J.J. Malago J.F.J.G. Koninkx R. Marinsek-Logar *Editors*

Probiotic Bacteria and Enteric Infections

Cytoprotection by Probiotic Bacteria



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Joshua J. Malago • Jos F. J. G. Koninkx R. Marinsek-Logar Editors

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Editors: Dr. Joshua J. Malago Department of Veterinary Pathology Faculty of Veterinary Medicine Sokoine University of Agriculture P.O. Box 3203, Chuo Kikuu, Morogoro Tanzania malagojj@yahoo.com

Dr. Jos F. J. G. Koninkx Division Pathology Department of Pathobiology Faculty of Veterinary Medicine Utrecht University Yalelaan 1, 3508 TD, Utrecht Netherlands j.f.j.g.koninkx@uu.nl Dr. R. Marinsek-Logar Zootechnical Department Biotechnical Faculty University of Ljubljana Domžale Slovenia romana.marinsek@bf.uni-lj.si

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Contents

Pa	rt I Introduction and History of Probiotics	1				
1	Probiotics: From the Ancient Wisdom to the Actual Therapeutical and Nutraceutical Perspective Giuseppe Caramia and Stefania Silvi					
Pa	rt II The Gut Microorganisms and Probiotics	39				
2	The Intestinal Microbiota and Probiotics Sofia D. Forssten, Sampo J. Lahtinen and Arthur C. Ouwehand	41				
3	Ecology of Probiotics and Enteric Protection Melanie Gagnon, Annina Zihler, Christophe Chassard and Christophe Lacroix					
Pa	rt III Pathophysiology of Enteric Disorders Due to Disturbed Microbiota	87				
4	Factors Causing Disturbances of the Gut Microbiota Joshua J. Malago and Jos F. J. G. Koninkx	89				
5	The Gut Microbiota, Probiotics and Infectious Disease Cormac G. M. Gahan, Gerald C. O'Sullivan and J. Kevin Collins					
Pa	rt IV Application of Molecular Biology and -omics of Probiotics in Enteric Protection	131				
6	Application of Molecular Biology and Genomics ofProbiotics for Enteric CytoprotectionSaloomeh Moslehi-Jenabian, Dennis Sandris Nielsen and Lene Jespersen	133				
7	Application of Probiotic Proteomics in Enteric Cytoprotection	155				

8	Promoting Gut Health with Probiotic Metabolomics Sebastiano Collino, François-Pierre J. Martin, Sunil Kochhar and Serge Rezzi				
Par	t V Probiotics, Gut Immunology and Enteric Protection	187			
9	Microbiota, Probiotics and Natural Immunity of the Gut Eduardo Jorge Schiffrin and Anne Donnet-Hughes	189			
10	Interaction of Probiotics with the Acquired Immune System in the Gut Rossana D'Arienzo, Kathryne B. Schwartz and Mauro Rossi				
11					
12	Modulation of Immune System by Probiotics to Protect Against Enteric Disorders Joshua J. Malago and Jos F. J. G. Koninkx				
Part VI Probiotics for Enteric Therapy 23					
13	Probiotic-Pathogen Interactions and Enteric Cytoprotection Joshua J. Malago and Jos F. J. G. Koninkx	289			
14	Bacteriocins of Probiotics and Enteric Cytoprotection Bojana Bogovič-Matijašić and Irena Rogelj	313			
15	 Probiotics in Clinical Practice as Therapeutics Against Enteric Disorders Ouafae Karimi and A. S. Peña 				
16	Potential Mechanisms of Enteric Cytoprotection by Probiotics: Lessons from Cultured Human Intestinal Cells Vanessa Liévin-Le Moal and Alain L. Servin	375			
17	Probiotics and Enteric Cancers Min-Tze Liong, Huey-Shi Lye, Siok-Koon Yeo, Joo-Ann Ewe, Lay-Gaik Ooi and Ting-Jin Lim	399			
Par	t VII The Future of Probiotics	427			
18	Designer Probiotics and Enteric Cytoprotection Adrienne W. Paton, Renato Morona and James C. Paton	429			
19	Future Prospects of Probiotics as Therapeutics Against Enteric Disorders E. P. Culligan, C. Hill and R. D. Sleator				
Ind	ex	465			

Contributors

Hans Christian Beck The Proteomics Group, Danish Technological Institute, Holbergsvej 10, 6000 Kolding, Denmark e-mail: hcb@teknologisk.dk

Giuseppe Caramia Hemeritus Head Physician of Paediatrics and Neonatology, Specialized Maternal-Infantil Hospital "G.Salesi", Via Toti 1, 60123 Ancona, Italy e-mail: caramiagm@libero.it

Christophe Chassard Laboratory of Food Biotechnology, Institute of Food Science and Nutrition, ETH Zurich, Schmelzbergstrasse 7, LFV C20 CH-8092 Zurich, Switzerland e-mail: christoph.chassard@ilw.agrl.ethz

J. Kevin Collins Department of Microbiology, University College Cork, Cork, Ireland Cork Cancer Research Centre, University College Cork, Cork, Ireland e-mail: microbiology@ucc.ie

Sebastiano Collino BioAnalytical Science, Metabolomics & Biomarkers, Nestlé Research Center, P.O. Box 44, 1000 Lausanne 26, Switzerland e-mail: Sebastiano.Collino@rdls.nestle.com

E. P. Culligan Alimentary Pharmabiotic Centre and Department of Microbiology, University College Cork, Cork, Ireland e-mail: eamonnculligan@gmail.com

Rossana D'Arienzo Institute of Food Sciences, CNR, via Roma 64, 83100 Avellino, Italy e-mail: rdarienzo@isa.cnr.it

Véronique Delcenserie Canadian Research Institute for Food Safety, University of Guelph, 43, McGilvray Street, Guelph, Ontario, Canada N1G 2W1 e-mail: vdelcens@uoguelph.ca

Anne Donnet-Hughes Nestlé Research Centre, P.O. Box 44, Vers-chez-les-Blanc, 1000 Lausanne, Switzerland e-mail: anne.donnet@rdls.nestle.com Joo-Ann Ewe School of Industrial Technology, Universiti Sains Malaysia, 11800 Penang, Malaysia e-mail: jooann_ewe@hotmail.com

Søren Feddersen Odense University Hospital, Department of Clinical Chemistry and Pharmacology, Sdr. Boulevard 29, DK-5000 Odense C, Denmark e-mail: soeren.feddersen@ouh.regionsyddanmark.dk

Sofia D. Forssten Danisco Health & Nutrition, Sokeritehtaantie 20, 02460 Kantvik, Finland e-mail: sofia.forssten@danisco.com

Melanie Gagnon Laboratory of Food Biotechnology, Institute of Food Science and Nutrition, ETH Zurich, Schmelzbergstrasse 7, 8092 Zurich, Switzerland e-mail: melanie.gagnon@ilw.agrl.ethz.ch

Cormac G. M. Gahan Department of Microbiology, University College Cork, Cork, Ireland e-mail: c.gahan@ucc.ie

C. Hill Alimentary Pharmabiotic Centre and Department of Microbiology, University College Cork, Cork, Ireland e-mail: c.hill@ucc.ie

Lene Jespersen Department of Food Science, Food Microbiology, Faculty of Life Sciences, University of Copenhagen, Rolighedsvej 30, 1958 Frederiksberg, Denmark e-mail: lj@life.ku.dk

Ouafae Karimi Department of Pathology, VU University Medical Center, Laboratory of Immunogenetics, P.O. Box 7057, 1007 MB Amsterdam, The Netherlands e-mail: ouafaekarimi@gmail.com

Sunil Kochhar BioAnalytical Science, Metabolomics & Biomarkers, Nestlé Research Center, P.O. Box 44, 1000 Lausanne 26, Switzerland e-mail: sunil.kochhar@rdls.nestle.com

Jos F. J. G. Koninkx Division Pathology, Department of Pathobiology, Faculty of Veterinary Medicine, Utrecht University, Yalelaan 1, 3508 TD Utrecht, The Netherlands e-mail: j.f.j.g.koninkx@uu.nl

Christophe Lacroix Laboratory of Food Biotechnology, Institute of Food Science and Nutrition, ETH Zurich, Schmelzbergstrasse 7, 8092 Zurich, Switzerland e-mail: christophe.lacroix@ilw.agrl.ethz.ch

Sampo J. Lahtinen Danisco Health & Nutrition, Sokeritehtaantie 20, 02460 Kantvik, Finland e-mail: sampo.lahtinen@danisco.com Vanessa Liévin-Le Moal Faculté de Pharmacie, Inserm Unité 756, 92296 Châtenay-Malabry, France e-mail: vanessa.lievin-le-moal@u-psud.fr

Ting-Jin Lim School of Industrial Technology, Universiti Sains Malaysia, 11800 Penang, Malaysia e-mail: ericlimtingjin@hotmail.com

Min-Tze Liong School of Industrial Technology, Universiti Sains Malaysia, 11800 Penang, Malaysia e-mail: mintze.liong@usm.my

Joshua J. Malago Department of Veterinary Pathology, Faculty of Veterinary Medicine, Sokoine University of Agriculture, 3203, Chuo Kikuu, Morogoro, Tanzania e-mail: malagojj@yahoo.com

Huey-Shi Lye School of Industrial Technology, Universiti Sains Malaysia, 11800 Penang, Malaysia e-mail: hueyshi_lye@yahoo.com

François-Pierre J. Martin BioAnalytical Science, Metabolomics & Biomarkers, Nestlé Research Center, P.O. Box 44, 1000 Lausanne 26, Switzerland e-mail: Francois-Pierre.Martin@rdls.nestle.com

Bojana Bogovič Matijašić Department of Animal Science, Chair of Dairy Science, Biotechnical Faculty, University of Ljubljana, Groblje 3, 1230 Domžale, Slovenia

e-mail: bojana.bogovic@bf.uni-lj.si

Renato Morona Research Centre for Infectious Diseases, School of Molecular and Biomedical Science, University of Adelaide, Adelaide, SA 5005, Australia e-mail: renato.morona@adelaide.edu.au

Saloomeh Moslehi-Jenabian Department of Food Science, Food Microbiology, Faculty of Life Sciences, University of Copenhagen, Rolighedsvej 30, 1958 Frederiksberg, Denmark e-mail: smje@life.ku.dk

Dennis Sandris Nielsen Department of Food Science, Food Microbiology, Faculty of Life Sciences, University of Copenhagen, Rolighedsvej 30, 1958 Frederiksberg, Denmark e-mail: dn@life.ku.dk

Lay-Gaik Ooi School of Industrial Technology, Universiti Sains Malaysia, 11800 Penang, Malaysia e-mail: ooilaygaik@yahoo.com

Gerald C. O'Sullivan Cork Cancer Research Centre, University College Cork, Cork, Ireland Deptment of Surgery, Mercy University Hospital, Cork, Ireland e-mail: gcosullivan@rcsi.ie Arthur C. Ouwehand Danisco Health & Nutrition, Sokeritehtaantie 20, 02460 Kantvik, Finland e-mail: arthur.ouwehand@danisco.com

Adrienne W. Paton Research Centre for Infectious Diseases, School of Molecular and Biomedical Science, University of Adelaide, Adelaide, SA 5005, Australia

e-mail: adrienne.paton@adelaide.edu.au

James C. Paton Research Centre for Infectious Diseases, School of Molecular and Biomedical Science, University of Adelaide, Adelaide, SA 5005, Australia e-mail: james.paton@adelaide.edu.au

A. S. Peña Department of Pathology, VU University Medical Center, Laboratory of Immunogenetics, P.O. Box 7057, 1007 MB Amsterdam, The Netherlands e-mail: pena.as@gmail.com

Jørgen Petersen The Proteomics Group, Danish Technological Institute, Holbergsvei 10, 6000 Kolding, Denmark e-mail: jpn@teknologisk.dk

Serge Rezzi BioAnalytical Science, Metabolomics & Biomarkers, Nestlé Research Center, P.O. Box 44, 1000 Lausanne 26, Switzerland e-mail: serge.rezzi@rdls.nestle.com

Irena Rogelj Department of Animal Science, Biotechnical Faculty, University of Ljubljana, Groblje 3, 1230 Domžale, Slovenia e-mail: irena.rogelj@bfro.uni-lj.si

Mauro Rossi Institute of Food Sciences, CNR, via Roma 64, 83100 Avellino, Italy e-mail: mrossi@isa.cnr.it

Denis Roy Departement of Food science and Nutrition, Institute of Nutraceutical and Functional Foods, Université Laval, 2440 Boul. Hochelaga, G1 V 0A6 QC, Canada

e-mail: denis.roy@inaf.ulaval.ca

Eduardo Jorge Schiffrin Nestlé Nutrition, R & D, Nestec Ltd, 22 Avenue Reller, 1800 Vevey, Switzerland e-mail: eduardo.schiffrin@nestle.com

Kathryne B. Schwartz Institute of Food Sciences, CNR, via Roma 64, 83100 Avellino, Italy e-mail: schwa21k@gmail.com

Alain L. Servin Faculté de Pharmacie, Inserm Unité 756, 92296 Châtenay-Malabry, France e-mail: alain.servin@u-psud.fr

Stefania Silvi Department of "Scienze Morfologiche e Biochimiche Comparate", University of Camerino, Via Gentile III da Varano, 62032 Camerino, Italy e-mail: stefania.silvi@unicam.it

R. D. Sleator Department of Biological Sciences, Cork Institute of Technology, Rossa Avenue, Bishopstown, Cork e-mail: roy.sleator@cit.ie

Siok-Koon Yeo School of Industrial Technology, Universiti Sains Malaysia, 11800 Penang, Malaysia e-mail: siokkoon85@yahoo.com

Annina Zihler Laboratory of Food Biotechnology, Institute of Food Science and Nutrition, ETH Zurich, Schmelzbergstrasse 7, 8092 Zurich, Switzerland e-mail: zihlera@ilw.agrl.ethz.ch

Part I Introduction and History of Probiotics

Chapter 1 Probiotics: From the Ancient Wisdom to the Actual Therapeutical and Nutraceutical Perspective

Giuseppe Caramia and Stefania Silvi

1.1 **Probiotics**

1.1.1 The Beginnings of Probiotics: The Fermented Milk

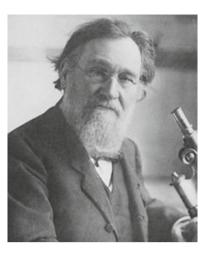
The recent history of probiotics began in the early 1900s. Thanks to Metchnikoff (1845–1916) (Fig. 1.1), professor of biology at the University of Odessa, who moved from Ukraine, his homeland, to Messina (Italy) for political reasons after the assassination of Czar Alexander II. In 1882 he discovered the mechanism of phagocytosis and cell-mediated immunity, for which he received the Nobel Prize in 1908, and in 1888, moved to Paris to work at the institute directed by Pasteur, pushed his research on the conditions and the organic alterations that promote aging. At Pasteur's death in 1895, he became the Director of the famous Pasteur Institute and continued his studies in various fields of knowledge and philosophy becoming famous among the general public for his books (*The Nature of Man*, 1904; *The Prolongation of Life*, 1906, etc.).

Starting from the studies of Pasteur on seething microorganisms, and of other researchers on the intestinal bacterial flora (Carre 1887; Tissier 1906), considering that the Caucasian shepherds had a longer average life than the inhabitants of Paris and, according to reports at the time, than the Americans (87 years against 48), he suggested that the shepherds' longevity depended on fermented milk, which they largely consumed, since it was a source of "good" and "anti-putrefactive" microorganisms. It was indeed known that the food wastes ferment in the colon due to some intestinal microorganisms and he was convinced that the putrefactive flora produces toxins, lethal in the long time.

G. Caramia (🖂)

- Hemeritus Head Physician of Paediatrics and Neonatology, Specialized Maternal-Infantil Hospital "G.Salesi", Via Toti 1, 60123 Ancona, Italy Tel.: +39 071 36938, +39 335 6166470
- e-mail: caramiagm@libero.it

Fig. 1.1 Elias Metchnikoff (1845–1916)



Really, the history of fermented milk and yogurt, with their excellent nutritional properties, was born together with man, in the earliest times of antiquity, most probably 500,000 years ago, when our ancient progenitors learned to light the fire defending themselves from the cold, keeping out animals, lighting the caverns, cooking the game and therefore many millennia before the beginning of the pasture and livestock. The use of fire, fermented milk and yogurt are thus part of human history and their role has been with humanity, to date, between legends and historical data (Flandrin and Montanari 1977; Perles 1977).

The need to keep such a precious food must have been felt since the beginning, and an ancient legend tells of a merchant who, having to cross the desert, brought some foods with him, including milk placed in a bag made with the dried stomach of a sheep. The enzymes remained on the wall of the sheep's stomach used as container, acidified milk and clotted its proteins in small lumps, giving rise to the curd and discovering cheese. The same phenomenon happened to the primordial yogurt derived from the acid fermentation of milk sugars. Thanks to the contamination with special milk enzymes, and a kind of liquid yogurt, used for many millennia by nomadic shepherds and people from the East. Certainly, it was used by the Indians and Sumerians in the fourth century BC, at the beginning of the Egyptian Civilization in the IV–III millennium BC, by the Phoenicians in the III–II millennium BC. The Bible, dated to the thirteenth century BC, reports that "Abraham offered to God, showed in an oakwood, fermented milk" (Genesis 18, 1–8) and Isaiah (VIII BC, 7:15) also says that "you will eat curdled milk and honey."

The Greek historian Herodotus (484–425 BC), Xenophon (430–355 BC), and Aristotle (384–322 BC) have spoken on the use of the yogurt (Bresciani 1977). At the time of the ancient Greeks and Romans, the consumption of fermented milk was recommended as a tonic, especially for children and convalescents, and the Greek physician Galen (129–216 AD), lived in the Imperial Rome, extensively spoke about the yogurt in one of his works, giving to it certain beneficial effects for both the liver and the stomach.

In the Middle Ages, fermented milk and cheese was mainly produced at the abbeys and convents, and they appear in the Crusaders' chronicles; later, we can find them in very distant populations such as Bulgarian shepherds, the Hindus, the Calmucchi, in France, at the court of Francis I (1494–1547), the Zulu, the Russians and other peoples of the Ottoman Empire that used yogurt, a term that derives from the Turkish *yogur* (kneading or mixing with a tool), as a panacea to purify the blood, to prevent tuberculosis, to solve some intestinal disorders and even to help sleeping.

It was known that fermentation is a very important aspect in the formation of yogurt, but the origin of such fermentation was still unclear.

The presence of invisible microorganisms (or micro-Dei), which can creep into our bodies causing diseases, is already present in trace in some Chinese legends and in ancient Egyptian medical texts. Afterwards, Marco Terenzio Varrone (116 BC–27 AD) before and Girolamo Fracastoro (1478–1553) later, talk about it openly. The existence of small organisms, called "animalculi", in the genesis of the diseases and of many other unclear phenomena, was firstly postulated by Lazzaro Spallanzani (1729–1799), who in 1780 coined and introduced into the medical literature the term "germ", so he is considered the founder of the experimental microbiology.

This was opposed to the "spontaneous generation" theory, for which the life is born in a "spontaneous" way from inert or inanimate matter by the effect of some "vital flows", a theory supported until then by the Aristotelian school disciples, by the Epicurean School, by famous philosophers of the Renaissance and in the eighteenth century by Georges-Louis Leclerc, Count of Buffon (1707–1788), and by John Turberville Needham (1713–1781). This dispute continued for many years and was finally permanently settled by Louis Pasteur (1822-1895) in 1864 which made light of that argument confirming the Spallanzani's thesis and thus winning the prize of the Science Academy of Paris for having clearly demonstrated the germs source. Pasteur arrived at such result, thanks to his studies on the fermentation of beer (1854), wine and vinegar (1861–1862) and on the deterioration of the wine by fungi or bacteria (1863–1864); findings confirmed in the following years by studies on silkworm disease (1865–1870), chickens cholera (1880), anthrax in bovines, sheep, horses (1881). In this route it was crucial, of course, the availability of the microscope, "small glasses to see minimal things nearly" that "multiplies things perhaps fifty thousand times" as his discoverer Galileo Galilei wrote (1564–1642) (Saggiatore: 1623), which significantly evolved over the past two centuries mainly thanks to Anton van Leeuwenhoek (1632–1723) and of his successors, thus triumphantly entering in the scientific research field (Caramia 2000).

1.1.2 From the Intuition to the Yogurt

Using bacterial strains selected from the milk of Caucasian and Bulgarian shepherds, through fermentation and acid coagulation of milk by the two microorganisms, *Streptococcus thermophilus* and *Lactobacillus delbruekii* subsp. *bulgaricus* (Fig. 1.2), is obtained a fermented milk, the "Lactobacilline", that in 1906 the

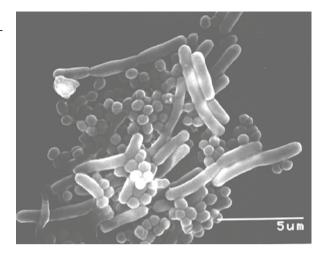


Fig. 1.2 Streptococcus thermophilus and Lactobacillus bulgaricus from yogurt matrix at scanning electron microscope. (By M. Benevelli—Dept. "Scienze degli alimenti", Bologna University, Italy)

French Society "Le Fermente" began to market and sold in pharmacies, according to the Metchnikoff's idea of helping children suffering from diarrhoea. The product obtained great success among the consumers: today French are the biggest consumers of yogurt compared with other European partners (including Italy), thanks also to the Greek entrepreneurs of Jewish origin, Isac and Daniel Carasso, who was born in Thessaloniki (in Spanish called Mr. Danone).

In 1907/1908 Metchnikoff in his book "The prolongation of life. Optimistic studies" confirms that not all microorganisms are harmful to human health and suggests that "The dependence of the intestinal microbes on the food makes it possible to adopt measures to modify the flora in our bodies and to replace the harmful microbes by useful microbes" (Metchnikoff 1907; Caramia 2008).

Some years later after his death, in 1925 it was sold a product called "yogurt" that rapidly spread in Europe and North America. However, there were also harsh critics since these microorganisms were not found in the faeces of "yogurt" consumers, than someone excluded any beneficial effect of the two seething bacteria. Metch-nikoff's intuition, based on empiricism, scientific observations and ingenious intuition, was then mocked by the scientific community, but the beneficial properties of yogurt remained in the collective imaginary, so its use was increasingly widespread.

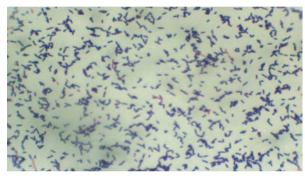
Always in the 20's, Minoru Shirota, a Japanese microbiologist at the University of Kyoto (Fig. 1.3), discovered that some bacteria of the intestinal flora contribute to bacterial pathogens defence. The following studies led to isolate and cultivate *Lactobacillus casei* (Lc) (Fig. 1.4), afterwards called Lc Shirota, and in 1935 in Japan began the production of a beverage containing this microorganism, called Yakult[®], that over the years was spread throughout the world.

An important contribution to the Metchnikoff's theory came in 1936 from two veterinarians, Zobell and Andersen, who suggested the existence in the large intestine of a "microbial film" made by the population of intestinal microorganisms adhering to the intestinal mucosa, which represents a "complex ecosystem with intensive metabolic activities".

Fig. 1.3 Minoru Shirota (1899–1982)



Fig. 1.4 Gram staining of *Lactobacillus casei Shirota*



1.1.3 The term "Probiotic" and its Technical-Scientific Evolution

Metchnikoff has the worth of having introduced the concept of probiotic microorganisms, from the Greek "pro-bios", for life, even if the origin of the term "probiotic" (to be distinguished from lactic ferments that are bacteria of not human origin and producing lactic acid) should be attributed for some to Kollath (1953) and for others to the German researcher Ferdinand Vergin, who in 1954 proposed to use the term "Probiotika" for the "active substances that are essential for a healthy development of life" (Vergin 1954).

In an article published in *Science* in 1962 two veterinarians, Lilly and Stillwell, very likely not knowing the Vergin's proposal, called "probiotics" the so-called "lactic ferments," that is "anaerobic bacteria able to produce lactic acid, starting from different dietary substrates, and to stimulate the growth of other microorganisms" (Lilly and Stillwell 1965).

The last term, also used in contrast to the antibiotic one (against life), which in 1960 was at its peak, thanks to the discovery and development of some important

new drugs with antibacterial action that changed the history of the anti-infective therapy, comes in the current use, not only in medicine. With the advance of knowledge on the physiological and therapeutic role of probiotics, the probiotic definitions became increasingly elaborate and exhaustive. So Parker in 1974 was the first man to use that term to identify the microorganisms- based supplements used for zootechnical feeding, defining them as: "organisms and substances which contribute to intestinal microbial balance" (Parker 1974). This new concept has been successful, especially through the work of a British microbiologist, Roy Fuller, specialized in the study of lactic acid bacteria, who in 1989 deleted from the definition the "substances" giving probiotic capabilities to microorganisms only: "a live microbial feed supplement which beneficially affects the host animal by improving its intestinal microbial balance" (Fuller 1989).

Few years later, two Dutch researchers, Haven and Huis in't Veld, extended the definition including in the beneficial action of the probiotic microorganisms the microflora of both the uro-genital and the upper respiratory system. The probiotics become then: "mono-or mixed cultures of live microorganisms which when applied to animal or man, beneficially affects the host by improving the properties of the indigenous microflora" (Huis in't Veld et al. 1994).

It is currently accepted the probiotic definition formulated in 2001 by FAO/ WHO "Live microorganisms which when administered in adequate amount confer a health benefit to the host" (FAO/WHO 2001). Respecting the "Guidelines on probiotics and prebiotics" their characteristics can be summarized as follows:

- Must not lose its properties during storage;
- Must be normally present in the human intestine;
- Must be able to survive, to overcome the gastric barrier, resisting to the action of digestive gastric juice, intestinal enzymes and bile salts and colonize the intestine: for this reason, the minimum effective dose, which is very indicative because it depends on the strain and preparation used, is 10⁷ CFU/day;
- Must be able to adhere to and to colonize the intestinal cells: the bacterial membrane structure is involved in the mechanism of adhesion and direct switch with the mucosa, the surface proteins and possibly also the secreted ones. In this respect should be reported the possible apoptotic induction on neoplastic cell lines, recently highlighted, which opens possible therapeutic implications;
- Must exert metabolic functions at the enteric level, with beneficial effects for human health, and antagonism against pathogenic microorganisms by producing antimicrobial substances;
- Should not cause immune or otherwise harmful reactions and then be considered as safe (GRAS status: generally recognized as safe);
- Resistance to antibiotics must be intrinsic or due to genetic mutations, whereas
 if it is caused by a horizontal gene transfer (i.e. transposons, genomic DNA segments that breaks off to join another, conjugative plasmids carrying genes for
 resistance, virulent or temperate phages) his choice becomes more problematic;
- Must also be administered in adequate doses and have a favourable cost-efficacy ratio.

1.2 Prebiotics

Prebiotics are predominantly dietary fibers, particularly soluble, also called "colonic food", consisting of specific carbohydrates. Increasingly used by the food industry (beverages, sweets) since 1980 for modifying viscosity, emulsification capacity, gel formation, freezing point and colour of foods, prebiotics have been widely studied since the early 90's, while the spread of the probiotics use, to provide the optimal nutrients to encourage growth of beneficial intestinal microflora (symbionts).

In 1995 Gibson and Roberfroid defined prebiotics as "non-digestible substances that when consumed provide a beneficial physiological effect on the host by selectively stimulating the favourable growth or activity of a limited number of indigenous bacteria in the colon, and thus improves host health" (Gibson and Robertfroid 1995).

As beneficial effect of health by "selective stimulation of the growth" and "activity of a limited number of colonic bacteria" are difficult to verify, in recent years the authors revisited their concept and defined prebiotics as: "a selectively fermented ingredient that allows specific changes, both in the composition and/or activity in the gastrointestinal microflora that confers benefits upon host well-being and health" (Gibson et al. 2004; Roberfroid 2007; Kelly 2008).

Based on the last definition, prebiotics may have the following characteristics (Gibson and Robertfroid 1995; Gibson et al. 2004; Roberfroid 2007; Kelly 2008; de Vrese and Schrezenmeir 2008):

- must pass, almost undamaged and in adequate amount, the digestive processes occurring in the first section of the digestive tract (mouth, stomach and small intestine);
- must be a nutritional fermentable substrate for intestinal microflora, in order to selectively stimulate the growth and/or metabolism of one or a few bacterial species;
- should positively change the bacterial flora in favour of the acidophile protective one (bifidobacteria, lactobacilli); and finally they should induce systemic or luminal effects that are positive for the human health.

Prebiotics are present in many edible plants such as chicory, artichoke, onions, leeks, garlic, asparagus, wheat, bananas, oats, soybeans and other legumes. Many commercial prebiotics are obtained from vegetable raw materials, while others are produced by enzymatic way through the hydrolysis of complex polysaccharides or the trans-glycosylation of mono- or disaccharides, a beneficial system for mass production starting from simple sugars (sucrose and lactose). Anyway, the addition of prebiotics in foods must comply with the ESPGHAN (European Society for Pae-diatric Gatroenterology Hepatology and Nutrition) recommendations (Aggett et al. 2003; Roberfroid 2007) including:

- standard methods for the analysis of carbohydrates content in food;
- right labels with the indication of quality and quantity carbohydrates content;
- international databases;
- · knowledge of the origin, specific effects and indications for the use of prebiotics.

The natural and commercial prebiotics consisting of oligo- and polysaccharides that are not, or only to a small extent, hydrolized by the digestive enzymes of the human upper intestinal tract and reach intact the colon where they are selectively fermented, particularly from indigenous and exogenous bifidobacteria and lactic acid bacteria, act as a fermentable carbon sources for the colonic microflora.

The most popular, most widely commercially available and the most researched prebiotic compounds are oligosaccharides oligofructose, fructooligosaccharides (FOS), metabolized by the β -fructofuranosidase (β -Fru) enzyme, the polysaccharide inulin, and partly the *trans*-galacto-oligo-saccharides (TOS) metabolized by the β -galactosidase (β -Gal) enzyme (Gibson and Robertfroid 1995; Bouhnik et al. 2006; Kolida and Gibson 2007; Roberfroid 2007; de Vrese and Schrezenmeir 2008; Kelly 2008). Oligofructose, fructooligosaccharides (FOS) (a mixture of oligosaccharides consisting of 3-10 carbohydrate monomers) and inulin (a mixture of fructooligo- and polysaccharides), are bifidogenic, but there is a great deal of intraindividual variability in bifidogenic and anaerobe responses to those inulin-type prebiotics (some experts consider oligofructose, FOS and inulin as synonymous terms for "inulin-type probiotics", oligo- or polysaccharide chains comprised primarily of linked fructose molecules, and inulin HP for the long-chain, high-molecular weight mixes of inulin-type fructans with a degree of polymerization (DP) > 10) (Roberfroid 2007; Kelly 2008, 2009). The effects on other gut microorganisms, as well as pathogenic organisms, are inconsistent but oligofructose and FOS show nutrition and health relevant properties like a low cariogenicity, a low calorimetric value and glycemic index, and a moderate sweetness (30-60% of the sucrose value = 1-2 kcal/g) (Kelly 2008). For this reason they are used as sweeteners in syrup, tablets or powder. Other candidates as prebiotics, for which there are already promising data, but for someone not yet sufficient, are the gluco-oligo-saccharides (GOS) which are oligo or polysaccharide chains comprised primarily of linked galactose units and which stimulate the growth of bifidobacteria and lactobacilli species, the soy-oligo-saccharides (SOS) raffinose and stachiose, metabolized by the α -galactosidase (α -Gal) enzyme, the iso-malt-oligo-saccharides and more (Roberfroid 2007; Kelly 2009; Bruzzese et al. 2009).

1.3 Synbiotics

An alternative chance to modulate or balance the intestinal microflora is the use of pro-and pre-biotic together making synbiotic compounds, that are alimentary or pharmaceutical preparations that containing either one or more probiotic strains and prebiotic ingredients, exploit the synergy between the microorganisms activity and their support for the benefit of the intestinal microflora and, consequently, of the whole body.

In 1995 Gibson and Roberfroid defined synbiotic as "a mixture of probiotics and prebiotics that beneficially affects the host by improving the survival and implantation of live microbial dietary supplements in the gastrointestinal tract, by selectively

stimulating the growth and/or activating the metabolism of one or a limited number of health-promoting bacteria, and thus improving host welfare".

The simultaneous administration of both probiotics and a substrate that they can metabolize gives to the administered strains greater opportunities for the colonization and survival of probiotic organisms in the colon of the host by increasing or prolonging their beneficial effects: this is really the best strategy for their integration, because it improves the survival (increasing the product shelf life) and on the other hand it provides a specific substrate for the resident bacterial flora.

Theoretically, the synbiotics have better beneficial effect on intestinal flora than pro- and prebiotics by lowering the pH, promoting growth of potentially protective bifidobacteria and inhibiting of potentially pathogenic microorganisms, stabilizing the intestinal environment and releasing short-chain organic acids.

Inulin-type probiotics, FOS or GOS, as well as their synbiotic combination with probiotic bacteria, *L. plantarum, L. paracasei* or *B. bifidum* strains, increased bifidobacteria and lactobacilli and inhibited various human- and animal pathogenic bacterial strains (*Clostridium sp., E. coli, Campylobacter jejuni, Enterobacterium sp., Salmonella enteritidis* or *S. typhimurium*) (Kanamori et al. 2004).

The most used and already marketed synbiotics regard mixtures of oligofructose, FOS, GOS, with probiotic bacterial strains of *L. plantarum*, *L. paracasei*, *L. rhamnosus*, *B. bifidum* or *B. lactis*.

1.4 Various Genera of Probiotics

The majority of probiotic microorganisms belong to the genera *Lactobacillus* (Figs. 1.5 and 1.6) and *Bifidobacterium* (Fig. 1.7). There are also other genera of bacteria and some yeasts widely used and reported in Table 1.1 (Baffoni and Biavati 2008). Lactobacilli and bifidobacteria are Gram-positive lactic acid-producing bac-

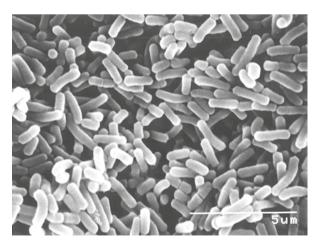


Fig. 1.5 Morphology of *Lac-tobacillus rhamnosus* at scanning electron microscope. (By M. Benevelli—Dept. "Scienze degli alimenti", Bologna University, Italy)

Fig. 1.6 Morphology of *Lac-tobacillus rhamnosus* from yogurt matrix at scanning electron microscope. (By M. Benevelli—Dept. "Scienze degli alimenti", Bologna University, Italy)

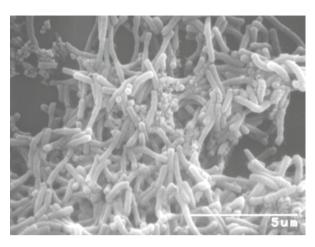
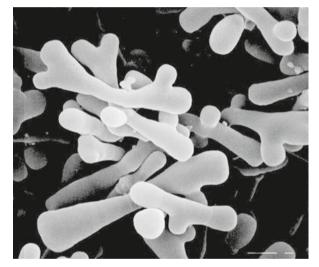


Fig. 1.7 Morphology of *Bifi- dobacterium* spp. at scanning electron microscope



teria that constitute a major part of the normal intestinal microflora in animals and humans. Lactobacilli are Gram-positive, non-spore forming rods or coccobacilli. They have complex nutritional requirements and are strictly fermentative, aerotolerant or anaerobic, aciduric or acidophilic. Lactobacilli are isolated from a variety of habitats where rich, carbohydrate-containing substrates are available, such as human and animal mucosal membranes, on plants or material of plant origin, sewage and fermented milk products, fermenting or spoiling food. Bifidobacteria constitute a major part of the normal intestinal microflora in humans throughout life. They appear in the faeces a few days after birth and increase in number thereafter. The number of Bifidobacteria in the colon of adults is 10^{10} – 10^{11} CFU/g, but this number decreases with age. Bifidobacteria are non-motile, non-spore forming, Gram-positive rods with varying cell morphology. Most strains are strictly anaerobic.

Lactobacillus	Bifidobacterium	Enterococcus	Streptococcus	Lactococcus			
L. acidophilus	B. adolescentis	E. faecalis	S. thermophilu	s L. lactis subsp. cremoris			
L. brevis	B. animalis	E. faecium		L. lactis subsp. lactis			
L. casei	B. bifidum						
L. curvatus	B. breve						
L. fermentum	B. infantis						
L. gasseri	B. longum						
L. johnsonii	B. thermophilum						
L. reuteri							
L. rhamnosus							
L. salivarius							
Propionibacter	ium	Yeast		Others			
P. freudenreich	ii	Kluyveromyces lactis L		Leuconostoc mesenteroides			
P. freudenreich	ii subsp. shermanii	Saccharomyces boulardii P		Pediococcus acidilactici			
P. jensenii		Saccharomyc	es cerevisiae				

Table 1.1 Microorganisms considered as probiotics. (By Baffoni and Biavati 2008, modified)

1.5 **Probiotics as Therapy**

The primordial milk enzymes at the beginning of last century, selected from the milk of the Caucasian and Bulgarian shepherds, have been sold according to the ideas of Metchnikoff and Tissier "to help children suffering from diarrhoea" and sold in pharmacies to bring "good and anti-putrefactive micro-organisms" because "not all microorganisms are harmful to human health".

In this light over the next few decades lactic acid bacteria with special features, now considered probiotics, kept the primary indication: the preventive-therapeutic use, particularly for some gastroenterological diseases, to try to restore and/or rebalance the functionality of microbiota, the intestinal mucosa and the immunological aspects, keeping in mind the indications listed in the guidelines about the evidence based medicine on the levels of scientific evidence and the strength of clinical recommendations.

1.5.1 Acute Infectious Diarrhoea

In most industrialized countries, acute infectious diarrhoea (AID) is now a minor disease because fatal cases are very exceptional. It is determined in about 70% by viral agents, such as rotavirus, which are responsible for 30–45% of all viral diarrhoea, calicivirus, including norwalk virus, enteric serotypes adenovirus 40 and 41, and Astrovirus; while among bacteria we should mention *Campylobacter jejuni* (main cause of diarrhoeal disease in adults in the US), *Salmonella, Shigella*, enteropathogens *Escherichia coli, and Yersinia enterocolitis*.

Firstly, it should be noted that not all probiotics are the same, because not all of them determine the same therapeutic effects, but, based on the levels of scientific evidence and the strength of clinical recommendations, it is believed appropriate to share in principle what was recently proposed by the ESPGHAN and by the European Society for Paediatric Infectious Disease (ESPID) and by many other scientists: "Probiotics may be an effective adjunct to the management of diarrhoea. However, because there is no evidence of efficacy for many preparations, we suggest the use of probiotic strains with proven efficacy and in appropriate doses for the management of children with acute gastroenteritis as an adjunct to rehydration therapy (levels of scientific evidence II and strength of clinical recommendations B). The following probiotics showed benefit in meta-analyses of RCTs: *Lactobacillus* GG (I, A), *L. reuteri* (I, A) and *Saccharomyces boulardii* (II, B)" (Floch et al. 2008; Guarino et al. 2008; Kligler and Cohrssen 2008).

In particular, *L. reuteri* has shown to shorten significantly the clinical course of rotavirus-induced gastroenteritis, as well as reducing incidence of acute diarrhoea (Figs. 1.8 and 1.9) (Shornikova et al. 1997a, b).

As for prevention of infectious diarrhoea, mostly of viral origin, which can be contracted at nursery schools, kindergartens or during hospitalization for other pathologies, it is not yet clear which probiotic or association of probiotics is more effective. Besides, the dose administered which must be equal to or greater than 5–10 billion CFU/day and the early initiation of therapy are important, so that the probiotic, with appropriate doses and immediately administered, may contrast the action of the pathogen (Floch et al. 2008; Guarino et al. 2008, 2009). More recently,

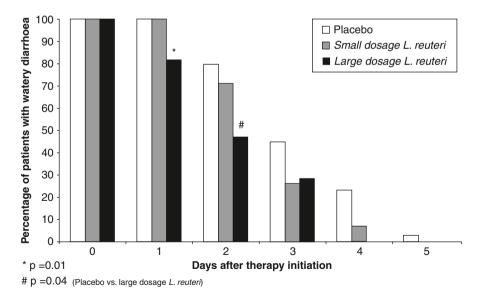


Fig. 1.8 Percentage of patients with persisting watery diarrhoea in the groups receiving placebo (n = 25) and small (n = 20) and large (n = 21) dosage of *L. reuteri* (Shornikova et al. 1997a)

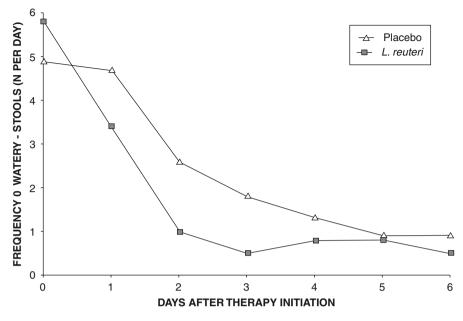


Fig. 1.9 Frequency of watery stools per 24-h period in patients receiving *L. reuteri* and placebo (Shornikova et al. 1997b)

Eom and colleagues showed the therapeutic effect of *L. reuteri*, administered at a dose of 2×10^8 CFU/die only, to significantly reduce acute diarrhoea in children (Eom et al. 2005).

1.5.2 Antibiotic Associated Diarrhoea

Antibiotics, aminopenicillins, cephalosporin, clindamycin etc., are much prescribed in all industrialized countries with several side effects especially in children: among which the most frequent is antibiotic associated diarrhoea (AAD). The resulting alteration of intestinal microflora reduces the development of anaerobic microflora, which leads to a reduced metabolism of carbohydrates and therefore to osmotic diarrhoea, favours the development of pathogens such as *Clostridium difficile, Salmonella, C. perfringens type A, Staphylococcus aureus* and *Candida albicans*. According to the recent studies, even in adults, there is level I of scientific evidence in favour of the use of probiotics in the treatment of AAD (Doron et al. 2008; Floch et al. 2008; Pham et al. 2008; Surawicz 2008). Therefore, there are grounds to recommend their use especially in risky cases, as in subjects where there is repeated use of antibiotics. This in an attempt to prevent inflammatory processes of the intestinal mucosa in children that can often lead to chronic inflammatory disease of the large intestine (Crohn's disease, ulcerative colitis, pouchitis) in subsequent years (Caramia 2008; Floch et al. 2008; Guandalini 2008). In a randomized, doubleblind, placebo-controlled pilot study, recently presented at the Clinical Nutrition Week 2009, patients receiving antibiotics were given *L. reuteri* (10⁸ CFU b. i. d.) or an identical placebo for 4 weeks. Patients treated with *L. reuteri* had a significantly lower incidence of diarrhoea (only 7.7%) compared to patients receiving placebo (50%) (Cimperman et al. 2009).

1.5.3 Clostridium difficile Associated Diarrhoea

The *Clostridium difficile* is the main cause of diarrhoea caused by antibiotics (CDAD) and of nosocomial colitis. It has been indicated as responsible for between 10% and 20% of all cases of diarrhoea caused by antibiotics, 60% of antibiotic-associated colitis and nearly all cases of pseudo membrane colitis. The diarrhoeal disease caused by *C. difficile* is determined only by the *C. difficile* strains producing the toxin A, who plays a mild cytotoxic activity and causes damage to the mucous, inflammation and intestinal secretion, and by toxin B, one of the most powerful cytotoxin, which determines loss of intracellular potassium, inhibition of protein synthesis and nucleic acids.

Unfortunately, the diversity of probiotics, their doses and the heterogeneity of studies make it difficult to recommend a definitive therapy, and also to indicate which probiotics to use as an antibiotic treatment and for prevention of *C. difficile* associated diarrhoea and/or colitis. For this reason, despite there are many promising data, the level of scientific evidence in favour of the use of probiotics or a combination of antibiotic and probiotic in the treatment of CDAD is currently of type II only (Doron et al. 2008; Floch et al. 2008; Guandalini 2008; Hookman and Barkin 2009; Yangco et al. 2009).

1.5.4 Infection Caused by Helicobacter pylori

Helicobacter pylori (HP) infection affects over 50% of the world's population and covers 80% of the population in the developing countries. HP infection is the main cause of peptic ulcer disease (70–90% of cases), lymphoma and in 1% of infected persons, leads to the development of gastric cancer with remarkable increase in mortality (Kelly and LaMont 2008; Jarosz et al. 2009). In developed countries, the infection starts in childhood, where it seems to have an incidence of 10–15%, then rapidly increasing during evolution (Sabbi et al. 2008). The transmission is orofaecal as the seed is located in the gingival bags and at the root of the tongue.

Several studies showed that patients treated with probiotics associated with the standard antibiotic therapy had higher rate of eradication with a minor number of side effects.

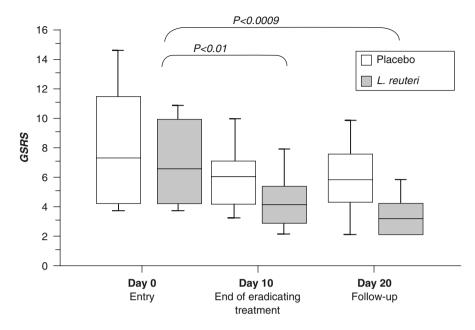


Fig. 1.10 Gastrointestinal symptom rating scale (GSRS) in children receiving *L. reuteri* or placebo. Comparison of continuous variables performed using Mann-Whitney. (Adapted from Lionetti et al. 2006)

A pilot study performed on 30 Hp positive adults treated with omeprazole + placebo or omeprazole + *L. reuteri* for 30 days, showed that 60% of patients in the *L. reuteri* group was eradicated, while none in the placebo group (Fig. 1.10) (Saggioro et al. 2005). Lionetti and colleagues in 2006 showed a reduction in gastro- intestinal symptoms by *L. reuteri* supplementation during and after the eradication therapy in a group of Hp infected children (Lionetti et al. 2006). Finally, Francavilla and colleagues in a recent pilot study conducted on 40 Hp positive adults, undergoing the standard eradication therapy, showed that the pre-administration of *L. reuteri* in the 4 weeks before treatment significantly reduces the bacterial load, and decreases the GI associated symptoms (Francavilla et al. 2008).

Excellent results are also reported by the administration of fermented milk enriched with probiotics (Tong et al. 2007; Sachdeva and Nagpal 2009). The HP infection in adults are related to the type and virulence of HP strain, the production of toxins A and B, the level of infection, the extent of inflammation, and the density of HP colonization. Therefore, there is a considerable interest in developing therapy to prevent HP infection. Probiotics intake, with suppression of HP, may have a favourable effect on HP infection and in decreasing the risk to develop related diseases; however, further, large, long-term and placebo-controlled studies are needed to confirm all those effects (Kelly and LaMont 2008; Selgrad and Malfertheiner 2008; McFarland 2009).

1.5.5 Traveller's Diarrhoea

The traveller's diarrhoea (TD) is the most common pathologic condition among travellers, and can occur in approximately 20–50% of the subjects during or immediately after a trip to a country with hot and humid weather conditions and inadequate sanitary conditions. More than one third of patients with TD had pathogennegative illness but in the case of food contaminated with pathogens, the most frequent are enterotoxigenic *Escherichia coli, Campylobacter jejuni, Shighella, Salmonella*, sometimes virus, such as *rotavirus, calicivirus, enterovirus*, or parassites such as *Giardia lambdia, Entamoeba histolytica, Cryptosporidium parvum,* and *Cyclospora cayetanensis* are less common causes of TD (McFarland 2007; Johnson et al. 2009).

Sometimes serious but of transitory nature, it is characterized by non formed or liquid evacuations, often accompanied by nausea, vomiting, abdominal cramps, tenesmus, in 10% of cases stool blood, occasionally fever. The evolution is favourable in 1–5 days, but sometimes it needs up to 10 days. It has also been reported that 2-10% of travellers with TD develop persistent diarrhoea and that about 10% of them suffer from post infectious irritable bowel syndrome (Caramia 2008).

In a recent meta-analysis of probiotics for the prevention of TD on randomized, controlled, blinded efficacy trials in humans diarrhoea, several probiotics, *S. boulardii* and a combination of *L. acidophilus* and *B. bifidum*, had significant efficacy (Takahashi et al. 2007; Guslandi 2008).

There are well founded reasons to believe that probiotics may be a safe and effective strategy to prevent TD, to which one can associate racecadotril an inhibitor of intestinal encefalinasis and therefore antisecretive and antidiarrhoeic action, but continued research are needed (Tormo et al. 2008; Vandenplas et al. 2009).

1.5.6 Necrotizing Enterocolitis

The necrotizing enterocolitis (NEC), inflammatory intestine disease with an incidence of 1 to 3/1,000 live births, is relatively common in preterm in the first six weeks of life, and leads to death in 15-30% of those with lower weight to 1,500 g and the 20-40% of cases requiring surgery.

The causes of the disease appear to be multifactorial and are represented by prematurity, hypoxia-ischemia of the intestine, extensive use of antibiotics, reduced exposure to maternal microflora and excessive exposure to the typical sections of neonatal intensive care (NICU) (Staphylococci, Enterobacteriaceae, enterococci, *Candida* spp.), or to the use of sterile food as an alternative to mother's milk.

Based on several multicentre, randomized, double blind investigations there are reasonable grounds to believe that despite the differences between tested probiotics, the beginning of administration, dose and duration of the treatment and groups of participants, the enteral supplementation of some probiotics may reduce the risk of severe NEC and mortality in premature newborn with weight >1,000 g. There is therefore a level I of evidence, even if a deeper understanding of NEC pathogenesis and the mechanisms by which probiotics prevent this disease is needed (Dani et al. 2002; Bin-Nun et al. 2005; Deshpande et al. 2007; Alfaleh and Bassler 2008; Lin et al. 2008; Caplan 2009).

1.5.7 Bacterial Vaginosis

Bacterial vaginosis (BV) is the most prevalent vaginal infection worldwide and is characterized by depletion of the indigenous lactobacilli. The vaginal microflora is composed by several species of lactobacilli (*L. acidophilus, L. brevis, L. casei* spp. *paracasei, L. rhamnosus*) named "Doderlein" microflora. They create on the mucosa a bacterial biofilm able to inhibit the growth and the adhesion of pathogenic bacteria. This microflora presents variations related to the life style (diet, stress, sexual habits, etc.) which may cause quali-quantitative modifications of the normal environment and the introduction of several kinds of pathogens such as *Gardnerella vaginalis* and *Candida albicans*.

Antimicrobial therapy is often ineffective while the probiotic approach, either topic or combined with the oral administration, gave interesting results. This has been confirmed by *L. rhamnosus* strain (Lcr35) showing the ability to adhere to cervical and vaginal cells and to affect the viability of two main vaginosis-associated pathogens, *Prevotella bivia, Gardnerella vaginalis*, as well as *C. albicans* (Coudeyras et al. 2008).

In a trial eighty-four patients with bacterial vaginosis were randomized to receive either oral metronidazole 500 mg twice a day for seven days, or one vaginal tablet containing freeze-dried *L. rhamnosus* once a week at bedtime for two months starting one week after the last antibiotic administration. Chi-squared analysis showed a significant difference between the two treatment groups at day 90 (p = 0.05). Safe and effective long-term vaginal administration of *L. rhamnosus* appears to be a useful complementary approach in the management of bacterial vaginosis (Marcone et al. 2008).

Recently sixty-four women diagnosed with BV were randomly assigned to receive a single dose of tinidazole (2 g) supplemented with either 2 placebo capsules or 2 capsules containing *L. rhamnosus* GR-1 and *L. reuteri* RC-14 every morning for the following 4 weeks. After a treatment of 28 days, the probiotic group had a significantly higher cure rate of BV (87.5%) than the placebo group (50.0%) (p = 0.001). In addition, according to the Gram-stain Nugent score, more women were assessed with "normal" vaginal microbiota in the probiotic group (75.0% versus 34.4% in the placebo group; p = 0.011) (Martinez et al. 2009a). This study shows that probiotic lactobacilli can provide benefits to women with BV and that probiotic capsules containing *L. rhamnosus* GR-1 and *L. reuteri* RC-14 can increase the effectiveness of an antifungal pharmaceutical agent (fluconazole) (Martinez et al. 2009b).

1.5.8 Irritable Bowel Syndrome

The irritable bowel syndrome (IBS), a disorder of the function of the intestine, affects 3–25% of the general population and is characterized by pain at the abdomen, constipation alternating with periods of diarrhoea, presence of air and a sense of abdominal bloating. This is due to psychological factors, including sexual abuse, because the central nervous system interacts with intestinal neurotransmitters and hormones, to dietary factors resulting in alteration of intestinal motility, and abnormal dismicrobism, colic fermentation by bacteria and sometimes a secondary gastrointestinal infection (e.g. *Campylobacter* or *Salmonella*) (Belaise et al. 2002; Mättö et al. 2005).

Recent reviews of the literature provide interesting data also on gut motility and pain perception, however the results, sometimes contradictory, are not yet very clear and the effectiveness of probiotics in IBS show a level of scientific evidence of type I (Kunze et al. 2009). Very likely the strength of the recommendations of type B is due to the limited number of patients treated with different probiotics (Jiménez 2009). It is therefore considered useful to continue with more appropriate and thorough studies on those probiotics that have shown a more promising result.

1.5.9 Crohn's Disease

Crohn's disease, a gastrointestinal disorder characterized by chronic inflammation or ulceration involving all layers of the intestinal wall, is one of a group of diseases called inflammatory bowel disease (IBD) such as Crohn's disease (CD), ulcerative colitis (UC) and pouchitis. IBD are most common in the developed countries of Europe, the US, and Scandinavia, and less in Southern and Eastern Europe, in Asia or Africa, and the diseases are relatively uncommon in Cuba and Central and South America (Loftus 2004). This has given rise to a number of theories regarding IBD etiology and the significance of diet high in refined sugar, meat, milk, eggs and low in fiber, fruit, and vegetables.

CD can affect any area from the mouth to the anus but often affects the lower part of the small intestine, the ileum, while the colon is the second most common site of involvement. Prevalence of Crohn's disease is approx 0.18% and is more frequent in young adults between 15 and 30 years of age (Wallace 2009).

The intestinal microbiota play an important role in the pathogenesis and maintenance of disease. Although marked alterations occur in faecal and mucosal bacterial communities in CD and others IBD, it is unclear whether they are responsible for causing disease, or are due to changes in the gut environment that result from inflammatory reactions and extensive tissue destruction that later concur to maintain the pathological condition.

A study has been recently conducted on a limited number of patients treated for 13.0 ± 4.5 months with a mixture of probiotics composed of *B. breve* 30×10^9 CFU/