Adele Mucci

List of Publications by Year
in descending order

13	500	933447 10	1125743 13
papers	citations	h-index	g-index

13	13	13	737
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Restored Macrophage Function Ameliorates Disease Pathophysiology in a Mouse Model for IL10 Receptor-deficient Very Early Onset Inflammatory Bowel Disease. Journal of Crohn's and Colitis, 2021, 15, 1588-1595.	1.3	10
2	Myeloid cellâ€based delivery of IFNâ€Î³ reprograms the leukemia microenvironment and induces antiâ€tumoral immune responses. EMBO Molecular Medicine, 2021, 13, e13598.	6.9	13
3	Effective hematopoietic stem cell-based gene therapy in a murine model of hereditary pulmonary alveolar proteinosis. Haematologica, 2020, 105, 1147-1157.	3.5	7
4	Pulmonary Transplantation of Human Induced Pluripotent Stem Cellâ€"derived Macrophages Ameliorates Pulmonary Alveolar Proteinosis. American Journal of Respiratory and Critical Care Medicine, 2018, 198, 350-360.	5.6	57
5	Impaired IFNÎ³-Signaling and Mycobacterial Clearance in IFNÎ³R1-Deficient Human iPSC-Derived Macrophages. Stem Cell Reports, 2018, 10, 7-16.	4.8	25
6	Hematopoietic stem cell gene therapy for IFNÎ³R1 deficiency protects mice from mycobacterial infections. Blood, 2018, 131, 533-545.	1.4	19
7	iPSC-Derived Macrophages Effectively Treat Pulmonary Alveolar Proteinosis in Csf2rb-Deficient Mice. Stem Cell Reports, 2018, 11, 696-710.	4.8	40
8	Human Effector Memory T Helper Cells Engage with Mouse Macrophages and Cause Graft-versus-Hostâ€"Like Pathology in Skin of Humanized Mice Used in a Nonclinical Immunization Study. American Journal of Pathology, 2017, 187, 1380-1398.	3.8	23
9	Lung surfactant metabolism: early in life, early in disease and target in cell therapy. Cell and Tissue Research, 2017, 367, 721-735.	2.9	50
10	Murine iPSC-Derived Macrophages as a Tool for Disease Modeling of Hereditary Pulmonary Alveolar Proteinosis due to Csf2rb Deficiency. Stem Cell Reports, 2016, 7, 292-305.	4.8	23
11	Gene Correction of Human Induced Pluripotent Stem Cells Repairs the Cellular Phenotype in Pulmonary Alveolar Proteinosis. American Journal of Respiratory and Critical Care Medicine, 2014, 189, 167-182.	5.6	85
12	Pulmonary transplantation of macrophage progenitors as effective and long-lasting therapy for hereditary pulmonary alveolar proteinosis. Science Translational Medicine, 2014, 6, 250ra113.	12.4	106
13	Promoter and lineage independent anti-silencing activity of the A2 ubiquitous chromatin opening element for optimized human pluripotent stem cell-based gene therapy. Biomaterials, 2014, 35, 1531-1542.	11.4	42