Rescue and repair during photoreceptor cell renewal mediated by docosahexaenoic acid-derived neuroprotectin D1

The Harvard community has made this article openly available. Please share how this access benefits you. Your story matters

| Citation          | Bazan, Nicolas G., Jorgelina M. Calandria, and Charles N. Serhan. 2010. “Rescue and Repair during Photoreceptor Cell Renewal Mediated by Docosahexaenoic Acid-Derived Neuroprotectin D1.” Journal of Lipid Research 51 (8): 2018–31. https://doi.org/10.1194/jlr.r001131. |
|-------------------|--------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| Citable link      | http://nrs.harvard.edu/urn-3:HUL.InstRepos:41483531                                                                                                                                               |
| Terms of Use      | This article was downloaded from Harvard University’s DASH repository, and is made available under the terms and conditions applicable to Other Posted Material, as set forth at http://nrs.harvard.edu/urn-3:HUL.InstRepos:dash.current.terms-of-use#LAA |
Rescue and repair during photoreceptor cell renewal mediated by docosahexaenoic acid-derived neuroprotectin D1

Nicolas G. Bazan, Jorgelina M. Calandria, and Charles N. Serhan

Neuroscience Center of Excellence and Department of Ophthalmology, School of Medicine, Louisiana State University Health Sciences Center, New Orleans, LA 70112; and Center for Experimental Therapeutics and Reperfusion Injury, Department of Anesthesiology, Perioperative and Pain Medicine, Brigham and Women’s Hospital and Harvard Medical School, Boston, MA 02115

Abstract Retinal degenerative diseases result in retinal pigment epithelial (RPE) and photoreceptor cell loss. These cells are continuously exposed to the environment (light) and to potentially pro-oxidative conditions, as the retina’s oxygen consumption is very high. There is also a high flux and to potentially pro-oxidative conditions, as the retina’s cells are continuously exposed to the environment (light) and RPE cells. Photoreceptor outer segment shedding and phagocytosis intermittently renews photoreceptor membranes. DHA is converted through 15-lipoxygenase-1 into neuroprotectin D1 (NPD1), a potent mediator that evokes counteracting cell-protective, anti-inflammatory, pro-survival repair signaling, including the induction of anti-apoptotic proteins and inhibition of pro-apoptotic proteins. Thus, NPD1 triggers activation of signaling pathway/s that modulate/s pro-apoptotic signals, promoting cell survival. This review provides an overview of DHA in photoreceptors and describes the ability of RPE cells to synthesize NPD1 from DHA. It also describes the role of neurotrophins as agonists of NPD1 synthesis and how photoreceptor phagocytosis induces refractoriness to oxidative stress in RPE cells, with concomitant NPD1 synthesis.—Bazan, N. G., J. M. Calandria, and C. N. Serhan. Rescue and repair during photoreceptor cell renewal mediated by docosahexaenoic acid-derived neuroprotectin D1. J. Lipid Res. 2010. 51: 2018–2031.

Supplementary key words age-related macular degeneration • retinal pigment epithelial cells • oxidative stress • neurotrophins • 15-lipoxygenase-1

Retinal degenerative diseases are a complex group of conditions with different etiologies that result in a common outcome: photoreceptor apoptotic cell death (1–5). Accordingly, there are differences in how these conditions evolve. For instance, in retinitis pigmentosa (RP), rod photoreceptor death initially occurs in the periphery, whereas in age-related macular degeneration (AMD), death is initiated in the macular zone and spreads in later phases throughout the retina (2, 5). RP is a collection of inherited blinding diseases caused by the mutation of a wide variety of genes resulting in more than 150 abnormalities of photoreceptor-specific proteins, including mutations of rhodopsin, peripherin, the β-subunit of cGMP phosphodiesterase, and retinal outer-segment membrane protein 1 (6–8).

Conversely, the etiology of AMD, which is the leading cause of blindness over the age of 65, is not as clear as that of RP. AMD is also a heterogeneous group of disorders, but the causes are proposed to be multifactorial, and the main known risk factors are both genetic and environmental (2, 5). There are two forms of AMD: the dry and the wet form. In the dry form, photoreceptors degenerate slowly...
and progressively, producing a thinning in the retinal layers and leaving deposits known as drusen. In the wet form, the less common of the two AMDs, the predominant feature is invasive choroidal neovascularization, which leads to severe vision loss (9).

AMD also affects the choriocapillaris and Bruch’s membrane, which separates the retinal pigment epithelium (RPE) from the blood vessels (10). The results from the Age-Related Eye Disease Study (NEI) show that the intake of high amounts of antioxidants and zinc can reduce the risk of developing advanced AMD by about 25%, implying an indirect role for oxidative stress in the pathogenesis. More specifically, the presence of oxidative damage markers in postmortem retinas of patients with geographic atrophy shows, at least in the dry form, oxidative stress is involved in the pathogenic mechanisms of AMD (11). In this manner, oxidative stress is enhanced and exaggerated and mitochondrial function is compromised (8–13), which leads to apoptotic cell death. Tunel studies performed on postmortem human retinas of patients presenting geographic atrophy and exudative forms of the disease show that apoptosis is the main mechanism of degeneration, not only for photoreceptors, but also for the RPE and inner retinal layers (12).

Initiation and progression of AMD involves the unsuccessful resolution of the inflammatory response. Single nucleotide polymorphisms occurring in the gene encoding factor H (CFH/HF1) (14–17) were proposed to be a major risk factor for AMD. Factor H is an inhibitor of the alternative pathway of complement system activation that, in this manner, oxidative stress is enhanced and exaggerated and mitochondrial function is compromised (8–13), which leads to apoptotic cell death. Tunel studies performed on postmortem human retinas of patients presenting geographic atrophy and exudative forms of the disease show that apoptosis is the main mechanism of degeneration, not only for photoreceptors, but also for the RPE and inner retinal layers (12).

Initiation and progression of AMD involves the unsuccessful resolution of the inflammatory response. Single nucleotide polymorphisms occurring in the gene encoding factor H (CFH/HF1) (14–17) were proposed to be a major risk factor for AMD. Factor H is an inhibitor of the alternative pathway of complement system activation that, as a result, has the ability to limit cell injury and inflammation (18, 19). Conversely, studies that focused on other regulatory proteins of the complement pathway, such as factor B and complement component 2 (C2) (18, 20), exhibit protective effects and reduce the risk of AMD to some extent. For example, the E318D variant of C2 (H10) as well as a variant in intron 10 of C2 and the R32Q variant of factor B (H7) confer a reduced risk of AMD.

Therefore, the identification of early pro-survival, anti-inflammatory signaling critical for the maintenance of photoreceptor cell integrity may be applicable for novel therapeutic intervention/s for slowing or halting disease progression.

THE PROTECTIVE ROLE OF THE RETINAL PIGMENT EPITHELIUM

As in the brain, retinal function relies on glycolysis and oxidative phosphorylation, which are coupled to the citric acid cycle. These processes require a sustained delivery of oxygen and glucose as well as adequate control of the enzymes that allow for equilibrated formation and consumption of ATP. The blood-retinal barrier actively promotes homeostasis by tightly controlling the stoichiometry and activity of a network of proteins that maintain ionic gradients and metabolic transport systems and preserve the environment through detoxifying mechanisms that scavenge and remove toxic molecules (Fig. 1) (21).

The outer blood-retinal barrier is composed of three distinctive structures: the fenestrated endothelium of the choriocapillaris, Bruch’s membrane, and the RPE (21). The outer blood-retinal barrier mediates the exchange of small molecules and solutes and other metabolites from the blood stream to the photoreceptor layer (22). RPE cells are the most restrictive layer of the three components of the outer blood-retinal barrier, preventing the passage of biomolecules based on size and charge and thus preserving a controlled environment for photoreceptors. The retinal pigment epithelium, like other epithelia, is a compact structure where cells communicate laterally through tight junctions. In addition, the retinal pigment epithelium participates in retinal maintenance (22) by participating in complement activation, which is a major pathological mechanism of AMD (23). The RPE and pigment epithelium present an elaborate transcellular transport system that actively participates in the transport of nutrients, cellular debris, and toxic molecules to allow for the continuation of retinal function (24). All-trans-retinal takes place between the photoreceptor outer segment and the RPE (24). All-trans-retinal is esterified with a fatty acid to be isomerized to 11-cis-retinol and hydrolyzed from the ester bond in RPE cells. Then 11-cis-retinol is oxidized to 11-cis-retinal, and the active chromophore then leaves the RPE apical surface and moves to the photoreceptor to regenerate rhodopsin; afterward, the cycle begins again (reviewed in 25 and 26). Failure of the retinal pigment epithelium to accomplish its function in the retinoid cycle leads to retinal degeneration. For instance, autosomal-recessive inherited retinitis punctata albescens has been associated with mutations in the RLBP1 gene that encodes the cellular retinaldehyde-binding protein, which carries 11-cis-retinol and 11-cis-retinaldehyde in the RPE and Müller cells. When cellular retinaldehyde-binding protein is mutated, it loses the ability to bind the second ligand, and, as a consequence, all-trans-retinyl esters accumulate and evolve to produce RPE atrophy, retinal pigmentary changes, and decreased blood vessel development (27).

Integral to the fragility of photoreceptor cells is their close relationship with RPE cells. In Stargardt’s disease, a juvenile form of AMD, RPE cell functional integrity is initially compromised, and in turn photoreceptors are damaged. Once RPE cells die, photoreceptor cells then succumb (28). In other words, similar to what was proposed to occur in the blood-brain barrier in the neurovascular...
hypothesis of Alzheimer’s disease (AD) (29), defective clearance of certain molecules across the blood-retinal barrier may initiate a series of faulty maintenance functions that could lead to a retino-vascular inflammatory response, contributing to the development of AMD.

RPE CELLS AND VASCULAR REMODELING

The significance of RPE cells in vascular remodeling is highlighted in several studies. For example, absence of RPE cells in mice expressing fibroblast growth factor (FGF) 9 directed by the tyrosinase-related protein 2 promoter (FGF9 transgenic mice) is due to forcing embryonic RPE cells to become neural retinal cells through the ectopic expression of FGF9 (30). These mice fail to form blood vessels in the choroidal layers adjacent to regions where RPE cells are absent; however, vessels are found near the patch where RPE cells are present at postnatal day 7, indicating the importance of these cells in vessel formation. Moreover, dependency between RPE and endothelial vascular cells continues during adulthood through regulation of neovascularization. Compelling evidence links RPE cells with the secretion of angiogenic-related factors. In particular, RPE from transgenic apolipoprotein E2 mice, which express human apolipoprotein E2 protein and whose eyes present features common to AMD patients, shows reciprocal unbalanced expression of pigment epithelium-derived factor (PEDF) and vascular endothelial growth factor (VEGF), indicating that neovascularization may be increased (31). Furthermore, autocrine VEGF signaling in RPE cells stimulates VEGF-related gene expression as well as PEDF modulation (32), which is a potent angiogenic inhibitor (33). Taken together, the balanced production and secretion of these factors contribute to the formation, maintenance, and remodeling of cells that surround the RPE layer and/or are in their vicinity.
(e.g., omega-3 and also omega-6), and exposure to light (41, 42). Recently it was shown that phagocytosis (24–48 h) of oxidized photoreceptor outer segments containing high oxidative products induces the downregulation of complement factor H in RPE cells, similar to the effect of pro-inflammatory cytokines tumor necrosis factor-α (TNFα) and interleukin (IL)-6 (43). The RPE complement regulatory system, in this manner, may be suppressed by pro-inflammatory conditions as well as phagocytosis of oxidized photoreceptor outer segments. Surprisingly, the process enhances refractoriness to oxidative stress-induced apoptosis in RPE cells (Fig. 2A) (44). The protective effect of photoreceptor outer segments is specific, because the phagocytosis of polystyrene microspheres by RPE cells does not lead to a protective response against oxidative stress. Furthermore, polystyrene microspheres failed to induce DHA release and activate synthesis of neuroprotectin D1 (NPD1); this will be discussed in the following section. Interestingly, photoreceptor outer segment-mediated RPE cell protection against oxidative stress, with concurrent activation of NPD1 synthesis, was shown in ARPE-19 cells (44), a spontaneously immortalized human cell line (45), as well as in low passage primary human RPE cells prepared from National Disease Research Interchange-supplied eyes (unpublished observations).

Fig. 2. Photoreceptor outer segment phagocytosis elicits protection in RPE cells subjected to oxidative stress. A: Quantitative analysis of Hoechst stained ARPE-19 cells indicates that photoreceptor outer segment phagocytosis significantly decreases the amount of apoptosis observed during oxidative stress. Phagocytosis of polystyrene microspheres during oxidative stress did not alter the amount of apoptosis observed during oxidative stress alone. Results represent averages ± SEM of repeats of two independent experiments. B: NPD1 changes as a function of time after photoreceptor outer segment phagocytosis or microspheres; effect of oxidative stress. NPD1 has been quantified in cells as well as in incubation media. Data represents average ± SEM of two independent studies. Statistical analysis is Student’s t-test. NS, not statistically significant.

DHA RELEASE AND NPD1 FORMATION

RPE cells respond to oxidative stress by activating synthesis of NPD1 from DHA (46). The name NPD1 was suggested based upon its neuroprotective bioactivity in oxidative stressed RPE cells and the brain (46, 47) and its potent ability to inactivate pro-apoptotic and pro-inflammatory signaling. D1 refers to its being the first identified stereoselective mediator derived from DHA. NPD1 can be formed from free (unesterified) DHA released from membrane phospholipids by a phospholipase A2 (PLA2) upon stimulation (Fig. 3A). DHA belongs to the essential omega-3 fatty acid family (derived from linolenic acid, 18:3, n-3). Photoreceptor cells are highly enriched in DHA, and they tenaciously retain DHA even during very prolonged periods of omega-3 fatty acid deprivation (41, 48, 49).

The amount of unesterified DHA simultaneously measured in RPE cells and in incubation media by MS/MS was found to be increased as a function of time during exposure to oxidative stress in RPE cells. Specifically, the free intracellular DHA pool size showed a moderate increase after 6 h when cells were subjected only to photoreceptor outer segment phagocytosis (44). Oxidative stress, however, strongly enhanced free DHA accumulation in a time-dependent fashion, peaking at 16 h (44). Interestingly, although the overall increase reached 10-fold, photoreceptor outer segment phagocytosis kept the DHA pool size at a constant 2.4-fold increased level. This implies that NPD1 synthesis does not result from the simple enhancement of the overall availability of free DHA upon phagocytosis. There is a general correlation between increases in free DHA pool size and in NPD1 synthesis. Photoreceptor outer segment phagocytosis stimulates NPD1 synthesis at 3–6 h in cells and accumulation in media after 16 h, while free DHA increases earlier and keeps accumulating up to 16 h. These enhancements in DHA and NPD1 pool size are much larger when photoreceptor outer segment phagocytosis takes place on RPE cells exposed to oxidative stress. Interestingly, microsphere phagocytosis does not cause enhanced changes in DHA and NPD1 (Fig. 2B). As such, a very specific free DHA pool may be the precursor for NPD1.

Therefore, the supply of DHA and the induction of NPD1 synthesis during photoreceptor outer segment phagocytosis represents a homeostatic regulatory event for RPE cell protection in conditions of oxidative stress challenge and, as a consequence, the fostering of photoreceptor cell integrity (44). In this context, not only is photoreceptor outer segment phagocytosis in RPE cells essential for photoreceptor cell function, but the survival of the RPE promoted by this process correlates with NPD1 synthesis.

ON THE STRUCTURE OF NPD1, BIOSYNTHESIS, AND STEREOCHEMICAL ASSIGNMENT

The results discussed in this review provide the biological basis for the important actions of NPD1 derived from DHA. In this section, we shall consider the results obtained...
Fig. 3. A: NPD1 biosynthesis. Representation of the oxygenation of DHA to form NPD1. PLA$_2$ releases DHA from the second carbon position of the phospholipids upon stimulation. 15-Lipoxygenase-1 catalyzes the synthesis of 17S-H(p)DHA, which is converted to a 16(17)-epoxide and then is enzymatically converted to NPD1. B: Comparison of NPD1/PD1 biosynthesis with that of 10S,17S-diHDDA isomer (see detailed discussion in the text).
from several independent lines of investigation required to address the structure of the potent bioactive NPD1. As with other bioactive mediators, such as the eicosanoids (50), it is important to establish the stereochemistry of the compound and/or mediator, because many structurally related products can be less active, inactive, or even in some cases display opposing biologic actions as a result of subtle changes in stereochemistry that are recognized in biologic systems. To confirm the proposed basic structure and establish the complete stereochemistry, these studies on the 10,17S-docosatriene termed NPD1 included results from biosynthesis studies, matching of materials prepared by total organic synthesis with defined stereochemistry, and the actions of these and related compounds in biological systems (42, 44, 46, 47, 51–55). We also considered the chronology in which these findings appear in the literature with the goal of providing a clear and rigorous account of the evidence that supports the structure and bioactions of NPD1/protectin D1 (PD1) for the readership of The Journal of Lipid Research. As interested JLR readers will surmise, investigations along these lines were essential to establish the complete structure of the potent NPD1/PD1 and related endogenous products biosynthesized from DHA in vivo/in situ because of the small amounts of NPD1 attainable from biological systems at the time, which precluded direct stereochemical analyses of the products identified in RPE cells. Thus, in this section, we focus on the evidence for NPD1/PD1 structural elucidation. JLR readers interested in the structural elucidation of the resolvins and their complete stereochemical assignments are directed to other recent reviews (56, 57).

In 1984, the first evidence was obtained for the conversion of DHA to mono-, di-, and tri-DHA-derived products, named docosanoids, in the retina (an integral part of the central nervous system) (58). Use of available inhibitors of the time suggested a role for lipoxigenase in the biosynthesis of these compounds. An initial step in docosanoid synthesis was envisioned to be the release of DHA from membrane phospholipids by PLA2, early demonstrated to be rapidly activated by ischemia or seizures (41). The structure of 10,17-docosatriene was first disclosed while reporting the characterization of the novel bioactive resolvins that were identified using a systems approach with resolving inflammatory exudates and LC-MS-MS-based lipidomics (51). These new compounds (resolvins and docosanienes) were biosynthesized from omega-3 essential fatty acids during the resolution phase of acute inflammatory reactions in vivo that promote resolution of inflammation in vivo [see Fig. 8 in reference (51) and related text]. Since the DHA-derived compounds we identified in resolving inflammatory exudates, additional evidence was obtained for their biosynthesis from murine brain and vascular endothelial cells for the new bioactive compounds (47). These investigations focused on aspirin and its impact in the biosynthesis of 17R-hydroxy-containing resolvins and related structures. The initial results indicated that DHA-derived products reduced cytokine IL-1β production by human glial cells stimulated with TNFa. In parallel, studies with human cells were carried out to reconstruct the potential biosynthetic routes involved in the biosynthesis of these mediators. In this context, hypoxic endothelial cells exposed to inflammatory stimuli in vitro converted DHA and eicosapentaenoic acid to intermediates that were taken up by human leukocytes and further converted to bioactive products that showed potent activities relevant to the control of inflammation (51, 59, 60).

Of interest to the present review, in these investigations without aspirin treatment, 17S-HDHA and corresponding 17S-hydroxy-containing di- and trihydroxy products were reported in murine exudates and isolated human cells (51). The formation of some of these compounds was modeled in vitro and by sequential lipoxigenase reactions. These products were investigated with 15-lipoxygenase and included the double dioxygenation products, namely 7S,17S-diHDHA and 10,17S-diHDHA, which were identified along with trihydroxy-containing products formed via epoxide-containing intermediates from DHA (51). The well-established lipoxigenase reaction mechanism suggested that new products 7S,17S-diHDHA and 10,17S-diHDHA, which could easily be made in vitro, each contained two diene conjugated double bond systems both in a trans, cis geometry. For example, the well-known 5S,12S-diHETE formed from arachidonate via double dioxygenation is an isomer of the potent chemoattractant LTB4 (50). This isomer 5S12S-diHETE separates in SP-HPLC from LTB4 and are very similar to each other, but 5S12S-diHETE shows little chemotactic activity compared with LTB4 (61).

DHA is also precursor to a novel family of endogenous docosanienes formed in blood, leukocytes, brain, and glial cells (46, 47, 52). The main bioactive member of the docosanienes from the 17S-hydroxy-containing docosanoids proved to be 10,17S-docosatriene, in addition to the resolvins (47, 52). Also, human polymorphonuclear leukocytes (PMN) convert 10,17S-docosatriene to its omega-22 hydroxy product with DHA as the precursor; this is likely an inactivation route for this compound (47). As with the new bioactive products from omega-3 precursors, the basic structures and proposed stereochemical assignments reported were based on results of biosynthesis studies and given as tentative stereochemical assignments, because matching studies with synthetic reference compounds of known stereochemistry were still underway (see below). At the time, some of the newly identified compounds were matched to reference compounds prepared with plant lipoxigenases, e.g., 17S-HDHA and 7S,17S-diHDHA, which matched to those profiled by LC-MS-MS in exudates and with isolated cells. It was clear that the 10,17S-docosatriene from exudates did not coelute with the major 10,17S-docosatriene produced in vitro with plant enzymes, suggesting it was an isomer; however, this system could be used to prepare related docosanienes via the LTA4 synthase activity of lipoxigenases to further probe the bioactions of the 10,17S-docosatriene while complete matching and total synthesis were in progress.

Of note, glial cells generate both 17S series resolvins and the 10,17S-docosatriene. Importantly, evidence for a novel omega-22-hydroxy-16,17S-trihydroxydocosatriene...
was obtained from these cells and human PMN, which suggest that the 10,17S-docosatriene biosynthesis is via a 16(17S)-epoxide-containing intermediate, because the identified vicinal diol could be a product of this epoxide intermediate. Hence, a series of alcohol trapping studies were undertaken to address the potential role of epoxide-containing intermediates in the biosynthesis of the bioactive compounds, in particular the 10,17S-docosatriene. Indeed, evidence for epoxide-containing intermediates in the biosynthesis of docosatrienes and 17S series resolvins was obtained from human PMN (52). In these incubations with human cells, two 16-OCH3 and two 10-OCH3 methoxy-trapping products, likely all-trans in their triene conjugation, were obtained, which implicates production of a 16(17S)-epoxide intermediate that was proposed in the biosynthetic pathway for the bioactive 10,17S-docosatriene (52). Also, 10,17S-docosatriene proved to display potent actions with human glial cells and produced 10,17S-docosatriene, which reduced IL-1β production at 1–50 nM and evoked ligand-operated extracellular acidification with glial cells in a microphysiometer (52). These findings indicated that DHA is precursor to novel potent protective mediators and that 10,17S-docosatriene carries potent anti-inflammatory activity in vivo and with human cells in vitro as well as activated surface receptors present on human glial cells to regulate their function. Thus, the basic structure of the novel 10,17S-docosatriene, later coined NPD1/PD1 (see below) due to its potent actions in vivo and in vitro in cell cultures, was established (47, 51, 52). This docosatriene also displayed potent anti-inflammatory actions, i.e., reducing PMN numbers in vivo and reducing the production of inflammatory cytokines by glial cells in vitro. Moreover, during the resolution phase of peritonitis, unesterified DHA levels increase in resolving exudates, where it appears to promote catabasis or the return to homeostasis following tissue insult via conversion to D-series resolvins and also 10,17S-docosatrienes (62) by shortening the resolution interval of an inflammatory response in vivo (63).

We then found that the DHA-derived 10,17S-docosatriene proved to be generated in vivo during experimental stroke in the ipsilateral cerebral hemisphere following focal ischemia and also demonstrated potent bioactions in this system, where it limits the entry of leukocytes, down-regulates cyclooxygenase-2 expression and nuclear factor κB activation, and decreases infarct volume (47). Next, we found that 10,17S-docosatriene is formed in the human retinal pigment epithelial cell line, ARPE-19, and introduced the term NPD1 based on its neuroprotective bioactivity (46, 47).

Taken together, these findings in ARPE-19 cells (46), inflammatory murine exudates, human PMN, glial cells, and the brain (47, 51, 52) underscored the need to establish the complete stereochemistry of endogenous, biologically generated, active 10,17S-docosatriene, namely the chirality of its carbon 10 position alcohol and its triene double geometry, which remained to be established (Fig. 3B). These involved the total organic synthesis and matching of both bioactivity and physical properties of the endogenous compound and those of the synthetic with established stereochemistry. In recognition of its wide scope of formation and uncovered actions, PD1 was introduced and used to denote the structure of this chemical mediator in the immune system. The prefix Neuro before PD1 (NPD1) was proposed to denote its biosynthesis and potent neuroprotective actions (53). It was also apparent that NPD1/PD1 was a member of a larger family of 17-hydroxy-containing docosatrienes, termed protectins.

Biosynthesis and function studies were undertaken with human T₈₂-skewed peripheral blood mononuclear cells (PBMC) that specifically express 15-lipoxygenase type 1 and convert DHA to the 10,17S-docosatrienes by serving as a 17-lipoxygenase with DHA as a substrate. When produced by these cells, PD1 promotes T cell apoptosis via the formation of lipid raft-encoded signaling complexes and reduces T-cell traffic in vivo. These results were consistent with the physical properties of NPD1. Matching materials prepared by total organic synthesis determined the complete stereochemistry of the PBMC DHA-derived product. NPD1/PD1 generated by human PBMC carried the complete stereochemistry of (10R,17S)-dihydroxydocosa-4Z,7Z,11E,13E,15Z,19Z-hexaenoic acid and was matched to the most potent bioactive product using several dihydroxytriene-containing, DHA-derived products isolated from human PBMC, human PMN, and murine exudates (53, 54).

During the course of these investigations, Butovich et al. (64) reported that NPD1 had the complete structure of 10S,17S-diHDDHA. This was based on results obtained with isolated lipooxygenase enzymes incubated with DHA without mammalian cell/tissue biosynthesis or authentic NPD1, as defined earlier in the literature (46, 51, 52). Importantly, neither the bioactivity of the product nor appropriate comparisons with authentic NPD1 was presented to support the conclusions in this report (64) in regards to the complete structure of NPD1. Note that 10S,17S-diHDDHA is an isomer of NPD1/PD1 (Fig. 3B).

With the preparation of six stereochemically defined, 10,17-dihydroxy-containing geometric isomers by total organic synthesis that had been initiated earlier, it was possible to match the stereochemistry and biological actions of the endogenously produced materials (54). In addition to PD1 formed from human leukocytes, additional isomers were identified in inflammatory exudates, including Δ15-trans-PD1 (isomer III), 10S,17S-dihydroxy-docosa-4Z,7Z,11E,13Z,15Z,19Zhexaenoic acid (isomer IV), and the expected double dioxygenation product 10S,17S-dihydroxy-docosa-4Z,7Z,11E,13Z,15E,19Zhexaenoic acid (isomer I), which are present in inflammatory exudates obtained from mice. ¹⁸O-labeling results provided evidence that this isomer I was a double dioxygenation product and that the 10 position of NPD1/PD1 originated from enzymatic conversion. Also, the rank order of activities was established between these isomers, and NPD1/PD1 proved to be most potent (55), with doses as low as 1–10 ng reducing murine peritonitis, followed by Δ15-trans-PD1 > 10S,17S-diHDDHA (isomer I). Hence, although the double dioxygenation product 10S,17S-diHDDHA was generated in
murine exudates, it was far less active than PD1/NPD1 both in vitro and in vivo.

The proposed biosynthetic route for NPD1/PD1 is shown in Fig. 3B from results previously reported (54). Following 17S HpDHA formation from 15-lipoxygenase action on DHA, an epoxide intermediate is formed that requires enzymatic transformation to obtain the correct double bond geometry, namely *cis,trans,trans* present in NPD1/PD1. This double bond geometry and chirality of the carbon 10 position and the *R*-configuration were established from the matching of synthetic compounds of defined chirality. Of interest, both 7S,17S diHDHA (resolvin D5) and 10S,17S diHDHA are double dioxygenation products, and at this point they appear to be less active than endogenous or synthetic NPD1/PD1. Without endogenous biosynthesis studies or assessment of biological actions, it is surprising that a claim could be made for assessment of the complete stereochemistry of NPD1 based only on nuclear magnetic resonance results obtained for 10S,17S diHDHA from plant lipoxygenase-prepared material with DHA (64).

With the stereochemistry of NPD1/PD1 established, its identification in human material was sought and found to be present in breath condensates from human asthmatics (65) and in the human brain, both under basal conditions and from patients with AD (66). In addition, PD1 was found to be a major product in bone marrow of female rats fed eicosapentaenoic acid and DHA (67). PD1 was generated in vivo during ischemia-reperfusion of renal tissues, where it has profound actions, namely reversing the deleterious consequences of ischemia-reperfusion in renal tissues (68), in agreement with our previous observations in the brain (47).

With the complete stereochemistry and synthetic compound in hand, it was possible to demonstrate for the first time that PD1 activates resolution programs in vivo and that it shortens the resolution time of experimental inflammation in animal models (63). With the total organic synthesis route of NPD1/PD1 in place, it was possible to radiolabel and purify *3H*-NPD1/PD1 made from the synthetic intermediate. With this radiolabel, it was then possible to define for the specific binding sites present with ARPE-19 cells (*K*<sub>d</sub> ~ 31 pM/mg cell protein) for *3H*-NPD1/PD1 as well as specific binding to human neutrophils that gave a *K*<sub>d</sub> of ~25 nM (55). Most importantly, critical information on NPD1 biosynthesis with ARPE-19 cells was obtained, namely identification of alcohol-trapping products, indicating the formation of a 16,17-sepoxide-containing intermediate from DHA in the biosynthesis of NPD1 (55). These results, as well as the rank order of potency established for NPD1/PD1 of defined stereocchemical analysis, indicated that the most potent of the isomers prepared was NPD1/PD1. Also in the ARPE-19 cells, NPD1 was more potent than either resolvin D1 or resolin E1. In other systems, resolin E1 and resolin D1 established higher potencies than PD1 (69).

Along with the complete stereocchemical identification of NPD1/PD1, recent studies using an unbiased LC-MS-MS profiling approach demonstrated that PD1 is made during the resolution of Lyme disease infections in mice (70). NPD1 inhibits retinal ganglion cell death (71), is renal protective (72), and regulates adiponectin (73). Of interest, the double dioxygenation product 10S,17S diHDHA isomer of NPD1/PD1 was recently shown to have actions on platelets, reducing platelet aggregation at 0.3 μM, 1 μM, and higher concentrations (74). In peritonitis, this isomer also showed biological activity but was less potent than NPD1/PD1 (54, 74). It is noteworthy that NPD1/PD1 and the resolins also are produced by trout tissues, including trout brain, from endogenous DHA, suggesting that these structures are highly conserved from fish to humans (75). This shows we still have much to learn regarding the bioactions and functions of NPD1/PD1, the D-series and E-series resolvins, and related products in human physiology and pathophysiology as well as in biological systems such as fish, where the actions of NPD1/PD1 remain to be fully appreciated.

NEUROTROPHINS INDUCE THE SYNTHESIS AND RELEASE OF NPD1 FROM HUMAN RPE CELLS

RPE cells also are capable of producing a wide variety of growth factors (76). These trophic factors support surrounding cells by paracrine and autocrine signaling and hence promote the communication and structure of the retina as well as photoreceptor survival (77, 78). It is important to note that neurotrophins enhance the production of NPD1 in RPE cells. In turn, NPD1 is released and serves as a lipid signaling messenger in its surroundings. This observation was made in human RPE cells grown to confluence using a specialized culture (79) that allows the cells to develop a high degree of differentiation, preserving the apical-basolateral polarization (Fig. 4A, C). Neurotrophins [pigment epithelium derived factor (PEDF), BDNF, ciliary neurotrophic factor, FGF, glial-derived neurotrophic factor (GDNF), leukemia inhibitory factor, NT3, or persephin], which have bioactivities that promote neuronal and/or photoreceptor cell survival, are also agonists of NPD1 synthesis (Fig. 4B), favoring the release of this lipid messenger through the cell’s apical surface (80). Among all the growth factors tested, PEDF is by far the most potent stimulator of NPD1 synthesis. PEDF, a member of the serine protease inhibitor (serpin) family, was identified in human RPE cells. If PEDF or ciliary neurotrophic factor are added to the incubation media, bathing the basolateral side in increasing concentrations, they evoke a lesser degree of NPD1 release on the apical side. Conversely, if these neurotrophins are added to the apical side, they exert concentration-dependent increases in NPD1 release only on the apical side (Fig. 4B, C) (80).

NPD1 BIOSYNTHESIS IS AN ENDOGENOUS RESPONSE TO OXIDATIVE STRESS

Previous studies have shown that the retina forms mono-, di-, and trihydroxy derivatives of DHA and that
lipoxigenase inhibitors block this synthesis, indicating an enzymatic process of a lipoxigenase nature (42). Although the stereochemistry and bioactivity of DHA-oxidized derivatives were not defined at the time of these observations, it was suggested that these lipoxigenase products might be neuroprotective (and therefore the name docosanoids was introduced) (42, 81). Liquid chromatography-photodiode array-electrospray ionization MS/MS-based lipidomic analysis was used to identify oxygenation pathways for the synthesis of the docosanoid NPD1 during brain ischemia-reperfusion (47) and the retinal pigment epithelium (46). Moreover, it was also found that RPE cells have the ability to synthesize NPD1 (46). Photoreceptors and RPE cells, although they contain phospholipids richly endowed with DHA (as docosahexaenoyl- or DHA-elongated fatty acyl-chains), display an undetectable quantity of unesterified (free) DHA [as is the case with unesterified arachidonic acid (AA)] under basal, nonstimulated conditions (82–86). This means that the pool size of unesterified DHA is tightly regulated by production (PLA₂), by its removal (e.g., by reacylation), and by peroxidation. Free DHA incorporated into membrane phospholipids first becomes the substrate of docosahexaenoyl-CoA synthesis for its channeling through acyltransferases, which incorporate this fatty acid into phospholipids (87–90). The RPE cell thus modulates the uptake, conservation, and delivery of DHA to photoreceptors (81). In addition, RPE cells utilize a specific DHA-phospholipid pool as a precursor for NPD1 synthesis. Then this stereospecific mediator is synthesized by 15-lipoxygenase-1 (15-LOX-1) (91) (Fig. 3A). In AD brain (short postmortem time), cPLA₂ and 15-LOX-1 expression changed in concert with NPD1-decreased content and DHA-enhanced pool size in the CA1 area of the hippocampus (66). In ARPE-19 cells (spontaneously transformed human RPE cells), IL-1β, oxidative stress, or the Ca²⁺ ionophore A23187 activates the synthesis of NPD1 (46). In turn, NPD1 might act in an autocrine fashion and/or diffuse through the interphotoreceptor matrix to act in a paracrine mode on photoreceptor cells and/or Müller cells (41).

15-LOX-1 DEFICIENCY IN RPE CELLS PROMOTES APOPTOSIS AND IT IS RESCUED SELECTIVELY BY NPD1

The increased availability of DHA is followed by NPD1 synthesis. This is of particular interest in the process of phagocytosis of photoreceptor outer segments, given that the endogenous pool of DHA is augmented upon activation of the RPE cell phagolysosomal system. In this context, it is known that NPD1 production and release is increased (Fig. 2). Recent evidence shows 15-LOX-1 as the enzyme that oxygenates DHA into NPD1 (91). In ARPE-19 cells where 15-LOX-1 protein expression was knocked down by 70% posttranscriptionally, the production of NPD1 was not increased, in contrast with normal cells (Fig. 5). In normal cells, NPD1 production was induced as early as 4 h after oxidative stress was applied. In this way, preexisting pools of 15-LOX-1 are activated upon stress stimulation (Fig. 5B). Thus, phagocytosis triggers the activation of 15-LOX-1 to oxygenate DHA into NPD1.

15-LOX-1, a nonheme iron-containing dioxygenase, stereospecifically inserts oxygen into AA, dually forming 15(S)-hydroxyeicosatetraenoic acid (HETE) and 12(S)-HETE as well as lipoxin A₄, a product of its joint activity with 5-LOX. 15-LOX-1 also has the capability to oxygenate linoleic acid into 13-hydroxyoctadecadienoic acid (92). Human 15-LOX-1 and 12-LOX are highly homologous proteins (65% identity) encoded by different genes, and their mRNAs are similar (70% identity). On the other hand, 15-LOX-2 is a different lipoxigenase that shares only 39% identity with human 15-LOX-1 (93). 15-LOX-2 and human 12-LOX differ from 15-LOX-1 not only in the ratio of 15-HETE and 12-HETE produced from AA but also in their localization (Fig. 5A). This means that they possess different selective product formations, and thus their activities contribute to different lipid mediators as well as display different substrate availability.

Silencing 15-LOX-1 also leads to enhanced susceptibility to apoptosis (Fig. 5C). It is worth noting that only NPD1 could rescue these cells from the exaggerated apoptosis experienced under oxidative stress conditions. This indicates that cells failing to produce this lipid mediator have an increased sensibility to oxidative stress caused by lack of the pro-survival signaling elicited by NPD1.

Fig. 4. Neurotrophins activate NPD1 synthesis in cultured primary human RPE cells. A: Zonula occludens-1 (ZO-1) antibody immunoreactivity (green) illustrates confluence of the monolayer polyhedric-shape of the cells. B: Differential ability of growth factors to selectively release NPD1 through the apical surface of the cell. Growth factors (20 ng/ml) were added to the apical medium. Apical and basal media were collected separately after 72 h and subjected to lipidomic analysis. Each bar is an average ± SEM of five independent wells. Statistical analysis was performed using Student’s t-test shows *P < 0.05. C: Schematic representation of the monolayer orientation within the insert. [Fig. 4 A, C, modified with permission from reference (40)].
The novel signaling pathway, resulting from the NPD1 action, may be initiated through the photoreceptor outer segment phagocytosis. A significant body of evidence has been accumulated that supports this idea.

FGF2 promotes bovine RPE cell survival through a sustained adaptive phenomenon that involves both FGF1-mediated activation of extracellular signal-regulated kinase and extracellular signal-regulated kinase 2-dependent Bcl-xL production (94). Bcl-xL may play a key role in integrating and transmitting exogenous FGF2 signals for RPE cell survival. Moreover, a well-organized signaling regulatory mechanism on the apical side of the RPE cell is reflected by the ability of neurotrophins to induce NPD1 synthesis and release (80). The response of human RPE cell monolayers in culture with NPD1 synthesis and release upon addition of certain neurotrophins to the apical side suggests sidedness of receptors for these ligands (80). Persephin is a novel neurotrophin with homology to GDNF (95, 96). Both persephin and GDNF are agonists of NPD1 synthesis and activators of its release from the apical surface of RPE cells (80). The same was true for leukemia inhibitory factor and FGF2 as well as for other neurotrophins (80). The finding that there is polarized (apically) neurotrophin/mediated NPD1 release has relevance for the initiation and progression of retinal degenerations. This is because when RPE cell polarization in the plane of the epithelium is disrupted, dysregulated growth factor secretion and pro-inflammatory signaling arises (2, 97, 98), thereby setting in motion pathological changes that include the proliferative component of macular degeneration: choroidal neovascularization (99–101).

Bcl-2 family proteins regulate apoptotic signaling at the mitochondrial level and at the endoplasmic reticulum. As a consequence, cytochrome c is released from mitochondria and caspase-3, a downstream effector of pro-apoptotic and anti-apoptotic Bcl-2 proteins, is activated (102). Also, oxidative stress-induced activation of caspase-3 in RPE cells is decreased by NPD1 (46). Apoptosis is an outcome of excessive oxidative stress in RPE cells, and NPD1 is effective in countering this oxidative stress-induced cell death (46). It is interesting that DHA itself inhibits apoptosis, concomitant with a remarkable, time-dependent formation of NPD1. Significantly, the potency of DHA for cytoprotection is much higher than that of added NPD1 (46), suggesting that NPD1 might exert its action close to the subcellular site of its synthesis. Importantly, these actions of DHA cannot be mimicked by other PUFAs (e.g.,...
20:4, n-6). Alternatively, it implies that other NPD-like mediators participate in promoting RPE cell survival, even though DHA fails to exert protection in 15-LOX-1-deficient cells, confirming that its pro-survival effect is mediated via NPD1. In RPE cells, cleavage of endogenous substrates by caspase-3 is enhanced by oxidative stress, as indicated by increased accumulation of poly(ADP-ribosyl) polymerases. NPD1 inhibits caspase-3 activation when added at the onset of oxidative stress (46), likely reflecting a downstream consequence of NPD1 modulation of Bcl-2 proteins.

A consequence of RPE cell damage and apoptosis is impaired photoreceptor cell survival, a dominant factor in AMD (103). The pigment lipofuscin, which increases in the RPE during aging, accumulates further during AMD. The progressively greater onslaught of photooxidative damage to the RPE affects photoreceptor survival. For example, in the juvenile form of macular degeneration known as Stargardt's disease, oxidative stress mediated by the lipofuscin fluorophore N-retinylidene-N-retinylethanolamide produces RPE damage; caspase-3 is part of the damaging cascade, whereas Bcl-2 exerts cellular protection (104). NPD1 downregulates lipofuscin fluorophore N-retinylidene-N-retinylethanolamide-mediated apoptosis induced by oxidative stress, restoring the integrity of the RPE and its relationship with the photoreceptor (80).

ANTI-APOPTOTIC AND ANTI-INFLAMMATORY BIOACTIVITY OF NPD1 IS MEDIATED IN PART THROUGH MODULATION OF GENE EXPRESSION

Photoreceptor outer segment phagocytosis is a mechanism that initiates NPD1 signaling. This signaling promotes modulation of protein expression and activity involved in counteracting apoptosis at the cell fate decision level. For instance, NPD1 promotes differential changes in expression of Bcl-2 family proteins, upregulating protective Bcl-2 proteins (Bcl-2, Bcl-xL, and Bfl-1/A1) and attenuating expression of proteins that challenge cell survival (e.g., Bax, Bad, Bid, and Bik). Thus, an NPD1-mediated and coordinated regulation of Bcl-2 protein availability for subsequent downstream signaling may be crucial for cell survival (46, 80). NPD1 regulates expression of the genes encoding death repressors and effectors of the Bcl-2 family of proteins, e.g., Bcl-2-associated athanogene domain 3, whose mRNA is upregulated in mouse retinas subjected to light damage (105, unpublished observations). Bcl-2-associated athanogene domain 3 is a co-chaperone protein involved in release of the refolded protein by the chaperone Hsp70 (106-108).

NPD1 activity lessens the effects of stressors such as oxidative stress not only by its production and release upon stimulation but also because its downstream signaling modulates gene transcription by compensating the activation or inhibition of gene expression. Thus, the consequence of the signaling is to return the system to normal levels. This is the case with NPD1 downregulation of the pro-apoptotic death-associated protein kinase 1, which displays increased expression under oxidative stress treatment (unpublished observations). In RPE cells, NPD1 downregulates the expression of pro-inflammatory genes, such as cyclooxygenase 2 (COX-2), which is induced by cytokines such as IL-1β (46, 109). In ischemia reperfusion-injured hippocampus, as well as in neural progenitor cells stimulated by IL-1β, NPD1 also inhibits COX-2 induction (46, 47). In brain ischemia reperfusion, NPD1 decreases infarct size and inhibits PMN infiltration (47). Moreover, our laboratory, through a genome-wide screen in human brain progenitor cells in culture (66), has identified other NPD1-targeted pro-inflammatory genes; they include IL-1β, cytokine exodus protein-1, and TNFα-inducible pro-inflammatory element (B94, TNFAIP2). NPD1 bioactivity acts as a modulatory signal that counteracts pro-inflammatory injury to the RPE, a condition involved in pathangiogenic signaling in the wet form of AMD and in proliferative vitreoretinopathy of diabetic retinopathy.

Oxidative stress enhances pro-inflammatory gene expression that leads to RPE cell injury. The inducible enzyme COX-2 is the rate-limiting step in prostaglandin synthesis and is involved in oxidative stress as well as cell function. COX-2 expression is regulated in RPE cells by photoreceptor outer-segment phagocytosis and by growth factors (110), and IL-1β activates expression of the proximal human COX-2 promoter. In the latter case, NPD1 potently counteracts the activation of the transcription mediated by IL-1β, displaying an IC_{50} of <5 nM (46).
CONCLUDING REMARKS

NP1D1 is a pro-survival, anti-inflammatory, and homeostatic mediator that, by acting on RPE cells, promotes photoreceptor cell integrity (42). NP1D1 synthesis agonists include neurotrophins and oxidative stress, and thus this lipid mediator modulates signaling pathways related with photoreceptor integrity (42). NPD1 synthesis agonists are lipid mediators that, by acting on RPE cells, promote photoreceptor cell integrity (42). Rattner, A., and J. Nathans. 2006. Macular degeneration: recent advances in the biological and clinical understanding of macular diseases. Curr. Opin. Investig. Drugs. 2006: 1020–1025. Spero, M. A., M. A. Hauser, S. Schmidt, W. K. Scott, P. Gollins, K. L. Spencer, Y. A. Zyczynski, T. L. McGee, and J. L. Haines. 2005. Complement factor B component 2 (C2) gene is associated with age-related macular degeneration. Nat. Genet. 38: 458–462. Several questions remain: What are the molecular mechanisms through which NP1D1 synthesis is stimulated? What are the early molecular NP1D1 activities/effectors in the signaling pathway? The evolving information on the bioactivity of NP1D1 in RPE/photoreceptor interactions suggests that DHA/NPD1 signaling may contribute to preventing and/or halting progression of retinal degenerative diseases.

REFERENCES

1. Papamarkos, D. S. 2002. The birth and death of photoreceptors: the Friedenwald lecture. Invest. Ophthalmol. Vis. Sci. 43: 1300–1309.
2. Rattner, A., and J. Nathans. 2006. Macular degeneration: recent advances and therapeutic opportunities. Nat. Rev. Neurosci. 7: 860–872.
3. Chang, G. Q., Y. Hsu, and F. Wang. 1993. Apoptosis: final common pathway of photoreceptor death in rd, rdS, and rhodopsin mutant mice. Neuron. 11: 593–605.
4. Porteria-Gualdi, C., C. H. Sung, J. Nathans, and R. Adler. 1994. Apoptotic photoreceptor death in mouse models of retinitis pigmentosa. Proc. Natl. Acad. Sci. USA. 91: 974–978.
5. Bird, A. C. 2003. The Bowman lecture: Towards an understanding of age-related macular disease. Eye (Lond.). 17: 457–466.
6. Liu, Q. J., and A. E. Pierro. 2004. The retinoid pigmentosa 1 protein is a photoreceptor microtubule-associated protein. J. Neurosci. 24: 6427–6436.
7. Dryja, T. P., T. L. McGee, E. Reichel, L. B. Hahn, G. S. Cowley, D. W. Vandell, M. A. Sandberg, and E. L. Berson. 1990. A point mutation of the rhodopsin gene in one form of retinitis pigmentosa. Nature. 343: 364–366.
8. Mendes, H. F. J., van der Spuy, J. P. Chapple, and M. E. Cheetham. 2005. Mechanisms of cell death in rhodopsin retinitis pigmentosa: implications for therapy. Trends Mol. Med. 11: 177–185.
9. de Jong, P. T. 2006. Age-related macular degeneration. N. Engl. J. Med. 355: 1474–1485.
10. Sreekumar, P. G., R. Kannan, J. Young, C. K. Spec, S. J. Ryan, and D. R. Hinton. 2005. Protection from oxidative stress by methionine sulfoxide reductases in RPE cells. Biochem. Biophys. Res. Commun. 334: 245–253.
11. Shen, J. K., A. S. Hackett, W. R. Bell, W. R. Green, and P. A. Campochiaro. 2007. Oxidative damage in age-related macular degeneration. Histol. Histopathol. 22: 1301–1308.
12. Dunaief, J. L., T. Dentschew, G. S. Ying, and A. H. Milam. 2002. The role of apoptosis in age-related macular degeneration. Arch. Ophthalmol. 120: 1435–1442.
13. Fisher, D., I. Kovacs, M. Artico, C. Cavallotti, A. Papale, and C. Balacco Gabrielli. 2006. Mitochondrial alterations of retinal pigment epithelium in age-related macular degeneration. Neurobiol. Aging. 27: 983–993.
14. Hageman, G. S., D. H. Anderson, L. V. Johnson, L. S. Hancox, A. J. Taiber, L. I. Hardisty, J. L. Hageman, H. A. Stockman, J. D. Borchardt, K. M. Gehrs, et al. 2005. A common haplotype in the complement regulatory gene factor H (HFI/CHF) predisphophoreceptor outer segment is associated with age-related macular degeneration. Proc. Natl. Acad. Sci. USA. 102: 7527–7532.
15. Klein, R., J. C. Zeiss, E. Y. Chew, J. Y. Tsai, R. S. Sackler, C. Haynes, A. K. Hening, J. L. SanGiovanni, S. M. Mane, S. T. Mayne, et al. 2005. Complement factor H polymorphism in age-related macular degeneration. Science. 308: 385–389.
16. Edwards, A. O., R. Ritter III, K. J. Abel, A. Manning, C. Panhuysen, and L. A. Farber. 2005. Complement factor H polymorphism and age-related macular degeneration. Science. 308: 421–424.
17. Haines, J. L., M. A. Hauser, S. Schmidt, W. K. Scott, L. M. Olson, P. Gollins, K. L. Spencer, S. Y. Kwan, M. Nourreddine, J. R. Gilbert, et al. 2005. Complement factor H variant increases the risk of age-related macular degeneration. Science. 308: 419–421.
18. Geidt, B., J. E. Meric-Bernstein, L. S. Hancox, A. J. Taiber, K. Gehrs, K. Cramer, J. Neel, J. Bergeron, G. R. Barile, et al. 2006. Variation in factor B (BF) and complement component 2 (C2) genes is associated with age-related macular degeneration. Nat. Genet. 38: 458–462.
19. Bok, D. 2005. Evidence for an inflammatory process in age-related macular degeneration gains new support. Proc. Natl. Acad. Sci. USA. 102: 7053–7058.
20. Spencer, K. L., M. A. Hauser, L. M. Olson, S. Schmidt, W. K. Scott, P. Gollins, A. Agarwal, E. A. Postel, M. A. Perick-Vance, and J. L. Haines. 2007. Protective effect of complement factor B and complement component 2 variants in age-related macular degeneration. Hum. Mol. Genet. 16: 1986–1992.
21. Pourmousavi, C. J., E. Vanruyten-Brendeels, C. E. Riva, S. H. Hardarrison, and E. Stefansson. 2008. Regulation of retinal blood flow in health and disease. Prog. Retin. Eye Res. 27: 284–330.
22. Strauss, O. 2005. The retinal pigment epithelium in visual function. Physiol. Rev. 85: 845–881.
23. Wang, J. S., and V. J. Kefalov. 2009. An alternative pathway mediates the mouse and human cone visual cycle. Curr. Biol. 19: 1665–1669.
24. Navid, A. S. C. Nicholas, and R. D. Hamer. 2006. A proposed role for all-trans retinal in regulation of rhodopsin regeneration in human rods. Vision Res. 46: 1449–1463.
25. Saari, J. C. 2000. Biochemistry of visual pigment regeneration: the Friedenwald lecture. Invest. Ophthalmol. Vis. Sci. 41: 337–348.
26. Lamb, T. D., and E. N. Pugh, Jr. 2004. Dark adaptation and the retinoid cycle of vision. Prog. Retin. Eye Res. 23: 307–380.
27. Besch, D., H. Jägle, H. P. N. Scholl, M. W. Seeliger, and E. Zrenner. 2003. Inherited multifocal RPD-diseases: mechanisms for local dysfunction in global retinoid cycle gene defects. Vision Res. 43: 3095–3108.
28. Cádiz, A. V., T. S. Aleman, M. Swider, S. B. Schwartz, J. D. Steinberg, A. J. Brucker, A. M. Maguire, J. Bennett, E. M. Stone, and S. G. Jacobson. 2004. Mutations in ABCA4 result in accumulation of lipofuscin before slowing the retinoid cycle: a reappraisal of the human disease sequence. Hum. Mol. Genet. 13: 525–534.
29. Zlokovic, B. V. 2005. Neurovascular mechanisms of Alzheimer’s neurodegeneration. Trends Neurosci. 28: 380–388.
30. Zhao, S., and P. A. Overbeek. 2001. Regulation of choroid development by the retinal pigment epithelium. Mol. Vis. 2: 277–282.
31. Lee, S. J., J. H. Kim, J. H. Kim, M. J. Chung, Q. Wen, H. Chung, K. W. Kim, and Y. S. Yu. 2007. Human apolipoprotein E2 transgenic mice show lipid accumulation first in retinal pigment epithelium and altered expression of VEGF and hFGF in the eyes. J. Microbiol. Biotechnol. 17: 1024–1030.
32. Gao, G., Y. Li, D. Zhang, S. Gee, C. Crosson, and J. Ma. 2001. Unbalanced expression of VEGF and PEDF in ischemia-induced retinal neovascularization. FEBS Lett. 499: 270–276.
33. Dawson, D. W., O. V. Volpert, F. Gillis, S. E. Crawford, H. Xu, W. Benedict, and N. P. Bouck. 2000. Pigment epithelium-derived factor: a potent inhibitor of angiogenesis. Science. 285: 245–248.
53. Ariel, A., P-L. Li, S. Wang, W-X. Tang, G. Fredman, S. Hong, R. Antony, K. Sheets, N. Petasis, C. N. Serhan, and N. G. Bazan. 2010. Neuroprotectin D1/protectin D1 stereoselective and specific binding with human retinal pigment epithelial cells and neutrophils. *Prostaglandins Leukot. Essent. Fatty Acids*, 82: 27–34.

54. Serhan, C. N. 2007. Resolution phases of inflammation: novel endogenous anti-inflammatory and pro-resolution lipid mediators and pathways. *Annu. Rev. Immunol.*, 25: 101–137.

55. Serhan, C. N., N. Chiang, and T. E. Van Dyke. 2008. Resolving inflammation: dual anti-inflammatory and pro-resolution lipid mediators. *Nat. Rev. Immunol.*, 8: 349–361.

56. Bazan, N. G., D. L. Birkle, and T. S. Reddy. 1984. Docosahexaenoic acid (22:6-3) is metabolized to lipoxigenase reaction products in the retina. *Biochem. Biophys. Res. Commun.*, 125: 741–747.

57. Serhan, C. N., C. B. Clish, J. Brannon, S. P. Colgan, N. Chiang, and K. Gronert. 2000. Novel functional sets of lipid-derivated mediators with antiinflammatory actions generated from omega-3 fatty acids via cylo-oxygenase 2-n-steroidal anti-inflammatory drugs and transcellular processing. *J. Exp. Med.*, 192: 1197–1204.

58. Marcheselli, V. L., P. K. V. Marcheselli, J. C. de Rivero Vaccari, W. C. Gordon, F. E. Jackson, and N. G. Bazan. 2007. Photoreceptor outer segment phagocytosis selectively attenuates oxidative stress.

59. Levy, B. D., P. Kohli, K. Gotlinger, O. Haworth, S. Hong, S. Kazani, E. Israel, K. J. Haley, and C. N. Serhan. 2007. Protectin D1 is generated in asthma and dampens airway inflammation and hyperresponsiveness. *J. Immunol.*, 178: 496–502.

60. Luik, W. J., J. G. Cuí, V. L. Marcheselli, M. Bodker, A. Botkjær, K. Gotlinger, C. N. Serhan, and N. G. Bazan. 2005. A role for docosahexaenoic acid-derived neuroprotectin D1 in neural cell survival and Alzheimer disease. *J. Clin. Invest.*, 115: 2774–2783.

61. Sam Giovanni, J., P., and E. Y. Chew. 2005. The omega-3 family of resolution and pro-resolving lipid mediators in bone marrow and their profile alteration with ovariec-
tomy and omega-3 intake. *Am. J. Hematol.*, 82: 437–445.

62. Levy, B. D., P. Kohli, K. Gotlinger, O. Haworth, S. Hong, S. Kazani, E. Israel, K. J. Haley, and C. N. Serhan. 2007. Protectin D1 is generated in asthma and dampens airway inflammation and hyperresponsiveness. *J. Immunol.*, 178: 496–502.

63. Blaho, V. A., M. W. Buczyński, C. R. Brown, and E. A. Dennis. 2009. Lipidomic analysis of dynamic eicosanoid responses during the induction and resolution of Lyme arthritis. *J. Biol. Chem.*, 284: 21599–21612.

64. Qin, Q., K. A. Patil, K. Gronert, and S. S. Sharma. 2008. Neuroprotectin D1 inhibits retinal ganglion cell death following axotomy. *Prostaglandins Leukot. Essent. Fatty Acids.*, 79: 201–207.

65. Hassan, I. R., and K. Gronert. 2009. Acute changes in dietary omega-3 and omega-6 polyunsaturated fatty acids have a pronounced impact on survival following ischemic renal injury and formation of renoprotective docosahexaenoic acid-derived protectin D1. *J. Immunol.*, 182: 3223–3232.

66. Gonzalez-Perez, A., R. Orrillo, N. Ferré, K. Gronert, B. Dong, E. Morán-Salvador, E. Titos, M. Martínez-Clemente, M. López-Parrera, V. Arroyo, et al. 2009. Obesity-induced insulin resistance and hepatic steatosis are alleviated by omega-3 fatty acids: A role for re-

67. Chen, P., B. Fenet, S. Michaud, N. Tomczyk, E. Vérecel, M. Lagarde, and M. Guichardant. 2009. Full characterization of PDX, a neuro-

2030 Journal of Lipid Research Volume 51, 2010
82. Bazan, N. G. 2003. Synaptic lipid signaling: significance of polyunsaturated acids and phospholipids in experimental models of retinal degeneration. Invest. Ophthalmol. Vis. Sci. 39: 592–602.

83. Aveldano, M. I. and N. G. Bazan. 1974. Displacement into incubation medium by albumin of highly unsaturated retina free fatty acids arising from membrane lipids. FEBS Lett. 40: 53–56.

84. Aveldano, M. I. and N. G. Bazan. 1975. Differential lipid deacylation and reacylation of interactions between the anti-apoptotic protein BAG-1 and Xie, S, Takayama, J, C. Reed, and K. R. Ely. 1998. Characterization of interactions between the anti-apoptotic protein BAG-1 and Hsc70 molecular chaperones. J. Biol. Chem. 273: 22506–22514.

85. Aveldano, M. I., and N. G. Bazan. 1997. A cell culture medium that supports the differentiation of human retinal pigment epithelium into functionally polarized monolayers. Mol. Vis. 7: 14–19.

86. Bhutto, I. A., D. S. McLeod, T. Hasegawa, S. Y. Kim, C. Merges, P. Tong, and A. G. Lutty. 2006. Pigment epithelium-derived factor (PEDF) and vascular endothelial growth factor (VEGF) in aged human choroid and eyes with age-related macular degeneration. Exp. Eye Res. 82: 341–353.

87. Belvisi, M. G., D. A. Shifrin, A. M. Corse, S. R. Bilak, and R. W. Kuncel. 1999. Neuroprotective utility and neurotrophic action of neurturin in photoreceptor outer segment retinal motor neurons: comparison with GDNF and persephin. Mol. Cell. Neurosci. 13: 326–336.

88. Bok, J., D. V. Vauhkonen, C. H. Hecquet, Y. Courtous, and F. Mascalari. 1999. Both FGFR1 and bcl-x synthesis are necessary for the reduction of apoptosis in retinal pigmented epithelial cells by FGFR2: role of the extracellular signal-regulated kinase 2. Oncogene. 18: 7584–7593.

89. Brash, A. R., W. E. Boecklin, and M. S. Chang. 1997. Discovery of a second 15S-lipoperoxidase in humans. Proc. Natl. Acad. Sci. USA. 94: 6148–6152.

90. Brykaert, M., X. Guillonneau, C. Hequet, Y. Courtous, and F. Mascalari. 1999. Both FGFR1 and bcl-x synthesis are necessary for the reduction of apoptosis in retinal pigmented epithelial cells by FGFR2: role of the extracellular signal-regulated kinase 2. Oncogene. 18: 7584–7593.

91. Brash, A. R., W. E. Boecklin, and M. S. Chang. 1997. Discovery of a second 15S-lipoperoxidase in humans. Proc. Natl. Acad. Sci. USA. 94: 6148–6152.

92. Calandria, J. M., V. L. Marcheselli, P. K. Mukherjee, J. Uddin, J. W. Winkler, N. A. Petasis, and N. G. Bazan. 2000. Selective survival rescue in 15-lipoxygenase-1 deficient retinal pigment epithelial cells by the novel docosahexaenoic acid-derived mediator, neuroprotectin D1. J. Biol. Chem. 275: 17877–17882.

93. Kühn, H., B. J. Thiele, A. Osterace-Lederer, H. Stender, H. Suzuki, T. Yoshimoto, and S. Yamamoto. 1993. Bacterial expression, purification, and functional characterization of recombinant rabbit reticuloctye 15-lipoxygenase. Biochim. Biophys. Acta. 1168: 75–78.

94. Sheets, K. G., Y. Zhou, M. E. Fertel, E. J. Knott, C. E. Regan, Jr., J. R. Ellison, W. C. Gordon, P. Gjorstrup, and N. G. Bazan. 2010. Neuroprotectin D1 attenuates laser-induced choroidal neovascularization in mice. Mol. Vis. 16: 320–329.