



# Gastrostomy Tube Feeding in Extremely Low Birthweight Infants: Frequency, Associated Comorbidities, and Long-term Outcomes

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**Objective** To assess the frequency of gastrostomy tube (GT) placement in extremely low birth weight (ELBW) infants, associated comorbidities, and long-term outcomes.

**Study design** Analysis of ELBW infants from 25 centers enrolled in the National Institute of Child Health and Human Development Neonatal Research Network's Generic Database and Follow-up Registry from 2006 to 2012. Frequency of GT placement before 18-22 months, demographic and medical factors associated with GT placement, and associated long-term outcomes at 18-22 months of corrected age were described. Associations between GT placement and neonatal morbidities and long-term outcomes were assessed with logistic regression after adjustment for center and common co-variables.

**Results** Of the 4549 ELBW infants included in these analyses, 333 (7.3%) underwent GT placement; 76% had the GT placed postdischarge. Of infants with GTs, 11% had birth weights small for gestational age, 77% had bronchopulmonary dysplasia, and 29% severe intraventricular hemorrhage or periventricular leukomalacia. At follow-up, 56% of infants with a GT had weight <10th percentile, 61% had neurodevelopmental impairment (NDI), and 55% had chronic breathing problems. After adjustment, small for gestational age, bronchopulmonary dysplasia, intraventricular hemorrhage/periventricular leukomalacia, poor growth, and NDI were associated with GT placement. Thirty-two percent of infants with GTs placed were taking full oral feeds at follow-up.

**Conclusions** GT placement is common in ELBW infants, particularly among those with severe neonatal morbidities. GT placement in this population was associated with poor growth, NDI, and chronic respiratory and feeding problems at follow-up. The frequency of GT placement postneonatal discharge indicates the need for close nutritional follow-up of ELBW infants. (*J Pediatr* 2019;214:41-6).

**Trial registration** [ClinicalTrials.gov](https://www.clinicaltrials.gov/ct2/show/study/NCT00063063): NCT00063063.

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Oral feeding difficulties are common among extremely low birth weight (ELBW) infants. Coordination of suck-swallow-breathe patterns are critical to feeding success, but these skills typically evolve and mature starting at 33-34 weeks of gestation.<sup>1-4</sup> This process can be delayed in those who require invasive medical interventions (endotracheal intubation and long-term nasogastric [NG] feedings).<sup>5,6</sup> ELBW infants with poor oral feeding skills have prolonged hospital stays, incur increased healthcare costs, and may be at increased risk for long-term deficits in feeding skills and neurodevelopment.<sup>7,8</sup>

ELBW infants with bronchopulmonary dysplasia (BPD) or severe neurologic injury are especially prone to ongoing feeding difficulties.<sup>9,10</sup> Infants with severe BPD have been found to have poorer coordination when feeding, poor endurance, and an inability to generate sucking pressures strong enough for successful oral feeding.<sup>11,12</sup> At follow-up visits, lower scores on neurodevelopmental testing in the cognitive and language domains have been demonstrated in ELBW infants with dysfunctional feeding behaviors.<sup>13</sup>

BPD	Bronchopulmonary dysplasia	NICHD	National Institute of Child Health and Human Development
ELBW	Extremely low birth weight		
GT	Gastrostomy tube	NJ	Nasogastric
IVH	Intraventricular hemorrhage	NRN	Neonatal Research Network
NEC	Necrotizing enterocolitis	PVL	Periventricular leukomalacia
NG	Nasogastric	SGA	Small for gestational age
NICU	Neonatal intensive care unit		

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Oral feeding difficulties often necessitate alternative feeding options before or after neonatal intensive care unit (NICU) discharge. There is some evidence in the literature supporting early discharge with home gavage feeding in stable premature infants who have difficulties establishing full oral feeds.<sup>14-16</sup> Nonetheless, the overall frequency of gastrostomy tube (GT) placement in neonates has increased with 1 study showing the rate of GTs doubling in very low birth-weight infants between 2000 and 2012.<sup>17</sup>

Although GT feeding in premature infants has been described in the literature, the frequency of use in ELBW infants and its association with other neonatal morbidities is unknown. The purpose of this study was to examine the frequency of GT placement in ELBW infants, to investigate the association with certain neonatal morbidities, and to evaluate longer-term growth and developmental outcomes.

## Methods

We performed a retrospective cohort analysis of all ELBW infants (birth weight <1000 g) who were enrolled in the National Institute of Child Health and Human Development (NICHD) Neonatal Research Network's (NRN) Generic Database (Clinicaltrials.gov: NCT00063063) between January 1, 2006 and December 31, 2012. Prior to discharge, all surgical procedures were noted. If an infant did not have a GT placed before discharge, then GT placement after discharge was determined by a question asked of the caregiver at follow-up. This was a simple yes or no question regarding whether the child had a gastrostomy tube or button placed. The timing of the follow-up evaluation changed during the research study period. Follow-up was performed between 18 and 22 months corrected age prior to July 1, 2012 and between 22 and 26 months corrected age afterward. Infants were excluded from the analysis if they had significant congenital heart disease, an upper airway or gastroenterology malformation, or a syndrome or chromosomal abnormality. Infants who developed short bowel syndrome during their NICU stay necessitating a GT were excluded as these infants often require GT management for reasons other than oral feeding difficulties. Infants dying before or after NICU discharge, those not completing the developmental follow-up visit, and those with missing outcome data at discharge and at follow-up also were excluded (Figure; available at [www.jpeds.com](http://www.jpeds.com)). Trained research personnel collected clinical data in a standardized manner.

Prevalence of GT placement prior to the follow-up visit, demographic and medical factors associated with GT placement, and associated outcomes (including respiratory, feeding, growth, and neurodevelopmental outcomes) were documented. Demographic factors included race, sex, and maternal education level. Medical factors included in the analyses were a birth weight small for gestational age (SGA), BPD, severe intraventricular hemorrhage (IVH), periventricular leukomalacia (PVL), and necrotizing enterocolitis (NEC). SGA was defined as a birth weight <10th percentile

for gestational age.<sup>18</sup> BPD was defined using the physiologic definition of infants at 36 weeks of postmenstrual age.<sup>19</sup> IVH included grade III or IV.<sup>20</sup> NEC was defined as stage IIA or greater on modified Bell staging criteria.<sup>21,22</sup> Infants diagnosed with surgical NEC leading to short bowel syndrome were excluded and not included in the primary analysis, however, information was gathered regarding the prevalence of GTs in this population. Longer-term outcomes evaluated were poor growth, neurodevelopmental impairment, "chronic feeding problems" (use of thickened feeds, abnormal swallowing, dysphagia, or documented aspiration at follow-up), and "chronic breathing problems" (use of oxygen, diuretics, or bronchodilators at follow-up). Poor growth, measured at follow-up on the National Center for Health Statistics (NCHS) growth curve, was defined as weight, length, or head circumference <10th percentile for corrected age.<sup>23</sup> Infants were evaluated between 18 and 26 months adjusted age using the Bayley Scales of Infant and Toddler Development-III and a standardized neurosensory examination. Neurodevelopmental impairment (NDI) was defined as any of the following: a cognitive composite score on the Bayley Scales of Infant and Toddler Development-III <85, moderate-to-severe cerebral palsy, gross motor function classification system level  $\geq 2$ , severe hearing impairment, or bilateral severe visual impairment.<sup>13</sup>

Descriptive statistics were calculated for baseline characteristics, medical factors, and outcomes at follow-up. Frequencies and percentages were reported for categorical variables with differences in characteristics between groups tested for by  $\chi^2$  tests or the Fisher exact test. Means, SDs, medians, and IQRs were reported for continuous variables with differences tested using the Wilcoxon test. Logistic regression models were used to assess associations between GT placement and neonatal morbidities and outcomes at follow-up. Models included 1 characteristic at a time as the primary independent variable with center, gestational age, SGA, BPD, NEC, and severe IVH/PVL as co-variables. ORs were estimated with statistical significance determined by Wald  $\chi^2$  tests. All analyses were conducted in SAS v 9.4. (SAS Institute, Cary, North Carolina).

Data were also gathered for infants discharged on NG or nasojejun (NJ) feedings. This information, however, was limited and only available from 2008 to 2011. The characteristics of infants discharged on NG/NJ vs GT tubes were obtained as well as the prevalence of those who were initially discharged with NG/NJ feeds but received a GT prior to follow-up.

## Results

A total of 4549 ELBW infants from 25 centers met the inclusion criteria for the study and were included in the analyses. Of these, 333 (7.3%) of the infants underwent GT placement. Among those with a GT, 77% had BPD (253/333), 29% (96/333) had a grade III or IV IVH or PVL, and 7% (22/333) had NEC (Table I). Variables that were found to be significantly

**Table I. Characteristics of patients with and without GT placement\***

Variables	Total N = 4549	No GT N = 4216	GT N = 333	P value
<b>Baseline characteristics</b>				
Gestational age, mean (SD)	25.73 (1.30)	25.76 (1.30)	25.43 (1.30)	<.01
SGA, n (%)	315 (7)	277 (7)	38 (11)	<.01
Male, n (%)	2245 (49)	2083 (49)	162 (49)	.86
White, n (%)	2390 (53)	2203 (53)	187 (58)	.13
Hispanic, n (%)	751 (17)	700 (17)	51 (16)	.54
High School degree, n (%)	2601 (76)	2417 (76)	184 (74)	.49
<b>Neonatal characteristics</b>				
Length of hospital stay (d), mean (SD)	112 (46)	108 (40)	169 (72)	<.01
NEC, n (%)	250 (5)	228 (5)	22 (7)	.38
BPD, n (%)	2288 (51)	2035 (49)	253 (77)	<.01
Severe IVH (grade III or IV) or PVL, n (%)	745 (16)	649 (15)	96 (29)	<.01
Weight z score at 36 wk, mean (SD)	-1.41 (0.83)	-1.39 (0.83)	-1.62 (0.87)	<.01
Length z score at 36 wk, mean (SD)	-1.94 (0.92)	-1.91 (0.92)	-2.26 (0.94)	<.01
Head circumference z score at 36 wk, mean (SD)	-1.34 (1.03)	-1.31 (1.03)	-1.71 (1.01)	<.01
<b>Outcomes at follow-up</b>				
Weight z score at follow-up, mean (SD)	-1.02 (1.32)	-0.98 (1.30)	-1.59 (1.51)	<.01
Length z score at follow-up, mean (SD)	-0.67 (1.31)	-0.62 (1.31)	-1.32 (1.18)	<.01
Head circumference z score at follow-up, mean (SD)	-0.55 (1.62)	-0.49 (1.60)	-1.32 (1.69)	<.01
Follow-up weight <10th %, n (%)	1852 (41)	1669 (40)	183 (56)	<.01
Follow-up height <10th %, n (%)	1255 (28)	1087 (26)	168 (51)	<.01
Follow-up head circumference <10th %, n (%)	1216 (27)	1063 (25)	153 (47)	<.01
Weight gain velocity <sup>†</sup> (g/mo), mean (SD)	414.53 (177.09)	416.24 (182.97)	394.78 (81.39)	<.01
Length gain velocity <sup>†</sup> (cm/mo), mean (SD)	1.88 (0.74)	1.88 (0.76)	1.82 (0.24)	<.01
Head circumference gain velocity <sup>†</sup> (cm/mo), mean (SD)	0.77 (0.30)	0.77 (0.31)	0.76 (0.15)	.31
NDI <sup>‡</sup> , n (%)	1420 (31)	1217 (29)	203 (61)	<.01
Moderate/severe cerebral palsy, n (%)	271 (6)	199 (5)	72 (22)	<.01
BSID III Cognitive <70, n (%)	424 (9)	315 (8)	109 (34)	<.01
BSID III Cognitive <85, n (%)	1300 (29)	1117 (27)	183 (56)	<.01
BSID III Language <70, n (%)	787 (18)	646 (16)	141 (44)	<.01
Chronic breathing problems, n (%)	1318 (32)	1139 (30)	179 (55)	<.01
Chronic feeding problems, n (%)	1281 (28)	1085 (26)	196 (59)	<.01

BSID, Bayley Scales of Infant and Toddler Development.

\*n (%) and mean (SD) scores were calculated based on nonmissing responses.

<sup>†</sup>Velocity is calculated from 36 weeks to follow-up.

<sup>‡</sup>NDI was defined as any of the following: a cognitive composite score on the BSID III <85, moderate-to-severe cerebral palsy, gross motor function classification system level  $\geq 2$ , severe hearing impairment, or bilateral severe visual impairment.

associated with GT placement were birth weight SGA, BPD, and IVH or PVL, and length of hospital stay after adjustment for center and neonatal morbidities. Demographic factors such as sex, race, and maternal education level were not significantly associated with GT placement. A diagnosis of NEC was not associated with GT placement in our cohort (Table II). For the 187 infants with surgical NEC, GT placement was significantly more likely in infants with short bowel syndrome (45%, 22/49) than in those without short bowel syndrome (9%, 12/138;  $P < .01$ ).

Most longer-term adverse outcomes were significantly more likely to occur in infants with GTs than in infants without GTs (Table I). At follow-up, GT placement was associated with poorer growth, NDI, cerebral palsy, and chronic breathing and feeding problems.

Thirty-two percent (108/333) of the infants with a GT were taking full oral feeds at follow-up. Hispanic ethnicity, as well as all components of NDI, and breathing status at follow-up were independently associated with the ability to attain full oral feedings by 2 years. No specific neonatal morbidities were associated with a resolved need for GT supplementation at follow-up (Table III).

Seventy-six percent (252/333) of those who underwent GT placement did so after discharge from the NICU. Significant differences between patients who had GT placed before and after discharge were slower growth of head circumference and increased chronic breathing and feeding problems among infants who underwent GT placement after discharge. There was no significant difference in weight gain velocity after discharge between these 2 groups (Table IV).

Of the 2271 ELBW infants discharged between 2008 and 2011, 4% (93/2271) were discharged with NG or NJ feeds and 13/93 (14%) of these went on to receive GTs. Infants discharged on NG feeds were more likely to subsequently have a GT placed if they had a severe IVH or poor length and head growth at follow-up and less likely to have a GT placed if male (Table V; available at [www.jpeds.com](http://www.jpeds.com)).

A fundoplication procedure was performed simultaneously with GT placement in 26% of the infants. There was wide center variation for GT placement and fundoplication rates. The rate of GT placement varied from 3% to 14% by center and fundoplication rates ranged from 0% to 6.4% among the centers (Table VI; available at [www.jpeds.com](http://www.jpeds.com)).

**Table II. Associations between patient characteristics and GT placement**

Variables	OR (95% CI)*	P value
Gestational age	0.91 (0.82, 1.00)	.06
SGA	1.92 (1.31, 2.84)	<.01
Male	0.89 (0.71, 1.13)	.34
White†	0.96 (0.75, 1.24)	.77
Hispanic	0.74 (0.51, 1.06)	.10
Maternal education (high school degree)‡	0.98 (0.71, 1.34)	.88
Length of hospital stay	1.02 (1.02, 1.02)	<.01
NEC	1.13 (0.70, 1.81)	.61
BPD	2.94 (2.20, 3.92)	<.01
Severe IVH (grade III or IV) or PVL	2.01 (1.54, 2.64)	<.01
Weight z score at 36 wk	0.65 (0.55, 0.76)	<.01
Length z score at 36 wk	0.67 (0.58, 0.78)	<.01
Head circumference z score at 36 wk	0.72 (0.64, 0.82)	<.01
Weight z score at follow-up	0.75 (0.69, 0.82)	<.01
Length z score at follow-up	0.77 (0.70, 0.84)	<.01
Head circumference z score at follow-up	0.77 (0.71, 0.83)	<.01
Follow-up weight <10th %	1.65 (1.30, 2.10)	<.01
Follow-up height <10th %	2.64 (2.07, 3.36)	<.01
Follow-up head circumference <10th %	2.02 (1.58, 2.59)	<.01
Weight gain velocity	1.00 (1.00, 1.00)	.25
Length gain velocity	0.96 (0.87, 1.05)	.36
Head circumference gain velocity	0.93 (0.71, 1.21)	.57
NDI	3.19 (2.49, 4.10)	<.01
Moderate/severe cerebral palsy	4.60 (3.25, 6.50)	<.01
BSID III Cognitive <70	4.93 (3.69, 6.58)	<.01
BSID III Cognitive <85	2.97 (2.31, 3.81)	<.01
BSID Language <70	3.53 (2.73, 4.56)	<.01
Chronic breathing problems	2.42 (1.89, 3.10)	<.01
Chronic feeding problems	4.06 (3.12, 5.28)	<.01

\*ORs were estimated using logistic regression models. Models included 1 characteristic at a time as the primary independent variable with center, gestational age, SGA status, physiological BPD, medically managed NEC, and severe IVH or PVL as covariates. Statistical significance was determined by Wald  $\chi^2$  tests.

†Reference all "non-White" other than Hispanic.

‡Reference those without high school degree.

**Table III. Demographics of those with or without GT at follow-up**

Variables	No tube feeding at follow-up n = 108	Tube feeding at follow-up n = 225	P value
SGA	12 (11)	26 (12)	.99
Male	53 (49)	109 (49)	.99
White	70 (67)	117 (53)	.02
Hispanic	29 (27)	22 (10)	<.01
High School degree	57 (72)	127 (75)	.64
Length of hospital stay	158.20 (62.83)	174.68 (75.11)	.08
NEC	9 (8)	13 (6)	.48
BPD	75 (71)	178 (80)	.07
Severe IVH (grade III or IV) or PVL	31 (29)	65 (29)	.99
Weight z score at 36 wk	-1.53 (0.86)	-1.66 (0.87)	.22
Length z score at 36 wk	-2.02 (0.88)	-2.38 (0.94)	.14
Head circumference z score at 36 wk	-1.58 (0.94)	-1.77 (1.04)	.38
Weight z score at follow-up	-1.72 (1.42)	-1.53 (1.55)	.17
Length z score at follow-up	-1.16 (1.11)	-1.40 (1.21)	.48
Head circumference z score at follow-up	-1.18 (1.57)	-1.39 (1.74)	.03
Follow-up weight <10th %	60 (57)	123 (55)	.81
Follow-up height <10th %	51 (48)	117 (53)	.48
Follow-up head circumference <10th %	43 (41)	110 (50)	.13
Weight gain velocity (g/mo)	381.73 (70.75)	401.04 (85.48)	.19
Length gain velocity (cm/mo)	1.80 (0.23)	1.84 (0.25)	.74
Head circumference gain velocity (cm/mo)	0.74 (0.14)	0.76 (0.15)	.11
NDI	47 (44)	156 (69)	<.01
Moderate/severe cerebral palsy	6 (6)	66 (29)	<.01
BSID III Cognitive <70	16 (15)	93 (43)	<.01
BSID III Cognitive <85	43 (40)	140 (65)	<.01
BSID III Language <70	27 (25)	114 (55)	<.01
Chronic breathing problems	39 (37)	140 (63)	<.01
Chronic feeding problems	40 (37)	156 (69)	<.01

## Discussion

GT placement is common among ELBW infants, especially among those with co-existing morbidities. The NICHD NRN collected information in their Generic Database from 2006-2012 on major surgeries, including GT placement before and after NICU discharge. The database provides a unique opportunity to assess the frequency of use of supplemental GT feeding in extremely preterm infants from multiple academic tertiary centers and to examine associations between this feeding strategy and long-term growth, respiratory, feeding, and developmental outcomes in extremely premature infants.

In our cohort, the majority of infants had their GTs placed after discharge (76%) suggesting that a large proportion of ELBW infants were first discharged from the NICU orally feeding but could not maintain these skills. Although there was no significant difference in weight gain velocity between the 2 groups, ELBW infants were more likely to have a GT placed after discharge if they had chronic respiratory or feeding problems. The high percentage of GT placement post-neonatal discharge indicates a need for close nutritional follow-up of ELBW infants, especially those needing respiratory support such as supplemental oxygen, bronchodilator or

diuretic therapy, as well as those with a history of dysphagia or who have risks for aspiration. In addition, oral feeding rehabilitation strategies prior to and following GT placement may be relevant in achieving better overall outcomes, both short- and long-term. Physiological basis for safe oral feeding practices that include volume tolerance and airway safety are both fundamental to successful oral feeding and prevention of need for a GT.<sup>1,24</sup> Further work is needed with standardization of oral feeding practices both in the NICU and postdischarge.

GT placement was most strongly associated with BPD followed by severe brain imaging abnormalities (IVH or PVL), then SGA. This is consistent with previous literature documenting the associations of severe BPD and feeding difficulties.<sup>9,11,12</sup> The association of significant neonatal brain injury and poor feeding skills is also not unexpected given the oromotor coordination and skills necessary to develop successful oral feeding abilities.

Longer-term outcomes associated with GT placement were assessed at follow-up. At 2 years of age, GT placement was associated with poor growth, NDI, and chronic respiratory and feeding problems. Although the reasons for failure to successfully transition to full oral feeds is poorly described in the literature, ongoing feeding difficulties recently have been described in ELBW infants from a 2006-2008 cohort,

**Table IV. Demographics of those who had GTs placed pre- and postdischarge**

Variables	GT		P value
	Predischarge n = 81	Postdischarge n = 252	
SGA	9 (11)	29 (12)	.99
Male	37 (46)	125 (50)	.53
White	47 (59)	140 (57)	.90
Hispanic	14 (18)	37 (15)	.60
High School degree	45 (79)	139 (73)	.39
Length of hospital stay (d)	163 (57)	171 (76)	.99
NEC	6 (7)	16 (6)	.80
BPD	58 (73)	195 (78)	.29
Severe IVH (grade III or IV) or PVL	20 (25)	76 (30)	.40
Weight z score at 36 wk	-1.59 (0.96)	-1.63 (0.84)	.39
Length z score at 36 wk	-2.20 (1.05)	-2.28 (0.90)	.89
Head circumference z score at 36 wk	-1.72 (1.10)	-1.70 (0.99)	.82
Weight z score at follow-up	-1.49 (1.39)	-1.62 (1.54)	.98
Length z score at follow-up	-1.24 (1.16)	-1.35 (1.19)	.30
Head circumference z score at follow-up	-1.04 (1.59)	-1.42 (1.71)	.19
Follow-up weight <10th %	44 (55)	139 (56)	.90
Follow-up height <10th %	36 (45)	132 (53)	.25
Follow-up HC <10th %	33 (41)	120 (49)	.30
Weight gain velocity (g/mo)	401.78 (83.40)	392.49 (80.77)	.38
Length gain velocity (cm/mo)	1.87 (0.26)	1.81 (0.23)	.76
Head circumference gain velocity (cm/mo)	0.79 (0.15)	0.75 (0.14)	.04
NDI	42 (52)	161 (64)	.07
Moderate/severe cerebral palsy	12 (15)	60 (24)	.09
BSID III Cognitive <70	24 (30)	85 (35)	.50
BSID III Cognitive <85	39 (49)	144 (59)	.12
BSID III Language <70	32 (40)	109 (46)	.37
Chronic breathing problems	32 (42)	147 (59)	.01
Chronic feeding problems	37 (46)	159 (63)	<.01

with 13% having dysfunctional feeding behaviors at 18-22 months of corrected age. These feeding problems were defined as tube feedings, choking or coughing with oral feeds, a history of aspiration, or difficulty swallowing.<sup>13</sup> Early preterm infants (<34 weeks) are also nearly twice as likely to have oromotor dysfunction and avoidant feeding behaviors at 3 and 12 months corrected age as compared to late preterm infants (34-37 weeks).<sup>4</sup> Rommel described 700 infants and young children referred for evaluation and treatment of a severe feeding disorder and reported an overrepresentation of premature infants, especially those born at <34 weeks of gestation or with lower birth weights for gestational age.<sup>25</sup>

Early feeding problems in premature infants may have significant consequences for growth and development. Overall, children with feeding problems are at risk for nutritional deficiencies and poor growth, regardless GT placement, as well as poorer cognitive, motor, and language outcomes than children without feeding problems. Mizuno and Ueda demonstrated an association between neonatal feeding difficulties and developmental problems at 18-month follow-up. In this study, the sensitivity and specificity of early feeding assessments were better predictors of neurodevelopmental outcomes than cranial ultrasound findings.<sup>26</sup> Adams-Chapman

et al showed an association between feeding difficulties and language delays in ELBW infants at 18- to 22-month follow-up.<sup>13</sup> Behavioral and emotional problems have also been described in children with feeding problems.<sup>27,28</sup> Parents often struggle to cope with feeding difficulties in premature infants, and feeding issues may be a primary concern for families after discharge.<sup>29,30</sup> Because of this, an increasing number of premature infants are referred for feeding therapy for both infant skill development and parent support, often continuing through school age.<sup>31</sup>

Given the risk of complications associated with surgically placing a gastrostomy tube in an ELBW infant, as well as possibly increasing length of stay, a better understanding of the duration of "oral feeding failure" in an ELBW infant at NICU discharge will guide providers in making recommendations for discharge feeding plans.<sup>32</sup> There is evidence suggesting that supplemental home nasogastric tube feeding may be a safe and effective means for treating oral feeding problems in premature infants, yet many centers do not consider this practice for discharge.<sup>14,15</sup> In a Cochrane review comparing early discharge home with gavage feeds and healthcare support with later discharge home when full oral feeds have been established, it was concluded there were not enough quality trials to make a practice recommendation.<sup>16</sup> However, several small studies suggest not only a reduced length of stay, but also a reduction in infection and improved breastfeeding in the home gavage group.<sup>15</sup> Although our database did not capture information regarding home NG feeding at NICU discharge in its entirety, the available data showed that 14% of these patients eventually received a GT. This study may provide preliminary evidence to support a clinical trial of home NG vs GT feeding in otherwise stable premature infants with oral feeding problems, which could then identify optimal discharge feeding plans for ELBW infants.

Although our study provided new information about ELBW infants who undergo GT placement, there were several limitations. Primarily, we were unable to determine the best method of feeding ELBW patients at discharge: prolonged hospitalization awaiting full oral feeding, NG tube, or GT. Optimistically, many babies discharged home with NG supplementation did not require GT placement, but we could not ascertain if complications occurred at home, if these children were readmitted to the hospital, or the time interval between discharge and GT placement. We had limited information on the infants discharged with NG feeds who did not progress to GT placement. The study was also limited by the changing criteria for age of follow-up during the study period (from 18-22 to 22-26 months adjusted age). Lastly, the method of ascertainment for GT placement after discharge depended on caregiver recall, which allows for potential bias, and it was not validated with postdischarge hospital records.

The high percentage of GT placement we observed postneonatal discharge indicates the need for close nutritional follow-up of ELBW infants. Further studies of nutritional interventions, including standardization of oral infant feeding

guidelines, requirements for and timing of GT placement, as well as safety and efficacy of home GT vs NG supplementation are needed to identify optimal discharge feeding plans for ELBW infants. ■

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## Data statement

Data sharing statement available at [www.jpeds.com](http://www.jpeds.com).

## References

- Jadcherla SR, Gupta A, Stoner E, Fernandez S, Shaker R. Pharyngeal swallowing: defining pharyngeal and upper esophageal sphincter relationships in human neonates. *J Pediatr* 2007;151:597-603.
- Jadcherla SR, Dail J, Malkar MB, McClead R, Kelleher K, Nelin L. Impact of process optimization and quality improvement measures on neonatal feeding outcomes at an all-referral neonatal intensive care unit. *J Parenteral Enteral Nutr* 2016;40:646-55.
- Mizuno K, Ueda A. The maturation and coordination of sucking, swallowing, and respiration in preterm infants. *J Pediatr* 2003;142:36-40.
- DeMauro SB, Patel PR, Medoff-Cooper B, Posencheg M, Abbasi S. Post-discharge feeding patterns in early- and late-preterm infants. *Clin Pediatr* 2011;50:957-62.
- Dodrill P, McMahan S, Ward E, Weir K, Donovan T, Riddle B. Long-term oral sensitivity and feeding skills of low-risk pre-term infants. *Early Hum Dev* 2004;76:23-37.
- Jadcherla SR, Wang M, Vijayapal AS, Leuthner SR. Impact of prematurity and co-morbidities on feeding milestones in neonates: a retrospective study. *J Perinatol* 2010;30:201.
- Lessen BS. Effect of the premature infant oral motor intervention on feeding progression and length of stay in preterm infants. *Adv Neonatal Care* 2011;11:129-39.
- Jadcherla SR, Khot T, Moore R, Malkar M, Gulati IK, Slaughter JL. Feeding methods at discharge predict long-term feeding and neurodevelopmental outcomes in preterm infants referred for gastrostomy evaluation. *J Pediatr* 2017;181:125-30.
- Jadcherla S. Dysphagia in the high-risk infant: potential factors and mechanisms. *Am J Clin Nutr* 2016;103:622S-8S.
- Jadcherla SR, Peng J, Moore R, Saavedra J, Shepherd E, et al. Impact of personalized feeding program in 100 NICU infants: pathophysiology-based approach for better outcomes. *J Pediatr Gastroenterol Nutr* 2012;54:62.
- Mizuno K, Nishida Y, Taki M, Hibino S, Murase M, Sakurai M, et al. Infants with bronchopulmonary dysplasia suckle with weak pressures to maintain breathing during feeding. *Pediatrics* 2007;120:e1035-42.
- Gewolb IH, Vice FL. Abnormalities in the coordination of respiration and swallow in preterm infants with bronchopulmonary dysplasia. *Dev Med Child Neurol* 2006;48:595-9.
- Adams-Chapman I, Bann CM, Vaucher YE, Stoll BJ. Association between feeding difficulties and language delay in preterm infants using Bayley Scales of Infant Development. *J Pediatr* 2013;3:680-5.
- Rosen D, Schneider R, Bao R, Burke P, Ceballos C, Hoffstadter-Thal K, et al. Home nasogastric feeds: feeding status and growth outcomes in a pediatric population. *J Parenteral Enteral Nutr* 2016;40:350-4.
- Meerlo-Habing ZE, Kusters-Boes EA, Klip H, Brand PL. Early discharge with tube feeding at home for preterm infants is associated with longer duration of breast feeding. *Arch Dis Child Fetal Neonatal Ed* 2009;94:F294-7.
- Collins CT, Makrides M, McPhee AJ. Early discharge with home support of gavage feeding for stable preterm infants who have not established full oral feeds. *Cochrane Database Syst Rev* 2003;CD003743.
- Hatch LD, Scott TA, Walsh WF, Goldin AB, Blakely ML, Patrick SW. National and regional trends in gastrostomy in very low birth weight infants in the USA: 2000–2012. *J Perinatol* 2018;38:1270.
- Olsen IE, Groveman SA, Lawson ML, Clark RH, Zemel BS. New intra-uterine growth curves based on United States data. *Pediatrics* 2010;125:e214-24.
- Walsh MC, Yao Q, Gettner P, Hale E, Collins M, Hensman A, et al. Impact of a physiologic definition on bronchopulmonary dysplasia rates. *Pediatrics* 2004;114:1305.
- Papile LA, Burstein J, Burstein R, Koffler H. Incidence and evolution of subependymal and intraventricular hemorrhage: a study of infants with birth weights less than 1500 gm. *J Pediatr* 1978;92:529-34.
- Bell MJ, Ternberg JL, Feigin RD, Keating JP, Marshall RI, Barton LE, et al. Neonatal necrotizing enterocolitis. Therapeutic decisions based upon clinical staging. *Ann Surg* 1978;187:1.
- Walsh MC, Kliegman RM. Necrotizing enterocolitis: treatment based on staging criteria. *Pediatr Clin N Am* 1986;33:179-201.
- Kuczumarski RJ, Ogden CL, Grummer-Strawn LM, Flegal KM, Guo SS, Wei R, et al. CDC growth charts: United States. *Adv Data* 2000;Jun:1-27.
- Kamitsuka MD, Nervik PA, Nielsen SL, Clark RH. Incidence of nasogastric and gastrostomy tube at discharge is reduced after implementing an oral feeding protocol in premature (<30 weeks) infants. *Am J Perinatol* 2017;34:606.
- Rommel N, De AM, Feenstra L, Veereman-Wauters G. The complexity of feeding problems in 700 infants and young children presenting to a tertiary care institution. *J Pediatr Gastroenterol Nutr* 2003;37:75-84.
- Mizuno K, Ueda A. Neonatal feeding performance as a predictor of neurodevelopmental outcome at 18 months. *Dev Med Child Neurol* 2005;47:299-304.
- Dahl M, Sundelin C. Early feeding problems in an affluent society: I. Categories and clinical signs. *Acta Paediatrica* 1986;75:370-9.
- Esparó G, Canals J, Jane C, Ballespi S, Vinas F, Domenech E. Feeding problems in nursery children: prevalence and psychosocial factors. *Acta Paediatrica* 2004;93:663-8.
- Kavanaugh K, Mead L, Meier P, Mangurten HH. Getting enough: mothers' concerns about breastfeeding a preterm infant after discharge. *J Obstet Gynecol Neonatal Nurs* 1995;24:23-32.
- Pridham K, Saxe R, Limbo R. Feeding issues for mothers of very low-birth-weight, premature infants through the first year. *J Perinatal Neonatal Nurs* 2004;18:161-9.
- Field D, Garland M, Williams K. Correlates of specific childhood feeding problems. *J Paediatr Child Health* 2003;39:299-304.
- Landisch RM, Colwell RC, Densmore JC. Infant gastrostomy outcomes: the cost of complications. *J Pediatr Surg* 2016;51:1976-82.

## Appendix

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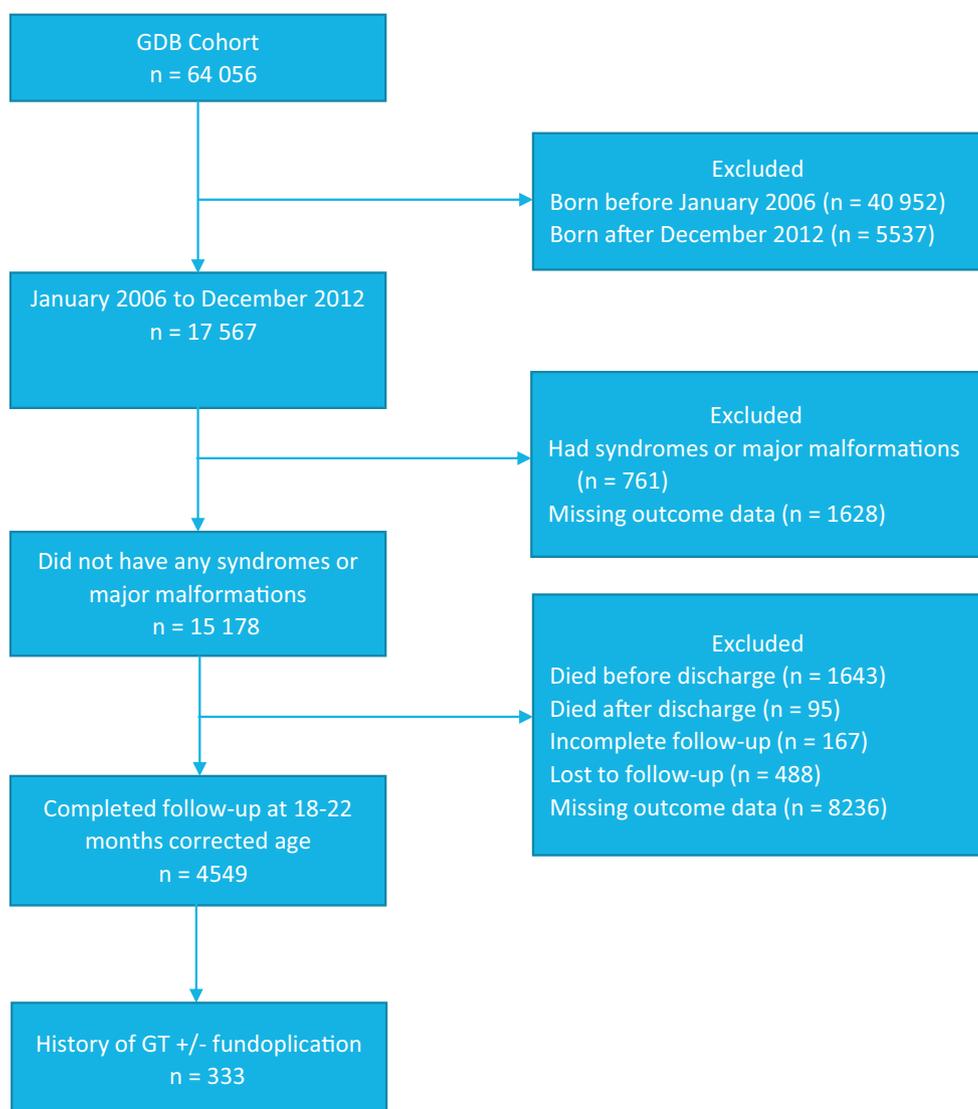
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(NRN) were transmitted to RTI International, the data coordinating center (DCC) for the network, which stored, managed and analyzed the data for this study. On behalf of the NRN, A.D. (DCC Principal Investigator) and D.K. (DCC Statistician) had full access to all the data in the study and take responsibility for the integrity of the data and accuracy of the data analysis. The authors declare no conflicts of interest.



**Figure.** Patient cohort.

**Table V. Characteristics of those discharged on NG with or without GTs placed postdischarge\***

Variables	Discharged on NG, no GT placed postdischarge n = 18	Discharged on NG, GT placed postdischarge n = 13	P value
SGA	2 (11)	2 (15)	.99
Male	12 (67)	3 (23)	.03
White	11 (61)	8 (62)	.99
Hispanic	1 (6)	2 (17)	.55
High school degree	14 (93)	5 (63)	.10
Length of hospital stay (d)	107 (9)	107 (10)	.36
NEC	2 (11)	0 (0)	.50
BPD	16 (89)	8 (62)	.10
Severe IVH (grade III or IV) or PVL	2 (11)	6 (46)	.04
Weight z score at 36 wk	-0.92 (0.41)	-1.27 (0.66)	.10
Length z score at 36 wk	-1.53 (0.64)	-1.94 (0.87)	.10
Head circumference z score at 36 wk	-0.88 (0.68)	-1.35 (0.80)	.18
Weight z score at follow-up	-0.47 (1.43)	-1.60 (1.55)	.10
Length z score at follow-up	-0.17 (0.77)	-1.15 (0.77)	<.01
Head circumference z score at follow-up	-0.07 (1.77)	-0.51 (1.11)	.03
Follow-up weight <10th %	5 (28)	7 (54)	.26
Follow-up height <10th %	1 (6)	6 (46)	.01
Follow-up head circumference <10th %	3 (17)	2 (17)	.99
Weight gain velocity (g/mo)	448.44 (108.42)	389.59 (83.20)	.36
Length gain velocity (cm/mo)	1.93 (0.22)	1.83 (0.16)	.10
Head circumference gain velocity (cm/mo)	0.78 (0.08)	0.78 (0.14)	.46
NDI <70	4 (22)	6 (46)	.25
NDI <85	7 (39)	6 (46)	.73
Moderate/severe cerebral palsy	2 (11)	1 (8)	.99
BSID III Cognitive <70	3 (17)	2 (15)	.99
BSID III Cognitive <85	6 (33)	5 (38)	.99
BSID III Language <70	5 (28)	2 (15)	.67
Chronic breathing problems	12 (67)	5 (38)	.16
Chronic feeding problems	9 (50)	10 (77)	.16

BSID, Bayley Scales of Infant and Toddler Development.

\*Information regarding discharge on NG/NJ feeding was only available from January 2008 through June 2011.

**Table VI. Rate of GT placement and fundoplication by center**

Centers	GT placement	Fundoplication
A	0/20 (0%)	0/20 (0%)
B	0/17 (0%)	0/17 (0%)
C	5/187 (3%)	2/187 (1%)
D	3/91 (3%)	0/91 (0%)
E	6/168 (4%)	2/168 (1%)
F	9/240 (4%)	2/240 (1%)
G	3/74 (4%)	1/74 (1%)
H	1/23 (4%)	1/23 (4%)
I	20/450 (4%)	3/450 (1%)
J	12/260 (5%)	6/260 (2%)
K	15/315 (5%)	2/315 (1%)
L	4/76 (5%)	3/76 (4%)
M	1/18 (6%)	1/18 (6%)
N	29/405 (7%)	12/405 (3%)
O	17/226 (8%)	8/226 (4%)
P	32/404 (8%)	7/404 (2%)
Q	19/232 (8%)	0/232 (0%)
R	10/119 (8%)	4/119 (3%)
S	9/96 (9%)	3/96 (3%)
T	34/342 (10%)	22/342 (6%)
U	22/201 (11%)	6/201 (3%)
V	26/231 (11%)	9/231 (4%)
W	5/37 (14%)	1/37 (3%)
X	50/312 (16%)	2/312 (1%)
Y	1/5 (20%)	0/5 (0%)