

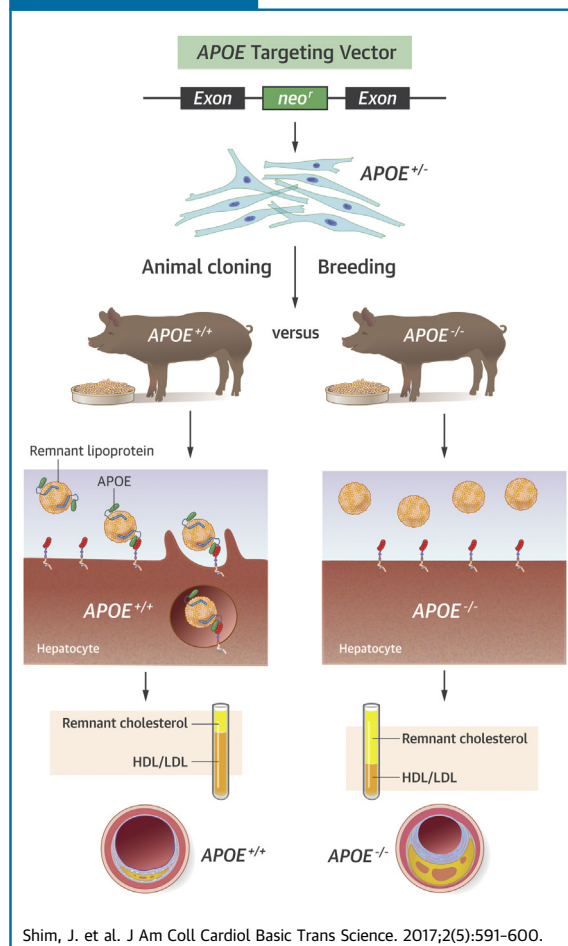
PRECLINICAL RESEARCH



Apolipoprotein E Deficiency Increases Remnant Lipoproteins and Accelerates Progressive Atherosclerosis, But Not Xanthoma Formation, in Gene-Modified Minipigs

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VISUAL ABSTRACT



HIGHLIGHTS

- *APOE*-deficient Yucatan minipigs were created by recombinant adeno-associated virus mediated gene targeting in porcine fibroblasts followed by somatic cell nuclear transfer.
- *APOE*^{-/-} minipigs displayed increased plasma cholesterol and accumulation of APOB48-containing chylomicron remnants on low fat-diet, which was significantly accentuated upon feeding a high-fat, high-cholesterol diet.
- *APOE*^{-/-} minipigs showed accelerated progressive atherosclerosis but not xanthoma formation indicating that remnant lipoproteinemia does not induce early lesions but is atherogenic in pre-existing atherosclerosis.

ABBREVIATIONS AND ACRONYMS

APOB = apolipoprotein B

APOE = apolipoprotein E

cDNA = complementary DNA

HFHC = high-fat
high-cholesterol

IDL = intermediate-density
lipoprotein

LAD = left anterior descending
(coronary artery)

LDL = low-density lipoprotein

LDLR = low-density lipoprotein
receptor

LF = low-fat

Neo = neomycin

rAAV = recombinant
adeno-associated virus

SMC = smooth muscle cell

VLDL = very-low-density
lipoprotein

SUMMARY

Deficiency of apolipoprotein E (APOE) causes familial dysbetalipoproteinemia in humans resulting in a higher risk of atherosclerotic disease. In mice, APOE deficiency results in a severe atherosclerosis phenotype, but it is unknown to what extent this is unique to mice. In this study, *APOE* was targeted in Yucatan minipigs. *APOE*^{-/-} minipigs displayed increased plasma cholesterol and accumulation of apolipoprotein B-48-containing chylomicron remnants on low-fat diet, which was significantly accentuated upon feeding a high-fat, high-cholesterol diet. *APOE*^{-/-} minipigs displayed accelerated progressive atherosclerosis but not xanthoma formation. This indicates that remnant lipoproteinemia does not induce early lesions but is atherogenic in pre-existing atherosclerosis. (J Am Coll Cardiol Basic Trans Science 2017;2:591-600) © 2017 The Authors. Published by Elsevier on behalf of the American College of Cardiology Foundation. This is an open access article under the CC BY-NC-ND license (<http://creativecommons.org/licenses/by-nc-nd/4.0/>).

The most efficient animal model of experimental atherosclerosis is the apolipoprotein E (APOE)-deficient mouse (1) which develops severe hypercholesterolemia and rapid, fibroatheromatous atherosclerosis on normal chow diet. APOE is synthesized in hepatocytes and several other cell types, and serves as a ligand for low-density receptor-related protein 1-mediated clearance of apolipoprotein B-48 (APOB-48)-containing chylomicron remnants and low-density lipoprotein receptor (LDLR)-mediated clearance of very-low-density lipoprotein (VLDL) and intermediate-density lipoprotein (IDL) (2). Due to hepatic editing of *Apob* mRNAs, mice secrete APOB-48-containing VLDLs, making them particularly dependent on APOE-mediated clearance mechanisms (3). However, loss of several other APOE-mediated functions may contribute to the severe atherosclerosis phenotype of the *Apoe*^{-/-} mice. APOE has been shown to dampen inflammation, inhibit smooth muscle cell (SMC) proliferation and migration, and to be involved in reverse cholesterol transport (4). Lipoproteins from *Apoe*^{-/-} mice also

more readily produce foam cells in vitro for reasons that are not well understood (5). In humans, genetic variants that increase remnant lipoproteins similar to those seen with APOE-deficiency not only increase the risk of ischemic heart disease but also the plasma level of C-reactive protein (CRP). This is not seen with gene variants that increase low-density lipoprotein (LDL); possibly reflecting a pro-inflammatory effect of remnant lipoproteins in the arterial wall (6).

Several lines of hypercholesterolemic minipigs exist with spontaneous or genetically engineered defects in LDLR-mediated hepatic LDL uptake (7). These include familial hypercholesterolemia-Bretoncelles-Meishan pigs with a spontaneous *LDLR* mutation (8), human *D374Y* gain-of-function proprotein convertase subtilisin/kexin type 9 (PCSK9) transgenic Yucatan minipigs (9), and most recently, *LDLR* knockout Yucatan minipigs (10). When fed a high-fat high-cholesterol diet, these lines accumulate LDL-sized and larger APOB-100-containing lipoproteins and they develop fibroatheromatous atherosclerotic plaques in major arteries. These lines are useful for many purposes, but rates of plaque progression are modest,

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All authors attest they are in compliance with human studies committees and animal welfare regulations of the authors’ institutions and Food and Drug Administration guidelines, including patient consent where appropriate. For more information, visit the JACC: Basic to Translational Science [author instructions page](#).

Manuscript received February 20, 2017; revised manuscript received June 1, 2017, accepted June 8, 2017.

especially in the coronary arteries, and the disease does not reach a symptomatic stage with plaque rupture or thrombotic events within time frames observed so far.

In the present study, we created Yucatan minipigs with targeted disruption of the *APOE* gene and assessed the impact of accumulating remnant lipoproteins on the development of atherosclerosis.

METHODS

Detailed methods are available in the [Supplemental Appendix](#).

ANIMALS. The Danish Animal Experiments Inspectorate approved all procedures involving animals. Yucatan minipigs were originally purchased from Moran Farm Enterprises (Ballina, Ireland) and a breeding colony was established and maintained at Aarhus University. All animals were housed in a specific pathogen-free stable facility under standard conditions. The minipigs were fed a low-fat (LF) standard pig diet until 8 weeks of age and subsequently a high-fat high-cholesterol (HFHC) diet consisting of standard pig feed comprising 20% (w/w) of lard and 2% cholesterol (Sigma-Aldrich, St. Louis, Missouri) until euthanization at the age of 16 weeks (*APOE*^{-/+}) or 52 weeks (*APOE*^{+/+} and *APOE*^{-/-}).

CONSTRUCTION OF RECOMBINANT ADENO-ASSOCIATED VIRUS/APOE KNOCKOUT TARGETING VECTOR. As human *APOE*, the porcine *APOE* gene consists of 4 exons and the porcine protein has an amino acid similarity of 70.3% compared to human *APOE* (11,12). The recombinant adeno-associated virus (rAAV)/*APOE* knockout (KO) vector was constructed as previously described (13,14). Two homology arms (~1 kb each), flanking exon 3 of the porcine *APOE* gene, were amplified by polymerase chain reaction (PCR) using Yucatan minipig genomic DNA as template. These 2 homology arms were subsequently linked to a neomycin (*neo*^r)/zeomycin (*zeo*^r) resistance gene cassette by a 3-way fusion PCR. The *neo*^r/*zeo*^r cassette, flanked by loxP sites, was comprised in a 4-kb *PvuI* fragment isolated from a plasmid called pNeDaKO-Neo (pronounced 'p-Need a Knockout-Neo'); a generous gift from Bert Vogelstein and Kenneth W. Kinzler, The Johns Hopkins University Medical Institutions, Baltimore, Maryland. The 3-way fusion PCR was performed as described by Kohli et al. (13) using a Platinum Pfx polymerase (Thermo Fisher Scientific, Waltham, Massachusetts) and the following PCR protocol: 1 cycle of 94°C for 1 min; 25 cycles of 94°C for 30 s, 59°C for 30 s, and 68°C for 4 min; 1 cycle of 68°C for 7 min. Fusion products were digested with *NotI* and ligated to a *NotI*-cleaved pAAV-MCS plasmid backbone

(Stratagene, San Diego, California, cat. no. 240071-5) containing an ampicillin (*amp*) resistance gene and the viral inverted terminal repeat sequences. The final rAAV/*APOE* KO plasmid construct was verified by *NotI* digestions and sequencing. Successful gene targeting using the rAAV/*APOE* KO vector results in deletion of 295 bp in the *APOE* gene including the entire exon 3. PCR primers for generating the rAAV/*APOE* KO construct are listed in [Supplemental Table S1](#).

CLONING AND EMBRYO TRANSFER. A detailed description of the methods used to clone piglets from genetically modified fibroblasts has been published previously (15). Briefly, after partial digestion of zona pellucida with 3.3 mg/ml pronase, oriented bisection of matured oocytes was performed manually under stereomicroscope with a microblade (AB Technology, Pullman, Western Australia, Australia). Each ooplast was attached with an *APOE* KO genetically modified fibroblast and fused in fusion medium (0.3 mol mannitol, 0.1 mmol MgSO₄ and 0.01% [w/v] polyvinyl alcohol) in a fusion chamber (BTX microslide 0.5-mm fusion chamber, model 450; BTX Instrument Division, Harvard Apparatus, Holliston, Massachusetts) with a single direct current pulse of 2.0 kV/cm for 9 μs. One hour later, each ooplast-*APOE* KO genetically modified cell pair was fused with another ooplast in activation medium (fusion medium supplemented with 0.1 mmol CaCl₂) by a single direct current pulse of 0.86 kV/cm for 80 μs. After incubation in porcine zygote medium 3 supplemented with 5 mg/ml cytochalasin B and 10 mg/ml cycloheximide for 4 h, the reconstructed embryos were cultured individually in well-of-the-wells made in 4-well dishes filled with porcine zygote medium 3 medium. Morulae and blastocysts were collected on days 5 and 6 and surgically transferred to Danish Landrace sows on day 4 or 5 after heat, which were observed 5 days after weaning. Five embryo transfers were performed with an average of 96 reconstructed embryos transferred per recipient sow (16). Successful pregnancies were diagnosed in 4 of the 5 recipient sows by ultrasonography on day 28. A total of 34 live-born piglets were delivered by caesarian section or vaginal birth 24 hours after induction with prostaglandin. Three of the cloned founders survived the neonatal period and were used for further breeding.

PATHOLOGY. Animals were intubated following sedation with 20 mg etomidate (Janssen-Cilag A/S, Birkerød, Denmark) intravenously, anesthetized using propofol (infusion rate 150 mg/h, B. Braun Melsungen AG, Melsungen, Germany) and fentanyl (infusion rate 1 mg/h, Hameln Pharmaceuticals Ltd., Gloucester, United Kingdom), and mechanically ventilated. To prevent clot formation, the animals were

anticoagulated with 10,000 IU heparin injection intravenously before being euthanized by exsanguination. Hearts, aortas and right iliofemoral arteries were excised and immersed in 4% phosphate-buffered formaldehyde for 24 h followed by immersion in cold phosphate-buffered saline.

Aortas were cut open longitudinally and divided into thoracic aorta and abdominal aorta at the site between the 6th and 7th intercostal arteries. Aortas and the right iliofemoral arteries were stained with Sudan IV (Sigma-Aldrich; 5 g/l in 96% ethanol for 5 min followed by 90 s washout in 96% ethanol) and *en face* images of the vessels were obtained using a digital scanner (Epson Perfection V600 Photo, Seiko Epson Corporation, Suwa, Nagano, Japan).

The percentage of intimal surface covered with lesions was determined by automated computer-assisted measurement of Sudan-IV stained (sudanolphilic) intima using ImageJ 1.48v (National Institutes of Health, Bethesda, Maryland). Lesions in the abdominal aorta were traced manually using ImageJ since most lesions were less sudanophilic.

Left anterior descending (LAD) coronary arteries were excised from the myocardium and the proximal 3 cm were divided into five 5-mm segments. The transversal segment, containing the most raised lesion (by macroscopic inspection) in the region proximal to the aortic trifurcation, was also obtained from each pig. Segments were paraffin-embedded and sectioned for histological analysis. LAD sections were stained with hematoxylin and eosin, and the lesions in the abdominal aorta sections were stained with elastin-trichrome.

Lesion sizes in the LAD were measured by computer-assisted planimetry. For the aortic plaques, maximum plaque-thickness was measured from the internal elastic lamina to the lumen.

We performed immunohistochemical staining for smooth muscle α -actin (SM α A) to detect SMCs in abdominal aorta lesions. Endogenous peroxidase activity was blocked by incubating the sections for 10 min with Hydrogen Peroxidase Block (Thermo Fisher Scientific) followed by blocking for nonspecific protein binding using Protein Block for 10 min. Sections were incubated with monoclonal mouse anti-human smooth muscle alpha actin antibody (Dako Denmark A/S, Glostrup, Denmark, cat. no. M0851, clone 1A4 [71 mg/l]) diluted 1:200 for 1 h at room temperature and visualized using horseradish peroxidase-conjugated secondary antibodies (Abcam, Cambridge, United Kingdom, EXPOSE detection IHC kit, cat. no. ab80436), 3,3'-diaminobenzidine for 7 min, and hematoxylin counterstain following the manufacturer's recommendations.

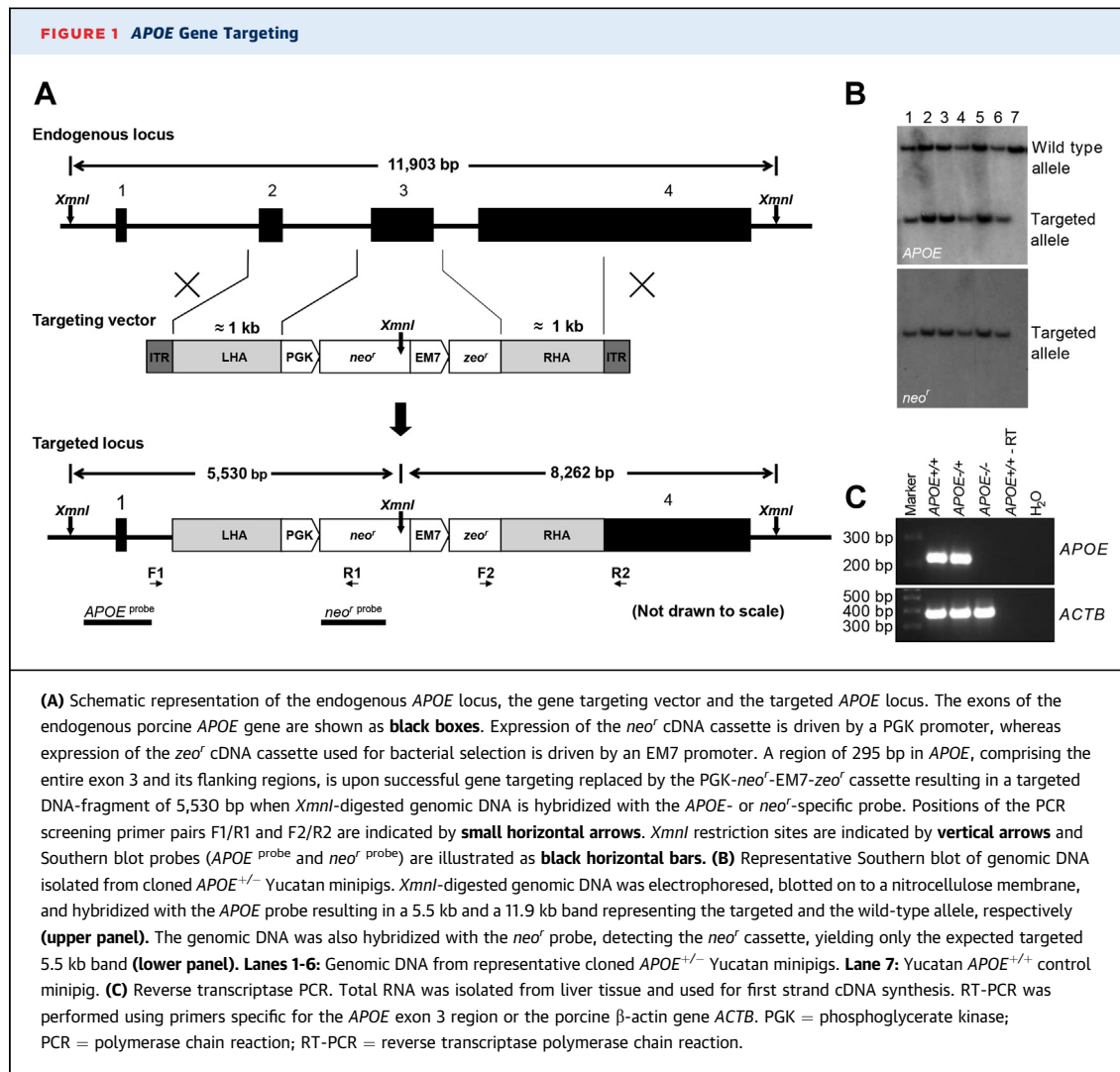
All microscopic analyses were performed blinded. Lesions were categorized according to the classification introduced by Virmani et al. (17) as normal, xanthoma, or progressive atherosclerotic lesions (pathological intimal thickening or fibroatheroma).

STATISTICS. All analyses were performed using Prism 6 (GraphPad Software Inc.). To compare 2 groups, we used the Student *t* test when data were normally distributed and Mann-Whitney U test when data were not normally distributed. Area under the curve was calculated by the trapezoidal method and compared to test for differences in time serial measurements. Plaque types were categorized as either progressive or nonprogressive types, and the frequency of progressive plaques was compared between groups using Fisher's exact test. $p < 0.05$ was considered significant. In all figures, an * indicates $p < 0.05$ and a ** indicates $p < 0.01$. Exact *p* values are given in figure legends.

RESULTS

TARGETED KO OF THE PORCINE APOE GENE. APOE comprises 2 structural domains separated by a hinge region. The N-terminal domain (amino acids 1-191) contains the receptor binding region and the C-terminal domain (amino acids 225-299) harbors the lipid binding region (18). We used rAAV-mediated gene targeting (14,19) to create Yucatan minipigs that lack APOE due to deletion of the entire exon 3 and disruption of the receptor binding region. An rAAV-APOE KO viral vector (Figure 1A) was generated to produce rAAV particles used for gene targeting in primary porcine fibroblasts isolated from newborn male Yucatan minipigs. The APOE^{+/-} gene targeted cell clones were validated by Southern blotting using an APOE-specific probe located upstream of the targeted region confirming deletion of exon 3, and a *neo*^r-specific probe detecting the *neo*^r region of the targeting vector to validate the absence of unwanted random integrations of the vector in the gene targeted cell clones.

CLONING OF APOE KO YUCATAN MINIPIGS. One of the APOE^{+/-} gene targeted cell clones was used as nuclear donor cells for somatic cell nuclear transfer by handmade cloning. All piglets were genotypically validated as APOE^{+/-} by Southern blotting (Figure 1B). Three APOE^{+/-} founders of the 34 cloned live-born male piglets survived the neonatal period. All 3 founders were healthy, and cytogenetical analysis performed on 1 of them revealed a seemingly normal karyotype (Supplemental Figure 1). The 3 APOE^{+/-} Yucatan boars were mated with wild-type Yucatan sows to obtain naturally bred F1 (APOE^{+/-}) individuals. The F1 minipigs were interbred to obtain F2

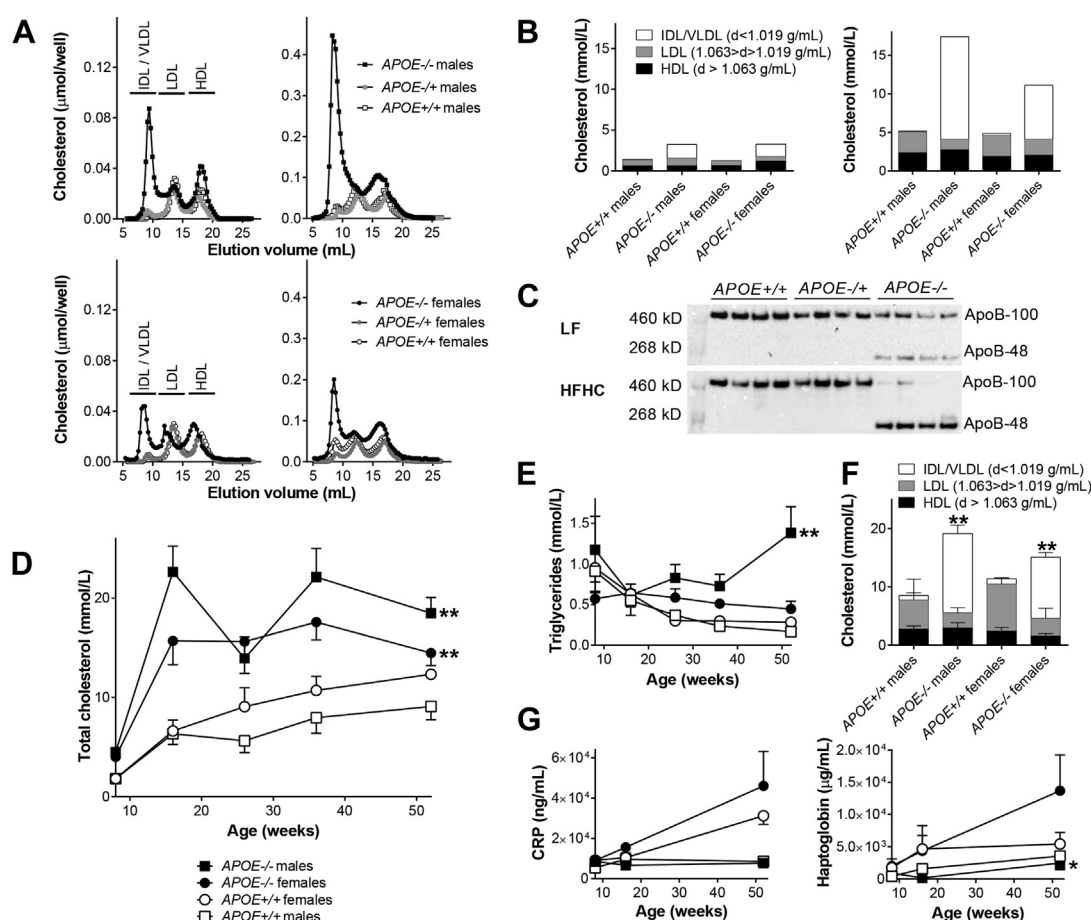


APOE^{+/+}, *APOE*^{+/-}, and *APOE*^{-/-} minipigs including 6 male *APOE*^{+/+}, 6 male *APOE*^{-/-}, 7 male *APOE*^{+/-}, 7 female *APOE*^{+/+}, 7 female *APOE*^{+/-}, and 5 female *APOE*^{-/-} minipigs which were included in the study. The expected level of *APOE* expression in the 3 different genotypes was validated by reverse transcriptase PCR performed on liver tissue isolated from the minipigs at euthanization (Figure 1C, Supplemental Figure 2).

ACCUMULATION OF CHYLOMICRON REMNANTS. At 8 weeks of age, when the animals were fed a LF standard diet, *APOE*^{-/-} minipigs exhibited significantly higher total cholesterol (4.28 ± 0.31 mmol/l) compared to both *APOE*^{+/-} (1.60 ± 0.08 mmol/l) and *APOE*^{+/+} (1.81 ± 0.08 mmol/l) ($p < 0.001$). Size-exclusion chromatography (Figure 2A, left column) and ultracentrifugation (Figure 2B, left column) showed that this was explained by accumulation of lipoproteins of IDL/VLDL size and density. The

APOE^{-/-} phenotype was intensified after 8 weeks on HFHC diet (Figure 2A, right column and Figure 2B, right column). Western blot of APOB showed accumulation of APOB-48 in *APOE*^{-/-} pigs on LF diet and accentuated APOB-48 accumulation after HFHC diet feeding (Figure 2C, Supplemental Figure 3). This indicates that the accumulating lipoproteins were chylomicron remnants because pigs, like humans, only produce APOB-48-containing lipoproteins in the small intestine (20).

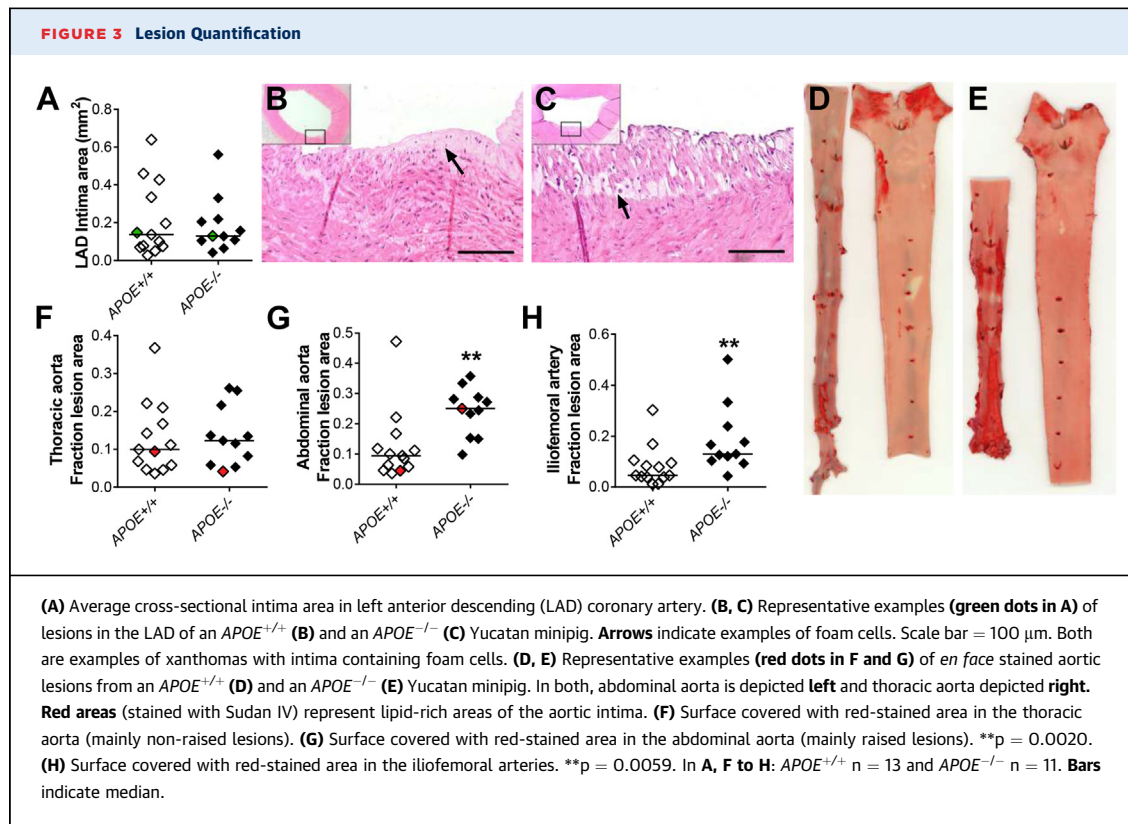
PERSISTENT INCREASE IN REMNANT CHOLESTEROL. To induce atherosclerosis, *APOE*^{+/+} and *APOE*^{-/-} minipigs were fed an HFHC diet until euthanization at the age of 52 weeks. At this age, mean weights were 55.2 ± 3.5 kg for male *APOE*^{+/+}, 48.7 ± 2.9 kg for male *APOE*^{-/-} ($p = 0.331$), 59.1 ± 3.1 kg for female *APOE*^{+/+}, and 58.8 ± 4.04 kg for female *APOE*^{-/-} minipigs ($p = 0.899$). Total cholesterol rose rapidly in *APOE*^{-/-}

FIGURE 2 Characterization of Cholesterol Profiles and Inflammatory Markers

(A) Size-exclusion chromatography on plasma pools of cholesterol in *APOE*^{+/+}, *APOE*^{+/-}, and *APOE*^{-/-} Yucatan male and female minipigs on low-fat (LF) diet (left column) and after 8 weeks on high-fat high-cholesterol (HFHC) diet (right column). (B) Ultracentrifugation characterization of cholesterol on plasma pools from *APOE*^{+/+} and *APOE*^{-/-} Yucatan minipigs on LF diet (left) and after 8 weeks on HFHC diet (right). (C) Western blot of APOB in plasma from *APOE*^{+/+}, *APOE*^{+/-}, and *APOE*^{-/-} Yucatan minipigs on LF diet (upper) and on HFHC diet (lower). (D) Enzymatic measurement of total cholesterol at 8 (on LF diet), 16, 26, 36, and 52 (all on HFHC diet) weeks of age. ****p** = 0.0022 for *APOE*^{+/+} male versus *APOE*^{-/-} male minipigs and **p** = 0.0051 for *APOE*^{+/+} female versus *APOE*^{-/-} female minipigs. (E) Enzymatic measurement of triglycerides at 8 (on LF diet), 16, 26, 36, and 52 (all on HFHC diet) weeks of age. ****p** = 0.0087 for *APOE*^{+/+} male versus *APOE*^{-/-} male minipigs. (F) Ultracentrifugation characterization of cholesterol in plasma from *APOE*^{+/+}, *APOE*^{+/-}, and *APOE*^{-/-} Yucatan minipigs on HFHC diet at 52 weeks of age. ****p** = 0.0022 for IDL/VLDL between *APOE*^{+/+} male and *APOE*^{-/-} male minipigs and **p** = 0.0025 for IDL/VLDL between *APOE*^{+/+} female and *APOE*^{-/-} female minipigs. **p** = 0.0303 for HDL and **p** = 0.0025 for LDL between *APOE*^{+/+} female and *APOE*^{-/-} female minipigs. (G) C-reactive protein (CRP) (left) and haptoglobin (right) levels in *APOE*^{+/+} and *APOE*^{-/-} Yucatan minipigs at 8, 16, and 52 weeks of age. ***p** = 0.0411 for *APOE*^{+/+} male versus *APOE*^{-/-} male minipigs. (D to G) *APOE*^{+/+} males **n** = 6, *APOE*^{+/+} females **n** = 7, and *APOE*^{-/-} males **n** = 6, and *APOE*^{-/-} females **n** = 5. Error bars indicate SEM. HDL = high-density lipoprotein; IDL = intermediate-density lipoprotein; VLDL = very-low-density lipoprotein.

minipigs after 8 weeks on HFHC diet and remained higher throughout the whole period at a level ≈ 18 mmol/l compared with *APOE*^{+/+} minipigs that reached a level ≈ 10 mmol/l (Figure 2D). The differences were statistically highly significant when comparing areas under the curve. Triglycerides were also numerically higher in *APOE*^{-/-} minipigs, although only statistically significant in male minipigs

(Figure 2E). Lipoprotein ultracentrifugation performed on plasma obtained at 52 weeks of age showed significantly increased IDL/VLDL sub-fractions of cholesterol in *APOE*^{-/-} compared with *APOE*^{+/+} minipigs (Figure 2F). Female *APOE*^{-/-} minipigs had significantly lower LDL and high-density lipoprotein (HDL) cholesterol compared with *APOE*^{+/+} female minipigs.



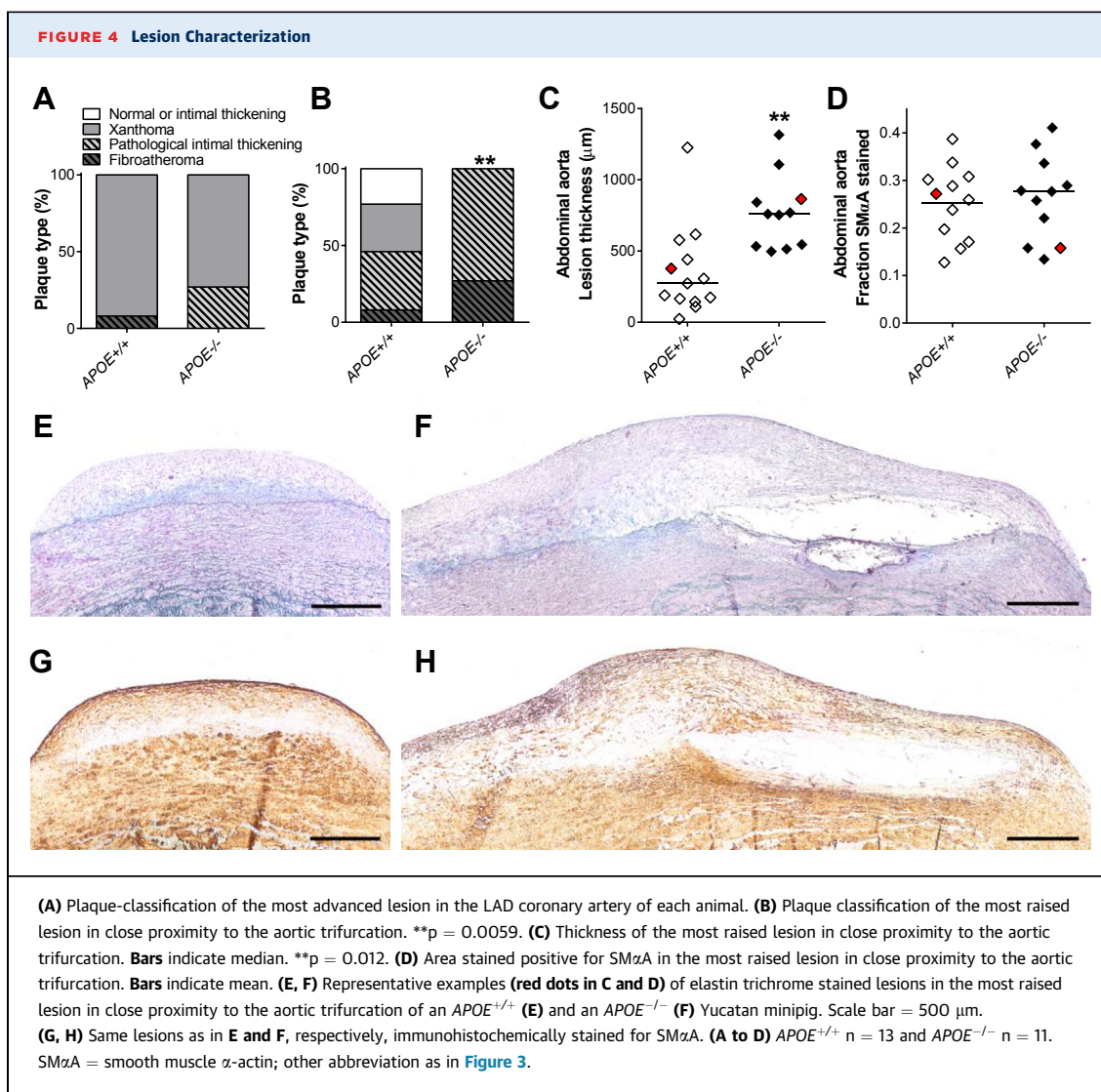
REMNANT HYPERLIPOPROTEINEMIA NOT ASSOCIATED WITH INCREASED INFLAMMATION. Recent genetic population studies in humans showed a link between gene variants causing higher levels of remnant lipoproteins and plasma levels of CRP in humans (6), but neither CRP nor haptoglobin, another acute phase reactant, was significantly different between $APOE^{-/-}$ and $APOE^{+/+}$ minipigs (Figure 2G). Although levels increased with age in female minipigs, values were lower than those seen in pigs with infectious disease (21).

ACCELERATED ATHEROSCLEROSIS. Atherosclerosis was analyzed in several arterial beds in the HFHC-fed $APOE^{-/-}$ and $APOE^{+/+}$ minipigs. Lesions in the LAD coronary artery were generally small and the mean intimal area was unaffected by the lack of APOE and the resulting remnant lipoproteinemia (Figures 3A to 3C). Thoracic aortas were largely covered by non-raised sudanophilic fatty streaks (Supplemental Figures 4A and 4B). The extent, judged by *en face* measurement (Figures 3D to 3F), did not differ between $APOE^{-/-}$ and $APOE^{+/+}$ minipigs. In contrast, in both the abdominal aortas and iliofemoral arteries where wild-type minipigs develop more advanced atherosclerosis, development of lesions were significantly accelerated in $APOE^{-/-}$ minipigs

showing 2.7-fold (Figure 3G) and 2.8-fold (Figure 3H, Supplemental Figures 4C and 4D) increases in median surface coverage in the 2 arterial beds, respectively.

We further characterized lesion types in histological sections from the LAD and abdominal aortas (Figure 4A). In serial sections of LADs, the most advanced lesion identified in each minipig was a xanthoma in all $APOE^{+/+}$ minipigs except 1 female that had a fibroatheroma and was an outlier in all atherosclerosis measurements. In $APOE^{-/-}$ minipigs, the most advanced lesion types were xanthomas (73%) and pathological intimal thickening (27%). The frequency of progressive lesions (pathological intimal thickening and fibroatheroma) was not significantly different between $APOE^{+/+}$ and $APOE^{-/-}$ minipigs.

Histology of the most raised lesion identified in the distal part of the abdominal aorta (Figures 4B, 4E, and 4F) showed normal or adaptive intimal thickening (23%), xanthomas (31%), pathological intimal thickenings (38%), and fibroatheromas (8%, $n = 1$) in $APOE^{+/+}$ minipigs. In $APOE^{-/-}$ minipigs, this was significantly skewed towards more advanced lesion stages with all showing progressive atherosclerotic lesions, 73% of which were pathological intimal thickenings and 27% of which were fibroatheromas.



Lesion thickness of the abdominal aorta plaque was also significantly increased in $APOE^{-/-}$ compared with $APOE^{+/+}$ minipigs (Figure 4C). All fibroatheromas were of the thick fibrous cap type and no thrombotic complications were observed.

Finally, as APOE has been shown to regulate SMC proliferation in mice, we addressed the possibility that exaggerated SMC accumulation could explain the acceleration of advanced atherosclerosis. Sections of lesions in the abdominal aortas were stained for smooth muscle α -actin, but the percentage of SM α A-positive area did not differ between $APOE^{-/-}$ and $APOE^{+/+}$ minipigs (Figures 4D, 4G, and 4H)—neither did it differ when correlated to lesion size (data not shown).

DISCUSSION

Defects or deficiency of APOE cause familial dysbetalipoproteinemia in humans, characterized by accumulations of VLDL, IDL, and chylomicron remnants and a higher risk of premature atherosclerotic disease (22). Whereas the human disease rarely becomes symptomatic before adulthood, the same genetic defect in mice is aggressive with rapid development of severe atherosclerosis. This is partly explained by the higher reliance on APOE for lipoprotein clearance in mice compared to humans (3), and consequently the much higher plasma cholesterol concentration resulting from its loss. However, it is also possible that a particular

atherogenicity of the accumulating lipoproteins or loss of intrinsic anti-atherosclerotic effects of APOE identified in the mouse may contribute (4). The *ApoE*^{-/-} mouse model is widely applied for experimental atherosclerosis because of its rapid lesion development, but it has an unusual etiology of atherosclerosis and its translational value is controversial (23).

In the present study, we created a model of dysbetalipoproteinemia in minipigs, where the distribution of lipoprotein classes and the lack of hepatic APOB editing are closer to human physiology, and examined the impact of remnant lipoproteinemia caused by *APOE* KO on the development of atherosclerosis. We found that, as in APOE-deficient humans, heterozygotes generally had normal lipid levels, but homozygotes showed large accumulations of APOB-48 remnant-type lipoproteins. Surprisingly, despite the severe remnant lipoproteinemia, which caused an approximate doubling of plasma total cholesterol, the effect on atherosclerosis development was modest and only present in the most atherosclerosis-susceptible arterial beds. APOE deficiency accelerated atherosclerosis in the abdominal aorta and iliofemoral arteries, which are predilection sites for progressive atherosclerosis in pigs and humans, but no effect was detected in thoracic aortas and the LAD where lesions were restricted to mainly xanthomas. In this way, the phenotype of *APOE*^{-/-} minipigs was clearly different from that of *D374Y-PCSK9* transgenic minipigs, which we have previously described using a similar diet composition, genetic background and follow-up time as in the present study. *D374Y-PCSK9* minipigs exhibited similar increases in plasma cholesterol, but cholesterol was mainly carried by APOB-100-containing LDLs and larger lipoproteins, and atherosclerosis was significantly accelerated in all arterial beds examined, including the thoracic aorta and the LAD (9). The coronary atherosclerosis phenotype of the *APOE*^{-/-} minipig model was also much milder than reported for several other porcine models, in which atherosclerosis is driven by increases in APOB100-containing lipoproteins, such as the diabetic Yorkshire model and pigs with spontaneous or induced defects in LDL-clearance (7,24). Part of the difference to those models may also lie in the induction of diabetes, differences in diet composition, including the use of cholic acid to accentuate atherosclerosis, and potential differences in atherosclerosis susceptibility of the background strain.

Our findings indicate that the remnant lipoproteinemia of *APOE*^{-/-} minipigs is not efficient in

initiating atherosclerosis. However, it may contribute to further progression of pre-existing lesions, either alone or together with the loss of direct anti-atherogenic effects of APOE. One explanation for the lack of effect in vascular beds with early atherosclerosis could be abolishment of APOE-mediated binding of lipoproteins to intimal proteoglycans. APOE, similar to APOB, contains motifs that facilitate lipoprotein retention in the vascular wall (25). Loss of this binding type may diminish the atherogenicity of APOE-deficient lipoproteins during initiation of atherosclerosis, but the effect is likely lost with further plaque development as has previously been shown for APOB (26).

STUDY LIMITATIONS. Further studies are needed to understand by which mechanisms APOE deficiency accelerates progressive atherosclerotic lesions. We addressed the possibilities that remnant lipoproteins could exert a pro-inflammatory effect in the body, as suggested by human genetic studies (6), or that loss of APOE would lead to increased SMC proliferation, but found no evidence that this was the underlying explanation.

CONCLUSIONS

We showed that targeted gene KO of *APOE* in minipigs causes severe remnant lipoproteinemia similar to human familial dysbetalipoproteinemia. We found that this increased progressive atherosclerotic lesions but not xanthoma formation. Further research in this model may pave the way for better understanding of the pathophysiological effects of remnant lipoproteinemia in atherosclerosis.

ACKNOWLEDGMENTS The authors would like to thank Bert Vogelstein and Kenneth W. Kinzler, Johns Hopkins University, Baltimore, Maryland, for kindly providing the pNeDaKO plasmid vector and Dr. R. Jude Samulski and the UNC Vector Core Facility, Chapel Hill, North Carolina, for the rAAV packaging. They would also like to thank Peter M. Kragh and Lisbeth D. Schrøder, Trine S. Petersen, Christian Knudsen, Lisa Maria Røge, Dorte Qualmann, Zahra P. Nasr, Anette M. Pedersen, Janne Adamsen, Klaus Villemoes, and Ruth Kristensen for skilled technical assistance.

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PERSPECTIVES

COMPETENCY IN MEDICAL KNOWLEDGE: Remnant cholesterol increases atherosclerosis and risk of ischemic heart disease. Existing large-animal models of dyslipidemia and atherosclerosis, both diet-induced as well as genetically modified animal models, display cholesterol profiles resembling familial hypercholesterolemia. We here report the first large-animal model of *APOE*-deficiency and remnant lipoproteinemia modelling the dyslipidemia seen in patients with familial dysbetalipoproteinemia.

TRANSLATIONAL OUTLOOK: The human-like features of this animal model, in terms of size, lipoprotein metabolism, and histopathological features of atherosclerosis, may pave the way for better understanding of dysbetalipoproteinemia and the mechanisms involved in atherogenesis of remnant lipoproteins observed in humans.

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KEY WORDS apolipoprotein E, atherosclerosis, pig, remnant cholesterol dysbetalipoproteinemia

APPENDIX For supplemental text, references, figures, and table, please see the online version of this article.