

The Influence of Milk Oligosaccharides on Microbiota of Infants: Opportunities for Formulas

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Abstract

In addition to a nutritive role, human milk also guides the development of a protective intestinal microbiota in the infant. Human milk possesses an overabundance of complex oligosaccharides that are indigestible by the infant yet are consumed by microbial populations in the developing intestine. These oligosaccharides are believed to facilitate enrichment of a healthy infant gastrointestinal microbiota, often associated with bifidobacteria. Advances in glycomics have enabled precise determination of milk glycan structures as well as identification of the specific glycans consumed by various gut microbes. Furthermore, genomic analysis of bifidobacteria from infants has revealed specific genetic loci related to milk oligosaccharide import and processing, suggesting coevolution between the human host, milk glycans, and the microbes they enrich. This review discusses the current understanding of how human milk oligosaccharides interact with the infant microbiota and examines the opportunities for translating this knowledge to improve the functionality of infant formulas.

INTRODUCTION

The process of bacterial colonization of the intestine begins naturally in a stepwise manner with three phases: delivery, breastfeeding, and weaning (Penders et al. 2006, Sherman et al. 2009). By the age of 18 months, the colonic bacterial microbiota is considered complete (Harmsen et al. 2000, Palmer et al. 2007, Rubaltelli et al. 1998). The gut microbiota of breastfed infants is modulated by human milk, the predominant diet of newborn infants. Milk is a truly unique food that has been shaped by mammalian evolution to provide both nutrition and protection to the developing infant, all at an energetic cost to the mother. As a result of this distinctive evolutionary tension, milk is unlikely to have retained superfluous contents that do not benefit the infant, as their presence comes at a cost to the mother. The benefits of breastfeeding in terms of infant development and protection have been well documented (Wu & Chen 2009). Various factors present in milk are known to modulate the developing microbiota within the infant gastrointestinal tract (GIT), including immunoglobulins, lactoferrin, lysozyme, bioactive lipids, leukocytes, and various milk glycans (glycolipids, glycoproteins, and free oligosaccharides) among others (Newburg 2005). Although some of these bioactive components are known for their functionality in reducing pathogens in the infant GIT, others are believed to encourage specific bacterial populations, such as bifidobacteria—a genus first identified more than 100 years ago in the feces of breastfed infants (Moro 1900, Tissier 1900). Such a prebiotic function of human milk was originally described by Gyorgy and coworkers (1954), who first identified *N*-acetyl-glucosamine (GlcNAc)-containing oligosaccharides, generally termed human milk oligosaccharides (HMOs), as the bifidus factor responsible for enrichment of bifidobacteria (Gauche et al. 1954, German et al. 2008).

HMOs are believed to have many roles in a developing infant in addition to putative prebiotic functions. HMOs may possess antiadhesive effects that reduce the binding of pathogenic bacteria to colonocytes (Lane et al. 2010). HMOs have modulating effects on immunologic processes at the level of gut-associated lymphoid tissue (Guarner 2009) and may also decrease intestinal permeability in preterm infants in a dose-related manner in the first postnatal month (Taylor et al. 2009). Others have suggested that HMOs are an important source of *N*-acetyl-neuraminic acid (NeuAc; sialic acid), an essential monosaccharide during the period of neonate brain development and myelination (Wang et al. 2001).

The newborn infant gastrointestinal tract is initially colonized by aerobic and facultatively anaerobic bacteria, often species of enterobacteria, enterococci, and staphylococci (Adlerberth & Wold 2009). As these initial bacteria consume oxygen present in the intestine, anaerobic genera such as bifidobacteria, clostridia, and bacteroides are enriched. A number of studies have demonstrated that bifidobacteria are overrepresented in the gastrointestinal microbiota of breastfed infants by comparison with adults (Favier et al. 2003, Harmsen et al. 2000, Mariat et al. 2009, Penders et al. 2006). More recently, some studies have illustrated the sporadic and individualized nature of microbial colonization of infants (Koenig et al. 2010, Palmer et al. 2007). These recent approaches have also challenged the often-reported observation of bifidobacterial dominance of the breastfed infant GIT. Unfortunately, technical biases in these studies derived from sequencing V2 16S rDNA region amplicons may have led to an underestimation of the actinobacterial clade—as has been observed in studies comparing different 16S variable region amplicons (Turnbaugh et al. 2006) or by comparing to complementary metagenomic approaches (Koenig et al. 2010). Regardless, these new approaches hold significant promise in more comprehensively characterizing the influence of breast milk on the developing infant gut microbiota and its inherent metabolic capacity.

In a recent review of a number of studies undertaken in the past 20 years, Adlerberth & Wold (2009) reported only minor differences in the levels of bifidobacteria present in formula-fed

infant feces compared with breastfed infant feces, contrasting a commonly held perception of a low bifidobacterial presence in formula-fed infants. Certain microbial clades, such as clostridia, bacteroides, and *Enterobacteriaceae*, are more often observed in formula-fed than breastfed infants, resulting in a common description of formula-fed infants as having a more adult-like gastrointestinal microbiota (Adlerberth & Wold 2009). Moreover, a recent study noted that the specific bifidobacterial species diversity present in formula-fed infants is more adult-like (Haarman & Knol 2005).

The World Health Organization has clearly identified breastfeeding as providing the optimum nutrition and protection for developing infants (WHO 2009). In spite of this advice, there remains a great need for optimal infant formulas as a substitute for breastfeeding in cases where the latter is simply not possible. Recent advances in our understanding of the complex functions of human milk create a conceptual path for the design of more functional formulas. Such formulas would not only provide the complex nutritional needs for infants but also facilitate the microbial successions witnessed in breastfed infant GIT.

A variety of options are available for specific modification of microbial colonization in infants fed formula. Besides adding live bacteria, such as bifidobacteria and lactobacilli as probiotic additives, prebiotic oligosaccharides can be added as substrates that arrive undigested to the colon (Boehm & Moro 2008) and stimulate the growth and/or metabolic activity of beneficial bacterial species such as bifidobacteria (Manning & Gibson 2004). Interestingly, the structure of HMOs, comprising more than 200 different molecular structures (Bode 2006, Coppa et al. 2004, Ninonuevo et al. 2006), differs significantly from plant-derived fructooligosaccharides (FOS) or enzymatically synthesized galactooligosaccharides (GOS) (Fanaro et al. 2005a). To date, the variable oligosaccharide content of human milk cannot be successfully reproduced on a large scale for inclusion in infant formulas. (German et al. 2008, Guarner 2009, Manning & Gibson 2004, Sarney et al. 2000, Taylor et al. 2009, Wang et al. 2001). This manuscript focuses on the latest discoveries in the field of milk oligosaccharides as well their potential use in infant formula. In addition, structure and effects of bovine milk oligosaccharides (BMOs) and plant-derived oligosaccharides are briefly compared with those of HMOs.

HUMAN MILK OLIGOSACCHARIDES

HMOs are the third most abundant component of human milk (Kunz et al. 2000). Among all of the components, such as proteins, lactose, and nucleotides, the HMO is the only component that has been demonstrated to play a significant role in the stimulation of the growth of specific bacteria (Coppa et al. 2006). HMO-like structures are also found as components of glycolipids and glycoproteins (Newburg 1999). There are approximately 200 known compositions incorporating ≥ 3 carbohydrate monomers via 13 possible glycosidic linkages (Kunz et al. 2000, Ninonuevo et al. 2006). The molecular structure of these oligosaccharides is highly variable; additionally, the composition and concentration change significantly during lactation (described below) (Boehm & Moro 2008). After ingestion, HMOs pass mainly unabsorbed through the small intestine into the colon, where they are fermented to short-chain fatty acids (SCFA) and lactic acids, creating an acidic environment (Ogawa et al. 1992).

HMO Composition and Structure

The composition of milk oligosaccharides, as well as other milk components, differs among mammalian species and also during the course of lactation. Oligosaccharides in human milk are characterized by an enormous structural diversity (Chaturvedi et al. 2001). HMOs are formed by

the attachment of a single glucose (Glc) molecule at the reducing end to galactose (Gal; bound to the Glc) to form a lactose core (Bode 2006). A linear chain is formed via β 1–3 linkage attached to the core structure of GlcNAc, whereas a branched chain results when two GlcNAcs are added on both the β 1–3 and β 1–6 positions (Wu et al. 2010). After addition of the GlcNAc, another Gal is added either at β 1–4 or β 1–3. The resulting GlcNAc and Gal disaccharide may repeat multiple times. There are at least 12 different types of glycosidic bonds described in HMO (Newburg et al. 2005, Kobata 2003).). The smallest oligosaccharides are generated either when fucose (Fuc) is added to lactose, thus generating the trisaccharide fucosyllactose (2'FL; Fuc α 1–2, Gal β 1–4Glc, and 3'FL; Gal β 1–4[Fuc α 1–3]Glc), or when NeuAc is added to lactose, generating the sialyllactoses (3'SL; NeuAc α 1–3Gal β 1–4Glc and 6'SL; NeuAc α 1–6Gal β 1–4Glc) (Espinosa et al. 2007). However, these small oligosaccharides are generally less abundant than the larger, more complicated structures.

The synthesis of these oligosaccharides within the lactating mammary gland is catalyzed by a number of specific glycosyltransferases, including galactotransferases, *N*-acetylglucosaminyltransferases, fucosyltransferases, and sialyltransferases, whose expression is required for the synthesis of various glycoconjugates that are normally found in both lactating and nonlactating mammary tissue (Kelder et al. 2001). Because the human intestine does not express the luminal enzymes to cleave the α -glycosidic linkages of Fuc and sialic acid, as well as β -glycosidic linkages in the core HMO molecule, these acids are resistant to enzymatic cleavages in the intestine (Engfer et al. 2000, Gnoth et al. 2000). As a result, HMOs can be detected in the feces of breastfed infants (Coppa et al. 2001). HMOs are also absorbed *in vivo* (Engfer et al. 2000, Gnoth et al. 2001, Kunz et al. 2000, Rudloff et al. 2006) through the intestinal wall in small amounts, possibly by receptor-mediated endocytosis (\sim 1% of intake) and can be detected in urine (Coppa et al. 1990, Coppa et al. 2001). In this process, HMOs are taken unmodified up via trans- and para-cellular pathways (Gnoth et al. 2001).

HMOs are especially rich in the type 1 oligosaccharides. Lacto-*N*-biose (LNB; Gal β 1–3GlcNAc) is a building unit of the three type 1 HMOs, such as lacto-*N*-tetraose (LNT; Gal β 1–3GlcNAc β 1–3Gal β 1–4Glc), lacto-*N*-fucopentaose I (LNFP; Fuc α 1–2Gal β 1–3GlcNAc β 1–3Gal β 1–4Glc), and lacto-*N*-difucohexaose I (Fuc α 1–2Gal β 1–3[Fuc α 1–4]GlcNAc β 1–3Gal β 1–4Glc). More detail on the structure and function of selected HMOs is available in a recent review by Bode (2006).

Although researchers began describing analytical methods for the isolation and characterizing of HMO over 40 years ago (Kobata & Ginsburg 1969), only recently has the technology advanced to precisely present and differentiate those complex glycans. Recently, Ninonuevo & Lebrilla (2009) discussed in detail the current methods for analysis of oligosaccharides in human milk. In recent years, more published data on the structure and function of HMOs allowed defining of the Lewis and Secretor blood groups corresponding to the specific HMOs produced by lactating mothers (Kunz & Rudloff 2008).

Lewis Blood Group and Secretor Status

Studies have shown that milk from different mothers may be qualitatively and quantitatively different with regards to its oligosaccharide content (Newburg 2000). A close relationship exists between HMO profiles, the structures of milk oligosaccharides, and the Lewis Blood Group and Secretor status (Kobata 1992, Thurl et al. 1997). The main criteria in the predicted variability of phenotypes seems to depend on the expression and activity of specific fucosyltransferases in the lactating mammary gland (Kunz & Rudloff 2008). Fucosyltransferase and fucosidase activities vary in milk specimens, both from different donors and from the same donors at different stages

of lactation (Wiederschain & Newburg 1996). Additionally, these differences might also apply to other glycoconjugates (e.g., glycoproteins that are constructed via the same glycosyltransferases that synthesize HMOs). In human milk from individuals with blood type Le(a – b +) (~70% of the population), HMO with α 1-2, α 1-3, and α 1-4-linked fucosyl residues occur (Kobata 1992). The second group, Le(a + b –) (20%) lacks compounds with α 1-2-linked fucosyl compounds. Finally, in the remaining 10% of the population (Le(a – b –)), oligosaccharides with α 1-4-linked fucose residues are missing. This system could potentially play a significant role in synchronizing select bacterial microbiota in infants with the mother's blood group. Population-based studies support this concept, and a high level of α 1-2-linked Fuc relative to total HMOs has been shown to correlate with a lowered incidence of infant diarrhea (Morrow et al. 2005).

Variations in HMO Production

HMOs are solely produced in the lactating mammary gland and vary over the course of lactation (Chaturvedi et al. 2001). HMOs achieve a maximum concentration in the colostrum (above 20 g L⁻¹) with 5–14 g L⁻¹ in mature milk (Kunz et al. 2000, Kunz et al. 1996). One limitation in the attempts to characterize HMO is a lack of availability in sufficient quantities and purity for in vitro and clinical studies. The majority of clinical studies using breast milk have to be interpreted carefully because assumed biological effects of HMO have also been credited to glycoproteins, glycolipids, and other milk constituents (Espinosa et al. 2007), and numerous other glycoconjugates might share the structural features with HMOs.

Given that various functions are associated with the diverse HMO structures, the details of variations in composition and differences of oligosaccharides among humans in remote populations need to be defined. Current analytical methods to characterize oligosaccharides in human milk include high-performance liquid chromatography (HPLC) (Chaturvedi et al. 1997, Leo et al. 2010), high pH anion exchange chromatography (HPAEC) (Thurl et al. 1996), capillary electrophoresis (CE) (Bao & Newburg 2008, Shen et al. 2000), and mass spectrometry (MS) (Albrecht et al. 2010, LoCascio et al. 2007, Marcobal et al. 2010, Niñonuevo & Lebrilla 2009). A method for precise quantification of consumption of individual HMOs named matrix-assisted laser desorption/ionization-Fourier transform ion cyclotron resonance mass spectrometry (MALDI-FTICR MS) allows detection of individual neutral oligosaccharides, which represent the majority of total HMOs (Ninonuevo et al. 2006).

Chaturvedi and coworkers have demonstrated that the ratio of fucosylated α 1-2-linked oligosaccharide concentrations to oligosaccharides devoid of α 1-2 linked Fuc changed during the first year of lactation from 5:1 to 1:1 (Chaturvedi et al. 2001). Furthermore, the concentrations of individual oligosaccharides varied substantially, both between the mothers and over the course of lactation. Those results suggest that the protective activities of HMOs might vary among the individuals and during the lactation. Interestingly, the attachment of Fuc was shown to be based on the Secretor status and Lewis blood group of the individual mother (Thurl et al. 1997). Another study analyzed the level of major neutral oligosaccharides for three consecutive days in human milk colostrums (Asakuma et al. 2007). Concentrations of 2'FL and lactodifucotetraose on day 1 were found to be substantially higher than those on day 2 and 3, whereas LNT concentration increased from day 1 to day 3.

Using HPLC-Chip/time-of-flight (TOF)-MS technology, Ninonuevo et al. (2008) reported significant variations in the oligosaccharide contents primarily with the minor HMO components, whereas there was a tendency to produce a single component in very large quantities among lactating mothers. In that study, the most abundant components were identified to be lacto-*N*-neotetraose (LNnT), LNT, and LNFP. These authors noted that LNnT,

LNT, and LNFP were also preferentially consumed by *Bifidobacterium longum* subsp. *infantis* (*B. infantis*), illustrating a unique correspondence between these most abundant oligosaccharides and the bacteria they enrich. Interestingly, adult-type bifidobacteria *Bifidobacterium adolescentis* and *Bifidobacterium animalis* do not degrade LNT (Xiao et al. 2010). It is generally accepted that the mother's diet, physiology, and feeding behavior may have an impact on the daily HMO production.

HMOs AS A DEFENSE MECHANISM

HMOs play a critical role in the infant's defense system, the development of a specific intestinal microbiota, and the inflammatory processes (Zopf & Roth 1996). Numerous local and systemic effects of HMOs have been described previously, including protective functions of HMOs against enteropathogens (Newburg et al. 2004a). Also, antipathogenic effects of fucosylated oligosaccharides, specifically those that contain the Fuc α 1–2 structural motif were elucidated (Newburg et al. 2004a, Newburg et al. 2004b, Ruiz-Palacios et al. 2003). For example, Ruiz-Palacios and coworkers have reported that infants fed with human milk having low concentrations of 2'FL may be more susceptible to diarrhea than babies fed the breast milk containing high concentrations of 2'FL (Ruiz-Palacios et al. 2003).

Prevention of Pathogen Adhesion

Other researchers reported that HMOs serve as soluble ligand analogs and block pathogen adhesion (Newburg et al. 2005). The chemical structures of HMO are homologous to the carbohydrate units of glycoconjugates, especially of glycolipids, on cell surfaces of mammalian epithelial cells. For example, binding of *Escherichia coli*, *Streptococcus pneumonia*, *Campylobacter jejuni*, *Helicobacter pylori*, and *Vibrio cholerae* was inhibited by the glycoconjugates present in HMOs (e.g., 2'fucosyllactosamine) (Bode 2009, Leach et al. 2005, Morrow et al. 2004, Newburg et al. 2005, Ruiz-Palacios et al. 2003). It is also possible that HMOs can have glycome-modifying effects through changing of the expression of intestinal epithelial cell surface glycans. Angeloni et al. (2005) demonstrated that Caco-2 cells change their surface glycan profile after the exposure to 3'SL, a constituent of HMOs. In that study, the expression of α 2–3- and α 2–6-linked sialic acid residues in Caco-2 cells was significantly downregulated. Thus, this particular HMO appears to modify the glycan content of the epithelial cell surface and the receptor sites for some pathogens. The same researchers further confirmed that the adhesion of enteropathogenic *E. coli* (EPEC) was reduced upon treatment with 3'SL. **Table 1** lists the HMOs that inhibit specific pathogens in vitro, ex vivo, or in vivo.

Role of Oligosaccharides in the Development of the Immune System

Previous research has demonstrated that HMOs directly affect the immune system (Eiwegger et al. 2004, Newburg 2009, Velupillai & Harn 1994). For example, HMOs have been shown to interact with selectins (Schumacher et al. 2006), integrins (Bode et al. 2004a), and toll-like receptors (Vos et al. 2007), as well as to affect leukocyte-endothelial cell and leukocyte-platelet interactions (Bode et al. 2004a, Bode et al. 2004b, Lasky 1995, McEver 1994, Schwertmann et al. 1996). Many sialylated and fucosylated HMOs may block the latter interactions by having significant effects on the progression of inflammatory responses (Kunz et al. 1999). A recent study has also shown that HMOs can inhibit transfer of HIV-1 virus to CD4+ lymphocytes (Hong et al. 2009). Furthermore, HMOs induce intracellular processes, including differentiation and apoptosis of intestinal epithelial cells. (Kuntz et al. 2009, Kuntz et al. 2008). Neutral HMO

Table 1 Pathogen inhibition by select HMOs in in vivo, ex vivo, and in vitro studies

Pathogen	HMO tested	Reference
Norwalk virus <i>Campylobacter jejuni</i> <i>Vibrio cholera</i> <i>Escherichia coli</i> (heat-stable enterotoxin)	Fucosylated oligosaccharides	(Ruvoën-clouet et al. 2006) (Morrow et al. 2004, Ruiz-Palacios et al. 2003) (Ruiz-Palacios et al. 2003) (Newburg et al. 1990, Newburg et al. 2004b)
<i>Streptococcus pneumoniae</i> Cholera toxin <i>E. coli</i> <i>Pseudomonas aeruginosa</i> <i>Aspergillus fumigatus</i> conidia Influenza virus Polymavirus <i>Helicobacter pylori</i>	Sialyllactose	(Leach et al. 2005) (Idota et al. 1995) (Virkola et al. 1993) (Devaraj et al. 1994) (Bouchara et al. 1997) (Gambaryan et al. 1997, Matrosovich et al. 1993) (Stehle et al. 1994) (Mysore et al. 1999)
HIV-1 <i>Streptococcus pneumoniae</i> Enteropathogenic <i>E. coli</i> (EPEC) <i>Haemophilus influenzae</i>	Oligosaccharides	(Hong et al. 2009) (Andersson et al. 1986, Idänpään-Heikkilä et al. 1997) (Cravioto et al. 1991) (Idänpään-Heikkilä et al. 1997)

structures, such as LNFP III and LNNt, affect murine IL-10 production (Velupillai and Harn 1994), suggesting that HMOs might be involved in the production of antiinflammatory mediators that suppress proinflammatory Th1 response in mice (Terrazas et al. 2001). In another study, HMOs affected Th1/Th2 skewing via production of cytokines as well as maturation and activation of human cord blood–derived T cells (Eiwegger et al. 2004). More recently, Eiwegger et al. (2010) demonstrated a novel, direct immunomodulatory effect of acidic fraction of HMO when compared with the same fraction from cow's milk. In this study, acidic HMOs stimulated production of IFN- γ and IL-10, directing the neonatal Th2-type T-cell phenotype toward a Th-0-type profile in cord blood–derived mononuclear cells. This effect also impacted Th-2-type immune response of allergen-specific T cells from peanut allergic individuals. Both results strongly suggest antiallergic properties of certain acidic HMOs.

HMOs AS GROWTH FACTORS FOR BIFIDOBACTERIA

A bifidobacterial presence in the feces of breastfed infants was described by Moro in 1900, who reported that human milk contains a growth factor for these bacteria (Moro 1900). Fifty years later, György identified the bifidus factor to be GlcNAc (previously named gynolactose) (György 1953, Hoover et al. 1953) using growth of *Bifidobacterium bifidum* subsp. *Pennsylvanicum*. In vitro studies have demonstrated that GlcNAc-containing oligosaccharides are indeed able to enhance the growth of this bifidobacteria, while other sugars showed less growth-promoting activity (Petschow & Talbott 1991). The bifidogenic effect in infants is often associated with a reduction of stool pH and changes in SCFA pattern (Rinne et al. 2005). The ability of selected bifidobacteria to consume prebiotic oligosaccharides from human milk is likely an essential trait enabling this genera to be one of the most abundant colonizers of the breastfed infant gut (LoCascio et al. 2009, LoCascio et al. 2007).

Ward et al. (2006) first demonstrated vigorous growth of *B. infantis* on HMOs as a sole carbon, whereas *Lactobacillus gasseri*, a common inhabitant of the adult intestine, grew poorly. Further analysis revealed bifidobacterial species-specific differences in HMO growth, with *B. infantis* reaching a cell density threefold higher than *B. longum*, *Bifidobacterium breve*, *B. bifidum*, and *B. adolescentis*

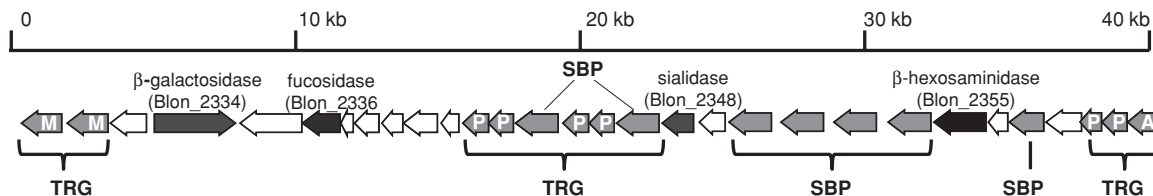


Figure 1

43 Kb gene cluster in *B. infantis* ATCC15697 containing glycosyl hydrolases and transport-related genes (TRG) required for importing and metabolizing HMOs. SBP: solute binding protein, M: major facilitator superfamily, P: ABC transporter permease component, and A: ABC transporter ATPase subunit. Adapted from Sela et al. (2008).

(Ward et al. 2007). Further work by LoCascio and colleagues (LoCascio et al. 2009, LoCascio et al. 2007) demonstrated preferential consumption by *B. infantis* of the smaller HMO species (degree of polymerization <7). These small HMO species represent the bulk of the HMOs present in pooled samples and are consistently presented over lactation (Ninonuevo et al. 2008). Tellingly, only *B. infantis* and *B. breve* could grow on the individual monosaccharide constituents of HMOs (Glc, Gal, GlcNAc, Fuc, NeuAc), suggesting another mechanism for these species to garner energy from these substrates within the intestine (Ward et al. 2007). Although growth of *B. bifidum* on HMO was less vigorous than *B. infantis*, direct consumption of HMO was observed (Ward et al. 2007). Interestingly, Fuc, GlcNAc, and NeuAc were not consumed by *B. bifidum* and remained in the media suggesting that this species is capable of deconstructing HMOs outside the cell to gain access to Glc and Gal constituents as growth substrates (Ward et al. 2007).

The recent genome sequence of *B. infantis* has enabled a more comprehensive analysis of the HMO growth phenotype by this species. Notably, Sela et al. (2008) described a 43 Kb gene HMO cluster containing the four glycosyl hydrolase activities needed to cleave HMO into its constituent monosaccharides (sialidase, fucosidase, galactosidase, and hexosaminidase) as well as an array of oligosaccharide transport-related genes (**Figure 1**). Proteomic analysis revealed that genes in this locus are induced upon growth on HMOs, suggesting that *B. infantis* imports the HMO, whereby the internalized oligosaccharides are catabolized by glycosidases prior to entry of the monosaccharides into the fructose-6-phosphate phosphoketolase central metabolic pathway (Sela et al. 2008). The transport proteins within the main HMO cluster contained six Family 1 extracellular solute binding proteins (SBP) predicted to bind oligosaccharides and to be a part of ABC transporters facilitating import and metabolism. A phylogenetic analysis of these six SBPs indicated a specific evolutionary divergence from other bifidobacterial SBP Family 1 proteins (Sela et al. 2008), suggesting a unique relationship to HMO metabolism. **Table 2** lists several bifidobacterial strains and corresponding genes related to HMO metabolism.

LoCascio et al. (2009) confirmed that the ability to consume HMOs is conserved in the entire *B. infantis* lineage, whereas other bifidobacteria isolated from infants reveal more strain-specific phenotypic variation. Recently, the same authors used comparative genomic hybridization to demonstrate a unique conservation of the HMO locus across the *B. infantis* subspecies. This work also revealed a mutant *B. infantis* strain (JCM1260) for which specific transporter genes within the main HMO cluster are absent (LoCascio et al. 2010). This mutant strain did not grow vigorously on HMOs unlike the other wild-type *B. infantis* strains (LoCascio et al. 2009), providing the first genetic evidence specifically linking the main HMO locus to the HMO growth phenotype. For a more detailed discussion of the phylogenomic aspects of HMO consumption in bifidobacteria, readers are referred to a recent review by Sela & Mills (2010).

Table 2 Presence of α -fucosidase, α -sialidase, and lacto-N-biose phosphorylase genes in sequenced bifidobacterial genomes

Sequenced bifidobacterial genomes	Fucosidases	Sialidases	Lacto-N-biose phosphorylases
<i>Bifidobacterium adolescentis</i> ATCC 15703; L2–32	–	–	–
<i>Bifidobacterium animalis lactis</i> AD011; HN019	–	–	–
<i>Bifidobacterium animalis</i> subsp. <i>lactis</i> BI-04; DSM 10140	–	–	–
<i>Bifidobacterium bifidum</i> NCIMB 41171	1	2	4
<i>Bifidobacterium breve</i> DSM 20213	–	1	2
<i>Bifidobacterium catenulatum</i> DSM 16992	–	–	–
<i>Bifidobacterium dentium</i> ATCC 27678	1	–	–
<i>Bifidobacterium dentium</i> Bd1	–	–	–
<i>Bifidobacterium gallicum</i> DSM 20093	–	–	–
<i>Bifidobacterium longum</i> DJO10A	–	–	2
<i>Bifidobacterium longum</i> NCC2705	–	–	1
<i>Bifidobacterium longum</i> subsp. <i>infantis</i> ATCC 15697	4	2	1
<i>Bifidobacterium longum</i> subsp. <i>infantis</i> ATCC 55813 ^a	–	–	1
<i>Bifidobacterium longum</i> subsp. <i>infantis</i> CCUG 52486 ^a	–	–	1
<i>Bifidobacterium pseudocatenulatum</i> DSM 20438	–	–	–

^aLoCascio et al. (2010) indicated these strains to be *B. longum* subsp. *longum*.

Nishimoto & Kitaoka (2007a) identified the novel degradation pathway in bifidobacteria specific for LNB, an HMO building block. Wada et al. (2008) further described the pathway in bifidobacteria involving both LNB and galacto-*N*-biose (GNB; Gal β 1–3GalNAc), a core structure of the mucin sugar that is present in the human intestine and milk (Lloyd et al. 1996). The latter pathway involves proteins and enzymes that are required for the uptake and degradation of disaccharides such as the GNB/LNB transporter (Suzuki et al. 2008, Wada et al. 2007), galacto-*N*-biose/lacto-*N*-biose I phosphorylase (GLNBP; LnpA) (Kitaoka et al. 2005, Nishimoto & Kitaoka 2007b), UDP-glucose-hexose 1-phosphate uridylyltransferase (GalT), and UDP-galactose

Table 3 Specific monosaccharide linkages in HMO, BMO, and commercial oligosaccharides. Adapted from Kuntz et al. (2009)

Glycans	Monosaccharide linkages
HMO	
Lacto-N-tetraose/hexaose/octaose/decaose	β 1–3, β 1–4, β 1–6
Fucosyllactose	α 1–2, α 1–3, α 1–4, β 1–3, β 1–4
Sialyllactose	α 2–3, α 2–6, β 1–4
Sialyl-lacto-N-tetraose	α 2–3, α 2–6, β 1–3, β 1–4
BMO	
Acetyllactosamine	β 1–4
Galactosyllactose	β 1–3, β 1–4, β 1–6
Acetylneuraminyllactose (Neu5Ac)	α 2–3, α 2–6, α 2–8, β 1–4
Glycolylneuraminyllactose (Neu5Gc)	α 2–3, α 2–6, β 1–4
Galactooligosaccharides (GOS)	α 1–2, α 1–4, α 1–6
Fructooligosaccharides (FOS)	β 1–2

epimerase (GalE). Other researchers confirmed that several bifidobacteria strains, including *B. longum* subsp. *longum*, *B. infantis*, *B. breve*, and *B. bifidum* were able to grow on LNB (Groschwitz et al. 2009), whereas none of the strains for *B. adolescentis*, *B. catenulatum*, *B. dentium*, *B. angulatum*, *B. animalis* subsp. *lactis*, and *B. thermophilum* showed any growth. The presence of the LnpA gene coincided with the LNB utilization in that study. Furthermore, previous studies have shown that some bifidobacterial strains have a unique pathway for the degradation of HMO, specifically with a type 1 chain (β -linked LNB) involving lacto-*N*-biosidase (LnbB). It has been suggested that the presence of the LnbB and GNB/LNB pathways in some bifidobacterial strains could provide a nutritional advantage for these organisms, thereby increasing their populations within the ecosystem of the breastfed newborns (Wada et al. 2008). Among mammalian milk oligosaccharides, those of *Homo sapiens* are especially rich in the type 1 LNB structure (Asakuma et al. 2007) and LnbB activity was found in the strains of *B. longum* and *B. bifidum* but not in *B. animalis* and *B. pseudolongum* (Wada et al. 2008). Indeed, LoCascio et al. (2007) and Ward et al. (2006) suggested that the ability to assimilate type 1 HMO is limited to certain species of bifidobacteria, e.g., *B. bifidum* and *B. infantis*.

Using 2'FL as a substrate, Katayama et al. (2008) examined bifidobacterial strains for the occurrence of fucosidase. Those researchers reported that several bifidobacteria strains, including *B. bifidum* JCM1254 and *B. longum* JCM1217, produce 1,2- α -L-fucosidase; this enzyme cleaves α -L-fucosyl residue bound to Gal through the α -1-2 linkages found at the nonreducing termini of HMO (Podolsky 1985, Song et al. 2002). Beside 2'-FL, this enzyme readily hydrolyzed lacto-*N*-fucopentaose I; however, it showed a very limited activity for α -(1-3)-linked L-fucosyl residues of 3-fucosyllactose and lacto-*N*-fucopentaose V, and had no action on the α -1-4 linkage and α -1-6 linkage.

The various catabolic strategies for HMO observed among different infant-borne bifidobacteria suggest different evolutionary adaptations to the gain a growth advantage from the same complex substrate (Sela & Mills 2010). *B. infantis* appears to internalize small HMO species (LoCascio et al. 2009, LoCascio et al. 2007), whereas *B. bifidum* exports enzymes to selectively remove LNB from the HMO structure and processes LNB intracellularly (Katayama et al. 2004). *B. breve* and *B. longum* subsp. *longum* are able to consume free LNT from the HMO pool (LoCascio et al. 2007, Ward et al. 2007). However, *B. breve* is also able to grow on the monosaccharide constituents of HMO (Ward et al. 2007). Given that these species are often isolated from the same infant feces, it is tempting to speculate that the various mechanisms may be linked to niche partitioning among bifidobacteria within the developing infant gastrointestinal tract.

ALTERNATIVE SOURCES FOR HMO-LIKE PREBIOTICS IN INFANT FORMULA: ANIMAL MILKS

Breastfed infants are better protected against several types of infections than formula-fed infants (Newburg 1997). Several researchers suggested supplementing infant formula with oligosaccharides similar to those found in human milk (McVeagh & Miller 1997, Motil 2000). Given that human milk is obviously not amenable to large-scale production, there is an urgent demand for alternative, yet functionally comparable, oligosaccharide sources from which to obtain sufficient amounts to perform clinical studies and examine the potential for use in infant nutrition. It was shown previously that the bioactivity of oligosaccharides from bovine and human milk is similar (Gopal & Gill 2000), and therefore BMOs could be used in milk products as bioactive components in human nutrition.

The oligosaccharides in milk of domestic animals, including bovine, differ in structure and have less complex structures with fewer isomers compared with HMOs, whereas the similarities

include β -glycosidic linkage of Gal and *N*-acetylhexosamine to lactose (Urashima et al. 2001). The linkages to Fuc are rare, whereas linkages of Gal or *N*-acetylglucosamine are dominant. The human intestine lacks enzymes able to hydrolyze all β -glycosidic linkages except the one in lactose. Thus, β -glycosidically bound Gal is the structural element that protects these molecules from digestion during passage through the small intestine (Boehm and Stahl 2007). Sialic acid is a major structural element in the BMOs (Kunz & Rudloff 1993, Saito et al. 1984); however, in contrast to HMO, *N*-acetylneuraminic acid (Neu5Ac) and *N*-glycolylneuraminic acid (Neu5Gc) are present (Bode 2006). It is important because differences in the chemical structure of sialic acid in human versus cow's milk are likely to influence bioavailability. It was previously reported that sialylated oligosaccharides also have a role in the initial stage of inflammation and may be effective against the influenza virus and ulcers caused by *Helicobacter pylori* (Parente et al. 2003). Unique differences between BMOs and HMOs were reported previously in terms of the size, type, and relative amounts (Tao et al. 2008). Bovine colostrums contain sialyl *N*-acetylglucosamine (Gal(β 1-4)GlcNAc) as well as 3'SL, whereas human milk contains only 3'SL (Martin-Sosa et al. 2003). Mature bovine milk also contains galactosylactoses at concentrations of 40 to 60 mg L⁻¹ (Davis et al. 1983). In a study by Tao et al. (2009), sialylated BMOs made up to 70% of colostrum and 50% of mature milk with the majority of the sialic acid Neu5Ac. Gopal & Gill (2000) reported 10 sialylated and eight neutral oligosaccharides in bovine milk and colostrums. Sialylacto-*N*-tetraose c (LSTc), 6'SL, and disialylacto-*N*-tetraose were the most representative constituents among sialyl-oligosaccharides (Coppa et al. 1999). Martin-Sosa et al. (2003) have shown that 3'SL was the most representative species in the bovine colostrums, although 6'SL remained at constantly high values during the lactation. The dominant neuramin lactose from human milk has the Neu5Ac acid linked to Gal via an α 2-6 bond, whereas the dominant bovine neuramin lactose is linked α 2-3.

It is possible that BMOs could have similar functions to HMOs in terms of pathogen deflection. The free trisaccharide, Gal α 1-3Gal β 1-4Glc, which is found in the bovine colostrum (Urashima et al. 1991), is thought to be an inhibitor of the binding of pathogenic organisms (e.g., *Clostridium difficile*) to the intestinal mucosa of newborn calves. Bovine colostrum also contains a potential prebiotic isoglobotriose (Gal α 1-3Gal β 1-4Glc), which has not been described in human milk.

Tao et al. (2008) integrated the nanoflow liquid chromatography (nanoLC) with MS in a HPLC-Chip/TOF-MS instrument to profile HMO and BMO. In that study, 40 BMOs were identified, most of which were sialylated; fucosylation was not observed in any of the samples. The same researchers demonstrated that anionic oligosaccharides are minor components in HMOs (<20%) but represent about 70% of the total oligosaccharides in bovine colostrums. Also, sialylated BMOs decreased dramatically during the first 24 hours of lactation, whereas neutral oligosaccharides increased (Nakamura et al. 2003).

Among the other milks from domesticated mammals, goat milk is especially rich in complex lactose-derived oligosaccharides. Interestingly, goat milk oligosaccharides (GMOs) have been shown to contain higher levels of oligosaccharides than bovine milks and are also reported to contain fucosylated species (Nakamura & Urashima 2004). Lara-Villoslada (Lara-Villoslada et al. 2006) recently demonstrated that GMOs reduced the inflammation and body weight loss in rats exposed to dextran sodium sulfate, a common model of colitis and inflammatory bowel disease. Other effects of GMOs included less severe colonic lesions, more favorable intestinal microbiota, and increased intestinal function. Thus, goat milk is another potential source of GMOs for human nutrition applications, including infant formulas (Martinez-Ferez et al. 2006).

Future Opportunities with Animal Milks

The milk contents of each mammalian species are precisely customized to meet the specific needs of the cognate newborns. Although recent data indicate significant differences among milk from domestic animals (Martinez-Ferez et al. 2006), milk and colostrum of domestic animals uniformly contain large amounts of sialyl oligosaccharides as well as many kinds of neutral oligosaccharides (Nakamura & Urashima 2004). Clearly, of the highest priority for infant formula is the search for the structural elements of HMO that are considered crucial to their biological effect and would serve as scientific basis for the selection of oligosaccharides from sources other than human. BMOs are particularly attractive candidates because the large size of the existing bovine dairy industry positions them as a readily available source for significant amounts of oligosaccharides with biological functions close to HMO. Recently, Barile et al. (2009) determined the composition of a variety of neutral and sialylated oligosaccharides in whey permeate using a MALDI-FTICR technique. Seven of the 15 oligosaccharides identified in that study had the same composition as some HMO structures and contained NeuAc. Those results suggest that whey permeate, a common waste stream in cheesemaking, could be a source of oligosaccharides with compositions similar to those present in human milk.

It has been previously demonstrated that sialylated oligosaccharides are important in brain development and increased immunity in infants (Boehm & Stahl 2007, Montserrat & Alicia 2001, Wang & Brand-Miller 2003). Bovine mature milk, which is used currently to produce infant formulas, has a relatively low sialic acid content (Carlson 1985, Neeser et al. 1991, Sánchez-Díaz et al. 1997, Wang et al. 2001). In humans, the sialyloligosaccharides range from 1 g L^{-1} in colostrums to $90\text{--}450 \text{ mg L}^{-1}$ in mature milk (Martin-Sosa et al. 2003, Martín-Sosa et al. 2004), whereas in bovine-based infant formulas the content of sialyloligosaccharides is as low as $15\text{--}35 \text{ mg L}^{-1}$ (Martin-Sosa et al. 2003, Wang et al. 2001). The majority of sialic acid in infant formulas is bound to protein (70%), followed by free oligosaccharides, with only 1% in the free form (Wang et al. 2001). Thus, infants fed bovine-based formulas receive significantly less sialic acid compared with breastfed infants. Several researchers attempted to concentrate and isolate the milk sialyloligosaccharides naturally present in whey (described in Barile et al. 2009). However, the exact number and type of monosaccharide residues forming sialyloligosaccharides in whey currently is not well known.

As mentioned above, both fucosylation and sialylation play an important role in the prevention of pathogens binding to the intestinal epithelia and promotion of the growth of beneficial bacteria. However, although HMO are highly fucosylated (Ninonuevo et al. 2006), the BMOs examined to date are not (Tao et al. 2009). The lack of fucosylation in BMOs is interesting given that the recent analysis of the bovine genome clearly indicates that the genetic capacity for creation of fucosylated oligosaccharides is present (Elsik et al. 2009). Certainly, more research on the molecular basis for the observed lack of fucosylation in bovine milk is warranted.

CURRENT COMMERCIAL OLIGOSACCHARIDES USED IN INFANT FORMULA

Among the rather large array of currently available and emerging prebiotics (Crittenden & Playne 2009), relatively few have been examined for use in infant formulas. Stemming from the common observation of bifidobacteria in the feces of breastfed infants, attempts have been made to reproduce this bifidogenic aspect in formulas by adding commercial prebiotics, in particular FOS and GOS, which are known to be broadly bifidogenic (Crittenden & Playne 2009).

Although HMOs are complex glycans composed of five different monosaccharides, FOS and GOS are much simpler structures. FOS are linear fructose polymers, whereas the basic structure of GOS incorporates lactose at the reducing end that is typically elongated with up to six Gal residues, which can contain different branching ([Gal(β 1–3/4/6)]1–6Gal(β 1–4)Glc). FOS can be commercially produced through the reverse reaction of fructanases and sucrases or via enzymatic hydrolysis of inulin (Espinosa et al. 2007). FOS produced by the first method lacks a reducing end and contains one Glc residue and two or more fructose moieties [short chain (sc) FOS; degree of polymerization (DP) 2–6] (Fanaro et al. 2005b), whereas hydrolysis of inulin produces free anomeric carbons and contains one fructose [long chain (lc) FOS; DP 7–60] (Roberfroid 2005). Commercial GOS preparations are mostly produced by enzymatic treatment of lactose with β -galactosidases from different sources, such as fungi, yeast, or bacteria, which results in a mixture of oligomers with various chain lengths (Park & Oh 2010).

Previous work suggested that the upper limit for the DP for GOS is eight (Macfarlane et al. 2008). However, recent work by Barboza et al. (2009) clearly demonstrated that there were oligosaccharides with a DP of up to fifteen. The same researchers reported in vitro growth behavior of different bifidobacterial strains of disaccharide- and monosaccharide-free fractions of GOS (pGOS). MALDI-FTICR MS analysis demonstrated that although all the strains tested were able to grow on the pGOS substrate, there were strain- and DP-specific bifidobacterial preferences for pGOS utilization. In general, the infant borne-isolates (*B. infantis* and *B. breve*) were able to consume the GOS species with DP ranging from three to eight more efficiently, while *B. adolescentis* and *B. longum* subsp. *longum* exhibited more differential consumption of select DP. Previously, GOS consumption with specific DP preferences had been determined only for *B. adolescentis* DSM 20083 (Van Laere et al. 2000). The selective consumption of certain GOS structures by different bifidobacterial species hints at the intriguing possibility of targeting GOS prebiotics to enrich select bifidobacterial species.

Falony et al. (2009) investigated FOS and inulin degradation by a wide range of *Bifidobacterium* species, focusing in particular on the presence of a preferential FOS breakdown mechanism. That study revealed the existence of a limited number of phenotypically distinct clusters among the tested bifidobacterial strains, however none of the species was able to degrade inulin or FOS completely. Noteworthy, common infant isolates *B. bifidum* and *B. breve* did not degrade inulin and FOS.

Perhaps the most studied prebiotic additive to infant formula is a GOS and FOS mixture, added at a 9:1 ratio (GOS:FOS) (Fanaro et al. 2005a). This particular ratio of prebiotics has been shown to increase bifidobacteria in infant feces (Boehm et al. 2002, Haarman & Knol 2005, Knol et al. 2005) and lower the incidence of pathogens (Knol et al. 2005). Other studies showed positive outcomes in terms of stool consistency and intestinal transit time with GOS/FOS (Mihatsch et al. 2006). Kapiki and colleagues showed that formula supplemented with FOS resulted in increased bifidobacteria and reduction in *E. coli* and enterococci (Kapiki et al. 2007). A recent study by Nakamura et al. (2009) demonstrated that fecal samples from infants fed formula supplemented with polydextrose, GOS, and lactulose (8 g L⁻¹) contained significantly less bifidobacteria (20.7%) than fecal samples from infants fed breast milk (83.5%). Interestingly, the same study also confirmed that the prebiotic blend may have a greater impact on infant fecal bacterial populations in younger than in older infants.

Future Opportunities with Commercial Prebiotics

Although the studies employing commercial prebiotic additions to infant formula look promising, at least in terms of enriching bifidobacteria, several questions remain. Growth on HMOs is

restricted to select bifidobacteria, primarily *B. infantis* and *B. bifidum*, species that possess the requisite genetic capacity (in particular fucosidase and sialidase functions) to deconstruct the HMO polymer. In contrast, FOS and GOS are more broadly utilized across the genus and thus may more nonspecifically enrich from this clade. Assuming HMOs evolved in concert with both the human host and the cognate infant-borne bifidobacteria (as postulated in Sela & Mills 2010), is a nonspecific enrichment of any bifidobacterial species, perhaps as the result of a GOS or FOS treatment, inherently of value to an infant? In other words, is any bifidobacteria resulting from a bifidogenic prebiotic a good outcome? As described above, the functions of HMOs are multifold, and it is unlikely that FOS and GOS possess similar developmental, immunological, or antiadherence functions. A recent study by Shoaf et al. (2006) demonstrated the ability of GOS to reduce enteropathogenic *E. coli* adherence to tissue culture cells. However, these authors noted the high level of GOS required to witness a significant reduction in *E. coli* adherence, and they speculated that this antiadherence activity might be enhanced by fucosylation or sialylation of the GOS.

Thus, one clear opportunity is to decorate existing prebiotics to obtain more HMO-like structures and functions. The technology for chemoenzymatic construction of complex carbohydrates has advanced tremendously enabling both decoration of existing structures and wholesale construction of HMO-like structures (Muthana et al. 2009). Following such a path, the generation of specifically designed, individual HMO structures would greatly enhance our ability to link biological function to specific glycan structural motifs. Perhaps more importantly, this would also set the conceptual stage for the creation of tailored synbiotic partners—very specifically designed and constructed HMOs paired with specific cognate bifidobacteria—to achieve a regulatable colonization of either the infant, or adult, GIT.

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Contents

Mammals, Milk, Molecules, and Micelles <i>P.F. Fox</i>	1
Dairy Products in the Food Chain: Their Impact on Health <i>Kirsty E. Kliem and D.I. Givens</i>	21
Avian Influenza: Public Health and Food Safety Concerns <i>Revis Chmielewski and David E. Swayne</i>	37
Molecular Design of Seed Storage Proteins for Enhanced Food Physicochemical Properties <i>Mary Rose G. Tandang-Silvas, Evelyn Mae Tecson-Mendoza, Bunzo Mikami, Shigeru Utsumi, and Nobuyuki Maruyama</i>	59
Minimization of <i>Salmonella</i> Contamination on Raw Poultry <i>N.A. Cox, J.A. Cason, and L.J. Richardson</i>	75
Nutrigenomics and Personalized Diets: What Will They Mean for Food? <i>J. Bruce German, Angela M. Zivkovic, David C. Dallas, and Jennifer T. Smilowitz</i>	97
Influence of Formulation and Processing on Absorption and Metabolism of Flavan-3-Ols from Tea and Cocoa <i>Andrew P. Neilson and Mario G. Ferruzzi</i>	125
Rheological Innovations for Characterizing Food Material Properties <i>H.S. Melito and C.R. Daubert</i>	153
Pomegranate as a Functional Food and Nutraceutical Source <i>Suzanne D. Johanningsmeier and G. Keith Harris</i>	181
Emerging Technologies in Food Processing <i>D. Knorr, A. Froehling, H. Jaeger, K. Reineke, O. Schlueter, and K. Schoessler</i>	203
Food Components with Anti-Obesity Effect <i>Kee-Hong Kim and Yeonbwa Park</i>	237

Rapid Detection and Limitations of Molecular Techniques <i>John J. Maurer</i>	259
Decontamination of Raw Foods Using Ozone-Based Sanitization Techniques <i>Jennifer J. Perry and Ahmed E. Yousef</i>	281
New Developments and Applications of Bacteriocins and Peptides in Foods <i>S. Mills, C. Stanton, C. Hill, and R.P. Ross</i>	299
The Influence of Milk Oligosaccharides on Microbiota of Infants: Opportunities for Formulas <i>Maciej Chichlowski, J. Bruce German, Carlito B. Lebrilla, and David A. Mills</i>	331
The Impact of Omic Technologies on the Study of Food Microbes <i>Sarah O'Flaherty and Todd R. Klaenhammer</i>	353
Synbiotics in Health and Disease <i>Sofia Kolida and Glenn R. Gibson</i>	373
Application of Sensory and Instrumental Volatile Analyses to Dairy Products <i>A.E. Croissant, D.M. Watson, and M.A. Drake</i>	395
Mucosal Vaccination and Therapy with Genetically Modified Lactic Acid Bacteria <i>Jerry Wells</i>	423
Hurdle Technology in Fruit Processing <i>Paula Luisina Gómez, Jorge Welte-Chanes, and Stella Maris Alzamora</i>	447
Use of FTIR for Rapid Authentication and Detection of Adulteration of Food <i>L.E. Rodriguez-Saona and M.E. Allendorf</i>	467

Errata

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