Takuji Masunaga

List of Publications by Year
in descending order

9 papers	149 citations	1478505 6 h-index	1474206 9 g-index
9 all docs	9 docs citations	9 times ranked	180 citing authors

#	Article	IF	Citations
1	Splice site mutation in <scp>COL</scp>7A1 resulting in aberrant inâ€frame transcripts identified in a case of recessive dystrophic epidermolysis bullosa, pretibial. Journal of Dermatology, 2018, 45, 742-745.	1.2	4
2	Compound heterozygosity for novel splice site mutations of ITGA6 in lethal junctional epidermolysis bullosa with pyloric atresia. Journal of Dermatology, 2017, 44, 160-166.	1.2	9
3	Japanese recurrent mutation c.6216+5G>T in COL7A1 leads to a mild phenotype of dystrophic epidermolysis bullosa. Journal of Dermatological Science, 2015, 80, 220-223.	1.9	2
4	Splicing abnormality of integrin Î²4 gene (ITGB4) due to nucleotide substitutions far from splice site underlies pyloric atresia-junctional epidermolysis bullosa syndrome. Journal of Dermatological Science, 2015, 78, 61-66.	1.9	6
5	Genotypeâ€“phenotype correlations in six Japanese patients with recessive dystrophic epidermolysis bullosa with the recurrent p.Glu2857X mutation. Journal of Dermatological Science, 2008, 52, 13-20.	1.9	12
6	Epidermal Basement Membrane: Its Molecular Organization and Blistering Disorders. Connective Tissue Research, 2006, 47, 55-66.	2.3	38
7	Pyloric atresia-junctional epidermolysis bullosa syndrome showing novel 594insC/Q425P mutations in integrin beta4 gene (ITGB4). Experimental Dermatology, 2004, 13, 61-64.	2.9	13
8	Differences in recurrent COL7A1 mutations in dystrophic epidermolysis bullosa: ethnic-specific and worldwide recurrent mutations. Archives of Dermatological Research, 2004, 295, 442-447.	1.9	33
9	Recurrent COL7A1 Mutations in Japanese Patients with Dystrophic Epidermolysis Bullosa: Positional Effects of Premature Termination Codon Mutations on Clinical Severity. Journal of Investigative Dermatology, 1999, 112, 991-993.	0.7	32