The very low genetic variability on Aceh Tamiang’s (Indonesia) population of Painted Terrapin (*Batagur borneoensis*) inferred by cytochrome oxidase I (CO I) and D-loop (control region)

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Abstract. Guntoro J, Wirdateti, Riyanto A. 2020. The very low genetic variability on Aceh Tamiang’s (Indonesia) population of Painted Terrapin (*Batagur borneoensis*) inferred by cytochrome oxidase I (CO I) and D-loop (control region). Biodiversitas 21: 2514-2520. Populations of *Batagur borneoensis* have been rapidly decreasing due to the harvesting of adults and eggs for food and the construction of beachfront property causing the loss of nesting areas. By the new Indonesian regulation, since 2018 this turtle listed in the protected animal. Meanwhile, IUCN placed as critically endangered which indicating a high risk of extinction in the wild in the near future (www.iucnredlist.org). We used cytochrome oxidase I (COI) and control region D-loop region to investigate intraspecific variations on Aceh Tamiang’s population of painted terrapin, *Batagur borneoensis*. DNA material was gathered from saliva collected from 90 juveniles in the reaching facility of Sukacita Lestari Indonesia Foundation which hatched from eggs collected from December 2015 to April 2016 from 30 nests on beach area at Aceh Tamiang. The population showed very low genetic variability (haplotype diversity, Hd = 0.457 based on COI and 0.405 based on D-loop; nucleotide diversity, π = 0.00089 based on COI and 0.00076 based on D-loop). So, we suggested that further study such as more exploration to find new wild populations and genetic study across wild populations should be done to reveal genetic variability and genetic structure which important to decide the conservation strategy. At the time for Aceh Tamiang’s population, the ranching conservation program should be maintained at least to keep the successful hatching from hunters and natural predators both during eggs laying and hatching.

**Keywords:** Aceh Tamiang, *Batagur borneoensis*, COI, D-loop, genetic variability, Indonesia

INTRODUCTION

Turtles particularly have value to humans, whether as food, medicine, pets, or as providers of ecological services, and their very slow recovery from (over) exploitation, turtles tend to be at the cutting edge of biodiversity decline, and an indicator of ecosystem degradation. The turtles distribution and their occurrence of deep phylogenetic lineages, is not uniform across the planet. Therefore, turtles clearly represent a global biodiversity conservation priority (Mittermeier et al. 2015). Mittermeier et al. (2015) also placed Indonesia in the turtle priority area of Sundaland hotspot based on species richness and endemicty.

Painted terrapin (*Batagur borneoensis*) is one of the non-marine Indonesian turtle which listed critically endangered (IUCN 2017) and listed in 25 most extinction in the world (Rhodin et al. 2011; Stanford et al. 2018). At the time, including in protected animals in Indonesia by Permenhut 106/2018. Populations of this turtle have been rapidly decreasing due to the harvesting of adults and eggs for food and the construction of beachfront property causing the loss of nesting areas (Stanford et al. 2018). Duli (2009) showed that no genetic differentiation between south and north Peninsular Malaysia population and until now nothing known on the dispersal pattern of this terrapin. Believed, that now to be restricted to small remnant populations in Malaysia and part of North Sumatra (Stanford et al. 2018). As a result of this population decline, *B. borneoensis* is listed by the IUCN as critically endangered, indicating a high risk of extinction in the wild in the near future (www.iucnredlist.org).

Aceh Tamiang beach is one of few known nesting areas of *Batagur borneoensis* in Indonesia (Guntoro 2012). Genetic variability is important that determine the availability or adaptation of the population from the environment changes (Moll and Moll 2004). In the small and limited population is believed to have low genetic variability. So, it is not excessive if information on the genetic variability of Aceh Tamiang’s population is urgent to be known, especially as consideration in conservation efforts.

The use of genetics to inform evidence-based management decisions for highly threatened species can improve conservation outcomes (Allendorf et al. 2010; Zhan et al. 2014; Corlett 2016; Ismail et al. 2016; Pierson et al. 2016; Alvarez et al. 2019). Almost half of freshwater turtle species worldwide are threatened, and many exhibit low levels of genetic variability, therefore the identification of genetic within natural populations is important for conservation strategies (Iblow et al. 2014). Mitochondrial DNA (mtDNA) and its loci are one of several approaches in genetic evaluation of wild animals, including in turtle
MATERIALS AND METHODS

Study area

The juveniles in the ranching facility of Sukacita Lestari Indonesia Foundation were hatched from eggs collected from 30 nests at riverbank in Ujung Aceh Tamiang, Aceh, Indonesia (N 04°25'12.29", E 098°16'49.35") December 2015 to April 2016 (Figure 1).

Figure 1. Map of Sumatra showing nest locality of eggs collected (yellow circle), Ujung Aceh Tamiang, Aceh, Indonesia

Biometric data

We measured each individuals juvenile before collected the buccal cells for DNA material sample. Measurement including stright median carapace length and stright maximum carapace width. We also recorded body weight.

DNA material sampling

DNA material gathered from buccal cells collected from 90 juveniles in the ranching facility from 30 nests of Sukacita Lestari Indonesia with origin mentioned above in study area. The collected samples as many as three animals of each nest by random. The buccal cells preserved in 95% ethanol prior to extraction.

DNA extraction and amplification

Genomic DNA was extracted from saliva using a commercial Qiagen DNeasy Tissue Kit Kit, following the manufacturer’s instructions. We amplified the D-loop region and COI gene mtDNA by polymerase chain reaction (PCR). The primer Bb dlF (5'-TCTCTACATGATTCACAGGATGCTTTTAACCGGTTGGTAG-3) using for D-loop (Hawkins 2010), and COI Tuntong F (5'-CGCGGAATTAAGCCAACCAGTGTTGCTTTAA-3) and COI Tuntong R (5'-TTGGTACAGGATTGGTGTCGTACCGG-3) for COI was designed. PCR amplifications were done with reaction mixture from KAPA Robust HotStart (KAPA Biosystems product) in 30 ul. Thermal cycling follows: (i) D-loop predenaturation at 95°C for 10 minutes, 35 denaturation cycles at 94°C for 30 seconds, annealing at 57°C for 20 seconds, extension at 72°C for 5 minutes, and final extension at 72°C for 10 seconds; (ii) COI predenaturation 94°C for 4 minutes, 35 denaturation cycles at 94°C for 45 seconds, annealing at 55°C for 35 seconds, extension at 72°C for 1 minute, followed by a final extension at 72°C for 10 minutes. Amplification was done used Biorad Thermo Cycler. PCR product was sequenced in First Base Co., Singapore, and Macrogen Co., Korea.

Data analysis

Biometric

We examined the body sizes and weight of 90 juveniles which hatched from 30 nests by using One-way ANOVA. The calculation was conduct by Minitab.

Molecular

The acquired sequences were edited using BioEdit software (Hall, 1999) and aligned using Clustal X (Thompson et al. 1994). For the haplotypes, we used DNaSP ver.5 (Librado and Rozas, 2009) to calculate haplotype diversity (h), Tajima’s test, Fu and Li’s test, and nucleotide diversity (π). Haplotype diversity can be interpreted as the probability that two individuals from a given sampling unit will have different haplotypes. The Kimura 2-parameter genetic distance (d) measure (Kimura 1980) was calculated to reveal the genetic diversity in population in MEGA-X (Kumar et al. 2018).

We calculated the genetic distance between all pairwise comparisons of wild-caught 90 individuals of 30 nestings. Relationships among mitochondrial haplotypes were
visualized by constructing a neighbor-joining tree and Maximum Likelihood using MEGA-X (Kumar et al. 2018). Branch points were tested with 5000 bootstraps using Kimura-2 parameter method.

RESULTS AND DISCUSSION

Morphometric and weight
The descriptive on carapace length, carapace width, and the weight of sample from each nest are presented in Table 1. These were no significant (p=1.000) differences between 90 juveniles which hatched from 30 nests.

Genetic variability
Both of genetic marker COI and D-loop showed the Aceh Tamiang’s population has very low genetic variability, is showing the lower genetic distance (d) of 90 juveniles from 30 nesting that was 0.001 ± 0.001; and between haplotype was 0.003 ± 0.001 (Table 2).

COI
The total number of the sequence length is 548 bp, and it found as many as 7 gaps in this alignment, so total number to analysis was 541 bp. The result of 90 individuals showing that number of variable sites (S) is 6 (1.12%), and sequence conservation (C) is 53.9 (89.9%). The haplotype diversity, Hd = 0.457 ± 0.037, this is indicated by a very low nucleotide variation that there are only 2 different sites in position 1 and to 2, i.e. base substitution from A to C or transversion. As a result, it’s revealing three distinct haplotypes, e.i. AA (29 individuals), CA (60 individuals), and AC (1 individual) (Table 3). Several sites insertion and deletion but not change the protein composition. The nucleotide diversity, π = 0.00089± 0.00008 with average number of nucleotide differences, k = 0.4717.

D-loop region
From augmented 578 bp of D-loop region that was extracted from 90 individuals revealed four distinct haplotypes from three number of variable sites (S = 0.52%) at position 412, 482, and 575, and sequence conservation (C) is 99.48% (575 sites). With only 22 individuals have different nucleotides with single base e.i. A to G, G to A and A to T. The haplotype diversity, Hd = 0.405 ± 0.058, it is revealing four haplotypes that are AGA (68 individuals); GGA (14 individuals); AGT (4 individuals); and AAA (4 individuals) (Table 3). The nucleotide diversity, π = 0.00076 ± 0.00012 with average of nucleotide differences, k = 0.4374.

Table 1. Recorded on range body sizes and weights and it’s mean ± standard deviation (SD) of juvenile’s Batagur borneoenesis from 30 nests in Aceh’s Tamiang nest area, Indonesia

| Nest | No. individuals | Weight | Median carapace length | Maximum carapace width |
|------|----------------|--------|------------------------|------------------------|
| 1    | 3              | 155-250 (198.3±48.04) | 9.4-10.8 (10.1±0.70) | 8.7-9.7 (9.2±0.50) |
| 2    | 3              | 155-195 (170.0±21.80) | 9.7-10.1 (9.8±0.23) | 8.8-9.4 (9.0±0.32) |
| 3    | 3              | 190-240 (211.7±25.66) | 10.2-11.1 (10.6±0.45) | 9.2-9.8 (9.5±0.30) |
| 4    | 3              | 200-275 (230.0±39.67) | 10.4-11.8 (10.9±0.81) | 9.4-10.4 (9.7±0.58) |
| 5    | 3              | 175-200 (19.0±13.23) | 9.8-10.4 (10.1±0.31) | 8.9-9.2 (9.1±0.17) |
| 6    | 3              | 175-200 (185.0±13.23) | 10.7-20.7 (13.7±0.69) | 9.3-9.8 (9.5±0.27) |
| 7    | 3              | 160-190 (173.3±15.28) | 9.6-10.5 (10.9±0.45) | 8.9-9.4 (9.2±0.29) |
| 8    | 3              | 150-155 (151.7±2.89) | 9.3-9.4 (9.4±0.06) | 8.6-8.8 (8.7±0.11) |
| 9    | 3              | 150-190 (166.7±20.82) | 9.3-10.0 (9.6±0.38) | 8.6-9.1 (8.8±0.25) |
| 10   | 3              | 170-190 (176.7±11.55) | 9.6-9.9 (9.8±0.15) | 8.8-9.8 (9.2±0.56) |
| 11   | 3              | 130-170 (155.0±21.79) | 8.9-9.7 (9.4±0.46) | 8.6-9.1 (8.8±0.25) |
| 12   | 3              | 180-190 (185.0±5.00) | 9.9-10.3 (10.1±0.21) | 9.9-9.3 (9.1±0.17) |
| 13   | 3              | 160-200 (176.7±20.82) | 9.7-10.7 (10.1±0.55) | 8.9-9.6 (9.1±0.50) |
| 14   | 3              | 160-190 (180.0±17.32) | 9.3-10.2 (9.9±0.52) | 8.5-9.2 (8.9±0.38) |
| 15   | 3              | 150-180 (160.0±17.32) | 9.2-10.2 (9.5±0.58) | 8.9-9.2 (8.6±0.60) |
| 16   | 3              | 160-180 (170.0±10.00) | 9.7-9.8 (9.8±0.06) | 8.8-9.0 (8.9±0.12) |
| 17   | 3              | 140-170 (156.7±15.28) | 9.7-10.0 (9.8±0.15) | 8.2-9.1 (8.7±0.46) |
| 18   | 3              | 150-190 (173.3±20.82) | 9.1-10.2 (9.8±0.64) | 8.6-9.8 (9.2±0.60) |
| 19   | 3              | 150-160 (150.0±10.00) | 9.2-9.7 (9.5±0.25) | 8.3-8.9 (8.6±0.31) |
| 20   | 3              | 150-180 (163.3±15.28) | 9.5-10.2 (9.7±0.40) | 8.9-9.1 (9.0±0.10) |
| 21   | 3              | 130-160 (143.3±15.28) | 9.9-9.5 (9.2±0.27) | 8.3-8.5 (8.4±0.10) |
| 22   | 3              | 120-150 (140.0±17.32) | 9.2-9.7 (9.4±0.27) | 8.5-8.8 (8.6±0.15) |
| 23   | 3              | 160-190 (173.3±15.28) | 9.4-10.3 (9.8±0.45) | 8.7-9.3 (9.0±0.30) |
| 24   | 3              | 140-150 (143.3±5.77) | 9.2-9.5 (9.4±0.15) | 8.2-8.7 (8.4±0.25) |
| 25   | 3              | 130-140 (133.3±5.77) | 9.2-9.7 (9.4±0.36) | 8.8-8.7 (8.4±0.36) |
| 26   | 3              | 140-160 (146.7±11.55) | 9.2-9.8 (9.5±0.31) | 8.4-8.7 (8.6±0.15) |
| 27   | 3              | 120-160 (136.7±20.82) | 8.7-9.7 (9.1±0.53) | 8.1-8.6 (8.3±0.30) |
| 28   | 3              | 120-150 (133.3±15.28) | 8.8-9.4 (9.2±0.32) | 8.1-8.7 (8.5±0.35) |
| 29   | 3              | 150          | 9.2-9.5 (9.3±0.15) | 8.5-8.7 (8.6±0.10) |
| 30   | 3              | 100-140 (123.3±20.82) | 8.4-9.5 (9.0±0.57) | 7.6-8.4 (8.1±0.44) |
| Overall | 3              | 100-271 (164.9±20.57) | 8.4-20.7 (9.8±1.28) | 7.6-10.4 (8.9±0.48) |
Table 2. The polymorphism data on COI and D-loop of Batagur borneoensis in Ujung Aceh Tamiang’s population (Aceh, Indonesia)

| mtDNA Markers | Distance (d) | Population | Haplotype | Haplotype diversity (Hd) | Nucleotide diversity (π) | Conserved region (C) |
|---------------|-------------|------------|-----------|--------------------------|--------------------------|----------------------|
| COI           | 0.001 ± 0.001 0.003 ± 0.002 | 0.457 ± 0.037 | 0.00089 ± 0.00008 | 0.989 |
| D-loop        | 0.001 ± 0.001 0.003 ± 0.001 | 0.405 ± 0.058 | 0.00076 ± 0.00012 | 0.995 |

Table 3. Haplotypes data of Batagur borneoensis in Ujung Aceh Tamiang’s population (Aceh, Indonesia) revealed based on COI and D-loop

| Nucleotide position | COI nucleotide | D-loop nucleotide |
|---------------------|----------------|------------------|
| 1                   | A              | A                |
| 2                   | A              | A                |
| 412                 | -              | A                |
| 482                 | -              | G                |
| 575                 | -              | AA               |
| Haplotypes          | AA             | CA               |
| No. of individuals in each haplotype | 29 | 60 |

Phylogenetic analysis

The topology of ML and NJ trees of haplotypes of Batagur borneoensis in Ujung Aceh Tamiang’s population based on COI presented in Figure 2, meanwhile based on D-loop region presented in Figure 3. In COI, both ML and NJ showed that three different haplotypes were group in 2 groups. As well as COI, phylogeny trees based on D-loop also show high levels of nucleotide similarity among populations with very low genetic distances. The trees divided into two major groups with a limited branching because almost all individuals have the same nucleotide. These groups separation is not very clear or unstable, because the bootstrap values were very low.

There were a total of three haplotypes on COI and four haplotypes on D-loop region with n = 90. Haplotype sequences were very similar to each other and were on average only 0.3% divergent (range 0.2%-0.4%, Kimura-2 parameter distance) on COI. As well as COI, haplotype sequences using D-loop were on average 0.3% (range 0.2%-0.4%, Kimura-2 parameter distance). This result is lower than the result from 30 individuals of B. borneoensis were a total number of 13 haplotypes from Malaysia. Haplotype sequences were very similar to each other and were on average only 0.5% divergent (range 0.2%-1.3%, Kimura-2 parameter distance) (Meredith, 2010). The value of Tajima’s test was -0.307 and Fu and Li’s test was 0.0497, which means that the genetic diversity in the population is not significant.

In our study, from 30 nests which each nest represented by 3 individuals. Based on haplotype distribution showed that in COI there were nine nests which contain all juveniles with same haplotype (1 nest has haplotype AA and 8 nests have haplotype CA), meanwhile, 21 nests have different haplotype (in this case has 2 haplotypes in each nest). In the control region, there were 20 nests has same haplotype which dominated by haplotype AGA (17 nests), GGA (2 nests), and AAA (1 nest), and 10 other nest has different haplotype.

The percentage of similarity on haplotype between nests was reached 66.7% based on D-loop and 30% based on COI. It is shown that the population of B. borneoensis on the Ujung Aceh Tamiang coast in a critical condition. The nucleotide diversity (π) is very low (Table 2) with average number of nucleotide differences (k) = 0.4716 in COI and 0.4374 in D-loop. This low nucleotide diversity suggested a population bottleneck in the B. borneoensis, possibly caused by the hunting for consumption, certain retails, habitat lost, and highest in breeding level on the restriction habitat. Based on these genes region, seen that the diversity of individuals was higher in D-loop, from ten different nests showed have three individuals with different haplotypes, even though the total number of nests that were higher monomorphic. In contrast, the COI marker shows that nests with different haplotypes have higher percentages, but individual diversity is lower in the same nest, which only has 2 haplotypes in each nest. Control region is a hypervariable region with high base substitution so that it is possible for individual diversity to be identified higher. D-loop, giving confidence to this condition, because in vertebrates this region well known as hypervariable or has a high base mutation both substitution, deletion, or base insertion so it is widely used to see the level of intra-species diversity (Sacccone et al. 1987). D-loop revealed very little variation (Tikochinski et al. 2012).

It also seems, that probably nest with same haplotype came from same male but it needs to be proven. Therefore, genetic fingerprinting may also important to be conducted. From this method, the sex ratios of breeding adults, mating patterns such as level of multiple paternity and male reproductive will be known (Komoroske et al. 2017). In B. borneoensis this information really doesn’t know as well as on other Sundaland freshwater turtles. Multiple paternity was recorded on Podocnemis erythrocephala from Brazil (Fantin et al. 2010), Elseya albagula from Australia (Todd...
et al. 2013), and captivity population of Testudo hermanni boettgeri (Farke et al. 2015). Most information multiple paternity limited on sea turtle such as in hawksbills (Phillips et al. 2013), loggerheads (Sari et al. 2017), leatherbacks (Stewart and Dutton 2011), olive ridleys (Jensen et al. 2006) and green turtle (Fitzsimmons 1998).

Referring to the results of the above research it is necessary to improve conservation management by selecting new brooders or new blood to increase population diversity. Bottleneck population indicates high levels of inbreeding so new blood is needed. Although B. borneoensis populations in Indonesia have historically spread across the east coast of Sumatra Island and western Kalimantan, it is currently thought that the remaining populations in Sumatra are only found on the coast of Langkat Regency and the Aceh Tamiang mangrove forest area (Guntoro, 2012). Thus it is necessary to know the quality of each population that is spread as a whole on the islands of Sumatra and Kalimantan which can be used as new blood. In 1985, B. borneoensis was reported to be very abundant in five rivers in Peninsular Malaysia with the Setiu River supporting the largest population (Sharma, 1997), but today all populations are declining precipitously (Norkarmila, 2009). This means that the decline in the quality of the population of B. borneoensis is almost everywhere in Indonesia and Malaysia.

**Figure 2.** Maximum-likelihood (A) and Neighbor-joining (B) dendrograms of three haplotypes of Batagur borneoensis in Ujung Aceh Tamiang’s population based on COI. Nest36.1 - individual 1 from nest 36. Haplotypes in parenthesis

**Figure 3.** Maximum-likelihood (A) and Neighbor-joining (B) dendrograms of four haplotypes of Batagur borneoensis in Ujung Aceh Tamiang’s population based on control region D-loop. Nest6.3 - individual 3 from nest 6. Haplotypes in parenthesis
A recent study by Duli (2009) on comparison of genetic variation at mitochondrial cytochrome b and D-loop for female *B. borneoensis* and its congener *B. baska* on the east and west coast of Peninsular Malaysia show the significant genetic structure between east and west coast for *B. baska* but not for *B. borneoensis*. Since lacks information on dispersal patterns of *B. borneoensis*, we agree to Hawkins (2010) that possibility of the adult has capable of long-distance dispersal resulting in low population differentiation. Dunson and Moll (1980) report that when hatching leaves the beach they can withstand seawater for up to weeks, Hawkins (2010) was hypothesised that hatched may disperse by ocean current.

Our finding in low genetic variability may relevant results of Spitzweg et al. (2018) that studied on captive breeding project of *B. baska* in Bangladesh and Australia. They conclude that given the long life expectancy of turtles, this situation suggests that the wild population had experienced a severe decline for long ago. The few survivors are largely related at the level of half-sibs (or first cousins, aunts/uncles-nieces/nephews, or grandparents-grandchildren). They propose conservation strategy with preserved the present genetic diversity to the greatest extent possible. The reproduction of closely related terrapins has to be avoided. They also suggested implementing the selective breeding and some mating combinations be avoided.

Based on those situations above, we suggested that further study such as more exploration to find new wild populations, genetic study across wild populations should be done to reveal genetic variability and genetic structure, and genetic fingerprinting which important to decide the conservation strategy. At this time, for Aceh Tamiang’s population, the ranching conservation program should be maintained at least to keep successful hatching from hunters and natural predators both during eggs and hatching.

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