Risks associated with suspected dysphagia in infants admitted to a neonatal intensive care unit in a South African public hospital

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Background. The prevalence of neonatal dysphagia is increasing, as medical advances contribute to the survival of critically ill and preterm infants. Additional factors such as low birth weight (LBW), gastro-oesoephageal reflux disorder, failure-to-thrive (FTT), and HIV may increase the complexity of dysphagia symptoms. Knowledge of context-specific risk factors for dysphagia may lead to an effective pathway of diagnosis and management in vulnerable neonates.

Objective. To describe the feeding characteristics and categories of underlying medical conditions in infants of gestational age 24 - 42 weeks. **Methods.** The study was a retrospective review of 231 purposively selected medical and speech-language therapy records. Participants had a mean stay of 28.5 days in a neonatal intensive care unit in a peri-urban public hospital and were referred for a swallowing and feeding assessment. An existing seven-category framework for the classification of suspected dysphagia was used.

Results. Most participants (90.0%) presented with multiple medical conditions. Underlying neurological conditions (48.5%) and feeding difficulties secondary to systemic illness (65.8%) contributed mostly to suspected dysphagia in the sample. It was found that 71.0% of infants presented with feeding difficulties secondary to other conditions such as LBW and prematurity, highlighting the need for an expanded dysphagia classification framework.

Conclusion. The results concur with the outcomes of previous studies and confirm the need for a unique classification framework in South Africa. Dysphagia is a complex condition and frequently cannot be attributed to a single risk factor.

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Dysphagia in children is ever-increasing, mostly due to the improved survival rate of infants and children with life-threatening conditions and multiple associated health problems. ^[1,2] Dysphagia, a swallowing disorder secondary to a problem in one or more of the four phases of swallowing, is managed by speech-language therapists (SLTs) who are qualified to assess the dysfunction and provide intervention. ^[3-5]

Infants with risks such as prematurity, congenital or acquired medical conditions, or those with prolonged stays in neonatal intensive care units (NICUs) are at greater risk of developing dysphagia and nutritional problems than typically developing, healthy neonates. [4,6] Also, infants requiring continued intervention for dysphagia are frequently those who were previously admitted to a NICU. [6] Infants with the most complex or severe medical conditions are most at risk of presenting with disorganised or dysfunctional feeding patterns. [4] When preterm infants present with disorganised feeding patterns, it is generally due to immaturity, whereas dysfunctional sucking patterns may be more severe and are usually associated with neurological involvement. [4] It therefore appears that a close relationship exists between dysphagia, the infant's medical diagnosis, associated conditions, and the severity thereof.

Limited research regarding risk factors associated with dysphagia in infants admitted to the NICU is currently available. [4,7,8] This is a concern in developing countries, where the burden of disease is high. [9] In a developing country such as South Africa (SA), numerous additional challenges, such as the effects of poverty and HIV, may contribute to dysphagia in infants. [9] HIV may affect all phases of swallowing as a result of oral thrush, odynophagia and gastro-oesophageal reflux disorder (GERD), which can lead to failure-to-thrive (FTT). [10] The prevalence of dysphagia and

GERD among children with HIV is poorly recorded, but frequently encountered in clinical practice and may contribute significantly to the morbidity of infants with exposure to the virus. [10] The incidence of dysphagia in typically developing children is estimated at 25 - 45%, and even higher, up to 80%, in children with developmental disabilities. [11] The prevalence of neonatal dysphagia is unknown, but represents a universal problem, as dysphagia may carry over to infancy and toddler-age groups. [12]

It is important to identify dysphagia as soon as possible after birth, while the infant is still in the hospital, so that the appropriate short- and long-term dietetic and SLT management and parent training can commence. [4] SLTs should be able to state when an infant is not ready for oral feeding and maximise oral feeding skills and safety in those infants who are ready to feed orally. [1,5] Left unidentified and untreated, dysphagia can lead to FTT, GERD, aspiration pneumonia and an inability to establish and sustain vital nutrition and hydration. [13]

The objectives of the study were to describe the feeding characteristics of infants admitted to a NICU and referred for suspected dysphagia in a public hospital, and to determine which medical conditions were associated with the participants. Identifying risk factors can contribute to a better understanding of infants with suspected dysphagia, which may lead to improved referral guidelines and SLT staff-planning to ensure adequate intervention for all. A holistic understanding of the diversity of context-specific risk factors, associated with suspected dysphagia in infants who are already compromised by medical conditions, may be attained.

Methods

The study was a retrospective review of medical and SLT records from 2010 to 2014 and included infants, aged 24 - 42 weeks' gestational age

(GA) at birth, who were admitted to a peri-urban public hospital. GA was determined using the mother's last menstrual period. Birth weight was defined as: normal (>2 500 g); low birth weight (LBW; <2 500 g), very LBW (<1 500 g) and extremely LBW (<1 000 g). All participants were referred by medical doctors, nurses, audiologists, SLTs, and dieticians for a clinical swallowing evaluation.

Participants

A total of 312 infants were referred for dysphagia assessments within the study period, of which 231 complete data sets were available. Inclusion criteria were that the infants had to present with symptoms of dysphagia, be referred for a feeding or swallowing evaluation by a healthcare professional, admitted to the NICU and assessed for dysphagia by a SLT. Common symptoms of dysphagia in the participants include oral phase symptoms such as absent oral reflexes, absent or poor primitive reflexes, weak suck, uncoordinated suck, immature biting, poor bolus propulsion and poor bolus containment. Abnormalities in triggering of the swallow include absent swallow, delayed trigger of swallow, suck-swallow-breathing (SSB) incoordination, and pharyngeal phase symptoms include laryngeal penetration, aspiration, choking, pharyngeal residue and nasopharyngeal reflux.^[14]

Common criteria by healthcare professionals for referral of infants and children for feeding and swallowing evaluation included: suckling and swallowing incoordination, weak suck, breathing disruptions or apnoea during feeding, excessive gagging or recurrent coughing during feeds, diagnosis of disorders associated with dysphagia or under-nutrition, severe irritability during feeding, history of recurrent pneumonia and feeding difficulty, concern for possible aspiration during feeds, lethargy or decreased arousal during feeds, tedious feeding times and nasopharyngeal reflux during feeding.

Materials

The materials used during data collection included the Neonatal Patient Discharge Report, which is available in electronic format from the local NICU database, and the SLT records, including a dysphagia assessment form. [15-17] Data were independently collected over five years by seven different SLTs who were trained to use the same data collection instrument. A classification framework described by Arvedson and Brodsky^[1] to determine the aetiology or risk factors of paediatric dysphagia was used to categorise each participant. These categories included conditions with neurological involvement (such as asphyxia and convulsions), anatomical and structural impairments (including laryngomalacia), genetic and chromosomal disorders (including trisomy 21), dysphagia secondary to systemic illness (including pneumonia), psychosocial factors (including oral deprivation) as well as dysphagia secondary to resolved medical conditions (including hospital-acquired infections). In a local study, Fourie^[7] expanded on the framework by adding a seventh category, 'other', as the prevalence of prematurity, LBW, GERD, and FTT is high in SA, and does not fit within any of the other six categories.^[7]

Procedures

The Research Ethics Committees of two different universities granted approval for the study. Data were manually captured from the printed records to an Excel spreadsheet and analysed using SPSS version 22 (IBM Corp., USA). Being a retrospective study, there was no direct contact with mothers or infants. Variables included those in Arvedson and Brodsky's^[1] framework for categorisation of risks for dysphagia, which was expanded by Fourie^[7] to include prenatal risks (such as age of mother, number of antenatal visits), perinatal risks (such as type of delivery, Apgar scores), and postnatal medical risks (such as enteral and parenteral feeding). Pivot tables were used to determine the distribution of each participant within the different categories of medical conditions associated with suspected dysphagia. Standard

deviations were calculated. Descriptive statistics were used to identify the feeding characteristics and risk factors.

Characteristic	n (%)*
Nationality of mother	
South African	204 (88.3)
Non-South African	27 (11.7)
Gender	
Male	111 (48.1)
Female	120 (51.9)
GA at birth (weeks), mean (SD)	34.9 (3.9)
Birth weight	
Normal	105 (45.5)
LBW	82 (35.5)
Very LBW	38 (16.4)
Extremely LBW	6 (2.6)
Time spent in NICU (days), mean (SD)	28.5 (36.9)
HIV status of mother (N=196)	
Negative	135 (69.6)
Positive	59 (30.4)
GA = gestational age; SD = standard deviation; LBW = low	birth weight;

*Unless otherwise specified.

Results

NICU = neonatal intensive care unit.

Participant description

The participant characteristics are described in Table 1. Most mothers were SA citizens. There were slightly more female (51.9%) than male (48.1%) participants, and the mean (standard deviation (SD)) GA of participants was 34.9 (3.9) weeks. The participants were mostly late preterm. More than half of the participants (54.5%) were LBW, very LBW or extremely LBW (Table 1). The mean (SD) stay in the NICU was 28.5 (36.9) days. The percentage of mothers who were HIV-positive and, by implication, had infants exposed to the virus, corresponds with the 2012 antenatal sentinel HIV prevalence survey. [18] The estimated HIV prevalence in the survey was 29.5% in pregnant women and in this study, 30.4% of mothers were HIV-positive. [18] It is not known how many of the mothers were receiving antiretroviral treatment.

Feeding characteristics of participants

Table 2 shows the feeding characteristics of the participants. All participants presented with one or more symptoms of dysphagia. This can explain the frequency of parenteral (14.4%) and enteral feeding (65.0%), as infants with dysphagia often require alternative feeding methods to obtain adequate nutrients and fluids.[1] Thirty-six (15.6%) participants presented with severe feeding difficulties or signs of aspiration. An instrumental assessment, a video fluoroscopic swallow study (VFSS) was only conducted in 14 (38.8%) of those participants. Instrumental assessments are recommended if there are concerns about risks for aspiration, safety of the airway, or possibilities of GERD. [1] The reasons why VFSS was not conducted in the 22 remaining participants referred for the procedure included: participants demised before VFSS could be conducted (n=7); VFSS screening machine was not functioning (n=1); participant was ventilated (n=1); participant was lethargic (n=1); unstable or desaturating during feeding (n=3); clinically

Characteristic	n (%)
Previous parenteral feeding	
Yes	33 (14.4)
No	198 (85.6
Previous enteral (NGT/OGT) feeding	
Yes	150 (65.0
No	61 (26.4)
Unknown	20 (8.6)
Referred for VFSS by doctors and SLTs (N=36))
Yes	6 (15.6)
No	30 (84.4)
VFSS conducted (in referrals)	
Yes	14 (38.8)
No	22 (61.2)
Manner of feeding at discharge	
Mixed	127 (55.1
Exclusive breastfeeding	69 (29.7)
Gastrostomy	14 (6.1)
Exclusive bottlefeeding	12 (5.2)
Cup	6 (2.6)
Syringe	3 (1.3)

aspirating but no suck/swallow palpable (*n*=4), clinical swallow present even though there were risks for aspiration (*n*=5). More than half of participants (55%) used a mixed manner of feeding such as breast- and cupfeeding, or cup- and syringe feeding. Only 29.7% of participants could breastfeed exclusively, which was related to preterm birth and LBW in most of the participants. The use of mixed feeding methods among the participants may indicate that the infants experienced breastfeeding difficulties, as establishing successful breastfeeding may be a challenge for many preterm infants and their mothers, owing to neonatal feeding difficulties.^[19] These difficulties may be due to incoordination of SSB as the suckling patterns of preterm infants often remain significantly less efficient than those of full-term infants at term age and beyond.^[2,19]

Furthermore, 6.1% of the participants required long-term tube feeding, such as a gastrostomy. In another study conducted in SA, it was found that infants and children requiring gastrostomies were likely to present with multiple diagnoses, of which neurological and/ or gastrointestinal impairments were the most prominent medical conditions.^[20]

Underlying medical conditions in participants

Underlying medical conditions in participants were classified according to the framework by Arvedson and Brodsky, [1] expanded by Fourie, [7] to determine the aetiology or risk factors of dysphagia (Table 3). Since most participants were not classified in a single category, and presented with multiple risks, the total in Table 3 does not add to 100%.

The majority of the infants (71.0%) presented with conditions that were not included in the risks of dysphagia described by Arvedson and Brodsky, ^[1] a classification system developed for conditions in a developed country such as the USA. The risks included FTT,

Table 3. Underlying medical conditions in participants (N=231)

Category	Description	n (%)
A	Neurological conditions	112 (48.5)
В	Anatomical and structural conditions	19 (8.2)
С	Secondary to systemic illness	152 (65.8)
D	Chromosomal (genetic) conditions	18 (7.8)
E	Psychosocial conditions	4 (1.7)
F	Secondary to resolved medical condition	32 (13.9)
G	Other (FTT, LBW, prematurity)	164 (71.0)
FTT = failure-to-thrive; LBW = low birth weight.		

GERD, LBW as well as HIV exposure. It was found that 65.8% of participants had feeding difficulties secondary to a systemic illness, such as respiratory distress syndrome, cardiac abnormalities, and pneumonia. This could be due to the fact that preterm infants with LBW are more at risk of developing systemic illnesses,^[13] and more so in a developing country, such as SA.^[7] Results indicated that 48.5% of participants had a condition with neurological involvement, such as asphyxia. The literature suggests that infants with neurological conditions, birth trauma, as well as pre- and perinatal asphyxia, are commonly found to have feeding difficulties.^[1]

The conditions that occurred the least in the participants were feeding difficulties secondary to resolved medical conditions (13.9%), including iatrogenic conditions such as hospital-acquired infections. A total of 8.2% of the participants presented with anatomical or structural conditions, such as cleft lip and palate, laryngomalacia and tracheo-oesophageal fistula, while only 7.8% of the 231 participants presented with genetic or chromosomal abnormalities, which included infants with trisomies 13, 18 and 21, and other syndromes. Only 1.7% of the participants presented with psychosocial conditions such as oral deprivation and under-nutrition due to social problems. When analysing the results, it became clear that a true profile of multiple underlying conditions to feeding difficulties in the participants could not have been obtained if single categories of risk were considered.

Combinations of risk conditions associated with suspected dysphagia

Combinations of risk categories in participants are described in Table 4. The results indicate that 90.0% of participants presented with multiple medical conditions, therefore revealing the complexity of combinations of different categories. A total of 36 different combinations were found, ranging from a single category to five different combinations. Most of the participants presented with two (50.2%) or three (28.1%) categories of risk factors and a total of 11.7% participants presented with four or five categories of risks. The minority (10.0%) of participants presented with a single category of risk for dysphagia. The results display the diversity and complexity of medical conditions in infants with symptoms of dysphagia. The results are in agreement with Jadcherla, ^[12] who states that neonatal dysphagia can rarely be associated with a single aetiology.

Discussion

Dysphagia symptoms were accompanied by multiple medical conditions in most of the participants. As was found in other local studies, [7,20] participants presented with a great variety of medical conditions and combinations of these conditions that either directly or indirectly affected their feeding ability. [7]

The high number of participants with neurological conditions in this sample can be explained by the fact that infants with neurological

Category	n (%)
Single risk category	23 (10.0)
Neurological	6 (2.6)
Anatomical	2 (0.9)
Dysphagia SSI	10 (4.3)
Other	5 (2.2)
Two risk categories	116 (50.2
Anatomical, genetic	1 (0.4)
Neurological, SSI	33 (14.3)
Neurological, other	7 (3.0)
SSI, other	65 (28.1)
Anatomical, SSI	4 (1.7)
SSI, SRMC	1 (0.4)
SSI, neurological	1 (0.4)
SSI, genetic	4 (1.7)
Three categories	65 (28.1)
Neurological, SSI, other	39 (16.9)
Neurological, SRMC, other	1 (0.4)
Anatomical, SSI, other	4 (1.7)
SSI, SRMC, other	10 (4.3)
SSI, genetic, other	2 (0.9)
Neurological, SSI, SRMC	4 (1.7)
SSI, psychosocial, SRMC	1 (0.4)
Neurological, anatomical, SSI	1 (0.4)
SSI, psychosocial, other	1 (0.4)
Neurological, other, SSI	1. (0.4)
Neurological, SSI, genetic	1 (0.4)
Four risk categories	22 (9.5)
Anatomical, SSI, genetic, other	3 (1.3)
Neurological, SSI, SRMC, other	9 (3.9)
Anatomical, SSI, SRMC, other	1 (0.4)
Neurological, anatomical, SSI, other	3 (1.3)
SSI, genetic, SRMC, other	1 (0.4)
Neurological, SSI, psychosocial, other	1 (0.4)
Neurological, Genetic, SSI, Other	1 (0.4)
SSI, Psychosocial, SRMC, Other	1 (0.4)
Neurological, SSI, Genetic, Other	2 (0.9)
Five risk categories	5 (2.2)
Neurological SSI, Genetic, SRMC, Other	1 (0.4)
Anatomical, SSI, Genetic, SRMC, Other	1 (0.4)
Neurological, Anatomical, SSI, Psychosocial, Other	2 (0.9)
Neurological, Anatomical, SSI, Psychosocial, Other	1 (0.4)
SI = secondary to systemic illness; SRMC = secondary to resolved	

conditions are commonly found to have feeding difficulties. [1,13] It is estimated that 85 - 90% of infants and children with neurological conditions, such as cerebral palsy, will present with dysphagia at some point in their lives. [1] Further, the high incidence of systemic illnesses, such as pneumonia, in paediatric populations with dysphagia is linked to specific diagnoses, such as trisomy 21, asthma, GERD, lower respiratory tract infection, and moist cough. [14] Literature indicated that paediatric patients with multisystem diagnoses, in addition to dysphagia, appear to be at greatest risk for developing pneumonia. [14] It is therefore evident that infants can present with multiple variations of swallowing impairments, such as those found in the participants of this study.[14]

The results indicate that the seven-category framework used for classification of risks for dysphagia in participants was successful to describe the complexities of different risk categories that may underlie neonatal dysphagia. Fourie^[7] found that 52% of participants had aetiological factors for dysphagia pertaining to the 'other' category. [7] In the current study, the high rate of 71.0% participants in the 'other' category included those with HIV exposure, as there was no dedicated category for infants exposed to HIV. Therefore, the results indicated a need for an expanded classification system and the importance of an additional risk category was highlighted. It is proposed that the framework as described by Arvedson and Brodsky should be expanded to an eight-category classification framework that includes a category for prematurity, LBW and related conditions (described by Fourie^[7] as 'other') as well as a category for infants exposed to HIV. HIV exposure in infants is associated with preterm birth.[21] As a result of prematurity and LBW, the infant is at risk for dysphagia after birth[19] and when HIV infection becomes apparent, feeding and swallowing can be affected due to encephalopathy. An additional category would provide information regarding feeding characteristics, and aid in early identification of dysphagia.

Conclusion

Dysphagia frequently occurs in infants and is highly complex in nature.[12] Within the context of a developing country, classifying dysphagia can be challenging and therefore an expanded framework may be beneficial. The eight-category framework can be used by healthcare personnel to refer infants for dysphagia assessment and intervention, and can be used by SLTs to identify infants at risk for dysphagia. Being a retrospective study, various limitations were present, including missing data as well as the restricted geographical location. The outcomes of the current study correspond with international research describing several risk factors for dysphagia related to the primary medical diagnosis and its sequelae, and may be present throughout the infants' hospitalisation. [8,12] Due to the increased survival rate of preterm infants and infants with complex medical conditions, it is suggested that more research regarding neonatal dysphagia in developing countries should be conducted.

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