

Corrections

DEVELOPMENTAL BIOLOGY. For the article “Genetic approaches identify adult pituitary stem cells,” by Anatoli S. Gleiberman, Tatyana Michurina, Juan M. Encinas, Jose L. Roig, Peter Krasnov, Francesca Balordi, Gord Fishell, Michael G. Rosenfeld, and Grigori Enikolopov, which appeared in issue 17, April 29, 2008, of *Proc Natl Acad Sci USA* (105:6332–6337; first published April 24, 2008; 10.1073/pnas.0801644105), the authors note the following: “We regret that we did not insert a reference to the important manuscript by Fauquier *et al.* [Fauquier T, Rizzoti K, Dattani M, Lovell-Badge R, Robinson ICAF (2008) *Proc Natl Acad Sci USA* 105:2907–2912], which published while our manuscript was under review, into the final galley revision of our manuscript. Fauquier *et al.* demonstrate that adult pituitary contains Sox2-positive cells that, upon cultivation *in vitro*, can generate differentiated pituitary lineages.”

www.pnas.org/cgi/doi/10.1073/pnas.0805200105

MEDICAL SCIENCES. For the article “Next generation of adeno-associated virus 2 vectors: Point mutations in tyrosines lead to high-efficiency transduction at lower doses,” by Li Zhong, Baozheng Li, Cathryn S. Mah, Lakshmanan Govindasamy, Mavis Agbandje-McKenna, Mario Cooper, Roland W. Herzog, Irene Zolotukhin, Kenneth H. Warrington, Jr., Kirsten A. Weigel-Van Aken, Jacqueline A. Hobbs, Sergei Zolotukhin, Nicholas Muzyczka, and Arun Srivastava, which appeared in issue 22, June 3, 2008, of *Proc Natl Acad Sci USA* (105:7827–7832; first published May 29, 2008; 10.1073/pnas.0802866105), the authors note that on page 7831, right column, in *Isolation of Nuclear and Cytoplasmic Fractions from HeLa Cells*, line 3, **V444F** should appear as **Y444F**. This error does not affect the conclusions of the article.

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MEDICAL SCIENCES. For the article “The lipofuscin fluorophore A2E perturbs cholesterol metabolism in retinal pigment epithelial cells,” by Aparna Lakkaraju, Silvia C. Finnemann, and Enrique Rodriguez-Boulan, which appeared in issue 26, June 26, 2007, of *Proc Natl Acad Sci USA* (104:11026–11031; first published June 19, 2007; 10.1073/pnas.0702504104), the authors note that the following grant was inadvertently omitted from the Acknowledgments: American Health Assistance Foundation Macular Degeneration Research Grant M2006-081 (to E.R.-B.).

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NEUROSCIENCE. For the article “Activity of human hippocampal and amygdala neurons during retrieval of declarative memories,” by Ueli Rutishauser, Erin M. Schuman, and Adam N. Mamelak, which appeared in issue 1, January 8, 2008, of *Proc Natl Acad Sci USA* (105:329–334; first published December 27, 2007; 10.1073/pnas.0706015105), the authors note that in Fig. 2, panels *E* and *F* were transposed. The corrected figure and its legend appear below.

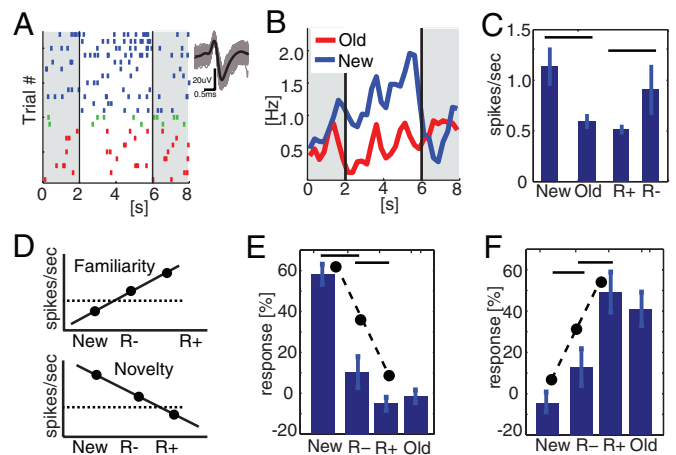


Fig. 2. Single-cell response. (A–C) Firing of a unit in the right hippocampus that increases its firing in response to new stimuli that were correctly recognized (novelty detector). (A) Raster of all trials during retrieval and the waveforms associated with every spike. Trials: new (blue), old and recollected (red, R⁺), and old and not recollected (green, R⁻). (B) Poststimulus time histogram. (C) Mean number of spikes after stimulus onset. Firing was significantly larger in response to new stimuli, and the neuron fired more spikes in response to stimuli that were later not recollected compared with stimuli that were recollected. (D) The hypothesis: The less that novelty neurons fire, the more likely it is that a stimulus will be recollected. The more that familiarity-detecting neurons fire, the more likely it is that a stimulus will be recollected. The dashed line indicates the baseline. (E and F) Normalized firing rate (baseline = 0) of all novelty (E) and familiarity-detecting (F) neurons during above-chance sessions (30-min R⁺). Novelty neurons fired more in response to not recollected items (R⁻), whereas familiarity neurons fired more in response to recollected items (R⁺). Errors are \pm SEM. Number of trials, from left to right: 388, 79, 259, and 338 (E) and 132, 31, 96, and 127 (F).

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The lipofuscin fluorophore A2E perturbs cholesterol metabolism in retinal pigment epithelial cells

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Proteins involved in cholesterol trafficking are known to contribute to the pathogenesis of atherosclerosis and Alzheimer's disease. Allelic variants in the cholesterol transporters apolipoprotein E and ATP-binding cassette protein A1 (ABCA1) have recently been associated with susceptibility to age-related macular degeneration (AMD). Histopathological analyses of eyes with AMD demonstrate the presence of cholesterol and cholesteryl ester deposits beneath the retinal pigment epithelium (RPE), implicating abnormal cholesterol trafficking in disease progression. Here, we show that A2E, a quaternary amine and retinoid by-product of the visual cycle, causes the accumulation of free and esterified cholesterol in RPE cells. The mechanism involves neither generalized alterations in late endosomal/lysosomal pH nor a direct inhibition of acid lipase activity. Rather, A2E prevents cholesterol efflux from these organelles, which in turn indirectly inhibits acid lipase, leading to a subsequent accumulation of cholesteryl esters. Transcriptional activation of the ABCA1 cholesterol transporter by agonists of the liver X receptor/peroxisome proliferator-activated receptor pathway relieves the A2E-induced block on cholesterol efflux and restores cholesterol homeostasis in RPE cells. Our data also demonstrate that A2E, which is a cone-shaped lipid, increases the chemical activity and displacement of cholesterol from model membranes, providing a biophysical mechanism for cholesterol sequestration in A2E-loaded cells. Although endogenously produced A2E in the RPE has been associated with macular degeneration, the precise mechanisms are unclear. Our results provide direct evidence that A2E causes aberrant cholesterol metabolism in RPE cells which could likely contribute to AMD progression.

drusen | lysosome | macular degeneration | phagocytosis | retina

Cholesterol modulates the biophysical properties of membranes and participates in lipid and protein trafficking. In the vertebrate retina, cholesterol stabilizes rhodopsin in photoreceptor outer segment (OS) disks and slows visual cycle kinetics (1). Photoreceptors continuously renew their OS by shedding distal disks in a circadian rhythm (2). Daily phagocytosis and processing of shed photoreceptor OS places a heavy metabolic burden on the underlying postmitotic retinal pigment epithelium (RPE) cells. Over time, incomplete digestion of internalized OS disks leads to the accumulation of autofluorescent lipid-protein aggregates called lipofuscin in RPE lysosomes. A major fluorophore of RPE lipofuscin is a hydrophobic quaternary amine called A2E (Fig. 1A), formed as a bis-retinal/ethanolamine Schiff base adduct (3, 4). A2E is a minor by-product of the visual cycle and progressively accumulates in RPE late endosomes/lysosomes (LE/Ly) [supporting information (SI) Fig. 7] throughout life (4, 5). With age, the RPE becomes less efficient in coping with the increasing burden of lysosomal A2E and its attendant toxicity. A causal role for A2E in RPE and photoreceptor damage was delineated in the *abcr*^{-/-} mouse model of Stargardt's macular degeneration, which shows accelerated A2E deposition in the RPE and subsequent vision loss (6, 7). A2E also accumulates in other animal models of inherited retinal degenerations (4).

Several hypotheses (reviewed in ref. 3) have been proposed to explain how A2E might damage the RPE, including lysosomal

dysfunction and blue-light mediated toxicity. We have previously demonstrated that A2E specifically delays the metabolism of phagocytosed OS lipids by the RPE, whereas lysosomal proteolysis is unaffected (8). Progressive lipid and cholesterol accumulation in the Bruch's membrane beneath the RPE has been identified as a contributing factor to age-related macular degeneration (AMD) (5). Apolipoproteins, cholesterol and cholesteryl esters (CE) are abundant in both drusen (lipid-protein aggregates in the Bruch's membrane) and other subRPE deposits in AMD eyes (9–11), suggesting compromised cholesterol homeostasis in the RPE. Moreover, cardiovascular disease and allelic variants in apolipoprotein E and ATP-binding cassette protein A1 (ABCA1) have been linked to AMD susceptibility in humans (12, 13).

However, little is known about factors that regulate cholesterol levels in the RPE and how these are altered in pathological conditions. Major subcellular sites of cholesterol metabolism are the endoplasmic reticulum, where it is synthesized, and LE/Ly, where CE internalized as part of lipoproteins or OS are degraded by acid lipase and free cholesterol is effluxed out by specific transporters. The objective of the present study was to investigate whether A2E interferes with cholesterol metabolism in the RPE and to identify the mechanism by which this might occur. We show that A2E, at levels similar to those found in aging human eyes, interferes with cholesterol efflux from LE/Ly, which in turn inhibits acid lipase and results in a feed-forward cycle of cholesterol and CE accumulation in RPE cells.

Results

A2E Causes Endogenous Cholesterol Accumulation in RPE Cells. To ensure the physiological relevance of our experiments, we established extracellular A2E concentrations required for RPE cells to accumulate A2E in the range found *in vivo* (14). In ARPE-19 cells, a single 15- μ M dose or three 5- μ M doses over 7 days resulted in intracellular A2E amounts in this range (SI Table 2). Thus, A2E accumulation is an additive process, and RPE cells lack the capacity to either degrade or efflux it out. Cells exposed to A2E had increased levels of both free cholesterol and CE compared with controls, whereas phospholipids and sphingomyelin were unchanged (Fig. 1B). Analysis of intracellular cholesterol distribution using the sterol-binding probe filipin showed a dramatic increase in

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The authors declare no conflict of interest.

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Abbreviations: AMD, age-related macular degeneration; RPE, retinal pigment epithelium; ABCA1, ATP-binding cassette protein A1; CE, cholesteryl esters; Bodipy-CE, Bodipy-labeled cholesteryl dodecanoate; LE/Ly, late endosomes/lysosomes; LXR, liver X receptor; OS, outer segment; PPAR, peroxisome proliferator-activated receptor; TFMUO, trifluoromethyl umbelliferyl oleate; DOTAP, dioleoyltrimethyl ammonium propane; LysoPC, lysophosphatidylcholine.

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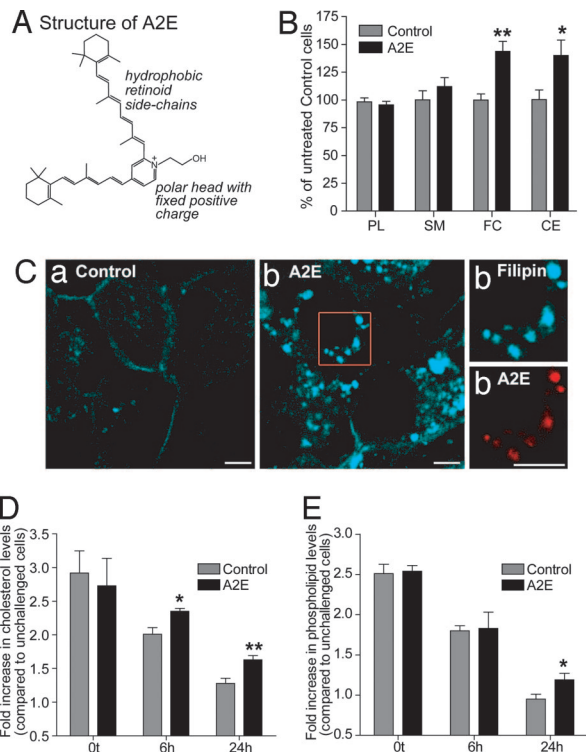


Fig. 1. A2E and RPE cholesterol and lipid levels. (A) A2E is a cone-shaped molecule with a small polar headgroup and large hydrophobic retinoid side chains. (B) Quantification of total cellular phospholipids (PL), sphingomyelin (SM), free cholesterol (FC), and CE 48 h after A2E loading in ARPE-19 cells. Significantly different from controls, *, $P < 0.05$; **, $P < 0.001$. (C) Filipin staining (blue) of intracellular cholesterol pools in control (a) and A2E-loaded (b) (red) cells. (Right) High-magnification images of area denoted by box in b. (Scale bar, 5 μm .) (D) Total cellular cholesterol levels after OS phagocytosis normalized to cholesterol levels in cells not fed OS. Significantly different from controls at corresponding time, *, $P < 0.05$; **, $P < 0.001$. (E) Total cellular phospholipid levels after OS phagocytosis normalized to phospholipid levels in cells not fed OS. Significantly different from controls at 24 h, *, $P < 0.05$.

filipin fluorescence in A2E-loaded cells (Fig. 1C). Brightly fluorescent filipin puncta colocalized with A2E, indicative of enlarged LE/Ly cholesterol pools. We observed a 1.56-fold increase in free cholesterol levels in the presence of A2E (Fig. 1B). Because LE/Ly contain $\approx 10\%$ of total cell cholesterol (15) and assuming that $\geq 50\%$ of the A2E-induced cholesterol accumulation occurs in LE/Ly, this translates to a 4- to 8-fold increase in the free cholesterol content of LE/Ly (see *SI Text* for detailed calculations).

To test whether A2E disrupts lipid metabolism after phagocytosis, we measured phospholipid and cholesterol levels after feeding bovine OS to ARPE-19 cells with or without A2E. We have previously shown that A2E affects neither the binding nor internalization of OS by RPE cells (8). Immediately after phagocytosis, total cholesterol levels in both control and A2E-loaded cells increased 3-fold relative to cells that were not fed OS. At 6 and 24 h after phagocytosis, there was significantly more cholesterol in cells with A2E compared with controls (Fig. 1D). We also observed an increase in total phospholipids in A2E-loaded cells 24 h after phagocytosis (Fig. 1E), in agreement with our published data that A2E delays the clearance of phagocytosed lipids (8).

A2E Does Not Alter LE/Ly pH. It has been suggested that A2E increases lysosomal pH and thus inactivates lysosomal enzymes (16). Because acid lipase has a pH optimum of 4–5, an increase in lysosomal pH would lead to inhibition of enzyme activity and an accumulation of both CE and cholesterol (17). However, Holz *et al.*

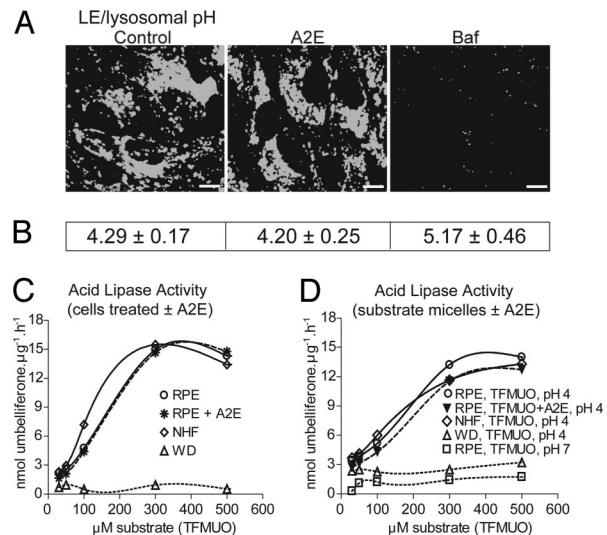


Fig. 2. LE/Ly pH and acid lipase activity are unaffected by A2E. (A) Qualitative estimate of LE/Ly pH with LysoSensor blue in ARPE-19 cells. Baf, bafilomycin A1. (Scale bar, 10 μm .) (B) Quantification of LE/Ly pH using LysoSensor yellow/blue. (C) Acid lipase activity using soluble protein extracts from either control or A2E-loaded cells as the enzyme source and TFMUO micelles as the substrate. (D) Acid lipase activity in soluble protein extracts from untreated cells as the enzyme source and TFMUO micelles with or without A2E as the substrate.

(16) did not quantify the intracellular concentration of A2E, which might have accumulated at nonphysiological levels. We used two independent assays to determine whether A2E, at physiological levels, increases LE/Ly pH. First, a qualitative estimate of LE/Ly pH using LysoSensor blue showed no difference in probe fluorescence between control and A2E-loaded ARPE-19 cells, whereas a short exposure to bafilomycin A1 decreased fluorescence, signifying an increase in pH (Fig. 2A). Second, quantification of LE/Ly pH with the ratiometric dye LysoSensor yellow/blue gave similar values for both control and A2E-laden cells (Fig. 2B), whereas treatment with bafilomycin A1 increased the pH by 1 unit. That A2E, at intracellular levels similar to that found *in vivo*, does not alter LE/Ly pH is in accordance with its fixed positive charge and lack of exchangeable protons (Fig. 1A) and supports *in vitro* data showing that A2E does not inhibit the activity of several lysosomal enzymes (3).

Acid Lipase Activity Is Not Directly Inhibited by A2E. We next tested the possibility that A2E might promote CE and cholesterol accumulation by directly inhibiting acid lipase activity (17). Acid lipase is a soluble LE/Ly enzyme that transiently associates with membranes to hydrolyze ester bonds at the lipid–water interface. Upon partitioning into LE/Ly membranes, lipophilic A2E could interfere with acid lipase activity by directly interacting with the enzyme’s catalytic domain or by preventing enzyme–substrate interaction via steric hindrance. To test this hypothesis, we used the fluorogenic substrate trifluoromethyl umbelliferyl oleate (TFMUO) to measure acid lipase activity in d407 protein lysates (18). Wolman’s disease fibroblasts that lack acid lipase activity and age-matched normal human fibroblasts were used as controls. Lack of substrate hydrolysis at pH 7 validated the assay. A2E did not affect the expression or degradation of acid lipase because 10 μg of soluble protein from either vehicle- or A2E-treated cells hydrolyzed TFMUO with similar kinetics (Fig. 2C). Further, A2E did not interfere with enzyme–substrate interactions, because there was no change in the hydrolysis rate when substrate micelles contained A2E (Fig. 2D). Thus, CE accumulation in RPE cells is not because of a direct inhibition of acid lipase by A2E.

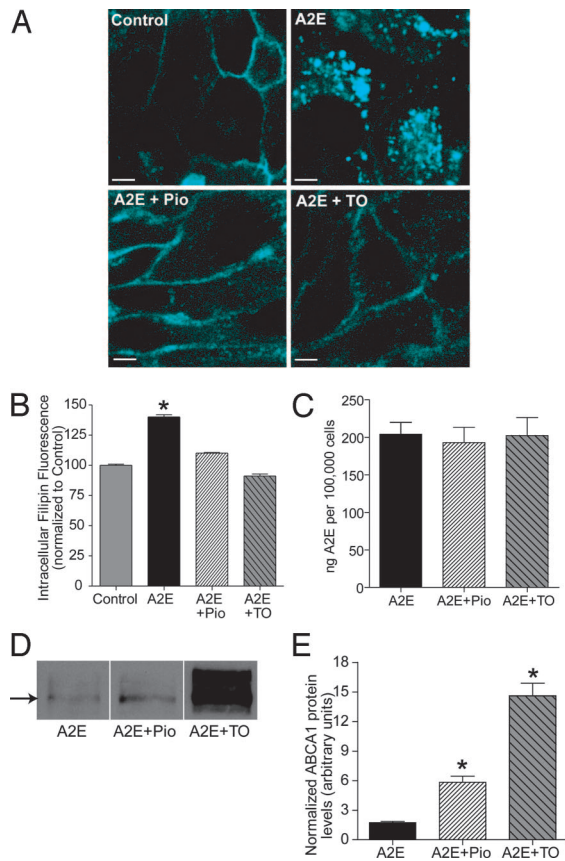


Fig. 4. LXR/PPAR agonists restore cholesterol homeostasis in A2E-laden cells. (A) Filipin staining of endogenous cholesterol in polarized ARPE-19 cells treated with vehicle (Control), A2E, A2E and pioglitazone (A2E+Pio), or A2E and the LXR agonist TO901317 (A2E+TO). (B) Quantification of intracellular filipin fluorescence. Data were normalized to vehicle-treated control cells whose intracellular fluorescence is set to 100%. Significantly different from all other conditions, *, $P < 0.01$. For Control, A2E and A2E+Pio, $n \geq 200$; for A2E+TO, $n = 80$. (C) Quantification of A2E levels after treatment with either Pio or TO compared with A2E alone. (D) Representative ABCA1 immunoblot of A2E-loaded cells treated with either Pio or TO. (E) Quantification of ABCA1 protein levels normalized to β -actin; *, significantly different from cells exposed to A2E alone, $P < 0.05$.

membrane architecture. Liposomes with A2E also increased cholesterol transfer to methyl- β -cyclodextrin (Fig. 6B), providing additional proof that A2E displaces membrane cholesterol and thus disturbs bilayer cholesterol organization.

Discussion

Phagocytosis and degradation of shed OS is an indispensable function performed by the RPE to maintain photoreceptor health and vision. Several lines of evidence suggest that OS phagocytosis is tightly coupled to RPE lipid and cholesterol metabolism: (i) Phagocytosis triggers expression of the AP-2 transcription factor (25), which regulates acid lipase expression (26); (ii) OS phagocytosis induces PPAR γ (25), which increases transcription of cholesterol transporters (21); and (iii) selective metabolism of OS lipids by the RPE leads to the conservation and recycling of docosahexaenoic acid (27). In addition, RPE cells express several receptors and cholesterol transporters (28) that participate in lipoprotein uptake and cholesterol efflux.

Because OS degradation takes place mainly in LE/Ly, and because our published data indicate that the lipofuscin fluorophore A2E in RPE LE/Ly interferes with the clearance of OS lipids (8), we investigated the effect of A2E on endogenous cholesterol and

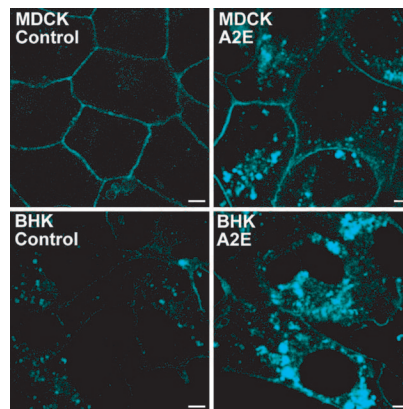


Fig. 5. Filipin staining of endogenous cholesterol in control or A2E-laden polarized MDCK and BHK-21 cells. (Scale bar, 5 μ m.)

lipid metabolism in the RPE. Our results show that A2E, at levels similar to those found *in vivo*, specifically causes cholesterol and CE accumulation in RPE cells. We did not observe an increase in LE/Ly pH in A2E-loaded cells, which is expected, because A2E is a quaternary amine that lacks exchangeable protons (Fig. 1A). Unlike classical lysosomotropic amines, the accumulation of A2E in LE/Ly is not because of “proton trapping” (29). As protons are neither consumed nor leaked out because of passive diffusion of the protonated base in A2E-loaded cells, there is no driving force for alkalinization. Moreover, a nonspecific increase in LE/Ly pH would lead to widespread inhibition of lysosomal enzymes, which has not been observed either *in vitro* or *in vivo* in the presence of A2E (3).

Using model lipid bilayers, we found that cone-shaped A2E competes with cholesterol for space under the phospholipid umbrella (22), displacing it from membranes. To undergo efflux from LE/Ly, cholesterol located in late endosomal internal membranes needs to be shuttled to the limiting membrane by nonvesicular mechanisms (30). Our data show that in RPE cells, cholesterol accumulates in the same organelles as A2E, indicating a block in LE/Ly cholesterol efflux. It is conceivable that once cholesterol displaced from A2E-containing membranes reaches a critical concentration, it is excluded from the pool destined for nonvesicular transport to the limiting membrane and thus gets trapped within the LE/Ly. Increased LE/Ly cholesterol levels result in an indirect inhibition of RPE acid lipase, in agreement with previous studies showing that late endosomal cholesterol efflux is necessary for acid lipase activity (19). The subsequent accumulation of CE along with free cholesterol generates a feed-forward cycle of further cholesterol accumulation, inhibition of acid lipase, and increased CE in the cell.

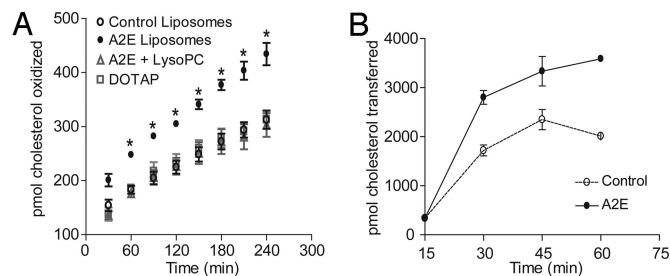


Fig. 6. A2E accelerates cholesterol displacement from lipid bilayers. (A) Influence of A2E on the accessibility of bilayer cholesterol to cholesterol oxidase. Liposomes (20 mol% cholesterol and 0.04 mol% of A2E, LysoPC or DOTAP) were incubated with cholesterol oxidase for the indicated times at 37°C. *, significantly different from all other liposomes at the corresponding time point; $P < 0.01$. (B) Kinetics of cholesterol transfer from control or A2E liposomes to cyclodextrin.

Table 1. Cholesterol dissociation rates from lipid bilayers

Liposome	K_{off} , h^{-1}	$t_{1/2}$, h	Percent cholesterol oxidized
Control	0.0134 ± 0.0003	51.66	1.95 ± 0.20
A2E	0.0222 ± 0.0003	31.20	3.98 ± 0.86
Control + LysoPC	0.0109 ± 0.0004	63.40	1.80 ± 0.39
A2E + LysoPC	0.0116 ± 0.0004	59.65	1.89 ± 0.33
DOTAP	0.0120 ± 0.0006	57.69	1.96 ± 0.21

Cholesterol efflux from LE/Ly to the plasma membrane is preferentially mediated by ABCA1 (31). Our data show that LRX/PPAR agonists induce ABCA1 expression, decrease cholesterol accumulation, and restore CE metabolism in RPE cells. Cholesterol accumulation in LE/Ly may also interfere with other degradative functions, such as the processing of OS lipids, as previously demonstrated by us (8) and confirmed in this study. Indeed, increased membrane cholesterol has been found to inhibit lysosomal phospholipase A1 (32).

The presence of cholesterol and CE deposits in the Bruch's membrane of AMD eyes (9, 10) and the correlation of genetic variations in cholesterol transporters with AMD susceptibility (12, 13) point to an age-related alteration in cholesterol homeostasis in the RPE that may contribute to AMD progression. In this context, it is noteworthy that cholesterol and CE deposits in the Bruch's membrane are thought to originate from the RPE, including incompletely processed OS and additional sources such as the RPE itself (9). Because our data demonstrate that A2E causes a >4-fold increase in LE/Ly free cholesterol (apart from CE accumulation), and because membranous debris beneath the RPE appear to originate as blebs and/or exocytosed organelles from the RPE basal surface (33, 34), it is tempting to speculate that the source of subRPE cholesterol/CE deposits could be LE/Ly internal membranes extruded from A2E-laden cells.

Our study provides evidence in support of the hypothesis that loss of cholesterol homeostasis in the RPE may play an important role in the pathogenesis of retinal dystrophy. Biochemical and histological analyses of cholesterol and phospholipid homeostasis in *abcr*^{-/-} mice would be useful in providing an *in vivo* correlate for our observations on A2E-induced cholesterol accumulation in RPE cells. However, these experiments are beyond the scope of the present study, because the main objective here was to identify the mechanism by which A2E can derail intracellular cholesterol trafficking pathways. At the cellular level, impaired cholesterol efflux from RPE LE/Ly may interfere with rhodopsin activation and photoreceptor membrane remodeling. Excess cholesterol, either within the RPE or extruded into the Bruch's membrane, could function as an atypical chronic inflammatory stimulus (35) that contributes to AMD. Finally, strategies that increase cholesterol efflux from the RPE may be a promising therapeutic approach in circumventing disease progression.

Materials and Methods

Cell Culture. Human ARPE-19 cells [American Type Culture Collection (ATCC), Bethesda, MD] were cultured in RPMI medium 1640 + 1% FBS for ≥ 6 weeks to establish polarity. Human d407 cells (a gift of R. C. Hunt, University of South Carolina, Columbia, SC) were grown in DMEM + 3% FBS for 48 h before A2E loading. Primary bovine RPE cells were isolated from freshly obtained eyes and cultured on laminin-coated dishes in DMEM/1% FBS (ATCC). Cells were plated at a density of 40,000 cells per cm^2 on 24-well plates for biochemical experiments. For confocal microscopy, cells were plated on laminin-coated Transwell filters (BD Biosciences, San Jose, CA) for ARPE-19 and bovine RPE or alcian blue-coated coverslips for d407 cells. Each experiment was performed in at least two different RPE cell types. Wolman's

disease fibroblasts and age-matched normal human fibroblasts (Coriell Cell Repositories, Camden, NJ) were cultured in DMEM/15% FBS. Madin–Darby canine kidney Type II cells were cultured on Transwell filters in DMEM/5% FBS, and BHK-21 cells were cultured on coverslips in alpha minimum essential medium (AMEM)/10% FBS.

A2E Synthesis and Loading. A2E was synthesized according to a published protocol (36) and purified by HPLC (>97%, electrospray ionization–MS). Cells were exposed to A2E (50 μM for d407 cells and 15 μM for ARPE-19 and bovine RPE cells) for 6 h in growth medium and chased overnight in fresh medium.

Lipid and Sterol Quantification. Cellular lipids were extracted in chloroform/methanol, and total lipid phosphorous was determined by a colorimetric assay after acid digestion (37). Sphingomyelin content was determined by the Amplex Red Sphingomyelinase assay (Molecular Probes, Eugene, OR). Samples were incubated with 100 μM Amplex red reagent, 2 units/ml HRP, 0.2 units/ml choline oxidase, 8 units/ml alkaline phosphatase, and 1 milliunit neutral sphingomyelinase. Free cholesterol and CE were quantified by the Amplex Red Cholesterol assay (Molecular Probes).

OS Phagocytosis. ARPE-19 cells were fed bovine OS for 4 h (8) and chased in fresh medium. After lipid extraction, total phospholipids and free cholesterol were quantified as above.

Bodipy-CE Hydrolysis. Vesicles with egg phosphatidylcholine, brain phosphatidylserine, cholesterol (Avanti Polar Lipids, Alabaster, AL), and Bodipy-CE (Molecular Probes) (molar ratio 42:42:15:0.5) were prepared by sonicating on ice for 10 3-min cycles (Branson Sonifier 350; Branson Ultrasonics, Danbury, CT). Cells were incubated with Bodipy-CE vesicles (45 min, 37°C) and chased in fresh medium. Undegraded Bodipy-CE was quantified by TLC in a mobile phase of hexane/isopropanol/acetic acid (93:6:1). Cells were treated with an acyl CoA: cholesterol acyl transferase inhibitor FR179254 (400 nM; Calbiochem, San Diego, CA) for a true estimate of Bodipy-CE accumulation. Other drugs used were U18666A (500 nM; Calbiochem), pioglitazone (10 μM ; Calbiochem), and TO901317 (1 μM ; Cayman, Ann Arbor, MI).

Filipin Staining and Quantification. Paraformaldehyde-fixed cells were stained with 50 $\mu g/ml$ filipin for 45 min followed by confocal microscopy. Intracellular filipin fluorescence was quantified by using Leica (Leica Microsystems, Wetzlar, Germany) software. For each cell, a region of interest (ROI) was drawn just within the plasma membrane, and the fluorescence intensity per pixel of the ROI per plane of the confocal image stack was quantified.

Late Endosomal/Lysosomal pH. Control or A2E-loaded cells were incubated with 7.5 μM LysoSensor blue (Molecular Probes, 5 min, 37°C), rinsed to remove excess dye, and imaged immediately. For quantitative determination of pH, cells were trypsinized, resuspended in recording medium (20 mM Hepes/4.5 g/liter glucose/1% FBS in Hanks' balanced salt solution) and incubated with 7.5 μM LysoSensor yellow/blue, as above. After centrifugation, cells were resuspended in Mes buffer, pH 7.0, and the fluorescence ratio at 436 and 525 nm was measured (λ_{ex} 380 nm). Calibration curves were established by using Mes buffers of defined pH values, and nigericin and monensin were used to dissipate intracellular pH gradients. Measured ratios for known pH values were fitted to a Boltzmann equation, and LE/Ly pH of control and A2E-loaded cells was calculated. Bafilomycin A1 (50 nM; Sigma, St. Louis, MO) was used as a positive control.

Acid Lipase Activity. Substrate micelles with 2 mM TFMUO (Sigma) and 3.25 mM egg phosphatidylcholine (EPC) were made in 2.5 ml of 2.4 mM sodium taurocholate by sonicating on ice for 5 min (18).

Under these conditions, the assay was linear up to 30 $\mu\text{g}/\text{ml}$ protein for ≥ 2 h at 37°C. A2E-containing micelles were prepared by adding 2.6 μg of A2E per milligram of lipid to the TFMUO/EPC mixture. The final assay solution contained 0–500 μM TFMUO, 150 μl of 0.2 M acetate buffer (pH 4), 10 μg of soluble protein extract from RPE cells, or Wolman's disease fibroblasts or normal human fibroblasts, in a total volume of 600 μl . After 1 h at 37°C, umbelliferone released upon hydrolysis of TFMUO was measured at 502 nm (λ_{ex} 385 nm). For experiments on cholesterol content and acid lipase activity, TFMUO micelles were prepared as above with increasing amounts of cholesterol (keeping TFMUO concentration in the micelles constant), and the assay was performed with 300 μM substrate.

Confocal Microscopy. Cells were imaged on a Leica TCS-SP2 laser-scanning confocal microscope using the 63×1.4 N.A. oil immersion objective. The argon/argon–krypton laser was used for Bodipy-CE, and the green helium–neon laser was used for A2E. In experiments with filipin, the 405 laser was used to excite both filipin and A2E, because A2E has an excitation peak at 418 nm, and images were captured at different emission windows (420–440 for filipin and 590–610 for A2E). For each set of experiments, the laser power, voltage, and offset were identical for a given fluorophore. All images were acquired with Leica software and processed identically for control and treated conditions.

Immunoblotting. Whole-cell lysates were prepared in buffer containing 1% Triton X-100, and proteins (≈ 50 μg) were electrophoresed on 7.5% SDS-polyacrylamide gels and transferred to nitrocellulose membranes. Blots were probed with rabbit polyclonal antibodies to either ABCA1 (Abcam, Cambridge, MA; 1:1,000) or actin (Santa Cruz Biotechnology, Santa Cruz, CA; 1:1,000). After chemiluminescence development, bands were quantified by using ImageJ (NIH, Bethesda, MD), and ABCA1 levels were normalized to that of β -actin.

A2E Quantitation. Cells were treated with A2E and agonists as above. Lipids were harvested, and A2E levels were quantified by TLC by using a mobile phase of chloroform/methanol/water (84:15:1, vol/vol).

Liposomes. Liposomes with a lipid composition similar to that of late endosomes (38) (dioleoylphosphatidylcholine, dioleoylphosphatidylethanolamine, sphingomyelin, phosphatidylserine, phosphati-

dylinositol, and cholesterol; 50:20:9:2:4:20 mol%) were prepared by extrusion in 10 mM Hepes/150 mM NaCl buffer (7.5 μmol of total lipid per milliliter). When required, A2E, lysophosphatidylcholine, or dioleoyltrimethyl ammonium propane (0.04 mol%) was added to the lipid mixture. Based on our data, ≈ 200 ng of A2E per milligram of total lipid is present in RPE cells; of total cellular lipid, ≈ 67 μg is present in LE/Ly (15). Because A2E is predominantly present in LE/Ly, this amounts to 200 ng of A2E/67 μg of total lipid or 0.04 mol%.

Cholesterol Oxidase Assay. Liposomes were diluted to 750 μM in buffer (0.5 M potassium phosphate/0.25 M NaCl, pH 7.4) and incubated with 2 units/ml HRP, 2 units/ml cholesterol oxidase, and 300 μM Amplex red reagent at 37°C. Fluorescence was measured at 590 nm (λ_{ex} 540 nm) on a Spectramax Gemini microplate reader (Molecular Devices, Sunnyvale, CA). Dissociation rates were calculated by fitting kinetic data to a first-order exponential equation (Prism; GraphPad, San Diego, CA).

Cholesterol Transfer to Cyclodextrin. Control or A2E-containing liposomes (375 μM) were incubated with 2.5 mM methyl- β -cyclodextrin at 37°C. Samples were filtered through Microcon YM-30 membranes (Millipore, Billerica, MA, 1,300 \times g, 15 min) to separate liposomes from cyclodextrin–cholesterol complexes (39). The amount of cholesterol transferred was quantified by the Amplex red cholesterol assay after chloroform/methanol extraction of filtrates and retentates.

Statistical Analysis. Data were analyzed by using either a two-tailed *t* test or one-way ANOVA followed by the Bonferroni multiple comparisons post-test (Prism). Unless otherwise stated, data are presented as mean \pm SEM of three or more independent experiments, with three to four replicates per condition per experiment.

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