Characterization of limes (Citrus aurantifolia) grown in Bhutan and Indonesia using high-throughput sequencing

Tshering Penjor¹,², Takashi Mimura¹, Ryoji Matsumoto¹, Masashi Yamamoto³* & Yukio Nagano⁴*

¹Department of Applied Biological Sciences, Saga University, Honjo, Saga, Japan, ²Renewable Natural Resources Research Centre Wengkhar, Mongar, Bhutan, ³Faculty of Agriculture, Kagoshima University, Korimoto, Kagoshima, Japan, ⁴Analytical Research Center for Experimental Sciences, Saga University, Honjo, Saga, Japan.

Lime [Citrus aurantifolia (Cristm.) Swingle] is a Citrus species that is a popular ingredient in many cuisines. Some citrus plants are known to originate in the area ranging from northeastern India to southwestern China. In the current study, we characterized and compared limes grown in Bhutan (n = 5 accessions) and Indonesia (n = 3 accessions). The limes were separated into two groups based on their morphology. Restriction site-associated DNA sequencing (RAD-seq) separated the eight accessions into two clusters. One cluster contained four accessions from Bhutan, whereas the other cluster contained one accession from Bhutan and the three accessions from Indonesia. This genetic classification supported the morphological classification of limes. The analysis suggests that the properties associated with asexual reproduction, and somatic homologous recombination, have contributed to the genetic diversification of limes.

Citrus species, which include mandarin (Citrus reticulata), sweet orange (C. sinensis), lemon (C. limon), and grapefruit (C. paradisi), have high economic and nutritional value. Lime (synonym: West Indian lime, Mexican lime, and key lime), C. aurantifolia (Cristm.) Swingle, is a popular ingredient in the dishes and drinks of many countries because of its aroma and acidity. Assam, which is a northeastern state of India, is one of the candidate origins of some Citrus species, because numerous varieties of citrus are grown in this area¹. Yunnan, which is a province located in southwestern China, is another candidate origin of some Citrus species². Thus, the region from northeastern India to southwestern China is ideal for studying the diversity of Citrus species. Bhutan is located within this region. However, to date, the genetic resources for Citrus species have not been well characterized in this country³. Thus, it was chosen as one of our two study sites. Limes are also widely cultivated and consumed in Indonesia, with varieties from this region being introduced to Saga University, Japan, as budwoods in 1988. Thus, we chose Indonesia as the second of our two study sites.

Lime is often considered to be a chance seedling, with citron (C. medica) and papeda possibly being its parents⁴–⁶. Furthermore, some Citrus species, including lime, typically reproduce asexually through nucellar embryony. Vegetative propagation by humans (e.g., grafting, budding, and layering) is another method for the asexual reproduction of Citrus species. Consequently, the heterozygous state of limes has been maintained over the course of their diversification. Therefore, it is important to identify variation in the heterozygosity of limes when studying their genetic diversity.

Recent advances in DNA sequencing have allowed the extensive use of short DNA fragments (single or multiple genes) to study the phylogenetic relationships of Citrus species. For example, by using these techniques, we have previously reported the phylogenetic relationships of Citrus and its relatives based on rbcL and matK gene sequences⁷,⁸. However, it is difficult to study intraspecies relationships from one or several DNA fragments. Restriction site-associated DNA sequencing (RAD-seq)⁹ may solve this problem. RAD tags are the DNA sequences that immediately flank a particular restriction site throughout the genome¹⁰. Recent developments in high-throughput sequencing have enabled us to read all RAD tags in parallel. In particular, RAD-seq has facilitated the rapid discovery of thousands of single-nucleotide variants (SNVs). In fact, this technique has facilitated the determination of relationships within species¹¹. However, this method has not been applied to the study of Citrus species or other asexually reproducing organisms.
In this study, we morphologically characterized limes grown in Bhutan. Furthermore, we analyzed five accessions among these limes by RAD-seq. For comparison, we analyzed three lime accessions that were previously introduced to Japan from Indonesia. Thus, in the current study, we studied the relationships of limes grown in Bhutan and Indonesia and the history of their genetic diversification.

**Results**

Between 2009 and 2011, we identified several local lime accessions throughout Bhutan (Supplementary Table S1). Table 1 summarizes the leaf and fruit characteristics of these accessions. However, these data were not previously introduced to Japan from Indonesia. Thus, in the current study, we studied the relationships of limes grown in Bhutan and Indonesia and the history of their genetic diversification.

Associated photographs are presented in Figure 1. All accessions possessed the characteristic features of limes (e.g., a fruit rounder than a lemon, a highly acidic juice with a distinctive aroma, and a polyembryonic seed). Bhutanese accessions were separated into two classes based on their characteristics. The fruit surfaces of Bhutan-09003, Bhutan-09005, and Bhutan-11014 were smooth, whereas those of the other Bhutanese accessions were slightly smooth to coarse in texture. Furthermore, the wing of the leaf of Bhutan-11014 was wider than that of the other accessions. Bhutan-09005 and Bhutan-11014 are grown in the southern part of the country, whereas the other accessions from Bhutan are grown in almost all parts of the country. For comparative purposes, we morphologically characterized four lime accessions that were introduced to Saga University, Japan, in 1988 as budwoods from Indonesia (Table 1; Fig. 2). The four Indonesian accessions had relatively similar characteristics. Furthermore, the fruit surfaces of the Indonesian accessions were smooth, and the wing of the leaves was wider than that of other Bhutanese accessions, except for Bhutan-11014. Thus, these leaf characteristics are similar to those of Bhutan-11014, while the fruit characteristics are similar to those of Bhutan-09003, Bhutan-09005, and Bhutan-11014.

For further analysis, including RAD-seq analysis, we selected five Bhutanese limes (i.e., Bhutan-09005, Bhutan-09015, Bhutan-09024, Bhutan-09027, and Bhutan-09030) and three Indonesian limes (i.e., Indonesia-88035, Indonesia-88045, and Indonesia-88065) based on the preliminary fruit diversity characteristics. Both types of fruit surface were selected for the Bhutanese accessions, while three out of the four accessions were randomly selected for the Indonesian accessions. In addition, when we started the RAD-seq analysis, the only available Bhutanese accession carrying fruit with smooth surfaces was Bhutan-09005 (out of a possible Bhutan-09003, Bhutan-09005, and Bhutan-11014).

Next, to further elucidate the difference among the morphological traits of these eight accessions, we measured the leaf characteristics using trees grown under similar (greenhouse) conditions (Fig. 3, Table 2). Because the leaf characteristics of Bhutan-09005 were not available in the preliminary characterization, it is important to obtain a measurement. The area of the leaf blade ranged from 200.7π mm² to 687.6π mm² and from 313.3π mm² to 651.8π mm² in the Bhutanese and Indonesian accessions, respectively. The wing area ranged from 2.01π mm² to 6.83π mm² and from 8.48π mm² to 17.70π mm² in the Bhutanese and Indonesian accessions, respectively. The wing area of the Indonesian accessions was larger than that of the Bhutanese accessions. Bhutan-09005 had the largest wing of the Bhutanese accessions. The wing index of this accession was 3.33, which was the highest among eight accessions. The indices of the remaining Bhutanese accessions (Bhutan-09015, Bhutan-09024, Bhutan-09027, and Bhutan-09030) and three Indonesian accessions were 0.42–0.61 and 1.66–2.93, respectively. Based on the leaf traits, particularly wing size, Bhutan-09005 was clearly distinguished from the other Bhutanese accessions, and more closely resembled the Indonesian accessions. Thus, the quantitative characterization of leaf traits classified the eight accessions into two clusters as follows (p < 0.01): (1) the Bhutanese cluster containing Bhutan-09015, Bhutan-09024, Bhutan-09027, and Bhutan-09030 accessions with small wings and (2) the Indonesian cluster containing Bhutan-09005, Indonesia-88035, Indonesia-88045, and Indonesia-88065 accessions with large wings.

We sequenced the EcoRI RAD-tag from each of the eight accessions. After quality filtering the sequencing data, we obtained ≥24.5 million RAD-tag sequences (~1 Gb sequences) from each sample (Table 3). We genotyped these eight accessions using the genome data of the sweet orange (327,944,670 bp) by analyzing the bowtie program (a program used to make short read alignments) and the pstacks program (a program used to build loci) of the Stacks pipeline. Because the de novo RAD assembly of short reads overassembles some paralogous loci, we only genotyped them in the presence of reference genome information. We showed the depth of RAD-seq coverage for each sample across the reference genome and the coverage histogram of each sample (Supplementary Table S1). The RAD-seq results showed that more than sufficient amount of sequencing data were obtained. Although the data from Indonesia-88045 contained a relatively higher amount of low-coverage reads (probably caused by the degradation of the genomic DNA), these low-coverage reads were removed by in silico analysis.

To compare the genotypes of the eight accessions, we analyzed the aligned data using the Stacks pipeline as follows: cstacks (a program used to create a catalog of loci), sstacks (a program used to match samples with those in the catalog), and genotypes (a program used to export a Stacks data set as a set of observed haplotypes at each locus or with the haplotypes encoded into genotypes). These in silico steps integrated the data from the eight lime accession samples, corrected false-negative heterozygous genotypes and false-positive homozygous genotypes, extracted the sites useful for genotyping, and efficiently excluded certain errors, such as contamination caused by improper PCR amplification or degradation of the genomic DNA. However, it should be noted that it is difficult to completely remove all errors. The genotypes program compared 78,527 loci (3,219,607 bp, including the EcoRI sequence ATTC). The 42,613 sites that were potentially useful for genotyping (Supplementary Fig. S2) were extracted from the output of the genotypes program using the Perl script. Therefore, ~1.3% of sites from the analyzed regions were useful for genotyping. Among these 42,613 sites, 4846 were parsimony-informative. We constructed a phylogenetic tree using the maximum-likelihood method. This phylogenetic tree clearly showed that the eight accessions could be separated into two clusters, the Bhutanese and Indonesian clusters, as defined above. This classification was in accordance with the results of our morphological characterizations (Fig. 1, 2, and 3; Tables 1 and 2). Furthermore, phylogenetic analysis divided the Indonesian cluster into two subclusters, Indonesian subcluster 1, containing Bhutan-09005 and Indonesia-88065, and Indonesian subcluster 2, containing Indonesia-88035 and Indonesia-88045. Among the 42,613 sites, the Bhutanese cluster, Indonesian subcluster 1, and Indonesian subcluster 2 carried 41,691, 41,810, and 41,827 identical genotypes, respectively. In other words, the numbers of the genotypes that did not match within the Bhutanese cluster, Indonesian subcluster 1, and Indonesian subcluster 2 were 922, 803, and 786, respectively. Among the 42,613 sites, the Indonesian cluster carried 37,612 identical genotypes. In other words, the numbers of the genotypes that did not match within the Indonesian cluster was 5001.

Based on the results of the genotypes program, we estimated the minimum number of heterozygous genotypes for each accession (Table 3). The analysis showed that the frequencies of the heterozyg-
## Table 1 | Leaf and fruit characteristics of the Bhutanese and Indonesian lime accessions

| Accession   | Leaf blade Length (mm) | Leaf blade Width (mm) | Wing Length (mm) | Wing Width (mm) | Skin color | Fruit surface | Flesh color | Puffing | Granulation | Peeling | Bitterness | Sweetness | Sourness | Diameter (mm) | Height (mm) | D/H index | pH of juice | Embryo color | Polyembryony Number |
|-------------|------------------------|-----------------------|------------------|-----------------|-------------|--------------|-------------|---------|-------------|---------|------------|-----------|----------|----------------|-------------|-----------|-------------|-------------|---------------------|
| Bhutan-09003 | -                      | -                     | -                | -               | Yellowish green | Smooth       | Green       | None    | None        | Difficult | -          | -         | High     | 36.6          | 38.5        | 94.8      | -           | -           | -                   |
| Bhutan-09005 | -                      | -                     | -                | -               | Yellowish green | Smooth       | Green       | None    | None        | Slightly difficult | None     | Low    | High     | 34.3          | 38.5        | 89.1     | -           | Cream, Pale green | Poly 5.2      |
| Bhutan-09015 | 65.1                   | 30.0                  | 5.6              | 1.5             | Green        | Slightly coarse | Cream       | None    | None        | Difficult | None       | Low     | High     | 40.4          | 39.2        | 103.2    | 2.4         | Pale green, Green | Poly 4.6      |
| Bhutan-09018 | 84.7                   | 38.3                  | 8.5              | 1.7             | Green        | Slightly smooth | Green       | None    | None        | Difficult | None       | Low     | Medium  | 34.7          | 35.9        | 96.7     | 2.5         | Pale Green | Poly 0.3      |
| Bhutan-09019 | 91.0                   | 45.2                  | 6.3              | 1.6             | Yellowish green | Medium       | Yellow     | None    | None        | Difficult | None       | Low     | Medium  | 42.7          | 44.1        | 96.8     | 2.5         | Pale Green | Poly 1.0      |
| Bhutan-09024 | 85.4                   | 37.6                  | 5.0              | 1.6             | Green        | Slightly smooth | Green       | None    | None        | Difficult | None       | Low     | Medium  | 30.4          | 29.8        | 102.0    | 2.4         | Pale Green | Poly 4.8      |
| Bhutan-09027 | 64.3                   | 30.1                  | 4.0              | 1.5             | Green        | Slightly smooth | Green       | None    | None        | Difficult | None       | Low     | High    | 39.6          | 40.5        | 97.8     | 2.5         | Cream, Pale green | Poly 6.4      |
| Bhutan-09030 | 78.2                   | 34.4                  | 4.6              | 1.7             | Yellowish green | Slightly coarse | Cream       | None    | None        | Difficult | None       | Low     | High    | 45.2          | 46.6        | 97.0     | 2.5         | Pale green, Green | Poly 15.3     |
| Bhutan-11006 | 75.7                   | 35.1                  | 4.4              | 2.0             | Yellow       | Slightly smooth | Medium     | Yellow  | None        | Difficult | None       | Low     | High    | 36.6          | 37.6        | 97.3     | 2.2         | Cream, Pale green | Poly 12.0     |
| Bhutan-11009 | 71.2                   | 33.5                  | 3.7              | 1.6             | Yellow       | Slightly smooth | Yellow     | None    | None        | Difficult | None       | Low     | High    | 42.2          | 44.0        | 95.9     | 2.3         | Cream       | Poly 11.4     |
| Bhutan-11014 | 64.1                   | 39.6                  | 8.8              | 5.5             | Yellow       | Smooth       | Cream      | None    | None        | Difficult | None       | Low     | High    | 39.7          | 44.9        | 88.4     | 2.1         | Cream       | Poly 6.8      |
| Bhutan-11018 | 80.6                   | 39.9                  | 4.2              | 1.8             | Green        | Slightly coarse | Green       | None    | None        | Difficult | None       | Low     | High    | 39.7          | 44.9        | 88.4     | 2.1         | Cream       | Poly 6.8      |
| Bhutan-11026 | 100.1                  | 49.0                  | 4.1              | 2.4             | Green        | Coarse       | Cream      | -       | -           | None       | Low        | None    | Low     | None          | None        | 54.6     | 103.8       | Cream       | Poly 11.5     |
| Bhutan-11030 | 98.9                   | 47.0                  | 4.3              | 2.1             | Green        | Coarse       | Cream      | -       | -           | None       | Low        | None    | Low     | None          | None        | 39.1     | 99.5        | Cream       | Poly 4.4      |
| Indonesia-88035 | 53.6                 | 32.6                  | 8.5              | 3.6             | Yellowish green | Smooth       | Yellowish green | None    | None        | Difficult | None       | Low     | High    | 33.7          | 38.0        | 88.7     | 2.2         | Cream, Pale green | Poly 7.2      |
| Indonesia-88045 | 65.2                 | 36.3                  | 6.6              | 3.1             | Yellow       | Smooth       | Yellow     | None    | None        | Difficult | None       | Low     | High    | 39.4          | 45.0        | 87.6     | 2.1         | Cream       | Poly 8.5      |
| Indonesia-88065 | 49.1                 | 26.0                  | 5.1              | 2.6             | Yellow       | Smooth       | Yellow     | None    | None        | Difficult | None       | Low     | High    | 37.7          | 39.5        | 95.4     | 2.1         | Cream       | Poly 11.0     |
| Indonesia-88076 | 52.0                 | 34.6                  | 5.3              | 3.1             | Yellow       | Smooth       | Yellow     | None    | None        | Difficult | None       | Low     | High    | 37.4          | 36.6        | 102.2    | 2.1         | Cream       | Poly 12.4     |
ous genotypes were at least 0.63–0.74%. Because of the hybrid origin of lime and the ability of limes to asexually reproduce, the heterozygosity of limes was maintained. The Bhutanese cluster contained more heterozygous genotypes than the Indonesian cluster. In the Indonesian cluster, Indonesian subcluster 1 contained more heterozygous genotypes than Indonesian subcluster 2.

We further classified the 42,613 sites into categories (i.e., Categories A–L) based on unique genotype combinations (Table 4; Supplementary Table S3). In Category A, we identified 7297/42,613 sites that contained unique heterozygous genotypes common to all eight accessions. In other words, M (A or C), R (A or G), W (A or T), S (C or G), Y (C or T), or K (G or T) were present in all eight accessions (Table 4; Supplementary Table S3).

In Category B, we identified 4026/42,613 sites that contained unique homozygous genotypes common to the Bhutanese cluster; other unique homozygous genotypes were common to the Indonesian cluster. For example, A and C were common to the Bhutanese and Indonesian clusters, respectively (Table 4; Supplementary Table S3). The classification of the eight accessions into two clusters may be primarily explained by the 4026 sites with unique homozygous genotypes.

In Category C, we identified 14,445/42,613 sites that contained unique heterozygous genotypes common to the Bhutanese cluster and unique homozygous genotypes that were common to the Indonesian cluster. For example, M and A were common to the Bhutanese and Indonesian clusters, respectively (Table 4; Supplementary Table S3). In comparison, we identified 10,912/42,613 sites in Category D that contained unique homozygous genotypes common to the Bhutanese cluster and unique heterozygous genotypes that were common to the Indonesian cluster. For example,
tions in the ancestor of each cluster may explain the observed changes of the homozygous genotypes into heterozygous genotypes. In Category E, we identified 1432/42,613 sites that contained unique heterozygous genotypes common to the Bhutanese cluster; identical heterozygous genotypes and/or homozygous genotypes derived from the heterozygous genotypes were common to the Indonesian cluster. For example, M was common to the Bhutanese cluster, while M and A were present in the Indonesian cluster (Table 4; Supplementary Table S3; Supplementary Fig. S3). We counted genotype matches between each accession in the Indonesian cluster (Table 5). The analysis showed that, in many cases, the genotypes found in Indonesian subcluster 1 were identical to each other, with a similar result being obtained for Indonesian subcluster 2. This result may explain why the phylogenetic analysis classified the Indonesian cluster into two subclusters. As shown in Table 6, among the 1432 sites classified into Category E, 755 sites contained heterozygous genotypes that were common to Indonesian subcluster 1 and unique homozygous genotypes that were common to Indonesian subcluster 2. For example, M and A were common to the Indonesian subclusters 1 and 2, respectively. Moreover, among the 1432 sites, 271 sites were contained unique homozygous genotypes that were common to Indonesian subcluster 1 and unique heterozygous genotypes that were unique to Indonesian subcluster 2. For example, A and M were common to Indonesian subclusters 1 and 2, respectively. Thus, the number of heterozygous genotypes in Indonesian subcluster 1 was approximately three-fold larger than that of Indonesian subcluster 2 in Category E. Because identical heterozygous genotypes are present in the Bhutanese cluster, these differences may be explained by the heterozygous genotypes changing into homozygous genotypes. Homologous recombination in the ancestors of the Indonesian cluster may, to some extent, explain the observed differences within the Indonesian cluster. Furthermore, these differences may, to some extent, be explained by the loss of the EcoRI sites (the emergence of false-positive homozygous genotypes).

In Category F, we identified 134/42,613 sites that contained unique heterozygous genotypes common to the Indonesian cluster; identical heterozygous genotypes and/or homozygous genotypes derived from the heterozygous genotypes were common to the Bhutanese cluster. For example, M and A were present in the Bhutanese cluster, while M was common to the Indonesian cluster (Table 4; Supplementary Table S3; Supplementary Fig. S4). Genotype matches between each accession in the Bhutanese cluster (Supplementary Table S4) had a narrow range (50–78 matches; 1.6-fold differences). This phenomenon may explain why the phylogenetic analysis did not classify the Bhutanese cluster into two subclusters. Because identical heterozygous genotypes are present in the Indonesian cluster, these differences may be explained by the heterozygous genotypes changing into homozygous genotypes. Like in Category E, both the homologous recombination in the ancestors of the Bhutanese cluster and the loss of EcoRI sites may explain the observed differences within the Bhutanese cluster.

In Category G, we identified 3087/42,613 sites that contained unique homozygous genotypes common to the Bhutanese cluster; unique heterozygous and homozygous genotypes derived from the heterozygous genotypes were present in the Indonesian cluster. For example, A was common to the Bhutanese cluster, while M and A were present in the Indonesian cluster (Table 4; Supplementary Table S3; Supplementary Fig. S5). We counted genotype matches between each accession belonging to the Indonesian cluster (Table 7). The analysis showed that, in many cases, genotypes found in Indonesian subcluster 1 were identical to each other, with the same result being obtained for Indonesian subcluster 2. This is the reason why the phylogenetic analysis classified the Indonesian cluster into two subclusters. As shown in Table 8, among the 3087 sites in Category G, 1573 sites contained heterozygous genotypes that were common to Indonesian subcluster 1 and unique homozygous genotypes that were common to Indonesian subcluster 2. Moreover, among the 3087 sites, 613 sites contained homozygous genotypes that were common to Indonesian subcluster 1 and unique heterozygous genotypes.

### Table 2 | Leaf characteristics of the nucellar seedlings of the five Bhutanese and three Indonesian lime accessions

| Accession     | Leaf blade | Wing | Wing index
|---------------|------------|------|-------------
|               | length (mm) | width (mm) | Area* (mm²) | length (mm) | width (mm) | Area* (mm²) |
| Bhutan-09015  | 51.1 ± 2.3 b | 26.2 ± 0.8 ab | 335.7 ± 24.0 ab | 5.7 ± 0.5 a | 1.4 ± 0.1 a | 2.01 ± 0.15 a |
| Bhutan-09024  | 66.1 ± 1.7 cd | 31.5 ± 1.1 bc | 522.1 ± 26.6 bc | 6.4 ± 1.0 a | 1.9 ± 0.1 ab | 3.09 ± 0.62 ab |
| Bhutan-09027  | 62.1 ± 2.30 bc | 32.2 ± 0.5 bc | 501.5 ± 22.3 bc | 6.7 ± 4.1 ab | 1.8 ± 0.1 ab | 3.11 ± 0.53 ab |
| Bhutan-09030  | 76.2 ± 5.0 d | 33.6 ± 1.9 bc | 687.6 ± 81.8 c | 5.8 ± 0.4 a | 1.9 ± 0.1 ab | 2.84 ± 0.33 ab |
| Bhutan-09005  | 36.7 ± 1.3 a | 22.4 ± 1.1 a | 207.0 ± 15.3 a | 9.2 ± 0.7 abc | 2.9 ± 0.4 bc | 6.83 ± 1.46 bc |
| Indonesia-88035 | 64.6 ± 1.9 cd | 37.4 ± 6.3 cd | 605.3 ± 34.2 cd | 13.7 ± 0.5 d | 5.1 ± 0.5 d | 17.70 ± 1.47 d |
| Indonesia-88045 | 64.5 ± 2.2 cd | 40.1 ± 2.5 d | 651.8 ± 34.2 c | 11.8 ± 1.5 cd | 3.7 ± 0.2 c | 10.76 ± 1.22 c |
| Indonesia-88065 | 47.5 ± 2.0 ab | 26.3 ± 0.9 ab | 313.3 ± 20.7 a | 11.0 ± 1.5 b | 3.1 ± 0.1 c | 8.48 ± 1.33 c |

*Leaf blade and wing area were estimated using the formula for calculating the ellipse area.

### Table 3 | Summary of the RAD-seq analysis

| Accession     | Number of sequencing reads after quality filtering | Minimum number of heterozygous genotypes estimated from 42,613 sites | Minimum frequency of heterozygous genotypes estimated from 42,613 sites |
|---------------|---------------------------------------------|-----------------------------------------------------------------|-------------------------------------------------------------------|
| Bhutan-09015  | 78,420,728                                  | 23,985                                                           | 0.74%                                                             |
| Bhutan-09024  | 67,963,194                                  | 23,902                                                           | 0.74%                                                             |
| Bhutan-09027  | 36,677,458                                  | 23,915                                                           | 0.74%                                                             |
| Bhutan-09030  | 45,902,996                                  | 23,960                                                           | 0.74%                                                             |
| Bhutan-09005  | 36,682,650                                  | 21,252                                                           | 0.68%                                                             |
| Indonesia-88035 | 24,485,978                              | 20,184                                                           | 0.63%                                                             |
| Indonesia-88045 | 48,889,661                                | 20,183                                                           | 0.63%                                                             |
| Indonesia-88065 | 28,103,773                                | 21,751                                                           | 0.68%                                                             |
zygous genotypes that were common to Indonesian subcluster 2. The number of heterozygous genotypes in Indonesian subcluster 1 was approximately three-fold larger than that of Indonesian subcluster 2 in Category G. When the Bhutanese cluster carried unique homozygous genotypes (e.g., A genotypes), changes from unique heterozygous genotypes (e.g., M genotypes) to homozygous genotypes identical to the Bhutanese cluster (e.g., A genotypes) occurred more frequently compared to those from heterozygous genotypes (e.g., M) to other homozygous genotypes (e.g., C genotypes) in the Indonesian cluster (2439 events vs. 609 events; estimated from Supplementary Table S3). Thus, the two types of changes did not arise equally (4-fold differences). Probably, the differences within the Indonesian cluster are not explained only by heterozygous genotypes changing into homozygous genotypes. Both heterozygous changes into homozygous genotypes and homozygous changes into heterozygous genotypes potentially contribute these differences. The former change may arise through homologous recombination and the loss of EcoRI sites, while the latter change may arise through de novo mutation.

In Category H, we identified 597/42,613 sites that contained unique homozygous genotypes common to the Indonesian cluster; unique heterozygous and homozygous genotypes derived from the heterozygous genotypes were present in the Bhutanese cluster. For

Table 4 | Summary of the analysis of the 42,613 sites genotyped in the current study

| Category | Number of sites |
|----------|----------------|
| Category A. | 7,297 |
| Unique heterozygous genotypes are common to all eight accessions. |
| Category B. | 4,026 |
| Unique homozygous genotypes are common to the Bhutanese cluster. |
| Category C. | 14,445 |
| Other unique homozygous genotypes are common to the Indonesian cluster. |
| Category D. | 10,912 |
| Unique heterozygous genotypes are common to the Bhutanese cluster. |
| Category E. | 1,432 |
| Unique homozygous genotypes are common to the Bhutanese cluster. |
| Category F. | 134 |
| Unique heterozygous genotypes are common to the Bhutanese cluster. |
| Category G. | 3,087 |
| Identical heterozygous genotypes and/or homozygous genotypes derived from the heterozygous genotypes are present in the Indonesian cluster. |
| Category H. | 597 |
| Unique homozygous genotypes are common to the Bhutanese cluster. |
| Category I. | 242 |
| Unique heterozygous genotypes and homozygous genotypes derived from the heterozygous genotypes are present in the Bhutanese cluster. |
| Category J. | 242 |
| Heterozygous and homozygous genotypes are distributed among eight accessions. |
| Category K. | 2 |
| Unique homozygous genotypes are common to the Bhutanese cluster. |
| Category L. | 197 |
| Different kinds of heterozygous genotypes are distributed among eight accessions. |
| Total number of genotypes. | 42,613 |
example, M and A were present in the Bhutanese cluster, while A was common to the Indonesian cluster (Table 4; Supplementary Table S3; Supplementary Fig. S6). Genotype matches between each accession belonging to the Bhutanese cluster (Supplementary Table S5) had a narrow range (213–342 matches; 1.6-fold differences). This phenomenon is the reason why the phylogenetic analysis did not classify the Bhutanese cluster into two subclusters. When the Indonesian cluster carried the unique homozygous genotypes (e.g., A), the number of changes from the unique heterozygous genotypes (e.g., M) to the homozygous genotypes in the Bhutanese cluster that were identical to the Indonesian cluster (e.g., A) was 385. In contrast, the number of changes from the heterozygous genotypes (e.g., M) to the other homozygous genotypes (e.g., C) was 204. Thus, fewer differences in the occurrence of the two types of changes were observed (about 2-fold difference) when compared to Category G. However, like Category G, both changes from heterozygous to homozygous genotypes and changes from homozygous to heterozygous genotypes might contribute to these differences.

We were able to roughly estimate the contributions of the loss of EcoRI sites (the emergence of false-positive homozygous genotypes) to genetic diversification within the cluster. For example, we considered the differences within the Indonesian cluster, in which more frequent changes occurred (Category E and G). Some of the changes in Category G may be explained by de novo mutation. These de novo mutations occurred in the 36 bp sequences adjacent to the EcoRI site. If we assume that the 3087 events observed in Category G were caused by de novo mutation only, the possible number of de novo mutations of the EcoRI sites (6 bp sequence) is 515 (= 3087/6). We observed 1432 events in Category E, which was about 3-fold larger.

| Table 5 | Number of genotype matches between each accession of the Indonesian cluster in Category E |
|---------|-------------------------------------------------------------|
|        | Bhutan-09005 | Indonesia-88035 | Indonesia-88045 |
| Indonesia-88035 | 163 |          |
| Indonesia-88045 | 144 | 1215 |
| Indonesia-88065 | 1270 | 180 | 194 |

| Table 7 | Number of genotype matches between each accession of the Indonesian cluster in Category G |
|---------|-------------------------------------------------------------|
|        | Bhutan-09005 | Indonesia-88035 | Indonesia-88045 |
| Indonesia-88035 | 437 |          |
| Indonesia-88045 | 313 | 2595 |
| Indonesia-88065 | 2575 | 508 | 464 |

| Table 6 | Comparison of genotypes among the Bhutanese cluster and the two Indonesian subclusters in Category E |
|---------|-------------------------------------------------------------|
| Bhutanese cluster | Indonesian subcluster 1 | Indonesian subcluster 2 | Number of the groupings |
| M | M | A | 26 |
| M | M | C | 34 |
| R | R | A | 120 |
| R | R | G | 110 |
| W | W | A | 45 |
| W | W | T | 57 |
| S | S | C | 37 |
| S | S | G | 29 |
| Y | Y | C | 100 |
| Y | Y | T | 105 |
| K | K | G | 44 |
| K | K | T | 48 |
| Total | 755 |

| BHUTANES.C CLUSTER | INDONESIAN SUBCLUSTER 1 | INDONESIAN SUBCLUSTER 2 | Number of the groupings |
|---------------------|-------------------------|-------------------------|-------------------------|
| A | M | A | 80 |
| A | M | C | 9 |
| C | M | A | 11 |
| C | M | C | 70 |
| A | R | A | 185 |
| A | R | G | 35 |
| G | R | A | 33 |
| G | R | G | 217 |
| A | W | A | 94 |
| A | W | T | 17 |
| T | W | A | 12 |
| T | W | T | 96 |
| C | S | C | 49 |
| C | S | G | 7 |
| G | S | G | 39 |
| G | S | C | 195 |
| G | Y | C | 29 |
| G | Y | T | 35 |
| T | Y | C | 192 |
| T | Y | T | 74 |
| G | K | G | 12 |
| T | K | G | 12 |
| Total | 1573 |

| Table 8 | Comparison of the genotypes among the Bhutanese cluster and the two Indonesian subclusters in Category G |
|---------|-------------------------------------------------------------|
| Bhutanese cluster | Indonesian subcluster 1 | Indonesian subcluster 2 | Number of the groupings |
| A | M | A | 80 |
| A | M | C | 9 |
| C | M | A | 11 |
| C | M | C | 70 |
| A | R | A | 185 |
| A | R | G | 35 |
| G | R | A | 33 |
| G | R | G | 217 |
| A | W | A | 94 |
| A | W | T | 17 |
| T | W | A | 12 |
| T | W | T | 96 |
| C | S | C | 49 |
| C | S | G | 7 |
| G | S | G | 39 |
| G | S | C | 195 |
| G | Y | C | 29 |
| G | Y | T | 35 |
| T | Y | C | 192 |
| T | Y | T | 74 |
| G | K | G | 12 |
| T | K | G | 12 |
| Total | 1573 |
than the hypothesized number. Thus, both homologous recombination and the loss of EcoRI sites may contribute the changes from heterozygous to homozygous genotypes.

In Category I, we identified 242/42,613 sites that contained heterozygous and homozygous genotypes that were distributed randomly among the eight accessions (Table 4; Supplementary Table S3: Supplementary Fig. 7). Some experimental or computational errors may have led to the identification of these sites. Genotype matches between each accession (Supplementary Table S6) had a relatively wide range (53–179 matches; 3.4-fold differences), but it was difficult to extract any clear trends from this analysis.

In Category J, we identified 242/42,613 sites that contained unique homozygous genotypes common to the Bhutanese cluster, and identical homozygous genotypes and other homozygous genotypes that were present in the Indonesian cluster (Table 4; Supplementary Table 3, Supplementary Fig. 8). We counted genotype matches between each accession belonging to the Indonesian cluster (Table 9). In most cases, genotypes found in Indonesian subcluster 1 were identical to each other, with the same result being obtained for Indonesian subcluster 2, which contributed to the classification of the two Indonesian subclusters.

In Category K, we identified 2/42,613 sites that contained two types of homozygous genotypes present in the Bhutanese cluster and unique homozygous genotypes that were common to the Indonesian cluster (Table 4; Supplementary Table S3; Supplementary Fig. S9). In Category L, we identified 197/42,613 sites that contained different types of heterozygous genotypes that were distributed among the eight accessions (Table 4; Supplementary Table S3; Supplementary Fig. S10). It was difficult to obtain useful information from these data.

### Discussion

Our analysis showed that none of the lime accessions assessed in the current study were fully homozygous, which may be explained by the hybrid origin of this species, as previously proposed4–6. As shown in Table 1, the seeds of these accessions are polyembryonic, which enables limes to maintain a heterozygous state by reproducing asexually. Thus, these observed heterozygositities support the proposed hybrid origin of limes and the ability of limes to reproduce asexually.

The genotypes shown in Category C and D mainly contributed the differences between the Bhutanese and Indonesian clusters (14,445 and 10,912 events, respectively). One cluster carried heterozygous genotypes, while the other cluster carried homozygous genotypes. These differences may arise through several mechanisms, including homologous recombination, the loss of EcoRI sites (the emergence of false-positive homozygous genotypes), and de novo mutations in the ancestor of each cluster. Sexual reproduction might contribute to homologous recombination in limes, which is one of the proposed mechanisms. Indeed, the lime has self-pollinating properties77. A previous study showed that both sexual and asexual reproduction in limes cultivated in Tunisia contributed to their genetic diversity78. However, further study is required to elucidate how sexual reproduction contributes to the genetic diversification of limes.

Our analyses separated the eight selected accessions into two clusters. In addition, the Indonesian cluster was further grouped into two subclusters. Thus, our analysis suggests that the Bhutanese cluster, and the Indonesian subclusters 1 and 2 may be independently selected. Because humans cultivate these eight accessions, humans may have played a role in these selection events. The ancestors of each cluster or subcluster may possess features that were beneficial for humans, which were subsequently selected for in subsequent generations. Alternatively, if humans did not select the ancestors of each cluster or subcluster, the ancestors may have acquired advantageous characteristics via natural selection.

Unexpectedly, the Indonesian cluster was distributed throughout both Indonesia and Bhutan. However, the origins of the Indonesian cluster are not clear. Bhutan-09005 is cultivated at Tsirang in southern Bhutan. This district is close to India, with the two countries having an active trading history. Therefore, the ancestor(s) of this tree may have had more opportunities to become distributed throughout various regions, unlike the four accessions of the Bhutanese cluster.

We showed that there were minimal differences among the genotypes within the Bhutanese cluster and within each Indonesian subcluster. Because 20,183 to 23,985 heterozygous genotypes were estimated from 42,613 sites (Table 3), crosses within the Bhutanese cluster or within each Indonesian subcluster are expected to result in the production of numerous genetic differences. Self-pollination would also result in the production of numerous genetic differences. It is possible that each cluster/subcluster was derived from a single tree by vegetative propagation. However, differences among the genotypes within the Bhutanese cluster or within each Indonesian subcluster remain, although some of these differences may have been generated from experimental or computational errors. During asexual reproduction, changes within a cluster or subcluster may have occurred. Some of the differences among the genotypes within the Bhutanese cluster or the two Indonesian subclusters (many events observed in Category E and F, and some events observed in Category G and H) may be explained by homologous recombination. Therefore, these observations strongly suggest that somatic homologous recombination, possibly by the mechanism of double strand break repair, may have contributed to the genetic diversification of limes. Furthermore, as observed in Category G and H, de novo mutation may have also contributed to this phenomenon.

Furthermore, small differences were obtained among the genotypes within the Indonesian cluster, namely those between each subcluster. Simple crosses within one subcluster are expected to result in the production of numerous genetic differences. The possibility of producing other types of subclusters by crossing is very low. Some differences among the genotypes between each subcluster (many events observed in Category E, and some events observed in Category G) may be explained by homologous recombination. Therefore, these observations suggest that somatic homologous recombination at larger regions may also contribute to genetic diversification within the Indonesian cluster. Furthermore, as observed in Category G, de novo mutation may have also contributed to this phenomenon.

Although we revealed the reproductive history of limes at the genomic level, the current study only analyzed a small number of accessions from just two countries. Therefore, larger sample numbers are required to elucidate the full evolutionary history of limes. Our analysis did not provide any information about the parental generation of the eight accessions. Analysis of other related Citrus species is required to address this question.

In conclusion, using high-throughput sequencing, we partially analyzed intraspecies relationships between limes. To the best of our knowledge, this is the first report about the characteristics of Citrus species using RAD-seq. Our analysis suggests part of the evolutionary history of limes and may link it to the past activities of humans.

### Methods

**Plant materials.** Preliminary characterization was conducted by investigating 14 local lime accessions in Bhutan between 2009 and 2011 (as part of a collaborative study on exploration and identification of wild citrus relatives in Bhutan). Leaf and fruit characteristics were measured just after sampling in the field at each location or purchasing at the market. The characteristics of four lime accessions, which were introduced as budwoods to Saga University, Japan, from Indonesia, were also

### Table 9

| genotype matches between each accession of the Indonesian cluster in Category J | Bhutan-09005 | Indonesia-88035 | Indonesia-88045 |
|---|---|---|---|
| Indonesia-88035 | 16 | | |
| Indonesia-88045 | 10 | 229 | |
| Indonesia-88065 | 225 | 13 | 7 |
measured. We classified the accessions based on the characteristics of the fruit surfaces and the wing of the leaves, because these characteristics are minimally influenced by environmental or growth conditions.

Quantitative characterization of the leaf wings was conducted. Specifically, the seeds were extracted from the fruit of five Bhutanese accessions that were collected (i.e., Bhutan-09005, Bhutan-09015, Bhutan-09024, Bhutan-09027, and Bhutan-09030). The seedlings implanted from these seeds were grown in a greenhouse. Nucellar seedlings and three Indian accessions (i.e., Indonesia-88035, Indonesia-88045, and Indonesia-88065) was investigated. Five leaves from one tree were used as the material for each accession. The length and width of the leaf blade and wing were measured. Leaf blade area and wing area were estimated using the formula for calculating the ellipse area: area = \( \pi \times \text{length} \times \frac{\text{width}^2}{2} \). The wing index was expressed as: (area of the wing) \times 100/(area of the leaf blade). For subsequent DNA analysis, the leaves collected from the five Bhutanese and three Indonesian accessions were used as the material.

**RAD-seq analysis.** Genomic DNA was extracted from the leaves using the DNeasy Plant Mini Kit (Qiagen, Valencia, CA, USA), with some modifications (i.e., the addition of 8 \( \mu \)l \( \beta \)-mercaptoethanol and 4 mg polyvinylpyrrolidone to 400 \( \mu \)l Buffer AP1). The quality of the isolated genomic DNA was checked by 1% agarose gel electrophoresis. EcoRI RAD-seq analysis was commissioned to BGI (Shenzhen, China) where the data were generated using HiSeq2000. At BGI, the raw data were modified using the following two steps: (1) the reads that were polluted by adapter sequences were deleted and (2) the reads were filtered to the low-quality bases (quality value \( <5 \)) or \( >10 \) N bases were removed. Each read received from BGI was a 41-bp single-end sequence [AATT (EcoRI site) \( +36 \) bp sequence]. Sequences are available at the DDBJ Sequence Read Archive (http://trace.ddbj.nig.ac.jp/dra/index_e.shtml; Accession no. DRA000989).

**Sequence analysis.** The data from BGI were analyzed using the Stacks software pipeline (version 1.09)[4][5]. The data were further quality-filtered by a process_shortreads program of Stacks (with the -c -q options). The reference genome of 8 \( \mu \)l low-quality bases (quality value \( <5 \)) or \( >10 \) N bases were removed. Each read received from BGI was a 41-bp single-end sequence [AATT (EcoRI site) \( +36 \) bp sequence]. Sequences are available at the DDBJ Sequence Read Archive (http://trace.ddbj.nig.ac.jp/dra/index_e.shtml; Accession no. DRA000989).

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Author contributions

T.P., R.M., M.Y., Y.N. conceived and designed the experiments. T.P., T.M., R.M., M.Y., Y.N. performed the experiments. T.P., R.M., M.Y., Y.N. analyzed the data. T.P., M.Y., Y.N. wrote the paper. All authors read and approved the final manuscript.

Additional information

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