Transgenic Overexpression of Tissue-Nonspecific Alkaline Phosphatase (TNAP) in Vascular Endothelium Results in Generalized Arterial Calcification

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Background—Ectopic vascular calcification is a common condition associated with aging, atherosclerosis, diabetes, and/or chronic kidney disease. Smooth muscle cells are the best characterized source of osteogenic progenitors in the vasculature; however, recent studies suggest that cells of endothelial origin can also promote calcification. To test this, we sought to increase the osteogenic potential of endothelial cells by overexpressing tissue-nonspecific alkaline phosphatase (TNAP), a key enzyme that regulates biomineralization, and to determine the pathophysiological effect of endothelial TNAP on vascular calcification and cardiovascular function.

Methods and Results—We demonstrated previously that mice transgenic for ALPL (gene encoding human TNAP) develop severe arterial medial calcification and reduced viability when TNAP is overexpressed in smooth muscle cells. In this study, we expressed the ALPL transgene in endothelial cells following endothelial-specific Tie2-Cre recombination. Mice with endothelial TNAP overexpression survived well into adulthood and displayed generalized arterial calcification. Genes associated with osteochondrogenesis (Runx2, Bglap, Sp51, Opg, and Col2a1) were upregulated in the aortas of endothelial TNAP animals compared with controls. Lesions in coronary arteries of endothelial TNAP mice showed immunoreactivity to Runx2, osteocalcin, osteopontin, and collagen II as well as increased deposition of sialoproteins revealed by lectin staining. By 23 weeks of age, endothelial TNAP mice developed elevated blood pressure and compensatory left ventricular hypertrophy with preserved ejection fraction.

Conclusions—This study presented a novel genetic model demonstrating the osteogenic potential of TNAP-positive endothelial cells in promoting pathophysiological vascular calcification. (J Am Heart Assoc. 2015;4:e002499 doi: 10.1161/JAHA.115.002499)

Key Words: arteriosclerosis • cardiovascular diseases • endothelium • enzymes • lesion

Vascular calcification is a pathological hallmark of advanced age, atherosclerosis, diabetes, and chronic kidney disease.1 Coronary artery calcification is an independent predictor of cardiovascular events,2–9 whereas calcification of peripheral arteries is associated with all-cause mortality.10 Consequently, calcification of vascular structures represents a significant medical problem.

Mechanisms of vascular calcification are complex; they typically require the relaxation of calcification inhibitors with concomitant increase in activators of biomineralization, both of which are tightly regulated at the cellular level.11 Transdifferentiation of vascular smooth muscle cells (VSMCs) is a well-characterized phenomenon that shares common molecular pathways with skeletal mineralization.12 During this process, VSMCs upregulate expression of osteogenic transcription factors and begin secreting matrix vesicles, membrane microparticles that initiate hydroxyapatite crystalization.13,14 Endothelial cells (ECs) could also promote vascular calcification, although evidence of this is still limited. Epidemiological studies suggest that circulating endothelial progenitor cells that express osteogenic markers (osteocalcin and/or alkaline phosphatase [AP]) correlate with vascular calcification in patients with atherosclerosis.15–18 Moreover, the
greater numbers of such cells are detected in patients with diabetes.19 Lineage-tracing studies show that ECs contribute to the osteoprogenitor pool in aortic lesions in mice with experimental vascular calcification associated with matrix γ-carboxyglutamic acid protein deficiency (Mgp knockout mice) and in the model of genetically determined diabetes (Akita mice).20 In a model of mitral valve calcification, ECs positioned along mechanically stretched valves express the osteoblastic marker osteocalcin, indicating their transdifferentiation.21 Outside of arterial and valvular examples, ECs isolated from solid tumors of the prostate, in which ectopic calcification colocalizes with endothelial marker CD31, clearly show ex vivo differentiation plasticity toward chondrocytes expressing cartilage-specific Col2a1 and Sox9 or osteoblasts expressing osteocalcin and osteopontin.22

Tissue-nonspecific AP (TNAP) is the key enzyme that regulates availability of extracellular pyrophosphate (PPi), an inhibitor of biomineralization.23 Reduction in extracellular PPi promotes vascular calcification in animals,24–26 and the decreased levels of circulating plasma PPi have been associated with human genetic or metabolic conditions that predispose affected persons to vascular calcification.25,27 Conversely, TNAP upregulation has been observed in calcified human arteries.28–30 Our recent study of transgenic mice overexpressing TNAP in VSMCs31 provided strong genetic evidence that TNAP is sufficient to induce severe vascular calcification and consequent cardiovascular complications—hypertension and heart failure—that drastically reduce the life span of affected animals. In this study, we used a similar strategy to manipulate expression of TNAP specifically in ECs to examine whether endothelial TNAP (eTNAP) might play a critical role in vascular calcification.

Materials and Methods

Animals

Transgenic mice expressing Cre recombinase driven by Tie2 gene promoter (Tie2-Cre) were obtained from The Jackson Laboratory (Bar Harbor, ME; stock 8863). Characterization of this mouse strain has shown panendothelial expression of the Cre transgene.32 HprtALPL knock-in mice were recently described.31 This mouse strain has a site-specific transgenic insertion of the ALPL gene into the Hprt locus on the X chromosome. The expression construct contains a ubiquitous CAG promoter, a floxed “stop” cassette, and ALPL cDNA encoding human TNAP. Breeding of homozygous HprtALPL/ALPL female mice with heterozygous Tie2-Cre male mice results in the excision of the stop cassette in 50% of male offspring and activation of constitutive transgene expression in the ECs. Using this breeding scheme, male mice were produced and genotyped by polymerase chain reaction with Cre-specific primers (forward: GCG GTC TGG CAG TAA AAA CTA TC; reverse: GTG AAA CAG CAT TGC TGT CAC TT) and an internal positive control primers (forward: CTA GGC CAC AGA ATT GAA AGA TCT; reverse: GTA GGT GGA AAT TCT AGC ATC ATC C). All animal studies were approved by the Sanford Research institutional animal care and use committee and complied with the National Institutes of Health guidelines for humane treatment of laboratory animals.

Whole-Body X-Ray Imaging

Whole body x-ray images were obtained using small animal imager Carestream In-Vivo Xtreme (Carestream Health). Mice were anesthetized by inhalation of 1.5% isoflurane in oxygen.

Blood Chemistry

Sera or heparin plasma were prepared from venous blood collected from right ventricles of anesthetized mice. All mice were fasted at least 5 hours before blood collection. AP activity, calcium, phosphorus, and creatinine were determined using ADVIA 1800 Clinical Chemistry analyzer (Siemens) with specific reagents: ALPAMP, CA_2, IP, and CREA. Inorganic PPi was measured in lithium heparin plasma according to the method described previously.33

Histology

AP activity was detected in fresh tissues embedded in optimal cutting temperature compound and cryosectioned at 5 μm. Sections were incubated in 5-bromo-4-chloro-3-indolyl-phosphate/nitro blue tetrazolium phosphatase substrate solution (KPL Inc.) for 5 minutes, rinsed in water, and fixed in 10% neutral buffered formalin. ECs were detected with FITC-labeled isoelectin B4 (IB4) from Bandeiraea simplicifolia (item FL1201; Vector Laboratories). Osteocalcin was detected with rabbit polyclonal antibody (item M173; Tanaka Bio Inc) followed by Alexa 488–labeled anti-rabbit IgG secondary antibody (item 711-546-152; Jackson ImmunoResearch). Runx2 was detected with rat monoclonal antibody (item MAB2006; R&D Systems) followed by Alexa 546–labeled anti-rat IgG antibody (item 712-586-153; Jackson ImmunoResearch). Collagen II was detected with sheep polyclonal antibody (item AF3615; R&D Systems) followed by Alexa 594–labeled anti-sheep IgG antibody (item 713-586-147; Jackson ImmunoResearch). Corresponding secondary antibodies alone served as negative controls for osteocalcin, Runx2, and collagen II staining. Osteopontin was detected with goat polyclonal antibody directly labeled with phycoerythrin (item IC808P; R&D Systems); phycoerythrin-labeled goat IgG (item IC108P; R&D Systems) served as a negative control.
acid was detected with FITC-labeled *Sambucus nigra* lectin (item FL1301; Vector Laboratories). For general histology, mice were perfused with neutral buffered formalin; tissues were dissected and stored in the same fixative (Formal-Fixx; Thermo Scientific). Tissues were embedded in paraffin, sectioned, and stained with hematoxylin and eosin, Alizarin Red, Masson’s trichrome, and von Kossa stains in the Sanford Research Molecular Pathology Core, following standard protocols.

**Real-Time Polymerase Chain Reaction**

RNA was isolated from the aortas using a Maxwell 16LEV simplyRNA Tissue kit (item AS1280; Promega). RNA quantity and quality were determined using a NanoDrop 1000 (Thermo Scientific). Equal amounts of RNA from each sample were converted to cDNA using GoScript reverse transcriptase (item A501C; Promega) with random hexamer primers. Gene expression was evaluated using primers designed by Integrated DNA Technologies (*Gapdh* Mm.PT.39a.1; *Alpl* Mm.PT.56a.8794492; *Sp7* Mm.PT.58.10898265; *Bmp2* Mm.PT.58.10419414; *Runx2* Mm.PT.58.41866893; *Sox9* Mm.PT.58.42739087; *Bglap* Mm.PT.58.9119501.g; *Spp1* Mm.PT.56.43709208; *Mgp* Mm.PT.58.32674923; *Acan* Mm.PT.58.10174685; *Col2a1* Mm.PT.58.41487458; *Opq* Mm.PT.56a.414946; *Tnfsf11* Mm.PT.55a.29202697) and SYBR green polymerase master mix reagent from Qiagen. Gene expression was normalized using the housekeeping gene *Gapdh* and expressed in arbitrary units, calculated as follows: $2^{\text{CtGene} - \text{CtGapdh}} \times 10^{5}$, where CtGene is the threshold cycle for target gene amplification, and CtGapdh is the threshold cycle for a reference gene (*Gapdh*).

**Physiology**

Echocardiography was performed using a Vevo2100 imaging system (FujiFilm Visual Sonics) equipped with an ultra–high-frequency linear array solid state transducer (MS550D 22 to 55 MHz). Mice were anesthetized with 0.8% to 1% isoflurane, and a parasternal short-axis view was obtained in B-mode and recorded in M-mode. M-mode echocardiograms were analyzed by tracing myocardial wall movement over 3 to 5 cardiac cycles to measure heart rate and left ventricular (LV) diameter in systole and in diastole and to calculate ejection fraction, cardiac output, and LV mass.

Blood pressure was measured directly with a pressure transducer (SPR-671; Millar Instruments). Right carotid artery catheterization was performed on mice anesthetized with 2% isoflurane, and data were recorded over 10 minutes at the maintenance level of 1.25% isoflurane in oxygen. On completion of data collection, mice were deeply anesthetized by 5% isoflurane inhalation prior to blood collection and euthanasia by exsanguination.

**Statistics**

Data were expressed as mean±SD or mean±SEM. GraphPad Prism version 5.0 (GraphPad Software) statistical software was used to analyze data. Two groups were compared by Student *t* test or nonparametric Mann–Whitney test when unequal variances were observed. One-way ANOVA models were calculated to compare multiple groups, followed by Bonferroni-adjusted multiple comparisons. The Kruskal–Wallis test was used to compare multiple groups when unequal variances were observed. This was followed by Dunn’s multiple comparisons test. The significance was accepted at *P*<0.05.

**Results**

**Panendothelial Expression of TNAP**

To test the hypothesis that upregulation of TNAP in ECs would be sufficient to cause vascular calcification, we used a conditional knock-in mouse model that overexpresses TNAP in a Tie2-Cre recombinase–dependent manner. Male offspring mice were used in all experiments to avoid random X chromosome inactivation that affects levels of X-linked *ALPL* transgene expression in female mice. The eTNAP mice were born at an expected Mendelian ratio (36 of 73 male pups) and survived into adulthood. To confirm TNAP overexpression, we examined the AP activity in eTNAP and wild-type (WT) mice aged 8 weeks by incubating freshly prepared cryosections of various tissues in chromogenic AP substrate followed by formalin fixation. Using this method, increased AP activity was detected in all tissues of eTNAP compared with WT mice (Figure 1A). AP activity in eTNAP mice was localized to the luminal side of the aorta and vascular networks of heart, lung, kidney, liver, small intestine, and pancreas. A diffuse pattern of AP activity was observed in the parenchyma of lymphoid organs (spleen, thymus, and lymph nodes) of eTNAP mice but was absent in WT mice. In the heart, AP activity, both endogenous (detected in WT mice) and overexpressed (present in eTNAP mice), mirrored the characteristic pattern of capillary network identified by IB4 staining34 (Figure 1B). To rule out possible side effects of eTNAP overexpression on skeletal development, whole-body x-ray imaging was performed at 8 weeks of age and showed the lack of detectable skeletal abnormalities in eTNAP mice (Figure 1C).

Serum AP activity was also greatly upregulated in eTNAP compared with WT mice in all age groups examined (aged 8, 13, and 23 weeks), whereas serum calcium, phosphorus, and creatinine were comparable within each age group (Table 1). Plasma PPi concentrations were not significantly different between eTNAP and WT mice aged 13 weeks (Table 1). In this study, plasma samples were not collected from mice aged 8
and 23 weeks, thus we could not perform the PPi assay in those age groups due to artificial increase of PPi, released from platelets during blood coagulation.

Histological Phenotype of eTNAP Mice

Histological examination (Alizarin Red staining) revealed the presence of calcified lesions in all tissues in eTNAP mice aged 13 weeks; no lesions were detected in WT mice (Figure 2). Calcified lesions in eTNAP mice were located in the arteries of the heart, kidney, mesentery, pancreas, and spleen as well as in the splenic capsule and lung parenchyma. Only minor lesions were detected in the aortas of eTNAP (distal thoracic aortas were sampled). No extraskeletal calcification was detected in preweaning-aged eTNAP mice (1 week) by Alizarin Red staining; a panel of tissues included heart, liver, kidney, pancreas, spleen.

Table 1. Blood Chemistry

| Parameter   | 8 Weeks | 13 Weeks | 23 Weeks |
|-------------|---------|----------|----------|
| AP, U/L     | WT (n=7) | eTNAP (n=7) | WT (n=8) | eTNAP (n=6) | WT (n=10) | eTNAP (n=7) |
| Ca, mg/dL   | 8.9±0.5  | 9.0±0.6  | 9.1±0.7  | 9.0±0.3  | 8.8±0.5  | 9.0±0.4  |
| Pi, mg/dL   | 6.5±1.5  | 6.8±1.1  | 6.1±0.9  | 5.4±0.6  | 7.1±1.3  | 6.2±0.6  |
| Creatinine, mg/dL | 0.17±0.07 | 0.14±0.05 | 0.13±0.04 | 0.11±0.03 | 0.11±0.04 | 0.11±0.02 |
| PPi, µmol/L | 1.7±1.0† | 1.7±0.8† | 1.7±1.0† |

Data are shown as mean±SD. AP indicates alkaline phosphatase; Ca, calcium; eTNAP, endothelial tissue-nonspecific alkaline phosphatase; P, phosphorus; PPi, pyrophosphate; WT, wild type.

*P<0.05.
*P<0.01 vs WT, Kruskal–Wallis test followed by Dunn’s post hoc pairwise comparisons within each age group. †n=4. ‡n=7.

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pancreas, spleen, and bone and served as a positive control (Figure 3).

The distribution and vascular localization of the lesions in different tissues of adult eTNAP mice were examined by hematoxylin and eosin staining. The 5 oldest (aged 24 weeks) eTNAP mice and 6 WT controls were perfusion fixed with formalin, and tissues were dissected after fixation. Three paraffin blocks were prepared from each mouse. The first block contained pancreas, mesentery, and spleen; the second block contained liver (left lateral lobe) and left kidney; and the third block contained mid–cross-section of the heart, left lung, and distal thoracic aorta. The presence of the lesions in different vascular beds and tissues in eTNAP mice was detected as basophilic inclusions in hematoxylin and eosin sections and confirmed by Alizarin Red staining. Consistent with observations in mice aged 13 weeks, calcium was absent in WT mice (not shown) and present in multiple tissues from eTNAP mice (histological findings are summarized in Table 2 and illustrated in Figure 4). In this sample, no lesions were detected in the aortas (n=3; aortas were not present in

Figure 2. Calcification in eTNAP mice aged 13 weeks. Alizarin Red staining of the aorta, heart, lung, liver, kidney, spleen, mesentery, pancreas, thymus, and intestine from WT and eTNAP mice. eTNAP indicates endothelial tissue-nonspecific alkaline phosphatase; WT, wild type.
the plane of section from 2 other mice). Arterial lesions were present in the mesenteric sections from all 5 mice, and calcification affected arteries of all sizes in every animal (Figure 4A through 4C). In addition, lesions were detected in the large veins of the mesentery (Figure 4D). In the pancreas, lesions were present in the arteries (Figure 4E and 4F) and were not detected in the veins of the same size (Figure 4F). Similarly, multiple lesions were detected in the arteries of the spleens of eTNAP mice (Figure 4G and 4H) and were not detected in the veins draining the spleen. Splenic stromal elements and capsules were also extensively calcified (Figure 4I and 4J). Coronary arteries were affected in every mouse (Figure 4K and 4L); no lesions were detected in the coronary veins. In the lung, neither pulmonary arteries nor veins were involved; however, multiple small round calcification nodules were present throughout the whole parenchyma (Figure 4M). In this histological sample of the kidneys, multiple small lesions were present; however, the sectioning plane did not capture any larger arteries or veins. The small size of the calcified lesions in the kidney did not allow for their clear vascular definition (Figure 4N and 4O). Based on our histological survey, we concluded that calcification primarily affected peripheral and coronary arteries of eTNAP mice.

In younger adult mice, we found that although small foci of calcification can be detected in the en face preparations of the aorta of eTNAP mice as early as 8 weeks of age (Figure 5A), coronary arteries were more extensively calcified compared with the aorta, as shown by Alizarin Red staining of the base of the heart at the level of the aortic sinus (Figure 5B). To investigate whether arterial lesions in eTNAP mice originate in direct contact with ECs, we examined the

Table 2. Tissues Affected by Calcification in eTNAP Mice Aged 24 Weeks (n=5)

| Tissue    | Arteries (Size) | Panel* | Veins (Size) | Panel* | Other Sites                  | Panel* |
|-----------|-----------------|--------|--------------|--------|------------------------------|--------|
| Aorta†    | n.d.            |        |              |        |                              |        |
| Mesentery | Large, medium, small | A, B, C | Large D      | n.d.   |                              | n.d.   |
| Pancreas  | Medium, small   | E, F   | n.d.         | n.d.   |                              |        |
| Spleen    | Medium, small   | G, H   | n.d.         | Stroma, capsule | I, J  |        |
| Heart     | Medium, small   | K, L   | n.d.         | n.d.   |                              |        |
| Lung      | n.d.            |        | n.d.         | Parenchyma | M   |        |
| Kidney    | Small‡          | N      | Small‡ O     | n.d.   |                              |        |

eTNAP indicates endothelial tissue-nonspecific alkaline phosphatase; n.d., not detected.
*shown in Figure 4.
†n=3.
‡Unclear morphological definition.
consecutive tissue sections of coronary arteries from mice aged 13 weeks by AP and IB4 staining (Figure 5C). Lesions in coronary arteries appeared as protrusions into the arterial lumen and were associated with AP- and IB4-positive cells. The presence of vascular calcification in eTNAP mice aged 13 weeks was confirmed by von Kossa staining, which showed a spectrum of phenotypes. The earliest lesions appeared to originate in the subendothelial space and were negative for von Kossa staining (Figure 5D, left panel); a granular pattern of von Kossa staining was detected in the matrix of intermediate lesions (Figure 5D, middle panel). Advanced lesions contained a von Kossa–positive core and, in some cases, were accompanied by calcification of the elastic lamina and medial layer (Figure 5D, right panel). In hematoxylin and eosin–stained sections, basophilic inclusions were detected in all arterial lesions, including early subendothelial and late

Figure 4. Histopathology of endothelial tissue-nonspecific alkaline phosphatase mice aged 24 weeks (hematoxylin and eosin staining). Arteries of the mesentery: large (A), medium (B), and small (C) diameter. D, Large vein in the mesentery. Arteries of the pancreas: medium (E) and small (F). Arteries of the spleen: medium (G) and small (H). Nonvascular lesions in the splenic stroma (I) and capsule (J). Coronary arteries: medium (K) and small (L). M, Parenchyma of the lung. N and O, Lesions in the kidney.
neointima-encapsulated lesions (Figure 5E). Masson’s trichrome staining highlighted the matrix core of the early lesions (light-blue staining) surrounded by cells of endothelial morphology (Figure 5F, top panels), whereas late lesions featured more complex morphology with foci of irregular matrix inclusions and collagen deposits (Figure 5F, bottom panels). In many histological sections at high magnification, lesions displayed lamellar characteristics of their core region (Figure 5B, 5D and 5F).

**Composition of Arterial Lesions in eTNAP Mice**

The composition of the matrix core was further examined for the presence of lipids and immunoreactivity for osteocalcin, a specific bone matrix protein. To detect lipid, cryosections of the hearts from WT and eTNAP mice aged 8 weeks were developed with lipophilic Oil Red O stain. No lipid inclusions were detected in the normal coronary arteries of WT or in association with the lesions in the arteries of eTNAP mice.
Immuno-fluorescent staining for osteocalcin specifically detected the presence this protein in the coronary lesions (both small and large) of eTNAP mice aged 8 weeks, whereas osteocalcin expression was undetectable in the normal coronary arteries of WT animals (Figure 6B).

To examine the activation of an osteochondrogenic transdifferentiation program in the arteries of eTNAP mice that might potentially explain their histological phenotype, we isolated RNA from the aortas of TNAP and WT mice aged 13 weeks and analyzed the expression of a set of specific osteochondrogenic markers (Table 3). Several markers of osteogenesis—Runx2, osteocalcin, osteopontin, and osteoprotegerin—were significantly upregulated at mRNA level in the aortas of eTNAP mice compared with WT mice. In addition, the expression of the Col2a1 gene, encoding cartilage-specific type II collagen, was also increased in eTNAP mice compared with controls. To confirm these observations at the protein level, we performed immunohistochemical analysis of coronary lesions from eTNAP mice using antibodies specific to Runx2, collagen II, and osteopontin. Runx2 staining was detected primarily in the arterial sections devoted to the nuclear DAPI stain, a region probably representing calcified VSMCs, and in the core of the lesions (Figure 7A). Collagen II was consistently detected in the core of coronary lesions (Figure 7B). Osteopontin staining appeared to localize to the periphery of coronary inclusions identified by differential interference contrast imaging (Figure 7C). Because osteopontin belongs to a group of sialylated matrix proteins (modified by carbohydrates, containing sialic acid), we used a specific reagent, fluorescently labeled S nigra lectin, to test for the presence of sialic acid in the arterial lesions. Coronary lesions detected by differential interference contrast imaging were reactive with S nigra lectin, whereas unaffected arteries and surrounding normal cardiac tissue were negative for S nigra lectin staining (Figure 7D). Similar to osteopontin staining, S nigra lectin staining was brighter at the periphery of the coronary inclusions.

**Cardiovascular Phenotype of eTNAP Mice**

Hemodynamic assessment was performed by catheterization of right carotid artery in eTNAP mice and control littermates aged 8, 13, and 23 weeks. Blood pressure was within the normal range and similar between groups at baseline (aged...
8 weeks) and at an intermediate time point (aged 13 weeks) despite the presence of disseminated arterial calcification in eTNAP mice. At age 23 weeks, however, blood pressure became significantly elevated in eTNAP mice compared with controls (systolic, 126 versus 109 mm Hg; diastolic, 90 versus 79 mm Hg; \( P \leq 0.01 \) for both) (Table 4). Analysis of cardiac function in eTNAP and WT animals was performed by M-mode echocardiography (Table 4). The calculated LV mass showed LV hypertrophy in an older group (aged 23 weeks) of eTNAP mice. No differences in end-diastolic or end-systolic LV diameters were observed between groups, and the ejection fraction normalized to body weight was not significantly different between eTNAP and WT mice, indicating that LV hypertrophy in eTNAP mice aged 23 weeks was compensatory in nature and in line with elevated blood pressure (Table 4).

**Discussion**

In this study, we tested the hypothesis that elevated TNAP activity in ECs might shift the balance toward calcification of vascular structures; however, we did not anticipate that calcification would preferentially affect arteries and be largely absent in veins and microvasculature. This observation, combined with the fact that endogenous TNAP is normally expressed in a subset of microvascular ECs, points to a restricted role of osteogenic (TNAP-positive) ECs in calcification of elastic arteries. Outside the arteries, calcification was observed in the lungs and splenic stroma and capsule, both tissues with high elastic content. Interestingly, haploinsufficiency of elastin was shown to reduce vascular calcification in one of the most severe models of vascular calcification, Mgp

**Table 3. Gene Expression in the Aortas of Mice Aged 13 Weeks**

| Gene     | WT (n=5) | eTNAP (n=8) | Fold Change* |
|----------|----------|-------------|--------------|
| Alpl     | 1484±256 | 2042±364    | 1.4          |
| Runx2    | 321±110  | 610±51      | 1.9†         |
| Osterix  | 11±4     | 23±8        | 2.1          |
| Sox9     | 473±145  | 966±204     | 2.0          |
| Bmp2     | 832±211  | 1366±191    | 1.6          |
| Osteocalcin | 15±5  | 52±11       | 3.5†         |
| Osteopontin | 78±16 | 963±212     | 12.3†        |
| Mgp      | 113 436±33 570 | 200 105±28 472 | 1.8          |
| Aggrecan | 1772±600 | 2846±357    | 1.6          |
| Col2a1   | 5±1      | 40±15       | 8.0†         |
| Opg      | 2668±846 | 624±1060    | 2.3†         |
| Rankl    | 0.06±0.03 | 0.13±0.03   | 2.2          |

Data are shown as arbitrary units, mean±SEM. eTNAP indicates endothelial tissue-nonspecific alkaline phosphatase; WT, wild type.

*eTNAP relative to WT.

†\( P < 0.05 \), Student t test, ‡\( P < 0.01 \), §\( P < 0.05 \), Mann-Whitney test.

**Figure 7.** Expression of Runx2, collagen II, osteopontin, and sialic acid in coronary lesions of eTNAP mice. A through D, coronary lesions were identified by DIC imaging. A, Consecutive sections of the hearts from eTNAP mice aged 8 weeks were stained with Runx2 antibody or secondary antibody only, DNA binding dye DAPI was used as counterstain. B, Lesions from eTNAP mice aged 8 weeks were stained with collagen II antibody or secondary antibody alone and counterstained with DAPI. C, Osteopontin expression was detected in the lesions from eTNAP mice aged 13 weeks by direct immunofluorescent staining; IB4 was used as counterstain. D, The presence of sialic acid was detected in the coronary lesions from eTNAP mice aged 13 weeks with fluorescein-labeled SNA lectin; DAPI was used as nuclear counterstain. Affected (left panels) and unaffected (right panels) coronary arteries from the same tissue section are shown. DIC indicates differential interference contrast; eTNAP, endothelial tissue-nonspecific alkaline phosphatase; IB4, isolectin B4; SNA, Sambucus nigra.
knockout mice\(^3\),\(^8\) pointing to the importance of the mineral scaffolding function of elastic structures when other procalci-fying conditions are equal.\(^3\) We can speculate that in our model of panendothelial TNAP-driven arterial calcification, the “wear and tear” of extracellular matrix, proximal to the endothelial layer, might play a role in the initiation of osteochondrogenic transdifferentiation and early vascular lesions. To this end, a recent study demonstrated that activation of a group of endothelial elastases and kallikreins is suf-ficient to induce endothelial–mesenchymal transitions of ECs and contributes to vascular calcification in Mgp knockout mice.\(^4\)

The eTNAP mice display massive upregulation of circulating TNAP activity, which raises a question about whether circulating TNAP should reduce the levels of plasma PPi, a known inhibitor of vascular calcification\(^2\),\(^4\)–\(^6\) and promote calcification independent of ECs. Previously, we reported findings from 2 nonvascular models of tissue-specific TNAP overexpression (under the control of Col1a1 or apoE promoters\(^4\)), in which we observed up to 20-fold higher average TNAP activity in plasma in the absence of vascular calcification. Moreover, when TNAP was expressed in VSMCs\(^3\),\(^1\) or in ECs (this study), different patterns of vascular calcification were observed despite equally elevated circulating TNAP activity. Consequently, it appears that tissue distribution of TNAP (as opposed to its activity in the blood) is responsible for vascular calcification. Conversely, no significant reduction in plasma PPi was detected in any of these studies,\(^3\),\(^1\),\(^4\) including the data presented in this report. TNAP is likely to be partially inhibited in whole plasma by the physiological concentrations of inorganic phosphate, whereas dilution of plasma in the conditions of clinical assay relieves this inhibition\(^4\) (but not the fold change). We can also speculate that the reduction in circulating PPi levels could be partially compensated by upregulation of PPi production, thus sustaining elevated TNAP activity. In our previous study, we detected higher expression of the PPi-producing enzyme ectonucleotide pyrophosphatase/phosphodiesterase 1 (Enpp1), in TNAP transgenic mice compared with WT mice.\(^3\) The lack of systemic PPi reduction in this model and others suggests that local tissue concentrations of PPi, (and the activity of TNAP) are more critical in regulating mineralization compared with their systemic levels.

Our initial assumption was that overexpression of TNAP and consequent reduction of tissue PPi would be sufficient to promote endothelial-driven vascular calcification. Unexpectedly, we observed that TNAP overexpression induced the formation of arterial lesions that were apparently initiated by subendothelial accumulation of nonmineralized material. This matrix served as a scaffold for calcification at the later stages of lesion progression; therefore, we could not exclude that transdifferentiation of ECs into a chondrogenic or osteogenic type is also required to promote vascular calcification, at least in this model of TNAP overexpression.

Despite TNAP overexpression in essentially every EC, the foci of calcification were rather infrequent, indicating that additional stimuli are required for ECs to initiate arterial lesions, even in the presence of high AP activity. A possible explanation is that ECs have to retain a certain level of pluripotency to progress toward transdifferentiation into the synthetic type capable of mounting an appropriate gene expression and secretion of matrix vesicles containing large amounts of substrates for mineral deposition. This theory would be in line with the recent epidemiological studies pointing out the important role of osteogenic endothelial progenitor cells in vascular calcification.\(^15\)–\(^18\)

| Parameter | 8 Weeks | 13 Weeks | 23 Weeks |
|-----------|---------|----------|----------|
| BW, g     | 30.0±2.0| 28.2±2.0 | 34.9±3.5 | 32.5±2.8 | 38.5±5.4 | 36.1±7.3 |
| HR, bpm   | 451±25  | 434±34   | 439±31   | 427±28   | 505±66   | 524±52   |
| SBP, mm Hg* | 101±4 | 99.7     | 102±7    | 107±11   | 109±6    | 126±8\(^7\) |
| DBP, mm Hg* | 75±3  | 73±5     | 75±5     | 81±10    | 79±5     | 90±7\(^7\) |
| LV Diam(s), mm | 2.82±0.25 | 2.95±0.33 | 2.98±0.29 | 2.92±0.25 | 2.89±0.35 | 2.91±0.21 |
| LV Diam(d), mm | 4.11±0.19 | 4.14±0.25 | 4.28±0.20 | 4.16±0.18 | 4.21±0.31 | 4.11±0.16 |
| EF, %     | 59.4±5.7 | 55.7±6.7 | 58.0±6.1 | 57.4±5.2 | 59.7±6.3 | 56.2±6.1 |
| CO/BW, mL/min per gram | 0.67±0.07 | 0.65±0.08 | 0.60±0.11 | 0.58±0.06 | 0.62±0.11 | 0.62±0.12 |
| LV/BW, mg/g | 3.23±0.33 | 3.25±0.42 | 3.12±0.40 | 3.23±0.54 | 3.01±0.49 | 3.79±1.19\(^1\) |

Data are shown as mean±SD. bpm indicates beats per minute; BW, body weight; CO, cardiac output; DBP, diastolic blood pressure; EF, ejection fraction; eTNAP, endothelial tissue-nonspecific alkaline phosphatase; HR, heart rate; LV Diam(d), diameter of left ventricle in diastole; LV Diam(s), diameter of left ventricle in systole; LV/BW, left ventricular mass/body weight ratio; SBP, systolic blood pressure; WT, wild type.

\(^*n=7\) to 9.  
\(^1\)*p=0.01, vs same age WT, 1-way ANOVA followed by Bonferroni-adjusted multiple comparisons.
In this study, we showed that a selective rise of TNAP expression in the ECs is sufficient to trigger arterial calcification. The phenotype of eTNAP mice, however, is substantially different from mice bearing TNAP in VSMCs. VSMC-driven TNAP expression results in extensive calcification of the aorta, which leads to early lethality, whereas panendothelial TNAP manifests primarily and systemically as disseminated lesions in the smaller diameter arteries. Lesions nucleate in the subendothelial space of medium-sized arteries and induce neointimal hyperplasia. Arterial remodeling, secondary to calcification, likely resulted in increased vessel wall stiffness, which can explain elevated blood pressure in older eTNAP mice. In this scenario, LV hypertrophy in eTNAP mice aged 23 weeks can be interpreted as a compensatory physiological response counteracting increased systemic vascular resistance. Decreased coronary perfusion due to partial occlusion of coronary arteries, however, can also play a role at this stage of cardiac remodeling.

Healthy absence of calcification in vasculature is maintained by a fine balance between activators and inhibitors of mineralization. These include factors promoting osteochondrogenic differentiation of cells residing in the vessel wall, enzymes and transporters maintaining in situ phosphorus and PPi levels, and extracellular matrix components that regulate calcium phosphate nucleation and crystal growth. We showed that ECs can induce vascular calcification and that upregulation of a single inducer of calcification, TNAP, is sufficient to promote the osteogenic potential of ECs.

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Disclosures
None.

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