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7 ELSSP

Introduction

In the United States, 1-2 children are born with hearing loss, per 1,000 births (CDC, 9 2018). This translates to 114,000 Deaf or Hard of Hearing (DHH) children born in the U.S. 10 per year (Martin, Hamilton, Osterman, & Driscoll, 2019). Of these 114,000, ~90\% will be 11 born to hearing parents (Mitchell & Karchmer, 2004), in a home where spoken language is 12 likely the dominant communication method. Depending on the type and degree of hearing 13 loss and whether the child uses amplification, spoken linguistic input will be partially or totally inaccessible. Some of these children will develop spoken language within the range of 15 their hearing peers (Geers, Mitchell, Warner-Czyz, Wang, & Eisenberg, 2017; Verhaert, 16 Willems, Van Kerschaver, & Desloovere, 2008), but many will face persistent spoken language deficits (Eisenberg, 2007; Luckner & Cooke, 2010; Moeller, Tomblin, Yoshinaga-Itano, Connor, & Jerger, 2007; Sarchet et al., 2014), which may later affect reading ability (Kyle & Harris, 2010) and academic achievement (Karchmer & Mitchell, 2003; Qi & Mitchell, 2012).

Despite many excellent studies examining language development in DHH children,
there is still a gap in the literature describing and analyzing spoken language development
across the full range of children receiving state services for hearing loss, with many studies
focusing in on specific subgroups (e.g. children under age X with Y level of hearing loss and
Z amplification approach, e.g. (Vohr et al., 2008; Yoshinaga-Itano, Sedey, Wiggin, & Mason,
2018)). In what follows, we first summarize the previous literature on predictors of spoken
language outcomes in DHH children. We then provide a brief overview of a common
vocabulary measure used in the current study, the MacArthur-Bates Communicative
Development Inventory (CDI). Finally, we turn to an empirical analysis of early vocabulary
in a wide range of young children receiving state services in North Carolina. We have two
broad goals in what follows. First, we aim to provide a comprehensive description of a
heterogeneous group of young children who receive state services for hearing loss. Second, we

³³ aim to connect the intervention approaches and child characteristics of this sample with

children's vocabulary, with the broader goal of considering the success of early diagnosis and

35 intervention initiatives.

36 Predictors of Language Outcomes

Though the literature points towards spoken language delays and deficits for DHH

children, this is a highly variable population with highly variable outcomes (Pisoni,

Kronenberger, Harris, & Moberly, 2018). Previous research indicates that gender (Ching et

al., 2013; C Kiese-Himmel & Ohlwein, 2002), additional disability (Ching et al., 2013;

Verhaert et al., 2008; Yoshinaga-Itano, Sedey, Wiggin, & Chung, 2017), degree and

configuration of hearing loss (Ching et al., 2013; de Diego-Lázaro, Restrepo, Sedey, &

Yoshinaga-Itano, 2018; Vohr et al., 2011; Yoshinaga-Itano et al., 2017), amplification (Walker

et al., 2015), communication (Geers et al., 2017), and early diagnosis/intervention

45 (Yoshinaga-Itano et al., 2017, 2018) predict language outcomes in DHH children.

Gender. For hearing children, the literature points to a female gender advantage in

early language acquisition. Girls speak their first word earlier (Macoby, 1966), have a larger

(Bornstein, Hahn, & Haynes, 2004; Fenson et al., 1994; Frank, Braginsky, Yurovsky, &

Marchman, 2017) and faster-growing vocabulary (Huttenlocher, Haight, Bryk, Seltzer, &

50 Lyons, 1991), and stronger grammatical and phonological skills (Lange, Euler, & Zaretsky,

51 2016; Özçalışkan & Goldin-Meadow, 2010). This finding appears to be consistent across

studies (Wallentin, 2009), various spoken languages (Frank, Braginsky, Marchman, &

Yurovsky, 2019), and gesture (Özçalışkan & Goldin-Meadow, 2010).

The DHH literature presents a more mixed (though rather understudied) picture. On

one hand, DHH girls, like hearing girls, have been found to have a larger spoken vocabulary

than DHH boys (Ching et al., 2013; C Kiese-Himmel & Ohlwein, 2002). However, in contrast

57 to their hearing peers, DHH children do not seem to show a gender-based difference for some

aspects of syntactic development (Pahlavannezhad & Tayarani Niknezhad, 2014).

Comorbidities. Additional co-occurring disabilities occur frequently in the DHH population, perhaps as much as three times more than in the hearing population (Pollack, 1997). Incidence estimates for co-occurring disabilities in DHH children range from 25-51% (Bruce & Borders, 2015; Guardino, 2008; Holden-Pitt & Diaz, 1998; Luckner & Carter, 2001; Picard, 2004; Schildroth & Hotto, 1996; Soukup & Feinstein, 2007), with approximately 8% children living with 2 or more co-occurring disabilities (Schildroth & Hotto, 1996).

Some of these conditions, particularly those which carry risk of developmental delay

(e.g., Down syndrome), result in language delays independent of hearing loss (Chapman,

1997; Kristoffersen, 2008; Weismer, Lord, & Esler, 2010), with cognitive ability more

predictive of language outcomes than presence or absence of a specific disability

(Meinzen-Derr, Wiley, Grether, & Choo, 2011; Sarant, Holt, Dowell, Richards, & Blamey,

2008). Disability and hearing loss likely each contribute to a given child's language

development (Ching et al., 2013; Rajput, Brown, & Bamiou, 2003; Van Nierop et al., 2016),

with differential effects of each (Vesseur et al., 2016). In some cases, additional disabilities

appear to interact with hearing loss to intensify developmental delays (Birman, Elliott, &

Gibson, 2012; Pierson et al., 2007).

Furthermore, incidence of hearing loss is higher among children born premature (defined as 37 weeks gestational age). Compared to an incidence 0.2% in full-term infants, incidence of hearing loss in extremely premature infants (defined as 33 weeks gestational age) ranges 2–11%, with increased prematurity associated with increased rates of hearing loss (Wroblewska-Seniuk, Greczka, Dabrowski, Szyfter-Harris, & Mazela, 2017).

Independently of hearing status, prematurity is linked to increased risk of language
delay and disorder (Barre, Morgan, Doyle, & Anderson, 2011; Carter & Msall, 2017; Cusson,
2003; Rechia, Oliveira, Crestani, Biaggio, & de Souza, 2016; Van Noort-van Der Spek,
Franken, & Weisglas-Kuperus, 2012; Vohr, 2014). Unfortunately, research on language
development in premature DHH children is scant (Vohr, 2016), so it remains unclear how

hearing loss and prematurity may interact within spoken language skills. One study of
premature infants finds that auditory brainstem response during newborn hearing screening
predicts language performance on the PLS-4 at age 3 (Amin, Vogler-Elias, Orlando, & Wang,
2014), suggesting a link between prematurity and hearing loss in early childhood, though
further research is needed in this domain. In extremely premature DHH children, incidence
of additional disabilities may be as high as 73% (Robertson, Howarth, Bork, & Dinu, 2009).
Indeed, pre-term infants with comorbidities have been found to be more likely to also have
hearing loss than those without comorbidities (Schmidt et al., 2003), further complicating
language development for this population.

Audiological Characteristics. Hearing loss varies in severity, ranging from slight to profound (Clark, 1981). More severe hearing loss (less access to spoken language) typically results in more difficulty with spoken language in infancy (Vohr et al., 2008), early childhood (Ching et al., 2010, 2013; Sarant et al., 2008; Sininger, Grimes, & Christensen, 2010; Tomblin et al., 2015) and school-age (Wake, Hughes, Poulakis, Collins, & Rickards, 2004). Although profound hearing loss is associated with more pronounced spoken language difficulty, even mild to moderate hearing loss is associated with elevated risk of language disorders (Blair, Peterson, & Viehweg, 1985; Delage & Tuller, 2007).

Hearing loss also varies in whether it affects one ear or both. Bilateral hearing assists 102 speech perception, sound localization, and loudness perception in quiet and noisy 103 environments (Ching, Van Wanrooy, & Dillon, 2007). The literature on hearing aids and 104 cochlear implants also points to benefits for bilateral auditory input (Lovett, Kitterick, 105 Hewitt, & Summerfield, 2010; Sarant, Harris, Bennet, & Bant, 2014; Smulders et al., 2016). At school-age, 3-6% of children have unilateral hearing loss (Ross, Visser, Holstrum, Qin, & Kenneson, 2010). Although children with unilateral hearing loss have one "good ear," even mild unilateral hearing loss has been tied to higher risk of language delays and educational 109 challenges relative to hearing children (C. Kiese-Himmel, 2002; Lieu, 2004, 2013; Lieu, 110 Tye-Murray, & Fu, 2012; Vila & Lieu, 2015). That is, just as in the bilateral case, more 111

severe hearing loss leads to greater deficits in language and educational outcomes for children with unilateral hearing loss (Anne, Lieu, & Cohen, 2017; Lieu, 2013).

Many DHH children receive hearing aids (HAs) or cochlear implants (CIs) to boost access to the aural world. These devices have been associated with better speech perception and spoken language outcomes (Niparko et al., 2010; Walker et al., 2015; Waltzman et al., 1997). In turn, aided audibility predicts lexical abilities with children in HAs (Stiles, Bentler, & McGregor, 2012).

For both hearing aids and cochlear implants, earlier fit leads to better spoken language 119 skills, if the amplification is effective. For hearing aids, some studies find that children with 120 milder hearing loss who receive hearing aids earlier have better early language achievement 121 than children who are fit later (Tomblin et al., 2015), but this finding does not hold for 122 children with severe to profound hearing loss (C. Kiese-Himmel, 2002; Watkin et al., 2007) 123 (for whom hearing aids are generally ineffective). Analogously, children who are eligible and 124 receive cochlear implants earlier have better speech perception and spoken language 125 outcomes than those implanted later (Artières, Vieu, Mondain, Uziel, & Venail, 2009; 126 Dettman, Pinder, Briggs, Dowell, & Leigh, 2007; Miyamoto, Hay-McCutcheon, Kirk, 127 Houston, & Bergeson-Dana, 2008; Svirsky, Teoh, & Neuburger, 2004; Yoshinaga-Itano et al., 128 2018), with best outcomes for children receiving implants before their first birthday 129 (Dettman et al., 2007). 130

Communication. Total Communication (TC) refers to communication that
combines speech, gesture, and elements of sign (but not a full sign language, such as
American Sign Language), sometimes simultaneously. Clinicians currently employ TC as an
alternative or augmentative communication method for children with a wide range of
disabilities (Branson & Demchak, 2009; Gibbs & Carswell, 1991; Mirenda, 2003).

Compared to total communication, DHH children using an exclusively oral approach have better speech intelligibility (Dillon, Burkholder, Cleary, & Pisoni, 2004; Geers et al.,

2017; Geers, Spehar, & Sedey, 2002; Hodges, Dolan Ash, Balkany, Schloffman, & Butts, 1999) and auditory perception (Geers et al., 2017; O'Donoghue, Nikolopoulos, & Archbold, 139 2000). That said, there is some debate as to whether an oral approach facilitates higher 140 spoken language performance, or whether children who demonstrate aptitude for spoken 141 language are steered towards the oral approach rather than TC (Hall, Hall, & Caselli, 2017). 142 1-3-6 Guidelines. Early identification (Apuzzo & Yoshinaga-Itano, 1995; Kennedy 143 et al., 2006; Robinshaw, 1995; White & White, 1987; Yoshinaga-Itano, Sedey, Coulter, & 144 Mehl, 1998; Yoshinaga-Itano et al., 2018) and timely enrollment in early intervention 145 programs (Ching et al., 2013; Holzinger, Fellinger, & Beitel, 2011; Vohr et al., 2008, 2011; 146 Watkin et al., 2007) are associated with better language proficiency. Indeed, DHH children who receive prompt diagnosis and early access to services have been found to meet age-appropriate developmental outcomes, including language (Stika et al., 2015).

In line with these findings, the American Academy of Pediatricians (AAP) has set an initiative for Early Hearing Detection and Intervention (EHDI). Their EHDI guidelines recommend that DHH children are screened by 1 month old, diagnosed by 3 months old, and enter early intervention services by 6 months old. We refer to this guideline as 1-3-6.

Meeting this standard appears to improve spoken language outcomes for children with HL (Yoshinaga-Itano et al., 2017, 2018) and the benefits appear consistent across a range of demographic characteristics.

At a federal level in the U.S., the Early Hearing Detection and Intervention Act of
2010 (Capps, 2009) was passed to develop state-wide systems for screening, evaluation,
diagnosis, and "appropriate education, audiological, medical interventions for children
identified with hearing loss," but policies for early diagnosis and intervention vary by state.
As of 2011, 36 states (including North Carolina, ("15A NCAC 21F .1201 - .1204," 2000)]
mandate universal newborn hearing screening (National Conference of State Legislatures,
2011). All states have some form of early intervention programs that children with hearing

loss can access (NAD, n.d.), but these also vary state-by-state. For instance, half of the states in the US do not consider mild hearing loss an eligibility criterion for early intervention (Holstrum, Gaffney, Gravel, Oyler, & Ross, 2008).

In evaluating the success of this initiative, the AAP (EHDI, n.d.) finds that about 70% of US children who fail their newborn hearing screening test are diagnosed with hearing loss before 3 months old, and that 67% of those diagnosed (46% of those that fail newborn hearing screening) begin early intervention services by 6 months old. These findings suggest that there may be breaks in the chain from screening to diagnosis and from diagnosis to intervention, and the effect may be further delays in language development for children not meeting these guidelines.

Quantifying vocabulary growth in DHH children

The MacArthur Bates Communicative Development Inventory (CDI, Fenson et al., 175 1994) is a parent-report instrument that gathers information about children's vocabulary 176 development. The Words and Gestures version of the form (CDI-WG) is normed for 8-18-month-olds, and includes 398 vocabulary items that parents indicate whether their child understands or produces, along with questions about young children's early communicative 179 milestones. The Words and Sentences version of the form (CDI-WS) is normed for 180 16-30-month-olds, and includes 680 vocabulary items that parents indicate whether their 181 child produces, along with some questions about grammatical development. The CDI has 182 been normed on a large set of participants across many languages (Anderson & Reilly, 2002; 183 Frank et al., 2017; Jackson-Maldonado et al., 2003). 184

The CDI has also been validated for DHH children with cochlear implants (Thal,
Desjardin, & Eisenberg, 2007). More specifically, in this validation, researchers asked parents
to complete the CDI, administered the Reynell Developmental Language Scales, and
collected a spontaneous speech sample. All comparisons between the CDI and the other

measures yielded significant correlations ranging from 0.58 to 0.93. Critically, the children in 189 this study were above the normed age range for the CDI, and thus this validation helps to 190 confirm that the CDI is a valid measurement tool for older DHH children. In further work, 191 Castellanos, Pisoni, Kronenberger, and Beer (2016) finds that in children with CIs, number 192 of words produced on the CDI predicts language, executive function, and academic skills up 193 to 16 years later. Building on this work, several studies have used the CDI to measure 194 vocabulary development in DHH children [Ching et al. (2013); Yoshinaga-Itano et al. (2017); 195 Yoshinaga-Itano et al. (2018); de Diego-Lázaro et al. (2018); Vohr et al. (2008); Vohr et al. 196 (2011); summarized in table XXX]. 197

• Goals and Predictions

This study aims to 1) characterize the demographic, audiological, and intervention
variability in the population of DHH children receiving state services for hearing loss; 2)
identify predictors of vocabulary delays; and 3) evaluate the success of early identification
and intervention efforts at a state level. We include two subgroups of DHH children
traditionally excluded from studies of language development: children with additional
disabilities and children with unilateral hearing loss (e.g., Yoshinaga-Itano et al., 2018).

For the first and third goal above, we did not have specific hypotheses and sought to provide descriptive information about a diverse sample of DHH children receiving state services. For the second, we hypothesized that male gender, more severe degree of hearing loss, bilateral hearing loss, no amplification use, prematurity, and presence of additional disabilities would predict larger spoken vocabulary delay. We did not have strong predictions regarding communication method, language background, or presence of other health issues (e.g., congenital heart malformation).

212 Methods

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Clinical evaluations were obtained through an ongoing collaboration with the North
Carolina Early Language Sensory Support Program (ELSSP), an early intervention program
serving children with sensory impairments from birth to 36 months. ELSSP passed along
deidentified evaluations to our team after obtaining consent to do so from each family. No
eligibility criteria beyond hearing loss and receiving an ELSSP evaluation were imposed,
given our goal of characterizing the full range of DHH children with hearing loss in North
Carolina.

The clinical evaluations included demographic and audiological information, CDI vocabulary scores, and the results of any clinical assessments administered (e.g., PPVT), all detailed further below. For some children (n=43), multiple evaluations were available from different timepoints. In these cases, only the first evaluation was considered for this study, due to concerns regarding within-subjects variance for statistical analysis.

While this collaboration is ongoing, we opted to pause for this analysis upon receiving 225 data from 100 children. Thus, the reported sample below consists of 100 children (56 male / 226 44 female) ages 4.20-36.17(M=21.30, SD=9.17). Race and SES information was not 227 available. Families were administered either the WG or WS version of the CDI based on 228 clinician judgement. Children who were too old for WG, but who were not producing many 229 words at the time of assessment, were often given WG (n=37). Families for whom Spanish 230 was the primary language (n = 14) completed the Spanish version of the CDI 231 (Jackson-Maldonado et al., 2003). 232

Children in this sample were coded as yes/no for cognitive development concerns (e.g.,
Down syndrome, global developmental delays; Cornelia de Lange syndrome), yes/no for
prematurity (i.e., more than 3 weeks premature), yes/no for health issues (e.g., heart defects,
kidney malformations, VACTERL association), and yes/no for vision loss (not corrected to

237 normal by surgery or glasses)

 \begin{table}

Condition	Specific Condition	n
Premature		17
	Extremely Premature	10
	NICU stay	16
Health Issues		34
	Heart	9
	Lung	5
	Illness	14
	Feeding Issues	13
	Pregnancy/Birth Complications	10
	Musculoskeletal	9
	Cleft Lip/Palate	4
	Other	14
Developmental Concerns		16
	Down Syndrome	5
	Chromosomal Issues	2
	Neural Tube Defects	2
	Other	9
Vision Loss		5
	Retinopathy of Prematurity	1
	Nearsightedness	1

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Degree of hearing loss was most often reported with a written description (e.g., "mild sloping 242 to moderate" or "profound high frequency loss"). We created 3 variables: hearing loss in the 243 better ear, hearing loss in the worse ear, and average hearing loss (average of better and 244 worse ear). Using the ASHA hearing loss guidelines, each of these was coded with a dB HL 245 value corresponding with the median dB HL for the level of hearing loss (e.g., moderate 246 hearing loss was coded as 48dB HL), and sloping hearing loss was coded as the average of 247 the levels (e.g. mild to moderate was coded as 40.5 dB HL). Participants were also coded for 248 unilateral or bilateral hearing loss; presence or absence of Auditory Neuropathy Spectrum 249 Disorder; sensorineural, conditive, or mixed hearing loss. Amplification was recorded as the 250 device the child used at the time of assessment-either hearing aid, cochlear implant, or none. 251

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\tab:amp_info)Audiological Characteristics of the Sample for Unilateral /
Bilateral Hearing Loss}

	n	Average HL (better ear)	Average HL (worse ear)	Average Age at Amplification
Hearing Aid (n=54)	11 / 43	3.89 / 47.92 dB	58.54 / 56.47 dB	9.73 / 8.75 months
Cochlear Implant (n=17)	0 / 17	NA / 85.6 dB	NA / 89.79 dB	NA / 14.12 months
No Amplification (n=27)	14 / 13	2.5 / NA dB	73.9 / NA dB	NA
Total (n=100)	25 / 73	3.11 / 57.37 dB	67.57 / 64.08 dB	NA

 $^{^{\}rm a}$ N.B. Age Amplification for children with CIs represents age at implantation $\label{eq:condition} \\ \mbox{end} \{ \mbox{table} \}$

Communication method was recorded as spoken language, total communication, or cued speech. One participant had a parent fluent in sign language, but the reported communication method in the home was total communication. No child in our sample used sign language. Participants were also coded as monolingual or multilingual based on whether families reported using more than one language at home. Total communication was not

counted as multilingualism.

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\caption{(#tab:comm_info)Language and communication characteristics of the sample}

Communication Method	English	Spanish	Hindi
Spoken Language (n=79)	68	10	1
Total Communication (n=18)	15	3	0
Cued Speech (n=1)	1	0	0

 $\ensuremath{\mathsf{end}}$

Age at screening was measured as the child's age in months at their first hearing screening. Age at screening was only available for (XXX) participants. If participants received their newborn hearing screening, age at screening was recorded as 0 (months). Age at diagnosis was taken as the age in months when children received their first hearing loss diagnosis. All children were enrolled in birth-to-three early intervention services through NC ELSSP, and the date of enrollment was listed on the clinician evaluation. From the clinician report, we calculated the number of hours of early intervention services received per month (including service coordination, speech therapy, and occupational therapy, among others). Because of the sparse data on screening age, if participants had an age at diagnosis <= 3 mo. and an age of intervention <= 6 mo., they were recorded as meeting 1-3-6. It is possible that a participant did not receive screening by 1 month, but did receive diagnosis by 3 months and services by 6 months. This special case would be coded as meeting 1-3-6 by our criteria.

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Diagnosis by 3 months	71.58%
Average Age Diagnosis (SD)	4.08 (7.27) months
Intervention by 6 months	39.18%
Average Age Intervention (SD)	11.16 (8.73) months
Meets 1-3-6	36.84%

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Variable	Scale	Range			
Age	Continuous	4.2-36 months			
Age at Amplification	Continuous	2-31 months			
Age at Diagnosis	Continuous	0-30 months			
Age at Implantation	Continuous	7-32 months			
Age at Intervention	Continuous	1-33 months			
Amplification	Categorical	Hearing Aid / Cochlear Implant / None			
Communication	Categorical	Spoken / Total Communication / Cued Speech			
Degree Hearing Loss (worse ear)	Continuous	17.75-100 dB HL			
Developmental Delay	Categorical	Yes / No			
Gender	Categorical	Female / Male			
Health Issues	Categorical	Yes / No			
Language in Home	Categorical	English / Other			
Laterality	Categorical	Unilateral / Bilateral			
Meets 1-3-6	Categorical	Yes / No			
Prematurity	Categorical	Full-term / Premature			
Services Received Per Month	Continuous	0-43 services per month			
Type of Hearing Loss	Categorical	Sensorineural / Conductive / Mixed			
CDI - Words Produced	Continuous	0-635 words			
\end{table}					

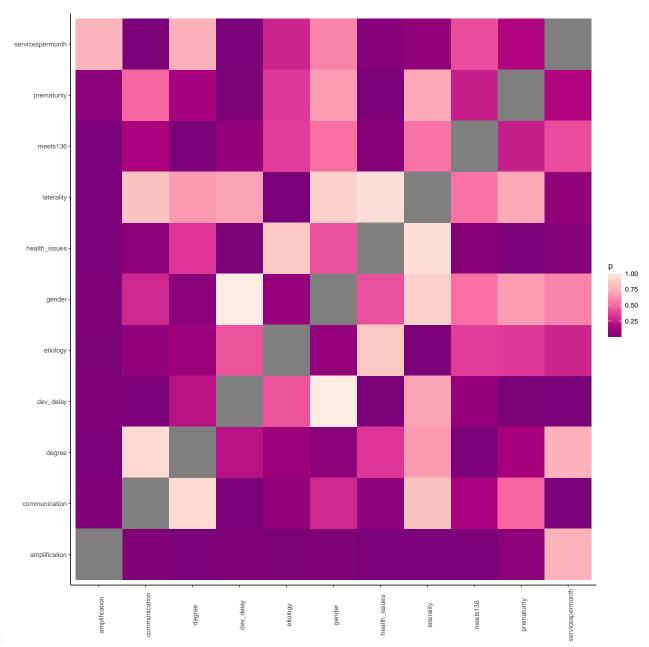
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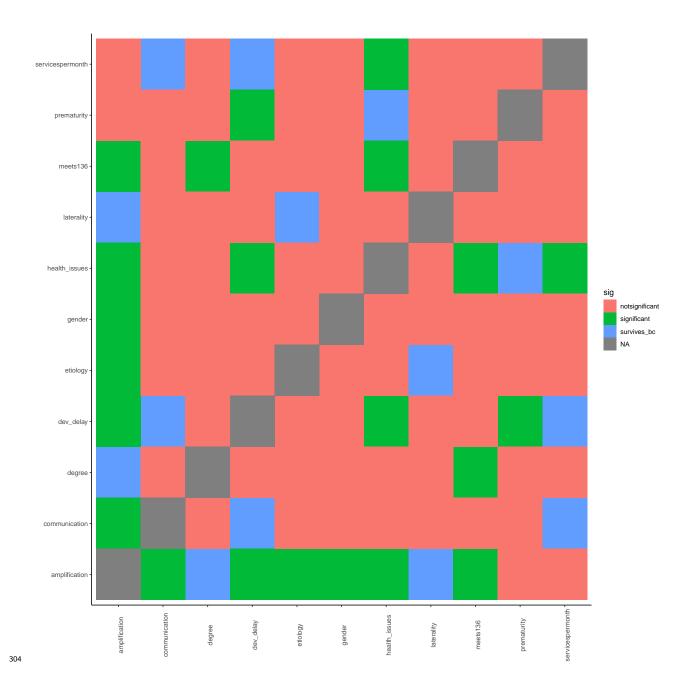
Results 287

All analyses were conducted in R. All code is available on Github. In the first section, we explore relationships among child demographic, audiological, and clinical variables. In the second section, we examine the influence of these factors on vocabulary development. In the third section, we describe the implementation of the EHDI 1-3-6 guidelines and predictors of early diagnosis and intervention.

Part I: Interactions Among Variables

Shapiro–Wilk tests revealed that all of our continuous measures (i.e. degree of hearing loss, services received per month, vocabulary delay) significantly differed from a normal distribution (ps <.05), so we used nonparametric tests to explore relationships among our variables. For categorical-categorical relationships, we used chi square tests; for continuous-categorical tests, we used mann-whitney U tests (2 levels for categorical variable) or kruskal-wallis tests (>2 levels for categorical variable; for continuous-continuous relationships, we used Of the fifty-five combinations of variables, p < .05 for sixteen, and seven survived bonferroni correction (p < 0.00). The full set of comparisons is shown in figure XXX.





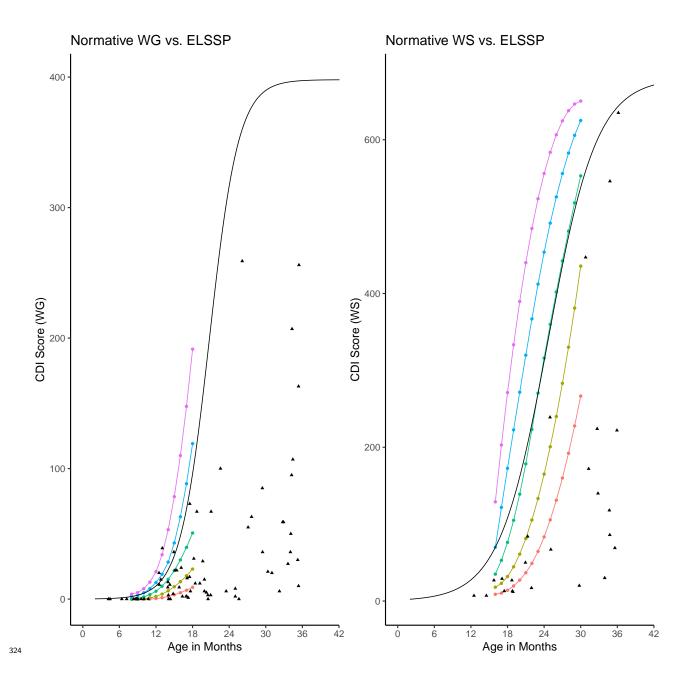
From this analysis, we found that children born premature were more likely to also have health issues (X2 (1, N = 96) = 25.69, p = 0.00.) Children with conductive hearing loss were more likely to have unilateral hearing loss (X2 (2, N = 86) = 14.84, p = 0.00). Children with unilateral hearing loss were unlikely to receive a cochlear implant and more likely to use no amplification (X2 (2, N = 96) = 17.19, p = 0.00). Children with more severe hearing loss were more likely to use a cochlear implant than children with milder hearing loss

 311 (H(2)=23.80, p=0.00). Children with developmental delays received more services per month 312 than typically developing DHH children (H()=134.50, p=0.00)and were more likely to use 313 total communication (X2 (2, N = 96)) = 19.38, p = 19.38). Children who used total 314 communication received more services per month than children using spoken language 315 (H(1)=15.60, p=0.00).

Part II: Influence on vocabulary

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We first constructed a binary logistic growth curve for vocabulary from the 50th percentile
data for typically developing children from Wordbank. With this function, each participant's
CDI score yielded a predicted age from the normative data. For each child, we subtracted
this predicted age (given the score) from the child's actual age to give us a measure of delay
in months. Descriptively, we found widespread vocabulary delays on both Words and
Gestures and Words and Sentences, with the majority of DHH children testing around or
below the 25th percentile for hearing children.

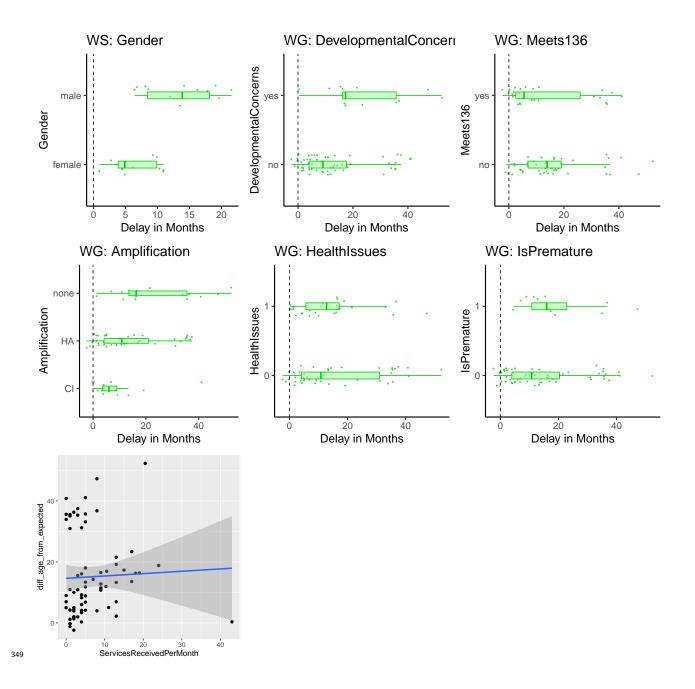


We next explored the effect of the different audiological, demographic, and intervention characteristics on vocabulary delay. Vocabulary delay did not meet the assumption of normality, so we used non-parametric tests for the following set of analyses.

Mann-Whitney-Wilcoxen tests were conducted to examine the effects of gender, laterality, developmental delay, health issues, prematurity, meeting 1-3-6 guidelines, and communication on vocabulary delay. We used kruskal-wallis tests for amplification and etiology, and Kendall's rank correlations for degree of hearing loss (worse ear) and services

received per month. These results are exploratory and descriptive, and their interpretation should be tempered accordingly.

Boys were significantly more delayed than girls on Words and Sentences but not Words and 334 Gestures. Children with developmental delays had larger vocabulary delays than children 335 without developmental delays on Words and Gestures. Because only one child with a 336 developmental delay took the Words and Sentences form, we did not perform the analysis for 337 Words and Sentences. Premature children and children with health issues had smaller 338 vocabularies than typically developing children on Words and Gestures but not Words and 339 Sentences. Children who met 1-3-6 guidelines had larger vocabulary than children who did 340 not on Words and Gestures but not Words and Sentences. On Words and Gestures but not 341 Words and Sentences, receiving more early intervention services was correlated with lower 342 vocabulary. We did not observe an effect of laterality, communication, degree, or etiology on vocabulary delay on either form of the CDI. For communication, we omitted cued speech from the analysis because only one child in our sample used this method of communication (shown on graph anyway for the curious). A kruskal-wallis test showed a significant effect of 346 amplification on vocabulary delay on Words and Gestures, such that children with no 347 amplification were more delayed than children without amplification. 348



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Variable	WG mean delays	WS mean delays	Method
Gender	Boy: 17.4282231239903; Girl: 12.0178307061822	Boy: 13.664443212515; Girl: 6.33347198690866	wilcox
Laterality	Unilateral: 13.2541771845055; Bilateral: 15.6575571166758	Unilateral: 7.71376632834074; Bilateral: 11.0552848390578	wilcox
Amplification	CI: 8.66855920541013; HA: 14.1473730177474, none: 23.0268553291739	CI: 19.6993965234822; HA: 8.13596277946654, none: 10.334020268944	kruskall
Health Issues	Yes: 14.4516825480366; No: 15.5235692630061	Yes: 8.18302524532061; No: 10.2696664076695	wilcox
Developmental Delay	Yes: 23.9379314219013; No: 13.1386097911128	Yes: 4.50279731078281; No: 10.0752357112171	wilcox
Prematurity	Premature: 19.3071057659814; Full-term: 14.263511986059	Premature: 8.91433657458079; Full-term: 9.98024748437499	wilcox
1-3-6 Guidelines	Meets: 12.7232472082048; Does not meet: 16.6412320183725	Meets: 10.753214167868; Does not meet: 9.42839653159108	wilcox
Communication	Spoken Language: 13.7969242301088; Total Communication: 21.1886273333096	Spoken Language: 10.1855168235174; Total Communication: 6.02078371954592	wilcox
Etiology	SNHL: 14.460852577098; Mixed: 18.7880795816761, Conductive: 16.4389485679933	SNHL: 8.52128484600432; Mixed: 15.7684319648513, Conductive: 8.04744160078312	kruskall
Degree	More severe: 15.2467016418905; Less severe: 15.2444169215399	More severe: 10.7924137424103; Less severe: 9.46051671496964	wilcox
Services Received Per Month	More services: 17.6409473630286; Less services: 13.7499928411619	More services: 12.4545331108601; Less services: 9.07717339077494	wilcox

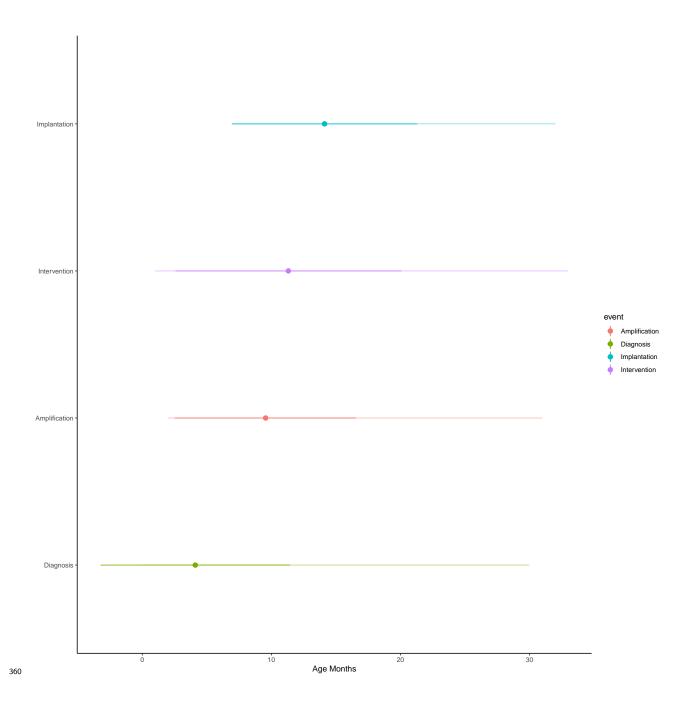
\end{table}

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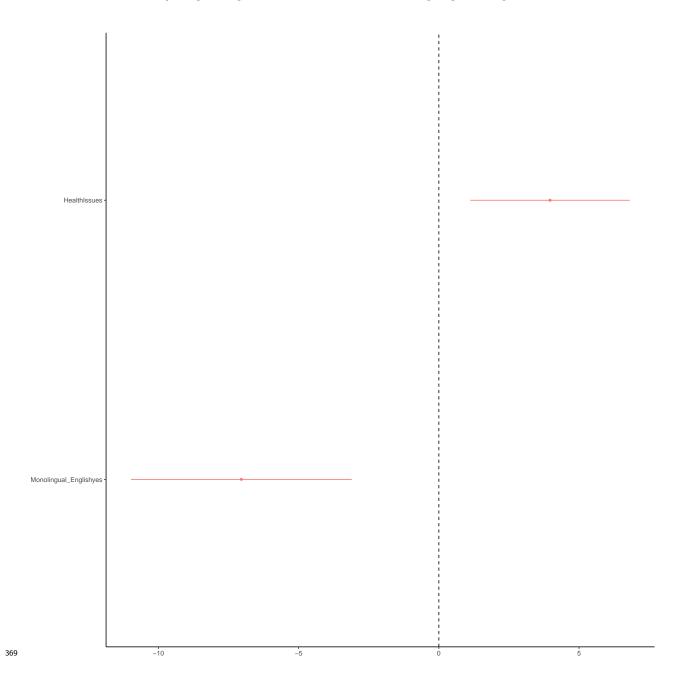
Part III: Meets136 success

Lastly, we looked at the ages at which children received diagnosis and intervention, and how this mapped onto the 1-3-6 guidelines. 36.84% of our sample met 1-3-6 guidelines for early diagnosis and intervention. Of children with comorbidities (i.e., developmental concerns, prematurity, health issues, vision loss), only 22% met 1-3-6 guidelines, compared to 45.76% of typically developing children. See table XXX for complete breakdown.



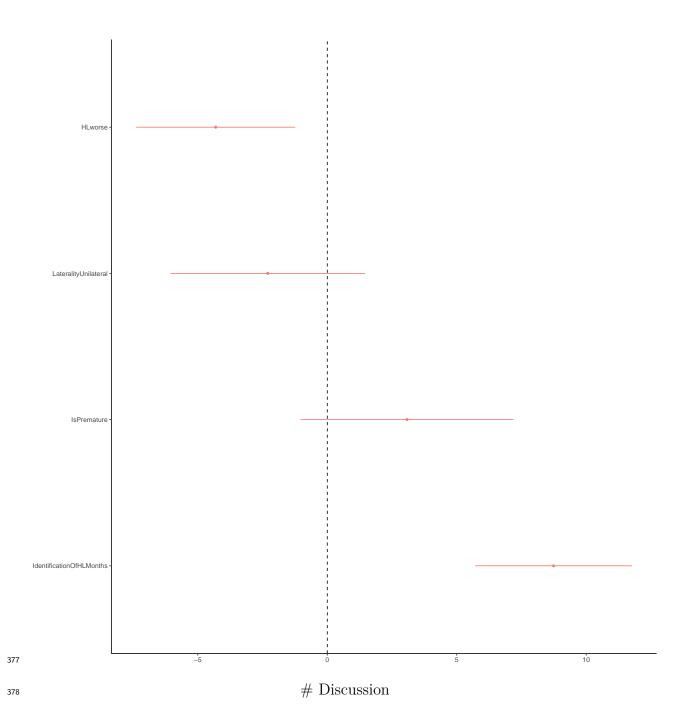
We created linear regression models for age at diagnosis and age at intervention. Models were paired down using stepwise regression by AIC using the stepAIC function (cite MASS package). For age at diagnosis, we included the set of child-specific factors that would be relevant before diagnosis of hearing loss. We began with: Age diagnosis \sim gender + laterality + degree (worse ear) + developmental delay + health issues + prematurity + laterality + language background + etiology The best fit model (R2=0.19 , p=0.00)included health

issues (beta weight = 4.14, p = 0.01) and language background (beta weight = -6.96, p = 0.00). Age diagnosis \sim health issues + language background



For age at intervention, we first included the variables potentially relevant prior to intervention: Age intervention \sim gender + degree (worse ear) + developmental delay + health issues + prematurity + laterality + language background + etiology + age diagnosis The best fit model (R2=0.41, p=0.00) included prematurity (beta weight = 4.05, p = 0.06),

laterality (beta weight = -2.44, p = 0.19), degree of hearing loss (beta weight = 3.96, p = 0.01), and age at diagnosis (beta weight = 0.62, p = 0.00). Age intervention \sim laterality + degree (worse ear) + prematurity + laterality + age diagnosis



Conclusion Conclusion

Footnotes: Despite exciting, increasing, and converging evidence for benefits of early sign
language exposure (e.g., Schick, De Villiers, De Villiers, & Hoffmeister, 2007; Clark et al.,
2016; Davidson, Lillo-Martin, & Pichler, 2014; Hrastinski & Wilbur, 2016; Magnuson, 2000;
Spencer, 1993), the majority of DHH children will not be raised in a sign language
environment. This is particularly true for North Carolina, which does not have a large
community of sign language users, relative to states like Maryland or areas like Washington
D.C. or Rochester, NY. For this reason, we focus on spoken language development.

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 $\label{thm:continuous} \begin{tabular}{ll} Table 1 \\ Summary of findings of CDI studies in DHH children \\ \end{tabular}$

Study	Population	Gender	1-3-6	Laterality	Degree	Amplification	Communication	Comorbidities
Ching et al., 2013	3 year old children receiving services in Australia	Female +	Did not study	Did not study	More severe -	No effect	No effect	Comorbidities -
Yoshinaga-Itano et al., 2017	8-39 month children with bilateral hearing loss	No effect	1-3-6 +	Did not study	More severe -	Did not study	Did not study	Comorbidities -
Yoshinaga-Itano et al., 2018	Children with cochlear implants	Did not study	1-3-6 +	Did not study	Did not study	Earlier CI activation +	Did not study	Did not study
De Diego-Lazaro et al., 2018	Spanish speaking children with bilateral hearing loss	No effect	Earlier intervention +	Did not study	Milder +	More functional hearing +	Did not study	Did not study
Vohr et al., 2011	18-24 month olds with hearing loss	Did not study	Earlier intervention +	Did not study	Milder +	Did not study	Did not study	NICU stay -; Comorbidities -

a + equals bigger vocab, - equals smaller vocab

Table 2

CDI details

CDI version	Average Age (SD)	Average Comprehension (SD)	Average Production (SD)	% Developmental Delays
WG (n=73)	20.07 (8.87) months	107 (99.9) words	32 (53.7) words	17.81%
WS (n=25)	26.05 (8) months	NA	132 (172.7) words	4%