

a- Royal Mitochondrial Lineage Mummies

| Mummy (tomb) | HVRI Diagnostic Mutation*, (Frequency, Experiments [†]) | Predicted Haplogroups [‡] | Sequences per Alignment | Min/ Max Coverage | Coverage |
|-------------------------|---|------------------------------------|-------------------------|-------------------|----------|
| Thuya (KV46) | 16224 (89% - 4 exps) 16311 (92% - 3 exps) | K | 87 | 4/57 | Full |
| KV35EL (Tiye) (KV35) | 16224 (90% - 5 exps) 16311 (100% - 2 exps) | K | 74 | 3/43 | Full |
| KV35YL (KV35) | 16224 (88% - 4 exps) 16311 (100% - 3 exps) | K | 72 | 6/55 | Full |
| KV55 (Akhenaten) (KV55) | 16224 (78% - 4 exps) 16311 (100% - 3 exps) | K | 137 | 27/69 | Full |
| Tutankhamun (KV62) | 16224 (96% - 5 exps) 16311 (98% - 3 exps) | K | 79 | 2/65 | Full |

b- Control Mummies

| Mummy (tomb) | HVRI Diagnostic Mutation*, (Frequency, No. of Trials [†]) | Predicted Haplogroups [‡] | Sequences per Alignment | Min/ Max Coverage | Coverage |
|----------------------|---|------------------------------------|-------------------------|-------------------|---------------------|
| Yuya (KV46) | 16224 (80% - 2 exps) 16311 (89% - 3 exps) | K | 64 | 2/58 | Full |
| Amenhotep III (KV35) | 16311 (100% - 1 exp) | H2b | 80 | 17/47 | Partial 16118-16409 |
| Sitra-In (KV60) | None | K excluded | 18 | - | Partial 16118-16232 |
| Hatshepsut (KV60) | None | K excluded | 20 | - | Partial 16118-16232 |

c- Staff Controls

| Staff Member | HVRI Diagnostic Mutation | Predicted Haplogroups [‡] |
|--------------|--|------------------------------------|
| Staff 1 | 16189; 16192; 16223; 16292; 16355; 16519 | L2a1 |
| Staff 2 | 16069; 16126; 16193; 16300; 16309 | J1d |
| Staff 3 | 16519 | H (CRS) |
| Staff 4 | 16304; 16519 | H5 |
| Staff 5 | 16217; 16309 | HV2 |
| Staff 6 | 16218; 16519 | H20 |
| Staff 7 | 16069; 16126; 16145; 16187; 16217; 16261; 16300; 16311 | J1b1 |
| Staff 8 | 16051; 16132; 16172; 16189; 16219; 16278 | U6 |
| Staff 9 | 16069; 16126; 16214; 16231; 16305 | J |
| Staff 10 | 16093; 16224; 16304; 16311 | K1a |
| Staff 11 | 16129; 16309; 16318; 16362; 16397 | U7 |

*Alterations observed in the mitochondrial DNA with a frequency higher than 75% were designated as diagnostic mutations.

[‡]The phylogenetic tree (<http://www.phylotree.org>) and the Genebase mtDNA Haplogroup Reference Guide (www.genebase.com) were used to determine mtDNA haplotypes (Van Oven & Kayser 2009, E386-E394).

[†]"Experiments" indicate the number of primer combinations used to generate differently sized PCR amplicons which harbour the diagnostic mutation.

[‡]Haplogroup determination for staff members was mainly based on the findings of Saunier *et al.* (2009, e97-e103) and Elmadawy *et al.* (2013, 338-341), in addition to general information from the van Oven & Kayser (2009, e386-e394) phylogenetic tree.

Table 4. Characterization of mitochondrial DNA of 18th Dynasty mummies