Therapeutic effects of placenta derived-, umbilical cord derived-, and adipose tissue derived-mesenchymal stem cells in chronic Helicobacter pylori infection: comparison and novel mechanisms

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(Rceived 7 September, 2020; Accepted 13 December, 2020)

Supported with significant rejuvenating and regenerating actions of mesenchymal stem cells (MSCs) in various gastrointestinal diseases including Helicobacter pylori (H. pylori)-associated gastric diseases, we have compared these actions among placenta derived-MSCs (PD-MSCs), umbilical cord derived-MSCs (UC-MSCs), and adipose tissue derived-MSCs (AD-MSCs) and explored contributing genes implicated in rejuvenation of H. pylori-chronic atrophic gastritis (CAG) and tumorigenesis. In this study adopting H. pylori-initiated, high salt diet-promoted gastric carcinogenesis model, we have administered three kinds of MSCs around 15–18 weeks in H. pylori infected C57BL/6 mice and sacrificed at 24 and 48 weeks, respectively, in order to either assess the rejuvenating capability or anti-tumorigenesis. At 24 weeks, MSCs all led to significantly mitigated atrophic gastritis, for which significant inductions of autophagy, preservation of tumor suppressive 15-PGDH, attenuated apoptosis, and efficient effectorcytosis was imposed with MSCs administration during atrophic gastritis. At 48 weeks, MSCs administered during H. pylori-associated atrophic gastritis afforded significant blocking the progression of CAG, as evidenced with statistically significant reduction in H. pylori-associated gastric tumor (p<0.05) accompanied with significant decreases in IL-1f, COX-2, STAT3, and NF-κB. Combined together with the changes of stanniocalcin-1 (STC-1), thrombospondin-1 (TSP-1), and IL-10 known as biomarkers reflecting stem cell activities at 48 weeks after H. pylori, PD-MSCs among MSCs afforded the best rejuvenating action against H. pylori-associated CAG via additional actions of effectorcytosis, autophagy, and anti-apoptosis at 24 weeks. In conclusion, MSCs, especially PD-MSCs, exerted rejuvenating actions against H. pylori-associated CAG via anti-mutagenesis of IL-10, CD-36, ATG5 and cancer suppressive influences of STC-1, TSP-1, and 15-PGDH.

Key Words: chronic atrophic gastritis, H. pylori, STC-1, TSP-1, mesenchymal stem cells

International Agency for Research on Cancer of World Health Organization defined H. pylori as class I carcinogen based on facts that H. pylori caused gastric carcinogenesis,(1) by which, reversely evidenced, the eradication of H. pylori could prevent metachronous gastric cancer after endoscopic resection of early gastric cancer. (2) If this is true, the eradication can be solution for prevention of gastric cancer, but intervention trials dealing with gastric cancer prevention by H. pylori eradication still need more evidences and confront additional risk of bacterial resistance. (3)

Therefore, non-microbial dietary or nutritional intervention has been considered as either alternate to eradication or the mechanistic provision of surrounding break up to clear mutagenic inflammation either capable of blocking field cancerization process or reverting into non-atrophic condition from precancerous atrophic gastritis. (4) Since the most human gastric cancers develop after long-term H. pylori infection according to the Professor Correa P’ pathway that the progression from chronic gastritis via atrophy and intestinal metaplasia to dysplasia or cancer, the strategy to detox from precancerous chronic atrophic gastritis (CAG) into non-atrophic condition can be another confidential hope for cancer prevention, though it still remains unclear whether CAG is a direct precursor of gastric cancer or merely a marker of high cancer risk. (5–7) In real world, antioxidants, anti-inflammatory drugs, and food therapy may contribute in the regression of CAG, especially better when used simultaneously with eradication therapy, “no biomarker for point of no return” or “no optimal timing for intervention” still remains obscure for clinical practice and should be further investigated. (8)

In this status, anticipation was paid to cell therapy, stem cells featured with self-renewal, cell proliferation, differentiation into specialized cell types, because they can provide the chance to revert protumor condition of gastric atrophy. Though the term “rejuvenation” is defined as the action or process of making someone or something look or feel better, younger, or more vital, in H. pylori-associated Correa P’s definition, the old hypothesis that CAG-intestinal metaplasia (IM)-dysplasia-carcinoma sequence because H. pylori infection may also trigger an autoimmune gastritis of the corpus mucosa, with CAG and IM, reaching to gastric cancer, (9,10) it means reverting into non-atrophy from atrophy is the way of escaping from the protumor condition and evidences that since severe CAG remained in the adjacent mucosa of the gastric cancer even after H. pylori eradication linked to gastric carcinogenesis, simple removal of carcinogen H. pylori limited the efficacy. (11–15)

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doi: 10.3164/jcbn.20-151
J. Clin. Biochem. Nutr. | Published online: 27 March 2021 | 1–15
In recent publications that we have shown newly rejuvinating and restorative actions of placenta derived-mesenchymal stem cells (PD-MSCs) against gastric damaging conditions such as \textit{H. pylori}-associated CAG, radiation-induced gastrointestinal (GI) injuries, GI damages after ischemia-reperfusion, and NSAIDs-induced GI damages\cite{15,16}, the curiosity emerged whether MSCs originated from other sources, e.g., umbilical cord derived (UC-MSCs) and adipose tissue-derived (AD-MSCs), bone marrow-derived MSCs, and other sources derived-MSCs are differed in these regenerative actions and search for additional beneficiary actions mechanisms beyond proliferative and restorative actions of stem cells was raised. In this study, we have compared the efficacy and mode of action according to origin of MSCs against chronic \textit{H. pylori}-associated CAG and gastric tumorigenesis model.

\textbf{Material and Methods}

\textbf{Cell culture.} PD-MSCs, UC-MSCs, and AD-MSCs were all provided from CHA University (Prof. Yong Soo Choi, CHA University, Seongnam, Korea). The MSCs line was cultured in \textit{α}-MEM medium containing 1 \textmu g/ml heparin, 25 \textmu g/ml fibroblast growth factor-4, 10\% (v/v) fetal bovine serum and 100 U/ml penicillin. Cells were maintained at 37°C in a humidified atmosphere containing 5\% CO_2.

\textbf{H. pylori culture.} \textit{H. pylori} strain ATCC43504 (American Type Culture Collection, \textit{cagA}+ and \textit{vacA} s1-m1 type strain) was used for \textit{in vitro} cell model and Sydney strain (SS1, a \textit{cagA}+,- \textit{vacA} s2-m2 strain adapted for mice infection) for \textit{in vivo} model (Fig. 1A and 4A). \textit{H. pylori} were cultured at 37°C in BBL Trypticase soy (TS) agar plate with 5\% sheep blood (TSAB); BD Biosciences, Franklin Lakes, NJ) under microaerophilic condition (BD GasPaK EZ Gas Generating Systems; BD Biosciences) for 3 days. The bacteria were harvested in clean TS broth, centrifuged at 3,000 \times g for 5 min, and resuspended in the broth at a final concentration of 1 \times 10^8 colony-forming units (CFUs)/ml. In all experiments, cultures grown for 72 h on TS agar plates were used.

\textbf{Animals and study protocol; \textit{H. pylori}-infected mice model.} Experimental protocol. Five-week-old male C57BL/6 mice (WT mice) were purchased from Orient (Seoul, Korea) and they were housed in a cage maintained in a 12 h/12 h of light/dark cycle under specific pathogen-free conditions (n = 120). They were fed sterilized commercial pellet diets (AIN-76A pellet diet, Biogenomics, Seoul, South Korea) and sterile water \textit{ad libitum}, and housed in an air-conditioned biohazard room at a temperature of 24°C. We divided mice into four groups: Group 1 (n = 20); WT mice as vehicle control group, Group 2 (n = 40); WT mice as \textit{H. pylori}-infected disease group, Group 3 (n = 10); WT mice as \textit{H. pylori}-infected disease group administrated with 1 \times 10^7/100 \mu l PD-MSCs, Group 4 (n = 20); WT mice as \textit{H. pylori}-infected disease group administrated with 1 \times 10^7/100 \mu l UC-MSCs; Group 4 (n = 20), and WT mice as \textit{H. pylori}-infected disease group administrated with 1 \times 10^7/100 \mu l AD-MSCs, half of all mice were sacrificed at 24 weeks and the remaining were sacrificed at 48 weeks, respectively. In detail, all groups were given intraperitoneal (i.p.) injections of pantoprazole, 20 mg/kg (Amore-Pacific Pharma, Seoul, Korea) as proton pump inhibitor, three times per week, to increase successful \textit{H. pylori} colonization through lowered gastric acidity, then, each mouse was intragastrically inoculated with a suspension of \textit{H. pylori} containing 10^8 CFUs/ml or with an equal volume (0.1 ml) of clean TS broth using gastric intubation needles. The \textit{H. pylori}-infected mice were fed a special pellet diet based on AIN-76A containing 7.5\% NaCl high salt diet (Biogenomics, Seongnam, Korea) for total 36 weeks (Fig. 1A and 4A) to promote \textit{H. pylori}-induced carcinogenic process in all infected animals. Randomized groups of mice (n = 10) sacrificed at 36 weeks of post \textit{H. pylori} infection, respectively based on our previous experience\cite{28,29} that CAG was generated at 24 weeks and gastric tumorigenesis was generated after 48 weeks. The body weight was checked in all mice every 3 days up to observational periods. The stomachs of mice were opened along the greater curvature and washed with ice cold PBS. The numbers of either erosions/ulcers or protruded nodules/masses were determined under the magnified photography (Fig. 1C and 4C). Stomachs were isolated and subjected to a histologic examination, ELISA, Western blotting, and RT-PCR. All animal studies were carried out in accordance with protocols approved by the Institutional Animal Care and Use Committee (IACUC) of CHA University CHA Cancer Institute after institutional review board of IACUC approval (IRB #17-1001).

\textbf{Gross lesion index.} After sacrificing the mice, the isolated stomachs were open along the greater curvature and washed in ice-cold saline. To investigate the degree of gross mucosal pathology, the mucosal sides of the stomachs were photographed using a digital camera and part of the mucosa was immediately fixed with 10\% formalin solution. The gross damage of the gastric mucosa was assessed by three gastroenterologists, who were blinded to the treatments, using a gross ulcer index\cite{30}.

\textbf{Index of histopathologic injury.} For histopathological analysis, the stomach was fixed in 10\% neutralized buffered formalin, processing using the standard method and embedded in paraffin. Sections of 4 \mu m thickness were then stained with hematoxylin and eosin. The glandular mucosae of corpus and antrum were examined histologically. The pathological changes of \textit{H. pylori}-infection, such as inflammatory cells infiltration, erosive lesions, ulceration, dysplasia, adenoma formation (precancerous lesion), were graded by three gastroenterologists (KB Hahn, JW Yoo, and JM Kim), who were blinded to the group, using an index of histologic injury defined. In this study, inflammation was defined as grade the infiltration of inflammatory cells, 0: none, 1: under the lamina propria, 2: half of mucosa 3: until the epithelial gland layer (all mucosa). The erosion was defined as proportion of erosive lesion, 0: none, 1: loss of epithelial gland layer (1/3 proportion), 2: two-three proportion of mucosa (2/3 proportion) 3: all mucosa (3/3 proportion).

\textbf{Immunohistochemical staining.} Immunohistochemistry was performed on replicate sections of mice gastric tissues. After deparaffinization, slides were dewaxed and rehydrated with graded alcohol, and boiled three times in 100 mM Tris buffered saline (pH 6) with 5\% urea in an 850 W microwave oven for 5 min each. And then, cooling in water for 15 min and washed in PBS, and slides were incubated overnight with the primary antibody at 4°C. Antibodies: F4/80 (1:500; eBioscience, San Diego, CA) or 15-PGDH (1:300; Dako, Santa Clara, CA) or Ki-67 (1:300; Santa Cruz Biotechnology, Santa Cruz, CA) in the presence of 1.0\% bovine serum albumin respectively. Slides incubated with secondary antibody (1:300) for 1 h at room temperature, and then with 40-6-diamidino-2-phenylindole (DAPI, 100 ng/ml) for 1 min at room temperature. And finally the slides were counterstained with hematoxylin (Sigma-Aldrich, St. Louis, MO). After incubation, a subsequent reaction was formed using a Vector kit (Vector Laboratories, Inc., Burlingame, CA). Finally, the slides were incubated with 3,3'-diaminobenzidine (Invitrogen Life Technologies, Carlsbad, CA) and counterstained with hematoxylin (Sigma-Aldrich).

\textbf{Terminal deoxynucleotidyl transferase-mediated dUTP nick-end labeling (TUNEL) staining.} Apoptosis was visualized using a terminal deoxynucleotidyl transferase (TdT) FRAGel DNA fragmentation detection kit (Oncogene Research Products, La Jolla, CA). To determine the apoptotic index in each group, TUNEL immune-stained sections were scanned under low-power magnification (\times100) to locate the apoptotic hotspots.

\textbf{RT-PCR.} Total RNA was isolated using the Trizol (Invitrogen, Carlsbad, CA). Trizol was added to 1.5 ml tube, which

\textbf{DOI:} 10.3164/cbn.20-151
A. Scheme for group, Group 1: normal control, Group 2: H. pylori-associated CAG disease control, Group 3: disease control treated with 1 × 10^6 cells/100 ml PD-MSCs, Group 4: disease control treated with 1 × 10^6 cells/100 ml UC-MSCs, and Group 5: disease control treated with 1 × 10^6 cells/100 ml AD-MSCs. (B) Body weight changes according to group. 24 weeks H. pylori infection (control group) showed significant decreases in mean body weights. (C) Representative photo of resected gross stomach according to group and mean gross lesion scores according to group. (D) Representative pathology of Group 2 showing CAG with some erosive changes and mean pathological scores according to group.

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1) Proton pump inhibitor (Pantoprazole): 20 mg/kg, intraperitoneal (i.p.) injection, three times
2) H. pylori: Sydney Strain 1 (SS1), 1.0 × 10^9 CFU/200 µl, broth administered via oral tube, 4 times
3) C57BL/6 mice (4~5 week-old age, n=10-20)
4) HSD (High salted diet); AIN-76A pellet diet with 7.5% salts
5) PD-MSCs, human placenta derived-mesenchymal stem cells, 1.0 × 10^6 cells/100 µl, 10 times, iv tail vein
6) UC-MSCs, human umbilical cord derived-mesenchymal stem cells, 1.0 × 10^6 cells/100 µl, 10 times, iv
7) AD-MSCs, human adipose tissue derived-mesenchymal stem cells, 1.0 × 10^6 cells/100 µl, 10 times, iv
were then incubated 10 min at 4°C and gently mixed with 100 µl chloroform (Merck, Rahway, NJ). After incubation for 10 min in ice, samples were centrifuged at 10,000 g for 30 min. Supernatants were extracted and mixed with 200 µl isopropanol (Merck), and mixtures were incubated at 4°C for 1 h. After centrifuging at 13,000 g for 30 min, pellets were washed with 70% (v/v) ethanol. After allowing the ethanol to evaporate completely, pellets were dissolved in 40 µl diethylene pyrocarbonate-treated water (Invitrogen Life Technologies). cDNA was prepared using reverse transcriptase originating from Murine Moloney leukemia virus (Promega, Madison, WI), according to the manufacturer’s instructions. The polymerase chain reaction (PCR) was performed over 25 cycles of: 94°C for 20 s, 58.5°C for 30 s, and 72°C for 45 s. Oligonucleotide primers were purchased from Bioneer (Daejeon, Korea). Oligonucleotide primers were as follows: for COX-2, sense 5′-GAA ATG GCT GCA GAG TGG GA-3′; for IL-β, sense 5′-GAG CAG CGG AGA TGA ACA AAA-3′ and antisense 5′-TGG GGA ACT CAT GAG ACT AAA-3′, for IL-8, sense 5′-GGG GCT TTG TCG TGC AAT AA-3′ and antisense 5′-GCA CAG GAG TGA GCA AAA A-3′, for IL-6, sense 5′-AGG GGA TTT CCA GAG AAG CC-3′ and antisense 5′-TGG ATG GTT GTC TGT CTT AG-3′; for tumor necrosis factor-alpha (TNF-α), sense 5′-ATG AGC ACA GAA AGG ATG ATC-3′ and antisense 5′-TAC AGG CTT GAG ACT CAT CGA ATT-3′, for IL-6, sense 5′-GGG GAT GAT GCT GTG GAC AA-3′ and antisense 5′-TTA CAG CAT ACG TGT TTA GGC GA-3′, for IFN-γ, sense 5′-ACA ATG AAG CCA GCT ACC AC-3′ and antisense 5′-TCA AAC TTT GCA GTC ATA CTC AT-3′, for IL-8, sense 5′-GGG GCT TTG TCG TGC AAT AA-3′ and antisense 5′-GCA CAG GAG TGA GCA AAA A-3′, for IL-6, sense 5′-GAG CAG CGG AGA TGA ACA AAA-3′ and antisense 5′-TGG GGA ACT CAT GAG ACT AAA-3′, for VEGF, sense 5′-CCC TTC CTC ATC TTC CTC TC-3′ and antisense 5′-CAC GTA CTA TGG GAG AGG G-3′, for IL-10, sense 5′-CCA GAT TTA CCT GGT AGA AG-3′ and antisense 5′-AGG TCG TGG AGT CCA GAC TC-3′, for IL-18, sense 5′-CTC CCC ACC TAA CTT GTA TG-3′ and antisense 5′-CCA GGA ACA ATG GCT GCC AT-3′, for TSP-1, sense 5′-GTT GCA TGT GTG TGG AAG CAC C-3′ and antisense 5′-ACC ACA ATG CAG ATC TGG CCA G-3′, for STC-1, sense 5′-TCT CTT GTG GAG TGC GTT-3′ and antisense 5′-GTC TTC CTC GGC ATT CGG-3′, for CD36, sense 5′-ACT CCA GAA CCC AGA CAA CCA C-3′ and antisense 5′-ACC AAG TAA GAC CAT CTC AAC CAG C-3′, for LRPI, sense 5′-GAG GAG TGC TGG GTG TAT GGC AC-3′ and antisense 5′-GAT GCC TGG TTG GAT GGT C-3′, for TGF-β, sense 5′-TGA GTG GCT GTC TTT GTA CG-3′ and antisense 5′-TCT CTG TGG AGC TGA AGC AA-3′, for HSP27, sense 5′-TGC CCT TCT CCC TAC TGC GG-3′ and antisense 5′-TCC AAT TGG GGC AGG GCC CT-3′, for bFGF, sense 5′-TAT GAA GGA AGA TGG AGG GC-3′ and antisense 5′-AAC AGG ATG AGC TTC TCT GC-3′, and for GAPDH, sense 5′-GGT GCT GAG TAT GTC GTG GA-3′ and antisense 5′-TTC AGC TCT GGG ATC ACC TT-3′.

**Western blotting.** Cells or resected gastric tissues were harvested and lysed in lysis buffer (Cell Signaling Technology) containing 1 mM phenylmethylsulfonyl fluoride (PMSF; Sigma Aldrich). After 30 min of incubation, samples were centrifuged at 12,000 g for 15 min 4°C. The supernatants were then collected and protein quantification was carried out with a Bio-Rad protein assay. Equal amounts of soluble protein (30 µg) were denatured by heating at 100°C for 3 min. Proteins were separated by sodium dodecyl sulphate-polyacrylamide gel electrophoresis (SDS-PAGE) and transferred to polyvinylidene fluoride membranes. The membranes were blocked in 5% BSA in PBST for 30 min. And then, the membranes probed initially with specific primary antibody, washed, incubated with peroxidase-conjugated secondary antibodies, and rewashed. The protein bands were detected by chemiluminescence (Supersignal; Pierce) exposure on chemiluminescence system (GE Healthcare, Buckinghamshire, UK). The general procedure for Western blot analysis of cultured mouse gastric mucosal cells was similar to the procedures described above. Antibodies used in the current study were cyclooxygenase 2 (COX-2), purchased from Thermogene, β-actin purchased from Sigma-Aldrich Biotechnology, 15-hydroxyprostaglandin dehydrogenase (15-PGDH), purchased from Cayman. Primary antibody against β-actin was purchased from Sigma-Aldrich Co., antibodies for lamin B from Santa Cruz Biotechnology, other antibodies for p-signal transducer and activator of transcription 3 (STAT3), total STAT3 from Cell Signaling Technology (Beverly, MA), horseradish peroxidase (HRP)-conjugated secondary antibody from Pierce Biotechnology (Rockford, IL). DL-dithiothreitol (DTT), TRIZol™, 4',6-diamidino-2-phenylindole (DAPI) from Invitrogen (Carlsbad, CA), and polyvinylidene difluoride (PVDF) membranes were supplied from Gelman Laboratory (Ann Arbor, MI). The ECL chemiluminescent detection kit was purchased from LPS solution (Daejeon, South Korea) and protein assay dye (Bradford) reagent was supplied by Bio-Rad Laboratories (Hercules, CA). Biinchonic acid (BCA) protein assay reagent was purchased from PierceBiotechnology (Rockford, IL). COX-2 nitric oxide synthase (iNOS), cytochrome c, survivin antibodies were purchased from Santa Cruz Biotechnology (Dallas, TX), phosphorylated STAT3, Bax, B-cell lymphoma 2 (Bel-2), cleaved caspase-3, cleaved caspase-8, poly-ADP-ribose polymerase (PARP), and Musashi-1 all from Cell Signaling Technology (Danvers, MA).

**Preparation of cytosolic and nuclear extracts.** After *H. pylori* infection, resected stomach tissues were washed twice with ice-cold 1× PBS and scraped in 1 ml of PBS, followed by centrifugation at 1,700 × g for 5 min at 4°C. Pellets were resuspended in hypotonic buffer A [10 mM N-2-hydroxyethylpiperazine-N'-2-ethanesulfonic acid (pH 7.9), 1.5 mM MgCl₂, 10 mM KCl, 0.5 mM DTT and 0.2 mM phenylmethylsulfonylfluoride (PMSF)] for 15 min on ice. Ten % Nonider P-40 was then added to final concentration of 0.1% for less than 3.5 min. The mixture was then centrifuged at 6,000 × g for 5 min at 4°C. Supernatant was collected as the cytosolic extract and stored at −80°C. The pellets were washed twice with hypotonic buffer A and resuspended again in hypertonic buffer C [20 mM N-2-hydroxyethylpiperazine-N'-2-ethanesulfonic acid (pH 7.9), 20% glycerol, 420 mM NaCl, 1.5 mM MgCl₂, 0.2 mM ethylenediaminetetraacetic acid, 0.5 mM DTT and 0.2 mM PMSF] for 1 h on ice and centrifuged at 18,000 × g for 15 min at 4°C. The supernatant containing nuclear proteins was collected and stored at −80°C. The protein concentrations of both fractions were determined by using the BCA protein assay reagent.

**Statistical analysis.** Results are expressed as the mean (SD). Statistical analyses were conducted with GraphPad Prism (GraphPad Software, La Jolla, CA) and SPSS software (ver. 12.0; SPSS Inc., Chicago, IL). Statistical significance between groups was determined by a multi-variate test, Kruskal-Wallis test. Differences between groups were evaluated using the paired-sample t test. Significance was set at p<0.05 and p<0.01, respectively, in two-tailed testing.

**Results**

**Comparison of the efficacy of PD-MSCs, UC-MSCs, and AD-MSCs against *H. pylori*-associated CAG; 24 weeks results.**

*Gross and pathological scores after MSCs administration.* Our group have established excellent animal model of *H. pylori*-induced CAG in mice as shown in Fig. 1A that after we have injected proton pump inhibitor (PPI), 20 mg/kg pantoprazole, to lower gastric acidity in order to facilitate *H. pylori* colonization
infiltrations after control weight, infection. Reduced
J.M. Park staining control subjected to pathological intraperitoneally. Relevant mice via showing were derived-mesenchymal cells, multiple weeks derived-mesenchymal stem cells, and AD-MSC (human adipose tissue-derived-mesenchymal stem cells) were delivered 10 times via mice tail vein during 15–20 weeks of *H. pylori* infection. The mice were subjected to sacrifice 24 weeks after *H. pylori* infection. In order to check exact colonization of *H. pylori*, six weeks after *H. pylori*, mice (*n = 5*) were randomly killed to confirm the successful colonization of *H. pylori* and all the mice tested were proven to be either positive CLO (rapid urease test) or positive Giemsa staining and control mice (*n = 5*) were subjected to pathological evaluation around 12–15 weeks in order to check the development of CAG. Since one of clinical manifestation of *H. pylori* infections denoting CAG development according to our previous study was the significant loss of body weight, we measured all body weight of mice. As shown in Fig. 1B, the mean body weight of Group 2, *H. pylori* alone infection control group, were significantly decreased compared to normal control (*p < 0.05*). When the mean body weights were compared between Group 2 and Group 3, Group 3 treated with PD-MSCs, were significantly different (*p < 0.05*), while lesser body weight reduction was seen in Group 4 and Group 5, but no statistical significance, signifying PD-MSCs among three kinds of MSCs significantly rejuvenated *H. pylori* infection-associated CAG. As seen in Fig. 1C, significant gross changes were noted in Group 2, showing irregular gastric surface, edematous gastric wall, thinned gastric wall, small protuberant gastric surfaces with erythematous and erosive mucosa. Based on scoring system, gross lesion index was significantly increased in Group 2 compared to Group 1 (*p < 0.01*), while these scores were significantly decreased in Group 3, Group 4, and Group 5 compared to Group 2 (*p < 0.05*, Fig. 1C). On Fig. 1D, representative pathology from Group 2 was presented, showing the development of moderate degree of CAG with cryptic gland loss with significant loss of parietal cells, multiple gastric erosions, and marked inflammatory cell infiltrations on to mucosal and submucosal area. Before separating pathological scores of gastric inflammation, atrophy, and ulceration, mean pathological scores according to group was shown in Fig. 1D, significant amelioration of pathological scores were noted with MSCs administration (*p < 0.01*).

DeCREASED impassory mediators after MSCs administration. Supported with the significant alleviation of pathological scores relevant to gastric inflammation after MSCs (Fig. 1D), we have measured the changes of TNF-α, IL-6, IFN-γ, IL-8, IL-1β, VEGF, and IL-10 mRNA via RT-PCR, all reported to be major mediators relevant to *H. pylori* infection. As seen in Fig. 2A, dissecting pathological changes according to group, Group 2 showed significant increases in these scores of gastric inflammation, while inflammatory scores were significantly decreased in group administered with MSCs (*p < 0.01*). With immunohistochemical staining with F4/80 to denote macrophage infiltration according to group since macrophages are responsible for inflammation after *H. pylori* infection, as seen in Fig. 2B, significantly increased F4/80 scores were noted in Group 2 (*p < 0.001*), but significantly decreased with MSCs administration (*p < 0.05*). As inflammatory mediators, TNF-α, IL-6, IFN-γ, IL-8, IL-1β, and VEGF mRNA were measured and compared according to group (Fig. 2C). All of these inflammatory mediators were significantly decreased after MSCs, PD-MSCs were best among MSCs. IL-10 as anti-inflammatory cytokines was significantly decreased in Group 2, but significantly increased in Group 3 and Group 5. Cox-2 and iNOS was significantly increased in Group 2, but these expressions were decreased in Group 3 (Fig. 2D) and nuclear translocation of STAT3 and NF-κB was significantly increased in Group 2, but significantly decreased in Group 3 and Group 5. All of these findings suggested significant anti-inflammatory actions of MSCs, especially better in PD-MSCs against *H. pylori* infection. As seen in Fig. 2E, PD-MSCs increased genes implication in effecorcytosis, CD-36, LDL receptor related protein 1 (LRP1), and IL-10 mRNA and the measurement of effecorcytosis using Jurkat T cells and Raw cells were shown, showing effecorcytosis were operated in the presence of PD-MSCs. Decreased cell death (apoptosis), but enhanced autophagy after MSCs administration. Supported with the significant alleviation of pathological scores relevant to gastric erosions and ulcers after MSCs (Fig. 1D), as seen in Fig. 3A, the mean scores of gastric erosions/ulcers according to group, these ulcer scores were significantly increased in Group 2, while the scores were significantly decreased in group treated with MSCs (*p < 0.05*). These erosions/ulcers scores were significantly well correlated with apoptotic index (Fig. 3B) and Bax expressions (Fig. 3C) according to group (*p < 0.01*), signifying anti-apoptotic actions after MSCs contributed to lower scores of gastric erosions/ulcers. The curiosity about autophagy arose because autophagy has been acknowledged as survival mechanisms against *H. pylori* infection. As seen in Fig. 3D and E, immunohistochemical staining of LC3B was performed to measure the autophagy phenomenon according to group. As results, mean expression of LC3B was significantly increased in Group 3, though significantly increasingly expressed in Group 2 (*p < 0.05*). These immunohistochemical staining of LC3B were further validated by Western blots. As seen in Fig. 3E, ATG5 and LC3B II were significantly increased in Group 3 (*p < 0.01*). Supported with pathological scoring such as lesser erosions/ulcers, we have traced the expressions of 15-PGDH, known as gene responsible for tumor suppressor and regeneration biomarker, and we found the expressions of 15-PGDH were significantly decreased in Group 2, but the expressions of 15-PGDH were significantly preserved in Group 3, Group 4, and Group 5, all treated with MSGs (Fig. 3E). These findings from Western blot were validated with immunohistochemical staining of 15-PGDH (data not shown, but immunohistochemical staining of 15-PGDH at 48 weeks was shown in Fig. 6C), showing the expressions of 15-PGDH were significantly decreased at 24 weeks of *H. pylori* infection, but their expressions were significantly preserved in Group treated with MSGs (*p < 0.05*, Fig. 3E).

Preventive effects of three kinds of MSCs against *H. pylori*-associated gastric tumorigenesis; 48 weeks results. Gross and pathological scores. As described before, our models, when sacrificed at 48 weeks of *H. pylori* infection (Fig. 4A), they developed significant gastric tumorigenesis as seen in Fig. 4C, multiple, various sized, scattered nodular masses were noted under the background of CAG. Gross and pathological lesion scores were significantly increased in Group 2 (*p < 0.005*), but the mean scores were significantly decreased in Group 3, Group 4, and Group 5, signifying MSCs administered during atrophic gastritis significantly blocked the progression in to gastric tumorigenic process (*p < 0.001*, Fig. 4C). Among pathological changes, atrophic gastritis, gastric mucosal erosions, gastric ulcers, gastritis cystica profunda, and tumorigenesis (gastric adenoma and gastric carcinoma), as seen in Fig. 5A and B, MSCs administration significantly mitigated these gastric tumorigenises.
Fig. 2. Changes of inflammatory mediators according to group, 24 weeks. (A) Mean pathological scores focused on inflammation according to group. (B) Mean immunohistochemical staining of F4/80 macrophagy denoting antibody. (C) RT-PCR for inflammatory mediators including TNF-α, IL-6, IFN-γ, IL-8, IL-1β, VEGF, and IL-10 mRNA. Right bar graph shows mean relative intensity of TNF-α, IL-6, and IFN-γ according to group (triplicate experiments). (D) Western blot for COX-2 and iNOS in whole extracts from each group. (E) Western blot for STAT3 and NF-κB in nuclear fractions of obtained tissues. (F) RT-PCR for efferocytosis engaged genes, CD-36, LRPI, IL-10, and TGF-β mRNA after PD-MSCs administration in different times, 12 h and 24 h. Right graph shows real accomplishment of efferocytosis with PD-MSCs administration.

doi: 10.3164/jcbn.20-151
Significant mitigation of H. pylori-associated inflammation with MSCs. Findings from previous in vivo model showing that MSCs significantly attenuated inflammatory condition led us to measure the changes of tumorigenesis-associated signaling in chronic H. pylori infection such as inflammasome, redox sensitive transcription factor, and IL-6 dependent STAT3.
**Fig. 4.** Influence of three kinds of MSCs, PD-MSCs, UC-MSCs, and AD-MSCs on *H. pylori*-initiated, high salt diet-promoted gastric tumorigenesis (48 weeks). (A) Scheme for group, Group 1: normal control, Group 2: *H. pylori*-associated CAG disease control, Group 3: disease control treated with 1 × 10^6 cells/100 μl PD-MSCs, Group 4: disease control treated with 1 × 10^6 cells/100 μl UC-MSCs, and Group 5: disease control treated with 1 × 10^6 cells/100 μl AD-MSCs was extended up to 48 weeks to compare the efficacy of anti-tumorigenesis via the administration of MSCs during 15–18 weeks of *H. pylori* infection. (B) Body weight changes according to group. (C) Representative photo of resected gross stomach according to group showing gross and pathological lesion scores according to group. Lower bar shows mean changes of scores according group, lower left gross lesion score and right pathological scores.
pathway. Since the expressions of NOD-, LRR-, and pyrin domain-containing protein 3 (NLRP3), apoptosis-associated speck-like protein containing caspase recruitment domain (ASC), IL-1β, component of inflammasome after H. pylori infection, 50 MOI H. pylori infection for 24 h led to significant induction of NLRP3, ASC, and IL-1β mRNA in in cell model (data not shown). As repeated in pathological scores according to group, MSCs-treated group showed significant decreases in gastric tumorigenesis (Fig. 5A), the oncogenic signals including COX-2, p-STAT3, and NF-κB were significantly increased in Group 2, but these levels were all significantly decreased in Group 3, Group 4, and Group 5, consistently showing anti-mutagenic action of MSCs administered during protumor CAG condition (Fig. 5B). Though debatable in its significance in GI carcinogenesis, inflammasome including IL-1β and IL-18 were all significantly decreased in MSCs administered group compared H. pylori control group (48 weeks).

Significant induction of tumor suppressive 15-PGDH with MSCs as anti-tumorigenesis mechanisms. 15-PGDH is gene showing significant anti-tumorigenesis in colon cancer as well as other GI cancers. Already in 24 weeks model, MSCs showed significant induction of 15-PGDH in chronic H. pylori infection and these changes were more prominently seen in 48 weeks model, as shown in Fig. 6 (Western blot according to group in Fig. 6B and immunohistochemical staining in Fig. 6C), signifying the significant induction of 15-PGDH might be one of critical biomarkers denoting rejuvenating and anti-mutagenic action of MSCs.

Significant induction of TSP-1 and STC-1 to restore H. pylori-associated gastric damages; rejuvenation with MSCs administration. Noted from other including our investigations that stem cells imposed significant regenerating and rejuvenating actions through the induction of thrombomodulin-1 (TSP-1) and sanniocalcin-1 (STC-1) in multiple cases of tissue damages, we have measured the expressions of TSP-1 and STC-1 after PD-MSCs in gastric epithelial cells, RGM-1 normal gastric mucosal cells. As seen in Fig. 7A, PD-MSCs significantly induced TSP-1 (from 3 h) and STC-1 (from 12 h) mRNA (p<0.01), which was noted in cell number dependent way under H. pylori infection (Fig. 7B). In order to imply the significance of these two
genes under *H. pylori* infection, we have checked cell viability under *H. pylori* infection (100 MOI, 24 h). As expected, *H. pylori* infection for 24 h led to significant reduction in cell viability assessed by MTT assay (*p*<0.01, Fig. 7C), but the presence of PD-MSCs (2 × 10⁶ cells) significantly preserved cell viability even under *H. pylori* infection (*p*<0.001). However, as seen in Fig. 7C, these rescuing actions of PD-MSCs were significantly abolished in case of either TSP-1 or STC-1 siRNA-transfected cells (*p*<0.05). Under same condition, we checked the expressions of TSP-1 and STC-1 mRNA and their expressions were significantly increased. Since *H. pylori* infection led to cytotoxicity via apoptosis, when we measured apoptotic executors, Bax, cleaved caspase-3, and cleaved PARP, *H. pylori* infection led to increases of these executor expressions, but PD-MSCs significantly decreased apoptotic executors. However, in cells transfected with either TSP-1 siRNA (Fig. 7D) or STC-1 siRNA, apoptotic executors were not decreased even after PD-MSCs. Another cytotoxic mechanism of *H. pylori* is through increased oxidative stress. On flow cytometric analysis for oxidative stress, *H. pylori* infection led to significant elevations of DCF-DA expressions (*p*<0.01), but ablated condition of either TSP-1 or STC-1 led to increased oxidative stress (Fig. 7E), combining together, led to conclusion that TSP-1 or STC-1 with MSCs, especially, PD-MSCs contributed to significant restoring action on *H. pylori*-associated CAG. These in vitro findings regarding TSP-1 and STC-1 were validated in the above in vivo models that the expressions of either TSP-1 or STC-1 were only significantly increased in Group 3, PD-MSCs administration, as shown by immunohistochemical staining (Fig. 8B and C) and RT-PCT (Fig. 8A).
Fig. 7. Influence of PD-MSCs on STC-1 and THP-1. (A) RT-PCR for the changes of stem cell related genes known as regeneration factors including TSP-1, STC-1, HSP27, and hFGF mRNA Right bar graph shows the relative intensity of TSP-1 (upper) and STC-1 (lower). (B) RT-PCR for TSP-1 and STC-1 mRNA after PD-MSCs in the absence or presence of H. pylori infection. (C) Cell viability after H. pylori infection according to STC-1 and TSP-1 status Significant loss of cell viability privilege after PD-MSCs in STC-1 or TSP-1 siRNA. (D) Changes of Bax, cleaved caspase-3, and cleaved PARP after H. pylori infection according to TSP-1 (Western blot on left) and STC-1 (Bar graph). (E) Flow cytometry after DCF-DA staining according to status; H. pylori alone, H. pylori in the presence of PD-MSCs, H. pylori in the presence of PD-MSCs in STC-1 siRNA transfected cells, and H. pylori in the presence of PD-MSCs in TSP-1 siRNA transfected cells.
Discussion

From the current investigation, we reconfirmed the significant rejuvenating action of stem cells, MSCs in the current study, against _H. pylori_-associated CAG as well as gastric tumorigenesis based on authentic renewing and regenerative action of stem cells with additional concerted actions of anti-inflammatory, antioxidative, and restorative action. As summarized in Fig. 9, MSCs, PD-MSCs, UC-MSCs, and AD-MSCs can either revert atrophic gastritis via 15-PGDH, STC-1, and TSP-1 or rejuvenating via autophagy, anti-apoptosis, and IL-10.

Various types of MSCs have been reported to be effective against tissue damages\(^1\) including human adipose tissue-derived MSCs,\(^18\) BM-MSCs,\(^19,20\) mesenchymal stromal cells,\(^21\) placental stromal cells,\(^22\) since stem cells afforded the multi-potent and multi-therapeutic effects for host defense and MSCs homed significantly to injured sites to signal local cells to mitigate inflammation and preserve innate organ function. In the literature, some papers deal with the comparative analysis of MSCs derived from amniotic membrane, umbilical cord, chorionic plate, placenta _decidua parietalis_, bone marrow-derived MSCs,\(^23-27\) placenta derived MSCs showed the best, safe, and low immunogenic advantages, optimal for clinical application. However, in this study, we have administrated via tail vein, but in previous study, we have found similar efficacy when administered via oral route.\(^28\)

Since 15-PGDH may function as a tumor suppressor through antagonizing oncogenic action of COX-2, 15-PGDH has been found to be down-regulated elevated levels of PGE\(_2\) in most tumors, as seen in current study that significantly decreased 15-PGDH was noted in either 24 or 48 weeks of _H. pylori_ infection, significant down-regulation was noted in control group 2, but significantly preserved or elevated expressions of 15-PGDH were observed in group treated with MSCs. Regarding the changes of decreased 15-PGDH in _H. pylori_ infection,\(^29\) these decreased expressions of 15-PGDH were reversed with successful _H. pylori_ eradication, in which suppressed 15-PGDH expressions were associated with TLR-4 and MyD88 expressions, phospho-ERK1/2, and EGFR receptor (EGFR)-Snail.\(^30\)

As restorative and regenerative contribution of MSCs, they secrete mitochondria related hormone named STC1 in a paracrine fashion, which improves the cell survival and regeneration\(^31\) in addition to anti-inflammatory effects via inducing uncoupling proteins to reduce oxidative stress,\(^32\) anti-apoptotic action,\(^33\) body fluid homeostasis,\(^34\) angiogenesis,\(^35\) macrophage polarization, and wound healing.\(^36,37\) Therefore, though STC-1 was originally identified as a calcium/phosphate-regulating hormone in bony fishes, the gene has been documented as key contributing mediator of MSCs in addition to ocular disease, renal disease, idiopathic pulmonary fibrosis, and other degenerative diseases.\(^38-40\) Interestingly, since STC-1,
unexpectedly, is not detected in the circulation under normal circumstances, STC-1 may play an autocrine/paracrine rather than a classic endocrine role in mammals. Therefore, with pleiotropic effects of STC-1 in the stomach, we speculated these paracrine/autocrine effects of STC-1 during atrophic gastritis by MSCs might play rejuvenating outcome. Conclusively, biological repertoires of STC-1 with PD-MSCs administration in our model was considerably larger than their role in fish as well as mineral metabolism.

TSP-1 plays major roles in tissue repair as a regulator of latent TGF-β activation and key player in wound healing and fibrosis relevant to TGF-β. Binding of the TSP to cell surface calreticulin in complex with LDLR-1 stimulates cell adhesion, cell migration, collagen expression and matrix deposition, thereby, altering endothelial cell–cell interactions and stimulating wound healing as well as regeneration via cell migration, well documented in corneal or gingival wound repair.

Though TSP-1 in tissue repair is well-known related to TGF-β-dependent mechanism, like STC-1, independently to TGF-β, they activated cell migration and regeneration. Taken together, in this investigation, for the first time, we identified that STC-1 and TSP-1 together with PD-MSCs concerted to rejuvenate procancerous CAG into non-atrophic condition.

In addition to the above cancer preventive and rejuvenating contribution of MSCs, especially PD-MSCs, in this study, we documented the autophagy induction and anti-apoptotic mechanism as featuring action of MSCs administration during CAG background. Though the role of autophagy in gastrointestinal diseases has been studied extensively, autophagy can be defined as double-sword phenomenon since autophagy is observed under various pathological processes of the GI tract as well as GI cancer, but autophagy can play an important role in the homeostasis as well as maintaining the integrity of intestinal epithelium.

In H. pylori infection, the significance of autophagy seems to be similar with double-edged sword, detrimental or beneficiary. From our investigation, we could document the autophagy induction as regenerating mechanism of PD-MSCs as rescuing from H. pylori-associated CAG (24 weeks). Though autophagy led to H. pylori persistence in the stomach as immune evasion strategy, autophagy accentuated anti-apoptotic mechanisms of MSCs as restoring strategy.

Lastly, IL-10 induction with 15-PGDH was identified as core action of MSCs. The fact that IL-10 gene promoter polymorphism, −819 C/T, was associated with the risk of gastric cancer and atrophic gastritis shed the importance of IL-10 in H. pylori infection. In this background, the induction of 15-PGDH also highlight the contributing role of MSCs. Hence, 15-PGDH, IL-10, and efferocytosis, regulatory T cells (Treg) cooperatively afforded rejuvenating outcome with MSCs in the current study, anti-inflammatory, tumor suppressive, and anti-mutagenic actions were operated with self-renewal and regenerative action of stem cells. Efferocytosis is the process of the recognition and removal of apoptotic inflammatory cells by tissue macrophages as non-professional phagocytes and can lead to the resolution of inflammation. During H. pylori infection, apoptotic neutrophils were detected within the cytoplasmic vacuoles of the foveolar cells of nine cases with chronic active or atrophic-gastritis.

As limitation of the current study, we did not perform in vivo animal model using STC-1 KO, TSP-1 KO, and IL-10 KO mice whether MSCs administration did not effect in this background, instead we validated these in vitro cell models. Also, though we did not experience of complications of MSCs administration in spite of tail vein administration in animal, embolism risk and immunogenic adverse effect was noted in clinical trials of stem cell therapy. In our similar experiment, we identified the administration of MSCs or their conditional media through oral route, similar outcome was noted. Conclusively, we anticipate beneficial efficacy of MSCs administration in H. pylori-associated CAG via endoscopic instillation just like rejuvenating therapy for aged skin, but further detailed investigation should be followed.
Author Contributions

Study concept and design: JMP and KBH; acquisition of data: YMH and JMP; analysis and statistical analysis: KBH; interpretation of data: YMH and JMP; drafting of manuscript: YMH and KBH. All authors approved the final version of this manuscript to be published.

Acknowledgments

This work was supported by Korean Society of Helicobacter and Upper GI Disease (WJ Ko).

Abbreviations

ASC apoptosis-associated speck-like protein containing caspase recruitment domain

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Conflict of Interest

No potential conflicts of interest were disclosed.
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