### MITOGENOME ANNOUNCEMENT

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# First complete mitochondrial genome of Phoenix dactylifera var. Khanezi

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#### ABSTRACT

In this study, we determined the complete mitochondrial (mt) genome of *Phoenix dactylifera* var. Khanezi. The results revealed a circular genome of 715,120 bp, having G + C content of 45.1%, containing 40 protein coding genes, 3 rRNA, and 18 rRNA genes. Evolutionary relationship analysis suggested that *P. dactylifera* var. Khanezi is more closely related to previously reported *P. dactylifera* var. Khalas.

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Date palm, *Phoenix dactylifera* L., belongs to *Arecaceae* and is an ecologically, culturally and economically important fruit tree in North Africa and Middle East (Moussouni et al. 2017). Date palm is a perennial, monocotyledon (2n = 36), dioecious, cross-pollinated tree that has been widely cultivated (Barrow 1998; Terral et al. 2012; Cherif et al. 2013). Morphological variations, which are heavily dependent on environmental factors and data variety, do exist among cultivars. These variations are reflected in the diversity of the chloroplast (cp) genome, and mt genomics as well.

The mitochondria play a vital role in plant growth and development (Ogihara et al. 2005). Recently, mt genomic gained much attention due to advancement in genomics, and mt DNA is considered as a significant and effective source of genetic variation among various species. The mt genomes of plant have big and complicated structure as compared to other eukaryotes (Li et al. 2009; Liu et al. 2011). Furthermore, angiosperm mt genomes are well known for uptake of foreign DNA by horizontal gene transfer (Goremykin et al. 2009) and for their very low mutation rate (Palmer and Herbon 1988). In the present study, we report for the first time the complete nucleotide sequence of *P. dactylifera* var. Khanezi mt genome (GenBank accession number: NC016740) and to infer its phylogenetic position on the basis of entire mito-genomes.

The *P. dactylifera* var. Khanezi (accession), plants were received from the GenBank at the University of Nizwa, Nizwa Oman. Young leaves were used to extract mitochondrial DNA by using the method described previously (Asaf et al. 2016;

Wu 2016) with little modification. The living material and DNA were stored in University of Nizwa, Nizwa, Oman. Geneious Pro v11.1 (http://geneious.com) was used to filter and assemble the raw alumina reads. All contigs were blasted using the NCBI database (http://www.ncbi.nlm.nih.gov/) for the annotation of mitochondrial sequences. Similarly, tRNA scan-SE software (http://lowelab.ucsc.edu/tRNAscan-SE/) was used to identify tRNAs. The entire mitochondrial genome was used to determine its phylogenetic position using Maximum persimony tree with 1000 bootstrap replications (Felsenstein 1985).

The mitochondrial genome of *P. dactylifera* var. Khanezi was assembled as a single circular molecule of 715,120 bp having 45.1% GC contents (deposited in GenBank under the accession MH176159). The largest part of the *P. dactylifera* mtDNA comprises the non-coding sequences (93.2%), which is slightly larger than the average non-coding sequences content (89.45%) in other angiosperm mt genomes (Chaw et al. 2008). Furthermore, the mt genome contains 67 genes encoding 24 transfer RNAs, 3 ribosomal RNAs and 40 protein coding genes. Moreover, the phylogenetic analysis revealed that mt genome of *P. dactelifera* var. Khanezi is closely related to *P. dactelifere* var. Khalas (NC016740; Figure 1) and *Cocos nucifera*. This study will help to understand the evolution of various date palm cultivars mitochondrial genome with related species.

### **Disclosure statement**

No potential conflict of interest was reported by the authors.

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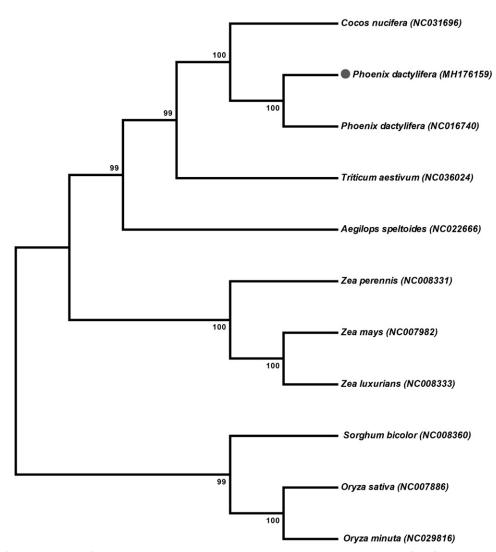


Figure 1. Phylogeny of the *Phoenix dactylifera* var. Khanezi mitochondrial genome with ten other related species (two from family Arecaceae and eight from family Poaceae). The phylogenetic tree was inferred using the Maximum parsimony method based on entire mitochondrial genomes of these species.

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