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1 The function and evolution of child-directed

2 communication

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23 Abstract

24 Humans communicate with small children in unusual and highly conspicuous ways (child-directed 25 communication (CDC)), which enhance social bonding and facilitate language acquisition. CDC-like inputs 26 are also reported for some vocally learning animals, suggesting similar functions in facilitating 27 communicative competence. However, adult great apes, our closest-living relatives, rarely signal to their 28 infants, implicating communication surrounding the infant as the main input for infant great apes and 29 early humans. Given cross-cultural variation in the amount and structure of CDC, we suggest that child-30 surrounding communication provides essential compensatory input when CDC is less prevalent— a 31 paramount topic for future studies.

32

33 Introduction

Human languages exhibit enormous variation at all linguistic levels, ranging from phonemes, the smallest meaning-distinguishing units, to morphemes, the smallest meaning-bearing units, to words, higher-level constructions, and rules of combination. Few, if any, of these features are under strong genetic control. As a consequence, linguistic units must be learned from scratch by every maturing individual: a process that, while often described as "effortless" [1], in fact takes many thousands of hours of exposure over multiple years. Inevitably, the communicative environment must provide the input required for learning a native language.

41

One prominent source of this input is a special speech register used by caregivers to address infants and
young children, frequently referred to as babytalk, motherese, parentese and, more recently, infantdirected or child-directed speech [2]. In this Essay, we use a more neutral term child-directed
communication (CDC, see Box 1) since there is lack of agreement of what constitutes infancy in humans,

46	and moreover, the input is modality independent (i.e., it is also encountered in sign languages [3,4]).
47	Such cross-modal prevalence has even been argued to support the notion that CDC is an automatic and
48	potentially species-wide trait [5]. Both in signed and spoken languages, CDC includes other multi-modal
49	features such as more exaggerated facial expressions [6], modified gestures [7] and motions in general,
50	with the latter known as motionese [8].
51	
52	Box 1. Definitions of key terms
53	Child-directed communication (CDC): All communication specifically directed at children, in which the
54	properties and structure of the signal often change in predictable ways, e.g. higher pitch, more
55	exaggerated gestures and more repetition. CDC supports language learning in children [2,9].
56	Child-surrounding communication (CSC): All communication that is perceptible to the child but not
57	directed at them.
58	Immature-directed communication: All communication specifically directed at the immature animal, as
59	indicated by the vocalizations or gestures being accompanied by body or head orientation towards the
60	immature animal, as well as a change in structural or acoustic features, for instance more repetition.
61	Natural pedagogy: The specific aspects of human communication that allow and facilitate the transfer of
62	generic knowledge to novices [10].
63	9-month revolution: A large set of cognitive and socio-cognitive skills that human infants typically
64	develop at around 9-12 months of age. Within this skill set they develop the ability to use gaze-
65	following, social referencing, pointing, joint attention and imitation to join the adult's attentional focus
66	[11]. They also become able to interpret adults' gestures as intentional acts [12].
67	Vocal learning: describes vocal production learning, which is traditionally defined as the production of
68	novel vocalizations as a result of learning from an acoustic signal [13]. Today, many dimensions and
69	degrees of vocal production learning are acknowledged [14]. Only few animal species are known to be

capable of vocal production learning (e.g. songbirds, hummingbirds, cetaceans and pinnipeds). In
contrast to vocal production learning stand usage and comprehension learning, which are more
common in a wide variety of species [15]. Usage learning is defined as learning to produce a signal in a
new context as a result of acoustic experience. Comprehension learning is defined as learning a new
meaning of a signal as a result of experience [13].

75

76 A second and much less researched source of input is child-surrounding communication (CSC, Box 1), 77 which includes all communication that is in perceptible proximity to, but not specifically directed 78 towards the child. Typically, this involves two or more individuals engaged in some type of social 79 interaction accompanied by a linguistic exchange. It may also include linguistic input from media 80 sources (e.g. TV, radio), but it remains unclear which impact this type of input might have on the child's 81 language development. CSC input is ubiquitous, and at least as omnipresent as CDC, yet we know much 82 less about its functional role in language acquisition. The few available studies on CSC suggest that it has 83 less impact than CDC on linguistic development in early ontogeny [16,17].

84

85 The reliance on child-directed communication for the acquisition of communicative competence may be 86 explained by three distinct evolutionary pathways (Fig 1). First, it might be shared with our closest living 87 relatives, the great apes. If this is the case we can assume that it is a feature that was also present in 88 early hominins (i.e. the "African Apes"; extant and extinct Homo, Pan and Gorillini genera). Second, it 89 may be derived in humans, and perhaps be one of the drivers of the evolution of language, potentially 90 as part of a wider change in cognitive architecture of early humans. This derived state can have arisen 91 uniquely in our ancestors or, third, it can be fully or partially shared with other, distantly related taxa, in 92 which case it arose via convergent evolution.

93

94	Fig 1. Evolutionary pathways of child-directed communication. A feature such as child-directed
95	communication with the function of aiding the acquisition of communicative competence can be (1)
96	ancestral: homologously derived among African great apes and thus also found in humans, (2) unique
97	among the great apes but convergently shared analogously with other, more distantly related species,
98	or (3) newly evolved within our own species. Red represents the presence of immature-directed
99	communication features. Outline credits: Human - T. Michael Keesey; Chimpanzee: Jonathan Lawley;
100	Bonobo: T. Michael Keesey; Gorilla: T. Michael Keesey (after Colin M.L. Burnett); Orangutan: Gareth
101	Monger; Gibbon: Kai R. Caspar; Tamarin: Yan Wong and T.F. Zimmerman; Zebra Finch: Jim Bendon
102	(photography) and T. Michael Keesey (vectorization); Bat: Yan Wong; Squamate: Ghedo and T. Michael
103	Keesey; Feline: Margot Michaud; Equine: T. Michael Keesey; Cetacean: Scott Hartman; Falcon: Liftarn;
104	Fish, macaque and baboon are uncredited. Link to creative commons license:
105	https://creativecommons.org/licenses/by-sa/3.0/. Link to public domain license:
106	https://creativecommons.org/publicdomain/zero/1.0/. Outlines were downloaded from
107	http://www.http://phylopic.org/. The layout of the figure was achieved in R (version 4.1.2, R
108	Development Core Team, 2012).
109	
110	
111	Current evidence suggests that in non-human primates in general (hereafter primates) the ability to
112	produce species-specific vocalizations develops with relatively little environmental contribution, i.e.,
113	irrespective of auditory input [18-21]. Instead, input seems to have more of a role in guiding vocal usage
114	and comprehension [22-25]. Nonetheless, at least some vocal production, flexibility does exist in
115	primates, although mainly in terms of socially driven vocal accommodation [22,26-32]. Although this
116	suggests a role for social input, how much of this is immature-directed communication (IDC) versus
117	immature-surrounding communication remains unclear [33]. So far, the few studies that have assessed

immature-directed vocalizations in great apes have yielded low rates (chimpanzees, *Pan troglodytes*[33]; bonobos, *Pan paniscus* [34]). A few studies have described vocalizations used by mothers in
chimpanzees [35] and orangutans [36]. However, this directed communication does not display any of
the features or functions of natural pedagogy. Overall, the current state-of-the-art suggests that
immature-directed input has only a small impact on great ape vocal ontogeny, if any. The preliminary
conclusion thus appears to be that most acoustic features of CDC are derived in humans. However, in
the structural domain some precursors of CDC might exist in apes.

125

126 However, a striking exception is found in the gestural domain. Orangutans [37], chimpanzees [38] and 127 bonobos [39] all use immature-directed gestures. Furthermore, one CDC-like feature, repetition, is 128 found in gorilla [40] and chimpanzee gestures [41]. The use of specific gestures and their repetition rates 129 by adult great apes towards immature individuals varies depending on the age and experience of the 130 immature animal, as in humans, suggesting functional significance in the acquisition of communicative 131 competence [40,41]. However, repetitions of gestures following lack of comprehension have also been 132 described in adult orangutans [42]. In addition, bonobos modify communication signals according to 133 recipient familiarity [43]. All of this suggests at least some shared cognitive features with humans. 134 Evidently, more research is needed to assess whether immature-directed gestures can be considered 135 the functional equivalent of CDC, especially in light of suggestions that at least part of the gestural 136 repertoire are the result of innovations and therefore have to be learned [44]. 137

138 If CDC is fully or at least partially derived in humans, this raises two important questions. First, which 139 elements of the broad bundle of features that make up human CDC were already present in the last 140 common ancestor? Identifying which elements were pre-existing (homologies: present in great apes), 141 which are found in other animals (analogies: convergently evolved), and which are new and uniquely

142	derived in our lineage would improve our understanding of how language acquisition evolved (Fig 1).
143	Second, as IDC in primates in general appears to be rare, primates must acquire the learnt part of their
144	communication from the communication that surrounds them, but is largely not directed at them. Has
145	this originally predominant source of input remained significant in humans, or has CDC replaced it (Fig
146	2)?
147	
148	Fig 2. Transition of child-surrounding to child-directed communication. The transition of the
149	importance of use of child-surrounding communication (CSC) to child-directed communication (CDC).
150	Darker color shows importance/presence and brighter color possible insignificance of CSC and CDC from
151	early hominins to extant humans.
152	
153	
154	In this Essay, we aim to address these two questions. In the first section, we deconstruct CDC into its
155	component parts and assess their proposed functions; we then ask for each of them whether
156	comparable phenomena exist in non-human animals (hereafter animals). In the second section, we
157	contrast CDC in humans with the lesser-studied CSC to shed light on the interplay between these two
158	forms of input and their respective roles in language acquisition. Answers to these questions should not
159	only improve our understanding of the development and acquisition of language but also its
160	evolutionary progression.
161	

162 The features and functions of CDC

163 CDC differs from adult-directed communication in a wide range of acoustic and structural features. This
164 has been observed in numerous cultures and is widely considered a universal of human language [9, 45,

165 46]. Over the past few decades, a plethora of studies have shown that features of CDC (Table 1) support 166 language acquisition by infants both in comprehension [47, 48] and production [49-51]. CDC is part of a 167 more general package of child-directed behaviors that serve to pass on cultural knowledge and skills to 168 the next generation, known as natural pedagogy [10] (Box 1). This active transmission process rests on 169 an (arguably) uniquely human capacity, ostension, which underlies pointing and results in gaze following 170 (often followed by joint attention on objects between caretaker and child [52] or a state of shared 171 intentionality more broadly [11]), as well as child-directed speech [10]. In this Essay, we argue that CDC 172 is a crucial part of this universal form of teaching. Such natural pedagogy is almost certainly derived 173 relative to the non-human great apes (hereafter great apes) and potentially evolved in relation to the 174 frequently highlighted shift in the breeding system from independent to more cooperative [53]. 175 Although the child-development literature may seem to suggest that natural pedagogy is primarily 176 aimed at preverbal infants and mainly geared toward teaching cultural knowledge, CDC is an obvious 177 and essential part of natural pedagogy extending well beyond early infancy. In fact, one might 178 hypothesize that CDC is a core feature enabling the transmission of language and as a consequence the 179 evolution of such a complex communication system.

181	Table 1. Known features of child-directed communication (C	CDC)	
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Type of feature	Known feature of CDC	Proposed function	Reference
Acoustic	Pitch variability	Attention grabbing	[54]
Acoustic	Lengthening of vowels and	Segmentation and	[55,56]
	pauses	discrimination of sounds	
Acoustic	Extended vowel triangle	Sound discrimination	[57]
Acoustic	Clear articulation	Facilitate comprehension	[46,58]
Acoustic	Increased voice-onset time	Sound discrimination	[59]

Acoustic	Slower speaking rate	Facilitate comprehension,	[54,60]
		segmentation	
Structural	Frequent repetitions	Structural generalization of word/unit classes	[61,62]
Structural	Short utterances	Facilitate comprehension	[63,64]
Structural	Low type/token ratio	Facilitate comprehension	[65,66]
Structural	Simplified syntax and	Facilitate comprehension	[63,65]
	semantics		
Structural	Frequent use of diminutives	Simplification of certain	[67,68]
		morphological aspects	
		(language specific)	
Structural	Frequent questions	Invite response, repetition,	[69,70]
		attention grabbing	
Structural	Variation sets	Structural generalization of	[71,72]
		word classes	
Structural	Scaffolding	Learning of word constructions	[73]

182 The first eight entries above the bold dividing line represent elements where a corresponding form

183 could possibly be present in animal vocal communication.

184

- 185
- 186 Adults and older children use the bundle of acoustic and structural features of CDC in varying

187 combinations when talking to infants and younger children (Table 1). For many of these features, there

188 is evidence that they facilitate the child's language learning.

- 190 Regarding the prosodic and acoustic features of the speech, CDC involves the production of higher and
- 191 more variable pitch [54], systematic lengthening of vowels and pauses [55,56,74] and an extended

192 "vowel triangle" or vowel hyperarticulation [57,75]. Studies have shown that these prosodic

modifications attract the child's attention [76] from an early age and that CDC is more salient to children
than adult-directed communication and is actually preferred by them [60,77-79]. Indeed,

195 neurobiological research has revealed that an infant's exposure to CDC in their first year of life results in

a higher brain activation in their left and right temporal areas compared with adult-directed speech [80].

197 These prosodic modifications also elicit increased infant vocal responses during their prelinguistic phase

198 [81], a form of active participation crucial to language acquisition [2]. Infants listening to CDC rather

than adult-directed speech also show greater sensitivity to syllable and vowel discrimination [75,82].

Lastly, caregivers tend to use exaggerated prosody to mark new or relevant vocabulary [74,83,84]. These

201 prosodic characteristics of CDC not only support the detection of word boundaries [85], but also word

comprehension [48,86] and production [49]. In sum, acoustic alternations of the speech signal appear to
 accelerate various aspects of language acquisition (see [87] for a review), suggesting that CDC serves as

an evolved teaching tool.

205

206 Regarding the structural features, CDC is characterized by short utterances [63,64], a low type/token 207 ratio [65,66], which indicates that caregivers use a simplified vocabulary, and the use of many questions 208 [69,70], diminutives [67,68] and repetitions [61,62]. One structural feature in particular is known to have 209 a significant role in the acquisition of language: frequency effects. The more frequently an element 210 occurs in the child's input, the faster it is expected to be learned [88,89]. Recent research has also 211 shown that frequent repetitions are structured in CDC. Repetitions of constructions at the beginning of 212 utterances (e.g., this is an X [62,90]) and discontinuous repetitions (e.g., I X you [91,92]) are ubiquitous 213 and support the generalization of word classes, such as nouns and verbs [93]. In addition, repetitive 214 structures or distribution of words surrounding specific verbs support the generalization of meaning 215 [94], and the high number of repetitions found in CDC are positively correlated with word

comprehension [95,96]. A specific form of repetitions frequently used in CDC is variation sets, successive
utterances with partial self-repetitions produced by caregivers [71,72], which themselves are positively
related to better linguistic outcomes in naturalistic longitudinal [97] and experimental settings [98].
These findings again support the hypothesis that CDC functions to accelerate language acquisition.

221 In addition to the prosodic and structural features of CDC, another important factor is the absolute 222 amount of linguistic input children receive. A number of studies have indicated that the amount of CDC 223 children experience is correlated with their later vocabulary development [16,99-102] and their word 224 processing skills [101]. The quality (variety of words and syntactic structures) of CDC also impacts 225 language development. Longitudinal studies have shown how input quality at an earlier stage of 226 development predicts subsequent diversity and variance in language outcome at a later stage of 227 development [103,104]. Quality and quantity may even have different roles during the child's language 228 development. For example, a longitudinal study of vocabulary acquisition revealed that input quantity 229 mattered most during the second year of development, whereas input quality was more important 230 during the third year [50]. The child's ability to profit from different properties of CDC might therefore 231 vary across development.

232

Most of the previously reviewed evidence is from children growing up in modern Western societies, characterized by child-rearing practices that are very different from what is typically seen in huntergatherer groups, our evolved and species-typical way of life [17]. In addition, there is substantial variation both within and across cultures in the amount of CDC that occurs and its features. Also important is that, in terms of sheer amount, there are linguistic communities in which children are only rarely directly addressed by their caregivers [105,106], suggesting that CDC is not essential for language acquisition, at least not as the main source of linguistic experience. A comparative study by Shneidman

240 et al. [16] demonstrated that for 1-year-old children growing up in a Yucatec Mayan community the 241 mean number of utterances a child encountered per hour amounted to approximately 400 utterances, 242 with only 20% of it being directed to the child. The US group of 1-year olds that served as a comparison 243 were exposed to approximately 900 utterances per hour, with more than 70% of these utterances being 244 directed. More recent studies from non-WEIRD (Western, Educated, Industrialized, Rich and Democratic 245 [107]) cultures confirmed that the amount of directed communication children are exposed to can vary 246 strongly (e.g., Netherlands: 303 vs. Mozambique: 58 utterances of CDC/30 min [108]; Tseltal: 3.63 min of 247 CDC/hour [109]; Tsimane: >1 min/daylight hour [17]; North American: 11.36 min of CDC/hour [110]), 248 raising questions about the relevance of CDC as the critical source of language acquisition. So far the 249 factors determining the amount of CDC are unclear. In particular the role of the child in the society 250 might be crucial, i.e., whether a society adapts situations to the child or expects to the child to adapt to 251 the situation [106,111].

252

Nonetheless, various studies revealed the presence of CDC features in non-WEIRD cultures (e.g. higher pitch [112]; slower speaking rate [113]; repetitions, diminutives and simpler syntax [114]). Overall the results suggest that both similarities (e.g. in pitch [113]) and differences [115] between WEIRD and non-WEIRD cultures do exist. However, not all CDC features can be found in every culture. In Quiché Mayan, for example, mothers do not seem to produce higher pitch when talking to their children, potentially because they must use this register when speaking to a person of higher status [116].

259

At this stage, it seems that the only universal characteristic of CDC is the presence of repetitive
 structural patterns in the input. Clearly, generalizations would be premature until more research reveals
 patterns linked to the social organization of a linguistic community. However, if one considers CDC as a

toolkit, the main features of CDC (Table 1) presumably change gradually as the infant progresses to

264	being a toddler and preschooler [117-120]. During the earliest stage before the 9-month revolution [12]
265	(see Box 1), acoustic and structural features appear to be very prominent, whereas structural features
266	seem to gain greater prominence at later stages (Table 1). Thus, initially the function of CDC may be to
267	establish and strengthen the social bond with infants, direct attention [121], introduce turn-taking via
268	proto-conversations [122], and scaffold the learning of the prosody, phonemes, morphemes and first
269	words of the local language. After the 9-month revolution, once joint attention, intention reading,
270	symbol recognition and rational imitation [11] have emerged, CDC may instead be geared more toward
271	the learning of vocabulary and grammar.
272	
273	A key next step in research would be to determine, for each culture, which features occur at what stage
274	in development and in which combination, and how these tools interact. CDC might turn out to be
275	heterogeneous across cultures. This variation might then be linked to the age at which children achieve
276	adult-level competence in the various components of language.
277	

278 The features and functions of immature-directed vocalizations

279 in animals

To identify both the evolutionary roots and adaptive functions of CDC in humans we must examine similar phenomena in animals. We already noted that preliminary work on great apes suggests our common ancestor featured few, if any, of the elements of CDC as listed in Table 1, at least in the vocal domain. However, it must be stressed that this absence may simply reflect a lack of focused research effort rather than actual absence. But if it is confirmed, this would suggest that surrounding vocalizations provide the primary input for the learned part of the vocal development in great apes and that CDC originated *de novo* in the human lineage (Fig 1), presumably linked to the emergence of natural
pedagogy, which may have preceded, and in fact facilitated, language evolution [53].

288

289 We now turn to possible convergent cases. First, we already discussed calls by great ape mothers, but 290 they also occur in other primates [123,124], as well as in many non-primate species, where mothers call 291 to their infants to retrieve them. Examples include domestic cats (*Felis silvestris catus* [125]), and 292 ungulates such as domestic sheep (Ovis aries [126]), cattle (Bos taurus [127]), goitred gazelles (Gazella 293 subgutturosa [128]) or saiga antelopes (Saiga tatarica tatarica [129]). Second, immature-directed calls 294 may serve to aid recognition of the mother's voice, as in domestic cats [125], Mexican free-tailed bats 295 (Tadarida brasiliensis mexicana [130]), fur seals (Arctocephalus tropicalis [131]), or domestic sheep 296 [126]. These examples show that even if IDC exists in an animal species, it is unlikely that these cases are 297 functionally equivalent to human CDC. 298 299 However, in a third category of species, we find immature-directed calls related to their capacity for 300 vocal accommodation (small alterations of vocalizations as a result of experience [132]) and vocal 301 learning (Box 1). Orcas (Orcinus orca) produce family-typical calls at higher rates after the birth of a calf 302 [133]. Likewise, common marmosets (*Callithrix jacchus*), which show evidence of accommodation 303 learning, and thus some level of vocal plasticity [134], modify call rates and repeat various different call 304 types before and after birth of infants [135]. In agile gibbons (Hylobates agilis), duetting by mothers with 305 inexperienced young has also been argued to represent IDC, serving to aid the acquisition of the species-

306 specific song [136]. In these cases, the calls may serve to acquire the group's vocal signature.

307

308 Finally, some cases show suggestive parallels to human CDC. In cooperatively breeding marmosets,

309 adults give contingent vocal feedback specifically to infants, which is suggested to impact vocal

310 ontogeny since infants exposed to more of such calls by adults produce and properly use adult-like calls 311 earlier [28,137], possibly owing to increased practice or because vocal feedback reduces stress [13]. This 312 contingent vocal feedback may help infants acquire the underlying rules of dyadic vocal communication 313 (i.e., turn-taking [138], but see [139]). Outside primates, in zebra finches, male tutors use a more 314 stereotypic song when they are near immature birds [140]. In greater sac-winged bats (Saccopteryx 315 *bilineata*), mothers adjust the pitch and timbre when they use immature-directed vocalizations [141]. 316 317 Despite these parallels, no study has asked exactly which features of the vocalizations (Table 1) are 318 essential and which functions they serve. It is therefore too early to conclude the common incidence of 319 CDC-like functions of immature-directed vocalizations in either primate or non-primate species

[28,40,140-142]. Systematic comparisons are needed to assess the extent of convergence and the
 determinants, but it remains plausible that IDC serves to facilitate the learning of vocal signatures (in
 accommodators) or call repertoires (in vocal learners sensu stricto), similar to the language-acquisition
 function of human CDC.

324

325 The function of CDC relative to CSC in humans

Although considerable attention has been paid to CDC and its structuring and function, comparatively less is known about the relative role of surrounding communication that children are exposed to (CSC). Indeed, in some linguistic communities surrounding communication is the primary source of input since adults rarely directly address infants (e.g. Kaluli and Samoan [106]; Yucatec Mayan [16], Tsimane [17]), at least in their first year of life. Despite these differences in input type, children still become competent native speakers [106,109,143,144]. This inevitably begs the question how important CDC actually is for speech development and suggests that CSC, though currently still under-researched, may have an equally important, perhaps compensatory role in facilitating language acquisition. In small-scale
societies, which arguably represent the more typical human condition, children are continuously
surrounded by individuals of all ages [145], suggesting that the amount and variation of CSC will be
higher than in WEIRD societies. To date, the few studies that to our knowledge have quantitatively
assessed this [17,109,146] have not revealed an effect of CSC on vocabulary development [16,101].
However, more work is needed to understand whether CSC supports the learning of other properties of
language such as grammatical features.

340

341 To obtain a full understanding of how communicative competence develops in both humans and 342 animals, it is critical to account for both sources of input — CDC and CSC — and the interplay between 343 them. Are both CSC and CDC essential for proper language learning, or are they to some extent 344 compensatory? If so, do the large amounts of CDC in WEIRD societies serve to compensate for the much 345 lower quantity of CSC? In animals, immature-surrounding vocalizations might well be the predominant 346 form of input, yet very little research has attempted to quantify their occurrence and assess their 347 influence on the development of communicative competence. Filling this gap should be a high priority 348 for research.

349

The question arises whether the relative amounts of CDC and CSC seen in humans are comparable to those found in great apes. The one study on chimpanzee infants suggests that immature-surrounding communicative events total approximately 15 gestures, 50 vocalizations and 3 gesture-call combinations per hour [147]. This is considerably more than what is known so far about the above mentioned low rate of immature-directed vocalizations. In all likelihood, therefore, immature-surrounding vocalizations were the most important source for the learnt part of the vocal system (usage and comprehension learning) in early hominins.

357

358 **Conclusion and future directions**

In human language learning, the amount and quality of CDC is one of the key facilitators of learning. But how the various features that make up CDC change with age, especially relative to the 9-month revolution, is not clear, and should be the target of future studies because they may vary in function from creating attachment, to establishing joint attention, to supporting specific details of language acquisition.

364

365 Despite its universality, research across and within cultures has shown enormous variation in a child's 366 exposure to directed communication. Studies of a few non-WEIRD societies show much lower rates of 367 CDC than found in the typical studies of WEIRD societies. This suggests that the amount of CDC children 368 are exposed to in WEIRD societies might be atypical for the rest of the world and most of human history. 369 Given the fact that all children learn the language of their culture, independent of culture-specific 370 variation in input, the role of CSC for language learning might have been underestimated. The increased 371 amount of CDC in WEIRD societies seems to result mainly in a refinement of skills, involving the size of 372 the vocabulary and the construction inventory involved. This raises the question how CDC produces this 373 refinement. Its impact may relate to the interactional situations in which it occurs. In these contexts 374 joint attention is the key component that actually facilitates learning [52,148,149]. Such joint-attentional 375 frames allow the reduction of interpretation space of form-meaning associations. Given the extreme 376 cross-linguistic variability of CDC we must ask the questions of whether and how much CDC is really 377 essential to language learning, whether CSC would do an equivalent job but just more slowly, or 378 whether CDC is essential at particular stages only. Daylong recordings in naturalistic conditions are likely 379 to provide answers to these questions.

381	To shed light on how CDC evolved, we examined research on our closest relatives, the great apes. So far
382	very little directed input to infants has been documented. Concerning the features of human CDC (Table
383	1), few have been found in ape communication, except for repetition of gestures. Repetition is arguably
384	the best predictor of language acquisition in human infants and children [88,89,150]. These findings
385	suggest that short-term repetitive use of communicative acts is potentially an ancestral feature of CDC.
386	We therefore propose more research is needed on structural repetition to complement the usual
387	emphasis on acoustic features of CDC.
388	
389	With regard to other animal species there is more evidence for immature-directed vocalizations in
390	species that engage in vocal learning. This supports the idea that CDC in hominins arose to support the
391	acquisition of highly culturally variable acoustic and structural features of language. However, much
392	more systematic comparisons are needed, which should indicate which of the features characterizing
393	human CDC are also found in these convergent cases. Obviously, more targeted work on great apes is a
394	high priority, if only to see whether repetition is the only CDC-like feature present and why gestures
395	appear to be the exception.
396	
397	In sum, the current state of research suggests that most features of human CDC have evolved anew in
398	our hominin ancestors. It serves to engage children in social interaction with caretakers and thus to
399	facilitate language acquisition, and in later phases more explicitly in the acquisition of semantics and
400	grammar. In other words, there is no doubt that CDC is an implicit teaching device. Doubt remains,
401	however, whether it is the only facilitator.
402	

404 **References**

- 406 1. Kuhl PK. Early language acquisition: cracking the speech code. Nat Rev Neurosci. 2004;5(11):831-43.
- 407 doi: 10.1038/nrn1533. PMID: 15496861.
- 408 2. Golinkoff RM, Can DD, Soderstrom M, Hirsh-Pasek K. (Baby) Talk to me: The social context of infant-
- directed speech and its effects on early language acquisition. Curr Dir Psychol Sci. 2015;24(5):339–
- 410 344. doi: 10.1177/0963721415595345.
- 411 3. Holzrichter AS, Meier RP. Child-directed signing in American sign language. In: Chamberlain C, C JPM,
- 412 Mayberry RI, editors. Language acquisition by eye. Hillsdale, NJ: Lawrence Erlbaum Associates; 2000.
 413 p. 25–40.
- 414 4. Masataka N. Motherese in a signed language. Infant Behav Dev. 1992;15(4):453–460. doi:
- 415 10.1016/0163-6383(92)80013-К.
- 416 5. De Boer B. Infant-directed speech and the evolution of language. In: Tallerman M, editor.
- 417 Evolutionary prerequisites for language. Oxford: Oxford University Press; 2005. p. 100–121.
- 418 6. Chong S, Werker JF, Russell JA, Carroll JM. Three facial expressions mothers direct to their infants.
- 419 Infant Child Dev. 2003;12(3):211–232. doi: 10.1002/icd.286.
- 420 7. Iverson JM, Capirci O, Longobardi E, Caselli MC. Gesturing in mother-child interactions. Cogn Dev.
- 421 1999;14(1):57–75. doi: 10.1016/S0885-2014(99)80018-5.
- 422 8. Brand RJ, Baldwin DA, Ashburn LA. Evidence for 'motionese': modifications in mothers' infant-
- 423 directed action. Dev Sci. 2002;5(1):72–83. doi: 10.1111/1467-7687.00211.
- 424 9. Soderstrom M. Beyond babytalk: Re-evaluating the nature and content of speech input to preverbal
- 425 infants. Dev Rev. 2007;27(4):501–532. doi: 10.1016/j.dr.2007.06.002.

- 426 10. Csibra G, Gergely G. Natural pedagogy. Trends Cogn Sci. 2009;13(4):148–153. doi:
- 427 10.1016/j.tics.2009.01.005.
- 428 11. Tomasello M. Constructing a language: A usage-based theory of language acquisition. Cambridge,
- 429 MA: Harvard University Press; 2003.
- 430 12. Tomasello M, Striano T, Rochat P. Do young children use objects as symbols? Br J Dev Psychol.

431 1999;17(4):563–584. doi.org/10.1348/026151099165483.

- 432 13. Janik VM, Slater PJ. The different roles of social learning in vocal communication. Anim Behav.
- 433 2000;60(1):1-11. doi: 10.1006/anbe.2000.1410. PMID: 10924198.
- 434 14. Vernes SC, Kriengwatana BP, Beeck VC, Fischer J, Tyack PL, Ten Cate C, Janik VM. The multi-
- 435 dimensional nature of vocal learning. Philos Trans R Soc Lond B Biol Sci. 25;376(1836):20200236.

436 doi: 10.1098/rstb.2020.0236. PMID: 34482723.

- 437 15. Janik VM, Slater PJ. Vocal learning in mammals. Adv Study Behav. 1997;26(1):59–100.
- 438 16. Shneidman LA, LA, Goldin-Meadow S. Language input and acquisition in a Mayan village: How
- 439 important is directed speech? Dev Sci. 2012;15(5):659–673. doi: 10.1111/j.1467-7687.2012.01168.x.
- 440 PMID: 22925514.
- 441 17. Cristia A, A, Gurven M, Stieglitz J. Child-directed speech is infrequent in a forager-farmer population:
- 442 A time allocation study. Child Dev. 2019;90(3):759–773. doi: 10.1111/cdev.12974. PMID: 29094348.
- 443 18. Snowdon CT. Language capacities of nonhuman animals. Am J Phys Anthropol. 1990;33(S11):215–
- 444 243. doi: 10.1002/ajpa.1330330510.

445	19. Talmage-Riggs G, Winter P, Ploog D, Mayer W. Effect of deafening on the vocal behavior of the
446	squirrel monkey (Saimiri sciureus). Folia Primatol. 1972;17(5-6):404–420. doi: 10.1159/000155458.
447	PMID: 4628332.

- 448 20. Winter P, Handley P, Ploog D, Schott D. Ontogeny of squirrel monkey calls under normal conditions
- 449 and under acoustic isolation. Vocal development in squirrel monkeys. Behaviour. 1973;47(3-4):230–
- 450 239. doi: 10.1163/156853973x00085. PMID: 4203637.
- 451 21. Hammerschmidt K, Freudenstein T, Jürgens U. Vocal development in squirrel monkeys. Behaviour.
 452 2001;138(9):1179–1204.
- 453 22. Seyfarth RM, Cheney DL. Vocal development in vervet monkeys. Anim Behav. 1986;34(6):1640–

454 1658.

455 23. Zuberbühler K. Interspecies semantic communication in two forest primates. Proc Biol Sci.

456 2000;267(1444):713-8. doi: 10.1098/rspb.2000.1061. PMID: 10821618.

457 24. Seyfarth RM, Cheney DL. Production, usage, and comprehension in animal vocalizations. Brain Lang.

458 2010;115(1):92–100. doi: 10.1016/j.bandl.2009.10.003. PMID: 19944456.

- 459 25. Wegdell F, Hammerschmidt K, Fischer J. Conserved alarm calls but rapid auditory learning in monkey
- 460 responses to novel flying objects. Nat Ecol Evol. 2019;3(7):1039–1042. doi: 10.1038/s41559-019-
- 461 0903-5. PMID: 31133723.
- 462 26. Zimmermann E. The vocal repertoire of the adult Senegal bushbaby (*Galago senegalensis*
- 463 *senegalensis*). Behaviour. 1985;94(3/4):212–233.
- 464 27. Fischer J, Hammerschmidt K. Towards a new taxonomy of primate vocal production learning. Philos
- 465 Trans R Soc Lond B Biol Sci. 2020;375(1789):20190045. doi: 10.1098/rstb.2019.0045. PMID:

466 31735147.

- 467 28. Takahashi DY, Fenley AR, Teramoto Y, Narayanan DZ, Borjon JI, Holmes P, et al. The developmental
- 468 dynamics of marmoset monkey vocal production. Science. 2015;349(6249):734–738. doi:
- 469 10.1126/science.aab1058. PMID: 26273055.
- 470 29. Watson SK, Townsend SW, Schel AM, Wilke C, Wallace EK, Cheng L, et al. Vocal learning in the
- 471 functionally referential food grunts of chimpanzees. Curr Biol. 2015;25(4):495-9. doi:
- 472 10.1016/j.cub.2014.12.032. PMID: 25660548.
- 473 30. Fischer J, Wheeler BC, Higham JP. Is there any evidence for vocal learning in chimpanzee food calls?
 474 Curr Biol. 2015;25(21):R1028-R1029. doi: 10.1016/j.cub.2015.09.010. PMID: 26528740.
- 475 31. Katsu N, Yamada K, Nakamichi M. Development in the usage and comprehension of greeting calls in
- 476 a free-ranging group of Japanese macaques (*Macaca fuscata*). Ethology. 2014;120(10):1024–1034.
- 477 doi: 10.1111/eth.12275.
- 478 32. Lameira AR, Hardus ME, Mielke A, Wich SA, Shumaker RW. Vocal fold control beyond the species-
- 479 specific repertoire in an orang-utan. Sci Rep. 2016;6:30315. doi: 10.1038/srep30315. PMID:
- 480 27461756.
- 481 33. Laporte MNC, Zuberbühler K. The development of a greeting signal in wild chimpanzees. Dev Sci.
- 482 2011;14(5):1220–1234. doi: 10.1111/j.1467-7687.2011.01069.x. PMID: 21884337.
- 483 34. Oller DK, Griebel U, Iver SN, Jhang Y, Warlaumont AS, Dale R, et al. Language origins viewed in
- 484 spontaneous and interactive vocal rates of human and bonobo infants. Front Psychol. 2019;10:729.
- 485 doi: 10.3389/fpsyg.2019.00729. PMID: 31001176.
- 486 35. Fröhlich M, Wittig RM, Pika S. Play-solicitation gestures in chimpanzees in the wild: flexible
- 487 adjustment to social circumstances and individual matrices. R Soc Open Sci. 2016;3(8):160278. doi:
- 488 10.1098/rsos.160278. PMID: 27853603.

- 489 36. Wich SA, Krützen M, Lameira AR, Nater A, Arora N, Bastian ML, et al. Call cultures in orang-utans?
- 490 PLoS One. 2012;7(5):e36180. doi: 10.1371/journal.pone.0036180. PMID: 22586464.
- 491 37. Knox A, Markx J, How E, Azis A, Hobaiter C, van Veen FJF, et al. Gesture use in communication
- 492 between mothers and offspring in wild orang-utans (*Pongo pygmaeus wurmbii*) from the Sabangau
- 493 Peat-Swamp Forest, Borneo. Int J of Primatol. 2019;40(3):393–416. doi.org/10.1007/s10764-019-
- 494 00095-w.
- 495 38. Fröhlich M, Wittig RM, Pika S. Should I stay or should I go? Initiation of joint travel in mother–infant
- 496 dyads of two chimpanzee communities in the wild. Anim Cogn. 2016;19(3):483–500. doi:
- 497 10.1007/s10071-015-0948-z. PMID: 26833496.
- 498 39. Halina M, Rossano F, Tomasello M. The ontogenetic ritualization of bonobo gestures. Anim Cogn.
 2013;16(4):653-66. doi: 10.1007/s10071-013-0601-7. PMID: 23370783.
- 500 40. Luef EM, Liebal K. Infant-Directed Communication in Lowland Gorillas (*Gorilla gorilla*): Do older
- animals scaffold communicative competence in infants? Am J Primatol. 2012;74(9):841-52. doi:
- 502 10.1002/ajp.22039. PMID: 22644596.
- 503 41. Fröhlich M, Müller G, Zeiträg C, Wittig RM, Pika S. Gestural development of chimpanzees in the wild:
- the impact of interactional experience. Anim Behav. 2017;134:271–282. doi:
- 505 10.1016/j.anbehav.2016.12.018.
- 506 42. Fröhlich M, van Schaik C. Social tolerance and interactional opportunities as drivers of gestural
- 507 redoings in orang-utans. Philos Trans R Soc B. Forthcoming 2022.
- 43. Genty E, Neumann C, Zuberbühler K. Bonobos modify communication signals according to recipient
- 509 familiarity. Sci Rep. 2015;5:16442. doi: 10.1038/srep16442. PMID: 26552655.

510	44. Tomasello M, Call J, Warren J, Frost GT, Carpenter M, Nagell K. The ontogeny of chimpanzee
511	gestural signals: A comparison across groups and generations. EOC. 1997;1(2):223–259. doi:
512	10.1075/eoc.1.2.04tom.
513	45. Ferguson CA. Talking to children: A search for universals. In: Greenberg JH, editor. Universals of

- 514 human language. Standford: Stanford University Press; 1978. p. 203–224.
- 515 46. Fernald A, Taeschner T, Dunn J, Papoušek M, De Boysson-Bardies B, Fukui I. A cross-language study

of prosodic modifications in mothers' and fathers' speech to preverbal infants. J Child Lang. 1989

517 ;16(3):477-501. doi: 10.1017/s0305000900010679. PMID: 2808569.

518 47. Singh L, Nestor S, Parikh C, Yull A. Influences of infant-directed speech on early word recognition.

519 Infancy. 2009;14(6):654–666. doi: 10.1080/15250000903263973. PMID: 32693515.

48. Song JY, Demuth K, Morgan J. Effects of the acoustic properties of infant-directed speech on infant

521 word recognition. J Acoust Soc Am. 2010;128(1):389-400. doi: 10.1121/1.3419786. PMID: 20649233.

49. Porritt LL, Zinser MC, Bachorowski JA, Kaplan PS. Depression diagnoses and fundamental frequency-

523 based acoustic cues in maternal infant-directed speech. Lang Learn Dev. 2014;10(1):51-67. doi:

524 10.1080/15475441.2013.802962. PMID: 24489521.

525 50. Rowe ML. Longitudinal investigation of the role of quantity and quality of child-directed speech in

526 vocabulary development. Child Dev. 2012;83(5):1762-74. doi: 10.1111/j.1467-8624.2012.01805.x.

527 PMID: 22716950.

51. Hartman KM, Ratner NB, Newman RS. Infant-directed speech (IDS) vowel clarity and child language
outcomes. J Child Lang. 2017;44(5):1140-1162. doi: 10.1017/S0305000916000520. PMID: 27978860.

530 52. Tomasello M, Farrar MJ. Joint attention and early language. Child Dev. 1986;57(6):1454-63. PMID:

531 3802971.

- 532 53. Burkart JM, Hrdy SB, Van Schaik CP. Cooperative breeding and human cognitive evolution. Evol
- 533 Anthropol. 2009;18(5):175–186. doi: 10.1002/evan.20222.
- 534 54. Fernald A, Simon T. Expanded intonation contours in mothers' speech to newborns. Dev Psychol.
- 535 1984;20(1):104–113. doi: 10.1037/0012-1649.20.1.104.
- 536 55. Ratner NB. Durational cues which mark clause boundaries in mother–child speech. J Phon.
- 537 1986;14(2):303–309. doi: 10.1016/S0095-4470(19)30670-9.
- 538 56. Uther M, Knoll MA, Burnham D. Do you speak E-NG-L-I-SH? A comparison of foreigner- and infant-
- 539 directed speech. Speech Commun. 2007;49(1):2–7. doi: 10.1016/j.specom.2006.10.003.
- 540 57. Kuhl PK, Andruski JE, Chistovich IA, Chistovich LA, Kozhevnikova EV, Ryskina VL, et al. Cross-language
- analysis of phonetic units in language addressed to infants. Science. 1997;277(5326):684–686. doi:
- 542 10.1126/science.277.5326.684. PMID: 9235890.
- 543 58. Fernald A. Intonation and communicative intent in mothers' speech to infants: Is the melody the
- 544 message? Child Dev. 1989;60(6):1497–1510. PMID: 2612255.
- 545 59. Englund KT. Voice onset time in infant directed speech over the first six months. First Lang.
- 546 2005;25(2):219–234. doi: 10.1177/0142723705050286.
- 547 60. Cooper RP, Aslin RN. Preference for infant-directed speech in the first month after birth. Child Dev.
- 548 1990;61(5):1584–1595. PMID: 2245748.
- 549 61. Papoušek M, Papoušek H, Haekel M. Didactic adjustments in fathers' and mothers' speech to their
- 550 3-month-old infants. J Psycholinguist Res. 1987;16(5):491–516.
- 551 62. Cameron-Faulkner T, Lieven E, Tomasello M. A construction based analysis of child directed speech.
- 552 Cogn Sci. 2003;27(6):843–873. doi: 10.1207/s15516709cog2706_2.

- 63. Snow CE. The development of conversation between mothers and babies. J Child Lang. 1977;4(1):1–
- 554 22. doi:10.1017/S030500090000453.
- 64. Martin A, Igarashi Y, Jincho N, Mazuka R. Utterances in infant-directed speech are shorter, not
 slower. Cognition. 2016;156:52–59. doi: 10.1016/j.cognition.2016.07.015.
- 557 65. Phillips JR. Syntax and vocabulary of mothers' speech to young children: Age and sex comparisons.
- 558 Child Dev. 1973;44(1):182–185.
- 66. Henninga A, Striano T, Lieven EVM. Maternal speech to infants at 1 and 3 months of age. Infant
 Behav Dev. 2005;28(4):519–536. doi: 10.1016/j.infbeh.2005.06.001.
- 561 67. Kempe V, Brooks PJ, Pirott L. How can child-directed speech facilitate the acquisition of
- 562 morphology? In: Research on child language acquisition: Proceedings of the 8th conference of the
- 563 International Association for the Study of Child Language. Cascadilla Press; 2001. p. 1237–1247.
- 564 68. Kempe V, Brooks PJ, Gillis S. Diminutives provide multiple benefits for language acquisition. In:
- 565 Savickienė I, Dressler WU, editors. The Acquisition of diminutives: A cross-linguistic perspective.
- 566 Amsterdam: John Benjamins B. V.; 2007. p. 319–342.
- 567 69. Newport E, Gleitman H, Gleitman L. Mother, I'd rather do it myself: Some effects and non-effects of
- 568 motherese. In: Snow CE, Ferguson CA, editors. Talking to children. Cambridge: Cambridge University
- 569 Press; 1977. p. 109–149.
- 570 70. Soderstrom M, Blossom M, Foygel R, Morgan JL. Acoustical cues and grammatical units in speech to
- 571 two preverbal infants. J Child Lang. 2008;35(4):869-902. doi: 10.1017/S0305000908008763. PMID:
- 572 18838016.
- 573 71. Küntay A, Slobin DI. Listening to a Turkish Mother: Some Puzzles for Acquisition. In: Slobin DI,
- 574 Gerhardt J, Kyratzis A, Guo J, editors. Social interaction, social context, and language: essays in
- honor of Susan-Ervin Tripp. Hillsdale, NJ: Lawrence Erlbaum Associates; 1996. p. 265–286.

- 576 72. Lester NA, Moran S, Küntay AC, Allen SEM, Pfeiler B, Stoll S. Detecting structured repetition in child-
- 577 surrounding speech: Evidence from maximally diverse languages. Cognition. 2022;221:104986. doi:
- 578 10.1016/j.cognition.2021.104986. PMID: 34953269.
- 579 73. Yurovsky D, Yu C, Smith LB. Statistical speech segmentation and word learning in parallel: scaffolding
- 580 from child-directed speech. Front Psychol. 2012;3:374. doi: 10.3389/fpsyg.2012.00374. PMID:

581 23162487.

582 74. Albin DD, Echols CH. Stressed and word-final syllables in infant-directed speech. Infant Behav Dev.

583 1996;19(4):401–418. doi.org/10.1016/S0163-6383(96)90002-8.

584 75. Trainor LJ, Desjardins RN. Pitch characteristics of infant-directed speech affect infants' ability to

585 discriminate vowels. Psychon Bull Rev. 2002;9(2):335-40. doi: 10.3758/bf03196290. PMID:

586 12120797.

- 587 76. Werker JF, McLeod PJ. Infant preference for both male and female infant-directed talk: A
- 588 developmental study of attentional and affective responsiveness. Can J Psychol. 1989;43(2):230-46.
- doi: 10.1037/h0084224. PMID: 2486497.
- 590 77. Cohen D, Cassel RS, Saint-Georges C, Mahdhaoui A, Laznik MC, Apicella F, et al. Do parentese

591 prosody and fathers' involvement in interacting facilitate social interaction in infants who later

- 592 develop autism? PLoS One. 2013;8(5):e61402. doi: 10.1371/journal.pone.0061402. PMID:
- 593 23650498.
- 594 78. Dunst CJ, Gorman E, Hamby DW. Preference for infant-directed speech in preverbal young children.
 595 CELL. 2012;5(1).
- 596 79. Fernald A. Four-month-old infants prefer to listen to motherese. Infant Behav Dev. 1985;8(2):181–

597 195. doi: 10.1016/S0163-6383(85)80005-9.

- 598 80. Naoi N, Minagawa-Kawai Y, Kobayashi A, Takeuchi K, Nakamura K, Yamamoto JI, et al. Cerebral
- responses to infant-directed speech and the effect of talker familiarity. NeuroImage.

600 2012;59(2):1735–1744. doi: 10.1016/j.neuroimage.2011.07.093. PMID: 21867764.

- 601 81. Niwano K, Sugai K. Acoustic determinants eliciting Japanese infants' vocal response to maternal
- 602 speech. Psychol Rep. 2002;90(1):83-90. doi: 10.2466/pr0.2002.90.1.83. PMID: 11899017.
- 603 82. Karzon RG. Discrimination of polysyllabic sequences by one- to four-month-old infants. J Exp Child
 604 Psychol. 1985;39(2):326-42. doi: 10.1016/0022-0965(85)90044-x. PMID: 3989467.

605 83. Bortfeld H, Morgan JL. Is early word-form processing stress-full? How natural variability supports

- 606 recognition. Cogn Psychol. 2010;60(4):241-66. doi: 10.1016/j.cogpsych.2010.01.002. PMID:
- 607 20159653.
- 608 84. Lyakso EE, Frolova OV, Grigorev AS. Infant vocalizations at the first year of life predict speech

609 development at 2-7 years: Longitudinal study. Psychology. 2014;5(12). doi:

- 610 10.4236/psych.2014.512154.
- 611 85. Thiessen ED, Hill EA, Saffran JR. Infant-directed speech facilitates word segmentation. Infancy.
- 612 2005;7(1):53–71. doi: 10.1207/s15327078in0701_5. PMID: 33430544.
- 613 86. Herold DS, Nygaard LC, Namy LL. Say it like you mean it: Mothers' use of prosody to convey word
 614 meaning. Lang Speech. 2012;55(Pt 3):423-36. doi: 10.1177/0023830911422212. PMID: 23094322.
- 615 87. Spinelli M, Fasolo M, Mesman J. Does prosody make the difference? A meta-analysis on relations
- 616 between prosodic aspects of infant-directed speech and infant outcomes. Dev Rev. 2017;44:1–18.
- 617 doi.org/10.1016/j.dr.2016.12.001.
- 618 88. Ambridge B, Kidd E, Rowland CF, Theakston AL. The ubiquity of frequency effects in first language
- 619 acquisition. J Child Lang. 2015;42(2):239-73. doi: 10.1017/S030500091400049X. PMID: 25644408.

- 620 89. Lieven E. Input and first language acquisition: Evaluating the role of frequency. Lingua.
- 621 2010;120(11):2546–2556. doi: 10.1016/j.lingua.2010.06.005.
- 622 90. Stoll S, Abbot-Smith K, Lieven E. Lexically Restricted Utterances in Russian, German, and English
- 623 child-directed speech. Cogn Sci. 2009;33(1):75-103. doi: 10.1111/j.1551-6709.2008.01004.x. PMID:
- 624 21585464.
- 91. Mintz TH. Frequent frames as a cue for grammatical categories in child directed speech. Cognition.
 2003;90:91–117. doi: 10.1016/s0010-0277(03)00140-9. PMID: 14597271.
- 627 92. Moran S, Blasi DE, Schikowski R, Küntay AC, Pfeiler B, Allen S, et al. A universal cue for grammatical
- 628 categories in the input to children: Frequent frames. Cognition. 2018;175:131–140. doi:
- 629 10.1016/j.cognition.2018.02.005. PMID: 29518682.
- 630 93. Hills T. The company that words keep: comparing the statistical structure of child-versus adult-
- 631 directed language. J Child Lang. 2013;40(3):586-604. doi: 10.1017/S0305000912000165. PMID:
- 632 22584041.
- 633 94. You G, Bickel B, Daum MM, Stoll S. Child-directed speech is optimized for syntax-free semantic
- 634 inference. Sci Rep. 2021;11(1):16527. doi: 10.1038/s41598-021-95392-x. PMID: 34400656.
- 635 95. Fernald A, Hurtado N. Names in frames: Infants interpret words in sentence frames faster than
- 636 words in isolation. Dev Sci. 2006;9(3):33–40. doi: 10.1111/j.1467-7687.2006.00482.x. PMID:
- 637 16669790.
- 638 96. Lew-Williams C, Pelucchi B, Saffran JR. Isolated words enhance statistical language learning in
- 639 infancy. Dev Sci. 2011;14(6):1323–1329. doi: 10.1111/j.1467-7687.2011.01079.x. PMID: 22010892.
- 640 97. Waterfall HR. A little change is a good thing: Feature theory, language acquisition and variation sets
- 641 [dissertation]. Chicago: University of Chicago; 2006.

- 642 98. Schwab JF, Lew-Williams C. Language learning, socioeconomic status, and child-directed speech.
- 643 Wiley Interdiscip Rev Cogn Sci. 2016;7(4):264-75. doi: 10.1002/wcs.1393. PMID: 27196418.
- 644 99. Hart B, Risley TR. Meaningful differences in the everyday experience of young American children.
- 645 Baltimore: Paul H Brookes Publishing; 1995.
- 646 100. Pan BA, Rowe ML, Singer JD, Snow CE. Maternal correlates of growth in toddler vocabulary
- 647 production in low-income families. Child Dev. 2005;76(4):763-82. doi: 10.1111/j.1467-
- 648 8624.2005.00876.x. PMID: 16026495.
- 649 101. Weisleder A, Fernald A. Talking to children matters: Early language experience strengthens
- 650 processing and builds vocabulary. Psychol Sci. 2013;24(11):2143-52. doi:
- 651 10.1177/0956797613488145. PMID: 24022649.
- 652 102. Newman RS, Rowe ML, Bernstein Ratner N. Input and uptake at 7 months predicts toddler
- 653 vocabulary: The role of child-directed speech and infant processing skills in language development. J
- 654 Child Lang. 2016;43(5):1158-73. doi: 10.1017/S0305000915000446. PMID: 26300377.
- 655 103. Huttenlocher J, Waterfall H, Vasilyeva M, Vevea J, Hedges LV. Sources of variability in children's
- 656 language growth. Cogn Psychol. 2010;61(4):343-65. doi: 10.1016/j.cogpsych.2010.08.002. PMID:
- 657 20832781.
- 658 104. Hirsh-Pasek K, Adamson LB, Bakeman R, Owen MT, Golinkoff RM, Pace A, et al. The contribution
- of early communication quality to low income children's language success. Psychol Sci.
- 660 2015;26(7):1071-83. doi: 10.1177/0956797615581493. PMID: 26048887.
- 661 105. Lieven EV. Crosslinguistic and crosscultural aspects of language addressed to children. In:
- 662 Gallaway C, Richards BJ, editors. Input and interaction in language acquisition. Cambridge:
- 663 Cambridge University Press; 1994. p. 56–73.

- 664 106. Ochs E, Schieffelin, Bambi B. Language acquisition and socializations: Three developmental
- stories and their implications. In: Shweder RA, LeVine RA, editors. Culture theory: Essays on mind,
- 666 self and emotion. Cambridge: Cambridge University Press; 1984. p. 276–320.
- 667 107. Henrich J, Heine SJ, Norenzayan A. Most people are not WEIRD. Nature. 2010;466(7302):29–29.
- 668 doi: 10.1038/466029a.
- 669 108. Vogt P, Mastin JD, Schots DM. Communicative intentions of child-directed speech in three
- 670 different learning environments: Observations from the Netherlands, and rural and urban
- 671 Mozambique. First Lang. 2015;35(4-5):341–358. doi: 10.1177/0142723715596647.
- 672 109. Casillas M, Brown P, Levinson SC. Early language experience in a Tseltal Mayan village. Child Dev.
- 673 2020;91(5):1819-1835. doi: 10.1111/cdev.13349. PMID: 31891183.
- 674 110. Bergelson E, Casillas M, Soderstrom M, Seidl A, Warlaumont AS, Amatuni A. What do North
- 675 American babies hear? A large-scale cross-corpus analysis. Dev Sci. 2019;22(1):e12724. doi:
- 676 10.1111/desc.12724. PMID: 30369005.
- 677 111. Keller H. Cultures of infancy. Mahwah, NJ: Lawrence Erlbaum Associates; 2013.
- 678 112. Sarvasy H, Elvin J, Li W, Escudero P. An acoustic analysis of Nungon vowels in child-versus adult-
- directed speech. In: Proceedings of the 19th International Congress of Phonetic Sciences Melbourne;
- 680 2019. p. 3155–3159.
- 681 113. Broesch TL, Bryant GA. Prosody in infant-directed speech Is similar across Western and
- 682 traditional cultures. J Cogn Dev. 2015;16(1):31–43. doi.org/10.1080/15248372.2013.833923.
- 683 114. Pye C. Quiché Mayan speech to children. J Child Lang. 1986;13(1):85-100. doi:
- 684 10.1017/s030500090000313. PMID: 3949901.

- Farran LK, Lee CC, Yoo H, Oller DK. Cross-cultural register differences in infant-directed speech:
 An initial study. PLoS One. 2016;11(3):e0151518. doi: 10.1371/journal.pone.0151518. PMID:
 26981626.
- 688 116. Ratner NB, Pye C. Higher pitch in BT is not universal: Acoustic evidence from Quiché Mayan. J
- 689 Child Lang. 1984;11(3):515-22. doi: 10.1017/s0305000900005924. PMID: 6501462.
- 690 117. Stern DN, Spieker S, Barnett RK, MacKain K. The prosody of maternal speech: Infant age and
 691 context related changes. J Child Lang. 1983;10(1):1-15. doi: 10.1017/s0305000900005092. PMID:
 692 6841483.
- 693 118. Vosoughi S, Roy D. A longitudinal study of prosodic exaggeration in child-directed speech in
- 694 Speech Prosody. In: 6th International Conference. SProSIG; 2012.
- 695 119. Kondaurova MV, Bergeson TR, Xu H. Age-related changes in prosodic features of maternal
- 696 speech to prelingually deaf infants with cochlear implants. Infancy. 2013;18(5):825–848. doi:
- 697 10.1111/infa.12010. PMID: 24244108.
- 698 120. Narayan CR, McDermott LC. Speech rate and pitch characteristics of infant-directed speech:
- Longitudinal and cross-linguistic observations. J Acoust Soc Am. 2016;139(3):1272-81. doi:
- 700 10.1121/1.4944634. PMID: 27036263.
- 701 121. Fernald A. Meaningful melodies in mothers' speech to infants. In: Papoušek H, Jürgens U,

702 Papoušek M, editors. Nonverbal vocal behaviour. Cambridge: Cambridge University Press; 1992. p.

- 703 262–282.
- 704 122. Gratier M, Devouche E, Guellai B, Infanti R, Yilmaz E, Parlato-Oliveira E. Early development of
- turn-taking in vocal interaction between mothers and infants. Front Psychol. 2015;6:1167. doi:
- 706 10.3389/fpsyg.2015.01167. PMID: 26388790.
- 707 123. Zimmermann E. Aspects of reproduction and behavioral and vocal development in Senegal
- 708 bushbabies (*Galago senegalensis*). Int J Primatol. 1989;10(1):1–16. doi.org/10.1007/BF02735700.

- 709 124. Biben M, Symmes D, Bernhards D. Contour variables in vocal communication between squirrel
 710 monkey mothers and infants. Dev Psychobiol. 1989;22(6):617-31. doi: 10.1002/dev.420220607.
 711 PMID: 2792572.
- 712 125. Szenczi P, Bánszegi O, Urrutia A, Faragó T, Hudson R. Mother-offspring recognition in the
- domestic cat: Kittens recognize their own mother's call. Dev Psychobiol. 2016;58(5):568-77. doi:
- 714 10.1002/dev.21402. PMID: 26935009.
- 715 126. Sèbe F, Duboscq J, Aubin T, Ligout S, Poindron P. Early vocal recognition of mother by lambs:
- contribution of low- and high-frequency vocalizations. Anim Behav. 2010;79(5):1055–1066. doi:
- 717 10.1016/j.anbehav.2010.01.021.
- 718 127. Padilla De La Torre M, Briefer EF, Ochocki BM, McElligott AG, Reader T. Mother-offspring
- recognition via contact calls in cattle, Bos taurus. Anim Behav. 2016;114:147–154. doi:
- 720 10.1016/j.anbehav.2016.02.004.
- 721 128. Volodin IA, Sibiryakova OV, Soldatova NV, Volodina EV. Acoustically different contact calls of
- mother and young goitred gazelle are equally individualized: Is this a common relationship in

723 ruminants? Behaviour. 2019;156(12):1185–1207. doi: 10.1163/1568539X-00003561.

- 724 129. Sibiryakova OV, Volodin IA, Frey R, Zuther S, Kisebaev TB, Salemgareev AR, et al. Remarkable
- vocal identity in wild-living mother and neonate saiga antelopes: A specialization for breeding in
- 726 huge aggregations? Sci Nat. 2017;104(3-4):11. doi: 10.1007/s00114-017-1433-0. PMID: 28243711.
- 130. Balcombe JP, McCracken GF. Vocal recognition in Mexican free-tailed bats: do pups recognize
 mothers? Anim Behav. 1992;43(1):79–87. doi: 10.1016/S0003-3472(05)80961-3.
- 729 131. Charrier I, Mathevon N, Jouventin P. Mother's voice recognition by seal pups. Nature.
- 730 2001;412(6850):873. doi: 10.1038/35091136. PMID: 11528465.

Ruch H, Zürcher Y, Burkart JM. The function and mechanism of vocal accommodation in humans
and other primates. Biol Rev Camb Philos Soc. 2018;93(2):996-1013. doi: 10.1111/brv.12382. PMID:
29111610.

734 133. Weiss BM, Ladich F, Spong P, Symonds H. Vocal behavior of resident killer whale matrilines with

- newborn calves: The role of family signatures. J Acoust Soc Am. 2006;119(1):627-35. doi:
- 736 10.1121/1.2130934. PMID: 16454316.
- 737 134. Zürcher Y, Willems EP, Burkart JM. Trade-offs between vocal accommodation and individual
- recognisability in common marmoset vocalizations. Sci Rep. 2021;11(1):15683. doi: 10.1038/s41598-
- 739 021-95101-8. PMID: 34344939.
- 740 135. Castro NA, Snowdon CT. Development of vocal responses in infant cotton-top tamarins.
- 741 Behaviour. 2000;137(5):629–646. doi: 10.1163/156853900502259.
- 742 136. Koda H, Lemasson A, Oyakawa C, Pamungkas J, Masataka N. Possible role of mother-daughter
- vocal interactions on the development of species-specific song in gibbons. PLoS One.
- 744 2013;8(8):e71432. doi: 10.1371/journal.pone.0071432. PMID: 23951160.
- 745 137. Takahashi DY, Liao DA, Ghazanfar AA. Vocal learning via social reinforcement by infant
- 746 marmoset monkeys. Curr Biol. 2017;27(12):1844-1852.e6. doi: 10.1016/j.cub.2017.05.004. PMID:
 747 28552359.
- 748 138. Chow CP, Mitchell JF, Miller CT. Vocal turn-taking in a non-human primate is learned during
- 749 ontogeny. Proc Biol Sci B. 2015;282(1807):20150069. doi: 10.1098/rspb.2015.0069. PMID:
- 25904663.
- 751 139. Takahashi DY, Fenley AR, Ghazanfar AA. Early development of turn-taking with parents shapes
- vocal acoustics in infant marmoset monkeys. Philos Trans R Soc Lond B Biol Sci.
- 753 2016;371(1693):20150370. doi: 10.1098/rstb.2015.0370. PMID: 27069047.

754	140.	Chen Y, Matheson LE, Sakata JT. Mechanisms underlying the social enhancement of vocal
755	lea	arning in songbirds. Proc Natl Acad Sci USA. 2016;113(24):6641-6. doi: 10.1073/pnas.1522306113
756	PN	MID: 27247385.

- 757 141. Fernandez AA, Knörnschild M. Pup directed vocalizations of adult females and males in a vocal
 758 learning bat. Front Ecol Evol. 2020;8:265. doi: 10.3389/fevo.2020.00265.
- 759 142. Koda H. Possible use of heterospecific food-associated calls of macaques by sika deer for
- foraging efficiency. Behav Processes. 2012;91(1):30-4. doi: 10.1016/j.beproc.2012.05.006. PMID:

761 22641112.

- 762 143. Brown P. The Cultural organization of attention. In: Duranti A, Ochs E, Schieffelin BB, editors.
- The handbook of language socialization. Amsterdam: John Benjamins B.V.; 2011. p. 29–55.
- 764 144. De León L. Language socialization and multiparty participation frameworks. In: A Duranti EO,
- 765 Schieffelin BB, editors. The handbook of language socialization. Amsterdam: John Benjamins B.V.;

766 2011. p. 81–111.

- 767 145. Ivey PK. Cooperative reproduction in Ituri forest hunter-gatherers: Who cares for Efe infants?
 768 Curr Anthropol. 2000;41(5):856–866. doi: 10.1086/317414
- 769 146. Casillas M, Brown P, Levinson SC. Early language experience in a Papuan community. J Child
- 770 Lang. 2021;48(4):792-814. doi: 10.1017/S0305000920000549. PMID: 32988426.
- 147. Hobaiter C, Byrne RW, Zuberbühler K. Wild chimpanzees' use of single and combined vocal and
- 772 gestural signals. Behav Ecol Sociobiol. 2017;71(6):96. doi: 10.1007/s00265-017-2325-1. PMID:
- 28596637.
- 148. Carpenter M, Nagell K, Tomasello M. Social cognition, joint attention, and communicative
- competence from 9 to 15 months of age. Monogr Soc Res Child Dev. 1998;63(4):i-vi, 1-143.

- 776 149. Scott K, Sakkalou E, Ellis-Davies K, Hilbrink E, Hahn U, Gattis M. Infant contributions to joint
- attention predict vocabulary development. In: CogSci 2013: the 35th Annual Conference of the
- 778 Cognitive Science Society. Cognitive Science Society; 2013. p. 3384–3389.
- 779 150. Rowland CF, Pine JM, Lieven EVM, Theakston AL. Determinants of acquisition order in wh-
- 780 questions: Re-evaluating the role of caregiver speech. J Child Lang. 2003;30(3):609-35. PMID:
- 781 14513470.