Dual-ion delivery for synergistic angiogenesis and bactericidal capacity with silica-based microsphere

Khaliun Boldbaatar\textsuperscript{a,b,\#}, Khandmaa Dashnyam\textsuperscript{a,b,c,\#}, Jonathan C. Knowles\textsuperscript{b,c,d,e}, Hae-Hyung Lee\textsuperscript{a,c,f}, Jung-Hwan Lee\textsuperscript{a, c,f,*}, Hae-Won Kim\textsuperscript{a,b,c,f,*}

\textsuperscript{a}Institute of Tissue Regeneration Engineering (ITREN), Dankook University, Cheonan 330-714, Republic of Korea
\textsuperscript{b}Department of Nanobiomedical Science & BK21 PLUS NBM Global Research Center for Regenerative Medicine, Dankook University, Cheonan 330-714, Republic of Korea
\textsuperscript{c}UCL Eastman-Korea Dental Medicine Innovation Centre, Dankook University, Cheonan, Republic of Korea
\textsuperscript{d}Division of Biomaterials and Tissue Engineering, Eastman Dental Institute, University College London, London, UK
\textsuperscript{e}The Discoveries Centre for Regenerative and Precision Medicine, Eastman Dental Institute, University College London, London, UK
\textsuperscript{f}Department of Biomaterials Science, College of Dentistry, Dankook University, Cheonan 330-714, South Korea

\#KB (khaliun1209@gmail.com) and KD (khandmaa@naver.com) contributed equally to this paper as first authors.

* Co-corresponding authors: JHL (ducious@gmail.com), HWK (kimhw@dku.edu)

Jung-Hwan Lee, DDS, PhD, Researcher, Institute of Tissue Regeneration Engineering (ITREN), Dankook University, Cheonan 330-714, Republic of Korea, Tel: +82 41 550 3081; Fax: +82 41 559 7839; E-mail: ducious@gmail.

Hae-Won Kim, PhD, Professor, Institute of Tissue Regeneration Engineering (ITREN), Dankook University, Cheonan, Republic of Korea, Tel: +82 41 550 3081; Fax: +82 41 559 7839; E-mail: kimhw@dku.edu
Abstract

Inhibition of bacterial growth with the simultaneous promotion of angiogenesis has been challenging in the repair and regeneration of infected tissues. Here, we aim to tackle this issue through the use of cobalt-doped silicate microspheres that can sustainably release dual ions (silicate and cobalt) at therapeutically-relevant doses. The cobalt was doped up to 2.5 wt% within a sol-gel silicate glass network, and microspheres with the size of ~300 µm were generated by an emulsification method. The cobalt and silicate ions released were shown to synergistically upregulate key angiogenic genes, such as HIF1-α, VEGF and the receptor KDR. Moreover, the incorporation of ions promoted the polarization, migration, homing and sprouting angiogenesis of endothelial cells. Neo-vascular formation was significantly higher in the dual-ion delivered microspheres, as evidenced in a chicken chorioallantoic membrane model. When cultured with bacterial species, the cobalt-doped microspheres effectively inhibited bacteria growth in both indirect or direct contacts. Of note, the bacteria/endothelial cell coculture model proved the efficacy of dual-ion releasing microcarriers for maintaining the endothelial survivability against bacterial contamination and their cell-cell junction. The current study demonstrates the multiple actions (proangiogenic and antibacterial) of silicate and cobalt ions released from microspheres, and the concept provided here can be extensively applied to repair and regenerate infected tissues as a growth factor- or drug-free delivery system.

Keywords: Ion delivery, cobalt/silicate, angiogenesis, antibacterial, multifunctional biomaterials

Statement of significance

While several ions have been introduced to biomaterials for therapeutic purposes, relaying the effects of antibacterial into tissue regenerative (e.g., angiogenesis) has been a significant challenge. In this study, we aim to develop a biomaterial platform that has the capacity of both ‘antibacterial’ and ‘proangiogenic’ from a microsphere sustainably releasing multiple ions (herein cobalt and silicate). Here, dual-actions of the microspheres revealed the stimulated endothelial functions as well as the inhibited growth of different
bacterial species. In particular, protecting endothelial survivability against bacterial contamination was reported using the bacterial/endothelial co-culture model. The current concept of drug-free yet multiple-ion delivery biomaterials can be applicable for the repair and regeneration of infected tissues with dual actions of angiogenesis and suppressing bacterial activity.

Graphical abstract

Dual ions (silicate and cobalt ions) released from microspheres promote angiogenesis while simultaneously inhibiting bacteria growth, as demonstrated by the salvaged survivability of endothelial cells against bacterial contamination. The multiple ion-delivery microspheres can be potentially useful for the repair and regeneration of infected tissues as a drug-free biomaterial platform.
Introduction

The delivery of therapeutics such as drugs or biomolecules that can promote tissue functions is considered an effective strategy to engineer biomaterials for regenerative medicine [1–3]. Recently, ions have been highlighted as the therapeutic components of biomaterials [4,5]. For example, treating damaged tissues with angiogenic ions (i.e., Si ions) was effective in accelerating tissue regeneration through neo-vascular formation [6,7]. In fact, myriad therapeutic ions (Ag\(^+\), Ca\(^{2+}\), Cu\(^{2+}\), Co\(^{2+}\), Mg\(^{2+}\), Sr\(^{2+}\), and Zn\(^{2+}\)) have been doped into biomaterials at proper doses to target other therapeutic functions such as stimulating cell proliferation, controlling cell differentiation, and relieving inflammation [8–11]. However, synergistic or combined effects from different ions delivered by biomaterials have not been investigated in detail, although this delivery can mimic the microenvironment in the human body, where many ions play roles in tissue regeneration [12].

A significant hurdle for tissue regeneration often arises from bacterial infection, which mitigates the tissue healing/repair process and results in limited regeneration [13]. Clinical therapies generally involve treatment with antimicrobial drugs or surgical removal of infected tissues [14,15]. However, complications are often encountered, including the insufficient function of administered or locally injected antibiotic drugs, incomplete removal of infected tissues due to anatomical complexities, and the limited bactericidal/bacteriostatic effects without additional therapeutic functionality [16].

In case of regeneration of infected tissue, the body naturally reacts to disease-causing bacteria by increasing local blood flow along with the stimulated angiogenesis intrinsically (from infected site to outer region) and extrinsically (from outer region to infected site) and recruiting immune cells to destroy the bacteria[17]. But, when initial angiogenesis is not processed in the infected site due to the destruction of proangiogenic precursors in surrounding area, the delay of tissue repair/regeneration is inevitable due to insufficient clearance of pathogens and consequent accumulation of toxins, deteriorating angiogenesis for the infected tissue regeneration [18]. Therefore, in case of accelerating regenerative potential in severely infected tissue, simultaneous angiogenic and antibacterial biomaterials are required, maximizing intrinsic angiogenesis by preserving
angiogenic precursors against toxins from bacterial overgrowth.

Therapeutic ion-incorporated materials can have a pivotal role in addressing the above issues. Metallic compound (e.g., Ag, Co, Cu, Ga, Mn, or Mg)-loaded biomaterials have been studied as promising antibacterial materials; sometimes, they are used with antibiotic drugs, which decreases the dose of the antibiotics and consequent adverse effects [19–23]. Among antibacterial metal compound, some metal (e.g. Co and Cu) ions are also well-known to promote angiogenesis via regulation of hypoxia-inducible (HIF)-1α, which can be synergistically activated by another angiogenic ion (i.e., Si ions) [24]. Thus, relaying the antibacterial and angiogenic activity into tissue regeneration process can be realized through the ionic delivery approach.

Here, we strategize the dual delivery of Si and Co ions to reap up the dual-functioning (pro-angiogenic and antibacterial functions). We design ion-doped glass microparticles by a sol-gel method that can deliver therapeutic ions (Si and Co ions) sustainably at therapeutic dose. While the Si ions released from the glass chemistry at a therapeutically relevant level, known to stimulate angiogenesis [25,26], the Co ions have also been implicated in stimulating angiogenesis [27,28]. Moreover, Co ions have been shown to have antibacterial activity, penetrating the bacterial wall of the microorganisms and inhibiting bacterial survivability and, thus, preserving angiogenic precursors for blood vessel formation [29,30]. The effects of the delivered ions on angiogenic behaviors are investigated in terms of polarization, migration, and tubular networking of endothelial cells in vitro as well as neo-blood vessel formation in vivo. In parallel, antibacterial effects and endothelial cells’ survivability are investigated using a cell-bacteria co-culture system. This approach, as the first report on the ionic-delivery of biomaterials for dual-functions, is envisaged to shed light on the development of a new class of therapeutic biomaterials for infected tissue regeneration.
2. Materials and methods

2.1 Preparation of the microcarriers

Tetraethyl orthosilicate (TEOS, Sigma-Aldrich, St. Louis, MO, USA) was used to form a silica network, and 0.1 M HCl was added to 10 ml of TEOS. Cobalt chloride (4%, Sigma-Aldrich) was separately diluted in deionized water. Cobalt chloride solution (at four different concentrations 0, 1, 2.5, and 5 wt%) was added to the silica sol, which was adjusted to pH values between 5.0-5.5 by the addition of 0.08 M ammonium hydroxide (Sigma-Aldrich). Cobalt-incorporated microcarriers were collected after stabilization, washed several times with ethanol and left overnight to dry. Then, 0%, 1%, 2.5%, and 5% cobalt incorporated in silica microcarriers were coded as Co0, Co1, Co2.5, and Co5.0. But, Co5.0 was excluded in further characterization due to their failure of cytocompatibility by cell viability test.

2.2 Microcarrier characterization and ion release profile

Scanning electron microscopy (SEM; JEOL JSM 6510, Tokyo, Japan) was used to observe the homogenous distribution of microspheres and mesoporous morphology of the surface. Cobalt ion incorporation was identified by the EDS method. Mesoporous characterization was quantified using the Quadrasorb SI automated surface area and pore size analyzer (Quadrasorb SI, Quantachrom instruments Ltd., Boynton Beach, FL, USA) using N2 gas adsorption/desorption with the Brunauer-Emmet-Teller (BET) theory. The total pore volume was determined from the adsorbed amount at maximum relative pressure (P/P0).

Cobalt and silicon release from microcarriers was measured in α-minimal essential medium (α-MEM LM 008-01, Welgene, Gyeongsan-si, Korea) solution for up to 7 days (10 mg/mL) at 37 °C. The samples were centrifuged, the supernatant was collected, and 10 µL of nitric acid was added to the supernatant to inhibit bacteria and free any bound cobalt and silicon prior to measurement. Inductively coupled plasma atomic emission spectroscopy (ICP-AES, OPTIMA 4300 DV, Perkin-Elmer, Waltham, MA, USA) was used for detection. 5 wt% microsphere group was excluded in this characterizing tests due to their failure of cytocompatibility by screening cell viability test.
2.3 Endothelial cell viability

First, 2.0 x 10^4 human umbilical vein endothelial cells (HUVECs, cat# PCS-100-010, ATCC, Manassas, VA, USA) at passages 4-5 were used to test cell compatibility for up to three days. Vascular cell basal medium (cat# PCS-100-030, ATCC) supplemented with endothelial cell growth kit-VEGF (cat# PCS-100-041, ATCC) was used as endothelial cell medium. Microcarriers were indirectly placed on the top of the well inserts (10 mg/mL), and cells were cultured on the bottom of a 24 well plate. At day 3, 100 µl of CCK-8 (cell counting kit-8, Dojindo Molecular Technologies Inc, Rockville, MD, USA) solution was added into each well and then incubated at 37°C for 2 h according to the manufacturer’s instructions. The supernatant was collected, and its absorbance was measured at 450 nm using a microplate reader (SpectraMax M2e, Molecular Devices, Sunnyvale, CA, USA). This test was used for screening experimental groups for previous physico-chemical characterization (section 2.2) and further biological study, excluding 5 wt % microsphere group.

2.4 Quantitative real-time polymerase chain reaction (qRT-PCR)

Pro-angiogenic gene expression was quantified by qRT-PCR. HUVECs (1.0x10^5) were indirectly treated with microcarriers for 24 h by using transwell insert (10 mg/mL). After removing the insert, the cells on the bottom were washed with cold phosphate-buffered saline (PBS), and total RNA was extracted by centrifuging and filter system according to the manufacturer’s instructions (GeneAll, Ribospin, Seoul, Korea). After purified mRNA was quantified (Nanodrop 2000 spectrophotometer; Thermo Scientific, Waltham, MA, USA), cDNA was synthesized by reverse-transcribing 1 µg of mRNA using the High-Capacity RNA-to-cDNA Kit (Thermofisher) and carried out by a thermal cycle (Model 2720, Applied Biosystems, Foster City, CA, USA). PCR amplification was performed using the SYBR green qPCR SuperMix reagent (Invitrogen) and analysed by a real-time PCR system (7300-RT-PCR; Applied Biosystems, Foster City, CA, USA) according to the manufacturers’ instructions [31]. GAPDH was used to normalize VEGF and HIF1-α genes with respect to control according to the previous literatures [32] using 2^(-∆∆Ct) value by StepOne (v2.3, Applied Biosystems, Foster City, CA, USA). Furthermore, different concentrations of cobalt chloride (6, 12, and 18 ppm) were used to study VEGF,
HIF1-α, and KDR gene expression to compare the effects of the free form of cobalt ions with the effects of the cobalt ions in the microcarriers. Cobalt chloride was dissolved in distilled water and added to the transwell plate with silica microcarriers, which were then used for the experiment.

2.5 Endothelial cell polarization study
HUVECs (5.0x10^3) were seeded on 24 well plates for 4 h for the cells to attach to the plate, and then microcarriers with transwell inserts (10 mg/mL) were placed there for 1 h. After removing transwell inserts, cells were fixed and stained with 4′,6-diamidino-2-phenylindole (DAPI, nucleus) and phalloidin (cytoskeleton). Each group of images was captured by confocal laser scanning microscopy (CLSM, LSM 700, Carl Zeiss, Oberkochen, Germany). Polarized distance was identified by measuring the distance between the centres of mass and the nucleus based on the captured images using Image J (ver. 1.52a, NIH, USA).

2.6 Ion attraction study
Microcarriers were positioned in the bottom of a 24 well plate (10 mg/mL), and 5.0 x 10^3 cells were seeded on the 8 µm pore-sized transwell insert for 4 h. Chemoattraction was examined after 4 h by staining the nucleus of the cells above the insert after scratching the bottom side. Attracted cell numbers were counted using ImageJ software and converted into a percentage.

2.7 Endothelial cell migration
Mitomycin B (Sigma) was used as an inhibitor of proliferation and mixed with non-supplemented HUVEC media. Moreover, 1.0x10^5 cells were seeded initially, and 24 h was waited for the cells to achieve confluence; mitomycin B was added to cells for 2 h. A 200 µL tip was used to scratch the confluent cell layer to observe cell migration. Microcarriers were placed on transwell insert plates (10 mg/mL) and transferred to the scratched cells. Images were taken at 6 h, 12 h, and 24 h by light microscopy (IX-71; Olympus, Tokyo, Japan).
2.8 Tubular network and formation ability

Tubulogenesis was determined using a Matrigel (356234, Corning, NY, USA) matrix. The Matrigel was prepared and coated onto 24-well plates according to the manufacturer’s instructions. HUVECs (3.0x10⁴) were seeded onto the Matrigel, and microcarriers (10 mg/mL) were delivered with the insert. Cells were photographed at each time point (6 h and 12 h) using light microscopy (IX-71; Olympus). Three samples were selected in each group, and random images were analyzed for the measurement of the number of branch points (nodes), mesh-like circles (circles), and total tubule length.

2.9 Endothelial cells culturing on Matrigel-coated microcarrier

Microcarriers were covered by Matrigel to study cell adhesion and spreading. First, Matrigel was diluted in α-MEM media, and microcarriers were soaked in the diluted Matrigel solution at 4°C for an hour on a shaker to obtain a homogenous coating. Next, Matrigel-coated microcarriers were collected and incubated at 37°C for 1 h to complete the gelation of the Matrigel. After the Matrigel-coated the microcarriers, 5.0 x 10⁴ cells were seeded directly onto the microcarriers and cultured for 6 h for the observation of cell attachment. Cell adhesion and spreading were observed by SEM and CLSM using the z-stack mode.

2.10 Angiogenesis in chick chorioallantoic membrane (CAM) model

Fertilized chicken embryos were pre-incubated at 37°C for 3 days under 60% humidity. At day 3, albumen was aspirated and returned for incubation until day 7. No-treatment, Co0 and Co2.5 groups were selected to study angiogenesis in the CAM model. Ten milligrams of microcarriers was placed on the chick membrane surface on day 7, and the samples were transferred to the rotating incubator. At day 11, the chick membrane was collected and fixed by neutralized buffered formalin for 24 h. Membranes were dried and visualized by Digital SLR camera (EOS 1000D, Canon, Tokyo, Japan). Tubule length, size, and the total junction of the vessel were further analyzed by AngioQuant software (Version 1.33, Mathwork Inc, Houston, TX, USA).

2.11 Antibacterial effect using bacteria monoculture system
Antibacterial tests were performed using two species of bacteria, *Escherichia coli* (*E. coli*, ATCC 25922) and *Streptococcus aureus* (*S. aureus*, ATCC 25923), representing gram-negative and gram-positive species, respectively. Bacteria (1 x 10^7/mL) in the log phase of growth were cultured with 10 mg of microcarriers or different concentrations of CoCl₂ solution in 24-well plates with inserts. For positive or negative control, each bacteria strain was cultured with vancomycin disk (30 µg, Thermo fisher, Waltham, MA, USA) or without any extrinsic biomaterials or chemicals respectively for further study.

After bacterial species were cultured for 4 h, 10% PrestoBlue (Molecular probes, Invitrogen, Waltham, USA) was added, and culturing continued for another 1 h. The liquid (100 µL) from each well (n=5) was transferred to each well of 96-well plates, and the absorbance at 570 nm was measured using a microplate reader (SpectraMax M2e) and then normalized to the absorbance at 600 nm. Previously, along with the PrestoBlue assay, colony-forming units (CFUs) from serially diluted bacterial species were counted to confirm the correlation between the normalized absorbance in the PrestoBlue assay and the microbial CFU numbers [27,28]. The assays were independently performed in triplicate.

The Live/Dead kit assay (Thermo fisher) was used to visualize bacterial survivability according to the manufacturer's protocols [23]. Briefly, bacteria (10^7 /ml) with microcarriers (10 mg) were incubated for 4 h and stained with Syto-9 and propidium iodide for direct visualization. To remove the autofluorescence backgrounds of microcarriers and bacterial media, centrifugation (10000 rpm, 3 min) was used for collecting bacteria species. The supernatant was discarded, and PBS (1 ml) with the above two fluorescent dyes was added to stain bacteria. Green fluorescence represents intact live species, and red represents dead species.

2.12 *Endothelial cell and bacteria co-culture*

Co-culture of HUVECs and *E. coli* was performed based on modified protocols, indicated in other literatures [33–35]. After HUVECs (2x10^4/100 µL) was seeded on the each well (96 well plate) and cultured for 24 h, 10 µL of *E. coli* (1 x 10^7 or 1 x 10^6/mL) in the log
phase of growth was added with 100 µL of extraction from microcarriers in HUVECs’ media (10 mg/mL) immersed for 24 h at 37 °C. After 6 h of co-culturing, live and dead (Thermo fisher, Waltham, MA, USA) or VE-cadherin staining (1:50, sc-6458, Santa cruz biotechnology, Dallas, TX, USA) was performed (n=3). Culturing HUVECs with extraction or HUVECs’ media without bacteria contamination was used as a control. Live and dead cells per image were counted from randomly selected five images in each well and averaged. Fluorescence of VE-cadherin normalized to control was analyzed from randomly selected five images in each well by Image J (NIH).

2.13 Statistics
Statistical analysis was performed with a one-way analysis of variance (ANOVA) with the Tukey honest significant differences test as a post hoc test (IBM SPSS Statistics v23.0; IBM Corp) (P=0.05). Normal distribution test was performed by Shapiro-Wilk test.
3. Results and discussion

3.1 Microcarrier characterization and ion release

The doping of Co into Si-based microspheres was aimed to achieve the synergistic effect of released Co$^{2+}$ and Si ions on angiogenesis as well as the antibacterial effect of Co$^{2+}$. After initial screening test, 5 wt% microsphere was excluded for further study, which are shown in detail below. Two different concentrations of Co (1 and 2.5 wt%) were evenly well doped into the Si-microsphere, as confirmed by EDS results (Supplementary data.1). Comparably sized microspheres (200–350 µm) with similar mesoporous structures (2-3 nm) (Fig. 1c) and high surface areas (450–620 m$^2$/g) (as detected by BET analysis) (Fig. 1b) were successfully produced by the sol-gel method. Doping of Co to the Si microspheres affected the structure of the microsphere to a limited extent except glass amorphous network (Supplementary data.1); it slightly increased the pore size and decreased the surface area without significant change of the total diameter, their distribution and amorphous structure along with the results from amorphous glass structure with a network modifier, enlarging 3D space in molecular level [1,36].

The Si$^{4+}$ and Co$^{2+}$ ions released in α-MEM were measured using ICP-AES for up to 3 days (Fig. 1c). The Co$^{2+}$ ions were released only from the 1% and 2.5% groups, as anticipated, in an initial burst release (~4 h) and highly sustained manner after that point (0% and 2.5% are representatively shown). In the 2.5% group, Co$^{2+}$ ions were released rather rapidly for up to 4 h (~9 ppm) and then continuously for up to 7 days (~15 ppm); thus the daily release in a week was ~1-2 ppm (per day). Loosely bound cobalt in the silica network as a modifier initially released in ionic form, and then cobalt in the inner part of network started to release in a sustained manner after that, accelerated by mesoporous structures of microspheres to approach the therapeutic concentration of ions. In a previous study, more than 2 ppm of Co$^{2+}$ ion release was shown to be toxic to the surrounding tissues [37]. Moreover, silicate ions were released at 2–3 ppm per day, which was reported to be nontoxic and in a therapeutic range [25,38].

To investigate cytocompatibility of microspheres in accelerated condition, enabling exclusion of toxic experimental group(s) for further biological studies, different concentration of microspheres (10–100 mg) were cultured indirectly (using a transwell...
insert) with relatively lower HUVECs numbers (2.0 x 10⁴/mL of 1 mL in 24 well) for 3 days, exhibiting no significant cellular toxicity in all groups except 5wt% (~30%) compared to control when used at 10 mg/mL of microsphere concentration. Thus, 0, 1, and 2.5 wt% at 10 mg/mL of microsphere concentration are selected for the following in vitro experiments (Supplementary data 2).

3.2 Cobalt- and silicate-ion synergism in angiogenesis-related gene expression

As a biological assay in detail, firstly, the synergistic role of the Co and Si ions in angiogenesis was investigated by the expression of angiogenic genes, quantified by qRT-PCR. VEGF, a major growth factor for angiogenesis, is well known to be upregulated by the activation of HIF1-α [39–41]. According to the literature, Co²⁺ ions can stimulate hypoxic conditions via direct binding to HIF1-α and stabilizing it, while Si⁴⁺ has been reported to promote HIF1-α expression by degrading the PHD2 enzyme [7], leading to the possibility of synergistically accelerating angiogenesis-related gene expression [27,42]. The Co2.5 group showed significantly higher expression of pro-angiogenic genes including VEGF, KDR and HIF1-α than the Co0 and Co1 groups, while Co0 also showed higher gene expression than the control, which means that Co²⁺ and Si ions synergistically promoted the expression of VEGF and KDR via upregulation of HIF1-α (Fig. 2). To confirm the role of Co ions, the free form of Co²⁺ ions (6, 12 and 18 ppm) was used to supplement Co0; these concentrations are around the range of the concentrations of Co ions released into same amount media from microspheres (10 mg/mL) for 24 h (Fig. 1c), and gene expression occurred in the same manner. All Co ion-treated groups showed a significant increase in gene expression, especially 12 ppm Co ions, which is the similar concentration of Co ions released into media in 24 h from the Co2.5 (~12 ppm). Because the Co2.5 group showed the greatest increase in angiogenic gene expression, 12 ppm Co ions was the optimal concentration for synergetic effects, matching results from other endothelial cells such as HMEC-1 and umbilical cord blood-derived CD133(+) cells [28,43].

3.3. Promotion of endothelial cell functions in vitro

The polarization of endothelial cells initiates endothelial cell migration for angiogenesis,
thus considered as the initial cell behavior for the formation of neo-blood vessels and mature vasculature [44,45]. Here the cellular polarization was determined in endothelial cells by measuring the shift in the cellular nucleus from the centre of mass at a single cell level (Fig. 3a). For single cell analysis, the twenty fold decreased cell number compared to that for gene expression study was seeded for 4 h and further co-cultured for 1 h with 10 mg of microspheres. Compared with the control group (w/o microspheres) all microsphere groups significantly promoted cell polarization, and the Co-releasing groups (Co1 and Co2.5) synergistically increased the cell polarization with respect to Co0. The cell homing behavior was then carried out to clarify the chemo (ion) attractant efficacy of the released Co2+ ion, an essential characteristic during angiogenic process for making tubular formation. After the small number of cells (5 x 10^3) initially placed in a transwell insert for 4 h then moved toward the ion-releasing microspheres in the 24 well bottom place, ion attracted cell numbers from insert top surface to microspheres in 24 well bottom were counted. Results showed that the number of cells migrated down increased in the order: control (~5%) < Co0 and Co1 (~60%) < Co2.5 (~85%) (Fig. 3b), which is similar to what was observed in cell polarization.

Next, the cell migration assay involving a scratch model was then performed to determine cellular motility (Fig. 4). In particular, HUVECs were treated with mitomycin B to inhibit proliferation. The cellular gathering into a scratched gap was captured at different time points (6, 12, and 24 h) to count the migrated cells (Fig. 4a). The migrated cell number increased with time (6 h < 12 h < 24 h) and the cell number became significantly different among groups with time (w/o Contr. < Co0 < Co1 < Co2.5, Fig. 4b).

After this, tubular formation on a Matrigel was determined for 12 h to demonstrate in vitro angiogenesis on biomimetic soft extracellular matrix condition. The greater vessel-like formation was found in the Co2.5 group than in the control and Co0 groups at each time point based on the image and quantitative analysis of node and circle numbers (6 and 12 h, P<0.05, Fig. 5a and b). After confirming the role of ions released from the microsphere, the endothelial cells were directly cultured on the microspheres after coated with Matrigel. The F-actin stained and electron microscopic images indicated that endothelial cells adhered and elongated much better on the surface of the co-delivery
microcarriers (Fig. 5c). Taken the results together, the synergetic stimulation of in vitro endothelial cell behaviors such as initial polarization poising, cell homing by ion attraction, cell migration and tubular formation were demonstrated by the simultaneous release of Co$^{2+}$ and Si$^{4+}$ ions from the microcarriers.

3.4. In vivo neo-vessel formation with microcarriers

Lastly, a CAM assay was performed to evaluate new blood vessel formation in the in vivo condition (Fig. 6). Microspheres of 10 mg were implanted for 4 days in the model, and the neo-vessel formation was imaged and analyzed. Optical micrographs showed a clear difference in blood vessel formation among the control, Co0 and Co2.5 groups, with noticeable observation of a number of neo-vessels branched from old vessels in the Co2.5 group. When the neo-vessel indices were quantified, the Co2.5 group exhibited significantly greater total vessel length, size, and junction (~3 times) than the Co0 and control group, demonstrating that Co$^{2+}$ and Si$^{4+}$ ions synergistically promoted the formation of vascular networks from existing vessels in in vivo systems.

Previously, we showed that the Si ion-releasing microspheres used at relatively high amount (100 mg) had pro-angiogenic effects (~1.5 times higher than control) in CAM assay [7]; strikingly, here we noticed that the Co/Si-releasing microspheres administered at only 10 mg (1/10 of the previous study) stimulated the neo-vessel formation of control by as high as ~3 times, and this level is comparable to that when VEGF was co-delivered with Si ion in the previous work. This fact suggests that the synergistic action of co-delivered ions might be effective in such a way that can replace the function of angiogenic growth factor(s), and more ambitiously, open a new strategy of growth factor-free therapeutic delivery system, which yet needs further comparative studies.

3.4 Antibacterial potential of the microcarriers

To examine the antibacterial effects of the Co/Si ion-delivering microspheres, two representative bacteria, *E. coli* (gram +ve) and *S. aureus* (gram -ve), that are prevalent bacterial strains in damaged tissues, were selected. The results showed that bacterial survivability was inhibited to almost below 50% in the Co2.5 group, demonstrating the
effectiveness of the Co$^{2+}$ ions released from microspheres (Fig. 7a). The use of free Co$^{2+}$ ions (6~18 ppm) from CoCl$_2$ similarly inhibited the bacterial viability. As the free Co$^{2+}$ ions have been used as positive control for strong antibacterial agent, the microspheres are considered to hold excellent antibacterial activity [29,30,46].

The live/dead staining of bacteria clearly showed the damage of cell membrane (Fig. 7b). More bacteria cells were damaged (red) (fewer live cells, in green) in the Co1 and Co2.5 groups than in the Co0 group. Because of the autofluorescence of microcarriers, detecting red signals in direct contact condition was difficult.

Supplementary data 3 shows more bacteria alive appeared in Co0 group compared to G1 and G2.5 groups. Taken from antibacterial results, the Co$^{2+}$ ions released from microcarriers were effective enough to inhibit bacterial survivability.

3.5 Salvage of endothelial cells from bacteria contamination

To further examine the multi-functional effects of the Co/Si-delivery microcarriers in a more clinically relevant pathogenic condition, HUVECs were intentionally contaminated with a model strain (E. coli), one of the prevalent bacterial strains found in tissue infection, and then cultured with the ionic extracts of microcarriers (Fig. 8a). In case of regeneration of infected tissue, the body naturally reacts to disease-causing bacteria by increasing local blood flow along with the stimulated angiogenesis intrinsically and extrinsically for rapidly recruiting immune cells to destroy/uptake the bacteria [17]. Thus, preserving proangiogenic cells in the infected site and forming blood vessel together with angiogenic precursors collected from surrounding tissue are suggested as a promising therapeutic strategy, especially for severely infected tissue [18]. Because, the delay of clearance of pathogens and consequent lack of proangiogenic cells via accumulated toxins deteriorates proper angiogenesis and further tissue repair/regeneration in infected site. Therefore, simultaneous angiogenic and antibacterial biomaterials are required for preserving angiogenic precursors against toxins from bacterial overgrowth and accelerating angiogenesis. The co-culture of bacteria/mammalian cells has recently been employed to mimic the clinically relevant bacterial infection-associated tissue environment [33–35]. When mammalian cells are in contact with certain number of bacterial strain, they lose viability and biological functionality due to micro-organism
induced toxins (through cell-bacteria direct interaction or paracrine signaling) [47]. Therefore, killing bacteria species or scavenging their dead body that can act microorganism associated molecular pattern and cause severe inflammation and malfunction of host cells is considered an effective strategy to inhibit bacterial growth and ultimately to regain tissue regenerative capacity [48]. In this study, exterminating model bacteria strain was aimed to preserve the survivability and functionality of endothelial cells. From the co-culture test, the G2.5 group enabled more persisting endothelial cell viability than the control and G0 group under the bacterial contaminations (Fig. 8b). Of note, the VE-cadherin was expressed at significantly higher levels in G2.5 group than others, implying the endothelial cells with G2.5 have the potential to form tubular networks of blood vessels as this VE-cadherin is known as an essential anatomical structure of endothelial cells that can form blood vessels. While the potential to preserving endothelial survivability was decreased with increasing the bacteria number, the role of Co/Si-releasing microspheres was evident across the conditions investigated.

Taken the bacterial studies together, the Co/Si-releasing microcarriers were proven to inhibit bacterial growth either in direct or indirect contact (monoculture) and even to salvage the endothelial cell survivability and possible functional activity from the bacterial contamination (co-cultured). Although this study could demonstrate some in vitro efficacy of the microcarriers, with its limited in vitro conditions, the extrapolation to clinical situations may be ambitious, warranting more in vivo studies in infected tissue models.

4. Conclusion
Multifunctional Co/Si-containing microcarriers were designed to deliver dual therapeutic ions that can ultimately promote angiogenesis while inhibiting bacterial activity. The results demonstrated synergistic stimulation of endothelial cell functions such as angiogenic gene expression, polarization, migration and tubular formation, through the Co$^{2+}$ and Si$^{4+}$ ions co-delivered from microspheres. Moreover, the viability of both gram-positive and gram-negative model bacterial strains was strongly prohibited, and the survivability and cell-cell junction of endothelial cells were significantly salvaged under bactericidal conditions made in a co-culture model. Although more clinically relevant in
vivo studies are required further, this study highlights the effective roles of multiple ions co-delivered from biomaterials played in antibacterial-proangiogenic actions and suggests possible applications in the infected tissues that require both inhibition of bacterial activity and vascularized tissue regeneration.

Acknowledgement
This work was supported by the National Research Foundation of Korea (NRF) grant funded by the Korean government (MSIP; Ministry of Science, ICT & Future Planning, 2018R1A2B3003446) and Global Research Development Center Program (2018K1A4A3A01064257).

Statement
The researcher claims no conflicts of interest

Appendix A. Supplementary data

Supplementary data 1. (a) Inner and outer part of microsphere was visualized by SEM and (b) their composition was characterized by EDS (n=3). Inner structure was revealed after being fractured by compressive force. Dotted line in (a) was analysed by EDS and results were shown in (b). Equal composition is revealed between inner and outer part of each microsphere (1 or 2.5%). (c) XRD results present amorphous structure of the silica network and there is no influence of the glass network due to lower concentration of cobalt.

Supplementary data 2. Different amounts of microcarriers were evaluated for their effects on cell proliferation by CCK assay (*P < 0.05 & **P < 0.01, n=5). Ten milligrams of microcarriers was selected for concentration used in further biological assay and 5% was excluded among experimental groups.

Supplementary data 3. Live and dead staining of bacteria using direct co-culture
conditions with microcarriers. Fewer live bacteria species were detected in the area surrounding the 2.5% microcarriers for both the *E. coli* and *S. aureus* species (n=3). Representative images with moving bacteria are shown.

**Supplementary table 1.** Detail of in vitro experiment
References


[10] C. Wu, J. Chang, Multifunctional mesoporous bioactive glasses for effective


Cobalt promotes angiogenesis via hypoxia-inducible factor and protects tubulointerstitium in the remnant kidney model, Lab. Investig. (2005).

doi:10.1038/labinvest.3700328.


doi:10.1074/jbc.270.52.31189.


Kreikemeyer, R. Bader, Co-Culture of S. epidermidis and Human Osteoblasts on


Figure 1. Characterization of Si (0%) and Si-Co (1% or 2.5%) microcarriers. (a) Spherical morphology, size distribution, and their composition were characterized by SEM with EDS. (b) BET measurements showed comparable pore size distributions and surface areas. (c) Degradable microspheres presented controlled ion release, studied by ICP-AES. Data are presented as the mean ± SD (n=3). Sharp (#) indicates a statistically significant difference compared to value in 0% group.
**Figure 2.** Effects of microcarriers and free Co ions on angiogenic gene expression in HUVECs. Gene expression of VEGF, KDR and HIF1-α was quantified by qPCR after 24 h of co-culturing with microcarriers (10 mg/mL) via incorporated or free cobalt ions (6~18 ppm) for comparison. Significant upregulation of angiogenic gene expression was revealed in microcarrier groups in the order of w/o < 0% < 1% < 2.5%, similar to that from free Co ions (12~18 ppm). Different letters in each graph indicate significant differences between groups at the level of P<0.05 (n=3).
Figure 3. Enhanced HUVEC polarization and cell homing effects by ion attraction in microcarriers. (a) Stained HUVECs (green (actin) and blue (DAPI)) were captured by fluorescent microscopy after 4h of co-culture with microspheres (10 mg/mL), and the distance between the nucleus centre and the centre of mass was quantified by ImageJ based on the captured image. Polarization was enhanced in the order of w/o < 0 and 1% < 2.5% (P<0.05). (b) Cell homing effects (1h) were negatively shown by the number of DAPI (blue)-stained HUVECs in the top of the insert (8 µm pore size). Fewer cells remained in the 2.5% group than in other groups, indicating that more cells migrated via pores to microspheres (10 mg/mL). Cells that migrated to the microsphere were quantified based on initial cell numbers and captured images. The 2.5% group exhibited greater cell homing than other groups (P<0.05). Different letters in each graph indicate significant differences between groups at the level of P<0.05 (n=3).
Figure 4. Enhanced HUVEC migration in the scratch assay. (a) Optical images of migrated cells from each group at different time points (6, 12 and 24 h) and (b) their quantification. Cells that migrated to the scratched area were quantified per unit area (mm²), revealing an acceleration of cell migration under microsphere co-culture conditions (10 mg/mL) after mitomycin B pretreatment (2 h) in the order of w/o < 0% < 1% < 2.5%. Different letters in each graph indicate significant differences between groups at the level of P<0.05 (n=3).
Figure 5. Effects of microcarriers (10 mg/mL) on HUVEC tubular network formation. (a) Optical images of the tubular network of endothelial cells with different groups at different time points (6 and 12 h). (b) Node number and circle number were assessed from optical images. Of the experimental groups, the 2.5% group showed the most excellent ability to form tubular networks. Different letters in each graph indicate significant differences between groups at the level of P<0.05 (n=3). (c) Cell delivery potential of microcarriers. Schematic illustration of the procedure coating the microcarrier with Matrigel to deliver endothelial cells (HUVECs). F-actin (green) and nucleus-stained (blue) images of endothelial cells delivered on the microcarriers with SEM morphology after 24 h.
Figure 6. Ex vivo pro-angiogenic capacity from microcarriers (10 mg) in the CAM assay. (a) Representative images of two groups are presented after 4 days of implantation, and (b) pro-angiogenic effects were quantified by total tubular lengths, sizes and junctions, revealing upregulation in the 2.5% group compared to other groups. Different letters in each graph indicate significant differences between groups at the level of P<0.05 (n=3).
Figure 7. Indirect monoculture conditions using transwells between bacteria strains and microcarriers indicated antibacterial effects of Co ion-releasing microcarriers on both gram-negative (E. coli) and gram-positive (S. aureus) model bacteria. (a) Inhibition of bacterial (1 x 10^7/mL) viability after 4 h of co-culturing with microcarriers (10 mg/mL) using transwells in a PrestoBlue assay. (b) Images of live (green) and dead (red) cells confirmed bactericidal effects of 2.5% microcarriers, similar to the Co ion control. Different letters in each graph indicate significant differences between groups at the level of P<0.05 (n=5).
Figure 8. Co-culture of bacteria (E. coli) and HUVECs in media supplemented with extraction from microcarriers. More live HUVECs with less dead numbers were detected in the 2.5% compared to contr. (w/o) and 0%. A number of live and dead HUVEC cells was counted based on the colour and more live HUVECs with less dead numbers were significantly detected in the 2.5% compared to contr. (w/o) and 0% in two different E. coli CFU condition. VE-cadherin intensity was highly conserved in 2.5% while compromised VE-cadherin expression was visualized and quantified in w/o and 0% group (P<0.05). Representative images are shown. Different letters in each graph indicate significant differences between groups at the level of P<0.05 (n=5). Different letters in a graph of ‘number of cells per image’ revealing comparison of live cells numbers between groups.
Supplementary data 1. (a) Inner and outer part of microsphere was visualized by SEM and (b) their composition was characterized by EDS (n=3). Inner structure was revealed after being fractured by compressive force. Dotted line in (a) was analysed by EDS and results were shown in (b). Equal composition is revealed between inner and outer part of each microsphere (1 or 2.5%). (c) XRD results present amorphous structure of the silica network and there is no influence of the glass network due to lower concentration of cobalt.
Supplementary data 2. Different amounts of microcarriers were evaluated for their effects on cell proliferation by CCK assay (*P < 0.05 & **P < 0.01, n=5). Ten milligrams of microcarriers was selected for concentration used in further biological assay and 5% was excluded among experimental groups.

Supplementary data 3. Live and dead staining of bacteria using direct co-culture conditions with microcarriers. Fewer live bacteria species were detected in the area surrounding the 2.5% microcarriers for both the E. coli and S. aureus species (n=3). Representative images with moving bacteria are shown.
### Supplementary table 1. Detail of in vitro experiment

<table>
<thead>
<tr>
<th>Experiment</th>
<th>Cell number</th>
<th>Culture condition</th>
<th>Well</th>
<th>Number</th>
<th>Note</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cell viability</td>
<td>2.0 x 10⁴</td>
<td>72 h</td>
<td></td>
<td>n=5</td>
<td>Maximizing cytotoxicity</td>
</tr>
<tr>
<td>qPCR</td>
<td>1.0 x 10⁵</td>
<td>24 h</td>
<td>24 well (2.0 cm²)</td>
<td>n=3</td>
<td></td>
</tr>
<tr>
<td>Cell migration</td>
<td>1.0 x 10⁵</td>
<td>24 h of pre-culture + 6, 12 or 24h</td>
<td>24 well (2.0 cm²)</td>
<td>n=3</td>
<td></td>
</tr>
<tr>
<td>Cell polarization</td>
<td>5.0 x 10³</td>
<td>4 h of pre-culture + 1h</td>
<td></td>
<td>n=3</td>
<td>For single cell analysis</td>
</tr>
<tr>
<td>Ion attraction</td>
<td>5.0 x 10³</td>
<td>4 h of pre-culture + 1h</td>
<td>Insert (0.33 cm²)</td>
<td>n=3</td>
<td>Considering insert area and maximizing ion attraction</td>
</tr>
<tr>
<td>Tubular formation</td>
<td>3.0 x 10⁴</td>
<td>6 and 12 h</td>
<td>24 well on matrigel</td>
<td>n=3</td>
<td>Optimizing cell number</td>
</tr>
<tr>
<td>Cell-bacteria co-culture</td>
<td>2.0 x 10⁴</td>
<td>24 h of pre-culture + 6 h with bacteria</td>
<td>96 well (0.33 cm²)</td>
<td>n=3</td>
<td>Optimizing cell number</td>
</tr>
</tbody>
</table>