Research Article

Substance P Administered after Myocardial Infarction Upregulates Microphthalmia-Associated Transcription Factor, GATA4, and the Expansion of c-Kit+ Cells

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Microphthalmia-associated transcription factor (MITF), a basic helix-loop-helix leucine zipper transcription factor, can govern gene expression by binding to E box elements in the promoter region of its target gene. Although high levels of MITF have been observed in cardiomyocytes and the heart, the role of MITF after myocardial infarction (MI) remains unclear. We investigated the association between substance P (SP)/neurokinin-1 receptor (NK1R) signaling and MITF expression after MI. Male Sprague-Dawley rats (8 weeks) were randomly divided in two groups: ischemia/reperfusion injury (I/R) and SP injection (5 nmol/kg, SP+I/R). At the end of 7 days, the left ventricle (LV; LV7daysI/R, LV7daysSP+I/R) and infarct-related areas (IA; IA7daysI/R, IA7daysSP+I/R) from the hearts were collected. Immunofluorescence staining demonstrated that the LV7daysSP+I/R had a larger population of c-Kit+ GATA4high cells, which markedly upregulated MITF, c-Kit, and GATA4. c-Kit+ cells in the explant-derived cells (EDCs) derived from IA7daysSP+I/R migrated more widely than EDCs IA7daysI/R. Immunofluorescence staining, western blot analysis, and qRT-PCR assay showed that SP-treated c-Kit+ cells exhibited a high expression of c-Kit, GATA4, and MITF. FTY720 (a MITF inhibitor), RP67580 (NK1R inhibitor), or both inhibited the migration and proliferation of c-Kit+ cells increased by SP and blocked the upregulation of c-Kit, GATA4, and MITF. Overall, we suggest that MITF might be a potential regulator in SP-mediated c-Kit+ cell expansion post-MI via c-Kit and GATA4.

1. Introduction

Heart failure (HF) is expected to grow as an important clinical and public health challenge. HF occurs in response to hypertension and myocardial infarction from coronary artery disease, consequently giving rise to left ventricular (LV) remodeling [1, 2]. Although the mechanisms underlying cardiac growth, cardiac hypertrophy, and LV remodeling are still poorly understood, their intricate interactions might involve endogenous resident c-Kit+ cells and cardiomyocytes lost through direct injury [3–5]. For example, endothelin-1, angiotensin II, natriuretic peptides, and/or α1 and β-adrenergic agonists might play a substantial modulatory role in cardiac remodeling [4]. Cardiac transcription factors (such as GATA4, Nkx2.5, TRX5, and MEF2) and chromatin remodeling enzymes are also involved in stress regulation of the adult heart [5, 6]. Therefore, gaining a more precise understanding of the molecular mechanisms governing transcriptional regulation during postinfarct LV remodeling would be helpful in the development of novel strategies for HF therapies.

Microphthalmia-associated transcription factor (MITF), a basic helix-loop-helix leucine zipper DNA-binding protein, is known as a master transcription factor that has been detected in various cell types, such as melanocytes, mast cells, and osteoclasts [7–10]. In case of the heart, MITF-H, the heart-specific isoform, is abundantly expressed in cardiomyocytes and is important in the hypertrophic response
[7–10]. MITF-mutated mice (MITF<sup>ce/ce</sup> mice with the ce/ce mutation lack the Zip domain of MITF) show dramatically reduced heart weight/body weight ratios, systolic function, and cardiac output compared to wild-type mice [7]. A previous study has demonstrated that MITF regulates cardiac hypertrophy through targeting miR-541 [10]. According to these studies, MITF is critical for the cardiac growth and hypertrophic response, though only a few genes and signaling pathways associated with MITF have been recently identified [7–10].

Neuropeptide substance P (SP) is an undecapeptide member of the tachykinin neuropeptide family [11–14]. Studies have found SP to function as a neuromodulator for many biological processes through neurokinin-1 receptor (NK<sub>1</sub>R) [11–14]. Previous research describes the important roles of SP/NK<sub>1</sub>R in the cardiovascular system [11–14]. Several studies note the cardioprotective effects of SP on ischemia/reperfusion injury (I/R) [11–14]. However, the role of SP/NK<sub>1</sub>R signaling during cardiac repair after I/R remains unclear. Our previous study demonstrates that SP prevents I/R by modulating stem cell mobilization [12]. The present study investigates how exogenous SP affects the healing of a damaged LV after I/R. To accomplish this, we analyzed cardiomyc cell-specific gene expression and related factors in the LV and infarct-related areas (IA) of I/R rats with or without SP treatment.

2. Materials and Methods

2.1. Materials. SP (6.7 μg/kg/0.1 ml) and FTY720 (a sphingosine 1-phosphate receptor agonist that inhibits the expression of MITF) [15] were purchased from Sigma (St. Louis, MO, USA). RP67580 (RP, a selective nonpeptide tachykinin NK<sub>1</sub>R antagonist, 1 mg/kg) was obtained from R&D Systems Inc. (Minneapolis, MN, USA). Phospho-MITF (ser180) antibody, Hoechst 33342, and DAPI were purchased from Thermo Fisher Scientific (Rockford, IL, USA). AccuPower<sup>®</sup> RocketScript<sup>™</sup> Cycle RT PreMix (dN12) and AccuPower<sup>®</sup> ProFi Taq PCR PreMix were purchased from Bioneer (Daejeon, Korea). SYBR<sup>®</sup> Green Mix was obtained from Applied Biosystems (Lincoln, CA, USA). Antibodies specific for c-Kit (sc5535), NK<sub>1</sub>R (sc15323), MITF (sc56725), and actin (sc47778) were obtained from Santa Cruz Biotechnology, Inc. (Santa Cruz, CA, USA). Antibodies that recognize GATA4 (ab86371) and Alexa Fluor<sup>®</sup> secondary antibodies were purchased from Abcam (Cambridge, UK). Another antigen specific for c-Kit (phosphor Tyr568/-Tyr570) was obtained from GeneTex (Iseworth, UK). Phospho-Akt (4060S) and Akt were purchased from Cell Signaling Technology (Beverly, USA).

2.2. Animal Models and Ex Vivo Explant Outgrowth Culture Assay. All the experiments on the eight-week-old male Sprague-Dawley (SD) rats were carried out according to the Institutional Animal Care and Ethics Committee of Kyung Hee Medical Center (KHMC-IACUC:2015-028). The SD rats were randomly divided into 2 groups (n = 22 each): I/R and I/R with 5 nmol/kg SP injection (SP+I/R). The left anterior descending coronary artery was occluded for 40 min followed by 7 days reperfusion with and without SP. The rats were euthanized, and the LV (LVID<sub>d</sub>/days, LVID<sub>s</sub>/days+i<sub>1</sub>) derived from heart samples were collected. IA tissue was cut into 1 to 2 mm fragments, washed with Ca<sup>2+</sup>/Mg<sup>2+</sup>-free PBS, and digested three times for 10 min with 0.2% trypsin and 0.1% collagenase at 37°C. IA fragments were incubated with complete medium (CM; Dulbecco’s modified Eagle’s medium supplemented with 10% ES cell grade FBS, 5% horse serum, 10 ng/ml LIF, 1% penicillin-streptomycin, fungizone, and gentamicin) at 37°C in a 5% CO<sub>2</sub> incubator. After 2 weeks, the attached cells, which migrated out and surrounded the explants, were analyzed by immunofluorescence staining (IFS) with c-Kit antibodies. The c-Kit<sup>+</sup> cells were purified by a magneto-activated cell sorting (MACS) system (Dynal Biotech, Oslo, Norway). Explant-derived cells (EDCs) were suspended in trypsin, incubated with a rabbit anti-c-Kit antibody (1:100), and separated using immunomagnetic microbeads (Dynal Biotech). c-Kit<sup>+</sup> cells were cultured for 1 month with CM at 37°C in a 5% CO<sub>2</sub> incubator.

2.3. c-Kit<sup>+</sup> Cell Proliferation Assay. The c-Kit<sup>+</sup> cell proliferation assay was assessed using an EZ-Cytox cell viability assay kit (DoGEN, Seoul, Korea) [16]. The c-Kit<sup>+</sup> cells were pre-treated with FTY720 (5 μM) or RP67580 (20 μg/ml) or both for 2 h. After SP treatment for 7 days, the cultured medium was removed. Cells were stained with EZ-Cytox solution for 1 h. Absorbance was determined at 490 nm using an ELISA reader (EMax; Molecular Devices, Sunnyvale, CA, USA).

2.4. c-Kit<sup>+</sup> Cell Migration Assay. The cell migration assay was performed using 0.8 μm pore size and 24-well transwell migration chambers coated with type IV collagen (10 μg/ml) as previously described [16]. 1 × 10<sup>4</sup> c-Kit<sup>+</sup> cells were seeded into the upper transwell chambers containing medium without FTY720 or RP67580 or both. The inner chamber was inserted into each well of 24-well plates containing 600 μl medium supplemented with or without SP at the indicated concentration in the presence or absence of FTY720 (5 μM) or RP67580 (20 μg/ml) or both. The chambers were then incubated for 24 h at 37°C in a 5% CO<sub>2</sub> incubator. The cells that migrated through to the outer side of the membrane were stained with a crystal violet staining solution. The absorbance was determined at 590 nm using an ELISA reader (EMax; Molecular Devices, Sunnyvale, CA, USA).

2.5. Quantitative Reverse-Transcription PCR (qRT-PCR). cDNA was synthesized using AccuPower<sup>®</sup> RocketScript<sup>™</sup> Cycle RT PreMix (dN12) (Bioneer, Daejeon, Korea). qRT-PCR assays were carried out with SYBR<sup>®</sup> Green Mix and the appropriate primers (Applied Biosystems) and were run on a StepOnePlus real-time PCR system (Applied Biosystems). The relative gene expression from all data was obtained using the ΔΔct method with normalization versus RPL-32 as previously described [16]. The primers used were as follows: c-Kit (forward: AGACGTACAGATCCAGAATG, reverse: TGCTCTTTGCTGTACCTTT); GATA4 (forward: ACCCTGGAGACACCCCAAT, reverse: GTAGAGGC ACAGGCGTTGC); MITF (forward: CATCAGCGATCTTG CTACGC, reverse: TGCATGAACTGGGCTGCTTG); and
26. IFS and Confocal Microscopy. The LVs were fixed with 4% paraformaldehyde (PFA) and paraffin embedded, sectioned into 7 μm thick sections, and stained with standard IFS methods as previously described [10, 17]. To detect the expression of c-Kit or MITF after SP treatment, c-Kit+ cells were treated with SP (10 nM). After SP treatment for 1 day, the cultured medium was removed. After fixing in 4% PFA at 4°C for 15 min, cells were washed with PBS, permeabilized with 5% BSA and 0.1% Triton X-100, and incubated with primary antibodies. After nuclear DAPI and Hoechst staining, immunostained cells were imaged using an inverted Zeiss Axio Observer Z1 confocal microscope with 405, 488, 514, 561, and 633 nm laser lines. All images were selected with sample identities blinded, and at least 20 random images were obtained from each well or group.

2.7. Western Blot Analysis. The frozen samples were disrupted using the TissueLyser II (Qiagen), after which an ice-cold PRP-PREP protein extraction solution with a protease inhibitor cocktail (INTrON Biotechnology, Inc., Seoul, Korea) was added, and the samples were homogenized with stainless steel beads (Qiagen, CA, USA). Protein concentration was assessed using a BCA kit (Thermo Scientific, Rockford, IL, USA). An equal amount of protein (80 μg) from each sample was loaded onto 10% to 12% SDS gel and transferred to a PVDF membrane (Merck Millipore, MA, USA). The membranes were blocked for 2 h at room temperature with 5% nonfat dry milk in PBS containing 0.1% Tween-20 and incubated with primary antibodies (1:1000 and 1:500, respectively) overnight at 4°C. After washing three times, the membranes were incubated with a horseradish peroxidase-conjugated secondary antibody (1:5000) at RT for 2 h and visualized with a chemiluminescence substrate.

2.8. Statistical Analysis. Student’s t-tests (for comparisons between two groups) or a one-way analysis of variance (ANOVA) (for comparisons among three or more groups) followed by Tukey post hoc tests was used for the statistical analyses. SPSS software ver. 17.0 (SPSS, Chicago, IL) was used. A value of P < 0.05 was considered significant. Data are expressed as means ± standard error (SE). *P < 0.05; **P < 0.01; ***P < 0.001 vs. corresponding controls. All error bars represent the standard deviation of three or more biological replicates.

3. Results

3.1. SP Increases the Endogenous Resident c-Kit+ GATAhigh Cells and Upregulates c-Kit, GATA4, and MITF in I/R-Injured LV. The mechanism by which SP contributes to the expansion of c-Kit+ cells after MI [14] was analyzed LV I/R and LV I/R treated with SP using IFS with c-Kit and GATA4 antibodies. Confocal images showed an increase of c-Kit+ GATA4high-expressed cells in LV I/R compared to LV I/R (Figure 1). The expression of c-Kit and GATA4 is important for promoting growth and migration of c-Kit+ cells [17]. In addition, several studies of I/R-injured hearts have suggested that the number of c-Kit+ cells increases due to pressure overload-induced cardiac hypertrophy [17]. Representative confocal images show the coexpression of MITF and GATA4 or c-Kit in LV I/R and LV I/R treated with SP, indicating a high level of MITF expression by SP treatment (Figures 2 and 3). MITF plays a key role in the pathway leading to cardiac growth and the hypertrophic response in the heart (6–10). However, very little is known about the molecular mechanism of MITF after MI. Next, we hypothesized that if SP/NK1R signaling leads to the activation of endogenous resident c-Kit+ GATA4high cells, the expression of c-Kit, GATA4, and MITF might be involved. To examine this, we separated IA 7 days I/R and IA 7 days SP+I/R from LV 7 days I/R and LV 7 days SP+I/R. The expression levels of c-Kit, MITF, and GATA4 at the mRNA and protein levels were measured by qRT-PCR and western blot analysis. These analyses revealed a significant increase in c-Kit, GATA4, and MITF expression in LV 7 days SP+I/R compared to LV 7 days I/R (Figure 4). In particular, GATA expression in LV 7 days SP+I/R at the mRNA level markedly increased (Figure 4(c)). These results provide evidence that the molecular pathways of SP might induce the expansion of endogenous resident c-Kit+ GATA4high, expressed cells, which are associated with c-Kit, GATA4, and MITF expression.

3.2. MITF Plays an Important Role in SP-Promoted Proliferation and Migration of c-Kit+ GATAhigh Cells. If SP/NK, R signaling positively regulates the endogenous resident c-Kit+ cells in IA through c-Kit, GATA4, and MITF, then many more c-Kit+ cells from IA I/R may migrate than from IA I/R. To test this, IA 7 days I/R and IA 7 days SP+I/R were separated IA 7 days I/R and IA 7 days SP+I/R in the absence and presence of additional SP treatment (Figure 5). To determine whether SP affects the expression of MITF and GATA4 in c-Kit+ cells, we purified c-Kit+ cells from IA 7 days I/R and IA 7 days SP+I/R fragments were cultured using an ex vivo explant outgrowth assay. Expanded c-Kit+ cells were characterized by confocal microscopy. Confocal images and FACS analysis demonstrated that migrated c-Kit+ cells from IA 7 days I/R and IA 7 days SP+I/R were significantly higher than from IA 7 days I/R in the absence and presence of additional SP treatment (Figure 5). To determine whether SP affects the expression of MITF and GATA4 in c-Kit+ cells, we purified c-Kit+ cells from IA 7 days I/R and IA 7 days SP+I/R using MACS methods with c-Kit antibodies. Purified c-Kit+ cells were treated with SP at the indicated time points. As shown in Figure 6(a), SP-treated c-Kit+ cells markedly activated the phosphorylation of Akt at 10 min and enhanced the phosphorylation of c-Kit and MITF at 8 h. The expression of c-Kit, GATA4, and MITF induced by SP subsequently increased at 8 h (Figure 6(a)). qRT-PCR showed an elevated expression of MITF at 16 h (Figure 6(b)). Confocal images further confirmed that the location and expression of MITF considerably overlapped with c-Kit or GATA4 expression (Figures 6(c–6(e)). We next used RP67589 and FTY720 to further verify whether the expansion of c-Kit+ cells was at least in part due to SP/NK, R signaling and MITF expression. These inhibitors both alone and in combination significantly inhibited the enhancement of cell proliferation and migration induced by SP (Figures 7(a) and 7(b)). In addition, these inhibitors alone and together prevented the upregulation of c-Kit, GATA4, and MITF in SP-treated c-Kit+ cells at mRNA.
and protein levels (Figure 7(c), Fig. S1 and Fig. S2). Therefore, these observations indicate that SP/NK,R signaling-accelerated c-Kit+ cell expansion might relate to c-Kit, GATA4, and MITF expression.

4. Discussion

In the present study, we have identified an abundance of c-Kit+ GATA4\textsuperscript{high} cells in IA\textsuperscript{7 days I/R} after SP treatment. The expression of MITF and GATA4 at the protein and mRNA levels in IA\textsuperscript{7 days SP+I/R} and SP-treated c-Kit+ cells was dramatically increased. Although the specific roles of MITF and GATA4 in cardiac repair have not yet been validated, several studies have proposed that the functional link between MITF and GATA4 might affect the c-Kit+ cells after MI. A previous study has reported that MITF transcriptionally promotes GATA4 expression by binding to the E box in the GATA4 promoter in cardiac hypertrophy, responding to stress from ischemia [9]. MITF can also interact directly with c-Kit signaling, expressed not only in cranial neural crest derivatives and mast cells but also in cardiomyocytes [18]. For example, a previous study has clearly demonstrated that c-Kit identifies cardiac progenitors of cardiac neural crest (CNC) origin using two recombinase systems with c-Kit\textsuperscript{CreERT2/+} and Wnt1-Flpe recombinase driver mouse lines [18]. The c-Kit+ CNC-derived cardiomyocytes in c-Kit\textsuperscript{CreERT2/+} lineage tracing are widespread in atrial and ventricular cardiomyocytes, as well as in pericardial, endocardial, and epicardial cells [18]. Interestingly, MITF is expressed in c-Kit+ CNCs and their cardiomyocytic derivatives [18].

GATA4 is the zinc finger transcription factor that has demonstrated the ability to activate Bcl2 at mRNA and protein levels in adaptive responses by cardiomyocytes, helping to improve cardiomyocyte survival [19]. In another study, GATA4 overexpression in transgenic mice upregulated the level of Bcl2 in the heart and increased coronary flow reserve, as well as heightened cardiac contractility by promoting
angiogenic factors such as VEGF [20]. In addition, a previous study generated a D3 mouse embryonic stem cell line with highly expressed GATA4 [21]. This cell line enhanced cardiomyocyte generation in differentiating embryoid bodies and elevated the expression of endogenous cardiac differentiation genes [21]. In another study, researchers created c-Kit+ cells with lentiviruses encoding four different cardiac transcription factors—GATA4, MEF3C, NKX2.5, and TBX5—a-alone or combined with each other [22]. No significant induction of endothelial cell markers was found in GATA4-overexpressing c-Kit+ cells [22].

Recently, the previous work of Dr. Piero Anversa, who studied c-Kit+ cardiac stem cells, has been retracted. In addition, the role of endogenous c-Kit cells is still questioned whether cardiac stem cells existed. In fact, the mechanisms underlying acute MI are very different from those governing the wound repair response to MI, so the assumption that substance P should contribute to cardiac regeneration after MI is simply unfounded. Although we demonstrated that the expression of c-Kit, MITF, and GATA4 induced by SP is associated with the resident c-Kit+ cell expansion in LV repair post I/R, the present study is not the major focus of c-Kit, but rather SP signaling in c-Kit+ cells. Of note, there is difference in c-Kit, MITF, or GATA4 cellular distribution between the IA7daysI/R and IA7daysSP+I/R groups. The distribution of c-Kit+, MITF+, and GATA4+ cells in the IA7daysI/R group is relatively well distributed throughout the infarct, whereas the highest concentrations of c-Kit+, MITF+, and GATA4+ cells in the IA7daysSP+I/R group are far more focal and located within vascular structures surrounded by rings of DAPI+ cells. As shown in Figure 7(d), our findings did not establish a causal relationship between the SP-mediated activation of c-Kit, GATA4, and MITF and the expansion of c-Kit+ GATA4high MITF+ cells in the post I/R myocardium. Nonetheless, we have clearly demonstrated by comprehensive analysis of cardiac specific markers between LV7daysI/R and LV7daysSP+I/R that SP, through GATA4 and MITF expression, can elevate an expansion of c-Kit+ GATA4high cells. Overall, our results suggest that MITF might play a transcriptional regulator role linking c-Kit+ GATA4high cells and SP-mediated c-Kit+ cell expansion post-MI.
Figure 3: Confocal analysis of MITF and GATA4 in LV
7 days I/R and c-LV
7 days SP+I/R. Confocal images of MITF staining (red fluorescence) in conjunction with GATA4 (green fluorescence) and nuclear staining (blue fluorescence) in sections of LV
7 days I/R and LV 7 days SP+I/R. Scale bars, 20 μm with a further magnified area indicated by the dotted white circle.

Figure 4: High expression of MITF, c-Kit, and GATA4 in IA
7 days SP+I/R. (a) Western blot analysis results for the IA
7 days I/R and IA
7 days SP+I/R rats. (b–d) qRT-PCR analysis of relative mRNA levels of the targeted genes and *P < 0.05 versus corresponding controls using Student’s t-test.
Figure 5: Expansion of c-Kit+ cells in IA7daysI/R and IA7daysSP+I/R with or without SP. (a) Confocal fluorescence images showing c-Kit-expressing (green) cells in each group. Hoechst 33342 (blue) allowed visualization of the nucleus in cells. Scale bars, 20 μm. Graphs of the number of c-Kit+ cells/mm² (%) in each group expressed as a percentage difference compared to the corresponding control. **P < 0.001 versus corresponding control using Student's t-test. (b) Representative dot plot of FACS analysis of EDCs derived from IA7daysI/R and IA7daysSP+I/R in the presence or absence of SP (10 nM) using c-Kit antibodies. Negative controls indicated as the assessment of nonspecific binding of Alexa Fluor®488 antibodies to c-Kit+ cells.

Figure 6: SP induces the expression of MITF in c-Kit+ cells. (a) Western blot analysis showing the expression of targeted proteins in SP-treated c-Kit+ cells over time. (b) qRT-PCR analysis of relative mRNA levels of MITF in c-Kit+ cells after SP treatment (10 nM) at the indicated time points. *P < 0.05 or **P < 0.01 versus corresponding controls using Student’s t-test. (c–e) Confocal microscopy images showing c-Kit, MITF, and GATA4 and dual staining of these targeted proteins in untreated and SP-treated c-Kit+ cells. DAPI (blue) allowed visualization of the nucleus in cells. Scale bars, 100 μm.
Abbreviations

CNC: Cardiac neural crest  
EDCs: Explant-derived cells  
HF: Heart failure  
IA: Infarct-related areas  
I/R: Ischemia/reperfusion injury  
LV: Left ventricular  
MI: Myocardial infarction  
MITF: Microphthalmia-associated transcription factor  
NK1R: Neurokinin-1 receptor  
qRT-PCR: Quantitative reverse-transcription PCR  
SP: Substance P.

Data Availability

(1) All data used to support the findings of this study are included within the article. (2) All data used to support the findings of this study are included within the supplementary information file. (3) All data used to support the

Conflicts of Interest

The authors declare that they have no competing interests.

Authors’ Contributions

YM Jeong, XW Cheng, and W Kim conceived, designed, improved, and corrected the style of the article; YM Jeong performed the experiments and wrote the paper.

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**Supplementary Materials**

Fig S1: the effects of FTY720 and RP67580 inhibitors on c-Kit cells with or without SP. (A) RT-PCR analysis of relative mRNA levels of the targeted genes in SP-stimulated c-Kit+ cells pretreated with one or both of these inhibitors. RPL32 is the loading control. (B) The plot densitometry graph of NK,R, MITF, GATA4, and c-Kit mRNA. Fig S2: FTY720 and RP67580 inhibit the expression of MITF, GATA4, and c-Kit protein in SP-treated c-Kit+ cells. The plot densitometry graph of NK,R, MITF, GATA4, and c-Kit protein that western blot analysis results for SP-stimulated c-Kit+ cells pretreated with one or both of these inhibitors. *P < 0.05 or **P < 0.01 versus corresponding controls using one-way analysis of variance (ANOVA) followed by Tukey’s post hoc test. (Supplementary Materials)

**References**


