

## Moin Mohamed's publications

Anti Vascular Endothelial Growth Factor therapies in Ophthalmology: Current Use, Controversies and the Future.

*Kwong TQ, Mohamed M.*

**Br J Clin Pharmacol. 2014 Mar 6**

Recessive mutations in SLC38A8 cause foveal hypoplasia and optic nerve misrouting without albinism.

*Poulter JA, Al-Araimi M, Conte I, van Genderen MM, Sheridan E, Carr IM, Parry DA, Shires M, Carrella S, Bradbury J, Khan K, Lakeman P, Sergouniotis PI, Webster AR, Moore AT, Pal B, Mohamed MD, Venkataramana A, Ramprasad V, Shetty R, Saktivel M, Kumaramanickavel G, Tan A, Mackey DA, Hewitt AW, Banfi S, Ali M, Inglehearn CF, Toomes C.*

**Am J Hum Genet. 2013 Dec 5;93(6):1143-50.**

Macular spectral domain optical coherence tomography findings in Tanzanian endemic optic neuropathy.

*Kisimbi J, Shalchi Z, Mahroo OA, Mhina C, Sanyiwa AJ, Mabey D, Mohamed M, Plant GT.*

**Brain. 2013 Nov;136(Pt 11):3418-26.**

A new recessively inherited disorder composed of foveal hypoplasia, optic nerve decussation defects and anterior segment dysgenesis maps to chromosome 16q23.3-24.1.

*Al-Araimi M, Pal B, Poulter JA, van Genderen MM, Carr I, Cudrnak T, Brown L, Sheridan E, Mohamed MD, Bradbury J, Ali M, Inglehearn CF, Toomes C.*

**Mol Vis. 2013 Nov 1;19:2165-72.**

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*Pelosini L, Hamilton R, Mohamed M, Hamilton AM, Marshall J.*

**Retina. 2013 Mar;33(3):548-58.**

Re: Single session of Pascal versus multiple sessions of conventional laser for panretinal photocoagulation in proliferative diabetic retinopathy: a comparative study.

*Jojo V, Mohamed M.*

**Retina. 2012 Sep;32(8):1697-8**

Mutations in NMNAT1 cause Leber congenital amaurosis and identify a new disease pathway for retinal degeneration

*Robert K Koeneke, Hui Wang, Jacek Majewski, Xia Wang, Irma Lopez, Huanan Ren, Yiyun Chen, Yumei Li, Gerald A. Fishman, Mohammed Genead, Jeremy Schwartzentruber, Naimesh Solanki, Elias I. Traboulsi, Jingliang Cheng, Clare V Logan, Martin McKibbin, Bruce E Hayward, David A Parry, Colin A Johnson, Mohammed Nageeb, FORGE Canada Consortium, James A Poulter, Moin D Mohamed, Hussain Jafri, Yasmin Rashid, Graham R Taylor, Vafa Keser, Graeme Mardon, Huidan Xu, Chris F Inglehearn, Qing Fu, Carmel Toomes and Rui Chen*

*Nat Genet. 2012 Sep;44(9):1035-9. doi: 10.1038/ng.2356. Epub 2012 Jul 2*

Central visual disturbance associated with transient disruption of photoreceptor inner-outer segment junction

*Mahroo OAR, Gavin EA, Mann S, Mohamed M*

*Can J Ophthalmol 2012 Oct;47(5):e19-20.*

Ethnic variations in the prevalence of diabetic retinopathy in people with diabetes attending screening in the United Kingdom (DRIVE UK).

*Sivaprasad S, Gupta B, Gulliford MC, Dodhia H, Mohamed M, Nagi D, Evans JR.*

*PLoS One. 2012;7(3):e32182.*

A case of African crystalline maculopathy.

*Dhital A, Mohamed M.*

*Eye (Lond). 2012 Apr;26(4):615-6*

Next generation sequencing identifies mutations in Atonal homolog 7 (ATOH7) in families with global eye developmental defects.

*Khan K, Logan CV, McKibbin M, Sheridan E, Elçioglu NH, Yenice O, Parry DA, Fernandez-Fuentes N, Abdelhamed ZI, Al-Maskari A, Poulter JA, Mohamed MD, Carr IM, Morgan JE, Jafri H, Raashid Y, Taylor GR, Johnson CA, Inglehearn CF, Toomes C, Ali M.*  
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*Ali M, Ramprasad VL, Soumitra N, Mohamed MD, Jafri H, Rashid Y, Danciger M, McKibbin M, Kumaramanickavel G, Inglehearn CF.*

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*M D Mohamed, M Gupta, M A Parsons, I G Rennie*

BJO. 2005, Jan; 89(1):14-16

A new phenotype of recessively inherited foveal hypoplasia and anterior segment dysgenesis maps to a locus on chromosome 16q23.2-24.2

*B Pal\*, M D Mohamed\*, T J Keen, G A Williams, J A Bradbury, E Sheridan, C F Inglehearn (\* joint first authors)*

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*S M Elsherbiny, M D Mohamed*

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