. Carcinogenesis, 2010, 31, 597-606.	1.3	96
3	Cooperation between integrin α5 and tetraspan TM4SF5 regulates VEGF-mediated angiogenic activity. Blood, 2009, 113, 1845-1855.	0.6	56
4	Transmembrane 4ÂL Six Family Member 5 Senses Arginine for mTORC1 Signaling. Cell Metabolism, 2019, 29, 1306-1319.e7.	7.2	50
5	Twist1 and AP-1 cooperatively upregulate integrin α5 expression to induce invasion and the epithelial–mesenchymal transition. Carcinogenesis, 2015, 36, 327-337.	1.3	47
6	TMPRSS4 upregulates uPA gene expression through JNK signaling activation to induce cancer cell invasion. Cellular Signalling, 2014, 26, 398-408.	1.7	32
7	TMPRSS4 induces invasion and proliferation of prostate cancer cells through induction of Slug and cyclin D1. Oncotarget, 2016, 7, 50315-50332.	0.8	31
8	TMPRSS4 promotes cancer stem–like properties in prostate cancer cells through upregulation of SOX2 by SLUG and TWIST1. Journal of Experimental and Clinical Cancer Research, 2021, 40, 372.	3.5	31
9	TMPRSS4 induces cancer cell invasion through pro-uPA processing. Biochemical and Biophysical Research Communications, 2014, 446, 1-7.	1.0	29
10	The extracellular loop 2 of TM4SF5 inhibits integrin $\hat{1}\pm 2$ on hepatocytes under collagen type I environment. Carcinogenesis, 2009, 30, 1872-1879.	1.3	25
11	Membrane Proteins Involved in Epithelial-Mesenchymal Transition and Tumor Invasion: Studies on TMPRSS4 and TM4SF5. Genomics and Informatics, 2014, 12, 12.	0.4	25
12	Anti-cancer activity of the novel 2-hydroxydiarylamide derivatives IMD-0354 and KRT1853 through suppression of cancer cell invasion, proliferation, and survival mediated by TMPRSS4. Scientific Reports, 2019, 9, 10003.	1.6	22
13	A New Nano-Platform of Erythromycin Combined with Ag Nano-Particle ZnO Nano-Structure against Methicillin-Resistant Staphylococcus aureus. Pharmaceutics, 2020, 12, 841.	2.0	21
14	Differential <scp>TM4SF5</scp> â€mediated <scp>SIRT1</scp> modulation and metabolic signaling in nonalcoholic steatohepatitis progression. Journal of Pathology, 2021, 253, 55-67.	2.1	20
15	Anti-cancer Activity of Novel TM4SF5-Targeting Antibodies through TM4SF5 Neutralization and Immune Cell-Mediated Cytotoxicity. Theranostics, 2017, 7, 594-613.	4.6	19
16	Preparation of siRNA encapsulated nanoliposomes suitable for siRNA delivery by simply discontinuous mixing. Biochimica Et Biophysica Acta - Biomembranes, 2018, 1860, 1318-1325.	1.4	16
17	Cooperation between ZEB2 and Sp1 promotes cancer cell survival and angiogenesis during metastasis through induction of survivin and VEGF. Oncotarget, 2018, 9, 726-742.	0.8	16
18	A New Surface Charge Neutralizing Nano-Adjuvant to Potentiate Polymyxins in Killing Mcr-1 Mediated Drug-Resistant Escherichia coli. Pharmaceutics, 2021, 13, 250.	2.0	15

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#	Article	IF	CITATIONS
19	TM4SF5-mediated liver malignancy involves NK cell exhaustion-like phenotypes. Cellular and Molecular Life Sciences, 2022, 79, 1.	2.4	13
20	A nontoxic biocompatible nanocomposite comprising black phosphorus with Au–γ-Fe ₂ O ₃ nanoparticles. RSC Advances, 2020, 10, 16162-16167.	1.7	9
21	TMEM52B suppression promotes cancer cell survival and invasion through modulating E-cadherin stability and EGFR activity. Journal of Experimental and Clinical Cancer Research, 2021, 40, 58.	3.5	8
22	Bovine Serum Albumin-Immobilized Black Phosphorus-Based γ-Fe2O3 Nanocomposites: A Promising Biocompatible Nanoplatform. Biomedicines, 2021, 9, 858.	1.4	6
23	Therapeutic effects of TM4SF5-targeting chimeric and humanized monoclonal antibodies in hepatocellular and colon cancer models. Molecular Therapy - Oncolytics, 2022, 24, 452-466.	2.0	3