Neonatal Feeding Tube Colonization and the Potential Effect on Infant Health: A Review

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Background: Infants in the neonatal intensive care unit (NICU) often require feeding tubes (FT) for weeks to months. Because FTs are in near constant contact with human milk and/or formula, rapid and extensive bacterial growth is possible. Due to their immature immunologic and gastrointestinal (GI) systems, infants may be at significant health risk due to FT colonization. In adults, length of time FTs remain in place (dwell time) affects the degree of colonization and biofilm formation which is important in infants whose tubes remain in place up to 30 days.

Objective: The purpose of this review was to describe and summarize the evidence regarding FT bacterial colonization in infants and identify gaps needing further investigation.

Methods: Medline, CINAHL, and Embase databases were searched for clinical and/or laboratory-based observational and randomized controlled studies investigating the presence of bacteria in neonatal FTs.

Results: This review of 10 studies found evidence that neonatal FTs may contain high quantities of potentially pathogenic and antibiotic resistant bacteria and longer dwell times may increase the bacterial load. Furthermore, evidence suggests FT colonization may be nosocomial in origin and contribute to adverse infant health. Feeding tubes are an unrecognized source of bacterial colonization which may increase morbidity in premature infants and thus the presence of bacteria in FTs is an important area of investigation in the nutritional care of vulnerable infants in the NICU.

Implications: Further appropriately powered studies which are clinically based, use appropriate analyses, and control for potential covariates are necessary to make clinical recommendations.

Keywords: infant, neonatal intensive care unit, feeding tube, contamination, colonization, enteral feeding, dwell time, infection risk
BACKGROUND

Due to prematurity and/or illness, infants in the neonatal intensive care unit (NICU) often require feeds to be provided via a feeding tube (FT). Feeding tubes are in near constant contact with formula or human milk providing ample nutrition for bacterial consumption and growth (1). Furthermore, frequent handling by health care workers may increase the risk of FT colonization which is especially important in infants in the NICU whose FTs are handled every 2–3 h in accordance with their feeding schedule (2). The presence of a FT is associated with an increased risk of late onset sepsis in preterm infants (3, 4) and it is possible this risk may be related to excessive FT colonization.

Previous research in adults indicates FTs are colonized with pathogenic bacteria and biofilm forms in the tube lumen (5). Biofilm formation is common in medical devices and significantly increases the risk of infection by enhancing bacterial resistance to antimicrobials and defending against environmental cleansing agents (6, 7). Because of their immature immunological and gastrointestinal (GI) systems, increased intestinal mucosal permeability, and often dysbiotic intestinal microflora, infants may be particularly vulnerable to FT colonization (2, 8, 9). Feeding administered through colonized FTs may introduce potentially pathogenic bacteria into the infant’s GI system increasing the risk of adverse health outcomes. Provision of contaminated feedings are known to cause neonatal infection further suggesting that bacteria entering the infant’s GI system through colonized FTs may be a risk for infection (2, 10).

The health risk associated with external devices such as central venous lines and urinary catheters is well-known to increase the longer the device remains in place (11). Because bacterial growth occurs rapidly in nutrient-rich environments, FT dwell time may influence bacterial growth. In adults, biofilm formation in FTs correlates with dwell time and develops in 60% of FTs in place ≤1 day, and 100% of tubes in place >1 day with dwell times >48 h resulting in dense biofilm formation (5, 12). In infants in the NICU, FTs often remain in place for up to 30 days likely increasing the risk for bacteria to multiply to potentially unsafe levels.

A systematic evaluation of available research on FT colonization in infants is lacking. Due to the potential health risk of colonized FTs in vulnerable infants in the NICU, the aim of this review is to describe and summarize the evidence regarding FT colonization in infants, to explore evidence on the association of FT dwell time and colonization and to demonstrate the need for further research.

METHODS

Clinical and laboratory studies investigating the presence of bacteria in infant FTs were included in this review. Clinical studies were limited to prospective observational studies and randomized control trials (RCTs).

Search Strategy

To identify eligible studies, a search was conducted of the following electronic databases: Medline (through PubMed), CINAHL, and Embase. Additionally, the reference lists of included articles were examined. Our search was limited to studies published between January, 2000 to December, 2020 in English and used a combination of different groups of keywords [free text and MeSH (Medical Subject Headings) terms] that described the population and outcomes of interest (Supplementary Table 1). All titles and abstracts identified by our search were screened for relevance by a single author (LAP). The full texts of those considered relevant were retrieved and evaluated for inclusion relevancy by three authors (LAP, MM, and KD) who independently performed data extraction and collected the following information: first author’s name, year of publication, study design, setting, and results.

A total of 216 articles were retrieved from the databases and reference lists of identified articles. After removing duplicates, 117 articles were screened by title and abstract. Of these, 104 were unrelated to FT colonization and excluded, leaving 13 articles potentially meeting inclusion criteria. After reviewing the full articles, 10 met criteria for inclusion in this review (Figure 1).

Data Analysis

We synthesized data obtained from the included studies and all discrepancies between reviewers regarding study selection process, data extraction, synthesis, and interpretation were discussed until a final agreement was reached. In the following sections we summarize and discuss the findings of these studies and discuss future research priorities.

RESULTS

Presence of Bacteria in FTs

All 10 included studies quantified or described the presence of bacteria in infant FTs (1, 2, 8, 9, 12–17). Five studies were laboratory-based which included studies that either analyzed inoculated FTs or performed additional analysis on specific bacteria isolated from FTs previously removed from infants in the NICU (1, 8, 13, 16, 17). Five were clinically based and included primary analysis of FTs removed from infants in the NICU (2, 9, 12, 14, 15). Two clinically-based studies included the same sample of infants and FTs (2, 13) and one laboratory-based studies (16) analyzed FTs obtained from clinically-based studies (9). Table 1 summarizes studies included for review.

Four clinically-based (2, 9, 12, 15) and three laboratory-based studies (1, 8, 17) described the presence of bacteria within the FT lumen. Hurrel et al. (9) prospectively analyzed 129 FTs for Enterobacteriaceae finding a maximum bacterial load of 10^7 CFU/tube while Mehall et al. (2) found 57% of 125 FTs removed from 50 infants to be contaminated (defined as >1,000 CFU of bacteria excluding Staphylococcus epidermidis and alpha-hemolytic Streptococcus) with a mean bacterial load of 908,173 CFU. Finally, Gomez et al. found dense biofilm but did not quantify how many FTs contained biofilm or the actual bacterial load (12) and Taft et al. using 16S rRNA sequencing to characterize the bacterial composition of biofilm found 100% of FTs contained bacteria (15). The three laboratory-based studies provided similar results (1, 8, 17). Hurrel et al. flushed FTs with formula inoculated with Enterobacteriaceae and found
biofilm formation of $10^5$-$10^6$ CFU/cm and a bacterial load up to $10^9$ CFU/cm (8). Similarly, Kim et al., inoculated FTs with Enterobacter Sakaszkii with biofilm formation after 12 h but only when incubated at 25 vs. 12°C (1). Finally, following inoculation of FTs with a 2:1 mixture of formula and infant saliva, Presti et al. found bacterial growth over time but did not report information regarding bacterial load (17).

Two clinically and one laboratory-based study analyzed residual fluid within FTs (9, 15, 16). Taft et al. found bacteria in the residual fluid of all FTs using 16S rRNA sequencing (15). Hurrell et al. found a peak bacterial load of $10^7$ CFU/ml with similar Enterobacteriaceae species as found in the FT lumen (9). Additional laboratory analysis of the residual fluid found 30 similar strains of Escherichia coli in the residual fluid and biofilm within the tube lumen (16).

Three studies examined fluid flushed through FTs (8, 12, 14). Petersen et al. found 89% of normal saline samples flushed through 94 FTs removed from 34 infants in the NICU contained $>1,000$ CFU/ml of bacteria with a mean bacterial load of $5.3 \log_{10}$ CFU/ml, a maximum bacterial load of $9.4 \log_{10}$ CFU/ml, and 55% contained either Enterobacteriaceae or Staphylococcus aureus (14). Similarly, Hurrell et al. found formula flushed through
Enterobacteriaceae inoculated FTs contained 10⁸-10⁹ CFU/ml of bacteria within 24 h (8). Finally, after leaving the external feeding system before entering the actual FT, mother’s own milk (MOM), donor human milk (DHM), and formula were found to contain bacteria (12). Because DHM and formula are considered sterile, these findings suggest bacteria in the external system may penetrate feedings as they pass through.

Identification of Bacteria in FTs

Five studies identified the type of bacteria present in analyzed samples (9, 12, 15, 16). Mehall et al. found the most common bacterial species in FT lumens included S. epidermidis, S. aureus, Enterococcus faecalis, Enterobacter cloacae, and Klebsiella (2), while Hurrell et al. found the most common Enterobacteriaceae isolates included Enterobacter cancerogenus, Seratia marcescens, Enterobacter hormaechei, E. coli, and Klebsiella pneumonia (9). When pulsed-field gel electrophoresis and seven-loci multilocus sequence typing was used to genotype E. coli in these samples, they clustered into five main pulsotypes known to cause serious and potentially fatal neonatal infection (16). It is important to note that FTs from 11 individual infants contained identical strains of E. coli K1 suggesting a common source of this pathogenic strain of bacteria. Similarly, after passing through the external feeding system, Staphylococcus was present in 93% of MOM feedings, 37% of DHM, and 11% of formula feedings (12). Finally, 16s rRNA gene sequencing found the most common phyla in biofilm were Proteobacteria, Firmicutes, and Actinobacteria and the most common genera were Staphylococcus, Enterococcus, and Enterobacteriaceae (15).

Antibiotic Resistant Bacteria

Antibiotic resistance was investigated in three studies (9, 13, 15). When susceptibility to methicillin and vancomycin on isolates of S. aureus and E. faecalis was performed on FTs from infants in the NICU, 12 of 23 S. aureus isolates were methicillin resistant but none of the 71 Enterococcus isolates were vancomycin resistant (13). Resistance was also found in Enterobacteriaceae isolates in an additional clinically-based study where all strains of Seratia marcescens were resistant to amoxicillin and amoxicillin-clavulanic acid, while E. hormaechei strains were resistant to ceftazidime (21%) and cefotaxime (23%), and four of 37 E. coli strains were resistant to ceftazidime and cefotaxime (9). Finally, using whole metagenome sequencing of six FTs, Taft et al. found a median of 9.5–11 transferable antimicrobial resistance genes (ARGs) depending on the section of tube analyzed (15).

Association of Dwell Time With FT Contamination

Six studies, four clinically (9, 12, 14, 15) and two laboratory-based (8, 17), explored bacterial growth in FT over time. Following inoculation with Enterobacteriaceae and flushing with infant formula, Hurrell et al., found FTs contained up to 10⁷ CFU/cm² of bacteria by 8 h and 10⁸-10⁹ CFU/cm² by 24 h with biofilm formation measuring 10⁵-10⁶ CFU/cm² after 24 h (8). The second laboratory study found the highest bacterial growth occurred over the first 3 h followed by 24 and 72 h after FTs were flushed with a mixture of formula and infant saliva, incubated for 72 h and then analyzed (17). The four clinically-based studies reported mixed results. Hurrell et al., found FTs with longer dwell times contained higher bacterial loads (p < 0.001) with a dwell time <6 h having the lowest bacterial counts compared to between 6 and >48 h (9). Similarly, analysis of six FTs found complex biofilm formation after 48 h dwell time and no biofilm in tubes with dwell times of <12 h (12). Increased Alpha diversity indicating an increased number of species was also observed in FTs with longer dwell times (15). In contrast, Petersen et al. reported no association between dwell time and the presence of bacteria or the presence of potentially pathogenic bacteria but comparisons were made between dwell time differences of <1 day which may be insufficient for differences in growth to occur (14).

Infant Outcomes Related to FT Colonization

One clinically-based and one laboratory based study investigated the association of FT colonization and adverse health outcomes (2, 13). Mehall et al. found only infants who received formula rather than human milk experienced feeding intolerance, with a greater incidence when fed through contaminated FTs (p < 0.05) (2). Of the 50 infants enrolled in the study, 10 had FTs containing >100,000 CFU/ml of gram negative rods and of those, seven developed necrotizing enterocolitis (NEC) with four requiring surgery. When surgery was required, bacteria cultured from the peritoneal fluid were identical to those found in each individual infant's FT. Furthermore, infants fed through colonized FTs experienced less mean weekly weight gain (178.9 vs. 94.2 g; p < 0.05). While colonized FTs were not associated with an increased risk of late onset sepsis, when infants were diagnosed with late onset sepsis, bacteria cultured from FTs were identical to the causative bacteria. In addition, organisms cultured from FTs of infants enrolled in the study were identical to those in blood cultures obtained from other infants in the NICU suggesting nosocomial spread of infection causing organisms. DNA pulse gel analysis of bacteria obtained from these FTs found the same species of MRSA as those infecting four infants residing in the same NICU but not enrolled in the study further suggesting nosocomial spread (13).

Potential Variables Affecting FT Colonization

Several studies included potential confounders for the presence of bacteria and bacterial growth in FTs including gestational and chronologic age, birthweight, and feeding regime, as well as exposure to histamine 2 (H2) blockers, probiotics, and antibiotics. Five studies (two including the same group of infants) reported gestational age and birthweight with mean gestational age and birthweight ranging from 27.2 to 32.3 weeks and 1,083 to 1,965 g, respectively (2, 12–15). However, only one included either as covariates finding greater microbial diversity in FTs from infants with more advanced gestational ages (15).

Three studies examined the effect of chronologic age on the presence of bacteria in FTs (9, 14, 15). Higher chronologic age was associated with greater amounts of Enterobacteriaceae (9),
TABLE 1 | Summary of included studies.

| Study            | Study design/Purpose                                                                 | Setting and participants                                                                 | Methodology                                                                 | Results                                                                                                                                 |
|------------------|---------------------------------------------------------------------------------------|------------------------------------------------------------------------------------------|----------------------------------------------------------------------------|----------------------------------------------------------------------------------------------------------------------------------------|
| Mehali et al.,   | Clinically-based, Prospective observational. Purpose: (1) to determine the           | Fifty infants with a mean gestational age of 28.5 weeks and mean birthweight of 1,387 g. | The lumen of 125 feeding tubes analyzed using standard culture techniques.  | Seventy-one (57%) of tubes were contaminated and only 8 (6%) were completely culture negative. The mean bacterial load was 908,173 CFU/cm.  |
| 2002 (2)         | determine the incidence of feeding tube contamination and (2) to determine           |                                                                                           | Contamination was defined as >1,000 CFU/cm excluding Staph epidermidis and    | Feeding tubes from infants treated with histamine type 2 blockers were more likely to be contaminated (85.4 vs. 48.5%; p < 0.05).           |
|                  | complications associated with feeding tube contamination.                           |                                                                                           | alpha Streptococcus.                                                       | There was no association between antibiotic therapy and feeding tube contamination.                                                   |
|                  |                                                                                       |                                                                                           | Culture results were compared with clinical cultures taken from infants in    | Feeding intolerance only occurred in infants fed formula through contaminated tubes. In infants fed formula, feeding intolerance occurred more |
|                  |                                                                                       |                                                                                           | the NICU who were not enrolled in the study who were diagnosed with late onset | often when feeding tubes were contaminated (75 vs. 0%; p < 0.05).                                                                |
|                  |                                                                                       |                                                                                           | sepsis using antibiotics susceptibility patterns and DNA electrophoresis.     | Ten infants had feeding tubes contaminated with >100,000 CFUs of Gram-negative  |
| Kim et al.,      | Laboratory-based. Purpose: (1) To examine the presence of nosocomial antibiotic-     | Hundred and twenty-five feeding tubes obtained from 50 infants previously analyzed by | Susceptibility to methicillin and vancomycin was determined on isolates of    | Feeding tube contamination was not associated with late onset sepsis. Organisms cultured from feeding tubes were identical to those          |
| 2006 (13)        | resistant pathogenic bacteria in feeding tubes. (2) To determine whether bacteria in | Mehali et al. (13).                                                                      | Staphylococcus aureus and Enterococcus faecalis using antibiotic susceptibility | subsequently causing infection both in infants who had the feeding tube and other  |
|                  | feedings tubes can cause infections in infants in the same NICU as infants with      |                                                                                            | patterns and DNA pulse gel electrophoresis.                                | infants in the NICU. DNA analysis indicated that the MRSA species was present in |
|                  | contaminated feeding tubes.                                                          |                                                                                            | Bacteria isolated from feeding tubes and from clinical cultures of infected   | at least one of the cultured feeding tubes suggesting nosocomial transmission. There was no vancomycin resistant Enterococcus.         |
|                  |                                                                                       |                                                                                            | infants were compared.                                                      | Twenty-three feeding tubes contained S aureus isolates, 12 of which were      |
| Presti and       | Laboratory-based. Purpose: To determine the effect of temperature and nutrients on  | Number of feeding tubes analyzed was not reported.                                      |                                                                                           | methicillin resistant. Four infants who did not have feeding tubes were        |
| Snyder, 2013     | biofilm formation by Enterobacter sakazkii in infant feeding tubes.                   |                                                                                            |                                                                                           | diagnosed with MRSA infection in the NICU. DNA analysis indicated that the    |
| (17)             |                                                                                       |                                                                                            |                                                                                           | MRSA species was present in at least one of the cultured feeding tubes      |
|                  |                                                                                       |                                                                                            |                                                                                           | suggesting nosocomial transmission.                                            |
|                  |                                                                                       |                                                                                            |                                                                                           | Entero bacter sakazkii attached to feeding tubes and as the biofilm aged, the |
|                  |                                                                                       |                                                                                            |                                                                                           | cells detached. Biofilm formation did not occur at 12°C and formed at 25°C   |
|                  |                                                                                       |                                                                                            |                                                                                           | only when immersed in formula in which it grew to 1.16–1.31 log CFU/cm².     |
|                  |                                                                                       |                                                                                            |                                                                                           | The most overall bacterial growth was seen at 3 h. Bacterial growth was       |
|                  |                                                                                       |                                                                                            |                                                                                           | significantly different based upon the design of the feeding tube cap at 3 (p= 0.04) and 24 h (p = 0.001) but not at 72 h.             |
|                  |                                                                                       |                                                                                            |                                                                                           | No difference in bacterial growth based on design of the distal end of the   |
|                  |                                                                                       |                                                                                            |                                                                                           | feeding tube or whether tubes were made of polyurethane or silicone.         |
|                  |                                                                                       |                                                                                            |                                                                                           | Bacteria were present in the hub, cap, and distal portion at all-time points  |
|                  |                                                                                       |                                                                                            |                                                                                           | but amounts were not reported.                                               |

(Continued)
| Study               | Study design/Purpose                                                                 | Setting and participants | Methodology                                                                 | Results                                                                                                                                 |
|--------------------|-------------------------------------------------------------------------------------|---------------------------|------------------------------------------------------------------------------|----------------------------------------------------------------------------------------------------------------------------------------|
| Hurrell et al.,    | Laboratory-based study, Purpose: To determine adherence to and growth of Enterobacteriaceae in feeding tubes. | Number of feeding tubes analyzed was not reported. | After being flushed with formula inoculated with Enterobacteriaceae, feeding tubes were then flushed with sterile formula. The feeding tube was examined for the presence of biofilm and viable counts reported. Formula flushed through the tubes was examined for bacteria using standard culturing techniques. | After 24 h, biofilm formation by Enterobacteriaceae was $10^{5}$-$10^{6}$ CFU/cm. Bacteria in the tube lumen grew to cell densities of $10^{5}$ CFU/cm within 8 h and $10^{3}$ CFU/cm within 24 h. Bacteria in formula flushed through the feeding tubes grew to $10^{5}$-$10^{9}$ CFU/ml within 24 h. |
| 2009 (8)           |                                                                                     |                           |                                                                              |                                                                                                                                          |
| Hurrell et al.,    | Clinically-based. Prospective observational. Purpose: To determine whether feeding tubes are colonized by Enterobacteriaceae and whether colonization is influenced by feeding regime. | Hundred and twenty-nine feeding tubes from 30 infants <1 week to >4 weeks of age collected from two NICUs. | The presence of Enterobacteriaceae in feeding tubes and residual liquid was determined using both standard culture and molecular methods and their antibiograms determined. | Enterobacteriaceae was isolated in 76% of feeding tubes including 52% of tubes from infants fed mother’s own milk (MOM); and 78–88% of tubes from infants receiving either a combination of MOM and formula or formula. Analysis of 104 tubes obtained from one NICU found feeding regime was significantly associated with Enterobacteriaceae colonization ($p < 0.0001$) with tubes from infants fed MOM or who were nil by mouth having the lowest bacterial counts. The most common isolates were Enterobacter hormaechei, Escherichia coli, and Klebsiella pneumonia. Residual fluid contained a peak bacterial count of $10^{7}$ CFU/ml and contained the same Enterobacteriaceae species as in the feeding tube. Chronological age associated with the number of bacteria with an increase seen after 2 weeks ($p < 0.001$). A dwell time <6 h was associated with lower bacterial counts ($p < 0.001$). Resistance to amoxicillin and third generation cephalosporins was found on antibiogram. Same strains of Escherichia coli were isolated from the residual liquid and the biofilm in 66% of feeding tubes which clustered into five pulsotypes. |
| 2009 (9)           |                                                                                     |                           |                                                                              |                                                                                                                                          |
| Alkeskas et al.,   | Laboratory-based. Purpose: To determine the diversity of Escherichia coli strains in previously isolated residual liquid from feeding tubes. | Sample included 30 strains of Escherichia coli isolated from residual liquid and biofilm obtained from 129 feeding tubes from 30 infants from two NICUs. Samples were collected in a previous prospective study (Hurrell et al., 2009a). | Strains of Escherichia coli were genotyped using pulsed-field gel electrophoresis, and seven-loci multilocus sequence typing. Potential pathogenicity of 30 virulence factors was determined by PCR-based assays and genome analysis. |                                                                                                                                          |
| 2015 (16)          |                                                                                     |                           |                                                                              |                                                                                                                                          |
| Petersen et al.,   | Clinically-based. Prospective observational. Purpose: To determine whether changing feeding tubes more frequently decreases contamination. | Ninety-four feeding tubes from 34 infants with a median gestational age of 30.1 weeks and median birthweight of 1,083 g. Median age of infant at collection was 37 days (1–119). | Saline flushed through feeding tube was analyzed using standard culture techniques. Potentially pathogenic bacteria were defined as either Gram-negative rods or Staphylococcus aureus. Biofilm within the feeding tube was visualized using scanning electron microscopy. | Eighty-nine percent of samples contained $>1,000$ CFU/ml of bacteria; 55% contained either Enterobacteriaceae or Staphylococcus aureus. Mean bacterial load was $5.3 \log_{10}$ CFU/ml and maximum was $9.4 \log_{10}$ CFU/ml. Median dwell time was 3.25 days (8 h–14.2 days). Neither the presence of bacteria (median 3.4 days if present vs. 3.2 days if not; $p = 0.18$) or presence of potentially pathogenic bacteria was associated with dwell time (median 3.7 days compared to 3.2 days; $p = 0.54$). Chronologic age was correlated with the presence of potentially pathogenic bacteria ($p = 0.036$). No difference in colonization of |
| 2016 (14)          |                                                                                     |                           |                                                                              |                                                                                                                                          |

(Continued)
TABLE 1 | Continued

| Study                  | Study design/Purpose                          | Setting and participants                                                                 | Methodology                                                                 | Results                                                                                                                                 |
|------------------------|----------------------------------------------|-------------------------------------------------------------------------------------------|------------------------------------------------------------------------------|----------------------------------------------------------------------------------------------------------------------------------------|
| Gomez et al., 2016 (12) | Clinically-based. Prospective observational.  | 135 samples from 26 infants <32 weeks or ≤1,200 g. Samples included the feeding system before entering the feeding tube. | Culture-based techniques were used to analyze samples. Scanning electron microscopy was used to visualize the biofilm present in six feeding tubes, connectors, and external feeding tubes from six infants. | bacteria (p = 0.33) or the presence of potentially pathogenic bacteria (p = 0.077) based on antibiotic exposure. Use of probiotics did not increase the risk of contamination. A dense biofilm was observed using scanning electron microscopy in the inner surface of the feeding tube. Staphylococcus was present in 93% of MOM samples compared to 37% of DHM and 11% of formula feedings. Enterococcus was the most common Gram-positive bacteria found in DHM (49%) and formula (27%). Significantly more Enterococcus (p = 0.004); Staphylococcus (p < 0.001); and Serratia (p = 0.05) was found in MOM samples compared to other feeding types. At a species level, Staphylococcus epidermidis was the most abundant organism in MOM, Enterococcus faecalis was the most abundant in DHM and formula. Observed dense bacterial biofilms in external feeding tubes, feeding tubes, and connectors. The biofilm was particularly complex when the dwell time was >48 h and none was present in feeding tube with dwell times <12 h. Alpha diversity increased with gestational age, day of life, tube dwell time, and human milk feedings. Eight phyla were detected including Actinobacteria, Bacteroidetes, Cyanobacteria, Firmicutes, Fusobacteria, Proteobacteria, Tenericutes, and Thermi. Four families had median relative abundance greater than zero including Enterococcaceae, Streptococcaceae, Staphylococcaceae, and Enterobacteriaceae. ARGs differed based upon feeding regime but not between tube sections. |
| Taft et al., 2019 (15)  | Clinically-based. Prospective observational.  | Ninety-seven feeding tubes from 47 infants with a mean gestational age of 32.3 weeks and a mean birthweight of 1,965 g. | Used 16S rRNA sequencing to characterize the bacterial composition of biofilms in different parts of feeding tubes (gastric, esophageal, pharyngeal) and to characterize bacteria in residual fluid. Whole metagenomics sequencing used to characterize antibiotic resistant genes (ARGs) present in a subset of feeding tubes from six infants. | >1,000 CFU/cm of bacteria compared to FTs obtained from infants who did not receive H2 blockers (85.4 vs. 48.5%; p < 0.05) while Taft et al. found no differences in alpha diversity when infants received H2 blockers (15). Two studies included information regarding probiotics (14, 15). Peterson et al. found no significant effect of probiotic use on the presence of bacteria in normal saline flushed through FTs (14) and while an additional study included information regarding whether or not infants received probiotics, they did not include probiotic exposure in their analysis (15). Antibiotic exposure was addressed in three studies and found to not affect the presence of >1,000 CFU/cm of bacteria (2), the presence of any bacteria, whether potentially pathogenic bacteria (defined as either Gram-negative rods or S. aureus) was present in saline flushed through FTs (14), or microbial diversity (15). However, definitions of exposure differed among studies and included exposure after birth (15), during the week tubes were removed (2), and while the tube was in place (14). |

an increased risk of colonization with potentially pathogenic organisms (14), and higher microbial diversity (15). However, the GI microbiome changes significantly following birth and differences may simply reflect the developing microbiome (18).

Feeding regime was included in three studies (9, 12, 15). Hurrel et al., found type of feeding was associated with the amount of Enterobacteriaceae (p < 0.0001) with infants receiving exclusive MOM feedings or nil by mouth having FTs with the lowest bacterial counts (9). While bacteria were found in all types of feeds, after passing through the external feeding system, compared to DHM or formula, MOM contained more Enterococcus (p = 0.004), Staphylococcus (p < 0.001), and Serratia (p = 0.05) (12). Finally, higher microbial diversity was found in FTs from infants fed either MOM or DHM (15).

Two studies reported whether infants received H2 blockers (2, 15) which could alter the microbial content of FTs by decreasing gastric acidity. Mehall et al. (2) found FTs from infants who received H2 blockers had an increased likelihood of containing
DISCUSSION

In this review, we found evidence that neonatal FTs contain bacteria which are potentially pathogenic and antibiotic resistant. FT colonization may allow bacteria to enter the infant's GI tract, potentially increasing the risk of complications including sepsis and NEC. Although external devices such as central venous lines and urinary catheters are known to increase morbidity, with risk increasing with longer dwell times, little attention has been paid to FTs as a possible risk to neonatal health (14). While the etiology of FT colonization is unknown, possible sources include bacteria present on the hands of parents and health care workers or in the infant's nasal or pharyngeal cavities (19). Furthermore, gastric bacteria may enter the FT during gastric residual evaluation or with gastroesophageal reflux (20). Because gastric pH is elevated in premature infants, gastric acidity may not impede microbial growth, potentially increasing the likelihood of FT colonization (21). Therefore, FT colonization is an important area of investigation in the care of infants in the NICU which has received little attention and for which further study is indicated.

Although evidence is limited, longer dwell times may be associated with higher levels of bacterial colonization. However, the optimal dwell time in neonates is unknown and while dwell times <12 h may limit bacterial growth, they are not clinically feasible due to workload and infant discomfort. The presence of bacteria for longer periods of time in a rich environment could enhance development of pathogens and horizontal gene transference, maximizing infant risk. A well-designed randomized controlled trial comparing more clinically appropriate dwell times is necessary to elucidate how often to change FTs and the effect of dwell time on infant health.

Inconsistencies between studies made comparisons challenging and included limiting analysis to specific families and genera of bacteria, analyzing different FT elements, and variations in type of analysis. In addition, inclusion of potentially confounding variables including gestational and chronologic age, birth weight, feeding regime, and exposure to medications/supplements were not consistent across studies potentially affecting results. Because human milk contains antimicrobial components and MOM contains an abundant and diverse microbiome (22), feeding both DHM and MOM likely affects the microbial contents of FTs (23, 24). However, no study analyzed bacteria in the feeding prior to infusion. Because higher gastric pH promotes bacterial growth, it is possible that H2 blocker exposure increases FT colonization yet was included as a covariate in only one study (2). Probiotics contain live bacteria and are increasingly administered to infants in the NICU, yet only two studies included probiotic exposure (14, 15). Finally, antibiotic exposure was only included in three studies, all of which differed in exposure definition (2, 14, 15). Future research which includes potential confounding variables is needed to elucidate their potential effect on FT colonization.

Because contaminated feedings are known to cause sepsis, meningitis, and NEC and bacteria known to cause these complications have been found in neonatal FTs (1, 9, 25), colonized FTs may increase the risk of infection. Furthermore, the presence of FTs is a known risk factor for sepsis (26, 27), and previous reports of identical strains of bacteria in the GI tract, blood, and FT of infants with late onset sepsis suggests FT colonization may increase the risk of sepsis in infants (2, 16). Finally, results of the study by Mehall et al. suggest FT colonization with pathogenic bacteria may be associated with NEC, feeding intolerance, and poor growth and that bacteria present in FTs may spread nosocomially (2, 13). Further research is necessary to determine the potential effect of FT colonization on neonatal health including whether the degree of colonization or type of bacteria influences risk of complications.

LIMITATIONS OF INCLUDED STUDIES

Selected studies are limited by small sample sizes, analysis of different FT elements, and different analysis methods. Lack of control for confounding factors could also have affected results. Finally, laboratory-based studies may not be representative of clinical settings and can’t be used to make clinical practice recommendations.

CONCLUSION

While this review found evidence that FT colonization may occur in infants in the NICU, information regarding potential effects on infant health is limited. Furthermore, while the degree of colonization, amount of biofilm formation, and number of Gram-negative bacteria present in FTs may be proportional to dwell time, insufficient evidence exists to inform clinical practice. Adequately powered studies using appropriate methodology and which consider relevant confounding variables are needed to fully understand to what extent FTs in the NICU contain bacteria, what level of colonization is considered safe and acceptable, type of bacteria which may be harmful, as well as factors which may increase or reduce the number of bacteria in FTs.

AUTHOR CONTRIBUTIONS

LP, MM, KD-S, MT, GL, and JN designed research. LP, MM, and KD-S conducted research and wrote the paper. LP had primary responsibility for final content. All authors read and approved the final manuscript and agree to be accountable for the content of the work.

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SUPPLEMENTARY MATERIAL

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fnut.2022.775014/full#supplementary-material
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