

# CITATION REPORT

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## Inherited epidermolysis bullosa: updated recommendations on diagnosis and classification

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#	Paper	IF	Citations
686	Er:YAG Laser Dental Treatment of Patients Affected by Epidermolysis Bullosa. <b>2014</b> , 2014, 421783		
685	Frequent Occurrence of Aplasia Cutis Congenita in Bullous Dermolysis of the Newborn. <b>2016</b> , 96, 784-7		4
684	Pain care for patients with epidermolysis bullosa: best care practice guidelines. <b>2014</b> , 12, 178		53
683	[Parallels between wound healing, chronic inflammatory skin diseases and neoplasia: clinical aspects]. <b>2014</b> , 65, 934-43		2
682	Small-molecule therapies for genetic skin fragility. <b>2014</b> , 22, 1724-5		2
681	Real-time PCR detection of the recessive dystrophic epidermolysis bullosa-associated c.2470insG mutation in unrelated Mexican families. <b>2014</b> , 45, 596-9		1
680	Novel KRT14 mutation causing epidermolysis bullosa simplex with variable phenotype. <i>Experimental Dermatology</i> , <b>2014</b> , 23, 684-7	4	7
679	Induced pluripotent stem cells from human revertant keratinocytes for the treatment of epidermolysis bullosa. <b>2014</b> , 6, 264ra164		95
678	Human COL7A1-corrected induced pluripotent stem cells for the treatment of recessive dystrophic epidermolysis bullosa. <b>2014</b> , 6, 264ra163		157
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669	Prevalence of anemia in patients with epidermolysis bullosa registered in Australia. <b>2015</b> , 1, 37-40		14
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497	Skin Cancer: Genetics, Immunology, Treatments, and Psychological Care. <b>2017</b> , 851-934		8
496	Loss of interaction between plectin and type XVII collagen results in epidermolysis bullosa simplex. <i>Human Mutation</i> , <b>2017</b> , 38, 1666-1670	4-7	6
495	Cut and Paste: Efficient Homology-Directed Repair of a Dominant Negative KRT14 Mutation via CRISPR/Cas9 Nickases. <b>2017</b> , 25, 2585-2598		49
494	Wound healing in epidermolysis bullosa. <i>British Journal of Dermatology</i> , <b>2017</b> , 177, e193-e195	4	6
493	Diagnosis of Inherited Epidermolysis Bullosa in Resource-Limited Settings: Immunohistochemistry Revisited. <b>2017</b> , 233, 326-332		5
492	Regeneration of the entire human epidermis using transgenic stem cells. <b>2017</b> , 551, 327-332		379
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490	Indikationserweiterungen von Rhabarberwurzel und Vitamin D. <b>2017</b> , 16, 6-7		

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488	Pain and quality of life evaluation in patients with localized epidermolysis bullosa simplex. <i>Orphanet Journal of Rare Diseases</i> , <b>2017</b> , 12, 119	4.2	15
487	Persistent elevation of serum interleukin-6 and serum amyloid A levels in patients with recessive dystrophic epidermolysis bullosa. <b>2017</b> , 27, 80-81		1
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485	Closure of a Large Chronic Wound through Transplantation of Gene-Corrected Epidermal Stem Cells. <b>2017</b> , 137, 778-781		75
484	The first familial cases of epidermolysis bullosa simplex, generalized severe with p.Asn176Ser in KRT5 revealing the clinical chronology. <b>2017</b> , 31, e251-e253		
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481	Molecular pathology of the basement membrane zone in heritable blistering diseases: The paradigm of epidermolysis bullosa. <i>Matrix Biology</i> , <b>2017</b> , 57-58, 76-85	11.4	44
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477	Focal adhesions in the skin: lessons learned from skin fragility disorders. <b>2017</b> , 27, 8-11		
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475	Manufacturing of Human Extracellular Vesicle-Based Therapeutics for Clinical Use. <i>International Journal of Molecular Sciences</i> , <b>2017</b> , 18,	6.3	142
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472	Novel and emerging therapies in the treatment of recessive dystrophic epidermolysis bullosa. <b>2017</b> , 6, 6-20		56

471	Rehabilitation von Kindern und Jugendlichen mit chronischen Hauterkrankungen. <b>2017</b> , 43, 366-379		
470	Nonsense variant in COL7A1 causes recessive dystrophic epidermolysis bullosa in Central Asian Shepherd dogs. <b>2017</b> , 12, e0177527		10
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466	Seven novel COL7A1 mutations identified in patients with recessive dystrophic epidermolysis bullosa from Mexico. <b>2018</b> , 43, 579-584		4
465	Genetic Diagnosis of Epidermolysis Bullosa: Recommendations From an Expert Spanish Research Group. <b>2018</b> , 109, 104-122		1
464	Advances on potential therapeutic options for epidermolysis bullosa. <b>2018</b> , 6, 283-293		1
463	Allogeneic fibroblast cell therapy in the treatment of recessive dystrophic epidermolysis bullosa. <b>2018</b> , 21, 8-11		4
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461	Epidermolysis bullosa: Molecular pathology of connective tissue components in the cutaneous basement membrane zone. <i>Matrix Biology</i> , <b>2018</b> , 71-72, 313-329	11.4	53
460	Deletion of a Pathogenic Mutation-Containing Exon of COL7A1 Allows Clonal Gene Editing Correction of RDEB Patient Epidermal Stem Cells. <b>2018</b> , 11, 68-78		27
459	Burnlike scars: A sign suggestive of KLHL24-related epidermolysis bullosa simplex. <b>2018</b> , 35, e193-e195		5
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456	EB2017-Progress in Epidermolysis Bullosa Research toward Treatment and Cure. <b>2018</b> , 138, 1010-1016		30
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454	Ophthalmologic Approach in Epidermolysis Bullosa: A Cross-Sectional Study With Phenotype-Genotype Correlations. <b>2018</b> , 37, 442-447		6

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447	The Conundrum of Allogeneic Bone Marrow Transplantation for Epidermolysis Bullosa. <b>2018</b> , 138, 1029-1031		1
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441	Gene editing for skin diseases: designer nucleases as tools for gene therapy of skin fragility disorders. <b>2018</b> , 103, 449-455		24
440	Cell-Cell Junctions Organize Structural and Signaling Networks. <i>Cold Spring Harbor Perspectives in Biology</i> , <b>2018</b> , 10,	10.2	146
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436	Natural history and clinical outcome of junctional epidermolysis bullosa generalized intermediate due to a LAMA3 mutation. <i>British Journal of Dermatology</i> , <b>2018</b> , 178, 973-975	4	5

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432	Functional therapies for cutaneous wound repair in epidermolysis bullosa. <b>2018</b> , 129, 330-343		9
431	Wound culture isolated antibiograms and caregiver-reported skin care practices in children with epidermolysis bullosa. <b>2018</b> , 35, 92-96		11
430	Recessive mutation in tetraspanin CD151 causes Kindler syndrome-like epidermolysis bullosa with multi-systemic manifestations including nephropathy. <i>Matrix Biology</i> , <b>2018</b> , 66, 22-33	11.4	39
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420	Advances in understanding the molecular basis of skin fragility. <b>2018</b> , 7, 279		2
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418	Bart's syndrome in a family affected three consecutive generations with mutation c.6007G>A in COL7A1. <b>2018</b> , 45, 1000-1002		4



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4 <sup>16</sup>	Case report of two siblings with a novel homozygous mutation in COL7A1 leads to recessive dystrophic epidermolysis bullosa: which type?. <b>2018</b> , 27, 138-141		
4 <sup>15</sup>	Collagen VII deficient mice show morphologic and histologic corneal changes that phenotypically mimic human dystrophic epidermolysis bullosa of the eye. <b>2018</b> , 175, 133-141		6
4 <sup>14</sup>	The development of induced pluripotent stem cell-derived mesenchymal stem/stromal cells from normal human and RDEB epidermal keratinocytes. <b>2018</b> , 91, 301-310		24
4 <sup>13</sup>	Successful Multidisciplinary Treatment of Chronic Facial Wounds in Junctional Epidermolysis Bullosa. <b>2018</b> , 98, 711-712		1
4 <sup>12</sup>	The value of ultrahigh resolution OCT in dermatology - delineating the dermo-epidermal junction, capillaries in the dermal papillae and vellus hairs. <b>2018</b> , 9, 2240-2265		26
4 <sup>11</sup>	Pediatric Nail Disorders and Selected Genodermatoses with Nail Findings. <b>2018</b> , 317-342		
4 <sup>10</sup>	Update on Genetic Conditions Affecting the Skin and the Kidneys. <b>2018</b> , 6, 43		6
4 <sup>09</sup>	Genotype-Phenotype Correlations of Dystrophic Epidermolysis Bullosa in India: Experience from a Tertiary Care Centre. <b>2018</b> , 98, 873-879		11
4 <sup>08</sup>	Epidemiology and Outcome of Squamous Cell Carcinoma in Epidermolysis Bullosa in Australia and New Zealand. <b>2018</b> , 98, 70-76		13
4 <sup>07</sup>	A cost-effective treatment model in dystrophic epidermolysis bullosa with congenital absence of skin. <b>2018</b> , 31, e12649		2
4 <sup>06</sup>	The Position of Targeted Next-generation Sequencing in Epidermolysis Bullosa Diagnosis. <b>2018</b> , 98, 437-440		20
4 <sup>05</sup>	A review of scoring systems for ocular involvement in chronic cutaneous bullous diseases. <i>Orphanet Journal of Rare Diseases</i> , <b>2018</b> , 13, 83	4.2	8
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4 <sup>02</sup>	Basement Membranes in Development and Disease. <b>2018</b> , 130, 143-191		64
4 <sup>01</sup>	Clinical subtypes and molecular basis of epidermolysis bullosa in Kuwait. <b>2018</b> , 57, 1058-1067		2
4 <sup>00</sup>	A rare case of skin blistering and esophageal stenosis in the course of epidermolysis bullosa - case report and literature review. <b>2018</b> , 18, 47		3

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397	Anesthetic management with subcostal transversus abdominis plane block in recessive dystrophic epidermolysis bullosa for peritoneal dialysis catheter replacement: a case report. <b>2018</b> , 4, 37		1
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395	APOBEC mutation drives early-onset squamous cell carcinomas in recessive dystrophic epidermolysis bullosa. <b>2018</b> , 10,		53
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393	A "late-but-fitter revertant cell" explains the high frequency of revertant mosaicism in epidermolysis bullosa. <b>2018</b> , 13, e0192994		14
392	Congenital and Hereditary Disorders of the Skin. <b>2018</b> , 1475-1494.e1		
391	Integra-Dermal Regeneration Template and Split-Thickness Skin Grafting: A Therapy Approach to Correct Aplasia Cutis Congenita and Epidermolysis Bullosa in Carmi Syndrome. <b>2018</b> , 8, 313-321		10
390	Toll-like receptor signalling induces the expression of serum amyloid A in epidermal keratinocytes and dermal fibroblasts. <b>2019</b> , 44, 40-46		8
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388	Prevalence and pathogenesis of osteopenia and osteoporosis in epidermolysis bullosa: An evidence-based review. <i>Experimental Dermatology</i> , <b>2019</b> , 28, 1122-1130	4	6
387	The direct binding of collagen XVII and collagen IV is disrupted by pemphigoid autoantibodies. <b>2019</b> , 99, 48-57		10
386	Basement membrane collagens and disease mechanisms. <b>2019</b> , 63, 297-312		30
385	Clinical efficacy of dialkylcarbamoylchloride-coated cotton acetate dressing versus combination of normal saline dressing and 2% mupirocin ointment in infected wounds of epidermolysis bullosa. <b>2019</b> , 32, e13047		1
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383	Biallelic KRT5 mutations in autosomal recessive epidermolysis bullosa simplex, including a complete human keratin 5 "knock-out". <i>Matrix Biology</i> , <b>2019</b> , 83, 48-59	11.4	8
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380	Assessment of the risk and characterization of non-melanoma skin cancer in Kindler syndrome: study of a series of 91 patients. <i>Orphanet Journal of Rare Diseases</i> , <b>2019</b> , 14, 183	4.2	9
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378	Improved Double-Nicking Strategies for COL7A1-Editing by Homologous Recombination. <b>2019</b> , 18, 496-507		17
377	Esophageal lichen planus: towards diagnosis of an underdiagnosed disease. <b>2019</b> , 54, 1189-1198		5
376	Cardiac transplant for epidermolysis bullosa simplex with KLHL24 mutation-associated cardiomyopathy. <i>JAAD Case Reports</i> , <b>2019</b> , 5, 912-914	1.4	4
375	Natural Exon Skipping Sets the Stage for Exon Skipping as Therapy for Dystrophic Epidermolysis Bullosa. <b>2019</b> , 18, 465-475		16
374	Bullous Diseases in Children: A Review of Clinical Features and Treatment Options. <b>2019</b> , 21, 345-356		6
373	Alopecia in Autoimmune Blistering Diseases: A Systematic Review of Pathogenesis and Clinical Features of Disease. <b>2019</b> , 5, 263-275		4
372	Fibroblast activation and abnormal extracellular matrix remodelling as common hallmarks in three cancer-prone genodermatoses. <i>British Journal of Dermatology</i> , <b>2019</b> , 181, 512-522	4	31
371	Ectodermal dysplasias: Classification and organization by phenotype, genotype and molecular pathway. <b>2019</b> , 179, 442-447		39
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369	[Hair anomalies in syndromic disorders]. <b>2019</b> , 70, 514-519		
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365	Occupational therapy for epidermolysis bullosa: clinical practice guidelines. <i>Orphanet Journal of Rare Diseases</i> , <b>2019</b> , 14, 129	4.2	6
364	Resting Energy Expenditure and Protein Balance in People with Epidermolysis Bullosa. <b>2019</b> , 11,		1

363	Pathogenesis and clinical features of alopecia in epidermolysis bullosa: A systematic review. <b>2019</b> , 36, 430-436		1
362	Clamping Together Hemidesmosomes and Latching Them in Place. <b>2019</b> , 27, 881-883		1
361	Attenuation of Severe Generalized Junctional Epidermolysis Bullosa by Systemic Treatment with Gentamicin. <b>2019</b> , 235, 315-322		13
360	Oleogel-S10 Phase 3 study "EASE" for epidermolysis bullosa: study design and rationale. <b>2019</b> , 20, 350		8
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357	Transmission electron microscopy. <b>2019</b> , 434-475		7
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355	MicroRNA-145-5p regulates fibrotic features of recessive dystrophic epidermolysis bullosa skin fibroblasts. <i>British Journal of Dermatology</i> , <b>2019</b> , 181, 1017-1027	4	9
354	Epidemiology and natural history of cutaneous squamous cell carcinoma in recessive dystrophic epidermolysis bullosa patients: 20 years' experience of a reference centre in Spain. <b>2019</b> , 21, 1573-1577		6
353	CRISPR/Cas9 gene editing for genodermatoses: progress and perspectives. <b>2019</b> , 3, 313-326		4
352	Clinically Relevant Correction of Recessive Dystrophic Epidermolysis Bullosa by Dual sgRNA CRISPR/Cas9-Mediated Gene Editing. <b>2019</b> , 27, 986-998		48
351	Orofacial management for epidermolysis bullosa during wisdom tooth removal surgery: A technical note. <b>2019</b> , 120, 467-470		3
350	Pediatric Oral Medicine. <b>2019</b> , 1641-1718		1
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348	Recessive Dystrophic Epidermolysis Bullosa and Pregnancy. <b>2019</b> , 110, 50-52		
347	Dermatologic Diseases. <b>2019</b> , 457-503		
346	Thymosin $\beta$ : potential to treat epidermolysis bullosa and other severe dermal injuries. <b>2019</b> , 29, 459-467		7

345	Epidermolysis Bullosa and Kindler Syndrome. <b>2019</b> , 907-942		
344	Autoimmune Bullous Diseases. <b>2019</b> , 868-897		3
343	Epidermolysis Bullosa-Associated Squamous Cell Carcinoma: From Pathogenesis to Therapeutic Perspectives. <i>International Journal of Molecular Sciences</i> , <b>2019</b> , 20,	6.3	32
342	Review of Keratin Disorders. <b>2019</b> , 1515-1523		
341	Previously Unreported Mutation in a Somali Patient with Dystrophic Epidermolysis Bullosa. <b>2020</b> , 10, 332-338		1
340	Epidermolysis Bullosa Patients' Perception of Surgical Wound and Scar Healing. <b>2019</b> , 45, 280-289		1
339	From Clinical Phenotype to Genotypic Modelling: Incidence and Prevalence of Recessive Dystrophic Epidermolysis Bullosa (RDEB). <b>2019</b> , 12, 933-942		4
338	Epidermolysis bullosa hereditaria. <b>2019</b> , 31, 397-409		
337	Toward treatment and cure of epidermolysis bullosa. <b>2019</b> ,		12
336	A sensitive and specific assay for the serological diagnosis of antilaminin 332 mucous membrane pemphigoid. <i>British Journal of Dermatology</i> , <b>2019</b> , 180, 149-156	4	28
335	Phenotypic Spectrum of Epidermolysis Bullosa: The Paradigm of Syndromic versus Non-Syndromic Skin Fragility Disorders. <b>2019</b> , 139, 522-527		30
334	Junctional Epidermolysis Bullosa: Allelic Heterogeneity and Mutation Stratification for Precision Medicine. <b>2018</b> , 5, 363		14
333	Assessment of the Timing of Milestone Clinical Events in Patients With Epidermolysis Bullosa From North America. <i>JAMA Dermatology</i> , <b>2019</b> , 155, 196-203	5.1	20
332	Retrospective longitudinal study of osteoporosis in adults with recessive dystrophic epidermolysis bullosa. <i>Clinical Case Reports (discontinued)</i> , <b>2019</b> , 7, 58-63	0.7	5
331	Manipulation of Transgene Expression in Fibroblast Cells by a Multifunctional Linear-Branched Hybrid Poly(βAmino Ester) Synthesized through an Oligomer Combination Approach. <b>2019</b> , 19, 381-391		27
330	Epidermolysis Bullosa Simplex with KLHL24 Mutations Is Associated with Dilated Cardiomyopathy. <b>2019</b> , 139, 244-249		16
329	Phenotypic Features of Epidermolysis Bullosa Simplex due to KLHL24 Mutations in 3 Italian Cases. <b>2019</b> , 99, 238-239		4
328	Epidermolysis bullosa-specific module of the Infants and Toddlers Dermatology Quality of Life (InToDermQoL) questionnaire. <b>2019</b> , 33, 612-617		13

327	Mutations in PLOD3, encoding lysyl hydroxylase 3, cause a complex connective tissue disorder including recessive dystrophic epidermolysis bullosa-like blistering phenotype with abnormal anchoring fibrils and type VII collagen deficiency. <i>Matrix Biology</i> , <b>2019</b> , 81, 91-106	11.4	28
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325	Inherited epidermolysis bullosa: description of clinical and subclinical morphological features with optical coherence tomography. <b>2019</b> , 33, e120-e123		2
324	Decorin counteracts disease progression in mice with recessive dystrophic epidermolysis bullosa. <i>Matrix Biology</i> , <b>2019</b> , 81, 3-16	11.4	25
323	Bart's Syndrome with Novel Frameshift Mutations in the COL7A1 Gene. <b>2019</b> , 38, 72-79		3
322	Epidermolysis bullosa simplex-generalized severe type due to keratin 5 p.Glu477Lys mutation: Genotype-phenotype correlation and in silico modeling analysis. <b>2019</b> , 36, 132-138		8
321	A novel de novo mutation p.Ala428Asp in KRT5 gene as a cause of localized epidermolysis bullosa simplex. <i>Experimental Dermatology</i> , <b>2019</b> , 28, 1131-1134	4	2
320	The Clinical Trials of Mesenchymal Stem Cell Therapy in Skin Diseases: An Update and Concise Review. <b>2019</b> , 14, 22-33		66
319	Genome-wide single nucleotide polymorphism-based autozygosity mapping facilitates identification of mutations in consanguineous families with epidermolysis bullosa. <i>Experimental Dermatology</i> , <b>2019</b> , 28, 1118-1121	4	13
318	Inherited epidermolysis bullosa: New diagnostics and new clinical phenotypes. <i>Experimental Dermatology</i> , <b>2019</b> , 28, 1146-1152	4	24
317	Life before and beyond blistering: The role of collagen XVII in epidermal physiology. <i>Experimental Dermatology</i> , <b>2019</b> , 28, 1135-1141	4	21
316	Recessive Dystrophic Epidermolysis Bullosa and Pregnancy. <b>2019</b> , 110, 50-52		2
315	Minor collagens of the skin with not so minor functions. <b>2019</b> , 235, 418-429		12
314	Clinical practice guidelines for laboratory diagnosis of epidermolysis bullosa. <i>British Journal of Dermatology</i> , <b>2020</b> , 182, 574-592	4	45
313	Skin and Lacrimal Drainage System. <b>2020</b> , 163-233.e10		
312	Amino acid charge and epidermolysis bullosa simplex severity: genotype-phenotype correlations. <b>2020</b> , 34, e87-e90		0
311	Tumour serine proteases C1r and C1s as novel biomarkers and therapeutic targets in invasive sporadic and recessive dystrophic epidermolysis bullosa-associated cutaneous squamous cell carcinoma. <i>British Journal of Dermatology</i> , <b>2020</b> , 182, 530-531	4	0
310	Foot care in epidermolysis bullosa: evidence-based guideline. <i>British Journal of Dermatology</i> , <b>2020</b> , 182, 593-604	4	9

309	Human Mesenchymal Stromal Cells Engineered to Express Collagen VII Can Restore Anchoring Fibrils in Recessive Dystrophic Epidermolysis Bullosa Skin Graft Chimeras. <b>2020</b> , 140, 121-131.e6		9
308	Surgical management of hand deformities in patients with recessive dystrophic epidermolysis bullosa. <b>2020</b> , 54, 33-39		0
307	Epidermolysis bullosa: diagnostic guidelines in the laboratory setting. <i>British Journal of Dermatology</i> , <b>2020</b> , 182, 526-527	4	1
306	Rare Vesiculopustular Eruptions of the Neonatal Period. <b>2020</b> , 47, 53-75		
305	Development of bullous pemphigoid in junctional epidermolysis bullosa. <b>2020</b> , 34, e146-e148		1
304	Absent Skin and Butterfly Children: A Sporadic Case of Bart Syndrome. <b>2020</b> , 217, 214-214.e1		
303	Morphological and morphometric analysis of cutaneous squamous cell carcinoma in patients with recessive dystrophic epidermolysis bullosa: a retrospective study. <b>2020</b> , 34, 1707-1714		3
302	Natural history of growth and anaemia in children with epidermolysis bullosa: a retrospective cohort study. <i>British Journal of Dermatology</i> , <b>2020</b> , 182, 1437-1448	4	16
301	EBGene trial: patient preselection outcomes for the European GENEGRAFT ex vivo phase I/II gene therapy trial for recessive dystrophic epidermolysis bullosa. <i>British Journal of Dermatology</i> , <b>2020</b> , 182, 794-797	4	15
300	[Dressing material in children with epidermolysis bullosa : A qualitative study on experiences of parents and affected children]. <b>2020</b> , 34, 156-165		2
299	Context-Dependent Strategies for Enhanced Genome Editing of Genodermatoses. <b>2020</b> , 9,		17
298	The challenges of living with and managing epidermolysis bullosa: insights from patients and caregivers. <i>Orphanet Journal of Rare Diseases</i> , <b>2020</b> , 15, 1	4.2	41
297	Toward Combined Cell and Gene Therapy for Genodermatoses. <i>Cold Spring Harbor Perspectives in Biology</i> , <b>2020</b> , 12,	10.2	13
296	Association of dystrophic epidermolysis bullosa and neuroblastoma in a newborn. <b>2020</b> , 61, 117-118		
295	Phase I/II open-label trial of intravenous allogeneic mesenchymal stromal cell therapy in adults with recessive dystrophic epidermolysis bullosa. <i>Journal of the American Academy of Dermatology</i> , <b>2020</b> , 83, 447-454	4.5	26
294	Homozygous Nonsense Mutation in DSC3 Resulting in Skin Fragility and Hypotrichosis. <b>2020</b> , 140, 1285-1288		4
293	Living with epidermolysis bullosa: Daily challenges and health-care needs. <b>2020</b> , 23, 368-376		6
292	Epidermolysis bullosa with pyloric atresia consistently demonstrates concurrent low intra-basal epidermal and lamina lucida cleavage planes: a survey of six cases. <b>2020</b> , 34, e200-e203		3

291	Squamous cell carcinoma in a pregnant woman with recessive dystrophic epidermolysis bullosa. <b>2020</b> , 2020, omaa059		0
290	Kindler Syndrome: A Multidisciplinary Management Approach. <b>2020</b> , 111, 775-780		0
289	Epidermolysis bullosa with congenital absence of skin: Review of the literature. <b>2020</b> , 37, 821-826		3
288	A nonsense variant in the KRT14 gene in a domestic shorthair cat with epidermolysis bullosa simplex. <b>2020</b> , 51, 829-832		1
287	Consensus reclassification of inherited epidermolysis bullosa and other disorders with skin fragility. <i>British Journal of Dermatology</i> , <b>2020</b> , 183, 614-627	4	179
286	A de novo mutation in KRT5 in a crossbred calf with epidermolysis bullosa simplex. <b>2020</b> , 34, 2800-2807		2
285	Clinical practice guidelines: Oral health care for children and adults living with epidermolysis bullosa. <b>2020</b> , 40 Suppl 1, 3-81		9
284	Beneficial Effect of Systemic Allogeneic Adipose Derived Mesenchymal Cells on the Clinical, Inflammatory and Immunologic Status of a Patient With Recessive Dystrophic Epidermolysis Bullosa: A Case Report. <b>2020</b> , 7, 576558		4
283	The Human Epidermal Basement Membrane: A Shaped and Cell Instructive Platform That Aging Slowly Alters. <b>2020</b> , 10,		15
282	Molecular basis of skin disease. <b>2020</b> , 463-494		
281	Inherited epidermolysis bullosa: update on the clinical and genetic aspects. <b>2020</b> , 95, 551-569		14
280	Transplantation of a New Biological Product in Rare Diseases, Such as Epidermolysis Bullosa: Response and Clinical Outcome. <b>2020</b> , 52, 2239-2243		5
279	EB (epidermolysis bullosa)-House Austria: Pioneering work for the care of patients with rare diseases. <b>2020</b> , 18, 1229-1235		1
278	A Neonate with Blisters. <b>2020</b> , 41, S27-S29		1
277	Emerging drugs for the treatment of epidermolysis bullosa. <b>2020</b> , 25, 467-489		2
276	Raloxifene and n-Acetylcysteine Ameliorate TGF-Signalling in Fibroblasts from Patients with Recessive Dominant Epidermolysis Bullosa. <b>2020</b> , 9,		5
275	Epidermolysis bullosa. <b>2020</b> , 6, 78		64
274	Anterior-segment spectral domain optical coherence tomography in epidermolysis bullosa. <b>2020</b> , 18, 912-919		0



273	Missense Variant in Australian Shepherd Dogs with Junctional Epidermolysis Bullosa. <b>2020</b> , 11,		4
272	Cells from discarded dressings differentiate chronic from acute wounds in patients with Epidermolysis Bullosa. <i>Scientific Reports</i> , <b>2020</b> , 10, 15064	4.9	6
271	Outcomes and Predictors for Re-stenosis of Esophageal Stricture in Epidermolysis Bullosa: A Multicenter Cohort Study. <b>2020</b> , 71, 310-314		5
270	Epidermolysis bullosa and the importance of evaluation of sleep complaints. <b>2020</b> , 56, 1477-1478		0
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268	Kindler Syndrome: A Multidisciplinary Management Approach. <b>2020</b> , 111, 775-780		
267	EB (Epidermolysis bullosa)-Haus Austria: Pionierarbeit für die Betreuung von Patienten mit seltenen Erkrankungen. <b>2020</b> , 18, 1229-1235		1
266	Phenotypic discordance between siblings with junctional epidermolysis bullosa-pyloric atresia. <b>2020</b> , 45, 793-795		0
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264	Next-generation sequencing through multigene panel testing for the diagnosis of hereditary epidermolysis bullosa in Chinese population. <b>2020</b> , 98, 179-184		4
263	A new mutation associated with Pierson syndrome. <b>2020</b> , 118, e288-e291		0
262	Epidermolysis bullosa: a 2020 perspective. <i>British Journal of Dermatology</i> , <b>2020</b> , 183, 603	4	2
261	Sporadic form of epidermolysis bullosa simplex with mottled pigmentation. <b>2020</b> , 95, 536-538		1
260	Laryngeal stenosis associated with epidermolysis bullosa simplex. <i>JAAD Case Reports</i> , <b>2020</b> , 6, 465-467	1.4	
259	Novel biallelic variants in COL7A1 cause recessive dystrophic epidermolysis bullosa. <i>Molecular Genetics &amp; Genomic Medicine</i> , <b>2020</b> , 8, e1347	2.3	1
258	The quality of life in epidermolysis bullosa (EB-QoL) questionnaire: Translation, cultural adaptation, and validation into the Farsi language. <b>2020</b> , 6, 301-305		1
257	A novel pathogenic mutation in the COL7A1 gene resulting in mild autosomal dominant bullous dermolysis of the newborn. <b>2020</b> , 37, 955-957		1
256	Leading edge: emerging drug, cell, and gene therapies for junctional epidermolysis bullosa. <i>Expert Opinion on Biological Therapy</i> , <b>2020</b> , 20, 911-923	5.4	5

255	Generation of a human induced pluripotent stem cell line (UQACi001-A) from a severe epidermolysis bullosa simplex patient with the heterozygous mutation p.R125S in the KRT14 gene. <b>2020</b> , 44, 101748		0
254	The photobiomodulation therapy together with the use of cord blood platelet gel could be safely suggested as primary treatment for oral lesions in patients with inherited epidermolysis bullosa. <b>2020</b> , 36, 318-321		2
253	Aberrant splicing as potential modifier of the phenotype of junctional epidermolysis bullosa. <b>2020</b> , 34, 2127-2134		4
252	Predictable CRISPR/Cas9-Mediated COL7A1 Reframing for Dystrophic Epidermolysis Bullosa. <b>2020</b> , 140, 1985-1993.e5		13
251	Efficacy and tolerability of the investigational topical cream SD-101 (6% allantoin) in patients with epidermolysis bullosa: a phase 3, randomized, double-blind, vehicle-controlled trial (ESSENCE study). <i>Orphanet Journal of Rare Diseases</i> , <b>2020</b> , 15, 158	4.2	3
250	Spontaneous Gene Mutation in Rhesus Macaques (): A Novel Nonhuman Primate Model of Epidermolysis Bullosa Simplex. <b>2020</b> , 57, 344-348		2
249	Peel Test to Assess the Adhesion Strength of the Dermal-Epidermal Junction in Tissue-Engineered Skin. <b>2020</b> , 26, 180-189		2
248	Reframing the Care of Children With Epidermolysis Bullosa Through the Lens of Medical Trauma. <b>2020</b> , 12, 16-23		0
247	Successful dapsone therapy in inherited epidermolysis bullosa. <b>2020</b> , 34, e333-e334		1
246	Family caregivers' lived experiences of caring for epidermolysis bullosa patients: A phenomenological study. <b>2020</b> , 29, 1552-1560		11
245	Digenic inheritance of KRT5 and KRT14 mutations in a family with epidermolysis bullosa simplex. <i>Australasian Journal of Dermatology</i> , <b>2020</b> , 61, e267-e269	1.3	0
244	[Ex vivo stem cell gene therapy of the skin : Ready for clinical use?]. <b>2020</b> , 71, 85-90		
243	A novel pathogenic FERMT1 variant in four families with Kindler syndrome in Argentina. <b>2020</b> , 37, 337-341		1
242	Hoarse cry in a newborn with epidermolysis bullosa simplex, generalized severe. <b>2020</b> , 37, 393-395		1
241	Surgical management of squamous cell carcinoma arising in patients affected by epidermolysis bullosa: a comparative study. <b>2020</b> , 17, 519-521		1
240	Genomics-based treatment in a patient with two overlapping heritable skin disorders: Epidermolysis bullosa and acrodermatitis enteropathica. <i>Human Mutation</i> , <b>2020</b> , 41, 906-912	4.7	7
239	Footprint-free gene mutation correction in induced pluripotent stem cell (iPSC) derived from recessive dystrophic epidermolysis bullosa (RDEB) using the CRISPR/Cas9 and piggyBac transposon system. <b>2020</b> , 98, 163-172		13
238	Novel homozygous deletion of the plakophilin-1 gene in a Chinese patient with ectodermal dysplasia-skin fragility syndrome. <b>2020</b> , 47, 779-781		

237	A cancer stem cell-like phenotype is associated with miR-10b expression in aggressive squamous cell carcinomas. <b>2020</b> , 18, 61			11
236	Quality of life in people with epidermolysis bullosa: a systematic review. <b>2020</b> , 29, 1731-1745			8
235	Molecular Therapeutics in Development for Epidermolysis Bullosa: Update 2020. <b>2020</b> , 24, 299-309			26
234	Absence of tongue papillae as a clinical criterion for the diagnosis of generalized recessive dystrophic epidermolysis bullosa types. <i>Journal of the American Academy of Dermatology</i> , <b>2020</b> , 83, 1815-1816	4.5	1816	3
233	Functional analysis of keratin filament network formation indicates clinical severity of epidermolysis bullosa simplex. <b>2020</b> , 34, e613-e616			0
232	Patient-reported outcomes and quality of life in recessive dystrophic epidermolysis bullosa: A global cross-sectional survey. <i>Journal of the American Academy of Dermatology</i> , <b>2021</b> , 85, 1161-1167	4.5		8
231	Genotype-phenotype correlations on epidermolysis bullosa with congenital absence of skin: A comprehensive review. <b>2021</b> , 99, 29-41			8
230	Intravenous Injection of Muse Cells as a Potential Therapeutic Approach for Epidermolysis Bullosa. <b>2021</b> , 141, 198-202.e6			8
229	Heterogeneous addiction to transforming growth factor-beta signalling in recessive dystrophic epidermolysis bullosa-associated cutaneous squamous cell carcinoma. <i>British Journal of Dermatology</i> , <b>2021</b> , 184, 697-708	4		3
228	Efficacy of allogeneic cord blood platelet gel on wounds of dystrophic epidermolysis bullosa patients after pseudosyndactyly surgery. <b>2021</b> , 29, 134-143			2
227	Transplantation of autologous single hair units heals chronic wounds in autosomal recessive dystrophic epidermolysis bullosa: A proof-of-concept study. <b>2021</b> , 30, 36-41			2
226	Targeting the Jak/Signal Transducer and Activator of Transcription 3 Pathway with Ruxolitinib in a Mouse Model of Recessive Dystrophic Epidermolysis Bullosa-Squamous Cell Carcinoma. <b>2021</b> , 141, 942-946			3
225	Novel insights into the epidemiology of epidermolysis bullosa (EB) from the Dutch EB Registry: EB more common than previously assumed?. <b>2021</b> , 35, 995-1006			11
224	Inherited epidermolysis bullosa: epidemiology and patient care in Slovenia with a review of the updated classification. <b>2021</b> , 30,			0
223	Cardiac Involvement in Epidermolysis Bullosa. <b>2021</b> , 33-48			
222	Reverse oblique proximal femoral fracture in dystrophic epidermolysis bullosa: challenges and recommendations. <b>2021</b> , 14,			1
221	A case of recessive dystrophic epidermolysis bullosa treated with a cultured epidermal autograft. <b>2021</b> , 48, e165-e166			1
220	Successful use of immunotherapy to treat advanced cutaneous squamous cell carcinoma in recessive dystrophic epidermolysis bullosa. <b>2021</b> , 14,			1

219	Dermatofibrosarcoma protuberans in a young patient with epidermolysis bullosa: a case report. <b>2021</b> , 21, 100		
218	Neonatal epidermolysis bullosa, step-by-step diagnostic procedure. <b>2021</b> , 36, 565-565		
217	Esophagitis. <b>2021</b> , 183-193		
216	Epigenetic and metabolic regulation of epidermal homeostasis. <i>Experimental Dermatology</i> , <b>2021</b> , 30, 1009-1022	4	3
215	Pediatric Nail Disorders. <b>2021</b> , 39, 231-243		2
214	A systematic literature review of the disease burden in patients with recessive dystrophic epidermolysis bullosa. <i>Orphanet Journal of Rare Diseases</i> , <b>2021</b> , 16, 175	4.2	9
213	Identification and Computational Analysis of Novel Pathogenic Variants in Pakistani Families with Diverse Epidermolysis Bullosa Phenotypes. <b>2021</b> , 11,		2
212	Gene Replacement Therapies for Genodermatoses: A. <i>Frontiers in Genetics</i> , <b>2021</b> , 12, 658295	4.5	2
211	Cutaneous Diseases of the Esophagus. <b>2021</b> , 789-803		
210	Variant NAXOS-Carvajal Syndrome with Rare Additional Features of Systemic Bulla and Brittle Nails: A Case Report and Literature Review. <b>2021</b> , 60, 1119-1126		0
209	Drug Development for Target Ribosomal Protein rpL35/uL29 for Repair of LAMB3R635X in Rare Skin Disease Epidermolysis Bullosa. <b>2021</b> , 34, 167-182		3
208	Dynamics and Emerging Trends in Genodermatology: A Scientometric Analysis. <b>2021</b> , 4, 67-69		
207	Pathogenetic Therapy of Epidermolysis Bullosa: Current State and Prospects. <b>2021</b> , 171, 109-121		0
206	Kidney and Urinary Tract Involvement in Epidermolysis Bullosa: Is Routine Follow-Up Necessary?. <b>2021</b> , 11, e2021051		0
205	Epidermolysis Bullosa in Spain: Observational Study of a Cohort of Patients Treated in a National Referral Center. <b>2021</b> , 112, 781-781		
204	Increased abundance of Cbl E3 ligases alters PDGFR signaling in recessive dystrophic epidermolysis bullosa.		
203	Novel mutations of epidermolysis bullosa identified using whole-exome sequencing in Indonesian Javanese patients. <b>2021</b> , 10, 88-94		0
202	Plantar involvement correlates with obesity, pain and impaired mobility in epidermolysis bullosa simplex: a retrospective cohort study. <b>2021</b> , 35, 2097-2104		0

201	Kindler epidermolysis bullosa-like skin phenotype and downregulated basement membrane zone gene expression in poikiloderma with neutropenia and a homozygous USB1 mutation. <i>Matrix Biology</i> , <b>2021</b> , 99, 43-57	11.4	0
200	Interventions for inherited forms of epidermolysis bullosa.		78
199	Prevalence and antimicrobial resistance profile of <i>Staphylococcus aureus</i> in inherited epidermolysis bullosa: a cross-sectional multicenter study in Brazil. <b>2021</b> , 60, 1126-1130		0
198	Epidermolysis bullosa dystrophica pretibialis - Clinical snapshot and management of a rare orphan disease. <b>2021</b> , 19, 983-986		
197	Single-keratinocyte transcriptomic analyses identify different clonal types and proliferative potential mediated by FOXM1 in human epidermal stem cells. <b>2021</b> , 12, 2505		5
196	Pediatric Oral Medicine. <b>2021</b> , 943-989		
195	Overview of complications associated with epidermolysis bullosa: A multicenter retrospective clinical analysis of 152 cases. <b>2021</b> , 56, 2392-2398		0
194	Hair follicle stem cell progeny heal blisters while pausing skin development. <b>2021</b> , 22, e50882		4
193	Mouse models for dominant dystrophic epidermolysis bullosa carrying common human point mutations recapitulate the human disease. <b>2021</b> , 14,		4
192	How to Deal with Skin Biopsy in an Infant with Blisters?. <b>2021</b> , 8, 159-175		0
191	Epidermolysis bullosa with pyloric atresia associated with compound heterozygous ITGB4 pathogenic variants: Minimal skin involvement but severe mucocutaneous disease. <b>2021</b> , 38, 908-912		0
190	Pruritic Lichenified Plaques on the Legs of a Man. <i>JAMA Dermatology</i> , <b>2021</b> , 157, 868-869	5.1	1
189	Case Report: Uncommon Association of and Gene Mutation in a Case of Epidermolysis Bullosa With Pyloric Atresia and Aplasia Cutis Congenita. <i>Frontiers in Genetics</i> , <b>2021</b> , 12, 641977	4.5	2
188	Epidermolysis Bullosa in Spain: Observational Study of a Cohort of Patients Treated in a National Referral Center. <b>2021</b> , 112, 781-781		
187	Epidermolysis bullosa dystrophica pretibialis - Klinischer Schnappschuss und Management einer seltenen Erkrankung. <b>2021</b> , 19, 983-986		
186	Novel and very rare causative variants in the COL7A1 gene of Vietnamese patients with recessive dystrophic epidermolysis bullosa revealed by whole-exome sequencing. <i>Molecular Genetics &amp; Genomic Medicine</i> , <b>2021</b> , 9, e1748	2.3	2
185	Evidence for cutaneous dysbiosis in dystrophic epidermolysis bullosa. <b>2021</b> , 46, 1223-1229		2
184	Canine junctional epidermolysis bullosa due to a novel mutation in LAMA3 with severe upper respiratory involvement. <b>2021</b> , 32, 379-e108		0

183	Oleogel Formulations for the Topical Delivery of Betulin and Lupeol in Skin Injuries-Preparation, Physicochemical Characterization, and Pharmacotoxicological Evaluation. <b>2021</b> , 26,		1
182	Genetic trend of the junctional epidermolysis bullosa in the German shorthaired pointer in Italy. <b>2021</b> , 8, e15		0
181	Assessing the quality of life in the families of patients with epidermolysis bullosa: The mothers as main caregivers.. <b>2021</b> , 7, 721-726		2
180	Hologene 5: A Phase II/III Clinical Trial of Combined Cell and Gene Therapy of Junctional Epidermolysis Bullosa. <i>Frontiers in Genetics</i> , <b>2021</b> , 12, 705019	4-5	1
179	The natural history of laryngo-onycho-cutaneous syndrome: A case series of six pediatric patients and literature review. <b>2021</b> , 38, 1094-1101		4
178	Characterization of mutant type VII collagens underlying the inversa subtype of recessive dystrophic epidermolysis bullosa. <b>2021</b> , 104, 104-111		0
177	Novel pathogenic variants in an Indian cohort with epidermolysis bullosa: Expanding the genotypic spectrum. <b>2021</b> , 64, 104345		1
176	Extracellular Vesicles as Therapeutic Tools for the Treatment of Chronic Wounds. <b>2021</b> , 13,		4
175	A Familial Form of Epidermolysis Bullosa Simplex Associated with a Pathogenic Variant in. <b>2021</b> , 12,		0
174	Surgical Treatment of Wounds Using Stem Cells in Epidermolysis Bullosa (EB).		1
173	Congenital myopathy and epidermolysis bullosa due to PLEC variant. <b>2021</b> ,		1
172	A non-viral and selection-free HDR approach with improved safety profile for dystrophic epidermolysis bullosa. <b>2021</b> , 25, 237-250		2
171	Highly branched poly(ε-amino ester)s for gene delivery in hereditary skin diseases. <b>2021</b> , 176, 113842		4
170	A Rare Skin Disorder with Bacteremia in a Neonate. <b>2021</b> , 22, e705-e708		
169	A Novel Mutation p.L461P in Causing Localized Epidermolysis Bullosa Simplex. <b>2021</b> , 33, 11-17		1
168	Epidermolysis Bullosa. <b>2021</b> , 435-446		
167	The relationship between quality of life and coping strategies of children with EB and their parents. <i>Orphanet Journal of Rare Diseases</i> , <b>2021</b> , 16, 53	4.2	1
166	Advances in gene editing strategies for epidermolysis bullosa. <b>2021</b> , 182, 81-109		3

165	Epidermolysis bullosa. Possible methods of treatment. <i>Klinicheskaya Dermatologiya I Venerologiya</i> , <b>2021</b> , 20, 22	0.3
164	Bullous Disorders of Childhood. <b>2016</b> , 205-220	1
163	Prenatal Diagnosis of Epidermolysis Bullosa: Current Aspects and Perspectives. <b>2015</b> , 239-248	1
162	Hereditäre Epidermolysen. <b>2018</b> , 823-838	2
161	Meeting Report: The First Global Congress on Epidermolysis Bullosa, EB2020 London: Toward Treatment and Cure. <b>2020</b> , 140, 1681-1687	8
160	Hemidesmosomes modulate force generation via focal adhesions. <b>2020</b> , 219,	39
159	Safety and early efficacy outcomes for lentiviral fibroblast gene therapy in recessive dystrophic epidermolysis bullosa. <b>2019</b> , 4,	30
158	Phase 1/2a clinical trial of gene-corrected autologous cell therapy for recessive dystrophic epidermolysis bullosa. <b>2019</b> , 4,	42
157	Recent advances in understanding and managing epidermolysis bullosa. <b>2018</b> , 7,	7
156	Gene editing in dermatology: Harnessing CRISPR for the treatment of cutaneous disease. <b>2020</b> , 9, 281	3
155	Combining GWAS and RNA-Seq Approaches for Detection of the Causal Mutation for Hereditary Junctional Epidermolysis Bullosa in Sheep. <b>2015</b> , 10, e0126416	10
154	Lysyl Hydroxylase 3 Localizes to Epidermal Basement Membrane and Is Reduced in Patients with Recessive Dystrophic Epidermolysis Bullosa. <b>2015</b> , 10, e0137639	18
153	Epidemiology and Providing of Healthcare for Patients with Inherited Epidermolysis Bullosa in the Russian Federation. <b>2018</b> , 73, 420-430	4
152	Challenges of the differential diagnosis between the subtypes of the junctional epidermolysis bullosa: presentation of two clinical cases. <b>2019</b> , 47, 83-93	1
151	Congenital epidermolysis bullosa: modern methods of diagnosis and therapy. Prospects for regenerative medicine. <i>Vestnik Dermatologii I Venerologii</i> , <b>2020</b> , 96, 10-17	0.4 3
150	Pediatric Ophthalmoplegia and Ptosis in Epidermolysis Bullosa Simplex Associated With Muscular Dystrophy. <b>2018</b> , 55, e26-e29	1
149	Health literacy in patients with epidermolysis bullosa in Iran. <b>2017</b> , 6, 105	1
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