

Foam cell-derived 4-hydroxynonenal induces endothelial cell senescence in a TXNIP-dependent manner

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Abstract

Vascular endothelial cell (VEC) senescence is considered an early event in the development of atherosclerotic lesions. Stressful stimuli, in particular oxidative stress, have been linked to premature senescence in the vasculature. Foam cells are a major source of reactive oxygen species and may play a role in the induction of VEC senescence; hence, we investigated their involvement in the induction of VEC senescence in a co-culture transwell system. Primary bovine aortic endothelial cells, exposed to the secretome of THP-1 monocyte-derived foam cells, were analysed for the induction of senescence. Senescence associated β -galactosidase activity and the expression of p16 and p21 were increased, whereas phosphorylated retinoblastoma protein was reduced. This senescent phenotype was mediated by 4-hydroxynonenal (4-HNE), a lipid peroxidation product secreted from foam cells; scavenging of 4-HNE in the co-culture medium blunted this effect. Furthermore, both foam cells and 4-HNE increased the expression of the pro-oxidant thioredoxin-interacting protein (TXNIP). Molecular manipulation of TXNIP expression confirmed its involvement in foam cell-induced senescence. Previous studies showed that peroxisome proliferator-activated receptor (PPAR) δ was activated by 4-hydroalkenals, such as 4-HNE. Pharmacological interventions supported the involvement of the 4-HNE-PPAR δ axis in the induction of TXNIP and VEC senescence. The association of TXNIP with VEC senescence was further supported by immunofluorescent staining of human carotid plaques in which the expression of both TXNIP and p21 was augmented in endothelial cells. Collectively, these findings suggest that foam cell-released 4-HNE activates PPAR δ in VEC, leading to increased TXNIP expression and consequently to senescence.

Keywords: atherosclerosis • senescence • foam cells • VEC • 4-HNE • PPAR δ • TXNIP

Introduction

Atherosclerosis is a chronic inflammatory disease affecting certain areas in the arteries, which are characterized by disturbed blood flow [1–3]. Impaired function of the vascular endothelium precedes the development of atherosclerotic lesions. Whilst this impairment is part of the natural ageing process [4–6], additional

pathophysiological factors promote and accelerate atherosclerosis, among which are hyperglycaemia, dyslipidaemia and hypertension [7–9]. These conditions may induce oxidative stress [10], endoplasmic reticulum (ER) stress [11] and inflammation [12, 13], which have emerged as major detrimental factors that induce endothelial cell dysfunction.

Cellular senescence has been linked to the development of vascular endothelial cell (VEC) dysfunction in atherosclerosis (reviewed in [14, 15]). Senescence, which was initially considered part of the

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ageing process of cultured cells, and termed 'replicative senescence', has been linked to progressive shortening of the telomeres due to successive cell divisions (reviewed in [16]). Senescence could also result from telomere-independent events, in which mitotically competent cells respond to stress, most often oxidative stress, by growth arrest. In contrast with replicative senescence, stress-induced premature senescence (SIPS) is potentially reversible [17]. Senescent cells are characterized by increased expression of the cyclin-dependent kinase inhibitors p21 and p16, and reduced phosphorylated pRb, which lead to cell cycle arrest at the G₀/G₁ checkpoint by inactivating members of the transcriptional factor family E2F [18]. In addition, senescent cells exhibit increased cell size and increased cytoplasmic activity of senescence-associated β -galactosidase (SA- β -Gal) [14, 15]. Senescent VEC were observed in areas of atherosclerotic lesions in the human aorta and coronary arteries, and in animal models of atherosclerosis [14, 19–23]. Cultured VEC were also shown to develop replicative- and stress-induced senescence [24], associated with changes in morphology and gene expression, causing impairment of important cell functions such as endothelium-dependent vasodilation, angiogenesis, vascular healing and increased expression of inflammatory mediators [14, 15].

The oxidized form of LDL-cholesterol (OxLDL) plays a central role in the pathophysiology of atherosclerosis, affecting key stages of this process. At the initial stage, it binds to scavenger receptors on VEC surface and induces the expression of adhesion molecules that facilitate monocyte adhesion and migration to the sub-endothelial layer [25]. OxLDL transverse the endothelial barrier and enters the sub-endothelium, where it transforms monocytes into macrophages, which become lipid-laden foam cells [26]. The latter contribute to the expansion of the lipid core in the atherosclerotic plaque and exacerbate oxidative stress. Indeed, foam cells produce high levels of superoxide, nitric oxide and hydrogen peroxide [27].

Cells exposed to high levels of reactive oxygen species (ROS) often respond by augmenting the generation of 4-hydroxynonenal (4-HNE), the non-enzymatic peroxidation product of n-6 polyunsaturated fatty acids (PUFA) [28–31]. In addition, 4-HNE induces the expression of monocyte chemoattractant proteins [32], alters the cell cycle [33, 34] and promotes apoptosis [35]. Moreover, it may induce ER stress and impair endothelial barrier and coagulation functions [36–38]. Cytotoxic effects of 4-HNE are attributed to its covalent interactions with nucleophilic moieties in proteins, DNA and phospholipids [39, 40]. Excessive accumulation of 4-HNE-modified proteins has been shown to inhibit the proteasome [41]. However, recent studies claim that low non-cytotoxic concentrations of 4-hydroxyalkenals, including 4-HNE and 4-hydroxydodecadienal, function as signalling molecules by activating the nuclear receptor peroxisome proliferator-activated receptor (PPAR) δ [31, 42–44]. Overall, the accumulation of foam cells in the sub-endothelial space may promote atherosclerosis by generating oxidative stress along with accumulation of lipid peroxidation products that affect VEC.

The thioredoxin (TRX) system regulates oxidative stress in various tissues, including the vascular endothelium [45]. This antioxidant system consists of TRX, TRX-reductase, NADPH and the natural TRX

inhibitor, the thioredoxin-interacting protein (TXNIP) [46]. Thioredoxin is ubiquitously expressed in VEC, where it scavenges ROS, prevents apoptosis and promotes angiogenesis [47, 48]. TXNIP impairs the reducing activity of TRX by interacting with the catalytic site of reduced TRX, leading to increased ROS accumulation [49]. Moreover, TXNIP has been shown recently to mediate endothelial cell inflammation in response to disturbed blood flow by increasing monocyte adhesion [50]. Lipid peroxidation products, such as 4-HNE and acrolein, inhibit TRX activity by binding to Cys73, which is placed out of the catalytic domain, thus inducing a conformational change; this has been associated with increased ROS production and enhanced monocyte adhesion to VEC [51]. These findings suggest that the TRX system is important for VEC function and protects against atherosclerosis.

The present study was designed to investigate the effects of foam cells-derived lipid peroxidation products on VEC senescence. Our findings show that 4-HNE produced by foam cells induces senescence in VEC by activating PPAR δ , resulting in increased TXNIP expression.

Materials and methods

Materials

All tissue culture media and reagents were purchased from Biological Industries (Beit Haemek, Israel). Transwell plates and inserts were from Nunc (Roskilde, Denmark). Calbiochem (Darmstadt, Germany) supplied GW501516 and 4-HNE; Sigma-Aldrich (Rehovot, Israel) the anti α -tubulin antibody, GSK0660, Oil red O, phorbol 12-myristate 13-acetate (PMA), WY14643 and troglitazone; Abcam (Cambridge, MA, USA) supplied the senescence-associated β -galactosidase detection kit and the following polyclonal antibodies: rabbit anti-p21, rabbit anti-p16 and anti-pRB (phospho-Ser⁷⁹⁵). Mouse anti-TXNIP was from MBL Ltd (Nagoya, Japan) and horseradish peroxidase-conjugated anti-rabbit IgG from Jackson ImmunoResearch (West Grove, PA, USA). Mirus Bio-Corporation (Madison, WI, USA) supplied the TransIT-LT1 reagent. Promega (Madison, WI, USA) supplied the dual-luciferase reporter assay system. The TXNIP small-interfering RNA (siRNA) was synthesized by GE Dharmacon (Lafayette, CO, USA). Qiagen (Hilden, Germany) supplied the negative (scrambled) control siRNA (allstars negative control). The pcDNA3 and pEGFP-N1 plasmids were kindly provided by Dr. R. Hertz (The Hebrew University Faculty of Medicine, Jerusalem, Israel). The pSVPORT1-hRXR vector and the 3xPPAR response element (PPRE)-TK-luciferase plasmid were courtesy of Dr. B.M. Spiegelman (Dana Farber Cancer Institute, Boston, MA, USA). The pCMX-hPPAR γ 1 and pCMX-hPPAR γ 2 plasmids and the respective empty plasmids were kindly provided by Dr. R. Evans (Howard Hughes Medical Institute, La Jolla, CA, USA). Dr. B. Staels (Institut Pasteur de Lille, Lille, France) kindly provided the pSG5 and pSG5-hPPAR α vectors. The PPAR δ plasmid was prepared by us, as described [44, 52]. The luciferase reporter construct encoding the human TXNIP promoter region and the plasmid encoding for human TXNIP were kindly provided by Dr. Anath Shalev (University of Wisconsin-Madison, Madison, WI, USA). All primers were synthesized by Sigma-Aldrich. FL-926-A16 was kindly provided by Flamma S.p.A (Chignolo d'Isola, Bergamo, Italy). Organic solvents were from Frutarom (Haifa, Israel) and Mal-

linckrodt Baker (Deventer, Holland). All chemicals and reagents for LC-MS/MS were of analytical-grade and purchased from Sigma-Aldrich Chemical Co. (Milan, Italy). High-performance liquid chromatography (HPLC)-grade water was prepared with a Milli-Q water purification system. Optimal cutting temperature (OCT) compound for cryosection was purchased from Bar-Naor Ltd. (Ramat Gan, Israel).

Preparation of oxidized lipoproteins

OxLDL was prepared at the Lipidomics Department, the Nicolae Simionescu Institute of Cellular Biology and Pathology (Bucharest, Romania). Briefly, native LDL was isolated from plasma of healthy donors by ultracentrifugation (Hematology Center, Bucharest, Romania). OxLDL was prepared *in vitro* by incubating nLDL under sterile conditions with 10 μ M copper chloride (24 hrs, at 37°C), in the absence of antioxidant protection. The oxidative reaction was stopped by addition of 1 mg/ml EDTA and after extensive dialysis against PBS, pH 7.4, 4°C, oxLDL was stored at 4°C, under sterile conditions. The copper-OxLDL was characterized as previously described [53].

Cell culture

Bovine aortic endothelial cell cultures

Primary cultures were isolated, cultured and characterized as described previously [54]. The cells were used up to passage 9, unless otherwise indicated.

THP-1 Monocyte

THP-1 cells, purchased from American Type Culture Collection (Rockville, MD, USA), were grown in suspension in complete RPMI 1640 medium according to the supplier's protocol. To induce differentiation to macrophages, the THP-1 monocytes (10^6 /ml) were exposed to 0.1 μ M of PMA for 24 hrs. Foam cell formation was induced by incubating the macrophages with 100 μ g/ml of OxLDL for 72 hrs.

Transwell cultures

Vascular endothelial cell were plated and grown to confluency in 6-well culture plates using complete DMEM. THP-1 cells were transformed into macrophages or foam cells on transwell insert membranes, as described above. The membranes were then washed three times with fresh DMEM and inserted into the wells over the apical side of the VEC monolayers. The co-culture was continued for the indicated periods. Cell viability was determined by Trypan blue exclusion assay.

Senescence-associated β -galactosidase assay

The SA- β -Gal activity was determined in VEC monolayers according to the manufacturer's instructions (Abcam). Briefly, cells were washed, fixed and stained with the X-Gal reagent at pH 6, followed by DAPI staining for visualization of cell nuclei. The percentage of SA- β -Gal positive cells relative to the total DAPI-positive nuclei was assessed by counting an area of 0.55 mm², per images (under $\times 10$ magnifying lens) obtained from three independent experiments using the Nikon confocal microscope software (NIS-Elements AR 4 30.0).

Western blot analysis

Vascular endothelial cell lysates were prepared and used for Western blot analysis. Briefly, cells were rinsed three times with PBS and lysed with NP-40 lysis buffer (Tris HCl 50 mM pH 7.5, NP-40 1%, sodium deoxycholate 0.25%, EGTA 1 mM, EDTA 1 mM, NaCl 150 mM, sodium orthovanadate 1 mM, sodium fluoride 1 mM, sodium β -glycerophosphate 10 mM, sodium pyrophosphate 5 mM, PMSF 1 mM, protease inhibitor cocktail 1%; Sigma). Bradford assay was used to determine protein concentrations in the lysates. Twenty μ g protein samplers were loaded per lane and separated by PAGE and transferred to nitrocellulose membranes. The membranes were blocked with TBST containing 5% (wt/v) bovine serum albumin. For the analysis of TXNIP PBS containing 10% (wt/v) low fat dry milk was used for blocking. Incubations with the various antibodies were according to the suppliers' protocols at the following dilutions: anti-p16 (1:500), anti-p21 (1:1000), anti-phosphorylated pRB (1:1000), anti-TXNIP (1:500), anti- α -tubulin (1:30,000).

HPLC analysis of 4-HNE

Polar lipids were extracted from macrophage or foam cell culture media and used to determine 4-HNE content by HPLC, as described before [42]. Briefly, medium was collected for analysis from confluent cell cultures maintained with serum-free RPMI 1640 medium for 16 hrs. This was important in order to prevent 4-HNE-protein adduct formation. Following extraction of polar lipids and HPLC, the elution peak of 4-HNE (223 nm) was detected at 4.2 min. The recovery of the standard, added to fresh samples prior to extraction, was 85–90%. Peaks in the HPLC profiles were monitored and quantified by using Clarity-Lite software (DataApex Co., Prague, Czech Republic).

LC-MS/MS analysis

Analysis of FL-926-A16 adducts with 4-HNE, acrolein, glyoxal and methylglyoxal in culture media of THP-1 macrophages or foam cells was performed as described previously [55]. The details of the analysis are given in the Supporting information section.

hTXNIP overexpression

Vascular endothelial cell plated in 6-well dishes were transfected at 60% confluency with the hTXNIP-lacZ expression plasmid or the empty plasmid [56] using TransIT-LT1 reagent, according to the manufacturer's instructions (Mirus Bio-Corporation). Briefly, DNA (2.5 μ g) was complexed with the transfection reagent for 20 min. in OptiMEM medium, then the mixture was added to cells in complete growth medium for an ON incubation. Cells were harvested after 3 days for analysis of TXNIP or after 7 days for analysis of senescence markers.

siRNA transfection

Vascular endothelial cell plated at 50% confluency in 6-well dishes were transfected with 25 nM of the siRNA for bovine TXNIP or

scrambled RNA sequences, using *TransIT*-TKO (Mirus Bio-Corporation). The RNA sequence was complexed with the transfection reagent for 20 min in OptiMEM and added to cells in complete growth medium. The medium was changed 24 hrs later, and the cells were co-cultured with foam cells in a transwell system for additional 72 hrs, as described above. The cells were then harvested and processed for Western blot analysis. The target sequence for bovine TXNIP silencing was 5'-CAGAAGUUGUCAUCAGUCA-3'.

Luciferase assay

Vascular endothelial cell cultures, at 60% confluency, were co-transfected with DNA complexed to *TransIT*-LT1 (Mirus Bio-Corporation) as described above. Analysis of PPAR and TXNIP activation were performed as previously described [44, 52]. Briefly, PPAR activation was monitored in cells that were transfected with 165 ng of expression vectors for human (h)PPAR- α , hPPAR- γ 1, hPPAR- γ 2 or hPPAR- δ , along with of the retinoid X receptor (hRXR, 82.5 ng), green fluorescent protein (82.5 ng) and Renilla luciferase (100 ng) expression vectors, and the 3XPPRE-TK-luciferase reporter (500 ng). For analysis of TXNIP activation, cells were co-transfected with a luciferase-reporter construct encoding the human TXNIP promoter region 1777 bp upstream of the ATG start codon (0.5 μ g) with renilla luciferase reporter plasmid (0.1 μ g). Luciferase-induced luminescence was determined by using the Mithras LB-940 luminometer (Berthold Technologies, Bad Wildbad, Germany); results were normalized to the Renilla luciferase activity, used as an internal control, according to the kit's instructions.

Immunofluorescent staining

Human atherosclerotic plaques removed from the carotid bifurcation of patients following endarterectomy and left internal mammary arteries (LIMA), removed during coronary artery bypass graft, were embedded in OCT compound and frozen sections prepared for immunofluorescence. Cryosections (7 μ m thick) were prepared and immunostained with monoclonal antibodies against Factor VIII (1:500; green) and either p21 (1:500, red) or TXNIP (1:100, red) for 16 hrs at 4°C; cell nuclei were visualized with DAPI staining. Secondary antibodies AlexaFluor488- and AlexaFluor647-labelled anti-rabbit and anti-mouse (1:800) were applied for 2 hrs at room temperature. Visualization was by confocal microscopy. The study was approved by the Hadassah Hebrew University Medical Center Institutional Review Board and each patient signed an informed consent.

Statistical analysis

Data are given as mean \pm SEM. One sample Student's *t*-test was used to validate statistical differences in experiments expressing data as percent of control. Statistical significance of differences between groups was determined by multiple comparison one-way ANOVA, followed by the Dunnett's test using the GraphPad Prism statistical software.

Results

Foam cells induce senescence in VEC

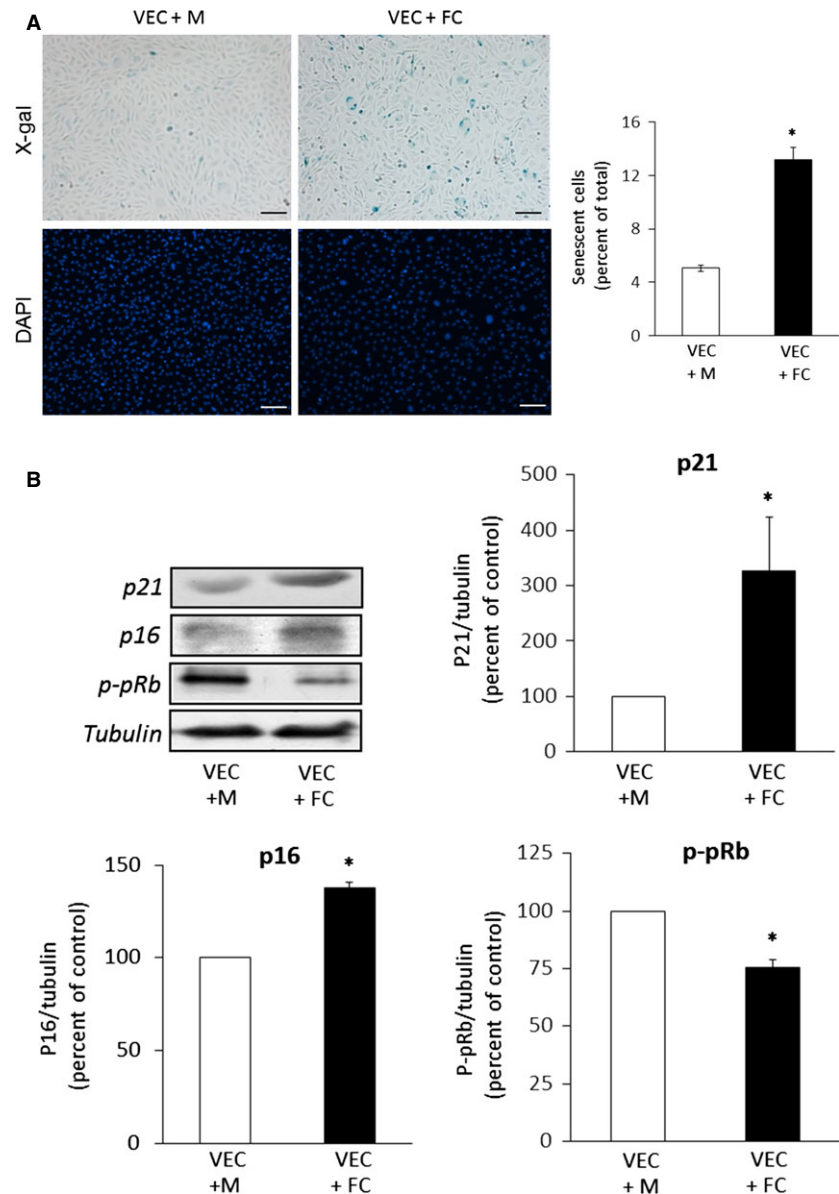
The capacity of foam cells to induce senescence in VEC was assessed in an *in vitro* co-culture transwell assay: THP-1 monocytes, seeded on transwell membranes, were first differentiated into macrophages by PMA treatment and then transformed into foam cells following exposure to OxLDL, as described in 'Materials and methods'. Positive Red oil O staining confirmed the formation of lipid-laden foam cells, whereas the control macrophages (PMA-treated only) incorporated negligible amount of the dye (Fig. S1). A significant increase in SA- β -gal staining and the enlarged morphology typical of senescent cells were observed in VEC co-cultured with foam cells (Fig. 1A). Western blot analysis (Fig. 1B) showed that VEC monolayer co-cultured with foam cells presented increased expression of p21 and p16 and reduced level of phosphorylated (Ser⁷⁹⁵) retinoblastoma protein (p-pRB). Together, these results show the ability of foam cells to induce senescence in VEC *via* secretion of soluble factor(s).

Senescence is induced by 4-HNE in VEC

Next, we studied whether 4-HNE was released from foam cells and could mediate senescence induction. HPLC analysis of polar lipid extracts of media collected from PMA-treated THP-1 cells and from foam-cell cultures showed increased secretion of 4-HNE (3.5 \pm 0.4-fold higher than the control, Fig. 2A). The effect of exogenously added 4-HNE on senescence in VEC was assessed following exposure to 1 and 10 μ M 4-HNE for 7 days. Treatment with 10 μ M 4-HNE increased the proportion of senescent cells, demonstrated by cellular enlargement and positive staining for SA- β -gal (Fig. 2B). This effect was accompanied by a 40 \pm 7.2% reduction in cell number per plate. In addition, increased expression of p16 and p21 and decreased p-pRB were most prominent at this concentration of 4-HNE (Fig. 2C). Taken together, these findings suggest that 4-HNE mediates foam cell-induced VEC senescence.

To further confirm this suggestion, we investigated whether the scavenging of 4-HNE in the transwell system could blunt foam cell-induced VEC senescence. FL-926-A16 is a potent scavenger of bioactive aldehydes, including 4-HNE (European Patent EP 2519507 B1, US Patent 8623900 B2 and [57]). Unlike the natural scavenger, L-carnosine, which is degraded by carnosinase in human cells [58], its alcohol derivative FL-926-A16 is resistant to degradation. LC-MS/MS analysis of culture medium from control (PMA-treated) macrophages and foam cells exposed to FL-926-A16 showed that the latter covalently bound 4-HNE (Fig. S2). Interestingly, this analysis also revealed that, in addition to 4-HNE, foam cells secreted glyoxal, methyl glyoxal and acrolein, which were also scavenged by FL-926-A16. When added to the medium of the co-culture system, FL-926-A16 reduced the level of p16 and p21 and increased p-pRB level (Fig. 3), supporting the suggestion that the scavenged molecules, including 4-HNE, induce senescence.

Fig. 1 Foam cell-induced senescence in VEC in the transwell co-culture system. **(A)** Senescence-associated β Galactosidase (SA- β -Gal) activity. VEC and THP-1 macrophages (M) or foam cells (FC) were maintained in transwell co-cultures for 72 hrs, as described under 'Materials and methods'. The VEC monolayers were then washed, fixed and stained with X-Gal or DAPI as described. SA- β -Gal positive cells and total cell number were quantified from three different fields in each well. Results are expressed as percent of SA- β -Gal positive VEC (mean \pm SEM, $n = 3$; $*P < 0.05$ for differences from VEC+M control); scale bar, 100 μ m. **(B)** Expression of senescence markers. At the end of the co-culture period, VEC were harvested, lysed and analysed by Western blot for the expression of p21, p16 and phospho-Ser795-pRB (p-pRB). Shown are representative Western blots and quantification of the different senescence markers. Results are expressed as mean \pm SEM, $n = 3$; $*P < 0.05$, for differences from the respective M-treated VEC (taken as 100%).



TXNIP mediates 4-HNE-induced senescence

It has been suggested that increased TRX activity protects VEC from atherogenesis [59]. TXNIP, the natural inhibitor of TRX, plays a prominent role in regulating TRX activity in VEC; high TXNIP levels were associated with endothelial cell dysfunction [49, 50, 60]. Figure 4A suggests the involvement of TXNIP in passage-induced replicative senescence in VEC. Indeed, overexpression of TXNIP in low passaged VEC cultures (transfected with the hTXNIP-lacZ expression plasmid) augmented the expression of p21 and reduced p-pRB levels (Fig. 4B). Next, we studied the effect of exogenously added 4-HNE on TXNIP expression and its impact on oxidative stress-induced senescence and found increased TXNIP

expression in VEC incubated with 10 μ M 4-HNE for 7 days (Fig. 4C). Importantly, TXNIP expression was increased in VEC co-cultured with foam-cells (Fig. 4D). The 4-HNE scavenger FL-926-A16 decreased TXNIP expression in VEC co-cultured with foam cells (Fig. 4E), supporting the hypothesis that lipid peroxidation products are involved in the up-regulation of TXNIP expression. Noteworthy, silencing of TXNIP in VEC co-cultured with foam cells prevented the induction of the senescence markers (Fig. 4F). Scrambled RNA sequences had no influence on TXNIP and senescence marker expression (Fig. S3). Collectively, these results suggest that 4-HNE up-regulates TXNIP expression and support a role for the latter in both replicative and stress-induced senescence in VEC.

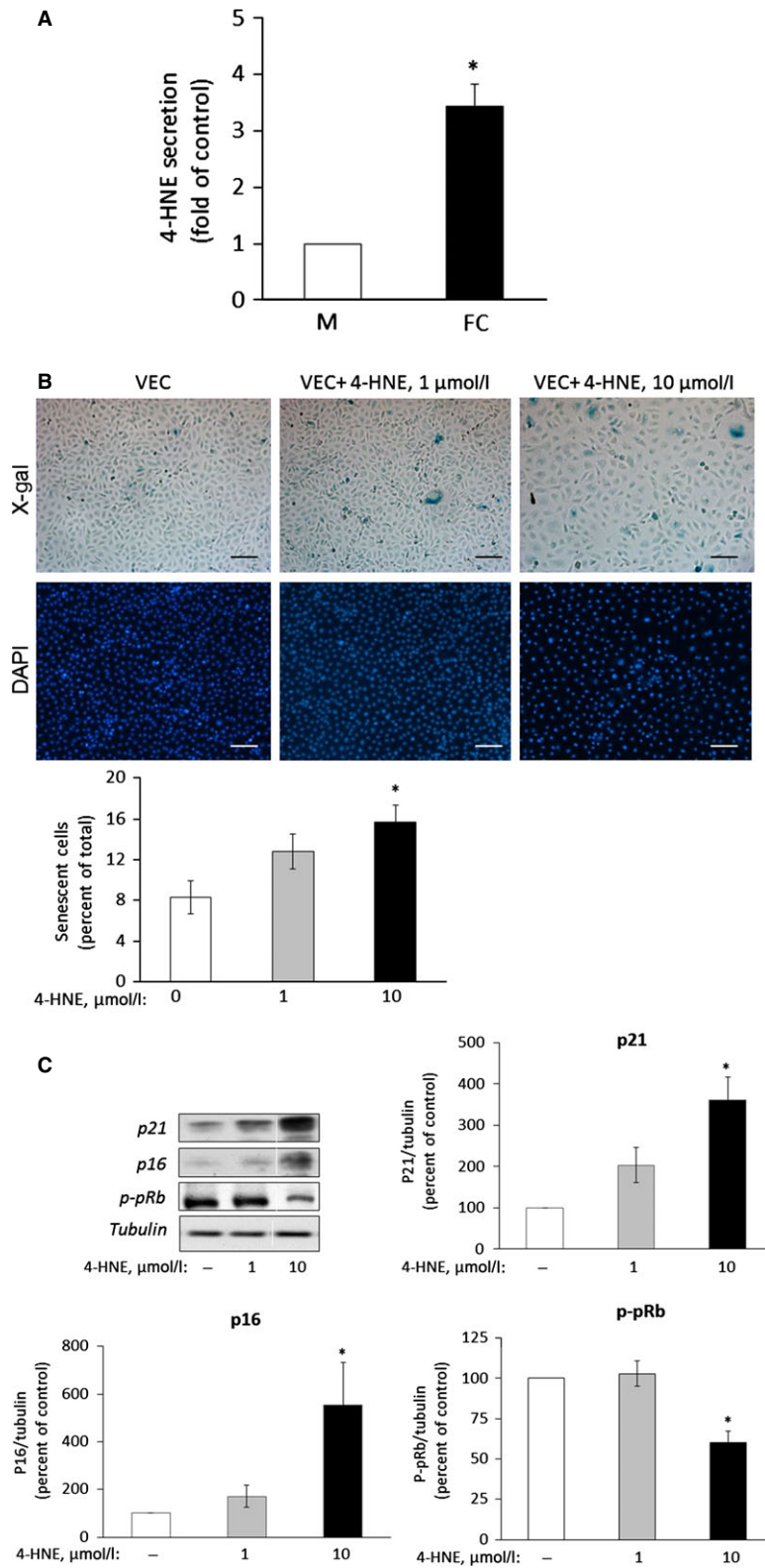
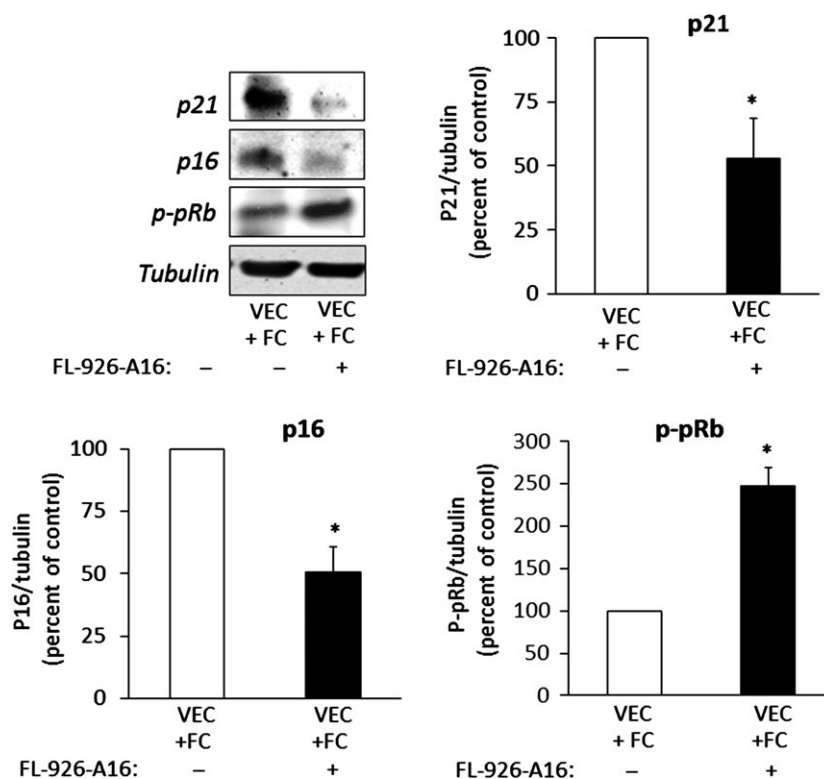


Fig. 2 The lipid peroxidation product 4-HNE induces senescence in VEC. **(A)** THP-1 macrophages (M) and foam cells (FC) were prepared as described under 'Materials and methods'. The culture medium was changed to serum free-medium for 16 hrs. The medium was then collected, cell debris was cleared off by centrifugation and lipids were extracted and taken for HPLC analysis of polar lipids. The level of 4-HNE, normalized to μg of protein, in the M extracts was taken as arbitrary unit (1). Results are expressed as mean \pm SEM, $n = 3$. * $P < 0.05$ for differences from the respective control. **(B)** Senescence-associated β -Galactosidase (SA- β -Gal) activity in 4-HNE-treated cells. VEC were treated daily with 1 or 10 μM HNE for 7 days. At the end of the incubation period, VEC were processed and analysed as described under the legend of Figure 1A. Results are expressed as percent of SA- β -Gal positive VEC (mean \pm SEM, $n = 3$; * $P < 0.05$, for differences from vehicle-treated controls); scale bar, 100 μm . **(C)** Expression of senescence markers. At the end of the incubation period with 4-HNE, the cells were harvested, lysed and processed as described under the legend of Figure 1B. Shown are representative Western blots and quantification of the various markers studied. Results are expressed as mean \pm SEM, $n = 3$; * $P < 0.05$, for differences from the respective 4-HNE untreated controls, taken as 100%.

Fig. 3 Scavenging of 4-HNE reduces the expression of senescence markers in VEC co-cultured with foam cells. THP-1 monocytes seeded in transwell inserts were transformed to foam cells (FC) as described under legend to Figure 1A, and then co-cultured with VEC for 72 hrs, in the absence or presence of 10 mM FL-926-A16, with daily addition of the scavenger. At the end of the incubation period, the cells were harvested, lysed and processed for Western blotting as described under the legend to Figure 1B. Shown are representative Western blots and quantification of the various senescence markers. Results are expressed as mean \pm SEM, $n = 3$; * $P < 0.05$ for differences from the untreated cultures.



The role of PPAR δ in foam cell-induced senescence

Others and we have shown that 4-HNE activates PPAR δ [42, 43]; hence we investigated whether this mechanism is also involved in 4-HNE-induced senescence in VEC. Figure 5A shows that the selective PPAR δ antagonist GSK0660 abolished 4-HNE effects on the expression of the different senescence markers and prevented the up-regulation of TXNIP. In addition, VEC were exposed for 48 hrs to selective agonists for the various PPAR receptors at their effective concentrations, as described by us before in the same VEC culture [44]. The selective PPAR δ agonist, GW501516, markedly augmented TXNIP expression, whereas, the PPAR α agonist Wy14643 and the PPAR γ agonist, troglitazone, did not modify significantly TXNIP expression (Fig. 5B). The hypothesis that 4-HNE activates PPAR δ in VEC was further investigated by assessing the transactivation of PPRE-coupled luciferase reporter [44]. Briefly, VEC were co-transfected with the PPAR response element (PPRE)-luciferase reporter vector (3xPPRE-TK-Luciferase) and a combination of a specific human PPAR vector and RXR expression vector, followed by a 48 hrs treatment with 4-HNE (Fig. 5C). A significant increase in luciferase activity in cells exposed to 10 μ M 4-HNE was observed only in cells over-expressing the human PPAR δ . The expression of PPAR α was not altered in the presence of 4-HNE, whereas the expression of PPAR γ_1 and PPAR γ_2 was somewhat reduced. To further test whether PPAR δ mediated the enhancement of TXNIP transcription by 4-HNE, VEC were transfected with the human TXNIP promoter (1777 bp upstream

of the ATG start codon)-luciferase vector. TXNIP promoter activity was increased by 4-HNE and reduced in the presence of the PPAR δ antagonist GSK0660 (Fig. 5D).

VEC senescence in human carotid atherosclerotic plaques

Finally, we verified the relevance of the above findings to human atherosclerosis. Figure 6 shows sections of a human carotid atherosclerotic plaque stained for the endothelial cell marker Factor VIII (green), and p21 or TXNIP (red) in the near-plaque area. The co-localization of TXNIP or p21 with Factor VIII in the shoulder region of the plaque suggests that endothelial cells expressing TXNIP are prone to senescence. No staining of TXNIP or p21 was observed in Factor VIII-positive cells from sections of LIMA, which are resistant to cholesterol buildup and atherosclerosis [61, 62] (Fig. 6, bottom panels).

Discussion

Cellular senescence has been linked to endothelial cell dysfunction and atherosclerosis [14, 63]. In the present study we investigated the mechanism involved in foam cell-induced VEC senescence. Using a co-culture system, in which VEC were exposed on the apical side to soluble factors released by foam cells [23, 25], we found that the latter produced soluble factors that promoted SIPS in VEC. Our findings

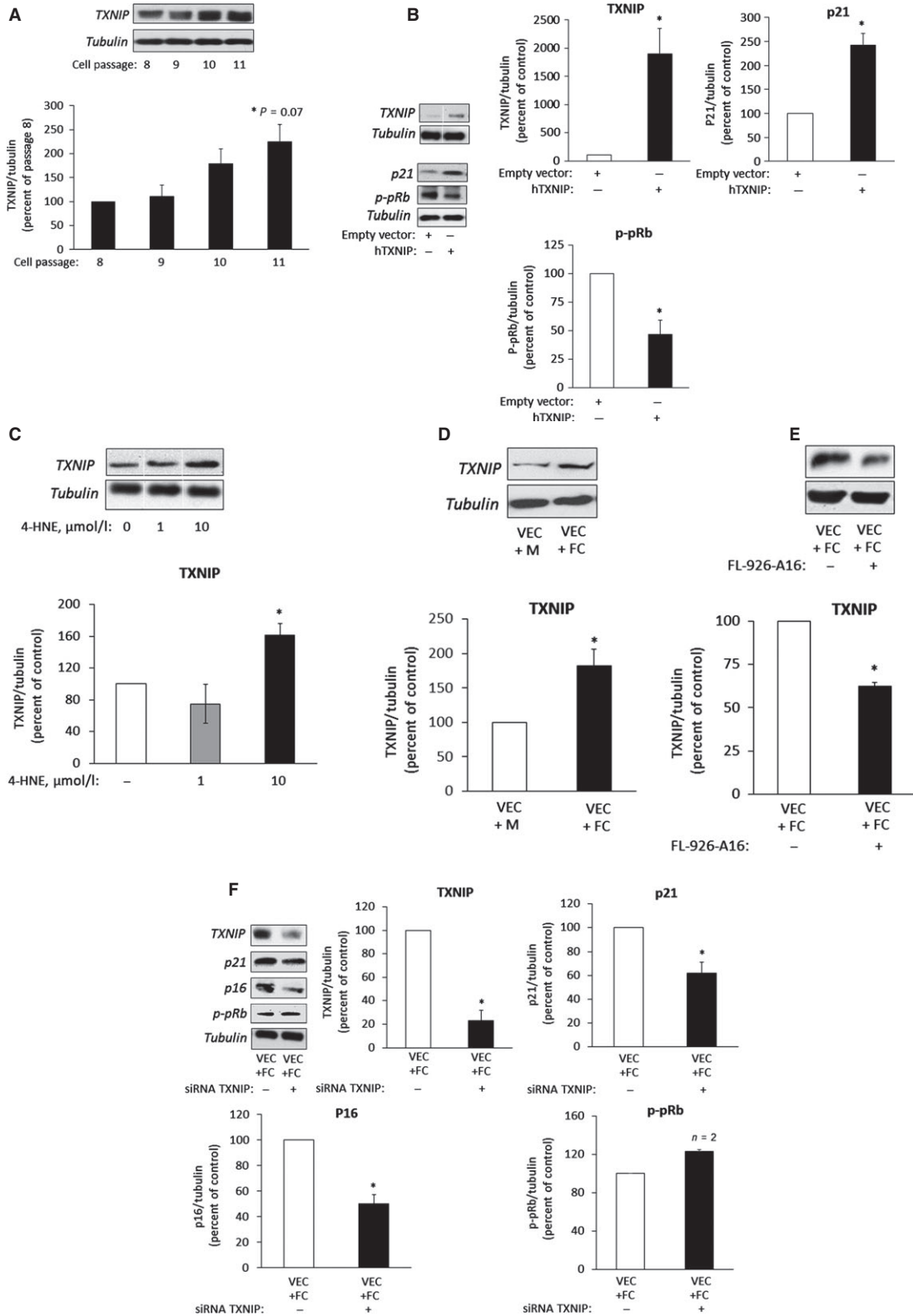


Fig. 4 The role of TXNIP in VEC senescence. **(A)** TXNIP expression in VEC at different passages was analysed by Western blot. $P < 0.07$, for differences from cells at passage 8 ($n = 3$). **(B)** VEC were transfected with the hTXNIP-lacZ expression plasmid or with an empty vector. Cells were then harvested and taken for Western blot analysis of TXNIP and senescence markers; results are expressed as mean \pm SEM, $n = 3$; $*P < 0.05$ for differences from cells transfected with the empty vector. **(C)** VEC cultures were treated with the indicated concentrations of 4-HNE for 3 days, with a daily change of medium. Cell lysates were prepared and used for Western blot analysis of TXNIP. The level of TXNIP in untreated controls was taken as 100%. Results are expressed as mean \pm SEM, $n = 3$; $*P < 0.05$ for differences from the control. **(D)** VEC co-cultured with THP-1 macrophages (M) or foam cells (FC) as described above, and analysed for TXNIP expression. TXNIP expression levels in the VEC + M group was taken as 100%. Results are expressed as mean \pm SEM, $n = 3$; $*P < 0.05$ for differences from the control groups. **(E)** VEC co-cultured with foam cells (FC) in the absence (control) or presence of 10 mM FL-926-A16, as described under the legend to Figure 3, and analysed for TXNIP expression. Results are expressed as mean \pm SEM, $n = 3$; $*P < 0.05$ for differences from the control groups. **(F)** VEC were transfected with TXNIP siRNA and taken 24 hrs later for co-culture with foam cells (FC) for additional 72 hrs. VEC were then lysed and taken for Western blot analysis of TXNIP or of the senescence markers. The level of TXNIP and the senescence markers in untransfected controls was taken as 100%. Representative Western blots quantification of TXNIP and of various senescence markers are shown. Results are expressed as mean \pm SEM, $n = 3$; $*P < 0.05$ for differences from respective controls ($n = 3$).

provide an important insight into the signal transduction pathway that culminated in endothelial cell senescence: foam cells derived from macrophages following exposure to OxLDL secreted lipid peroxidation products, including the nucleophilic aldehyde 4-HNE. The latter interacted with VEC and activated the nuclear factor PPAR δ , which transactivated the TXNIP promoter. Consequently, TXNIP expression was increased, promoting senescence in the endothelial cells (see model in Fig. 7).

Non-enzymatic lipid peroxidation of PUFA and their metabolites, promoted by ROS, is known to generate various α,β -unsaturated aldehydes [29, 31]. Moreover, since these products are often more stable than ROS, they can diffuse into other tissues and propagate oxidative injury. Among these, 4-HNE is considered one of the most abundant and bioactive species. Its plasma concentration in healthy individuals is estimated to be around 1 μ M, whereas in disease states it may reach much higher levels [64]. Lipid peroxidation products are known to affect the redox balance during ageing and are involved in VEC dysfunctions, including apoptosis, monocyte adhesion to VEC and inflammation; the latter has been suggested recently to result from 4-HNE-induced ER stress [37]. Our study shows that exposure of macrophages to OxLDL results in the formation of lipid-laden foam cells that produce several lipid peroxidation products including 4-HNE, which induced VEC senescence at non-cytotoxic concentrations. The stable L-carnosine derivative FL-926-A16, which scavenges lipid peroxidation products, attenuated foam cell-induced VEC senescence. Collectively, these observations suggest that lipid peroxidation products, such as 4-HNE, mediate the effect of foam cells on VEC senescence. Among the known properties of 4-HNE is its biphasic activity: at low concentrations it presents a signalling function *via* activation of the nuclear factor PPAR δ , whereas at high concentrations it acts as a cytotoxic agent [31, 42, 43]. Interestingly, our previous studies showed that endothelial cells do not produce detectable amounts of 4-HNE [44]. Therefore, it appears that exogenous 4-HNE that diffuses from foam cells is involved in VEC senescence.

An important target of 4-HNE in VEC is the TRX system which is suggested to be involved in the etiology of several human diseases, including atherosclerosis [46]. It has been

shown before that 4-HNE interacts with and attenuates the function of the TRX system in various cells, including VEC [51, 65, 66]. The present study shows that 4-HNE may also affect the TRX system indirectly by augmenting the expression of its natural inhibitor, TXNIP. Recent studies have shown that TXNIP promoted endothelial inflammation and increased leucocyte adhesion to endothelial cells under conditions of disturbed blood flow that characterize blood vessel regions prone to atherosclerosis [50]. The present study shows that increased TXNIP expression in VEC, either following treatment with 4-HNE or by overexpression of human TXNIP, enhanced VEC senescence. Importantly, the silencing of *Txnip*, which partially inhibited foam cell-induced senescence in VEC, suggests that TXNIP is a downstream target of 4-HNE in this process.

Others and we have previously shown that 4-HNE is an activator of the nuclear factor PPAR δ [42, 43]. Using both pharmacological and molecular tools we now show that PPAR δ activation is also involved in 4-HNE-induced VEC senescence. Treatment with 4-HNE of VEC transfected with the different human PPAR isotypes showed specific activation of PPAR δ , but not PPAR α . Interestingly, 4-HNE reduced the activity of PPAR γ 1 and γ 2. In this context, it should be noted that reduced PPAR γ activity has been linked to the induction of senescence in photo-irradiated human fibroblasts [67]. In addition, GSK0660, a selective PPAR δ antagonist [68], attenuated the effects of 4-HNE on senescence and TXNIP expression. These findings led to the hypothesis that TXNIP is a putative target for PPAR δ in VEC. Indeed, the transactivation assay in the presence of 4-HNE and the antagonist GSK0660 support the involvement of PPAR δ in this process.

TXNIP has been linked to atherosclerosis by several *in vivo* and *in vitro* studies [50, 69]. Here, we show by immunofluorescence staining of sections from atherosclerotic arteries that both TXNIP and the p21 senescence marker were present in the endothelial cells near the plaque area, supporting the relationship between TXNIP and senescence *in vivo*. Endothelial cells of LIMA, which are not prone to atherosclerosis, stained neither for TXNIP nor for p21.

What could be the outcome of premature senescence on the vascular endothelium? As opposed to age-related senescence,

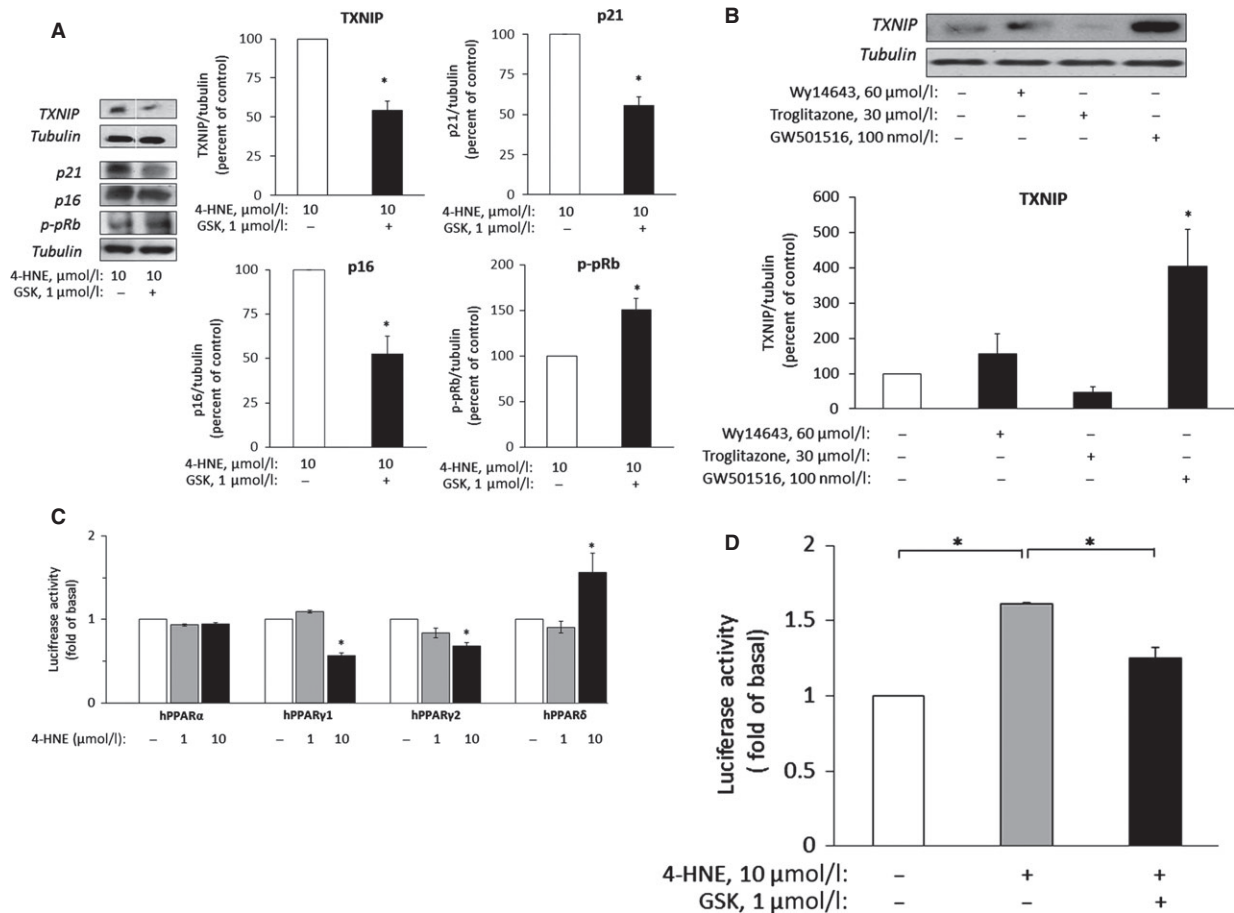


Fig. 5 PPAR δ modulates the expression of TXNIP and senescence markers in VEC. **(A)** VEC were treated with 10 μM 4-HNE without (control) or with 1 μM GSK0660 (GSK) for 7 days with daily changes of medium. The expression of TXNIP and the senescence markers was determined in lysed cells by Western blot analysis. The level of TXNIP and the senescence markers in control lysates was taken as 100%. $*P < 0.05$ for differences from the respective controls (mean \pm SEM, $n = 3$). **(B)** VEC cultures were maintained for 48 hrs without (control) or with the indicated agonists for PPAR α , $-\gamma$ and $-\delta$. Cells were then taken for Western blot analysis of TXNIP expression. $*P < 0.05$, for differences from control cells treated with vehicle only. **(C)** VEC cultures were transfected with the following expression vectors: pSG5-hPPAR α , pCMX-hPPAR γ 1, pCMX-hPPAR γ 2 or pCDNA-hPPAR δ and co-transfected with pSVPORT-hRXR, pEGFP-N1 plasmid, 3xPPRE-TK-Luciferase reporter plasmid and Renilla luciferase plasmid. The cells were incubated for 48 hrs. During the last 24 hrs of incubation the cultures were incubated without (control) or with 1 or 10 μM 4-HNE. The cells were then harvested, lysed and the relative luciferase activity was determined and normalized as described under 'Materials and methods'. $*P < 0.05$, for differences from the respective controls, taken as 1 unit (mean \pm SEM, $n = 3$). **(D)** VEC cultures were co-transfected with a luciferase reporter construct containing the human TXNIP promoter and with the Renilla luciferase reporter plasmid, as described under 'Materials and methods'. After 24 hrs of incubation the medium was changed and cells were incubated for additional 24 hrs without (control) or with 10 μM 4-HNE in the absence or presence of 1 μM GSK0660. Cells were then lysed and taken for the luciferase activity assay as described above. The relative luciferase activity was normalized to untreated control. $*P < 0.05$, for differences from the indicated groups (mean \pm SEM, $n = 3$).

involving shortening of the telomeres, which is irreversible, telomere-independent SIPS is potentially a reversible process. Brodsky *et al.* [70] have shown that ebselen, a peroxynitrite scavenger and antioxidant, prevented and reversed senescence in early-passaged human umbilical vein endothelial cell cultures. This also occurred in ebselen-fed Zucker diabetic fatty rats, in which the symptoms of vasculopathy were reduced, accompanied

by prevention and reversal of endothelial cell senescence in their aorta. Similarly, *N*-acetyl-cysteine reversed stress-induced senescence in bone marrow-derived hematopoietic stem-cells, further supporting the hypothesis that SIPS is reversible [17, 71]. The association between senescence and atherosclerosis and the potential reversibility of telomere-independent SIPS, as opposed to telomere dependent replicative senescence, suggest that the

Fig. 6 Double immunofluorescence staining for Factor VIII and p21 or TXNIP in sections of human carotid plaque or from LIMA. Human atherosclerotic plaques were removed from the carotid bifurcation of patients by endarterectomy and frozen in OCT as described in 'Materials and methods'. Immunostaining of tissue sections with monoclonal antibodies against Factor VIII (green) and either p21 or TXNIP (red) was performed. Cell nuclei were stained with DAPI. Merged images in the right panels demonstrate co-localization of p21 and TXNIP with Factor VIII (top panels). LIMA sections were used as negative control; scale bar 20 μ m. The images are representative of three samples of human carotid plaques and two samples of LIMA.

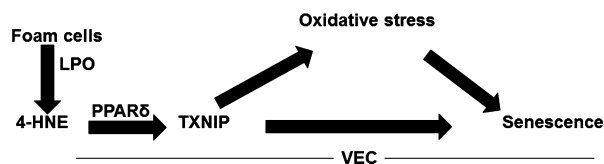
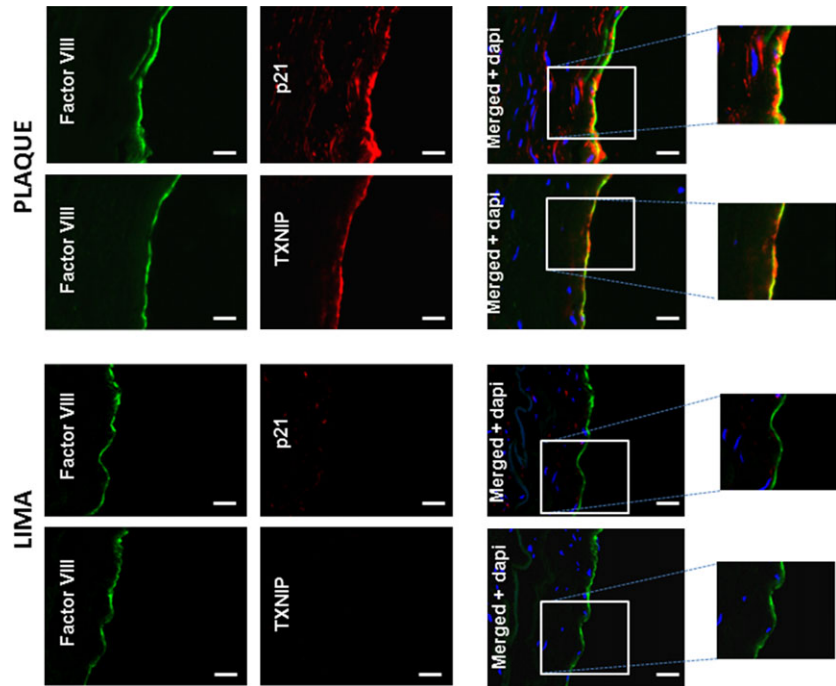


Fig. 7 Suggested model of foam cell-induced VEC senescence. Foam cell-derived lipid peroxidation products, among which 4-HNE, reach VEC and activate PPAR δ and subsequently increase the expression of TXNIP which augments oxidative stress and VEC senescence.

former may be more relevant in the context of vascular pathology. Therefore, better understanding of the mechanisms of stress-induced senescence is important for devising means to control atherosclerosis by reversing or attenuating this process in VEC. Our study shows that in addition to the well-known effects of oxidative stress, 4-HNE, TXNIP and PPAR δ mediate SIPS, and thus may serve as potential targets for the prevention of VEC senescence. The modulation of these targets attenuated foam cell-induced SIPS *in vitro*. Specifically, we showed that attenuation of SIPS could be achieved by: (i) scavenging of foam-cell derived lipid peroxidation products, in particular the long-lived bioactive aldehyde 4-HNE; (ii) reduction of TXNIP expression and (iii) inhibition of PPAR δ activity. In conclusion, better characterization of the newly identified pathway(s) of 4-HNE-induced premature senescence in VEC (Fig. 7) may provide new insight into the molecular mechanisms of this process and identify potential targets for therapeutic intervention.

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Conflicts of interest

Erol Cerasi is the founder and unpaid manager of Diab R&D. All other authors declare no conflict of interest and are not employed by Diab R&D.

Supporting information

Additional Supporting Information may be found in the online version of this article:

Data S1 Foam cell-derived 4-hydroxynonenal induces endothelial cell senescence in a TXNIP-dependent manner.

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