

The classification of inherited epidermolysis bullosa (EB)
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Citation Report

#	ARTICLE	IF	CITATIONS
2	Epidermolysis Bullosa: Prospects for Cell-Based Therapies. <i>Journal of Investigative Dermatology</i> , 2008, 128, 2140-2142.	0.3	24
4	Dystrophic epidermolysis bullosa pruriginosa is not associated with frequentFLGgene mutations. <i>British Journal of Dermatology</i> , 2008, 159, 464-469.	1.4	31
5	Classifying epidermolysis bullosa. <i>Journal of the American Academy of Dermatology</i> , 2008, 59, 1075-1076.	0.6	4
7	Update on diagnosis and therapy of inherited epidermolysis bullosa. <i>Expert Review of Dermatology</i> , 2008, 3, 721-733.	0.3	10
8	Epidermolysis Bullosa: The Pediatricians Role. <i>Current Pediatric Reviews</i> , 2008, 4, 250-257.	0.4	1
9	Epidermolysis bullosa simplex: a paradigm for disorders of tissue fragility. <i>Journal of Clinical Investigation</i> , 2009, 119, 1784-1793.	3.9	174
10	Dominant-negative Effects of COL7A1 Mutations Can be Rescued by Controlled Overexpression of Normal Collagen VII. <i>Journal of Biological Chemistry</i> , 2009, 284, 30248-30256.	1.6	40
11	Cetuximab Therapy of Metastasizing Cutaneous Squamous Cell Carcinoma in a Patient with Severe Recessive Dystrophic Epidermolysis Bullosa. <i>Dermatology</i> , 2009, 219, 80-83.	0.9	72
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17	Progress in Heritable Skin Diseases: Translational Implications of Mutation Analysis and Prospects of Molecular Therapies*. <i>Acta Dermato-Venereologica</i> , 2009, 89, 228-235.	0.6	27
18	Genodermatoses. <i>Medicine</i> , 2009, 37, 298-302.	0.2	0
19	Connective Tissue and Related Disorders and Preterm Birth: Clues to Genes Contributing to Prematurity. <i>Placenta</i> , 2009, 30, 207-215.	0.7	93
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21	Severe keratin 5 and 14 mutations induce down-regulation of junction proteins in keratinocytes. <i>Experimental Cell Research</i> , 2009, 315, 2995-3003.	1.2	50

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165	A novel KRT5 mutation, p.Lys199Asn, is associated with three subtypes of epidermolysis bullosa simplex phenotypes in a single Chinese family. <i>Journal of Dermatological Science</i> , 2011, 64, 241-243.	1.0	1
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