Inactivated probiotic *Bacillus coagulans* GBI-30 induces complex immune activating, anti-inflammatory, and regenerative markers in vitro

**Objective:** The aim of this study was to document the immune activating and anti-inflammatory effects of inactivated probiotic *Bacillus coagulans* GBI-30, 6086 (Staimune™) cells on human immune cells in vitro.

**Methods:** In vitro cultures of human peripheral blood mononuclear cells (PBMC) from healthy blood donors were treated with inactivated *B. coagulans* GBI-30, 6086 cells for 24 hours. After incubation, the PBMC were stained with fluorochrome-labeled monoclonal antibodies for CD3, CD56, and CD69 to monitor cellular activation by flow cytometry. The culture supernatants were tested for cytokine profile using a 27-plex Luminex array, including pro- and anti-inflammatory cytokines, chemokines, and growth factors.

**Results:** Inactivated *B. coagulans* GBI-30, 6086 cells induced the CD69 early activation marker on CD3<sup>+</sup>CD56<sup>-</sup> T lymphocytes, CD3<sup>+</sup>CD56<sup>+</sup> NKT cells, CD3<sup>-</sup>CD56<sup>+</sup> NK cells, and also some cells within the CD3<sup>-</sup>CD56<sup>-</sup> non-T non-NK cell subset. Culture supernatants showed robust increases in the immune-activating cytokines IL-1β, IL-6, IL-17A, and TNF-α. IFN-γ levels were increased, along with three chemokines, MCP-1, MIP-1α, and MIP-1β. The two anti-inflammatory cytokines IL-1ra and IL-10 showed increases, as well as the G-CSF growth factor involved in repair and stem cell biology. In contrast, GM-CSF levels showed a mild decrease, showing a highly selective growth factor response.

**Conclusion:** The inactivated *B. coagulans* GBI-30, 6086 cells activated human immune cells and altered the production of both immune activating and anti-inflammatory cytokines and chemokines. Of special importance is the novel demonstration of a selective upregulation of the G-CSF growth factor involved in postinjury and postinflammation repair and regeneration. This suggests that important immunogenic cell wall components, such as lipoteichoic acid, are undamaged after the inactivation and retain the complex beneficial biological activities previously demonstrated for the cell walls from live *B. coagulans* GBI-30, 6086 (GanedenBC<sup>30</sup>) probiotic bacteria.

**Keywords:** anti-viral, anti-inflammatory, cytokines, growth factors, lipoteichoic acid, inactivated bacillus coagulans GBI-30, 6086, Staimune

**Plain language summary**

This study was done to test a new consumable health product, made by inactivating a specific strain of beneficial probiotic gut bacteria. We have previously shown that this strain of gut bacteria activates human immune cells and helps mature certain immune cells that are of importance for detecting foreign antigens. The inactivating process allows the bacterium to be used in broader applications, such as foods, where a living bacterium could spoil the food or give it a very limited
shelf life. It was important to show that the inactivated bacteria had similar properties to the live bacteria.

To test this, we used blood samples from healthy humans and isolated a part of the white blood cells that include immune cells and stem cells. The blood samples contained the same types of cells as in the blood circulation in our intestinal walls, where antigens from the gut are presented to immune cells. We cultured the cells with the inactivated bacterial cells for 24 hours. We examined the immune cells for their activation status. We tested the liquid culture medium for secreted biomarkers.

We found that the inactivated bacterial cells had similar effects as the live ones with respect to immune activation and anti-inflammatory effects. We also found novel effects that showed that the human cells secreted a growth factor important for tissue repair after trauma and injury.

Introduction

The gut microbial community, “gut microbiome”, has a vast impact on the health of the human host. An integral collaboration exists between microbial forms colonizing the gut and our immune function, metabolism, and brain function. The interaction between gut microbes and host cells and tissue takes place in several ways, including via bacterial cell wall components and secreted metabolites. The most immediate and direct interaction between microbes and host cells involves the outer layers of bacterial cell wall components engaging with receptors on immune cells, such as dendritic cells directly sampling antigens in the gut lumen. This interaction presents different types of bacterial cell wall components of Gram-positive versus Gram-negative bacteria. Gram-negative bacteria present lipopolysaccharides (LPSs), which are recognized by toll-like receptor-4 (TLR-4), whereas the outer cell walls of Gram-positive bacteria present teichoic acid and lipoteichoic acid to immune cells, recognized by TLR-2. For many pathogenic bacteria, lipoteichoic acid is associated with virulence, whereas lipoteichoic acid from beneficial probiotic bacteria trigger complex beneficial immune modulation. Lipoteichoic acid has been widely used as a model TLR-2 ligand to explore a wide variety of immune activating mechanisms at the cellular and molecular level, and interestingly has proven to exert both proinflammatory and anti-inflammatory activities in vitro. The structural complexity of lipoteichoic acid is said to impact the host immune response. The chemical composition of this acid differs between microbes and between strains of similar microbes. This is of high importance in triggering diverse effects on host cells and is likely one of the key factors in the highly selective immune-modulating effects induced by different microbial strains.

The consumption of beneficial probiotic bacteria is associated with a range of health benefits tied to inflammation regulation, such as gastrointestinal disease, respiratory tract infections, neuro-immune and neuropsychiatric disorders, satiety and psychosocial behavior in obese individuals, and alleviation of symptoms of anxiety and depression, as a result of the extensive communication between the gastrointestinal and central nervous systems, also referred to as the “gut–brain axis”. While the consumption of probiotic bacteria is considered highly safe, there are many useful applications for inactivated probiotic strains, such as increased shelf life, as well as usefulness in many types of food products where metabolically active, living bacteria may spoil the appearance of the food. Inactivated probiotic bacteria also have a use in specific clinical situations involving immune-compromised individuals where there could potentially be a risk of translocation of gut bacteria into the bloodstream. Inactivated probiotic bacteria can be produced, for example by heating, leaving the outer bacterial cell wall as the main mechanism of interaction with host immune cells. Heat-killed Lactobacillus plantarum L-137 (HK L-137) has been widely studied over the past decades for its effects in rodents and humans. Animal studies have shown that consumption of HK L-137 offers protection against influenza virus infection, associated with increased production of interferons, suggesting a general support of antiviral immune defense activity. Clinical trials have shown that the daily intake of HK L-137 supports a healthy immune function, including enhanced acquired immune responses and TH1-related immune function, and reduced incidence of upper respiratory tract infections. Consumption was also associated with improved oral health. The heat-killed bacterial cells are also known to exert antiallergic and antitumor effects, in part due to the potent induction of IL-12 and interferons by lipoteichoic acids on the bacterial cell wall surfaces. Furthermore, the L-137 strain has higher levels of lipoteichoic acids exposed on the surface, with higher amounts of alanine, than the closely related L. plantarum JCM1149 strain; this correlates with the higher induction of IL-12 by the L-137 strain than by the JCM1149 strain.

Another group of lactic acid-producing probiotic bacteria includes several unique strains of the spore-forming Bacillus coagulans (previously classified as L. sporogenes).
The teichoic acid from *B. coagulans* walls has a higher lipid content than most Gram-positive bacteria, and is a glycerophosphate polymer substituted with two neutral sugars, glucose and galactose. It is unique in lacking amino acid substituents, otherwise considered a characteristic of teichoic acids.\(^{26}\)

Cell walls from the live *B. coagulans* GBI-30, 6086 strain (GanedenBC\(^{26}\)) have demonstrated immune modulating and anti-inflammatory effects in vitro.\(^{27}\) Our team previously showed that immune-modulating effects of the BC30 strain were associated both with the cell wall fraction and with the metabolites produced by the live bacteria in vitro.\(^{28}\) The probiotic strain was further shown to prolong the survival and reduce symptoms in mice infected with *Clostridium difficile*.\(^{29,30}\) Clinical studies showed that consuming BC30 helped alter the gut microbiome by increasing the numbers of beneficial bacteria,\(^{31}\) and ex vivo testing of blood from elderly humans who had consumed BC30 for 28 days showed increased anti-inflammatory cytokine responses.\(^{32}\) Results from a recent clinical trial suggest that the consumption of BC30 supports exercise performance and helps reduce exercise-induced muscle damage.\(^{33}\)

Recently, inactivated *B. coagulans* GBI-30, 6086 cells have been produced for oral consumption. The work presented here was undertaken to document whether the immune activating and anti-inflammatory properties associated with the cell walls of the live BC30 bacteria were protected in the inactivated product. An important focus for this work was to document the biological activities of the inactivated *B. coagulans* GBI-30 bacterial cells when presented to human immune cells in a cell culture system that allows cross-talk between antigen-presenting monocytes and dendritic cells with lymphocytes and natural killer cells, and thus mimics events in the gut mucosal immune tissue.

**Materials and methods**

**Reagents**

Phosphate-buffered saline, Roswell Park Memorial Institute 1640 (RPMI-1640) medium, penicillin–streptomycin 100×, IL-2, and LPS were purchased from Sigma-Aldrich Co. (St. Louis, MO, USA). CD69 fluorescein isothiocyanate, CD56 phycoerythrin, CD3 peridinin chlorophyll protein, CD25 brilliant violet 421 and heparin Vacutainer vials were purchased from BD Biosciences (Franklin Lakes, NJ, USA). The Bio-Plex Pro\(^{TM}\) human cytokine 27-Plex was purchased from Bio-Rad Laboratories Inc. (Hercules, CA, USA).

**Inactivated *B. coagulans* GBI-30 (Staimune\(^{TM}\))**

Inactivated *B. coagulans* GBI-30, 6086 cells were provided by Ganeden Inc. (Mayfield Heights, OH, USA). Bacterial cell numbers were 15 billion (1.5×10\(^{10}\)) CFU per gram prior to inactivation. Inactivated bacteria were diluted in physiological saline and added to human peripheral blood mononuclear cell cultures at doses from 0.78×10\(^{-9}\)–100×10\(^{-9}\) inactivated cells/mL cell culture.

**Peripheral blood mononuclear cell cultures**

Healthy human volunteers between the age of 50 and 60 years served as blood donors upon informed consent, as approved by the Sky Lakes Medical Center Institutional Review Board (FWA 2603). Freshly drawn peripheral venous blood samples in sodium heparin were layered onto Lympholyte-Poly and centrifuged for 35 minutes at 450 g. The upper peripheral blood mononuclear cells (PBMC)-rich interface was harvested using sterile transfer pipettes into new vials and washed twice with 10 mL PBS, without calcium or magnesium, by centrifugation for 10 minutes. The cells were resuspended into RPMI 1640 with 10% fetal bovine serum, l-glutamine, and antibiotics (penicillin and streptomycin) to a cell density of 10\(^{6}\)/mL. Triplicate cultures were established for each of the eight doses of inactivated GBI-30 tested. Untreated cell cultures (negative controls) were established in hexaplicate. Two sets of positive control cultures were established in triplicates, one set using LPS (10 ng/mL) and another set using IL-2 (100 IU/mL) to activate the immune cells by two different pathways.

**Cytokine testing**

Supernatants were harvested from the human immune cell 24-hour cultures, and the levels of 27 cytokines, chemokines, and growth factors were analyzed. Testing was performed on culture supernatants from cell cultures treated with the six higher doses of inactivated GBI-30. The following markers: IL-1β, IL-1ra, IL-2, IL-4, IL-5, IL-6, IL-7, IL-8, IL-9, IL-10, IL-12 (p70), IL-13, IL-15, IL-17, eotaxin, basic FGF, G-CSF, GM-CSF, Interferon-gamma (IFN-γ), IP-10, MCP-1, MCAF, MIP-1α, MIP-1β, PDGF-BB, RANTES, TNF-α, and VEGF were quantified using Bio-Plex protein arrays (Bio-Rad Laboratories Inc.) and utilizing xMAP technology (Luminex, Austin, TX, USA).

**Statistical analysis**

Averages and SDs for each data set were calculated using Microsoft Excel. Statistical analysis was performed using
the 2-tailed, independent t-test. Statistical significance was indicated when \( P<0.05 \) and a high level of significance when \( P<0.01 \).

**Results**

**Immune cell activation**

Immune cell activation by inactivated *B. coagulans* GBI-30 cells was determined by measuring cell-surface expression of the activation marker CD69. The gating on immune cells with different forward and side scatter properties allowed analysis of CD69 expression on lymphocytes versus monocytes/macrophages (Figure 1). Treatment of both cell types with inactivated GBI-30 cells for 24 hours resulted in activation across a broad dose range. The results for CD69 expression on lymphocytes showed that even at the lowest dose the CD69 expression was not returning to baseline, and suggests that much lower doses would still have been able to activate lymphocytes (Figure 1A). In contrast, the most robust and statistically significant activation of monocytes was in a narrower dose range, returning toward baseline at the lowest dose shown (Figure 1B).

![Figure 1](https://www.dovepress.com/)

**Figure 1** Expression of the CD69 cellular activation marker on lymphocytes and monocytes.

**Notes:** *P<0.05; **P<0.01. CD69 expression on lymphocytes (A) and monocytes (B) in human PBMC cultures treated for 24 hours with serial dilutions of inactivated *Bacillus coagulans* GBI-30 cells. Mean fluorescence intensity for CD69 expression is shown. Data presented as mean ± SD from triplicate cultures and represent one of three separate experiments using PBMC from three different healthy human donors. For lymphocytes, the mean ± SD for the controls were: UT 1,432±32, LPS 1,635±33, and IL-2 1,910±115. For monocytes, the mean±SD for the controls were: UT 9,448±373, LPS 12,803±286, and IL-2 11,494±1,455.

**Abbreviations:** LPS, lipopolysaccharide; PBMC, peripheral blood mononuclear cells; UT, untreated.
The use of fluorescently labeled antibodies to CD3, CD56, and CD69 allowed the monitoring of changes to lymphocyte subsets, including CD3+ T lymphocytes, CD3+ CD56+ Natural Killer T (NKT) cells, CD3-CD56+ NK cells, and non-T non-NK lymphocytes. Treatment of PBMC cultures with inactivated GBI-30 cells for 24 hours led to the activation of T lymphocytes, NKT cells, NK cells, and non-T non-NK cells (Figure 2). T lymphocyte activation was seen across a broad dose range, and at the third-lowest dose (3.13x10^6 bacteria/mL), the CD69 expression was as robust as for LPS-induced CD69 expression (Figure 2A). The T lymphocyte activation remained highly significant even at

**Figure 2** Expression of the CD69 cellular activation marker on immune cell subsets.

Notes: *P*<0.05; **P**<0.01. CD69 expression on T lymphocytes (A), NKT cells (B), NK cells (C), and non-T non-NK cells (D) in human PBMC cultures treated for 24 hours with serial dilutions of inactivated Bacillus coagulans GBI-30 cells. Mean fluorescence intensity for CD69 expression is shown. Data presented as mean ± SD from triplicate cultures and represent one of three separate experiments using PBMC from three different healthy human donors. For T lymphocytes, the mean ± SD for the controls were: UT 1,192 ± 23, LPS 1,346 ± 20, and IL-2 1,638 ± 97. For NKT cells, the mean ± SD for the controls were: UT 1,908 ± 36, LPS 2,687 ± 126, and IL-2 3,669 ± 493. For NK cells, the mean ± SD for the controls were: UT 2,880 ± 259, LPS 4,770 ± 503, and IL-2 8,930 ± 1,613. For non-T non-NK cells, the mean±SD for the controls were: UT 2,364 ± 146, LPS 3,393 ± 137, and IL-2 2,935 ± 184.

**Abbreviations:** LPS, lipopolysaccharide; PBMC, peripheral blood mononuclear cells; UT, untreated.
the lowest dose of inactivated GBI-30. Activation of NKT cells was also observed across the entire dose range, and at some doses, the activation was highly significant, compared to untreated control cultures (Figure 2B). NK cell activation was most robust at the lower doses and less prominent at higher doses (Figure 2C). The CD69 expression in the non-T non-NK lymphocyte population was seen for a broad dose range as well, returning to baseline at the lowest dose tested (Figure 2D). LPS was used as a positive control (10 ng/mL) and resulted in an increase in CD69 expression on all cell types. IL-2 was used as a second positive control (100 IU/mL) and also showed an increase in CD69 on all cell types.

Please note that occasionally, a large variation in CD69 expression was seen within one set of triplicate cultures, as reflected by large error bars, and in some cases, an average response in the triplicate set appears out of line with the overall dose response. We suggest that this is due to the nature of the test product, where the inactivated bacterial cells may clump within a culture well instead of dispersing, thus not providing optimal interaction between bacterial cells and PBMC within those wells. An example is the very low CD69 expression on non-T non-NK lymphocytes at the dose of 12.5×10^6 inactivated GBI-30/mL, where only one of the three triplicate culture wells showed the CD69 expression level comparable to the dose above and the dose below (Figure 2D).

**Imune-activating cytokines** *B. coagulans*

Supernatants from the PBMC cultures exposed to inactivated *B. coagulans* GBI-30 cells for 24 hours were simultaneously assayed for the levels of 27 different cytokines, chemokines, and growth factors, using a magnetic bead-based array and Luminox xMAP technology. Increases in the levels of cytokines with various immune activating and regulating properties were seen. This included a robust upregulation of certain proinflammatory cytokines, including IL-1β, IL-6, IL-17, and TNF-α (Figure 3). Increases were also seen for the cytokines IL-4, IL-7, IL-8, IL-9, and IL-12p70 (data not shown). Furthermore, increases were seen for four biomarkers involved in antiviral immune defense activity, namely IFN-γ, and the three chemokines MCP-1, MIP-1α, and MIP-1β (Figure 4).

**Anti-inflammatory cytokines**

In parallel to increases in immune-activating, proinflammatory cytokines, higher doses of the inactivated *B. coagulans* GBI-30 cells also triggered robust increases in the two anti-inflammatory cytokines IL-1ra and IL-10 (Figure 5). The increases were comparable between these cytokines, with an approximal 300-fold increase above the levels in untreated cell cultures.

**Growth factors**

The exposure of human PBMC to inactivated GBI-30 triggered changes in growth factor production (Figure 6). Highly selective changes were seen for the growth factors granulocyte-colony stimulating factor (G-CSF) and granulocyte macrophage-colony stimulating factor (GM-CSF), both having differential effects on stem cell biology (Figure 6A and B), where treatment of PBMC with inactivated GBI-30 led to a very strong increase in G-CSF production, in contrast to a mild reduction in GM-CSF production.

**Discussion**

The work reported here on the inactivated *B. coagulans* GBI-30, 6086 cells is a direct extension of previous work by our team, involving the live *B. coagulans* GBI-30, 6086. Our previous results have demonstrated that the cell wall from BC30 has complex biological properties, including both immune-activating and anti-inflammatory properties.

The current work showed that the inactivated *B. coagulans* GBI-30 cells triggered an increase in the CD69 activation marker on multiple human immune cell types, whereas previous work focused only on NK cells. CD69 is an early activation marker on many cell types, and is directly involved in the molecular apparatus responsible for NK cell-mediated killing of virus-infected and transformed cells. In addition, CD69 has a broad range of functions, including T-cell/B-cell interactions, homing of cells into appropriate tissue environments, and the process of generating and maintaining immunological memory. The ability of inactivated GBI-30 to induce CD69 on multiple cell types suggests a broad effect involving cells from both the innate and the adaptive immune systems. Further work will need to evaluate the various cell types in the non-T non-NK subset of cells, which includes dendritic cells, B lymphocytes, circulating hematopoietic, mesenchymal, and endothelial stem cells. Previous work on live BC30 showed that exposure of human mononuclear phagocytes triggered a favorable maturation of antigen-presenting cells toward both macrophage and dendritic cell phenotypes. This is important, since dendritic cells are first-line antigen-presenting cells in gut mucosal immune tissue, capable of unique surveillance activity and antigen recognition across intact epithelial barriers, and therefore may represent the cell type that initially encounters consumed BC30 cells in the gut mucosa in vivo.
Inactivated *B. coagulans* GBI-30 cells triggered robust increases in the production of multiple cytokines, chemokines, and growth factors. The current data span a broader range of cytokines, chemokines, and growth factors than previously tested for live BC30. This has helped confirm the potent immune-activating properties of BC30. It has also helped demonstrate novel biological properties associated with antiviral and regenerative functions. The increases in the biomarkers IFN-γ, MCP-1, MIP-1α, and MIP-1β, involved in antiviral immune defense mechanisms and cellular recruitment, are of special importance. IFN-γ has direct antiviral properties, activates macrophages, and enhances NK cell killing activity of transformed cells. The three chemokines facilitate recruitment of immune cells to sites of inflammation, whether caused by injury or by infection. The increase in two anti-inflammatory cytokines, IL-1ra and IL-10, points to the complexity of inactivated GBI-30 immune modulation.

With the purpose of resolving the initial proinflammatory immune activation and limiting the inflammatory process in space and time.

Several growth factors known to play specific roles in endogenous regeneration were upregulated in the PBMC cultures. Highly selective, contrasting effects were seen for the two stem cell growth factors G-CSF and GM-CSF. Inactivated *B. coagulans* GBI-30-treated PBMC cultures showed a robust increase in G-CSF levels, reaching over 7,000-fold above untreated cultures at the highest dose, in contrast to mildly reduced GM-CSF levels. This is an important differentiation, since G-CSF supports stem cells to produce neutrophils, whereas GM-CSF promotes the production of multiple cell types, including eosinophils, involved in immune defense against multicellular parasites and also in inflammation in allergies and asthma. In addition, G-CSF is used therapeutically to support stem cell mobilization, migration, and tissue repair. Therefore, the selective

**Figure 3** Changes in proinflammatory cytokine levels in human PBMC cultures.

**Notes:** *P*<0.05; **P**<0.01. Changes in cytokine levels in human PBMC cultures treated for 24 hours with serial dilutions of inactivated *Bacillus coagulans* GBI-30 cells are shown as percent change from untreated cell cultures. Data presented as mean ± SD from duplicate testing of culture supernatants from one of three separate experiments using PBMC from three different healthy human donors.

**Abbreviation:** PBMC, peripheral blood mononuclear cells.
The effect of inactivated *B. coagulans* GBI-30 cells on growth factor production may be directly beneficial in repair and regeneration of the gut mucosal tissue, for example in situations of ulceration. The effect may also affect systemic changes via many cell types, including inflammation-modulating mesenchymal stem cells. Mesenchymal stem cells are able to sense signals from, and migrate into injured, inflamed, and ischemic tissue. They can cross the blood–brain barrier and contribute to repair of brain injuries such as stroke. Interestingly, it has been shown that the treatment of mesenchymal stem cells from healthy human donors with IL-1, TNF-α, and IFN-γ production contributed to a strong increase in G-CSF production by the mesenchymal stem cells and subsequently the increase in G-CSF reprogrammed LPS-activated microglial cells to secrete fewer inflammatory mediators. This sequential process may apply to events when mononuclear cells, which include multiple types of stem cells, are exposed to inactivated *B. coagulans* GBI-30 cells. Further work is needed to document whether a cascade of events is triggered by inactivated GBI-30, where an initial immune-activating signal upregulated the production of proinflammatory cytokines, followed by anti-inflammatory processes intended to resolve the inflammation, combined with reparative growth factors.

Three observations from the current work reported here are of key importance in the validation of similar biological activities of the inactivated *B. coagulans* GBI-30 cells, compared to the immune-modulating properties of cell wall fractions from living BC30: 1) Both can increase the CD69 activation marker on lymphocytes, 2) both are able to increase production of the proinflammatory cytokine IL-6, and 3) both are capable of increasing the anti-inflammatory cytokine IL-10. This suggests that cell wall components, including lipoteichoic acid, have remained at least partially preserved by the inactivation process.

A direct dose comparison to our previous in vitro work is not feasible. Previous work on the cell wall’s immune-
activating properties was performed on material that went through repeated freeze/thaw and bead-milling cycles, thus breaking down the cell walls into smaller particles, each capable of engaging appropriate receptors on the surface of the PBMC. In contrast, the current work involved the addition of intact bacterial cells to the human immune cell cultures.27

The complex properties of inactivated GBI-30 suggest possible multifaceted clinical responses after consumption, involving immune activation, anti-inflammatory effects, and effects involving stem cell mobilization, homing, and reprogramming involved in accelerated repair. The direct effects of inactivated GBI-30 are expected to translate to immune activation at the level of the gut mucosa and trigger rapid systemic effects. This is different from a study on the live BC30 where ingested spores will give rise to living bacteria that can colonize the intestinal tract, and where an important part of the biological effects is due to secreted bacterial metabolites. Future work on inactivated B. coagulans GBI-30 cells should include a human clinical study to examine acute effects, using the study design previously published by our team on efficacious immune-modulating natural products.44,45

Figure 5 Changes in anti-inflammatory cytokine levels in human PBMC cultures treated for 24 hours.
Notes: *P<0.05; **P<0.01. The first sentence should read: Changes in IL-1ra (A) and IL-10 (B) cytokine levels in human PBMC cultures treated for 24 hours with serial dilutions of inactivated Bacillus coagulans GBI-30 cells are shown as percent change from untreated cell cultures. Data presented as mean ± SD from duplicate supernatants from one of three separate experiments using PBMC from three different healthy human donors.
Abbreviation: PBMC, peripheral blood mononuclear cells.

Figure 6 Changes in growth factor levels in human PBMC cultures treated for 24 hours.
Notes: **P<0.01. Changes in G-CSF (A) and GM-CSF (B) growth factor levels in human PBMC cultures treated for 24 hours with serial dilutions of inactivated Bacillus coagulans GBI-30 cells are shown as percent change from untreated cell cultures. Data presented as mean ± SD from duplicate supernatants from one of three separate experiments using PBMC from three different healthy human donors.
Abbreviations: G-CSF, granulocyte-colony stimulating factor; GM-CSF, granulocyte macrophage-colony stimulating factor; PBMC, peripheral blood mononuclear cells.
as well as nutraceutical products that have effects on human stem cell biology.46,47

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Author contributions
All authors contributed toward data analysis, drafting and critically revising the paper and agree to be accountable for all aspects of the work.

Disclosure
HAC, SF, and DK are employees of the sponsor of the study, Ganeden Inc. SF has also been a consultant for Ganeden Inc. GSJ reports no conflicts of interest in this work.

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