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Abstract

Leuconostoc mesenteroides H40 (H40) was isolated from kimchi, and its probiotic properties and neuroprotective effect was evaluated in oxidatively stressed SH-SY5Y cells. H40 was stable in artificial gastric conditions and can be attached in HT-29 cells. In addition, H40 did not produce β-glucuronidase and showed resistant to several antibiotics. The conditioned medium (CM) was made using HT-29 cells refined with heat-killed probiotics (probiotic-CM) and heated yogurts (Y-CM) to investigate the neuroprotective effect. Treatment with H40-CM not only increased cell viability but also significantly improved brain derived neurotropic factor (*BDNF*) expression and reduced the *Bax/Bcl-2* ratio in oxidatively stress-induced SH-SY5Y cells. Besides, probiotic Y-CM significantly increased *BDNF* mRNA expression and decreased *Bax*/*Bcl-2* ratio. The physicochemical properties of probiotic yogurt with H40 was not significantly different from the control yogurt. The viable cell counts of lactic acid bacteria in control and probiotic yogurt with H40 was 8.66 Log CFU/mL and 8.96 Log CFU/mL, respectively. Therefore, these results indicate that H40 can be used as prophylactic functional dairy food having neuroprotective effects.

Keywords probiotics, *Leuconostoc mesenteroides*, neuroprotective effect, probiotic yogurt, oxidative stress

Introduction

Probiotics are living microbes that deliver health benefits to the host when ingested in adequate amounts by the FAO/WHO (Chamber et al., 2019). The common probiotic strains mainly belong to the *Lactobacillus*, *Leuconostoc*, *Pediococcus*, and *Bifidobacterium* species and are widely used in many probiotic products (O'Toole et al., 2017). Probiotics have many reported health benefits such as improvement of cognitive function (Ton et al., 2020), antioxidant (Jang et al., 2018; Yu et al., 2019b), anti-inflammatory (Yu et al., 2019c), antihypertensive (Klippel et al., 2016), or cholesterol lowering (Ishimwe et al., 2015) activities. To utilize this functionality, probiotics are also used as medical or food additives.

The brain, which is rich in phospholipids, is an organ with high oxygen demand and is vulnerable to the effects of reactive oxygen species (ROS) (Dussert et al., 2006). ROS is an essential byproduct of aerobic metabolism (Wang and Michaelis, 2010). However, excessive ROS levels cause cell damage by oxidizing cellular biomolecules, including nucleic acids, proteins, and lipids (Lobo et al., 2010). ROS can contribute to pathologies, such as cancer (Lee et al., 2014), cardiovascular disease (Elahi et al., 2009), diabetes, and aging (Pamplona and Barja, 2006).

The bidirectional signaling connecting the brain and the gastrointestinal tract is crucial for maintaining homeostasis and is regulated the neural, hormonal, and immunological levels (Ghaisas et al., 2016; Wang and Kasper, 2014). Probiotics have recently become a target as live bacterial cell biotherapies for neurodegenerative disease (Quigley, 2017; Wang et al., 2016). *Clostridium butyricum* can exert neuroprotective effects against ischemia/reperfusion injury mice through antioxidant and anti-apoptosis mechanisms (Sun et al., 2016). *Lactobacillus buchneri* KU200793 showed neuroprotective effect using SH-SY5Y cells induced with 1-methyl-4-phenylpyridinium (MPP^+) (Cheon et al., 2020).

Brain derived neurotropic factor (BDNF) expression occurs in the brain, and low secretion of BDNF influences human memory and hippocampal functions (Egan et al., 2003). BDNF is medicated by extracellular signal-regulated kinase (ERK) 1/2, ERK5, and phosphatidylinositol-3 kinase (PI3k) pathways in cortical neurons to promote neuronal survival (Liu et al., 2003). Oxidative stress may induce mitochondrial dysfunction and deficiency in protein aggregation and ultimately cause nerve cell death (Lobo et al., 2010). The mitochondrial apoptotic pathways are mediated through the Bcl-2 family proteins, which include Bax that promotes pro-apoptotic mitochondrial permeability and anti-apoptotic Bcl-2 that inhibits apoptotic effects (Azmi et al., 2013). The *Bax*/*Bcl-2* ratio is a determining factor in the regulation of apoptotic cell death.

Leuconostoc mesenteroides is bacteria sometimes related to fermentation under salinity and low temperature in fermented foods (Yoon et al., 2018). *L. mesenteroides* is an obligate heterofermentative lactic acid bacterium that is mostly used in dairy fermentation. *L. mesenteroides* has been studied as a probiotic strain that facilitates the removal of Pb (II) toxicity (Yi et al., 2017) and inhibits biofilm formation against *Listeria monocytogenes* (Shao et al., 2019). However, the neuroprotective effects of *L. mesenteroides* have not been studied. Therefore, the aims of this study were to demonstrate the probiotic properties and neuroprotective effect of *L. mesenteroides* H40 isolated from kimchi and confirm this effect in yogurt fermented using *L. mesenteroides* H40.

Materials and Methods

Bacterial strains and culture condition

Lactobacillus fermentum KU200060, *Lactobacillus brevis* KU200080, and *Leuconostoc mesenteroides* H40 were isolated from kimchi with salted water, mustard leaf (*Brassica juncea*) kimchi, and Chinese cabbage kimchi using lactobacilli MRS medium (MRS; BD Biosciences, Franklin Lakes, USA) and identified by 16S rRNA analysis (Bionics, Seoul, Korea). *Lactobacillus rhamnosus* GG (Cell Biotech., Ltd., Gimpo, Korea) was used as a control strain. Bacteria were propagated and maintained in MRS medium at 37°C for 24 h.

Cell culture condition

The HT-29 (human colon adenocarcinoma, KCLB 30038) and SH-SY5Y (human neuroblastoma, KCLB 22266) cells were used for this study. The cells were grown in Roswell Park Memorial Institute (RPMI) 1640 medium (Gibco, Grand Island, USA) and Dulbecco's Modified Eagle's Medium (HyClone Laboratories Inc., Logan, USA), respectively. All media were accompanied with 10% (v/v) fetal bovine serum (Gibco) and 1% (v/v) penicillin/streptomycin (Gibco). The cells were maintained at 37°C in 5% CO_2. The cultured cells were maintained to monolayer.

Tolerance to artificial gastric conditions

To measure the stability against gastric conditions, artificial gastric juice and bile salts were followed the methods by Yang et al. (2019). The tested strains were incubated in MRS broth at 37°C for 18 h. Initial cells were inoculated at the concentration of 1×10^7 CFU/mL. Artificial gastric conditions were dealt on 0.3% pepsin (Sigma-Aldrich, St. Louis, USA) adjusted to pH 2.5 at 37°C for 3 h. Artificial bile conditions were used 0.3% oxgall (BD Biosciences) at 37°C for 24 h. After incubation, the survival rate was determined by calculating viable cells on MRS plates.

Adhesion ability to HT-29 cells

The adhesion ability of isolated strains was examined using HT-29. HT-29 cells (1×10^5 cells/mL) was planted in a 24-well cell culture plate and incubated at 37°C (Lee et al., 2015). After 24 h, isolated strains (1×10^7 CFU/mL) were inoculated and incubated in HT-29 cells at 37°C for 2 h. Non-adherent bacteria were washed three times using PBS buffer (Gibco), 1% Triton X-100 (Sigma-Aldrich) solution was used for separate the adherent bacteria. The number of adherent bacteria was determined by dilution and plating on MRS plates.

Enzyme production

To measure of enzyme production, the API ZYM kit (BioMerieux, Lyon, France) were used as manufacture's guideline. Each strain at 10^6 CFU/mL was put in each cupule and incubated at 37°C for 4 h. After incubation, zym A and B reagents put in each cupule, and represented as production concentration (between 0 and ≥40 nM).

Antibiotic resistance

Antibiotic resistance was followed Clinical and Laboratory Standards Institute guideline (CLSI, 2012). One hundred microliters of each lactic acid bacteria (LAB) strains (1×10^7 CFU/mL) was inoculated onto MRS agar and paper disc were put on agar plate. Used antibiotics were ampicillin (10 μg), gentamycin (10 μg), kanamycin (30 μg), ciprofloxacin (5 μg), chloramphenicol (30 μg), streptomycin (10 μg), tetracycline (30 μg), and doxycycline (30 μg). After incubation at 37°C for 24 h, the inhibitory diameter zone was calculated and compared to the cut-off value (>20 mm, susceptible; 15-19 mm, intermediate; ≤14, resistant) by represented in CLSI.

Conditioned medium (CM) from HT-29 cells

The CM was prepared using HT-29 cells following the method of Park et al. (2017) with minor modifications. For CM preparation, each sample of LAB strains and yogurt was heated at 121°C for 15 min and stored at -80°C upto use. HT-29 cells were inoculated into 6-well plates to 1.0×10^6 cells/well and incubated to a confluent monolayer. After incubation, cells were handled with heat-killed LAB (8 Log CFU/mL) or heated yogurt for 24 h. CM treated PBS (Gibco) instead to samples were used as control. The mixture was centrifuged (12,000×g, 10 min) and the supernatant was assembled using a syringe filter (0.45 μm pore size, Millipore Sigma, Burlington, MA, USA).

Protective effect on oxidative stress-induced apoptosis

To confirm the protective effect on oxidative stress-induced apoptosis, oxidative stress was induced utilizing H_2O_2 (Junsei Chemical, Tokyo, Japan) or $NaAsO_2$ (Sigma-Aldrich). The SH-SY5Y cells (100 μL, 1.0×10^5 cells/well) were inoculated in 96-well plate with of 50 μM H_2O_2 (20 μL) or 10 μM $NaAsO_2$ (20 μL) for 20 h after pretreatment with 80 μL of sample (CM) for 4 h. After incubation, the media were eliminated, and the cells were incubated with 5 mg/mL MTT solutions (100 μL) for 1 h. After incubation, the liquid was removed and DMSO (100 μL) was added to each well. Absorbance was gauged at 540 nm utilizing microplate reader. The cell viability (%) was calculated as follows:

$$\text{Cell viability (\%)} = \frac{\text{Absorbance of sample}}{\text{Absorbance of control}} \times 100$$

Yogurt production, physicochemical composition, and viable cell counts of LAB

Yogurt was prepare from whole milk (Seoul Milk Co., Ltd., Seoul, Korea) purchased from a local market. The milk was heated at 90°C for 10 min and cooled to 40°C using water bath. An overnight culture of *L. mesenteroides* H40 was centrifuged (14,000×g, 10 min, 4°C) and the cells were washed twice with PBS (Gibco). Then, the pasteurized milk was

inoculated with ABT-B commercial yogurt starter culture containing *Lactobacillus acidophilus*, *Lactobacillus delbrueckii* subsp. *bulgaricus*, *Bifidobacterium longum*, and *Streptococcus thermophilus* (Samik Dairy Co., Ltd., Gimje, Korea) or a mixed culture (1:1) of *L. mesenteroides* H40 and ABT-B commercial yogurt starter culture. The inoculated mixture was incubated at 40°C to pH 4.5. Then, the yogurt samples fortified with *L. mesenteroides* H40 were ripened for 24 h in the refrigerator, and its physicochemical properties were analyzed. Composition and pH of yogurt was analyzed using Milko Scan Minor (Foss, Hillerod, Denmark) and a pH-meter (WTW inoLab 7110, Weilheim, Germany), respectively. Titratable acidity was assessed according to AOAC International (1999) by titration with sodium hydroxide using phenolphthalein. Measurements of viscosity were performed with Brookfield DV-E Viscometer (Brookfield Eng. Lab. Inc., Middleboro, USA) using spindle No. 3 at 50 rpm. Viable cell counts of LAB in yogurt samples was confirmed using decimal dilutions, spread-plated on MRS medium, and incubation at 37°C for 48 h.

BDNF, _Bax_, and _Bcl-2_ expression on oxidative stress-induced apoptosis in SH-SY5Y cells

SH-SY5Y cells (1.0×10^6 cells/well) were seeded on 6-well plate and incubated to form a confluent monolayer. After incubation, the cells were treated with 800 μL of CM for 4 h. To induce oxidative stress, 200 μL of H_2O_2 (50 μM) or $NaAsO_2$ (10 μM) was added for 20 h. Total RNA was isolated using the RNeasy Mini total RNA isolation kit (Cheon et al., 2020; Park et al., 2017).

Real-time polymerase chain reaction

The RNA quality was quantified using microplate reader (Multiscan^{TM} Go, Thermo Fisher Scientific, Waltham, USA). cDNA was manufactured using cDNA synthesis kit

(Thermo Fisher Scientific). Semi-quantitative real-time PCR was performed according to the PikoReal 96 system (Thermo Fisher Scientific). The reactants contained SYBR Green master mix, primer (Table 1), cDNA, and RNase free water. Further, 20 μL of the mixture was amplified as 95°C for 2 min as initial denaturation; 40 cycles of 95°C for 5 s as denaturation; 60°C for 15 s as annealing and extension. The results were analyzed by ΔΔCt method using the melt curve analysis method.

Statistical analysis

All tested data are represented as mean±standard deviation by three replicates. One-way analysis of variance (ANOVA) was utilized to verify significant differences. The mean values were used for the Duncan's multiple range test to perform post-hoc verification ($p<0.05$).

Results and Discussion

Tolerance to artificial gastric conditions and Adhesion to HT-29 cells

L. fermentum KU200060, *L. brevis* KU200080, and *L. mesenteroides* H40 was isolated from various kimchi for probiotic use. *L. rhamnosus* GG, *L. fermentum* KU200060, *L. brevis* KU200080, and *L. mesenteroides* H40 was confirmed probiotic properties (Table 2; $p<0.05$). These strains showed high tolerance to artificial gastric conditions. *L. rhamnosus* GG and *L. mesenteroides* H40 decreased to 8.51 Log CFU/mL and 7.17 Log CFU/mL in acidic conditions, however increased to 8.58 Log CFU/mL and 8.26 Log CFU/mL in bile conditions, respectively. *L. fermentum* KU200060 and *L. brevis* KU200080 showed strong acid tolerance having 8.29 Log CFU/mL and 7.91 Log CFU/mL, however decreased to 7.41 Log CFU/mL

and 7.75 Log CFU/mL in bile conditions, respectively. *L. plantarum* Ln1 and KCTC 3108 showed similar trends having decrease in acidic conditions and remaining in bile conditions (Jang et al., 2018).

L. rhamnosus GG, *L. fermentum* KU200060, *L. brevis* KU200080, and *L. mesenteroides* H40 showed 2.34%, 1.18%, 3.42%, and 2.86% adhesion rate to HT-29 cells. Especially, *L. brevis* KU200080 and *L. mesenteroides* H40 showed a higher adhesion rate than *L. rhamnosus* GG. Jang et al. (2018) showed lower 2.19% adhesion rate of *L. plantarum* KCTC 3108. Adhered probiotic strains may be temporary colonization and influence host health trough adjustment of intestinal microflora (Jang et al., 2019; Yu et al., 2019a).

Enzyme production

β-Glucuronidase can be produced by the human intestine microbiota and liberate toxin and mutagen in liver (Dabek et al., 2008). Therefore, isolated strains were confirmed nonproduction of β-glucuronidase using API ZYM kit (Table 2). *L. rhamnosus* GG produced 30 nM of leucine aryamidase, 30 nM of valine arylamidase, 20 nM of naphthol-AS-BI-phosphohydrase, 20 nM of β-galactosidase, and 30 nM of β-glucosidase. *L. fermentum* KU200060 produced 30 nM of α-galactosidase and ≥40 nM of β-galactosidase. *L. brevis* KU200080 produced 20 nM of β-galactosidase, 30 nM of β-glucosidase, and 30 nM of leucine arylamidase. *L. mesenteroides* H40 produced 20 nM of α-glucosidase and 30 nM of β-glucosidase. α-Galactosidase and β-galactosidase can act the use of indigestible carbohydrates of raffinose family oligosaccharides and milk products, respectively. In addition, β-glucosidase may influence bioavailability by the cleavage of glycosidic bonds in ginsenoside, isoflavone, and phenolic compounds (Son et al., 2018). Produced enzyme by these isolated strains may be useful for carbohydrate digestion.

Antibiotic resistance

L. fermentum KU200060 and *L. mesenteroides* H40 are resistant to gentamycin, kanamycin, and ciprofloxacin. *L. rhamnosus* GG and *L. brevis* KU200080 are resistant to gentamycin, kanamycin, streptomycin, and ciprofloxacin. Among tested antibiotics, most *Lactobacillus* sp. are intrinsically resistant to aminoglycoside (gentamycin, kanamycin, and streptomycin), inhibitors of nucleic acid synthesis (ciprofloxacin) (Campedelli et al., 2015). Therefore, isolated strains showed a potential of safe probiotic strains in a view of antibiotic resistance.

Protective effects of probiotics-CM on oxidative stress-induced apoptosis in SH-SY5Y cells

H_2O_2 and $NaAsO_2$ converts to a highly reactive toxic hydroxyl radical (Pardillo-Díaz et al., 2016), causing damage by reducing antioxidant enzymes in brain (Herrera et al., 2013). Additionally, gut microbiota influence the neurophysicals at the base of the gut-brain axis (Park et al., 2017). The modulatory effect of probiotics in intestinal microbiota was demonstrated by increased a ratio of Firmicutes to Bacteriodes and it can relieve inflammation by cytokine expression (Martin et al., 2018). Therefore, the CM using HT-29 cells with probiotics was used for neuroprotective effects.

Oxidative stress was induced in SH-SY5Y cells using H_2O_2 or $NaAsO_2$, and cell viability was confirmed by MTT assay (Fig. 1). During the induction of oxidative stress by H_2O_2, the cell viability of SH-SY5Y cells was 53.5% (Fig. 1A; $p<0.05$). The cell viability of the probiotics-CM for *L. rhamnosus* GG, *L. fermentum* KU200060, *L. brevis* KU200080, and *L. mesenteroides* H40 was 70.7%, 49.7%, 65.0%, and 69.9%, respectively. *L. rhamnosus* GG, *L. brevis* KU200080, and *L. mesenteroides* H40 showed a protective effect compared to H_2O_2 treated cells (53.5%).

During induction of oxidative stress by $NaAsO_2$, the cell viability of SH-SY5Y cells was 55.8% (Fig. 1B; $p<0.05$). The cell viability of the probiotics-CM of *L. rhamnosus* GG, *L. fermentum* KU200060, *L. brevis* KU200080, and *L. mesenteroides* H40 was 55.3%, 49.2%, 55.3%, and 70.7%, respectively. Only *L. mesenteroides* H40 showed a protective effect compared to $NaAsO_2$ treated cells (55.8%).

Among these strains, *L. mesenteroides* H40 has highest cell viability in SH-SY5Y cells using both H_2O_2 and $NaAsO_2$. Cheon et al. (2020) showed the cell viability of *L. rhamnosus* GG (72.0%), *L. fermentum* KU200060 (60.2%), *Lactobacillus delbrueckii* KU2000171 (66.8%), and *L. buchneri* KU200793 (73.4%) with MPP^+ as Parkinson-inducing toxin having oxidative phosphorylation (Cheon et al., 2020). Therefore, *L. rhamnosus* GG and *L. mesenteroides* H40 was demonstrated neuroprotective effects against oxidative stress.

BDNF mRNA expression and anti-apoptotic effects of probiotics-CM on oxidative stress-induced apoptosis in SH-SY5Y cells

The gut-brain axis (GBA) is bi-directional communication network encompassing the autonomic nervous system (ANS), the central nervous system (CNS), and the enteric nervous system (ENS). These complex network was influenced by gastrointestinal tract (Kennedy et al., 2016; Ranuh et al., 2019). Among serum response factor, BDNF have known as regulator of the synaptic protein and precursors for appropriated neuronal function, survival, and apoptosis (Numakawa et al., 2010). Decreased *BDNF* mRNA expression confirms brain related diseases such as Alzheimer's disease, Parkinson's disease, and depression. Increased ratio of *Bax/Bcl-2* induced apoptosis.

BDNF mRNA expression and *Bax*/*Bcl-2* ratio is shown in Fig. 2. Treatment with H_2O_2 reduced *BDNF* mRNA expression by 0.73-fold compared with that in H_2O_2 nontreated cells (Fig. 2A; $p<0.05$). *L. rhamnosus* GG and *L. mesenteroides* H40 showed 0.80- and 0.85-fold

276 *BDNF* mRNA expression, respectively. The *Bax*/*Bcl-2* ratio in H_2O_2 nontreated cells was
277 1.00-fold, whereas H_2O_2 increased the ratio of 2.69-fold (Fig. 2B; $p<0.05$). Treatment with *L.*
278 *rhamnosus* GG and *L. mesenteroides* H40 reduced the *Bax*/*Bcl-2* ratio to 2.24- and 2.03-fold,
279 respectively.

280 Treatment with $NaAsO_2$ reduced 0.76-fold *BDNF* mRNA expression compared with that
281 in the control without $NaAsO_2$ treatment (Fig. 2C; $p<0.05$). *L. rhamnosus* GG and *L.*
282 *mesenteroides* H40 represented 0.95- and 1.08-fold *BDNF* mRNA expression, respectively.
283 The *Bax*/*Bcl-2* ratio in $NaAsO_2$ nontreated cells was 1.00-fold, while $NaAsO_2$ increased 2.24-
284 fold in $NaAsO_2$ treated cells. Treatment with *L. rhamnosus* GG increased 2.61-fold, while
285 treatment with *L. mesenteroides* H40 reduced 1.46-fold (Fig. 2D; $p<0.05$).

286 *L. mesenteroides* H40 can increase *BDNF* mRNA expression and reduce apoptosis of
287 SH-SY5Y cells oxidatively stressed using both H_2O_2 and $NaAsO_2$. The difference of
288 neuroprotective effect of *L. rhamnosus* GG and *L. mesenteroides* H40 depends on strain and
289 oxidant.

290

291 **Physicochemical property and LAB cell counts of control and probiotic yogurt**

292 Yogurt is a major probiotic carrier to consumers without side-effect. Each yogurt was
293 manufactured using the following: 1) ABT-B commercial starter culture (control yogurt) and
294 2) ABT-B commercial starter mixed with *L. mesenteroides* H40 (probiotic yogurt). The fat,
295 protein, lactose, total solids, and acidity content are shown in Table 3 ($p<0.05$). Probiotic
296 yogurt made using *L. mesenteroides* H40 had 2.96% fat, 3.23% protein, 6.16% lactose, and
297 27.33% total solids. In addition, probiotic yogurt was not significantly different from control
298 yogurt. However, probiotic yogurt exhibited significantly higher viscosity than control yogurt.
299 Texture of stirred yogurt is the result of both acid aggregation of casein micelles by ropy
300 strains during incubation (Zhao et al., 2016). The viable cell counts of lactic acid bacteria in

control and probiotic yogurt with H40 was 8.66± Log CFU/mL and 8.96± Log CFU/mL, respectively (data not shown).

Protective effects of Y-CM oxidative stress-induced apoptosis in SH-SY5Y neuroblastoma cells

Y-CM was manufactured with HT-29 cells and yogurt, and its neuroprotective effect was assessed in SH-SY5Y cells (Table 4; $p<0.05$). The treatment of H_2O_2 reduced cell viability of SH-SY5Y cells to 55.5%. However, cell viability of control yogurt CM (CY-CM) and probiotic yogurt CM (PY-CM) was 72.2% and 114.8%, respectively. Under treatment with $NaAsO_2$, cell viability of positive control was 51.4% and that of CY-CM and PY-CM was 49.9% and 109.5%, respectively. The PY-CM using *L. mesenteroides* H40 showed high cell viability in oxidatively stressed SH-SY5Y cells in both H_2O_2 and $NaAsO_2$ treatment. When compare Fig. 1 and Table 4, PY-CM showed higher cell viability than *L. mesenteroides* H40. These results showed that the PY-CM effectively protected the cells from oxidative damage caused by H_2O_2 and $NaAsO_2$.

H_2O_2 treatment resulted in a 0.78-fold increase in *BDNF* mRNA expression compared with that in the H_2O_2 nontreated cells (Table 4; $p<0.05$). CY-CM and PY-CM increased *BDNF* mRNA expression by 0.83- and 1.12-fold, respectively. The ratio of *Bax*/*Bcl-2* ratio in H_2O_2 nontreated cells was 1.00, while H_2O_2 increased to 2.69-fold. The treatment with CY-CM and PY-CM reduced the *Bax*/*Bcl-2* ratio to 2.05- and 1.24-fold, respectively.

The treatment with $NaAsO_2$ reduced *BDNF* mRNA expression by 0.76-fold compared to that in $NaAsO_2$ nontreated cells (Table 4; $p<0.05$). CY-CM and PY-CM treatment resulted in a 1.03- and 1.18-fold *BDNF* mRNA expression, respectively. The ratio of *Bax*/*Bcl-2* ratio in $NaAsO_2$ nontreated cells was 1.00, while $NaAsO_2$ treatment increased this to 2.24-fold in

$NaAsO_2$ treated cells. The treatment with CY-CM and PY-CM reduced to 1.88- and 1.32-fold, respectively. Thus, PY-CM can reduce apoptosis of H_2O_2 or $NaAsO_2$ stressed SH-SY5Y cells.

The treatment with PY-CM significantly increased *BDNF* mRNA expression and reduced apoptosis on SH-SY5Y mRNA. In addition, mRNA expression was markedly higher based on the yogurt type than with the strain alone or with the yogurt and probiotic strain. These synergistic effects originate from whey protein containing methionine, lysine, and proline, which are associated with apoptosis (Lee and Hur, 2019). Therefore, PY-CM using *L. mesenteroides* H40 has potent neuroprotective effect in preventing the oxidative stress induced by H_2O_2 and $NaAsO_2$.

Conclusions

L. mesenteroides H40 was isolated from kimchi and its probiotic property was demonstrated through stability in gastric conditions, adhesion to intestinal cells, enzyme production, and safe antibiotic resistance. For neuroprotective effect, *L. mesenteroides* H40-CM confirmed an increase of cell viability with increase of *BDNF* mRNA expression and decrease of the *Bax/Bcl-2* ratio in oxidatively stress-induced SH-SY5Y cells. Compared to probiotic strain and yogurt type, the probiotic yogurt showed a higher neuroprotective effect than the strain alone. Therefore, these results suggested a potential of prophylactic therapy as probiotics and functional dairy products. In addition, the cognitive function-related experiments will need to be performed in humans for efficacy verification.

Conflict of Interest

The authors declare no potential conflict of interest.

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Table 1. Primer sequence for neuroprotective effect used in semi-quantitative real-time PCR

Gene		Primer sequence	References
GAPDH	Forward	5' GAGTCAACGGATTTGGTCGT 3'	Park et al., 2017
	Reverse	5' GACAAGCTTCCCGTTCTCAG 3'	
BDNF	Forward	5' CAAACATCCGAGGACAAGGTGG 3'	
	Reverse	5' CTCATGGACATGTTTGCAGCATCT 3'	
Bax	Forward	5' GTGGTTGCCCTCTTCTACTTTGC 3'	
	Reverse	5' GAGGACTCCAGCCACAAAGATG 3'	
Bcl-2	Forward	5' CGGCTGAAGTCTCCATTAGC 3'	
	Reverse	5' CGGCTGAAGTCTCCATTAGC 3'	

BDNF, brain-derived neurotrophic factor; *Bax*, *Bcl-2*-associated X protein; *Bcl-2*, B-cell lymphoma.

476 **Table 2. Probiotic properties of isolated strains**

Treatment	LGG[a]	200060	200080	H40
Tolerance to artificial gastric conditions (Viable cell number (Log CFU/mL))				
Initial cell number	8.55±0.01[a]	8.26±0.01[b]	7.79±0.01[c]	8.17±0.03[b]
0.3% (w/v) pepsin, pH 2.5, 3 h	8.51±0.05[a]	8.29±0.01[ab]	7.91±0.00[b]	7.17±0.01[c]
0.3% (w/v) oxgall, 24 h	8.58±0.03[a]	7.41±0.01[bc]	7.75±0.02[b]	8.26±0.01[a]
Adhesion rate (%)[b]	2.34±0.26[c]	1.18±0.08 [d]	3.42±0.49[a]	2.86±0.16[b]
β-Glucuronidase (nM)	0	0	0	0
Antibiotic resistance	Gentamycin, kanamycin, streptomycin, ciprofloxacin	Gentamycin, kanamycin, ciprofloxacin	Gentamycin, kanamycin, streptomycin, ciprofloxacin	Gentamycin, kanamycin, ciprofloxacin

477 [1]LGG, *L. rhamnosus* GG; 200060, *L. fermentum* KU200060; 200080, *L. brevis* KU200080;
478 H40, *L. mesenterodies* H40

479 [2]Adhesion rate = (adhered bacteria to HT-29 cells after 2 h)/(initial bacteria)×100

480 Data are represented as the mean±standard deviation of triplicate experiments. Means within
481 a row with same superscript differ ($p<0.05$).

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Table 3. Physicochemical properties of control and probiotic yogurt

Physicochemical properties	Yogurt type	
	Control yogurt[1]	Probiotic yogurt[2]
Fat (%)	2.96±0.05[a]	2.96±0.20[a]
Protein (%)	3.16±0.20[a]	3.23±0.11[a]
Lactose (%)	6.23±0.11[a]	6.16±0.15[a]
Total solid (%)	27.66±0.15[a]	27.33±0.28[a]
Titratable acidity	0.82±0.01[a]	0.81±0.01[a]
pH	4.33±0.04[a]	4.26±0.03[a]
Viscosity (cP)	1,724.20±15.60[a]	2,048.30±7.30[b]

[1]Control yogurt, yogurt manufactured by ABT-B starter.

[2]Probiotic yogurt, yogurt manufactured by ABT-B starter and *L. mesenteroides* H40.

Data are represented as the mean±standard deviation of triplicate experiments. Means within a row with same superscript differ ($p<0.05$).

Table 4. Neuroprotective effect of control and probiotic yogurt

Characteristics	Oxidant	
	H_2O_2 (50 μM)	$NaAsO_2$ (10 μM)
Cell viability (%)		
Non-treatment	100±5.43[c]	100±11[b]
Control	55.45±0.78[a]	51.37±3.5[a]
CY-CM[1]	72.21±1.55[b]	49.92±1.50[a]
PY-CM[2]	114.76±9.30[d]	109.5±11.5[b]
BDNF expression (fold of control)		
Non-treatment	1.00±0.06[b]	1.00±0.07[b]
Control	0.73±0.02[a]	0.76±0.03[a]
CY-CM[1]	0.78±0.12[a]	1.03±0.02[b]
PY-CM[2]	1.05±0.05[b]	1.18±0.02[b]
Bax/Bcl-2 ratio		
Non-treatment	1.00±0.08[a]	1.00±0.07[a]
Control	2.69±0.08[c]	2.25±0.04[b]
CY-CM[1]	2.05±0.07[b]	1.88±0.21[b]
PY-CM[2]	1.24±0.13[a]	1.32±0.17[a]

[1]CY-CM, conditioned medium using yogurt manufactured by ABT-B starter.

[2]PY-CM, conditioned medium using yogurt manufactured by ABT-B starter and *L. mesenteroides* H40.

Data are represented as the mean±standard deviation of triplicate experiments.

Means within same characteristics a row with same superscript differ ($p<0.05$).

Figure legends

Fig. 1. Cell viability of conditioned medium using lactic acid bacteria (LAB-CM) in SH-SY5Y cells with oxidative stress induced by (A) H_2O_2 (50 μM) and (B) $NaAsO_2$ (10 μM). LGG, *L. rhamnosus* GG; 200060, *L. fermentum* KU200060; 200080, *L. brevis* KU200080; H40, *L. mesenteroides* H40. Error bars indicate standard deviation from three independent experiments. Different letters on each bar represent significantly different ($p<0.05$).

Fig. 2. mRNA expression levels of *BDNF* and apoptosis-related genes on oxidatively stressed SH-SY5Y cells treated with conditioned medium using lactic acid bacteria (LAB-CM). (A) *BDNF* mRNA expression and (B) *Bax/Bcl-2* ratio in oxidative stress-induced SH-SY5Y cells induced by H_2O_2 (50 μM). (C) *BDNF* mRNA expression and (D) *Bax/Bcl-2* ratio in oxidatively stress-induced SH-SY5Y cells induced by $NaAsO_2$ (10 μM). LGG, *L. rhamnosus* GG; H40, *L. mesenteroides* H40. Error bars indicate standard deviation from three independent experiments. Different letters on each bar represent significantly different ($p<0.05$).

513 **Fig. 1.**

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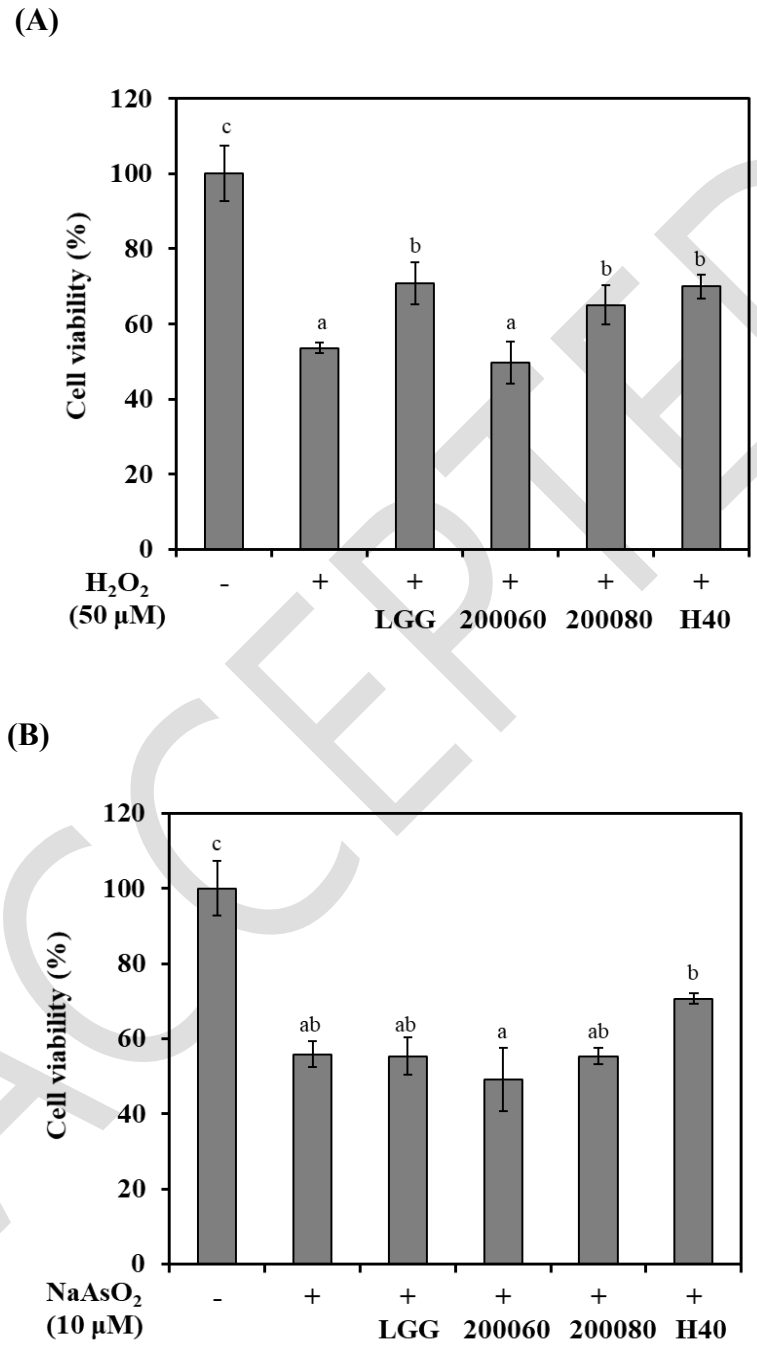

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Fig. 2.