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Submitted on 17 Jul 2013

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Genetic diversity and population structure of an insular tree, *Santalum austrocaledonicum* in New Caledonian archipelago

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Abstract

We present a study of the genetic diversity and structure of a tropical tree in an insular system. *Santalum austrocaledonicum* is endemic to the archipelago of New Caledonia and is exploited for oil extraction from heartwood. A total of 431 individuals over 17 populations were analysed for eight polymorphic microsatellite loci. The number of alleles per locus ranged from 3 to 33 and the observed heterozygosity per population ranged from 0.01 in Maré to 0.74 in Ile des Pins. The genetic diversity was lowest in the most recent islands, the Loyautés, and highest in the oldest island, Grande Terre, as well as the nearby small Ile des Pins. Significant departures from panmixia were observed for some loci–population combinations (per population $F_{IS} = 0–0.03$ on Grande-Terre and Ile des Pins, and 0–0.67 on Loyautés). A strong genetic differentiation among all islands was observed ($F_{ST} = 0.22$), and the amount of differentiation increased with geographic distance in Iles Loyauté and in Grande Terre. At both population and island levels, island age and isolation seem to be the main factors influencing the amount of genetic diversity. In particular, populations from recent islands had large average $F_{IS}$ that could not be entirely explained by null alleles or a Wahlund effect. This result suggests that, at least in some populations, selfing occurred extensively. Conclusively, our results indicate a strong influence of insularity on the genetic diversity and structure of *Santalum austrocaledonicum*.

Keywords: conservation, gene flow, insularity, nuclear microsatellites, population genetic structure, sandalwood

Received 28 September 2004; revision received 12 January 2005; accepted 8 March 2005

Introduction

Island systems have long fascinated biologists, in particular since Darwin’s theory of evolution by natural selection led to consider them as ‘evolutionary laboratories’ (Darwin 1859). Islands are strongly attractive environments for studying evolution for various reasons: they present discrete entities; despite their small size, they contain a variety of habitats; and they are often geologically dynamic (Emerson 2002). From a genetic point of view, at the within-island level, populations have been characterized as depauperate, because of possible recency of the founding event, isolation from source population, and stochastic processes due to their limited size (Carlquist 1980; Crawford *et al*. 1987, 1988, 1990; Brauner *et al*. 1992; Elisens 1992; Kwon & Morden 2002). At the among-island level, the presence of oceanic barriers restricts gene flow between populations. Hence populations from different islands are expected to be strongly differentiated, at least at neutral loci.

Although general expectations exist on the impact of island systems on genetic diversity structuring, especially those concerning natural selection and random drift (Barton 1989), empirical evidence is still lacking to confirm or attenuate the general expected pattern on plant species. In particular, the genetic structure of forest tree species is poorly documented in fragmented habitats (Savolainen
and have been found in some systems (Young & Clarke 2000). In most cases, a loss of genetic variability and increased genetic differentiation of subpopulations through drift are predicted and have been found in some systems (Young et al. 1996; Newman & Pilson 1997; Young & Clarke 2000). Hamrick et al. (1992) and Hamrick & Godt (1996) studied the genetic diversity in continental forest tree species and underlined the factors influencing differentiation between populations (mating system, life history, distribution area, etc.), but did not consider the influence of insularity (e.g. isolation and size of populations).

In this study we address a number of questions related to the genetic diversity patterns of an insular forest tree species, Santalum austrocaledonicum (sandalwood) in the archipelago of New Caledonia. Santalum austrocaledonicum is, like all Santalaceae (Malécot et al. 2004) a hemiparasitic forest tree species (Nasi & Ehrhart 1996) endemic to New Caledonia and Vanuatu. It reaches 8 m in height and 30 cm in trunk diameter. It grows in lowlands, preferentially in open areas (Quémin 1988; Ehrhart 1998). As a hemiparasitic plant species, it cannot grow out of the vicinity of other species, particularly nitrogen-fixing species like Acacia and Casuarina (Quémin 1988; Radomiljac & McComb 1997; Ehrhart 1998). It occurs on all islands of the archipelago at altitudes lower than 200 m (Quémin 1988) as isolated trees or patches of trees of various sizes. Sandalwood reproduction is still poorly documented. The seeds are fleshy and small (about 1 cm) and seem to be disseminated by frugivorous birds (Columba vitiensis was seen eating fruits) that can travel among islands (Gibbs et al. 2001), and potentially bats (Cox et al. 1991). Santalum austrocaledonicum has been one of the most exploited sandalwood species since the 19th century. Essential oils extracted from heartwood are used in medicine and the perfume industry. Still exploited, some populations, particularly those of the Iles Loyauté, are seriously threatened.

Using eight microsatellite markers specifically developed for the species studied, we compared the pattern of genetic diversity among islands of the archipelago. We then tried to relate the amount of genetic diversity to the size and isolation of islands, and to look for isolation-by-distance patterns among and within islands. We also asked whether there is any evidence for an impact of the last maximum glaciations on the structure of the genetic diversity. The New Caledonia archipelago is a good system to address these questions, as it consists of six islands of various sizes and at various distances from the largest and oldest of them, Grande-Terre. Each island is large enough that it can itself be subdivided into a few geographic regions.

Material and methods

Sampling methodology, DNA extraction and genetic analysis

Leaves of Santalum austrocaledonicum were collected on individuals growing in natural stands throughout the different islands: Grande Terre (island size: 16 350 km²), Ile des Pins (152 km²), Iles Loyauté (Ouvéa (132 km²), Lifou (1196 km²), Maré (650 km²)) (Fig. 1). Population sampling areas are shown in Table 1. On Grande Terre, the largest island, all known populations were sampled. These populations were far from each other (at least 25 km) and had a small number of individuals so that the sampling was exhaustive. Ouen Toro was an exception as it was densely populated, so in this population individuals sampled were only a subset of the total population. Only populations with more than 10 individuals were conserved for the genetic analysis: Pindaï, Malhec, Paita, Hienghène, Tiéa and Ouen Toro. Hienghène was the only population known on the east coast, where the climate is wetter and the vegetation more luxurious than on the west coast.

On other islands, the situation was quite different: individuals were scattered over the whole area, so like in Ouen Toro, the samples, collected everywhere in the islands, represented a subset of the total population. In order to have population areas more comparable with those of Grande Terre, and to avoid a potential effect of substructuring on the structure indices, we tried to determine subpopulations in these islands according to the spatial distribution of individuals. Samples from Loyauté islands were divided in three populations: north (N), midlands (M) and south (S) for Ouvéa and Lifou, north (N), southeast (SE) and southwest (SW) for Maré. Ile des Pins was subdivided into two areas, north (N) and south (S).

Leaves were collected between February 1998 and November 2003, but each individual, identified by its geographical coordinates, was only sampled once. Leaf specimens were placed in sealed plastic bags containing silica gel, until DNA extraction. Total DNA was extracted from 100 mg of dry leaf material using a MATAB method derived from Bousquet et al. (1990), with one additional chloroform–isoamyl alcohol (24:1) extraction. The genetic analysis was done using eight nuclear microsatellites: mSaCIRE09, mSaCIRH09, mSaCIRG01, mSaCIRH11, mSaCIRG10, mSaCIRF04, mSaCIRF10 and mSaCIRH10, designed specifically for Santalum austrocaledonicum. Their characteristics and the methods used to obtain them are described elsewhere (Bottin et al. in press).

Selection of individuals for the analysis

As sandalwood can reproduce by suckering (Quémin 1988), sampling of several individuals in a restricted area
runs the risk of collecting the same genetic individuals. In order to avoid this problem we excluded from the following analyses all individuals (but one) with the same genotypes that were less than 100 m apart. We first identified similar genotypes by constructing neighbour-joining (NJ) trees for each population with Darwin 4-4 (Perrier & Jaquemoud-Collet 2003). We then examined their localization on geographical maps where we had placed all the individuals using MapInfo software. Out of 541 individuals, about 100 were eliminated this way, suggesting that suckering occurs frequently. The total number of individuals analysed per population is presented in Table 1.

Analysis of genetic diversity and departure from random mating

Allele frequencies, observed number of alleles per locus \((A)\), observed heterozygosity \((H_O)\) and expected heterozygosity \((H_E)\) (Nei 1978) per population were computed with Genetix 4.03 (Belkhir et al. 2001).

The inbreeding coefficient \((F_{IS})\) was estimated for each population and departure from Hardy–Weinberg equilibrium was assessed also using Genetix 4.03 by a permutation test with 5000 permutations. \(P\) values were corrected using sequential Bonferroni procedure (Rice 1989).

To check if the differences in sample sizes and the various spatial scales over which individuals were pooled into ‘populations’ affected the diversity estimates, we calculated the allelic richness per population and island taking into account the dependence on sample size with an adaptation of the rarefaction index of Hurlbert (1971) (El Mousadik & Petit 1996), named ‘\(R\)’, using FSTAT 2.9.3.2 (Goudet 1995). The principle is to estimate the expected number of alleles in a subsample of \(2n\) genes, given that \(2N\) genes have been sampled \((N > n)\). In FSTAT, \(n\) is fixed as the smallest number of individuals typed for a locus in a sample.

Analyses of population differentiation

Differentiation among all samples and all sample pairs was tested using probability tests (Fisher exact tests), as described by Raymond & Rouset (1995). Wright’s
F-statistics $F_{ST}$ (Wright 1951) were estimated for all populations and all population pairs by a ‘weighted’ analysis of variance (Cockerham 1973; Weir & Cockerham 1984) with GENEPOP. To investigate the genetic structure of populations, we ran an analysis of molecular variance using ARLEQUIN (Schneider et al. 2000) with 1000 permutations (AMOVA, Excoffier et al. 1992) which tests a particular genetic structure by partitioning the total variance into covariance components due to intrapopulation differences, interindividual differences, and/or interpopulation differences. We tested two kinds of structure: among island and among populations within each island. We also tested the differentiation between the ‘Grande Terre and Ile des Pins’ group and the ‘Ile Loyauté’ group.

To relate the dispersal ability of *S. austrocaledonicum* to its geographical distribution, we ran Mantel tests (ManTEL 1967) implemented in GENEPOP 3.4. The procedure assesses the significance of the correlation between pairwise $F_{ST}/(1 − F_{ST})$ estimates and the logarithm of the Euclidian distance (in kilometres) between pairs of localities (Rousset 1997), with the Spearman rank coefficient as statistical test, using 5000 random permutations of the matrix. Two tests were used: between populations in Grande Terre and Ile des Pins, and between populations of Iles Loyauté.

Pairwise genetic distances between pairs of populations were computed using Cavalli-Sforza’s chord measure (Cavalli-Sforza & Edwards 1967), obtained from the GENDIST program (PHYLIP, version 3.6, Felsenstein 1989, 1993). The distance tree was constructed using the NJ method of Saitou & Nei (1987) using the NEIGHBOR program of PHYLIP. The robustness of each node was evaluated by bootstrapping data over locus for 1000 replications using the SEQBOOT program of PHYLIP 3.6 and the consensus tree obtained by SEQBOOT (PHYLIP 3.6) was displayed with TREEVIEW software (Page 1996).

**Results**

**Within-population genetic diversity and departure from random mating**

The eight microsatellite loci were polymorphic and the number of alleles per locus ranged from 3 for *mSaCIRG01* to 33 for *mSaCIRG10*. Mean numbers of alleles per locus per population ($\mu$) ranged from 1.38 in Maré South-east (allelic richness or $R$ in this population: 1.19) to 7.88 in Ile des Pins North ($R$: 5.14). Forty-two alleles were private, the largest number in one population being 9 in Ile des Pins North. The Pearson correlation coefficient between the number of alleles per locus ($\mu$) and the allelic richness corrected with a rarefaction index ($R$) (Table 2) was 0.96 and was highly significant (Spearman correlation coefficient: 0.935***), which demonstrated a strong relationship between these two parameters.

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**Table 1** Characteristics of the populations of *Santalum austrocaledonicum* in the archipelago of New Caledonia: number of individuals sampled (no. of individuals), sampling area (in Grande Terre individuals are aggregated in populations, whereas they are quite isolated in Loyauté islands), and geographical coordinates of the populations

<table>
<thead>
<tr>
<th>Population</th>
<th>No. of individuals</th>
<th>Sampling area [island size] (km$^2$)</th>
<th>Latitude</th>
<th>Longitude</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ouen Toro</td>
<td>22</td>
<td>0.19</td>
<td>22°30'</td>
<td>166°45</td>
</tr>
<tr>
<td>Pindai</td>
<td>26</td>
<td>3.43</td>
<td>21°31'</td>
<td>164°96</td>
</tr>
<tr>
<td>Malek</td>
<td>37</td>
<td>0.87</td>
<td>20°19'</td>
<td>164°10</td>
</tr>
<tr>
<td>Paita</td>
<td>53</td>
<td>0.67</td>
<td>22°09'</td>
<td>166°22</td>
</tr>
<tr>
<td>Hienghène</td>
<td>20</td>
<td>0.57</td>
<td>20°43'</td>
<td>164°55</td>
</tr>
<tr>
<td>Tiéa</td>
<td>10</td>
<td>0.05</td>
<td>21°08'</td>
<td>164°56</td>
</tr>
<tr>
<td><strong>Grande Terre</strong></td>
<td><strong>168</strong></td>
<td><strong>5.8 [16 350]</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>IP north</td>
<td>26</td>
<td>32.25</td>
<td>22°35'</td>
<td>167°28</td>
</tr>
<tr>
<td>IP south</td>
<td>35</td>
<td>25.25</td>
<td>22°39'</td>
<td>167°28</td>
</tr>
<tr>
<td><strong>Ile des Pins</strong></td>
<td><strong>61</strong></td>
<td><strong>57.5 [152]</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Lifou north</td>
<td>36</td>
<td>275.28</td>
<td>20°42'</td>
<td>167°13</td>
</tr>
<tr>
<td>Lifou middle</td>
<td>42</td>
<td>271.81</td>
<td>20°58'</td>
<td>167°04</td>
</tr>
<tr>
<td>Lifou south</td>
<td>15</td>
<td>96.95</td>
<td>21°01'</td>
<td>167°22</td>
</tr>
<tr>
<td><strong>Lifou</strong></td>
<td><strong>93</strong></td>
<td><strong>644 [1196]</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Mare northwest</td>
<td>21</td>
<td>125.32</td>
<td>21°24'</td>
<td>167°52</td>
</tr>
<tr>
<td>Mare southwest</td>
<td>18</td>
<td>118.91</td>
<td>21°35'</td>
<td>167°53</td>
</tr>
<tr>
<td>Mare East</td>
<td>22</td>
<td>89.18</td>
<td>21°33'</td>
<td>168°05</td>
</tr>
<tr>
<td><strong>Maré</strong></td>
<td><strong>61</strong></td>
<td><strong>333.4 [650]</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Ouvea north</td>
<td>21</td>
<td>45.87</td>
<td>20°27'</td>
<td>166°36</td>
</tr>
<tr>
<td>Ouvea middle</td>
<td>15</td>
<td>21.40</td>
<td>20°38'</td>
<td>166°34</td>
</tr>
<tr>
<td>Ouvea south</td>
<td>12</td>
<td>20.58</td>
<td>20°43'</td>
<td>166°25</td>
</tr>
<tr>
<td><strong>Ouvéa</strong></td>
<td><strong>48</strong></td>
<td><strong>87.85 [132]</strong></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Considering each locus in each population separately, the distribution of allele frequencies was highly unbalanced (results not shown). In 67% of the situations, one allele presented a frequency higher than 0.5.

Observed and expected (in parenthesis) heterozygosity values ranged from 0.01 (0.03) in Maré southeast up to 0.74 (0.74) in Ile des Pins North. The pattern of variation for $A$ and $H_O$ differed among populations within each island (Table 2). For instance within Grande Terre, the mean number of allele varied from 5.13 to 7.5 and mean observed (respectively expected) heterozygosity from 0.61 (0.62) to 0.75 (0.72). Concerning Hardy–Weinberg equilibrium, all results per population are given in Table 2.

A significant heterozygote deficit was detected in each island except Ile des Pins. $F_{IS}$ values varied widely among islands, ranging from 0.14 in Ouvéa to 0.62 in Maré. Within islands, significant $F_{IS}$ values after Bonferroni correction were all positive and varied weakly among populations.

**Analyses of population differentiation**

All $F_{ST}$ values were significant. $F_{ST}$ was 0.22 among islands and 0.35 among populations (Table 3). Between the two groups ‘Grande Terre — Ile des Pins’ and ‘Îles Loyauté’, the $F_{ST}$ was 0.24 and in this case $F_{ST}$ was 0.39 among populations. Global $F_{ST}$ among populations within islands was 0.12, it was around 0.20 within Grande Terre and Maré, around 0.07 within Lifou and Ouvéa and around 0.01 within Ile des Pins. All pairwise $F_{ST}$ values were significant at the level $\alpha = 0.05$ except for the pair Maré north and Maré southeast ($F_{ST} = 0.01, P = 0.19$) and for the pair Ile des Pins North and South ($F_{ST} = 0.02, P = 0.10$) (Table 3).

Populations from Grande Terre and Ile des Pins and from Îles Loyauté were separated by the NJ tree (Fig. 2), and populations of Ile des Pins were clearly related to the group of populations of Grande Terre. The populations of Lifou had an intermediate position between the populations of Maré, Ouvéa and Grande Terre. The populations of the north of Grande Terre (Malek and more specially Hienghène) were closer to Îles Loyauté than the other Grande Terre populations.

Mantel tests revealed a significant correlation between geographical and genetic distances considering all the populations ($r = 0.31, P < 0.05$) (Fig. 3a). There was a significant association among Îles Loyauté populations ($r = 0.65, P < 0.001$) and among Grande Terre populations ($r = 0.6, P < 0.05$) (Fig. 3b). When Ile des Pins was included in the model with Îles Loyauté or with Grande Terre, the coefficient was not significant ($P > 0.05$ in both cases).
Table 3 Results of the AMOVA testing the genetic structure among islands, among two groups (‘Grande Terre-Ile des Pins’ and ‘Iles Loyauté’) and among population within islands. For each analysis, we figured the percentage of the total differentiation attributable to the variation among groups (if there are some), among populations and within populations. Moreover, we figured the $F_{ST}$ between population within island.

<table>
<thead>
<tr>
<th>Group</th>
<th>Source of variation</th>
<th>d.f.</th>
<th>Percentage of variation</th>
<th>$F_{ST}$</th>
</tr>
</thead>
<tbody>
<tr>
<td>Among islands</td>
<td>Among islands</td>
<td>4</td>
<td>22.11</td>
<td>0.35***‡</td>
</tr>
<tr>
<td></td>
<td>Among populations within islands</td>
<td>12</td>
<td>12.36‡</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Within populations</td>
<td>845</td>
<td>65.53</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Total</td>
<td>861</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Between ‘Grande Terre-Ile des Pins’ and ‘Iles Loyauté’ groups</td>
<td>Among groups</td>
<td>1</td>
<td>23.58</td>
<td>0.39***‡</td>
</tr>
<tr>
<td></td>
<td>Among populations within groups</td>
<td>15</td>
<td>15.47‡</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Within populations</td>
<td>845</td>
<td>60.96</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Total</td>
<td>330</td>
<td>19.51</td>
<td>0.20***</td>
</tr>
<tr>
<td></td>
<td>Total</td>
<td>335</td>
<td>80.49</td>
<td></td>
</tr>
<tr>
<td>Within Grande Terre</td>
<td>Among populations</td>
<td>5</td>
<td>1.36</td>
<td>0.01***</td>
</tr>
<tr>
<td></td>
<td>Within populations</td>
<td>120</td>
<td>98.64</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Total</td>
<td>121</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Within Ile des Pins</td>
<td>Among populations</td>
<td>1</td>
<td>1.36</td>
<td>0.01***</td>
</tr>
<tr>
<td></td>
<td>Within populations</td>
<td>120</td>
<td>98.64</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Total</td>
<td>121</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Within Mare</td>
<td>Among populations</td>
<td>2</td>
<td>19.77</td>
<td>0.2***</td>
</tr>
<tr>
<td></td>
<td>Within populations</td>
<td>119</td>
<td>80.23</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Total</td>
<td>121</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Within Lifou</td>
<td>Among populations</td>
<td>2</td>
<td>6.69</td>
<td>0.07***</td>
</tr>
<tr>
<td></td>
<td>Within populations</td>
<td>183</td>
<td>93.31</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Total</td>
<td>185</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Within Ouvea</td>
<td>Among populations</td>
<td>2</td>
<td>6.68</td>
<td>0.07***</td>
</tr>
<tr>
<td></td>
<td>Within populations</td>
<td>93</td>
<td>93.32</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Total</td>
<td>95</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

‡The $F_{ST}$ represents the differentiation among populations within the total population.
‡The percentage represents the differentiation among populations within island (group).

Discussion

Diversity

In this study we used different parameters to assess genetic diversity. The parameters using the number of alleles ($A$ and $R$) are complementary of those using allele frequencies ($H$), specially for analyses raising conservation issues (El Moussadik & Petit 1996). Here we used the allelic richness corrected by the rarefaction index ($R$) to take into account the differences in sample size of the populations (from 10 to 53 individuals). Our result showed a very high correlation between $A$ and $R$ (Pearson coefficient: 0.96, Spearman coefficient: 0.935) demonstrating that the correction with the rarefaction index has no effect on diversity assessment. This also suggests that rare alleles (which strongly influence measures of allelic richness) have not a more scattered distribution than the other alleles.

Many tree species exhibit regional structuring (Morand et al. 2002), particularly when gene flow is limited by barriers such as mountains or oceans. Our analysis confirmed this principle, as it revealed great differences in diversity parameters between islands throughout the archipelago. The broad range in observed heterozygosity ($H_o$) values resulted from the broad variation in the mean number of alleles per locus ($A$) and follows the pattern found in other microsatellite studies of tropical tree species. For example, Dayanandan et al. (1999) found $A$ values between 2 and 15, and $H_o$ values between 0.13 and 0.93 in Carapa guianensis.

Our diversity parameters ($A = 2–16$, mean = 15.37; $H_S = 0.14–0.79$, mean = 0.66) were comparable to those of other tree species analysed with microsatellites. They were higher than those of Vitellaria paradoxa ($A = 3.4–4.2$, $H_E = 0.38–0.44$) (Kelly et al. 2004), Vouacapoua americana ($A = 3.2–5.1$, $H_S = 0.34–0.52$) (Dutech et al. 2004), Grevillea macleayana ($A = 3.2–4.2$, $H_E = 0.42–0.53$) (England et al. 2002), and lower than those of Melaleuca alternifolia ($A = 20–27$, $H_E = 0.13–0.92$) (Rossetto et al. 1999) and Symphonia globulifera (mean $A = 3.7–16$, mean $H_E = 0.67–0.85$) (Aldrich et al. 1998).

Populations of Iles Loyauté had a lower genetic diversity, particularly Maré (mean observed heterozygosity $H_o = 0.156$, mean number of alleles $A = 2$), compared to Ile des Pins ($H_o = 0.74$, $A = 10.25$) and Grande Terre ($H_o = 0.69$, $A = 16$). Genetic variability was not correlated positively with population size ($r = -0.47$, not significant with a two-tailed Pearson correlation test with $a = 0.05$; $P > 0.025$), it was lower in Iles Loyauté than in Grande Terre and Ile des Pins (for example, mean number of allele per locus per km$^2$).
was 0.006 in Maré and 2.16 in Grande Terre). The higher genetic diversity in Grande Terre can be related to its higher number of populations and its larger area. But it can also simply result from the history of sandalwood colonization throughout the archipelago. Grande Terre and Ile des Pins are the most ancient islands of the archipelago and in major part comprise sedimentary and volcanic formations from the Permian (225–280 million years ago) to the Tertiary (1.5–65 Ma). Their same geological origin and their connection during the last glaciation between 14 000 and 9000 years bp (Stevenson et al. 2001) could explain their similarity concerning genetic diversity, which is also reflected by their proximity in the genetic tree. The Eastern arc of the Iles Loyauté rests on old volcanoes, that were gradually drowned under the sea, whereas the coral grew in height, forming a ring which evolved into an atoll when the volcanic island disappeared under the sea. In the Quaternary period (1.8 Ma) these filled lagoons were raised, giving the current limestone islands (Picard 1999).

Given the different ages of the islands, it is very likely that sandalwood arrived first on Grande Terre and Ile des Pins, then differentiated into the species *Santalum austrocaledonicum*, and then colonized the Loyautés. Frankham (1997) established comparisons between mainland and island diversity of many organisms, and found that in most cases mainland populations were more diverse. Given their large size, Grande Terre and Ile des Pins can be likened to mainland, and lower diversity in the Iles Loyauté can be explained by the island colonization process and restricted gene flow.

The very low diversity in Mare ($A = 2$, $H_E = 0.14$) can be explained by a stepping-stone model (Kimura & Weiss 1964) if we consider that there was no colonization from the east side of Grande Terre to the Iles Loyauté, and that there was two paths of colonization, one from north Grande Terre to Ouvéa then Lifou, and then Maré, the other from south Grande Terre and Ile des Pins to Maré. Hence Maré could be the end of the path and would have received less genetic diversity.

**Heterozygote deficit**

Our study revealed a strong heterozygote deficit in each island, and high values of $F_{IS}$ particularly in Iles Loyauté. At least four explanations may account for this deficit.

The first is the occurrence of null alleles (alleles that are never amplified because of mutations in the flanking primer sequences (Callen et al. 1993)). This could explain the departure from Hardy–Weinberg equilibrium of loci *mSaCIRE09*, *mSaCIRG10* and *mSaCIRF04*. But this seems unlikely because amplification failures that would reflect null/null homozygotes were rare (maximum of 5.6% for...
locus G-10). One means to test their absence would be to design sets of new primers located upstream and downstream from the original ones, to see if the individuals previously scored as homozygote remain as such when using the new primers, as did Gibbs et al. (1997). We decided not to search for null alleles as the analyses gave similar results with and without the three loci previously mentioned.

A second explanation is the Wahlund effect, which occurs when a subdivided population contains fewer heterozygotes than predicted despite the fact that all subdivisions are in Hardy–Weinberg equilibrium. This effect may explain a part of the heterozygote deficit of the whole island population. For example, Ouvéa shows a high and significant $F_{ST}$ (0.14*) (high heterozygote deficit), but when divided into three populations, two of them do not show heterozygote deficit (Ouvéa Midland $F_{IS} = -0.03$ ns, Ouvéa South $F_{IS} = 0.1$ ns). Similarly, in Grande Terre ($F_{IS} = 0.17**$), in which populations are more isolated, all populations but Pindai ($F_{IS} = 0.09$*) showed no significant departure from panmixia.

Two characteristics of *S. austrocaledonicum* may contribute to the creation of a Wahlund effect. First, the suckering from adventitious buds on roots following disturbance, which has permitted sandalwood to subsist after the dramatic cuttings in the 19th century and the frequent fires in the archipelago. This asexual way of reproduction leads to clones in a small perimeter. We clearly observed this phenomenon throughout the archipelago, as close individuals often had the same genotype (cf. Material and methods).

Mating among individuals belonging to the same clone is akin to selfing, and will similarly create departure from Hardy–Weinberg expectations.

Second, a low seed and/or pollen dispersal can create a clustering of genetically related trees around a mother tree. This is the case for *Vouacapoua americana* (Dutech et al. 2004) whose seeds are heavy (around 30 g) and dispersed by small rodents which bury them usually less than 10 m from their source. Unlike *V. americana*, *S. austrocaledonicum*'s seeds are not very large (8 mm × 5 mm) so they can easily be disseminated by birds over large distances. However, if pollinators are lacking, it is very likely that trees reproduce by selfing. Selfing will also happen when tree density is low and pollinators tend to stay on the same tree. Selfing seems to be the most logical explanation for heterozygote deficiency in situations such as Maré where mean observed heterozygosity is extremely low, $F_{IS}$ is very strong and significant (0.62), and subdivision of the island into three populations does not improve the results ($F_{ST}$ all significant and around 0.5).

The Wahlund effect can be not only spatial but also temporal (Morand et al. 2002): when flowering dates are consistently different among trees, as with *S. austrocaledonicum*, reproduction is restricted to the individuals flowering at the same time, hence creating a cluster of trees. However, this hypothesis assumes that ‘populations’ of trees flowering at the same time keep the same flowering dates from one year to another and are of finite size. Moreover, it supposes that these populations have evolved different allelic frequencies at microsatellite loci, or that there was an initial disequilibrium between neutral markers and loci involved in homogamy that are affected by a heterozygote deficit.

**Differentiation between populations**

The degree of differentiation between populations is both influenced by drift, which increases differentiation, and gene flow, which reduces it. Gene flow through pollen is expected to be low in *S. austrocaledonicum* as it is insect-pollinated, but seed dispersal through bird ingestion can occur over long distances and may allow gene flow between islands.

Both $F_{ST}$ values and percentage of variance obtained with the AMOVA indicated a strong differentiation between islands ($F_{ST} = 0.22***$) and a lower differentiation between populations within island (intra island $F_{ST} = 0.16$ representing 12% of the total variance). This result was expected in an island system, where ocean barriers limit gene flow.
between populations (MacArthur & Wilson 1967), and confirms the first results obtained for tree species with similar geographical patterns, for example *Pterocarpus officinalis*, which revealed a strong differentiation between islands with AFLP markers (Rivera-Ocasio *et al*. 2002).

As expected with the isolation of its populations, Grande Terre had a high $F_{ST}$ (0.2***). The high $F_{ST}$ value in Maré (0.2***) was harder to explain, but the possible more inbred mating system in this island may have influenced the calculation as effective size is reduced in selfing populations.

Because of its large size, Grande Terre was the most suitable island for comparisons of population structure with continental species.

Our within-island results are similar to those of *Santalum spicatum* which had an $F_{ST}$ of 0.09 (Byrne *et al*. 2003), but the markers used for this analysis were RFLP, which may affect the result compared to what would be obtained with microsatellites, RFLP marker mutation rate being lower than that of microsatellite markers.

To our knowledge, there is no study of continental tree species using microsatellites with similar patterns of dispersal. The majority of trees studied are insect pollinated, but have seed dispersal over small distances from the mother tree (barochorous, zochorous by small animals), or anemochorous over a small distance, so their gene flow is potentially lower than that of species dispersed by birds, hence their $F_{ST}$ is expected to be higher. However, examples of those tree species studied on a similar geographical scale showed a lower differentiation than on Grande Terre: with *Caryocar brasiliense*, Collevatti *et al*. (2001) found an $F_{ST}$ of 0.11 and *Swietenia macrophylla* exhibited a $F_{ST}$ of 0.1 in the Brazilian Amazon (Lemes *et al*. 2003), and 0.11 in Central America (Novick *et al*. 2003). This result indicates that gene flow in *S. austrocaledonicum* on Grande Terre is lower than expected considering its large dispersal distance.

The tree representing genetic distances between populations confirmed the emergence of two distinct groups: Iles Loyauté and Grande Terre/Ile des Pins. Ile des Pins populations were genetically very close to Grande Terre ones, as already suspected from their geographical connectivity during the last glaciations (Stevenson *et al*. 2001), but could be explained simply by their geographical proximity.

The case of Hienghène is interesting, as it is the only island already suspected from their geographical connectivity during the last glaciations (Stevenson *et al*. 2001). This result indicates that in *S. austrocaledonicum* conservation strategies, like the creation of reserves, could be designed to preserve large areas to minimize the loss of diversity due to genetic drift, and to conserve maximally the regional genotypic diversity.

Our study allows us to define two molecular ESUs (evolutionary significant units, Ryder 1986) that we can consider as provenance zone (in the forestry sense) for *S. austrocaledonicum*: 'Grande Terre and Ile des Pins' and 'Iles Loyauté', which are differentiated by an $F_{ST}$ of 0.24. The term ESU was devised in a practical way to approach the conservation of genetic resources, given the broad recognition of the importance of genetic diversity in conservation policy and the frequent inadequacy of existing taxonomy to describe it (Moritz 1994). The criterion for identification of an ESU has been defined as reciprocal monophyly for organelle haplotypes and significant divergence of allele frequencies at nuclear loci (Moritz 1994). Crandall *et al*. (2000) have pointed out that ESUs based on neutral molecular criteria will not address many of the real problems of conservation. Ecological factors, such as frequency-dependent mating and pollinator interactions, must also be taken into account. It is likely that the best strategy lies in concordance between neutral genetic and adaptive information (Moritz 2002) but, as a starting point, identification of molecular ESUs provides a valuable practical framework (Cavers *et al*. 2003). The study of the genetic diversity of chloroplast microsatellites we are now conducting on *S. austrocaledonicum* will permit us to define these molecular ESUs.

Acknowledgments

We would like to thank Alexandre Vaillant and Pierre Sire for laboratory work and Alexandre Lagrange for field work in New Caledonia. Many thanks go to IAC (Institut Agronomique néo-Calédonien) and to the Development Services of the Provinces of Islands, North and South. We also acknowledge MEED (Ministère de l’Ecologie et du Développement Durable) for financial support.

References


These results are part of Lorraine Bottin’s PhD thesis on the analysis of genetic diversity of Santalum austrocaledonicum. This study was one of the tasks of the sandalwood project founded by the French Ministry of Ecology and Sustainable Development involving Isabelle Olivieri working on evolutionary genetics at ISEM, University of Montpellier. Laboratory work and analyses were done in Forest Department of CIRAD where Jean-Marc Bouvet is the responsible of the ‘Forest genetics’ research unit. Daniel Verhaegen is a molecular biologist and Jacques Tassin is an ecologist positioned in New-Caledonia working in the same research unit.