International Congress Series 1261 (2004) 428-430





# Mitochondrial DNA profiling of modern Malay and Orang Asli populations in peninsular Malaysia

Z. Zainuddin<sup>a</sup>, W. Goodwin<sup>b,\*</sup>

<sup>a</sup>Department of Forensic Medicine and Science, University of Glasgow, Glasgow, UK <sup>b</sup>Department of Forensic and Investigative Science, University of Central Lancashire, Preston, PR1 2HE, UK

**Abstract.** A study was undertaken to assess the utility of mtDNA as a forensic tool in the Malay Peninsular. Two populations, modern Malay and Orang Asli, were sampled and their mitochondrial DNA (mtDNA) analysed. Comparing the different haplogroups that were found, there are clear differences between the two populations. As a forensic tool, the use of mtDNA in the Orang Asli is limited as many individuals share the same haplotype, reducing the power of discrimination. In the Malay population, the power of discrimination is sufficiently high to make it a valuable tool. © 2003 Elsevier B.V. All rights reserved.

Keywords: mtDNA; Hypervariable region I; Malay; Orang Asli

# 1. Introduction

Mitochondrial DNA has proven to be a useful tool for human identification as well as understanding modern human evolution and migrations. Malaysian populations are underrepresented in the published data, and therefore, before using mtDNA as a forensic tool in Malaysia, it is important to study the population. This is particularly so when there are distinct populations within the main population, such as the Orang Asli. In addition to providing useful information for forensic studies, this information will also provide information on the relationships of the Malay population with neighboring populations.

# 2. Materials and methods

In total, 102 samples were collected from 11 states of Peninsular Malaysia and 58 samples from two populations of Orang Asli (Kensiu and Jahai) were collected. DNA was extracted using Puregene<sup>®</sup> DNA Extraction kit. The HVI region was amplified using L15926 and H16431 primers and sequenced. Polymorphisms were reported by

<sup>\*</sup> Corresponding author. Tel.: +44-1772-894254; fax: +44-1772-894981.

E-mail address: whgoodwin@uclan.ac.uk (W. Goodwin).

Table 1				
Population	Samples	Haplotypes <sup>a</sup>	Power of exclusion <sup>b</sup>	
Malay	102	73 (6)	0.0203	
Orang Asli	58	12 (18)	0.1961	

<sup>a</sup> Total number of haplotypes observed (most frequently occurring haplotype in brackets).

<sup>b</sup> The probability that two samples chosen at random would have the same haplotype.

aligning the sequence obtained to the Anderson Reference Sequence [1]. RFLP analysis of the coding region was performed to examine the following mutations: +10394/+10397 *Ddel/AluI* [2,3], +13262 *AluI* [4], -5176 *AluI* [4,5], -12406 *HincII* [6], +12308 *Hinf*I [7], -1715 *DdeI* [7,8] and -7025 *AluI* [5,8]. The haplogroups were determined using a combination of the HVI sequence and RFLP analysis (haplogroup nomenclature was in accordance with the above references and M. Richards, personal communication).

#### 3. Results

Analysis of the hypervariable region I was used to place the samples into haplogroups. A summary of the mtDNA diversity is shown in Table 1.

Additional information from RFLP sites located outside of the hypervariable regions was used, in some cases, to place the samples into more highly resolved haplogroups. The different haplogroups along with their defining polymorphisms and the frequencies of these haplogroups in the tested populations are shown in Table 2.

From the data, it is clear that there are large differences between the modern Malay and the Orang Asli populations, both in the power of discrimination and in the haplogroups composition.

Haplogroup	Defining polymorphisms	Population frequencies	
		Malay	Orang Asli
M <sup>a</sup>	+10394/+10397 <i>DdeI/Alu</i> I	39.12	5.17
M21a	+10394/+10397 DdeI/AluI	5.88	34.48
M21b	+10394/+10397 DdeI/AluI	_	1.72
M-D	+10394/+10397 DdeI/AluI, -5176 AluI	1.96	_
M-C	+10394/+10397 DdeI/AluI, +13262 AluI	0.98	_
B4	9-bp deletion	7.84	_
B5	9-bp deletion	10.78	6.90
Е	+10394/+10397 DdeI/AluI, -7598 HhaI	1.96	_
F1a	- 10394/ - 10397 DdeI/AluI, - 12406 HincII	16.67	_
N9a	– 10394/ – 10397 <i>DdeI/Alu</i> I	2.94	3.45
R	– 10394/ – 10397 <i>DdeI/Alu</i> I	7.84	_
R21	+10394/-10397 DdeI/AluI, -1715 DdeI	1.96	48.28
Y	+10394/-10397 DdeI/AluI	1.96	_

Table 2 The distribution of haplogroups within the modern Malay and Orang Asli populations

<sup>a</sup> Includes all variations of the M super haplogroup that are not specified.

### 4. Discussion

As in other populations, mtDNA can be a powerful tool for human identification when used in the Malay population. However, the differences between the Malay and Orang Asli populations highlight the need for care in interpreting any results in a forensic context. Further studies examining the nuclear autosomal and Y-chromosome loci will provide an improved understanding of the genetic composition in different populations in Peninsular Malaysia.

#### References

- [1] S. Anderson, A.T. Bankier, B.G. Barrell, M.H. de Bruijn, A.R. Coulson, J. Drouin, I.C. Eperon, D.P. Nierlich, B.A. Roe, F. Sanger, P.H. Schreier, A.J.H. Smith, R. Staden, I.G. Young, Sequence and organisation of the human mitochondrial genome, Nature 290 (1999) 457–465.
- [2] A. Torroni, T.G. Schurr, M.F. Cabell, M.D. Brown, J.V. Neel, M. Larsen, D.G. Smith, C.M. Vullo, D.C. Wallace, Asian affinities and continental radiation of the four founding Native American mtDNAs, Am. J. Hum. Genet. 53 (1993) 563–590.
- [3] T. Kivisild, H.-V. Tolk, J. Parik, Y. Wang, S.S. Papiha, H.-J. Bandelt, R. Villems, The emerging limbs and twigs of the East Asian mtDNA tree, Mol. Biol. Evol. 19 (10) (2002) 1737–1751.
- [4] A. Torroni, T.G. Schurr, C.-C. Yang, E.J.E. Szathmary, R.C. Williams, M.S. Schanfield, G.A. Troup, W.C. Knowler, D.N. Lawrence, K.M. Weiss, D.C. Wallace, Native American mitochondrial DNA analysis indicates that the Amerind and Nadene populations were founded by two independent migrations, Genetics 130 (1992) 153–162.
- [5] A. Torroni, D.C. Wallace, Mitochondrial DNA variation in human populations and implications for detection of mitochondrial DNA mutations of pathological significance, J. Bioenerg. Biomembranes 26 (1994) 261–271.
- [6] T. Kivisild, K. Kaldma, M. Metspalu, J. Parik, S. Papiha, R.Villems, The place of the Indian mitochondrial DNA variants in the global network of maternal lineages and the peopling of the old world, in: S. Papiha, R. Deka, R. Chakraborty (Eds.), Genomic Diversity: Applications in Human Population Genetics, Kluwer Academic/Plenum Publisher, New York, 1999, pp. 135–152.
- [7] A. Torroni, K. Huoponen, P. Francalacci, M. Petrozzi, L. Morelli, R. Scozzari, D. Obinu, M.L. Sarontaus, D.C. Wallace, Classification of European mtDNAs from an analysis of three European populations, Genetics 144 (1996) 1835–1850.
- [8] V. Macaulay, M. Richards, E. Hickey, E. Vega, F. Cruciani, V. Guida, R. Scozzari, B. Bonne-Tamir, B. Sykes, A. Torroni, The emerging tree of West Eurasian mtDNAs: a synthesis of control-region sequences and RFLPs, Am. J. Hum. Genet. 64 (1999) 232–449.

#### 430