

## Documents

Jenuit, M.<sup>a</sup>, Zainuddin, Z.Z.<sup>b</sup>, Payne, J.<sup>b</sup>, Ahmad, A.H.<sup>b c</sup>, Yusof, A.M.<sup>d e</sup>, Isa, M.L.M.<sup>d e</sup>, Ibrahim, M.<sup>f</sup>

**Establishment And Cryopreservation Of Fibroblast Cell Line From A Sumatran Rhinoceros (*Dicerorhinus Sumatrensis*)**

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<sup>a</sup> Department of Biotechnology, Kulliyah of Science, International Islamic University Malaysia, Jalan Sultan Ahmad Shah, Bandar Indera Mahkota, Kuantan, Pahang 25200, Malaysia

<sup>b</sup> Borneo Rhino Alliance Berhad, Faculty of Science and Natural Resources, Universiti Malaysia Sabah, Jalan UMS, Kota Kinabalu, Sabah 88400, Malaysia

<sup>c</sup> Faculty of Sustainable Agriculture, Universiti Malaysia Sabah, Locked Bag No 3, Sandakan, Sabah 90509, Malaysia

<sup>d</sup> Department Basic Medical Sciences, Kulliyah of Nursing, International Islamic University Malaysia, Jalan Sultan Ahmad Shah, Bandar Indera Mahkota, Kuantan, Pahang 25200, Malaysia

<sup>e</sup> IIUM Molecular and Cellular Biology Research Cluster (iMoleC), International Islamic University Malaysia, Jalan Sultan Ahmad Shah, Bandar Indera Mahkota, Kuantan, Pahang 25200, Malaysia

<sup>f</sup> Department of Physics, Faculty of Science, Benha University, Egypt, Benha 13518, Egypt

**Abstract**

Cell lines have been established to preserve the genetic material of endangered animals. This study aims to establish, characterize and authenticate fibroblast cells derived from the kidney tissue of a Sumatran rhinoceros' carcass. The primary cultures were obtained using the mixed enzymatic-explant method, supplemented with complete media and maintained at 37°C with 5% CO<sub>2</sub> in an incubator. Following routine trypsinization, viability and growth curves were generated through the Trypan Blue counting method. Cellular senescence was quantified by Sa-P-gal staining assay and G-banding for karyotyping. As a result, the cell derivation had generated 81 frozen stocks. The viability of cells at P5 and P10 showed reasonable recovery after six months. Cell population doubling time at P5 was 20.45 hours, while it was 22.35 hours at P10 and P15. The senescence level significantly increased from P5 to P10, and was especially significant at P15. Genetic stabilities were considered stable at P5 and P10, with frequency of over 70 %. In conclusion, this study was able to derive a primary fibroblast culture from the preserved tissue of a Sumatran rhinoceros, with certain changes in morphology, senescence level, growth curves and cell viability as the number of passages increased © 2021, Penerbit UMT, All Rights Reserved.

**Author Keywords**

characterization; cryopreservation; fibroblast cells establishment; Sumatran rhinoceros; Sustainability

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**Correspondence Address**

Isa M.L.M.; Department Basic Medical Sciences, Malaysia; email: lokman@iium.edu.my

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