Nuclear factor-κB signaling negatively regulates high glucose-induced vascular endothelial cell damage downstream of the extracellular signal-regulated kinase/c-Jun N-terminal kinase pathway

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Abstract. Diabetes mellitus (DM)-induced high blood sugar severely damages vascular endothelial cells (VECs), which are in direct contact with the blood. Diabetic complications cause difficulties in skin wound healing and VECs are important for this process. Previous studies demonstrated that high blood sugar delayed the repair of wounded VECs, but the underlying mechanism has remained elusive. To explore the effects of diabetic conditions on VEC damage, cells were incubated in a medium with high glucose and then subjected to RNA-sequencing based transcriptome analysis. The results revealed that numerous biological processes were altered by HG stress, including extracellular matrix-receptor interaction, NOD-like receptor signaling and the nuclear factor (NF)-κB pathway. HG treatment increased the levels of phosphorylated inhibitor of NF-κB (IκB-α), the key NF-κB signaling regulator as well as the transcripts of plasminogen activator inhibitor-1 and interleukin-8, two inflammatory response markers. Treatment with extracellular signal-regulated kinase (ERK)- and c-Jun N-terminal kinase (JNK)-specific inhibitors U0126 and sp600125, respectively, led to the activation of IκB-α; however, the inhibitor of IκBα phosphorylation Bay11-7082 did not affect ERK and JNK activity, suggesting that ERK/JNK signaling occurs upstream of NF-κB in VECs. The present study provided useful information regarding the effects of diabetes on VECs, which may provide approaches for therapies of diabetes-associated complications in the future.

Introduction

With exponential increases in the number of individuals affected by diabetes mellitus (DM), this disease represents a great medical and socioeconomic challenge worldwide. Despite various treatment strategies adopted, vascular complications represent a major problem causing morbidity and mortality of diabetic patients. Early in the course of diabetes, intracellular hyperglycemia causes endothelial dysfunction and endocrine disorders (1-3). The overall result is tissue dysfunction, which results in the formation of non-healing ulcers.

Skin wound repair involves a series of coordinated processes such as cell proliferation and migration, collagen deposition and remodeling, wound contraction and angiogenesis. These mechanisms involve various cell types, mostly including fibroblasts/myofibroblasts, keratinocytes and endothelial cells (4,5). Due to their incapacity to regulate glucose influx, vascular endothelial cells (VEC) represent an important target for DM-induced damage. The hyperglycemia in diabetic patients has resulted in difficulties in wound healing, particularly altered angiogenesis and extracellular matrix remodeling (6), while the nature of the linkage has remained elusive. Extensive studies have described alterations in cell migration associated with diabetic conditions. For instance, Lerman et al (7) demonstrated that fibroblasts from diabetic mice migrated 75% less than those from normoglycemic mice and displayed a constant response to hypoxia, a condition commonly present in chronic wounds. Recently, an inhibition phenomenon similar to that in mice was observed in keratinocytes cultured in a high-glucose (HG) environment (8), which suggested that HG has a significant role in delaying wound repair. Mitogen-associated protein kinases (MAPKs) were identified to be involved in reactive oxygen species (ROS) production upon HG stress (9). However, these studies failed to address the detailed cellular regulatory mechanisms involved in this phenomenon.
The present study analyzed the effects of HG on transcriptome changes in VECs via RNA-sequencing (seq) analysis. In addition, the association between NF-κB and two MAPK members, extracellular signal-regulated kinase (ERK) and c-Jun N-terminal kinase (JNK), were analyzed using pharmacological inhibitors. RNAs differentially expressed by VEGs under HG conditions and their associated biological processes were identified, and the association between NF-κB and MAPK pathways under diabetic conditions was analyzed.

Materials and methods

VEC culture. The EA.hy926 human endothelial cell line was obtained from the Cell Bank of the Chinese Academy of Sciences (Shanghai, China). These cells were immediately expanded and frozen so that the cell line could be restarted every 3 months. Cells were cultured in low-glucose (LG) Dulbecco's modified Eagle's medium (GE Healthcare Life Sciences, Little Chalfont, UK) supplemented with 10% fetal bovine serum (GE Healthcare Life Sciences), 100 µg/ml streptomycin and 100 U/ml penicillin. The cultures were maintained at 37°C in a humidified chamber with 5% CO₂ and 95% air, and the medium was replaced every 2 days. The cultured cells were digested and passed with 0.25% trypsin (Gibco; Thermo Fisher Scientific, Inc., Waltham, MA, USA) after the cells reached ~90% confluency. The cells were then seeded at 2.5x10⁴ cells/well on 6-well plates or at 1x10⁴ cells/well on 48-well plates and cultured for 24 h. The EA.hy926 cells in the LG or HG group were treated with 5.5 or 35 mM glucose, respectively, for 3 days. The cells were cultured in 6 well plates, and treated with 1 µM Bay117082 (cat. no. 19542-67-7; Sigma-Aldrich; Merck KGaA, Darmstadt; Germany), U0126 (cat. no, 109511-58-2; Sigma-Aldrich; Merck KGaA), sp600125 (cat. no. 129-56-6; Sigma-Aldrich; Merck KGaA) and 1 µM RNA polymerase chain reaction (PCR).

Total RNA was extracted from VECs from the LG and HG groups. The cell monolayer was rinsed with ice-cold PBS. The cells then lysed by adding 1 ml TRIzol reagent (Invitrogen; Thermo Fisher Scientific, Inc.) per each 3.5-cm diameter 6-well dish, and 0.2 ml chloroform was added. Total RNA (1 µg) was reverse-transcribed by using a GoScript Reverse Transcription kit (A5003; Promega Corp., Madison, WI, USA) following the manufacturer's instructions. The kit included Buffer, dNTP, Oligo (dT), RNase inhibitor and reverse transcriptase. The reverse transcription reaction was subjected to the following conditions: 65°C for 5 min, 42°C for 15 min, 75°C for 15 min. The SYBR-Green kit (Bio-Rad Laboratories, Inc., Hercules, CA, USA) was used for quantitative PCR assay according to the manufacturer's instructions (5 µl of 2X SYBR-Green buffer, 1 µl of each 5 µM primer and 3 µl of H₂O to total volume of 10 µl for each PCR reaction). qPCR was performed with the following thermocycling conditions: 95°C for 3 min; 40 cycles for 95°C for 30 sec, 58°C for 30 sec, 72°C for 30 sec; and 72°C for 10 min; followed by a holding temperature of 8°C. The transcript levels were normalized against those of GAPDH using the 2^(-ΔΔCt) method (11). The sequences of primers used for PCR are listed in Table I.

Analysis of the pathway. Pathway analysis was performed to determine the significant pathways enriched by differentially expressed genes under HG according to the Kyoto Encyclopedia of Genes and Genomes (KEGG; http://www.genome.jp/kegg/), Biocarta (https://cgap.nci.nih.gov/Pathways/BioCarta_Paths) and Reactome (http://www.reactome.org/) databases. Fisher's exact test was performed followed by Benjamini-Hochberg multiple testing correction for isolating significantly changed pathways, and the threshold of significance was defined by the P-value (P<0.05) and false-discovery rate (10).

Total RNA extraction, complementary DNA synthesis and polymerase chain reaction (PCR). Total RNA was extracted from the VECs of the HG and LG groups. The cell monolayer was rinsed with ice-cold PBS. The cells then lysed by adding 1 ml TRIzol reagent (Invitrogen; Thermo Fisher Scientific, Inc.) per each 3.5-cm diameter 6-well dish, and 0.2 ml chloroform was added. Total RNA (1 µg) was reverse-transcribed by using a GoScript Reverse Transcription kit (A5003; Promega Corp., Madison, WI, USA) following the manufacturer's instructions. The kit included Buffer, dNTP, Oligo (dT), RNase inhibitor and reverse transcriptase. The reverse transcription reaction was subjected to the following conditions: 65°C for 5 min, 42°C for 15 min, 75°C for 15 min. The SYBR-Green kit (Bio-Rad Laboratories, Inc., Hercules, CA, USA) was used for quantitative PCR assay according to the manufacturer's instructions (5 µl of 2X SYBR-Green buffer, 1 µl of each 5 µM primer and 3 µl of H₂O to total volume of 10 µl for each PCR reaction). qPCR was performed with the following thermocycling conditions: 95°C for 3 min; 40 cycles for 95°C for 30 sec, 58°C for 30 sec, 72°C for 30 sec; and 72°C for 10 min; followed by a holding temperature of 8°C. The transcript levels were normalized against those of GAPDH using the 2^(-ΔΔCt) method (11). The sequences of primers used for PCR are listed in Table I.

Western blot analysis. For extraction of total protein, the cells were lysed in an ice-cold lysis solution (7 M urea, 2 M thiourea, 2% CHAPS detergent, 40 mM Trizma base, 40 mM dithiothreitol, 1% protease inhibitor) and the lysates were centrifuged for 15 min at 15,000 x g at 4°C. All reagents were sourced from Sigma-Aldrich. The protein concentration was measured via Nanodrop 100 (Thermo Fisher Scientific Inc.). A total of 20 µg protein from each sample was separated by 10% SDS-PAGE and then electrotransferred onto Immobilon-P Transfer Membranes (Millipore, Tokyo, Japan). The membranes were incubated in 1X Tris-buffered saline containing 5% skimmed milk and 0.05% Tween-20 for 1 h at room temperature (25°C) and blotted with primary antibodies at overnight. For the primary antibodies, anti-phosphorylated inhibitor of NF-κB (p-IκB-α) antibody (1:2,000 dilution; cat. no. 39A1431; Abcam, Cambridge, MA, USA), anti-IκB-α antibody (1:2,000 dilution; cat. no. ab7217; Abcam), anti-p-p38 mitogen-activated protein kinase (SAPK) JNK (Thr183/Tyr185) antibody (1:1,000 dilution; cat. no. 4668; Cell Signaling Technology, Inc., Danvers, MA, US), anti-SAPK/JNK (1:1,000; cat. no. 9252; Cell Signaling Technology, Inc.), anti-p-p44/42 MAPK (ERK1/2

### Table I. Primer sequences used for polymerase chain reaction.

<table>
<thead>
<tr>
<th>Primer</th>
<th>Sequence (5’-3’)</th>
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<tbody>
<tr>
<td>PAI-1 F</td>
<td>GAGACTGAAGTCGACCTCAG</td>
</tr>
<tr>
<td>PAI-1 R</td>
<td>CTTGCCTATGATGCTCCTC</td>
</tr>
<tr>
<td>GAPDH F</td>
<td>GACCTGCGCTCTAGAAGAAC</td>
</tr>
<tr>
<td>GAPDH R</td>
<td>CTGTAGCAAAATCTGTTGC</td>
</tr>
<tr>
<td>IL-8 F</td>
<td>GGTGACGTTGTGCAAGAG</td>
</tr>
<tr>
<td>IL-8 R</td>
<td>TTCCTTGGCTCCACAGACA</td>
</tr>
<tr>
<td>Caspase-3 F</td>
<td>TGTAGTACATGCGCTGTC</td>
</tr>
<tr>
<td>Caspase-3 R</td>
<td>GTGTGCCACCTTCTCGATTAC</td>
</tr>
<tr>
<td>CCL13 F</td>
<td>CGTCCCATCTACTTGCTG</td>
</tr>
<tr>
<td>CCL13 R</td>
<td>TCAAGTCTCAGGTTGTCG</td>
</tr>
</tbody>
</table>

F, forward; R, reverse; IL, interleukin; CCL, chemokine ligand 13; PAI-1, plasminogen activator inhibitor-1.
(Thr202/Tyr204) (1:1,000 dilution; cat. no. 9101; Cell Signaling Technology, Inc.), anti-p44/42 MAPK (ERK1/2) (1:1,000 dilution; cat. no. 9102; Cell Signaling Technology, Inc.) and anti-GAPDH antibody (1:2,000 dilution; cat. no. ab8245; Abcam) were used. The membranes were incubated for 1 h at room temperature (25˚C) with an anti‑mouse or anti‑rabbit horseradish peroxidase‑linked secondary antibody (1:2,000 dilution; cat. no. 7074; Cell Signaling Technology, Inc.). Reaction products were visualized by detection of chemiluminescence using the ECL Western Blotting Detection System (GE Healthcare Life Sciences). The images were captured on the X‑ray films (Kodak; Rochester, NY, USA).

Statistical analysis. Statistical calculations were performed by using GraphPad Prism 5 software (GraphPad, Inc., La Jolla, CA, USA). All values are expressed as the mean ± standard error. Significant differences between groups were analyzed by using the Student's t-test.

Results

Identification of HG-associated transcriptome in fibroblasts. To identify genes and pathways upregulated under HG, RNA-seq experiments were performed by using human VECs incubated under LG (5.5 mM) and HG (35 mM) conditions for 3 h. The RNA-seq results revealed that 721 genes were differentially expressed (>1.5 fold-change; P<0.05) in the HG-treated VECs compared with those incubated under LG conditions (Fig. 1A). To verify the RNA-seq data, the HG-induced expression of inflammatory and apoptotic marker genes (IL-8, PAI-1, CCL13 and Caspase-3) were further monitored by reverse-transcription quantitative PCR. Three inflammation-associated genes (IL-8, PAI-1 and CCL13) and an apoptosis-associated gene (Caspase-3) were induced by HG stimulation (Fig. 1B), which was similar to the RNA-seq results. Furthermore, the 721 genes altered by HG stress were subjected to pathway analysis. The results demonstrated that these genes were enriched various canonical pathways, the top 15 of which are listed in Fig. 2 according to their rank-ings, including alcoholism as well as NF-κB and extracellular matrix (ECM)-receptor interaction signaling pathways. NOD-like receptor signaling was identified in VEC cell with HG stress, which was previously identified as a target pathway in HG-stressed fibroblast cells (12). These data indicated that HG regulates a large number of genes involved in different biological processes.

Figure 1. HG altered the transcriptome profile in VECs. (A) Heat map diagram of the genes whose expression levels in VECs were altered after HG (35 mM) treatment for 3 h. Gene expression is indicated by a pseudocolor scale with red denoting higher and green denoting lower gene expression levels. (B) Reverse-transcription quantitative polymerase chain reaction analysis was performed to verify the expression levels of IL-8, PAI-1, CCL13 and Caspase-3. GAPDH was used as an internal control. *P<0.05 vs. Low Glc. Low Glc/Con, low glucose (5.5 mM); High Glc/HG, high glucose (35 mM); VECs, vascular endothelial cells; IL, interleukin; CCL, chemokine ligand 13; PAI-1, plasminogen activator inhibitor-1.

Association between NF-κB and ERK/JNK signaling in VECs. ERK and JNK are well-known MAPKs, which are involved in diverse aspect of biology, including HG-mediated damage of VECs (9). In fibroblasts, NF-xB signaling was identified to function upstream of JNK (13). To address the association between ERK/JNK and NF-xB signaling, pharmacological and biochemical tests were utilized. As a key regulator, IκBα activity was analyzed after HG stress by analyzing the levels p-IκBα and total IκBα. Western blot analysis indicated HG-induced p-IκBα levels, particularly after 1 h of HG treatment, while total IκBα levels were not affected (Fig. 3A). In addition, the VECs were treated with the inhibitor of IκBα phosphorylation Bay117082 and the p-ERK, ERK, p-JNK and JNK levels were monitored by western blot analysis. The results indicated that Bay117082 did not change ERK and JNK activities (Fig. 3B). In a different experiment, U0126 and sp600125, respective inhibitors of ERK and JNK, were applied and the
levels of p-IκBα and total IκBα were examined. U0126 and sp600125 treatment induced changes in p-IκBα, while IκBα levels were maintained (Fig. 3C). These results suggested that NF-κB signaling functioned downstream of ERK and JNK, and that ERK and JNK negatively regulated the NF-κB pathway.

Discussion

VECs are the layer of blood vessels that is in direct contact with the blood, and is important for maintaining vascular tone and function, in part by the synthesis and release of certain substances in the vasculature, such as nitric oxide. VEC dysfunction contributes to the pathogenesis of vascular diseases in diabetics. While the mechanisms of VEC dysfunction in diabetes mellitus have remained to be clarified, one possibility is increased inactivation of endothelium-derived nitric oxide by ROS acting as signaling intermediates (14,15). 

In vitro, the high ambient glucose was observed to affect endothelial and other vascular cells at the cellular level (16), delay VEC replication (17) and cause excessive cell death (18). However, the molecular mechanisms of HG-induced damage in VECs require further investigation.
By detecting changes in the transcriptome in VECs induced by HG, 721 differentially expressed genes were uncovered and a possible association between the NF-κB pathway and HG treatment in VECs was identified. Inflammatory target genes (CCL13, PAI-1 and IL-8) associated with the activation of the NF-κB pathway were upregulated by HG, indicating that HG stress activates NF-κB target genes. In addition, at the protein level, an effect of HG on the activation of IκBα, which is the most characterized and studied NF-κB regulator, was observed. In other words, HG activated NF-κB target genes and the activation of the NF-κB regulator-IκBα to promote the inflammatory response in VECs. PAI-1, a pro-inflammatory cytokine located downstream of NF-κB (19), was found to be activated by HG.

It is thought that elevated levels of pro-inflammatory cytokines and proteases are present in chronic wounds due to abnormal immune responses (20). NF-κB is considered to be a regulator of the inflammatory response, which controls numerous disease states through regulating exaggerated inflammatory responses associated with certain diseases, including pulmonary diseases such as acute lung injury and acute respiratory distress syndrome (21). In Drosophila, the duration of JNK activation by lipopolysaccharide is directly controlled by the NF-κB protein Relish (22). The results of the present study contribute to the understanding of the molecular mechanisms regarding the interaction of NF-κB and ERK/JNK signaling in VECs under diabetic conditions. The JNK and ERK inhibitors activated IκBα as HG did. However, supplementation with NF-κB inhibitor Bayl17082 did not affect ERK and JNK activities, implying that ERK/JNK may function upstream of NF-κB signaling in VECs. In addition, NOD-like receptor signaling was altered upon HG stress in VECs. Decreased NOD-like signaling in human fibroblast cells with HG stress were also observed in a previous study (12). Furthermore, genetic variation of NOD1 and NOD2 has been associated with type II DM (23), suggesting the importance of NOD-like signaling in DM-mediated stress. Previous results have suggested that ECM remodeling is important for wound repair, which is in agreement with previous findings (24). EMC receptor interaction changes induced by HG stress identified in transcriptome analysis imply that HG-mediated inhibition of wound repair in VECs may be caused by ECM and receptor interaction, subsequently resulting in signaling inhibition. The present study provided useful information for understanding the mechanisms of HG stress on VECs and the results will be important for the further study of diabetes-associated complications.

Acknowledgements

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References