

# Bone Morphogenetic Protein 4 Promotes Vascular Smooth Muscle Contractility by Activating MicroRNA-21 (miR-21), which Down-regulates Expression of Family of Dedicator of Cytokinesis (DOCK) Proteins<sup>\*[5]</sup>

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**Background:** miR-21 expression is regulated by BMP4 and plays a critical role in vSMC phenotype regulation.

**Results:** Affinity purification of mRNAs associated with miR-21 yielded nearly all members of the DOCK superfamily.

**Conclusion:** miR-21 targets multiple members of the DOCK superfamily and modulates the activity of Rac1 small GTPase to regulate vSMC phenotype.

**Significance:** This study identified novel targets of miR-21 using a biochemical method.

The bone morphogenetic protein 4 (BMP4) signaling pathway plays a critical role in the promotion and maintenance of the contractile phenotype in vascular smooth muscle cell (vSMC). Misexpression or inactivating mutations of the BMP receptor gene can lead to dedifferentiation of vSMC characterized by increased migration and proliferation that is linked to vascular proliferative disorders. Previously we demonstrated that vSMCs increase microRNA-21 (miR-21) biogenesis upon BMP4 treatment, which induces contractile gene expression by targeting programmed cell death 4 (*PDCD4*). To identify novel targets of miR-21 that are critical for induction of the contractile phenotype by BMP4, biotinylated miR-21 was expressed in vSMCs followed by an affinity purification of mRNAs associated with miR-21. Nearly all members of the dedicator of cytokinesis (DOCK) 180-related protein superfamily were identified as targets of miR-21. Down-regulation of *DOCK4*, *-5*, and *-7* by miR-21 inhibited cell migration and promoted cytoskeletal organization by modulating an activity of small GTPase. Thus, this study uncovers a regulatory mechanism of the vSMC phenotype by the BMP4-miR-21 axis through DOCK family proteins.

MicroRNAs (miRNAs)<sup>2</sup> regulate basic cellular activities, such as differentiation, proliferation, and development, by sup-

pressing gene expression post-transcriptionally (1, 2). In mammals, the active strand miRNA sequence of ~22 nucleotides recognizes partially complementary miRNA recognition element (MRE) or seed-matched sequences located mostly in the 3'-UTR of mRNAs (3, 4). Seed-matched sites are defined as Watson-Crick consecutive base pairing between mRNAs and the miRNA positioned at 2–7 nucleotides from the 5'-end. Gene suppression by miRNAs is thought to occur either by inhibiting translation, mediating mRNA destabilization, or a combination of both. Aberrant miRNA expression is associated with various developmental abnormalities and human diseases (5, 6). miR-21 is involved in tumorigenesis and cardiovascular function (7–12). Elevated expression of miR-21 is observed in various types of tumors, and suppression of miR-21 in tumors leads to reduced proliferation and migration, suggesting that miR-21 acts as an oncogene (13–22). Higher levels of miR-21 are observed after multiple cardiac stresses, including acute pressure overload by transaortic constriction, chronic calcineurin activation, infusion of angiotensin II, and myocardial infarction (10, 23, 24). Cardiac hypertrophy and fibrosis in response to pressure overload after transaortic constriction are prevented by knockdown of miR-21 using antisense RNA oligonucleotides against miR-21 (11), suggesting a critical role for miR-21 in cardiac hypertrophy and fibrosis.

One of the effects of BMP and transforming growth factor- $\beta$  (TGF- $\beta$ ) signaling is to promote the contractile phenotype in vSMC. We identified miR-21 as a critical downstream mediator

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<sup>[5]</sup> This article contains supplemental Figs. S1–S4, Tables S1 and S2, and a list of primers.

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<sup>2</sup> The abbreviations used are: miRNA, microRNA; DOCK, dedicator of cytokinesis; miR-21, microRNA-21; vSMC, vascular smooth muscle cell; BMP, bone mor-

phogenetic protein; PDCD4, programmed cell death 4; MRE, miRNA recognition element; pri-miR-21, primary transcripts of miR-21; PASM, pulmonary artery smooth muscle cell; cel-miR-67, control *C. elegans* miR-67 mimic; bio-miR-21, 3'-biotinylated miR-21 mimic; bio-cel-miR-67, 3'-biotinylated control *C. elegans* miR-67 mimic; qRT, quantitative reverse transcription; anti-miR-21, antisense oligonucleotides against miR-21; ZEB, zinc finger homeodomain enhancer-binding protein.

of this effect (8). Upon TGF- $\beta$  or BMP treatment, the R-Smad proteins, which are signal transducers of TGF- $\beta$  and BMPs, associate with primary transcripts of miR-21 (pri-miR-21) in complex with Drosha to promote pri-miR-21 to precursor miR-21 processing and elevated miR-21 expression (8, 25). By inducing miR-21, both BMP4 and TGF- $\beta$  lead to knockdown of a negative regulator of contractile gene expression, programmed cell death 4 (*PDCD4*) (8). We have demonstrated previously that platelet-derived growth factor-BB (PDGF-BB), a potent inducer of the synthetic phenotype in vSMCs, induces miR-221, which in turn promotes (i) down-regulation of contractile genes and (ii) cell proliferation through down-regulation of two target genes, *c-kit* and *p27Kip1*, respectively (26). Although down-regulation of *PDCD4* by miR-21 is critical for the regulation of vSMC contractile gene expression, it remained unclear whether other effects of TGF- $\beta$  or BMP on vSMC are also mediated by miR-21 and which miR-21 targets are critical mediators of such activities.

There are several approaches to identify targets of miRNAs. The most common uses bioinformatics algorithms to predict potential targets based on the existence of partial complementary MREs in the 3'-UTR. Another approach is to identify mRNAs that are down-regulated when a miRNA is overexpressed (8, 27, 28). miRNA targets can also be enriched by co-immunoprecipitation with exogenously expressed tagged Argonaute (Ago) or GW182 proteins in cells overexpressing miRNA (29, 30). Some studies used stable isotope labeling with amino acids in culture to identify proteins that are down-regulated upon overexpression of specific miRNAs (31, 32).

In this study, we expressed biotinylated miR-21 in vSMCs and isolated mRNAs that are associated with miR-21 using streptavidin beads followed by microarray analysis for identification of targets. Multiple members of the *DOCK* family, which are guanine nucleotide exchange factors for Rac and/or Cdc42 (33), were found to be associated with miR-21 in cells. *DOCK* family genes are not predicted as targets of miR-21 by conventional target prediction algorithms, and little is known about their function in vSMC. Identification of 3'-UTR sequences, which are critical for recognition by miR-21, revealed that miR-21 regulates *DOCK* genes by base pairing at sites other than the seed sequence. We also demonstrate a function for *DOCK* family members in the regulation of vSMC phenotype by BMP4.

## EXPERIMENTAL PROCEDURES

**Cell Culture**—Human primary pulmonary artery smooth muscle cells (PASCs) were purchased from Lonza (CC-2581) and maintained in Sm-GM2 medium (Lonza) containing 5% fetal bovine serum (FBS). COS7 and rat pulmonary artery smooth muscle PAC1 cells were maintained in Dulbecco's modified Eagle's medium (DMEM) supplemented with 10% FBS. Cells were cultured at 37 °C in the presence of 5% CO<sub>2</sub>.

**Isolation of miR-21 Targets**—3'-Biotinylated miR-21 mimic (bio-miR-21) and 3'-biotinylated control *Caenorhabditis elegans* miR-67 mimic (bio-cel-miR-67) were synthesized by Dharmacon. PASCs were transfected with bio-miR-21 or bio-cel-miR-67 mimic at a final concentration of 30 nM using RNAiMax (Invitrogen). Twenty-four hours later, the cells were

trypsinized and lysed in TNE buffer (20 mM Tris (pH 7.5), 100 mM KCl, 5 mM MgCl<sub>2</sub>, 0.3% Nonidet P-40, RNase OUT (Invitrogen), protease inhibitor mixture (Roche Applied Science)). The streptavidin-coated magnetic beads (Invitrogen) were incubated with cell lysates at 4 °C for 4 h followed by washing with a washing buffer (same as TNE buffer except containing 200 mM KCl). Total mRNAs bound to the streptavidin beads (pull-down RNA) were isolated using TRIzol LS (Invitrogen) according to the manufacturer's instructions. For microarray analysis, isolated mRNAs were subjected to Affymetrix human U133plus 2.0 array at the Clinical Microarray Core, University of California, Los Angeles. Fold enrichment of mRNAs by miR-21 over control cel-miR-67 was calculated.

**Quantitative Reverse Transcription-PCR (qRT-PCR)**—Total RNAs were isolated using TRIzol (Invitrogen) and reverse transcribed using a first strand cDNA synthesis kit (Invitrogen) according to the manufacturer's instructions. qRT-PCR analysis was performed in triplicates using the SYBR Green Master Mix (Applied Biosystems). The mRNA level was normalized to glyceraldehyde-3-phosphate dehydrogenase (GAPDH). For a quantitation of miRNAs, a TaqMan microRNA assay kit (Applied Biosystems) was used according to the manufacturer's instructions, and results were normalized to U6 snRNA. Primers for qRT-PCR analysis are listed in the supplemental information.

**Luciferase Reporter Constructs**—The full-length 3'-UTR sequences were cloned by RT-PCR from mRNAs isolated from PASCs. Predicted MRE sequences were cloned into the pISO vector (Addgene) containing the luciferase gene. Primers for cloning are listed in the supplemental information.

**Luciferase Assay**—COS7 cells were transfected with luciferase reporter constructs using FuGENE 6 (Roche Applied Science) and a  $\beta$ -galactosidase ( $\beta$ -gal) expression plasmid as an internal transfection control. Twenty-four hours later, cells were transfected using RNAiMax (Invitrogen) with 5 nM miR-21 mimic or control miRNA mimic. Luciferase assays were carried out, and luciferase activities were normalized to  $\beta$ -gal activities.

**RNA Interference**—Small interfering RNA (siRNA) duplexes were synthesized by Thermo Scientific. The target nucleotide sequences for *DOCK4* siRNA (5'-GUGUGAUGGCGUGA-CAGAUU-3'), *DOCK5* siRNA (5'-AUCCAUUGCUAUA-GAAGAAU-3'), and *DOCK7* siRNA (5'-AAGACGUU-CAAUGUCAAUAGAUU-3') were used. Negative control siRNA (Qiagen) was used as a control.

**In Vitro Migration Assay**—PASCs transfected with miR-21 mimic, antisense oligonucleotides against miR-21 (anti-miR-21), or siRNAs were plated in 6-well plates, and three scratch wounds were generated with a 200- $\mu$ l disposable pipette tip. Scratch wounds were photographed over 9 h with a Nikon inverted microscope with an attached digital camera, and their widths were quantitated with ImageJ software.

***DOCK7* Expression Plasmid**—*DOCK7* transcript with deleted 3'-UTR (*DOCK7*<sup>wo3'-UTR</sup>) was cloned into pcDNA3.1(+) by RT-PCR from mRNAs isolated from PASCs. For PCR amplification, primers 5'-ATGGCTAGCATGGCCGAGCGCCGCGCC-3' and 5'-CGTTCTAGATTAGAGATCCATTTTGCG-3' were used.

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**Collagen Matrix Contraction Assay**—A collagen matrix contraction assay was performed as described (34). PASCs transfected with miR-21 mimic, control miRNA mimic, anti-miR-21, or siRNAs were embedded in attached collagen matrices followed by 3 nM BMP4 treatment for 24 h. Twenty-four hours after detachment of the gel from the plate, the gel surface area was quantitated with the ImageJ program.

**Detection of Active Rac1**—To detect active GTP-bound Rac1, cell lysates of PASCs transfected with control miRNA mimic or miR-21 mimic were incubated with 20  $\mu$ g of GST-Pak1-PBD (Thermo Scientific), which binds specifically to the GTP-bound form of Rac1, in the presence of glutathione-agarose beads. To measure the total amount of Rac1, Western blot was also performed with anti-Rac1 (Thermo Scientific). The band intensity in the Western blot was quantified, and the levels of Rac1-GTP were normalized to the amount of total Rac1.

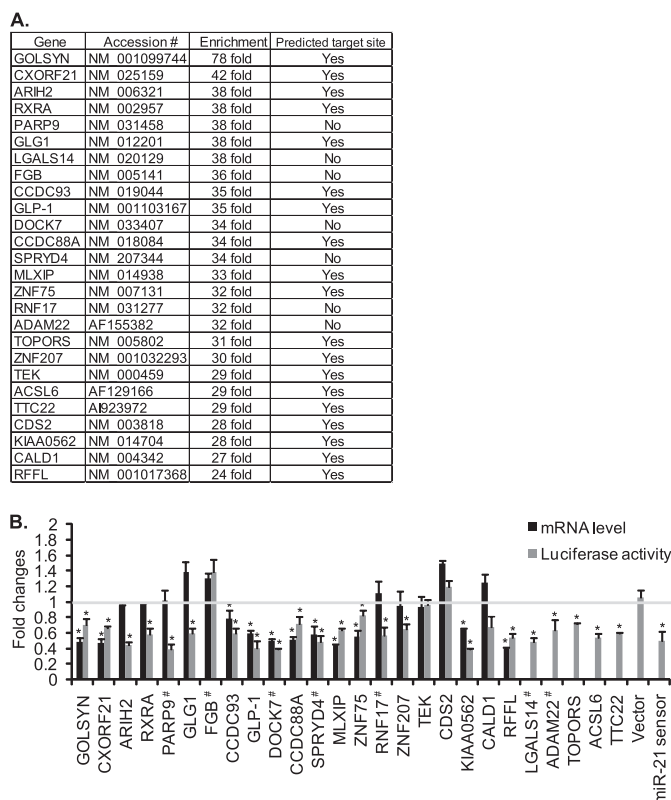
**Immunoblotting**—Cells were lysed in TNE buffer, and total cell lysates were separated by SDS-PAGE, transferred to PVDF membranes (Millipore), immunoblotted with antibodies, and visualized using an enhanced chemiluminescence detection system (Amersham Biosciences). Antibodies used for immunoblotting were: anti-GAPDH antibody (2E3-2E10, Abnova) and anti-human DOCK7 antibody (35). Protein bands were quantitated with the imaging analysis software ImageJ.

**Statistical Analysis**—The results presented are an average of at least three experiments, each performed in triplicate with standard errors. Statistical analyses were performed by analysis of variance followed by Tukey's multiple comparison test or by Student's *t* test as appropriate using Prism 4 (GraphPad Software Inc.). *p* values of <0.05 were considered significant and are indicated with asterisks.

## RESULTS

**Identification of Target mRNAs for miR-21**—To further understand the physiological role of miR-21 in vSMCs, targets of miR-21 were isolated using a modified method developed previously (36, 37). PASCs were transfected with bio-miR-21 or bio-cel-miR-67 mimic as a negative control. mRNAs associated with bio-miRNA mimic were isolated following affinity purification using streptavidin beads. mRNA targets were identified by comparing isolated pulldown RNAs from cells transfected with bio-miR-21 mimic with those of cells transfected with control mimic by human gene microarray analysis. There are currently 33 genes that are validated as targets of miR-21 according to the miRecords web site. 30 of 33 validated miR-21 targets were enriched more than 2-fold by miR-21 pulldown (supplemental Table S1). None of the housekeeping gene transcripts were enriched (supplemental Table S2). Among previously validated miR-21 targets, *E2F1*, *GLCCII*, and *SLC16A10* were not enriched significantly in PASCs (supplemental Table S1). However, *GLCCII* cannot be found on the microarray, and *SLC16A10* is not expressed in PASCs. Thus, the bio-miR-21 pulldown technique was able to detect all validated targets except one and has a potential to identify novel targets.

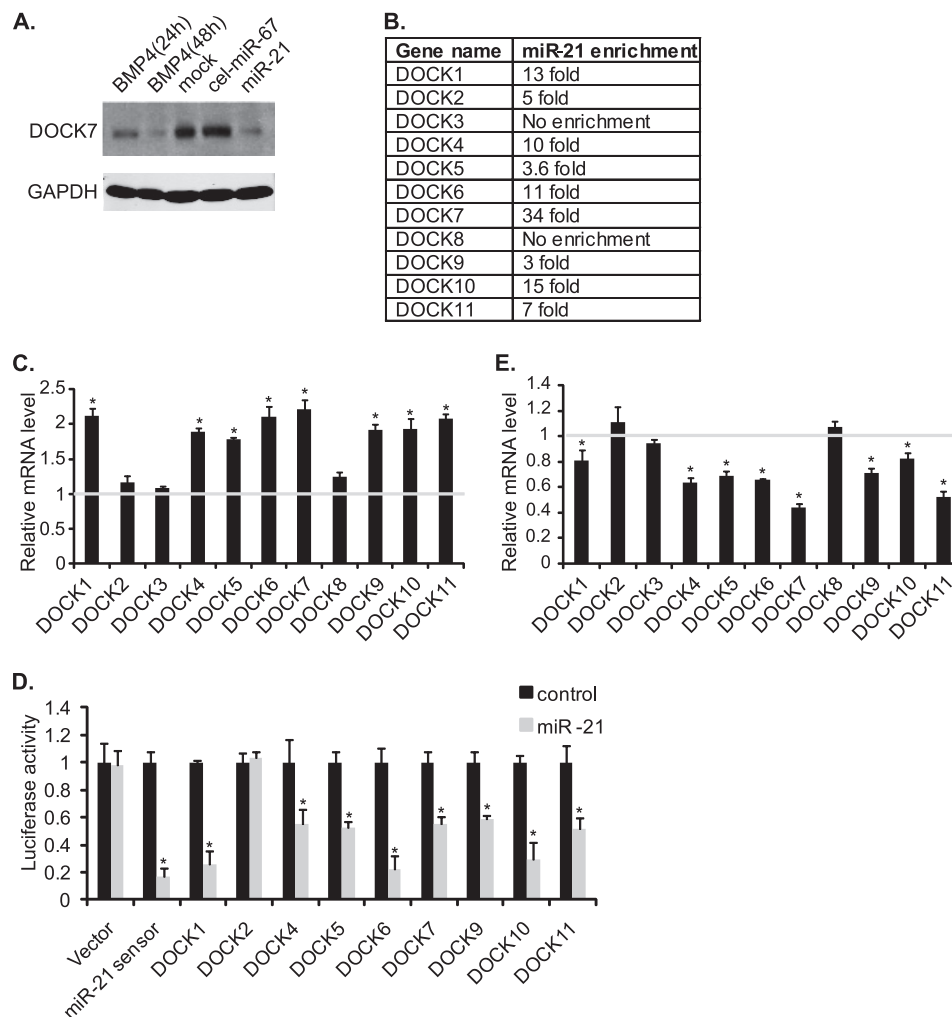
Of >9000 gene transcripts that were enriched by  $\geq 2$ -fold over control in bio-miR-21 pulldown samples, we examined 26 genes that were not identified previously as miR-21 targets and



**FIGURE 1. Genes enriched by miR-21 pulldown.** A, the 26 most highly enriched genes by bio-miR-21 pulldown in PASCs. The presence of predicted target sites for miR-21 was identified by using PITA and RNA22. Accession indicates NCBI Reference Sequence accession numbers. B, overexpression of miR-21 down-regulates genes that are highly enriched by bio-miR-21 pulldown. Total RNAs were harvested from PASCs transfected with control or miR-21 mimic, and mRNA levels of the indicated genes relative to *GAPDH* were measured by qRT-PCR. Fold change of mRNA levels of each gene in cells overexpressing miR-21 mimic in comparison with control (cel-miR-67) mimic-transfected cells is presented (black bars). COS7 cells transfected with luciferase construct containing full-length 3'-UTR or MRE of the indicated genes were transfected with control (cel-miR-67) or miR-21 mimic. # indicates the luciferase reporter constructs containing a full-length 3'-UTR, whereas the rest of the constructs contain only predicted miR-21 MREs. A luciferase vector carrying two copies of a sequence complementary to miR-21 (*miR-21 sensor*) and a luciferase vector without the 3'-UTR sequence (*Vector*) were used as positive and negative controls, respectively. Luciferase activity was then measured in triplicates. Fold changes by miR-21 mimic transfection in comparison with cel-miR-67 mimic transfection are presented (gray bars). Data represent mean  $\pm$  S.E., *p* < 0.01.

had the highest enrichment in bio-miR-21 pulldown samples ( $\geq 24$ -fold over bio-cel-miR-67 pulldown samples) (Fig. 1A). To investigate whether expression of these genes is down-regulated by miR-21, the full-length 3'-UTRs or predicted MREs of all 26 genes was inserted at the 3'-end of the luciferase gene, and luciferase reporter activity was measured upon overexpression of miR-21 or control cel-miR-67 mimic. As miR-21 MRE or seed sequence is not predicted by target prediction algorithms in the 3'-UTR of transcripts encoding *PARP9*, *LGALS14*, *FGB*, *DOCK7*, *SPRYD4*, *RNF17*, and *ADAM22*, full-length 3'-UTRs were examined. For the rest of the genes, predicted miR-21 MREs or seed regions were examined. As a positive control, a miR-21 sensor luciferase construct, which contains two copies of a sequence complementary to miR-21, was used. The miR-21 sensor activity was reduced to  $47 \pm 6\%$  when miR-21 mimic was expressed (Fig. 1B). Luciferase reporter activities of 22 genes





**FIGURE 2. miR-21 regulates multiple members of DOCK family proteins.** *A*, PSMCs were treated with 3 nM BMP4 (24 or 48 h) or transfected with control mimic (cel-miR-67) or miR-21 mimic. Total cell lysates were subjected to immunoblotting with anti-DOCK7 or anti-GAPDH (loading control) antibody. *B*, all members of the DOCK family except DOCK3 and DOCK8 were enriched 3–34-fold upon bio-miR-21 pulldown in comparison with control (bio-cel-miR-67) pulldown. *C*, total RNA was harvested from PSMCs transfected with control or anti-miR-21 oligonucleotides, and mRNA levels of the indicated DOCK genes relative to GAPDH were measured by qRT-PCR. Data represent fold changes of the mRNA levels in anti-miR-21-transfected cells in comparison with control oligonucleotide-transfected cells  $\pm$  S.E. from triplicates.  $*$ ,  $p < 0.01$ . *D*, COS7 cells were transfected with a luciferase reporter containing 3'-UTR of DOCK1, -2, -4, -5, -6, -7, -9, -10, or -11 gene transcript and cel-miR-67 (control) or miR-21 mimic. Relative luciferase activities were examined. The miR-21 sensor and an empty luciferase vector (Vector) were used as controls. Data represent fold changes in the luciferase activity in miR-21-mimic-transfected cells compared with control-mimic-transfected cells  $\pm$  S.E. from triplicates.  $*$ ,  $p < 0.01$ . *E*, PSMCs were treated with 3 nM BMP4 for 24 h. mRNA levels of the indicated DOCK genes relative to GAPDH were measured by qRT-PCR. Data represent fold changes of the mRNA level in BMP4-treated cells compared with mock-treated cells  $\pm$  S.E. from triplicates.  $*$ ,  $p < 0.01$ .

were significantly reduced, suggesting that  $\sim 85\%$  (22 of 26) of miR-21 pulldown mRNAs, which are enriched  $\geq 24$ -fold over control, are miR-21 targets (Fig. 1*B*, gray bars). Among these 22 genes, qRT-PCR analysis demonstrated that mRNAs for 11 genes (*GOLSYN*, *CXORF21*, *CCDC93*, *GLP-1*, *DOCK7*, *CCDC88A*, *SPRYD4*, *MLXIP*, *ZNF75*, *KIAA0562*, and *RFFL*) were significantly reduced by miR-21, indicating that these transcripts undergo destabilization of mRNA by miR-21 (Fig. 1*B*, black bars). Levels of expression of four genes (*FGB*, *TEK*, *CDS2*, and *CALD1*) were unchanged both by luciferase assay and at the mRNA level (Fig. 1*B*). Thus, using this method, we identified 22 novel targets of miR-21 in PSMCs (Fig. 1*B*).

**miR-21 Regulates Expression of DOCK Family Proteins**—One of the newly identified miR-21 targets is DOCK7, a member of the DOCK family of guanine nucleotide exchange factors for the Rho family of GTPases (Fig. 1*A*). Immunoblot analysis indi-

cated that the endogenous DOCK7 protein in PSMCs was down-regulated upon miR-21 overexpression in comparison with control cel-miR-67 (Fig. 2*A*). Furthermore, cells stimulated with BMP4 for 48 h exhibited reduced DOCK7 levels, similar to what was seen in miR-21-transfected cells, confirming that miR-21 leads to down-regulation of DOCK7 in PSMCs (Fig. 2*A*). Other members of the DOCK family genes, DOCK1, -2, -4, -5, -6, -9, -10, and -11, were also significantly enriched in miR-21 pulldowns (Fig. 2*B*). To determine whether miR-21 also targets other members of the DOCK family and mediates destabilization of their mRNAs similarly to DOCK7 (Fig. 1*B*), qRT-PCR analysis was performed upon overexpression of anti-miR-21, which inhibits miR-21 activity in PSMCs (Fig. 2*C*). Significant up-regulation of the mRNA levels of DOCK1, -4, -5, -6, -7, -9, -10, and -11 was observed after anti-miR-21 transfection, suggesting that these DOCK family mem-

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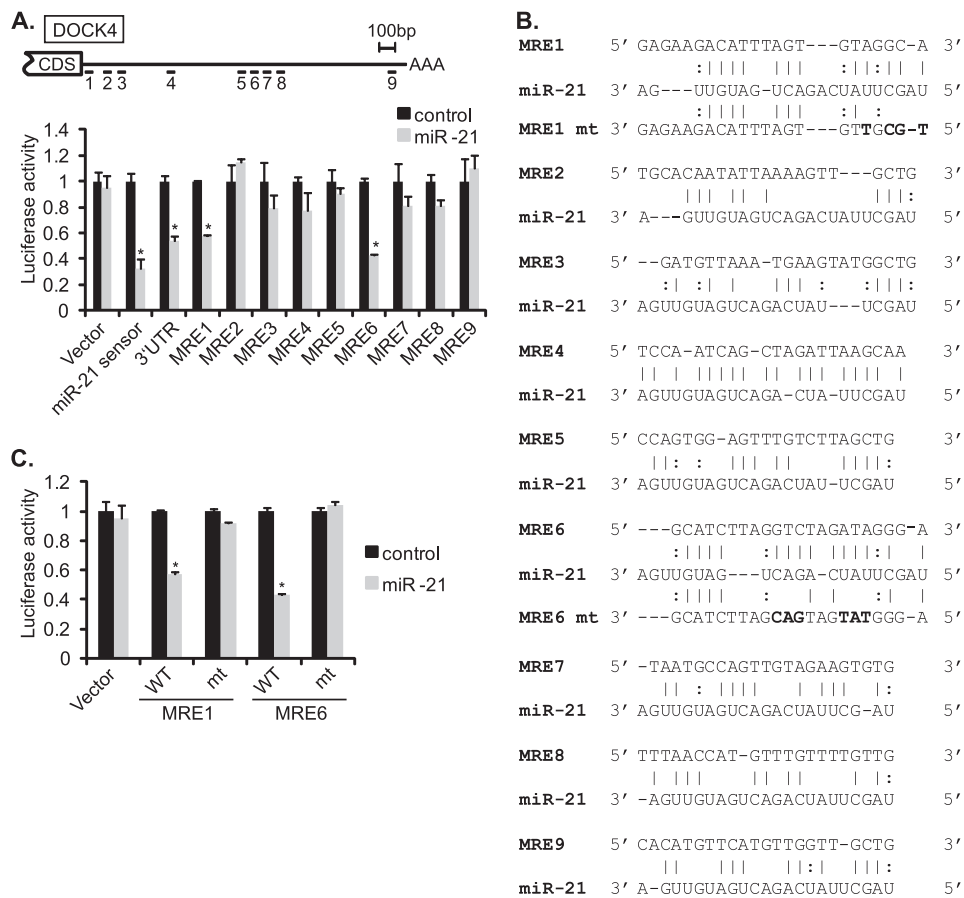


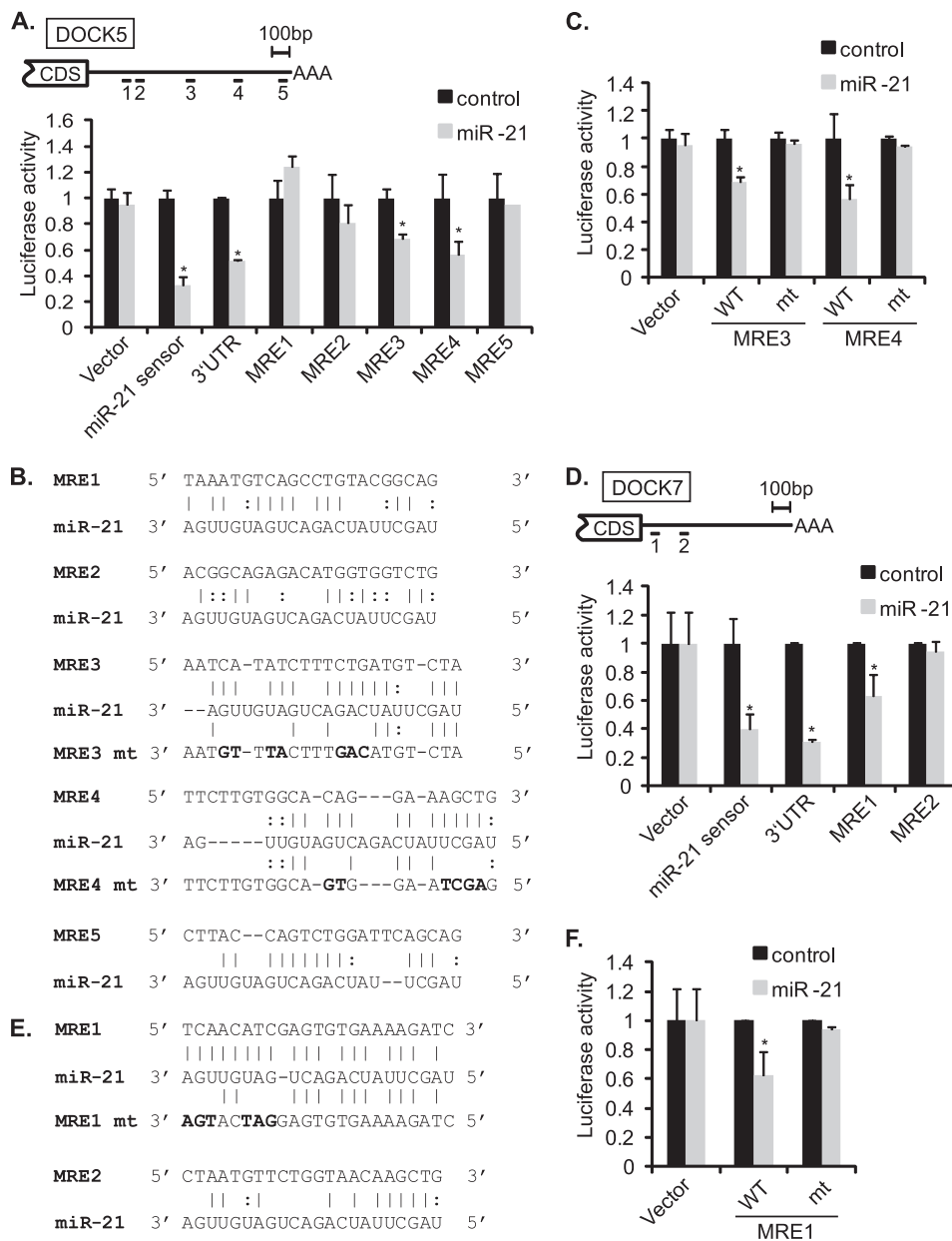
FIGURE 3. **DOCK4** is direct target of miR-21. *A*, schematic diagram of predicted miR-21 MREs in the 3'-UTR of *DOCK4* transcripts (top panel). CDS and AAA stand for protein coding sequence and poly(A) tail, respectively. Luciferase activity was examined in COS7 cells using the luciferase vector containing each of nine MREs or the full-length 3'-UTR of *DOCK4* mRNAs. Luciferase activity of each construct upon overexpression of miR-21 mimic relative to control mimic is shown (bottom panel). Data represent mean  $\pm$  S.E. \*  $p < 0.001$ . *B*, sequences of miR-21 MREs predicted by the RNA22 program in the 3'-UTR of *DOCK4*. Perfect base matches are indicated by a vertical line; G:U pairs are indicated by a colon (:). Mutations introduced in MRE1 or MRE6 to disrupt a base pairing with the miR-21 sequence are indicated in bold letters. *C*, luciferase activities of constructs with wild type or mutated *DOCK4* MRE1 (*MRE1 mt*) or MRE6 (*MRE6 mt*) were examined in COS7 cells by transfecting cel-miR-67 (control) or miR-21 mimic. Data represent mean  $\pm$  S.E. \*,  $p < 0.0001$ .

bers are down-regulated by endogenous miR-21 (Fig. 2C). To confirm that miR-21 targets members of the DOCK family, we measured the expression of constructs containing the 3'-UTR of *DOCK1*, -2, -4, -5, -6, -7, -9, -10, or -11 downstream of the luciferase reporter gene in the presence of miR-21 mimic (Fig. 2D). Consistent with the results of anti-miR-21 in Fig. 2C, the luciferase activity of the *DOCK1*, -4, -5, -6, -7, -9, -10, and -11 3'-UTR constructs were significantly reduced upon expression of miR-21 mimic, demonstrating that, in addition to *DOCK7*, *DOCK1*, -4, -5, -6, -9, -10, and -11 are targets of miR-21 (Fig. 2D). Although *DOCK2* transcripts were enriched in miR-21 pulldowns 5-fold over cel-miR-67 (Fig. 2B), the luciferase assay indicated that *DOCK2* is not a target of miR-21 (Fig. 2D). Finally, treatment of PSMCs with BMP4, which is known to induce miR-21 by  $\sim 2$ -fold (8), resulted in decreases in *DOCK1*, -4, -5, -6, -7, -9, -10, and -11 transcripts, further confirming that these members of the DOCK family are regulated by the BMP4-miR-21 axis (Fig. 2E and supplemental Fig. S4).

**Identification of miR-21 Recognition Sequence in 3'-UTR of *DOCK4*, -5, and -7 Transcripts**—*DOCK4*, -5, and -7 are known as guanine nucleotide exchange factors for Rac1 GTPases and play a role in cell migration and cytoskeletal organization in neuronal and tumor cells (33, 38). *DOCK4*, -5, and -7 were not

predicted as miR-21 targets by the algorithms whose prediction is based on a seed-matched sequence and cross-species conservation of the seed sequence. RNA22, which predicts targets based on a pattern-based method (39), found nine potential MREs in the 3'-UTR of *DOCK4* that are partially complementary to the miR-21 sequence (Fig. 3, A and B). To determine whether any of these predicted MREs are responsible for knockdown by miR-21, luciferase reporters containing individual MREs were transfected into COS7 cells together with miR-21 or control cel-miR-67 mimic. miR-21 significantly reduced the luciferase activity of MRE1 ( $43 \pm 1\%$ ) and MRE6 ( $58 \pm 1\%$ ) (Fig. 3A). Mutations in MRE1 or MRE6 (Fig. 3B, *mt*), which disrupted critical base pairing with miR-21, abrogated the inhibition of luciferase activity by miR-21 (Fig. 3C), suggesting that MRE1 and MRE6 are critical target sites for recognition of *DOCK4* mRNAs by miR-21.

There are five potential miR-21 MREs in the *DOCK5* 3'-UTR based on prediction by RNA22 (Fig. 4, A and B). miR-21 significantly reduced the luciferase activity of MRE3 ( $32 \pm 4\%$ ) and MRE4 ( $44 \pm 10\%$ ) (Fig. 4A). Mutations in MRE3 or MRE4 abolished the regulation by miR-21 (Fig. 4B, *mt*), indicating an essential role of these two MREs in the regulation of *DOCK5* mRNA by miR-21 (Fig. 4C). Although *DOCK7* does not contain



**FIGURE 4. DOCK5 and DOCK7 are miR-21 targets.** *A*, schematic diagram of predicted miR-21 MREs in the 3'-UTR of *DOCK5* transcripts (top panel). CDS and AAA stand for protein coding sequence and poly(A) tail, respectively. Luciferase activity was examined in COS7 cells using the luciferase vector containing each of five MREs or the full-length 3'-UTR of *DOCK5* mRNAs. miR-21 sensor and an empty luciferase vector (Vector) were examined as positive and negative controls, respectively. Luciferase activity of each construct upon overexpression of miR-21 mimic relative to control mimic is shown (bottom panel). Data represent mean  $\pm$  S.E. \*,  $p < 0.001$ . *B*, sequences of miR-21 MREs predicted by the RNA22 program in the 3'-UTR of *DOCK5*. Perfect base matches are indicated by a vertical line; G:U pairs are indicated by a colon (:). Mutations introduced in the MRE3 or MRE4 to disrupt a base pairing with miR-21 sequence are indicated in bold letters. *C*, luciferase activities of constructs with wild type or mutated *DOCK5* MRE3 (MRE3 mt) or MRE4 (MRE4 mt) were examined in COS7 cells by transfecting cel-miR-67 or miR-21 mimic. Data represent mean  $\pm$  S.E. \*,  $p < 0.001$ . *D*, schematic diagram of potential miR-21 MREs in the 3'-UTR of *DOCK7* transcripts (top panel). Luciferase activity was examined in COS7 cells using the luciferase vector containing each of two MREs or the full-length 3'-UTR of *DOCK7* mRNAs. miR-21 sensor and an empty luciferase vector (Vector) were examined as controls, respectively. Luciferase activity of each construct upon overexpression of miR-21 mimic relative to control mimic is shown (bottom panel). Data represent mean  $\pm$  S.E. \*,  $p < 0.01$ . *E*, sequences of miR-21 MREs in the 3'-UTR of *DOCK7*. Perfect base matches are indicated by a vertical line; G:U pairs are indicated by a colon (:). Mutations introduced in the MRE1 to disrupt a base pairing with the miR-21 sequence are indicated in bold letters (MRE1 mt). *F*, luciferase activities of constructs with wild type or mutated *DOCK7* MRE1 were examined in COS7 cells by transfecting cel-miR-67 or miR-21 mimic. \*,  $p < 0.01$ .

a seed match or predicted MRE in the 3'-UTR, it contains two sequence elements that are highly complementary to miR-21; MRE1 and MRE2 (Fig. 4, D and E). Luciferase activity of the construct containing MRE1, but not MRE2, was significantly reduced by miR-21 ( $38 \pm 2\%$ ) (Fig. 4D). Mutations in MRE1 (Fig. 4E, MRE1 mt) blocked down-regulation by miR-21, suggesting a critical role of MRE1 for recognition of *DOCK7* by

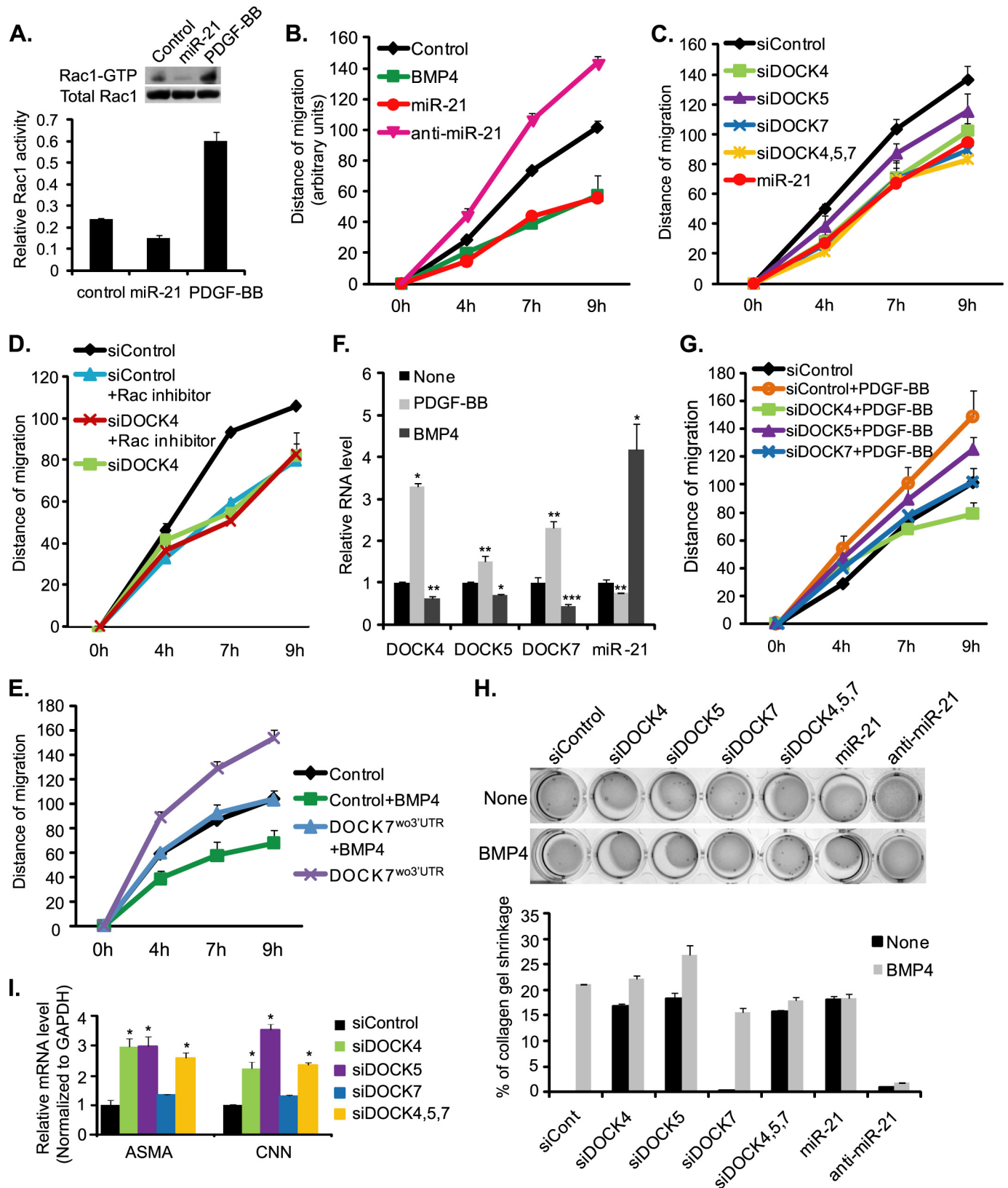
miR-21 (Fig. 4F). Thus, we identified sequence element(s) critical for recognition by miR-21 in all three *DOCK* gene transcripts.

**Down-regulation of DOCKs by miR-21 Inhibits Rac1 Activity and Functions**—As *DOCK4*, *-5* and *-7* are linked to the activation of the Rac1 GTPase (33, 38), we hypothesized that miR-21 might modulate vSMC migration and cytoskeletal organization

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by inhibiting Rac1 activity. To examine whether miR-21 modulates Rac1 GDP/GTP exchange activity in vSMCs, the amount of active Rac1 was measured by pull-down assays using the GST-Pak1-PBD domain fusion protein, which specifically binds

active Rac1. Treatment with PDGF-BB, which is known to activate Rac1, was included as a positive control and triggered a ~3-fold increase in active Rac1 (Fig. 5A). miR-21 reduced the level of active Rac1 by ~50% compared with control cel-miR-





67-transfected cells without affecting the amount of total Rac1 (Fig. 5A), suggesting that elevation of miR-21 decreases active GTP-bound Rac1. Next, we examined a role for miR-21 in cell migration by performing a scratch wound assay (34). To this end, a wound was created in a monolayer of PASCs, and images were captured at the beginning and at regular intervals during cell migration to close the wound. Similarly to the cells treated with BMP4 (Fig. 5B, green line), the migration distance was reduced to  $47 \pm 3\%$  9 h after a scratch wound was generated in miR-21-transfected cells (Fig. 5B, red line) compared with control miRNA-transfected cells (Fig. 5B, black line). In contrast with the miR-21 mimic, transfection of anti-miR-21 increased cell migration (Fig. 5B, magenta line), indicating that a differential level of expression of miR-21 determines vSMC migration. When *DOCK4*, -5, and -7 were down-regulated individually or simultaneously by siRNAs by  $54 \pm 3$ ,  $59 \pm 15$ , and  $64 \pm 4\%$ , respectively (supplemental Fig. S1A), cell migration was significantly reduced comparably with that of miR-21-overexpressing cells (Fig. 5C), suggesting that *DOCK4*, -5, and -7 regulate vSMC motility. This result also indicates a redundant function among *DOCK4*, -5, and -7 for the regulation of cell migration. When PASCs were treated with NSC23766, an inhibitor of Rac1 GDP/GTP exchange activity (38), cell migration was reduced similarly to that in *DOCK4*, -5, and -7 knockdown cells (Fig. 5D). Furthermore, exogenous expression of *DOCK7* mRNAs (*DOCK7*<sup>wo3'-UTR</sup>), which have the 3'-UTR deleted and are resistant to miR-21, in rat pulmonary artery smooth muscle PAC1 cells blocked BMP4-induced inhibition of migration (Fig. 5E, green square versus blue triangle). Thus, regulation of *DOCK4*, -5, and -7 levels is critical for the migratory function in vSMCs. Altogether, these results indicate that the GDP/GTP exchange activity of Rac1 is critical for vSMC migration, and the BMP4-miR-21 axis suppresses vSMC migration by down-regulation of DOCK proteins.

As PDGF-BB antagonizes BMP4 activity and promotes vSMC migration (34), we hypothesized that activation of Rac1 by PDGF-BB in PASCs (see Fig. 5A) might be in part due to down-regulation of miR-21, leading to increased expression of DOCK. qRT-PCR analysis demonstrated that miR-21 expression was reduced by 50% at 9 h (supplemental Fig. S3) and 26% at 24 h of PDGF-BB treatment and that *DOCK4*, -5, and -7 mRNAs were induced by  $\sim 1.5$ –3-fold by PDGF-BB (Fig. 5F). Furthermore, knockdown of *DOCK4*, -5, or -7 greatly reduced

PDGF-BB-mediated cell migration (Fig. 5G), suggesting that regulation of *DOCK4*, -5, and -7 levels is critical for the promigratory function of PDGF-BB in vSMCs.

Next, *in vitro* collagen gel contraction assays were performed in PASCs transfected with miR-21 mimic, anti-miR-21, or siDOCKs to evaluate the effect of miR-21-mediated down-regulation of DOCKs in the cytoskeletal organization of PASCs. Similarly to BMP4-treated cells, transfection of miR-21 mimic reduced gel size by  $\sim 20\%$  compared with control samples (Fig. 5H). Overexpression of miR-21 mimicked the effect of BMP4 and induced contraction, whereas transfection of anti-miR-21 abolished BMP4-induced contraction (Fig. 5H), indicating a critical role of miR-21 expression in contraction of the collagen gel. Down-regulation of *DOCK4* or *DOCK5* by siRNA also reduced the size of collagen lattice by  $18 \pm 3$  and  $20 \pm 4\%$ , respectively (Fig. 5H), which is similar to that of miR-21-transfected cells, suggesting that down-regulation of *DOCK4* or *DOCK5* by miR-21 is sufficient to mediate vSMC contraction. Knockdown of *DOCK7* showed no effect on contraction (Fig. 5H). Consistently, knockdown of *DOCK4* or *DOCK5* elevated the levels of contractile genes (*ASMA* and *CNN*), whereas siDOCK7 had no effect (Fig. 5I), suggesting a specific role of *DOCK4* and -5 in the regulation of contractile gene expression and contractility. Treatment of cells with the Rac1 inhibitor NSC23766 did not affect the contraction of the collagen lattice (supplemental Fig. S2). Thus, *DOCK4* and *DOCK5* promote cytoskeletal remodeling in a Rac1-independent manner. These results indicate that miR-21 modulates cell migration and contractility through down-regulation of DOCK family members in vSMCs. Therefore, the miR-21-DOCK axis regulates critical characteristics of the vSMC phenotype, such as motility and cytoskeletal organization, in response to BMP4 and PDGF signaling (Fig. 6).

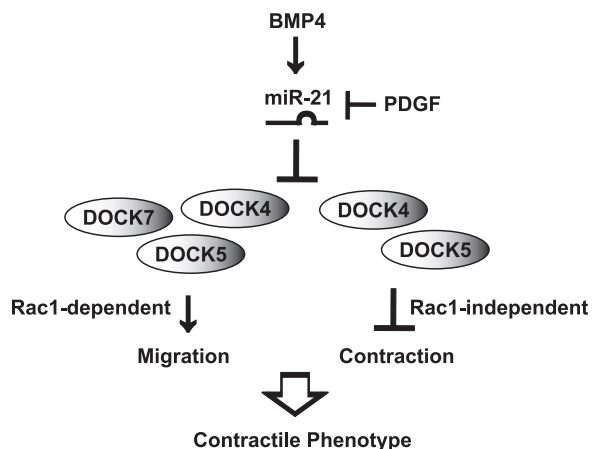
## DISCUSSION

In this study, we identified the DOCK family members as miR-21 targets downstream of BMP4 and provided evidence that their down-regulation by miR-21 is critical for promoting the contractile vSMC phenotype through inhibiting cell migration and promoting contractility (Fig. 6). The role of DOCK proteins in the regulation of cell motility, cytoskeletal organization, phagocytosis, and polarity has been elucidated in various cell types, especially in the nervous system (33). However,

**FIGURE 5. miR-21-mediated inhibition of DOCKs modulates vSMC motility.** A, PASCs were transfected with control cel-miR-67 mimic or miR-21 mimic. Active GTP-bound Rac1 was pulled down by GST-Pak1-PBD fusion protein and visualized by immunoblotting with anti-Rac1 antibody (top panel, *Rac1-GTP*). Immunoblotting of total cell lysates with anti-Rac1 antibody is shown to demonstrate total amount of Rac1 (top panel, *Total Rac1*). The level of Rac1-GTP relative to total Rac1 is presented as a relative Rac1 activity (bottom panel). B, PASCs transfected with control cel-miR-67 mimic (Control; black diamonds), miR-21 mimic (*miR-21*; red circles), or anti-miR-21 (*anti-miR-21*; magenta triangles) were subjected to the scratch wound assay. As a control, PASCs were treated with 3 nM BMP4 (green squares). The distance of migration was measured using ImageJ over 9 h after a scratch wound was introduced. C, PASCs transfected with 25 nM non-targeting control siRNA (siControl); siRNA against *DOCK4* (siDOCK4), *DOCK5* (siDOCK5), or *DOCK7* (siDOCK7); or miR-21 mimic (*miR-21*) were subjected to the scratch wound assay. D, PASCs transfected with siControl or siDOCK4 were subjected to the scratch wound migration assay in the presence or absence of 50  $\mu$ M Rac1 inhibitor (NSC23766). E, rat PASC PAC1 cells were transfected with control vector (Control) or *DOCK7* expression construct carrying the *DOCK7* cDNA deleted in the 3'-UTR (*DOCK7*<sup>wo3'-UTR</sup>) followed by the scratch wound assay in the presence or absence of 3 nM BMP4. F, levels of *DOCK4*, -5, or -7 mRNA relative to *GAPDH* or miR-21 relative to U6 snRNA were measured by qRT-PCR in PASCs stimulated with 20 ng/ml PDGF-BB or 3 nM BMP4 for 24 h. Data represent mean  $\pm$  S.E. \*,  $p < 0.0001$ ; \*\*,  $p < 0.01$ ; \*\*\*,  $p < 0.001$ . G, PASCs transfected with siRNA (siControl, siDOCK4, siDOCK5, or siDOCK7) were subjected to the scratch wound assay in the presence or absence of 20 ng/ml PDGF-BB. Data represent mean  $\pm$  S.E. H, PASCs were transfected with siControl, siDOCK4, siDOCK5, siDOCK7, miR-21 mimic, or anti-miR-21 followed by stimulation with 3 nM BMP4 or no stimulation (None) for 24 h. The PASC embedded collagen lattices were photographed by using a digital camera to measure gel contraction. The relative size of collagen gel was quantitated using ImageJ. Data represent the mean percentage of the collagen gel shrinkage compared with unstimulated control. I, levels of contractile genes.  $\alpha$ -Smooth muscle actin (*ASMA*) and calponin1 (*CNN*) mRNA levels relative to *GAPDH* were measured by qRT-PCR in PASCs transfected with siControl (siCont), siDOCK4, siDOCK5, or siDOCK7. Data represent mean  $\pm$  S.E. \*,  $p < 0.01$ .



## Direct Identification of miR-21 Targets in vSMCs



**FIGURE 6. Schematic representation of cellular effects of miR-21-mediated down-regulation of DOCK4, -5, or -7 in vascular smooth muscle cells.** BMP4 induces miR-21 and inhibits migration and increases contractility of vSMCs. miR-21 targets *DOCK4*, -5, and -7 to control vSMC migration by modulating the activity of Rac1, whereas *DOCK4* and -5 modulate contractility of vSMC via a Rac1-independent mechanism.

no study on DOCK proteins has been performed in vascular cells previously. We previously reported that DOCK7 protein is asymmetrically distributed in unpolarized hippocampal neurons and becomes enriched in axons during the development of this process (35). However, the mechanism underlying the regulation of *DOCK7* in hippocampal neurons is unclear. It is interesting to speculate that such specific localization of *DOCK7* might be due to selective expression of miR-21.

Using a miRNA pulldown method, we were able to identify *DOCK* family genes as targets of miR-21. Over 29,000 genes are predicted as targets of miR-21 by at least one of 11 commonly used target prediction algorithms according to the miRecords web site. Thus, experimental methods to purify/identify targets of specific miRNAs are needed. *In vivo* study of targets of several miRNAs, including miR-1, miR-16, miR-30a, miR-155, and let-7b, using pulsed stable isotope labeling with amino acids in culture analyses in HeLa cells that overexpress these miRNAs indicates that roughly 300 proteins are down-regulated by at least 30% (32). The same study estimated that about 60% of the 300 down-regulated proteins contain seed-matched sites (Watson-Crick consecutive base pairing between mRNAs and the miRNA at positions 2–7 counted from the 5′-end of the miRNA) in the 3′-UTR regions of transcripts (32). Among the 33 previously validated miR-21 targets, only eight targets are predicted by a seed sequence-based algorithm. Non-seed-based algorithms tend to predict too many sites (on average ~3,000–11,000 MREs per miRNA), which defeats the purpose of using target prediction algorithms. Thus, the biochemical enrichment of miRNA targets presented in this study is useful. We identified ~26 genes that are ≥24-fold enriched in miR-21 pulldowns over control cel-miR-67 in PSMCs. We tested 28 predicted MREs located in the 3′-UTR of mRNAs associated with bio-miR-21, including 16 predicted MREs or sequences partially complementary to miR-21 localized in the 3′-UTR of *DOCK4*, -5, or -7 transcripts. Independently, miR-34a target analysis was performed in HCT116 and K562 cells using the bio-miR-34a pulldown method and confirmed that genes enriched in bio-miR-34a pulldowns by as little as 2.5-fold over

control miRNA pulldowns in both cells have 3′-UTRs that are regulated directly by miR-34a as measured by the luciferase reporter activity assay (40). Thus, additional miR-21-regulated genes are likely, and further verification and investigation of genes that are enriched in bio-miR-21 pulldowns in PSMC are warranted to understand the biological functions of miR-21 in vSMCs.

Six of the 22 miR-21-associated mRNAs that are enriched >24-fold over control miRNAs (*PARP9*, *DOCK7*, *ADAM22*, *RNF17*, *LGALS14*, and *SPRYD4*) do not contain miR-21 seed sequence or predicted MREs. RNAhybrid predicted some of the *DOCK* genes; however, this algorithm predicts on average ~11,000 target sites per miRNA. MRE sites in *DOCK4*, *DOCK5*, and *DOCK7* 3′-UTRs identified in this study contain complementary sequences of 6–11-mers starting somewhere between position 1 and 15. These results suggest that base pairing between a miRNA and target mRNA can be important downstream of the seed site.

It is intriguing that miR-21 targets multiple transcripts belonging to the same family. There are a few other examples of a single miRNA targeting multiple genes of the same family. miR-200 targets the zinc finger homeodomain enhancer-binding protein (ZEB) family of transcription repressors; ZEB1 and ZEB2 regulate the epithelial-to-mesenchymal transition (41). miR-164 targets the NAC domain transcription factors CUP-SHAPED COTYLEDON1 (*CUC1*) and *CUC2* that establish axillary meristems of leaves in plant (42). In both cases, products of the same gene family share a redundant function as in the case of *DOCK4*, -5, and -7 and Rac1 regulation in the regulation of cell migration (33). Although examples of a single miRNA targeting multiple genes in the same signaling pathway or with related biological functions have been reported previously, our results are striking because they suggest that a single miRNA targets multiple transcripts of the same gene family. In the case of miR-21 MREs identified in the 3′-UTR of human *DOCK7* (MRE1), an 8-mer at the 5′-end is 100% conserved in mouse. For human *DOCK4*, a 6-mer complementary sequence starting at position 3 in MRE1 is also 100% conserved in mouse, whereas in MRE6, only six of the 11 residues complementary to miR-21 starting at position 3 are conserved in the mouse 3′-UTR sequence. However, two MREs found in *DOCK5* are not conserved in mouse, suggesting that only some MREs are evolutionally conserved.

Increased proliferation and migration of vSMCs are associated with various stimuli, including activation of PDGF signaling, inhibition of BMP/TGFβ or Notch signaling, hypoxia treatment, mechanical stretch, and activation of Rac1 (43, 44). However, a role for *DOCK4*, -5, or -7 has not been characterized in vSMCs previously. Our study illustrates both common and distinct roles of *DOCK* family of proteins in the regulation of vSMC phenotype (Fig. 6). *DOCK4* is known to mediate cell migration promoted by the PDGF signal in NIH3T3 cells (45). In vSMCs, BMP and PDGF signals antagonize each other and modulate vSMC phenotype during vascular injury repair or pathogenesis of vascular proliferative diseases (46). Thus, the BMP4-miR-21-*DOCK4* axis is a novel mechanism of antagonism between BMP and PDGF signaling pathways in addition to the miR-24-Trb3 axis, which has been described previously

(34). Interestingly, cytoskeletal remodeling is mediated by DOCK4 and DOCK5, but not by DOCK7, through a mechanism that is independent of Rac1. As DOCK4 and -5 contain multiple Src homology 3 domains that are not found in DOCK7 (33), we speculate that the Src homology 3 domain might play a critical role in the cytoskeletal remodeling through docking other signaling molecule(s).

miR-21 is elevated in most human cancers and has been shown to function as an oncogene by promoting cell growth and migration (17). Indeed, many validated targets of miR-21 are tumor suppressors, including *PDCD4* (47), *PTEN* (48), and tropomyosin 1 (*TPM1*) (21). Homozygous deletion of *DOCK4* gene occurs during tumor progression in the *TP53-NF2* mouse model and is associated with increased invasion of tumor cells (49). Furthermore, a missense mutation (Pro-1718 to Leu) of *DOCK4* that results in defective activation of the Rap1 GTPase (49) has been identified in human prostate and ovarian cancers. Thus, in this context, *DOCK4* is a novel target of miR-21 with tumor suppressor function. It is intriguing, however, that miR-21, which promotes invasion and migration of cancer cells (50), inhibits migration in vSMCs. This observation is indicative of a cell type-specific function of miR-21. A potential mechanism of the cell type-specific function of miRNAs involves how different sets of genes are targeted by a miRNA due to (i) a cell type-specific expression of mRNA targets and/or (ii) a cell type-specific recognition/targeting of mRNA targets by a miRNA. A potential advantage of the miRNA target pull-down approach is that it allows identification of miRNA targets in a context-dependent manner. In addition to the relevance to vSMC biology, further investigation of the targets identified by the miR-21 pull-down may provide insights for understanding the sequence requirements for specific recognition of mRNA targets by miRNA and facilitate the development of more accurate target prediction algorithms.

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