Aberrant promoter methylation of multiple genes in VSMC proliferation induced by Hcy

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Abstract. Vascular smooth muscle cell (VSMC) proliferation is a primary pathological event in atherosclerosis (AS), and homocysteine (Hcy) is an independent risk factor for AS. However, the underlying mechanisms are still lagging. Studies have used the combination of methylation of promoters of multiple genes to diagnose tumors, thus the aim of the current study was to investigate the role of methylation status of several genes in VSMCs treated with Hcy. CpG islands were identified in the promoters of platelet-derived growth factor (PDGF), p53, phosphatase and tensin homologue on chromosome 10 (PTEN) and mitofusin 2 (MFN2). Hypomethylation was observed to occur in the promoter region of PDGF, hypermethylation in p53, PTEN and MFN2, and hypomethylation in two global methylation indicators, aluminium (Alu) and long interspersed nucleotide element-1 (Line-1). This was accompanied by an increase in the expression of PDGF, and reductions of p53, PTEN and MFN2, both in mRNA and protein levels. An elevation of S-adenosylmethionine (SAM) and a reduction of S-adenosylhomocysteine (SAH) and the SAM/SAH ratio were also identified. In conclusion, Hcy impacted methylation the of AS-associated genes and global methylation status that mediate the cell proliferation, which may be a character of VSMCs treated with Hcy. The data provided evidence for mechanisms of VSMCs proliferation in AS induced by Hcy and may provide a new perspective for AS induced by Hcy.

Introduction

Previous studies have demonstrated that elevated levels of homocysteine (Hcy) have been implicated as an independent risk factor for atherosclerosis (AS) (1-3). Studies have focused on the involvement of Hcy on the dysfunction and injury of vascular cells, such as vascular smooth muscle cells (VSMCs) (4,5). It has been previously observed that Hcy induced VSMCs proliferation, however the underlying mechanisms remain to be fully elucidated (6).

Hcy is a non-protein, sulfur containing amino acid, which is a metabolic intermediate of the methionine cycle. It is a precursor of S-adenosylmethionine (SAM), the unique methyl group donor for DNA methylation (7). Subsequent to transfer of the methyl group, SAM is transformed into S-adenosylhomocysteine (SAH), which can be hydrolyzed to form Hcy (8). DNA methylation refers to the addition of a methyl group to the 5 position of cytosine in the context of a CpG dinucleotide. Increasing evidence indicates that human diseases, including AS, are either caused or impacted by abnormal methylation (9). A previous study suggested that Hcy may also be involved in disturbing the expression of AS-associated genes through the interference of DNA epigenetic phenotype modification, such as DNA methylation (10). An additional previous study demonstrated that abnormal DNA methylation of genes including peroxisome proliferator-activated receptor α , apolipoprotein E (ApoE) and genomic DNA contribute to the development of AS induced by Hcy (11,12). Studies have identified DNA methylation profiling of the vascular tissues in the setting of AS or atherosclerotic plaques (13,14). However, the methylation status of several key genes that characterized the VSMC proliferation induced by Hcy in single VSMCs remains to be elucidated. In addition, it remains unclear whether Hcy affected the methylation status and caused proliferation.

In the present study, platelet-derived growth factor (PDGF), p53, phosphatase and tensin homologue on chromosome 10 (PTEN) and mitofusin 2 (MFN2), which are associated with cell proliferation regulation, and two global methylation indicators, aluminium (Alu) and long interspersed nucleotide element-1 (Line-1) were selected to analyze the methylation

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status in the VSMCs treat with Hcy, which may clarify the mechanisms of VSMCs proliferation in AS induced by Hcy and provide a potential diagnostic marker for AS induced by Hcy.

Materials and methods

Cell culture. Primary culture of VSMCs was obtained from the media of the umbilical vein of human as previously described (15). Experiments were approved by the Ningxia Medical University Medical Ethical Committee (Yinchuan, China). Cells were cultured in Dulbecco's modified Eagle's medium-Han's F12 media (DMEM-F12; Gibco; Thermo Fisher Scientific, Inc., Waltham, MA, USA) supplemented with 20% fetal bovine serum (FBS; Gibco; Thermo Fisher Scientific, Inc.), 100 U/ml penicillin and 100 U/ml streptomycin (Gibco; Thermo Fisher Scientific, Inc.) at 37°C in an incubator with 5% CO₂. When cells grew to 80% confluence, serum was media-deprived for 24 h to become synchronous, then 5% FBS (Gibco; Thermo Fisher Scientific, Inc.) for another 24 h before Hcy addition. Hcy was applied at the concentrations of 30, 50, 100, 200 and 500 μM folate (Sigma-Aldrich; Merck KGaA, Darmstadt, Germany), and 5 μM 5-aza-2'-deoxycytidine (AZC; Sigma-Aldrich; Merck KGaA), respectively.

Nested methylation-specific-polymerase chain reaction (nMS-PCR) for methylation analysis. The detection of methylation levels was conducted as previously described (16). The summary of primers and product sizes of the nMS-PCR assays are presented in Table I. The PCR products were separated by electrophoresis through a 2% agarose gel containing ethidium bromide. DNA bands were visualized by Gel Documentation and Analysis System ChemiDoc XRS (Bio-Rad Laboratories, Inc., Hercules, CA, USA) and calculated by the formula: Methylation % = methylation/(methylation + unmethylated) × 100%.

Reverse transcription-quantitative PCR (RT-qPCR) for measurement of mRNA expression. The total RNA was isolated using TRIzol reagent (Invitrogen; Thermo Fisher Scientific, Inc.). And the RNA (2 ml) was reverse transcribed using the Revert Aid First Strand cDNA synthesis kit (Fermentas; Thermo Fisher Scientific, Inc., Pittsburgh, PA, USA) for 60 min at 42°C and 70°C for 5 min. The primer nucleotide sequences of PDGF are as presented in Table II. The RT-qPCR was conducted with the FTC-3000 Real-Time PCR detection system (Funglyn Biotech, Inc., Toronto, ON, Canada). The qPCR reaction system as follows: 25 μl 2X SYBR mixture, 1 μl forward primer, 1 μl reverse primer, 2 μl cDNA and then RNase-free water up to 50 μl. The thermocycling conditions were as follows: Initial activation at 95°C for 5 min, followed by 30 cycles of 95°C for 20 sec, annealing temperatures for 20 sec (as stated in Table II) and at 72°C for 30 sec. The mRNA level of each gene was acquired from the value of the quantification cycle (Cq) of the qPCR as associated to that of glyceraldehyde-3-phosphate dehydrogenase (GAPDH) using the following equation (17): $\Delta Cq = Cq_{(GAPDH)} - Cq_{(sample)}$. Final results, expressed as N-fold differences in the target gene expression and relative to the calibrator termed 'N_{target}', were determined according to the following equation: $N_{target} = 2^{Cq_{(sample)} - Cq_{(calibrator)}}$, where Cq values

of the calibrator and sample were determined by subtracting the Cq value of the target gene from the Cq value. Each qPCR assay was performed in duplicate and included positive, negative and no template reagent controls.

Western blotting assay for measurement of proteins expression. Whole-cell proteins were extracted with cell lysis buffer (KeyGene, Shanghai, China) that included the protease inhibitor phenylmethanesulfonyl fluoride (KeyGene) at 4°C for 30 min, then were separated by 12% sulfate-polyacrylamide gel electrophoresis. The proteins and the pre-stained marker (Fermentas; Thermo Fisher Scientific, Inc.) were then transferred onto an Immobilon-P transfer polyvinylidene fluoride membrane (EMD Millipore, Billerica, MA, USA) with a Semi-dry Transfer Cell (model 755; Bio-Rad Laboratories, Inc.) for 90 min, allowing the pre-stained marker to be completely transferred from the gel to the membrane. The gel was then discarded and the membrane was incubated at room temperature in 5% non-fat milk prepared with phosphate-buffered saline with 0.05% v/v Tween-20 (PBS-T) buffer. A total of 2 h later, the membrane was cut as required, placed in a suitable hybridization bag with 1 ml of primary antibody [all primary antibodies were diluted to 1:1,000 with 1% non-fat milk; anti-PDGF (cat. no. ab23914), anti-p53 (cat. no. ab26), anti-PTEN (cat. no. ab32199), anti-MFN2 (cat. no. ab56889) and anti-β-actin (cat. no. ab8227) antibodies were obtained from Abcam, Cambridge, UK] and were incubated at 4°C. The membrane was washed three times with PBS-T and the mouse anti-goat horseradish peroxidase-conjugated secondary antibody (cat. no. sc-2354; 1:2,000; Santa Cruz Biotechnology, Inc., Dallas, TX, USA) was added for 2 h at room temperature. Blots were developed on X-ray film (Kodak, Shanghai, China). The protein bands were visualized and analyzed using the Gel Documentation and Analysis System ChemiDoc XRS system with Image Lab software (version 4.1; Bio-Rad Laboratories, Inc.), and calculated by the gray value of the bands.

SAM and SAH concentrations examined by HPLC. The concentrations of SAM and SAH were determined by high performance liquid chromatography (HPLC). Samples were loaded into a C18 column (Shimadzu, Tokyo, Japan), run by a Hitachi L2000 HPLC system (HPLC; D-2000 Elite HPLC; Hitachi High Technologies, Tokyo, Japan). Absorption of eluted compounds was monitored at (λ) ex=254 nm. Elution of SAM and SAH was achieved at a flow rate of 1.0 ml/min with the mobile phase ammonium format solution. Chromatograms were recorded using a D-2000 Elite integrator. SAM and SAH standards were used to identify the elution peaks, and the SAM and SAH values of the tissues were calculated with the standard curve.

Statistical analysis. Each experiment was repeated 3 times. Results were expressed as the mean ± standard deviation. One-way analysis of variance was used to compare the means of multiple groups, followed by Newman-Keuls test. P≤0.05 was considered to indicate a statistically significant difference.

Results

The bioinformatics analysis of promoters in PDGF, p53, PTEN and MFN2. As a key epigenetic modification of the genome,

Table I. Summary of nested methylation-specific-polymerase chain reaction primers.

Gene	Forward primer sequence (5'→3')	Reverse primer sequence (5'→3')	Size (bp)	Annealing temperature (°C)
PDGF-O	AAGGTTGTTTTATTTATTTTTTGT	AACTACAACTACAACACTCTCCAAT	305	52
PDGF-M	GTTTGTGTTGTTTTTTTTCGTATTC	CTACTCCGATTTTCTCTTTACAACG	193	59
PDGF-U	GGTTTGTGTTGTTTTTTTGTGTATTT	ACTCCAATTTTCTCTTTACAACAAA	192	57
p53-O	GTTTTGGTTTGAAGGATAGTAGTT	AAAAACCCCTAAAACTTAATAAAAAAC	404	55
p53-M	TTAGTTTTAGTTAGGATGGTTTCGA	GAAAAATAAACCCGAAATCCCG	212	56
p53-U	ATTAGTTTTAGTTAGGATGGTTTTGA	CAAAAAATAAACCAAAATCCCAC	214	55
PTEN-O	TTGGAAAGTTTTTTAATTAGGGATA	ATTTCAAAAACCCAAAAAACAC	445	55
PTEN-M	GTGATTTTTTTCGGAAAGTAGTTTC	TAAAAACCCGACAAAATAAATCG	211	57
PTEN-U	ATTTTTTTTGAAAGTAGTTTTGA	AAAAACCCAACAAAATAAATCACC	207	56
MFN2-O	ATAGAATGTAAATTTGGATTTTAGA	ACTAATAAACCCCTAAACCCAACC	355	53
MFN2-M	TTTGTTCGTTTTTTTAGTTTCG	CTAAACCCAACCGACTCG	210	55
MFN2-U	TTTGTGTTGTTTTTTTAGTTTTGG	TAAACCCAACCAACTCACC	209	55
Alu-O	TATTTTGGTTAATAAGGTGAAATTT	TCCAACAACATAAAAAAACTTTTT	256	55
Alu-M	GGGCGTTTGTAGTTTTAGTTATTC	TAAAACGAAATCTCGCTCTATCG	143	56
Alu-U	GGTGTGTTGTAGTTTAGTTATTTGG	TTAAACAAAATCTCACTCTATCACC	143	56
Line-1-O	TTTATTAGGGAGTGTTAGATAGTGGG	TACCCAAACAAACCTAAACAATAAC	359	57
Line-1-M	TATTAGGGAGTGTTAGATAGTGGGC	AACCCGATTTTCCAAATACGT	162	59
Line-1-U	TTAGGGAGTGTTAGATAGTGGGTGT	AAAAAACCCAATTTTCCAAATACA	164	59

bp, base pairs; PDGF, platelet-derived growth factor; O, outer primers; M, methylated-specific primers; U, unmethylated-specific primers; PTEN, phosphatase and tensin homologue on chromosome 10; MFN2, mitofusin 2; Alu, aluminium; Line-1, long interspersed nucleotide element 1.

Table II. Summary of reverse transcription-quantitative polymerase chain reaction primers.

Gene	Forward primer sequence (5'→3')	Reverse primer sequence (5'→3')	Size (bp)	Annealing temperature (°C)
PDGF	CCACTCGATCCGCTCCTTTGA	GAACCCAGGCTCCTTCTTCCAC	150	60
p53	CGTGTGTTGTCCTGTCTTG	TGCTCGCTTAGTGCTCCCT	105	58
PTEN	AAGACCATAACCCACCACAGC	ACCAGTTCGTCCCTTTCCAG	125	57
MFN2	TACTGCTGCTCAACTGC	AACCAACCGGCTTTATTC	188	55

PDGF, platelet-derived growth factor; PTEN, phosphatase and tensin homologue on chromosome 10; MFN2, mitofusin 2; bp, base pairs.

DNA methylation occurs almost exclusively in the context of CpG dinucleotides, particularly in the CpG islands (18). Ranging from 0.5-5 kb and occurring on average every 100 kb, CpG islands are GC rich (50-70%) and have a ratio of CpG:GpC of at least 0.6 (19). Collectively, CpG islands account for 1-2% of the genome and their location is primarily in the 5' regulatory regions of all housekeeping genes as well as up to 40% of tissue-specific genes (20). CpG methylation is associated with gene silencing, and to identify whether CpG islands exist in the chosen genes PDGF, p53, MFN2 and PTEN, the 5'-flanking region of the genes was analyzed using an online search engine (www.urogene.org/cgi-bin/methprimer/methprimer.cgi). The criteria of the CpG island is a CpG-rich region length more than 100 bp, GC percentage >50%, and observe/expect ratio >0.60. As presented in Fig. 1, at least one CpG island

was identified in the DNA promoter regions of all the 4 genes. In the PTEN gene promoter, 6 CpG islands were observed, which indicated that they had greater potential to be modified by DNA methylation and regulated their expression.

PDGF, P53, PTEN, MFN2, Alu and line-1 methylation levels in VSMCs. To ascertain whether DNA methylation occurred in these genes, the methylation status of genes was analyzed by nMS-PCR. For PDGF, as presented in Fig. 2A, the methylation status of PDGF is increased, particularly, in the 100 μ M Hcy group, in which it was increased 7.6-fold, compared with the untreated group ($P<0.01$). For p53, as presented in Fig. 2B, p53 methylation levels were increased in a dose-dependent manner with the Hcy concentration. For PTEN and MFN2, the methylation levels were elevated by treatment with Hcy

Figure 1. The bioinformatics analysis of promoters in PDGF, P53, PTEN and MFN2. The properties of CpG islands in the proximal region of the PDGF promoter were examined using the CpG Island Search engine (www.urogene.org/cgi-bin/methprimer/methprimer.cgi). At least one CpG island exists in the promoter region in all of the four genes. The light blue regions and the blue lines represent the CpG islands. PDGF, platelet-derived growth factor; PTEN, phosphatase and tensin homologue on chromosome 10; MFN2, mitofusin 2.

(Fig. 2C and D), however, it was not dose-dependent. Notably, the most marked effect was observed in the 100 μ M group. When treated with folate, the methylation levels of PDGF, PTEN and MFN2 were decreased compared with that of the 100 μ M group ($P<0.05$).

It was known that Hcy impacted the gene-specific methylation, however it remained unclear whether it affected the global methylation status, Alu and LINE-1 methylation status is considered as a good indicator of global methylation (21), thus these methylation levels were detected in VSMCs treated by Hcy. The results indicated that both Alu and LINE-1 were in a hypo-methylated state (Fig. 2E and F; $P<0.01$). No significant difference was observed in 100 μ M group, however the

results in the 50 μ M group were significant. In addition, the folate group exhibited an antagonistic effect ($P<0.05$).

AZC, an inhibitor of DNA methylation, was added, and co-incubated VSMCs were treated with different concentrations of Hcy, then the methylation levels of Alu and Line-1 were measured with nMS-PCR. When compared with untreated cells, the methylation levels of Alu and Line-1 were suppressed in the Hcy treated groups. These data indicated that Hcy impacted the methylation status of these genes and that this may contribute to the proliferation of VSMCs induced by Hcy.

PDGF, P53, PTEN and MFN2 expression levels in Hcy-treated VSMCs. It was identified that Hcy affected the methylation

Figure 2. The methylation levels of PDGF, P53, PTEN, MFN2, Alu and Line-1. The methylation status was detected in VSMCs treated with different concentrations of Hcy by nMS-PCR. (A) The methylation of PDGF was detected by nMS-PCR in VSMCs, following co-incubation with different concentrations of Hcy for 72 h; (B) The methylation of P53 was detected by nMS-PCR in VSMCs, following co-incubation with different concentrations of Hcy for 72 h; (C) The methylation of PTEN was detected by nMS-PCR in VSMCs, following co-incubation with different concentrations of Hcy for 72 h; (D) The methylation of MFN2 was detected by nMS-PCR in VSMCs, following co-incubation with different concentrations of Hcy for 72 h; (E) The methylation of Alu was detected by nMS-PCR in VSMCs, following co-incubation with different concentrations of Hcy for 72 h; (F) The methylation of Line-1 was detected by nMS-PCR in VSMCs, following co-incubation with different concentrations of Hcy for 72 h. M, methylation; U, unmethylated; Folate group, 30 μ M folate in 100 μ M Hcy; AZC group, 5 μ M AZC in 100 μ M Hcy. Data are presented as the mean $\pm$ standard deviation from 3 independent experiments performed in triplicate. * P <0.05, ** P <0.01, vs. the control group; # P <0.05, ## P <0.01, vs. the 100 μ M Hcy group. PDGF, platelet-derived growth factor; PTEN, phosphatase and tensin homologue on chromosome 10; MFN2, mitofusin 2; Alu, aluminium; Line-1, long interspersed nucleotide element-1; nMS-PCR, nested methylation-specific-polymerase chain reaction; VSMCs, vascular smooth muscle cells; Hcy, homocysteine; AZC, 5-aza-2'-deoxycytidine.

levels of genes, however whether this further impacted the expression remained unclear. In order to investigate whether this occurred in VSMCs treated by Hcy, the mRNA and protein levels of the genes were assayed by RT-qPCR and western blotting. As presented in Fig. 3A, PDGF mRNA and protein expression levels were increased in a dose-dependent manner. p53, PTEN and MFN2 were decreased in the 50,

100, 200 and 500 μ M Hcy groups (Fig. 3B-D). The expression levels of p53, PTEN and MFN2 consistently exhibited the greatest effects in the 100 μ M group. In addition, more folate suppressed the changes induced by Hcy. When treated with AZC, the antagonist of DNA methylation, the expression of PDGF, p53, PTEN and MFN2 increased significantly (P <0.05). These data suggested that Hcy affected the

Figure 3. The expression of PDGF, P53, PTEN and MFN2 in Hcy-treated VSMCs. VSMCs were exposed to 0, 50, 100, 200 and 500 μ M of Hcy for 72 h, mRNA and protein were extracted from VSMCs and detected by reverse transcription-quantitative polymerase chain reaction and western blotting, respectively. Expression levels of (A) PDGF, (B) p53, (C) PTEN and (D) MFN2 in VSMCs co-incubated with Hcy for 72 h are presented. Folate group, 30 μ M folate in 100 μ M Hcy; AZC group, 5 μ M AZC in 100 μ M Hcy. Data are presented as the mean $\pm$ standard deviation from 3 independent experiments performed in triplicates. * P <0.05 and ** P <0.01, vs. the control group; # P <0.05 and ## P <0.01, vs. the 100 μ M Hcy group. PDGF, platelet-derived growth factor; PTEN, phosphatase and tensin homologue on chromosome 10; MFN2, mitofusin 2; Hcy, homocysteine; VSMCs, vascular smooth muscle cells.

expression of the genes, which may be involved in the DNA methylation process.

SAM and SAH levels in VSMCs treated with Hcy. Similar to Hcy, SAM and SAH are intermediates in the methionine metabolism, and are important factors in the transmethylation process (22). The SAM and SAH levels were measured using HPLC in the VSMCs, which were treated with Hcy. As presented in Fig. 4, the results demonstrated that the intracellular levels of SAH were significantly increased, compared with the control group, while the concentrations of SAM and the ratio of SAM/SAH were decreased, when Hcy concentrations increased (P <0.05; Fig. 4). SAH levels increased and SAM levels decreased (P <0.05), when VSMCs were treated with Hcy.

Discussion

VSMCs are major components of the arterial wall, and abnormal proliferation of VSMCs serves a pivotal role in the pathogenesis of AS (23). As an important independent risk factor of AS, Hcy stimulated the VSMCs proliferation, which was identified in previous studies (6,24). Cellular proliferation is associated with the regulation of key genes. Thus in the present study, 4 genes were selected for analysis; PDGF, p53, PTEN and MFN2. PDGF is a potent mitogen, which serves crucial roles in developmental and physiological processes,

in addition to being directly implicated in proliferative disorders, and it is a potent stimulator of VSMC growth (25). p53 is a key tumor suppressor and a key regulator of various signaling pathways, including cell-cycle regulation, induction of apoptosis, development and differentiation (26). *In vivo*, p53 expression is negatively correlated with markers of cell proliferation in human atherosclerosis; adenoviral expression of p53 reduces cellular proliferation in the rat carotid artery (27), and by contrast, siRNA of p53 increases cellular proliferation (28). In p53^{-/-}/ApoE^{-/-} mice, p53 have been demonstrated to increase aortic plaque formation, with increased rates of cell proliferation and reduced rates of apoptosis in brachiocephalic artery plaques (29). PTEN is well known tumor suppressor and is additionally involved in regulation of a variety of physiological and pathological processes, including cell proliferation, differentiation, apoptosis, adhesion and migration (30). PTEN was expressed endogenously in VSMCs, and overexpression of PTEN significantly inhibited basal and PDGF-mediated VSMC proliferation and migration (31). PTEN overexpression in VSMCs using adenoviral transfection resulted in inhibition of cell proliferation and migration induced by angiotensin II (32). MFN2 was first cloned from VSMCs in spontaneous hypertensive rats using the differential display technique (33), which restrained the proliferation of VSMCs mediated by inhibition of extracellular signal-related kinase/mitogen-activated protein kinase signaling and subsequent cell-cycle arrest. The expression changes of these key

Figure 4. The SAM and SAH levels in VSMCs treated by Hcy. (A) A chromatogram of SAM and SAH of all groups detected by high-performance liquid chromatography, chromatograms were recorded by an integrator. a, Untreated group; b, 50 μ M Hcy group; c, 100 μ M Hcy group; d, 200 μ M Hcy group; e, 500 μ M Hcy group; and f, folate group. (B) The concentrations of SAM and SAH in VSMCs. (C) The ratio of SAM and SAH in VSMCs. Folate group, 30 μ M folate in 100 μ M Hcy. The data were obtained using automatic peak area integration. Data are presented mean $\pm$ standard deviation from 3 independent experiments performed in triplicate. * $P < 0.05$ and ** $P < 0.01$, vs. the control group; # $P < 0.05$ vs. the 100 μ M Hcy group. SAM, S-adenosylmethionine; SAH, S-adenosylhomocysteine; VSMCs, vascular smooth muscle cells; Hcy, homocysteine.

genes may involve in the imbalance of cell proliferation. The present study indicated an elevation of PDGF, and a reduction of p53, PTEN and MFN2 in VSMCs treated with Hcy, which indicated that Hcy impacted the expression of these genes and resulted in the proliferation of VSMCs.

DNA methylation is an important epigenetic modification at the transcriptional level, which commonly occurs at the CpG dinucleotides of gene promoter regions, particularly in the promoter-associated CpG islands (34). Consistent with these important roles, a growing number of human diseases have been identified to be associated with aberrant DNA methylation. In the present study, CpG islands were searched for in the promoters of the selected genes, PDGF, p53, PTEN and MFN2, using two CpG island search engines. It was identified that at least one CpG island exists in the promoter regions of all 4 genes. In the promoter region of PTEN, 6 CpG islands were identified, which indicated that they have potential to be modified by DNA methylation and regulate their expression.

It was previously reported that Hcy caused cell-type-specific hypomethylation in endothelial cells, VSMCs and foam cells (35) and it was hypothesized that Hcy metabolism and methylation may be associated, resulting in tissue-specific pathology in AS. In addition,

DNA methylation also impacts the selected genes PDGF, p53, PTEN and MFN2. In the present study, the key genes associated with cell proliferation regulation were selected in order to investigate the methylation changes in the single VSMCs. In the VSMCs treated with Hcy, hypermethylation or hypomethylation was observed to occur in the promoters of the selected genes, of which PDGF was hypomethylated and p53, PTEN, MFN2 were hypermethylated. Furthermore, the expression levels were altered from normal to aberrant levels. When treated with AZC, an antagonist of DNA methylation, both the methylation status and the expression of the four genes was altered. These results demonstrate that Hcy affected the expression of the genes, which may be involved in the DNA methylation process.

In order to try to confirm the methylation status characteristics in the VSMCs affected by Hcy, we detected global methylation levels were measured. In general, Alu and LINE-1 elements are methylated in somatic tissues, however are hypomethylated in human cancer (36). In the present study, Hcy was observed to reduce the methylation levels of Alu and LINE-1. In the VSMCs, Hcy not only affected the methylation levels of particular genes, however additionally impacted the global methylation status. Hcy-associated methylation changes

are also suggested as a biomarker for use in the prevention and therapy of AS in the future.

In the methionine cycle, SAM is the key methyl donor for cellular methylation events including DNA methylation, and this process produces SAH. SAH is converted to Hcy in a reversible reaction, and elevated levels of SAH and concomitant increases in the SAH/SAM ratio are suggested to inhibit cellular methylation reactions (37). The present study observed an increase of SAM and a decrease in the SAM/SAH ratio, which demonstrated that Hcy amassment accumulated in the methionine cycle, and caused the SAM, SAH and SAM/SAH ratio changes, then affected methylation process. Furthermore, in the cycle, folate is integrally involved in both substrate synthesis and product removal via its role in methionine synthesis from Hcy. Folate insufficiency leads to a decrease in SAM synthesis (38), which may compromise SAM-dependent methylation reactions, however additionally also leads to an increase in cellular concentrations of SAH by promoting Hcy accumulation via reversal of SAH hydrolase. In the present study, folate was added to reduce the Hcy effect. The results indicated an antagonistic effect against Hcy both in the methylation levels and in the gene expression levels. These data indicated that Hcy may impact the methylation status partly through the methionine cycle.

In conclusion, Hcy impacted the methylation status of the genes involved in the cell proliferation, leading to a loss of cellular control in VSMCs, and induced the proliferation of VSMCs. Hcy not only affected the methylation levels of special genes, however additionally impacted the global methylation status, which may be a characteristic of VSMCs treated with Hcy. The data provided evidence for the mechanisms of VSMC proliferation in AS induced by Hcy and may provide a potential diagnostic marker for AS induced by Hcy.

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