SENSORY EVALUATION OF ICE CREAM MADE WITH PREBIOTIC INGREDIENTS SUBSTITUTED FOR SUGAR

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SENSORY EVALUATION OF ICE CREAM MADE WITH PREBIOTIC INGREDIENTS SUBSTITUTED FOR SUGAR

By

Jennifer M. Wood

A THESIS

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SENSORY EVALUATION OF ICE CREAM MADE WITH PREBIOTIC INGREDIENTS SUBSTITUTED FOR SUGAR

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University of Nebraska 2011

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Prebiotics are non-digestible food ingredients that beneficially affect the host by selectively stimulating growth and/or activity of beneficial bacteria in the colon. The project objective was to determine consumer acceptability of ice cream with prebiotic ingredients substituted for part of the sugar and to determine sensory attributes of sweetness, smoothness, and vanilla flavor. A commercial ice cream mix was made substituting 0%, 10%, 20%, or 30% of the sugar for either Fructooligosaccharides (FOS) or inulin. Sensory analyses were conducted using 95 non-trained panelists. Overall consumer acceptability and sensory attributes were measured on a 175 mm anchored hedonic scale. When 10% and 20% inulin ice cream were compared to the control (0%), no significant differences in sweetness, smoothness, vanilla flavor or overall acceptability were found (P < 0.05). The 30% inulin ice cream was significantly less sweet than the control and 10% and 20% inulin ice cream, less smooth and less vanilla flavor than the control, and less acceptable than the control and 10% inulin ice cream (P < 0.05). For 10% and 20% FOS ice cream, no significant differences were found in sweetness, smoothness, vanilla flavor or overall acceptability compared to the control (P < 0.05).
The 30% FOS ice cream was significantly less sweet than 10% FOS ice cream but not significantly different than the control or 20% FOS ice cream (P < 0.05). The 30% FOS ice cream was significantly less smooth than the control, 10% and 20% FOS (P < .05). Vanilla flavor was not significantly different. Overall acceptability was significantly less for 30% FOS compared to the control and 10% FOS ice cream but not significantly different than 20% FOS (P < .05). These results suggest that FOS and inulin may be acceptable ingredients in ice cream when substituted for up to 20% of the sugar.
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Chapter 1. Review of Literature

Health Concerns

Western society is plagued with many degenerative diseases which may be directly related to food intake (Jenkins and others 1980, Kirtchevsky and Tepper 1995). Consumers have a desire for disease prevention. In a survey conducted by Gilbert and Sloan (1998), consumers indicated that an improved immune system was a major health concern (Gilbert and Sloan 1998, Niness 1999).

The microflora of the human gut is a major part of one’s immune system and there is a growing interest in the role diet plays in the microbial composition of the intestinal tract and the related health benefits. The human colonic flora consists of more than 400 bacterial species of which anaerobes outnumber aerobes (Bohunik 1996, Eckburg and others 2005). There are two categories of colonic bacteria: 1) those that are desirable and create unfavorable conditions for growth of pathogenic bacteria, such as *Bifidobacterium*, *Eubacterium* and *Lactobacillus*, and 2) those that are undesirable or promote an environment favorable for increased proliferation of pathogenic bacteria, such as *Clostridia, Escherichia, Salmonella and Campylobacter* (Roberfroid 2001, Gibson and Wang 1994, Ziemer and Gibson 1998). Factors that contribute to the health of the human gastrointestinal tract are diet, gastrointestinal pH and host health, as well as competitive inhibition and metabolic interactions among the various bacterial species.
Functional Foods May Improve Health

Functional foods provide a unique opportunity to contribute to well-being and health because they provide benefits beyond adequate nutritional effects and may reduce the risk of disease (Roberfroid 2002). The term, functional food, is used to describe nutrients that have an effect on physiologic processes that is separate and distinct from those associated with their role as nutrients (Koletzko and others 1998, Roberfroid 2000). Researchers in the last few decades have demonstrated the health promoting benefits of prebiotics, probiotics and symbiotic foods as functional foods. Symbiotic foods combine probiotics and prebiotics in a food product, which together may promote health benefits for the consumer or create increased functionality or desired qualities in the end food product. In addition, the selection of food systems for the delivery of prebiotics and probiotics are important factors to consider for viability and synergistic effect in functional foods.

Probiotics

A probiotic is defined as a live microbial feed supplement, which beneficially affects the host animal by improving its intestinal microbial balance (Fuller 1989). Probiotics must be able to be prepared on a large scale and must remain as a viable, live organism under storage and be able to survive the intestinal ecosystem (Fuller 1991, 1992). Probiotics have been consumed by human beings in the form of fermented foods for thousands of years and have been considered helpful for bodily ailments and longevity (Lourens-Hattingh and others 2001, Rasic 2003). Today, it is well accepted that daily intake of probiotics helps maintain balanced intestinal flora and prevents gastrointestinal disorders.
*Lactobacillus* and *Bifidobacterium* and other species of microorganisms have been used as probiotics for years, primarily in fermented dairy products (Boyle and Tang 2006). There are concerns regarding the safety of probiotics outside the genera of *Lactobacillus* and *Bifidobacterium*, since some of the other genera contain many pathogenic species, particularly *Enterococcus*. Fermenting organisms such as *Streptococcus salivarius* subsp. *thermophilus* and *Lactobacillus delbrueckii* subsp. *bulgaricus* used in yogurt production, are generally considered fermenting organisms rather than probiotics by most scientists because they are not expected to survive and grow in the host’s intestinal tract (Senok and others 2005). Probiotic microorganisms may produce many health-promoting metabolites and may suppress colonization of pathogenic organisms and metabolites that are detrimental to human health.

*Lactobacillus* and *Bifidobacteria* are probiotics isolated from dairy products or from human or animal intestinal tracts. In the U.S., probiotics are available in conventional foods such as probiotic-cultured yogurt, dietary supplements and medical foods. Probiotic foods worldwide also include cheese, juice, infant formula and cereals (Sanders 2006).

**Understanding Bifidobacteria**

*Bifidobacteria* are a natural part of the bacterial flora in the human body and have a symbiotic bacteria-host relationship with humans. They primarily exist in the large intestine and are one of the major genera of colonic microflora. First isolated in 1899 from a healthy breast-fed infant by Tissier of the Pasteur Institute in France,
Bifidobacteria are gram-positive, non-motile, often branched, anaerobic bacteria (Ishibashi and others 1997). Currently about 30 Bifidobacteria species are known. The Bifidobacterium species that inhabit the human intestinal tract are rather distinct from those that inhabit the intestines of animals (Mitsuoka 1984). The representative species of human origin include B. longum, B. breve, B. bifidum, B. adolescentis and B. pseudocatenuatum. Representatives of animal-derived species include B. pseudolongum, B. thermophilus and B. animalis.

How Colonic Bifidobacteria are Established

Before birth, the human fetus is germ-free and intestinal bacteria do not exist. When humans are born, they are populated with mostly anaerobic bacteria from their mother’s vaginal and fecal floral. From the time of birth, bacteria begin to colonize the intestinal tract forming a complex system of intestinal microflora. Colonization is initially determined by feeding modem, primarily breastfeeding versus formula feeding, but is further determined through use of antibiotics and other pharmaceutical agents throughout life. Bifidobacteria constitute over 95% of the intestinal flora in breast-fed infants (Yoshioka and others 1991). They report that the number of Bifidobacteria in bottle-fed infants is lower than that in breast-fed infants; nevertheless, even in bottle-fed infants, Bifidobacteria remain the predominant bacteria.

Breast-fed infants have an increased resistance to infections compared to formula fed infants. This may partly be attributed to the predominating Bifidobacteria in the intestinal
microflora of breast-fed infants. Bacteria, as measured in the feces of breast-fed infants are almost exclusively *Bifidobacteria* (Ishibashi and others 1997, Yoshioka and others 1991). Human breast milk naturally contains prebiotic fructooligosaccharides (FOS), which may contribute to the predominate *Bifidobacteria* in the intestines of breastfed infants and perhaps their increased resistance to infections (Rotimi and Duerden 1981, Gnoth and others 2000, Dai and Walker 2000). This is supported by research showing that supplementation of prebiotics to formula-fed infants proliferates *Bifidobacteria* as well as improves other health outcomes, such as reduced incidence of fever and gastrointestinal infections (Rao and others 2009). *Bifidobacteria* gradually decrease in number from the time of weaning as numerous different types of bacteria are introduced with a more complex diet. By adulthood, *Bifidobacteria* constitute only 25% of the colonic bacteria yet they remain an important component of the intestinal microflora. *Bifidobacteria* are regarded as a marker of the stability of the human intestinal microflora with optimal levels as high as one-third of total intestinal microflora (Boeckner and others 2001).

**The Role of Bifidobacteria in Human Health**

*Bifidobacteria* aid in digestion, boost immunity, are associated with a lower incidence of allergies (Björkstén and others 2001), and may inhibit some types of tumor growth (Guarnier and Malagelada 2003). *Bifidobacteria* may also help to lower cholesterol by converting it to a less absorbable form known as coprostanol, which may decrease cholesterol absorption from the intestinal tract (Tahri and others 1996). In addition, an
increased proportion of *Bifidobacteria* in the gut may crowd out undesirable pathogenic bacteria thus reducing the chance of diarrhea and infection (Gibson and Roberfriod 1995, Walker 2000). Pathological bacteria such as *E. coli* and *Clostridium perfringens* have been associated with diarrhea, infections, liver damage and putrefaction of the intestinal contents. Proliferation of these pathogenic bacteria may be further decreased through selective inhibition by increasing the proliferation of *Bifidobacteria* (Bernet 1994, Gibson and others 1995). Further beneficial effects from increased proliferation of *Bifidobacteria* may include increased phagocytosis of *E. coli* (Schiffrin and others 1997). In addition the health promoting effects prompted by bifidobacteria and other healthful bacteria may be due to the growth inhibition of harmful bacteria, stimulations of immune function, lowering of gas distention problems, and improved digestion/absorption of essential nutrients and synthesis of vitamins (Gibson 1995, Gibson and others 1995).

Being mainly saprophytes, probiotic bacteria depend on fermentable carbohydrates in the colon. *Bifidobacteria* metabolize non-digestible oligosaccharides (NDOs), such as inulin-type fructooligosaccharides (FOS) to produce vitamins and short-chain fatty acids (SCFAs), predominantly lactate and acetate. These short chain fatty acids are the main source of energy utilized by the epithelial cells in the colonic mucosa. Short-chain fatty acids lower intestinal pH, which has been shown to increase the absorption of calcium and magnesium (Bosscher and others 2006). Lower pH may also inhibit the growth of pathogenic bacteria such as *E. coli* and *Cl. perfringens*, which are completely inhibited at pH 5.0 and 4.5 (Wang and Gibson 1993).
The Role of Prebiotics in Human Health

Research findings suggest dietary habits influence the prevalence of *Bifidobacteria* in the colon. A prebiotic is a non-digestible food ingredient that beneficially affects the host by selectively stimulating the growth and/or the activity of one or a limited number of bacterial species in the colon, thus improving host health or colonic microflora composition (Collins and Gibson 1999, Gibson and others 1994a, Gibson and others 1995, Boeckner and others 2001). Non-digestible oligosaccharides (NDOs) such as inulin-type fructooligosaccharides (FOS) resist hydrolysis and digestion in the upper gastrointestinal tract but are hydrolyzed and fermented in the large bowel by colonic bacteria, such as *Bifidobacteria* (Kleesen and others 1997). These prebiotics may change the composition of fecal bacteria by 1) increasing beneficial bacteria, such as *Lactobacilli* and *Bifidobacteria*, which help modulate the activity of the immune system, and by 2) decreasing pathogenic organisms, such as *Clostridia* and protein-degrading bacteriodes, which may produce tumor-promoters from metabolites of proteins that escaped digestion in the upper gut (Rastall 2005, Sanders 2006).

Fructan polysaccharides are classified as fiber and are a storage polysaccharide predominately made up of chains of fructose. Various types of fructan polysaccharides are linked in a beta 2-1 configuration (Figure 1.1). These fructooligosaccharides (FOS) differ in degree of polymerization (DP), which is the number of fructose units in the chain. Inulin has the longest DP ranging from two to 60 (Gibson and others 1994a, Roberfriod 1998). Inulin is the naturally occurring storage oligomer of fructose found in
Jerusalem artichoke, asparagus, garlic, banana, rye, wheat, chicory and onion.

Commercial inulin is obtained through hot water extraction from chicory root.

Oligofructose is not found naturally in food; rather it is created from partial enzymatic hydrolysis of inulin resulting in a shorter DP ranging from two to 20. The term non-digestible oligosaccharide (NDO) is used to describe short-chain carbohydrates that are not digested in the small intestine and enter the large intestine unaltered. Inulin, fructooligosaccharides, oligofructose, oligosaccharides, lactulose and galactooligosaccharides are NDOs that are well-known prebiotics. The term “oligosaccharide” refers to a short chain of sugar molecules (“oligo” means “few” and “saccharide” means “sugar.”). Fructooligosaccharides (FOS) and inulin, which are found in many vegetables, consist of short chains of fructose molecules. Oligofructose is obtained through partial hydrolysis of inulin. Oligofructose was introduced as a synonym for fructooligosaccharides in 1989 (Coussement, 1999). Galacto-oligosaccharides (GOS), which also occur naturally, consist of short chains of galactose molecules.
Both oligofructose and inulin are composed of the same beta 2-1 linkages between fructose molecules and carry essentially the same nutritional benefits (Roberfriod 1998). The energy released from FOS is primarily due to the production of short chain fatty acids and lactate, which are metabolized to contribute 1.5 kcal/g of energy (Nines 1999). Research findings suggest that both inulin and oligofructose beneficially and selectively
feed anaerobic *Bifidobacteria* in the colon but the degree of polymerization may be a factor in gastrointestinal side effects such as flatulence, as well as bifidogenic properties.

The average American consumes between 2.6 and 2.5 grams per day of inulin and oligosaccharides, respectively, with a range of 1.3 to 3.5 g from commonly consumed foods, such as bananas, onions, and wheat (Moshfegh and others 1999). Some research findings indicate less than 10 grams per day to be ineffective in promoting desired colonic microbial changes. However, gastrointestinal disturbances, such as increased flatuance and bloating, as well as the potential for diarrhea have been reported in response to doses of ten to 20 grams, the amount often reported for desired changes in colonic microflora composition (Bouhnik and others 1997, Hunter and others 1999, Menne and others 2000, Kleesen and others 1997, Kruse and others 1999).

Therefore, further research has been conducted to determine a level that induces the desired bifidogenic stimulation with the least amount of gastrointestinal disturbances, thus increasing the chance of compliance. Gastrointestinal disturbances may reduce consumption of functional foods containing FOS. Fermentation of FOS by colonic bacteria may be influenced by chain length, with longer chains being fermented in more distal regions with perhaps less gaseous response (Perrin and others 2002). Some earlier reports have indicated eight to ten grams per day of FOS may be the optimum daily amount to reduce the side effects and still provide health benefits. Bouhnik and others (2004) reported that the most commonly used non-digestible carbohydrates such as
inulin-type FOS measurably raised fecal counts of *Bifidobacterium* at reasonable dose ranges of 2.5 to 5 grams per day within 7 days of administration.

Inulin and oligofructose are prebiotics that are also classified as dietary fibers. Fiber is a general term for substances which resist hydrolysis in the stomach and the small intestine (Trowell 1974). Inulin and oligofructose have been given GRAS (generally recognized as safe) status in the United States since 2002 (FDA 2003). Beyond their bifidogenic properties, these fibers have fecal bulking effects and have been shown to increase mineral absorption, increase B vitamin synthesis, increase short chain fatty acid (SCFA) production, increase immune function and improve blood lipids (Jenkins and others 1999).

Currently, inulin and oligofructose are used worldwide as a fat-replacer and sugar substitute in foods. They are also used to increase the fiber content of foods and as prebiotics (Sangeetha and others 2005a, 2005b). Both inulin and oligofructose are widely used in a variety of food products, both for their technological and nutritional attributes (Roberfroid 2002). Adding prebiotics to commonly consumed foods may help increase the daily consumption of prebiotic ingredients. However, in addition to the gastrointestinal side effects caused with increased consumption, the degrees of polymerization (DP) or chain lengths of FOS products contribute slightly different attributes to finished food products. Although inulin-type FOS may impart sweetness to
food products, it may also change the texture, water activity and color of the food product.

**Bifidogenic properties of inulin-type FOS**

Inulin-type FOS is the most studied prebiotic because of its bifidogenic properties (Tomamatsu 1994). Bouhnik and others (1999) reported that as the amount of short chain FOS consumed increased, fecal *Bifidobacteria* increased. Numerous researchers reported that a dose of ten to 20 grams per day of NDOs such as inulin-type FOS are required to produce beneficial changes in colonic bacteria through increased proliferation of *Bifidobacteria* (Bouhnik and others 1997, Hunter and others 1999, Menne and others 2000, Kleesen and others 1997, Kruse and others 1999).

The chain length of NDOs affects the rate and ability of *Bifidobacteria* to ferment the NDOs (Perrin and others 2002, Gibson and Wang 1994). Growth rates of *Bifidobacteria* cultured on either oligofructose or inulin were obtained and increased growth rate was shown on oligofructose versus the inulin, which has a longer chain length (Wang and Gibson 1993, Gibson and Wang 1995). Therefore, these researchers concluded that oligofructose is the preferred source of carbon and energy for *Bifidobacteria*. Roberfroid and others (1998) also demonstrated through in vitro fermentation of inulin, that all fructans are bifidogenic but those molecules with a chain DP > 10 are fermented on average half as quickly as molecules with a DP < 10.
FOS in infant formula is bifidogenic

Rao and others (2009) reviewed 11 trials that compared infants (minimum age 28 days) receiving formula milk supplemented with or without prebiotics and fed for a minimum of 2 weeks. In these trials, outcomes included stool colony counts of *Bifidobacteria*, *Lactobacilli*, and enteric pathogenic bacteria such as *E. coli*, as well as stool pH, consistency, frequency, anthropometry and symptoms of tolerance. Of these 11 trials, significant increases in *Bifidobacteria* after prebiotic supplementation were found in six trials and a trend toward increased *Bifidobacteria* counts were reported in two trials. A significant reduction in stool pH in the prebiotic-supplemented group was reported. Prebiotic supplemented infants had softer, more frequent stools, similar to those of breastfed infants. Weight gain was slightly higher in the prebiotic supplemented group compared to the controls. Tolerance of prebiotic supplementation was found in all but one of the eight trials. Intolerance was defined as excessive vomiting, diarrhea, regurgitation, and excessive irritability. In one study, infants who received prebiotic supplementation had more frequent diarrhea (18% vs. 4%), irritability (16% vs. 4%) and eczema (18% vs. 7%) compared to the controls, suggesting that more research is needed in this area prior to recommending routine prebiotic supplementation of formula-fed infants. However, a number of studies support the use of prebiotics due to positive alteration of fecal microflora.
**Fructooligosaccharide’s effect on calcium absorption**

Findings by Briel and others (1995) reveal dietary fiber plays a significant role in human health. However, questions remain whether dietary fiber impairs mineral absorption, similar to the effect of phytate on zinc and iron absorption. Inulin-type FOS are soluble fibers containing negligible amounts of phytate.

Inulin-type FOS are easily fermented by colonic bacteria and may stimulate absorption of several minerals and improve mineralization of bone. Although much of the research on the functional effects of NDOs, such as inulin-type FOS, is based on animal experiments, findings support the effect of NDOs on increased absorption of calcium, magnesium, zinc, and iron (Scholz-Ahren and others 2001, van den Heuval and others 1998).

Increased calcium and magnesium absorption may be partially explained by the increased production of short-chain fatty acids, fermentation end products, which lower the pH of fecal content. Lower fecal pH increases mineral solubility leading to improved mineral absorption (Remesy and others 1993, Bouhnik and others 1997, Wang and others 1993, Campbell and others 1997). These short-chain fatty acids may further influence mineral absorption by forming complexes with the minerals, leading to an increase in their uptake by the intestinal cells (Trinidad and others 1993, 1997). Some researchers speculate that the metabolic byproduct, butyrate, may stimulate the intestinal epithelium and increase its absorptive capacity (Topping and Clifton 2001). However, in humans, the effects of NDOs on fecal pH and the ratio of SCFAs is less conclusive perhaps due to the shorter
duration of human studies and the less pronounced stimulation of mineral absorption by SCFAs. Mineral absorption as well as fecal fermentation may also be dose dependent on prebiotics and calcium intake, as well as the duration of the study, a finding supported by Ohta and others (1995), Remesy and others (1993), Coudray and others (2005).

Favorable effects of NDOs on mineral metabolism in humans occur under conditions of increased calcium requirements such as in adolescence and post menopause (Maha Tahiri and others 2003, Abrams and others 2005). Adolescence is a time when increased calcium absorption is important to support growth and optimize bone mineral density (BMD) for later life. NDOs increase calcium and mineral absorption efficiency in people with low calcium intakes or increased calcium needs which is supported by research (Coudray and others 2005). Results reported by Bosscher and others (2006) from a one-year intervention trial on pre-pubertal girls and boys (n=100) indicated significantly increased calcium absorption in the group receiving a mixture of synergistically active oligofructose and long-chain inulin (8 g/d) after 8 weeks. The effect of increased calcium absorption lasted during the entire intervention period resulting in improved bone mineral content (BMC) and significantly increased BMD during periods of rapid growth compared to the control group.

Research conducted by Van den Heuvel and others (1999) further support the beneficial effects of oligofructose on calcium absorption in adolescents. In their study, 12 male adolescents aged 14-16 years received 15 grams oligofructose or sucrose (control
treatment) daily distributed over 3 meals per day for 9 days. Treatments were administered in a randomized, double-blind crossover design separated by a 19 day washout period. An increase in the amount of dietary calcium that was absorbed, also known as true fractional calcium absorption (%), was reported in the adolescents after consumption of oligofructose. The authors concluded that 15 grams of oligofructose per day stimulates fractional calcium absorption in male adolescents.

Genetic variances, including specific vitamin D receptor gene polymorphisms, may alter the effect of prebiotic consumption initially, however the research indicated that long term (1 year) consumption of both short and long-chain inulin-type fructans significantly increased calcium absorption and enhanced bone mineralization during pubertal growth (Abrams and others 2005).

Benefits of increased intestinal calcium absorption would be important for postmenopausal women whose intestinal calcium absorption naturally declines. However, Tahiri and others (2003) reported consumption of short-chain fructooligosaccharides (scFOSs) did not increase calcium absorption for women who were not on hormone replacement therapy. The researchers reported that the effects of calcium absorption with dietary scFOSs may be dependent on factors other than lifestage (age) and dietary calcium content intake such as impaired vitamin D status or reduced intestinal responsiveness to calcium needs, as is common with increasing age. However, results from a subgroup of women in this study who have had menopause for > 6 years (n=6)
suggested some benefit of dietary scFOSs on calcium absorption in the late postmenopausal phase. The small subject number warrants further investigation.

Van den Heuvel and others (1998) studied the effect of inulin, FOS and galactooligosaccharides (GOS) on true intestinal absorption of iron and calcium in men. Mineral absorption was measured by using double stable-isotope techniques. In this study, 12 healthy non-anemic males subjects, 20-30 years of age received four treatments with a consistent basal diet supplemented with 15 grams per day of either inulin, FOS, GOS, or not supplemented (control) for 21 days for each treatment according to a randomized, crossover design with no wash-out period noted. Iron absorption was measured during the last 7 days of treatment (days 15-21) and calcium absorption was measured on day 21 of each treatment period. They reported no significant differences among treatments. The researchers concluded that 15 grams per day inulin, FOS, or GOS did not have a negative effect on iron and calcium absorption in young healthy men. Their findings are supported by Coudray and others (1997) who indicated both soluble inulin and partly-soluble sugar beet fiber improved calcium absorption and calcium balance without significantly altering magnesium, iron or zinc absorption.

**The effect of FOS on Lipid Metabolism**

Dietary fiber plays a role in lipid metabolism and may reduce coronary artery disease (CAD) risk, depending on the type and source of fiber. Non-fermentable fiber, such as wheat-bran, reduces colonic transit time, but has little effect on lipid metabolism,
whereas fermentable fibers, such as pectins and β-glucans, reduce serum cholesterol but do not affect transit time (Truswell 1992). Because the fructooligosaccharides, inulin and oligofructose, are classified as dietary fiber, research has been conducted on their lipid-lowering effects. Fiordaliso and others (1995) demonstrated lipid-lowering effects of oligofructose in the serum of rats. Oligofructose supplementation has also been shown to prevent accumulation of cholesterol in liver tissue of rats as well protecting against an increase in free cholesterol concentration induced by high-fat diets (Kok and others 1996). Research conducted by Delzenne and others (1993) revealed a large decrease in triglycerides in test animals supplemented for 30 days at a dose of 20g/100g oligofructose. Although total cholesterol did not change in this study, the ratio of HDL to LDL cholesterol improved. This finding is supported by research from Levrat and others (1991) with rats fed inulin 10% by weight, possibly due to short chain fatty acid production present in the cecum of rats fed inulin. However in research reported by Vanhoof and Schrijver (1995), hypercholesterolemic rats fed inulin had no significant effect on plasma or liver cholesterol concentrations. Differences in normo- and hypercholesterolemic rats make conclusions difficult.

Because animal studies revealed some evidence of lipid-lowering effects of FOS, attention has been given to studies in humans. However, research has been limited and somewhat conflicting. A review conducted by Chowla and others (2010) concluded that soluble fibers such as pectin, oat bran, guar gum, and psyllium had a small but significant decreasing effect on total and LDL-cholesterol levels within a practical range of intake
Inulin appeared to have a similar effect on blood lipids when consumed by hyperlipidemic adults. Therefore, the authors concluded that preliminary evidence exists for a hypotriglyceridemic effect of FOS, but at the present stage of knowledge, it is not possible to conclude a hypocholesterolemic effect.

**Prebiotics and Probiotics in Food**

Probiotics have been consumed by human beings in the form of fermented foods for thousands of years (Cross and others 2001). Elie Metchnikoff, a Russian scientist first observed that consuming large amounts of fermented milk products containing *Lactobacilli* prolonged life and gave the first scientific explanation for the beneficial effects of lactic acid bacteria present in fermented milk (Rasic 2003). It is currently accepted that daily intake of probiotic-rich foods improves and helps maintain well-balanced intestinal flora, and prevents gastrointestinal disorders (Lavermicocca, 2006).

For probiotic bacteria to exert positive health benefits, these bacteria have to establish themselves in the gastrointestinal tract. Diet and food substrates are one of the most influential factors in regulating colonization of microorganisms in the gastrointestinal tract. Food not only acts as a buffer enhancing survival of bacteria through the stomach, food may also contain functional components or ingredients, such as prebiotics, which may alter probiotic survival and functionality in the gut. The food, or substrate, also plays a role in probiotic survival during product manufacture and storage.
The growing interest in the complex interactions between diet, gut microbiology and health has encouraged research and development of dietary strategies to increase proliferation of beneficial bacteria in the gut beyond the ability of the traditional American diet. Research findings suggest that symbiotic foods could play a significant role in modulating gut bacteria by increasing survival rates for probiotics during manufacture and storage as well as selectively feeding beneficial bacteria once established in the gut. Therefore, manufactures are interested in manufacturing symbiotic foods that provide greater viability of probiotic organisms while maintaining quality sensory properties in the finished product.

Food, particularly dairy products are considered an ideal vehicle for delivering probiotic bacteria to the human gastrointestinal tract (Granato and others 2010). Fat content, protein content and type, sugars and pH of food products are factors that could affect probiotic growth and survival in food. Therefore, product formulation can be manipulated to aid efficacy. There is potential for increasing the synergistic effect of foods when combining probiotics and prebiotics in the diet, separately or as symbiotics. Some foods may naturally contain ingredients that function as prebiotics, increasing the efficacy of probiotics, whereas other foods can be fortified with prebiotics during the manufacturing process. It is important to develop probiotic, prebiotic and symbiotic products with are part of day-to-day normal diet to maintain minimum therapeutic level easily.
Technical and nutritional properties of inulin

Inulin and oligofructose pose both nutritional and functional attributes that are useful in formulating innovative healthful products for consumers. The difference in chain length between inulin and oligofructose account for their distinctly different functional attributes. Inulin has a longer chain length and is less soluble than oligofructose. The ability of inulin to form inulin microcrystals when added to milk or water is useful in producing a smooth creamy texture in food. Inulin has been used successfully to replace fat in food products because it provides a fat-like mouth feel (Niness 1999). Inulin-type fructans also help stabilize foams (Cummings and Roberfroid 1997). In addition, inulin is used by the food industry as a water binder, emulsifier, stabilizer and texturizer. Traditionally, inulin has been used to replace fat in dressings, spreads, dairy products and frozen desserts.

Technical and Nutritional Properties of Oligofructose

Oligofructose is composed of shorter-chain oligomers as compared to inulin and has functional qualities similar to sucrose and glucose syrups. It is more soluble than sucrose and provides 50% of the sweetness of table sugar (Niness 1999). This quality contributes to the body of dairy products and to the humectancy of soft baked goods. It can provide crispiness to low-fat cookies and work as a binder in energy bars in the same way sucrose (sugar) functions. However, oligofructose has the added benefit of fewer calories than sucrose and does not affect serum glucose levels in diabetic patients nor does it increase insulin production or glucagon secretion (Beringer and Wenger 1995). Oligofructose is
often used in combination with high intensity sweeteners to replace sugar and provide a well-balanced flavor profile while reducing the aftertaste of artificial sweeteners.

Oligofructose also provides fiber enrichment in products giving manufacturers additional marketing venues. Oligofructose is successfully used in cereals, fruit preparations for yogurt, frozen desserts, cookies and nutritional dairy products (Niness 1999).

Oligofructose lowers the freezing point in frozen desserts and acts as a stabilizer similar to inulin, a quality that may be beneficial in preventing ice crystal formation during the hardening process and recrystallization during transport (Akalin and others 2002).

**Fructooligosaccharides in Dairy manufacture**

Most foods containing probiotic bacteria are found the refrigerated section of supermarkets as bacteria are easily destroyed by heat and other processing conditions. The dairy sector has a major advantage in the development of innovative new probiotic, prebiotic and symbiotic foods. Dairy products incorporated with probiotic bacteria are gaining popularity and comprise approximately 65% of the world functional food market (Agrawal 2009). *Lactobacillus* and *Bifidobacterium* are the most commonly used species of bacteria used in dairy products for probiotic effects (Sexelin and others 2005).

Ensuring a high survival rate of probiotic bacteria is important as standards requiring a minimum of \(10^6\) to \(10^7\) CFU/g of *Lactobacillus acidophilus* and/or *Bifidobacteria* in fermented dairy products have been introduced by several food organizations worldwide (Shah 2000). Therefore, research has been conducted on symbiotic dairy products to promote increased viability of probiotics.
Yogurt as a probiotic carrier

Fermented milk products, such as yogurt, are a common carrier for probiotic delivery. For fermentation of yogurt, *Streptococcus thermophilus* and *Lactobacillus delbrueckii* ssp. *Bulgarius* are used. *Bifidobacteria* are sometimes added to yogurt and other fermented dairy products as a probiotic with varying viability over a range of shelf lives. Hamann and Marth (1984) reported that *Streptococcus thermophilus* and *Lactobacillus delbrueckii* ssp. *bulgaricus* survive in yogurt until the use by date. However, Birollo and others (2000) observed that in whole set yogurt the level of *Streptococci* decreased approximately 1 log cycle at 6 °C in 60 days shelf life, while they remained viable and even reproduce in skimmed set yogurt under the same conditions. Higher concentrations of milk fat in yogurt produce an inhibitory effect for probiotic cultures, particularly *B. bifidum* BBI in yogurt (Vinderola and others 2000). Research has suggested plain yogurts retain higher levels of *L. acidophilus* over the shelf life compared to yogurts containing fruit purees other than mango or strawberry. This may be partially due to the differences in pH with the presence of fruit pulp (Kailasapathy 2008). Shah and others (1995) demonstrated *L. acidophilus* is affected by the low pH of the environment. Therefore, ingredients that lower the pH of yogurt may have an effect on the viability of probiotics. However, ascorbic acid was reported to improve viability of *L. acidophilus* in yogurts but did not have an effect on *Bifidobacteria* (Dave and others 1997). Dave and Shah (1997) reported that *Bifidobacteria*, when added to the commercial starter culture, decline rapidly after the initial inoculation of yogurt. Viability of probiotic bacteria depend on many factors: species of the probiotics, incubation time and temperature, pH of the final
product, storage temperature, and the length of time from inoculation to consumption (Akin and others 2007).

**Probiotic Yogurt has Beneficial Health Effects**

Consuming *bifidobacteria*-containing yogurt (B longum BB536) compared to a standard yogurt containing no *Bifidobacteria* created significant positive changes in the relative percentage of *Bifidobacterium* amongst other main bacteria in the intestinal flora (Yaeshima 1997). A simultaneous decrease in ammonia, which is representative of putrefactive product, was also shown as well as an increased frequency of defecation in subjects with a tendency toward constipation (Yaeshima 1997). Yogurts containing other strains of *Bifidobacteria*, *B breve M-16V* and *B. longum BB536* fed to very small infants of 1000 g or less enhanced early colonization of *Bifidobacteria* and reduced necrotizing enterocolitis and other intestinal tract infections (Akiyama and others 1994).

Gibson and Roberfroid (1995) reported that *Bifidobacteria* can survive the intestinal ecosystem, as do some of the fermenting organisms. Although the use of human-origin species as food supplement seems to be the reasonable and correct choice (Ishibashi and others 1997), *B. animalis* is often identified as the species used in various yogurts, possibly due to its acid-resistant nature. Animal derived species have not been isolated from the human intestinal tract, perhaps due to the inability to colonize the host intestinal tract. Recently, the *B. animalis* strains isolated from probiotic yogurts were reported to have genetic differences compared to *B animalis* originating from animal intestinal tracts.
and these strains have been renamed *B. lactis* (Meile and others 1997). This species is not an inhabitant of the human intestinal tract.

**FOS added to yogurt increases probiotic viability**

The addition of prebiotics as a food ingredient may increase viability of probiotics added to food products. Donker and others (2007) demonstrated that the addition of “Hi-maize” or amylase maize starch (a natural dietary fiber/resistant starch) and inulin resulted in improved growth, viability and organic acid production of *L. acidophilus* and *L. casei* in set-type yogurts. Both cultures demonstrated better retention of viability with inulin compared to Hi-maize however, proteolytic activity remained higher in both cultures with “Hi-maize” or no supplement as compared to inulin. Positive effects of adding Hi-maize, inulin and FOS on viability of *L. acidophilus*, *L. Casei*, *Lactobacillus rhamnosus* and *Bifidobacterium* spp. were also observed in yogurt reported by Capela and others (2006). These authors reported fructooligosaccharide as most effective in retaining viability of probiotics as compared to the other prebiotics.

In a study by Akalin and others (2004), the addition of FOS in yogurt increased viability of both *Bifidobacterium animalis* and *Bifidobacterium longum* from inoculation through storage. The level of *Bifidobacterium animalis* remained higher than the recommended level of one million cells throughout storage at 4°C for 28 days. *Bifidobacterium longum* did not remain viable at the same high levels as *Bifidobacterium animalis*, yet counts were at the same level after 28 days of storage with FOS as were found at 7 days of
storage without the addition of FOS demonstrating that FOS increased viability of both strains. Shin and others (2000) found that the viability of commercial *Bifidobacterium* *spp.* in skim milk improved by 55.7% after 4 weeks of refrigerated storage when fructooligosaccharides (FOS) were added.

**Prebiotic, Probiotic and Symbiotic Ice Cream**

Ice cream is a popular product and is consumed by people of all ages. In 2003, 5333 million liters of ice cream were produced in the USA, which accounts for 10% of the total milk production and 16% of processed milk. In Canada, 380 million liters of milk were used to produce desserts, of which 79% was used for the production of ice cream (Goff and Griffiths 2006). Because of the popularity and the beneficial composition of ice cream, it could be classified as a commonly consumed food that manufacturers could use for delivery of probiotics, prebiotics, and symbiotics.

Ice cream and frozen dairy desserts have the potential as probiotic carriers. Ice cream is a frozen mixture of milk, sweeteners, stabilizers, emulsifiers and flavoring agents (Marshal and others 2003). The category includes plain ice cream, reduced-fat, low-fat, nonfat, fruit and nut ice creams, puddings, variegated, mousse, sherbet, frozen yogurt and other frozen products. The fact that ice cream has a low storage temperature and less risk of temperature abuse may lead to a higher viability of probiotics at the time of consumption (Cruz and others 2009).
The dairy industry has found probiotic cultures to be a tool for the development of new functional products (Champagne and others 2005). Probiotic bacteria have been incorporated into fermented and non-fermented ice cream which may be an acceptable vehicle for delivery of the probiotic microorganism in the human diet (Akin and others 2007, Hekmat and McMahon 1992, Kailasapathy and Sultana 2003, Ravula and Shah 1998). The matrix of milk fat, milk proteins, lactose and other compounds endogenous to ice cream make it a good vehicle for probiotic cultures. Commercial production of probiotic enriched ice cream and fermented frozen desserts are gaining popularity (Kailasapathy and Sultan, 2003) despite the loss of viability of the microorganisms in the finished products possibly due to acidity, freeze injury and oxygen toxicity (Hekmat and McMahon 1992; Ravula and Shah 1998). The species of bacteria most commonly used in dairy products for probiotic effect are *Lactobacillus* and *Bifidobacterium* (Saxelin and others 2005). The viability of added probiotic bacteria depends on the dose level, temperature, type of dairy foods and presence of air (Homayouni, Ehsani, Azizi, yarmand, and Razavi, 2006a). Other parameters that may affect probiotic viability are probiotic strain, freeze time and temperature, pH, temperature abuse and time to consumption. Freezing and thawing causes various degrees of damage to cells, which include microorganism death through inhibition of their development, reduction, and/or interruption of metabolic activity (Davies and Obafemi 1985).

Prebiotics have been used to increase the viability of probiotic bacteria in dairy products and are also used as texturizing agents in low-fat foods such as ice cream (Devereux and
others 2003). Studies have reported on the functionality of inulin as a fat replacer in reduced fat ice cream (Schaller-Povolny and Smith 2001) and in yog-ice cream (El-Nagar and others 2002). In addition, Homayouni and others (2008) reported encapsulation of probiotic bacteria Lactobacillus casei (Lc-01) and Bifidobacterium lactis (Bb-12) can increase the survival rate of probiotic bacteria in symbiotic ice cream containing 1% resistant starch, without effecting sensory properties.

The pH of non-fermented ice cream is approximately 6.9, which provides conditions for survival of the probiotic bacteria (Christiansen and others 1996) although Reza and others (2010) suggest the freezing process, storage conditions, probiotic strain and whether or not microencapsulation is used all effect survival. Research conducted by Hekmat and McMahon (1992) concluded that probiotic ice cream at 5.5 pH had better overall acceptance than ice cream prepared at 6.0 pH. In their study, 88 untrained judges were asked to indicate their most and least preferred samples and evaluate flavor, texture, and overall acceptance of strawberry-flavored iced cream using a hedonic scale of 1 to 9. The authors did note that the preference of ice cream at the varying pH levels was affected by the panelist’s pattern of yogurt consumption, although not significantly. The authors further determined that the pattern of frozen yogurt consumption was not a predictor of consumer preference due to the varying acidity (pH4.5-6.7) of frozen yogurt depending on whether the yogurts are fully fermented or not.
Agitation during freezing incorporates air, thus conferring the desirable smoothness and softness to the frozen products. This also has an effect on probiotic survival, being disadvantageous to oxygen susceptible probiotic species. Inulin and oligofructose have different effects on the overrun in ice cream, which indicates increased air incorporation in the ice cream product, with inulin having a greater capacity than oligofructose. Being strictly anaerobic, *bifidobacterium spp.* are more sensitive to oxygen than *L. acidophilus* (Talwalkar and Kallaspathy 2003).

Akalin and Erisir (2008) compared the rheological characteristics of regular and low-fat ice cream and probiotic ice cream as well as the survival of probiotic starter culture. The authors found high apparent viscosity in the probiotic ice cream mix containing oligofructose or inulin which can be explained by the interactions of dietary fiber and liquid components of the ice cream mix, with the highest mean viscosity in the mix containing inulin. This finding is supported by Schmidt and others (1993) who found ice cream mixes containing carbohydrate-based fat replacers exhibit a viscous behavior because of the capability for imbibing water, which would increase the viscosity of the system. Schaller-Povolny and Smith (2001) had similar findings in ice cream replacing 100% of the corn syrup with inulin in reduced-fat ice cream. Similar results were reported by El-Nager and others (2002) and Akin (2005) for yog-ice cream and probiotic-fermented ice cream, respectively. Inulin is highly hygroscopic causing it to bind water and form a gel-like network.
Akalin and Erisir (2008) also reported a direct correlation between firmness and melting behavior. They found all probiotic ice creams were firmer than regular ice cream. The addition of oligofructose or inulin increased the firmness in probiotic ice cream ($P < 0.05$). The ice cream supplemented with inulin was significantly firmer than other products throughout the storage except the last day ($P < 0.05$). The ability of inulin to bind water molecules and form a particle gel network can improve the firmness of the products (Franck 2002).

In ice cream production, ice crystal size may increase up to 40% during the hardening process. Recrystallization happens during storage, whereby small ice crystals melt and large crystals grow simultaneously, a problem that is exacerbated with temperature fluctuations causing a course grainy texture in ice cream. Small crystals, with a slightly lower melting point, are more sensitive to temperature fluctuations than larger crystals (Marshal and others 2003). Inulin and oligofructose may act as a stabilizing agent to help control ice recrystallization, therefore, first dripping time of ice cream supplemented with these prebiotics can be improved by the recrystallization process as storage time increases. The inulin-supplemented group had the lowest change in melting properties and the longest first dripping time as well as the most increase in firmness ($P < 0.05$) most likely due to the high molecular weight and hygroscopic properties of inulin. First dripping time is the time for the ice cream at room temperature to begin dripping. In this research, first dripping and complete melting times were measured according to Guven and Karaca (2002). 25 g of tempered samples were left to melt (at room temperature, 20
C) on a 0.2 cm wire mesh screen above a beaker. First dripping and complete melting times of samples were determined as seconds. These observations are consistent with those of El-Natar and others (2002) who demonstrated that inulin supplementation reduced the melting rate and increased firmness in yog-ice cream. Akin (2005) also reported that addition of inulin retarded the melting time of probiotic-fermented ice cream. These authors found that the highest values for apparent viscosity, overrun, and firmness and the most remarkable improvement in the meltdown characteristics were obtained in the mix or ice cream containing probiotics and inulin (P < 0.05).

Melting properties were improved by oligofructose and inulin (P < 0.05) with melting times decreasing in all samples as storage time increased. The first dripping times were longer in probiotic ice creams supplemented with oligofructose and inulin in comparison to the control sample with inulin increasing the first dripping time more than oligofructose. Ice creams resulting in higher overruns have a high amount of air and tend to melt slowly as air cells act as an insulator (Marshall and others 2003). In their study, the addition of *L. Acidophilus* La-5 and *B. animalis* Bb-12 did not significantly affect overrun values, which is supported by the work of Alamprese and others (2002).

When compared to the control, the viable counts for both *L. acidophilus* La-5 and *B. animalis* Bb-12 significantly increased in the probiotic ice cream mix by addition of oligofructose (P < 0.05) due the possible prebiotic effect oligofructose. The viable counts of probiotics declined in all samples as a result of freezing, most likely due to the freeze
injury of cells leading to eventual death of the cells. This may have been exacerbated by the incorporation of air into the ice cream. The incorporation of air is essential to obtain the desired overrun in ice-cream: however, excess oxygen will affect the growth of micro-aerophilic *Lactobacillus acidophilus* and anaerobic *Bifidobacteria* (Kailasapathy and Sultana 2003). Viable counts continued to decline through storage; however the mixing and freezing had a greater effect on counts than storage (P < 0.05). A similar finding was reported by Hagen and Narvhus (1999) for different microorganisms with different production technologies and pH. However, Davidson and others (2000) did not report significant changes on culture bacteria during storage. Hekmat and McMahon (1992), and Haynes and Playne (2002) reported satisfactory survival of probiotic bacteria in frozen dairy desserts. The authors in this study reported *B. animalis* Bb-12 survived better than *L. Acidophilus* La-5 in ice cream over 90 days, yet counts of *B. animalis* Bb-12 only remained higher than the recommended minimum limit of $10^6$ CFU during storage in the ice cream supplemented with oligofructose.

Akalin and Erisir (2008) concluded that the best improvement in textural characteristics in terms of firmness, melting properties and first dripping time was obtained in probiotic ice cream with inulin during storage (P < 0.05). Survival of probiotic bacteria were significantly enhanced with oligofructose (P < 0.05) and the recommended minimum limit of $10^6$ CFU/g were maintained for *B. animalis* BB-12 in only probiotic ice cream with oligofructose during storage.
Akin and others (2006) studied the effects of inulin and sugar on physical and sensory characteristics of probiotic ice cream containing *Streptococcus salivarius* spp. *thermophilus, Lactobacillus delbrueckii* ssp. *bulgaricus, Lactobacillus acidophilus* LA-14 and *Bifidobacterium lactis* BL-01. The addition of sugar at concentrations of 15, 18, and 21% were used and they found that viability of organisms remained highest at a sugar concentration of 18%. The addition of inulin stimulated the growth of *L. acidophilus* and *B. lactis* improving the viability of both of these organisms. *Streptococcus thermophilus* was the most stable in all samples of probiotic ice cream with > 10⁷ CFU/g throughout the storage period. *Lactobacillus delbrueckii* ssp. *bulgaricus* was reduced by 1.5 log cycles. *Lactobacillus acidophulous* and *Bifidobacterium lactis* decreased to 10⁵ CFU/g in the control samples, whereas the counts were 10⁶ CFU/g in the samples supplemented with inulin. These results suggest that the addition of inulin stimulated the growth of *L. acidophilus* and *B. lactis*, which resulted in improved viability of these organisms.

Atkin and others (2006) also found that increasing sugar concentration led to products with better physical and sensory properties. The initial pH of milk (6.59-6.62) decreased to 5.8- 6.0 during probiotic ice cream making. The acidity increased as the inulin level increased (p < 0.01). The authors attributed this to inulin stimulating the metabolic activities of the starter bacteria resulting in improved development of acidity. The effect of sugar level and addition of inulin on acetaldehyde was insignificant (P < 0.05). Overrun values rose from 34.0 to 37.5% as sugar content increased from 15% to 21% (P
< 0.01). Similar results were found in frozen yogurts by Guven and Karaca (2002). These authors found that the addition of inulin had an insignificant effect on overrun values of the ice-cream samples (P < 0.05). This may be partially due to the small percentage of inulin added to the mix in this study.

First dripping times were increased as sugar content increased in the samples (P < 0.05). The complete melting time was related to sugar content (P < 0.01) with sugar having a negative impact. The addition of inulin at 1% had an insignificant effect on the first dripping time however, addition of 2% inulin to ice cream led to an increase in first dripping time. The results indicated that increased additions of inulin to ice-cream mixes increased complete melting times. This may be attributed to the stabilizing ability of inulin due to binding water. The viscosity of the samples increased as the sugar content increased (P < 0.01). The addition of inulin caused an increase in the viscosity (P < 0.05). Similar results were reported by El-Nagar and others (2002).

Ten panelists using a sensory rating scale of 1-10 for flavor and taste and 1-5 for consistency, color and appearance assessed organoleptic properties in the study by Atkin and others (2006). This external panel of non-smokers used for sensory evaluation was very familiar with dairy products and were checked on the basis of sensory acuity and consistency. The properties evaluated included six attributed for flavor and taste (a) (no criticism: 10, cooked flavor:9-7 lack of sweetness and too sweet: 9-7, lack of flavor: 9-6 yogurt/probiotic flavor: 8-6, acidic/sour: 8-6, rancid and oxidized: 6-1, and others 5-1),
(b) eight characteristic of body and texture (no criticism: 5, crumbly: 4-2, course: 4-1, weak: 4-1, gummy: 4-1, fluffy: 3-1, sandy: 2-1) and (c) four terms describing color and appearance (no criticism: 5, dull color: 4-1, non-uniform color: 4-1, unnatural color: 3-1).

Di Criscio and others (2010) reported on probiotic, prebiotic and synbiotic ice cream. For their study, three types of ice cream were produced by adding *Lactobacillus casei* DSM 20011 and *Lactobacillus rhamnosus* DSM 20021, prebiotic inulin, or a combination of both. Two different mixes were evaluated, vanilla and fruit, however vanilla only was used for the prebiotic and symbiotic ice creams. Prebiotic ice creams included inulin at 2.5%, 5%, and 10% of the ice cream mix. Synbiotic ice cream were produced with *Lb. casei* and 3% inulin, *Lb. casei* and 6% inulin, *Lb rhamnosus* and 3% inulin and *Lb rhamnosus* and 6% inulin. Microbial counts, pH, acidity and physical and functional properties were evaluated. Microbial analyses were carried out 0-1 d, and after 1, 3, 7 and 16 weeks of frozen storage. Sensory assessment was conducted using a semi-trained panel of 10 judges. Five terms were used: flavor, homogeneity, color, taste and consistence. A scale of 1 to 3 was used for flavor and homogeneity. A scale of 1 to 4 was used for consistence, and a scale of 1 to 5 for taste and color. The samples stored at -20°C after 7 days were removed from the freezer and tempered for 5 minutes at 20°C before sensory analysis. In probiotic ice creams, firmness was not significantly influenced by the presence of microorganisms. Probiotic vanilla ice cream was only slightly lower in taste intensity compared with the control vanilla ice cream. Prebiotic ice
cream with 5 and 10% inulin were significantly firmer by about 50 and 25% respectively (p < 0.05). The authors related this to changes in freezing points because of higher solute concentrations together with the gelling properties of inulin and the increased water binding, which improve viscosity and modifies the rheology of the mix (El-Nager and others 2002). The prebiotic ice cream with the best firmness characteristic was that with 2.5% inulin, for which no significant differences were found compared with the control ice cream. Overrun values were about 15% lower in ice cream with 5 and 10% inulin compared to the control and 2.5% inulin. No significant differences were found between control samples and samples with 2.5% inulin in melting rate values. Melting rate values were significantly higher in samples with 5% inulin compared to 2.5% inulin and the controls. However the values stabilized with 10% inulin. El-Nagar and others (2002) reported the addition of 5% inulin significantly increases the rate of meltdown, which decreases from 5 to 9% because of the formation of a cohesive network able to bind water, thus reducing the mobility of water molecules among other molecules of the mix.

In sensory analysis of prebiotic ice creams, the addition of 10% inulin lowered acceptability of taste and flavor, consistency and homogeneity. The physical analysis was consistent with the sensory analysis for consistency and homogeneity. The researchers reported that ice cream with 2.5% inulin was the best ice cream with values similar to the control ice cream for all tested parameters (P < 0.05). For symbiotic ice cream, the addition of microorganisms and inulin did not significantly affect ice creams for consistency, taste, intensity, or homogeneity. Poor results were observed for color, which appeared more opaque, whereas better evaluations were expressed for flavor, because of
higher sweetness intensity. Ice creams with 6% inulin were less icy compared with those with 3% inulin. These results were in accordance with Schaller-Povolny and Smith (1999) who observed a cryoprotectant effect of inulin that helps in reduction of ice crystal growth. The best symbiotic ice cream was that with *Lb. rhamnosus* and 6% inulin, followed by ice cream with *Lb. casei* and 6% inulin. The authors concluded that ice cream with high inulin doses (10%) altered sensorial and physical properties of prebiotic ice creams, however formulations with 2.5% inulin (which did not adversely affect ice cream characteristics), it is possible to intake 5 g/d, assuming intake of 80 g ice cream, would thus provide the needed amount for beneficial effects on intestinal microorganisms and that it should be possible to produce a functional ice cream (symbiotic) with inulin (minimum 3%) and potentially probiotic microorganisms.

Stiff competition exists in today’s market as savvy shoppers want functional foods that may help stimulate the immune system or prevent disease, but also demand products that taste good. Food manufacturers need solid scientific evidence about ingredients and their functional qualities for product development. Because ice cream is a good source of calcium and minerals as well as a commonly consumed food, inulin and oligofructose could be used in ice cream production to increase daily intake of prebiotics. In addition, inulin-type FOS added to ice cream could potentially increase the calcium and mineral absorption as well as improve functional qualities. Therefore, more research is needed to determine the acceptable amount of inulin-type FOS, in varying DP, which could be added to ice cream to create a functional food product with acceptable sensory properties.
Research suggests inulin, in water-based foods such as dairy products, when used as a fat replacer, gives a fat-like mouth feel and texture (Izzo and Franck 1998, Zimeri and Kokini 2003). Limited research is available regarding FOS as a replacement for sugar in ice cream production.

Non-digestible Oligosaccharides (NDO), such as inulin and fructooligosaccharides (FOS), have potential for functional food products, however there is question as to how much can be added to food products without compromising sensory attributes.
Objectives

The goal of this research project was to determine the maximum amount of inulin and oligofructose that can replace sugar in a standardized ice cream recipe while maintaining acceptable sensory qualities. The specific research objectives were:

1. To use an ice cream formulation that is used commercially as the control.
2. To replace part of the sugar in the control with either inulin (DP > 23) or oligofructose (DP 3-10) at 10, 20, and 30% of the sugar.
3. Conduct sensory evaluation of the ice cream samples with 10, 20, or 30% of the sugar replaced with varying amounts of inulin or oligofructose then to compare these samples with the control sample. Sensory evaluation measurements include sweetness, smoothness, taste and overall acceptability of the ice cream products using hedonic scales with a 175 mm anchored line.
4. Determine the effect of adding oligofructose and inulin on water activity, texture, and color in the control samples and experimental samples using physical tests.
Chapter 2. Materials and Methods

Preliminary Ice Cream Production

The University of Nebraska’s basic ice cream formulation (Table 2.1) was used to determine the feasibility of small-scale production of ice cream as the control formulation using a 1000 g sample. Preliminary research was conducted to determine what levels of inulin and fructooligosaccharide (FOS) were to be used in the product used for sensory evaluation. Four 1000 g treatment batches were made substituting 5, 15, 50 and 100% of sugar in the control formulation (Table 2.2) with inulin (Raftiline HP; DP >23; New Century, KS). Three 1000 g treatment samples were produced substituting 10, 50, and 100% of sugar in the control recipe (Table 2.3) using FOS (Raftilose P95; DP of 2-10; Orafti Active Food Ingredients, Malvern, PA). A control batch was also made. All samples were produced weighing liquid and dry ingredients in separate containers. The liquid and dry ingredients were then mixed together and the ice cream mix was quickly poured into the freezing chamber of a consumer ice cream maker (Hamilton Beach Ice Cream Maker). Frozen ice cream samples were placed in 1-quart disposable plastic containers containing 520g and two (2 oz.) containers to determine how full to fill the smaller containers to allow for expansion for the samples to be used for sensory evaluation. After freezing the samples in a consumer refrigerator, subjective taste and texture comparisons were made on these eight batches of ice cream.
Table 2.1 University of Nebraska ice cream formula

<table>
<thead>
<tr>
<th>Ingredient</th>
<th>Amount (g)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Whole Milk</td>
<td>596</td>
</tr>
<tr>
<td>Cream</td>
<td>202</td>
</tr>
<tr>
<td>Sugar</td>
<td>100</td>
</tr>
<tr>
<td>NFDM*</td>
<td>50</td>
</tr>
<tr>
<td>CSS**</td>
<td>45</td>
</tr>
<tr>
<td>Summit Stabilizer</td>
<td>7</td>
</tr>
<tr>
<td>Total</td>
<td>1000</td>
</tr>
</tbody>
</table>

*Non-fat dry milk
**Corn syrup solids
Table 2.2  Ice cream formulation for the control and the treatments made with inulin substituted for sugar in the preliminary testing.

<table>
<thead>
<tr>
<th>Ingredients</th>
<th>Control (g)</th>
<th>Ice cream with 5% sugar replaced with inulin (g)</th>
<th>Ice cream with 15% sugar replaced with inulin (g)</th>
<th>Ice cream with 50% sugar replaced with inulin (g)</th>
<th>Ice Cream with 100% sugar replaced with inulin (g)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Whole Milk</td>
<td>596</td>
<td>596</td>
<td>596</td>
<td>596</td>
<td>596</td>
</tr>
<tr>
<td>Cream</td>
<td>202</td>
<td>202</td>
<td>202</td>
<td>202</td>
<td>202</td>
</tr>
<tr>
<td>Sugar</td>
<td>100</td>
<td>95</td>
<td>85</td>
<td>50</td>
<td>0.0</td>
</tr>
<tr>
<td>NFDM</td>
<td>50</td>
<td>50</td>
<td>50</td>
<td>50</td>
<td>50</td>
</tr>
<tr>
<td>CSS</td>
<td>45</td>
<td>45</td>
<td>45</td>
<td>45</td>
<td>45</td>
</tr>
<tr>
<td>Summit Stabilizer</td>
<td>7</td>
<td>7</td>
<td>7</td>
<td>7</td>
<td>7</td>
</tr>
<tr>
<td>Inulin</td>
<td>0</td>
<td>5</td>
<td>15</td>
<td>50</td>
<td>100</td>
</tr>
<tr>
<td>FOS</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>Total</td>
<td>1000</td>
<td>1000</td>
<td>1000</td>
<td>1000</td>
<td>1000</td>
</tr>
</tbody>
</table>
Table 2.3  Ice cream formulation for the control and the treatments made with FOS substituted for sugar in the preliminary testing.

<table>
<thead>
<tr>
<th>Ingredients</th>
<th>Control (g)</th>
<th>Ice cream with 10% sugar replaced with FOS (g)</th>
<th>Ice Cream with 50% sugar replaced with FOS (g)</th>
<th>Ice cream with 100% sugar substituted with FOS (g)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Whole Milk</td>
<td>596</td>
<td>596</td>
<td>596</td>
<td>596</td>
</tr>
<tr>
<td>Cream</td>
<td>202</td>
<td>202</td>
<td>202</td>
<td>202</td>
</tr>
<tr>
<td>Sugar</td>
<td>50</td>
<td>90</td>
<td>50</td>
<td>0</td>
</tr>
<tr>
<td>NFDM</td>
<td>45</td>
<td>50</td>
<td>50</td>
<td>50</td>
</tr>
<tr>
<td>CSS</td>
<td>45</td>
<td>45</td>
<td>45</td>
<td>45</td>
</tr>
<tr>
<td>Summit Stabilizer</td>
<td>7</td>
<td>7</td>
<td>7</td>
<td>7</td>
</tr>
<tr>
<td>Inulin</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>Oligofructose</td>
<td>0</td>
<td>10</td>
<td>50</td>
<td>100</td>
</tr>
<tr>
<td>Total</td>
<td>1000</td>
<td>1000</td>
<td>1000</td>
<td>1000</td>
</tr>
</tbody>
</table>

Based on initial observations, further preliminary testing was conducted replacing 10, 20 and 30% of sugar with inulin (Table 2.4) and FOS (Table 2.5) recording freezing times (Table 2.9). The temperature of the ice cream mix, ice in the freezers, and the salt were not standardized in the preliminary testing.
Table 2.4 Ice cream formulation for control A and treatments made with inulin substituted for sugar in preliminary testing

<table>
<thead>
<tr>
<th>Ingredient</th>
<th>Control 1 (g)</th>
<th>Ice cream with 10% sugar replaced with inulin (g)</th>
<th>Ice cream with 20% sugar replaced with inulin (g)</th>
<th>Ice cream with 30% sugar replaced with inulin (g)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Whole Milk</td>
<td>595.5</td>
<td>595.5</td>
<td>595.5</td>
<td>595</td>
</tr>
<tr>
<td>Cream</td>
<td>202</td>
<td>202</td>
<td>202</td>
<td>202</td>
</tr>
<tr>
<td>Sugar</td>
<td>100</td>
<td>90</td>
<td>80</td>
<td>70</td>
</tr>
<tr>
<td>NFDM</td>
<td>50</td>
<td>50</td>
<td>50</td>
<td>50</td>
</tr>
<tr>
<td>CSS</td>
<td>45</td>
<td>45</td>
<td>45</td>
<td>45</td>
</tr>
<tr>
<td>Stabilizer</td>
<td>7</td>
<td>7</td>
<td>7</td>
<td>7</td>
</tr>
<tr>
<td>Vanilla extract</td>
<td>0.5</td>
<td>0.5</td>
<td>0.5</td>
<td>0.5</td>
</tr>
<tr>
<td>Inulin</td>
<td>0</td>
<td>10</td>
<td>20</td>
<td>30</td>
</tr>
<tr>
<td>FOS</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>Total</td>
<td>1000</td>
<td>1000</td>
<td>1000</td>
<td>1000</td>
</tr>
</tbody>
</table>
**Table 2.5** Ice cream formulation for control B and treatments made with FOS substituted for sugar in preliminary testing

<table>
<thead>
<tr>
<th>Ingredient</th>
<th>Control B (g)</th>
<th>Ice cream with 10% sugar replaced with FOS (g)</th>
<th>Ice cream with 20% sugar replaced with FOS (g)</th>
<th>Ice cream with 30% sugar replaced with FOS (g)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Whole Milk</td>
<td>595.5</td>
<td>595.5</td>
<td>595.5</td>
<td>595</td>
</tr>
<tr>
<td>Cream</td>
<td>202</td>
<td>202</td>
<td>202</td>
<td>202</td>
</tr>
<tr>
<td>Sugar</td>
<td>100</td>
<td>90</td>
<td>80</td>
<td>70</td>
</tr>
<tr>
<td>NFDM</td>
<td>50</td>
<td>50</td>
<td>50</td>
<td>50</td>
</tr>
<tr>
<td>CSS</td>
<td>45</td>
<td>45</td>
<td>45</td>
<td>45</td>
</tr>
<tr>
<td>Stabilizer</td>
<td>7</td>
<td>7</td>
<td>7</td>
<td>7</td>
</tr>
<tr>
<td>Vanilla extract</td>
<td>0.5</td>
<td>0.5</td>
<td>0.5</td>
<td>0.5</td>
</tr>
<tr>
<td>Inulin</td>
<td>0</td>
<td>10</td>
<td>20</td>
<td>30</td>
</tr>
<tr>
<td>FOS</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td><strong>Total</strong></td>
<td><strong>1000</strong></td>
<td><strong>1000</strong></td>
<td><strong>1000</strong></td>
<td><strong>1000</strong></td>
</tr>
</tbody>
</table>

**Ice Cream Production for Sensory Evaluation**

Ice cream was made by replacing 10, 20, or 30% (10, 20, or 30 g of the 100 g of the sugar in a 1000 g batch of ice cream) with inulin (**Table 2.6**) or fructooligosaccharides (FOS) (**Table 2.7**), modifying the University of Nebraska-Lincoln Dairy Store vanilla ice cream formulation. Two control batches of ice cream, without any addition of inulin or FOS were also made as controls (**Tables 2.6 and 2.7**).
Corn syrup solids (36 DE; DRI-SWEET 36) were obtained from Germantown Summit, Roquette, IA. The stabilizer was obtained from Danisco Cultor USA, Inc., New Century, KS. Inulin (Raftiline HP; DP >23) and fructooligosaccharide (FOS) (Raftilose P95; DP of 2-10) were obtained from Orafti Active Food Ingredients, Malvern, PA. The remaining ingredients were purchased from a local grocery store. Liquid ingredients (milk and cream) and dry ingredients were weighed and placed in two separate containers, then the liquid and dry ingredients were mixed together. This ice cream mix was heated on a stove top to a temperature of 71°C (160°F) to solubilize the stabilizer. In commercial ice cream production, raw milk would be used and a pasteurization process would occur at this stage. After the ice cream mix was combined, each 1000g batch was poured into the freezing chamber of a consumer ice cream maker (Hamilton Beach Ice Cream Maker). Salt and ice were measured and placed around the freezing chamber and kept constant between the samples. The initial amount of ice added to the freezer was 3.45 kg with an additional amount of 1.09 kg added during the freezing process. The initial amount of salt added to the freezer was 0.55 kg with an additional amount of 0.18 kg added when the second amount of ice was added to the freezer. Freeze times were recorded from the time the ice cream freezer started until the time the freezer stopped churning (Tables 2.10). The frozen ice cream was then placed into 2-ounce plastic portion cups (PL2, Solo Cup Co., IL). Each treatment group and the two control batches were assigned random numbers and the samples were immediately placed in a blast freezer (U.S Cooler, IL) and stored until one prior to the sensory evaluation. One day prior to sensory analysis, the containers were transferred to a consumer freezer (Westinghouse) set at -20°C (0°F).
Table 2.6 Ice cream formulation for control 1 and treatments made with inulin substituted for sugar in sensory

<table>
<thead>
<tr>
<th>Ingredient</th>
<th>Control 1 (g)</th>
<th>Ice cream with 10% sugar replaced with inulin (g)</th>
<th>Ice cream with 20% sugar replaced with inulin (g)</th>
<th>Ice cream with 30% sugar replaced with inulin (g)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Whole Milk</td>
<td>595.5</td>
<td>595.5</td>
<td>595.5</td>
<td>595</td>
</tr>
<tr>
<td>Cream</td>
<td>202</td>
<td>202</td>
<td>202</td>
<td>202</td>
</tr>
<tr>
<td>Sugar</td>
<td>100</td>
<td>90</td>
<td>80</td>
<td>70</td>
</tr>
<tr>
<td>NFDM</td>
<td>50</td>
<td>50</td>
<td>50</td>
<td>50</td>
</tr>
<tr>
<td>CSS</td>
<td>45</td>
<td>45</td>
<td>45</td>
<td>45</td>
</tr>
<tr>
<td>Stabilizer</td>
<td>7</td>
<td>7</td>
<td>7</td>
<td>7</td>
</tr>
<tr>
<td>Vanilla extract</td>
<td>0.5</td>
<td>0.5</td>
<td>0.5</td>
<td>0.5</td>
</tr>
<tr>
<td>Inulin</td>
<td>0</td>
<td>10</td>
<td>20</td>
<td>30</td>
</tr>
<tr>
<td>FOS</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td><strong>Total</strong></td>
<td><strong>1000</strong></td>
<td><strong>1000</strong></td>
<td><strong>1000</strong></td>
<td><strong>1000</strong></td>
</tr>
</tbody>
</table>
Table 2.7  Ice cream formulation for control 2 and the treatments made with FOS substituted for the sugar in sensory.

<table>
<thead>
<tr>
<th>Ingredient</th>
<th>Control 2 (g)</th>
<th>Ice cream with 10% sugar replaced with FOS (g)</th>
<th>Ice cream with 20% sugar replaced with FOS (g)</th>
<th>Ice cream with 30% sugar replaced with FOS (g)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Whole Milk</td>
<td>595.5</td>
<td>595.5</td>
<td>595.5</td>
<td>595.5</td>
</tr>
<tr>
<td>Cream</td>
<td>202</td>
<td>202</td>
<td>202</td>
<td>202</td>
</tr>
<tr>
<td>Sugar</td>
<td>100</td>
<td>90</td>
<td>80</td>
<td>70</td>
</tr>
<tr>
<td>NFDM</td>
<td>50</td>
<td>50</td>
<td>50</td>
<td>50</td>
</tr>
<tr>
<td>CSS</td>
<td>45</td>
<td>45</td>
<td>45</td>
<td>45</td>
</tr>
<tr>
<td>Summit Stabilizer</td>
<td>7</td>
<td>7</td>
<td>7</td>
<td>7</td>
</tr>
<tr>
<td>Vanilla Extract</td>
<td>0.5</td>
<td>0.5</td>
<td>0.5</td>
<td>0.5</td>
</tr>
<tr>
<td>Inulin</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>FOS</td>
<td>0</td>
<td>10</td>
<td>20</td>
<td>30</td>
</tr>
<tr>
<td>Total</td>
<td>1000</td>
<td>1000</td>
<td>1000</td>
<td>1000</td>
</tr>
</tbody>
</table>

Recruitment

Approval of the project was obtained from the University of Nebraska Institutional Review Board (Appendix A). A flier, (Appendix B), was used to recruit University of Nebraska-Lincoln faculty and students for the sensory analysis. An email, (Appendix C), was also sent out to faculty and students in the Departments of Food Science and Nutrition and Health Sciences at the University of Nebraska-Lincoln.

Physical Analysis of Ice Cream

Ice cream samples were analyzed for physical characteristic (color and water activity).
For color, colorimeter (Minolta CR-300, Konica Minolta Sensing Americas Inc., NJ) was used. The L value signifies lightness or darkness, the b value signifies yellow or blue, and the a value signifies red or green. For water activity, Pawkit water activity meter (Decagon Devices Inc., WA) was used with a 0.5 oz sample and three replications were conducted for each sample at room temperature, approximately 20°C (68°F).

**Sensory Testing**

Participants were asked to read and sign an informed consent form, *(Appendix D)* prior to the sensory evaluation. Participants were then asked to taste four ice cream samples consisting of a control sample and three samples of ice cream containing either inulin or FOS replacing sugar at 10, 20, or 30%. Ice cream samples were presented all at one and in no specific order, with random numbers written on the containers. Four different sensory forms *(Appendix E)* were presented to each participant, stapled together, with the control number on the top sheet and then in order by 10, 20 or 30% of the inulin or FOS, with only the random numbers listed. A glass of water was included on the tray. Participants were asked to taste the ice cream samples in the order of the sensory forms place on the tray. Participants were asked to rate the ice cream for the sensory attributes of sweetness, smoothness, vanilla flavor, and overall acceptability for each sample on a 175 mm anchored hedonic scale *(Appendix E)*. Participants were instructed to cleanse the pallet with water between tasting each sample. Participants could write additional comments on the bottom of the sensory instrument.
**Statistical Analysis**

Data were entered into an Excel spreadsheet and uploaded to SAS (version 9.1).

Statistical analysis was conducted using SAS (version 9.1) to determine the means and standard deviation of the physical data and sensory data. An ANOVA was conducted to determine statistical significance and Least Square Means test was conducted to determine differences between means.
Chapter 3. Results and Discussion

Preliminary Research Results and Discussion

Eight initial batches of ice cream were produced using 0, 5, 15, 50 and 100% of the sugar substituted with inulin or 10, 50 and 100% of the sugar substituted with FOS and one control for the preliminary research. Subjective taste and texture comparisons were made on the initial eight batches of ice cream produced in preliminary research. The results are provided in Table 3.1. Freeze times recorded during the second phase of the preliminary ice cream production are listed in Table 3.2.
Table 3.1 Subjective taste and texture results of preliminary ice cream samples using inulin and FOS.

<table>
<thead>
<tr>
<th>Percent Sugar Substitution</th>
<th>Comments</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>Smooth and creamy, yet very bland in flavor.</td>
</tr>
<tr>
<td>5% Inulin</td>
<td>When compared to control, this has a rougher feel to the tongue and is perceived as slightly less sweet.</td>
</tr>
<tr>
<td>15% Inulin</td>
<td>When compared to the control, this has a rougher feel to the tongue and seems very bland.</td>
</tr>
<tr>
<td>50% Inulin</td>
<td>When compared to the other batches, this batch was perceived as smoother than the ones with less inulin substituted for sugar. It left a “greasy” feeling as if to coat the roof of the mouth. It was very bland with not much flavor.</td>
</tr>
<tr>
<td>100% Inulin</td>
<td>This batch was very hard, nearly impossible to scoop out of the container. Researchers commented that it took a long time to freeze. It was so hard that researchers could not taste a sample during the time frame of the other ice cream samples.</td>
</tr>
<tr>
<td>10 % FOS</td>
<td>As compared to the sample with inulin substituted for sugar, this batch initially felt smoother to the tongue. However it was not perceived as smooth as the control samples. No comments were made that it tasted less sweet than the control.</td>
</tr>
<tr>
<td>50% FOS</td>
<td>As compared to the lower percentage of FOS substituted for sugar, this sample felt slightly rougher to the tongue, however it was not perceived as distinctly different in taste from the 10% Raffinose substitution.</td>
</tr>
<tr>
<td>100% FOS</td>
<td>This sample did not have a sweet flavor. It was not as hard as the 100% inulin substitution.</td>
</tr>
</tbody>
</table>
Table 3.2 Freeze times recorded in minutes during second phase of preliminary ice cream production

<table>
<thead>
<tr>
<th>Prebiotic</th>
<th>Control Batches</th>
<th>Ice Cream with 10% sugar replacement</th>
<th>Ice Cream with 20% sugar replacement</th>
<th>Ice Cream with 30% sugar replacement</th>
</tr>
</thead>
<tbody>
<tr>
<td>FOS</td>
<td>(A) 39</td>
<td>44</td>
<td>47</td>
<td>52</td>
</tr>
<tr>
<td>Inulin</td>
<td>(B) 46</td>
<td>40</td>
<td>47</td>
<td>50</td>
</tr>
</tbody>
</table>

In the preliminary testing, researchers subjectively noted that as the percentage of sugar replaced with inulin increased, the ice cream took longer to freeze and it became harder during freezer storage, however the amount of ice and salt, nor the temperature of the mix in preliminary testing were standardized between the batches and the freeze times were not recorded. Ice cream with 100% of the sugar substituted with inulin took longer than the other treatment samples to thaw as well, a characteristic that could be beneficial in commercial production to decrease the impact of temperature abuse during transit times.

During the second round of preliminary testing, two control samples and six treatment samples were then produced substituting either inulin or FOS for 10, 20 and 30% of the sugar while recording freeze times. Freeze times are listed in Table 3.2 and were timed from when the ice cream freezer was plugged into the electrical outlet until the time that the ice cream freezer stopped churning. However, the freeze time for control B may not accurate due to the higher temperature of the ice cream mix at the time it was placed in the ice cream maker. All other ice cream samples were produced heating the liquid mixture to 160°F. Ice and salt were not kept constant during this phase of research. Subjective taste tests revealed that the ice cream with up to 30% of sugar replaced with
FOS or inulin was perceived smooth and acceptable. Therefore, we determined to substitute either inulin or FOS for 10, 20 and 30% of the sugar for the samples for the sensory evaluation and to keep the salt and ice constant for accurate freeze times for the final samples for sensory evaluation. Vanilla was also added to the formulation to make the ice cream less bland in flavor.

**Final Research Discussion**

**Freeze Times**

Freeze times were recorded during ice cream production (Table 3.3). Freeze times were shorter with the addition of inulin or FOS as compared to the controls. This was not an objective of this research project, but freeze times were recorded because of preliminary findings. The reduction in freeze time found with the addition of inulin or FOS would be consistent with principle of the addition of a large polymer to a liquid to raise the freezing point of that liquid. This finding may be related to the subjective comments during the preliminary trials that the addition of inulin or FOS made the ice cream harder, which in turn took longer to melt and was more difficult to scoop (Table 3.1).

Freeze times decreased with the addition of both inulin and FOS substituted for sugar, which may be attributed to increased freezing temperature for ice cream. This may partially be due to the increased viscosity caused by the addition of dietary fiber to the liquid components of the ice cream mix. Ice cream mixes containing carbohydrate-based fat replacers exhibit a viscous behavior because of the capability for imbibing water,
which would increase the viscosity of the system (Schmidt and others 1993). Akalin and others (2008) found ice cream containing inulin had higher viscosity than ice cream containing oligofructose and suggested this might be due to the higher molecular weight of inulin. Inulin is also highly hygroscopic and would, therefore, bind water and form a gel-like network that, in addition to the other components in the ice cream, may affect rheology of the mix. Similar results were reported by El-Nagar and others (2002) and Akin (2005) for yog-ice cream and probiotic-fermented ice cream, respectively.

Previous studies indicated that inulin and FOS added to ice cream may increase the firmness of ice cream containing probiotic bacteria *Lactobacillus acidophilus* La-5 and *Bifidobacterium animalis* Bb-12 (Akalin and others 2002). These researchers demonstrated that probiotic ice cream containing 4% inulin and 4% oligofructose (compared to 1, 2 and 3% inulin and 1, 2 and 3% oligofructose used in this project; 10, 20 and 30 g in 1000 gram batches) increased firmness throughout storage except for the 1st day. In their research, inulin exhibited a higher firmness than oligofructose. Akalin and others (2002) also demonstrated that both inulin and oligofructose added to ice cream improved the melting properties of ice cream. The ice cream remained firm longer at room temperature, with inulin exhibiting the lowest change in melting properties.
Table 3.3 Freeze times recorded in minutes during final ice cream production

<table>
<thead>
<tr>
<th>Prebiotic</th>
<th>Control Batches</th>
<th>Ice Cream with 10% sugar replacement</th>
<th>Ice Cream with 20% sugar replacement</th>
<th>Ice Cream with 30% sugar replacement</th>
</tr>
</thead>
<tbody>
<tr>
<td>FOS</td>
<td>42</td>
<td>35</td>
<td>27</td>
<td>33</td>
</tr>
<tr>
<td>Inulin</td>
<td>37</td>
<td>30</td>
<td>30</td>
<td>30</td>
</tr>
</tbody>
</table>

Color

Color measurements for the control samples (n = 12) were compared to samples containing 10, 20, or 30% of sugar substituted with either inulin or FOS (n = 6 for each treatment) (Table 2.11). Ice cream with inulin substituted for 30% of the sugar was significantly darker compared to all treatment samples including the control. Ice cream with sugar substituted with 20% inulin and 20% FOS were significantly darker than the control, however ice cream with 30% of sugar substituted with FOS was not significantly darker than the control. It appears that both inulin and FOS darken the color of the products at 20% but this trait may not appear at higher concentrations of FOS where it continues to darken with additional inulin. There were no significant differences in color measurements between the control and treatment samples for yellow/blue (P ≤ 0.05 and the red/green (a) values.
Water Activity

No significant differences were found in water activity between the control and treatment samples (Table 2.11). Water activity is affected by temperature with increasing water activity for increasing temperature. Water activity measurements were made controlling for temperature for all the ice cream samples.
Table 3.4 Color and Water Activity of ice cream made with inulin and FOS substituted for 10, 20, and 30% of sugar. Means and standard deviations listed.

<table>
<thead>
<tr>
<th></th>
<th>Color</th>
<th>Water Activity</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>L (Lightness/darkness)</td>
<td></td>
</tr>
<tr>
<td>Control n=12</td>
<td>84.44 ± 1.49abc</td>
<td>5.91 ± 1.16</td>
</tr>
<tr>
<td>Ice cream with 10% sugar replaced with inulin (n = 6)</td>
<td>83.30 ± 1.15g</td>
<td>5.58 ± 0.68</td>
</tr>
<tr>
<td>Ice cream with 20% sugar replaced with inulin (n = 6)</td>
<td>82.73 ± 1.95bh</td>
<td>5.30 ± 1.65</td>
</tr>
<tr>
<td>Ice cream with 30% sugar replaced with inulin (n = 6)</td>
<td>80.96 ± 1.39cdegh</td>
<td>6.08 ± 0.72</td>
</tr>
<tr>
<td>Ice cream with 10% sugar replaced with FOS (n = 6)</td>
<td>83.19 ± 1.31d</td>
<td>5.62 ± 0.67</td>
</tr>
<tr>
<td>Ice cream with 20% sugar replaced with FOS (n = 6)</td>
<td>82.54 ± 0.65se</td>
<td>5.75 ± 0.94</td>
</tr>
<tr>
<td>Ice cream with 30% sugar replaced with FOS (n = 6)</td>
<td>83.53 ± 0.63f</td>
<td>5.95 ± 0.45</td>
</tr>
</tbody>
</table>

 Superscripts that are the same in each column are significantly different (P < 0.05). No Subscript indicates no significant differences.
**Sensory Analysis**

Ninety-five participants evaluated the ice cream samples made with either 10, 20 or 30% of the sugar substituted with inulin or FOS and compared these treatments with a control sample. Overall consumer acceptability and sensory attributes (sweetness, smoothness, and flavor) were measured on a 175 mm anchored hedonic scale. Results of the sensory analysis are listed in Table 3.5 and Table 3.6. No significant differences were found between the two control samples (Table 3.7). When ice cream with 10% and 20% of the sugar replaced with inulin was compared to the control (0%), no significant differences in sweetness, smoothness, vanilla flavor or overall acceptability were found (P < 0.05). The ice cream with 30% of the sugar substituted with inulin was significantly less sweet than the control and the ice cream with 10% and 20% of the sugar substituted with inulin. It was also less smooth and less vanilla flavor than the control, and less acceptable than the control and the ice cream with 10% of the sugar replaced with inulin (P < 0.05). For ice cream with 10% and 20% of the sugar substituted with FOS, no significant differences were found in sweetness, smoothness, vanilla flavor or overall acceptability compared to the control (P < 0.05). Ice cream with 30% of the sugar substituted with FOS was significantly less sweet than ice cream with 10% of the sugar substituted with FOS, but not significantly different than the control (0%) or ice cream with 20% of the sugar substituted with FOS (P < 0.05). Ice cream with 30% of the sugar substituted with FOS was significantly less smooth than the control (0%), and ice cream with 10% and 20% of the sugar substituted with FOS (P < 0.05). Vanilla flavor was not significantly different between the control (0%) and ice cream with 10, 20 or 30% of the sugar substituted with FOS. Overall acceptability was significantly less for ice cream with 30% of the sugar substituted with FOS compared to the
control (0%) and ice cream with 10% of the sugar substituted with FOS, but not significantly different than ice cream with 20% of the sugar substituted with FOS (P < 0.05).

**Table 3.5** Overall acceptability and attribute ratings for ice cream with inulin substituted for sugar at 10, 20 and 30%. Means and standard deviations listed.

<table>
<thead>
<tr>
<th>Attribute</th>
<th>Control 1</th>
<th>10% Inulin</th>
<th>20% Inulin</th>
<th>30% Inulin</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sweetness</td>
<td>9.9 ± 3.4&lt;sup&gt;a&lt;/sup&gt;</td>
<td>9.6 ±3.3&lt;sup&gt;b&lt;/sup&gt;</td>
<td>8.9 ± 3.3&lt;sup&gt;c&lt;/sup&gt;</td>
<td>7.2 ± 3.7&lt;sup&gt;abc&lt;/sup&gt;</td>
</tr>
<tr>
<td>Smoothness</td>
<td>9.8 ± 3.5&lt;sup&gt;a&lt;/sup&gt;</td>
<td>9.2 ± 3.9</td>
<td>9.3 ± 3.9</td>
<td>8.2 ± 3.9&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
<tr>
<td>Vanilla Flavor</td>
<td>10.6 ± 3.8&lt;sup&gt;a&lt;/sup&gt;</td>
<td>10.4 ± 3.7</td>
<td>10.2 ± 3.2</td>
<td>8.9 ± 3.9&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
<tr>
<td>Overall Acceptability</td>
<td>11.0 ± 3.8&lt;sup&gt;a&lt;/sup&gt;</td>
<td>10.7 ± 3.6&lt;sup&gt;b&lt;/sup&gt;</td>
<td>9.7 ± 3.6</td>
<td>8.4 ± 3.3&lt;sup&gt;ab&lt;/sup&gt;</td>
</tr>
</tbody>
</table>

<sup>a, b, c</sup> Superscripts that are the same in each row show significantly different outcomes from each other.

**Table 3.6** Overall acceptability and attribute ratings for ice cream with FOS substituted for sugar at 10, 20, and 30%. Means and standard deviations listed.

<table>
<thead>
<tr>
<th>Attribute</th>
<th>Control 2</th>
<th>10% FOS</th>
<th>20% FOS</th>
<th>30% FOS</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sweetness</td>
<td>8.63 ± 3.7</td>
<td>9.40 ± 3.2&lt;sup&gt;a&lt;/sup&gt;</td>
<td>8.74 ± 3.6</td>
<td>7.57 ± 3.6&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
<tr>
<td>Smoothness</td>
<td>10.10 ± 3.6&lt;sup&gt;a&lt;/sup&gt;</td>
<td>10.96 ± 3.3&lt;sup&gt;b&lt;/sup&gt;</td>
<td>10.58 ± 4.1&lt;sup&gt;c&lt;/sup&gt;</td>
<td>8.44 ± 3.6&lt;sup&gt;abc&lt;/sup&gt;</td>
</tr>
<tr>
<td>Vanilla Flavor</td>
<td>9.71 ± 4.0</td>
<td>9.73 ± 3.4</td>
<td>8.69 ± 4.2</td>
<td>8.50 ± 4.2</td>
</tr>
<tr>
<td>Overall Acceptability</td>
<td>10.76 ± 4.1&lt;sup&gt;a&lt;/sup&gt;</td>
<td>11.06 ± 3.2&lt;sup&gt;b&lt;/sup&gt;</td>
<td>10.55 ± 3.7</td>
<td>9.21 ± 3.9&lt;sup&gt;ab&lt;/sup&gt;</td>
</tr>
</tbody>
</table>

<sup>a, b, c</sup> Subscripts that are the same in each row show significantly different outcomes from each other P < 0.05. No subscript indicates no significant difference.
Table 3.7 Comparison of the 2 control samples. Means and standard deviations listed.

<table>
<thead>
<tr>
<th></th>
<th>Control 1</th>
<th>Control 2</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>(n = 44)</td>
<td>(n = 51)</td>
<td></td>
</tr>
<tr>
<td>Sweetness</td>
<td>9.87 ± 3.4</td>
<td>8.63 ± 3.7</td>
<td>P = .08</td>
</tr>
<tr>
<td>Smoothness</td>
<td>9.83 ± 3.5</td>
<td>10.10 ± 3.6</td>
<td>P = .72</td>
</tr>
<tr>
<td>Vanilla Flavor</td>
<td>10.64 ± 3.8</td>
<td>9.71 ± 4.0</td>
<td>P = .24</td>
</tr>
<tr>
<td>Overall Acceptability</td>
<td>11.03 ± 3.8</td>
<td>10.76 ± 4.1</td>
<td>P = .72</td>
</tr>
</tbody>
</table>

No significant difference were indicated (P = 0.05)

During the sensory evaluation, participants had the opportunity to make comments on their sensory forms. The number of participants who made similar comments is listed in parenthesis behind the comment. One participant mentioned the 10% FOS sample was difficult to scoop, but no participants mentioned grittiness or ice crystals. In the 20% FOS sample, one participant mentioned it was smoother than the 30% FOS sample, other comments included: their favorite of the four (1), creamy and good texture (2), creamy but gritty (1), sugar appears to be crystallizing out (1). In the 30% FOS sample, comments included: not very creamy and after-taste (1), gritty or icy (3), didn’t taste right (1), not as good as control (1). Comments on the 10% inulin sample included: very smooth (1), still too sweet (1), funny texture (1), keeper and best flavor (1), coated mouth (1), gritty (1), a little blah (1), hard (1), needs more sweetness (1). In the 20% inulin sample, comments included: sweeter (2), after-taste (1), gritty or grainy (4), hard (1), best sample or keeper (6). Comments on the 30% inulin samples included: clumpy (9), didn’t care for or not great (3), fluffy (1), very smooth and creamy (2), the best one (1). Comments on the control for the inulin included: very good and not too sweet (1), after-taste (2), sweet (3), strange film (1),
favorite sample (3), gritty (2). Lum and Albrecht (2007) reported on sensory evaluation conducted on ice cream with 10% of sugar replaced with inulin or FOS. Seventy-one participants evaluated the ice cream and liked the ice cream made with inulin or FOS equally well (p ≤ 0.05). However, the participants liked the control sample better than ice cream with 10% of the sugar substituted with either inulin or FOS (p ≤ 0.001).

Di Criscio and others (2010) reported ice cream with 2.5% inulin did not alter physical or sensory characteristics of ice cream significantly; however ice cream with high inulin (10%) altered the physical and sensory characteristics of ice cream. Our results support ice cream with up to 2% inulin or FOS (20% of the sugar substituted with inulin or FOS) is acceptable.

**Conclusion**

Inulin and FOS are potential ingredients for use in ice cream; they may however, have an effect on physical and sensory attributes when substituted for sugar in ice cream. Future research should include tests for hardness. Considerations for ice cream producers:

- The color of ice cream may be affected (slightly more green) when more than 30% of sugar is substituted for inulin in ice cream
- Inulin and FOS may darken the color of ice cream when substituted for greater than 20% of the sugar.
- Inulin or FOS substituted for up to 20% of sugar in ice cream does not affect sweetness, smoothness, vanilla flavor or overall acceptability
These results suggest that inulin and FOS may be acceptable ingredients in ice cream when substituted for up to 20% of sugar. However, more research is needed to confirm these results.
References


Lum, AK, and Albrecht JA. 2008. Sensory evaluation of ice cream made with prebiotic ingredients. RURALS. http://digitalcommons.unl.edu/cgi/viewcontent.cgi?article=1033&context=rurals


Appendix A - IRB Approval Letter
January 30, 2008

Jennifer Wood
Dr. Julie Albrecht
324 Church Street South
Peterson, MN 55962

IRB#2007-12-8619 EP

TITLE OF PROJECT: The Investigation of Fructooligosacharides in Ice Cream Production

Dear Jennifer:

This letter is to officially notify you of the approval of your project by the Institutional Review Board (IRB) for the Protection of Human Subjects. It is the Board's opinion that you have provided adequate safeguards for the rights and welfare of the participants in this study. Your proposal seems to be in compliance with this institution's Federal Wide Assurance 00002258 and the DHHS Regulations for the Protection of Human Subjects (45 CFR 46).

Date of EP Review: 01/28/08

You are authorized to implement this study as of the Date of Final Approval: 01/28/08

This approval is Valid Until: 01/27/09

We wish to remind you that the principal investigator is responsible for keeping this Board informed of any changes involved with the procedures or methodology in this study. You should report any unanticipated problems involving risks to the participants or others to the Board. For projects which continue beyond one year from the starting date, the IRB will request continuing review and update of the research project. Your study will be due for continuing review as indicated above. The investigator must also advise the Board when this study is finished or discontinued by completing the enclosed Protocol Final Report form and returning it to the Institutional Review Board.

1. Uploaded on NUgrant is the IRB approved Informed Consent form for this project. Please use this form when making copies to distribute to your participants. If it is necessary to create a new informed consent form, please send us your original so that we may approve and stamp it before it is distributed to participants.

We wish to remind you that the principal investigator is responsible for reporting to this Board any of the following events within 48 hours of the event:

- Any serious event (including on-site and off-site adverse events, injuries, side effects, deaths, or other problems) which in the opinion of the local investigator was unanticipated, involved risk to subjects or others, and was possibly related to the research procedures;
- Any serious accidental or unintentional change to the IRB-approved protocol that involves risk or has the potential to recur;
- Any publication in the literature, safety monitoring report, interim result or other finding that indicates an unexpected change to the risk/benefit ratio of the research;
- Any breach in confidentiality or compromise in data privacy related to the subject or others; or
- Any complaint of a subject that indicates an unanticipated risk or that cannot be resolved by the research staff.

If you have any questions, please contact Shirley Horstman, IRB Administrator, at 472-9417 or email shorstmanl@unl.edu.

Sincerely,

Dan R. Hoyt, Chair
for the IRB
Appendix B – Recruitment Flier

**Attention Students and Faculty!**

You are invited to participate in a Research Study on the Addition of Fructooligosaccharides in Ice Cream

**When?** Spring Semester 2008 *(exact date yet to be decided)*

**Where?** Sensory Analysis Laboratories in the Departments of Nutrition and Health Sciences and Food Science and Technology

**Details:** You must be at least 19 years of age to participate. You will be tasting vanilla flavored ice cream samples and asked to fill out a sensory analysis questionnaire.

**What you will get for participating:** a gift certificate for the UNL Dairy Store or a candy bar
Appendix C – Recruitment Email sent to students, staff and faculty

Dear Students, Staff and Faculty,

You are invited to participate in a study on the Investigation of Fructooligosaccharides in Ice Cream.

During this study you will be asked to taste several samples of vanilla flavored ice cream that have various amounts of two types of Fructooligosaccharides added. As compensation for participating, you will receive a gift certificate for the UNL Dairy Store.

The study is being held in the sensory analysis laboratory in the Department of Nutrition and Health Sciences on (date) and in the Department of Food Science and Technology on (date).
Appendix D - Informed Consent Form
Informed Consent Form

Title of Project: The Investigation of Fructooligosaccharides in Ice Cream Production

Purpose of the Research: Fructooligosaccharides (FOS) are non-digestible carbohydrates that selectively feed the beneficial bacterial in your gut. It is also found in commonly consumed foods such as bananas, onions, garlic, wheat, asparagus, barley, wheat, and tomatoes. FOS is derived from chicory root for commercial use and is used in place of part of the sugar in the ice cream samples in this study. The optimal daily intake of FOS to stimulate growth of beneficial bacteria in the colon as well as enhance the absorption of bone enhancing minerals is 8 to 10 grams per day. Average daily consumption for Americans is currently 1 to 4 grams per day. To increase the daily consumption of FOS, we are investigating the possibility of adding FOS to ice cream, as ice cream is a commonly consumed food. We are investigating adding two types of FOS with varying degrees of polymerization (carbohydrate chain length) at various levels in place of sugar. Our goal is to determine how much FOS can be added to the ice cream and still maintaining acceptable sensory qualities. FOS and inulin are currently added to some commercial yogurt products.

Procedures: To participate in this study, you will first be asked to read and sign this informed consent form. Then you will be given several samples of vanilla flavored ice cream and asked to fill out sensory analysis forms identifying qualities such as sweetness, smoothness, flavor, and overall acceptability. You will also be given the opportunity to add any additional comments about ice cream.

Risks: The known risks associated with the consumption of FOS are increased flatulence and the urgency to defecate due to increased fecal bulk similar to consuming high fiber foods.

Benefits: The benefit of participating in this study is that you have helped the University of Nebraska-Lincoln further scientific research that benefits the health of the consumers.

Confidentiality: Information collected in this study will be kept strictly confidential. No contact information will be collected. The results of the study will be published in a graduate thesis and submitted for publication in scientific journals and presented at conferences were appropriate. Personal information will not be collected and individual initials ____
data will not be identifiable. Information collected will be stored in a locked cabinet in
104 RLH in the Department of Nutrition and Health Sciences for up to two years.

**Compensation:** When you complete this study, you will receive a gift certificate for the
UNL Dairy store.

**Opportunity to ask questions:** You may ask any questions about this study and have those
questions answered before agreeing to participate in the study or at any time during the
study.

**Initial Consent:** You are free to decide not to participate in this study or to withdraw at any
time without negatively affecting your relationship with the University of Nebraska. Your
decision will not result in any loss of benefits to which you are otherwise entitled. you are
free to call the investigators at any time if you have any questions about your rights as a
research participant that have not been answered by the investigator or to report any
concerns about the study. You may contact the University of Nebraska – Lincoln,
Institutional Review Board whose telephone number is 402-472-6965. By signing this form,
you indicate that you have read and understand the information presented and all questions
have been answered. You will be given a copy of this consent for you to keep.

**Signature of Participant**

Names and Phone Numbers of the Researchers:

Julie A. Albrecht, Ph.D
University of Nebraska-Lincoln
Office: (402) 472-8884

Jennifer M. Wood, RD, LD
University of Nebraska-Lincoln
(507) 875-2705
Appendix E – Sensory Scale Used for Research

Directions

1. Discard any food, gum, or tobacco from your mouth and rinse your mouth out with water.

2. Taste each product in the order listed. Taste the first product and place an X on the line to indicate its level of sweetness, smoothness, flavor, and overall acceptability.

Example:

| not sweet | X | very sweet |

3. You may swallow the sample or spit the sample out in the cups provided. Rinse your mouth out with water before tasting the next sample.

4. After tasting a product, wait one minute (by setting the timer) to sample the next product.

5. Repeat steps #2 and #3 until all the products are sampled and all the scales are filled out.
Appendix E

Sensory Scale

Sweetness

- not sweet
- very sweet

Smoothness

- not smooth
- very smooth

Flavor

- not vanilla
- very vanilla

Overall Acceptability

- not acceptable
- very acceptable
Additional Comments:________________________________________________________________________
________________________________________________________________________
________________________________________________________________________
________________________________________________________________________