PNIE PROCEEDINGS OF THE NATIONAL INSTITUTE OF ECOLOGY OF THE REPUBLIC OF KOREA

Original Article

PNIE 2022;3(3):149-153
https://doi.org/10.22920/PNIE.2022.3.3.149
pISSN 2765-2203, eISSN 2765-2211

Emergence and Structure of Complex Mutualistic Networks

KyoungEun Lee[1], Nam Jung[1], Hyun Min Lee[1], Seung Eun Maeng[2], Jae Woo Lee[3]*

[1]Team of Vulnerable Ecological Research, Division of Climate and Ecology, Bureau of Conservation and Assessment Research, National Institute of Ecology, Seocheon, Korea
[2]Bizdata R&D Center, Chuncheon, Korea
[3]Department of Physics, Inha University, Incheon, Korea

ABSTRACT

The degree distribution of the plant-pollinator network was identified by analyzing the data in the ecosystem and reproduced by a model of the growing bipartite mutualistic networks. The degree distribution of pollinator shows power law or stretched exponential distribution, while plant usually shows stretched exponential distribution. In the growth model, the plant and the pollinator are selected with probability P_P and P_A=1–P_P, respectively. The number of incoming links for the plant and the pollinator is l_P and l_A, respectively. The probability that the link of the plant selects the pollinator of the existing network given as $A_{k_i}=k_i^{\lambda_A}/\sum_i k_i^{\lambda_A}$, and the probability that the pollinator selects the plant is $P_{k_i}=k_i^{\lambda_P}/\sum_i k_i^{\lambda_P}$. When the nonlinear growth index is λ_X=1 (X=A or P), the degree distribution follows a power law, and if $0\leq\lambda_X<1$, the degree distribution follows a stretched exponential distribution. The cumulative degree distributions of plants and pollinators of 14 empirical plant-pollinators included in Interaction Web Database were calculated. A set of parameters (P_A,P_P,l_A,l_P) that reproduces these cumulative degree distributions and a growth index λ_X (X=A or P) were obtained. We found that animal takes very heterogenous connections, whereas plant takes a more flexible connection network.

Keywords: Degree of distribution, Mutualistic network, Plant–pollinator network, Power law, Stretched exponential distribution

Introduction

Ecosystems exhibit complex relationships as numerous species interact with each other. In ecological networks, the species is represented by a node, and the relationships between the species are represented by links. The connecting links in the prey–predator food web represent the relationship between the prey and predator. In addition to the prey–predator network, there are various other network types, such as mutualistic networks and parasite–host networks (Bascompte, 2009; Bascompte *et al*., 2003; de Lima Filho *et al*., 2021; Dunne *et al*., 2002; Luz *et al*., 2021; McLeod & Leroux, 2021; Montoya *et al*., 2006; Strydom *et al*., 2021). In a mutualistic network, such as a seed disperser network or a plant–pollinator network, plants and animals are mutually supportive because they can profit from each other (Cohen, 2020; Hwang *et al*., 2008; Lee *et al*., 2012; Maeng & Lee, 2011; Olesen *et al*., 2007). However, in the parasite–host network, parasites

Received November 12, 2021; **Revised** January 14, 2022; **Accepted** January 17, 2022

***Corresponding author:** Jae Woo Lee
e-mail jaewlee@inha.ac.kr
https://orcid.org/0000-0003-0622-4649

benefit from the host. When a parasite parasitizes a host, the two species become linked.

Mutualistic relationships in ecosystems are classified into four types, namely seed dispersal, pollination, resource harvesting, and protection (Boucher, 1985). Plants that spread pollen via animals or insects use less energy to produce pollen, but they use considerable energy to produce colorful flowers and scents that attract animals and insects. Plants offer benefits to pollinators, including pollen, oil, resin, nectar, and fragrance; a common example is bees and flowers. Bees fly from one flower to another, gathering nectar. When they land on the flower, the bees obtain pollen as it rubs off onto their hairy bodies. When they land on the next flower, some of the pollen is rubbed off, thus pollinating the plant. In this mutualistic relationship, the bees consume the nectar, and the plants successfully reproduce.

In an ecological network, the degree of a node represents the number of connecting links. The degree of distribution for ecological networks characterizes the properties of ecological systems. When the degree of distribution function for the ecological network was investigated, various network types were identified. The cumulative degree of distribution function is defined as the integral of the degree distribution in the range of a specific lower degree to the maximum degree. The cumulative degree distribution functions identified in ecosystems have typical functions such as power law, exponential function, truncated scale-free function, stretched exponential function, uniform function, and irregular distribution (Lee *et al*., 2012; Maeng & Lee, 2011; Montoya *et al*., 2006). Table 1 summarizes the cumulative degree of distribution functions found in ecological networks.

Ecological networks have highly different characteristics compared to other complex networks. The total number of nodes in ecological networks is small because the ecosystems being examined by ecologists are relatively small (McLeod & Leroux, 2021; Montoya *et al*., 2006; Strydom *et al*., 2021). Given that ecosystems have a range of relationships between species, the characteristics of the connecting lines are distinct. Because species within an ecosystem strongly compete or cooperate with each other, the clustering coefficient is small compared with other networks. This clustering coefficient refers to the proportion of connections among the nearest neighbors of a node that are actually realized, compared with the number of all possible connections. A distinct hierarchy of predator relationships exists in the food chain. When the habitat environment in an ecosystem changes, the linkage or link strength between the species changes dynamically. Ecological networks are both robust and fragile. Some ecosystems maintain a high level of robustness against invasive species, whereas others are more vulnerable to rapid changes across the entire ecosystem (Maeng *et al*., 2019; Montoya *et al*., 2006).

In an ecological network, some species behave like wild bald eagles with many connections and are called generalists. However, some species are specialists with highly specific diets and a small number of prey species being consumed. Specialists have a fragile network structure because their survival cannot be guaranteed when connected species disappear, but they do have the advantage of being able to monopolize their prey. Research is currently underway to deepen our understanding of the expression principle of the structure of an ecological network. (Maeng *et al*., 2012; 2013; 2019). According to Maeng *et al*. (2019) in the case of a mutualistic network, the dependence of the cumulative degree of distribution function differs between plants and animals. The cumulative degree of distribution function of the plant–pollinator network has been recorded in coastal forests (Maeng & Lee, 2011).

Materials and Methods

Datasets on prey–pollinator networks

We examined datasets on plant–pollinator networks and suggested a model for growing them. We used plant–pollinator data from the Interaction Web Database (http://www.ecologia.ib.usp.br/iwdb). After analyzing the structure of the network, we proposed a model that reproduces the structure of the observed mutualistic networks. We summarized the 14 plant–pollinator networks analyzed in this study. The number of nodes and links in the ecological network was small and the network was sparse.

Nonlinear evolutionary model of a mutualistic network

The principle driving the generation of the degree of distribution function in a mutualistic network is not well understood. Here, we developed a growth network model to reproduce the degree of distribution function observed in plant–pollinator networks. The degree of distribution function follows a power law or a stretched exponential distribution. Therefore, we considered the linear or nonlinear preferential attachment in bipartite-growing mutu-

Table 1. Types of cumulative degree of distribution in the ecological networks

Function	Cumulative degree distribution
Power law	$P_c(k) \sim k^{-\gamma}$
Exponential function	$P_c(k) \sim \exp(-ak)$
Truncated power law	$P_c(k) \sim k^{-\gamma}\exp(-k/\gamma)$
Stretched exponential function	$P_c(k) \sim \exp(-ak^{\beta})$
Uniform function	$P_c(k) \sim constant$
No functional form	Irregularly distributed

alistic networks. Fig. 1 shows a model of a growing bipartite mutualistic network. The network growth begins with the initial core network. During each update, we selected a plant with the probability q_p and a pollinator with the probability q_A=1–q_p. The incoming node has links l_P for the plant and links l_A for the pollinator.

We controlled the connecting probability of the incoming links when a node such as a plant or a pollinator becomes attached to the existing network node. We then examined the linear and nonlinear preferential attachments for new incoming links. The linear preferential attachment introduced by Barabasi and Albert (1999), known as the Barbasi–Albert model, exhibits a scale-free degree distribution. The nonlinear preference attachment model can explain the stretched exponential function of the degree of distribution (Krapivsky & Redner, 2001; Maeng *et al*., 2019). We applied the general nonlinear preferential attachment of incoming links for plants and pollinators. The incoming links for the plant or pollinator were chosen as the target nodes according to the nonlinear preferential attachment with an attaching probability of $A_{k_i}=k_i^{\lambda_A}/\sum_i k_i^{\lambda_A}$ for the pollinators and $P_{k_i}=k_i^{\lambda_P}/\sum_i k_i^{\lambda_P}$ for the plants. We then repeated the attachment process to reach the target network size.

Results and Discussion

Solutions for a nonlinear evolutionary model of a mutualistic network

The growing mutualistic network follows the power law of the degree of distribution as $\lambda_{A\ or\ P}$=1. When the parameter is less than one (0<λ<1), the degree of distribution of the plant or pollinator shows a stretched exponential function expressed as $P(k)\sim\exp(-k^{1-\lambda_j})$, where j=A or P. We set a master equation for the mean number of plants and pollinators. Maeng *et al*. (2012) obtained the degree of distribution for the growing networks. For λ_X=1, the degree of distribution follows the power law:

$$P_X(k)\sim k^{-\gamma_X}$$

where the power-law exponent is obtained as $\gamma_X = 2+\frac{P_X l_X}{P_Y l_Y}$ and X=(A or P). For λ_X<1, the degree distribution shows a stretched exponential function such that

$$P_X(k)\sim\exp\left(-\frac{\mu_X\, k^{1-\lambda_X}}{1-\lambda_X}\right)$$

where $\mu_X = 1+\frac{P_X l_X}{P_Y l_Y}$.

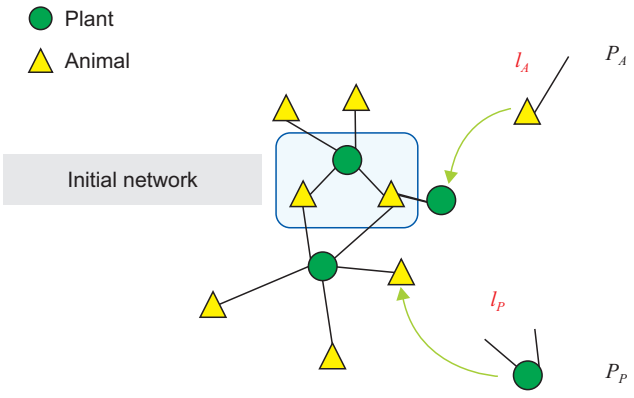

Fig. 1. Model of a growing bipartite network. The growth of the network starts from the initial core network. During each update, we selected a plant with a probability P_p and a pollinator with a probability P_A=1– P_p. The incoming node has links l_p for the plant and links l_A for the pollinator.

Structure of growing bipartite mutualistic networks

We generated the growing bipartite mutualistic networks from the initial core networks for a given set of parameter sets (P_A,P_P,l_A,l_P), as shown in Fig. 1. We simulated a mutualistic network using linear or nonlinear growth exponents for preferential attachment. Fig. 2 shows the cumulative degree of distribution of the simulated growing bipartite mutualistic network with nonlinear exponents for λ_A=1.0 for the plant and λ_P=0.5 for the animal and for the given set of parameter sets ($P_A=\frac{2}{3}, P_P=\frac{1}{3}, l_A=1, l_P=2$). The cumulative degree of distribution shows the power law for the animal with λ_A=1.0 and the stretched exponential distribution for the plant with λ_P=0.5. The simulated results were consistent with the analytical prediction.

Applications of the model for growing bipartite networks to real mutualistic networks

We applied the model of the growing bipartite mutualistic network to 14 empirical plant–pollinator networks. The incoming number of links l_P for plants is much larger than that of the animal l_A. There were found to be more plants than animals in the mutualistic networks, as shown in Table 2; the nonlinear growth exponents show significant asymmetry. In most of the networks in Table 2, we recorded the relation λ_A>λ_P except in the case of the Hocking network. This implies a significantly strong competition between a new animal and existing pollinators, in contrast with the relatively weak competition between plants. The restriction on the number of available plant species is a more important factor in shaping the mutualistic community than the restriction on the animal species available, which is likely related to the difference in their survival and reproduction rates. Plants with a large degree of distribution have the advantage of high abundance, and are screened by the competition between

animals characterized by $\lambda_P<1$, which leads the degree of distribution to take the stretched exponential form. This is not the case for "Hocking" in Fig. 3, where it has been reported in Hocking (1968) that competition between plants is more significant than that between the pollinators, implying $\lambda_A>\lambda_P$.

In summary, we determined the degree of distribution for the plant–pollinator network and introduced a growing model for bipartite mutualistic networks. We can reproduce the power law or stretched exponential degree of distribution for a mutualistic network for plants and animals. Thus, we report that competition among animals is stronger than that among plants.

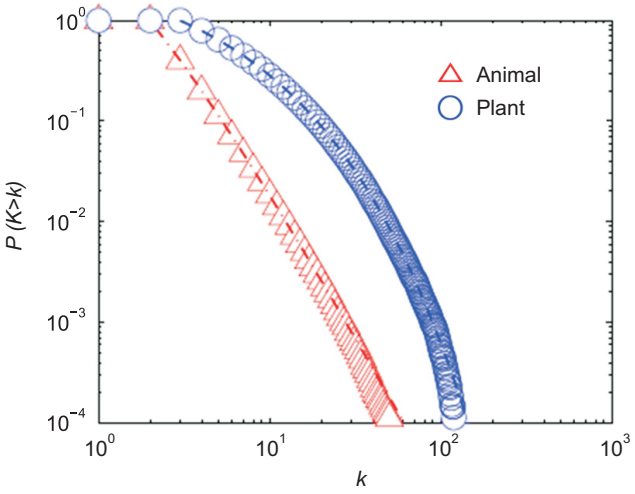

Fig. 2. Simulation of a growing bipartite mutualistic network using the growing exponents λ_A=1.0 for the plant and λ_P=0.5 for the animal and for the given set of parameter sets ($P_A=\frac{2}{3}, P_P=\frac{1}{3}, l_A=1, l_P=2$). We plotted the cumulative degree of distribution for the simulated results (symbols) and the analysis results (the dashed and dot-dashed line).

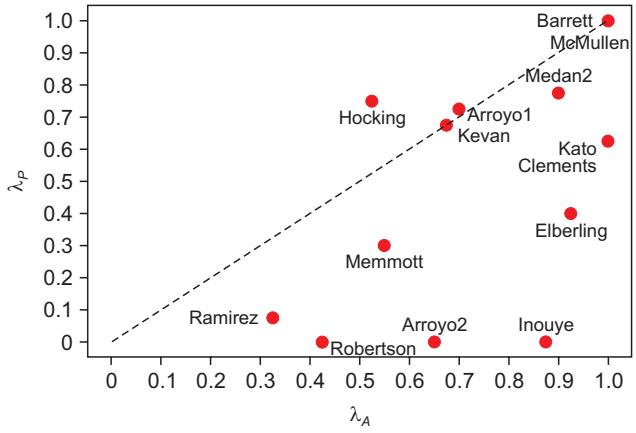

Fig. 3. Plot of the growth exponent λ_A for the plant versus λ_A for the empirical mutualistic networks. The nonlinear growing exponents (λ_A, λ_P) are obtained by fitting the cumulative degree of distribution of the empirical plant and pollinator, respectively.

Table 2. Fourteen plant–pollinator networks

Network	Animals	Plants	Links	Reference
Arroyo1	87	98	371	Arroyo *et al.* (1982)
Arroyo2	43	62	199	Arroyo *et al.* (1982)
Barret	12	102	167	Barrett and Helenurm (1987)
Clements	96	275	923	Clements and Long (1923)
Elberling	23	118	238	Elberling and Olesen (1999)
Hocking	29	86	184	Hocking (1968)
Inouye	42	91	281	Inouye and Pyke (1988)
Kato	93	679	1,206	Kato *et al.* (1990)
Kevan	32	115	312	Kevan (1970)
McMullen	106	54	204	McMullen (1993)
Medan	23	72	125	Medan *et al.* (2002)
Memmott	25	79	299	Memmott (1999)
Ramirez	33	53	109	Ramirez and Brito (1992)
Robertson	456	1,428	15,255	Robertson (1928)

The number of nodes and links depends on the mutualistic networks.

Conflict of Interest

The authors declare that they have no competing interests.

Acknowledgments

This study was supported by a National Research Foundation of Korea (NRF) grant funded by the Korean government (Grant No. NRF-2020R1A2C1005334).

References

Arroyo, M.T.K., Primack, R., and Armesto, J. (1982). Community studies in pollination ecology in the high temperate Andes of Central Chile. I. pollination mechanisms and altitudinal variation. *American Journal of Botany,* 69, 82-97.

Barabasi, A.L., and Albert, R. (1999). Emergence of scaling in random networks. *Science (New York, N.Y.),* 286, 509-512.

Barrett, S.C.H., and Helenurm, K. (1987). The reproductive biology of boreal forest herbs. I. Breeding systems and pollination. *Canadian Journal of Botany*, 65, 2036-2046.

Bascompte, J. (2009). Mutualistic networks. *Frontiers in Ecology and the Environment,* 7, 429-436.

Bascompte, J., Jordano, P., Melián, C.J., and Olesen, J.M. (2003). The nested assembly of plant-animal mutualistic networks. *Proceedings of the National Academy of Sciences of the United States of America,* 100, 9383-9387.

Boucher, D.H. (1985). *The Biology of Mutualism: Ecology and Evolution.* London: Oxford University Press.

Clements, F.E., and Long, F.L. (1923). *Experimental Pollination an Outline of the Ecology of Flowers and Insects.* Washington, D.C.: Carnegie Institution of Washington.

Cohen, J.E. (2020). Species-abundance distributions and Taylor's power law of fluctuation scaling. *Theoretical Ecology*, 13, 607-614.

de Lima Filho, J.A., Vieira, R.J.A.G., de Souza, C.A.M., Ferreira, F.F., and de Oliveira, V.M. (2021). Effects of habitat fragmentation on biodiversity patterns of ecosystems with resource competition. *Physica A Statistical Mechanics and its Applications,* 564, 125497.

Dunne, J.A., Williams, R.J., and Martinez, N.D. (2002). Food-web structure and network theory: the role of connectance and size. *Proceedings of the National Academy of Sciences of the United States of America,* 99, 12917-12922.

Elberling, H., and Olesen, J.M. (1999). The structure of a high latitude plant-flower visitor system: the dominance of flies. *Ecography,* 22, 314-323.

Hocking, B. (1968). Insect-flower associations in the high arctic with special reference to nectar. *Oikos,* 19, 359-387.

Hwang, J.K., Lee, K.E., Maeng, S.E., and Lee, J.W. (2008). Scaling behaviors of plant-pollinator mutualistic networks. *Journal of the Korean Physical Society,* 53, 3151-3155.

Inouye, D.W., and Pyke, G.H. (1988). Pollination biology in the Snowy Mountains of Australia: comparisons with montane Colorado, USA. *Australian Journal of Ecology,* 13, 191-205.

Kato, M., Kakutani, T., Inoue, T., and Itino, T. (1990). Insect-flower relationship in the primary beech forest of Ashu, Kyoto: an overview of the flowering phenology and the seasonal pattern of insect visits. *Contributions from the Biological Laboratory, Kyoto University,* 27, 309-376.

Kevan, P.G. (1970). *High arctic insect-flower visitor relations: the inter-relationships of arthropods and flowers at Lake Hazen, Ellesmere Island, Northwest Territories, Canada*. (Doctoral dissertation). University of Alberta, Edmonton.

Krapivsky, P.L., and Redner, S. (2001). Organization of growing random networks. *Physical Review E Statistical Nonlinear and Soft Matter Physics,* 63(6 Pt 2), 066123.

Lee, D.S., Maeng, S.E., and Lee, J.W. (2012). Scaling of nestedness in complex networks. *Journal of the Korean Physical Society*, 60, 648-656.

Luz, F.A., Goetz, A.P.M., and Mendonça, M.d.S. (2021). What drives gallers and parasitoids interacting on a host plant? A network approach revealing morphological coupling as the main factor. *Ecological Entomology,* 46, 334-341.

Maeng, S.E., and Lee, J.W. (2011). Asymmetric network properties of bipartite ecological networks. *Journal of the Korean Physical Society,* 58, 851-854.

Maeng, S.E., Lee, J.W., and Lee, D.S. (2012). Interspecific competition underlying mutualistic networks. *Physical Review Letters,* 108, 108701.

Maeng, S.E., Lee, J.W., and Lee, D.S. (2013). Impact of compatibility on the organization of mutualistic networks. *Physical Review E Statistical Nonlinear and Soft Matter Physics,* 88, 022804.

Maeng, S.E., Lee, J.W., and Lee, D.S. (2019). Competition-induced increase of species abundance in mutualistic networks. *Journal of Statistical Mechanics*, 2019, 033502.

McLeod, A.M., and Leroux, S.J. (2021). Incongruent drivers of network, species and interaction persistence in food webs. *Oikos,* 130, 1726-1738.

McMullen, C.K. (1993). Flower-visiting insects of the Galapagos Islands. *Pan-Pacific Entomologist,* 69, 95-106.

Medan, D., Montaldo, N.H., Devoto, M., Maniese, A., Vasellati, V., Roitman, G.G., *et al*. (2002). Plant-pollinator relationships at two altitudes in the Andes of Mendoza, Argentina. *Arctic Antarctic and Alpine Research,* 34, 233-241.

Memmott, J. (1999). The structure of a plant-pollinator food web. *Ecology Letters,* 2, 276-280.

Montoya, J.M., Pimm, S.L., and Solé, R.V. (2006). Ecological networks and their fragility. *Nature,* 442, 259-264.

Olesen, J.M., Bascompte, J., Dupont, Y.L., and Jordano, P. (2007). The modularity of pollination networks. *Proceedings of the National Academy of Sciences of the United States of America,* 104, 19891-19896.

Ramirez, N., and Brito, Y. (1992). Pollination biology in a palm swamp community in the Venezuelan Central Plains. *Botanical Journal of the Linnean Society,* 110, 277-302.

Robertson, C. (1928). *Flowers and Insects: Lists of Visitors of Four Hundred and Fifty-Three Flowers.* Carlinville: Science Press Printing Company.

Strydom, T., Catchen, M.D., Banville, F., Caron, D., Dansereau, G., Desjardins-Proulx, P., *et al*. (2021). A roadmap towards predicting species interaction networks (across space and time). *Philosophical Transactions of the Royal Society of London Series B Biological Sciences,* 376, 20210063.