The Distributions and Boundary of Two Distinct, Local Forms of Japanese Pond Frog, *Pelophylax porosus brevipodus*, Inferred From Sequences of Mitochondrial DNA

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The Nagoya Daruma pond frog *Pelophylax porosus brevipodus* is distributed in western Japan and is traditionally divided into two local forms: the Okayama form in the west and the Nagoya form in the east. These two forms are genetically differentiated, but have never been defined taxonomically because their distributions are unclear to date. To complete the distributions and identify the boundary of the two forms, we genetically investigated 16 populations including eight populations located within the unexamined area. We found that the distributional boundary is located within a small area of Hyogo Prefecture where haplotypes of mitochondrial cytochrome b (*cytb*) and D-loop region corresponding to the two forms co-existed. On the other hand, the polymorphic site of the nuclear gene SOX3 revealed introgression over the boundary into Okayama *cytb* clade. These results suggest that the two forms were geographically isolated from each other in the past, and secondarily contacted and then accepted one-way introgression. As a next step of the research, taxonomic approach is expected to define the two forms.

Keywords: Japanese pond frog, cytochrome b, D-loop, SOX3, two major forms

INTRODUCTION

Two pond frog species live in the Japanese islands, *Pelophylax nigromaculatus* and *Pelophylax porosus*. The latter species is endemic to Japan and is called the Daruma pond frog. It is similar to a traditional Japanese Daruma doll with its round shape. This species is comprised of two subspecies: *P. p. porosus* (Tokyo Daruma pond frog), which is distributed in eastern Japan, and *P. p. brevipodus* (Nagoya Daruma pond frog), which is distributed in western Japan. *P. p. brevipodus*
is traditionally divided into two distinct, local forms called the Okayama form in the west and the Nagoya form in the east (Ito, 1941; Moriya, 1954; Kawamura, 1962; Matsui and Hikida, 1985). They are genetically differentiated from each other by their external morphologies (Figure 1), mating calls, sex chromosomes, allozymes and mitochondrial genes (Moriya, 1951, 1954; Nishioka et al., 1992; Nishioka and Sumida, 1994; Ueda, 1994; Sumida et al., 1998, 2000a,b; Komaki et al., 2015). However, the two forms have never been defined taxonomically because their distributions are unclear. Since the genetic researches on the two local forms to date were always restricted to several representative populations, the area covering around 150 km between the two forms remains unstudied. It is still unknown whether the two forms are geographically separated or distributed sympatrically with mutual genetic introgression. Such information is definitely necessary for judging taxonomic positions of the two forms. Recently, the geographic populations of the Okayama form have been declining and are concerned about their possible extinction (Okochi et al., 1997). The degradation is especially severe in the western edge of the distribution, Hiroshima Prefecture, where only a few tiny populations have survived (Naito et al., 2014). Conservation of the population and environment is an urgent issue and taxonomic definition of the form is expected to assist the conservation activities.

In this study, we collected samples of the two major forms in western Japan and investigated sequences of mitochondrial and nuclear genes in order to assess whether the two forms are separated geographically or are distributed sympatrically with mutual genetic introgression. In particular, the eight populations in Okayama and Hyogo Prefectures are located between the known distributions of the two forms and were genetically examined for the first time in 63 years since the primary morphological study of Moriya (1954) (Figure 2).

MATERIALS AND METHODS

Frogs

The number of frogs of Pelophylax porosus brevipodus, P. p. porosus and P. nigromaculatus used for sequence analyses are listed in Table 1 and their collecting locations are shown in Figure 2. We collected three frogs of P. p. brevipodus each from Aichi and Gifu Prefectures, and reared them at our laboratory, while all other tissue samples were taken from the toe-clips in the fields, and stored in 100% ethanol until use. The frogs were thereafter released to the fields. Animal care and experimental procedures were conducted under approval of the Committee for Ethics in Animal Experimentation at Hiroshima University (Permit Number: G13-3).

DNA Extraction and PCR Amplification

Genomic DNA was extracted from the tissue samples using DNeasy blood and tissue kit (QIAGEN) according to the manufacture’s instruction. Mitochondrial cytochrome b and nuclear SOX3 fragments were amplified in 50 µl solution including 1.0 µl of DNA solution, 0.2 µl GXL Taq polymerase (TaKaRa), 5 µl of 10× Buffer, 4 µl of 2.5 mM dNTP, and 1 µl of 12.5 mM primers at 98°C for 5 s followed by 30 cycles of 98°C...
**TABLE 1** | Populations, haplotype of mitochondrial cytochrome b, D-loop region, and genotype of nuclear SOX3.

| Locality No. | Species | Population | Prefecture | City | Town or area | No. of frogs ($\text{♂}, \text{♀}, \text{juvenile}$) | cytb haplotype | Repeats in D-loop region | SOX3 (233rd) |
|--------------|---------|------------|------------|------|--------------|-----------------|----------------|--------------------------|---------------|
| 1            | Pelophylax | K-Nagoya | Aichi | Kita-Nagoya | Shikatsu | 3 (2,1,0) | A1, A3 | AB | GG |
| 2            | porosus | Gifu | Gifu | Gifu | | 3 (1,2,0) | A2, A3, A4 | | |
| 3            | brevipodus | Iga A | Mie | Iga | (A) | 4 (1,3,0) | B1 | | |
| 4            | brevipodus | Iga B | Mie | Iga | (B) | 2 (0,2,0) | B3 | | |
| 5            | Kobe-O | Hyogo | Kobe | Oshibedani | | 9 (7,2,0) | B1 | AB | |
| 6            | Kobe-H | Hyogo | Kobe | Hirano-machii | | 10 (3,2,5) | B1 | | |
| 7            | Kakogawa-YA | Hyogo | Kakogawa | Yahata (A) | | 30 (3,6,21) | B2, B1, C3 | | |
| 8            | Kakogawa-YB | Hyogo | Kakogawa | Yahata (B) | | 6 (1,1,4) | B1, C3 | | |
| 9            | Kakogawa-I | Hyogo | Kakogawa | Inami | | 10 (2,5,3) | B1, C3 | ABABAB, ABA | | |
| 10           | Kobe-O | Hyogo | Kobe | Oshibedani | | 9 (7,2,0) | B1 | AB | |
| 11           | Okayama-S | Okayama | Okayama | Seto | | 11 (5,5,1) | C1, C3 | ABA | TT |
| 12           | Okayama-N | Okayama | Okayama | Nodono | | 20 (5,11,4) | C1 | | |
| 13           | Kurashiki | Mabi | Kurashiki | Mabi | | 8 (1,2,5) | C1 | | |
| 14           | Fukuyma | Hiroshima | Fukuyma | Kannab | | 8 (3,5,0) | C1 | | |
| 15           | Miyoshi-Y | Okayama | Miyoshi | Kisa, Yasuda | | 10 (0,0,10) | C1 | | |
| 16           | Miyoshi-K | Okayama | Miyoshi | Kisa, Kaitahara | | 10 (0,0,10) | C1 | | |

**RESULTS**

**Mitochondrial Cytochrome b**

We collected samples consisting of 156 specimens (38 males, 55 females, and 63 juveniles) from 16 populations covering their present habitat in western Japan (Figure 2 and Table 1). We determined the nucleotide sequences of 566 base pairs of the mitochondrial cytochrome-b gene. Ten haplotypes were identified and the gene tree was constructed using the maximum likelihood (ML) method (Figure 3 and Table 1). The haplotypes formed two distinct clades, which are designated PB-N and PB-O because they correspond to the Nagoya form and Okayama form, respectively. The genetic ($p$) distance between the two clades was 0.055 and that between the two subspecies was 0.054, suggesting that the genetic relationships among $P. p. porosus$ and the two local forms of $P. p. brevipodus$ are within almost equal range of each other. Notably, two haplotypes of PB-O and PB-N were detected in Kakogawa-YA, -YB and -I populations of Hyogo Prefecture (7, 8, and 9 in Figures 2–4), which were located immediately east over the Kakogawa River. Three of 30, one of six, and three of ten specimens examined in the populations had PB-O haplotypes, while the others possessed 10 s, 64°C for 40 s, and 72°C for 60 s. The amplified product was purified using GFX PCR DNA and Gel band purification kit (GE Healthcare), and was used for nucleotide sequence determination with 3130XL sequencing machine (ABI).

Mitochondrial fragments including D-loop region (300∼500 bp) were amplified and purified by the above methods, and were cloned into pUC118 vector using Mighty cloning kit (TaKaRa) with competent cell DH5α (Ecos, Nippon gene) according to the manufacturers' instructions. One to three colonies were picked up and the nucleotide sequences were determined by the method described above. Gene trees were constructed based on the nucleotide sequence of cytochrome-b gene by the methods of maximum likelihood (ML), neighbor joining (NJ) and maximum parsimony (MP) methods using Mega 7 software (Kumar et al., 2016). $p$-distance was also calculated using the above software. Primers used are forward 5′-CCA TGC ACT ACA CAG CCG ACA-3′ and reverse 5′-AGG TTT TTG CGA TAG GCC GGA-3′ for cytochrome b (designed in this study using software Genetix ver. 7.3, Genetix corp.), S1 5′-GTG CGC TCC TCC TGC TCT TCT TTT-3′ and A1 5′-TCC TCA AGT TTT CTG CAT TCT GAT-3′ for SOX3 (Miura et al., 2016), and F23 5′-ATG AAT GCT ATA ATG ACA TAA TGT-3′ and R21 5′-TGC TGG TGG TCT AGA CCG CCA GTG GAG GCC TGT-3′ for D-loop region (Sumida et al., 2000a). The sequences of ten haplotypes (A1–4, B1–3, and C1–3) of cytochrome b have been deposited with the DDBJ Data Libraries under the accession numbers LC217488-LC217457, and the sequences of SOX3 (Kurashiki and Iga populations), under the accession numbers LC316654 and LC316655, respectively.
Repeated Sequence in D-loop Region

The D-loop region of the mitochondrial genome includes a highly repeated sequence. We cloned this region and determined the nucleotide sequences of specimens from eight populations of *P. p. brevipodus* (Table 1). The repeated region comprised of repeats of two kinds of 17-bp units designated types A and B of which nucleotides at the 9th and 10th positions were different: AG and GT, respectively (Figure 3 and Supplementary Table 1). The repeated pattern was different among populations (Figure 3 and Supplementary Table 1). Pattern AB was specific to the Kita-Nagoya population (population No. 1 in Table 1 and Figure 3), while pattern ABA was observed in the Kakogawa I2, Okayama-S1 and three Hiroshima populations (9, 11, and 14–16). In the Iga, Kobe-O and Kakogawa I1 populations (4, 6, and 9), the observed patterns were ABAB, AB, and ABABAB, respectively. All the repeat patterns were thus classified into two types: AB (or repeats of AB) and ABA. The two types corresponded with the two major clades of cytochrome b: PB-N with AB or repeats of AB and PB-O with ABA, respectively. In the Kakogawa-I population (9), the specimen with the PB-N cyt-b haplotype had the ABABAB pattern while that with the PB-O haplotype possessed the ABA pattern.

SOX3

The sequence of 860 base pairs of the nuclear SOX3 gene was determined for 140 specimens from 16 populations. The nucleotide at position 233 varied by population (Figure 4 and Table 1). In the six eastern populations (1–6: Kita-Nagoya, Gifu, Iga-A, Iga-B, Kobe-O, and Kobe-H), all specimens were homozygous for guanine. On the other hand, in the five western populations (11, 13–16: Okayama, Kurashiki, and three of Hiroshima Prefecture), all were homozygous for thymine. In the other five populations (7–10, 12) located at the intermediate regions, the specimens were heterozygous or homozygous for guanine or thymine (Figure 4). The frequency of guanine in these populations varied from 10 to 41%.

DISCUSSION

Based on the mitochondrial cytochrome b, the two major *P. p. brevipodus* forms of Okayama and Nagoya were identified as distinct clades, and two major types of the D-loop region supported the cytb clades. The genetic distance (p-distance, 0.055) between the two forms was very similar to that (0.054) between the two subspecies. These genetic relationships are well supported by another study that used mitochondrial and nuclear genes (Komaki et al., 2015). The distribution boundary between the two forms was for the first time found in this study. It is located at a very small area that included the Kakogawa populations (7–9 in Figures 2, 4) of Hyogo Prefecture and was where two haplotypes of the Okayama and Nagoya forms co-existed. This shows that the two forms were geographically isolated from each other in the past and have secondarily contacted at the small area after they were genetically differentiated. The molecular clock based on cytochrome b and seven nuclear genes estimates that the two forms were separated from each other around 1.3
that separates the two forms of Pelophylax porosus brevipodus, or no geographic event that actually occurred 1.3 MYA is known. However, some geographic barrier must have existed in the past and prevented crossings across the boundary area, because many other animals, such as grasshopper, harvestman, frog, landing snail, and monkey, are likewise genetically differentiated between the west and the east of the boundary region (Tsursushi et al., 1991; Kawakami, 1999; Nishi and Sota, 2005; Kawamoto et al., 2007; Nichizawa et al., 2011). Conversely, it was found that nuclear gene SOX3 showed introgression over the boundary from eastern Nagoya form into the western Okayama form. The genetic affinity between the two forms is also confirmed by the results of artificial crossings in the study of Moriya (1960a,b), showing fertile hybrids between the two forms. However, it was quite difficult in this study to recognize the genetic introgression in external morphology: for example, a central line on the back, which was normally observed in 44% frogs of Nagoya form, was not found in any populations of the Okayama form (except one specimen in Kakogawa I population, No. 9 on the map). A deeper analysis on nuclear genomes of the two forms focusing on the populations around the boundary is required to verify the on-going introgression of the genomes.

CONCLUSION

It is evident that the two local forms had been once isolated from each other and accumulated their genetic differences, and thereafter they have secondarily contacted immediately east over the Kakogawa River (Figure 4) and possibly the Okayama form is now accepting introgression from the Nagoya form. We speculate that the ancestral lineage of the Okayama form remains around the eastern edge of the range (Hiroshima Prefecture and the western region of Okayama Prefecture in Figure 2) where population declining and extinction are concerned. At a next step of the research, taxonomic definition of the two forms are expected (for example, name of Okayama Daruma pond frog is given to the Okayama form), because they are precisely identified and the geographic boundary between the two forms is very clear based on the mitochondrial DNA. Unfortunately, the previous study on morphology (Moriya, 1954) used no statistical analyses and examined only one population of the Okayama form, and the previous mating call analysis (Ueda, 1994) was restricted to just one or two populations of each form, which are located at the extremes in distribution. Hence, a future taxonomic approach needs to consider the distribution range for choosing populations and complete investigation on the morphology and mating calls of the two forms.

AUTHOR CONTRIBUTIONS

IM and MO designed the study and wrote the manuscript. YN, MO, and IM performed the experiments and analyses. TD, KI, YY, TF, and J-iN collected the specimens and discussed about the results. All authors read and approved the final manuscripts.

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SUPPLEMENTARY MATERIAL

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fgene.2018.00079/full#supplementary-material

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Conflict of Interest Statement: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

The handling Editor declared a past co-authorship with the authors MO and IM.

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