

Dan Deng

List of Publications by Year in descending order

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Version: 2024-02-01

15
papers

343
citations

1307594

7
h-index

996975

15
g-index

16
all docs

16
docs citations

16
times ranked

572
citing authors

#	ARTICLE	IF	CITATIONS
1	The clinical efficacy and safety of anti-IL17E therapy in recessive dystrophic epidermolysis bullosa. <i>Clinical Genetics</i> , 2022, 101, 110-115.	2.0	5
2	Benzothiazole-decorated iridium-based nanophotosensitizers for photodynamic therapy of cancer cells. <i>Dalton Transactions</i> , 2022, 51, 3666-3675.	3.3	7
3	Crisaborole: an emerging therapy for prurigo pigmentosa. <i>Journal of the European Academy of Dermatology and Venereology</i> , 2022, 36, .	2.4	4
4	Detection and characterization of low-level mosaicism among clinically unaffected parents of sporadic epidermolysis bullosa simplex cases. <i>British Journal of Dermatology</i> , 2022, 187, 441-443.	1.5	2
5	Differential susceptibility to SARS-CoV-2 in the normal nasal mucosa and in chronic sinusitis. <i>European Journal of Immunology</i> , 2022, , .	2.9	3
6	The first case of Chinese phacomatosis pigmentokeratolica diagnosed by a missense HRAS mosaicism. <i>Journal of Dermatology</i> , 2022, 49, 921-924.	1.2	4
7	Roles of the H19/microRNA-675 axis in the proliferation and epithelial-mesenchymal transition of human cutaneous squamous cell carcinoma cells. <i>Oncology Reports</i> , 2021, 45, .	2.6	16
8	Cutaneous metastases from triple primary extramammary Paget's disease. <i>JDDG - Journal of the German Society of Dermatology</i> , 2020, 18, 1169-1172.	0.8	3
9	Next-generation sequencing through multigene panel testing for the diagnosis of hereditary epidermolysis bullosa in Chinese population. <i>Clinical Genetics</i> , 2020, 98, 179-184.	2.0	10
10	CHILD syndrome mimicking verrucous nevus in a Chinese patient responded well to the topical therapy of compound of simvastatin and cholesterol. <i>Journal of the European Academy of Dermatology and Venereology</i> , 2018, 32, 1209-1213.	2.4	11
11	Comparison of Autologous, Allogeneic, and Cell-Free Scaffold Approaches for Engineered Tendon Repair in a Rabbit Model-A Pilot Study. <i>Tissue Engineering - Part A</i> , 2017, 23, 750-761.	3.1	21
12	A novel mutation in TRPV3 gene causes atypical familial Olmsted syndrome. <i>Scientific Reports</i> , 2016, 6, 21815.	3.3	36
13	Less is more: New biomimetic approach to control spatial and temporal cell loading for tissue engineering. <i>Journal of Biomedical Materials Research - Part A</i> , 2014, 102, 4108-4117.	4.0	3
14	Repair of Achilles tendon defect with autologous ASCs engineered tendon in a rabbit model. <i>Biomaterials</i> , 2014, 35, 8801-8809.	11.4	99
15	Engineering human neo-tendon tissue in vitro with human dermal fibroblasts under static mechanical strain. <i>Biomaterials</i> , 2009, 30, 6724-6730.	11.4	118