1,2-Fucosyllactose Does Not Improve Intestinal Function or Prevent Escherichia coli F18 Diarrhea in Newborn Pigs

Cilieborg, Malene Skovsted; Sangild, Per Torp; Jensen, Michael Ladegaard; Østergaard, Mette Viberg; Christensen, Lars; Rasmussen, Stine Ostenfeldt; Mørbak, Anne Louise; Jrgensen, Claus Bttcher; Bering, Stine Brandt

Published in:
Journal of Pediatric Gastroenterology and Nutrition

Link to article, DOI:
10.1097/MPG.0000000000001276

Publication date:
2017

Document Version
Publisher's PDF, also known as Version of record

Link back to DTU Orbit

Citation (APA):
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ABSTRACT

Objectives: Infectious diarrhea, a leading cause of morbidity and deaths, is less prevalent in breastfed infants compared with infants fed infant formula. The dominant human milk oligosaccharide (HMO), α1,2-fucosyllactose (2′-FL), has structural homology to bacterial adhesion sites in the intestine and may in part explain the protective effects of human milk. We hypothesized that 2′-FL prevents diarrhea via competitive inhibition of pathogen adhesion in a pig model for sensitive newborn infants.

Methods: Intestinal cell studies were coupled with studies on cesarean-delivered newborn pigs (n = 24) without (control) or with inoculation of enterotoxigenic Escherichia coli F18 (7.5 × 10^9/day for 8 days) fed either no (F18) or 10 g/L 2′-FL (2FL-F18).

Results: In vitro studies revealed decreased pathogen adhesion to intestinal epithelial cells with 2′-FL (5 g/L; P < 0.001). F18 pigs showed more diarrhea than control pigs (P < 0.01). Administration of 2′-FL to F18 pigs failed to prevent diarrhea, although the relative weight loss tended to be reduced (~19 vs –124 g/kg, P = 0.12), higher villi were observed in the distal small intestine (P < 0.05), and a trend toward increased proportion of mucosa and activities of some brush border enzymes in the proximal small intestine. In situ abundance of α1,2-fucose and E. coli was similar between groups, whereas sequencing showed higher abundance of Enterobacteriaceae in F18, Enterococcus in control and Lachnospiraceae in 2FL-F18 pigs.

Conclusions: 2′-FL inhibited in vitro adhesion of E. coli F18 to epithelial cells, but had limited effects on diarrhea and mucosal health in newborn pigs challenged with E. coli F18.

Key Words: α1,2-fucosyllactose, E. coli, human milk oligosaccharide, infection, newborn pigs

What Is Known

- Breastfeeding reduces the risk of infectious diarrhea.
- Human milk oligosaccharides prevent bacteria-induced diarrhea in older infants but limited information is available from newborns.
- Piglets are sensitive to enteric infections, making them good models to investigate dietary effects on the gut in sensitive newborn infants.

What Is New

- α1,2-Fucosyllactose reduced Escherichia coli F18 epithelial adhesion, but failed to prevent diarrhea in newborn F18-challenged pigs. Marginal improvements were seen for intestinal structure and function.
- α1,2-Fucosyllactose supplementation has limited protective effects on the newly colonized, immuno-compromised newborn intestine. Benefits of human milk oligosaccharide may be highly age, diet, and dose dependent.

Compared with formula feeding, breastfeeding reduces the risk of infectious diarrhea, a worldwide problem and a leading cause of infant deaths, especially in the third world (1).

Received September 30, 2015; accepted May 21, 2016. From the *Comparative Pediatrics and Nutrition, Faculty of Health and Medical Sciences, University of Copenhagen, Copenhagen, the National Veterinary Institute, Technical University of Denmark, Lyngby, the Department of Food Science, Faculty of Science, University of Copenhagen, Frederiksberg, and the §Arla Foods Amba, Brabrand, Viby J, Denmark.

Address correspondence and reprint requests to Stine B. Bering, PhD, Comparative Pediatrics and Nutrition, Department of Veterinary Clinical and Animal Sciences, Faculty of Health and Medical Sciences, University of Copenhagen, 68 Dylægevej, DK-1870 Frederiksberg C, Denmark (e-mail: sbb@sund.ku.dk).
Infectious Diarrhea in Newborn Pigs

Inhibiton of E. coli F18 Adhesion to PSlc1 Cells

PSIc1 cells (BioNutriTech, Lunel, France) originating from jejunal epithelium of an adult pig were cultured in DMEM supplemented with 2% fetal bovine serum (FBS), 2 mmol/L Glutamax, 40 U/mL penicillin, and 40 μg/mL streptomycin (all from Gibco, Life Technologies, Carlsbad, CA) at 37°C and 5% CO2 in a humidified atmosphere. At 90% to 95% confluence, cells were harvested by trypsinization using 10 Trypsin:EDTA (Gibco, Life Technologies) and cultured in T75 culture flasks and 12-well plates (TPP, Trasadingen, Switzerland) for genotyping and adhesion assay, respectively. TaqMan single nucleotide polymorphism genotyping was performed as previously described (18) to investigate guanine (G)/adenine (A) polymorphism at nucleotide 307 of the FUT-1 gene, in which M307G/G or M307G/A represents susceptibility to E. coli F18 adhesion (15).

First, it was tested if E. coli F18 and the control E. coli ATCC 25922 strain adhered to PSIc1 cells. After cultivation in 12-well plates, each well of confluent PSIc1 cells was added 300 μL E coli solutions of OD0.5 to OD0.0005, giving a cell:bacteria ratio of 1:1000 to 1:1. Cells and E. coli were then coincubated at 37°C and 5% CO2 in a humidified atmosphere in medium without antibiotics. After 2 to 3 hours, each well was washed twice in cold PBS, and cells were harvested in 1 mL PBS, vortexed thoroughly and plated in × 10 dilutions on LB agar plates for bacterial enumeration. To visualize adhesion of E. coli F18, 3 hours coincubation and subsequent washing in PBS was performed as above on 8 well CultureSlides (BD Falcon, WVR, Herlev, Denmark) in a cell:bacteria ratio of 1:100. Bacteria were stained with LIVE/DEAD BacLight stain (Molecular Probes, Life Technologies) according to the manufacturer’s protocol and visualized using a florescence microscope (Carl Zeiss, Oberkochen, Germany).

To test the antiadhesive effects of 2'-FL, the PSIc1 cells and E. coli F18 or E. coli ATCC 25922 (both at OD0.5) were incubated as above, but before incubation the medium was supplied with 2'-FL (produced utilizing technology from Glycosyn LLC, Medford, MA, by DuPont Nutrition and Biosciences ApS, Copenhagen, Denmark) in concentrations of 1 or 5 μL/L compared with a control only added saline. Lactose and maltose were used as control sugars at equivalent molar concentrations. Bacteria were enumerated on agar plates as above. The level of inhibition was calculated based on 8 to 14 replicates for E. coli F18 and 4 replicates for E. coli ATCC 25922.

Pig Studies

All animal procedures were approved by the Danish National Committee on Animal Experimentation.

Experiment 1: 2'-FL Tolerance Study

A blinded study was set up to test pig tolerability toward 2'-FL to verify that large doses of 2'-FL would not induce osmotic diarrhea. Two sows (Danish Landrace × Large White × Duroc) were selected for homozygosity (M307G/G) of the FUT-1 gene to ensure at minimum heterozygosity of the offspring (18). Thirty pigs were delivered at term by cesarean section and transferred to individual temperature controlled incubators and, still under anesthesia, fitted with a vascular catheter in the umbilical aorta and an orogastric catheter. During the first 24 hours, pigs were infused with plasma from the sow (4–7 mL/kg) to ensure passive
immunization and with parenteral nutrition (7 mL/kg/h of Kabiven and Vamin, Fresenius Kabi, Bad Homburg, Germany) adjusted to the nutrient requirements of pigs. The above procedures has been thoroughly described previously (20). To standardize the initial gut colonization within each litter, each pig received an oral fecal suspension from the sow on day 1 (1 mL with totally 2 × 10^9 cfu). A baseline blood sample was taken from the umbilical catheter and analyzed for blood gases on a GEM-Premier3000 (Instrumentation Laboratory, Zaventem, Brussels, Belgium). From day 2, full enteral feeding was given as boluses of 15 mL/kg every 2 hours with the exception of a 4 hours break during night, reaching a daily dose of 150 mL/kg/day. Pigs were fed a milk replacer (MILEX, Arla Foods, Viby, Denmark) with added protein (50 g/L Lacprodan Df-9224, Arla Foods) as previously described (18). Based on birth weight and sex, the pigs were allocated into 4 diet groups, supplementing the milk replacer with 2'-FL to a total concentration of 10 g/L (FL-10, n = 7), 5 g/L (FL-5, n = 8), 2 g/L (FL-2, n = 7), or no 2’-FL (controls, n = 8).

Body weight and temperature was recorded every morning before feeding, and the consistency and amount of feces was scored morning and evening, where 0 was absence of feces, 1 was normal feces, 2 was pasty feces, 3 was droplets of watery feces, 4 was moderate amounts of watery feces, and 5 was extensive amounts of watery feces. Also, the level of hydration, as assessed by skinfold test and piglet physical activity, was assessed twice daily. In case of severe dehydration and apathy, the pigs were euthanized before the scheduled euthanasia and tissue collection on day 5. Intestinal permeability was determined as the post-mortem urine ratio of lactulose and mannitol after oral administration of the 2 sugars (0.5 and 0.3 g/kg, respectively) 3 hours before euthanasia (21). At euthanasia, blood gases were measured as above, and 10 cm sections of proximal and distal small intestine were sampled for determination of the wet and dry proportion of mucosa (21).

Experiment 2: E. coli F18 Dose-Response Study

The optimal E. coli F18 inoculation dose was then identified in a dose-response study. Thirty-one term pigs were delivered by cesarean section from 2 sows, fitted with orogastric and umbilical catheters, immunized with sow’s serum, and nourished by parenteral nutrition as described for experiment 1. For the following 4 days feeding, the pigs were fed 15 mL/kg three hours (Miles formula) as above supplemented with 23 g/L Lacprodan, Df9224) until euthanasia on day 5. The pigs were randomly assigned to 4 different treatment groups, controls (CON, n = 8) or 3 doses of F18, low dose (LOW, 1 × 10^7 CFU/day, n = 9), medium dose (MED, 2 × 10^7 CFU/day, n = 7), and high dose (HIGH, 8 × 10^7 CFU/day, n = 7). E. coli F18 was cultured overnight as described above, and harvested with 10 mL PBS. From the first day and throughout the experiment all LOW, MED, and HIGH pigs were inoculated daily with 1 mL F18 suspension. The suspension was given through the orogastric catheter followed 5 mL of the Miles formula. Initial gut colonization was standardized by giving 1 mL of maternal fecal suspension, as described in experiment 1. Body weight was recorded daily and fecal consistency and amount was scored twice daily as described for experiment 1. Likewise, clinical condition, hydration, intestinal permeability, and mucosal proportions were measured. Sections of distal small intestine were formalin fixed and paraffin embedded for fluorescence in situ hybridization (FISH) with an E. coli–specific 16 s rRNA probe (S-S-E. coli-116L, 5’TCCATAAAGCGCTGCTGGCCG 3’-3’ labeled with fluorescein isothiocyanate derivative (CY3, red signal) to evaluate abundance of E. coli in the proximal and distal small intestine. In short, 3 μm tissue sections were mounted on glass slides (Superfrost/Plus slides, Menzel-Gläser, Braunschweig, Germany), deparaffinized in xylene, dehydrated in ethanol, hybridized overnight with the probe at 49 °C using Shandon Coverplates and Sequenza immunostaining workstations (Thermo Scientific, Waltham, MA) followed by 3 × washing in hybridization buffer (Tris 0.1 mol/L, NaCl 0.9 mol/L, 0.1% SDS, pH 7.2) and 3 × washing in washing buffer (Tris 0.1 mol/L, NaCl 0.9 mol/L, pH 7.2). An Axioimager M1 epifluorescence microscope (Carl Zeiss) was used for visualization and images were obtained using an AxioCam MRm version 3 monochrome camera and multidimensional acquisition in the software AxiosVision version 4.5 (Carl Zeiss). Based on the abundance of E. coli, each pig was given a standardized score where 1 = no bacteria, 2 = few bacteria, 3 = some bacteria, and 4 = many bacteria in the tissue section.

Experiment 3: 2’-FL F18 Infection Study

Finally, an intervention study was performed to test whether 2’-FL inhibited E. coli F18 infectious diarrhea. Twenty-five pigs were delivered at term by cesarean section from 2 FUT-1 homozygous sows (Danish Landrace x Large White x Duroc). As described above, the pigs were prepared with umbilical and orogastric tubes, infused with parenteral nutrition and sow’s plasma, inoculated with maternal feces, and from day 2 transferred to full enteral feeding with boluses of milk replacer. Based on birth weight and sex, the pigs were allocated into 3 groups. One group was inoculated daily with 7.5 × 10^7 E. coli F18 (F18, n = 9), 1 group was given the same dose of E. coli F18 plus 10 g/L 2’-FL in the milk replacer as described in the tolerance study above (2FL-F18, n = 8) and 1 control group received only milk replacer (control, n = 8). E. coli F18 was cultured overnight at 10% CO2 on Iso-sentistest agar with Alizarin yellow as described above, and harvested with 10 mL PBS, of which each pig received 1 mL. On day 1, E. coli F18 was given 1 mL of the respective diets, while inoculations the following days were followed by enteral feeding. The sensitivity toward diets and pathogen inoculation were followed until euthanasia on day 8, unless severe dehydration and apathy required euthanasia beforehand.

Body weight was recorded every second day and clinical evaluation was assessed as above. Fecal consistency scores were evaluated morning and evening using a scoring system, where 0 = absence of feces, 1 = normal feces, 2 = pasty feces, and 3 = watery feces. Also, the level of hydration, as assessed by skinfold test and piglet physical activity, was assessed twice daily. In case of severe dehydration and apathy, the pigs were euthanized before scheduled euthanasia and tissue collection on day 5. Intestinal permeability was determined as the post-mortem urine ratio of lactulose and mannitol after oral administration of the 2 sugars (0.5 and 0.3 g/kg, respectively) 3 hours before euthanasia (21). At euthanasia, blood gases were measured as above, and 10 cm sections of proximal and distal small intestine were sampled for determination of the wet and dry proportion of mucosa (21).
light for 1 hour and finally washed in 1× PBS. All procedures were performed at room temperature. Sections were evaluated under microscope as above and given a score where 0 = no staining, 1 = few positive stained goblet cells, 2 = several positive stained goblet cells, mainly in the crypt area or moderate luminal staining, 3 = abundant positive stained goblet cells along the entire mucosal lining, and 4 = abundant positive stained goblet cells along the entire mucosal lining together with extensive luminal staining.

Statistical Analyses

Data analyses were performed in R version 3.0.1. Treatment effects on diarrhea and weight development were analyzed using the linear function of repeated measurement analysis with litter and pig as random variable and the factors treatment and time as fixed variables. In vitro adhesion data and treatment effects on relative weight gain, intestinal parameters, blood parameters, FISH, and lectin hybridization from the pig studies were tested using the linear function of mixed model analysis with sow as random variable and treatment, birth weight and sex as fixed variables. In case of significance, pairwise analysis using the glht function was employed to compare effects of the different treatments. Normal distribution of all measured parameters was tested by plotting the model residuals and by the qq-plot using the qnorm function. All values given in the report are adjusted means and standard error of mean. Data are considered significant for \( P < 0.05 \).

RESULTS

In Vitro Adhesion

Genotyping of PSIc1 cells showed homozygosity (M307\( ^{G\rightarrow G} \)) for the F18 susceptible genotype of FUT1 (data not shown). Both \( E\ coli \) F18 and ATCC 25922 adhered to the PSIc1 cells, which was confirmed on culture slides by fluorescence microscopy for \( E\ coli \) F18 (Fig. 1A). High doses of 2'-FL (5 g/L) effectively inhibited \( E\ coli \) F18 adhesion (\( P < 0.001 \); Fig. 1B), whereas inhibition of \( E\ coli \) F18 adhesion was less effective but significant when lactose and to a lesser extend maltose (not shown) was added (\( P < 0.001 \) and 0.05, respectively). Adhesion of \( E\ coli \) ATCC 25922 was not affected by 2'-FL or any of the control sugars (Fig. 1B).

Experiment 1: 2'-FL Tolerance Study

One control pig and 1 FL-5 pig were euthanized before completion of the protocol due to respiratory problems or apparent sepsis. Fecal scores from these 2 pigs were included in the final calculations. Although 3 FL-10 pigs (43%), 1 FL-5 (13%) and 2 FL-2 pigs (29%) had lost weight at euthanasia relative to their birth weight, there was generally a slight increase in weight during the experiment, with no difference between groups at any time point (\( P = 0.3–0.7 \); Supplemental Fig. 1A, http://links.lww.com/MPG/A704). Fecal consistency was noted from day 2 where feces first appeared and thus resulted in 6 observations in total per pig. Pigs from all treatment groups developed diarrhea (score 3–5) at some point; 3 controls (38%), 3 FL-2 pigs (43%), 4 FL-5 (50%) and 2 FL-10 pigs (29%), and mean fecal scores generally increased during the 5-day period with no difference between groups (\( P = 0.3–0.8 \); Supplemental Fig. 1B, http://links.lww.com/MPG/A704). Intestinal permeability, expressed as the lactulose/mannitol ratio, did not differ among groups (0.02–0.12 across groups). There were no group differences in dry weight mucosal proportions in the proximal small intestine or distal small intestine (75 ± 2% and 73 ± 2%, respectively).

The rectal temperature increased from 36.2 ± 0.0°C after delivery to 39.1 ± 0.1°C on the following day in which it remained stable until euthanasia. There was no difference between treatment groups and all pigs remained within the normal range during the experiment. Blood gas values showed no differences among groups, except that FL-2 pigs had lower hematocrit values than FL-10 pigs (22.6 ± 1.0% vs 26.5 ± 0.9%, \( P < 0.05 \)). Across all groups, mean hematocrit decreased from 35.2 ± 0.6% at birth to 25.1 ± 0.8% at euthanasia (\( P < 0.05 \)). At birth, mean \( pO_2 \) and \( pO_2/G \) was higher than \( PO_2 \) at euthanasia (119.5 ± 17.3 vs 52.3 ± 5.6 mmHg and 52.5 ± 1.5 vs 47.6 ± 1.8 mmHg, respectively, \( P < 0.05 \)), whereas there was no overall difference in \( p\)H (\( 7.47 ± 0.01 \) vs 7.41 ± 0.03 at birth and euthanasia, respectively, \( P = 0.13 \)). Blood glucose concentration increased from birth to euthanasia (2.6 ± 0.2 vs 5.2 ± 0.4 mmol/L, respectively, \( P < 0.05 \))

 Experiment 2: \( E\ coli \) F18 Dose-Response

The incidence of diarrhea increased with time after inoculation (\( P < 0.01 \)), and was dose dependent with more diarrhea in HIGH versus CON (\( P < 0.05 \)), and a tendency to more diarrhea in MED versus CON and HIGH versus LOW pigs (both \( P < 0.07 \); Fig. 2). Body weight gain was lower in the HIGH group compared with CON (\( P < 0.01 \)), but hydration score (1.06 ± 0.01), mucosal proportion (77.8 ± 3.9%), and intestinal permeability (0.017 ± 0.004) did not differ among groups. The abundance of \( E\ coli \) in the proximal (1.1 ± 0.4) and distal small intestine (1.8 ± 0.15) as measured by FISH analyses did not differ between groups.

FIGURE 1. Microscopic visualization of \( Escherichia\ coli\ F18\) adhesion to PSI cells (A) where bacteria are stained with a red fluorescence dye. Mean relative adhesion of \( E\ coli\ F18\) and control \( E\ coli\ ATCC\ 25922\) to PSI cells in vitro after addition of 2'-FL and lactose (1 and 5 g/L, B). Different superscript letters indicate significant differences (\( P < 0.05 \)).
antiadhesive potential of the intervention. Also the high Escherichia coli F18 dose was chosen to ensure consistent diarrhea development (experiment 2). All 9 F18-pigs, 6 out of 8 2FL-F18 pigs and 2 of 8 control pigs were euthanized before completion of the protocol because of extensive diarrhea and poor clinical condition (Fig. 3A). Nevertheless, there was no difference in the mean age at euthanasia among the treatment groups (154 ± 9, 141 ± 9, and 132 ± 8 hours for control, 2FL-F18, and F18 pigs, respectively, P = 0.20). At euthanasia, control and 2FL-F18 pigs tended to show less relative weight loss than F18 pigs (P = 0.08; Fig. 3B). All pigs developed diarrhea during the experiment, but overall both F18 and 2FL-F18 pigs had more severe diarrhea, and earlier onset of diarrhea, than controls (mean score 2.0 for F18 and 2FL-F18 pigs vs 1.2 in control pigs, P < 0.01; Supplemental Fig. 2, http://links.lww.com/MPG/A705).

The rectal temperature increased from 36.0 ± 0.0 C after delivery to 38.5 ± 0.1 C the following day, where it remained stable until euthanasia. There was no difference between treatment groups and all pigs remained within the normal range with no evidence of fever (sepsis). Because of poor clinical status at euthanasia, blood gasses were not obtained from 6 F18 and 6 2FL-F18 pigs and 2 controls. Plasma glucose concentrations were elevated in F18 pigs, compared with 2FL-F18 and controls (41.0 ± 2.8 % vs 23.2 ± 2.2 % and 28.5 ± 1.2 %, respectively, P < 0.01). Blood pH was lowest for F18 pigs (7.17 ± 0.11 vs 7.40 ± 0.09 and 7.27 ± 0.06 for F18, 2FL-F18, and controls, respectively), but the differences were not significant (P = 0.34 and 0.73, respectively).

As a result of severe dehydration, urine was only obtained from a few pigs, and lactulose and mannitol were therefore not analyzed for permeability measures. By histological evaluation, mucosal damage in terms of destroyed villi with total or partly sloughing of enterocytes was observed in 25%, 14%, and 44% of controls, 2FL-F18, and F18 pigs, respectively (P = 0.32). F18 pigs had shorter villi than 2FL-F18 pigs and tended to have shorter villi than controls (P < 0.05 and P = 0.06, respectively, Table 1). Dry weight mucosal proportions in the proximal and distal small intestine was not different between groups (P = 0.47 and 0.41, respectively, Table 1).

The activities of 6 brush border enzymes in the proximal and distal small intestine were not different among groups (Table 1); however, the 2FL-F18 pigs showed the highest mean values for all 3 disaccharidases and aminopeptidase N in the proximal intestine, relative to F18 pigs, but the differences did not reach significance (Table 1, P = 0.07–0.37). Values for aminopeptidase A and N and dipeptidyl peptidase IV were higher in the distal small intestine than in proximal intestine (P < 0.05 for all 3 enzymes). FISH analyses revealed low total abundance of Enterococcus in the distal small intestine (Fig. 4A), where 36% of the tissues were negative for Enterobacteriaceae genus. There were no differences in mean Enterococcus abundance among the treatment groups (Fig. 4B), and no difference in numbers of tissues negative for Enterobacteriaceae genus (22%, 57%, and 38% for F18, 2FL-F18, and controls, respectively, P = 0.3–0.8)) or numbers of tissues with high abundance (score 3–4; 29%, 22%, and 50% for F18, 2FL-F18, and controls, respectively). Furthermore there was no significant correlation between FISH score and mean fecal scores (P = 0.26). The F18 pigs had higher relative abundance of an unclassified Enterobacteriaceae genus, compared with control pigs (red bar, P < 0.01; Fig. 4C), whereas Enterococcus tended to be highest for control pigs (dark blue bar, P = 0.058). Compared with 2FL-F18 pigs, the control pigs had higher Enterococcus levels (P < 0.05).

The 2FL-F18 pigs had an increased relative abundance of an unclassified Lachnospiraceae genus, compared with F18 pigs (purple bar, P < 0.05).
Lectin hybridization revealed modest to high levels of α-1,2-fucosyltransferase activity in the distal intestine (Fig. 4D), with strong staining of goblet cells, indicating endogenous production of α-1,2-fucose. A few pigs from all 3 treatment groups only displayed positive staining in the lumen and there were only few observations of α-1,2-fucose associated with the cell surface. The average score was not affected by treatment (2.0 ± 0.7, 2.5 ± 0.7, and 2.3 ± 0.7 for controls, 2FL-F18, and F18, respectively). There were no significant correlations between these scores but a negative correlation was observed between α-1,2-fucose score and E. coli FISH score (P < 0.01, r = −0.33, data not shown).

### DISCUSSION

Breastfeeding is strongly associated with a reduced risk of infectious diarrhea, especially in developing countries (1), and the effect may be due to numerous bioactive compounds in human milk. A compound that may have a significant effect is 2′-FL, the major HMO in human milk, which has antiadhesive effects on the intestinal epithelium due to its structural homology with intestinal adhesins sites for pathogenic bacteria (9,10). This has been demonstrated for C. jejuni at a dose of 10 g/L in mice and also for human intestinal biopsies (9). Absence of 2′-FL may have more important effects in vivo data indicate that the antiadhesive effect of 2′-FL occurred via F18 fimbria-dependent mechanism. Importantly, the effect was dose dependent, with no effect of 2′-FL at the low 1.0 g/L concentration. The 2′-FL concentrations applied in vitro relates to the physiological concentrations found in human milk, but may not be correctly translated to the in situ situation, in which dilution by gastric and intestinal secretions and the mucus layer would reduce the level of 2′-FL in contact with the epithelium, depending on the infants health status. Preterm infants with very immature gastrointestinal functions and low digestibility may have reduced capacity to secrete these fluids and in situ concentrations may come closer to the raw milk concentrations. Considering the 5.0 g/L as a maximum of physiological relevant concentration, and the lacking effects at 1.0 g/L, the in vitro studies may confirm the minor effects of 2′-FL in the pig studies, even though the concentrations represent physiological levels. The dose-dependent effect implies that optimization of the dosage for in vivo administration is crucial to exert protection from E. coli F18 infections. Therefore, we first tested 2′-FL in a dose-response newborn pig study comparing a low (1 g/L), medium (5 g/L), and high (10 g/L) dose of 2′-FL without E. coli F18 infection. Considering a potential risk of osmotic diarrhea, the high dose was taken from the upper range of physiological concentrations in human milk (up to 8 g/L) (25–27). Even using a relatively high dose of 2′-FL, diarrhea was not prevented in newborn pigs inoculated with E. coli F18. The explanation for the lack of effect may be related to the very immature state of the intestine and its colonization but also the dosing of 2′-FL and E. coli F18. The 10 g/L 2′-FL used for the formula may not result in a comparable high concentrations at the epithelial surface since dilutions due to both endogenous mucus production and other epithelial secretions occur. In our in vitro studies on cells devoid of these physiological characteristics a dose of minimum 5 g/L was needed for significant antiadhesive effects and ex vivo studies in mice and human intestinal biopsies required a dose of 10 g/L to inhibit adhesion of C. jejuni (9). The observed diarrhea in the 2FL-F18 group may be partly an osmotic diarrhea because HMOs are generally indigestible and become fermented by bacteria. Excessive fermentation increases the osmolality of the luminal contents and thereby induce osmotic diarrhea (28). It is therefore possible that the diarrhea observed in the 2FL-F18 and F18 pigs was of a different type, being more

### TABLE 1. Villus height, mucosal proportion, and activity of intestinal brush border enzymes in the proximal (Prox) and distal (Dist) small intestine of control, 2FL-F18, and F18 pigs

<table>
<thead>
<tr>
<th></th>
<th>Control</th>
<th>2FL-F18</th>
<th>F18</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Prox</td>
<td>Dist</td>
<td>Prox</td>
</tr>
<tr>
<td>Villus, μm</td>
<td>358.3 ± 18.8</td>
<td>351.3 ± 14.1</td>
<td>303.5 ± 14.4*</td>
</tr>
<tr>
<td>Mucosa, %</td>
<td>50.0 ± 5.9</td>
<td>60.4 ± 1.6</td>
<td>60.0 ± 1.3</td>
</tr>
<tr>
<td>Sucrase, U/g</td>
<td>0.34 ± 0.16</td>
<td>0.25 ± 0.04</td>
<td>0.58 ± 0.17</td>
</tr>
<tr>
<td>Maltrase, U/g</td>
<td>2.01 ± 0.60</td>
<td>2.02 ± 0.19</td>
<td>3.90 ± 0.05</td>
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<tr>
<td>Lactase, U/g</td>
<td>5.15 ± 1.77</td>
<td>3.48 ± 0.68</td>
<td>8.54 ± 1.86</td>
</tr>
<tr>
<td>ApN, U/g</td>
<td>2.23 ± 0.58</td>
<td>3.53 ± 0.62</td>
<td>3.00 ± 0.60</td>
</tr>
<tr>
<td>ApA, U/g</td>
<td>0.88 ± 0.30</td>
<td>1.79 ± 0.28</td>
<td>1.31 ± 0.31</td>
</tr>
<tr>
<td>DPPIV, U/g</td>
<td>0.73 ± 0.13</td>
<td>2.23 ± 0.58</td>
<td>1.07 ± 0.14</td>
</tr>
</tbody>
</table>

Values are Mean ± standard error of mean.

*Villus height in F18 is significantly different from 2FL-F18 (P < 0.05). ApA = aminopeptidase A; ApN = aminopeptidase N; DPPIV = dipeptidyl peptidase IV.
osmotic for the 2FL-F18 pigs and more infectious for the F18 group, in the latter group leading to more mucosal disruption as shown for infectious diarrhea induced by ETECs (29). On the contrary, the tendencies for 10 g/L 2'-FL to reduce the weight loss, blood parameters, and mucosal lesions in F18-infected, formula-fed piglets may result from direct host tissue responses via immune modulation, or modulation of epithelial glycosylation processes, as shown for 2'-FL and other HMOs (6,24).

We based the E coli F18 dose of 7.5 × 10¹⁰ cfu/day on findings from experiment 2 in which a high dose (8 × 10⁹ cfu/day) more effectively induced diarrhea than lower doses. A previous study indicated an apparent upper tolerability limit of 2.6 × 10¹¹ cfu/day (unpublished results). In the present 2'-FL intervention study, diarrhea was more severe in both inoculated groups compared with control pigs. FISH analysis, however, did not reveal higher intestinal abundance of total E coli in the inoculated groups of experiments 2 and 3. Still, data from this and an earlier study (18) document a robust and consistent pathogen-specific induction of diarrhea by E coli F18 in newborn, cesarean-delivered pigs. Replacement of formula with bovine colostrum did not prevent E coli F18 infection in

![Graph showing mean FISH score](image1)

**FIGURE 4.** Representative tissue from the distal small intestine showing high abundance of Escherichia coli (red signal) visualized by FISH (A), and abundance of E colin the treatment groups in the 2′FL F18 infection study (experiment 3) (B, means and standard error of mean). Average relative abundance distribution of the major genera in colon contents of control, F18, and 2FL-F18 pigs by 16S rRNA gene MiSeq-based high-throughputs sequencing (C). Tissue sections from the distal small intestine hybridized with fluorescence- (fluorescein isothiocyanate [FITC]) labeled lectin (ulex europaeus agglutinin I [UEA-I]) specific for α-1,2-fucose (D). The sections shown indicate tissues with fucosylation scores 1 (most left), 2, 3, and 4 (most right). FISH = fluorescence in situ hybridization.
this model (unpublished observations), despite the higher contents of oligosaccharides in colostrum versus mature milk. Both sow’s and cow’s milk contain significantly lower amounts of fucosylated oligosaccharides than human milk (27), and the contents of 2′-FL in sow’s milk has not yet been reported. Regardless, it appears that the intestine of suckling pigs is not likely to be highly dependent on soluble 2′-FL from milk for natural pathogen protection. Conversely, our in situ hybridization analyses with the α-1,2-fucosyl—specific lectin, ulex europaeus agglutinin I, revealed high endogenous secretion of α-1,2-fucose in the pig small intestine. The physiological effect of these may possibly override the modest additional effect of 2′-FL supplemented via the formula diet. In contrast to humans, pigs may therefore rely more on endogenous α-1,2-fucose, explaining the lacking or modest effects of 2′-FL supplementation on enteric F18 infection in this study. Although the endogenous α-1,2-fucose could not prevent E. coli F18 diarrhea in our newborn pig model, there was a negative correlation between the level of endogenous α-1,2-fucose and the abundance of E. coli in general and this may reflect some effect of α-1,2-fucose on intestinal colonization.

Our in vitro studies showed that 2′-FL at a concentration of 5 g/L inhibited E. coli F18 adhesion to porcine intestinal epithelial cells. The lacking correlation between the E. coli FISH score and diarrhea score may suggest that even low titers of E. coli F18 can induce diarrhea. The binding of E. coli F18 to luminal H-2 epitopes may also not be the only mechanism whereby the pathogens adhere to the epithelium and become virulent. In weanling pigs, E. coli F18 diarrhea was most prevalent in susceptible pigs being homo- or heterozygote for the functional FUT-1 gene, but also unsusceptible pigs developed diarrhea (14). A similar finding was reported ex vivo (16) and indicates that pathogenesis may occur by other means than by binding specifically to α-1,2-fucosylated receptors. Other receptors than H-2 may allow E. coli F18 epithelial adhesion and increase the efficiency of inhibition of E. coli F18 adhesion and thus decreased prevalence of diarrhea may result from supplementation with multiple HMOs and antimicrobial milk factors.

The increased luminal abundance of an unclassified Enterobacteriaceae genus in the F18 pigs (compared with controls) most likely includes the inoculated E. coli F18. The abundance of Enterobacteriaceae in the 2FL-F18 pigs was intermediate between values in the other 2 groups, but not significantly different from either of them, indicating that 2′-FL supplementation tend to reduce the abundance of E. coli F18. The unclassified Lachnospiraceae family was relatively more abundant in 2FL-F18 pigs, supporting the increase of this group in responses to HMO supplementation, including 2′-FL, in newborn pigs (30). The control group showed the highest abundance of Enterococcus, despite that these were increased in abundance after 2′-FL supplementation in preterm pigs (31). From the above, it seems that inoculation with E. coli F18 may override any potential effect of 2′-FL on Enterococcus density.

From the present in vitro and in vivo studies we conclude that 2′-FL reduces E. coli F18 adhesion in the intestine of newborn, cesarean-derived, formula-fed pigs. The employed dose of 2′-FL was not sufficient to prevent diarrhea when E. coli F18 was inoculated at a dose of 7.5 × 10^6 cfu/day. Tendencies to improve intestinal morphology and health in the 2FL-F18 pigs suggest only a modest beneficial effect of 2′-FL. It remains to be investigated whether an increased dosage of 2′-FL, or a longer time of exposure in a more mature intestine, more effectively prevents pathogen-induced intestinal infections in early life of pigs and infants.

Acknowledgments: We acknowledge Mette Schmidt for assistance with cesarean section, Elin Skytte for assistance with animal procedures and laboratory analyses, Joanna Z. Amenuvor and Annie R. Pedersen for assistance with tissue preparation for in situ analyses, and finally Christian Ritz for assistance with statistical analyses in R.

REFERENCES
15. Meijerink E, Fries R, Vo P, et al. Two or (1,2) fucosyltransferase genes on porcine Chromosome 6q11 are closely linked to the blood group inhibitor (S) and Escherichia coli F18 receptor (EClF18) loci. Mamm Genome 1997;8:736–41.

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