

Pro-Intensity is a powerful advanced combination of **16 selected probiotic strains** (10 human, 2 plant and 4 dairy), bovine **colostrum** (high proline-rich polypeptide content), **inulin** (chicory), **A.O.S.** (larch) and xylooligosaccharides (XOS). It provides a minimum of 20 billion viable bacteria per capsule with GPSTM enteric coating for optimal protection against the stomach's acid secretions.

Ingredients: Potato starch, bacterial culture (20 billion live active, healthy cells per capsule, see nutritional information), bovine colostrum (from *Bos taurus*) (milk), inulin (from chicory root, *Cichorium intybus*), arabinogalactan (from *Larix laricinia*), xylooligosaccharides, L-ascorbic acid (vitamin C), anti-caking agent (magnesium salts of fatty acids and silicon dioxide), GPSTM enteric coated vegetable capsule (glazing agent: hydroxypropylmethylcellulose; aqueous enteric-coating solution; purified water).

Nutritional information:	1 capsule (392 mg)	Size and format:
Lactobacillus rhamnosus UB5115 ¹	7,427 billion CFU	30 enteric-coated vegetable capsules
Lactobacillus crispatus UB4719 ¹	1,903 billion CFU	
Lactobacillus casei UB1499 ¹	1,887 billion CFU	
Bifidobacterium animalis ssp. lactis UB3963 ¹	1,427 billion CFU	
Lactobacillus gasseri UB8141 ¹	1,427 billion CFU	Recommended daily dose: 1–2 capsules daily. If you are taking antibiotics, take this product at least 2–3 hours before or after taking them. Do not exceed the stated recommended daily dose. Store preferably refrigerated.
Bifidobacterium bifidum UB4280 ¹	951 million CFU	
Bifidobacterium breve UB8674 ¹	315 million CFU	
Bifidobacterium longum ssp. infantis UB9214 ¹	315 million CFU	
Bifidobacterium longum ssp. longum UB7691 ¹	315 million CFU	
Lactobacillus acidophilus UB5997 ¹	26 million CFU	
Lactobacillus salivarius UB4198 ²	1,427 billion CFU	
Lactobacillus plantarum UB2783 ²	73 million CFU	
Lactobacillus johnsonii UB3394 ³	1,903 billion CFU	
Lactobacillus helveticus UB7229 ³	539 million CFU	
Lactobacillus paracasei UB1978 ³	52 million CFU	
Lactococcus lactis LL-23 ³	13 million CFU	
Colostrum (high content of proline-rich polypeptides	25 mg	
Inulin	10 mg	
Arabinogalactan (AOS)	10 mg	
Xylooligosaccharides (XOS)	10 mg	
Vitamin C (L-ascorbic acid)	6 mg (15% NRV*)	
Source of strains: ¹ human / ² plant / ³ dairy.		
CFU: Colony-Forming Unit Cells		
NRV: Nutrient Reference Value in %.		

The **GPS[™] enteric coating** protects contents from stomach acids and delivers 100% potency to the intestines.

Indications and uses:

- Crohn's disease
- Hypercholesterolaemia
- Improves the immune system and digestive function
- Antibiotic-associated diarrhoea

- Ulcerative colitis
- Diabetes mellitus
 - Inflammatory bowel disease

Cautions:

Consult a health-care practitioner before using if you have fever, vomiting, bloody diarrhoea, or severe abdominal pain. Discontinue use if symptoms of digestive upset (diarrhoea) persist or worsen beyond 3 days. Consult a health-care practitioner if you have an immune-compromised condition (e.g. lymphoma or AIDS).



DETAILS:

PRO-INTENSITY contains a selection of scientifically proven probiotic strains, supplementary prebiotics of natural origin and a colostrum extract. It is the latest development in probiotic supplementation and is the ideal product to improve the immune system and digestive function, being essential for good health and disease resistance.

Each capsule contains more than 20 billion live cells, with a strain selection of 16 beneficial strains, including 10 of human origin. Although the origin of a probiotic strain is not the sole criterion for efficacy, strains of human origin exhibit the ability to colonise at multiple sites in the gastrointestinal tract.

The activity of the strains helps to boost immune function, disease resistance, optimal digestion and absorption of nutrients, improved vitamin synthesis, better lactose tolerance, and improved gastrointestinal transit.

Bovine colostrum from high quality sources has a high proline-rich polypeptide content. Specific immunoglobulins (IgG) and growth factors (IGF) in bovine colostrum exert their beneficial effects on the intestine, with IgGs destroying pathogenic bacteria and IGFs improving the intestinal mucosa lining.

The enteric coating of the capsule protects the product from gastric juices and ensures 100% potency.

INGREDIENTS:

<u>LACTOBACILLUS RHAMNOSUS</u>: this product contains the UB5115 human strain. It is one of the most widely researched probiotic species due to its tolerance to acidic conditions. This product contains more than 7.4 billion colony-forming units (CFUs) from this species.

It colonises the intestinal membranes, providing numerous health benefits: it increases lactic acid production, actively suppressing the growth of harmful bacteria such as *Salmonella* ⁽¹⁾. It is effective in preventing antibiotic-associated diarrhoea ⁽²⁾ and *Clostridium difficile*-associated diarrhoea ⁽³⁾. It strengthens the immune system and is a good coadjuvant for the influenza vaccine ⁽⁴⁾. It improves intestinal barrier function for the relief of autoimmune diseases such as arthritis ⁽⁵⁾ and allergies ⁽⁶⁾. It improves the blood lipid profile ⁽⁷⁾ and reduces cholesterol ⁽⁸⁾. It may prevent or relieve symptoms of post-partum depression and anxiety ⁽⁹⁾, regenerate the vaginal flora in women by reducing colonisation by oral bacteria and fungi ⁽¹⁰⁾, and may reduce the prevalence of gestational diabetes mellitus ⁽¹¹⁾. In children, it reduces the frequency and duration of diarrhoea and vomiting ⁽¹²⁾, rotavirus diarrhoea ⁽¹³⁾, and antibiotic-associated diarrhoea ⁽¹⁴⁾. It reduces the incidence of atopic dermatitis^(15, 16). Drinking milk supplemented with *L. rhamnosus* reduces the risk of tooth decay in children ⁽¹⁷⁾.

<u>LACTOBACILLUS CRISPATUS</u>: this product contains the UB4719 human strain. Numerous studies have shown its considerable potential for maintaining the health of the female reproductive system, helping to prevent recurrent urinary tract infections, as well as bacterial vaginosis and candidiasis ⁽¹⁸⁻²⁰⁾. It is also able to modulate the immune system ⁽²¹⁾ and reduce allergic symptoms in mice ⁽²²⁾.

<u>LACTOBACILLUS CASEI</u>: this product contains the UB1499 human strain. It reduces the duration and incidence of infections such as bronchitis, pneumonia and rhinopharyngitis ⁽²³⁻²⁵⁾. Regarding intestinal infections, it boosts immunity against bacterial infections (e.g. *Escherichia coli*) and viral infections (e.g. influenza vaccinations) ⁽²⁶⁻²⁹⁾.

In children, it improves allergic rhinitis symptoms ⁽³⁰⁾, helps eradicate *Helicobacter pylori* in conjunction with antibiotic therapy ⁽³¹⁾, is effective against viral diarrhoea⁽³²⁾, and reduces the general incidence of infections ⁽³³⁾.

<u>BIFIDOBACTERIUM ANIMALIS subsp. LACTIS</u>: this product contains the UB3963 human strain. It helps reduce constipation and bloating in children and adolescents with irritable bowel syndrome ⁽²⁴⁾. It boosts the immune system by increasing levels of NK (natural killer) cells and polymorphonuclear leukocytes ⁽²⁵⁾. It helps to repair the permeability of the intestinal barrier by enhancing apical junction proteins and goblet cell population ⁽²⁶⁾. It reduces abdominal visceral fat in overweight people with metabolic disorders and has beneficial effects on weight control and metabolic health ^(27, 28). It also improves glucose intolerance in animals ⁽²⁹⁾.

Pro-Intensity Code: FE2018 - 30 enteric-coated vegetable capsules



LACTOBACILLUS GASSERI: this product contains the UB8141 human strain. It improves functional dyspepsia by improving gastric microbiota by helping to suppress *Helicobacter pylori* in the stomach ⁽³⁰⁾. It is also a predominant species in the vaginal flora, inhibits the adherence of pathogenic bacteria and helps in the prevention and treatment of bacterial vaginosis ⁽³¹⁾. It has antimicrobial activity through the production of bacteriocins ^(32, 33), improves symptoms such as diarrhoea in Irritable Bowel Syndrome ^(34, 35), helps boost the immune system ⁽³⁶⁾ and may help regulate allergic response ⁽³⁷⁾. Its effect on weight control has been studied in recent years. It has a reducing effect on abdominal adiposity, body weight and other measures of obesity and helps to regulate blood lipids (triglycerides, cholesterol), suggesting its beneficial impact on metabolic disorders ⁽³⁸⁻⁴⁰⁾.

<u>BIFIDOBACTERIUM BIFIDUM</u>: this product contains the UB4280 human strain. They are found in the mucosal lining of the last part of the small bowel and are the predominant strains that colonise the large bowel and support bowel health, hygiene, and functionality. They reduce serum cholesterol and dissolve bile salts ^(41, 42). *B. bifidum* has been shown to exert antibacterial activity against *Helicobacter pylori* ^(43,44), reduce apoptosis in the intestinal epithelium of children with necrotising enterocolitis ⁽⁴⁵⁾, regulate the immune system response⁽⁴⁶⁻⁴⁸⁾, reduce the duration and severity of colds ⁽⁴⁷⁾, provide anti-inflammatory activity in chronic diseases of the large bowel (e.g. irritable bowel syndrome) ^(49, 50), and reduce the incidence of radiotherapy-induced diarrhoea in cervical cancer patients ⁽⁵¹⁾.

<u>BIFIDOBACTERIUM BREVE</u>: this product contains the UB8674 human strain. It maintains colon homeostasis by reducing inflammation through induction of intestinal IL-10-producing Tr1 cells ⁽⁵²⁾. It protects colon function, relieves constipation, and reduces gas, bloating, and diarrhoea ^(52,53). It improves ulcerative colitis symptoms ⁽⁵⁴⁾. It also stimulates the immune system^(53,55), inhibits *Escherichia Coli* ⁽⁵⁶⁾ and suppresses the Candida fungus ⁽⁵⁷⁾. It reduces fat, liver function, and systemic inflammation in people prone to obesity ⁽⁵⁸⁾. In neonates, it improves gastrointestinal problems by stabilising the intestinal flora ⁽⁵⁹⁾ and reduces the incidence of necrotising enterocolitis ⁽⁶⁰⁾. In children with coeliac disease, it reduces the pro-inflammatory cytokine TNF-alpha ⁽⁶¹⁾. It improves adverse effects in chemotherapy patients, such as fever, infections, and intestinal disorders ⁽⁶²⁾.

<u>BIFIDOBACTERIUM LONGUM subsp. INFANTIS</u>: this product contains the UB9214 human strain. It is the dominant probiotic inhabiting the distal part of the small bowel and colon. It is one of the first species to colonise the infant gastrointestinal tract ⁽⁶³⁾ and is critical in adults for intestinal health and immune system function ⁽⁶⁴⁾. It is extremely good at surviving stomach and bile acids ⁽⁶⁵⁾ and is typically able to adhere to intestinal tissues ⁽⁶⁶⁾. It produces acetic acid and inhibits pathogenic bacteria ⁽⁶⁷⁾. It produces bacteriocins, which act against *Salmonella, Shigella*, and *E. coli* ^(68, 69). It relieves many symptoms of Irritable Bowel Syndrome (IBS) (e.g. pain, bloating), normalises bowel movements, and regulates the IL-10/IL-12 ratio ⁽⁷⁰⁻⁷²⁾. It reduces systemic pro-inflammatory biomarkers in chronic inflammatory diseases such as ulcerative colitis, chronic fatigue syndrome, and psoriasis, demonstrating that the immunomodulatory effects of microbiota are not limited to the mucosa but encompass the systemic immune system (⁽⁷³⁾. It can alleviate symptoms of untreated coeliac disease ⁽⁷⁴⁾.

<u>BIFIDOBACTERIUM LONGUM</u> subsp. LONGUM: this product contains the UB7691 human strain. A protein factor produced by *B. longum* inhibits the adhesion of the enterotoxigenic strain of *Escherichia coli* ⁽⁷⁵⁾. It has antiinflammatory properties and is indicated for gastrointestinal disorders such as ulcerative colitis ⁽⁷⁶⁾, antibioticassociated diarrhoea ^(77, 78), Irritable Bowel Syndrome ⁽⁷⁹⁾, and seasonal allergies ^(80, 81). It aids the formation of lactic acid and formic acid, lowering the pH of the intestines and preventing the proliferation of harmful bacteria ⁽⁸²⁾. It is also a significant producer of B vitamins ⁽⁸³⁾.

LACTOBACILLUS ACIDOPHILUS: this product contains the UB5997 human strain. It improves the general symptoms of patients with Irritable Bowel Syndrome ⁽⁸⁴⁾. It helps to maintain an acidic environment in the intestinal tract by preventing the growth of harmful bacteria and reduces antibiotic-associated diarrhoea ⁽⁸⁵⁾. It reduces total plasma cholesterol and low-density lipoprotein (LDL) cholesterol ^(86, 87). It helps to improve digestive health by maintaining the intestinal barrier, restoring intestinal flora, improving digestion, boosting the immune system, and supporting beneficial bacteria that thrive in the colon ⁽⁸⁸⁾. It helps to improve symptoms of allergic rhinitis ⁽⁸⁹⁾, hay fever ⁽⁹⁰⁾, and atopic dermatitis ⁽⁹¹⁾.

<u>LACTOBACILLUS SALIVARIUS</u>: this product contains the UB5997 plant strain. It inhibits the growth and activity of harmful pathogenic bacteria, including *Helicobacter pylori* ^(92,93) and *Salmonella* ⁽⁹⁴⁾. It helps to break down undigested proteins and deactivate toxins produced by intestinal putrefaction ⁽⁹⁵⁾. It improves the lipid (cholesterol) profile and reduces inflammation, tumour necrosis factor, and *Escherichia coli* populations ⁽⁹⁶⁾. When used in combination with prebiotics (fructooligosaccharides), it is effective in reducing the symptoms of atopic dermatitis in children ⁽⁹⁷⁾ and adults ⁽⁹⁸⁾.

New Roots

<u>LACTOBACILLUS PLANTARUM</u>: this product contains the UB2783 plant strain. It acts against unwanted bacteria by improving the symptoms of Irritable Bowel Syndrome, such as excessive gas, bloating and abdominal discomfort ⁽⁹⁹⁻

¹⁰³), and ulcerative colitis ^(104, 105). It regulates immune response and is beneficial in the treatment of atopic dermatitis in children ⁽¹⁰⁶⁾. It has immunostimulatory effects in the elderly, reducing the number of infections ⁽¹⁰⁷⁾. It improves gastrointestinal symptoms during antibiotic therapy ⁽¹⁰⁸⁾. It reduces cardiovascular risk factors and may be useful as a protective agent in the primary prevention of atherosclerosis in smokers ⁽¹⁰⁹⁾. In adults with hypercholesterolaemia, it lowers cholesterol and high blood pressure, which, as a result, may reduce the risk of cardiovascular diseases ⁽¹¹⁰⁾. It improves symptoms of lactose intolerance, such as diarrhoea and flatulence, in combination with another probiotic ⁽¹¹¹⁾. Together with other *Lactobacillus* species, it can restore the vaginal flora by improving the pH and diagnosis of bacterial vaginosis when administered orally ⁽¹¹²⁾.

<u>LACTOBACILLUS JOHNSONII</u>: this product contains the UB3394 dairy strain. It has several benefits, such as in *Helicobacter pylori* gastritis ⁽¹¹³⁾, regulates immune response ⁽¹¹⁴⁾, may help in the control of diabetes ⁽¹¹⁵⁾, is helpful against vaginal infections ⁽¹¹⁶⁾, and improves allergic rhinitis in children ⁽¹¹⁷⁾.

<u>LACTOBACILLUS HELVETICUS</u>: this product contains the UB7229 dairy strain. It protects the gastrointestinal tract, strengthening the systemic humoral and intestinal mucosal immune response in elite athletes ⁽¹¹⁸⁾. It has been shown to cause an antidepressant effect in animals, probably due to the microbiota-gut-brain axis connection ⁽¹¹⁹⁾. Fermented milk with *L. helveticus* improves cognitive function ⁽¹²⁰⁾ and lowers blood pressure ⁽¹²¹⁾. In animals, it increases bone density and bone mineral content ⁽¹²²⁾, and in post-menopausal women, it has a positive effect on calcium metabolism ⁽¹²³⁾. It controls unwanted intestinal micro-organisms and bacteria (*Salmonella enteritidis, Campylobacter jejuni, Escherichia coli, Candida albicans*, etc.), regulates immune response and reduces lactose intolerance ⁽¹²⁴⁾.

<u>LACTOBACILLUS PARACASEI</u>: this product contains the UB1978 dairy strain. It significantly enhances the specific immune response in healthy people who have received the influenza vaccine ⁽¹²⁵⁾. It improves digestive function ⁽¹²⁶⁾ and symptoms (especially eye symptoms) in patients with allergic rhinitis treated with oral antihistamines ⁽¹²⁷⁾. It is also effective against *Staphylococcus aureus, Escherichia coli*, and *Salmonella* infections ⁽¹²⁸⁻¹³⁰⁾. It relieves the frequency and duration of acute diarrhoea in children ⁽¹³¹⁾. It improves neurocognitive function in patients with chronic fatigue syndrome when used in combination with other probiotics ⁽¹³²⁾.

<u>LACTOCOCCUS LACTIS</u>: this product contains the LL-23 dairy strain. It produces bacteriocins such as lacticin, nisin and lactococcin ⁽¹³³⁾. Nisin is the best studied compound in this group. Nisin is a so-called lantibiotic bacteriocin with a broad spectrum of antimicrobial activity and an immunomodulatory effect ⁽¹³⁴⁾. One of the most important properties of nisin is its activity against Gram-positive bacteria and bacterial spores such as *Clostridium difficile* ⁽¹³⁵⁾. *Lactococcus lactis* also boosts antiviral immunity by reducing cold and flu symptoms ^(136, 137), may help lower blood pressure ⁽¹³⁸⁾, and may help reduce intestinal inflammation ⁽¹³⁹⁾, among other properties ⁽¹⁴⁰⁾.

<u>L. lactis LL-23 strain</u>: together with other probiotics, it reduces inflammatory markers in people with rheumatoid arthritis⁽¹⁴¹⁾, and also together with other probiotics and diet helps to significantly reduce abdominal fat and increased antioxidant enzyme activity⁽¹⁴²⁾.

<u>CALOSTRO</u>: contains high levels of proline-rich polypeptides (PRP's) that help reduce the inflammatory response responsible for some of the symptoms associated with Irritable Bowel Syndrome and Leaky Gut Syndrome (intestinal dysbiosis). It contains a high proportion of immunoglobulin (IgG), antimicrobial factors (lactoferrin), immunomodulatory polypeptides, anti-inflammatory cytokines, growth factors and other bioactive compounds that promote immune response, inhibit excessive production of "reactive oxygen species" and act in synergy as prebiotics for the intensive growth of specific probiotic strains. Growth factors are involved in the regeneration and proliferation of the intestinal epithelium for proper intestinal absorption and permeability ⁽¹⁾. Proline-rich polypeptides are one of the most important components of colostrum because of their ability to modulate the immune system and regulate the production of certain cytokines, signalling molecules that control the inflammatory process ^(143, 144).

Clinical studies show that bovine colostrum regulates the immune response after exercise ^(145, 146), reduces muscle damage and inflammation after exercise ⁽¹⁴⁷⁾, has a protective effect on the respiratory tract mucosa ^(148, 149), is effective in HIV treatment-associated diarrhoea ⁽¹⁵⁰⁾, reduces the duration and severity of rotavirus diarrhoea ⁽¹⁵¹⁾, and prevents gastrointestinal damage (increased permeability) caused by non-steroidal anti-inflammatory drugs ⁽¹⁵²⁾. The lactoferrin it contains inhibits the growth of various pathogenic micro-organisms such as *Helicobacter pylori* ⁽¹⁵³⁾.



<u>INULIN</u>: It is a fructooligosaccharide (FOS) of plant origin, extracted from the root of chicory (*Cichorium intybus*). It acts as a prebiotic, creating the right environment for probiotics or beneficial micro-organisms to reproduce faster and in greater numbers ⁽¹⁵⁴⁻¹⁵⁶⁾. It increases the population of *Bifidobacterium* probiotics in the colon and reduces toxic metabolites and harmful enzymes. It prevents pathogenic and autogenous diarrhoea and constipation and protects liver function ⁽¹⁵⁷⁾.

<u>ARABINOGALACTAN</u>: it is an arabino-oligosaccharide (AOS) of plant origin from the larch tree (*Larix laricina*). It is an excellent prebiotic that increases the production of short-chain fatty acids (mainly butyrate), which acts as an energy substrate for the epithelial cells of the colon and protects the intestinal mucosa. It activates the immune response and selectively stimulates the growth and activity of probiotic bacteria ⁽¹⁵⁸⁾. It is useful in fighting infections due to its ability to decrease bacterial adherence ^(159, 160). In addition, it lowers the intestinal pH and improves mineral absorption ⁽¹⁶⁰⁻¹⁶³⁾.

<u>XYLOOLIGOSACCHARIDES (XOS)</u>: are xylan-derived oligosaccharides with a prebiotic effect stimulating the selective growth of beneficial bacteria. XOS also have other beneficial health effects. These positive effects are related to the optimisation of colon functions, as well as the metabolism (increasing or changing the composition of short-chain fatty acids), antioxidant properties, immunostimulation, reduction of triglycerides and cholesterol, reduction of procarcinogenic enzymes, etc. ⁽¹⁶⁴⁻¹⁶⁶⁾.

References

1) De Keersmaecker, Sigrid CJ, et al. "Strong antimicrobial activity of *Lactobacillus rhamnosus* GG against *Salmonella typhimurium* is due to accumulation of lactic acid." FEMS microbiology letters 259.1 (2006): 89-96.

2) Szajewska, H., and M. Kołodziej. "Systematic review with meta-analysis: Lactobacillus rhamnosus GG in the prevention of antibiotic-associated diarrhoea in children and adults." Alimentary pharmacology & therapeutics 42.10 (2015): 1149-1157.

3) Goldenberg, Joshua Z., et al. "Probiotics for the prevention of *Clostridium difficile*-associated diarrhea in adults and children." The Cochrane Library (2013). 4) Davidson, Lisa E., et al. "*Lactobacillus* GG as an immune adjuvant for live-attenuated influenza vaccine in healthy adults: a randomized double-blind placebocontrolled trial." European journal of clinical nutrition 65.4 (2011): 501-507.

5) Baharav, Ehud, et al. "Lactobacillus GG bacteria ameliorate arthritis in Lewis rats." The Journal of nutrition 134.8 (2004): 1964-1969.

6) Thomas, Debra J., et al. "Lactobacillus rhamnosus HN001 attenuates allergy development in a pig model." PLoS One 6.2 (2011): e16577.

7) Kekkonen, Riina A., et al. "Effect of probiotic Lactobacillus rhamnosus GG intervention on global serum lipidomic profiles in healthy adults." World journal of gastroenterology: WJG 14.20 (2008): 3188.

8) Costabile, Adele, et al. "Effect of soluble corn fibre with *Lactobacillus rhamnosus* GG and the pilus-deficient derivative GG-PB12 on faecal microbiota, immune function and metabolism in healthy elderly (Saimes study)." Frontiers in Immunology 8 (2017): 1443.

9) Slykerman, R. F., et al. "Effect of Lactobacillus rhamnosus HN001 in pregnancy on postpartum symptoms of depression and anxiety: a randomized doubleblind placebo-controlled trial." EBio- Medicine 24 (2017): 159-165.

10) Reid, Gregor, et al. "Oral use of Lactobacillus rhamnosus GR-1 and L. fermentum RC-14 significantly alters vaginal flora: randomized, placebo- controlled trial in 64 healthy women." Pathogens and Disease 35.2 (2003): 131-134.

11) Wickens, Kristin L., et al. "Early pregnancy probiotic supplementation with *Lactobacillus rhamnosus* HN001 may reduce the prevalence of gestational diabetes mellitus: a randomized controlled trial." British Journal of Nutrition 117.6 (2017): 804-813.

12) Basu, Sriparna, et al. "Effect of Lactobacillus rhamnosus GG in persistent diarrhea in Indian children: a randomized controlled trial." Journal of clinical gastroenterology 41.8 (2007): 756-760.

13) Szymanski, H., et al. "Treatment of acute infectious diarrhoea in infants and children with a mixture of three *Lactobacillus rhamnosus* strains–a randomized, double-blind, placebo-controlled trial." Alimentary pharmacology & therapeutics 23.2 (2006): 247-253.

14) Ruszczynski, M., A. Radzikowski, and H. Szajewska. "Clinical trial: effectiveness of *Lactobacillus rhamnosus* (strains E/N, Oxy and Pen) in the prevention of antibiotic-associated diarrhoea in children." Alimentary pharmacology & therapeutics 28.1 (2008): 154-161.

Code: FE2018 - 30 enteric-coated vegetable capsules



15) Wu, Yi-Jie, et al. "Evaluation of efficacy and safety of *Lactobacillus rhamnosus* in children aged 4–48 months with atopic dermatitis: An 8-week, doubleblind, randomized, placebo-controlled study." Journal of Microbiology, Immunology and Infection 50.5 (2017): 684-692.

16) Kalliomäki, Marko, et al. "Probiotics in primary prevention of atopic disease: a randomized placebo-controlled trial." The Lancet 357.9262 (2001): 1076-1079.

17) Kaye, Elizabeth Krall. "Daily Intake of Probiotic Lactobacilli May Reduce Caries Risk in Young Children." Journal of Evidence Based Dental Practice 17.3 (2017): 284-286.

18) Stapleton, Ann E., et al. "Randomized, placebo-controlled phase 2 trial of a *Lactobacillus crispatus* probiotic given intravaginally for prevention of recurrent urinary tract infection." Clinical infectious diseases 52.10 (2011): 1212-1217.

19) Hemmerling, Anke, et al. "Phase 1 dose-ranging safety trial of *Lactobacillus crispatus* CTV-05 (LACTIN-V) for the prevention of bacterial vaginosis." Sexually transmitted diseases 36.9 (2009): 564.

00) Wang, Shuai, et al. "Antimicrobial compounds produced by vaginal *Lactobacillus crispatus* are able to strongly inhibit *Candida albicans* growth, hyphal formation and regulate virulence-related gene expressions." Frontiers in microbiology 8 (2017): 564.

21) Eslami, Solat, et al. "Lactobacillus crispatus strain SI-3C-US induces human dendritic cells (DCs) maturation and confers an anti-inflammatory phenotype to DCs." Apmis 124.8 (2016): 697-710.

22) Tobita, Keisuke, Hiroyuki Yanaka, and Hajime Otani. "Anti-allergic effects of Lactobacillus crispatus KT-11 strain on ovalbumin-sensitized BALB/c mice." Animal science journal 81.6 (2010): 699-705.

23) Guillemard, E., et al. "Consumption of a fermented dairy product containing the probiotic *Lactobacillus casei* DN-114 001 reduces the duration of respiratory infections in the elderly in a randomised controlled trial." British journal of nutrition 103.1 (2010): 58-68.

24) Cobo Sanz, JMa, J. A. Mateos, and A. Muñoz Conejo. "Efecto de Lactobacillus casei sobre la incidencia de procesos infecciosos en niños/as." Nutrición Hospitalaria 21.4 (2006): 547-551.

25) Turchet, P., et al. "Effect of fermented milk containing the probiotic *Lactobacillus casei* DN-114001 on winter infections in free-living elderly subjects: a randomised, controlled pilot study." The journal of nutrition, health & aging 7.2 (2003): 75-77.

26) Isolauri, Erika, et al. "Improved immunogenicity of oral D x RRV reassortant rotavirus vaccine by Lactobacillus casei GG." Vaccine 13.3 (1995): 310-312.

27) Matsuzaki, T., et al. "The effect of oral feeding of *Lactobacillus casei* strain Shirota on immunoglobulin E production in mice." Journal of Dairy Science 81.1 (1998): 48-53.

28) Ingrassia, Isabelle, Antony Leplingard, and Arlette Darfeuille-Michaud. "Lactobacillus casei DN-114 001 inhibits the ability of adherent-invasive Escherichia coli isolated from Crohn's disease patients to adhere to and to invade intestinal epithelial cells." Applied and environmental microbiology 71.6 (2005): 2880-2887.

29) Boge, Thierry, et al. "A probiotic fermented dairy drink improves antibody response to influenza vaccination in the elderly in two randomised controlled trials." Vaccine 27.41 (2009): 5677-5684.

30) Giovannini, Marcello, et al. "A randomized prospective double blind controlled trial on effects of long-term consumption of fermented milk containing *Lactobacillus casei* in pre-school children with allergic asthma and/or rhinitis." Pediatric research 62.2 (2007): 215-220.

31) Sýkora, Josef, et al. "Effects of a specially designed fermented milk product containing probiotic *Lactobacillus casei* DN-114 001 and the eradication of *H. pylori* in children: a prospective randomized double-blind study." Journal of clinical gastroenterology 39.8 (2005): 692-698.

32) Guarino, Alfredo, et al. "Oral bacterial therapy reduces the duration of symptoms and of viral excretion in children with mild diarrhea." Journal of pediatric gastroenterology and nutrition 25.5 (1997): 516-519.

33) Merenstein, D., et al. "Use of a fermented dairy probiotic drink containing *Lactobacillus casei* (DN-114 001) to decrease the rate of illness in kids: the DRINK study A patient-oriented, double-blind, cluster-randomized, placebo-controlled, clinical trial." European journal of clinical nutrition 64.7 (2010): 669-677.

24) Basturk, Ahmet, Reha Artan, and Aygen Yilmaz. "Efficacy of synbiotic, probiotic, and prebiotic treatments for irritable bowel syndrome in children: a randomized controlled trial." Turk J Gastroenterol 27.5 (2016): 439-443.

25) Miller, Larry E., Liisa Lehtoranta, and Markus J. Lehtinen. "The effect of *Bifidobacterium animalis* ssp. *lactis* HN019 on cellular immune function in healthy elderly subjects: systematic review and meta-analysis." Nutrients 9.3 (2017): 191.

26) Martín, Rebeca, et al. "*Bifidobacterium animalis* ssp. *lactis* CNCM-12494 restores gut barrier permeability in chronically low-grade inflamed mice." Frontiers in microbiology 7 (2016): 608.

27) Takahashi, Shota, et al. "Effect of *Bifidobacterium animalis* ssp. *lactis* GCL2505 on visceral fat accumulation in healthy Japanese adults: a randomized controlled trial." Bioscience of microbiota, food and health 35.4 (2016): 163-171.

28) Uusitupa, Henna-Maria, et al. "Bifidobacterium animalis subsp. lactis 420 for Metabolic Health: Review of the Research." Nutrients 12.4 (2020): 892.

29) Stenman, L. K., et al. "Potential probiotic Bifidobacterium animalis ssp. lactis 420 prevents weight gain and glucose intolerance in diet-induced obese mice." Beneficial microbes 5.4 (2014): 437-445.

30) Koga, Yasuhiro, et al. "Probiotic *L. gasseri* strain (LG21) for the upper gastrointestinal tract acting through improvement of indigenous microbiota." BMJ open gastroenterology 6.1 (2019): e000314.

31) Lin, Ta-Chin, et al. "Improvement of Bacterial Vaginosis by Oral Lactobacillus Supplement: A Randomized, Double-Blinded Trial." Applied Sciences 11.3 (2021): 902.

32) Ishikawa, Takumi, et al. "Antibacterial activity of the probiotic candidate *Lactobacillus gasseri* against methicillin-resistant *Staphylococcus aureus*." Asian Pacific Journal of Dentistry 20.1 (2020): 1-8.

33) Kobayashi, R., et al. "Oral administration of *Lactobacillus gasseri* SBT2055 is effective in preventing *Porphyromonas gingivalis*-accelerated periodontal disease." Scientific reports 7.1 (2017): 1-10.

34) Shin, Suk Pyo, et al. "A double blind, placebo-controlled, randomized clinical trial that breast milk derived-*Lactobacillus gasseri* BNR17 mitigated diarrheadominant irritable bowel syndrome." Journal of clinical biochemistry and nutrition 62.2 (2018): 179-186.

35) Suzuki, Takayoshi, et al. "Yogurt containing *Lactobacillus gasseri* mitigates aspirin-induced small bowel injuries: a prospective, randomized, double-blind, placebo-controlled trial." Digestion 95.1 (2017): 49-54.

36) Nishihira, Jun, et al. "Lactobacillus gasseri SBT2055 stimulates immunoglobulin production and innate immunity after influenza vaccination in healthy adult volunteers: a randomized, double-blind, placebo-controlled, parallel-group study." Functional Foods in Health and Disease 6.9 (2016): 544-568.

37) Nishihira, Jun, et al. "Lactobacillus gasseri potentiates immune response against influenza virus infection." Immunity and Inflammation in Health and Disease. Academic Press, 2018. 249-255.

38) Kadooka, Y., et al. "Regulation of abdominal adiposity by probiotics (*Lactobacillus gasseri* SBT2055) in adults with obese tendencies in a randomized controlled trial." European journal of clinical nutrition 64.6 (2010): 636-643.

39) Kim, Joohee, et al. "Lactobacillus gasseri BNR17 supplementation reduces the visceral fat accumulation and waist circumference in obese adults: a randomized, double-blind, placebo-controlled trial." Journal of medicinal food 21.5 (2018): 454-461.

40) Wang, Chen, et al. "The effect of probiotic supplementation on lipid profiles in adults with overweight or obesity: A meta-analysis of randomized controlled trials." Journal of Functional Foods 86 (2021): 104711.

41) Klaver, F. A., and Roelof Van Der Meer. "The assumed assimilation of cholesterol by *Lactobacilli* and *Bifidobacterium bifidum* is due to their bile saltdeconjugating activity." Applied and Environmental Microbiology 59.4 (1993): 1120-1124.

42) Zanotti, Ilaria, et al. "Evidence for cholesterol-lowering activity by *Bifidobacterium bifidum* PRL2010 through gut microbiota modulation." Applied microbiology and biotechnology 99.16 (2015): 6813-6829.

43) Shirasawa, Y., et al. "*Bifidobacterium bifidum* BF-1 suppresses Helicobacter pylori-induced genes in human epithelial cells." Journal of dairy science 93.10 (2010): 4526-4534.

Pro-Intensity

Code: FE2018 - 30 enteric-coated vegetable capsules



44) Chenoll, E., et al. "Novel probiotic *Bifidobacterium bifidum* CECT 7366 strain active against the pathogenic bacterium *Helicobacter pylori*." Applied and environmental microbiology 77.4 (2011): 1335-1343.

45) Khailova, Ludmila, et al. "Bifidobacterium bifidum reduces apoptosis in the intestinal epithelium in necrotizing enterocolitis." American Journal of Physiology-Gastrointestinal and Liver Physiology 299.5 (2010): G1118-G1127.

46) Fu, Yu-Rong, et al. "Effects of *Bifidobacterium bifidum* on adaptive immune senescence in aging mice." Microbiology and immunology 54.10 (2010): 578-583.

47) De Vrese, Michael, et al. "Probiotic bacteria reduced duration and severity but not the incidence of common cold episodes in a double blind, randomized, controlled trial." Vaccine 24.44 (2006): 6670-6674.

48) Park, Ji-Hee, et al. "Encapsulated Bifidobacterium bifidum potentiates intestinal IgA production." Cellular immunology 219.1 (2002): 22-27.

49) Guglielmetti, Simone, et al. "Randomised clinical trial: *Bifidobacterium bifidum* MIMBb75 significantly alleviates irritable bowel syndrome and improves quality of life—a double-blind, placebo-controlled study." Alimentary pharmacology & therapeutics 33.10 (2011): 1123-1132.

50) Kim, Namju, et al. "Oral feeding of *Bifidobacterium bifidum* (BGN4) Prevents CD4+ CD45RB high T cell-mediated inflammatory bowel disease by inhibition of disordered T cell activation." Clinical Immunology 123.1 (2007): 30-39.

51) Chitapanarux, Imjai, et al. "Randomized controlled trial of live Lactobacillus acidophilus plus Bifidobacterium bifidum in prophylaxis of diarrhea during radiotherapy in cervical cancer patients." Radiation Oncology 5.1 (2010): 31.

52) Jeon, Seong Gyu, et al. "Probiotic Bifidobacterium breve induces IL-10-producing Tr1 cells in the colon." PLoS pathogens 8.5 (2012): e1002714.

53) Tabbers, M. M., et al. "Is *Bifidobacterium breve* effective in the treatment of childhood constipation? Results from a pilot study." Nutrition journal 10.1 (2011): 19.

54) Ishikawa, Hideki, et al. "Beneficial effects of probiotic bifidobacterium and galacto-oligosaccharide in patients with ulcerative colitis: a randomized controlled study." Digestion 84.2 (2011): 128-133.

55) Mullié, Catherine, et al. "Increased poliovirus-specific intestinal antibody response coincides with promotion of *Bifidobacterium longum-infantis* and *Bifidobacterium breve* in infants: a randomized, double-blind, placebo-controlled trial." Pediatric research 56.5 (2004): 791-795.

56) Sheehan, Vivien M., et al. "Improving gastric transit, gastrointestinal persistence and therapeutic efficacy of the probiotic strain *Bifidobacterium breve* UCC2003." Microbiology 153.10 (2007): 3563-3571.

57) Mendonça, Fabio Henrique Boarini Pacheco, et al. "Effects of probiotic bacteria on Candida presence and IgA anti-Candida in the oral cavity of elderly." Brazilian dental journal 23.5 (2012): 534-538.

58) Minami, Jun-ichi, et al. "Oral administration of *Bifidobacterium breve* B-3 modifies metabolic functions in adults with obese tendencies in a randomised controlled trial." Journal of nutritional science 4 (2015).

59) Kitajima, Hiroyuki, et al. "Early administration of *Bifidobacterium breve* to preterm infants: randomised controlled trial." Archives of Disease in Childhood-Fetal and Neonatal Edition 76.2 (1997): F101-F107.

60) Braga, Taciana Duque, et al. "Efficacy of *Bifidobacterium breve* and *Lactobacillus casei* oral supplementation on necrotizing enterocolitis in very-low-birth-weight preterm infants: a double-blind, randomized, controlled trial-." The American journal of clinical nutrition 93.1 (2010): 81-86.

61) Klemenak, Martina, et al. "Administration of *Bifidobacterium breve* Decreases the Production of TNF-alfa in Children with Celiac Disease." Digestive diseases and sciences 60.11 (2015): 3386-3392.

62) Wada, Mariko, et al. "Effects of the enteral administration of *Bifidobacterium breve* on patients undergoing chemotherapy for pediatric malignancies." Supportive care in cancer 18.6 (2010): 751-759.

63) He, Fang, et al. "Comparison of mucosal adhesion and species identification of bifidobacteria isolated from healthy and allergic infants." Pathogens and Disease 30.1 (2001): 43-47.

64) Ishibashi, N., T. Yaeshima, and H. Hayasawa. "Bifidobacteria: their significance in human intestinal health." Malaysian Journal of Nutrition 3.2 (1997): 149-159.

65) Sun, Wenrong, and Mansel W. Griffiths. "Survival of bifidobacteria in yogurt and simulated gastric juice following immobilization in gellan–xanthan beads." International Journal of Food Microbiology 61.1 (2000): 17-25.

66) Bernet, Marie-Francoise, et al. "Adhesion of human bifidobacterial strains to cultured human intestinal epithelial cells and inhibition of enteropathogencell interactions." Applied and environmental microbiology 59.12 (1993): 4121-4128.

67) Gibson, G. R., and Xin Wang. "Regulatory effects of bifidobacteria on the growth of other colonic bacteria." Journal of Applied Microbiology 77.4 (1994): 412-420.

68) Cheikhyoussef, Ahmad, et al. "Antimicrobial activity and partial characterization of bacteriocin-like inhibitory substances (BLIS) produced by *Bifidobacterium infantis* BCRC 14602." Food Control 20.6 (2009): 553-559.

69) Cheikhyoussef, Ahmad, et al. "Bifidin I–A new bacteriocin produced by *Bifidobacterium infantis* BCRC 14602: Purification and partial amino acid sequence." Food Control 21.5 (2010): 746-753.

70) Whorwell, Peter J., et al. "Efficacy of an encapsulated probiotic *Bifidobacterium infantis* 35624 in women with irritable bowel syndrome." The American journal of gastroenterology 101.7 (2006): 1581-1590.

71) Brenner, Darren M., and William D. Chey. "Bifidobacterium infantis 35624: a novel probiotic for the treatment of irritable bowel syndrome." Reviews in gastroenterological disorders 9.1 (2009): 7-15.

72) O'Mahony, Liam, et al. "Lactobacillus and bifidobacterium in irritable bowel syndrome: symptom responses and relationship to cytokine profiles." Gastroenterology 128.3 (2005): 541-551.

73) Groeger, David, et al. "Bifidobacterium infantis 35624 modulates host inflammatory processes beyond the gut." Gut microbes 4.4 (2013): 325-339.

74) Smecuol, Edgardo, et al. "Exploratory, randomized, double-blind, placebo-controlled study on the effects of *Bifidobacterium infantis* natren life start strain super strain in active celiac disease." Journal of clinical gastroenterology 47.2 (2013): 139-147.

75) Fujiwara, Shigeru, et al. "Proteinaceous factor (s) in culture supernatant fluids of bifidobacteria which prevents the binding of enterotoxigenic *Escherichia* coli to gangliotetraosylceramide." Applied and environmental microbiology 63.2 (1997): 506-512.

76) Furrie, Elizabeth, et al. "Synbiotic therapy (*Bifidobacterium longum*/Synergy 1) initiates resolution of inflammation in patients with active ulcerative colitis: a randomised controlled pilot trial." Gut 54.2 (2005): 242-249.

77) Orrhage, K., B. Brismar, and C. E. Nord. "Effect of supplements with *Bifidobacterium longum* and *Lactobacillus acidophilus* on the intestinal microbiota during administration of clindamycin." Microbial Ecology in Health and Disease 7.1 (1994): 17-25.

78) Koning, Catherina JM, et al. "The effect of a multispecies probiotic on the intestinal microbiota and bowel movements in healthy volunteers taking the antibiotic amoxycillin." The American journal of gastroenterology 103.1 (2008): 178-189.

79) Ortiz-Lucas, María, et al. "Effect of probiotic species on irritable bowel syndrome symptoms: A bring up to date meta-analysis." Rev Esp Enferm Dig 105.1 (2013): 19-36.

80) Xiao, Jin-zhong, et al. "Clinical efficacy of probiotic *Bifidobacterium longum* for the treatment of symptoms of Japanese cedar pollen allergy in subjects evaluated in an environmental exposure unit." Allergology international 56.1 (2007): 67-75.

81) Takahashi, N., et al. "Immunostimulatory oligodeoxynucleotide from *Bifidobacterium longum* suppresses Th2 immune responses in a murine model." Clinical & Experimental Immunology 145.1 (2006): 130-138.

82) Makras, Lefteris, and Luc De Vuyst. "The in vitro inhibition of Gram-negative pathogenic bacteria by bifidobacteria is caused by the production of organic acids." International Dairy Journal 16.9 (2006): 1049-1057.

83) LeBlanc, J. G., et al. "B-Group vitamin production by lactic acid bacteria–current knowledge and potential applications." Journal of Applied Microbiology 111.6 (2011): 1297-1309.

84) Sinn, Dong Hyun, et al. "Therapeutic effect of *Lactobacillus acidophilus*-SDC 2012, 2013 in patients with irritable bowel syndrome." Digestive diseases and sciences 53.10 (2008): 2714-2718.

Pro-Intensity

Code: FE2018 - 30 enteric-coated vegetable capsules



85) Gao, Xing Wang, et al. "Dose–response efficacy of a proprietary probiotic formula of *Lactobacillus acidophilus* CL1285 and *Lactobacillus casei* LBC80R for antibiotic-associated diarrhea and *Clostridium difficile*-associated diarrhea prophylaxis in adult patients." The American journal of gastroenterology 105.7 (2010): 1636-1641.

86) Ooi, L-G., et al. "Lactobacillus acidophilus CHO-220 and inulin reduced plasma total cholesterol and low-density lipoprotein cholesterol via alteration of lipid transporters." Journal of dairy science 93.11 (2010): 5048-5058.

87) Rerksuppaphol, Sanguansak, and Lakkana Rerksuppaphol. "A randomized double-blind controlled trial of *Lactobacillus acidophilus* plus *Bifidobacterium bifidum* versus placebo in patients with hypercholesterolemia." Journal of clinical and diagnostic research: JCDR 9.3 (2015): KC01.

88) Bader J, et al. "Processing, consumption and effects of probiotic microorganisms." Encyclopedia of Life Support Systems. (2012).

89) Ishida, Y., et al. "Clinical effects of Lactobacillus acidophilus strain L-92 on perennial allergic rhinitis: a double-blind, placebo-controlled study." Journal of Dairy Science 88.2 (2005): 527-533.

90) Ishida, Yu, et al. "Effect of milk fermented with Lactobacillus acidophilus strain L-92 on symptoms of Japanese cedar pollen allergy: a randomized placebocontrolled trial." Bioscience, biotechnology, and biochemistry 69.9 (2005): 1652-1660.

91) Torii, Shinpei, et al. "Effects of oral administration of Lactobacillus acidophilus L-92 on the symptoms and serum markers of atopic dermatitis in children." International archives of allergy and immunology 154.3 (2011): 236-245.

92) Ryan, Kieran A., et al. "Strain-specific inhibition of Helicobacter pylori by Lactobacillus salivarius and other lactobacilli." Journal of Antimicrobial Chemotherapy 61.4 (2008): 831-834.

93) Kabir, A. M., et al. "Prevention of Helicobacter pylori infection by lactobacilli in a gnotobiotic murine model." Gut 41.1 (1997): 49-55.

94) Riboulet-Bisson, Eliette, et al. "Effect of Lactobacillus salivarius bacteriocin Abp118 on the mouse and pig intestinal microbiota." PLoS One 7.2 (2012): e31113.

95) European Food Safety Authority (EFSA). "Scientific Opinion on the safety and efficacy of *Lactobacillus salivarius* (CNCM I-3238) n *Lactobacillus casei* (ATCC PTA-6135) as silage additives for all species." EFSA Journal 10.9 (2012): 2884.

96) Rajkumar, Hemalatha, et al. "Effect of probiotic *Lactobacillus salivarius* UBL S22 and prebiotic fructo-oligosaccharide on serum lipids, inflammatory markers, insulin sensitivity, and gut bacteria in healthy young volunteers: A randomized controlled single-blind pilot study." Journal of Cardiovascular Pharmacology and Therapeutics 20.3 (2015): 289-298.

97) Wu, K-G., T-H. Li, and H-J. Peng. "Lactobacillus salivarius plus fructo-oligosaccharide is superior to fructo-oligosaccharide alone for treating children with moderate to severe atopic dermatitis: a double-blind, randomized, clinical trial of efficacy and safety." British Journal of Dermatology 166.1 (2012): 129-136. 98) Drago, L., et al. "Effects of Lactobacillus salivarius LS01 (DSM 22775) treatment on adult atopic dermatitis: a randomized placebo-controlled study." International Journal of Immunopathology and Pharmacology 24.4 (2011): 1037-1048.

99) Niedzielin, Krzysztof, Hubert Kordecki, and Boz ena Birkenfeld. "A controlled, double-blind, randomized study on the efficacy of *Lactobacillus plantarum* 299V in patients with irritable bowel syndrome." European Journal of Gastroenterology & Hepatology 13.10 (2001): 1143-1147.

100) Kim, H. Jae, et al. "A randomized controlled trial of a probiotic, VSL# 3, on gut transit and symptoms in diarrhoea-predominant irritable bowel syndrome." Alimentary Pharmacology & Therapeutics 17.7 (2003): 895-904.

101) Nobaek, Sören, et al. "Alteration of intestinal microflora is associated with reduction in abdominal bloating and pain in patients with irritable bowel syndrome." The American Journal of Gastroenterology 95.5 (2000): 1231-1238.

102) Nikfar, Shekoufeh, et al. "Efficacy of probiotics in irritable bowel syndrome: a meta-analysis of randomized, controlled trials." Diseases of the Colon & Rectum 51.12 (2008): 1775-1780.

103) Ducrotté, Philippe, Prabha Sawant, and Venkataraman Jayanthi. "Clinical trial: Lactobacillus plantarum 299v (DSM 9843) improves symptoms of irritable bowel syndrome." World Journal of Gastroenterology: WJG 18.30 (2012): 4012.

104) Kumar, CSV Satish, et al. "Protective effect of Lactobacillus plantarum 21, a probiotic on trinitrobenzenesulfonic acid-induced ulcerative colitis in rats." International Immunopharmacology 25.2 (2015): 504-510.

105) Bibiloni, Rodrigo, et al. "VSL# 3 probiotic-mixture induces remission in patients with active ulcerative colitis." The American Journal of Gastroenterology 100.7 (2005): 1539-1546.

106) Han, Youngshin, et al. "A randomized trial of *Lactobacillus plantarum* CJLP133 for the treatment of atopic dermatitis." Pediatric Allergy and Immunology 23.7 (2012): 667-673.

107) Mane, J., et al. "A mixture of *Lactobacillus plantarum* CECT 7315 and CECT 7316 enhances systemic immunity in elderly subjects. A dose-response, double-blind, placebo-controlled, randomized pilot trial." Nutricion Hospitalaria 26.1 (2011).

108) Lönnermark, Elisabet, et al. "Intake of Lactobacillus plantarum reduces certain gastrointestinal symptoms during treatment with antibiotics." Journal of Clinical Gastroenterology 44.2 (2010): 106-112.

109) Naruszewicz, Marek, et al. "Effect of *Lactobacillus plantarum* 299v on cardiovascular disease risk factors in smokers." The American Journal of Clinical Nutrition 76.6 (2002): 1249-1255.

110) Costabile, Adele, et al. "An in vivo assessment of the cholesterol-lowering efficacy of *Lactobacillus plantarum* ECGC 13110402 in normal to mildly hypercholesterolaemic adults." PLoS One 12.12 (2017): e0187964.

111) Roškar, Irena, et al. "Effects of a probiotic product containing *Bifidobacterium animalis* subsp. *animalis* IM386 and *Lactobacillus plantarum* MP2026 in lactose intolerant individuals: Randomized, placebo-controlled clinical trial." Journal of Functional Foods 35 (2017): 1-8.

112) Strus, Magdalena, et al. "Studies on the effects of probiotic *Lactobacillus* mixture given orally on vaginal and rectal colonization and on parameters of vaginal health in women with intermediate vaginal flora." European Journal of Obstetrics & Gynecology and Reproductive Biology 163.2 (2012): 210-215.

113) Cruchet, Sylvia, et al. "Effect of the ingestion of a dietary product containing *Lactobacillus johnsonii* La1 on Helicobacter pylori colonization in children." Nutrition 19.9 (2003): 716-721.

114) Marcial, Guillermo E., et al. "Lactobacillus johnsonii N6. 2 modulates the host immune responses: a double-blind, randomized trial in healthy adults." Frontiers in immunology 8 (2017): 655.

115) Lau, Kenneth, et al. "Inhibition of type 1 diabetes correlated to a *Lactobacillus johnsonii* N6. 2-mediated Th17 bias." The Journal of Immunology 186.6 (2011): 3538-3546.

116) Joo, Hyun-Min, et al. "Lactobacillus johnsonii HY7042 ameliorates Gardnerella vaginalis-induced vaginosis by killing Gardnerella vaginalis and inhibiting NF-κB activation." International immunopharmacology 11.11 (2011): 1758-1765.

117) Lue, Ko-Haung, et al. "A trial of adding *Lactobacillus johnsonii* EM1 to levocetirizine for treatment of perennial allergic rhinitis in children aged 7–12 years." International journal of pediatric otorhinolaryngology 76.7 (2012): 994-1001.

118) Michalickova, Danica M., et al. "Lactobacillus helveticus Lafti L10 Supplementation Modulates Mucosal and Humoral Immunity in Elite Athletes: A Randomized, Double-Blind, Placebo-Controlled Trial." The Journal of Strength & Conditioning Research 31.1 (2017): 62-70.

119) Liang, S., et al. "Administration of Lactobacillus helveticus NS8 improves behavioral, cognitive, and biochemical aberrations caused by chronic restraint stress." Neuroscience 310 (2015): 561-577.

120) Chung, Young-Chul, et al. "Fermented milk of *Lactobacillus helveticus* IDCC3801 improves cognitive functioning during cognitive fatigue tests in healthy older adults." Journal of Functional Foods 10 (2014): 465-474.

121) Jauhiainen, Tiina, et al. "Lactobacillus helveticus fermented milk lowers blood pressure in hypertensive subjects in 24-h ambulatory blood pressure measurement." American Journal of Hypertension 18.12 (2005): 1600-1605.

122) Narva, Mirkka, et al. "Effects of long-term intervention with *Lactobacillus helveticus*-fermented milk on bone mineral density and bone mineral content in growing rats." Annals of Nutrition and Metabolism 48.4 (2004): 228-234.

123) Narva, Mirkka, et al. "The effect of *Lactobacillus helveticus* fermented milk on acute changes in calcium metabolism in postmenopausal women." European journal of nutrition 43.2 (2004): 61-68.

124) Taverniti, Valentina, and Simone Guglielmetti. "Health-promoting properties of Lactobacillus helveticus." Frontiers in Microbiology 3 (2012).

Pro-Intensity

Code: FE2018 - 30 enteric-coated vegetable capsules



125) 103) Rizzardini, Giuliano, et al. "Evaluation of the immune benefits of two probiotic strains *Bifidobacterium animalis* ssp. *lactis*, BB-12[®] and *Lactobacillus paracasei* ssp. *paracasei*, *L. casei* 431[®] in an influenza vaccination model: a randomised, double-blind, placebo-controlled study." British Journal of Nutrition 107.6 (2012): 876-884.

126) Riezzo, G., et al. "Randomised clinical trial: efficacy of *Lactobacillus paracasei*-enriched artichokes in the treatment of patients with functional constipation–a double-blind, controlled, crossover study." Alimentary Pharmacology & Therapeutics 35.4 (2012): 441-450.

127) Costa, D. J., et al. "Efficacy and safety of the probiotic *Lactobacillus paracasei* LP-33 in allergic rhinitis: a double-blind, randomized, placebo-controlled trial (GA2LEN Study)." European Journal of Clinical Nutrition 68.5 (2014): 602-607.

128) Bendali, Farida, Nassim Madi, and Djamila Sadoun. "Beneficial effects of a strain of *Lactobacillus paracasei* subsp. *paracasei* in *Staphylococcus aureus*induced intestinal and colonic injury." International Journal of Infectious Diseases 15.11 (2011): e787-e794.

129) Tsai, Yueh-Ting, Po-Ching Cheng, and Tzu-Ming Pan. "Immunomodulating activity of *paracasei* subsp. *paracasei* NTU 101 in enterohemorrhagic *Escherichia coli* O157H7-infected mice." Journal of Agricultural and Food Chemistry 58.21 (2010): 11265-11272.

130) Jankowska, Alicja, et al. "Competition of Lactobacillus paracasei with Salmonella enterica for adhesion to Caco-2 cells." BioMed Research International 2008 (2008).

131) Passariello, A., et al. "Randomised clinical trial: efficacy of a new synbiotic formulation containing *Lactobacillus paracasei* B21060 plus arabinogalactan and xilooligosaccharides in children with acute diarrhoea." Alimentary Pharmacology & Therapeutics 35.7 (2012): 782-788.

132) Sullivan, Åsa, Carl E. Nord, and Birgitta Evengård. "Effect of supplement with lactic-acid producing bacteria on fatigue and physical activity in patients with chronic fatigue syndrome." Nutrition Journal 8.1 (2009): 4.

133) Al-Omari, Aisha W., Ikhlas Ramadan Matter, and Alaa Hussein Almola. "An overview of Bacteriocins." Samarra Journal of Pure and Applied Science 4.2 (2022): 58-72.

134) Małaczewska, Joanna, and Edyta Kaczorek-Łukowska. "Nisin—A lantibiotic with immunomodulatory properties: A review." Peptides 137 (2021): 170479. 135) Le Lay, Christophe, et al. "Nisin is an effective inhibitor of *Clostridium difficile* vegetative cells and spore germination." Journal of medical microbiology 65.2 (2016): 169-175.

136) Shibata, Takeo, et al. "Lactococcus lactis JCM5805 activates anti-viral immunity and reduces symptoms of common cold and influenza in healthy adults in a randomized controlled trial." Journal of Functional Foods 24 (2016): 492-500.

137) Thu, Nghiem Nguyet, et al. "Impact of Infectious Disease after *Lactococcus lactis* Strain Plasma Intake in Vietnamese Schoolchildren: A Randomized, Placebo-Controlled, Double-Blind Study." Nutrients 14.3 (2022): 552.

138) Beltrán-Barrientos, Lilia M., et al. "Randomized double-blind controlled clinical trial of the blood pressure–lowering effect of fermented milk with *Lactococcus lactis*: A pilot study." Journal of Dairy Science 101.4 (2018): 2819-2825.

139) Liu, Meiling, et al. "Protective effects of a novel probiotic strain, *Lactococcus lactis* ML2018, in colitis: in vivo and in vitro evidence." Food & function 10.2 (2019): 1132-1145.

140) Khemariya, Priti, et al. "Probiotic Lactococcus lactis: A review." Turkish Journal of Agriculture-Food Science and Technology 5.6 (2017): 556-562.

141) Cannarella, Ligia Aparecida Trintin, et al. "Mixture of probiotics reduces inflammatory biomarkers and improves the oxidative/nitrosative profile in people with rheumatoid arthritis." Nutrition 89 (2021): 111282.

142) Gomes, Aline Corado, et al. "The additional effects of a probiotic mix on abdominal adiposity and antioxidant Status: a double-blind, randomized trial." Obesity 25.1 (2017): 30-38.

143) Godhia, Meena L., et al. "Colostrum-its Composition, Benefits as a Nutraceutical-A Review." Curr Res Nutr Food Sci J 1.1 (2013): 37-47.

144) Fortín, A.M., et al. "Determinación de la calidad del calostro bovino a partir de la densidad y de la concentración de IgG y del número de partos de la vaca y su efecto en el desarrollo de los terneros hasta los 30 días de edad." BS thesis. Zamorano: Escuela Agrícola Panamericana, 2012, 2009.

145) Shing, C.M. "Effects of bovine colostrum supplementation on immune variables in highly trained cyclists." J Appl Physiol 102.3 (2007): 1113-22.

146) Jones, A.W., et al. "The effects of bovine colostrum supplementation on in vivo immunity following prolonged exercise: a randomised controlled trial." Eur J Nutr (2017): 1-10.

147) Kotsis, Yiannis, et al. "A low-dose, 6-week bovine colostrum supplementation maintains performance and attenuates inflammatory indices following a Loughborough Intermittent Shuttle Test in soccer players." Eur J Nutr (2017): 1-15.

148) Crooks, Christine, et al. "Effect of bovine colostrum supplementation on respiratory tract mucosal defenses in swimmers." Int J Sport Nutr Exerc Metab 20.3 (2010): 224-235.

149) Jones, A.W., et al. "Effects of bovine colostrum supplementation on upper respiratory illness in active males." Brain Behav Immun 39 (2014): 194-203. 150) Kaducu, F.O., et al. "Effect of bovine colostrum-based food supplement in the treatment of HIV-associated diarrhea in Northern Uganda: a randomized

controlled trial." Indian Journal of Gastroenterology 30.6 (2011): 270-276. 151) Mitra, AK., et al. "Hyperimmune cow colostrum reduces diarrhoea due to rotavirus: a double-blind, controlled clinical trial." Acta Paediatrica 84.9 (1995): 996-1001.

152) Playford, Raymond J., et al. "Co-administration of the health food supplement, bovine colostrum, reduces the acute non-steroidal anti-inflammatory drug-induced increase in intestinal permeability." Clinical Science 100.6 (2001): 627-633.

153) Dzik, Sara, et al. "Properties of bovine colostrum and the possibilities of use." Polish Annals of Medicine 24.2 (2017): 295-299.

154) Institute of FoodTechnologists (IFT). What are fructooligosaccharides and how do they provide digestive, immunity and bone health benefits?. ScienceDaily (2013).

155) Gibson, Glenn R. "Dietary modulation of the human gut microflora using the prebiotics oligofructose and inulin." The Journal of nutrition 129.7 (1999): 14385-1441s.

156) Flamm, Gary, et al. "Inulin and oligofructose as dietary fiber: a review of the evidence." Critical reviews in food science and nutrition 41.5 (2001): 353-362.

157) Cardarelli, Haíssa R., et al. "Inulin and oligofructose improve sensory quality and increase the probiotic viable count in potentially synbiotic petit-suisse cheese." LWT-Food Science and Technology 41.6 (2008): 1037-1046.

158) Robinson, Ramona R., Joellen Feirtag, and Joanne L. Slavin. "Effects of dietary arabinogalactan on gastrointestinal and blood parameters in healthy human subjects." Journal of the American College of Nutrition 20.4 (2001): 279-285.

159) Gibson, Glenn R. "Dietary modulation of the human gut microflora using the prebiotics oligofructose and inulin." The Journal of nutrition 129.7 (1999): 14385-1441s.

160) Flamm, Gary, et al. "Inulin and oligofructose as dietary fiber: a review of the evidence." Critical reviews in food science and nutrition 41.5 (2001): 353-362.

161) Van Loo, Jan, et al. "On the presence of inulin and oligofructose as natural ingredients in the western diet." Critical Reviews in Food Science & Nutrition 35.6 (1995): 525-552.

162) Niness, Kathy R. "Inulin and oligofructose: what are they?." The Journal of nutrition 129.7 (1999): 1402S-1406s.

163) Rao, A. V. "Dose-response effects of inulin and oligofructose on intestinal bifidogenesis effects." The Journal of nutrition 129.7 (1999): 1442S-1445s. 164) Samanta, A. K., et al. "Xylooligosaccharides as prebiotics from agricultural by-products: Production and applications." Bioactive Carbohydrates and Dietary Fibre 5.1 (2015): 62-71.

165) Wang, Jing, et al. "Wheat bran xylooligosaccharides improve blood lipid metabolism and antioxidant status in rats fed a high-fat diet." Carbohydrate Polymers 86.3 (2011): 1192-1197.

166) Palaniappan, Ayyappan, Usha Antony, and Mohammad Naushad Emmambux. "Current status of xylooligosaccharides: Production, characterization, health benefits and food application." Trends in Food Science & Technology 111 (2021): 506-519.