



Transitioning from gavage to full oral feeds in premature infants: When should we discontinue the nasogastric tube?

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Abstract

Background The optimal timing for discontinuation of nasogastric (NG) tube in premature infants transitioning to oral feeding is not known.

Objective To determine whether early removal of NG-tube is appropriate in low-risk premature infants.

Methods Prospectively collected data of premature infants started on oral feeds at ≤ 34 weeks gestation were reviewed. Infants were categorized into ‘early’ or ‘late’ NG-removal groups based on the proportion of oral intake in the preceding 2-days, i.e., 60–79% or 80–100% of the total volume, respectively.

Results In total 50 infants in early group vs. 43 in late group. Both groups had similar oral intake and weight change in the subsequent 2-days post-NG removal. The days from NG-removal to target oral volume, and to hospital discharge trended shorter in early vs. late group.

Conclusions Discontinuing NG-tube when the oral feeding competency reaches $\sim 75\%$ of prescribed feeding volume is safe and appropriate in low-risk premature infants.

Introduction

Successful transition from full gavage to independent oral feedings is a criterion of hospital discharge in premature infants [1]. Delay in achieving full oral feeding milestone is one of the main reasons for delays in hospital discharge [2–4], and adverse neurodevelopmental outcome in premature infants [5]. During this period of functional immaturity of coordinating sucking, swallowing, and breathing [6], many premature infants require enteral feeding, usually by a nasogastric (NG) tube, until they achieve adequate oral feeding skills. The optimal timing for discontinuation of NG tube feeding in premature infants transitioning to oral feeding is not known, and thus associated with practice variation.

Some of the cue-based oral feeding protocols suggest to keep the NG feeding tube until the infant have appropriate weight gain for 48 hours without supplementation by NG tube [7–9]. However, having a NG tube is not without problems. In presence of NG tube especially with size $>5F$, premature infants have increased nasal airway resistance [10, 11], and reduced tidal volume and minute ventilation during sucking, which can affect their oral feeding performance [12]. Its presence can potentially increase the gastroesophageal reflux (GER) events depending on the size of NG tube [13, 14]. Use of NG tube is also associated with significant nutrient loss, especially fat, fat soluble vitamins, and minerals because fat adheres to tubing surfaces [15], and is a potential source of infection as it is easily colonized by pathogenic bacteria/fungi [16].

If earlier removal of NG tube is safe and feasible, it can potentially reduce these complications and length of hospital stay in premature infants. Our study objective was to determine whether early removal of NG tube is safe and appropriate in relatively low-risk premature infants transitioning to oral feeding. We hypothesize that earlier removal of NG tube would lead to earlier transition to independent oral feeding resulting in decreased length of hospital stay.

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Methods

The study was conducted at the all-referral level IV neonatal intensive care unit (NICU) at Nationwide Children's Hospital, Columbus, Ohio. As part of an ongoing quality improvement (SIMPLE feeding initiatives) [17], focused on prevention and early identification of feeding related problems in premature infants, data of all infants admitted at ≤ 34 weeks' post-menstrual age (PMA) were collected prospectively. NG tube sizes 6.5F or 8.0F was used for enteral feeding and feedings were typically given intermittently over 10–30min every 3 h. There was no unit guideline regarding the timing of NG tube removal during the study period from Jan 2011 to Dec 2017. In general, the decision to remove NG tube occurs when the oral feeding volume reaches 60–100% of prescribed enteral feeding depending on the individual provider comfort level.

To meet the objective of this study, SIMPLE feeding program database was reviewed for 'low-risk' premature infants which we defined as infants born ≤ 32 weeks' gestation and had started on cue based oral feeds at ≤ 34 weeks' PMA. Infants on invasive respiratory support ($>2L$ of nasal cannula support, CPAP or ventilator support), gastrointestinal surgical conditions, intraventricular hemorrhage $>$ grade 2, and major congenital anomalies were excluded. Among eligible infants, the day of NG tube removal was recorded and the percentage of oral feeding volume (OFV) in the preceding two days of NG tube removal was calculated from the electronic medical records. Infants were categorized into 'early' and 'late' NG removal groups. In early group, NG tube was removed when the mean OFV was between 60–79% of prescribed feeding volume in the preceding 2 days, while in late group, it occurred at 80–100%. Infants with mean OFV less than 60% in the preceding 2 days of NG removal were excluded considering their relatively quicker progression and more mature oral feeding skills. For both groups, characteristics before and at NG removal were collected from the SIMPLE feeding program database and electronic medical records. The clinical outcomes including time from NG removal to reach the target full oral feeding volume (ability to take PO of all prescribed enteral feeding volume, which generally ranges from 130 to 150 ml/kg) and to NICU discharge, and growth metrics were compared between the two groups. OFV in infants who are on direct breast feeding was considered as 150 ml/kg after the NG removal. The project was approved by the Nationwide Children's Hospital Institutional Review Board.

Statistics

Appropriate bivariate analysis was performed to identify the unadjusted differences between the cases and controls.

Student's *t*-test was used for parametric continuous variables, and the Mann–Whitney U test was used for nonparametric continuous variables. Chi-square tests and Fisher exact tests were used for categorical variables. All quantitative data were expressed as the mean \pm standard deviation (SD), or median with inter-quartile range. A $p \leq 0.05$ was considered to be statistically significant. The statistical software IBM SPSS Statistics version 23 (SPSS, Chicago, IL) was used for the statistical analysis of the data.

Results

Of the 93 eligible infants during the study period, 50 infants were in early NG removal group, the rest 43 were in late NG removal group. Early group had similar perinatal and birth characteristics, compared to the late group except that late group were exposed to more number of CPAP days (Table 1). At NG tube removal, early group had similar corrected gestational age, and body weight compared to the late group (Table 1). Early group had similar direct breast feeding rate at NG removal compared to late group (30.0 vs. 16.3%, $p = 0.12$). The mean OFV % in the preceding two days of NG removal was 72.3 ± 5.8 in the early group vs. 87.3 ± 4.0 in the late group ($p < 0.001$). The oral feeding volume and weight change in the subsequent two days after NG removal were similar in both groups (Fig. 1, Table 2). Supplemental oxygen use at initial oral feeding, at NG tube removal, at 36 weeks' gestation, and at NICU discharge were similar in both groups. Majority of the infants reached full oral target volume before discharge (90% in early vs. 88.4 in late group, $p = 0.80$). Early group showed a trend for lower number of days from the NG removal to reach oral target volume (median days of 2.0 vs 2.5, $p = 0.13$, mean days 2.4 ± 1.9 vs. 4.4 ± 5.8 , $p = 0.051$), and to NICU discharge (mean days of 10.9 vs. 13.5, $p = 0.21$), compared to late group. None required replacing NG tube in both groups, and none required tube feeding at discharge. The NICU discharge outcomes including BPD (oxygen requirement at 36 week PMA), length of hospital stay (LOHS), OFV, and discharge weight were similar in both groups (Table 2). A subgroup analysis of infants born ≤ 28 weeks and > 29 weeks also did not show differences between early and late groups (Table 3).

Discussion

The purpose of this study was to determine whether early removal of NG tube is safe and appropriate in relatively

Table 1 Characteristics before and at NG removal in early and late NG removal groups

Characteristics	Early group (n = 50)	Late Group (n = 43)	P value
Before NG tube removal			
Birth gestation (weeks)	28.9 ± 2.5	28.3 ± 2.3	0.24
Birth weight (g)	1333.4 ± 422.2	1214.7 ± 365.0	0.16
≤ 28 weeks gestation (%)	20 (40%)	24 (55.8)	0.15
Male gender (%)	28 (56.0)	55.8	0.99
Small for gestational age (%)	1 (2.0)	1 (2.3)	1.00
Large for gestational age (%)	8 (16.0)	4 (9.5)	0.37
Born by Caesarian section (%)	35 (70.0)	29 (67.4)	0.79
Maternal diabetes (%)	2 (4.0)	4 (9.3)	0.41
Intubation (%)	29 (58.0)	28 (65.1)	0.48
Intubation days*	1 (0–2)	1 (0–2)	0.49
CPAP (%)	41 (82.0)	34 (79.1)	0.72
CPAP days	11.0 ± 14.2	17.5 ± 16.5	0.05
IVH grade 1 or 2 (%)	12 (24.0)	11 (25.6)	0.86
PMA at initial oral feeding	33.3 ± 0.5	33.6 ± 0.6	0.03
Human milk at initial PO (%)	36 (72.0)	24 (55.8)	0.10
Oxygen use at initial PO (%)	21 (42.0)	25 (58.1)	0.12
At NG tube removal			
PMA (weeks)	36.2 ± 2.3	36.1 ± 1.9	0.89
Weight (g)	2446.2 ± 457.9	2370.4 ± 471.8	0.44
Weight <10 th percentile (%)	7 (14)	5 (11.9)	1.00
Human milk use (%)	16 (32.0)	16 (37.2)	0.60
Direct breast feeding (%)	15 (30)	7 (16.3)	0.12
Oral feeding calories/oz	24.1 ± 1.3	24.7 ± 1.8	0.11
Oxygen use (%)	23 (46.0)	16 (37.2)	0.39
OFV % 2-days prior	68.6 ± 13.2	84.6 ± 8.6	0.00
OFV % 1-day prior	76.0 ± 5.8	90.0 ± 8.4	0.00
OFV % preceding 2-days	72.3 ± 5.8	87.3 ± 4.0	0.00

Values stated as mean ± SD or as stated otherwise. *Median with interquartile range

NG nasogastric, IVH Intraventricular hemorrhage, PMA Postmenstrual age, PO Per Oral, OFV Oral Feeding Volume

low-risk premature infants transitioning to oral feeding. By using a prospectively collected clinical database of premature infants enrolled in the SIMPLE feeding program at our all-referral NICU, our study suggests that discontinuing NG tube when the oral feeding competency reaches ~75% of prescribed enteral feeding volume in the preceding two days is safe and appropriate in low-risk premature infants. The practice of keeping the NG tube until these infants reach their full targeted oral feeding

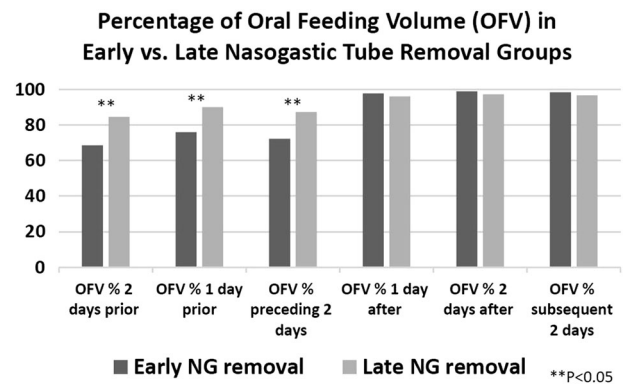


Fig. 1 Percentage of oral feeding volume (OFV) in early vs. late nasogastric tube removal groups

volume is not associated with any significant clinical benefits.

In routine clinical practice, the decision to remove the NG tube is often based on the oral feeding performance of infants in the preceding days. The relationship between chronological age and functional maturity of the oral feeding skills is not always linear, and is often influenced by growth metrics and associated co-morbidities [18]. Because of the positive association between postnatal growth and neurodevelopmental outcomes [19], potential for faltering growth with early NG tube removal is often a concern among neonatal providers. However, our data of relatively stable low-risk premature infants who have made steady oral feeding progression suggest that discontinuing NG tube when the OFV reaches about ¾ of the prescribed enteral feeding volume was not associated with significant reduction in time to achieve targeted oral feeding volume or subsequent postnatal growth until NICU discharge. Even though the late group infants were exposed to more number of CPAP days suggesting more severe early lung disease, the use of supplemental oxygen need during the oral feeding period, at 36 weeks, and at hospital discharge were similar between the two groups. Similar to our study, Fucile et al. [20], reported that setbacks in oral feeding progression are least likely to occur when the oral feeding attempt frequency reaches 6–8/day vs. 1–2 or 3–5 per day. In our center, we generally consider hospital discharge, once infant is 48h off-NG tube feeding if there are no other medical or social reasons. However, infants in both our study groups were discharged from the NICU about 11–13 days after the NG tube removal. This suggests that attaining oral feeding competency is not the only reason that was delaying NICU discharge in our study population. Considering our NICU being an all-referral center, the complexity of referred infants’ initial NICU course, incidence of clinical BPD at 36 weeks PMA in about 1/3 of study infants, and other unmeasured medical and social variables may have

Table 2 Outcomes after NG tube removal and at NICU discharge in early and late NG removal groups

Outcomes	Early Group (n = 50)	Late Group (n = 43)	P value
After NG tube removal			
OFV % 1-day after	97.8 ± 7.2	96.1 ± 5.4	0.20
OFV % 2-days after	99.0 ± 9.6	97.3 ± 7.7	0.35
OFV % subsequent 2-days	98.4 ± 7.1	96.7 ± 5.9	0.22
Δ Weight 1-day after (g)	28.2 ± 34.2	30.4 ± 40.1	0.78
Δ Weight in 2-days after (g)	56.3 ± 55.8	67.1 ± 44.0	0.31
Target full OFV (ml/kg)	144.0 ± 7.8	145.3 ± 7.3	0.39
Reached target OFV (%)	45 (90.0)	38 (88.4)	0.80
Days to reach target OFV*	2.0 (1–3)	2.5 (1–5)	0.13
Days from initial PO to NG removal	19.8 ± 14.5	19.0 ± 13.6	0.80
At NICU Discharge			
BPD (at 36 weeks PMA, %)	15 (30.0)	19 (44.2)	0.16
OFV at discharge	151.7 ± 14.7	154.1 ± 19.3	0.49
Days from NG removal to discharge	10.9 ± 7.1	13.5 ± 11.6	0.21
Length of NICU stay (days)	64.4 ± 28.6	63.2 ± 34.9	0.86
Discharge weight (g)	2715.6 ± 529.4	2742.0 ± 564.5	0.82
PMA at discharge (weeks)	37.7 ± 2.7	38.0 ± 2.5	0.58
Discharged on oxygen (%)	11 (22.0)	17 (39.5)	0.07

Values stated as mean ± SD or as stated otherwise. *Median with interquartile range

NG nasogastric, PMA Postmenstrual age, PO Per Oral, OFV Oral Feeding Volume, BPD Bronchopulmonary dysplasia

Table 3 Subgroup analysis stratified by GA (GA ≤ 28 vs. >29 weeks) of characteristics in the early and late NG removal groups

Characteristics	≤28 weeks		P value	>29 weeks		P value
	Early Group (n = 20)	Late Group (n = 24)		Early Group (n = 30)	Late Group (n = 19)	
Birth gestation (weeks)	26.1 ± 1.4	26.6 ± 1.6	0.26	30.7 ± 0.9	30.4 ± 0.9	0.20
PMA at initial PO (weeks)	33.5 ± 0.4	33.7 ± 0.6	0.19	33.2 ± 0.5	33.4 ± 0.5	0.26
PMA at NG removal (weeks)	37.4 ± 2.8	36.8 ± 2.2	0.42	35.3 ± 1.5	35.2 ± 0.7	0.72
OFV % 2-days prior	69.9 ± 16.5	84.3 ± 9.2	.002	67.6 ± 10.7	85.0 ± 7.9	<0.001
OFV % 1-day prior	74.4 ± 12.4	91.2 ± 8.7	<0.001	77.1 ± 10.2	88.6 ± 7.9	<0.001
OFV % preceding 2-days	72.2 ± 5.6	87.7 ± 4.3	<0.001	72.4 ± 6.0	86.8 ± 3.7	<0.001
OFV % 1-day after	96.6 ± 8.4	95.2 ± 5.9	0.53	98.6 ± 6.2	97.1 ± 4.6	0.38
OFV % 2-days after	99.4 ± 7.8	96.8 ± 8.4	0.59	98.8 ± 10.7	98.0 ± 7.0	0.77
OFV % subsequent 2-days	97.9 ± 6.4	96.0 ± 6.4	0.32	98.7 ± 7.7	97.6 ± 5.3	0.58
Δ Weight 1-day after (g)	32.3 ± 38.3	35.5 ± 40.1	0.79	25.5 ± 31.5	23.6 ± 40.2	0.06
Δ Weight in 2-days after (g)	51.8 ± 67.1	71.1 ± 45.3	0.26	59.3 ± 47.8	61.7 ± 42.9	0.07
Target full OFV (ml/kg/day)	142.0 ± 8.3	143.1 ± 8.1	0.62	145.3 ± 7.3	148.2 ± 5.1	0.12
Days to reach target OFV*	1.5 (1.0–2.0)	4.0 (1.0–6.3)	0.06	2.0 (1.0–3.3)	2.0 (1.0–3.5)	0.78
BPD (at 36 weeks PMA, %)	12 (60.0)	13 (54.2)	0.70	3 (10.0)	6 (31.6)	0.07
OFV at discharge (ml/kg/day)	148.6 ± 13.6	156.9 ± 22.3	0.13	153.7 ± 15.3	150.5 ± 14.3	0.46
Days from NG removal to discharge	14.1 ± 8.8	14.5 ± 9.9	0.89	8.8 ± 4.9	12.2 ± 13.5	0.31
PMA at discharge (weeks)	39.5 ± 3.0	38.9 ± 2.4	0.49	36.6 ± 1.6	36.9 ± 2.2	0.51

Values stated as mean ± SD or as stated otherwise. *Median with interquartile range

NG nasogastric, PMA Postmenstrual age, PO Per Oral, OFV Oral Feeding Volume, BPD Bronchopulmonary dysplasia

affected the timing of NICU discharge. However, considering our specific inclusion and exclusion criteria and that all our study patients were started on oral feeding ≤ 34 weeks' PMA suggest that our study cohort represents the relatively low-risk premature infants in a contemporary academic NICU hospital setting. Though the described approach and conclusions are limited to the select inclusions and exclusions of our study population, we believe, these findings may be relevant to most level-II NICUs or step-down units where the majority of such low-risk premature infants are admitted.

This study was observational single-center study; consist of a heterogeneous group of relatively stable premature infants born at different gestational ages, but appropriate PMA for oral feeding initiation and maintenance. The primary outcome of the study (Days from NG removal to target full oral feeds, median days of 2.0 vs. 2.5, $p = 0.13$, mean days 2.4 ± 1.9 vs. 4.4 ± 5.8 , $p = 0.051$) showed a trend for lower number of days in the early group, compared to the late group. Assuming parametric distribution of the variable, the difference between the groups has a small to medium effect size of 0.46. With this effect size, a statistically significant difference in days from NG removal to target full oral feeds between the two groups can be obtained from a sample size of 148 (74 in each group) with 80% power ($\alpha = 0.05$, $\beta = 0.20$). The decision to remove NG tube was not based on a standardized unit practice, but was based on the individual provider discretion, which can be a limitation in that it adds to the subjectivity in approaches. Variability in individual views among a large number of NICU providers potentially may have influenced the study group determination and the outcomes observed. Generalization of these findings to premature infants with significant comorbidities or anomalies must be applied with caution, as our study participants did not include such patients.

In summary, lack of consensus regarding the timing of NG tube removal in relation to oral feeding competency in premature infants can be associated with practice variation among individual providers. Our study suggests that discontinuing NG tube when the oral feeding competency reaches $\sim 75\%$ of prescribed enteral feeding volume in the preceding two days is safe and appropriate in stable low-risk premature infants. Keeping NG tube longer is not associated with any additional clinical benefits. As there is no harm in discontinuing NG tube at $\sim 75\%$ oral feeding competency, as evidenced by similarities in growth and LOHS, at least in monitored clinical settings, this practice can be safe and may have potential benefits in lowering resource utilization, potentially reduce the length of hospitalization, increase opportunities for parent participation, all resulting in cutting down the healthcare costs.

Compliance with ethical standards

Conflict of interest The authors declare that they have no conflict of interest.

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