Unexpected Role of the Copper Transporter ATP7A in PDGF-Induced Vascular Smooth Muscle Cell Migration


Rationale: Copper, an essential nutrient, has been implicated in vascular remodeling and atherosclerosis with unknown mechanism. Bioavailability of intracellular copper is regulated not only by the copper importer CTR1 (copper transporter 1) but also by the copper exporter ATP7A (Menkes ATPase), whose function is achieved through copper-dependent translocation from trans-Golgi network (TGN). Platelet-derived growth factor (PDGF) promotes vascular smooth muscle cell (VSMC) migration, a key component of neointimal formation.

Objective: To determine the role of copper transporter ATP7A in PDGF-induced VSMC migration.

Methods and Results: Depletion of ATP7A inhibited VSMC migration in response to PDGF or wound scratch in a CTR1/copper-dependent manner. PDGF stimulation promoted ATP7A translocation from the TGN to lipid rafts, which localized at the leading edge, where it colocalized with PDGF receptor and Rac1, in migrating VSMCs. Mechanically, ATP7A small interfering RNA or CTR small interfering RNA prevented PDGF-induced Rac1 translocation to the leading edge, thereby inhibiting lamellipodia formation. In addition, ATP7A depletion prevented a PDGF-induced decrease in copper level and secretory copper enzyme precursor prolylsyl oxidase (Pro-LOX) in lipid raft fraction, as well as PDGF-induced increase in LOX activity. In vivo, ATP7A expression was markedly increased and copper accumulation was observed by synchrotron-based x-ray fluorescence microscopy at neointimal VSMCs in wire injury model.

Conclusions: These findings suggest that ATP7A plays an important role in copper-dependent PDGF-stimulated VSMC migration via recruiting Rac1 to lipid rafts at the leading edge, as well as regulating LOX activity. This may contribute to neointimal formation after vascular injury. Our findings provide insight into ATP7A as a novel therapeutic target for vascular remodeling and atherosclerosis. (Circ Res. 2010;107:787-799.)

Key Words: vascular remodeling ■ vascular smooth muscle ■ migration ■ copper transporter ■ platelet-derived growth factor

Copper, an essential micronutrient, plays an important role in physiological repair processes including wound healing and angiogenesis, as well as various pathophysiologies including tumor growth, neurodegenerative disease, and atherosclerosis.1-7 Copper levels are significantly increased in cancer and atherosclerosis.1,7 Copper uptake is mainly mediated by the copper transporter (CTR)1 copper importer, which is involved in embryonic development.1,15 Once copper enters the cell via
Here, we demonstrate the novel role of ATP7A in PDGF-induced VSMC migration. PDGF stimulation promotes ATP7A translocation from the TGN to lipid rafts which localize at the leading edge in migrating VSMC, thereby promoting lamellipodia formation through recruiting Rac1, in a CTR1-dependent manner. This is associated with a decrease in cellular copper and secretory copper enzyme precursor prolysyl oxidase in caveolae/lipid rafts, which may contribute to activation of LOX. In vivo, ATP7A expression is markedly increased and copper accumulation is observed at neointimal VSMC in wire injury model. These findings provide insight into ATP7A as potential therapeutic targets for vascular remodeling and development of atherosclerosis.

Methods
An expanded Methods section is available in the Online Data Supplement at http://circres.ahajournals.org.

Detergent-Free Purification of Caveolin-Rich Membrane Fractions
Caveolae/lipid raft fractions were separated by the sodium carbonate–based detergent-free method.

Copper Measurements
Copper contents were analyzed by inductively coupled plasma mass spectrometry (ICP-MS) using a PlasmaQuad3, as reported previously.26

Vascular Injury
Animal protocols were approved by the Animal Care and Use Committee of the University of Illinois at Chicago and University of Minnesota. Wire-induced injury of the carotid artery in apolipoprotein (Apo)E-deficient atherosclerotic mice was performed, as reported previously.27

Synchrotron X-Ray Fluorescence Microscopy
Sections (5-μm thick) of formalin-fixed, paraffin-embedded, wire-injured femoral artery were used. For x-ray imaging, the sections (5-μm thick) of formalin-fixed, paraffin-embedded, wire-injured femoral artery were prepared on silicon nitride windows (Silson), as reported previously.28

Results
Expression of the ATP7A Copper Transporter and the CTR1 Copper Importer in VSMCs
RT-PCR analysis of rat, mouse, and human aortic smooth muscle cells (RASMs, HASMs, and MASMs) detected both ATP7A and CTR1 expression (Figure 1A). Western analysis with anti-ATP7A antibody showed expression of ATP7A with a 178-kDa protein in whole cell lysates of RASMs, HASMs, and MASMs (Figure 1B).16,17 By contrast, CTR1 protein was not detected in whole cell lysates, but in caveolae/lipid rafts fraction, by specific anti-CTR1 antibody, as shown later (Figure 5A; Online Figure V, A and B).

ATP7A Is Involved in PDGF-Induced VSMC Migration in a Copper-Dependent Manner
We next examined the role of ATP7A in VSMC migration. Modified Boyden chamber assays demonstrated that knockdown of endogenous ATP7A expression with small interfering (si)RNA significantly inhibited PDGF-stimulated migration in VSMCs (Figure 1C and Online Figure I). Because the
copper transporter function of ATP7A is dependent on the
delivery of copper from the extracellular space via CTR1.29,30
we also investigated the role of CTR1. Transfection of
VSMCs with CTR1 siRNA significantly reduced endogenous
CTR1 expression (Online Figures I [A] and V [B]). PDGF-
induced cell migration was significantly inhibited by CTR1
siRNA (Figure 1C) as well as copper chelators tetrathiomo-
lybdate (TTM) (cell-permeable) and bathocuproine disulfon-
ate (BCS) (cell-impermeable) (Figure 1D), suggesting that
this response is copper-dependent. Wound scratch assay of
confluent monolayer of VSMCs in the presence of PDGF also
showed that ATP7A siRNA as well as CTR1 siRNA or BCS
significantly inhibited directional cell migration in response
to wound injury (Figure 1E). In contrast, transfection of
ATP7A siRNA did not have significant effect on PDGF-
induced VSMC proliferation (Online Figure II). These results
suggest that ATP7A is involved in PDGF-stimulated VSMCs
migration in a copper-dependent manner.

PDGF Stimulation Promotes ATP7A
Translocation to the Leading Edge in a
Copper-Dependent Manner
To gain insight into the mechanism by which ATP7A
mediates VSMC migration in response to PDGF, we exam-
ined the subcellular localization of ATP7A after wound
scratch in the presence of PDGF. In confluent monolayers of
VSMC before wounding or in migrating VSMCs away from
the scratched area, ATP7A was found predominantly in
perinuclear regions (Figure 2A). In contrast, ATP7A accu-
mulated and colocalized with F-actin at the leading edge in
actively migrating VSMCs, but did not colocalize with actin
stress fibers in the cell body (Figure 2A). We confirmed the
specificity of the ATP7A staining by ATP7A siRNA, as
shown later (Figure 3D). Furthermore, nonimmune IgG
(control) showed no staining (data not shown). Pretreatment
of VSMCs with copper chelators BCS or TTM as well as
CTR1 siRNA markedly inhibited wound-induced translocat-
tion of ATP7A toward the leading edge (Figure 2B). Similarly, in untreated VSMC, ATP7A was found predominantly in perinuclear regions, and colocalized with syntaxin 6, a TGN marker (Online Figure III, B). PDGF stimulation rapidly promoted ATP7A translocation from the Golgi to the plasma membrane leading edge with peak at 5 minutes, which gradually returned to the perinuclear region within 30 minutes (Figure 2C; Online Figure III, A). This effect was observed in VSMCs from other species, such as MASMs, suggesting that function of ATP7A is similar in VSMC across the different species (Online Figure III, C). Furthermore, this response was inhibited by copper chelators as well as knockdown of CTR1 (Figure 2D). Of note, the location of syntaxin 6 was not altered by PDGF in VSMCs (Online Figure III, B), suggesting that PDGF-induced ATP7A translocation was not attributable to the general effect on Golgi structure. Taken together, these findings suggest that PDGF stimulation promotes ATP7A translocation to the leading edge, in a copper-dependent manner, thereby stimulating VSMC migration.

### ATP7A Is Involved in PDGF-Stimulated Lamellipodia Formation and Rac1 Translocation in a CTR1-Dependent Manner in VSMCs

To assess further the mechanism by which ATP7A is involved in VSMC migration, we examined whether ATP7A is involved in actin reorganization. Knockdown of ATP7A by siRNA significantly impaired wound scratch- (Figure 3A) and PDGF- (Figure 3B) stimulated lamellipodia formation at the leading edge in VSMCs as visualized by phalloidin staining. These effects were also prevented by siRNA knockdown of the CTR1. Because Rac1 plays a role in lamellipodia formation and cell migration, we next examined the role of ATP7A in Rac1 activation and translocation in PDGF-stimulated VSMCs. PDGF stimulation increased active, GTP-bound form of Rac1 within 1 minute, which was not affected by either ATP7A or CTR1 siRNA (Online Figure IV). Immunofluorescence analysis showed that PDGF stimulation promoted translocation of Rac1 to the leading edge where it colocalized with ATP7A (Figure 3C). Coimmuno-
precipitation analysis showed that ATP7A associated with Rac1 in the basal state, which was further enhanced after PDGF stimulation (Online Figure IV, C), suggesting that ATP7A recruits Rac1 to the lipid rafts via binding to Rac1 in response to PDGF directly or indirectly. Furthermore, Rac1 trafficking to the leading edge was inhibited by either ATP7A or CTR1 siRNA (Figure 3D). Thus, the copper transporter ATP7A is involved in PDGF-stimulated translocation of Rac1 to the leading edge, but not Rac1 activation, in a CTR1-dependent manner, which may contribute to lamellipodia formation and VSMC migration.

**ATP7A Colocalizes With PDGFR at the Leading Edge in a CTR1-Dependent Manner in PDGF-Stimulated VSMCs**

Because we found that ATP7A is a downstream mediator for PDGFR signaling linked to VSMC migration, we next examined the relationship between ATP7A and PDGFR in PDGF-stimulated VSMCs. Immunofluorescence analysis showed that ATP7A colocalized with PDGFR at the leading edge after PDGF stimulation (Figure 4A). Of note, both CTR1 and ATP7A siRNAs inhibited PDGFR-induced ATP7A movement to the leading edge without affecting PDGFR localization (Figure 4B). Coimmunoprecipitation analysis further confirmed that PDGF stimulation promoted PDGFR association with ATP7A, which was somewhat inhibited by CTR1 siRNA (Figure 4C). Under this condition, PDGF-induced PDGFR autophosphorylation was not affected by CTR1 siRNA (Figure 4C). Taken together, these results suggest that PDGF stimulation induces colocalization and association of ATP7A with PDGFR at the leading edge in a CTR1-dependent manner, and that CTR1-ATP7A pathway is downstream of PDGFR activation.

**PDGF Promotes ATP7A Recruitment to the Caveolae/Lipid Rafts, Where PDGFR, Rac1, and CTR1 Are Localized, in a CTR1-Dependent Manner**

To gain further insight into the subcellular compartments in which ATP7A is localized, we performed detergent-free sucrose gradient fractionation in VSMCs, as previously described. Western analysis of sequential fractions from the gradient showed that ATP7A was found in both caveolin-enriched lipid rafts fractions and noncaveolin/lipid rafts fractions which mainly contain paxillin (Figure 5A). By contrast, the CTR1 copper importer was predominantly found in caveolae/lipid rafts. The specificity of the CTR1 antibody, which detects glycosylated CTR1, was verified by CTR1-deficient mouse embryonic fibroblast cells and VSMCs.
transfected with CTR1 siRNA (Online Figure V, A and B). Localization of ATP7A and CTR1 in caveolae/lipid rafts was further confirmed in various VSMCs, including RASM, HASM, and MASM (Online Figure V, C) and using detergent-free OptiPrep gradient cell fractionation (Online Figure V, D). Figure 5B shows that PDGF stimulation for 5 minutes promoted recruitment of ATP7A, CTR1 and Rac1 to the caveolae/lipid rafts fractions without affecting PDGFR localization, which was associated with an increase in PDGFR phosphorylation in these fractions. Significantly, CTR1 siRNA prevented PDGF stimulation for 5 minutes promoted recruitment of ATP7A, CTR1 and Rac1 to the caveolae/lipid rafts fractions without affecting PDGFR localization, which was associated with an increase in PDGFR phosphorylation in these fractions. Significantly, CTR1 siRNA prevented PDGF-induced translocation of ATP7A and Rac1 to the caveolae/lipid rafts without affecting PDGFR phosphorylation (Figure 5B). These suggest that PDGF-induced PDGFR autophosphorylation occurs in caveolae/lipid rafts, which in turn promotes recruitment of ATP7A and Rac1 to these specialized microdomains where PDGFR is localized, in a CTR1-dependent manner.

ATP7A Is Translocated to the Lipid Rafts Localized at the Leading Edge in PDGF-Stimulated VSMCs

Lipid rafts have been shown to be localized at the leading edge during cell migration.35 We thus examined the spatial relationships among ATP7A, lipid rafts and leading edge after PDGF stimulation using immunofluorescence analysis. As shown in Figure 5C and Online Figure VI (B), ATP7A colocalized with cholera toxin subunit B (CTxB), a lipid raft marker,35 at the leading edge in PDGF- and wound scratch-stimulated VSMCs. Furthermore, CTxB accumulated and colocalized with F-actin at the leading edge in actively migrating VSMCs (Online Figure VI, A and C). Disruption of lipid rafts by cholesterol binding reagent, methyl-β-cyclodextrin, completely abrogated the ATP7A localization at the leading edge and CTxB staining (Figure 5D) as well as lamellipodia formation (Online Figure VI, A). These results suggest that PDGF stimulates translocation of ATP7A to the lipid rafts localized at the leading edge, thereby promoting lamellipodia formation.

PDGF Stimulation Reduces Copper Content in Caveolae/Lipid Raft Fractions in VSMCs

Because ATP7A is involved in exporting copper to the extracellular space, we next examined the effects of PDGF on copper levels in VSMCs. At first, we performed $^{64}\text{Cu}$
efflux and $^{64}$Cu uptake experiments in cultured VSMCs and found that PDGF stimulation had no effects on either response (Online Figure VII). Similarly, ICP-MS analysis of total cell lysates of VSMCs also showed that copper contents were not changed after PDGF stimulation (Figure 6A). Because ATP7A is recruited to the caveolae/lipid rafts in response to PDGF, we next measured the copper content in the lipid rafts and nonlipid raft fractions. Intriguingly, we found that the caveolae/lipid rafts fraction contains much higher amounts of copper than nonlipid raft fractions. Moreover, PDGF stimulation significantly reduced copper content in caveolae/lipid rafts in an ATP7A-dependent manner (Figure 6C and Online Figure VIII). These results suggest that PDGF-induced recruitment of ATP7A to the lipid rafts may contribute to copper export to the extracellular space in these specialized microdomains. This was not detected by global measurements of $^{64}$Cu efflux and $^{64}$Cu uptake in whole cells. The plasma membrane localization of ATP7A and CTR1 in PDGF-stimulated VSMCs was confirmed by using cell surface biotinylation assays (Online Figure IX).

We next examined whether PDGF-induced decrease in copper content in caveolae/lipid rafts may reflect the secretion of copper-binding proteins, such as Pro-LOX which obtain copper from ATP7A. We found that pro-LOX was localized in caveolae/lipid rafts in the basal state, and PDGF stimulation promoted the reduction of pro-LOX level in an ATP7A dependent manner in these fractions (Figure 6D and 6E; Online Figure X). After secretion, Pro-LOX is processed to a mature active nonglycosylated LOX and a glycosylated propeptide LOX-PP. Of note, LOX and LOX-PP were not found in caveolae/lipid rafts in the basal state or after PDGF treatment (Figure 6D and 6E). Taken together, these findings suggest that PDGF-stimulated ATP7A recruitment into the caveolae/lipid rafts may be required for the secretion of copper-dependent enzymes such as pro-LOX localized in these specific compartments.

It has been reported that ATP7A is involved in LOX activity and that LOX activity is involved in PDGF-induced VSMC migration in addition to its effect on extracellular matrix remodeling. We thus examined whether LOX activity is involved in copper transporter ATP7A-dependent PDGF-induced VSMC migration. ATP7AsiRNA significantly inhibited LOX activity in VSMCs treated with PDGF (Online Figure XII, A). Furthermore, treatment of a specific chemical LOX activity inhibitor, βaminopropion-
itrile (BAPN), significantly inhibits PDGF-induced VSMC migration (Online Figure XII, B), which is consistent with the report by Lucero et al.\(^{24}\) These findings suggest that LOX activity is involved in PDGF-induced, ATP7A-dependent VSMC migration.

**Figure 6.** PDGF stimulation reduces copper content and Pro-LOX in lipid rafts fractions in VSMCs. A through C, Copper contents were measured by ICP-MS in whole cells (A) or caveolae/lipid rafts or noncaveolae/lipid rafts (B and C) in RASMs with or without PDGF stimulation for 5 minutes. Equal amounts of proteins in caveolae/lipid rafts (fractions 4 to 5) or noncaveolae/lipid rafts (fractions 9 to 13) were obtained by sucrose gradient fractionation as described for Figure 5. \(^*P < 0.05\) vs nonlipid rafts or unstimulated cells (means ± SD, n = 3).

**D,** Identification of Pro-LOX, but not LOX, in caveolae/lipid rafts in VSMCs. RASMs were fractionated by sucrose gradient centrifugation, followed by immunoblotted with anti–LOX-PP (which detects both Pro-Lox and LOX-PP), anti-LOX, or anti-caveolin-1 antibodies.

**E,** Effect of PDGF treatment on Pro-LOX level in caveolae/lipid rafts fractions in VSMCs. Equal amounts of proteins in caveolae/lipid rafts (fractions 4 to 5) were immunoblotted with anti–LOX-PP, -LOX, or –caveolin-1 antibodies in RASMs with or without 50 ng/mL PDGF for 5 minutes.

**ATP7A Is Upregulated in Neointimal Formation in Response to Vascular Injury In Vivo**

To determine the functional significance of ATP7A in VSMC migration in vivo, we examined the role of ATP7A in neointimal formation using a mouse wire injury model.
Immunohistochemical analysis showed that ATP7A protein expression was robustly increased in neointimal VSMC and endothelial cells lining the lumen in the injured vessel of ApoE-deficient atherosclerotic mice (Figure 7A). Immunofluorescence analysis demonstrated that ATP7A was colocalized with α-smooth muscle actin, a VSMC marker, in the neointima formed in response to injury (Figure 7B). We next examined if induction of ATP7A precedes VSMC migration after injury to address the cause-effect of ATP7A in VSMC migration in vivo. As shown in Online Figure XI, we observed increase in ATP7A expression at 3 days after wire injury in the vessels of ApoE-deficient atherosclerotic mice, whereas there was no VSMC migration and neointima formation at this time point as reported by Linder et al.37 Thus, these findings suggest that ATP7A expression is increased before the onset of VSMC migration following vascular injury and this upregulation is also observed in neointimal VSMCs. Finally, we examined spatial distribution of copper in neointima using synchrotron-based x-ray fluorescence microscopy (SXFM).28 As shown in Figure 7C, copper accumulation was observed at neointimal lesions in wire injury model. These results suggest that ATP7A may be involved in neointima formation in response to vascular injury in vivo.

**Discussion**

A role for copper in tissue repair, neointima thickening and atherosclerosis has been suggested; however, the underlying detailed mechanisms remain unknown.3–7,9–11 Here we provide novel evidence that the ATP7A copper transporter is involved in PDGF-stimulated VSMC migration, which is critical for neointimal formation and vascular remodeling (Figure 8). We found that stimulation with PDGF promotes ATP7A translocation from TGN to the leading edge. This PDGF-induced ATP7A movement toward the site of actin remodeling is inhibited by cell permeable and impermeable copper chelators as well as depletion of the CTR1 copper importer. It has been shown that relocation of ATP7A from the Golgi is triggered by increased cytoplasmic copper as well as by estrogen, insulin, NMDA (N-methyl-D-aspartate) activation, hypoxia, and cytokines.2,16,38,39 However, our data are the first demonstration that the ATP7A
copper transporter translocates to the leading edge during growth factor-stimulated cell migration. It has been reported that ATP7A trafficking is in part dependent on its catalytic activity, which requires copper binding to conserved cysteines within its N-terminal region.17–19 Extracellular copper influx is mainly mediated through CTR1, the copper importer, which is also the major source of “bioavailable” copper for ATP7A via the Atox1 copper chaperone.29,30 Indeed, Atox1 siRNA prevented PDGF induced ATP7A translocation to the leading edge, suggesting that Atox1 is involved in PDGF-induced translocation of ATP7A (data not shown). Thus, our findings are consistent with the possibility that ATP7A obtains copper via Atox1 by stimulating extracellular copper uptake through CTR1 in response to PDGF, thereby promoting ATP7A trafficking from the Golgi to the leading edge in VSMCs. However, we failed to detect a PDGF-induced increase in 64Cu uptake in whole cell labeling of VSMCs with 64Cu in this study. This may be attributable to the possibility that CTR1-mediated copper uptake occurs in the specific plasma membrane compartment, as discussed below. It is not known how ATP7A is translocated to the leading edge after PDGF stimulation in a copper-dependent manner. Phosphorylation of ATP7A has been shown to be involved in copper-induced translocation of this protein.40 Thus, it is possible that PDGF stimulation may induce phosphorylation of ATP7A, which may contribute to its translocation to the leading edge. Taken together, our findings provide a novel linkage between copper homeostasis and PDGF-induced VSMC migration.

Migrating cells form protrusions, such as lamellipodia, by reorganizing of the actin cytoskeleton at the leading edge, which plays an important role in cell migration. The small G protein family Rac1 plays a key role in lamellipodia formation11 and PDGF-stimulated cell migration.13 In the present study, we demonstrate that ATP7A is involved in PDGF-induced lamellipodia formation and Rac1 translocation to the leading edge without affecting Rac1 activity, in a copper- and CTR1-dependent manner. Coimmunoprecipitation analysis showed that ATP7A associated with Rac1 in the basal state, which was further enhanced after PDGF stimulation (Online Figure IV, C), suggesting that ATP7A recruits Rac1 to the lipid rafts via binding to Rac1 in response to PDGF directly or indirectly. In line with our data, previous reports show that trafficking of ATP7A from the TGN to the cell surface is associated with actin networks and involves Rac1, which is distinct from the major constitutive secretory pathway.16,41 Of note, PDGF stimulation also promotes colocalization and association of ATP7A with PDGFR at the leading edge. However, CTR1 siRNA prevents ATP7A translocation to the leading edge without affecting PDGFR localization and its autophosphorylation. These results suggest that CTR1-ATP7A-Rac1 pathway is downstream of PDGFR activation involved in lamellipodia formation and VSMC migration.

Caveolae and lipid rafts are specialized membrane microdomains that are highly enriched in cholesterol and sphingolipids and function as platforms for assembly of signaling molecules including G proteins and receptors involved in various functions.35 In chemoattractant-stimulated cancer cells, lipid rafts are accumulated at the leading edge.35 Using sucrose gradient fractionation, the present study shows that PDGF stimulation promotes recruitment of ATP7A and Rac1 to the caveolae/lipid rafts where CTR1 and PDGFR are localized, in a CTR1-dependent manner. By contrast, CTR1 siRNA has no effect on PDGF-induced PDGFR tyrosine phosphorylation as well as localization of caveolin-1 and PDGFR in caveolae/lipid rafts fractions, suggesting that the structure of caveolae/lipid rafts is intact in CTR1 depleted VSMCs. Immunofluorescence analysis also reveals that the marker for lipid rafts, CTxB,35 colocalizes with ATP7A at the leading edge of migrating VSMCs. We confirmed the plasma membrane localization of ATP7A and CTR1 in PDGF-stimulated VSMCs using cell surface biotinylation assays. PDGF-induced Rac1 translocation to the lipid rafts has been reported in Rat-1B cells;13 however, our data provides the first evidence that copper transporters CTR1 and ATP7A are localized in these specialized plasma membrane microdo-
mains. Functional significance of copper-dependent ATP7A trafficking into lipid rafts is demonstrated by showing that disruption of lipid rafts by the cholesterol-binding reagent, methyl-β-cyclodextrin, completely inhibits PDGF-induced ATP7A translocation to the leading edge as well as lamellipodia formation. Moreover, PDGF stimulation rapidly decreases the copper content of caveolae/lipid rafts in an ATP7A-dependent manner, as measured by ICP-MS. This PDGF effect is associated with recruitment of ATP7A to lipid rafts where CTR1 is predominantly found. The rapid kinetics of ATP7A is reminiscent of copper-induced ATP7A recruitment into a fast-recycling pool located in close proximity with the plasma membrane.2,16,17,42 Hung YH et al have reported that the copper content of lipid rafts is paradoxically decreased, when cytoplasmic copper is increased.43 Thus, it is conceivable that PDGF may stimulate copper uptake via CTR1 to provide copper to ATP7A, which in turn promotes ATP7A translocation from the Golgi to the caveolae/lipid rafts to export copper to extracellular space in the specific compartments. Interestingly, we could not detect any changes in copper levels in response to PDGF in total cell lysates. This may support the suggestion that global measurements of 64Cu efflux and 64Cu uptake in whole cells would fail to detect compartmentalized changes of copper distribution observed in the present study.

The physiological consequence of copper export via ATP7A is not merely the elimination of excess cellular copper, but to supply adequate copper to the developing fetus as gestation progresses,16 or to provide copper as part of a neuronal protective mechanism.2 Using SXFM, Finney et al reported that copper is transported from intracellular compartments to the tips of endothelia during capillary growth, which is required for capillary tube formation.28 In the present study, we demonstrate that the copper-dependent Pro-LOX, which is secreted and activated in the extracellular space,25 is localized at caveolae/lipid rafts in the basal state. PDGF stimulation significantly reduces Pro-LOX levels in caveolae/lipid rafts fractions in an ATP7A-dependent manner. This may reflect the secretion of pro-LOX after obtaining copper from the recruited ATP7A in these specialized compartments. Importantly, recent evidence suggests that LOX and LOX-PP, both of which are produced from Pro-LOX through proteolysis in the extracellular space, have opposite effects on cell migration.23–25 Furthermore, and consistent with our results, ATP7A has been shown to deliver copper to some secretory copper enzymes in the post-Golgi vesicles rather than in the TGN, where copper loading normally takes place.38,44 Taken together, these findings suggest that copper-dependent ATP7A trafficking to lipid rafts in response to PDGF may not simply reflect export copper to the extracellular space, but also facilitate copper-loading to pro-LOX in post-Golgi compartments such as lipid rafts, thereby promoting VSMC migration. To support this, we found that ATP7A siRNA significantly inhibited LOX activity in VSMCs treated with PDGF and that the chemical LOX activity inhibitor β-aminopropionitrile significantly blocked PDGF-induced VSMC migration.

A functional role of the copper transporter ATP7A in VSMC migration in vivo is underscored by upregulation of ATP7A expression and copper accumulation (shown by SXFM) in intimal a-smooth muscle actin positive cells. Of note, ATP7A expression is increased before the onset of VSMC migration after vascular injury and this upregulation was also observed in neointimal VSMCs at later time. These findings suggest that ATP7A may play a role in VSMC migration in vivo. Previous reports indicate that copper plays an important role in various cardiovascular diseases and cancer. High serum copper levels are associated with an increased future risk of coronary heart disease.4,5,7 Atherosclerotic lesions have higher copper levels than normal tissues.9 Neointimal thickening after vascular injury is inhibited by copper chelation11 and increased by copper cuffs and stents.10 Copper deficiency therapies prevent tumor progression in clinical trials.3 Interestingly, we found that ATP7A is involved in PDGF-stimulated decrease in copper level and Pro-LOX in caveolae/lipid rafts as well as PDGF-stimulated increase in LOX activity in VSMCs. Furthermore, we showed that inhibition of LOX activity blocks PDGF-induced VSMC migration as reported previously.24 Given that LOX activity is also essential for vascular extracellular matrix maturation, these results suggest that ATP7A copper transporter may promote extracellular matrix deposition, as well as neointimal formation after vascular injury via regulation of LOX activity.45 In addition, one of the major copper- and ATP7A-dependent enzymes regulating ROS metabolism is SOD3, but not either SOD1 or SOD2.21 We thus investigated the contribution of SOD3 but found that SOD3 siRNA had no effect of PDGF-induced directional VSMC migration (Online Figure XII, C). X-ray fluorescence microscopy scans do not clearly show codistribution of copper levels with ATP7A. This may be attributable to the possibility that copper secreted by ATP7A is more widely distributed because of the diffusible nature as ions, and that other copper binding proteins may be involved in neointima formation. Taken together with in vitro data, our findings indicate that upregulation of ATP7A and copper accumulation in neointimal VSMCs may contribute to neointimal formation and vascular remodeling in vivo at least in part by regulating VSMC migration and LOX activity. Because our vascular injury data implicates but does not conclude that ATP7A may play a role in VSMC migration in vivo, further investigation will be required using ATP7A mutant mice46 in future study.

In conclusion, the present study uncovers a novel function of ATP7A as a regulator for PDGF-induced VSMC migration via recruiting Rac1 to lipid rafts at the leading edge in a copper dependent manner, as well as controlling LOX activity, which may contribute to vascular remodeling (Figure 8). They also indicate an important new role for lipid rafts in organizing the protein components involved in copper homeostasis and signaling. Our findings also provide insight into ATP7A as a potential therapeutic target for various pathophysiologies, such as atherosclerosis, postangioplasty restenosis, diabetes, and cancer, which are associated with dysregulation of cell migration.
Acknowledgments

We thank Dr Dennis Thiele (Duke University, Durham, NC) for the generous gift of CTR1<sup>−/−</sup> and CTR1<sup>+/−</sup> mouse embryonic fibroblasts.

Sources of Funding

This work was supported by NIH grants R01 HL070187 (T.F.), R01 HL077524 (to M.U.-F.), and R01 HL080569-01 (to Y.H.); American Heart Association Postdoctoral Fellowship 09POST2250151 (to N.U.); Ruth L. Kirschstein-National Service Research Award (Kirschstein-NRSA) T32 training grant (to G-F.C.); the Uehara Memorial Foundation (to J.O.); and American Heart Association Grant 1-10-BS-76 (to Y.H.); and American Heart Association Grant 10GRNT4400005 (to M.U.-F.), and R01 HL080569-01 (to Y.H.). Use of the Advanced Photon Source at Argonne National Laboratory was supported by the US Department of Energy, Office of Science, Office of Basic Energy Sciences, under contract no. DE-AC02-06CH11357.

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**Novelty and Significance**

**What Is Known?**

- Copper is an essential micronutrient in all living organisms and involved in physiological repair processes, as well as various pathophysiological conditions including atherosclerosis.
- Copper deficiency therapies prevent neointimal thickening in response to vascular injury.
- Platelet-derived growth factor (PDGF) stimulates vascular smooth muscle cell (VSMC) migration, which promotes neointimal formation and vascular remodeling after vascular injury.

**What New Information Does This Article Contribute?**

- Copper transporter ATP7A plays an important role in PDGF-induced VSMC migration in a copper-dependent manner.
- PDGF stimulation promotes ATP7A translocation from the trans-Golgi network (TGN) to the lipid raft microdomains which localize at the leading edge, thereby promoting lamellipodia formation through recruiting Rac1 as well as increasing lysyl oxidase (LOX) activity.
- In vivo, ATP7A expression is upregulated and copper accumulation is observed in neointimal VSMC in response to vascular injury.

Copper, an essential micronutrient, has been implicated in vascular remodeling in response to injury by unknown mechanisms. Because excess copper is toxic, bioavailability of copper is tightly controlled not only by the copper importer CTR1, but also by the copper exporter ATP7A. Function of ATP7A is achieved through copper-dependent translocation from TGN. We investigated a role of ATP7A in PDGF-induced VSMC migration, a key component of neointimal formation after vascular injury. Here we show that depletion of ATP7A inhibits PDGF-induced VSMC migration in a copper/CTR1 dependent manner. Mechanistically, PDGF stimulation promotes ATP7A translocation from TGN to the lipid raft microdomains which localize at the leading edge in migrating VSMCs, thereby promoting lamellipodia formation through recruiting Rac1, as well as regulating LOX activity. In vivo, ATP7A is markedly increased in neointimal VSMC in wire injury model, in which copper accumulation is observed. Our findings uncover an unexpected role of ATP7A as a regulator of vascular migration in response to injury and provide insight into copper transporters as potential therapeutic targets for vascular remodeling and atherosclerosis. Furthermore, our studies will suggest a novel linkage between copper homeostasis and vascular migration.
Unexpected Role of the Copper Transporter ATP7A in PDGF-Induced Vascular Smooth Muscle Cell Migration


Circ Res. 2010;107:787-799; originally published online July 29, 2010; doi: 10.1161/CIRCRESAHA.110.225334

Circulation Research is published by the American Heart Association, 7272 Greenville Avenue, Dallas, TX 75231
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Print ISSN: 0009-7330. Online ISSN: 1524-4571

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SUPPLEMENTAL MATERIAL

Detailed Methods

Materials: Antibodies to PDGFR-β, Rac1 (C-11, for immunoprecipitation) and paxillin were from Santa Cruz. Antibodies to caveolin-1 and Rac1 (for immunofluorescence) were from BD Biosciences Pharmingen. Rac1 antibody (for immunoblotting) and Rac/cdc42 Assay Reagent were from Millipore. Antibody against ATP7A was from Sigma-Aldrich, and antibodies against LOX-PP or LOX were from Novus Biologicals. Ctrl1 antibody has been described previously. Transwell 24-well plates were from BD Biosciences. Enhanced chemiluminescence (ECL) Western Blotting Detection Reagents and nitrocellulose membranes (Hybond-ECL) were obtained from Amersham Biosciences Corp. Oligofectamine, Opti-MEM I Reduced-Serum Medium, Alexa Fluor 568 phalloidin and Vybrant Lipid Raft Labeling Kit were from Invitrogen Corp. All other chemicals and reagents were from Sigma.

Cell Culture: Vascular smooth muscle cells (VSMCs) were isolated from male Sprague-Dawley rat thoracic aortas by enzymatic digestion. Cells were grown in Dulbecco's modified Eagle's medium (DMEM) supplemented with 10% bovine serum and used at between passage 7 and 15. Cells were starved for 24 hours with serum free media before PDGF stimulation. For some experiments, VSMCs from other species, such as human aortic smooth muscle cells (HASMs) and mouse aortic smooth muscle cells (MASMs), were used. HASMs (Clonetics Corp) were cultured in smooth muscle basal medium (Clonetics) and 5% FBS. Experiments were performed using cells between passages 4 to 8.

MASMs were isolated from 8- to 10-week-old C57BL/6 mice (Jackson Laboratories) and cultured as previously described with modifications. After removal of the connective tissue and blood, two aortas were incubated in an enzyme solution containing collagenase II (1.5 mg/mL), elastase (type II, 0.5 mg/mL), and trypsin inhibitor (type I-S, 1mg/mL) at 37°C. Cells were rinsed and plated in Dulbecco’s modified Eagle’s medium containing 10% FCS, glutamine (2 mmol/L), streptomycin (100 mg/mL), and penicillin (100 U/mL), and amphotericin B (25 mg/mL). The SMC identity of the cells was confirmed by staining with SMC-specific α-actin monoclonal antibody (clone 1A4, Sigma Chemicals Inc). Experiments were performed using cells between passages 4 to 8.

siRNA Transfection: siRNAs were obtained from Applied Biosystem. VSMCs were seeded into culture dishes one day prior to transfection. Transfection of siRNA (30 nM) was performed using Oligofectamine (Invitrogen) according to the manufacturer’s protocol. PDGF stimulation was performed 48 h after transfection.

Reverse transcription-polymerase chain reaction (RT-PCR): Total RNA was isolated from VSMCs using TRI reagent (Molecular Research Center, Inc) according to the manufacturer’s instructions, and first stranded cDNA was synthesized with a High-Capacity cDNA Reverse Transcription Kits. The PCR was performed according to the manufacturer’s protocol using ABI PRISM® 7000 Sequence Detection System (Applied Biosystems, CA) and QuantiFast SYBR Green PCR Kit (Qiagen, CA). PCR was performed with a 5 min preincubation at 95°C followed by 40 cycles of 10 s at 95°C and 30 s at 60°C. PCR products were subjected to melting curve analysis, using the ABI PRISM® 7000 Sequence Detection System, to exclude amplification of unspecific products, and separated on 2.5% agarose gel and stained with ethidium bromide. All real-time PCR primers were purchased from predesigned primers of QuantiTect primer assays (Qiagen).
**Scratch Wound Assay:** ATP7A, CTR1, or control siRNA-transfected VSMCs or BCS treated VSMCs were cultured on 6-well plates. Confluent cells were scraped using sterilized 10-μL pipette tips and washed with 0.1% serum media, and stimulated with 50 ng/ml of PDGF for 24 h, as previously described⁵. Images were captured immediately at 0 h and at 24 h after the wounding.

**Modified Boyden Chamber Assay:** Migration assays using VSNCs with the Modified Boyden Chamber method were conducted in duplicate 24-well transwell chambers as described previously⁵. The upper insert (8-μm pores coated with 0.1% gelatin) containing VSMCs suspensions (5×10⁴ cells) transfected with ATP7A siRNA, CTR1 siRNA or control siRNA were placed in the bottom 24-well chamber containing fresh media with 0.2% FBS and stimulants. The chamber was incubated at 37°C for 8 h. The membrane was fixed and stained using Diff-Quick. Four random fields at × 200 magnifications were counted.

**Confocal Immunofluorescence Microscopy in VSMCs:** VSMCs on glass coverslips were rinsed quickly in ice-cold PBS, fixed in freshly prepared 4% paraformaldehyde in PBS for 10 min at room temperature, permeabilized in 0.05% Triton X-100 in PBS for 5 min, and rinsed sequentially in PBS, 50 m mol/L NH₄Cl and PBS for 10 min each. After incubation for 1 h in blocking buffer (PBS+3% BSA), cells were incubated with ATP7A antibody, PDGFR antibody, FITC-conjugated Rac1 antibody, Alexa Fluor 568-conjugated phallolidin for 18 h at 4°C, rinsed in PBS/BASA, and then incubated in either FITC-conjugated goat anti-IgY, rhodamine TRITC-conjugated donkey-IgY or Alexa Fluor 546-conjugated goat anti-rabbit IgG for 1 h at room temperature. Lipid rafts were stained with Vybrant Alexa Fluor 555 Lipid Raft Labeling kit. Cells on coverslips were mounted onto glass slides in Vectashield (Vector Laboratories) and examined using the 488 and 543 nm lines of the argon ion and green HeNe lasers. Controls with no primary antibody showed no fluorescence labeling and single label controls were performed in double labeling experiments.

For immunofluorescence staining of lipid rafts and F-actin in VSMCs, VSMCs on glass coverslips were stained with Vybrant Alexa Fluor 555 Lipid Raft Labeling kit, and then fixed in freshly prepared 4% paraformaldehyde in PBS for 10 min at room temperature, permeabilized in 0.05% Triton X-100 in PBS for 5 min, and rinsed sequentially in PBS, 50 m mol/l NH₄Cl and PBS for 10 min each. After incubation for 1 h in blocking buffer (PBS+3% BSA), cells were incubated with Alexa Fluor 488-conjugated phallolidin for 1 h at room temperature. Cells on coverslips were mounted onto glass slides in Vectashield and examined using the 488 and 543 nm lines of the argon ion and green HeNe lasers.

**Immunoprecipitation and Immunoblotting:** Growth-arrested cells were stimulated with PDGF at 37°C, and cells were lysed with 500 μl of ice-cold lysis buffer, pH 7.4 (50 mM HEPES, 5 mM EDTA, 100 mM NaCl), 1% Triton X-100, 60 mM n-Octyl-β-D-glucopyranoside, protease inhibitors (10 μg/ml aprotinin, 1 mM phenylmethylsulfonyl fluoride, 10 μg/ml leupeptin) and phosphatase inhibitors (50 mM sodium fluoride, 1 mM sodium orthovanadate, 10 mM sodium pyrophosphate). For immunoprecipitation cell lysates (1000 μg) were precipitated with antibody overnight at 4°C and then incubated with 20 μl of protein A/G-agarose beads for 1.5 h at 4°C. Cell lysates (25 μg) or immunoprecipitates were separated using SDS-polyacrylamide gel electrophoresis and transferred to nitrocellulose membranes, blocked overnight in PBS containing 5% nonfat dry milk and 0.1% Tween 20, and incubated for overnight with primary antibodies as described previously. After incubation with secondary antibodies, proteins were detected by ECL chemiluminescence.
**Rac1 Activity Assay:** Rac activity was assessed with activity assay kit (Upstate Biotechnology) using a GST-conjugated PAK-1 protein-binding domain peptide, PAK-1 PBD, which binds only to Rac-GTP, according to manufacturer’s instructions.

**Detergent-free Purification of Caveolin-rich Membrane Fractions:** Caveolae/lipid raft fractions were separated by the sodium carbonate-based detergent-free method. Briefly, VSMCs were scraped into 2 ml of 500 mM sodium carbonate containing protease inhibitors (pH 11). Homogenization was carried out sequentially in the following order using a loose-fitting Dounce homogenizer (10 strokes) and a sonicator (four 20-s bursts). The homogenates were adjusted to 45% sucrose by adding 90% sucrose in a buffer containing 25 mM Mes (pH 6.5) and 0.15 M NaCl and placed at the bottom of an ultracentrifugation tube. A 5–35% discontinuous sucrose gradient was formed above and centrifuged at 39,000 rpm at 4°C for 16–20 h in a Beckman SW-40Ti rotor. From the top of the tube, 13 fractions were collected. Caveolae/lipid raft fractions were accumulated in fractions 4–6. These fractions contained <5% of total tissue protein content.

In some experiments, detergent-free OptiPrep gradient cell fractionation was also performed, as previously described. Briefly, plasma membranes were isolated by Percoll gradient fractionation and then caveolae membranes were separated by two OptiPrep density gradients, a continuous 23% to 10% gradient followed by centrifugation against a discontinuous gradient to concentrate the lightest material. Proteins from each fraction were analyzed by Western Blotting.

**Copper Measurements by Inductively coupled plasma mass spectrometry (ICP-MS):** Samples were diluted to 2.0 ml with deionized water containing 5% v/v nitric acid. The copper contents were analyzed by ICP-MS, using a PlasmaQuad3, as previously described. Copper concentrations were calculated from calibration curves, and values for water blank were subtracted.

**Radioactive $^{64}$Cu Uptake and Efflux Studies:** For copper uptake experiments, fresh serum free DMEM medium of 1.8 ml was added to the growth-arrested VSMCs, as described previously. Cells were incubated for 30 min at 37 °C. Copper uptake was initiated by adding 200 µl of 10×CuCl$_2$ solution (20 µM) containing trace levels of $^{64}$Cu (MIR radiological sciences, Washington University Medical School) and 50 ng/ml PDGF. Cells were incubated for 5 min (non-specific binding to the cells) or 30 min at 37 °C, and copper transport was terminated by the addition of ice-cold stop buffer (150 mM NaCl, 5 mM KCl, 2.5 mM MgCl$_2$, 25 mM Heps, pH 7.4, and 10 mM Na$_2$EDTA), after which cells were washed 3 additional times with ice-cold stop buffer. Cells were lysed using 0.1 M NaOH, and aliquots were counted using a liquid scintillation-counter (Beckman-Coulter LS6500). All transport determinations were carried out in triplicate. To exclude non-specific Cu uptake, the count of 5 min was subtracted from the count of 30 min. Copper uptake was then expressed as pmol of copper taken up by the cells per mg of protein after determination of the protein content of the cell monolayer by the Bio-Rad protein assay (Bio-Rad #500-0006). For copper efflux experiments, cells were pulsed with 2 µM CuCl$_2$ solution containing trace levels of $^{64}$Cu for 2 h, followed by three quick rinses with prewarmed serum free DMEM containing 2 µM CuCl$_2$ solution to remove any residual copper, and then incubated with prewarmed serum free DMEM containing 2 µM CuCl$_2$ solution and 50 ng/ml PDGF for several chase periods. Culture medium was collected at different time intervals and counted as above.

**Biotinylation of cell surface ATP7A and CTR1:** The cell surface ATP7A and CTR1 was biotinylated following exposure to VSMCs to 50 ng/ml of PDGF for indicated time. Biotinylation of cells was carried out using the cell-impermeable, thiol-cleavable Sulfo-NHS-SS-biotin (Pierce) reagent to label cell surface proteins, as previously described. All of the biotinylation procedures
were carried out at 4 °C. Briefly, cell-surface proteins were labeled for 45 min with 0.2 mg/ml cleavable water-soluble cell-impermeant sulpho-NHS-SS-biotin (Pierce Endogen) in PBS supplemented with 0.5 mM MgCl$_2$ and 1 mM CaCl$_2$. The unbound sulfo-NHS-SS-biotin was quenched by 50 mM Tris (pH 8.0) containing 100 mM NaCl. The cells were lysed in 500 μl of ice-cold lysis buffer, pH 7.4 (50 mM HEPES, 5 mM EDTA, 100 mM NaCl), 1% Triton X-100, 60 mM n-Octyl-β-D-glucopyranoside, protease inhibitors (10 μg/ml aprotinin, 1 mM phenylmethylsulfonyl fluoride, 10 μg/ml leupeptin) and phosphatase inhibitors (50 mM sodium fluoride, 1 mM sodium orthovanadate, 10 mM sodium pyrophosphate), and precipitated on streptavidin–agarose beads (Pierce Endogen) overnight. The beads were washed three times in lysis buffer, and were eluted in loading buffer. Proteins were subjected to Western immunoblot analysis to detect biotinylated ATP7A and CTR1 using anti-ATP7A or CTR1 antibodies.

**Vascular Injury and Morphometric Analysis:** Animal protocols were approved by the Animal Care and Use Committee of the University of Illinois at Chicago and University of Minnesota. Male ApoE deficient mice were fed a western diet for 2 weeks, followed wire injury, then continued western diet for 4 more weeks, euthanized, and collected injured arteries.

Wire-induced injury of the carotid artery in ApoE deficient atherosclerotic mice were performed, as described$^{12}$. Briefly, mice were anesthetized using an intraperitoneal injection of ketamine (80 mg/kg body weight) and xylazine (5 mg/kg) (Phoenix Scientific, Inc., St. Joseph, MO). After midline neck incision, the left external carotid artery was tied off distally and a 0.014-inch flexible angioplasty guide wire was advanced by 1 cm along the common carotid artery via transverse arteriotomy. Complete and uniform endothelial denudation was achieved by five passes with a rotating motion. At different time points after injury, mice were anesthetized and perfused in situ using 4% paraformaldehyde at 100 mm Hg for tissue fixation. Injured arteries were excised and embedded in paraffin.

**Immunohistochemical analysis:** Paraffin-embedded tissue sections were stained with anti-ATP7A antibody (1:200), followed by a HRP-conjugated donkey anti-chicken IgY (1:100; Jackson ImmunoResearch). Staining was developed using the DAB Kit (Vector Laboratories, Burlingame, CA).

**Confocal Immunofluorescence Microscopy in tissues:** Paraffin-embedded tissue sections were stained with anti-ATP7A antibody (1:200) and anti-actin (for a smooth muscle isoform of actin) antibody (1:500), followed by a FITC-conjugated goat anti-IgY (1:100) and Rhodamine Red-conjugated goat anti-mouse IgG (1:50).

**Synchrotron X-ray Fluorescence Microscopy:** Sections (5 μm thick) of formalin-fixed paraffin-embedded wire-injured femoral artery were used. For X-ray imaging, the sections were mounted intact on silicon nitride windows (area, 2 x 2 mm; thickness, 200 nm) manufactured by Silson (Blisworth, U.K.) and attached by brief heating to 55°C, as previously described$^{13}$. Specimens were imaged with the scanning X-ray fluorescence microprobe at beamline 2-ID-E of the Advanced Photon Source (Argonne, IL). Undulator-generated x-rays of 10-keV incident energy were monochromatized with a single bounce Si <111> monochromator and focused to a measured spot size of 0.3 x 0.5 μm using Fresnel zone plate optics (X-radia, Concord, CA). Sections were raster-scanned in steps of 4.0 μm, and fluorescence spectra were collected for 1- to 2-sec dwell times by using a single-element silicon drift detector (Vortex-EX, SII Nanotechnology, CA). Quantitation and image-processing of the X-ray fluorescence (XRF) data sets was performed with MAPS software$^{14}$. Quantitation of elemental content was achieved by fitting XRF spectra at each pixel, and comparing
against a calibration curve derived from measurements of thin-film standards NBS-1832 and NBS-1833 (National Bureau of Standards, Gaithersburg, MD).

**Lysyl oxidase (LOX) activity assay.** LOX activity in cultured medium was measured by a high-sensitivity fluorescence assay as previously described \(^{15}\). Protein samples was incubated in the presence and absence of 500 µmol/L BAPN at 37°C for 30 minutes with 1 U/mL horseradish peroxidase, 10 µmol/L Amplex red (Molecular Probes), and 10 mmol/L 1,5-diaminopentane in 1.2 mol/L urea and 0.05 mol/L sodium borate (pH 8.2). The reaction was stopped on ice, and differences in fluorescence intensity (563-nm excitation wavelength and 587-nm emission wavelength) between samples with and without BAPN were determined. Results were normalized by cell protein content.

**Statistical analysis:** Statistical analysis was performed using the one-way analysis of variance (ANOVA), with the Scheffe test for post hoc comparisons when significance was determined by ANOVA or Student’s \(t\)-test at the analysis indicated in the figures. The accepted level of significance was set at \(p < 0.05\).
Supplemental Reference


Online Figure I: siRNA-mediated knockdown of endogenous ATP7A and CTR1 expression in VSMCs. 

A, Real-time PCR analysis of total RNA extracted from RASMs transfected with either ATP7A, CTR1, or control siRNA. B, Western-blot analysis of total cell lysates from RASMs transfected with either ATP7A or control siRNA, using anti-ATP7A (upper) and anti-actin (lower) antibodies. Actin was used as a loading control.
Online Figure II: Effect of ATP7A siRNA on PDGF-induced VSMC proliferation. Cell proliferation stimulated with or without 50 ng/ml of PDGF in RASMs was determined by cell number after plating cells in 0.1 % bovine serum containing culture medium for 72 hour, as previously described. Values are the mean ± S.D. for 3 independent experiments. NS, not significant.
Online Figure III: PDGF stimulation promoted ATP7A translocation from transGolgi network (TGN) to the plasma membrane leading edge in VSMCs. **A**, Growth-arrested RASMs were stimulated with 50 ng/ml PDGF for indicated time, and cells were stained for ATP7A. **B**, Growth-arrested RASMs were stimulated with or without 50 ng/ml PDGF, and cells were co-stained for ATP7A and syntaxin 6, a TGN marker. Small white arrowheads point to the plasma membrane leading edge, while white arrows point to TGN (PDGF (-)cells). Results are representative of 3 independent replicates of immunofluorescence images.
Online Figure III: C, ATP7A is translocated to the leading edge in MASMs stimulated by PDGF. Growth-arrested MASMs were stimulated with or without 50 ng/ml PDGF for 5 min in a similar fashion to Figure 2. Cells were co-stained with ATP7A and phalloidin. Small white arrowheads point to the leading edge and large arrows point to direction of migration.
Online Figure IV: PDGF stimulation increased active, GTP-bound form of Rac1 within 1 min, which was not affected by either ATP7A or CTR1 siRNA. A, Effect of PDGF on Rac1 activation. Growth-arrested RASMs were stimulated with 50 ng/mL of PDGF for indicated times (min). Lysates bound to p21-activated kinase 1 agarose were immunoblotted with anti-Rac antibody. B, Effect of ATP7A or CTR1 siRNA on PDGF induced Rac1 activation. Growth-arrested RASMs transfected with ATP7A, CTR1, or control siRNA were stimulated with 50 ng/mL of PDGF for 2 min. Lysates bound to p21-activated kinase 1 agarose were immunoblotted with anti-Rac antibody. C, PDGF stimulation promoted Rac1 association with ATP7A in a time-dependent manner. Growth-arrested RASMs was stimulated with 50 ng/mL of PDGF for indicated times (min). Lysates were IP with anti-Rac1 antibody, followed by IB with ATP7A.
Online Figure V: The ATP7A copper transporter and the CTR1 copper importer are localized in caveolae/lipid raft fraction which contains PDGFR and Rac1 in VSMCs. A and B, The CTR1 copper importer is predominantly found in caveolae/lipid rafts, as assessed by CTR1 specific antibodies. A, Identification of the CTR1 copper importer in caveolin-enriched lipid rafts fractions in mouse embryonic fibroblasts isolated from control, but not from CTR1 deficient mice. Cells were fractionated to 13 fractions by sucrose gradient centrifugation, followed by immunoblotting with anti-CTR1 antibody against either the intracellular carboxyl-terminal tail or the intracellular loop. B, Identification of the CTR1 copper importer in caveolin-enriched lipid rafts fractions in RASMs transfected with control siRNA, but not with CTR1 siRNA, using CTR1 specific antibodies as described above. C, Identification of both ATP7A and CTR1 in caveolae/lipid raft fraction in various VSMCs, including RASMs, HASMs, and MASMs. D, Identification of both ATP7A and CTR1 in caveolae/lipid raft fraction in RASMs, using detergent-free OptiPrep gradient cell fractionation method, as previously described.
Online Figure VI: ATP7A is translocated to the lipid rafts localized at the leading edge in actively migrating VSMCs, thereby promoting lamellipodia formation. A, CTxB accumulated and colocalized with F-actin at the leading edge in PDGF stimulated VSMCs. Growth-arrested RASMs were treated with or without 10 mM of methyl-β-cyclodextrin for 2 h, and stimulated with 50 ng/mL of PDGF for 5 min. VSMCs were stained with phalloidin (green) and cholera toxin subunit B (CTxB) (red). All fluorescence images were taken at 5 different fields/well, and the cell images are representative of more than 3 different experiments. B and C, ATP7A is translocated to the lipid rafts localized at the leading edge in VSMCs after wound scratch, thereby promoting lamellipodia formation. Growth-arrested confluent monolayer of RASMs were stained with either anti-ATP7A antibody (green) and cholera toxin subunit B (CTxB)-Alexa 555 (red) (B) or phalloidin (green) and cholera toxin subunit B (CTxB)-Alexa 555 (red) (C) after wound scratch in the presence of PDGF for 18 hours. Small white arrowheads point to the leading edge and large arrows point to direction of migration. All fluorescence images were taken at 5 different fields/well, and the cell images are representative of more than 3 different experiments.
Online Figure VII: Copper-uptake and efflux assessed by $^{64}$Cu copper transport assays were not altered by PDGF in VSMCs. A, $^{64}$Cu uptake assays in PDGF stimulated VSMCs. Growth-arrested RASMs were treated with 2 μM CuCl$_2$ containing trace levels of $^{64}$Cu and 50 ng/ml PDGF. Cells were incubated for 5 min (non-specific binding to the cells) or 30 min at 37 °C. Uptaken $^{64}$Cu was counted using a $\gamma$-counter. B, $^{64}$Cu efflux assays in PDGF stimulated VSMCs. Growth-arrested RASMs were pulsed with 2 μM CuCl$_2$ solution containing trace levels of $^{64}$Cu for 2 h, followed by rinses with prewarmed serum free DMEM containing 2 μM CuCl$_2$ solution, and then incubated with prewarmed serum free DMEM containing 2 μM CuCl$_2$ solution and 50 ng/ml PDGF for several chase periods. Culture medium was collected at different time intervals and counted using a $\gamma$-counter. The results shown here are the means ± S.D. for a single experiment and are representative of those obtained from three independent experiments.
Online Figure VIII: PDGF stimulation reduces copper content in caveolae/lipid raft fractions in an ATP7A-dependent manner in VSMCs. Copper contents were measured by inductively coupled plasma mass spectrometry (ICP-MS) in caveolae/lipid rafts in RASMs transfected with ATP7A or control siRNA, followed by stimulated with or without PDGF for 5 min. Equal amount of proteins in caveolae/lipid rafts (fraction 4-5) were obtained by sucrose gradient fractionation as described in Figure 5. (Mean ±S.D. n=3) *p<0.05 vs. unstimulated cells.
Online Figure IX: Plasma membrane localization of ATP7A and CTR1 in PDGF-stimulated VSMCs using cell surface biotinylation assays. The cell surface ATP7A and CTR1 was biotinylated following exposure to RASMs to 50 ng/mL of PDGF for indicated time. Biotinylated proteins were precipitated using streptavidin-agarose beads and separated by SDS/PAGE as described in the Method section. Biotinylated ATP7A and CTR1 were detected by Western-blots using anti-ATP7A and CTR1 antibodies.
Online Figure X: PDGF stimulation decreases Pro-LOX level in lipid raft fractions in an ATP7A-dependent manner in VSMCs. Equal amounts of proteins in caveolae/lipid rafts (fraction 4-5) were immunoblotted with anti-LOX-PP or caveolin-1 antibodies. RASMs were transfected with CTR1 or control siRNA, followed by stimulated with or without 50 ng/ml of PDGF for 5 min. Sucrose gradient centrifugation was performed as described in Figure 5. Right panel shows averaged data for ATP7A protein, expressed as % of unstimulated, control siRNA-treated VSMCs (means ± S.D, n=3). *p<0.05 vs. control siRNA-treated cells.
Online Figure XI. Immunofluorescence analysis for uninjured (control) or injured (3 days after) carotid arteries of ApoE-deficient atherosclerotic mice. Vessels were co-stained with anti-ATP7A (green) and α-smooth muscle actin (red) antibodies.
Online Figure XII:  A, Role of ATP7A in lysyl oxidase (LOX) activity in VSMCs. RASMs were transfected with control siRNA or ATP7A siRNA with or without PDGF treatment. LOX secreted in the culture medium was collected at 48-hour after addition of PDGF (50ng/ml). Its activity was measured by the oxidation of Amplex Red (Invitrogen) in the presence of 1,5-diaminopentane substrate, as previously described15. The results are presented as mean ± SE from 3 separate experiments. * P < 0.01 vs. control siRNA cells with or without PDGF treatment. B, Role of LOX activity in VSMC migration. RASMs were treated with beta-aminopropionitrile (BAPN)(100uM for 24h), an irreversible inhibitor of lysyl oxidase activity18. Cell migration was assessed by the modified Boyden chamber assay after stimulation with or without 50 ng/ml of PDGF for 8 hours. Bar graph represents averaged data, expressed as cell number per field. *p<0.05 vs. untreated cells. Values are the mean ± S.D. for 3 independent experiments. C, Role of SOD3 in VSMC migration. Wound scratch assay was performed in confluent monolayers of RASMs transfected with siRNA (two kinds of SOD3 siRNAs, or control siRNA) in the presence of PDGF (50 ng/ml), as described in Figure 1E. Images were captured immediately after rinsing at 0 h and at 24 h after the wounding in the cells. Bar graph represents averaged data, expressed as cell number per field. *NS vs. control siRNA-treated cells. Values are the mean ± S.D. for 3 independent experiments.