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Prevalence and risk-factors for age-related macular degeneration in a population-based cohort study of older adults in Northern Ireland using multi-modal imaging- NICOLA Study.

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Running head: AMD epidemiology in NICOLA

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Critically revised the manuscript: DMW, NBQ, KAM, BH, LS, AJMcK, JW, MT, SC, BMcG, ISY, FK, TP and UC.

Statistical analyses were performed by DMW.

All authors read and approved the final manuscript. REH and DMW had full access to all study data and takes responsibility for the integrity of the data and the accuracy of the data analysis. UC takes the responsibility of overall content as the guarantor.

Abstract

Purpose: To report prevalence and risk factor associations for AMD and AMD features from multi-modal retinal grading in a multidisciplinary longitudinal population-based study of aging in Northern Ireland.

Study-Design: Population based longitudinal cohort study.

Methods: Retinal imaging at the NICOLA study health assessment included stereo CFP (Canon CX-1, Tokyo, Japan) and SD-OCT ((HRA+OCT; Heidelberg Engineering, Heidelberg, Germany). Medical history and demographic information was obtained during a home interview. Descriptive statistics were used to describe the prevalence of AMD and individual AMD features. Multiple imputation followed by multiple regression modelling was used to explore risk factor associations including relationships with AMD genetic risk score.

Results: Retinal images from 3386 participants were available for analysis. Mean age of the sample was 63.4 (sd. 9.01 range. 36-99). Population weighted prevalence of AMD using colour grading in those over 55 was: No drusen:60.4%, drusen<63µm:15.9%, drusen 63-125µm:13.7%, drusen>125µm or pigmentary changes: 8.3%, late AMD:1.6%. Prevalence of AMD features in those over 55 years was: OCT drusen 27.5%, complete outer retinal pigment epithelium and outer retinal atrophy (cRORA) on was OCT 4.3%, reticular drusen 3.2% and subretinal drusenoid deposits 25.7%. The genetic risk score was significantly associated with drusen and cRORA but less so for SDD alone and non-significant for hyperpigmentation or vitelliform lesions.

Conclusions: Multi modal imaging-based classification has provided evidence of some divergence of genetic risk associations between classical drusen and SDD. Our findings support an urgent review of current AMD severity classification systems.

o What is already known on this topic

Age-related macular degeneration is of significant public health concern due to the impact of vision loss on quality of life together with an increasingly aged population. Substantial advances in retinal imaging has furthered our understanding of the condition through the use of optical coherence tomography (OCT) images yet most epidemiological studies rely solely on colour fundus photographs for assessment.

o What this study adds

The Northern Ireland Cohort for the Longitudinal Aging Study (NICOLA) included multi-modal retinal imaging (colour fundus photography, OCT and ultra-widefield retinal imaging enabling an unprecedented assessment of AMD including individual retinal features of the condition. The results show disparities between assessments using different imaging modalities highlighting the importance of using multi modal imaging for future studies of prevalence and incidence.

o How this study might affect research, practice or policy

This study supports the incorporation of OCT based features into future grading schemes or severity staging systems and the need for longitudinal data on progression of OCT based features to better understand their significance in progression to late-stage disease.

Precise

Prevalence and Risk factors for age-related macular degeneration using multi modal retinal imaging provides additional information beyond colour photography alone particularly for early AMD features. New AMD severity classification systems are urgently needed.

Introduction

Epidemiological studies of age-related macular degeneration (AMD) have traditionally relied on colour fundus photography to identify the characteristics used to classify each eye into disease severity stages. Landmark studies such as the Beaver Dam Eye study¹, the Rotterdam Eye Study² and the Blue Mountain's Eye Study³ have identified the key phenotypic features and revealed the natural history of progression and associations with risk factors especially age, smoking, and genetic risk⁴⁻⁷. Although studies showed the importance of drusen size and hyperpigmentation in influencing the risk of progression, the role of other color defined features such as reticular pseudo drusen⁸ (RPD; a characteristic form of extracellular drusenoid deposits) and small drusen remains controversial. This is mainly because color grading can over-estimate presence of small drusen⁹ and underestimate presence of reticular pseudo-drusen (RPD)¹⁰. Optical Coherence tomography (OCT permits the distinction of nodular (classic) drusen from RPD through characterization of layer location.

The establishment of a new population based longitudinal study of ageing in Northern Ireland (The Northern Ireland Cohort for the Longitudinal Study of Aging-NICOLA) offered an opportunity to assess the retina using multi-modal retinal imaging (colour fundus photography (CFP), OCT, ultra-widefield imaging (UWFI)) approach, define previously unexplored phenotypes from an epidemiological perspective and study risk factor associations with an extended range of markers of systemic health

Methods

NICOLA study overview

The Northern Ireland Cohort Study for the Longitudinal Study of Ageing (NICOLA) is a multidisciplinary prospective population-based cohort study. Wave 1 of the study commenced in December 2013 and ended in April 2018. In total, 8452 persons completed the home-based computer assisted personal interview (CAPI). Of these, 3420 (40.5%) attended for the health assessment, which consisted of anthropometric, cardiac, respiratory, cognitive, and ophthalmic tests (Figure 1).

Ethical approval was obtained from the School of Medicine, Dentistry and Biomedical Sciences Ethics Committee, Queen's University Belfast (Ref: 12/23). The study adhered to the Declaration of Helsinki and all participants gave informed written consent. Details on image acquisition, image grading, physical data collection, self-report, blood-based biomarkers, genetic analysis, outcome categorisation and covariate selection can be found in the supplementary methods online.

Statistical analysis

All statistical analysis was conducted using R (version 3.6.0).¹¹

Prevalence analysis

Prevalence of AMD was estimated, weighting records by the population distribution of age and sex as recorded in the 2011 Census for Northern Ireland. For the eye level analysis, the frequency, prevalence and 95% confidence intervals (CI) for each AMD

feature was calculated. To facilitate comparison with prior studies¹² estimates are provided by age category and with a separate estimate for participants aged 55 and above.

Imputation and Risk factor association

Missing risk factor values were imputed using Multivariate Imputation by Chained Equations (MICE) using the *R* package *mice* (version 3.8.0)¹³, which yielded a set of five datasets in which the missing values had been imputed probabilistically. The regression models outlined below were fitted using each of the imputed datasets and estimates were pooled across imputations. A single estimate was generated for each parameter of interest that encompassed the additional uncertainty introduced by imputation. Further details of the imputation procedure are given in supplementary methods.

Associations between AMD features and risk factors were explored using logistic regression. A series of regressions were fitted for each AMD feature with the feature as the response variable. First, a set of univariate models was fitted, the sole predictor in each being one of the candidate risk factors. A second set was adjusted with the addition of age (linear and quadratic terms) and sex. Risk factors receiving some statistical support in the age-sex adjusted models ($P < 0.1$) were included in a multivariable model. Finally, a saturated model including all risk factors was fitted.

Where an AMD feature was particularly rare, inclusion of all the selected variables occasionally resulted in unstable estimates or a failure of multivariable or saturated models to converge. In these instances, problematic combinations of predictors were identified, and models refitted after removing these risk factors. For the analysis of person level AMD risk factors (dichotomised AMD stage) an additional model was fitted excluding the GRS to explore the extent to which associations with other risk factors differed in the absence of the GRS.

Results

Cohort Characteristics

Of the 3420 participants who attended the health assessment, retinal imaging was performed in 3386 (99%) (Figure 1). Images were available for color 6259 eyes (92%), OCT 6601 (96%) and UWFI 6138(89%). Images were ungradable in approximately 121 (4%) (color), 42(1%) (OCT) and 172 (5%) (UWFI) eyes. The number of individuals for whom both eyes were gradable was: color 2994 (88%), OCT 3257(95%) and UWFI 2934 (85%). More than 85% of participants had gradable images from both eye on all 3 imaging modalities. Supplementary Table 1 shows proportions of participants with ungradable images by imaging modality. The mean (SD) age of participants with gradable color images was 63.5(8.9) years (supplementary table 1) and over 99.9% were Caucasian.

Prevalence by Beckman stage

The prevalence of stage 0 and stage 1 (representing no features of AMD or small hard drusen only) declined from 62.7% and 23.7% to 54.3 % and 10.9% respectively (Table 1). For stage 2 the prevalence rose steadily from 9.6% to 17.3 % in those in age band

75 to 84 but was 7.1% in the oldest age band. For participants in stage 3, the prevalence increased from 4.0% to 16.9%. Stage 4 representing the most advanced cases of AMD was rare in the two younger age bands, was infrequent in the age groups between 65 and 84 and occurred at 10.9% in the oldest age group. Prevalence rates were similar across sexes (Supplementary table 2).

Prevalence by graded features

The prevalence of features of AMD based on color or OCT grading is shown in Table 2. With color detection the prevalence of focal hyperpigmentation increased from 1.0 to 9.6% with age. Small drusen <63 μm rose from 7.9% in the youngest to 22.9% in those older than 85 years. The prevalence of drusen >125 μm rose steadily from 1.3% to 9.6% doubling by each age category but the oldest two age bands were similar. RPD frequencies were similar for the two youngest age bands but showed steady rises in the three oldest age categories. On OCT SDD were observed frequently on OCT in all age bands compared to RPD on color. These differentials were striking 18.0% on OCT versus 1.1% on color in the youngest age category and 53.6% versus 12.0% in those > 85 years. The prevalence of GA was low on color and only seen in the two oldest age bands at 0.5% and 1.2%. This contrasted with the higher proportions of cRORA on OCT, which rose from 2.5% from those under 55 year to 16.5% in those aged 85 and above. The prevalence of all the graded features was not significantly different between men and women (Supplementary table 3).

Prevalence of AMD features on UWFI color imaging

The prevalence of hard and soft drusen and hyperpigmentation in UWFI color images by age groups are shown in table 3.

Small hard drusen were highly prevalent in the central (62.8%), mid (86.8%) and far periphery (75.8%) of the fundus with the highest prevalence's seen in the mid region but no significant differences were seen with age and sex (sex data not shown). The prevalence of soft drusen rose monotonically with age in the central fundus, with similar rises in the mid and far periphery. Hyperpigmentation increased with age in all 3 regions, but the prevalence by age group and rise with increasing age group was most marked in the far periphery. Table 4 shows the relationship between Beckman stage and the UWFI grades. The UWFI based prevalence of soft drusen in central, mid and far regions rose with increasing Beckman stage.

The prevalence of hard drusen in all UWFI regions was similar across all Beckman stages. The prevalence of hyperpigmentation increased with increasing Beckman stage.

Genetic Risk Score (GRS)

The GRS ranged from -3.0 to 5.1 (SD 1.1) and was normally distributed. In controls (Beckman 0,1) the mean GRS was 0.4 (SD 1.1). For cases (Beckman 2,3,4) the GRS score was higher 0.7 (SD 1.2) (supplementary table 4). Mean GRS (SD) for AMD features detected on color were hyperpigmentation 0.4 (1.2), drusen (any size) 0.6 (1.2), drusen>63 μm 0.7 (1.2), drusen >125 μm 1.0 (1.4), reticular drusen 1.0 (1.2), geographic atrophy 1.6(0.8). The mean GRS for AMD features detected on OCT were classic drusen

0.8 (1.2), SDD 0.7(1.2), SDD only 0.5 (1.1), cRORA 0.8 (1.3), CNV 1.4 (1.1) and vitelliform lesion 0.1 (1.1).

Risk factor associations

Comparison between stage 0 or 1 versus stage 2, 3 and 4 (supplementary table 4) showed that only the genetic risk score (OR 1.25 95% CI 1.15-1.35, $p < 0.001$) and serum HDL remained highly statistically significant in the fully adjusted model. Self-reported chronic lung disease (OR 1.71 95%CI 1.11-2.64, $p = 0.014$), Parkinson's disease (OR 5.10 95% CI 1.39-18.74, $p = 0.014$) and the category of elevated hypertension (OR 1.44 95%CI 1.06-1.96 $p = 0.020$) just reached significance in the fully adjusted model. The association with hsCRP was higher in the model with GRS removed (OR 1.09 95%CI 1.00-1.19 $p = 0.045$) than when it was included (OR 1.08 95%CI 0.99-1.18 $p = 0.065$).

Risk factor associations with individual features detected by color grading are shown in supplementary Table 5. The GRS was consistently associated with classical drusen and RPD. The OR for classical drusen $< 63 \mu\text{m}$ was 1.24 (1.13, 1.37) and increased to 1.52 (1.27, 1.81) for drusen $> 125 \mu\text{m}$. The OR for RPD was 1.47 (1.23, 1.75). There was no significant association for genetic risk score and hyperpigmentation. The presence of a thickened choroid was significantly associated with drusen of any size.

Supplementary Table 6 shows the risk factor associations with individual grading features based on OCT. A consistent and highly significant association was seen between the genetic risk score and classical drusen with an OR of 1.33 (1.24, 1.43 $p < 0.001$). Eyes with SDD with classical drusen also had a statistically significant association OR 1.35 (1.26, 1.44, $p < .001$). For eyes with SDD without classical drusen the association with the genetic risk score lost significance 1.15 (1.02, 1.28, $p = 0.017$) but the direction of risk remained unchanged. The OR for cRORA was statistically significant $p = 0.003$ at 1.26 (1.08, 1.46). Classical drusen were significantly associated with thick choroid OR 1.84 (1.42, 2.37, $p < .001$) and RPD with thin choroid with an OR of 1.69 (1.27, 2.23 $p < 0.001$).

Discussion

NICOLA ascertained the prevalence of features of early AMD through traditional color imaging and extended this to include UWFI and SD OCT, permitting us to distinguish between participants based on individual characteristics. We showed clear and steady almost monotonic age-related rises in the prevalence of Beckman stages 2, 3 representing early and intermediate AMD respectively (Table 1). These findings are in accord with prior epidemiological studies.^{2 3}

In the eye level analysis, differences by age group in the frequencies of drusen were seen on comparing features detected by color versus OCT. Notably, on color, the prevalence of drusen (any size) was higher than that seen on OCT in the younger age groups and although it rose with age, the increase was shallow. On OCT the prevalence of any drusen which was around 16% in the youngest age band, rose steadily to 54% in the oldest age band. These color versus OCT discrepancies mainly occurred in the

detection of drusen $< 63\mu\text{M}$. Our data suggests that small drusen were more likely to be graded as present in younger age groups and likely represented an over calling of this feature by the graders.^{9 14} By contrast, small drusen when present were more likely to be missed on color grading in the older age groups in whom there is a higher prevalence of lens opacities with resultant degradation of image quality interfering with the detection process.

We observed good correspondence for prevalence of drusen $>125\ \mu\text{M}$ on comparing detection by *color* or by OCT. We demonstrated poor agreement for the prevalence RPD comparing detecting by *en face* technologies versus its OCT correlate of SDD, a finding which is in keeping with previous studies.^{10 15 16} We expected that the OCT correlate of pseudo drusen; subretinal drusenoid deposits would be detected at higher frequency than that reported by *en face* imaging but the magnitude of the difference was surprising. Even though the prevalence on *en face* imaging of RPD was marginally lower in NICOLA at 3.2% than the Rotterdam study (4.9%) or that reported from another UK cohort (5.06%)¹⁷ it was still much higher than that of a large community-based-cohort-study in Australia¹⁸ (0.41%). Studies that have either used OCT alone or in conjunction with other imaging modalities have generally reported high prevalence rates of SDD. Alienor found the prevalence of SDD to be 13.4% using multimodal imaging¹⁹ in contrast to the Alstar study who reported a prevalence of 32% in their clinic based enrolment cohort²⁰ using multimodal imaging. In NICOLA we found the prevalence to be 25.7% in those aged 55 and older. We contend that NICOLA offers better representation of the true population prevalence of SDD in older adults as its community-based sampling strategy is less likely to be biased in the direction of persons with other ocular morbidities which are common in clinic-based samples.

As with other epidemiological studies the prevalence of large areas of atrophy visible on color images representing GA was low, precluding generation of robust estimates of prevalence and risk factor associations for this late stage of AMD. Nonetheless we were able to characterize in detail the presence of cRORA an OCT based definition of focal atrophy in the outer retina,²¹ using SD OCT and indeed NICOLA is the first epidemiological study to record its prevalence. The proportion of eyes exhibiting cRORA even in the younger age groups was around 2.5% and this rose steadily with age allowing us to provide robust estimates by age-band for focal outer retinal atrophy. This information will be particularly useful for sample size calculations when developing protocols for GA interventional trials. In this context we recognise that while it is presently unknown if cRORA is a robust precursor of geographic atrophy, longitudinal case series²² and data from clinical trials²³ that have enrolled participants with early AMD who have progressed to GA strongly support this view.

Peripheral retinal changes have been reported in several clinical AMD cohorts^{24 25} and one other population-based study²⁶. The prevalence in the periphery of small hard drusen, soft drusen and pigmentary irregularities in the UWFI images in the NICOLA population were in accord with the high rates of abnormalities reported in these prior studies. It was notable that increasing Beckman severity stages was mirrored by increases in soft drusen and hyperpigmentation in the central, mid and far periphery. By

contrast the prevalence of hard drusen was similar in the central mid and far periphery across all Beckman severity stages suggesting that these are a ubiquitous finding. Histological studies however dispute these findings and show that the pathology visible in the periphery is not the same as that in the macula.²⁷ Widefield OCT images should be prioritised in future studies as these may help resolve these disparities in clinical cohorts.

Various risk factors for intermediate and late AMD have been identified in longitudinal epidemiological studies²⁸ and clinical cohorts.^{29 30} The fully adjusted multivariate regression model revealed that age and the GRS were the only highly statistically significant associations with the Beckman severity stage person level classification. Associations between AMD and chronic lung disease is relatively novel though has recently been reported from a population based retrospective study in Taiwan³¹ so deserves further exploration. Some factors such as physical activity that were significant in univariate are likely to be highly collinear with age hence drop from age-adjusted models. The OR's and risk estimates for the GRS and Beckman stages of intermediate and late AMD are in accord those of the Eye-risk consortium which recently reported a similar mean GRS from their large, pooled analysis of cross-sectional data from the European Eye Epidemiology Consortium.³²

In the AMD feature level analysis, the GRS was strongly associated with classical drusen, on separating eyes with SDD from eyes with both SDD and classical drusen, the association with the GRS lost significance despite previous studies showing a significant association between SDD and the two major AMD risk loci independent of drusen presence (ARMS2 positively associated and CFH Y402H negatively associated).³³ We also observed that GRS was not significantly associated with hyperpigmentation when this feature present in the absence of drusen. This is not surprising since focal hyperpigmentation on its own can represent pathology such as past inflammation.

All types of drusen were associated with a thicker choroid which is in keeping with many previous studies³⁴⁻³⁶ whereas those with SDD alone had a significantly thinner choroid. A detailed study of the relationship between choroidal thickness, choroidal vascularity index and SDD presence by Keenan et al (2020)³⁵ reported a biphasic alteration in choroidal dimensions across the disease spectrum with those with large drusen showing increased choroidal thickness and increased choroidal vascularity index (CVI) whereas the same parameters in those with advanced AMD in the fellow eye were no different to controls. Those with just SDD had significantly thinner choroid and reduced choroidal vascularity index.³⁵ Keenan et al propose CVI as a potential biomarker of aging given its significant and negative correlation with age and our data on SDD alone suggests that it too may be more reflective of ubiquitous aging rather than AMD per se. It is also interesting to note SDD alone also had a lower GRS than the other drusen related features though still significantly associated in keeping with recent genetic studies showing significant associations with the major AMD susceptibility loci.³⁷

Limitations

Our study suffers from several limitations. Firstly, the response rate for those who attended the health assessment was moderate, but we implemented appropriate weighting strategies to mitigate the effects of such bias. Nonetheless when weighted and unweighted prevalence estimates were compared (Table 1) the differences were minor suggesting that the cohort demographic structure closely matched that of the general population. Secondly, we investigated a large number of potential risk factors a process which can increase the risk of false positive results. We therefore took a highly conservative approach in the creation of the multivariable models and interpreted the findings in terms of effect size and biological plausibility rather than explicit p-value cut-offs. Thirdly, we did not grade for intraretinal hyperreflective foci an OCT feature, which is now considered a biomarker of deteriorating RPE health and a predictor for progression to late AMD³⁸.

Conclusions

This study provides further insight into the prevalence and risk factors of AMD and AMD features using multiple imaging modalities. Interestingly the correlation between the Beckman Classification and our findings from UWF imaging provide evidence that on a pragmatic level that the former continues to have validity. It highlights the benefits of using a multi modal approach in future epidemiological studies but also the challenges in interpreting findings that can be compared with previous colour only studies. New severity stage systems which incorporate AMD based OCT features are urgently needed.

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References

1. Klein R, Klein BE, Moss SE. Diabetes, hyperglycemia, and age-related maculopathy. The Beaver Dam Eye Study. *Ophthalmology* 1992;99(10):1527-34. doi: 10.1016/s0161-6420(92)31770-1 [published Online First: 1992/10/01]
2. Vingerling JR, Dielemans I, Hofman A, et al. The prevalence of age-related maculopathy in the Rotterdam Study. *Ophthalmology* 1995;102(2):205-10. doi: 10.1016/s0161-6420(95)31034-2 [published Online First: 1995/02/01]

3. Mitchell P, Smith W, Attebo K, et al. Prevalence of age-related maculopathy in Australia. The Blue Mountains Eye Study. *Ophthalmology* 1995;102(10):1450-60. doi: 10.1016/s0161-6420(95)30846-9 [published Online First: 1995/10/01]
4. Buitendijk GHS, Rochtchina E, Myers C, et al. Prediction of age-related macular degeneration in the general population: the Three Continent AMD Consortium. *Ophthalmology* 2013;120(12):2644-55. doi: 10.1016/j.ophtha.2013.07.053 [published Online First: 2013/10/15]
5. Joachim N, Colijn JM, Kifley A, et al. Five-year progression of unilateral age-related macular degeneration to bilateral involvement: the Three Continent AMD Consortium report. *Br J Ophthalmol* 2017;101(9):1185-92. doi: 10.1136/bjophthalmol-2016-309729 [published Online First: 2017/01/22]
6. Joachim N, Kifley A, Colijn JM, et al. Joint Contribution of Genetic Susceptibility and Modifiable Factors to the Progression of Age-Related Macular Degeneration over 10 Years: The Three Continent AMD Consortium Report. *Ophthalmol Retina* 2018;2(7):684-93. doi: 10.1016/j.oret.2017.10.019 [published Online First: 2019/05/03]
7. Saunier V, Merle BMJ, Delyfer MN, et al. Incidence of and Risk Factors Associated With Age-Related Macular Degeneration: Four-Year Follow-up From the ALIENOR Study. *JAMA Ophthalmol* 2018;136(5):473-81. doi: 10.1001/jamaophthalmol.2018.0504 [published Online First: 2018/03/30]
8. Mimoun G, Soubrane G, Coscas G. [Macular drusen]. *J Fr Ophtalmol* 1990;13(10):511-30. [published Online First: 1990/01/01]
9. Graham KW, Chakravarthy U, Hogg RE, et al. IDENTIFYING FEATURES OF EARLY AND LATE AGE-RELATED MACULAR DEGENERATION: A Comparison of Multicolor Versus Traditional Color Fundus Photography. *Retina* 2018;38(9):1751-58. doi: 10.1097/IAE.0000000000001777 [published Online First: 2017/08/24]
10. Hogg RE, Silva R, Staurengi G, et al. Clinical characteristics of reticular pseudodrusen in the fellow eye of patients with unilateral neovascular age-related macular degeneration. *Ophthalmology* 2014;121(9):1748-55. doi: 10.1016/j.ophtha.2014.03.015 [published Online First: 2014/05/27]
11. Team RC. R: a language and environment for statistical computing. Vienna, Austria.: R Foundation for Statistical Computing 2019.
12. Colijn JM, Buitendijk GHS, Prokofyeva E, et al. Prevalence of Age-Related Macular Degeneration in Europe: The Past and the Future. *Ophthalmology* 2017;124(12):1753-63. doi: 10.1016/j.ophtha.2017.05.035 [published Online First: 2017/07/18]
13. van Buuren S, Groothuis-Oudshoorn, K. . mice: Multivariate Imputation by Chained Equations in R. *Journal of Statistical Software* 2011;45:1-67.
14. Farinha C, Cachulo ML, Coimbra R, et al. Age-Related Macular Degeneration Staging by Color Fundus Photography vs. Multimodal Imaging-Epidemiological Implications (The Coimbra Eye Study-Report 6). *J Clin Med* 2020;9(5) doi: 10.3390/jcm9051329 [published Online First: 2020/05/07]
15. Wu Z, Ayton LN, Luu CD, et al. Reticular Pseudodrusen in Intermediate Age-Related Macular Degeneration: Prevalence, Detection, Clinical, Environmental, and Genetic Associations. *Invest Ophthalmol Vis Sci* 2016;57(3):1310-6. doi: 10.1167/iovs.15-18682 [published Online First: 2016/03/22]
16. De Bats F, Mathis T, Mauget-Faysse M, et al. Prevalence of Reticular Pseudodrusen in Age-Related Macular Degeneration Using Multimodal Imaging. *Retina* 2016;36(1):46-52. doi: 10.1097/IAE.0000000000000648 [published Online First: 2015/06/20]
17. Wilde C, Poostchi A, Mehta RL, et al. Prevalence of reticular pseudodrusen in an elderly UK Caucasian population-The Bridlington Eye Assessment Project (BEAP): a cross-sectional study (2002-2006). *Eye (Lond)* 2018;32(6):1130-37. doi: 10.1038/s41433-018-0049-8 [published Online First: 2018/03/02]
18. Finger RP, Chong E, McGuinness MB, et al. Reticular Pseudodrusen and Their Association with Age-Related Macular Degeneration: The Melbourne Collaborative Cohort Study. *Ophthalmology* 2016;123(3):599-608. doi: 10.1016/j.ophtha.2015.10.029 [published Online First: 2015/12/19]

19. Chan H, Cougnard-Gregoire A, Delyfer MN, et al. Multimodal Imaging of Reticular Pseudodrusen in a Population-Based Setting: The Alienor Study. *Invest Ophthalmol Vis Sci* 2016;57(7):3058-65. doi: 10.1167/iovs.16-19487 [published Online First: 2016/07/02]
20. Zarubina AV, Neely DC, Clark ME, et al. Prevalence of Subretinal Drusenoid Deposits in Older Persons with and without Age-Related Macular Degeneration, by Multimodal Imaging. *Ophthalmology* 2016;123(5):1090-100. doi: 10.1016/j.ophtha.2015.12.034 [published Online First: 2016/02/15]
21. Sadda SR, Guymer R, Holz FG, et al. Consensus Definition for Atrophy Associated with Age-Related Macular Degeneration on OCT: Classification of Atrophy Report 3. *Ophthalmology* 2018;125(4):537-48. doi: 10.1016/j.ophtha.2017.09.028 [published Online First: 2017/11/07]
22. Jaffe GJ, Chakravarthy U, Freund KB, et al. Imaging Features Associated with Progression to Geographic Atrophy in Age-Related Macular Degeneration: CAM Report 5. *Ophthalmol Retina* 2020 doi: 10.1016/j.oret.2020.12.009 [published Online First: 2020/12/22]
23. Guymer RH, Wu Z, Hodgson LAB, et al. Subthreshold Nanosecond Laser Intervention in Age-Related Macular Degeneration: The LEAD Randomized Controlled Clinical Trial. *Ophthalmology* 2019;126(6):829-38. doi: 10.1016/j.ophtha.2018.09.015 [published Online First: 2018/09/24]
24. Writing Committee for the OPRs, Domalpally A, Clemons TE, et al. Peripheral Retinal Changes Associated with Age-Related Macular Degeneration in the Age-Related Eye Disease Study 2: Age-Related Eye Disease Study 2 Report Number 12 by the Age-Related Eye Disease Study 2 Optos PEripheral Retina (OPERA) Study Research Group. *Ophthalmology* 2017;124(4):479-514 87. doi: 10.1016/j.ophtha.2016.12.004 [published Online First: 2017/01/17]
25. Tan CS, Heussen F, Sadda SR. Peripheral autofluorescence and clinical findings in neovascular and non-neovascular age-related macular degeneration. *Ophthalmology* 2013;120(6):1271-7. doi: 10.1016/j.ophtha.2012.12.002 [published Online First: 2013/02/26]
26. Lengyel I, Csutak A, Florea D, et al. A Population-Based Ultra-Widefield Digital Image Grading Study for Age-Related Macular Degeneration-Like Lesions at the Peripheral Retina. *Ophthalmology* 2015;122(7):1340-7. doi: 10.1016/j.ophtha.2015.03.005 [published Online First: 2015/04/15]
27. Rudolf M, Clark ME, Chimento MF, et al. Prevalence and morphology of druse types in the macula and periphery of eyes with age-related maculopathy. *Invest Ophthalmol Vis Sci* 2008;49(3):1200-9. doi: 10.1167/iovs.07-1466 [published Online First: 2008/03/11]
28. Joachim N, Mitchell P, Kifley A, et al. Incidence and progression of geographic atrophy: observations from a population-based cohort. *Ophthalmology* 2013;120(10):2042-50. doi: 10.1016/j.ophtha.2013.03.029 [published Online First: 2013/05/28]
29. Grunwald JE, Daniel E, Huang J, et al. Risk of geographic atrophy in the comparison of age-related macular degeneration treatments trials. *Ophthalmology* 2014;121(1):150-61. doi: 10.1016/j.ophtha.2013.08.015 [published Online First: 2013/10/03]
30. Wu Z, Luu CD, Hodgson LAB, et al. Prospective Longitudinal Evaluation of Nascent Geographic Atrophy in Age-Related Macular Degeneration. *Ophthalmol Retina* 2020;4(6):568-75. doi: 10.1016/j.oret.2019.12.011 [published Online First: 2020/02/24]
31. Bair PJ, Hsia NY, Lin CL, et al. Population-based retrospective cohort study on risk of age-related macular degeneration in people with chronic obstructive pulmonary disease. *Sci Rep* 2021;11(1):15079. doi: 10.1038/s41598-021-94657-9 [published Online First: 2021/07/25]
32. Colijn JM, Meester-Smoor M, Verzijden T, et al. Genetic Risk, Lifestyle, and Age-Related Macular Degeneration in Europe: The EYE-RISK Consortium. *Ophthalmology* 2020 doi: 10.1016/j.ophtha.2020.11.024 [published Online First: 2020/12/01]
33. Lin LY, Zhou Q, Hagstrom S, et al. Association of Single-Nucleotide Polymorphisms in Age-Related Macular Degeneration With Pseudodrusen: Secondary Analysis of Data From the Comparison of AMD Treatments Trials. *JAMA Ophthalmol* 2018;136(6):682-88. doi: 10.1001/jamaophthalmol.2018.1231 [published Online First: 2018/05/26]

34. Lee J, Kim M, Lee CS, et al. Drusen Subtypes and Choroidal Characteristics in Asian Eyes with Typical Neovascular Age-Related Macular Degeneration. *Retina* 2020;40(3):490-98. doi: 10.1097/IAE.0000000000002419 [published Online First: 2018/12/15]
35. Keenan TD, Klein B, Agron E, et al. Choroidal Thickness and Vascularity Vary with Disease Severity and Subretinal Drusenoid Deposit Presence in Nonadvanced Age-Related Macular Degeneration. *Retina* 2019 doi: 10.1097/IAE.0000000000002434 [published Online First: 2019/01/22]
36. Velaga SB, Nittala MG, Vupparaboina KK, et al. Choroidal Vascularity Index and Choroidal Thickness in Eyes with Reticular Pseudodrusen. *Retina* 2019 doi: 10.1097/IAE.0000000000002667 [published Online First: 2019/10/22]
37. Domalpally A, Agron E, Pak JW, et al. Prevalence, Risk, and Genetic Association of Reticular Pseudodrusen in Age-related Macular Degeneration: Age-Related Eye Disease Study 2 Report 21. *Ophthalmology* 2019;126(12):1659-66. doi: 10.1016/j.ophtha.2019.07.022 [published Online First: 2019/09/29]
38. Nassisi M, Lei J, Abdelfattah NS, et al. Risk Factors for Development of Late Age-Related Macular Degeneration in the Fellow Eyes of Patients Enrolled in the HARBOR Study. *Ophthalmology* 2019;