

# David T Woodley

## List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

69  
papers

6,055  
citations

40  
h-index

74  
g-index

74  
ext. papers

6,743  
ext. citations

7.7  
avg, IF

5.13  
L-index

#	Paper	IF	Citations
69	Evaluation of Systemic Gentamicin as Translational Readthrough Therapy for a Patient With Epidermolysis Bullosa Simplex With Muscular Dystrophy Owing to PLEC1 Pathogenic Nonsense Variants.. <i>JAMA Dermatology</i> , <b>2022</b> ,	5.1	2
68	Immunoglobulin and Complement Immunohistochemistry on Paraffin Sections in Autoimmune Bullous Diseases: A Systematic Review and Meta-analysis. <i>American Journal of Dermatopathology</i> , <b>2021</b> , 43, 689-699	0.9	2
67	Association between vaccination and autoimmune bullous diseases: A systematic review. <i>Journal of the American Academy of Dermatology</i> , <b>2021</b> ,	4.5	7
66	Over-expression of stromal periostin correlates with poor prognosis of cutaneous squamous cell carcinomas. <i>Experimental Dermatology</i> , <b>2021</b> , 30, 698-704	4	4
65	Characterization of mutant type VII collagens underlying the inversa subtype of recessive dystrophic epidermolysis bullosa. <i>Journal of Dermatological Science</i> , <b>2021</b> , 104, 104-111	4.3	0
64	Multidisciplinary care of epidermolysis bullosa during the COVID-19 pandemic-Consensus: Recommendations by an international panel of experts. <i>Journal of the American Academy of Dermatology</i> , <b>2020</b> , 83, 1222-1224	4.5	5
63	Gentamicin Induces Laminin 332 and Improves Wound Healing in Junctional Epidermolysis Bullosa Patients with Nonsense Mutations. <i>Molecular Therapy</i> , <b>2020</b> , 28, 1327-1338	11.7	13
62	Clinical outcomes of patients with pemphigus treated by the same physicians in a public safety net healthcare system vs. a private healthcare system. <i>British Journal of Dermatology</i> , <b>2019</b> , 181, 850-852	4	
61	Improving Hairdressers' Knowledge and Self-efficacy to Detect Scalp and Neck Melanoma by Use of an Educational Video. <i>JAMA Dermatology</i> , <b>2018</b> , 154, 214-216	5.1	6
60	Gentamicin induces nonsense mutation readthrough and restores functional laminin 332 in junctional epidermolysis bullosa. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , <b>2018</b> , 115, E6536-E6545	11.5	32
59	Distinct Fibroblasts in the Papillary and Reticular Dermis: Implications for Wound Healing. <i>Dermatologic Clinics</i> , <b>2017</b> , 35, 95-100	4.2	53
58	Gentamicin induces functional type VII collagen in recessive dystrophic epidermolysis bullosa patients. <i>Journal of Clinical Investigation</i> , <b>2017</b> , 127, 3028-3038	15.9	61
57	Keratinocyte Migration and a Hypothetical New Role for Extracellular Heat Shock Protein 90 Alpha in Orchestrating Skin Wound Healing. <i>Advances in Wound Care</i> , <b>2015</b> , 4, 203-212	4.8	24
56	Intravenously Administered Recombinant Human Type VII Collagen Derived from Chinese Hamster Ovary Cells Reverses the Disease Phenotype in Recessive Dystrophic Epidermolysis Bullosa Mice. <i>Journal of Investigative Dermatology</i> , <b>2015</b> , 135, 3060-3067	4.3	19
55	Reduced Toxicity Conditioning and Allogeneic Hematopoietic Progenitor Cell Transplantation for Recessive Dystrophic Epidermolysis Bullosa. <i>Journal of Pediatrics</i> , <b>2015</b> , 167, 765-9.e1	3.6	21
54	Definitions and outcome measures for mucous membrane pemphigoid: recommendations of an international panel of experts. <i>Journal of the American Academy of Dermatology</i> , <b>2015</b> , 72, 168-74	4.5	93
53	Aminoglycosides restore full-length type VII collagen by overcoming premature termination codons: therapeutic implications for dystrophic epidermolysis bullosa. <i>Molecular Therapy</i> , <b>2014</b> , 22, 1741-52	11.7	44

52	Omalizumab therapy for bullous pemphigoid. <i>Journal of the American Academy of Dermatology</i> , <b>2014</b> , 71, 468-74	4.5	109
51	De novo anti-type VII collagen antibodies in patients with recessive dystrophic epidermolysis bullosa. <i>Journal of Investigative Dermatology</i> , <b>2014</b> , 134, 1138-1140	4.3	30
50	Identification of the critical therapeutic entity in secreted Hsp90 that promotes wound healing in newly re-standardized healthy and diabetic pig models. <i>PLoS ONE</i> , <b>2014</b> , 9, e113956	3.7	24
49	Topical application of recombinant type VII collagen incorporates into the dermal-epidermal junction and promotes wound closure. <i>Molecular Therapy</i> , <b>2013</b> , 21, 1335-44	11.7	44
48	Epidermolysis bullosa acquisita: autoimmunity to anchoring fibril collagen. <i>Autoimmunity</i> , <b>2012</b> , 45, 91-101		49
47	Epidermolysis bullosa acquisita. <i>Clinics in Dermatology</i> , <b>2012</b> , 30, 60-9	3	119
46	Successful treatment of bullous pemphigoid with omalizumab. <i>Archives of Dermatology</i> , <b>2012</b> , 148, 1241-3		42
45	Epidermolysis Bullosa Acquisita <b>2011</b> , 113-136		
44	Bone marrow transplantation for recessive dystrophic epidermolysis bullosa. <i>New England Journal of Medicine</i> , <b>2010</b> , 363, 629-39	59.2	262
43	Injection of recombinant human type VII collagen corrects the disease phenotype in a murine model of dystrophic epidermolysis bullosa. <i>Molecular Therapy</i> , <b>2009</b> , 17, 26-33	11.7	107
42	TANGO1 facilitates cargo loading at endoplasmic reticulum exit sites. <i>Cell</i> , <b>2009</b> , 136, 891-902	56.2	254
41	151A Traffic Control Role for TGF-Beta in Skin Cell Motility During Wound Healing. <i>Wound Repair and Regeneration</i> , <b>2008</b> , 13, A28-A48	3.6	
40	The classification of inherited epidermolysis bullosa (EB): Report of the Third International Consensus Meeting on Diagnosis and Classification of EB. <i>Journal of the American Academy of Dermatology</i> , <b>2008</b> , 58, 931-50	4.5	690
39	Characterization of molecular mechanisms underlying mutations in dystrophic epidermolysis bullosa using site-directed mutagenesis. <i>Journal of Biological Chemistry</i> , <b>2008</b> , 283, 17838-45	5.4	26
38	Duplicating autoimmune bullous diseases by passively transferring autoantibodies into animals. <i>Journal of Investigative Dermatology</i> , <b>2008</b> , 128, E25-7	4.3	1
37	Intravenously injected human fibroblasts home to skin wounds, deliver type VII collagen, and promote wound healing. <i>Molecular Therapy</i> , <b>2007</b> , 15, 628-35	11.7	88
36	Autoimmunity to type VII collagen: epidermolysis bullosa acquisita. <i>Clinical Reviews in Allergy and Immunology</i> , <b>2007</b> , 33, 78-84	12.3	42
35	The role of IgE anti-basement membrane zone autoantibodies in bullous pemphigoid. <i>Archives of Dermatology</i> , <b>2007</b> , 143, 249-50		7

34	The cartilage matrix protein subdomain of type VII collagen is pathogenic for epidermolysis bullosa acquisita. <i>American Journal of Pathology</i> , <b>2007</b> , 170, 2009-18	5.8	53
33	Induction of epidermolysis bullosa acquisita in mice by passive transfer of autoantibodies from patients. <i>Journal of Investigative Dermatology</i> , <b>2006</b> , 126, 1323-30	4.3	80
32	Evidence that anti-type VII collagen antibodies are pathogenic and responsible for the clinical, histological, and immunological features of epidermolysis bullosa acquisita. <i>Journal of Investigative Dermatology</i> , <b>2005</b> , 124, 958-64	4.3	88
31	Injection of recombinant human type VII collagen restores collagen function in dystrophic epidermolysis bullosa. <i>Nature Medicine</i> , <b>2004</b> , 10, 693-5	50.5	111
30	Intradermal injection of lentiviral vectors corrects regenerated human dystrophic epidermolysis bullosa skin tissue in vivo. <i>Molecular Therapy</i> , <b>2004</b> , 10, 318-26	11.7	70
29	Epidermolysis bullosa: then and now. <i>Journal of the American Academy of Dermatology</i> , <b>2004</b> , 51, S55-7	4.5	8
28	Normal and gene-corrected dystrophic epidermolysis bullosa fibroblasts alone can produce type VII collagen at the basement membrane zone. <i>Journal of Investigative Dermatology</i> , <b>2003</b> , 121, 1021-8	4.3	93
27	Restoration of type VII collagen expression and function in dystrophic epidermolysis bullosa. <i>Nature Genetics</i> , <b>2002</b> , 32, 670-5	36.3	144
26	The recombinant expression of full-length type VII collagen and characterization of molecular mechanisms underlying dystrophic epidermolysis bullosa. <i>Journal of Biological Chemistry</i> , <b>2002</b> , 277, 2118-24	5.4	59
25	The first international consensus on mucous membrane pemphigoid: definition, diagnostic criteria, pathogenic factors, medical treatment, and prognostic indicators. <i>Archives of Dermatology</i> , <b>2002</b> , 138, 370-9		526
24	The carboxyl terminus of type VII collagen mediates antiparallel dimer formation and constitutes a new antigenic epitope for epidermolysis Bullosa acquisita autoantibodies. <i>Journal of Biological Chemistry</i> , <b>2001</b> , 276, 21649-55	5.4	58
23	Epidermolysis bullosa acquisita: update and review. <i>Clinics in Dermatology</i> , <b>2001</b> , 19, 712-8	3	43
22	Autoantibodies to type VII collagen have heterogeneous subclass and light chain compositions and their complement-activating capacities do not correlate with the inflammatory clinical phenotype. <i>Journal of Clinical Immunology</i> , <b>2000</b> , 20, 416-23	5.7	18
21	NC1 domain of type VII collagen binds to the beta3 chain of laminin 5 via a unique subdomain within the fibronectin-like repeats. <i>Journal of Investigative Dermatology</i> , <b>1999</b> , 112, 177-83	4.3	92
20	Alpha 2 beta 1 integrin mediates dermal fibroblast attachment to type VII collagen via a 158-amino-acid segment of the NC1 domain. <i>Experimental Cell Research</i> , <b>1999</b> , 249, 231-9	4.2	27
19	Epitope spreading: lessons from autoimmune skin diseases. <i>Journal of Investigative Dermatology</i> , <b>1998</b> , 110, 103-9	4.3	249
18	Interactions of the amino-terminal noncollagenous (NC1) domain of type VII collagen with extracellular matrix components. A potential role in epidermal-dermal adherence in human skin. <i>Journal of Biological Chemistry</i> , <b>1997</b> , 272, 14516-22	5.4	143
17	Development of an ELISA for rapid detection of anti-type VII collagen autoantibodies in epidermolysis bullosa acquisita. <i>Journal of Investigative Dermatology</i> , <b>1997</b> , 108, 68-72	4.3	107

16	Ultraviolet A irradiation upregulates type VII collagen expression in human dermal fibroblasts. <i>Journal of Investigative Dermatology</i> , <b>1997</b> , 108, 125-8	4-3	34
15	Colchicine for epidermolysis bullosa acquisita. <i>Journal of the American Academy of Dermatology</i> , <b>1996</b> , 34, 781-4	4-5	85
14	Type VII collagen specifically binds fibronectin via a unique subdomain within the collagenous triple helix. <i>Journal of Investigative Dermatology</i> , <b>1994</b> , 103, 637-41	4-3	29
13	Epidermal growth factor (EGF) promotes human keratinocyte locomotion on collagen by increasing the alpha 2 integrin subunit. <i>Experimental Cell Research</i> , <b>1993</b> , 209, 216-23	4-2	146
12	Labeling of fractured human skin with antibodies to BM 600/nicein, epiligrin, kalinin and other matrix components. <i>Journal of Dermatological Science</i> , <b>1993</b> , 5, 97-103	4-3	11
11	Identification and partial characterization of a novel 105-kDalton lower lamina lucida autoantigen associated with a novel immune-mediated subepidermal blistering disease. <i>Journal of Investigative Dermatology</i> , <b>1993</b> , 101, 262-7	4-3	69
10	Immunofluorescence on Salt-Split Skin for the Diagnosis of Epidermolysis Bullosa Acquisita. <i>Archives of Dermatology</i> , <b>1990</b> , 126, 229		24
9	Laminin inhibits human keratinocyte migration. <i>Journal of Cellular Physiology</i> , <b>1988</b> , 136, 140-6	7	147
8	Increased frequency of HLA-DR2 in patients with autoantibodies to epidermolysis bullosa acquisita antigen: evidence that the expression of autoimmunity to type VII collagen is HLA class II allele associated. <i>Journal of Investigative Dermatology</i> , <b>1988</b> , 91, 228-32	4-3	125
7	Clearing of epidermolysis bullosa acquisita with cyclosporine. <i>Journal of the American Academy of Dermatology</i> , <b>1988</b> , 19, 937-42	4-5	89
6	Organ-specific, phylogenetic, and ontogenetic distribution of the epidermolysis bullosa acquisita antigen. <i>Journal of Investigative Dermatology</i> , <b>1986</b> , 86, 376-9	4-3	37
5	Epidermolysis bullosa acquisita antigen, a major cutaneous basement membrane component, is synthesized by human dermal fibroblasts and other cutaneous tissues. <i>Journal of Investigative Dermatology</i> , <b>1986</b> , 87, 227-31	4-3	27
4	Evidence that anti-basement membrane zone antibodies in bullous eruption of systemic lupus erythematosus recognize epidermolysis bullosa acquisita autoantigen. <i>Journal of Investigative Dermatology</i> , <b>1985</b> , 84, 472-6	4-3	135
3	Identification of the skin basement-membrane autoantigen in epidermolysis bullosa acquisita. <i>New England Journal of Medicine</i> , <b>1984</b> , 310, 1007-13	59-2	447
2	Epidermolysis bullosa acquisita--a pemphigoid-like disease. <i>Journal of the American Academy of Dermatology</i> , <b>1984</b> , 11, 820-32	4-5	220
1	Localization of basement membrane components after dermal-epidermal junction separation. <i>Journal of Investigative Dermatology</i> , <b>1983</b> , 81, 149-53	4-3	161