






## Review

## Plant–insect chemical communication in ecological communities: An information theory perspective

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**Abstract** Cross-species communication, where signals are sent by one species and perceived by others, is one of the most intriguing types of communication that functionally links different species to form complex ecological networks. Global change and human activity can affect communication by increasing fluctuations in species composition and phenology, altering signal profiles and intensity, and introducing noise. So far, most studies on cross-species communication have focused on a few specific species isolated from ecological communities. Scaling up investigations of cross-species communication to the community level is currently hampered by a lack of conceptual and practical methodologies. Here, we propose an interdisciplinary framework based on information theory to investigate mechanisms shaping cross-species communication at the community level. We use plants and insects, the cornerstones of most ecosystems, as a showcase and focus on chemical communication as the key communication channel. We first introduce some basic concepts of information theory, then we illustrate information patterns in plant–insect chemical communication, followed by a further exploration of how to integrate information theory into ecological and evolutionary processes to form testable mechanistic hypotheses. We conclude by highlighting the importance of community-level information as a means to better understand the maintenance and workings of ecological systems, especially during rapid global change.

**Key words:** chemical communication, ecological networks, functional traits, information theory, plant–insect interaction, volatile organic compounds, Zipf's law.

## 1 Introduction

Communication is prevalent in nature. For example, honeybees waggle to send information that guides other bees in the colony (Von Frisch, 1974); birds sing to mate or alert others in the flock (Freeberg, 2008); and flowers exhibit colors and scents that can attract pollinators or deter herbivores (Schoonhoven et al., 2005). Cross-kingdom (Plantae and Animalia) plant–insect communication is of extreme interest not only because of its ubiquity and fundamental roles in both natural and agricultural systems (Seastedt & Crossley, 1984; Strong et al., 1984; Isbell et al., 2011; Potts et al., 2016; Ollerton, 2021) but also due to the extraordinarily diverse communication mediators

shaped by hundreds of millions of years of co-evolution (Ehrlich & Raven, 1964).

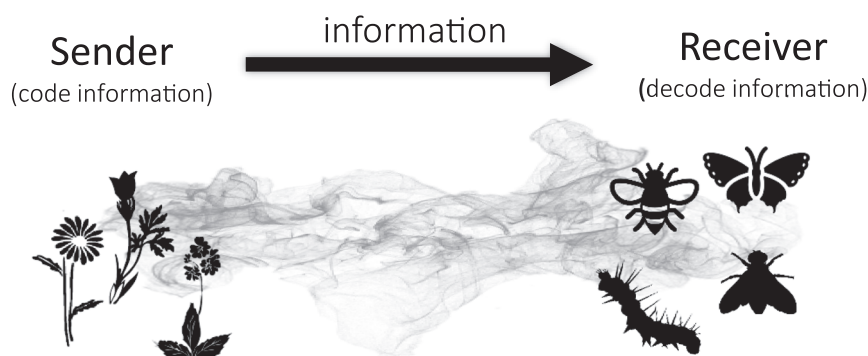
Chemical communication is one of the most ancient and pivotal means for plants and insects. In fact, chemistry underlies color (pigments), shape (genetic encoding in nucleic acids and chemical inducers such as hormones), and scent (volatile organic compounds, or VOCs). Yet, chemical mediators of communication are most often considered from among the vast number of secondary metabolites produced by plants, with an estimate of 200 000 compounds including VOCs that have been extracted and identified (Kessler & Kalske, 2018; Robbins, 2000), which are not strictly required for plant growth and development but may have important defensive and attractive functions (Fraenkel, 1959; Ehrlich

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& Raven, 1964; Pichersky & Gershenzon, 2002). Insects, with their numerous chemoreceptors, are very sensitive to many plant chemicals (Schoonhoven et al., 2005; Kaupp, 2010; Hansson & Stensmyr, 2011) and depend on chemical signals for fundamental activities, such as foraging (Schiestl, 2010), mating (Alexander et al., 1997), and oviposition (Renwick & Chew, 1994). Presumably, plant-feeding insects and their natural enemies have been under strong selective pressure to perceive and interpret plant chemicals that are relevant for these essential activities. So far, most plant-insect chemical communication studies have focused on specific species, isolating them from the community context in which they are naturally embedded. However, scaling up plant-insect communication from the species to the community level can be possible through the lens of information theory and communication systems (Shannon, 1948).

Communication can be considered as a process of information transfer from a sender who encodes the information (e.g., visual, vocal, or olfactory), to a receiver who decodes the information (Fig. 1). Depending on how well

the information is reproduced at the receiver's end, we can quantify the clarity or ambiguity of the communication (Shannon, 1948). That is, the better a message is reproduced, the greater amount of information (clarity) it contains, and the lower its entropy (ambiguity, uncertainty). This mathematical theory of communication is called information theory. Figure 1 summarizes key information quantities for a vector of signals (entropy and self-information of each signal) and for an object-signal matrix (conditional entropy and mutual information). Information theory was originally developed for man-made communication systems, such as telecommunication, and later applied to a wide range of communication-related fields, for example, cybernetics (Gabor, 1954), cryptography (Ahlsweide & Csiszár, 1993), linguistics (Zipf, 1949; Ferrer-i-Cancho & Solé, 2003), and neurobiology (Sharpee et al., 2014). Information theory brings at least two important new features that make it a promising tool for various systems and scales. First, information theory translates diverse communication signals into information. Thus, it unites different signals by extracting their information content without being overwhelmed by their different identities. Second, information



### Vector

For a vector  $S$  with  $n$  signals:  $S = (S_1, S_2, \dots, S_n)$ ,

Probability of each signal is:  $(p_1, p_2, \dots, p_n)$ ,

Using information theory, we can calculate:

**Entropy ( $H$ )** of the information, which describes the uncertainty (or the ambiguity) of the signals:

$$H = - \sum_{i=1}^n p_i \log p_i$$

The information each signal contains is called

**Self information ( $I$ ):**

$$I(S_i) = \log \frac{1}{p_i}$$

### Matrix

	Signal1	...	Signal n
Object1	$OS_{11}$	$\cdots$	$OS_{1n}$
...	$\vdots$	$\ddots$	$\vdots$
Object m	$OS_{m1}$	$\cdots$	$OS_{mn}$

(Object-Signal matrix)

For a matrix with  $n$  signals and the corresponding  $m$  objects, we can calculate:

**Conditional entropy** quantifies the uncertainty (or the amount of information needed) about one variable given another variable.

$$H_n(O|S) = \sum_{j=1}^n p(S_j) H_n(O|S_j),$$

$$\text{Where } p(S_j) = \frac{\sum_{i=1}^m OS_{i,j}}{\sum_{i=1, j=1}^m OS_{i,j}},$$

$$H_n(O|S_j) = - \sum_{i=1}^m p(O_i|S_j) \log_{2n} p(O_i|S_j),$$

$$p(O_i|S_j) = \frac{OS_{i,j}}{\sum_{i=1}^m OS_{i,j}}.$$

**Mutual information** quantifies mutual dependence between two variables. In our example of object-signal matrix, it quantifies the amount of information obtained about the object by observing the signals.

$$I(O, S) = H_m(O) - H_n(O|S),$$

$$\text{Where } H_m(O) = - \sum_{i=1}^m p(O_i) \log_{2n} p(O_i),$$

$$p(O_i) = \frac{1}{m},$$

$$I(O, S) = I(S, O).$$

**Fig. 1.** Information transferring process in communication and formula quantifying information in communication systems.

theory measures the reproducibility of information between two states based on probabilistic concepts (Boso & Tartakovsky, 2018). This nonparametric perspective allows the scaling of information across different dimensions.

Not surprisingly, information theory has long been proposed as a conceptual and quantitative framework to study information transfer across ecological systems (Margalef, 1968; Ulanowicz, 2001; O'Connor et al., 2019). However, concrete applications to plant–insect communication are just starting to emerge (Zu et al., 2020). In this paper, we illustrate the application of information theory as a means to increase our understanding of plant–insect communication. In particular, we propose to consider plant VOCs as communication signals and explore the emerging information patterns of plant–insect chemical communication. We then discuss potential future directions to uncover the underlying mechanisms of plant–insect communication by the integration of information theory with ecological and evolutionary processes. Note that here we use “signals” in an information theory framework to simply refer to *all* plant VOCs that form the repository (regardless of whether they have demonstrated attractant or deterrent functions, or not).

## 2 Patterns of Plant–Insect Chemical Communication

### 2.1 Zipf's law

Zipf's law (Zipf, 1932, 1949) in linguistics describes that the frequency of a word decays rapidly proportional to its rank, or frequency of usage, following a power-law distribution ( $p(r) = ar^{-k}$ , where  $r$  denotes rank). Taking the English language for example, “the” is the most frequent word in corpora, occurring at a rate of 6% in British National Corpus, followed by “of,” which only occurs half as often (3%) (Leech et al., 2014). And there is a large number of words that occur in very low frequency, leading to a heavy tail distribution. The power-law decay distribution is also related to the Pareto principle (or 80-20 rule) known in social sciences: that is, roughly 80% of outcomes result from 20% of causes (Pareto, 1964). Importantly, the power-law exponent describing the speed of decay (i.e., the shape of this tailed distribution) and the redundancy of information in communication systems is around  $k = 1$  in many human languages (Zipf, 1949; Piantadosi, 2014). Zipf's law has also been explored in various biological systems (Semple et al., 2021).

If we consider a plant chemical “language,” we can think of plants emitting different chemical “words” to their environment. Studies on plant VOCs have revealed that some VOCs are more frequent than others (Knudsen et al., 2006; Farré-Armengol et al., 2020), but the detailed information structures have not been explicitly studied. To explore the redundancy of plant chemical language and whether the frequency of its vocabulary (VOCs) follows Zipf's law, we gathered the data from the only four community-level plant VOC studies so far, three of which focused on floral VOCs in plant-pollinator networks (Filella et al., 2013; Kantsa et al., 2017; Burkle & Runyon, 2019) and one on leaf VOCs in a plant-herbivore (caterpillar) network

(Zu et al., 2020). Additionally, we also use a review study by Farré-Armengol et al. (2020), who compiled a floral VOC dataset from 305 species. In the review study, the authors also categorized chemical groups of VOCs, families of plant species, and pollination systems of these plant species.

By analyzing the distributions of VOCs in these studies and in different categories (chemical groups, plant families, pollination groups), we found that most cases follow a heavy-tail distribution where a few VOCs are predominant whereas many other VOCs occur rarely (Fig. 2). In five cases (out of the 16 examined cases and categories), a power-law distribution fits either the whole data (in one community study PV\_k, and the group of N-, S-containing compounds, Fig. 2, Table 1) or part of the data (in Orchidaceae, insect-pollinated flowers, and the overall review data, Fig. 2, Table 1). The fitting was performed using the R package “powerLaw” (Clauset et al., 2009; Gillespie, 2017). Many of the remaining cases appear to have power-law-like distributions, but a fit cannot be statistically supported (Table S1). Indeed, although power-law distributions are thought to be common, it is not easy to achieve a strict fit to many empirical data (Clauset et al., 2009) partially due to limited sample size and sample bias on abundant vs. rare signals.

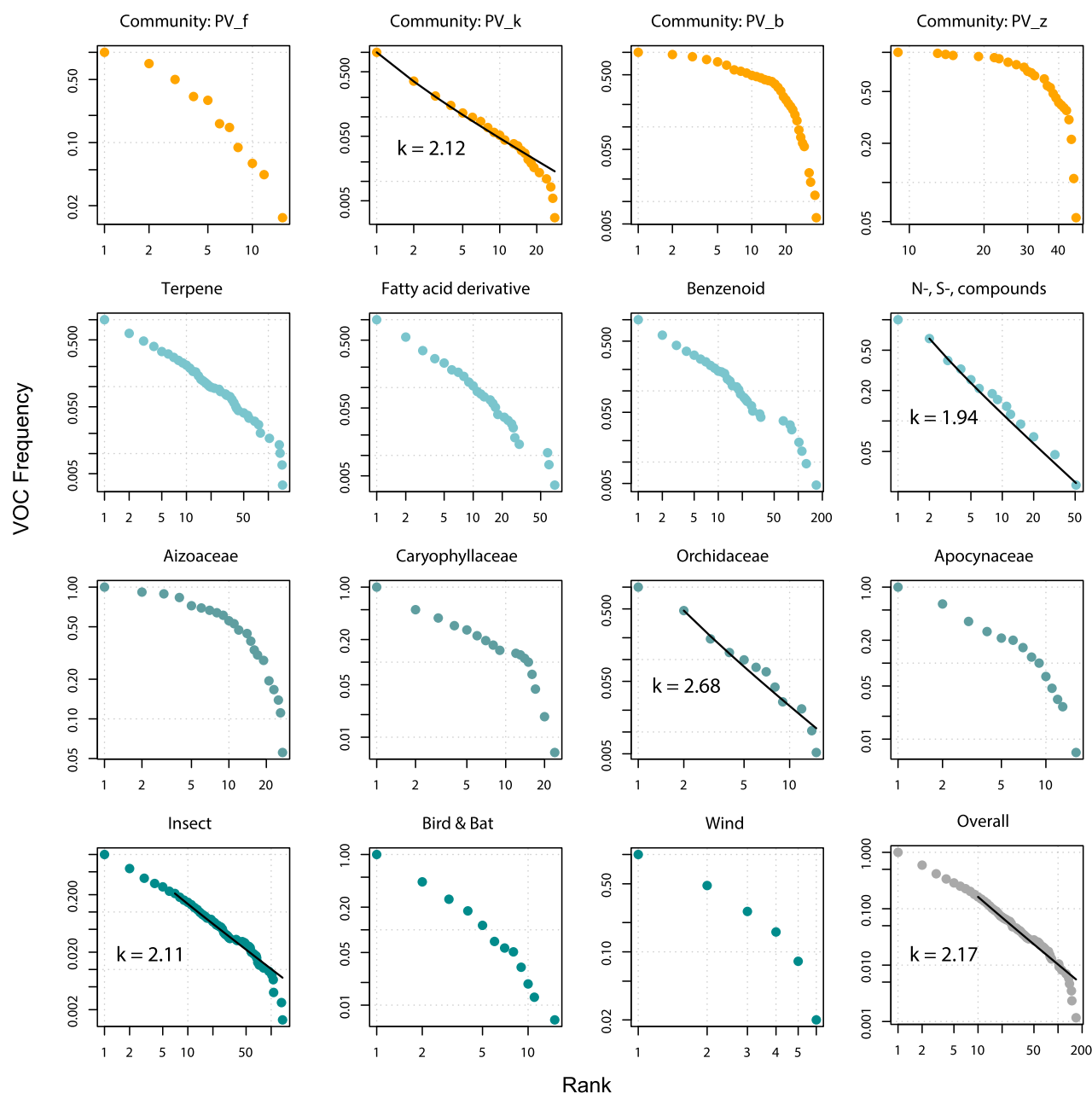
In the five cases in which a full or partial fit can be made, the slopes of the distribution range from  $k = 1.94$  to  $k = 2.68$ , which is steeper than observed in human languages. These patterns suggest that plant VOC languages may be more redundant than human language. Following information theory, the entropy of VOCs in all these cases is very high, ranging from 0.78 to 0.99 ( $H(V)$  in Table 1A).

Note that throughout the whole paper, we used standardized entropy so that it ranges between 0 and 1, where the higher the value (closer to 1), the higher the uncertainty. Entropy calculation formulas can be found in Fig. 1.

### 2.2 Coding, decoding, and interactions

In the previous section, we only focus on the putative signals (VOCs) themselves. In this section, we place these into the context of communication: the coding and decoding of signals, and species interactions. Following the work by Zu et al. (2020), we can use the associations of VOCs (V) with plants (P) in a community to describe the plant coding process (PV-matrix), the associations of VOCs with insects (or animals, A) to describe insect decoding process (AV-matrix), and associations of animals with plants (AP-matrix) for insect-plant interactions. Empirically, to screen how each insect reacts to each of the VOCs in the community requires extensive experimental manipulations under both controlled and natural conditions. Therefore, it is challenging to generate the whole insect decoding matrix from field and experimental work.

From the four community-level plant VOC studies that documented a PV-matrix, three of them (Burkle & Runyon, 2019; Kantsa et al., 2019; Zu et al., 2020) also document insect-plant interaction networks (AP-matrix) in the same community. We found that plant-coding patterns vary from community to community (Fig. 3, Table 1). In the case of the leaf VOC community (PV\_z), VOCs are more commonly shared among plants ( $H(P|V) = 0.94$ ), whereas in



**Fig. 2.** Volatile organic compound (VOC) frequency vs rank (log-log scale) from four community-level study papers (top row) and a review paper (second to fourth rows) that summarized VOC frequency in different chemical groups (second row), four representatives of plant families (third row), and plants with different pollination groups and the overall review data (fourth row). Black lines in the cases of “Community: PV\_k,” “N-, S-, compounds,” “Orchidaceae,” “Overall” indicate that data can be described by power-law distribution ( $k$  values represent the exponent of the power-law distribution  $p(r) = ar^{-k}$ ), although log-normal distribution can describe the data similarly well (tests see Table S1). N-, S-, compounds: nitrogen or sulfur-containing volatile compounds. Number of sampled plant species ( $N_{\text{plant}}$ ) and number of VOCs ( $N_{\text{VOC}}$ ) in each case can be found in Table 1.

two cases of flower VOCs (PV\_f, PV\_k), there are more unique VOCs ( $H(P|V) = 0.41$ ,  $0.51$  respectively), and the third case of flower VOCs is in between ( $H(P|V) = 0.75$ ). We also analyzed plant coding patterns based on the review study and found these to vary from category to category (Table 1A), with the overall (from the whole dataset) value of

around 0.5 indicating some specialization ( $H(P|V) = 0.53$ ,  $H(V|P) = 0.41$ , Table 1A).

We observe specialized patterns for all these insect-plant interactions as revealed by entropy values much less than 1 ( $H(A|P)$  around 0.5,  $H(P|A)$  around 0.2, Table 1B, Fig. 4). The asymmetric values of  $H(A|P)$  and  $H(P|A)$  indicate that insects are

**Table 1** Entropy measures based on (A) plant-VOC associations (PV-matrix based) and (B) insect-plant networks (AP-matrix based) in the four community-level studies papers and a review paper that compiled plant-VOC publications**A** Plant-VOC associations (PV-matrix based)

Paper	Type	Category	Matrix	N <sub>Plant</sub>	N <sub>VOC</sub>	Power Law*	H(V)	H(P V)	H(V P)
Filella et al. 2013	Plant-pollinator	PV_f	PV	21	68	No	0.93	0.51	0.49
Kantsa et al. 2017, 2018	Plant-pollinator	PV_k	PV	41	366	Yes	0.90	0.41	0.48
Burkle & Bunyon 2019	Plant-pollinator	PV_b	PV	40	165	No	0.94	0.75	0.73
Zu et al. 2020	Plant-herbivore	PV_z	PV	47	56	No	0.99	0.94	0.92
Farré-Armengol et al. 2020	Review	Terpene	PV	270	296	No	0.80	0.58	0.33
Farré-Armengol et al. 2020	Review	Fatty acid derivative	PV	237	274	No	0.86	0.40	0.23
Farré-Armengol et al. 2020	Review	Benzenoid	PV	246	211	No	0.78	0.60	0.31
Farré-Armengol et al. 2020	Review	N-, S-, compounds	PV	105	43	Yes	0.79	0.53	0.15
Farré-Armengol et al. 2020	Review	Alzooceae	PV	27	36	No	0.93	0.83	0.73
Farré-Armengol et al. 2020	Review	Caryophyllaceae	PV	25	160	No	0.89	0.61	0.63
Farré-Armengol et al. 2020	Review	Orchidaceae	PV	22	192	Cut off	0.94	0.35	0.51
Farré-Armengol et al. 2020	Review	Apocynaceae	PV	17	150	No	0.92	0.55	0.65
Farré-Armengol et al. 2020	Review	insect	PV	222	753	Cut off	0.84	0.53	0.42
Farré-Armengol et al. 2020	Review	Bird & Bat	PV	25	157	No	0.93	0.35	0.47
Farré-Armengol et al. 2020	Review	Wind	PV	9	50	No	0.95	0.40	0.59
Farré-Armengol et al. 2020	Review	Overall	PV	296	857	Cut off	0.84	0.53	0.41

**B** Insect-plant networks (AP-matrix based)

Paper	Type	Category	Matrix	N <sub>Animal</sub>	N <sub>plant</sub>	H(P)	H(A P)	H(P A)
Kantsa et al. 2017, 2019	Plant-pollinator	AP_k	AP	168	38	0.86	0.56	0.15
Burkle & Bunyon 2019	Plant-pollinator	AP_b	AP	76	40	0.89	0.49	0.18
Zu et al. 2020	Plant-herbivore	AP_z	AP	28	20	0.94	0.42	0.19

$H(V)$  entropy of VOCs.  $H(\cdot|\cdot)$  conditional entropy. Calculation formulas see Fig. 1. \*Details of statistical tests for power-law, see Table S1. VOC, volatile organic compound.

more specialized on plants than plants specialized on insects ( $H(P|A)$  are lower). In addition, the interaction patterns seem to be indistinguishable between mutualistic (pollinator-plant) and antagonistic (herbivore-plant) networks based on these few studies. Indeed, a global analysis of more than 250 mutualistic and antagonistic networks has revealed that the network structures (e.g., generality, nestedness) *per se* cannot separate mutualistic from antagonistic networks (Michalska-Smith & Allesina, 2019) unless one includes environmental factors (Song & Saavedra, 2020).

### 3 From patterns to mechanisms

We have shown the information structures of plant–insect chemical communications as revealed by the apparent power-law behavior in some cases (but where these conclusions are limited by limited data availability), and by the ubiquitous decrease in entropy of VOC associations to plants as opposed to total entropy of VOCs. The latter is not surprising, as not all plants produce all VOCs. Interestingly, the conditional entropy of VOCs ( $H(P|V)$ ) involved in plant-herbivore interactions seems to be generally higher than the entropy of those involved in plant-pollinator interactions, but this is not reflected in the conditional entropy of plant-animal associations ( $H(A|P)$ )—which is similar among these interaction types. This suggests that plants try to communicate

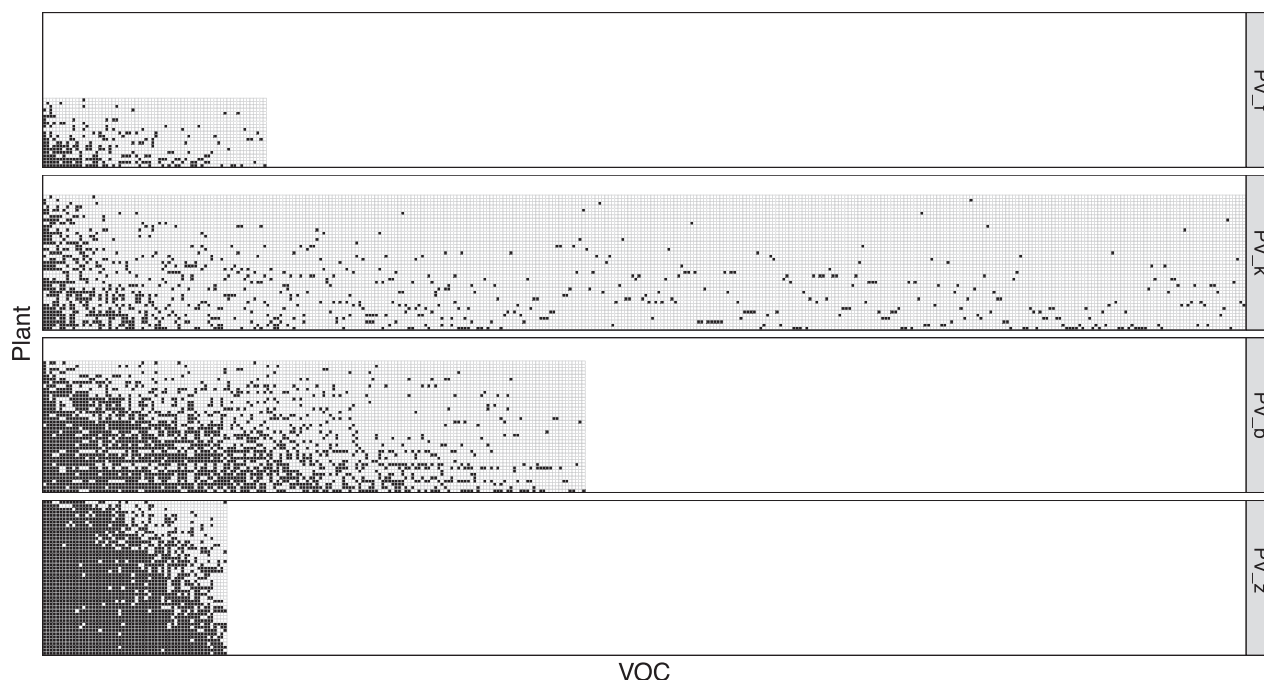
more clearly with their pollinators than with their herbivores, while herbivores must try harder than pollinators to decode information from their host plants.

While we recognize that further conclusions are likewise currently limited by a shortage of data, we can still ask why plant–insect chemical communication is structured in the way it is, and not otherwise. In this section, we highlight two approaches to investigate the potential mechanisms driving these chemical communication patterns between plants and insects. Recall the difficulties of building a complete decoding matrix (how each insect decodes each of the VOCs) empirically. Here, we theoretically construct insect decoding from two approaches: (i) using a top-down logic assuming that the insect can decode all the VOCs from a plant as long as they can feed on that plant; that is,  $AV = AP \times PV$  and (ii) use a bottom-up logic by hypothesizing the relationships between VOC abundance and functionality (i.e., effects on insects). In both directions, we integrate information theory into ecological and evolutionary theories to generate testable hypotheses.

#### 3.1 Top-down: Interactions shape communication

This direction follows the inspiration of studies exploring how Zipf's law patterns emerge in linguistics. Various hypotheses have been proposed to explain this intriguing pattern in human languages. Among these, the “least effort” hypothesis (Zipf, 1949; Ferrer-i-Cancho & Solé, 2003) is a compelling hypothesis that nicely recovers the Zipf's law





**Fig. 3.** Patterns of plant-volatile organic compound (VOC) associations (PV-matrix) in the four community studies (details see top four rows in Table 1A). Each row and column represents plant species and VOCs, respectively. Black and white squares show that a given VOC was present or absent in a given plant, respectively.

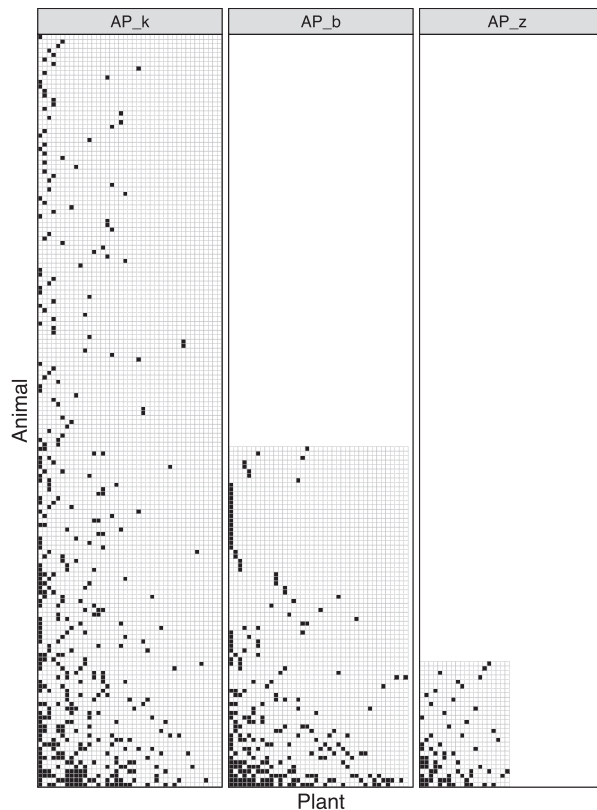
distribution as a natural consequence from the conflicting interests between speakers (who are thought to aim for “brevity and phonological reduction” to minimize their effort to speak) and listeners (who should desire “explicitness and clarity” to minimize their effort to understand). The successful application of information theory in studying the structure and emergence of human language has inspired studies on animal vocal communication (e.g., reviews in McCowan et al., 1999, 2008; Kershenbaum et al., 2021), and indicates its potential to be extended in studying the chemical “language” in plant–insect communities (Zu et al., 2020).

Indeed, Zu et al. (2020) aimed to borrow the framework to test whether plant–herbivore chemical communication patterns can be explained by conflicting interests of plants (speakers) and herbivores (listeners). It is well-known that the chemical arms race between plants and herbivores has been playing out for hundreds of millions of years, where herbivores must constantly adapt to plant defense chemicals while plants must keep producing novel chemicals (Ehrlich & Raven, 1964). Zu and colleagues (Zu et al., 2020) translated this chemical arms race into an information arms race: plants code the (chemical) information in a way (changes in PV-matrix) to make the decoding (AV given by  $AP \times PV$ ) difficult (high entropy) for herbivores, whereas herbivores interact with plants in a way (changes in AP-matrix) to make the decoding easier (low entropy). With repeated cycles of information optimization by plants and herbivores, they found that an equilibrium stage is reached in which a plant–VOC redundancy matrix and herbivore–plant specialization matrix have emerged. These information patterns at the equilibrium stage matched their field data collected in a tropical dry forest.

This is a successful case of integrating other known evolutionary and ecological theories into information theory to build a framework that aims to disentangle mechanisms driving plant–insect interactions and communication at the community level. In general, there are three steps to construct communication frameworks for a given system. First, we must define who are the speakers and who are the listeners. Second, we must define proper fitness functions for both parties based on their ecological and evolutionary relationships. Third, we must define the rules of optimization for each party in the information game. Both the second and the third steps require very careful examination by integrating evolutionary and ecological processes. We also want to emphasize that the “fitness” function at the community level, which defines success in terms of passing into the next “round” in a game, is not the same as the Darwinian fitness proxies that biologists normally use to assess individual success in survival and reproduction but rather refers to broader benefits, which can be assessed at the level of communities rather than individuals. This corresponds as well to the broader use of the term “signal” introduced at the beginning of this paper. This approach is compatible with the evaluation of individual-level fitness but aims at criteria that can be used at the community level. We hope more studies will be inspired by this information perspective to provide insights into other plant–insect chemical communication types (e.g., plant–pollinator, plant–herbivore–parasitoid).

### 3.2 Bottom-up: Connect VOC frequency with “information functionality”

A different, bottom-up approach would connect VOC frequency with information functionality to study



**Fig. 4.** Patterns of insect-plant interactions (AP-matrix) in the three community studies (details see Table 1B). Each row and column represents insect species and plant species, respectively. Black and white squares show that a given insect was, or was not observed feeding on a given plant, respectively.

plant–insect communication. Given that there are rare and common VOCs (frequency), it is natural to ask whether rare VOCs or common VOCs can be decoded more easily by insects (functionality) and how much information can be gained by decoding the VOCs.

Shannon's original mathematical theory of communication (Shannon, 1948) was mostly based on the “Inverse Relationship Principle” (D'Alfonso, 2011), which states that the less probable a signal is, the more information it bears. In our plant–insect communication systems, insects face an environment with VOCs at different frequencies. On the one hand, specializing in decoding rare VOCs brings advantages if substantial information is thereby gained to access a niche that can be exploited by specialists. However, specialization comes with risks because rare VOCs (emitted by only one plant species or a few phenotypes) can go extinct more easily, locally or globally. On the other hand, decoding common VOCs is likely to be safer but less informative, meaning that insects might need to decode more signals to gather enough information to identify their plant hosts. Which strategy do insects use and how will different strategies affect the interaction network structure? For example, will positive-frequency-dependent decoding structure: that is, preferentially

decoding common VOCs, lead to more generalized plant–insect interactions; whereas negative-frequency dependence, preferentially decoding rare VOCs, may result in more specialized interactions? How could this be understood in terms of the co-evolution of plant VOCs and insect olfaction?

Within this framework, one can hypothesize different specific relationships between VOC frequency and information functionality (VA-matrix), to test how these give rise to plant–insect interaction network structures (using  $PA = PV \times VA$ ). Specifically, information functionality can be modeled with two elements: one, how easily a VOC can be decoded (VA\_func); two, how much information can be gained by decoding the VOC (V\_info). Therefore, the information functionality matrix (VA) can be treated as an information-weighted decoding matrix, generated by  $(VA = V\_info \times VA\_func)$ . We can use a drift process mimicking frequency-independent decoding, which might act as a control for testing the strength of frequency-dependent decoding occurring in the empirical data.

## 4 Conclusion

We have found from the limited studies that plant VOC frequencies follow a heavy-tail distribution with a few predominant compounds and many rare compounds; all the three documented plant–insect interaction matrices are highly specialized, whereas plant–VOC coding patterns seem to vary from case to case, with different plant communities and perhaps, interaction types, resulting in signals that are encoded more or less redundantly. We also provide a top-down and a bottom-up direction for constructing theoretical frameworks to integrate information theory in studies of plant–insect chemical communication. By proposing both top-down and bottom-up frameworks, we hope we can draw attention to, and inspire more studies on the information perspective of trophic networks in ecosystems.

In the face of more serious and more frequent threats to ecological systems, plant–insect interactions are vulnerable, yet particularly important aspects of ecosystem functioning: insect pollinators are vital for agricultural productivity as well as for the survival of wild plant communities (Ollerton, 2021; Wei et al., 2021); herbivores support all higher trophic levels that depend on plants (Price et al., 1980; Harvey et al., 2003; Moreira et al., 2016), and their predators and parasitoids perform pest control services (Schmidt et al., 2003). We still struggle to control these interactions with relatively crude tools that carry heavy collateral damage, such as physically uprooting and transporting communities of pollinators to agricultural fields and broadly prescribed spraying of pesticides. Integrating information theory has the potential to help us gain a better understanding of the underlying mechanisms, and thus promises novel insights to better understand and manage these interactions. For example, novel chemicals of exotic plants have been suggested to be the main mechanism for invading the local communities (Cappuccino & Arnason, 2006; Macel et al., 2014). From the perspective of information theory, we can ask what are the effects of invasive species on the information landscape of the current community and whether some information

structures are more resilient than others. We can furthermore ask how climate change, pollution (including pesticide), and habitat fragmentation will affect communication signal profiles, dynamics, and signal-to-noise ratios and thus rewire plant–insect networks. Overall, understanding the effects of communication on species interactions and vice versa has the potential to improve our ability to assess impacts of anthropogenic disturbances on ecosystems, and assist conservation and restoration practices. For instance, instead of focusing on taxonomic diversity, chemical diversity and chemical information structure may be shown to be more crucial and informative for ecosystem biodiversity (Schuman et al., 2016).

Information approaches can provide the scalability to integrate further dimensions to achieve a better and potentially more unified understanding of information flows and their effects in ecosystems (Deng et al., 2021). For instance, we can expand these frameworks by including different sets of VOCs (e.g., from flowers, leaves, and roots); different sets of partners and trophic levels (e.g., pollinators, microbes, parasitoids, and predators); and even different information forms (e.g., visual, vocal). Information theory abstracts all kinds of signals (or functional traits) as information and thus has the potential to gather the multi-layer functional maps into a unified framework. In summary, integrating information theory to existing theories in ecology and evolution has the potential to unveil central biological mechanisms driving the formation and maintenance of the functionality of species interactions and thus of entire ecosystems (Margalef, 1968; Kessler & Kalske, 2018; O'Connor et al., 2019), as has recently been argued more generally for plant chemistry (Walker et al., 2022).

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## Conflicts of Interest

The authors declare no competing financial interest.

## Author Contributions

Pengjuan Zu initiated and performed the study. All authors contributed with ideas and wrote the manuscript.

## Data Availability Statement

The data and R code supporting the results can be found at Zenodo: <https://zenodo.org/record/6308472>

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## Supplementary Material

The following supplementary material is available online for this article at <http://onlinelibrary.wiley.com/doi/10.1111/jse.12841/supinfo>:

**Table S1.** Statistical results of power-law tests for the 16 cases in Fig. 2.