


Correction

Correction: Littink, K. W.; et al. Autosomal Recessive *NRL* Mutations in Patients with Enhanced S-Cone Syndrome. *Genes* 2018, 9, 68

Karin W. Littink ¹, Patricia T.Y. Stappers ¹, Frans C.C. Riemsdag ^{1,2}, Herman E. Talsma ^{1,2}, Maria M. van Genderen ², Frans P.M. Cremers ^{3,4}, Rob W.J. Collin ^{3,4}  and L. Ingeborgh van den Born ^{1,*}

¹ The Rotterdam Eye Hospital, 3011 BH Rotterdam, The Netherlands; k.littink@oogziekenhuis.nl (K.W.L.); p.stappers@oogziekenhuis.nl (P.T.Y.S.); FRiemsdag@gmail.com (F.C.C.R.); HTalsma@bartimeus.nl (H.E.T.)

² Bartiméus Center for Complex Visual Disorders, 3703 AJ Zeist, The Netherlands; mvgenderen@bartimeus.nl

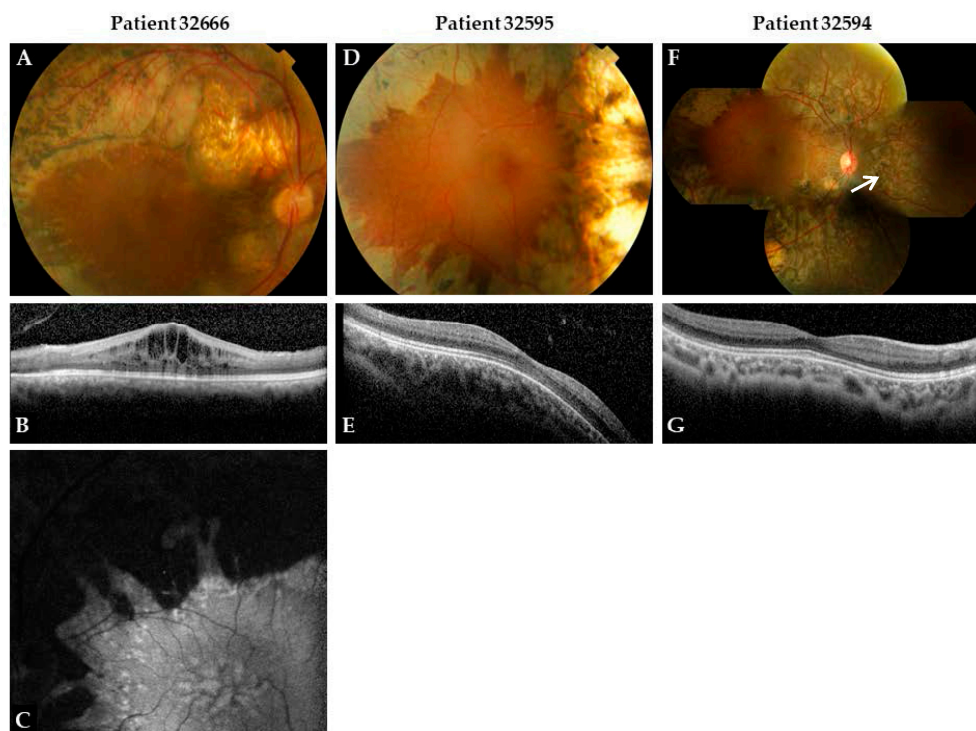
³ Department of Human Genetics, Radboud University Medical Center, 6525 GA Nijmegen, The Netherlands; Frans.Cremers@radboudumc.nl (F.P.M.C.); Rob.Collin@radboudumc.nl (R.W.J.C.)

⁴ Donders Institute for Brain, Cognition and Behaviour, Radboud University Medical Center, 6525 GA Nijmegen, The Netherlands

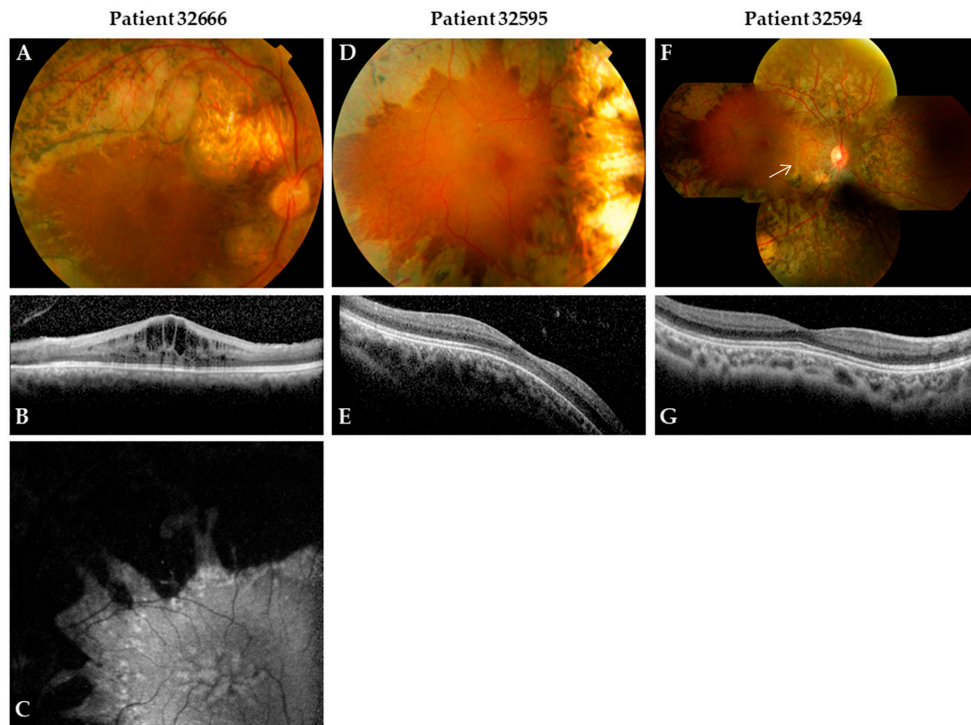
* Correspondence: born@eyehospital.nl; Tel.: +31-10-4017777

Received: 2 March 2018; Accepted: 5 March 2018; Published: 7 March 2018

The authors wish to make the following correction to this paper [1]. Due to mislabeling, replace:



with



The authors would like to apologize for any inconvenience caused to the readers by these changes.

Reference

1. Littink, K.W.; Stappers, P.T.Y.; Riemsdag, F.C.C.; Talsma, H.E.; van Genderen, M.M.; Cremers, F.P.M.; Collin, R.W.J.; van den Born, L.I. Autosomal Recessive *NRL* Mutations in Patients with Enhanced S-Cone Syndrome. *Genes* **2018**, *9*, 68. [[CrossRef](#)] [[PubMed](#)]



© 2018 by the authors. Licensee MDPI, Basel, Switzerland. This article is an open access article distributed under the terms and conditions of the Creative Commons Attribution (CC BY) license (<http://creativecommons.org/licenses/by/4.0/>).