Speech intelligibility measurement

A latent variable approach on utterances' transcriptions

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Abstract

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1. Introduction

Intelligible speech can be defined as the extent to which the elements in an speaker's acoustic signal, e.g. phonemes or words, can be correctly recovered by a listener [47, 73, 70, 36]. Because intelligible spoken language requires all core components of speech perception, cognitive processing, linguistic knowledge, and articulation to be mastered [36], its attainment carries an important societal value, as it is a milestone in children's language development, the ultimate checkpoint for the success of speech therapy, and has been qualified as the 'gold standard' for assessing the benefit of cochlear implantation [13].

The literature suggest two perspectives from which *speech intelligibility* can be assessed: the message and listener's perspective [4, 5]. The first, also known as acoustic studies, is focused on assessing separately particular characteristics of the speech samples, e.g. their pitch, duration or stress (supra segmental characteristics), or the articulation of vowels and consonants (segmental characteristics) [63]. Whereas the second, also known as perceptual studies, is centered on making holistic assessments of the speech stimuli, e.g. measure their perceived quality [4, 5]. On both instances, the stimuli (children's utterances) can be generated from reading at loud, contextualized utterances, or spontaneous speech tasks[1].

Furthermore, perceptual studies can use multiple approaches to measure intelligibility, but they can be largely grouped into two: objective and subjective ratings [45]. In *objective rating* methods, listeners transcribe the children's utterances orthographically or phonetically, and use such information to construct a score. In that sense, in the transcription task, intelligibility can be inferred from the extent a set of transcribers can identify the word contained in an utterance [5]. In contrast, under *subjective rating* methods, listeners directly infer the utterance's intelligibility score through specific procedures, e.g. absolute holistic, analytic, or comparative judgments, among others.

It is easy to deduce that *objective rating* methods produce more valid[2] and reliable[3] scores than its *subjective* counterpart, and as a result, are usually used as an objective measure of intelligibility [5, 27].

Accompanying the intelligibility assessment methods, the literature supply a myriad of factors that are thought also contribute to the (under)development of intelligible spoken language [57, 6, 40, 28]. Among these are audiology related factors, such chronological age, age at implantation, the duration of device use, *hearing age*, bilateral or contralateral cochlear implantation, and the children's preoperative and postoperative hearing levels. On the other hand, there are also child related factors, such as the cause of the hearing impairment (genetic, infections), additional disabilities (mental retardation, speech motor problems), and gender. Finally, there are also environmental factors, such as communication modality.

Considering all of the above, this paper seeks to investigates the speech intelligibility levels of normal hearing (NH) versus hearing-impaired children with cochlear implants (HI/CI). For that purpose, ten utterances recordings, from thirty two NH and HI/CI children, were selected from a large corpus of *spontaneously spoken speech* collected by the CLiPS research center. Additionally, we set up an experiment, where one hundred language students transcribed each stimuli to the Qualtrics environment [77]. Finally, the transcriptions were transformed into an entropy measure per utterance, which served as our outcome variable.

We believe this paper make three specific contributions to the understanding of the factors that drive the intelligibility of spoken language. First, we develop a novel analysis using a latent variable approach [26]. More specifically, we model *speech intelligibility* as a latent variable that can be inferred from the entropy replicates. This method offers three specific benefits. On the one hand, the method 'constructs' an intelligibility score, which in turn, allow us to test different hypothesis and even make individual comparisons at the appropriate level. On the other hand, it allow us to control for different sources of variation. This is particularly important as, by failing to account for the appropriate hierarchies in the data, we could be 'manufacturing' false confidence in the parameter estimates, leading us to incorrect inferences [53]. Finally, the method also provides a 'criterion' on how reliable are the entropy replicates to measure speech intelligibility.

Second, we use Directed Acyclic Graph (DAG) [59, 17] to depict all the relevant variables though to influence speech intelligibility. We describe in detail our causal and non-causal hypothesis, and supplement our description with a causal diagram. The benefit of the method lies, not only, in that it makes the assumptions of our hypothesis more transparent, but also allow us to derive statistical procedures from the aforementioned causal assumptions [53, 79, 62].

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[1]ordered on increasing level of ecological validity [32, 25]

[2]the extent to which scores are appropriate for their intended interpretation and use [50, 68].

[3]the extend to which a measure would give us the same result over and over again [68], i.e. measure something, free from error, in a consistent way.

Third and final, we wrap the analysis procedure under the Bayesian framework, providing the assumptions, and the steps required to reproduce the computational implementation of the models.

2. Materials and Methods

We set up an experiment where speech samples were transcribed by a group of listeners. The current section succinctly describes the participating children, the stimuli used, and the experimental setup, while also delve into the causal and statistical framework of analysis.

2.1. Children

Thirty two children were selected using a large corpus of *spontaneously spoken speech*, collected by the Computational Linguistics, Psycholinguistics and Sociolinguistics research center (CLiPS). The selection followed a two step procedure [27]. First, a sample of sixteen hearing-impaired children (ten boys, six girls) were selected based on the quality of their registered stimuli. Second, an additional matched sample of sixteen normal hearing children was also selected (six boys, ten girls), and served as a comparison group.

For the first group, all the hearing-impaired children with cochlear implants (HI/CI) were native speakers of Belgian Dutch, living in Flanders, the Dutch speaking area of Belgium. They were all raised orally using monolingual Dutch, with a limited support of signs. All of the children were screened by the Universal Neonatal Hearing Screening (UNHS), using an automated auditory brainstem response hearing tests for newborns, and all received the cochlear implantation before the age of two. Their medical and audiological records did not ascertain any additional health or developmental issues. Hence, no known additional comorbidities were though present. Finally, at the date of the measurement, they were all enrolled in the mainstream educational system.

For the second group, the sixteen normal hearing children (NH) were closely matched to the HI/CI group based on chronological age. All children were also native speakers of Belgian Dutch, and enrolled in the mainstream educational system. None reported hearing loss or additional disabilities, judged from the UNHS screening procedure and their respective parental report.

The characteristics of the selected children is detailed in Table 3, in the supplementary section A.1.

2.2. Stimuli

The stimuli consisted of children's utterances, i.e. sentences of similar length, recovered from previously mentioned CLiPS corpus. More specifically, we use a portion of the corpus that consisted of ten utterances recordings for each of the thirty two selected children, adding to a total of 320 stimuli.

The stimuli were documented when the child was telling a story cued by the picture book 'Frog, where are you' [52] to a caregiver 'who does not know the story'.

The recordings were orthographically transcribed with the CLAN editor in CHAT format [51]. The quality of the stimuli was ensured by selecting utterances with no syntactically ill-formed or incomplete sentences, any background noise, cross-talk, long hesitations, revisions or non-words [5]. The aforementioned transcriptions were used only in the selection process of the stimuli for the experiment.

2.3. Experimental setup

The experiment was setup to perform a transcription task in the Qualtrics environment [77]. One hundred language students from the University of Antwerp participated. The participants were native speakers of Belgian Dutch, without any particular experience with the speech of hearing-impaired children.

The participants and stimuli were divided into five groups, where each group of 20 students transcribed 64 stimuli on their series. The stimuli were presented to the listeners in a random order. As a result, the setup produced 20 transcriptions per utterance, adding to a total of 6400 transcriptions. The steps that comprised the task are detailed in the supplementary section A.2.1.

The data resulting from the transcription task was then processed and converted into one entropy measure per utterance (H), which served as our outcome variable.

The entropy measure is bounded in the unit interval $[0, 1]$, and it was used as a quantification of (dis)agreement between listeners' transcriptions: utterances yielding a high degree of agreement between transcribers were considered highly intelligible, and therefore, registered a lower entropy ($H \to 0$). In

contrast, utterances yielding a low degree of agreement were considered as exhibiting low intelligibility, and therefore, registered a higher entropy $(H \rightarrow 1)$ [5, 27]. The procedure followed to calculate the entropies is detailed in the supplementary section A.2.2.

2.4. Causal framework

The analysis was informed by a preliminary work aimed at describing the causal and non-causal factors influencing speech intelligibility. More specifically, the current research uses a Directed Acyclic Graph (DAG) [59, 17] to describe all the relevant variables though to influence intelligibility. A DAG is a type of *structural causal model* that can be represented, among other ways, by a *causal diagram*.
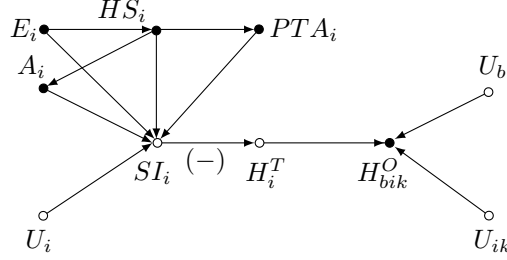

Figure 1: DAG: causal diagram describing the relationships among the analyzed variables. Open circles describe latent/unobserved variables, full circles describe observed variables.

Figure 1 shows the *causal diagram* for our research hypothesis. In the figure, H^O_{bik} denote the *observed* entropy replicates nested within children and experimental blocks, where $k = 1, \ldots, 10$ utterances, $i = 1, \ldots, 32$ children, and $b = 1, \ldots, 5$ blocks. Moreover, H^T_i and SI_i denotes the child's *true* entropy and speech intelligibility scores, respectively. In addition, HS_i denotes the children's hearing status group, A_i their *hearing age*, PTA_i their unaided post-implant pure tone average, and E_i the etiology of the disease that led to the hearing impairment.

Three main features can be emphasized from the figure. First, the children's *speech intelligibility* and *true* entropy scores are thought to be latent/unobservable variables [26] (drawn with open circles, see supplementary section A.3 about the appropriate interpretations of the scores). The figure also shows the scores are thought to be inferable from the *observed* entropy replicates. More specifically, we are asserting the *observed* entropy replicates H^O_{bik} represent multiple realizations of a child's *true* entropy H^T_i. Finally, as expected from our the theory, we assume the *intelligibility* score SI_i is inversely/negatively related to the *true* entropy H^T_i, i.e. the lower the intelligibility the higher the entropy, and vice versa.

Second, the figure reflects the expected multilevel structure of variability present in our data. This is particularly important as, by failing to account for the appropriate dependencies in the data, we could be 'manufacturing' false confidence in the parameter's estimates, leading us to incorrect inferences [53].

Based on the experimental setup described in section 2.3, we anticipated the ten utterances, originated from each of the thirty two children, were also observed within a group of transcribers (series) assigned to the observation, i.e. our data has a hierarchy with children, replicates and block levels (U_i, U_{ik} and U_b, respectively).

Considering the latter, we expect that if the experiment was 'set up right', the block random effects would explain a small amount of variability in the data, and its inclusion/exclusion in the model would not change the parameter estimates. Moreover, we expect a larger variability between children's *speech intelligibility*, at least larger than the block random effects. Several evidence suggest this is particularly true among HI/CI children [80, 60, 54, 11, 78, 58, 36]. Finally, we did not had any comparable expectation for the variability in the replicates, as this feature has not been investigated before.

Third, the figure shows the assumed relationship among the relevant variables [57, 6, 40, 28], and how these influence the children's intelligibility of speech. Furthermore, it also reveals we assume the variables are independent, beyond the described relationships. Here follows a description of our causal hypothesis for the relevant variables.

For **hearing status** (HS_i), it is clear that its inclusion directly corresponds with the main purpose of the current research endeavor. i.e. compare the *speech intelligibility* among NH and HI/CI children. Yet, we do not have a clear expectation about the levels among the groups. In terms of intelligibility, previous literature suggest that some HI/CI children manage to 'catch up' with their NH counterparts

[74, 42, 7, 37, 9, 21, 75], implying the two group attain similar levels. However, other studies also indicate HI/CI children never reach their NH analogues [56, 11, 14, 38, 36, 24, 41], therefore implying different levels of *intelligibility* among the groups.

For **Hearing age** (A_i), we expect the variable to be one of the main responsible for the increase in children's *speech intelligibility*, no mater the children's hearing status [5]. Several studies provide evidence that *intelligibility* increases with age [15, 16, 32, 33, 2, 8, 45]. *Hearing age* is a composite variable constructed by combining the *chronological age* for the NH group, and the *device length of use* for the HI/CI group [27] (see table 3 in supplementary section A.1). The variable tries to approximate the amount of time a child has been actively hearing and developing his(her) language.

However, no short of evidence has been presented in favor of using others surrogate measures, like *chronological age* [34, 42, 41] or *age at implantation* [57, 6, 9, 21]. We argue that the feasibility of using any other proxy measure, largely depends on the assumed reliability of the surrogate, to approximate the variable of interest. In that sense, although we recognize *hearing age* is not a 'perfect' proxy [27], we argue it is the most appropriate to test our hypothesis, based on the relevant literature and its assumed reliability to capture children's language development (although the latter has not been tested). Furthermore, we believe the variable serve two additional purposes: (i) control for sampling bias (expanded in supplementary section A.4), and (ii) de-confound the parameter estimates of *hearing status* [17].

Lastly it is important to highlight that, for modeling purposes, using more than one of the aforementioned proxies in tandem is not recommended. It is apparent from the previous description, the three surrogate measures share high similarities in their data construction. This in turn, could cause problems in the modeling procedure, as including variables that provide 'similar information' might lead to a statistical problem known as multicollinearity, in which our estimates get biased and less precise [29], leading us to incorrect inferences and conclusions.

For **pure tone average** (PTA_i), we expect it to have a small or null effect on *speech intelligibility*, as the empirical evidence seem to suggest [5]. *Pure tone average* is the child's subjective hearing sensitivity, aided or unaided, by their hearing apparatus.

However, beyond the empirical evidence, the variable was included for two additional reasons. First, given that previous modeling efforts did not capture the full data hierarchy, it is possible that the effects of PTA on *speech intelligibility* has been largely overlooked. Although evidence seem to suggest the variable has no effect, it is also sensible to think that HI/CI children with severe hearing loss, as accounted by the variable, might develop their language at a slower rate. This is especially true, if we consider the signal provided by the cochlear implant is still degraded compared to normal hearing scenarios [22]. Lastly, the second reason for its inclusion lies in the possibility that the variable might be useful to de-confound the parameter estimates of *hearing status* [17], a purpose shared with the *hearing age* variable.

In the case of the **Etiology** of the disease that led to the hearing impairment (E_i), we expect it to have a differential effect on *speech intelligibility*, within the HI/CI group. However, since the severity of the etiology cannot be easily ascertain nor ordered, we cannot foresee the direction of such effects, i.e. genetic factors not necessarily lead to worse levels of language development and intelligibility, than factors related to infections.

The previous assumption does not have a correspondence with the empirical evidence, were the variable was deemed unimportant [5]. However, we argue that it is possible its effects have also been largely overlooked, due to the lack of statistical control on the data variability hierarchy, similar to the PTA case. Moreover, as with its predecessors, the variable might also be useful to de-confound the parameter estimates of *hearing status* [17], assuming our DAG is appropriate.

Finally, it is important to highlight the reason for the absence of other variables, deemed relevant by the literature, in our causal hypothesis.

For the **type of cochlear implantation**, i.e. bilateral or contralateral, the variable was not included because we did not expect it to be related to other variables in the DAG, i.e. the decision on receiving one or the other is solely based on the intelligibility outcome, no matter how the latter is measured. This in turn means that its inclusion/exclusion would not confound our estimates. Additionally, given that most of the children underwent through sequential bilateral implantation (eleven in total), we anticipated the effect of variable already permeates the HI/CI sample.

For the case of **additional disabilities**, e.g. mental retardation or speech motor problems, given that the children did not reported any additional comorbidities, the variable was deemed unnecessary.

In the case of **environmental factors**, such as communication modality, the variable was also deemed unnecessary, given that HI/CI children were raised orally using monolingual Dutch with a limited support of signs, a scenario similar to the NH group (see section 2.1).

Last but not least, ***gender*** was not included in our hypothesis, as no theoretical nor empirical evidence have been found on its effects [5].

As expected, it is possible that other unobserved confounding variables are not accounted by our assumptions, and therefore, our causal diagram. This is true for any type of social, behavioral and educational research. However, we argue that the additional transparency of our approach, and its ability to derive statistical procedures from causal assumptions, is its main strength [53, 79, 62].

2.5. Statistical analysis

Using the DAG, described in the previous section, we describe the algebraic formalism for our multiple Bayesian probabilistic models [46]. Each model targeted a different manner in which our research hypothesis could be investigated.

In general terms, every model was composed of two parts: a latent measurement model [26] and a structural equation model [44]. For the former, we represented the child's *speech intelligibility* as a latent variable, inversely related to the child's latent *true* entropy. Moreover, we modeled the latter as a variable that can be inferred from the observed entropy replicates in the following way:

$$H_{bik}^{O} \sim \text{BetaProp}\left(\bar{H}_{bi}, M_i\right) \tag{1}$$

$$\bar{H}_{bi} = \text{logit}^{-1}(a_b - SI_i) \tag{2}$$

$$H_i^T = \text{logit}^{-1}(-SI_i) \tag{3}$$

where BetaProp(μ, θ) defines the beta-proportion distribution with parameters μ and θ [30, 48]; with $\mu \in [0, 1]$ and $\theta > 0$. Furthermore, $\bar{H}_{bi}$ represented the average entropy measure nested within blocks and children ($b = 1, \ldots, 5$ and $i = 1, \ldots, 32$, respectively), while M_i denotes the 'sample size' of the distribution, possibly nested within children. Furthermore, H_{bik}^O, H_i^T and SI_i denotes the children's entropy replicates, *true* entropy and *speech intelligibility* score, respectively. Finally, a_b represents the block random effects, and $\text{logit}^{-1}(x) = \exp(x)/(1 + \exp(x))$ the inverse-logit transformation.

From the previous algebraic structure we can notice. First, we modeled the average entropy of the replicates with $\mu = \bar{H}_{bi}$, which is a non-linear transformation of the block random effects and the children's *speech intelligibility*. Second, the children's *true* entropy H_i^T is inversely and non-linearly related to the *speech intelligibility* SI_i. Third and last, we captured the variability of the entropy replicates using $\theta = M_i$. See supplementary section A.5.1 for a detailed overview of the implications of using this approach.

For the second component we used a structural model [44]. In this portion, different iterations of our research hypothesis were proposed. Hereby we present the general representation, from which the others can be derived:

$$SI_i = a_i + \alpha + \alpha_{E[i], HS[i]} + \beta_{A, HS[i]}(A_i - \bar{A}) + \beta_{P, HS[i]} sPTA_i \tag{4}$$

where E_i and HS_i are defined as in the previous section, A_i is the *hearing age* in years, while $sPTA_i$ is the standardized version of the unaided post-implant pure tone average PTA_i. Moreover, a_i denoted the children's random intercepts, and α the fixed intercept. On the other hand, $\alpha_{E[i], HS[i]}$ denoted the *intelligibility* levels per etiology and hearing status group, while $\beta_{A, HS[i]}$ denoted the evolution of *intelligibility* per unit of *hearing age*, for each hearing status group. Finally, $\beta_{P, HS[i]}$ described the evolution of *intelligibility* per unit of the standardized pure tone average, for each hearing status group.

In the previous algebraic structure it is important to highlight, all parameters are estimated in the logit scale and centered at $sPTA_i = 0$ and $\bar{A} = 5$, the latter denoting the minimum *hearing age* in the sample. This is done to facilitate the interpretation of the parameters.

Thirteen statistical models were derived from the previous general description. Each model expressed one specific way in which our research hypothesis could be investigated. Table 1 provides an overview of the parameterization for the full set of proposed models. The models were characterized by two aspects: (i) the use of interactions and the selection of interacting variables, and (ii) the 'sample size' set to capture the replicates variability. For the former, the options were: intercept only (model 1), and multivariate regression with no interactions (models 2 to 4), with age and hearing status interaction only (models 5, 8, and 11), with etiology and hearing status interaction only (models 6, 9, and 12), and a full interaction model comprising the previous two (models 7, 10 and 13). In a similar fashion, for the latter the options were: one fixed 'sample size' (models 1, 2, 5, 6, and 7), one estimated 'sample size' (models 3, 8, 9, and

10), and finally, one 'sample size' per child (models 4, 11, 12, and 13), which we dubbed *robust* models (see supplementary section A.5.2, where we expand about the need for *robust* models).

Notice all proposals used block and children's random effects (a_b and a_i, respectively), as well as the fixed effect intercept (α). Moreover, the models with one fixed 'sample size' used a value of ten, corresponding to number of utterances per child (see supplementary section A.5.1 on its implications). Finally, all models contemplated the interaction of the standardized pure tone average and the hearing status. The latter was imposed, because the collection of data did not contemplated the measurement of the pure tone average for the NH group. Therefore, the only interpretable parameter is the one estimated for the HI/CI children.

It is important to highlight that for all model implementations, the visual inspection of trace, trace-rank and autocorrelation plots was performed. Additionally, we also evaluated the Gelman-Rubin diagnostic and effective number of samples [39]. In all cases, the plots and statistics indicated the parameters achieved convergence, good mixing and lack of serial autocorrelation, all necessary requirements to be able to interpret the model estimates [53].

Additional details concerning the Bayesian implementation of the models, such as the representation of variability in a beta-proportion distribution, the parameters' priors and hyper-priors, the estimation procedure, the data pre-processing phase, and simulation studies are expanded in the supplementary section A.5.

Model	Name	Parameters						
		M_i	a_b	a_i	α	$\alpha_{E[i],HS[i]}$	$\beta_{A,HS[i]}$	$\beta_{P,HS[i]}$
1	Intercept only (fixed 'size')	10	a_b	a_i	α	$-$	$-$	$-$
2	No interaction (fixed 'size')	10	a_b	a_i	α	$\alpha_{HS[i]}$	β_A	$\beta_{P,HS[i]}$
3	No interaction (one 'size')	M	a_b	a_i	α	$\alpha_{HS[i]}$	β_A	$\beta_{P,HS[i]}$
4	No interaction (robust)	M_i	a_b	a_i	α	$\alpha_{HS[i]}$	β_A	$\beta_{P,HS[i]}$
5	Age interaction (fixed 'size')	10	a_b	a_i	α	$\alpha_{HS[i]}$	$\beta_{A,HS[i]}$	$\beta_{P,HS[i]}$
6	Etiology interaction (fixed 'size')	10	a_b	a_i	α	$\alpha_{E[i],HS[i]}$	β_A	$\beta_{P,HS[i]}$
7	Full interaction (fixed 'size')	10	a_b	a_i	α	$\alpha_{E[i],HS[i]}$	$\beta_{A,HS[i]}$	$\beta_{P,HS[i]}$
8	Age interaction (one 'size')	M	a_b	a_i	α	$\alpha_{HS[i]}$	$\beta_{A,HS[i]}$	$\beta_{P,HS[i]}$
9	Etiology interaction (one 'size')	M	a_b	a_i	α	$\alpha_{E[i],HS[i]}$	β_A	$\beta_{P,HS[i]}$
10	Full interaction (one 'size')	M	a_b	a_i	α	$\alpha_{E[i],HS[i]}$	$\beta_{A,HS[i]}$	$\beta_{P,HS[i]}$
11	Age interaction (robust)	M_i	a_b	a_i	α	$\alpha_{HS[i]}$	$\beta_{A,HS[i]}$	$\beta_{P,HS[i]}$
12	Etiology interaction (robust)	M_i	a_b	a_i	α	$\alpha_{E[i],HS[i]}$	β_A	$\beta_{P,HS[i]}$
13	Full interaction (robust)	M_i	a_b	a_i	α	$\alpha_{E[i],HS[i]}$	$\beta_{A,HS[i]}$	$\beta_{P,HS[i]}$

Table 1: Proposed statistical models. Parameterizations.

3. Results

3.1. About our hypothesis

The current research used the Information-Theoretic Approach [1, 12] for model selection and inference. The application of the approach required: (i) the expression of the research hypothesis into statistical models, (ii) the selection of the most plausible models, and (iii) to produce inferences based on one or multiple selected models. The first requirement of the approach is covered in sections 2.4 and 2.5, and expanded in supplementary section A.5. Here we use the results of the second requirement, detailed in the supplementary section A.5.6, to produce the final inferences.

As detailed in the supplementary section, the final conclusions of our research will be drawn from the comparisons of two models: (i) a no interaction model, with one estimated 'sample size' (model 3), and (ii) a full interaction model, with one estimated 'sample size' (model 10). The former is selected as it is the model with highest probabilistic support. The latter is considered because it encompasses the remaining highest supported models. Furthermore, no *robust* model is inspected, as we prefer a more parsimonious depiction of our hypothesis. Table 2 summarizes the parameters posterior estimates and contrasts of interest, for the aforementioned models.

| | | | CI | | HPDI | | | |
Parameter	Mean	SD	2.5%	97.5%	2.5%	97.5%	n eff.	Rhat
Model 3: No interaction (one 'size')								
parameters:								
a	0.179	0.189	-0.197	0.554	-0.198	0.552	3'677.705	1.001
bP[2]	-0.117	0.166	-0.439	0.207	-0.440	0.202	1'738.855	1.000
bA	0.432	0.141	0.154	0.715	0.170	0.723	1'815.243	1.001
aHS[1]	0.284	0.235	-0.178	0.761	-0.203	0.728	2'719.833	1.000
aHS[2]	0.116	0.217	-0.303	0.537	-0.304	0.537	2'646.671	1.000
constrast:								
aHS[2]-aHS[1]	-0.168	0.246	-0.661	0.371	-0.650	0.324	n.a.	n.a.
Model 10: Full interaction (one 'size')								
parameters:								
a	0.217	0.179	-0.142	0.562	-0.118	0.580	3'902.629	0.999
bP[2]	-0.122	0.173	-0.457	0.220	-0.460	0.216	1'659.639	1.002
bAHS[1]	0.435	0.157	0.127	0.745	0.123	0.741	1'477.460	1.000
bAHS[2]	0.237	0.177	-0.119	0.596	-0.089	0.615	1'844.785	1.001
aEHS[1,1]	0.183	0.278	-0.358	0.726	-0.349	0.729	2'524.451	1.000
aEHS[2,2]	0.212	0.241	-0.259	0.684	-0.259	0.684	2'418.275	0.999
aEHS[3,2]	0.077	0.245	-0.402	0.547	-0.383	0.561	3'015.559	0.999
aEHS[4,2]	0.007	0.269	-0.522	0.530	-0.547	0.499	3'268.043	1.000
contrast:								
bAHS[2]-bAHS[1]	-0.197	0.208	-0.600	0.213	-0.590	0.223	n.a.	n.a.
aEHS[2,2]-aEHS[1,1]	0.029	0.352	-0.673	0.723	-0.714	0.682	n.a.	n.a.
aEHS[3,2]-aEHS[1,1]	-0.106	0.365	-0.823	0.607	-0.822	0.609	n.a.	n.a.
aEHS[4,2]-aEHS[1,1]	-0.176	0.380	-0.906	0.568	-0.928	0.535	n.a.	n.a.
aEHS[3,2]-aEHS[2,2]	-0.135	0.299	-0.719	0.453	-0.730	0.439	n.a.	n.a.
aEHS[4,2]-aEHS[2,2]	-0.205	0.344	-0.888	0.453	-0.902	0.430	n.a.	n.a.
aEHS[4,2]-aEHS[3,2]	-0.070	0.344	-0.744	0.612	-0.735	0.617	n.a.	n.a.

CI = compatibility interval

HDPI = highest posterior density interval

n eff. = effective number samples

Rhat = Gelman-Rubin diagnostic

n.a. = not available / not applicable

[1] = NH children, [2] = HI/CI children

[1,i] = NH children, [2,i] = genetic, [3,i] = CMV infection, [4,i] = unknown etiology

Table 2: Selected statistical models: results.

Before providing any parameter interpretation, it is important to highlight an statistical issue that will permeate all of our inferences. Given the large amount of variability registered at the children and replicates levels, the models will not be able to produce unequivocal null hypothesis rejections, i.e. CI and HPDI not including the zero value (expanded in section 3.2). However, from supplementary section A.5.5, we can be certain that our models can affirm the estimate values with at least 60% power, depending on the size of the estimates; i.e. our models can correctly estimate values different from zero, when such alternative hypothesis is true. Therefore we will continue interpreting the parameters, but the reader should cautious into understanding the CI and HPDI indicate that our data size still might not be enough, to ultimately define the direction of the effects, and further evidence might still be needed.

First, about ***hearing status***, model three reveals that HI/CI children have a modest lower level of *speech intelligibility*, compared to their NH counterparts (aHS[2]-aHS[1]). The result then seem to support previous evidence on the matter [56, 11, 14, 38, 36, 24, 41]. However, model ten paints a more nuanced story considering the ***Etiology*** of the disease. The contrasts of the model reveal that HI/CI children with genetic etiology manage to reach similar levels of *intelligibility* as NH children (aEHS[2,2]-aEHS[1,1]). Nevertheless, the same cannot be said when the hearing impairment is caused either by a

CMV infection (`aEHS[3,2]-aEHS[1,1]`) or other unknown causes (`aEHS[4,2]-aEHS[1,1]`).

About the former result, we theorize a intuitive explanation. Since hearing loss is a brain issue, rather than ear issue [31], as children with genetic causes grow developing their language using the hearing apparatus, they grow constructing effective auditory neural pathways that benefit from the new hearing input from the start, even when that input is slightly degraded [22]. However, the same cannot be said for other etiology status. What is hinted is that, a child with other etiology status might need to 're-wire' his(her) auditory neural connections in order to fully take advantage of the new hearing signal. This is particularly important, if we consider these children begin their life hearing normally, and then loosing the ability to process sound during an important maturimg stage of the brain cortex (first 12 months) [31]. This result is particularly important, as it might reveal the factors that caused the hearing impairment matter, even when the children receive the new hearing input signal at a young age.

In relation to **hearing age**, model three reveals what we initially expected: the factor is one of the main determinants for the increase in children's *speech intelligibility*, showing moderate effects (`bA`) [18, 64]. However, model ten offers preliminary evidence contrary to what was previously found [5], indicating there is a difference in the evolution of *intelligibility* between HI/CI and NH children (`bAHS[1]`, `bAHS[2]`, and `bAHS[2]-bAHS[1]`). We intuit the latter result have two possible complementary explanations, either: (i) the children develop their language differently at different stages of their (*hearing*) age [31], or (ii) given that the hearing signal provided by the apparatus is degraded [22], HI/CI children might take longer to achieve similar levels of *intelligibility* than their NH counterparts.

Finally, for **pure tone average**, we see both models support our initial hypothesis: HI/CI children with severe hearing loss, as accounted by the variable, develop their language at a slower rate than their NH counterparts (`bP[2]`). But more interestingly, this results seem to couple rather well with the second intuition described in the previous paragraph, i.e. the more degraded the signal a child receives, either because of the apparatus or the severity of their hearing-impairment, the slower the *intelligibility* evolves. Ultimately, notice from the perspective of effect sizes [18, 64], the estimates can be considered small in magnitude, but present nonetheless (as proved by our power analysis).
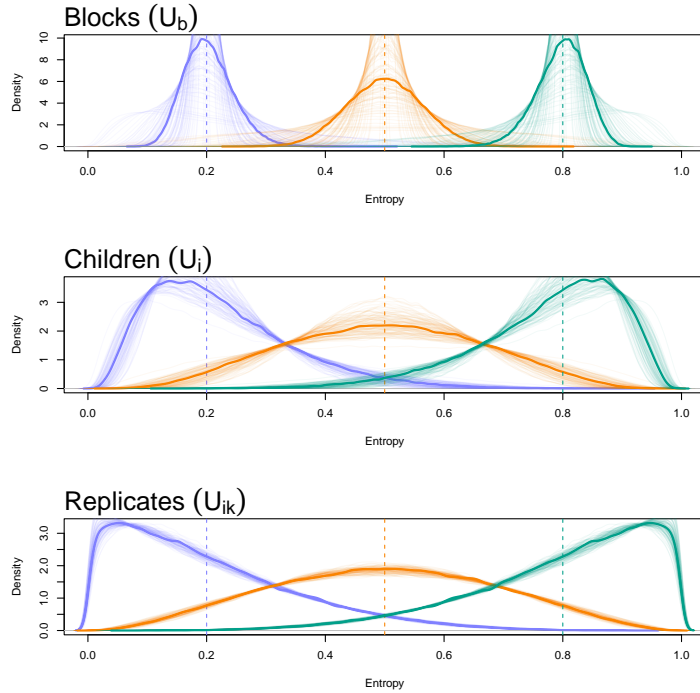

Figure 2: Posterior predictive: hierarchy of variability in the data. Distributions are plotted at the entropy replicates scale, considering three different average entropy values: $\mu = 0.2$, $\mu = 0.5$, and $\mu = 0.8$ (discontinuous lines). Thick solid lines represent the marginal distribution, thin solid lines depict 100 random posterior samples.

3.2. About the hierarchy of variability

As expected, three hierarchies of variability were present in our data: the blocks and children's random effects, as well as, the variability of the entropy replicates.

Evidence from the posterior estimates reveal the block random effects explained a small amount of variability in the data (top panel of figure 2), and its inclusion/exclusion in the model did not change the parameter estimates. The previous implies the experiment was correctly 'set up', as the series in which the utterances were transcribed did not explain a significant amount of variation, nor its exclusion biased the parameter estimates.

On the contrary, we observe a significantly larger variability between children's *speech intelligibility*. More precisely, more than three times the block effects variability (middle panel of figure 2). The previous corroborated preliminary evidence on the matter [80, 60, 54, 11, 78, 58, 36, 5]. Additionally, it implies that given such a large amount of between variability, the statistical models might have a harder time delimiting the *hearing status* groups location on the intelligibility scale.

Finally, for the variability of the entropy replicates, the posterior estimates reveal a reasonable finding: the amount of variability at the replicates level is even larger than the one observed at the children level (bottom panel of figure 2). The latter implies there is significant error in measuring speech intelligibility with the entropy replicates. This further emphasize the difficulty of producing unequivocal inferences, in respect to the intelligibility levels of the *hearing status* groups.
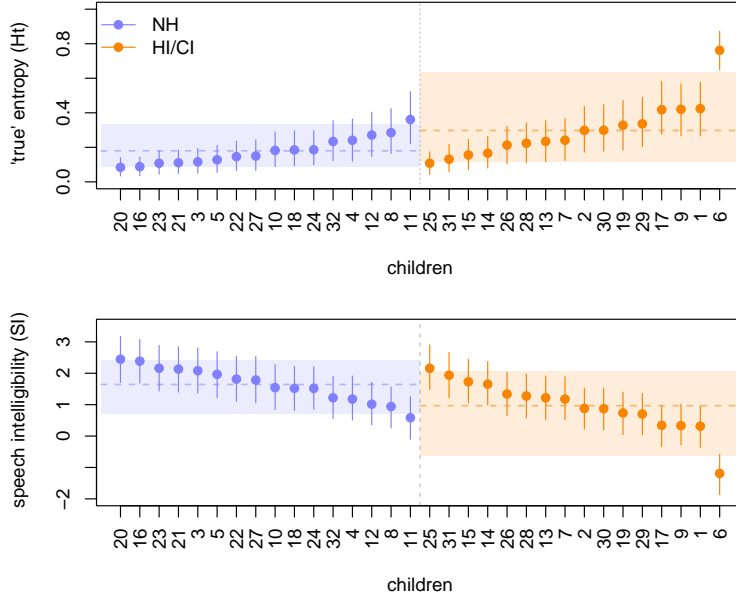

Figure 3: Model 10, posterior predictive: *true* entropy and *speech intelligibility* scales. Colored circles represent mean values per *hearing status* group. Thin vertical continuous lines depict 95% highest posterior density intervals (HDPI). Thin horizontal discontinuous lines represent the marginal average for the *hearing status* group. Shaded area represent the 95% confidence interval for the marginal averages.

3.3. The speech intelligibility scale

As previously stated, one of the main benefits of the current methodology is that we can 'construct' the *intelligibility* score of the sampled children, which in turn, allow us to test different hypothesis and even make individual comparisons at the appropriate level. See supplementary section A.3 for the appropriate interpretation of the scores.

Figure 3 depicts the estimated *true* entropy and *speech intelligibility* scores per child. Three points can be highlighted from the figure. First, from the horizontal discontinuous lines, we can clearly assess that on average NH children have a modest higher level of intelligibility (lower entropy) than the HI/CI counterparts. However, given the children's variability, we cannot reject the statistical hypothesis that

HI/CI have similar levels than the NH children, as one can notice from the overlapping shaded areas of both groups.

Second, the figure reveals there is some inherited uncertainty on the score estimates, resulting from using entropy replicates as a tool for measuring *speech intelligibility*. The latter can be observed from the vertical lines, describing the highest posterior density intervals for the *intelligibility* of each child. We argue this last trait is one of the main strengths of the method, as it allow us to not only make individual comparisons, but also consider their inherited (un)certainty.

Third and final, the non-linear relationship between *speech intelligibility* and the *true* entropy scale is clearly noticed in the HPD intervals, where lowest/highest entropy values are paired with narrower intervals (see top panel of figure 3). This behavior is expected under non-linear statistical models. But more importantly, under our particular model, it also means that we can be more certain of the children's *true* entropy values at the extremes (either zero or one), rather than in the middle part of the scale. Good examples of the former are found in child 20, and 25 at the lowest part of the scale, and on child 6 at the highest end.

3.4. Posterior predictive

The posterior predictive results give us with a tool to assess how good our models were to reconstruct the observed data. Figure 4 shows our full interaction model managed to 'recreate' our data, while also provided the (un)certainty around the entropy replicates. The latter was shown in two ways: (i) through the point estimates and HPD intervals for the *true* entropies (red points and horizontal lines), and (ii) though the expected estimated marginal and sampled distributions (thick and thin solid distribution lines, respectively).

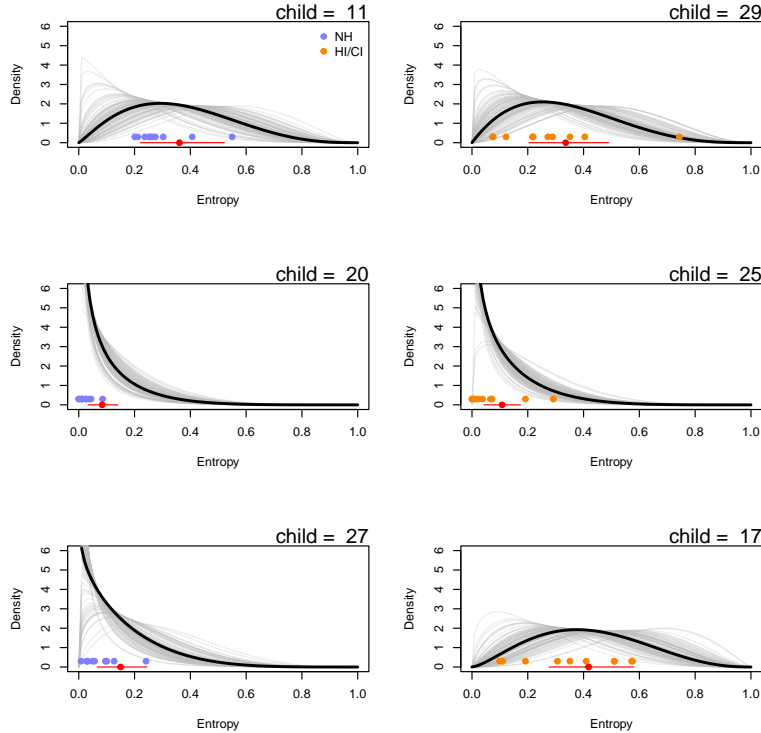Figure 4: Model 10, posterior predictive: entropy replicates, *true* entropy, and distributions. Colored points represent the entropy replicates per *hearing status* group. Red points with lines represent the mean *true* entropy with 95% highest posterior density interval (HPDI). Thick solid line represents the marginal distribution, thin solid lines depicts 100 random posterior samples for the distributions.

3.5. Influential observations

Finally, implementing the statistical models under the Bayesian framework provide us with one last advantage: we were able to identify influential observations that might sway our estimates, under our model framework [53]. We argue this method is preferable, as the identification of influential observations through preliminary or univariate procedures might lead to erroneous exclusion of information, ultimately damaging our research inferences. See supplementary section A.5.2 for more details.

Figure 5 provides an overview of the identified high-leverage observations (pair child, utterance). Model ten reveals some observations for children 8, 10, and 15 can be deemed prominent (the first two were NH children, while the last was a HI/CI analogue).

After a careful inspection, the only similitude these observations shared was that all of them registered the lowest replicate values observed in the sample (0.007 for child 8, and zero for the remaining two). Therefore, it makes sense our model found these points 'surprising', as they are mostly out of the expected range for the outcome. However, notice additionally that even with their presence, no further evidence was found in favor of *robust* models over other more parsimonious iterations of our research hypothesis (see supplementary section A.5.6). As a result, none of these observation were excluded from the analysis, but they were identified nonetheless.
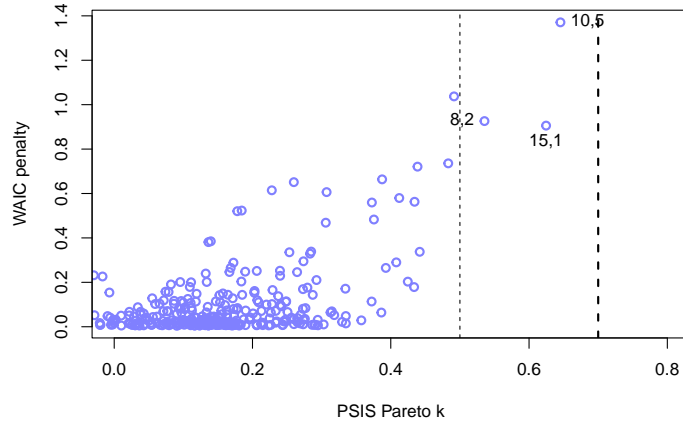

Figure 5: Model 10, influential observations. Pairs child, utterance are reported for specific observations. Thin and thick discontinuous lines indicate the minimum and extreme recommended threshold of identification (0.5 and 0.7, respectively).

4. Discussion

As stated in previous paragraphs, the main focus of the present research was to investigate the speech intelligibility levels of normal hearing (NH) versus hearing-impaired children with cochlear implants (HI/CI). For that purpose, we set up an experiment where ten utterances, from thirty two children, were transcribed and transformed into entropy measures. Finally, the entropy replicates served as our outcome variable.

Rather than use the common multilevel approach, the current research proposed a novel method of statistical analysis. First, we modeled the *speech intelligibility* as a latent variable that can be inferred from the entropy replicates [26, 44]. The previous method not only allow us to 'construct' an *intelligibility* score, but also to control for all the levels of variability present in our data. We argue the former was important to make comparisons at the appropriate levels, while the latter, to not 'manufacture' false confidence in our parameter estimates, leading us to incorrect inferences [53]. Second, we used Directed Acyclic Graphs (DAG) [59, 17] to describe our causal and non-causal hypothesis about the *intelligibility* of speech, and determine which variables were relevant for its development. Finally, we implemented the method under the Bayesian framework, where we provided the steps and code for its implementation.

The statistical results highlighted XXX points. First, and most important, considering the variability present in our data, we were not able to produce unequivocal null hypothesis rejections. The latter was a result of the large variability registered at the children and replicates levels, which impeded our ability

to delimit the *hearing status* group location on the *intelligibility* scale. However, based on simulation studies, we were able to say our models could affirm the estimated values, as long as the estimates were different from zero (power analysis). In that sense, although we continue interpreting the parameters estimates, we believe that to ultimately define the direction of the effects, a larger sample size might still be needed.

Considering the previous, we still believe our parameter estimates provided preliminary evidence about our causal hypothesis. On the one hand, we found there is a difference in *intelligibility* among the *hearing status* groups, in favor of the NH children. However, the evidence indicated the difference is only observed between NH and HI/CI children with etiologies other than the genetic one. Our intuition indicates that this might be due to the need of a 're-wiring' of the child's neural hearing patterns, something that might be inherently done in children with genetic etiology. As it is point out by Flexer [31]: "the more delayed the age of acquisition of a skill, the farther behind children are in the amount of cumulative practice they have had to perfect that skill. The same concept holds true for cumulative auditory practice". It is fair to say then, the researchers believe it would be interesting to further inspect this hypothesis with the support of Magnetic Resonance Imaging (MRI).

In contrast, *hearing age* was clearly identified as one of the main determinant in the evolution of *speech intelligibility*. Moreover, the evidence seem to indicate there is a different evolution between NH and HI/CI children, contrary to what was previously found [5]. We intuit the latter result have two possible complementary explanations, either: (i) the children develop their language differently, at different stages of their (*hearing*) age, as it is implied by Flexer [31] in the previous paragraph, or (ii) given that the hearing signal provided by the apparatus is degraded [22], HI/CI children might take longer to achieve similar levels of *intelligibility* than their NH counterparts. As with the previous case, the researchers believe it would be interesting to assess the latter hypothesis. For that purpose, a further collection of data would be required, where we identify the levels of degradation of the hearing signals in each of the cochlear implants.

4.1. Perspective of future research

5. Acknowledgments

6. Author contributions

All authors contributed to the development of the causal hypothesis. Jose Rivera performed the statistical analysis, Sven de Maeyer supervised the production of the documents and statistical results, and Steven Gillis collected the data.

7. Financial support

8. Conflicts of interest

The authors declare they have no conflict of interest.

9. Research transparency and reproducibility

The model simulation procedures and testing that support the findings of this study are openly available at `https://github.com/jriveraespejo/PhD_UA_paper1`.

Due to the privacy and confidentiality of subjects, the data set in which the model was implemented cannot be put online.

A. Supplementary

A.1. Children characteristics

Table 3 shows the detailed information of the sampled children. The referred table includes the variable used for the matching procedure, i.e. chronological age, while also additional variables thought to be relevant for our hypothesis. No other variables are included as no known additional comorbidities, beside their hearing impairment, are suspected. Additionally, notice the pure tone average (PTA) data was not collected for the NH children.

Child	Gender	Chronological age (y;m)	Device length of use (y;m)	Hearing age (y;m)	Etiology	PTA (dB.) unaided	aided
	HI/CI children						
1	female	05;07	05;00	05;00	Genetic	120	19
2	male	06;04	05;09	05;09	CMV	106	23
3	male	06;07	05;10	05;10	Genetic	114	35
4	female	06;10	06;00	06;00	Unknown	120	20
5	female	07;00	06;03	06;03	CMV	115	25
6	male	07;00	05;08	05;08	Genetic	93	32
7	female	07;00	06;08	06;08	Genetic	117	17
8	female	07;00	05;05	05;05	Unknown	112	42
9	male	07;00	05;05	05;05	CMV	120	15
10	female	07;01	05;11	05;11	Genetic	120	35
11	male	07;01	05;07	05;07	Genetic	113	42
12	male	07;02	06;05	06;05	Genetic	120	37
13	male	07;08	06;10	06;10	CMV	114	27
14	male	07;09	06;02	06;02	CMV	120	35
15	male	08;07	07;10	07;10	CMV	120	33
16	male	08;08	09;09	09;09	Genetic	95	27
	NH children						
17	female	06;05	n.a.	06;05	n.a.	n.a.	n.a.
18	female	06;06	n.a.	06;06	n.a.	n.a.	n.a.
19	female	06;07	n.a.	06;07	n.a.	n.a.	n.a.
20	female	06;09	n.a.	06;09	n.a.	n.a.	n.a.
21	female	06;09	n.a.	06;09	n.a.	n.a.	n.a.
22	male	06;09	n.a.	06;09	n.a.	n.a.	n.a.
23	male	06;09	n.a.	06;09	n.a.	n.a.	n.a.
24	male	06;10	n.a.	06;10	n.a.	n.a.	n.a.
25	female	07;01	n.a.	07;01	n.a.	n.a.	n.a.
26	male	07;01	n.a.	07;01	n.a.	n.a.	n.a.
27	male	07;04	n.a.	07;04	n.a.	n.a.	n.a.
28	female	07;08	n.a.	07;08	n.a.	n.a.	n.a.
29	male	07;08	n.a.	07;08	n.a.	n.a.	n.a.
30	female	07;09	n.a.	07;09	n.a.	n.a.	n.a.
31	female	08;00	n.a.	08;00	n.a.	n.a.	n.a.
32	female	08;01	n.a.	08;01	n.a.	n.a.	n.a.

(y;m) = (years;months)

n.a. = not applicable / not available

Table 3: Characteristics of selected children.

A.2. Experiment details

A.2.1. Transcription task

The setting for the transcription task comprised the following steps [4, 5]:

1. the listener took a seat in front of a computer screen, located at the campus' computer laboratory.

2. the listener opened Qualtrics [77] and select the transcription task.

3. the listener read two set of instructions presented on the computer screen about:

   a) *how to perform the task,*

   b) *the aspects not considered for the task.*

4. the listener hear the stimuli through high quality headphones, set at a comfortable volume.

5. the listener wrote the orthographic transcriptions of the utterances, in a free text field in the software environment.

A.2.2. Entropy calculation

The outcome from the transcription task was obtained following a two step procedure [5]. First, we aligned the participant's orthographic transcriptions, at the utterance level, in a column-like grid structure similar to the one presented in Table 4. This step was repeated for every one of the 6400 transcriptions. Lastly, we computed the entropy measure of the aligned transcriptions as in Shannon [65]:

$$H = H(\boldsymbol{p}) = \frac{-\sum_{i=1}^{n} p_i \cdot \log_2(p_i)}{\log_2(N)} \tag{5}$$

where H is bounded in the unit interval $[0, 1]$, n denotes the number of word occurrences within each utterance, p_i the probability of such word occurrence, and N the total number of aligned transcriptions per utterance.

Transcription	Utterance				
number	1	2	3	4	5
1	de	jongen	ziet	een	kikker
	the	boy	see	a	frog
2	de	jongen	ziet	de	[X]
	the	boy	sees	the	[X]
3	de	jongen	zag	[B]	kokkin
	the	boy	saw	[B]	cook
4	de	jongen	zag	geen	kikkers
	the	boy	saw	no	frogs
5	de	hond	zoekt	een	[X]
	the	dog	searches	a	[X]
Entropy	0	0.3109	0.6555	0.8277	1

[B] = blank space, [X] = unidentifiable word

Table 4: Alignment and entropy calculation. Extracted from Boonen et al. [5], and slightly modified with illustrative purposes.

Entropy was used as a quantification of (dis)agreement between listeners' transcriptions, i.e. utterances yielding a high degree of agreement between transcribers were considered highly intelligible, and therefore registered a lower entropy ($H \to 0$). In contrast, utterances yielding a low degree of agreement were considered as exhibiting low intelligibility, and therefore registered a higher entropy ($H \to 1$) [5, 27].

To exemplify relevant scenarios for the procedure, we generate the entropy for utterances 2, 4 and 5 present in Table 4. To make the example easy to calculate, we assume our data consisted only of five transcriptions in total ($N = 5$).

For the second utterance, we observe that four transcriptions identify it with the word *jongen*, while the last with the word *hond*. Therefore, we registered two word occurrences ($n = 2$), with probabilities $\boldsymbol{p} = (p_1, p_2) = (4/5, 1/5)$, and entropy measure equal to:

$$
\begin{aligned}
H &= \frac{-\sum_{i=1}^{2} p_i \cdot \log_2(p_i)}{\log_2(5)} \\
&= \frac{-\left[0.8 \log_2(0.8) + 0.2 \log_2(0.2)\right]}{\log_2(5)} \\
&\approx 0.3109
\end{aligned}
$$

For the fourth utterance, we observe that two transcriptions identify it with the word *een*, one with *de*, one with *geen*, and one with a blank space [B]. Notice the blank space was not expected in such position, therefore, it was considered as a different word occurrence. As a result, the scenario had four word occurrences ($n = 4$), with probabilities $\boldsymbol{p} = (p_1, p_2, p_3, p_4) = (2/5, 1/5, 1/5, 1/5)$, and entropy measure equal to:

$$
\begin{aligned}
H &= \frac{-\sum_{i=1}^{4} p_i \cdot \log_2(p_i)}{\log_2(5)} \\
&= \frac{-\left[0.4 \log_2(0.4) + 3 \cdot 0.2 \log_2(0.2)\right]}{\log_2(5)} \\
&\approx 0.8277
\end{aligned}
$$

Finally, for the fifth utterance, we observe that all transcriptions identify it with different words. Notice that when a transcriber did not managed to identify (part of) the complete utterance, (s)he was instructed to write [X] to replace it. However, if more than one transcriber used [X] for an unidentifiable word, each was considered as being different from one another. The latter is done to avoid the artificial reduction of the entropy measure, as [X] values already indicate the lack of intelligibility of the word. Therefore, for the fifth utterance we registered five word occurrences ($n = 5$), with probabilities $\boldsymbol{p} = (p_1, \ldots, p_5) = (1/5, \ldots, 1/5)$, and entropy measure equal to:

$$
\begin{aligned}
H &= \frac{-\sum_{i=1}^{5} p_i \cdot \log_2(p_i)}{\log_2(5)} \\
&= \frac{-5 \cdot 0.2 \log_2(0.2)}{\log_2(5)} \\
&= 1
\end{aligned}
$$

A.3. About speech intelligibility

As described in the introduction, intelligible speech can be defined as the extent to which the elements in an speaker's acoustic signal, e.g. phonemes or words, can be correctly recovered by a listener [47, 73, 70, 36]. More specifically, in the context of the transcription task, speech intelligibility can be inferred from the extent a set of transcribers can identify the word contained in an utterance [5].

Therefore in this paper, through the implementation of our proposed model, *speech intelligibility* is interpreted as a latent trait of individuals (hypothetical construct), which underlies the probability of observing a set of entropy replicates; that in turns, describes the ability of transcribers to identify the words in an utterance. Henceforth, statements such *'speech intelligibility is influenced by'* can be read as *'the probability of observing a set of entropy replicates for each individual in the sample is influenced by'*. Similar interpretation can be extended to the (latent) *true* entropy measures.

Despite this practical approach, we emphasize we did our best to ensure the construct validity of our study, by ensuring the transcription task was well understood and appropriately performed by the transcribers.

We then expect speech intelligibility, as measured by our model, to reflect the (general) intelligibility of speech possessed by individuals, but do not deal with general epistemological considerations on the connection between the two.

A.4. Sampling bias

As it happens in most observational, and some experimental studies, ours can also be a potential victim of sampling bias. While stratifying on the selection variables can help to balance the samples, and even 'correct' the estimates [17, 20], as we do here by controlling for *hearing age*. Given the sample's selection and matching procedures, we cannot ensure the HI/CI nor the NH groups are representative of their respective populations.

Nevertheless, we argue that by controlling for other relevant confounders, the qualitative results presented in this study holds. However, we cannot discard the presence of unobservable variables that could bias our results, and in that sense, inferences beyond this particular set of children must be taken with care.

A.5. Model details

A.5.1. Variability in the beta-proportion distribution

Figure 6 shows the implications of different 'sample sizes' (M_i) on the dispersion of the beta-proportion distribution [48]. The panels show different average entropies: the middle panel assumes $\mu = 0.5$, the left $\mu = 0.2$, and the right $\mu = 0.8$ (as shown by the discontinuous lines).

In all three panels we notice two prevalent pattern: (i) the higher the 'sample size', the less dispersed is the distribution, and (ii) as expected from non-linear models, the behavior of the dispersion depends on the location of the distribution, i.e. their average value.

Concerning the first patterns, we expect that if the posterior estimates for M_i reach lower values, it would imply the entropy replicates H^O_{bik} had high dispersion. In contrast, higher M_i estimates would imply a lower dispersion in the replicates. In this particular case, a good comparison point is the number of utterances/replicates. If $M_i < 10$, it would imply that our effective 'sample size' (measuring points) is less than the actual number of replicates, and therefore, we would need more replicates to provide an accurate measure of the (latent) *true* entropy. In contrast, $M_i > 10$ would imply the opposite.

As a final point, it is important to highlight that as the approach uses the 'sample size' parameter to model the replicates' heterogeneity, we are effectively estimating a measurement error model [10] on entropy values.
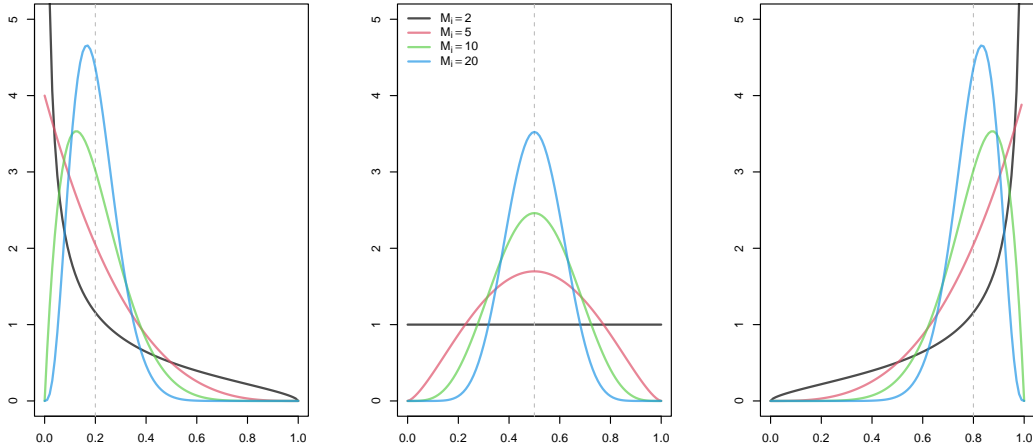

Figure 6: Variability in a beta-proportional distribution. Discontinuous lines describe the average value for the distribution (μ), solid lines describe the distribution assuming different $\theta = M_i$.

A.5.2. Data pre-processing

Besides the exclusion of corrupted observations, e.g. no available transcription, no other information was excluded before the modeling process. This decision departs from what has been done in previous research

[4, 69, 5]. The reason is that we believe the identification of influential observations, through preliminary or univariate procedures, might lead to erroneous exclusion of information, ultimately biasing our results. We believe the identification of *outliers* should not be done outside the context of a full modeling effort [53], as what can behave as an *outlier* based on a univariate analysis, might behave as expected under the appropriate model.

Considering the previous, instead of eliminating information based on preliminary analysis, we proposed a set of models that can be qualified as *robust* against influential observations. Considering the flexibility of the Bayesian framework, the proposal of such models was a trivial task (see Table 1).
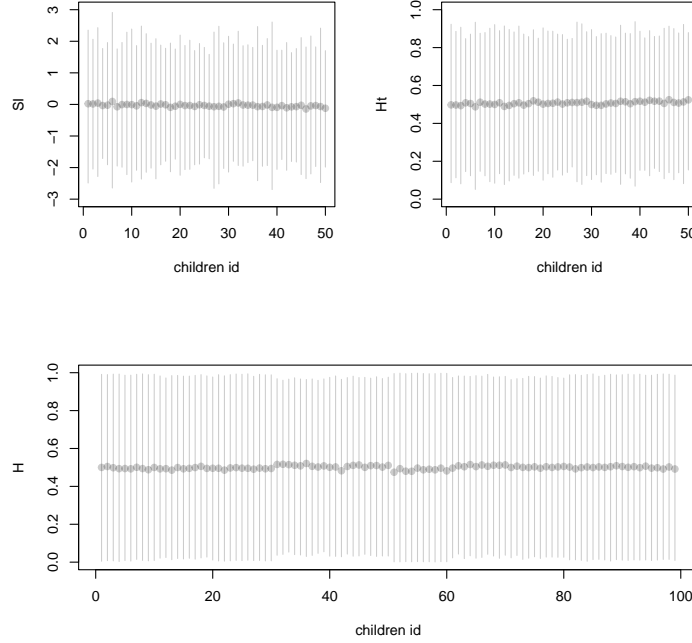

Figure 7: Prior distribution implications: speech intelligibility, *true* entropy and observed entropy scales. Circles represent mean values, lines depict the 95% compatibility intervals.

A.5.3. Priors and hyper-priors

For all models, the selection of priors and hyper-priors was done through prior predictive simulation [53]. The selected priors were considered mildly informative and regularizing.

Figure 7 shows the implication of our assumptions on the three outcome scales of interest: the *speech intelligibility*, the *true* entropy, and the *observed* entropy replicates. Notice, no undesirable assumption has crept in any of the scales. Consequently, the estimates are free to visit a wide range of results, while also establishing a low probability of reaching impossible outcomes.

Hereby follows a description of the priors:

$$M_i \sim \text{Log-Normal}(\mu_M, \sigma_M) \tag{6}$$
$$a_b \sim \text{Normal}(\mu_b, \sigma_b) \tag{7}$$
$$a_i \sim \text{Normal}(\mu_i, \sigma_i) \tag{8}$$
$$\alpha \sim \text{Normal}(0, 0.2) \tag{9}$$
$$\alpha_{E[i], HS[i]} \sim \text{Normal}(0, 0.3) \tag{10}$$
$$\beta_{A, HS[i]} \sim \text{Normal}(0, 0.3) \tag{11}$$
$$\beta_{P, HS[i]} \sim \text{Normal}(0, 0.3) \tag{12}$$

while the hyper-priors were defined as follows:

$$\mu_M \sim \text{Normal}(0, 0.5) \tag{13}$$
$$\sigma_M \sim \text{Exponential}(1) \tag{14}$$
$$\mu_b \sim \text{Normal}(0, 0.2) \tag{15}$$
$$\sigma_b \sim \text{Exponential}(1) \tag{16}$$
$$\mu_i \sim \text{Normal}(0, 0.2) \tag{17}$$
$$\sigma_i \sim \text{Exponential}(1) \tag{18}$$

A.5.4. Estimation procedure

The proposed models in Table 1 were estimated under the Bayesian framework. More specifically, we used the No-U-Turn Hamiltonian Monte Carlo algorithm (No-U-Turn HMC) [3, 23, 43, 55] implemented in `Stan` [67]. We used four chains of 2000 iterations each, from which the first 1000 were discarded as a warm-up required by the software, leaving 1000 effective iterations.

Additionally, all procedures were performed in `R` 4.1.2 [61] and its integration packages [66], in a 64-bit system with a MINGW 32-bit cross-compiler (`x86_64-w64-mingw32/x64`).
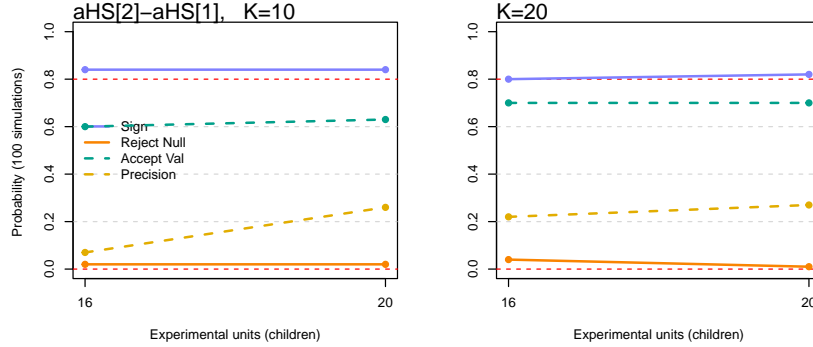

Figure 8: Group contrasts: power on small hearing group differences (≈ 0.15)

A.5.5. Simulation

Prior to fitting real data, all the proposed models were tested on simulated data. The simulation was done with two purposes: (i) to test the models' ability to recover the 'real' parameter values [35], and (ii) to test the power of our data size and assumed hypothesis [48]. On both instances, we used the *equivalent prior sampling* method [76], informed by the appropriate children population [19].

The results of our simulations highlight three key points. First, the probabilistic implementations work as intended, i.e. the statistical models manage to recover the 'real' estimate values, if enough sample size is available.

Second, considering our data size (sixteen children per group), if the assumed group difference is small (approximately 0.15), we can be certain that at least 60% of the time our models will affirm the predicted value, i.e. our model can correctly estimate values different from zero, when such alternative hypothesis is true (see green discontinuous line in figure 8). Moreover, the previous power threshold surpasses 80% if we assume medium group differences (approximately 0.4, not shown). In the case of other parameters of interest, even if we assume small effects, the models always affirm the predicted value with power above 80%, e.g. for $bA = 0.15$ or $bP = 0.1$.

Third, with our current data size, the statistical precision of our estimates do not achieve similar power (see yellow discontinuous line in figure 8). The latter implies that with sixteen children per group, the standard error of our estimates will not allow us to produce definite conclusions, e.g. we will not be certain if the two groups are not different because they are truly not statistically different or because the sample size is not enough to produce unequivocal inferences. A great part of this result is due to the assumed variability at the children and replicates levels, which we assumed moderate. However, further

model testing indicated that such unequivocal results can be accomplished, if we have sample sizes similar to the full population size (70 children per group).

For more details on the assumed effects and the simulation code, refer to the GitHub repository: `https://github.com/jriveraespejo/PhD_UA_paper1`

A.5.6. Model selection

As stated in previous sections, the current research used the Information-Theoretic Approach [1, 12] for model selection. The application of the approach required: (i) the expression of the research hypothesis into statistical models, (ii) the selection of the most plausible models, and (iii) to produce inferences based on one or multiple selected models. Since the first step is covered in sections 2.4 and 2.5, and expanded in supplementary section A.5, here we proceed with the intermediate step.

We used the widely applicable information criterion (WAIC) [72], and the Pareto-smoothed importance sampling cross-validation (PSIS) [71] as the criteria to select among competing models. Two reasons justify our decision. First, both criteria allow us to use all the information of our Bayesian models, i.e. the posterior distribution of the parameters. Last, and more important, both criteria provide us with the best approximations for the out-of-sample (cross-validated) deviance [53]. The deviance is the best approximation for the Kullback-Liebler (KL) divergence [49], i.e. a measure of how far a model is from describing the *true* distribution of our data. In that sense, by comparing the deviance between competing models, we can measure which model is the farthest from '*perfect predictive accuracy*' for our data [53].

Table 5 and 6 report the criteria for all the proposed models, sorted by the appropriate statistic. Three patterns are visible from the tables. First, we notice that around 63% of the evidence (WE) support two types of models: (a) no interaction and one estimated 'sample size', and (b) three (similar) interaction models with one estimated 'sample size'. Furthermore, 35% of the evidence is condensed in the diverse types of *robust* models, while the remaining 2% is negligible dispersed in the remaining ones.

Second, comparing the differences in WAIC/PSIS from the 'best' model (dWAIC and dPSIS, respectively), we arrive at the same conclusion as the previous paragraph, i.e. the model with no interaction and one estimated 'sample size' is the best model, followed by the interaction models with one 'sample size', and a bit further away, by the *robust* models. One important point to highlight is that, even when there seem to be a hierarchy of fit among the models, effectively, all of these are indistinguishable from each other, as it can be inferred from their precision estimate (dSE). However, we can also observe, a more precise support for models with any type of interaction and one estimated 'sample size', than for the *robust* models.

Third, we observe a greater over-fitting penalty (pWAIC and pPSIS, respectively) for models with a fixed 'sample size' than any other model. This is quite particular, as one would expect that a model with less parameters would over-fit less the data. However in this case, it implies the complexity of the data requires more parameters, especially considering there might be some highly influential observations that might sway the estimates of the less parameterized models (see section 3.5).

Model	Name	WAIC	SE	dWAIC	dSE	pWAIC	WE
3	No interaction (one 'size')	-621.0	42.99	0.0	–	31.1	0.18
10	Full interaction (one 'size')	-621.0	42.98	0.1	0.78	31.2	0.18
9	Etiology interaction (one 'size')	-620.7	43.07	0.4	0.48	31.2	0.15
8	Age interaction (one 'size')	-620.2	42.91	0.8	0.76	32.1	0.12
11	Age interaction (robust)	-620.1	42.70	0.9	2.21	34.2	0.12
12	Etiology interaction (robust)	-619.9	42.77	1.1	2.06	34.4	0.10
4	No interaction (robust)	-619.2	42.81	1.8	2.18	34.8	0.07
13	Full interaction (robust)	-618.8	42.65	2.2	2.17	35.0	0.06
5	Age interaction (fixed 'size')	-578.2	52.51	42.8	16.85	50.7	0.00
6	Etiology interaction (fixed 'size')	-578.2	52.63	42.8	16.95	50.7	0.00
2	No interaction (fixed 'size')	-577.6	52.59	43.5	16.97	51.0	0.00
1	Intercept only (fixed 'size')	-576.8	52.98	44.3	17.40	51.8	0.00
7	Full interaction (fixed 'size')	-575.8	52.58	45.3	16.96	52.1	0.00

SE	WAIC approximate standard error
dWAIC	difference in WAIC from the best model
dSE	standard error for the difference in WAIC from the best model
pWAIC	WAIC over fitting penalty
WE	weight of evidence

Table 5: Fit of statistical models: WAIC.

Model	Name	PSIS	SE	dPSIS	dSE	pPSIS	WE
3	No interaction (one 'size')	-619.8	42.98	0.0	–	31.8	0.22
10	Full interaction (one 'size')	-619.7	42.97	0.0	0.82	31.8	0.22
9	Etiology interaction (one 'size')	-619.2	43.05	0.5	0.51	32.0	0.17
8	Age interaction (one 'size')	-619.0	42.89	0.7	0.81	31.8	0.15
11	Age interaction (robust)	-617.8	42.70	2.0	2.19	35.4	0.08
12	Etiology interaction (robust)	-617.7	42.77	2.1	2.07	35.5	0.08
4	No interaction (robust)	-616.5	42.81	3.3	2.19	36.1	0.04
13	Full interaction (robust)	-616.3	42.64	3.4	2.13	36.2	0.04
6	Etiology interaction (fixed 'size')	-576.0	52.63	43.7	17.00	51.7	0.00
5	Age interaction (fixed 'size')	-575.9	52.54	43.9	16.94	51.9	0.00
2	No interaction (fixed 'size')	-575.3	52.59	44.4	17.03	52.1	0.00
1	Intercept only (fixed 'size')	-574.6	52.97	45.1	17.44	52.9	0.00
7	Full interaction (fixed 'size')	-573.8	52.61	46.0	17.01	53.1	0.00

SE	WAIC approximate standard error
dPSIS	difference in PSIS from the best model
dSE	standard error for the difference in PSIS from the best model
pPSIS	PSIS over fitting penalty
WE	weight of evidence

Table 6: Fit of statistical models: PSIS.

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