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# THE DOCTORAL RESEARCH ABSTRACTS

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**Title :** EVALUATION OF THE SIMPLE SEQUENCE REPEAT (SSR) GENOTYPING OF *Elaeis oleifera* GERMPLASM

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Special attention has been given to the second species of oil palm, *Elaeis oleifera* as it possess several interesting agronomic traits such as slow growth, higher oil unsaturation and disease resistance. Studying the variability of *E. oleifera* germplasm is therefore very important as it serves as a tool to select source of genetic diversity for the oil palm conservation programme. The objectives of this study were 1) to identify the polymorphic *E. oleifera* gSSRs for *E. oleifera*, 2) to measure the information content of *E. oleifera* gSSRs, 3) to unravel the genetic diversity of *E. oleifera* germplasm, 4) to determine the genetic differentiation of *E. oleifera* germplasm and 5) to assess the genetic structure of *E. oleifera* germplasm. MPOB has developed a collection of simple sequence repeats (SSRs) from *E. oleifera* genome. Initially, optimization of PCR conditions for analysis of the *oleifera* samples using *E. oleifera* gSSRs was carried out. A total of 316 *E. oleifera* gSSRs were tested to evaluate their usefulness to assess the genetic diversity and population structure of *E. oleifera* populations. The PCR conditions were optimized while keeping the original DNA concentration, annealing temperature (TA) and other reagent constant. Out of 316 *E. oleifera* gSSRs screened, 270 produced amplicons and of these numbers, 140 were polymorphic and potentially useful for diversity analysis. The modified PCR condition increases the success of amplifying *E. oleifera* gSSRs in the *E. oleifera* DNA samples analyzed. The PCR methods together with the polymorphic *E. oleifera* gSSRs were applied in genotyping the entire *E. oleifera* populations. A set of 21 polymorphic *E. oleifera* gSSRs was analyzed on a total of 214 *E. oleifera* genomic DNA belonging to eight germplasm originated from four countries in Central and South America (Columbia, Panama, Costa

Rica and Honduras). The analysis covered on genetic diversity and genetic structure of eight *E. oleifera* populations, inferences from 21 polymorphic *E. oleifera* gSSRs. The average observed heterozygosity across population ( $H_o=0.249$ ) was less than the expected heterozygosity ( $H_e=0.363$ ). The highest population diversity was obtained in population 08 from Columbia ( $H_e=0.460\pm 0.055$ ,  $I=0.870\pm 0.121$ ). Eight of 21 polymorphic *E. oleifera* gSSRs were informative with  $PIC>0.5$ , where sMo00131 is the most informative ( $PIC=0.853$ ). The populations analysed showed great genetic differentiation ( $F_{ST}=0.223$ ). The Nei genetic distance showed the highest genetic distance was between population 01 from Columbia and population 02 from Costa Rica (0.555) while the lowest was between population 02 and population 03 from Honduras (0.019). The eigenvalues of PCoA plot showed that the first two components explained 38.70% of the total variation, which roughly ordinated the *E. oleifera* individuals into three major groups. Construction of neighbour-joining (NJ) tree separated *E. oleifera* individuals into two clusters. Model based clustering revealed that *E. oleifera* population has the highest  $\Delta K$  when K was set to 7. The present study provides a diverse pattern of genetic diversity and the existence of genetic differentiation among *E. oleifera* germplasm. This study highlights the potential contribution of genetic variation of the *E. oleifera* collection analyzed using *E. oleifera* gSSRs for germplasm conservation and for utilization in breeding programs. Further conservation should focus on more populations with less number of palms per population development of core collection.