## Anthony J Aldave

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	IC3D Classification of Corneal Dystrophies—Edition 2. Cornea, 2015, 34, 117-159.	0.9	425
2	The Boston Type I Keratoprosthesis. Ophthalmology, 2009, 116, 640-651.	2.5	286
3	The IC3D Classification of the Corneal Dystrophies. Cornea, 2008, 27, S1-S42.	0.9	277
4	International Results with the Boston Type I Keratoprosthesis. Ophthalmology, 2012, 119, 1530-1538.	2.5	158
5	Report of the Eye Bank Association of America Medical Advisory Board Subcommittee on Fungal Infection After Corneal Transplantation. Cornea, 2013, 32, 149-154.	0.9	106
6	Etiology of Global Corneal Blindness and Current Practices of Corneal Transplantation: A Focused Review. Cornea, 2018, 37, 1198-1203.	0.9	92
7	NoVSX1Gene Mutations Associated with Keratoconus. , 2006, 47, 2820.		87
8	Outcomes After DSEK in 101 Eyes With Previous Trabeculectomy and Tube Shunt Implantation. Cornea, 2014, 33, 223-229.	0.9	81
9	Corneal Endothelium in Patients with Anterior Uveitis. Ophthalmology, 2016, 123, 1637-1645.	2.5	74
10	Transcriptomic Analysis of Cultured Corneal Endothelial Cells as a Validation for Their Use in Cell Replacement Therapy. Cell Transplantation, 2016, 25, 1159-1176.	1.2	58
11	Autosomal Recessive CHED Associated With Novel Compound Heterozygous Mutations in SLC4A11. Cornea, 2007, 26, 896-900.	0.9	57
12	Posterior polymorphous corneal dystrophy is associated with <i>TCF8</i> gene mutations and abdominal hernia. American Journal of Medical Genetics, Part A, 2007, 143A, 2549-2556.	0.7	57
13	Corneal Endothelial Cell Loss 3 Years After Successful Descemet Stripping Automated Endothelial Keratoplasty in the Cornea Preservation Time Study. JAMA Ophthalmology, 2017, 135, 1394.	1.4	49
14	Safety of Concurrent Boston Type I Keratoprosthesis and Glaucoma Drainage Device Implantation. Ophthalmology, 2017, 124, 12-19.	2.5	44
15	Classification of Posterior Polymorphous Corneal Dystrophy as a Corneal Ectatic Disorder Following Confirmation of Associated Significant Corneal Steepening. JAMA Ophthalmology, 2013, 131, 1583.	1.4	41
16	Phenotypic and functional characterization of corneal endothelial cells during in vitro expansion. Scientific Reports, 2020, 10, 7402.	1.6	41
17	Multifunctional ion transport properties of human SLC4A11: comparison of the SLC4A11-B and SLC4A11-C variants. American Journal of Physiology - Cell Physiology, 2016, 311, C820-C830.	2.1	40
18	A Clinical and Histopathologic Examination of Accelerated TGFBIp Deposition After LASIK in Combined Granular-Lattice Corneal Dystrophy. American Journal of Ophthalmology, 2007, 143, 416-419.	1.7	37

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19	Long-Term Visual Outcomes, Complications, and Retention of the Boston Type I Keratoprosthesis. Cornea, 2018, 37, 3-10.	0.9	36
20	Identification of mutations in UBIAD1 following exclusion of coding mutations in the chromosome 1p36 locus for Schnyder crystalline corneal dystrophy. Molecular Vision, 2007, 13, 1777-82.	1.1	31
21	Posterior Amorphous Corneal Dystrophy Is Associated with a Deletion of Small Leucine-rich Proteoglycans on Chromosome 12. PLoS ONE, 2014, 9, e95037.	1.1	28
22	Recurrent corneal erosion syndrome. British Journal of Ophthalmology, 2019, 103, 1204-1208.	2.1	27
23	Epithelial Debridement and Bowman's Layer Polishing for Visually Significant Epithelial Irregularity and Recurrent Corneal Erosions. Cornea, 2009, 28, 1085-1090.	0.9	26
24	Corneal Blindness in Asia: A Systematic Review and Meta-Analysis to Identify Challenges and Opportunities. Cornea, 2020, 39, 1196-1205.	0.9	26
25	Outcomes of the Boston Type I Keratoprosthesis as the Primary Penetrating Corneal Procedure. Cornea, 2018, 37, 1400-1407.	0.9	25
26	Linkage of Posterior Amorphous Corneal Dystrophy to Chromosome 12q21.33 and Exclusion of Coding Region Mutations in <i>KERA</i> , <i>LUM</i> , <i>DCN</i> , and <i>EPYC</i> , 2010, 51, 4006.		24
27	Development of a nomogram for femtosecond laser astigmatic keratotomy for astigmatism after keratoplasty. Journal of Cataract and Refractive Surgery, 2016, 42, 556-562.	0.7	24
28	Transcriptomic Profiling of Posterior Polymorphous Corneal Dystrophy. , 2017, 58, 3202.		24
29	Corneal dystrophies. Nature Reviews Disease Primers, 2020, 6, 46.	18.1	24
30	Comparison of Endothelial Keratoplasty Techniques in Patients With Prior Glaucoma Surgery: A Case-Matched Study. American Journal of Ophthalmology, 2019, 206, 94-101.	1.7	21
31	The Genetics of the Corneal Dystrophies. Developments in Ophthalmology, 2011, 48, 51-66.	0.1	20
32	ZEB1 insufficiency causes corneal endothelial cell state transition and altered cellular processing. PLoS ONE, 2019, 14, e0218279.	1.1	20
33	Confirmation of the OVOL2 Promoter Mutation c307T>C in Posterior Polymorphous Corneal Dystrophy 1. PLoS ONE, 2017, 12, e0169215.	1.1	20
34	Newly Reported p.Asp240Asn Mutation in UBIAD1 Suggests Central Discoid Corneal Dystrophy Is a Variant of Schnyder Corneal Dystrophy. Cornea, 2010, 29, 777-780.	0.9	19
35	Posterior polymorphous corneal dystrophy 3 is associated with agenesis and hypoplasia of the corpus callosum. Vision Research, 2014, 100, 88-92.	0.7	18
36	Complications Related to a Cosmetic Eye-Whitening Procedure. American Journal of Ophthalmology, 2014, 158, 967-973.	1.7	17

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37	Alterations in GRHL2-OVOL2-ZEB1 axis and aberrant activation of Wnt signaling lead to altered gene transcription in posterior polymorphous corneal dystrophy. Experimental Eye Research, 2019, 188, 107696.	1.2	16
38	Infectious keratitis after lamellar keratoplasty. Survey of Ophthalmology, 2021, 66, 623-643.	1.7	14
39	No Pathogenic Mutations Identified in the TGFBI Gene in Polymorphic Corneal Amyloid Deposition. Cornea, 2006, 25, 413-415.	0.9	13
40	Variant lattice corneal dystrophy associated with compound heterozygous mutations in the <i>TGFBI</i> gene. British Journal of Ophthalmology, 2017, 101, 509-513.	2.1	13
41	Elucidating the molecular basis of PPCD: Effects of decreased ZEB1 expression on corneal endothelial cell function. Molecular Vision, 2017, 23, 740-752.	1.1	13
42	Exclusion of Positional Candidate Gene Coding Region Mutations in the Common Posterior Polymorphous Corneal Dystrophy 1 Candidate Gene Interval. Cornea, 2009, 28, 801-807.	0.9	12
43	Long-term outcomes of the Boston type I keratoprosthesis in eyes with previous herpes simplex virus keratitis. British Journal of Ophthalmology, 2018, 102, 48-53.	2.1	12
44	COVID and the Cornea: From Controversies to Consensus. Cornea, 2021, 40, 809-816.	0.9	12
45	Whole Exome Sequencing and Segregation Analysis Confirms That a Mutation in COL17A1 Is the Cause of Epithelial Recurrent Erosion Dystrophy in a Large Dominant Pedigree Previously Mapped to Chromosome 10q23-q24. PLoS ONE, 2016, 11, e0157418.	1.1	10
46	Identification of novel PIKFYVE gene mutations associated with Fleck corneal dystrophy. Molecular Vision, 2015, 21, 1093-100.	1.1	10
47	Autosomal Dominant Cornea Plana is not Associated with Pathogenic Mutations inDCN, DSPG3, FOXC1, KERA, LUM,orPITX2. Ophthalmic Genetics, 2007, 28, 57-67.	0.5	9
48	Hereditary Benign Intraepithelial Dyskeratosis: Report of a Case and Re-examination of the Evidence for Locus Heterogeneity. Ophthalmic Genetics, 2016, 37, 1-5.	0.5	9
49	Identification of the First <i>De Novo UBIAD1</i> Gene Mutation Associated with Schnyder Corneal Dystrophy. Journal of Ophthalmology, 2016, 2016, 1-9.	0.6	9
50	Identification of Potentially Pathogenic Variants in the Posterior Polymorphous Corneal Dystrophy 1 Locus. PLoS ONE, 2016, 11, e0158467.	1.1	9
51	Punctiform and Polychromatic Pre-Descemet Corneal Dystrophy: Clinical Evaluation and Identification of the Genetic Basis. American Journal of Ophthalmology, 2020, 212, 88-97.	1.7	6
52	POSTOPERATIVE POSTERIOR SEGMENT COMPLICATIONS AFTER BOSTON TYPE 1 KERATOPROSTHESIS. Retina, 2021, 41, 2499-2509.	1.0	6
53	Identification of presumed pathogenic KRT3 and KRT12 gene mutations associated with Meesmann corneal dystrophy. Molecular Vision, 2015, 21, 1378-86.	1.1	5
54	Multimodal Imaging of Pre-Descemet Corneal Dystrophy Associated With X-Linked Ichthyosis and Deletion of the STS Gene. Cornea, 2020, 39, 1442-1445.	0.9	4

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#	Article	IF	CITATIONS
55	Sutured Custom Foldable Silicone Artificial Iris Implantation Combined With Intraocular Lens Implantation and Penetrating Keratoplasty. Cornea, 2020, Publish Ahead of Print, 1236-1247.	0.9	4
56	Confirmation and refinement of the heterozygous deletion of the small leucine-rich proteoglycans associated with posterior amorphous corneal dystrophy. Ophthalmic Genetics, 2018, 39, 419-424.	0.5	3
57	ldentification of a Novel Missense <b><i>KRT12</i></b> Mutation in a Vietnamese Family with Meesmann Corneal Dystrophy. Case Reports in Ophthalmology, 2020, 11, 120-126.	0.3	3
58	Idiopathic Vitritis after Boston Type 1 Keratoprosthesis Implantation: Incidence, Risk Factors and Outcomes in a Multicentric Cohort. Ocular Immunology and Inflammation, 2020, , 1-7.	1.0	3
59	Role of Antifungal Prophylaxis After Receipt of a Positive Donor Rim Fungal Culture: The Case for Treatment. Cornea, 2021, 40, 1093-1095.	0.9	3
60	Infectious Keratitis After Boston Type 1 Keratoprosthesis Implantation. Cornea, 2021, Publish Ahead of Print, 1298-1308.	0.9	2
61	Late Onset Interface Calcium Deposition After Laser In Situ Keratomileusis. Cornea, 2021, Publish Ahead of Print, 116-120.	0.9	1
62	Infectious keratitis in Vietnam: etiology, organisms, and management at Vietnam National Eye Hospital. International Journal of Ophthalmology, 2022, 15, 128-134.	0.5	1
63	Identification of A Novel <i>TGFBI</i> Gene Mutation (p.Serine524Cystine) Associated with Late Onset Recurrent Epithelial Erosions and Bowman Layer Opacities. Ophthalmic Genetics, 2020, 41, 639-644.	0.5	0
64	Atypical Corneal Deposits after Deep Anterior Lamellar Keratoplasty. Ophthalmology, 2020, 127, 466.	2.5	0
65	Corneal ectasia associated with posterior lamellar opacification. Ophthalmic Genetics, 2021, 42, 486-492.	0.5	0
66	Confirmation of association of <i>TGFBI</i> p.Ser591Phe mutation with variant lattice corneal dystrophy. Ophthalmic Genetics, 2022, , 1-4.	0.5	0