

Academic Staff Resume

Name: Kang Zhang
Title: Chair Professor
Dept. SKLplanet



Office : P26
Tel. : 00853-6555 8758
E-mail : kang.zhang@gmail.com

Academic Qualification:

Ph.D. in Harvard University, Cambridge, MA, Genetics
MD in Harvard University Medical School - M.I.T., Health Sciences and Technology, Cambridge, MA, Medicine
Bachelor in Sichuan University, Sichuan, China, Biochemistry

Teaching Area

08/2019-present, Chair Professor of the Faculty of Medicine, Macau University of Science and Technology
07/2008-07/2019, full professor with tenure, University of California San Diego
03/2002-6/2006, Tenure Track-Assistant Professor; 07/2006, Promotion, Associate Professor with tenure, University of Utah, Depart. of Ophthalmology/Visual Sciences
07/1999-09/2000, Instructor, Johns Hopkins University, Wilmer Eye Institute

Research Area

Ophthalmology
Genetics
Stem cells
Biomaterials
Artificial intelligence
Precision medicine

Working Experience

2019 – pres Chair Professor, Director of Biomedicine and Innovation Center, The Faculty of Medicine, Macau University of Science and Technology
2019 – pres Ophthalmologist, University Hospital
07/2008-08/2019 Staff Physician, Veteran Affairs San Diego Health System
2014 – 2019 □Co-Director, Biomaterials and Tissue Engineering Center, Institute of Engineering in Medicine, University of California San Diego, La Jolla, CA
2009 – 2013 □Founding Director, Institute for Genomic Medicine, University of California San Diego, La Jolla, CA
2006 – 2008□Associate Professor with tenure, University of Utah, Department of Ophthalmology and Visual Science, John Moran Eye Center, Salt Lake City, UT
2002 – 2006 Attending Staff Physician, Veteran's Administration Medical Center, Salt Lake City, UT
2002 -2006□Assistant Professor, University of Utah, Department of Ophthalmology and Visual Science, John Moran Eye Center, Salt Lake City, UT

2002 - 2003 FELLOW, University of Utah School of Medicine, Salt Lake City, UT, Vitreoretinal Diseases and Surgery
10/2000-02/2002, Assistant Staff, Cleveland Clinic Foundation, Cole Eye Institute
1999 -2000Instructor, Johns Hopkins University, School of Medicine, Wilmer Eye Institute, Baltimore, MD
1996 - 1999RESIDENT, Johns Hopkins University, Wilmer Eye Institute, Baltimore, MD, Ophthalmology
1995 - 1996 INTERN, Presbyterian/St. Lukes Hospital, Denver, CO, Medicine

Research Projects

FDCT: 0035/2020/A: Key techniques for diagnosing new coronavirus 2019-ncov infection based on artificial intelligence
FDCT-NSFC: 0007/2020/AFJ: Induction of conjunctival stem cells into limbal-like stem cells and corneal reconstruction
FDCT:20200530132841618/A2: New Artificial Intelligence Algorithm and Its Application in Electronic Medical Record System

Professional Certification and Awards

Highly Cited Researcher 2020 (Cross-Field)
Highly Cited Researcher 2019 (Cross-Field)
The Ophthalmologist World 100 Power List (2016, 2018)
America's Top Ophthalmologists, Consumer's research Council of America (2011)
NIH Director's Transformative R01 Program (with a perfect review score 10, 2010)
Senior Investigator Award, Research to Prevent Blindness (2010)
Outstanding Achievement Award, Chinese Ophthalmological Society, 2009
Burroughs Wellcome Fund Clinical Scientist Award in Translational Research, 2008
Lew R. Wasserman Merit Award, Research to Prevent Blindness, 2006
Macular Vision Research Award, 2002
Ruth Steinbach Fund for Macular Degeneration, 2001
Charles Schepens Award for Excellence in Retina Research. 2001
Johns Hopkins Medical Institutions Clinician Scientist Award. 1999
Stark Research Award in Ophthalmology, Wilmer Eye Institute, Johns Hopkins University. 1998
Fellow's Research Forum, runner-up. 1997
Knights Templar Eye Foundation Research Award. 1996
Magna Cum Laude, Harvard Medical School. 1995
First Bower Award for Research on Macular Degeneration Wills Eye Hospital, Philadelphia. 1993
Reed Scholar, M.I.T.. 1993 -1994
Johnson & Johnson Scholar, M.I.T.. 1991 -1992
Chinese National Scholarship for Graduate Studies Abroad. 1984
Highest Scholarships and High Distinction in Biochemistry, Sichuan Univ. 1980 -1984

Professional Society Membership

The Royal Society of Medicine
American Academy of Ophthalmology
American Institute for Medical and Biological Engineering
American Association for the Advancement of Science
American Society for Clinical Investigation
Association for Research in Vision and Ophthalmology
Association of American Physicians
International Society for Stem Cell Research

Journal Articles:

Yang JY, Wang W, Chen ZM, Lu SY, Yang FL, Bi ZF, Bao LL, Mo F, Li X, Huang Y, Hong WQ, Yang Y, Zhao Y, Ye F, Lin S, Deng W, Chen H, Lei H, Zhang ZQ, Luo M, Gao H, Zheng Y, Gong YQ, Jiang XH, Xu YF, Lv Q, Li D, Wang MN, Li FD, Wang SY, Wang GP, Yu P, Qu YJ, Yang L, Deng HX, Tong AP, Li J, Wang ZL, Yang JL, Shen GB, Zhao ZW, Li YH, Luo JW, Liu HQ, Yu WH, Yang ML, Xu JW, Wang JB, Li HY, Wang HX, Kuang DX, Lin PP, Hu ZT, Guo W, Cheng W, He YL, Song XR, Chen C, Xue ZH, Yao SH, Chen L, Ma XL, Chen SY, Gou ML, Huang WJ, Wang YC, Fan CF, Tian ZX, Shi M, Wang FS, Dai LZ, Wu M, Li G, Wang GY, Peng Y, Qian ZY, Huang CH, Lau J YN, Yang ZL, Wei YQ, Cen XB, Peng XZ, Qin C, **Zhang K**, Lu GW & Wei XW. (2020). A vaccine targeting the RBD of the S protein of SARS-CoV-2 induces protective immunity. *Nature*, 586, 572-577. DOI: 10.1038/s41586-020-2599-8. Corresponding author.

Xu X, Sun J, Nie S, Li HY, Kong YZ, Liang M, Hou JL, Huang XZ, Li DF, Ma T, Peng JQ, Gao SK, Shao Y, Zhu H, Lau J YN, Wang GY, Xie CB, Jiang L, Huang AL, Yang ZL, **Zhang K** & Hou FF. (2020). Seroprevalence of Immunoglobulin M and G Antibodies Against SARS-CoV-2 in China. *Nature Medicine*, 26, 1193–1195. DOI: 10.1038/s41591-020-0949-6. Corresponding author.

Zhang K, Liu XH, Shen J, Li ZH, Sang Y, Wu XW, Zha YF, Liang WH, Wang CD, Wang K, Ye LS, Gao M, Zhou ZG, Li L, Wang J, Yang ZH, Cai HM, Xu J, Yang L, Cai WJ, Xu WQ, Wu SX, Zhang W, Jiang SP, Zheng LH, Zhang X, Wang L, Lu L, Li JM, Yin HP, Wang W, Li O, Zhang C, Liang L, Wu T, Deng RY, Kang W, Zhou Y, Chen T, Lau J YN, Fok M, He JX, Lin TX, Li WM & Wang GY. (2020). Clinically Applicable AI System for Accurate Diagnosis, Quantitative Measurements and Prognosis of COVID-19 Pneumonia Using Computed Tomography. *Cell*. 181, 1423-1433. DOI: 10.1016/j.cell.2020.04.045. Corresponding and lead author.

Jiang H, Ou ZY, He YY, Yu MX, Wu SQ, Li G, Zhu J, Zhang R, Wang JY, Zheng LH, Zhang XH, Hao W, He LY, Gu XQ, Quan QL, Zhang E, Luo HY, Wei W, Li ZH, Zang GX, Zhang C, Poon T, Zhang D, Ziyar I, Zhang RZ, Li OL, Cheng LH, Shimizu T, Cui XP, Zhu JK, Sun X & **Zhang K**. (2020). DNA methylation markers in the diagnosis and prognosis of common leukemias. *Signal Transduction and targeted Therapy*, 5, 1-10. DOI: 10.1038/s41392-019-0090-5. Corresponding author

Xu Y, Li XF, Zhu B, Liang HY, Fang CX, Gong Y, Guo QZ, Sun X, Zhao DY, Shen J, Zhang HY, Liu HS, Xia HM, Tang JL, **Zhang K** & Gong ST. (2020). Characteristics of pediatric SARS-CoV-2 infection. *Nature Medicine*, 26, 1462-1465. DOI: 10.1038/s41591-020-0307-0. Corresponding author.

Huiying Liang, , Kang Zhang and Huimin Xia.(2019). Evaluation and accurate diagnoses of pediatric diseases using artificial intelligence. *Nature Medicine*. 25(3):433-438. *co-corresponding authors.

Kermany D, Goldbaum M, Cai W, Valentim CCS, Liang H, Baxter SL, McKeown A, Yang G, Wu X, Yan F, Dong J, Prasadha MK, Pei J, Ting M, Zhu J, Li C, Hewett S, Dong J, Ziyar I, Shi A, Zhu J, Ming C, Fu X, Duan Y, Hoang D, Rutgard J, Zhang R, Wang W, Hou R, Zhang D, Zhang E, Zhang C, Hao X, Xiong W, **Zhang, K.** (2017). Gene and mutation independent therapy via CRISPR-Cas9 mediated cellular reprogramming in rod photoreceptors. *Cell Research*. 27(6):830-833.

Xu RH, Wei W, Krawczyk M, Wang W, Luo H, Flagg K, Yi S, Shi W, Quan Q, Li K, Zheng L, Zhang H, Caughey BA, Zhao Q, Hou J, Zhang R, Xu Y, Cai H, Li G, Hou R, Zhong Z, Lin D, Fu X, Zhu J, Duan Y, Yu M, Ying B, Zhang W, Wang J, Zhang E, Zhang C, Li O, Guo R, Carter H, Zhu JK, Hao X, **Zhang K.** (2017). Circulating tumour DNA methylation markers for diagnosis and prognosis of hepatocellular carcinoma. *Nat Materials*. 16(11):1155-1161.

Hao X, Luo H, Krawczyk M, Wei W, Wang W, Wang J, Flagg K, Hou J, Zhang H, Yi S, Jafari M, Lin D, Chung C, Caughey BA, Li G, Dhar D, Shi W, Zheng L, Hou R, Zhu J, Zhao L, Fu X, Zhang E, Zhang C, Zhu JK, Karin M, Xu RH, **Zhang K.** (2017). DNA methylation markers for diagnosis and prognosis of common cancers. *Proc Natl Acad Sci USA*. Proc Natl Acad Sci U S A. 114(28):7414-7419.

Suzuki K, Tsunekawa Y, Hernandez-Benitez R, Wu J, Zhu J, Kim EJ, Hatanaka F, Yamamoto M, Araoka T, Li Z, Kurita M, Hishida T, Li M, Aizawa E, Guo S, Chen S, Goebel A, Soligalla RD, Qu J, Jiang T, Fu X, Jafari M, Esteban CR, Berggren WT, Lajara J, Nuñez-Delicado E, Guillen P, Campistol JM, Matsuzaki F, Liu GH, Magistretti P, Zhang K, Callaway EM, **Zhang K,** and Belmonte JCI. (2016). In vivo genome editing via CRISPR/Cas9 mediated homology-independent targeted integration. *Nature*. 540:144-149.

Hodgson N, Wu F, Zhu J, Wang W, Ferreyra H, **Zhang K,** Wang J. Economic and Quality of Life Benefits of Anti-VEGF Therapy. (2016). *Molecular pharmaceuticals*. 13(9):2877-80.

Wang WQ, Wang FH, Qin WX, Liu HY, Lu B, Chung C, Zhu J, Gu Q, Shi W, Wen C, Wu F, **Zhang K,** Sun XD. (2016). Joint Antiangiogenic Effect of ATN-161 and Anti-VEGF Antibody in a Rat Model of Early Wet Age-Related Macular Degeneration. *Molecular pharmaceuticals*. 13(9):2881-90.

Zhu W, Ma X, Gou M, Mei D, **Zhang K,** Chen S. (2016). 3D printing of functional biomaterials for tissue engineering. *Current opinion in biotechnology*. 40:103-12.

Bhisitkul RB, Desai SJ, Boyer DS, Sadda SR, **Zhang K.** Fellow Eye Comparisons for 7-Year Outcomes in Ranibizumab-Treated AMD Subjects from ANCHOR, MARINA, and HORIZON (SEVEN-UP Study). (2016). *Ophthalmology*. 123(6):1269-77

Bailey JN, Loomis SJ, Kang JH, Allingham RR, Gharahkhani P, Khor CC, Burdon KP, Aschard H, Chasman DI, Igo RP Jr, Hysi PG, Glastonbury CA, Ashley-Koch A, Brilliant M, Brown AA, Budenz DL, Buil A, Cheng CY, Choi H, Christen WG, Curhan G, De Vivo I, Fingert JH, Foster Ghazi NG, Abboud EB, Nowilaty SR, Alkuraya H, Alhommadi A, Cai H, Hou R, Deng WT, Boye SL, Almaghamsi A, Al Saikhan F, Al-Dhibi H, Birch D, Chung C, Colak D, LaVail MM, Vollrath Fritsche, LG, Igl, W, Bailey, JNC, Grassmann, F, Sengupta, S, Bragg-Gresham, JL, Burdon, KP, Hebbbring, SJ, Wen, C, Gorski, M, Kim, IK, Cho, D, Zack, D, Souied, E, Scholl, HPN, Bala, E, Lee, KE, Hunter, DJ, Sardell, RJ, Mitchell, P, Merriam, JE, Cipriani, V, Hoffman, JD, Schick, T, Gross, AM, Jaeger, PA, Kreisberg, JF, Licon, K, Jepsen, KL, Khosroheidari, M, Morsey, BM, Swindells, S, Shen, H, Ng, CT, Flagg, K, Chen, D, **Zhang, K,** Fox, HS, Ideker, T. (2016). Methylome-wide Analysis of Chronic HIV Infection Reveals Five-Year Increase in Biological Age

Lin H, Ouyang H, Zhu J, Huang S, Liu Z, Chen S, Cao G, Li G, Signer R, Xu Y, Chung C, Zhang Y, Lin D, Patel S, Wu F, Cai H, Hou J, Wen C, Jarafi M, Liu X, Luo L, Zhu J, Qiu A, Hou R, Chen B, Chen J, Granet D, Heichel C, Shang F, Li X, Krawczyk M, Skowronska-Krawczyk D, Wang Y, Shi W, Chen D, Zhong Z, Zhong S, Zhang L, Chen S, Morrison SJ, Maas RL, **Zhang K***, Liu Y*. (2016). Lens Regeneration using endogenous stem cells with gain of visual function. *Nature* 531:323-328.

Liu Y, Wu F, Lu L, Lin D, **Zhang K**. (2015). VIDEOS IN CLINICAL MEDICINE. Examination of the Retina. *N Engl J Med*. 373:e9. doi: 10.1056

Skowronska-Krawczyk D, Zhao L, Zhu J, Weinreb RN, Cao G, Luo J, Flagg K, Patel S, Wen C, Krupa M, Luo H, Ouyang H, Lin D, Wang W, Li G, Xu Y, Li O, Chung C, Yeh E, Jafari M, Ai M, Zhong Z, Shi W, Zheng L, Krawczyk M, Chen D, Shi C, Zin C, Zhu J, Mellon PL, Gao W, Abagyan R, Zhang L, Sun X, Zhong S, Zhuo Y, Rosenfeld MG, Liu Y, **Zhang K** (2015). P16INK4a Upregulation Mediated by SIX6 Defines Retinal Ganglion Cell Pathogenesis in Glaucoma. *Molecular Cell*. 59:931-40.

Hu, CM, Fang, RH, Wang, KC, Luk, BT, Thamphiwatana, S, Dehaini, D, Nguyen, P, Angsantikul, P, Wen, CH, Kroll, AV, Carpenter, C, Ramesh, M, Qu, V, Patel, S, Zhu, J, Hofman, FM, Chen, Zhao L, Chen, XJ, Zhu, J, Xi, YB, Yang, X, Hu, LD, Ouyang, H, Patel, SH, Jin, X, Lin, D, Wu, F, Flagg, K, Cai H, Li, G, Cao, G, Lin, Y, Chen, D, Wen, C, Chung, C, Wang, YD, Qiu, A, Yeh, E, Salvo J, Lyubasyuk V, Xu M, Wang H, Wang F, Nguyen D, Wang K, Luo H, Wen C, Shi C, Lin Huang LZ, Li YJ, Xie XF, Zhang JJ, Cheng CY, Yamashiro K, Chen LJ, Ma XY, Cheung CM, Wang YS, Zhang CF, Bai YJ, Hou J, Chen XL, Qi Y, Li SS, Sun YY, Mei JP, Cheng Y, Yu WZ, Hu XB, Zhuang FF, Fan L, Lu Y, Sun XH, Zhu XJ, Shen DF, Chan CC, Zhao MW, Yoshimura N, Bhisitkul RB, Mendes TS, Rofagha S, Enanoria W, Boyer DS, Sadda SR, **Zhang K**. Macular Atrophy Progression and 7-Year Vision Outcomes in Subjects From the ANCHOR, MARINA, and HORIZON Studies: the SEVEN-UP Study. (2015). *Am J Ophthalmol*. 159:915-924.

Huu VA, Luo J, Zhu J, Zhu J, Patel S, Boone A, Mahmoud E, McFearin C, Olejniczak J, de Gracia Lux C, Lux J, Fomina N, Huynh M, **Zhang K**, Almutairi A. (2015). Light-responsive nanoparticle depot to control release of a small molecule angiogenesis inhibitor in the posterior segment of the eye. *J Control Release*. 200:71-7.

Copp JA, Fang RH, Luk BT, Hu CM, Gao W, Zhang K, Zhang L. (2014). Clearance of pathological antibodies using biomimetic nanoparticles. *Proc Natl Acad Sci U S A*. 111:13481-6.

Kang X, Xu H, Teng S, Zhang X, Deng Z, Zhou L, Zuo P, Liu B, Liu B, Wu Q, Wang L, Hu M, Dou H, Liu W, Zhu F, Li Q, Guo S, Gu J, Lei Q, Lü J, Mu Y, Jin M, Wang S, Jiang W, Liu K, Wang C, Li W, **Zhang K**, Zhou Z. (2014). Dopamine release from transplanted neural stem cells in Parkinsonian rat striatum in vivo. *Proc Natl Acad Sci U S A*. 111:15804-9.

Wang F, Wang Y, Zhang B, Zhao L, Lyubasyuk V, Wang K, Xu M, Li Y, Wu F, Wen C, Bernstein PS, Lin D, Zhu S, Wang H, **Zhang K**, Chen R. (2014). A missense mutation in HK1 leads to autosomal dominant retinitis pigmentosa. *Invest Ophthalmol Vis Sci*. 55:7159-64.

Chi W, Li F, Chen H, Wang Y, Zhu Y, Yang X, Zhu J, Wu F, Ouyang H, Ge J, Weinreb RN, **Zhang K**, Zhuo Y. (2014). Caspase-8 promotes NLRP1/NLRP3 inflammasome activation and IL-1 β production in acute glaucoma. *Proc Natl Acad Sci U S A*. 111:11181-6.

Ouyang, H, Xue, Y, Lin, Y, Zhang, X, Xi, L, Patel, S, Gen Li, Jing Luo, Wei Jiang, Yang Y, Li, H, Zhang, M, Cai, G, Yeh, E, Pei, M, Cao, G, Zhang, L, Yu, B, Chen, S, Fu, XD, Liu, Y, **Zhang K**. (2014). WNT7A and PAX6 define corneal epithelium homeostasis and pathogenesis. *Nature* 511:358-61.

Liao C, Yin, A, Peng CF, Fu F, Yang JX, Li R, Chen YY, Luo DH, Zhang YL, Ou YM, Li J, Wu J, Mai MQ, Hou R, Wu F, Luo H, Li DZ, Liu, HL, Zhang XZ, **Zhang K**. (2014). Noninvasive prenatal diagnosis of common aneuploidies by semiconductor sequencing, PNAS, 111:7415-20.

Gou, M, Qu, X, Zhu, W, Xiang, M, Yang, J, **Zhang, K**, Wei Y & Chen S (2014). Bio-inspired detoxification using 3D-printed hydrogel nanocomposites. Nature Communications, 5:3774.

Yehoshua Z, Alexandre de Amorim Garcia Filho C, Nunes RP, Gregori G, Penha FM, Moshfeghi AA, **Zhang K**, Sadda S, Feuer W, Rosenfeld PJ. (2014). Systemic Complement Inhibition with Eculizumab for Geographic Atrophy in Age-Related Macular Degeneration: The COMPLETE Study. Ophthalmology. 121(3):693-701.

Wang F, Wang H, Tuan HF, Nguyen DH, Sun V, Keser V, Bowne SJ, Sullivan LS, Luo H, Zhao L, Wang X, Zaneveld JE, Salvo JS, Siddiqui S, Mao L, Wheaton DK, Birch DG, Branham KE, Heckenlively JR, Wen C, Flagg K, Ferreyra H, Pei J, Khan A, Ren H, Wang K, Lopez I, Qamar R, Zenteno JC, Ayala-Ramirez R, Buentello-Volante B, Fu Q, Simpson DA, Li Y, Sui R, Silvestri G, Daiger SP, Koeneke RK, **Zhang K***, Chen R*. (2014) Next generation sequencing-based molecular diagnosis of retinitis pigmentosa: identification of a novel genotype-phenotype correlation and clinical refinements. Hum Genet. 133(3):331-45. *co-corresponding authors.

Zhao JJ, Ouyang H, Luo J, Patel S, Xue Y, Quach J, Sfeir N, Zhang M, Fu X, Ding S, Chen S, **Zhang K**. (2014). Induction of retinal progenitors and neurons from mammalian Muller glia under defined conditions. J Biol Chem. 289(17):11945-51.

Yu FX, Luo J, Mo JS, Liu G, Kim YC, Meng Z, Zhao L, Peyman G, Ouyang H, Jiang W, Zhao J, Chen X, Zhang L, Wang CY, Bastian BC, **Zhang K***, Guan KL*. (2014). Mutant Gq/11 promote uveal melanoma tumorigenesis by activating YAP. Cancer Cell. 25:822-30.

Wang D, Zhou J, Hou X, Nguyen DH, Cao G, Li G, Qiu G, **Zhang K**, Zhang M, Su Z. (2015). CETP Gene may be Associated with Advanced Age-related Macular Degeneration in the Chinese Population. Ophthalmic Genet. 36(4):303-8

Loomis SJ, Kang JH, Weinreb RN, Yaspan BL, Cooke Bailey JN, Gaasterland D, Gaasterland T, Lee RK, Lichter PR, Budenz DL, Liu Y, Realini T, Friedman DS, McCarty CA, Moroi SE, Olson L, Schuman JS, Singh K, Vollrath D, Wollstein G, Zack DJ, Brilliant M, Sit AJ, Christen WG, Fingert J, Kraft P, **Zhang K**, Allingham RR, Pericak-Vance MA, Richards JE, Hauser MA, Haines JL, Pasquale LR, Wiggs JL. (2014). Association of CAV1/CAV2 genomic variants with primary open-angle glaucoma overall and by gender and pattern of visual field loss. Ophthalmology. 121:508-16.

Garcia Filho CA, Yehoshua Z, Gregori G, Nunes RP, Penha FM, Moshfeghi AA, **Zhang K**, Feuer W, Rosenfeld PJ. (2014). Change in drusen volume as a novel clinical trial endpoint for the study of complement inhibition in age-related macular degeneration. Ophthalmic Surg Lasers Imaging Retina. 45(1):18-31.

Luo J, Baranov P, Patel S, Ouyang H, Quach J, Wu F, Qiu A, Luo H, Hicks C, Zeng J, Zhu J, Lu J, Sfeir N, Wen C, Zhang M, Reade V, Patel S, Sinden J, Sun X, Shaw P, Young M, **Zhang K**. (2014). Human Retinal Progenitor Cell Transplantation Preserves Vision. J Biol Chem. 289(10):6362-71.

X. Qu, W. Zhu, S. Huang, J. Y. Li, S. Chien, K. Zhang, S.C. Chen, (2013). Relative impact of uniaxial alignment vs. form-induced stress on differentiation of human adipose derived stem cells. Biomaterials, 11:9812-9818.

Zhao L, Patel SH, Pei J, **Zhang K**. (2013). Antagonizing Wnt pathway in diabetic retinopathy. Diabetes. 62(12):3993-5.

Luo J, Zhao L, Chen AY, Zhang X, Zhu J, Zhao J, Ouyang H, Luo H, Song Y, Lee J, Patel SH, Shaw PX, Sadda S, Zhuo Y, Rosenfeld MG, **Zhang K.** (2013). TCF7L2 Variation and Proliferative Diabetic Retinopathy. *Diabetes*. 62(7):2613-7.

Zhan X, Larson DE, Wang C, Koboldt DC, Sergeev YV, Fulton RS, Fulton LL, Fronick CC, Branham KE, Bragg-Gresham J, Jun G, Hu Y, Kang HM, Liu D, Othman M, Brooks M, Ratnapriya R, Boleda A, Grassmann F, von Strachwitz C, Olson LM, Buitendijk GH, Hofman A, van Duijn CM, Cipriani V, Moore AT, Shahid H, Jiang Y, Conley YP, Morgan DJ, Kim IK, Johnson MP, Cantsilieris S, Richardson AJ, Guymer RH, Luo H, Ouyang H, Licht C, Pluthero FG, Zhang MM, **Zhang K**, Baird PN, Blangero J, Klein ML, Farrer LA, DeAngelis MM, Weeks DE, Gorin MB, Yates JR, Klaver CC, Pericak-Vance MA, Haines JL, Weber BH, Wilson RK, Heckenlively JR, Chew EY, Stambolian D, Mardis ER, Swaroop A, Abecasis GR. (2013) Identification of a rare coding variant in complement 3 associated with age-related **macular degeneration**. *Nat Genet*. 45(11):1375-9.

Helgason H, Sulem P, Duvvari MR, Luo H, Thorleifsson G, Stefansson H, Jonsdottir I, Masson G, Gudbjartsson DF, Walters GB, Magnusson OT, Kong A, Rafnar T, Kiemeny LA, Schoenmaker-Koller FE, Zhao L, Boon CJ, Song Y, Fauser S, Pei M, Ristau T, Patel S, Liakopoulos S, van de Ven JP, Hoyng CB, Ferreyra H, Duan Y, Bernstein PS, Geirsdottir A, Helgadottir G, Stefansson E, den Hollander AI, **Zhang K**, Jonasson F, Sigurdsson H, Thorsteinsdottir U, Stefansson K. (2013). A rare nonsynonymous sequence variant in C3 is associated with high risk of age-related macular degeneration. *Nat Genet*. 45(11):1371-4.

Hu CM, Fang RH, Luk BT, Chen KN, Carpenter C, Gao W, **Zhang K**, Zhang L. (2013). 'Marker-of-self' functionalization of nanoscale particles through a top-down cellular membrane coating approach. *Nanoscale*. 5(7):2664-8.

Ouyang H, Zhuo Y, **Zhang K.** (2013). WNT signaling in stem cell differentiation and tumor formation. *J Clin Invest*. 123(4):1422-4.

Fritsche LG, Chen W, Schu M, Yaspan BL, Yu Y, Thorleifsson G, Zack DJ, Arakawa S, Cipriani V, Ripke S, Igo RP Jr, Buitendijk GH, Sim X, Weeks DE, Guymer RH, Merriam JE, Francis PJ, Hannum G, Agarwal A, Armbricht AM, Audo I, Aung T, Barile GR, Benchaboune M, Bird AC, Bishop PN, Branham KE, Brooks M, Brucker AJ, Cade WH, Cain MS, Campochiaro PA, Chan CC, Cheng CY, Chew EY, Chin KA, Chowers I, Clayton DG, Cojocaru R, Conley YP, Cornes BK, Daly MJ, Dhillon B, Edwards AO, Evangelou E, Fagerness J, Ferreyra HA, Friedman JS, Geirsdottir A, George RJ, Gieger C, Gupta N, Hagstrom SA, Harding SP, Haritoglou C, Heckenlively JR, Holz FG, Hughes G, Ioannidis JP, Ishibashi T, Joseph P, Jun G, Kamatani Y, Katsanis N, N Keilhauer C, Khan JC, Kim IK, Kiyohara Y, Klein BE, Klein R, Kovach JL, Kozak I, Lee CJ, Lee KE, Lichtner P, Lotery AJ, Meitinger T, Mitchell P, Mohand-Said S, Moore AT, Morgan DJ, Morrison MA, Myers CE, Naj AC, Nakamura Y, Okada Y, Orlin A, Ortube MC, Othman MI, Pappas C, Park KH, Pauer GJ, Peachey NS, Poch O, Priya RR, Reynolds R, Richardson AJ, Ripp R, Rudolph G, Ryu E, Sahel JA, Schaumberg DA, Scholl HP, Schwartz SG, Scott WK, Shahid H, Sigurdsson H, Silvestri G, Sivakumaran TA, Smith RT, Sobrin L, Souied EH, Stambolian DE, Stefansson H, Sturgill-Short GM, Takahashi A, Tosakulwong N, Truitt BJ, Tsironi EE, Uitterlinden AG, van Duijn CM, Vijaya L, Vingerling JR, Vithana EN, Webster AR, Wichmann HE, Winkler TW, Wong TY, Wright AF, Zelenika D, Zhang M, Zhao L, **Zhang K**, Klein ML, Hageman GS, Lathrop GM, Stefansson K, Allikmets R, Baird PN, Gorin MB, Wang JJ, Klaver CC, Seddon JM, Pericak-Vance MA, Iyengar SK, Yates JR, Swaroop A, Weber BH, Kubo M, Deangelis MM, Léveillard T, Thorsteinsdottir U, Haines JL, Farrer LA, Heid IM, Abecasis GR; AMD Gene Consortium. (2013). Seven new loci associated with age-related macular degeneration. *Nat Genet*. 45(4):433-9.

Zhu J, Nguyen D, Ouyang H, Zhang XH, Chen XM, **Zhang K.** (2013). Inhibition of RhoA/Rho-kinase pathway suppresses the expression of extracellular matrix induced by CTGF or TGF- β in ARPE-19. *Int J Ophthalmol.* 6(1):8-14. doi: 10.3980/j.issn.2222-3959.2013.01.02. Epub 2013 Feb 18.

Rofagha S, Bhisitkul RB, Boyer DS, Sadda SR, **Zhang K;** SEVEN-UP Study Group. (2013). Seven-Year Outcomes in Ranibizumab-Treated Patients in ANCHOR, MARINA, and HORIZON: A Multicenter Cohort Study (SEVEN-UP). Seven-Year Outcomes in Ranibizumab-Treated Patients in ANCHOR, MARINA, and HORIZON: A Multicenter Cohort Study (SEVEN-UP). *Ophthalmology.* pii: S0161-6420(13)00331-X. doi: 10.1016/j.opthta.2013.03.046. [Epub ahead of print]

Nguyen DH, Luo J, **Zhang K,** Zhang M. (2013). Current therapeutic approaches in neovascular age-related macular degeneration. *Discov Med.* 2013 Jun;15(85):343-8.

Zhou J, Wang D, Zhang J, Zhang M, Lu F, Qiu G, Zhao L, Nguyen DH, Luo H, Cao G, Zhang W, Jiang W, Li G, **Zhang K,** Zhang M, Su Z. (2013). RAD51 gene is associated with advanced age-related macular degeneration in Chinese population. *Clin Biochem.* pii: S0009-9120(13)00335-4. doi: 10.1016/j.clinbiochem.2013.07.002. [Epub ahead of print]

Pasquale LR, Loomis SJ, Weinreb RN, Kang JH, Yaspan BL, Bailey JC, Gaasterland D, Gaasterland T, Lee RK, Scott WK, Lichter PR, Budenz DL, Liu Y, Realini T, Friedman DS, McCarty CA, Moroi SE, Olson L, Schuman JS, Singh K, Vollrath D, Wollstein G, Zack DJ, Brilliant M, Sit AJ, Christen WG, Fingert J, Kraft P, **Zhang K,** Allingham RR, Pericak-Vance MA, Richards JE, Hauser MA, Haines JL, Wiggs JL. (2013). Estrogen pathway polymorphisms in relation to primary open angle glaucoma: An analysis accounting for gender from the United States. *Mol Vis.* 19:1471-81.

Zhao, L, Grob, S, Avery, Kimura, Pieramici, D, Lee, J, Rabena, M, Hughes, G, Ortiz, S, Tornambe, P, Goldbaum, M, Ferreyra, H, Kozak, I, and **Zhang, K.** (2013). Common Variant in *VEGFA* and Response to Anti-VEGF Therapy for Exudative Age-Related Macular Degeneration. *Lu, B., C. Morgans, S. Girman, J. Luo, J. Zhao, H. Du, S. Lim, S. Ding, C. Svendsen, K. Zhang, and S. Wang* (2013). Neural stem cells derived by small molecules preserve vision. *Translational Vision Science & Technology.* T: January 2013, Vol. 2, No. 1.

Du H, Sun X, Guma M, Luo J, Ouyang H, Zhang X, Zeng J, Quach J, Nguyen DH, Shaw PX, Karin M, Zhang K. (2013). JNK inhibition reduces apoptosis and neovascularization in a murine model of age-related macular degeneration. *Proc Natl Acad Sci U S A.* 110:2377-82.

Xue, Y-C., Ouyang, K., Huang, J., Zhou, Y., Ouyang, H., Li, H., Wang, G., Wu, Q., Wei, C., Bi, Y., Jiang, L., Cai, Z., Sun, H., **Zhang, K.,** Zhang, Y., Chen, J., and Fu, X-D. (2012) Direct conversion of fibroblasts to neurons by reprogramming PTB-regulated microRNA circuits. *Cell* 152:82-96.

Hannum G, Guinney J, Zhao L, Zhang L, Hughes G, Sadda S, Klotzle B, Bibikova M, Fan JB, Gao Y, Deconde R, Chen M, Rajapakse I, Friend S, Ideker T, **Zhang K.** (2013). Genome-wide Methylation Profiles Reveal Quantitative Views of Human Aging Rates. *Mol Cell,* 49:359-67.

Zhuo Y, Luo H, Zhang K. (2012). Leber hereditary optic neuropathy and oxidative stress. *Proc Natl Acad Sci U S A.* 109(49):19882-3.

Zhao J, Sun W, Cho HM, Ouyang H, Li W, Lin Y, Do J, Zhang L, Ding S, Liu Y, Lu P, **Zhang K.** (2013). Integration and long distance axonal regeneration in CNS from transplanted primitive neural stem cells. *J Biol Chem.* 288(1):164-8. doi: 10.1074/jbc.M112.433607.

Fang RH, Chen KN, Aryal S, Hu CM, **Zhang K**, Zhang L. (2012). Large-scale synthesis of lipid-polymer hybrid nanoparticles using a multi-inlet vortex reactor. *Langmuir*. 39:13824-9.

Deng WT, Dinculescu A, Li Q, Boye SL, Li J, Gorbatyuk MS, Pang J, Chiodo VA, Matthes MT, Yasumura D, Liu L, Alkuraya FS, **Zhang K**, Vollrath D, LaVail MM, Hauswirth WW. (2012). Tyrosine-mutant AAV8 delivery of human MERTK provides long-term retinal preservation in RCS rats. *Invest Ophthalmol Vis Sci*. 53(4):1895-904.

Wiggs JL, Yaspan BL, Hauser MA, Kang JH, Allingham RR, Olson LM, Abdrabou W, Fan BJ, Wang DY, Brodeur W, Budenz DL, Caprioli J, Crenshaw A, Crooks K, Delbono E, Doheny KF, Friedman DS, Gaasterland D, Gaasterland T, Laurie C, Lee RK, Lichter PR, Loomis S, Liu Y, Medeiros FA, McCarty C, Mirel D, Moroi SE, Musch DC, Realini A, Rozsa FW, Schuman JS, Scott K, Singh K, Stein JD, Trager EH, Vanveldhuisen P, Vollrath D, Wollstein G, Yoneyama S, **Zhang K**, Weinreb RN, Ernst J, Kellis M, Masuda T, Zack D, Richards JE, Pericak-Vance M, Pasquale LR, Haines JL. (2012). Common variants at 9p21 and 8q22 are associated with increased susceptibility to optic nerve degeneration in glaucoma. *PLoS Genet*. 8(4):e1002654.

Ulmer M, Li J, Yaspan BL, Ozel AB, Richards JE, Moroi SE, Hawthorne F, Budenz DL, Friedman DS, Gaasterland D, Haines J, Kang JH, Lee R, Lichter P, Liu Y, Pasquale LR, Pericak-Vance M, Realini A, Schuman JS, Singh K, Vollrath D, Weinreb R, Wollstein G, Zack DJ, **Zhang K**, Young T, Allingham RR, Wiggs JL, Ashley-Koch A, Hauser MA. (2012). Genome-wide analysis of central corneal thickness in primary open-angle glaucoma cases in the NEIGHBOR and GLAUGEN consortia. *Invest Ophthalmol Vis Sci*. 53(8):4468-74

Wiggs JL, Hauser MA, Abdrabou W, Allingham RR, Budenz DL, Delbono E, Friedman DS, Kang JH, Gaasterland D, Gaasterland T, Lee RK, Lichter PR, Loomis S, Liu Y, McCarty C, Medeiros FA, Moroi SE, Olson LM, Realini A, Richards JE, Rozsa FW, Schuman JS, Singh K, Stein JD, Vollrath D, Weinreb RN, Wollstein G, Yaspan BL, Yoneyama S, Zack D, **Zhang K**, Pericak-Vance M, Pasquale LR, Haines JL. (2012). The NEIGHBOR Consortium Primary Open-Angle Glaucoma Genome-wide Association Study: Rationale, Study Design, and Clinical Variables. *J Glaucoma* 22(7):517-525.

Pasquale LR, Loomis SJ, Kang JH, Yaspan BL, Abdrabou W, Budenz DL, Chen TC, Delbono E, Friedman DS, Gaasterland D, Gaasterland T, Grosskreutz CL, Lee RK, Lichter PR, Liu Y, McCarty CA, Moroi SE, Olson LM, Realini T, Rhee DJ, Schuman JS, Singh K, Vollrath D, Wollstein G, Zack DJ, Allingham RR, Pericak-Vance MA, Weinreb RN, **Zhang K**, Hauser MA, Richards JE, Haines JL, Wiggs JL. (2013). CDKN2B-AS1 Genotype-Glaucoma Feature Correlations in Primary Open-Angle Glaucoma Patients From the United States. *Am J Ophthalmol*. 155(2):242-252.e5.

Shaw P, Zhang L, Zhang M, Du H, Zhao L, Lee C, Grob S, Lim SL, Hughes G, Lee J, Bedell M, Nelson MH, Lu F, Krupa M, Luo J, Ouyang H, Tu Z, Su Zhiguang, Zhu J, Wei X, Feng Z, Duan Y, Yang Z, Ferreyra H, Bartsch DU, Kozak I, Zhang L, Lin F, Sun H, Feng H, **Zhang K**. (2012). Complement factor H genotypes impact risk of age-related macular degeneration by interaction with oxidized phospholipids. *PNAS* 109:13757-13762.

Harkewicz R, Du H, Tong Z, Alkuraya H, Bedell M, Sun W, Wang X, Hsu YH, Esteve-Rudd J, Hughes G, Su Z, Zhang M, Lopes VS, Molday RS, Williams DS, Dennis EA, Zhang K. (2011). Essential role of ELOVL4 in very long chain fatty acid synthesis and retinal function. *J Biol Chem*.

Sun F, Park KK, Belin S, Wang D, Lu T, Chen G, Zhang K, Yeung C, Feng G, Yankner BA, He Z. (2011). Sustained axon regeneration induced by co-deletion of PTEN and SOCS3. *Nature* 480:372-375.

Zhang L, Lim SL, Du H, Zhang M, Kozak I, Hannum G, Wang X, Ouyang H, Hughes G, Zhao L, Zhu X, Lee C, Su Z, Zhou X, Shaw R, Geum D, Wei X, Zhu J, Ideker T, Oka C, Wang N, Yang Z, Shaw PX, Zhang K. HTRA1 regulates angiogenesis through TF- β family member GDF6. *J Biol Chem*. 2012 Jan 6;287(2):1520-6. Epub 2011 Nov 2.

Nacev BA, Low WK, Huang Z, Su TT, Su Z, Alkuraya H, Kasuga D, Sun W, Träger M, Braun M, Fischer G, Zhang K, Liu JO. A calcineurin-independent mechanism of angiogenesis inhibition by a nonimmunosuppressive cyclosporin A analog. *J Pharmacol Exp Ther*. 2011 Aug;338(2):466-75. Epub 2011 May 11.

Carbone MA, Chen Y, Hughes GA, Weinreb RN, Zabriskie NA, Zhang K, Anholt RR. Genes of the unfolded protein response pathway harbor risk alleles for primary open angle glaucoma. *PLoS One*. 2011;6(5):e20649. Epub 2011 May 31

Jones A, Kumar S, Zhang N, Tong Z, Yang JH, Watt C, Anderson J, Amrita, Fillerup H, McCloskey M, Luo L, Yang Z, Ambati B, Marc R, Oka C, Zhang K, Fu Y. (2011). Increased expression of multifunctional serine protease, HTRA1, in retinal pigment epithelium induces polypoidal choroidal vasculopathy in mice. *Proc Natl Acad Sci U S A*. 2011 Aug 20;108(35):14578-82. Epub 2011 Aug 15

Luo H, Hu X, Liu X, Ma X, Guo W, Qiu C, Wang Y, Wang Q, Zhang X, Zhang W, Hannum G, Zhang K, Liu X, Li T (2012). Hair cortisol level as a biomarker for altered hypothalamic-pituitary-adrenal activity in female adolescents with posttraumatic stress disorder after the 2008 Wenchuan earthquake. *Biol Psychiatry*, 1;72(1):65-9.

Zhang K*, Harada Y, Wei X, Shukla D, Rajendran A, Tawansy K, Bedell M, Lim S, Shaw PX, He X*, Yang Z* (2011). An essential role of the cysteine-rich domain of FZD4 in Norrin/Wnt signaling and familial exudative vitreoretinopathy. *J Biol Chem*. 25;286(12):10210-5.

Li W, Sun W, Zhang Y, Wei W, Ambasudhan R, Xia P, Talantova M, Lin T, Kim J, Wang X, Kim WR, Lipton SA, Zhang K*, Ding S*. (2011) Rapid induction and long-term self-renewal of primitive neural precursors from human embryonic stem cells by small molecule inhibitors. *Proc Natl Acad Sci U S A*. 108:8299-304. *co-corresponding author.

Kim J, Efe JA, Zhu S, Talantova M, Yuan X, Wang S, Lipton SA, Zhang K, Ding S. (2011). Direct reprogramming of mouse fibroblasts to neural progenitors. *Proc Natl Acad Sci U S A*. 108:7838-43.

Korn BS, Zhang K. . Carotid-cavernous sinus fistula. *N Engl J Med*. 2011 Feb 24;364(8):

Zhang, K*, Hopkins, JJ, Heier, JS, Birch, DG, Halperin, LS, Albini, TA, Brown, DM, Jaffe, GJ, Tao, W, and Williams, GA. (2011). Ciliary neurotrophic factor delivered by encapsulated cell intraocular implants for treatment of geographic atrophy in age-related macular degeneration. *PNAS* 108(15):6241-5. *corresponding author.

Chen, Y, Zeng, J, Zhao, C, Wang, K, Trood, E, Buehler, J, Weed, M, Bernstein, PS, Kasuga, D, Hughes, G, Fu, V, Lee, C, Crocker, M, Bedell, M, Salasar, F, Goldbaum, M, Ferrayra, H, Yang, Z, and **Zhang, K** (2010). Assessing Susceptibility to Age-Related Macular Degeneration with Genetic Markers and Environmental Factors. *Archives of Ophthalmology, Archives of Ophthalmol*. 2011;129(3):344-351.

Zhu, S, Li, W., Zhou, H., Wei, W., Ambasudhan, R., Lin, T., Kim, J., **Zhang, K.**, Ding, S. (2010). Reprogramming of Human Primary Somatic Cells by OCT4 and Chemical Compounds. *Cell Stem Cells* 7: 651-655

Yang X, Yamada K, Katz B, Guan H, Wang L, Andrews C, Zhao G, Engle EC, Chen H, Tong Z, Kong J, Hu C, Kong Q, Fan G, Wang Z, Ning M, Zhang S, Xu J, Zhang K. KIF21A mutations in two Chinese families with congenital fibrosis of the extraocular muscles (CFEOM). *Mol Vis*. 2010 Oct 13;16:2062-70.

Zhou H, Li W, Zhu S, Joo JY, Do JT, Xiong W, Kim JB, **Zhang K**, Schöler HR, Ding S. (2010). Conversion of mouse epiblast stem cells to an earlier pluripotency state by small molecules. *J Biol Chem*. 285(39):29676-80. Epub 2010 Aug 12.

Zeng J, Chen Y, Tong Z, Zhou X, Zhao C, Wang K, Hughes G, Kasuga D, Bedell M, Lee C, Ferreyra H, Kozak I, Haw W, Guan J, Shaw R, Stevenson W, Weishaar P, Nelson M, Tang L, **Zhang K** (2010). Lack of association of CFD polymorphisms with advanced age-related macular degeneration. *Molecular Vision*, in press.

Niu, Y, Yu Y, Bernat, A, Yang, S, He, X, Guo, X, Chen, D, Chen, Y, Ji, S, Si, W, Li, Y, Tan, T, Wei, Q, Wang, H, Shi, L, Guan, J, Zhu X, Afanassieff, M, Savatier, P*, **Zhang, K***, Zhou, Q*, Ji, W*. (2010) Transgenic rhesus monkeys produced by gene transfer into early cleavage stage embryos using a simian immunodeficiency virus-based vector. *PNAS*, in press. *co-corresponding authors.

Riazuddin S A, Iqbal M, Wang Yue, Masuda T, Chen Y, Bowne S, Sullivan L S, Waseem N H, Bhattacharya S, Daiger S P, **Zhang K**, Khan S N, Riazuddin S, Hejtmancik J F, Sieving P A, Zack D J, Katsanis N. A splice-site mutation in a retina-specific exon of BBS8 causes nonsyndromic retinitis pigmentosa. *American journal of human genetics* 2010;86(5):805-12.

Neale B M, Fagerness J, Reynolds R, Sobrin L, Parker M, Raychaudhuri S, Tan P L, Oh E C, Merriam J E, Souied E, Bernstein P S, Li B, Frederick J M, **Zhang K**, Brantley M A, Lee A Y, Zack D J, Campochiaro B, Campochiaro P, Ripke S, Smith R T, Barile G R, Katsanis N, Allikmets R, Daly M J, Seddon J M. Genome-wide association study of advanced age-related macular degeneration identifies a role of the hepatic lipase gene (LIPC). *Proceedings of the National Academy of Sciences of the United States of America* 2010;107(16):7395-400.

Michaelides M, Gaillard MC, Escher P, Tiab L, Bedell M, Borruat FX, Barthelmes D, Carmona R, Zhang K, White E, McClements M, Robson AG, Holder G, Bradshaw K, Hunt DM, Webster A, Moore AT, Schorderet D, Munier FL. The PROM1 mutation p.R373C causes an autosomal dominant bull's eye maculopathy associated with rod, rod-cone and macular dystrophy. *Invest Ophthalmol Vis Sci*. 2010 Apr 14. [Epub ahead of print]

Yang Z, Tong Z, Chen Y, Zeng J, Lu F, Sun X, Zhao C, Wang K, Davey L, Chen H, London N, Muramatsu D, Salazar F, Carmona R, Kasuga D, Wang X, Bedell M, Dixie M, Zhao P, Yang R, Gibbs D, Liu X, Li Y, Li C, Li Y, Campochiaro B, Constantine R, Zack DJ, Campochiaro P, Fu Y, Li DY, Katsanis N, Zhang K. (2010). Genetic and functional dissection of HTRA1 and LOC387715 in age-related macular degeneration. *PLoS Genet*. 6(2):e1000836.

Chen H, Zhang M, Tang S, London NR, Li DY, Zhang K. (2010). Slit-robo signaling in ocular angiogenesis. *Adv Exp Med Biol*. 664:457-63.

Kuny S, Gaillard F, Mema SC, Freund PR, Zhang K, Macdonald IM, Sparrow JR, Sauvé Y. (2010). Inner retina remodeling in a mouse model of stargardt-like macular dystrophy (STGD3). *Invest Ophthalmol Vis Sci.* 1:2248-62.

Bedell M, Harkewicz R, Wang X, Zhang K. (2010). Focus on molecules: ELOVL4. *Exp Eye Res.*:476-7.

Takeuchi T, Hayashi T, Bedell M, Zhang K, Yamada H, Tsuneoka H. (2009). A novel haplotype with the R345W mutation in the EFEMP1 gene associated with autosomal dominant drusen in a Japanese family. *Invest Ophthalmol Vis Sci.* 2010 Mar;51(3):1643-50.

Crow RW, Katz BJ, Warner JE, Alder SC, Zhang K, Schulman S, Digre KB. (2009). Giant cell arteritis and mortality. *J Gerontol A Biol Sci Med Sci.* 64:365-9.

Jiao X, Yang Z, Yang X, Chen Y, Tong Z, Zhao C, Zeng J, Chen H, Gibbs D, Sun X, Li B, Wakins WS, Meyer C, Wang X, Kasuga D, Bedell M, Pearson E, Weinreb RN, Leske MC, Hennis A, DeWan A, Nemesure B, Jorde LB, Hoh J, Hejtmancik JF, **Zhang K.** (2009). Common variants on chromosome 2 and risk of primary open-angle glaucoma in the Afro-Caribbean population of Barbados. *Proc Natl Acad Sci U S A.* 106:17105-10. Epub 2009 Sep 24.

Li F, Marchette LD, Brush RS, Elliott MH, Le YZ, Henry KA, Anderson AG, Zhao C, Sun X, Zhang K, Anderson RE. DHA does not protect ELOVL4 transgenic mice from retinal degeneration. *Mol Vis.* 2009 Jun 13;15:1185-93.

Jones CA, Nishiya N, London NR, Zhu W, Sorensen LK, Chan AC, Lim CJ, Chen H, Zhang Q, Schultz PG, Hayallah AM, Thomas KR, Famulok M, Zhang K, Ginsberg MH, Li DY. Slit2-Robo4 signaling promotes vascular stability by blocking Arf6 activity. *Nat Cell Biol.* 2009 Oct 18. [Epub ahead of print]

Brown DM, Michels M, Kaiser PK, Heier JS, Sy JP, Ianchulev T; ANCHOR Study Group. (2009). Ranibizumab versus verteporfin photodynamic therapy for neovascular age-related macular degeneration: Two-year results of the ANCHOR study. *Ophthalmology.* 116(1):57-65.

Li F, Marchette LD, Brush RS, Elliott MH, Le YZ, Henry KA, Anderson AG, Zhao C, Sun X, Zhang K, Anderson RE. (2009) DHA does not protect ELOVL4 transgenic mice from retinal degeneration. *Mol Vis.* 13;15:1185-93.

Yamashita, T, Liu, J, Gao, J, LeNoue, S, Wang, C, Kaminoh, J, Bowne, S, Sullivan, L, Daiger, S, **Zhang, K,** Fitzgerald, M, Kefalov, V, and Zuo, J. (2009). Essential and synergistic roles of RP1 and RP1L1 in rod photoreceptor axoneme and retinitis pigmentosa. *Journal of Neuroscience,* 29:9748-60.

Khurana RN, Sun X, Pearson E, Yang Z, Harmon J, Goldberg MF, Zhang K. A reappraisal of the clinical spectrum of North Carolina macular dystrophy. *Ophthalmology.* 2009 Oct;116(10):1976-83. Epub 2009 Jul 18.

Yang Z, Stratton C, Francis PJ, Kleinman ME, Tan PL, Gibbs D, Tong Z, Chen H, Constantine R, Yang X, Chen Y, Zeng J, Davey L, Ma X, Hau VS, Wang C, Harmon J, Buehler J, Pearson E, Patel S, Kaminoh Y, Watkins S, Luo L, Zabriskie NA, Bernstein PS, Cho W, Schwager A, Hinton DR, Klein ML, Hamon SC, Simmons E, Yu B, Campochiaro B, Sunness JS, Campochiaro P, Jorde L, Parmigiani G, Zack DJ, Katsanis N, Ambati J, Zhang K. (2008). Toll-like receptor 3 and geographic atrophy in age-related macular degeneration. *New England Journal of Med.* 14:1456-63. Epub 2008 Aug 27.

Yang Z, Chen Y, Lillo C, Chien J, Yu Z, Michaelides M, Klein M, Howes KA, Li Y, Kaminoh Y, Chen H, Zhao C, Chen Y, Al-Sheikh YT, Karan G, Corbeil D, Escher P, Kamaya S, Li C, Johnson S, Frederick JM, Zhao Y, Wang C, Cameron DJ, Huttner WB, Schorderet DF, Munier FL, Moore AT, Birch DG, Baehr W, Hunt DM, Williams DS, **Zhang K**. (2008). Mutant prominin 1 found in patients with macular degeneration disrupts photoreceptor disk morphogenesis in mice. *Journal Clinical Investigation* 118:2908-2916.

Tong Z, Yang Z, Patel S, Chen H, Gibbs D, Zeng J, Yang X, Ma X, Harmon J, Pearson E, Beuhler J, Luo L, Hau VS, Kaminoh Y, Zabriskie NA, Sun JK, Prakash M, Haman R, Tonna S, Constantine R, Ronquillo CC, Sadda SV, Avery RL, Brand JM, London N, King GL, Bernstein PS, Watkins S, Genetics of Diabetes and Diabetic Complication Study Group, Jorde LB, Li DY, Aiello LP, Pollak MR, **Zhang K**. (2008). Promoter polymorphism of the Erythropoietin gene in severe diabetic eye and kidney complications. *Proc Natl Acad Sci*. 105:6998-7003. Epub 2008 May 5.

Tonna SJ, Needham A, Polu K, Uscinski A, Appel GB, Falk RJ, Katz A, Al-Waheeb S, Kaplan BS, Jerums G, Savige J, Harmon J, Zhang K, Curhan GC, Pollak MR. (2008). NPHS2 variation in focal and segmental glomerulosclerosis. *BMC Nephrol*. 2008 Sep 29;9:13.

Tonna S, Dandapani SV, Uscinski A, Appel GB, Schlöndorff JS, Zhang K, Denker BM, Pollak MR. (2008). Functional genetic variation in aminopeptidase A (ENPEP): lack of clear association with focal and segmental glomerulosclerosis (FSGS). *Gene*. 410(1):44-52. Epub 2007 Dec 3.

Luo L, Harmon J, Yang X, Chen H, Patel S, Mineau G, Yang Z, Constantine R, Buehler J, Kaminoh Y, Ma X, Wong TY, Zhang M, Zhang K. (2008) Familial aggregation of age-related macular degeneration in the Utah population. *Vision Res*. 48(3):494-500.

Jiang L, Wheaton D, Bereta G, Zhang K, Palczewski K, Birch DG, Baehr W. (2008). A novel GCAP1(N104K) mutation in EF-hand 3 (EF3) linked to autosomal dominant cone dystrophy. *Vision Res*. 48:2425-32.

Chen Y, Jiang D, Yu L, Katz B, Zhang K, Wan B, Sun X. (2008). CYP1B1 and MYOC mutations in 116 Chinese patients with primary congenital glaucoma. *Arch Ophthalmol*. 126(10):1443-7.

Jones, CA, London, NR, Chen, H, Park, KW, Sauvaget, D, Rebecca A. Stockton, RA., Wythe, J. D., Suh, W, Larrieu-Lahargue, F, Mukouyama, Y, Lindblom, P, Seth, P, Frias, A, Nishiya, N, Ginsberg, M, Gerhardt, H, **Zhang, K***, and Li, D.Y.* (2008). Robo4 stabilizes the vascular network by inhibiting pathologic angiogenesis and endothelial hyperpermeability. *Nature Medicine* 14, 448-453. *co-corresponding authors.

Kleiman M E, Yamada K, Kakeda A, Chandrasekaran V, Nozaki M, Baffi J Z, Albuquerque R J C, Yamasake S, Itaya M, Pan Y, Appukuttan B, Gibbs D, Yang Z, Kariko K, Ambati B, Wilgus T A, DiPietro L A, Sakurai E, **Zhang K**, Smith T R, Taylor E W, Ambati J. (2008). Sequence- and target-independent suppression of angiogenesis by siRNA via TLR3. *Nature* 452, 591-597.

Yang X, Zabriskie NA, Hau VS, Chen H, Tong Z, Gibbs D, Farhi P, Katz BJ, Luo L, Pearson E, Goldsmith J, Ma X, Kaminoh Y, Chen Y, Yu B, Zeng J, **Zhang K***, and Yang, Z*. (2008). Genetic association of LOXL1 gene variants and exfoliation glaucoma in a Utah cohort. *Cell Cycle* 15 (4):521-4. *co-corresponding authors.

Yang Z, Tong Z, Chorich LJ, Pearson E, Moore A, Hunt DM, **Zhang K**. (2008) Clinical feature and fine mapping of North Carolina Macular Degeneration *Vision Research* 48(3):470-7.

Lu F, Hu J, Zhao, P, Lin Y, Yang Y, Liu X, Fan Y, Chen B, Liao S, Du, Q, Lei C, Cameron J, **Zhang K***, Yang Z*. (2007). HTRA1 variant increases risk to neovascular age-related macular degeneration in Chinese population. *Vision Research* 47:3120-3123. *co-corresponding authors.

Gibbs D, Yang Z, Constantine R, Cameron J, Chen H, DeWan A, Ma X, Yang X, Jorgenson A, Zeng J, Harmon J, Tong Z, Hoh J, **Zhang K**. (2008). Further mapping of 10q26 supports strong association of an HTRA1 promoter polymorphism as a causal variant in age-related macular degeneration. *Vision Research*, 48(5):685-9

Chen H, Yang Z, Gibbs D, Yang X, Hau V, Zhao P, Ma X, Zeng J, Luo L, Pearson E, Constantine R, Kaminoh Y, Harmon J, Tong Z, Stratton CA, Cameron JD, Tang S, **Zhang, K**. (2008). Association of HTRA1 polymorphism and bilaterality in advanced age related macular degeneration. *Vision Research*, 48(5):690-4.

Kaiser PK, Brown DM, **Zhang K**, Hudson HL, Holz FG, Shapiro H, Schneider S, Acharya NR (2007) Ranibizumab for Predominantly Classic Neovascular Age-related Macular Degeneration: Subgroup Analysis of First-year ANCHOR Results. *Am J Ophthalmol*. 144:850-857.

Chan CC, Shen D, Zhou M, Ross RJ, Ding X, **Zhang K**, Green WR, Tuo J. (2007) Human Htra1 in the archived eyes with age-related macular degeneration. *Trans Am Ophthalmol Soc*. 105:92-7

Yu J, Wiita P, Kawaguchi R, Honda J, Jorgensen A, **Zhang K**, Fischetti VA, Sun H. (2007) Biochemical analysis of a common human polymorphism associated with age-related macular degeneration. *Biochemistry*, 46(28), 8451-61.

Tuo J, Bojanowski CM, Zhou M, Shen D, Ross RJ, Rosenberg KI, Cameron DJ, Yin C, Kowalak JA, Zhuang Z, **Zhang K**, Chan CC. (2007) Murine ccl2/cx3cr1 deficiency results in retinal lesions mimicking human age-related macular degeneration. *Invest Ophthalmol Vis Sci*, 48(8), 3827-36.

Li W, Chen Y, Cameron DJ, Wang C, Karan G, Yang Z, Zhao Y, Pearson E, Chen H, Deng C, Howes K, **Zhang K**. (2007) Elovl4 haploinsufficiency does not induce early onset retinal degeneration in mice. *Vision Res*, 47(5), 714-22.

Fu L, Garland D, Yang Z, Shukla D, Rajendran A, Pearson E, Stone EM, **Zhang K**, Pierce EA. (2007) The R345W Mutation in EFEMP1 is Pathogenic, and Causes AMD-Like Deposits in Mice. *Hum Mol Genet*. 16, 3411-22.

Cameron DJ, Yang Z, Gibbs D, Chen H, Kaminoh Y, Jorgensen A, Zeng J, Luo L, Brinton E, Brinton G, Brand JM, Bernstein PS, Zabriskie NA, Tang S, Constantine R, Tong Z, **Zhang K**. (2007) HTRA1 Variant Confers Similar Risks to Geographic Atrophy and Neovascular Age-related Macular Degeneration. *Cell Cycle*, 6(9), 1122-5.

Cameron DJ, Tong Z, Yang Z, Kaminoh J, Kamiyah S, Chen H, Zeng J, Chen Y, Luo L, **Zhang K**. (2007) Essential role of Elovl4 in very long chain fatty acid synthesis, skin permeability barrier function, and neonatal survival. *Int J Biol Sci*, 3(2), 111-9.

Alvarez BV, Vithana EN, Yang Z, Koh AH, Yeung K, Yong V, Shandro HJ, Chen Y, Kolatkar P, Palasingam P, **Zhang K**, Aung T, Casey JR. (2007) Identification and characterization of a novel mutation in the carbonic anhydrase IV gene that causes retinitis pigmentosa. *Invest Ophthalmol Vis Sci*, 48(8), 3459-68.

- Yang Z, Camp NJ, Sun H, Tong Z, Gibbs D, Cameron DJ, Chen H, Zhao Y, Pearson E, Li X, Chien J, Dewan A, Harmon J, Bernstein PS, Shridhar V, Zabriskie NA, Hoh J, Howes K, **Zhang K.** (2006) A variant of the HTRA1 gene increases susceptibility to age-related macular degeneration. *Science*, 314(5801), 992-3. (see *Science New Focus*, same issue; *Science* *Research of the Year*, 2006)
- Rosenfeld PJ, Brown DM, Heier JS, Boyer DS, Kaiser PK, Chung CY, Kim RY for the MARINA Study Group. (2006) Ranibizumab for Neovascular Age-Related Macular Degeneration, *New Eng J of Med*, 355(14): 1419-1431.
- Brown DM, Kaiser PK, Michels M, Goubrane MD, Heier, JS, Kim RY, Sy JP, Schneider S for the ANCHOR Study Group. (2006) Ranibizumab versus Vertiporfin for Neovascular Age-Related Macular Degeneration. *New Eng J of Med*, 355(14): 1432-1444.
- Wang C, Katz BJ, Turner MJ, Hu J, **Zhang K.** (2006) Intravitreal triamcinolone acetate for the treatment of diffuse diabetic macular oedema--a case report. *Ann Acad Med Singapore*, 35(2), 112-4.
- Wang C, Hu J, Bernstein PS, Teske MP, Payne M, Yang Z, Li C, Adams D, Baird JH, **Zhang K.** (2006) Intravitreal injection of triamcinolone acetate for macular edema due to retinitis pigmentosa and other retinal diseases. *Adv Exp Med Biol*, 572309-14.
- Tong Z, Yang Z, Meyer JJ, McInnes AW, Xue L, Azimi AM, Baird J, Zhao Y, Pearson E, Wang C, Chen Y, **Zhang K.** (2006) A novel locus for X-linked retinitis pigmentosa. *Ann Acad Med Singapore*, 35(7), 476-8.
- Sauve Y, Karan G, Yang Z, Li C, Hu J, **Zhang K.** (2006) Treatment with carbonic anhydrase inhibitors depresses electroretinogram responsiveness in mice. *Adv Exp Med Biol*, 572439-46.
- Nozaki M, Raisler BJ, Sakurai E, Sarma JV, Barnum SR, Lambris JD, Chen Y, **Zhang K**, Ambati BK, Baffi JZ, Ambati J. (2006) Drusen complement components C3a and C5a promote choroidal neovascularization. *Proc Natl Acad Sci U S A*, 103(7), 2328-33.
- Magnusson KP*, Duan S, Sigurdsson H, Petursson H, Yang Z, Zhao Y, Bernstein PS, Ge J, Jonasson F, Stefansson E, Helgadóttir G, Zabriskie NA, Jonsson T, Björnsson A, Thorlacius T, Jonsson PV, Thorleifsson G, Kong A, Stefansson H, **Zhang K***, Stefansson K, Gulcher JR*. (2006) CFH Y402H confers similar risk of soft drusen and both forms of advanced AMD. *PLoS Med*, 3(1), 109-114. *Co-corresponding authors.
- Li Y, Wang G, Dong B, Sun X, Turner MJ, Kamaya S, **Zhang K.** (2006) A novel mutation of the VMD2 gene in a Chinese family with best vitelliform macular dystrophy. *Ann Acad Med Singapore*, 35(6), 408-10.
- Katz BJ, Zhao Y, Warner JE, Tong Z, Yang Z, **Zhang K.** (2006) A family with X-linked optic atrophy linked to the OPA2 locus Xp11.4-Xp11.2. *Am J Med Genet A*, 140(20), 2207-11.
- Katz BJ, Zhang K. (2006) Juxtapapillary hemangioblastoma as a manifestation of von Hippel-Lindau disease. *Pediatr Int*, 48(4), 423-5.
- Katz BJ, Yang Z, Payne M, Lin Y, Zhao Y, Pearson E, Duan S, Kamaya S, Karan G, **Zhang K.** (2006) Fundus appearance of choroideremia using optical coherence tomography. *Adv Exp Med Biol*, 57257-61.

Chen H, Chen Y, Horn R, Yang Z, Wang C, Turner MJ, **Zhang K.** (2006) Clinical features of autosomal dominant retinitis pigmentosa associated with a Rhodopsin mutation. *Ann Acad Med Singapore*, 35(6), 411-5.

Bidinost C, Matsumoto M, Chung D, Salem N, **Zhang K**, Stockton DW, Khoury A, Megarbane A, Bejjani BA, Traboulsi EI. (2006) Heterozygous and homozygous mutations in PITX3 in a large Lebanese family with posterior polar cataracts and neurodevelopmental abnormalities. *Invest Ophthalmol Vis Sci*, 47(4), 1274-80.

Yang Z, Alvarez BV, Chakarova C, Jiang L, Karan G, Frederick JM, Zhao Y, Sauve Y, Li X, Zrenner E, Wissinger B, Hollander AI, Katz B, Baehr W, Cremers FP, Casey JR, Bhattacharya SS, **Zhang K.** (2005) Mutant carbonic anhydrase 4 impairs pH regulation and causes retinal photoreceptor degeneration. *Hum Mol Genet*, 14(2), 255-65.

Pauer GJ, Xi Q, Zhang K, Traboulsi EI, Hagstrom SA. (2005) Mutation screen of the membrane-type frizzled-related protein (MFRP) gene in patients with inherited retinal degenerations. *Ophthalmic Genet*, 26(4), 157-61.

Li C, Kosmorsky G, **Zhang K**, Katz BJ, Ge J, Traboulsi EI. (2005) Optic atrophy and sensorineural hearing loss in a family caused by an R445H OPA1 mutation. *Am J Med Genet A*, 138(3), 208-11.

Karan G, Yang Z, Howes K, Zhao Y, Chen Y, Cameron DJ, Lin Y, Pearson E, **Zhang K.** (2005) Loss of ER retention and sequestration of the wild-type ELOVL4 by Stargardt disease dominant negative mutants. *Mol Vis*, 11657-64.

Karan G, Lillo C, Yang Z, Cameron DJ, Locke KG, Zhao Y, Thirumalaichary S, Li C, Birch DG, Vollmer-Snarr HR, Williams DS, **Zhang K.** (2005) Lipofuscin accumulation, abnormal electrophysiology, and photoreceptor degeneration in mutant ELOVL4 transgenic mice: a model for macular degeneration. *Proc Natl Acad Sci U S A*, 102(11), 4164-9.

Jiang L, Katz BJ, Yang Z, Zhao Y, Faulkner N, Hu J, Baird J, Baehr W, Creel DJ, **Zhang K.** (2005) Autosomal dominant cone dystrophy caused by a novel mutation in the GCAP1 gene (GUCA1A). *Mol Vis*, 11143-51.

Berthelemy-Okazaki N, Zhao Y, Yang Z, Camp NJ, Farnham J, Parker D, Tsuruda J, Macdonald J, **Zhang K**, Cannon-Albright LA. (2005) Examination of ELN as a candidate gene in the Utah intracranial aneurysm pedigrees. *Stroke*, 36(6), 1283-4.

Yang Z, Li Y, Jiang L, Karan G, Moshfeghi D, O'Connor S, Li X, Yu Z, Lewis H, Zack D, Jacobson S, **Zhang K.** (2004) A novel RDS/peripherin gene mutation associated with diverse macular phenotypes. *Ophthalmic Genet*, 25(2), 133-45.

Xu Q, Wang Y, Dabdoub A, Smallwood PM, Williams J, Woods C, Kelley MW, Jiang L, Tasman W, **Zhang K**, Nathans J. (2004) Vascular development in the retina and inner ear: control by Norrin and Frizzled-4, a high-affinity ligand-receptor pair. *Cell*, 116(6), 883-95.

Toomes C, Bottomley HM, Scott S, Mackey DA, Craig JE, Appukuttan B, Stout JT, Flaxel CJ, **Zhang K**, Black GC, Fryer A, Downey LM, Inglehearn CF. (2004) Spectrum and frequency of FZD4 mutations in familial exudative vitreoretinopathy. *Invest Ophthalmol Vis Sci*, 45(7), 2083-90.

Toomes C, Bottomley HM, Jackson RM, Towns KV, Scott S, Mackey DA, Craig JE, Jiang L, Yang Z, Trembath R, Woodruff G, Gregory-Evans CY, Gregory-Evans K, Parker MJ, Black GC, Downey LM, **Zhang K**, Inglehearn CF. (2004) Mutations in LRP5 or FZD4 underlie the common familial exudative vitreoretinopathy locus on chromosome 11q. *Am J Hum Genet*, 74(4), 721-30.

Payne M, Yang Z, Katz BJ, Warner JE, Weight CJ, Zhao Y, Pearson ED, Treft RL, Hillman T, Kennedy RJ, Meire FM, **Zhang K**. (2004) Dominant optic atrophy, sensorineural hearing loss, ptosis, and ophthalmoplegia: a syndrome caused by a missense mutation in OPA1. *Am J Ophthalmol*, 138(5), 749-55.

Maugeri A, Meire F, Hoyng CB, Vink C, Van Regemorter N, Karan G, Yang Z, Cremers FP, **Zhang K**. (2004) A novel mutation in the ELOVL4 gene causes autosomal dominant Stargardt-like macular dystrophy. *Invest Ophthalmol Vis Sci*, 45(12), 4263-7.

Karan G, Yang Z, **Zhang K**. (2004) Expression of wild type and mutant ELOVL4 in cell culture: subcellular localization and cell viability. *Mol Vis*, 10:248-53.

Hunt DM, Wilkie SE, Newbold R, Deery E, Warren MJ, Bhattacharya SS, **Zhang K**. (2004) Dominant cone and cone-rod dystrophies: functional analysis of mutations in retGC1 and GCAP1. *Novartis Found Symp*, 255:37-49; discussion 49-50, 177-8.

Berry V, Yang Z, Addison PK, Francis PJ, Ionides A, Karan G, Jiang L, Lin W, Hu J, Yang R, Moore A, **Zhang K**, Bhattacharya SS. (2004) Recurrent 17 bp duplication in PITX3 is primarily associated with posterior polar cataract (CPP4). *J Med Genet*, 41(8), e109.

Zhang XM, Yang Z, Karan G, Hashimoto T, Baehr W, Yang XJ, **Zhang K**. (2003) Elov14 mRNA distribution in the developing mouse retina and phylogenetic conservation of Elov14 genes. *Mol Vis*, 9:301-7.

Yang Z, Lin W, Moshfeghi DM, Thirumalaichary S, Li X, Jiang L, Zhang H, Zhang S, Kaiser PK, Traboulsi EI, **Zhang K**. (2003) A novel mutation in the RDS/Peripherin gene causes adult-onset foveomacular dystrophy. *Am J Ophthalmol*, 135(2), 213-8.

Kniazeva M, Sieber M, McCauley S, **Zhang K**, Watts JL, Han M. (2003) Suppression of the ELO-2 FA elongation activity results in alterations of the fatty acid composition and multiple physiological defects, including abnormal ultradian rhythms, in *Caenorhabditis elegans*. *Genetics*, 163(1), 159-69.

Johnson S, Halford S, Morris AG, Patel RJ, Wilkie SE, Hardcastle AJ, Moore AT, **Zhang K**, Hunt DM. (2003) Genomic organisation and alternative splicing of human RIM1, a gene implicated in autosomal dominant cone-rod dystrophy (CORD7). *Genomics*, 81(3), 304-14.

Zhang K, Garibaldi DC, Li Y, Green WR, Zack DJ. (2002) Butterfly-shaped pattern dystrophy: a genetic, clinical, and histopathological report. *Arch Ophthalmol*, 120(4), 485-90.

Yang Z, Peachey NS, Moshfeghi DM, Thirumalaichary S, Chorich L, Shugart YY, Fan K, **Zhang K**. (2002) Mutations in the RPGR gene cause X-linked cone dystrophy. *Hum Mol Genet*, 11(5), 605-11.

Zhang K, Kniazeva M, Han M, Li W, Yu Z, Yang Z, Li Y, Metzker ML, Allikmets R, Zack DJ, Kakuk LE, Lagali PS, Wong PW, MacDonald IM, Sieving PA, Figueroa DJ, Austin CP, Gould RJ, Ayyagari R, Petrukhin K. (2001) A 5-bp deletion in ELOVL4 is associated with two related forms of autosomal dominant macular dystrophy. *Nature Genet*, 27(1), 89-93.

Wilkie SE, Li Y, Deery EC, Newbold RJ, Garibaldi D, Bateman JB, Zhang H, Lin W, Zack DJ, Bhattacharya SS, Warren MJ, Hunt DM, **Zhang K**. (2001) Identification and functional consequences of a new mutation (E155G) in the gene for GCAP1 that causes autosomal dominant cone dystrophy. *Am J Hum Genet*, 69(3), 471-80.

Li Y, Marcos I, Borrego S, Yu Z, **Zhang K**, Antinolo G. (2001) Evaluation of the ELOVL4 gene in families with retinitis pigmentosa linked to the RP25 locus. *J Med Genet*, 38(7), 478-80.

Bernstein PS, Tammur J, Singh N, Hutchinson A, Dixon M, Pappas CM, Zabriskie NA, **Zhang K**, Petrukhin K, Leppert M, Allikmets R. (2001) Diverse macular dystrophy phenotype caused by a novel complex mutation in the ELOVL4 gene. *Invest Ophthalmol Vis Sci*, 42(13), 3331-6.

Ayyagari R, **Zhang K**, Hutchinson A, Yu Z, Swaroop A, Kakuk LE, Seddon JM, Bernstein PS, Lewis RA, Tammur J, Yang Z, Li Y, Zhang H, Yashar BM, Liu J, Petrukhin K, Sieving PA, Allikmets R. (2001) Evaluation of the ELOVL4 gene in patients with age-related macular degeneration. *Ophthalmic Genet*, 22(4), 233-9.

Stefko ST, **Zhang K**, Gorin MB, Traboulsi EI. (2000) Clinical spectrum of chromosome 6-linked autosomal dominant drusen and macular degeneration. *Am J Ophthalmol*, 130(2), 203-8.

Kniazeva M, Traboulsi EI, Yu Z, Stefko ST, Gorin MB, Shugart YY, O'Connell JR, Blaschak CJ, Cutting G, Han M, **Zhang K**. (2000) A new locus for dominant drusen and macular degeneration maps to chromosome 6q14. *Am J Ophthalmol*, 130(2), 197-202.

Allikmets R. (2000) Further evidence for an association of ABCR alleles with age-related macular degeneration. The International ABCR Screening Consortium. *Am J Hum Genet*, 67(2), 487-91.

Zhang K, Kniazeva M, Hutchinson A, Han M, Dean M, Allikmets R. (1999) The ABCR gene in recessive and dominant Stargardt diseases: a genetic pathway in macular degeneration. *Genomics*, 60(2), 234-7.

Zhang K, Jampel HD. (1999) Discordance of primary infantile glaucoma in monozygotic twins. *Am J Ophthalmol*, 128(1), 97-8.

Zhang K, Garibaldi DC, Kniazeva M, Albin T, Chiang MF, Kerrigan M, Sunness JS, Han M, Allikmets R. (1999) A novel mutation in the ABCR gene in four patients with autosomal recessive Stargardt disease. *Am J Ophthalmol*, 128(6), 720-4.

Kniazeva MF, Chiang MF, Cutting GR, Zack DJ, Han M, **Zhang K**. (1999) Clinical and genetic studies of an autosomal dominant cone-rod dystrophy with features of Stargardt disease. *Ophthalmic Genet*, 20(2), 71-81.

Kniazeva M, Chiang MF, Morgan B, Anduze AL, Zack DJ, Han M, **Zhang K**. (1999) A new locus for autosomal dominant stargardt-like disease maps to chromosome 4. *Am J Hum Genet*, 64(5), 1394-9.

Hata Y, Duh E, **Zhang K**, Robinson GS, Aiello LP. (1998) Transcription factors Sp1 and Sp3 alter vascular endothelial growth factor receptor expression through a novel recognition sequence. *J Biol Chem*, 273(30), 19294-303.

Zhang K, Bither PP, Park R, Donoso LA, Seidman JG, Seidman CE. (1994) A dominant Stargardt's macular dystrophy locus maps to chromosome 13q34. *Arch Ophthalmol*, 112(6), 759-64.

Rutledge BJ*, Zhang K*, Bier E, Jan YN, Perrimon N. (1992) The *Drosophila* *spitz* gene encodes a putative EGF-like growth factor involved in dorsal-ventral axis formation and neurogenesis. *Genes Dev*, 6(8), 1503-17. co-first authors.

Zhang K, Smouse D, Perrimon N. (1991) The crooked neck gene of *Drosophila* contains a motif found in a family of yeast cell cycle genes. *Genes Dev*, 5(6), 1080-91.

Zhang K, Chaillet JR, Perkins LA, Halazonetis TD, Perrimon N. (1990) *Drosophila* homolog of the mammalian *jun* oncogene is expressed during embryonic development and activates transcription in mammalian cells. *Proc Natl Acad Sci U S A*, 87(16), 6281-5.

Perkins LA, Doctor JS, **Zhang K**, Stinson L, Perrimon N, Craig EA. (1990) Molecular and developmental characterization of the heat shock cognate 4 gene of *Drosophila melanogaster*. *Mol Cell Biol*, 10(6), 3232-8.

Kapler GM, **Zhang K**, Beverley SM. (1990) Nuclease mapping and DNA sequence analysis of transcripts from the dihydrofolate reductase-thymidylate synthase (R) region of *Leishmania major*. *Nucleic Acids Res*, 18(21), 6399-408.

Kapler GM, **Zhang K**, Beverley SM. (1987) Sequence and S1 nuclease mapping of the 5' region of the dihydrofolate reductase-thymidylate synthase gene of *Leishmania major*. *Nucleic Acids Res*, 15(8), 3369-83.

Books & Book Chapters:

Cameron DJ, Yang Z, Tong Z, Zhao Y, Praggastis A, Brinton E, Harmon J, Chen Y, Pearson E, Bernstein PS, Brinton G, Li X, Jorgensen A, Schneider S, Gibbs D, Chen H, Wang C, Howes K, Camp NJ, Zhang K. (2008) 10q26 is associated with increased risk of age-related macular degeneration in the Utah population. In Anderson RE, LaVail MM, Hollyfield JG (Eds.), *Recent Advances In Retinal Degeneration*. New York: Plenum Publishing Corporation.

Chen H, Yang Z, Zhao P, Zhang K (2008). Genetics of AMD. in Kaiser PK, Sears J, Singh R (Eds), *Macular Degeneration – State of the Art*. London, Remedica Medical Education and Publishing Limited

Moshfeghi DM, Yang Z, Faulkner ND, Karan G, Hirumalaichary S, Jiang L, Tsai T, Zhang K. (2005). Choroidal Neovascularization in Patients with Adult-onset Foveomacular Dystrophy Caused by Mutations in the RDS/Peripherin Gene: Report of 3 Cases. In Hollyfield JG, Anderson RE, LaVail MM (Eds.), *Retinal Degeneration: Clinical and Laboratory Applications*. New York: Plenum Publishing Corporation.

Katz BJ, Yang Z, Payne M, Zhang K. (2005). Fundus Appearance of Choroideremia. In Hollyfield JG, Anderson RE, LaVail MM (Eds.), *Retinal Degeneration: Clinical and Laboratory Applications*. New York: Plenum Publishing Corporation.

Zhang K, Carr RE, Sunness JS. (2005). Hereditary choroidal dystrophies. In Ryan SJ (Ed.), *Retina*.

Hu J, Payne M, Zhang K. (2005). Intravitreal Injection of Triamcinolone Acetonide for Macular Edema Due to Retinitis Pigmentosa and Other Retinal Diseases. In Hollyfield JG, Anderson RE, LaVail MM (Eds.), *Retinal Degeneration: Clinical and Laboratory Applications*. New York: Plenum Publishing Corporation.

Sauve Y, Karan G, Yang Z, Li C, Hu J, Zhang K. (2005). Treatment with Carbonic Anhydrase Inhibitors Depresses Electroretinogram Responsiveness in Mice. In Hollyfield JG, Anderson RE, LaVail MM (Eds.), *Retinal Degeneration: Clinical and Laboratory Applications*.

Zhang K, Carr RE, Sunness JS. (2001). Hereditary choroidal dystrophies. In Ryan SJ (Ed.), Retina.

Li Y, Lam L, Yu Z, Yang Z, Bither P, Zhang K. (2000). Clinical spectrum of autosomal dominant Stargardt's macular dystrophy with a mutation in ELOVL4. In Hollyfield JG, Anderson RE, LaVail MM (Eds.), Retinal Degeneration: Clinical and Laboratory Applications. New York: Plenum

Garibaldi GC, Yang Z, Li Y, Yu Z, Zhang K. (2000). The role of fatty acids in the pathogenesis of retinal degeneration. In Hollyfield JG, Anderson RE, LaVail MM (Eds.), Retinal Degeneration: Clinical and Laboratory Applications. New York: Plenum Publishing Corporation.

Zhang, K. (Ed.) Ophthalmic Disease Mechanisms and Drug Discovery. (2016). New Jersey: World Scientific.

Liu, Y and Zhang, K. Ed. Ocular Stem Cell Therapy. World Scientific Publishing, in preparation.

Conference Papers:

Gonzales, JA; Tawansy, KA; Simon, A; Zhang K,(2005). Chorioretinal coloboma in juvenile X-linked retinoschisis. KINVESTIGATIVE OPHTHALMOLOGY & VISUAL SCIENCE, 46

Jiang, L; Bottomley, HM; Yang, Z; Zhao, Y; Lee, H; Tawansy, K; Tasman, W; Nathans, J; Toomes, C; Zhang K, (2004).Novel mutations in FZD4 with familial exudative vitreoretinopathy. KINVESTIGATIVE OPHTHALMOLOGY & VISUAL SCIENCE. 45:U944-U944

Bottomley, HM; Downey, LM; Mackey, DA; Craig, JE; Gregory-Evans, CY; Woodruff, G; Black, GCM; Zhang, K; Inglehearn, CF; Toomes, (2004), Mutation screening in FZD4 and LRP5 in familial exudative vitreoretinopathy patients. CINVESTIGATIVE OPHTHALMOLOGY & Zhao, Y; Jiang, L; Yang, Z; Li, X; Zhang, K (2004). Identification of mutations in GCAP1 in autosomal dominant cone dystrophy. INVESTIGATIVE OPHTHALMOLOGY & VISUAL SCIENCE. 45:U942-U942

Locke, KG; Karan, G; Yang, Z; Zhang, K; Birch, (2004). Human mutant ELOVL4 causes retinal dystrophy in transgenic mice. DGINVESTIGATIVE OPHTHALMOLOGY & VISUAL SCIENCE. 45:U715-U715

Karan, G; Maugeri, A; Yang, Z; Jiang, L; Li, X; Cremers, F; Zhang, K (2004). Functional consequences of a novel mutation in ELOVL4 causing macular dystrophy. INVESTIGATIVE OPHTHALMOLOGY & VISUAL SCIENCE. 45:U714-U714

Cameron, DJ; Karon, G; Yang, Z; Li, X; Zhang, K; Vollmer-Snarr, (2004). A2E accumulation associated with Elov14 and macular degeneration. HRINVESTIGATIVE OPHTHALMOLOGY & VISUAL SCIENCE. 45:U714-U714

Yang, Z; Payne, M; Weight, C; Pearson, ED; Zhao, Y; Li, X; Faulkner, N; Katz, B; Warner, J; Zhang, K (2004). Dominant optic atrophy, sensorineural hearing loss, ptosis, and ophthalmoplegia: A syndrome caused by a mutation in OPAL1. INVESTIGATIVE OPHTHALMOLOGY & Paillart, C; Zhang, K; Rebrik, T; Baehr, W; Korenbrot, (2004). Differential expression of CNG channels in fish photoreceptors. JINVESTIGATIVE OPHTHALMOLOGY & VISUAL SCIENCE. 45:U541-U541

Toomes, C; Bottomley, HM; Jackson, RM; Towns, KV; Mackey, DA; Craig, JE; Gregory-Evans, K; Downey, LM; Zhang, K; Inglehearn, (2004). Mutations in LRP5 or FZD4 underlie the common FEVR locus on chromosome 11q13. CFINVESTIGATIVE OPHTHALMOLOGY & VISUAL SCIENCE. 45:U382-U382

Zhang, K; Zhang, C; Hofeldt, KJ; Tso, MOM; Lei, (2004). Minocycline preserves rod and cone function in light-induced retinal degeneration. BINVESTIGATIVE OPHTHALMOLOGY & VISUAL SCIENCE. 45:U347-U347

Hu, J; Blom, E; Bernstein, P; Teske, M; Blanchard, D; Yang, Z; Zhang, K(2003). Intravitreal injection of triamcinolone acetonide for macular edema due to various retinal vascular diseases. INVESTIGATIVE OPHTHALMOLOGY & VISUAL SCIENCE. 44:U426-U426

Zhang, H; Liu, XH; Zhang, K; Chen, CK; Frederick, JM; Prestwich, GD; Baehr, (2003). The putative cGMP phosphodiesterase delta subunit ("PDE delta") functions as a prenyl binding protein (PBP). INVESTIGATIVE OPHTHALMOLOGY & VISUAL SCIENCE. 44:U357-U357

Jin, Y; Jiang, L; Yang, Z; Li, X; Hoffman, R; Hu, J; Zhang, K(2003). Clinical and genetic studies of an autosomal dominant form of congenital cataract. INVESTIGATIVE OPHTHALMOLOGY & VISUAL SCIENCE. 44:U302-U302

Young, ML; Fan, K; Yang, Z; Zhang, K(2002). Age-related macular degeneration: Clinical characteristics and genetic linkage study in a dominant family. INVESTIGATIVE OPHTHALMOLOGY & VISUAL SCIENCE. 43:U659-U659

Fan, KK; Kenney, GE; Kitsos, G; Yang, Z; Economou-Petersen, E; Grigoriadou, M; Zack, DJ; Psilas, K; Petersen, MB; Zhang, K(2002). Autosomal dominant Stargardt macular dystrophy: Clinical features and linkage analysis in a large Greek family. INVESTIGATIVE OPHTHALMOLOGY & VISUAL SCIENCE. 43:U658-U658