Neighbor-Stranger Discrimination of Yellow-throated Buntings (*Emberiza elegans*) and Gray-headed Buntings (*Emberiza fucata*) to Playback of Song

Bo-Yeon Hwang and Shi-Ryong Park*

Department of Biology Education, Korea National University of Education, Cheongwon, Chungbuk 363-791, Korea

Songs of the Yellow-throated Bunting (Emberiza elegans) and the Gray-headed Bunting (Emberiza fucata) in allopatric populations in Gangnae-meon, Cheongwongun, Chungbuk in Korea, were recorded during the breeding season and analyzed in sound spectroraphs. Males of E. elegans and E. fucata were tested to investigate whether territorial males can discriminate between neighbor and stranger based on playback of natural and artificial song repertoires. In addition, E. fucata was stimulated by playback of only the individually specific section as well as of only the posterior portion of the song. Males of E. elegans were able to discriminate individually between neighbor and stranger in response to natural song repertoires, but they did not respond to playback of the artificial song repertoires of neighbor and stranger. Males of E. fucata were able to discriminate individually between neighbor and stranger in response to natural, artificial song repertoires, and the anterior section of the song, while males did not respond to playback of the posterior section of the song.

KEY WORDS: Emberiza elegans, Emberiza fucata, Neighbor-Stranger Discrimination, Natural Song Repertoires, Artificial Song Repertoires.

A bird that can discriminate between its neighbors' vocalizations can save time and energy by gauging its territorial response according to the threat posed by a particular individual neighbor. Birds should gain from learning to recognize their neighbors by voice and learn not to respond to conspecifics who are established for the whole breeding season and pose no immediate threat. Strange males should, on the other hand, be recognized as threats and responded accordingly. Differential responses of established birds are adaptive in reducing energy expenditure to stimuli with little significance to them (Falls, 1982). Recognition of individual birds by vocalization

that one aspect of song complexity, song repertoires, may adversely affect neighbor recognition (Wiley and Wiley, 1977, Krebs and Kroodsma 1980, Falls 1982). Field experiments have confirmed that males of *E. elegans* could discriminate individually between neighbor and stranger (Sung and Park, 1993).

requires adequate identifying information and an

ability on the part of receivers to detect and learn

individual differences (Falls, 1982, Elfström,

Emberiza fucata has a single song type. Each

^{1990).}One possible function of song complexity is to render the singer more readily identifiable to neighboring conspecifics (Beecher and Stoddard 1990), however several authors have suggested

^{*}To whom correspondence should be addressed.

males has a distinctive section at the anterior part of the song (Kim and Park, 1993). Therefore Kim and Park (1993) predicted that this part was used as an important discriminative factor for individual recognition.

In this study, we conducted four playback experiments. The first, we directly examined if the males of E. elegans could discriminate individually between natural song repertoires of neighbor and stranger. The second, we examined if they are able to discriminate individually between artificial song repertoires (loop playback of one song) of neighbor and stranger. The third, we investigated if males of E. fucata could discriminate individually between natural song repertoires of each neighbor and stranger. The fourth, we also tested if these males are able to discriminate individually between artificial song repertoires of neighbor and stranger, and if so, whether they were able to discriminate individually between neighbor and stranger using only the individually specific section (fixed part of the song of each male) of the song or only the plastic part of the song of each male.

Materials and Methods

Field work and habit

This study was performed from March to June in 1993 and 1994. Songs of males Yellow-throated Bunting (*E. elegans*) (n=5) and Grayheaded Bunting (*E. fucata*) (n=4) were recorded in Gangnae-meon, Cheongwon-gun, Chungbuk in Korea. Before the playback experiment, each individual's territory and its boundary were determined by observations.

Equipment

Kowa telescope (8×42) and BUSHNELL spacemaster II telescope (70mm) were used to observe the bird's behavior. All recordings were made using Uher 4000 tape recorder (tape speed of 19 cm/s) with an AKG C1000S microphone mounted in a 50 cm diameter parabolic reflector. For the playback experiments, we used the Uher reel-to-reel tape recorder and a Sony TCM-5000EV cassette tape recorder with a remote (20m) JBL proIII loudspeaker. We made the

sonagrams on a Kay Electric Company DSP Sonagraph Model 5500 and computer interface data processing.

Stimuli tape preparation

We made every attempt to place the microphone close to the bird, ie. less than 10 m to minimize attenuation and degradation of the sound quality. Our playback tapes were of three kinds: 1) the natural song sequence of neighbor and stranger (both E. elegans and E. fucata), each cassette tape being 3 mins long; 2) the artificial song sequence of neighbor and stranger (both E. elegans and E. fucata), each stimulus songs was presented for 3mins (loop playback of one song). One song type of the natural song repertoires was chosen and digitized on a computer disk using computer interface. This information was then sent to a DSP Sona-graph. This stimulus song was rescaled to approximate the song intervals of each male, by repeating one song of each male continuously; 3) anterior and posterior portions of the song of neighbor and stranger (E. fucata only) with each cassette tape of 3 mins.

Playback experiments: design and procedure

All trials were conducted in semiopen coniferous and mixed forests. Each territorial male (*E. elegans* and *E. fucata*) was exposed to playback of songs of conspecific neighbors and strangers. For playback a loudspeaker was placed in a tree 1.5 to 2 m above ground. The loudspeaker was oriented toward the territorial center of each test bird. The experimenter monitored the playback experiments from about 20 m away. The volume of playback was adjusted to 90dB at 1 m from the speaker using a precision integrating sound level meter.

Different trials with the same bird were at least 2 days apart and no adjacent area was visited every other day. The order in which birds were tested each day and the order in which test songs and speaker locations were used with each bird on different days were randomized for disturbing song habituation. Because the birds sang most actively near sunrise and sunset, playback experiments were generally conducted within 3 hours (06:30-

09:30) after sunrise and within 2 hours (16:30-18:30) before sunset with light to moderate winds. If the weather deteriorated or if no response occurred, the experiment was abandoned but repeated on another day. When subject individuals for playback experiment were chosen, one male of *E. fucata* was excluded because it occurred in a territory too far from others.

Measures of response

The measures of response were of three kinds: 1) Latency time from the start of playback to the first response, either by vocalizations or by approach within 15 m radius circle of the loudspeaker. 2) The closest distance of a bird to the loudspeaker, estimated in centimeters. 3) Staying time measured as the time that a bird spent within 15 m of the loudspeaker.

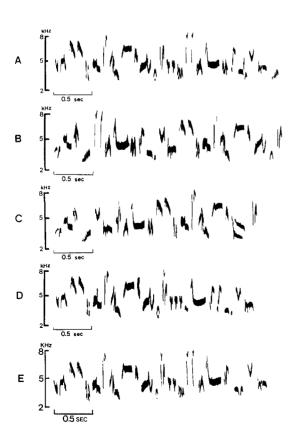


Fig. 1. Serial song sequences of a male Yellow-throated Bunting (*Emberiza elegans*). The order of alphabet indicates the order of continual song.

Sonagram interpretation and data treatment

All comparisons were made between responses to each stimulus song presentations (neighbor and stranger song, natural and artificial song, anterior and posterior song and so on). Quantitative statistics of song of each species were used for ONE-WAY ANONA test and mean test. Wilcoxon matched pairs signed rank test and Mann-Whitney U test (Siegel 1956) were used to determine

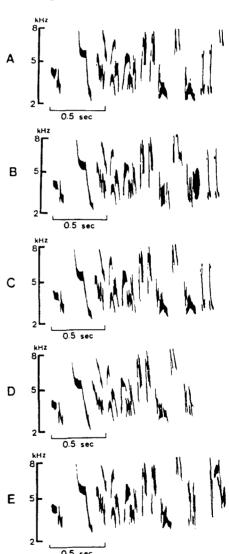


Fig. 2. Serial song sequences of a male Gray-headed Bunting (*Emberiza fucata*). The order of alphabet indicates the order of continual song.

Table 1. Comparison of the response to playback of neighbor's and stranger's natural and artificial song repertoires of Yellow-throated Bunting (*Emberiza elegans*).

	Natural song repertoires of Neighbor			Natural song repertoires of Stranger			Artificial song repertoires of Neighbor	Artificial song repertoires of Stranger
	latency time (sec)	closest distance to speaker (cm)	xtaying time within 15 m radius circle (sec)	latency time (sec)	closest distance to speaker (cm)	staying time within 15 m radius circle (sec)		
Α	83	1330	205	54	810	391	no response	no response
В	57	1730	159	37	515	270		
С	115	910	298	96	780	453		
D	34	1310	132	18	300	737		
Е	89	1160	41	78	515	127		

All test birds could discriminate individually between neighbor and stranger to playback of natural song repertoires. On the other hand, they didn't respond to playback of artificial song repertoires of neighbor and stranger.

Table 2. Comparison of mean values of the three response measurements from playback experiments with male Grey-headed Bunting (*Emberiza fucata*).

	Natural song repertoires							
	latency ti	me (sec)	closest distance	to speaker(cm)	staying time within 15 m radius circle (sec)			
	neighbor	stranger	neighbor	stranger	neighbor	stranger		
Α	80.3±15.5*	49.6±10.4	1206.6±261.0	340.0±206.6	500.6±245.7	683.6±361.4		
В	83.6±38.4	49.3±26.5	931.7±307.5	118.6±64.4	496.3±30.6	787.0±247.3		
C	114.0±62.5	30.3 ± 23.3	856.6±332.7	360.0±98.4	252.3±56.3	599.6±241.0		
D	103.6±27.2	34.0±11.7	1211.6±443.9	276.6±136.9	208.3±91.4	261.3±145.7		
	ANOVA test; P<0.05		P<0	0.01	P<0.01			
	Artificial song repertoires							
	latency time (sec)		closest distance	to speaker (cm)	staying time within 15 m radius circle (sec)			
					10 III Idala	circle (sec)		
	neighbor	stranger	neighbor	stranger	neighbor	stranger		
A	neighbor 95.3±64.0	stranger 75.0±31.3	neighbor 1180.0±407.0	stranger 270.0±193.1				
A B					neighbor	stranger		
	95.3±64.0	75.0±31.3	1180.0±407.0	270.0±193.1	neighbor 211.0±122.7	stranger 280.6±35.5		
В	95.3±64.0 66.7±12.6	75.0±31.3 44.6±8.9	1180.0±407.0 1016.6±253.8	270.0±193.1 240.0±195.1	neighbor 211.0±122.7 212.6±40.1	stranger 280.6±35.5 338.0±45.9		

^{*;} Standard deviation (S.D.)

All test birds could individually discriminate between neighbor and stranger to playback of natual and artificial song repertoires of *Emberiza fucata*.

relative differences in response among the treatments when each measure of response was treated separately.

Results

Males of Yellow-throated Buntings sing 15-65 songs per bout and a bout usually consists of 3-8 song types. Each song types was repeated 2-20 times before switching to a different song type.

Table 3. Comparison with P values of Mann-Whitney U test between the response of natural and artificial song repertoires in Gray-headed Bunting (*Emberiza fucata*).

measured factors	stimulus song	neighbor	stranger	
latency time	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	0.1060 n.s.	0.8852 n.s.	
closest distance to speaker		0.4354 n.s.	0.7948 n.s.	
staying time within 15	m radius circle	0.0004*	0.0001*	

Note; n.s. (not significant), *; significant P<0.001

The individual response of natural and artificial song repertoires was not significantly different with latency time and closest distance to speaker in measured factors, but the difference of staying time within 15 m radius circle was significantly different in context to each case.

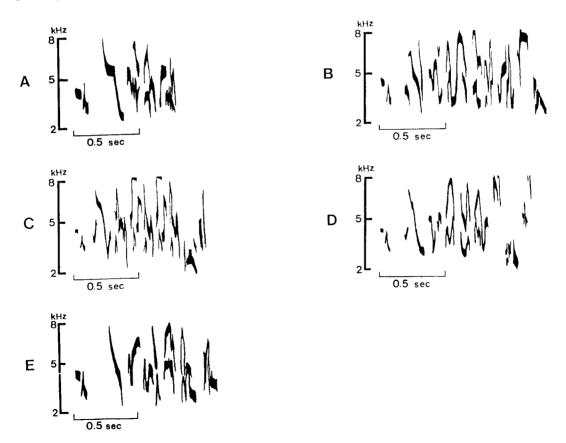


Fig. 3. Individual specific sections (anterior section) of song in the five males' Gray-headed Bunting (Emberiza fucata).

Sonagrams of serial song sequence in *E. elegans* are shown in Fig. 1. Males of Gray-headed Buntings sing 30-80 songs per bout and a bout usually consists of one song type. Sonagrams of serial song sequences in *E. fucata* are shown in Fig. 2.

Experiment 1.

Neighbor-stranger discrimination to playback of natural and artificial song repertoires in Yellow-throated Bunting and Grayheaded Bunting.

Emberiza elegans exhibited significantly different responses to the natural songs of neighbors and strangers for each measurement (Wilcoxon matched pairs signed rank test (n=5); latency time P<0.0431, closest distance P<0.0431, staying time within 15 m P<0.0431). Emberiza elegans was also exposed artificial song

repertoires of neighbor and stranger. But none of the subjects responded to the stimulus songs (Table 1).

Emberiza fucata exhibited significantly different responses to the natural songs of neighbors and strangers for each measurement (Wilcoxon matched pairs signed rank test (n=4); latency time P<0.0022, closest distance P<0.0022, staying time within 15 m P<0.0022). Emberiza fucata was also exposed artificial song repertoires of neighbor and stranger. Emberiza fucata discriminated individually between neighbor and stranger for each measurement (Wilcoxon matched pairs signed rank test (n=4); latency time P<0.0022, closest distance P<0.0022, staying time within 15 m P<0.0022) (Table 2).

Comparing results with natural song repertoires and those with artificial song repertoires, the staying time within 15 m radius circle was

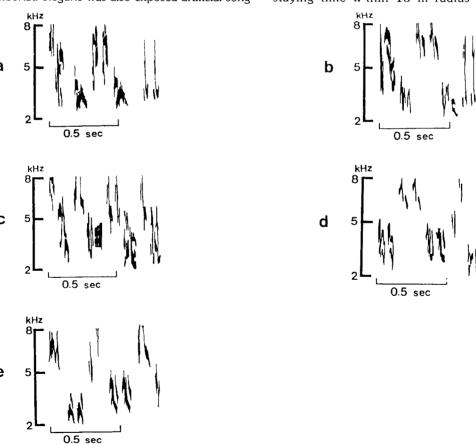


Fig. 4. Plastic sections (posterior section) of song in the five males' Gray-headed Bunting (Emberiza fucata).

significantly different (Table 3).

Experiment 2.

Neighbor-stranger discrimination to playback of anterior and posterior sections of the song in Gray-headed Bunting.

The individually specific sections of the song of five males of *E. fucata* are shown in Fig. 3 and the other sections of the song in Fig. 4.

When *E. fucata* males were exposed to the anterior section of the songs of neighbors and strangers, they exhibited significantly different responses of neighbors and strangers for each measurement (Wilcoxon matched pairs signed rank test (n=4); latency time P<0.0117, closest distance P<0.0117, staying time within 15 m P<0.0117). *Emberiza fucata* was also exposed to the posterior section of the songs of neighbors and strangers. But none of the subjects responded to the stimulus songs (Table 4).

Discussion

The results of our experiments are consistent with the hypothesis that territorial male Yellow-throated Buntings and Gray-headed Buntings

discriminate between songs of neighbor and stranger. As in the studies of neighbor-stranger discrimination in other species (Weeden and Falls, 1959; Emlen, 1971; Krebs, 1971; Goldman, 1973; Brooks and Falls, 1975; Kroodsma, 1976; Harris and Lemon, 1976; Wiley and Wiley 1977, Wunderle 1978, Searcy et al., 1981, Slater, 1981; Sung and Park, 1993), playback of stranger songs elicited stronger responses than did playback of songs of neighbors.

The most direct way to demonstrate individual recognition by sound is to compare responses of birds to recordings of different individuals (Falls, 1982). We showed that E. elegans and E. fucata could in fact discriminate neighbor from strangers on the basis of song. In this study, these two species were clearly different in their ability to make individual discrimination. Emberiza elegans could discriminate individually between the natural song repertoires of neighbor and stranger. Although the song repertoires of this species ranged from three to eight discrete song types, when we executed to loop playback of a randomly chosen song from a male, none of the males responded to the stimulus song. We thought that E. elegans did not recognize the individual's information with just one song type. These

Table 4. Comparison between the response of neighbor-stranger discrimination to playback of only the anterior and posterior section of the song in Gray-headed Bunting (*Emberiza fucata*).

	Anterior section of neighbor's song			Anterior section of stranger's song			Posterior section of neighbor's song	Posterior section of stranger's song
	latency time (sec)	closest distance to speaker (cm)	staying time within 15 m radius circle (sec)	latency time (sec)	closest distance to speaker (cm)	staying time within 15 m radius circle (sec)		
Α	96.0±	1155.0±	308.0±	45.0±	377.5±	312.5±	no response	no response
	12.72*	205.06	90.51	22.63	392.44	142.13		
В	60.5±	912.5±	233.5±	45.0±	422.5±	486.5±		
	4.95	144.96	30.41	5.66	152.03	16.26		
С	87.0±	1020.0±	454.5±	29.5±	84.0±	542.5±		
	56.67	438.41	381.13	30.41	65.05	324.56		
D	103.5±	1187.5±	288.0±	16.5±	310.0±	414.0±		
	53.03	371.23	121.62	6.36	127.28	106.07		

^{*;} Standard deviation (S.D.) Note: testing was used to ANOVA test.

findings indicate that males of *E. elegans* must attain familiarity with most or all of the song types in their neighbors' repertoires. Thus, *E. elegans* need to hear multiple song types in order to make this discrimination, as some have suggested (e.g., Falls 1982, Weary *et al.*, 1987).

Emberiza fucata could discriminate between neighbors and strangers to playback of natural and artificial song repertoires. In contrast with *E. elegans*, these results might be due to continuously repeated song sequence in *E. fucata*. In fact, natural and artificial song repertoires of *E. fucata* are different in only the posterior section of the continuous song sequence. Therefore, although they listen to the only one song, males may be able to acquire needed information for individual recognition.

Searching for discriminant elements within a single song type for neighbor-stranger discrimination, we also performed a playback experiment using the individually specific section and the other section, individually common section. *Emberiza fucata* could discriminate between neighbor and stranger to playback of only the anterior section of the song, while none of the males responded to only the posterior section of the song. This result proposes that anterior section of the song play, an important role for individual recognition.

We think that our experiment of neighbor recognition in *E. fucata* supports the 'releasing' hypothesis in the context of four hypotheses of how species-specific information is conveyed in birds. The 'releasing' hypothesis states that only certain acoustic components of the song are used for species identification (Date *et al.*, 1991).

Conclusionally, our experiments propose that the ability of song recognition between two species, *E. fucata* and *E. elegans* is different, and such a difference might be due to song repertoire size and song variation in a bout.

Acknowledgements

We thank our research assistants for help in the field. We also thank Dr. J. Bruce Falls in the Department of Zoology, University of Toronto and

Dr. R. E. Lemon in the Department of Biology, McGill University, Canada, for commenting on earlier drafts of the manuscript.

References

- Beecher, M.D. and P.K. Stoddard, 1990. The Role of Bird Song and Calls in Individual Recognition: Contrasting Field and Laboratory Perspectives, In: Comparative Perception (Stebbins, W.C. and M.A. Berkeley, eds.). John Wiley and Sons, New York, Vol. 2, pp. 375-408.
- Brooks, R.J. and J.B. Falls, 1975. Individual recognition by song in white-throated sparrows. I. Discrimination of songs of neighbors and strangers. *Can. J. Zool.* **53:** 879-888.
- Date, E.M., R.E. Lemon, D.M. Weary, and A.K. Richter, 1991. Species identify by birdsong: discrete or additive information? *Anim. Behav.* 41: 111-120.
- Emlen, S.T., 1971. The role of song in individual recognition in the Indigo Bunting. Z. Tierpsychol. 28: 241-246.
- Elfström, S.T., 1990. Individual and species-specific song patterns of rock and meadow pipits: physical characteristics and experiments. *Bioacoustics* **2:** 277-301.
- Falls, J.B., 1982. Individual Recognition by Sound in Birds, In: Acoustic Communication in Birds. (Kroodsma, D.E. and E.H. Miller, eds.). Academic Press, New York, Vol. 2, pp. 237-278.
- Goldman, P., 1973. Song recognition by Field Sparrow. Auk 90: 106-117.
- Harris, M.A. and R.E. Lemon, 1976. Response of male Song Sparrows *Melospiza melodia* to neighbouring and non-neighbouring individuals. *Ibis* 118: 421-424.
- Kim, K.W. and S.R. Park, 1993. Intraindividual and interindividual variation of stereotyped songs in Grayheaded Bunting (Emberiza fucata). Korean J. Zool. 36: 476-486.
- Krebs, J.R., 1971. Territory and breeding density in the Great tit, *Parus major L. Ecology* **52**: 2-22.
- Krebs J.R. and D.E. Kroodsma, 1980. Repertoires and territory defense. *Nature* 271: 539-542.
- Kroodsma, D.E., 1976. The effect of large song repertoires on neighbor 'recognition' in male Song Sparrows. Condor 78: 97-99.
- Searcy, W.A., P.D. McArthur, S.S. Peters, and P. Marler, 1981. Response of male Song and Swamp Sparrows to neighbour, stranger and self songs. *Behaviour* 77: 152-166.
- Siegel, S., 1956. Nonparametric Statistics for the

Behavioral Sciences, McGraw-Hill, New York.

Slater, P.J.B., 1981. Chaffinch song repertoires: Observations, experiments and a discussion of their significance. Z. Tierpsychol. 56: 1-24.

Sung, H.C. and S.R. Park, 1993. Neighbor recognition by song in the Yellow-throated Bunting (*Emberiza elegans*). Korean J. Behav. Biol. 2: 45-54.

Weary, D.M., R.E. Lemon, and E.M. Date, 1987.
Neighbour-stranger discrimination by song in the veery, a species with song repertoires. Can. J. Zool.
65: 1206-1209.

Weeden, J.S. and J.B. Falls, 1959. Differential

responses of male ovenbirds to recorded songs of neighboring and more distant individuals. *Auk* **76**: 343-351.

Wiley R.H. and M.S. Wiley, 1977. Recognition of neighbors duets by stripe-backed wrens (Campylorhynchus nuchalis). Behaviour **62:** 10-34.

Wunderle, J.M., 1978. Differential response of territorial Yellow-throats to the songs of neighbors and non-neighbors. Auk **95:** 389-395

(Accepted November 30, 1995)

노랑턱멧새(Emberiza elegans)와 붉은뺨멧새(Emberiza fucata)에서 Song의 Playback을 통한 이웃-낯선 개체의 인식 황보연·박시룡(한국교원대학교 생물교육과)

번식시기동안 상호 이소종인 노랑턱멧새와 붉은뺨멧새의 song을 충청북도 청원군 강내면에서 녹음하여 분석하였다. 두종의 수컷들을 대상으로 playback실험을 실시하였다. 노랑턱멧새와 붉은뺨멧새가 각각 자연적 그리고 인공적인 song repertoires로 playback을 하였을 때 이웃-낯선개체를 인식하는지의 여부를 조사하였다. 또한 붉은뺨멧새에 대해 전반부의 song과 후반부의 song을 playback하여 이웃-낯선개체의 인식 여부도 조사하였다. 노랑턱멧새는 자연적인 song repertoires로 이웃-낯선개체를 인식하였으나, 인공적인 song에 대해서는 반응하지 않았다. 붉은뺨멧새는 자연적. 인공적, 그리고 song의 전반부를 playback하였을 때 이웃-낯선개체를 구별하였다. 반면에 song의 후반부를 playback하였을 때는 반응하지 않았다.