Omega-3 fatty acids supplementation protects the retina from age-associated degeneration in aged C57BL/6J mice

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ABSTRACT

Objective To evaluate the therapeutic effects of omega-3 (ω3) fatty acids in the retina of aged mice when the blood arachidonic acid (AA)/eicosapentaenoic acid (EPA) ratio is maintained between 1.0 and 1.5.

Methods and analysis Aged (24-month-old) wild-type C57BL/6J mice were allocated to two groups: ω3 treated and untreated. Treatment with ω3 was by daily gavage administration of EPA and docosahexaenoic acid for 60 days. Gas chromatography was used to identify and quantify fatty acids in the blood and retina. To count lipofuscin granules and measure the photoreceptor layer, eyecups were examined histologically using transmission electron microscopy and light microscopy. We also analysed eyecups using mass spectrometry-based proteomics.

Results AA levels were lower, and EPA levels were higher, in the blood and retinas of the ω3-treated group than in the untreated group, resulting in a lower AA/EPA ratio. The ω3-treated group also showed significantly fewer lipofuscin granules and a thicker outer nuclear layer than the untreated group. Proteomic analysis revealed significantly greater expression of myelin basic protein, myelin regulatory factor-like protein, myelin proteolipid protein and glial fibrillar acidic protein in the ω3-treated group than in the untreated group. Three different pathways were significantly affected by ω3 treatment: fatty acid elongation, biosynthesis of unsaturated fatty acids and metabolic pathways.

Conclusion Two months of ω3 supplementation (when the blood AA/EPA~1.0–1.5) in aged mice reduced lipofuscin granule formation in the retina and protected the photoreceptor layer, suggesting that ω3 supplementation slows normal age-related retinal degeneration.

INTRODUCTION

Visual function declines with age, as observed in the majority of apparently ‘disease-free’ individuals who experience some degree of age-associated changes in vision.1 These changes include reductions in visual acuity,2 3 colour perception,4 5 dark adaptation and contrast sensitivity.6 7 The prevention of age-related loss of vision or enhancement of visual acuity in elderly people would significantly improve their quality of life, in addition to improving their function in activities of daily living.

Age-associated changes in visual acuity can be explained by two different theories: the ‘biological clock theory’ and the ‘tear
Retinas in young and aged individuals and in patients with AMD showed multiple changes in the morphology and distribution of astrocytes, which are glial cells that communicate with neurons and the surrounding connective tissue. In addition to being involved in numerous supportive processes, astrocytes have a high antioxidant content, as well as the potential to protect neurons from free radicals.

Aged retinas demonstrated fewer astrocytes and spaces with no glial fibrillar acidic protein (GFAP) reactivity in the nerve fibre layer (due to the disappearance of the processes from the astrocytes located in this layer). Furthermore, astrocytes from aged retinas have more cytoplasmic organelles and glial filaments than astrocytes from younger retinas. The number of chambers in the astroglial plexus reduces increasingly with age, as demonstrated by a comparison of persons aged 60–79 and persons aged over 80. Large numbers of reactive astrocytes were reported in all patients with AMD, which revealed that these astrocytes phagocytosed the residue from ganglion cells that died via apoptosis/necrosis. More lipofuscin deposits were observed in the aged retinas than in the younger retinas; however, pathological retinas from patients with AMD had even more deposits of lipofuscin granules.

Similarities in the physiology and cell biology of ageing in humans and rodents have been observed; for example, 2.5-year-old mice show decreases in the maximum voltage obtained from electroretinograms, while opsins gene transcription, cone density and increment thresholds are unchanged even in extremely old mice, just as in humans. The number of rods and the length of their outer segment decreases with age, as does the total level of retinal opsin. The scotopic visual acuity, contrast sensitivity and a and b waves from rod electroretinogram recordings are also reduced in aged mice.

Samuel et al compared all major retinal neuronal types in young adult (3–5 months old) and aged (24–28 months old) C57BL/6J mice. The study revealed that the total retinal area expanded with age, whereas the retinal ganglion cell dendritic arbours, and the size and complexity of the retinal ganglion cell axonal arbours, decreased with age. This finding suggests that the transmission of visual signals to the brain is compromised. In addition, age-related structural changes were observed in the amacrine and bipolar cells. Taking into account the morphological and molecular changes that occur in aged C57BL/6J mice, and the similarities they have with human aged eyes, we considered these mice an appropriate animal model for the purpose of the present study.

The protective effects of omega-3 (ω3) polyunsaturated fatty acids (PUFAs) against retinal damage have been demonstrated in several disease states, including pathologies associated with inflammation, ischaemia, light and oxygen. Furthermore, the treatment of retinal degeneration with ω3 PUFAs has shown promise in animal models, through the reduction of A2E accumulation and preservation of retinal functions.
ω3 PUFAs are known to reduce inflammation, primarily through resolvins that are derived from eicosapentaenoic acid (EPA, C20:5 n−3) and docosahexaenoic acid (DHA, C22:6 n−3), and through anti-inflammatory mediators. DHA activates a number of nuclear hormone receptors that modulate proinflammatory genes, including the peroxisome proliferator-activated receptor alpha (PPARα). EPA-rich diet reduces choroidal neovascularisation in C57BL/6J mice, through a suppression of intercellular adhesion molecule, monocyte chemotactic protein 1, vascular endothelial growth factor and interleukin 6. Proinflammatory eicosanoids, including prostaglandins and leukotrienes that are involved in leucocyte chemotaxis and inflammatory cytokine production, are generated from the omega-6 (ω6) PUFA arachidonic acid (AA, C20:4 n−6). ω3 PUFAs also have the capacity to affect production and activation of angiogenic growth factors, AA-based vasoregulatory eicosanoids and metalloproteinases. Finding the right balance between the proinflammatory and anti-inflammatory mediators produced by ω3 and ω6 PUFAs plays a key role in the resolution of inflammatory responses and therefore possibly in age-associated vision loss.

The therapeutic potential of ω3 PUFAs in eye pathologies has also been demonstrated by our research team in several animal models. The fish oil formulation that we use in our studies is rich in EPA. Previous reports demonstrated that EPA competes with AA and significantly inhibits AA oxidation in vitro, thus reducing activity in the inflammatory pathways involving prostaglandins D, E and F. The activity of ω3 was tested in a rat model of anterior ischaemic optic neuropathy. Our findings showed that the positive effect of ω3 on the retina is mediated by several different actions, including preventing retinal ganglion cell apoptosis, decreasing inflammatory cell infiltration and regulating macrophage polarisation, which decreases cytokine-induced injury of the optic nerve.

Similarly, Prokopiou et al reported the effects of ω3 PUFAs in the CCL2−/− mouse model of retinal degeneration. A protective response, perhaps with a regenerative potential, was observed in ω3-treated animals, with a pronounced increase in the size of the photoreceptor outer nuclear layer (ONL) and a reduction in interleukin 18. Kalogerou et al demonstrated the effectiveness of ω3 PUFAs in the DBA/2J glaucoma model; retinal ganglion cells were protected against cell death and a reduced inflammatory response was observed.

Recently, the therapeutic potential of ω3 was also reported by Prokopiou et al in the ABCA4−/− mouse model of Stargardt disease. In particular, significantly less A2E and fewer lipofuscin granules were observed in ω3-treated animals than in untreated animals, and the photoreceptor ONL was preserved. Proteomic analysis revealed lower levels of complement component 3 in the treatment group, indicative of a decreased complement-induced inflammatory response. Importantly, in all these studies, the levels of particular PUFAs (including EPA and AA) were examined during treatment, allowing the treatment dosage to be adjusted in order to achieve and maintain a blood AA/EPA level of ~1.0–1.5. We emphasise the importance of detecting these fatty acids (FAs) in our previous reports. Lower levels of EPA and higher levels of AA, in either circulating blood or the retina, are associated with some retinopathies, including AMD. Monitoring such systemic biomarkers during experimental animal investigations or in human trials could provide a good indication of disease progression and treatment success.

We hypothesise that ω3 PUFAs would be beneficial for disease-free individuals who experience some degree of age-associated changes in vision. Therefore, in the present study, we investigated how the administration of ω3 PUFAs (primarily EPA and some DHA), to 2-year-old C57BL/6J wild-type mice, affects RPE cells and photoreceptors.

**MATERIALS AND METHODS**

Animals

Young (3-month-old) and aged (24-month-old) C57BL/6J wild-type mice (inbred at the animal facility at the Cyprus Institute of Neurology and Genetics) of equal amount of both sexes were used in this study; the mice were kept under standard housing conditions on a 12 hours/dark/light cycle with food and water provided ad libitum. Standard diet was provided (Mucedola, Milano, Italy). Mean body weight of animals prior to treatment was 28.5±2.5 g. The general appearance of the mice was monitored daily, and body weight measurements were recorded weekly.

**Study design**

The animals were allocated to three different groups (n=15/group): young untreated; aged untreated and aged treated with ω3 PUFAs. The aged mice were randomly allocated to the latter two groups. The animals in the treatment group received the supplement daily by gavage for 60 days. The total oral dosing volumes of the gavage administration did not exceed 10 mL/kg (~0.3 mL/mouse). The ω3 treatment was a fish oil formulation that contained 571 mg/mL EPA and 114 mg/mL DHA (Eyetas, TAS Ophthalmos L.C., Nicosia, Cyprus); therefore, each treatment administration had 172 mg EPA (Eyetas, TAS Ophthalmos L.C., Nicosia, Cyprus); therefore, each treatment administration had 172 mg EPA and 34 mg DHA per 0.3 mL. The supplement dosage was adjusted in order to achieve the blood AA/EPA within the range of 1.0–1.5. At the end of the study, the mice were euthanised by cervical dislocation.

**FA analysis**

To determine the AA/EPA ratio before, during and on completion of treatment, we collected blood samples (n=5 mice) from the tail of each mouse in each group on day zero (D0), day 30 (D30) and day 60 (D60). Blood samples from young mice were only collected on D60 for comparison purposes. Additionally, on study
completion (D60), both retinas from three animals per group were collected and analysed for the same purpose. Samples were collected on Whatman (GE Healthcare, Life Sciences, Buckinghamshire, UK) filter paper and stored at −20°C until ready for analysis. Samples were processed as previously described, briefly, they were separated into organic and aqueous phases, where the organic phase was collected, redissolved in n-hexane and analysed using a gas chromatography flame ionisation detector.

An Agilent GC-6890 system (Hewlett Packard Agilent Technologies, California, USA) was equipped with a flame ionisation detector. The conditions, temperatures of the injection port and detector were set as previously described. The injection volume was 1 mL in the splitless injection mode. A capillary column (DB-23 fused-silica capillary, 30 m×0.25 mm internal diameter, 0.25 µm film thickness; Supelco, Bellefonte, Pennsylvania, USA) was used.

**Transmission electron microscopy (TEM)**

Electron microscopy was used to quantify lipofuscin granules in the RPE cells. For this analysis, the cornea and lens from five animals per group were removed, and the dissected eyecups were fixed, dehydrated and embedded in an epon and araldite resin mixture (Agar Scientific, Essex, UK), as previously described. Ultrathin (80 nm) sections were prepared using a Reichert Jung ultramicrotome (Leica, Wetzlar, Germany). Sections with a silver–gold interference colour were mounted on 200-mesh copper grids (Agar Scientific) and contrasted with uranyl acetate and lead citrate. Images were obtained on a JEM 1010 TEM (JEOL, Tokyo, Japan) equipped with a Mega View III digital camera (Olympus, Hamburg, Germany).

As the appearance of lipofuscin is somewhat variable, we used a previous morphological definition to identify lipofuscin granules. Melanosomes that were fused with lipofuscin were considered melanolipofuscin granules and were included in the combined category of ‘lipofuscin and melanolipofuscin’. We used ImageJ software (http://imagej.nih.gov/ij, National Institutes of Health, Bethesda, Maryland, USA) for lipofuscin and melanolipofuscin quantification. For each image, we determined the total area of RPE cytoplasm as previously reported. The extent of the lipofuscin/melanolipofuscin area is expressed as the number of square micrometres occupied by lipofuscin per 1000 µm² of RPE cytoplasm.

**Light microscopy**

To assess how the treatment affected retinal photoreceptors (ie, ONL thickness), we examined the same eyes that were used for the quantification of lipofuscin granules, under a Leica DMi8 light microscope (Leica). ONL measurements were made at different locations around the entire retinal section (centre, middle and periphery). ImageJ was used to calculate the ONL thickness at each location, and the mean ONL thickness around the entire retina was compared among the different groups.

**Mass spectrometry-based proteomics**

To further assess the effect of treatment, proteomic analysis was conducted on eyecup extracts. The cornea and lens from five eyes per group (one eye/mouse) were removed, and the dissected eyecups were immediately snap frozen and stored at −80°C until needed. The frozen eyecups were analysed as previously described. Briefly, the samples were lysed, and once centrifuged, the supernatant was collected and the proteins were precipitated in acetone overnight and subsequently resuspended in urea buffer. The protein concentration was determined using the bicinchoninic acid protein assay, and the samples were further processed using a modified filter-aided sample preparation protocol. Peptides were analysed on a Waters Synapt G2Si high-definition mass spectrometer (Waters) operated in ion-mobility mode using an ultradefinition mass spectrometry approach.

Each sample was analysed in triplicate. The raw mass spectrometry data were analysed using Progenesis QI for proteomics V.3.0 (Waters) and proteins identified using the UniProt mouse reference proteome database (http://www.uniprot.org/, January 2018 version, containing 16 889 reviewed proteins). The PANTHER Classification System V.12.0 (http://pantherdb.org/) was used to determine the functional classification of the identified proteins. Proteins showing an increase (>1.5-fold) or a decrease (<0.66-fold) were further analysed using DAVID Bioinformatics Resources V.6.8 (https://david.ncifcrf.gov/).

**Statistical analysis**

All statistical analyses were performed using GraphPad Prism statistical software (GraphPad Software, USA, V.5). Data are expressed as the means±SEMs. A two-tailed Student’s t-test or one-way analysis of variance with Tukey’s post hoc test was used to evaluate the differences between groups. Statistical significance was set at p<0.05.

**Patient and public involvement**

Patients were not involved in this study.

**RESULTS**

**General observations**

No abnormalities were noted in the general appearance of the mice, and no significant changes in body weight were found over time in any of the groups. No sex differences were observed at any stage of the analysis.

**ω3 PUFAs increase blood EPA and decrease blood AA**

Blood was collected from mice in different groups (n=5) on D0 (prior to the initiation of treatment), on D30, and at the end of the study on D60, to determine the percentage of total ω3 PUFAs (EPA and DHA) and ω6 PUFAs (AA and dihomo-γ-linolenic acid, DGLA; figure 1). Blood samples from young mice were only collected on D60, since no differences were observed at these ages (1, 2 and...
3 months old) in previous pilot studies we performed. The most important finding was that EPA level as a percentage of total blood FAs was about six times higher in the \(\omega_3\)-treated group (6.55±0.15, p<0.0001) than in the aged untreated group (1.13±0.15) on D30, a difference that was maintained to D60. The percentage of AA was lower on D30 in the aged treated group (7.30±0.07, p<0.001) than in the aged untreated group (8.96±0.05), and this difference was even greater on D60 (5.44±0.05, p<0.001 vs 8.61±0.20). Therefore, on D30, the blood AA/EPA ratio was more than seven times lower in the \(\omega_3\)-treated animals (1.1±0.02, p<0.0001) than in the aged untreated (7.98±0.26) animals. At the end of the study on D60, the AA/EPA ratio in the \(\omega_3\)-treated mice remained unchanged from that on D30 and was still lower than the AA/EPA ratio in the young untreated mice (4.56±0.05, p<0.001). In the \(\omega_3\)-treated mice, a small reduction was observed in the percentage of DHA and DGLA on D30 and D60. Therefore, \(\omega_3\)-treated mice demonstrated lower AA and higher EPA levels in the blood, favourable of the desired AA/EPA ratio (1.0–1.5).

\(\omega_3\) PUFAs increase retinal EPA levels and decrease retinal AA levels

Retinas from each group (n=3) were collected at the end of the study on D60, and the constituent FAs (EPA, DHA, DGLA and AA) were quantified as a percentage of total FA content using gas chromatography (figure 2). We observed similar trends in the retinas as in the blood. Specifically, the percentage of retinal EPA in the aged \(\omega_3\)-treated group (1.75±0.04, p<0.01) was significantly higher than that in the aged untreated group (1.23±0.00). Retinal EPA was undetectable in the young untreated animals. Conversely, the AA level in the treatment group (5.58±0.01, p<0.05) was significantly lower than that in the aged untreated group (6.56±0.28), whereas the AA

![Figure 1](http://bmjophth.bmj.com/)

**Figure 1** (A)–(E) Fatty acid analysis. Percentages of total (A) arachidonic acid (AA), (B) eicosapentaenoic acid (EPA), (C) docosahexaenoic acid (DHA) and (D) dihomo-\(\gamma\)-linolenic acid (DGLA) in blood samples collected on D0, D30 and D60. (E) The AA/EPA ratio in each group at D0, D30 and D60. The data represent the means ± SEMs (n = 5 mice per group). ANOVA, analysis of variance.

![Figure 2](http://bmjophth.bmj.com/)

**Figure 2** Fatty acid analysis of retina samples collected on D60. The percentage of (A) arachidonic acid (AA), dihomo-\(\gamma\)-linolenic acid (DGLA), eicosapentaenoic acid (EPA) (ND: not detected) and (B) docosahexaenoic acid (DHA) is expressed as the mean ± SEM (n=3 per group). ANOVA, analysis of variance.
level in the young untreated animals (6.93±0.04) was similar to that in the aged untreated animals, but significantly higher than that in the treated animals (p<0.01). DHA was the most abundant FA in the retina, with a level much higher than that of any other FA, as previously reported.56 A significantly lower level of DHA was observed in the aged ω3-treated group (24.25±0.64) compared with the aged untreated (30.86±0.02, p<0.001) and young untreated (29.70±0.15, p<0.001) groups (figure 2B).

ω3 PUFAs reduce the number of lipofuscin granules

The RPE layer was examined using electron microscopy to assess the ultrastructural differences between the young untreated, aged untreated and aged ω3-treated animals (figure 3). Melanosomes (figure 3A, blue arrows) were abundant in all groups and were uniformly electron dense, not fused with other organelles, and spindle-shaped, ovoid or round. An accumulation of irregularly shaped, electron-dense material in the cytoplasm of RPE cells was mostly noted in the aged untreated animals (figure 3A, red arrows). This type of organelle might be similar to typical lipofuscin granules. Melanolipofuscin granules (figure 3A, yellow arrows) were more frequently observed and were therefore included in the quantification analysis (figure 3B). The total area (µm²) occupied by lipofuscin and melanolipofuscin per 1000 µm² section of RPE cytoplasm was significantly lower in aged mice treated with ω3 than in aged untreated mice. The data represent the means ± SEMs (n=5 per group). ANOVA, analysis of variance.

ω3 PUFAs and the photoreceptor layer

The ONL thickness represents the long-term health of photoreceptor nuclei; therefore, we captured images (figure 4A) and quantified the ONL thickness in the three groups (figure 4B). In the young untreated mice, the ONL thickness was 201.0±35.0 µm, whereas in the aged untreated mice, it was significantly thinner: 114.5±7.7 µm (p<0.05). The ONL in ω3-treated aged
mice was significantly greater in thickness (188.0±26.0 µm, p<0.05) than in the aged untreated mice, and did not significantly differ in thickness to the ONL of young untreated mice.

ω3 PUFAs and proteomic analysis
Proteomic analysis enabled us to identify and quantify 2786 proteins in the eyecups. The levels of approximately 49 of these proteins differed significantly (p<0.05) among the three groups. According to PANTHER functional classification, the total number of function hits was 33, as shown in figure 5, with the majority of these proteins exhibiting catalytic activity (including isomerase, ligase, acetyltransferase and decarboxylase activity). A smaller proportion of the proteins that differed significantly between groups has binding activity or is involved in processes such as transport (ie, membrane organisation, cation and lipid transportation).

Notably, among the proteins that differed significantly within the groups, there were some that may play a key role in the progression of age-associated degeneration. In particular, myelin basic protein (MBP), myelin proteolipid protein (MPP) and myelin regulatory factor-like protein (MRFLP) were analysed separately. Figure 6 shows the fold change defined as the level of each protein detected in each group divided by the measured level of that protein in the young untreated group. Overall, significantly greater fold changes in MBP, MPP and MRFLP levels were observed in the aged ω3-treated group (2.17±0.57, 1.89±0.32 and 1.49±0.22, respectively; p<0.05) than in the aged untreated group (0.57±0.06, 0.71±0.04 and 0.47±0.06, respectively). The fold changes in the levels of MBP and MPP in the ω3-treated group were also significantly higher than those in the young untreated group (p<0.05).

In addition to differences between the groups in myelin-related proteins, we also determined the fold change of GFAP, an astrocyte marker, in the three groups (figure 7). There was a two-fold reduction in the level of GFAP in the aged untreated mice (0.55±0.05, p<0.05) compared with that in the young untreated mice (1.00±0.11). Interestingly, there was a twofold increase in the ω3-treated group (1.18±0.10, p<0.01) compared with the aged untreated group. The fold change in GFAP level in the ω3-treated group was comparable to that in the young untreated group.

To identify specific pathways that were significantly altered following treatment with ω3, further analysis of the proteins that were expressed differently between the groups was performed. Specifically, three pathways demonstrated a difference between the treatment and control groups: pathways involved in FA elongation (acyl-CoA thioesterase 1, Acot2, Acot3, Acot5, p=9.5E−06), biosynthesis of unsaturated FAs (Acot1, Acot2, Acot3, ...
of ω3 PUFAs in eye diseases. The positive effect of ω3 treatment was investigated in animal models of retinal and optic nerve pathologies and in clinical studies of AMD and dry eyes. Although the results were not conclusive, we investigated the effect of ω3 PUFAs in age-associated retinal degeneration in healthy mice. We hypothesised that the most beneficial effect would be achieved when using a dosage that maintains an AA/EPA ratio between 1.0 and 1.5 and an appropriate formulation of ω3 PUFAs (EPA:DHA=5:1).

Our findings show that treatment with ω3 PUFAs, when AA/EPA is maintained between 1.0 and 1.5, reduced the accumulation of lipofuscin granules and increased the level of myelin-related proteins and the number of astrocytes (as indicated by an increase in GFAP). Demonstrating a protective effect against photoreceptor damage would require a considerably longer experiment than our resources permitted; even so, compared with no treatment in aged mice, 2 months of treatment with ω3 PUFAs in aged mice resulted in a notable preservation of the photoreceptor layer. If this effect can be confirmed in the long term, supplementation with ω3 PUFAs could be considered a preventive measure for age-associated vision loss. In particular, EPA appears to have a critical role in protecting the retina, since its percentage was significantly increased in the retina, whereas the percentage of DHA was not.

Serhan et al have extensively studied the mechanism of action of ω3 PUFAs. The processes involved are complex, and primarily implicate the generation of anti-inflammatory mediators—for example, resolvins, which are generated from the ω3 PUFAs EPA and DHA via cyclooxygenase/lipoxygenase pathways. Resolvins promote the control of inflammation in neural tissues and reduce the levels of proinflammatory eicosanoids generated from AA. Although the level of resolvins in blood and retina would have been beneficial for the purpose of this study, several attempts were made to detect them without any success, perhaps due to their short half-life.

During the ω3 supplementation treatment and at completion of the study, we observed that the AA/EPA ratio was maintained at an average of 1.0–1.5, which was within our target range. The reduction in the ratio was due to the increase in EPA (from the fish oil formulation), and due to the reduction in AA observed in both the blood and the retinal tissue. Our present and previous studies also show that EPA successfully incorporates into the retinal tissue. DHA is known to be expressed abundantly in photoreceptors; this was confirmed by the high levels of DHA compared with the levels of the other FAs in the retina.

Additional mechanisms unrelated to inflammation might be involved in the mechanism of action of ω3 PUFAs. Our retinal proteomic analysis showed that three different pathways were significantly affected by treatment: FA elongation, biosynthesis of unsaturated FAs and metabolic pathways. These alterations were not surprising, since supplementation with dietary ω3 PUFAs is known to increase the elongation of particular FAs.

In addition, the present data suggest that treatment with ω3 PUFAs may confer regenerative potential through the stimulation of myelin-related proteins, such as MBP, MRFLP and MPP. Our findings (increased MBP, MRFLP and MPP protein levels) agree with previous reports of myelogenesis being affected by environmental interferences including nutrition, with ω3-rich diets inducing an increased myelin synthesis. Further evidence demonstrated that EPA stimulates the expression of myelin proteins in the rat brain. Specifically, a single dose of EPA or DHA increased the MBP, MPP and myelin oligodendrocyte protein mRNA levels in nearly all brain regions in 2-day-old rats; this effect was more pronounced in EPA-treated rats. In contrast, AA was shown to decrease the phosphorylation of MBP in cultured rat oligodendrocytes via either a diffusible paracrine signal or a signal transduction mechanism, thus reducing MBP’s activity. Furthermore, Salvati et al reported that EPA but not AA stimulates the expression of proteolipid protein via cAMP-mediated pathways in a time-dependent and concentration-dependent manner in C6 glial cells. Our results confirm that ω3 PUFAs stimulate proteins essential for myelination in the retina, and suggest that they might furthermore prevent retinal degeneration.

Our proteomic analysis revealed that GFAP protein expression was elevated after ω3 treatment, indicating that the treatment increased the number of astrocytes. As
confirmed in our animal model, astrocytes show a degenerative profile during normal ageing, implying a reduction in antioxidant and protective benefits to the host. Thus, by increasing astrocyte abundance via ω3 PUFAs administration, neuroprotection of the eye may be established. A similar effect was recently demonstrated by treating anterior horn cells in the spinal cord of male albino rats with ω3 PUFAs plus aspartame; rats treated with aspartame alone showed a higher rate of cell death and fewer GFAP-positive astrocytes. In addition, Zgórsyńska et al showed that the incubation of rat primary astrocytes with DHA and EPA improves the antioxidant defense in astrocytes via an Nrf2-dependent mechanism, determined by the ratio of DHA to EPA incorporated into membrane phospholipids. Together, these findings suggest that supplementation with ω3 PUFAs increases the number of astrocytes in the retina and may protect neurons against age-associated retinal degeneration.

Interest in how dietary ω3 PUFAs influence several physiological (eg, ageing) or pathological (eg, cardiovascular disease) conditions has grown considerably over the past decade. Numerous clinical studies have focused mainly on the anti-inflammatory properties of ω3 PUFAs as well as on their neuroprotective and regenerative effects.

In particular, one review analysed human brain MRI studies and experimental studies and confirmed that ω3 PUFAs can protect the brain during ageing by preventing peroxidation, inflammation and excitotoxicity; modulating neurotransmission; and increasing neurogenesis, thereby improving the brain’s ‘coaching’ mechanisms during ageing.

We have established here that ω3 PUFAs have protective and therapeutic potential against age-related decline in visual function in 2-year-old-aged C57BL/6J mice. This was demonstrated by trained mice showing a lower number of lipofuscin granules, preservation of the retinal structure (ie, the photoreceptor layer), greater expression of myelin-related proteins and a higher level of GFAP expression (indicating an increased number of astrocytes). Although future work would be valuable, our results suggest that ω3 PUFAs may be considered a protective dietary supplement against the vision loss caused by normal ageing.

Contributors EP and PK designed and performed the experiments. MK, LP, CG and KS contributed to some of the experimental procedures. TG supplied the reagents/materials/analysis tools. TG and KK provided guidance and support. EP analysed the data and wrote the manuscript. All contributing authors have read and approved the final version of the manuscript.

Funding This research was supported by Ophthalmos Research and Educational Institute (Nicosia, Cyprus) and was not supported by any specific grant from funding agencies in the public, commercial, or not-for-profit sectors.

Competing interests TG has a patent on the use of omega-3 fatty acids in eye diseases.

Patient consent for publication Not required.

Ethics approval The animal care and experimental procedures were performed in accordance with the EU Directive 2010/63/EU for animal experiments and approved by the Cyprus Government’s Chief Veterinary Officer.

Provenance and peer review Not commissioned; externally peer reviewed.

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