

See discussions, stats, and author profiles for this publication at: <https://www.researchgate.net/publication/326621068>

The association between blood vitamins D and E with age-related macular degeneration: A pilot study

Article in *Interventional Medicine and Applied Science* · July 2018

DOI: 10.1556/1646.10.2018.22

CITATIONS

0

READS

51

8 authors, including:



Rezvan Hashemi

Tehran University of Medical Sciences

19 PUBLICATIONS 126 CITATIONS

SEE PROFILE



Mahin Bandarian

Tehran University of Medical Sciences

5 PUBLICATIONS 36 CITATIONS

SEE PROFILE



Elahe Abedi-Taleb

4 PUBLICATIONS 5 CITATIONS

SEE PROFILE



Leila Khedmat

Baqiyatallah University of Medical Sciences

24 PUBLICATIONS 47 CITATIONS

SEE PROFILE

Some of the authors of this publication are also working on these related projects:



The Association of General and Central Obesity with Major Dietary Patterns of Adult Women Living in Tehran, Iran [View project](#)



the relationship between pregnant mothers intake on newborn weight [View project](#)

The association between blood vitamins D and E with age-related macular degeneration: A pilot study

ASGHAR MOLLAZADEH JELODAR^{1,2,*}, REZVAN HASHEMI¹, MAHIN BANDARIAN²,
ELAHE ABEDI-TALEB³, HASSAN KHOJASTEH⁴, LEILA KHEDMAT⁵, ELNAZ ASADOLLAHI⁶,
MINA BEYTOLLAHI⁷

¹Department of Ophthalmology, Ziaean Hospital, Tehran University of Medical Sciences, Tehran, Iran

²Department of Geriatric Medicine, Ziaean Hospital, School of Medicine, Tehran University of Medical Sciences, Tehran, Iran

³Department of Obstetric and Gynecology, Ziaean Hospital, School of Medicine, Tehran University of Medical Sciences, Tehran, Iran

⁴Ziaean Hospital, Tehran University of Medical Sciences, Tehran, Iran

⁵Department of Community Medicine, School of Medicine, Tehran University of Medical Science, Tehran, Iran

⁶Ophthalmology Research Center, Farabi Eye Hospital, Tehran University of Medical Sciences, Tehran, Iran

⁷Department and Faculty of Medical Sciences and Technology, Islamic Azad University, Tehran, Iran

*Corresponding author: Asghar Mollazadeh Jelodar; Department of Geriatric Medicine, Ziaean Hospital, School of Medicine, Tehran University of Medical Sciences, Tehran 1366736511, Iran; Phone: +98 91 2387 3622; Fax: +98 55 751 333;

E-mail: dr.as.mollazadeh@gmail.com

(Received: February 27, 2018; Revised manuscript received: March 21, 2018; Accepted: April 16, 2018)

Abstract: *Background:* This study was aimed to evaluate the association of serum vitamins D and E level with age-related macular degeneration (AMD). *Methods:* This pilot study was performed in two groups of 15 patients in treatment group and 15 patients in control group. Measurements of blood factors [such as C-reactive protein (CRP) and high-density lipoprotein (HDL)] were performed after 12 h of fasting. To measure vitamins D and E, the serum was isolated from 5 cc blood samples. *Results:* HDL was higher in the control group as compared with the AMD group. However, no significant difference was found between the two groups ($p = 0.08$). On the other hand, serum vitamin E in the AMD group was remarkably higher as compared to the control group ($p < 0.002$). However, no significant difference was found in serum vitamin D levels between the two groups ($p = 0.662$). Our findings also revealed that there was no statistically significant relationship between BMI and AMD. Moreover, no significant correlation was determined between serum CRP and AMD ($p = 0.96$). *Conclusions:* Our data indicated that none provides evidence for associations between AMD and serum vitamin D levels. The association between vitamin D and AMD requires further investigations in a large population studies, to elucidate whether vitamin D deficiency can be an important risk factor for AMD.

Keywords: vitamins D and E, serum, healthy lifestyles, blood factors, age-related macular degeneration

Introduction

Age-related macular degeneration (AMD) is a progressive disease that may be associated with blurred or no vision in the center of the visual field [1]. Macular degeneration causes the macula to lose its natural function. It can be damaged by several factors, such as diabetes, genetic disease, and the consumption of medications, including chloroquine and hypertension [2]. The prevalence of this disease in the United States is high, and more than

14% of people aged 80 years and older are wholly involved with it [3]. It has also been estimated in 2000 that more than 9 million people have been infected with AMD [4]. The disease is expected to double in 2020 [5]. The prevalence of this disease in a study between the ages of 40 and 65 years old was 4.7%, which is lower than the western countries and is higher than the eastern countries [6]. Common risk factors for macular degeneration include smoking [7], high blood pressure [8], obesity [9], whiteness [10], low high-density lipoprotein (HDL) and

This is an open-access article distributed under the terms of the [Creative Commons Attribution-NonCommercial 4.0 International License](https://creativecommons.org/licenses/by-nc/4.0/), which permits unrestricted use, distribution, and reproduction in any medium for non-commercial purposes, provided the original author and source are credited, a link to the CC License is provided, and changes – if any – are indicated.

high low-density lipoprotein (LDL) [11], C-reactive protein (CRP) [12], exposed to light sunshine [13], genetics and family history, and immune and inflammatory disorders [7]. Apolipoprotein E (ApoE) is the main apolipoprotein of the CNS, which plays a role in the transmission of lipids and cholesterol in the body. Three forms E2, E3, and E4 are known for the ApoE allelic variants associated with AMD. E4 allele has anti-AMD effects, whereas E2 allele increases the risk of the disease, E3 allele does not have an effect on the chance of developing [14]. Studies have focused on the role of some nutritional compounds on macular degeneration [15–19]. Therefore, vitamins and minerals can reduce the complications of the disease and improve vision [20]. Particularly, foods rich in antioxidants, such as vitamin E, play an important role in inflammation and oxidative stress in this pathway [16, 21]. Omega-3, a member of the family of polyunsaturated fatty acids, is not synthesized in the body; therefore, the intake of this fatty acid is important. In this group, docosahexaenoic acid and eicosapentaenoic acid have beneficial effects on the body. These compounds contain a small percentage of the tissue fatty acids, but they are abundant in the retina, so their deficiency may alter the function of the retina [22, 23]. The beneficial effects of omega-3 have been implicated in inflammatory diseases [24, 25], which is also beneficial in its anti-inflammatory function in the retina. Inflammation creates a redistribution of the choroid arteries in the wet form of AMD [26]. Several studies have shown anti-inflammatory effects of vitamin D [27–29]. Therefore, it is likely that vitamin D deficiency is also related to the incidence of AMD.

If macular degeneration is not prevented, severe vision impairment or vision loss can occur, requiring various treatments, such as Avastin injection and photocoagulation laser. These treatments do not have a satisfactory result for the patient, despite the high financial burden for the patient. On the other hand, there is no disease for the dry type. Regarding the increase in the aging population of our country and the huge costs that the disease can bring to the country, it is necessary to examine the various factors that can help reduce the disease. In Iran, there is little information about dietary intake, the nutritional status of individuals, and their impact on AMD. Moreover, regarding the levels of vitamins D and E, the intake of omega-3 and lipid profiles in AMD patients has not been studied. Therefore, in this study, the relationship between serum levels of vitamins D and E, the intake of omega 3 sources, and the lipid profiles in macular degeneration have been studied in order to use this information to prevent these patients.

Materials and Methods

Ethical committee

All procedures performed in studies involving human participants were in accordance with the ethical

standards of the institutional and/or national research committee and with the 1964 Helsinki Declaration and its later amendments or comparable ethical standards and informed consent was received from all the participants in the study. The study protocol was approved by the Ethics Committee of Tehran University of Medical Sciences.

Patients study

This case–control study was conducted in two hospitals affiliated to Tehran University of Medical Sciences (Ziaeean Hospital and Farabi Hospital). People over the age of 50 have been screened for ophthalmology, patients with AMD in the case group, and those who are complaining of visual impairment and not having AMD are included in the study.

Inclusion criteria of case group:

1. Getting newly discovered AMD without treatment
2. Age over 50 years

Inclusion criteria of control group:

1. Not having AMD
2. Age over 50 years
3. People with complaints of reduced vision

Exclusion criteria:

Use of vitamin D supplementation with therapeutic doses.

From each person, general information including age, gender, history of previous illnesses, history of drug use and supplementation, history of smoking, and exposure to sunlight was obtained. The height of the people was measured in the form of standing with no shoes and with a seca wall-mounted stadiometers. The weight of subjects was measured as fasting with a minimum dress and no shoe with a seca scale and a precision of 500 g. For both the case and control groups, the Food Frequency Questionnaire was completed by a nutrition expert. In this questionnaire, 117 questions have been used to examine food intake in the elderly population of Iran by Hashemi et al. [6].

Laboratory information

All the participants were asked to come to the hospital laboratory on the next day after 12 h of fasting. In the hospital, blood sample (10 ml) was next obtained. The measurements of CRP and HDL were performed and eventually the results were reported weekly from laboratories. To measure vitamins D and E from 5 cc blood samples, the serum was isolated and maintained until use in a –20 freezer for 1 month. Serum samples are packaged into foil to measure vitamin E. The measurement of vitamin D is done using the ELISA kit and vitamin E by HPLC technique.

Method for calculating sample size

The sample size in this study was calculated to compare the mean of vitamin D levels between the two groups. If the standard deviation of vitamin D in each group is 18 (ng/ml) [30] to detect a difference of 10 ng/ml between the two groups, with a probability of 95% and a probability of committing a type I error (5%) for each group, 85 samples are required based on the following formula:

$$n = \frac{(Z_{1-\frac{\alpha}{2}} + Z_{1-\beta})^2 \times 2\sigma^2}{\Delta^2} = \frac{(1.96 + 1.64)^2 \times 2 \times 18^2}{10^2} = 85.$$

Statistical analyses

The *t*-test was used to compare the mean of vitamins D and E between AMD and control groups. Covariance analysis was also applied when considering the effects of confounding factors, such as the dietary intake of these compounds and the duration of exposure to sunlight.

Results

This is a pilot study and it is part of a major study

This is a pilot study conducted in two groups: treatment group with 15 subjects and control group with 15 subjects. The findings showed that HDL was higher in the control group than the other group, but there was no significant difference between the two groups (*p* = 0.08). In addition, serum vitamin E level in case group was significantly higher than control group (*p* < 0.002) (Table I).

However, serum level of vitamin D did not show significant difference between the two groups (*p* = 0.662). The study also revealed that there is no statistically significant relationship between BMI and AMD. In addition, there was no significant association between serum level of CRP and AMD (*p* = 0.96) (Table I).

Discussion

Our findings indicated that serum vitamin E level in AMD group was significantly higher than control group. Conversely, HDL level was higher in the control group than the AMD group, but there was no significant difference between the two groups.

In this study, there was no significant correlation between serum vitamin D level and AMD.

This result is in agreement with previous findings that indicated no significant correlation between blood 25-hydroxyvitamin D levels and advanced AMD [31], but no early AMD. Parekh has investigated the association between serum vitamin D levels and AMD on 7,752 people. The prevalence of AMD in this study was 11%, where vitamin D deficiency was determined to be inversely linked to early AMD but not advanced AMD [31].

Our finding is contrary to those of a previous study, which revealed significant correlation between blood vitamin D levels and AMD [32]. Of course, these results can be related to the low sampling rate, which is one of the limitations of this study as a pilot study and requires a comprehensive investigation. Therefore, limiting any definitive conclusions in terms of the vitamin D effects on AMD patients

A cross-sectional study was conducted in Korea to investigate the relationship between serum vitamin D

Table I | Number of clinical characteristics on means 25-hydroxyvitamin D [25(OH)D] and vitamin E status and other factors in patients in subgroups of age-related macular degeneration

Blood factors	Case (n = 15) Mean (SD)	Control (n = 15) Mean (SD)	<i>p</i>
Vitamin E	17.7 (4.9)	11.34 (4.6)	<0.002
Vitamin D	37.04 (12.6)	41.34 (17.7)	0.662
BMI	26.7 (5.03)	30.53 (29.82)	0.735
CRP	1.07 (0.25)	1.07 (0.26)	0.960
ESR	17.87 (11.7)	13.50 (11.75)	0.581
FBS	102.13 (29.30)	122.79 (53.70)	0.038
LDL	110.20 (34.82)	91.92 (16.12)	0.031
HDL	42.67 (11.78)	50.85 (8.84)	0.086
CHOL	189.13 (50.33)	168.71 (19.40)	0.002
TG	130.0 (49.85)	122.0 (62.79)	0.405

SD: standard deviation; BMI: body mass index; CRP: C-reactive protein; LDL: low-density lipoprotein; HDL: high-density lipoprotein; ESR: erythrocyte sedimentation rate; FBS: fasting blood sugar; TG: triglyceride; CHOL: cholesterol

levels and AMD. About 17,045 people who referred to the hospital during the period from 2008 to 2012 were enrolled. Therefore, after measuring the serum levels of vitamin D, they were subjected to eye examinations and imaging. Of those, 1,163 were diagnosed with AMD at an early stage and 115 subjects were advanced AMD patients. The mean serum vitamin D level in women was 17.5 ng/ml and in men was 20 ng/ml. The findings of mentioned survey demonstrated that AMD disease at advanced levels was significantly associated with a decrease in blood 25-hydroxyvitamin D levels in men [30, 32].

Tan et al. [33] examined the association between the intakes of fatty acids with the incidence of AMD for 10 years. In the mentioned study, 2,454 AMD patients participated in the study, which were initially evaluated after 5 years and finally after 10 years. The photos were taken from retina and the Food Frequency Questionnaire was also obtained. The results showed that the use of fish in a meal during a week reduces the risk of developing AMD. Moreover, nutrients use of 1–2 units per week resulted in a reduction in AMD [33]. Another study in 2006 evaluated the relationship between fish consumption, omega-3, and smoking among 681 twins, of which 222 were twins with AMD and 459 without AMD. The results showed that the risk of developing AMD increased by smoking, while fish consumption (at least twice a week) and omega-3 would reduce its risk [34].

As matter of fact, high level of blood 25-hydroxyvitamin D was inversely linked to late AMD in men but not women. A study in North Carolina was conducted in 2015 to compare the levels of vitamin D in neovascular AMD (146 individuals), non-neovascular AMD (260 individuals), and non-AMD control (100 individuals). This study showed that vitamin D in neovascular AMD was significantly lower than the other two groups [30], where vitamin D deficiency was more prevalent in neovascular AMD patients. A cross-sectional study in Denmark (2013) has been aimed at investigating the relationship between AMD disease and serum vitamin D levels. In the mentioned study, 129 subjects with different degrees of AMD and 49 controls were enrolled. The results showed that there is a significant difference between serum levels of vitamin D in control subjects (75.6 nmol/L) and AMD patients with grade 5 (47.3 nmol/L) [35]. Furthermore, the association of serum vitamin D level with AMD in the early stages was studied in 1,313 women with AMD and 1,287 without AMD [36]. Another study was conducted in Italy in 2002, in which 49 patients with AMD and 46 controls were included to compare vitamins C, E, and beta-cryptoxanthin levels. The results revealed a significant decrease in these compounds in the AMD group compared to the control group [37]. A study reported that serum levels of zinc and vitamin E in the AMD group were significantly lower than that of the control group [38].

In general, it should be noted that the sun exposure and food intake during the recent weeks, instead of the year, will increase the chance of random measurement error. Various factors may affect the outcome of the study, including unknown, unmeasured healthy lifestyles risk, or protective factors that are more common among people whose high levels of vitamin D have compared to low serum vitamin D levels [39, 40]

On the other hand, our data showed that serum vitamin E in the AMD group was significantly higher than the control group. In contrast with these findings, a study by Christen et al. [41] on 117 cases of AMD in the vitamin E group has reported that vitamin E treatment had no large beneficial or harmful effect on risk of advanced AMD. Furthermore, another study by Timms et al. [42] revealed that vitamin D intake decreases CRP, a marker of systemic inflammation. Moreover, the two studies by Seddon et al. [43, 44] have found associations between markers of inflammation (such as CRP) and AMD, but Klein et al. [45] have suggested that there was no statistically significant association between CRP and AMD. In parallel, our data showed that no significant correlation was determined between serum CRP and AMD. In addition, McKay et al.'s [46] study on 4,753 participants have been observed that no association was found with vitamin D and early or late AMD or neovascular AMD. There was no association between insufficient or deficient status with early or late AMD. This is in agreement with our finding.

In conclusion, this study conducted in a small sample as pilot study of the Iran population and none provides evidence for associations between AMD and serum vitamin D levels. In addition, this study does not show any relationship between serum CRP and AMD levels. Our results warrant reconsidering the existence of an association between vitamin D and the occurrence and progression of AMD using a large population.

* * *

Funding sources: This study had no funding body.

Authors' contribution: AMJ, RH, MB, EA-T, HK, LK, EA, and MB participated in conceiving and designing the study, and participated in drafting the manuscript, statistical analyses, administrative, technical, and material support. All authors read and approved the final manuscript.

Conflict of interest: None.

References

1. Van Lookeren Campagne M, LeCouter J, Yaspan BL, Ye W: Mechanisms of age-related macular degeneration and therapeutic opportunities. *J Pathol* 232, 151–164 (2014)
2. Centers for Disease Control and Prevention (2000): Measuring healthy days. CDC, Atlanta

3. Fard ZG, Schneider S, Hudson JL, Habibi M, Pooravari M, Heidari ZH: Early maladaptive schemas as predictors of child anxiety: The role of child and mother schemas. *Int J Appl Behav Sci* 1, 9–18 (2015)
4. Friedman DS, O'Colmain BJ, Munoz B, Tomany SC, McCarty C, De Jong P, Nemesure B, Mitchell P, Kempen J: Prevalence of age-related macular degeneration in the United States. *Arch ophthalmol* 122, 564–572 (2004)
5. World Health Organization (2010): Prevention of blindness and visual impairment. Priority eye diseases-corneal opacities. Retrieved from <http://www.who.int/blindness/causes/priority/en/index9.html>. Accessed on: July 2014
6. Hashemi H, Ghafari E, Khabazkhoob M, Noori J, Taheri A, Eshghabadi A, Khodabandeh A, Emamian MH, Shariati M, Fotouhi A: Age-related macular degeneration in an Iranian population. *Iran J Ophthalmol* 26, 203–211 (2014)
7. Lambert NG, Singh MK, ElShelmani H, Mansergh FC, Wride MA, Padilla M, Keegan D, Hogg RE, Ambati BK: Risk factors and biomarkers of age-related macular degeneration. *Prog Retin Eye Res* 54, 64–102 (2016)
8. Churchill AJ, Carter JG, Lovell HC, Ramsden C, Turner SJ, Yeung A, Escardo J, Atan D: VEGF polymorphisms are associated with neovascular age-related macular degeneration. *Hum Mol Genet* 15, 2955–2961 (2006)
9. Seddon JM, Cote J, Davis N, Rosner B: Progression of age-related macular degeneration: Association with body mass index, waist circumference, and waist-hip ratio. *Arch Ophthalmol* 121, 785–792 (2003)
10. Chang MA, Bressler SB, Munoz B, West SK: Racial differences and other risk factors for incidence and progression of age-related macular degeneration: Salisbury Eye Evaluation (SEE) Project. *Invest Ophthalmol Vis Sci* 49, 2395–2402 (2008)
11. Reynolds R, Rosner B, Seddon JM: Serum lipid biomarkers and hepatic lipase gene associations with age-related macular degeneration. *Ophthalmology* 117, 1989–1995 (2010)
12. Seddon JM, George S, Rosner B, Rifai N: Progression of age-related macular degeneration: Prospective assessment of C-reactive protein, interleukin 6, and other cardiovascular biomarkers. *Arch Ophthalmol* 123, 774–782 (2005)
13. Coleman HR, Chan C-C, Ferris FL, Chew EY: Age-related macular degeneration. *Lancet* 372, 1835–1845 (2008)
14. Baird PN, Guida E, Chu DT, Vu HT, Guymer RH: The $\epsilon 2$ and $\epsilon 4$ alleles of the apolipoprotein gene are associated with age-related macular degeneration. *Invest Ophthalmol Vis Sci* 45, 1311–1315 (2004)
15. Cho E, Seddon JM, Rosner B, Willett WC, Hankinson SE: Prospective study of intake of fruits, vegetables, vitamins, and carotenoids and risk of age-related maculopathy. *Arch Ophthalmol* 122, 883–892 (2004)
16. Ciulla TA, Walker JD, Fong DS, Criswell MH: Corticosteroids in posterior segment disease: An update on new delivery systems and new indications. *Curr Opin Ophthalmol* 15, 211–220 (2004)
17. Seddon JM, Ajani UA, Sperduto RD, Hiller R, Blair N, Burton TC, Farber MD, Gragoudas ES, Haller J, Miller DT: Dietary carotenoids, vitamins A, C, and E, and advanced age-related macular degeneration. *JAMA* 272, 1413–1420 (1994)
18. Seddon JM, Gensler G, Klein ML, Milton RC: C-reactive protein and homocysteine are associated with dietary and behavioral risk factors for age-related macular degeneration. *Nutrition* 22, 441–443 (2006)
19. Van den Langenberg GM, Mares-Perlman JA, Klein R, Klein BE, Brady WE, Palta M: Associations between antioxidant and zinc intake and the 5-year incidence of early age-related maculopathy in the Beaver Dam Eye Study. *Am J Epidemiol* 148, 204–214 (1998)
20. Group A-REDSR: A randomized, placebo-controlled, clinical trial of high-dose supplementation with vitamins C and E, beta carotene, and zinc for age-related macular degeneration and vision loss: AREDS report no. 8. *Arch Ophthalmol* 119, 1417 (2001)
21. van Leeuwen R, Boekhoorn S, Vingerling JR, Witteman JC, Klaver CC, Hofman A, de Jong PT: Dietary intake of antioxidants and risk of age-related macular degeneration. *JAMA* 294, 3101–3107 (2005)
22. SanGiovanni JP, Chew EY: The role of omega-3 long-chain polyunsaturated fatty acids in health and disease of the retina. *Prog Ret Eye Res* 24, 87–138 (2005)
23. SanGiovanni JP, Chew EY, Clemons TE, Davis MD, Ferris F 3rd, Gensler GR, Kurinij N, Lindblad AS, Milton RC, Seddon JM, Sperduto RD, Age-Related Eye Disease Study Research Group: The relationship of dietary lipid intake and age-related macular degeneration in a case-control study: AREDS report No. 20. *Arch Ophthalmol* 125, 671–679 (2007)
24. Turner D, Zlotkin SH, Shah PS, Griffiths AM: Omega 3 fatty acids (fish oil) for maintenance of remission in Crohn's disease. *Cochrane Database Syst Rev* CD006320 (2009)
25. James M, Proudman S, Cleland L: Dietary n-3 fats as adjunctive therapy in a prototypic inflammatory disease: Issues and obstacles for use in rheumatoid arthritis. *Prostaglandins Leukot Essent Fatty Acids* 68, 399–405 (2003)
26. Ozaki E, Campbell M, Kiang A-S, Humphries M, Doyle SL, Humphries P: Inflammation in age-related macular degeneration. *Adv Exp Med Biol* 801, 229–235 (2014)
27. Manolagas S, Provvedini D, Murry E, Tsoukas C, Deftos L: The antiproliferative effect of calcitriol on human peripheral blood mononuclear cells. *J Clin Endocrinol Metab* 63, 394–400 (1986)
28. D'Ambrosio D, Cippitelli M, Cocciolo MG, Mazzeo D, Di Lucia P, Lang R, Sinigaglia F, Panina-Bordignon P: Inhibition of IL-12 production by 1, 25-dihydroxyvitamin D₃. Involvement of NF-kappaB downregulation in transcriptional repression of the p40 gene. *J Clin Invest* 101, 252 (1998)
29. Müller K, Gram J, Bollerslev J, Diamant M, Barington T, Hansen M, Bendtzen K: Down-regulation of monocyte functions by treatment of healthy adults with 1 α , 25 dihydroxyvitamin D₃. *Int J Immunopharmacol* 13, 525–530 (1991)
30. Itty S, Day S, Lyles KW, Stinnett SS, Vajzovic LM, Mruthyunjaya P: Vitamin D deficiency in neovascular versus nonneovascular age-related macular degeneration. *Retina* 34, 1779–1786 (2014)
31. Parekh N, Chappell RJ, Millen AE, Albert DM, Mares JA: Association between vitamin D and age-related macular degeneration in the Third National Health and Nutrition Examination Survey, 1988 through 1994. *Arch Ophthalmol* 125, 661–669 (2007)
32. Kim EC, Han K, Jee D: Inverse relationship between high blood 25-hydroxyvitamin D and late stage of age-related macular degeneration in a representative Korean population association of 25-hydroxyvitamin D with AMD. *Invest Ophthalmol Vis Sci* 55, 4823–4831 (2014)
33. Tan JS, Wang JJ, Flood V, Mitchell P: Dietary fatty acids and the 10-year incidence of age-related macular degeneration: The Blue Mountains Eye Study. *Arch Ophthalmol* 127, 656–665 (2009)
34. Kertes PJ: Cigarette smoking, fish consumption, omega-3 fatty acid intake, and associations with age-related macular degeneration: The US Twin Study of Age-Related Macular Degeneration. *Evid Based Ophthalmol* 8, 50–52 (2007)
35. Singh A, Falk MK, Subhi Y, Sørensen TL: The association between plasma 25-hydroxyvitamin D and subgroups in age-related macular degeneration: A cross-sectional study. *PLoS One* 8, e70948 (2013)
36. Millen AE, Volland R, Sondel SA, Parekh N, Horst RL, Wallace RB, Hageman GS, Chappell R, Blodi BA, Klein ML, Gehrs KM, Sarto GE, Mares JA, CAREDS Study Group: Vitamin D status and early age-related macular degeneration in postmenopausal women. *Arch Ophthalmol* 129, 481–489 (2011)
37. Simonelli F, Zarrilli F, Mazzeo S, Verde V, Romano N, Savoia M, Testa F, Vitale DF, Rinaldi M, Sacchetti L: Serum oxidative and antioxidant parameters in a group of Italian patients with age-related maculopathy. *Clin Chim Acta* 320, 111–115 (2002)

38. Ishihara N, Yuzawa M, Tamakoshi A: Antioxidants and angiogenic factor associated with age-related macular degeneration (exudative type). *Nippon Ganka Gakkai Zasshi* 101, 248–251 (1997)
39. Klein ML, Mauldin WM, Stoumbos VD: Heredity and age-related macular degeneration: Observations in monozygotic twins. *Arch Ophthalmol* 112, 932–937 (1994)
40. Smith W, Mitchell P: Family history and age-related maculopathy: The Blue Mountains Eye Study. *Aust N Z J Ophthalmol* 26, 203–206 (1998)
41. Christen WG, Glynn RJ, Chew EY, Buring JE: Vitamin E and age-related macular degeneration in a randomized trial of women. *Ophthalmology* 117, 1163–1168 (2010)
42. Timms PM, Mannan N, Hitman GA, Noonan K, Mills PG, Syndercombe-Court D, Aganna E, Price CP, Boucher BJ: Circulating MMP9, vitamin D and variation in the TIMP-1 response with VDR genotype: Mechanisms for inflammatory damage in chronic disorders? *Quart J Med* 95, 787–796 (2002)
43. Seddon JM, George S, Rosner B, Rifai N: Progression of age-related macular degeneration: Prospective assessment of C-reactive protein, interleukin 6, and other cardiovascular biomarkers. *Arch Ophthalmol* 123, 774–782 (2005)
44. Seddon JM, Gensler G, Milton RC, Klein ML, Rifai N: Association between C-reactive protein and age-related macular degeneration. *JAMA* 291, 704–710 (2004)
45. Klein R, Klein BE, Marino EK, Kuller LH, Furberg C, Burke GL, Hubbard LD: Early age-related maculopathy in the cardiovascular health study. *Ophthalmology* 110, 25–33 (2003)
46. McKay GJ, Young IS, McGinty A, Bentham GC, Chakravarthy U, Rahu M, Seland J, Soubrane G, Tomazzoli L, Topouzis F, Vioque J, de Jong PT, Fletcher AE: Associations between serum vitamin D and genetic variants in vitamin D pathways and age-related macular degeneration in the European Eye Study. *Ophthalmology* 124, 90–96 (2017)