Guardian of Ancient Egypt

Studies in Honor of Zahi Hawass Volume I

> edited by Janice Kamrin Miroslav Bárta Salima Ikram Mark Lehner Mohamed Megahed

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Maternal and Paternal Lineages in King Tutankhamun's Family

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Abstract

In this study, analysis of the mitochondrial and Y-chromosomal haplogroups was used to provide information about the phylogenetic groups of Tutankhamun's family members, and their presence among the reported contemporary Egyptian population data. The mitochondrial and Y-chromosomal DNA analysis of Tutankhamun's family confirms our previous data of the royal family pedigree, with multiple controls authenticating all results. The proposed sibling relationship between Tutankhamun's parents, KV55 (Akhenaten) and KV35YL, is further supported. The royal lineage is composed of the Y-chromosome haplogroup R1b and the mitochondrial haplogroup K. Population genetics point to a common origin at ca. 14. 000–28. 000 years before present locating to the Near East.

Introduction

The analysis of mitochondrial DNA (mtDNA) has been an approach of interest to study the history of populations' migration and phylogeny (Macaulay et al. 1999, 232–249; Cavalli-Sforza and Feldman 2003, 266–275; González et al. 2006, 124; Torroni et al. 2006, 339–345; Baca and Molak 2008, 39–61). Compared to nuclear DNA, the hypervariable regions of mtDNA accumulate mutations rapidly. Being non-recombining and maternally inherited, mtDNA has become the molecular marker of choice to investigate genetic flow, to study human evolution, and to construct distinct phylogenetic groups (Giles et al. 1980, 6715–6719; Richards et al. 1996, 185–203; Macaulay et al. 1999, 232–249; Cavalli-Sforza and Feldman 2003, 266–275; Pakendorf and Stoneking 2005, 165–183; González et al. 2006, 124; Torroni et al. 2006, 339–345).

However, mtDNA identifies only the female perspective of a population and is liable to genetic selection (Pakendorf and Stoneking 2005, 165–183). Therefore, it has been integrated and supported with the study of paternally inherited Y-chromosomal DNA. This latter has been used to establish a global Y-chromosome phylogenetic tree analogous to that of the mitochondrial one (Jobling and Tyler-Smith 2003, 598–612; Baca and Molak 2008, 39–61).

One of the most renowned royal families in ancient Egypt was that of the late 18th Dynasty (ca. 1550–1295 B.C.); including the pharaoh Akhenaten and his successor, Tutankhamun. The identity of the mummies, the relationship between them, and the description of inherited genetic diseases and pathological hallmarks in this dynasty have not been fully elucidated. Also, it is still not known to what extent endogamy has contributed to the early demise of the young king Tutankhamun and the consequent ending of the 18th Dynasty. Genetic fingerprinting together with statistical analysis was used in our previous study, to reveal the paternal and maternal autosomal lineages of Tutankhamun's family (Hawass et al. 2010, 638-647). The inherited autosomal half-allele, the partial Y-chromosomal data, and the possible trio (mother, father and child) figuring of Tutankhamun's family warranted the assembly of the most possible pedigree of the king's direct lineages. Yuya and Thuya were confirmed as the great-grandparents of Tutankhamun (generation I). Amenhotep III and KV 35EL were identified as his grandparents (generation II), and the KV 55 male and the younger lady in KV 35 (KV35YL) were found to be his sibling parents (generation III). Consolidating these findings with the historical records, KV 55 could be identified as Akhenaten and KV 35EL as Queen Tiye (Hawass et al. 2010, 638-647; Habicht et al. 2016, S216-S231).

To expand the knowledge of the royal pedigree, another study was conducted to investigate the family maternal and paternal lineages. The current study employed analysis of the mitochondrial and Y-chromosomal haplogroups to test our previous results, (Hawass et al. 2010, 638–647) and to learn more about the genealogy of Tutankhamun's family.

Materials and Methods

DNA extraction, authentication and contamination monitoring:

Strict adherence to requirements of ancient DNA authentication, including laboratory design and practice standards, was adopted for proof of reliability and quality of the results (Richards et al. 1995, 291–299; Roberts and Ingham 2008, 600–613; Hawass et al. 2010, 638–647; Hawass et al. 2012, e8268; Keller et al. 2012, 698; Khairat et al. 2013, 309–325). DNA sampling and extraction were done following the published protocols for ancient DNA (Scholz and Pusch 1997, 61–64; Hawass et al. 2010, 638–647; Hawass et al. 2012, e8268). Sampling, DNA extraction, amplification, and sequencing were carried out in distinct working areas. Surfaces and equipment were continuously decontaminated with DNase and bleach and exposed to UV irradiation. Procedural components were strictly controlled and tested for the presence of contaminants. Negative controls were processed alongside each procedure to detect possible contamination with exogenous DNA sources (Tully et

al. 2001, 83–91; Pusch and Bachmann 2004, 957–964; Gilbert et al. 2005, 541–544; Lamers et al. 2009, 40–55; Haak et al. 2010, e1000536; Hawass et al. 2010, 638–647; Fulton 2012, 1–11; Khairat et al. 2013, 309–325). DNA fingerprinting for the lab members was carried out to exclude the presence of staff DNA contamination. Reproducible results were verified through bacterial cloning of amplicons (Pääbo et al. 2004, 645–679; Fernández et al. 2014, e1004401) and through the analyses of different biopsies per mummy, where 3–6 bone samples were taken from various body areas of each studied mummy (Hawass et al. 2010, 638–647). Recruitment of a dedicated laboratory separated from the original one, run by an independent research team, was used for replication purposes (Gilbert et al. 2005, 541–544; Lamers et al. 2009, 40–55; Hawass et al. 2010, 638–647; Fulton 2012, 1–11; Hawass et al. 2012, e8268).

Mummies and subjects:

Ten mummies were investigated in this study (Table 1), including Tutankhamun, his identified parents, grandparents, and great grandparents. In addition to the analyzed 10 mummies, the published data of a father-son pair [Ramesses III and Unknown man E/Pentaware] (Hawass et al. 2012, e8268) were included for haplotype comparison.

Y-STR profile analysis:

The AmpFISTR Yfiler PCR amplification Kit (Applied Biosystems, Foster City, California) was used to analyze the Y-chromosome specific haplotype profiles. The kit included sixteen human Y-chromosome DNA markers with sizes ranging between 99 bp and 324 bp, and a mean amplification size of 200 bp. All markers were tetranucleotide repetitive microsatellites with the exception of DYS392, DYS438 and DYS448, which displayed, respectively, trinucleotide, pentanucleotide and hexanucleotide repeat motifs. DNA templates were amplified according to the manufacturer's protocol with some modifications that involved decreasing the concentration of primers by 40% and increasing the AmpliTaq DNA polymerase concentration by 50%. Detection of the amplified PCR products was carried out using the ABI Genetic Analyzer 3130, Data collection Software v3.0 and GeneMapper ID v3.2 (Applied Biosystems).

The majority rule was employed for STR allele designation, where a predominant peak was generated at least 10 times [i.e. each exceeding the threshold of 50 relative fluorescence units] per mummy, using various bone samples from different body areas and monitored for slippage (Hawass et al. 2010, 638–647). A final consensus profile of each investigated mummy was constructed using the reproducible data of the generated partial profiles (Cowen et al. 2011, 400–406).

Amplification of the mitochondrial hypervariable region I, cloning and Sanger sequencing:

Specific primers were designed to amplify part of the hypervariable region I (HVRI) encompassing positions 16056 to 16409 (Table 2). These mitochondrial segments were amplified in a 50 µL PCR mix containing 1X AmpliTaq Gold® PCR Master Mix (Applied Biosystems), 20 pmol of each primer,

| age, yage, yMembers of Tutankhamun LineageThuya F Thuya F Thuya F Yuya M Yus <td< th=""><th>Estimated Tomb</th><th>Status</th></td<> | Estimated Tomb | Status |
|---|----------------------|--|
| Iutankhamun LineageF50-60N F 50-60N F 50-60N F 50N F 50N T T N T N T <th>age, y</th> <th></th> | age, y | |
| $ \begin{array}{c ccccc} & F & 50-60 \\ & M & 35-45 \\ & 19 \\ & 19 \\ & & 19 \\ & & 19 \\ & & & 19 \\ \hline {\bf p} \\ & & & & \\ \hline {\bf p} \\ & & & & \\ \hline {\bf p} \\ & & & & \\ \hline {\bf p} \\ & & & & \\ \hline {\bf p} \\ & & & & \\ \hline {\bf p} \\ & & & & \\ \hline {\bf p} \\ & & & & \\ \hline {\bf p} \\ & & & & \\ \hline {\bf p} \\ & & & & \\ \hline {\bf p} \\ & \\ \hline \hline \hline \hline {\bf p} \\ \hline $ | eage | |
| $ \begin{array}{ c c c c c c c c c c c c c c c c c c c$ | 50-60 KV46 | Great grandmother of Tutankhamun, mother of Tiye |
|) F 50 M ≥ 50 M ≥ 50 M $35-45$ M 19 M 19 M 19 M 13 M 2 M 2 M 2 M $35-45$ M 2 M 2 M 2 M 2 M 3 M 2 M 2 M 2 M 2 M 2 M 3 M 2 M 2 M 3 M 3 | 50-60 KV46 | Great grandfather of Tutankhamun, father of Tiye |
| M ≥50 M ≥50 F 25-35 M 35-45 M 19 M 19 M 19 M ≈30 F ≈50 | 50 KV35 | Grandmother of Tutankhamun, mother of KV35YL & |
| M ≥50 aten) M 35-45 M 19 M 19 P ≈30 F ≈50 | | Akhenaten |
| | ≥50 KV35 | Grandfather of Tutankhamun, father of KV35YL & |
| - E 25-35 Akhenaten) M 35-45 namun M 19 I Group CCG61065 M ≈30 F ≈50 | | Akhenaten |
| Akhenaten) M 35-45 A 35-55 A | 25-35 KV35 | Mother of Tutankhamun |
| namun M 19 I Group CCG61065 M ≈30 F ≈50 | 35-45 KV55 | Father of Tutankhamun |
| I Group CCG61065 M ≈30 F ≈50 | 19 KV62 | Son of Akhenaten and KV35YL |
| CCG61065 M ≈30 F ≈50 | | |
| - ≈50 | ≈30 TT320 | Thought to be Thutmose I, unidentified 18th dynasty royal |
| (orig | ≈50 KV60 | Royal wet nurse of Hatshepsut |
| | (original tomb KV20) | |
| Hatshepsut F ≈50 | ≈50 KV60 | Daughter of Thutmose I, early 18 th dynasty royal |
| (orig | (original tomb KV20) | |

Table 1. Characterization of royal mummies under investigation

| Name | Primer Pair | Sequence (5'-3') | Annealing Temperature | Product Size(bp) |
|------|-------------|--------------------------|--------------------------|---------------------|
| m10 | L-15931 | ACCAGTCTTGTAAACCGGAG | 59°C | 250 |
| | H-16141 | ATGTACTACAGGTGGTCAAG | | |
| m8 | L-16056 | GAAGCAGATTTGGGTACCAC | 59°C | 220 |
| | H-16232 | GCTTTGGAGTTGCAGTTGATGTGT | | |
| m24 | L- 16059 | AAGCAGATTTGGGTACCACCCA | TDP*62-57-52 | 97 |
| | H-16111 | GGTACCGTACAATATTCATGGTG | (5,5,30) | |
| m3 | L-16118 | TACATTACTGCCAGCCACCAT | 62°C | 159 |
| | H-16232 | GCTTTGGAGTTGCAGTTGATGTGT | | |
| m11 | L-16118 | TACATTACTGCCAGCCACCAT | 55°C | 225 |
| | H-16302 | TGGCTTTATGTACTATGTAC | | |
| m4 | L-16118 | TACATTACTGCCAGCCACCAT | 59°C | 140 |
| | H-16217a | TGTGTGATAGTTGAGGGTTG | | |
| m18 | L-16126 | CTGCCAGCCACCATGAATATTG | 60°C | 185 |
| | H-16266 | TAGGTTTGTTGGTATCCTAGTGG | | |
| m23 | L-16129 | GCCAGCCACCATGAATATTGTAC | TDP62-57-52 | 134 |
| | H- 16217b | TGATGTGTGATAGTTGAGGGTTG | (5,5,30) | |
| m26 | L-16129 | GCCAGCCACCATGAATATTGTAC | 64°C | 150 |
| | H-16232 | GCTTTGGAGTTGCAGTTGATGTGT | | |
| m12 | L-16171 | CCACCTGTAGTACATAAAAACCCA | 63°C | 109 |
| | H-16232 | GCTTTGGAGTTGCAGTTGATGTGT | | |
| m6 | L-16210 | CCCCATGCTTACAAGCAAGT | 52°C | 132 |
| | H-16302 | TGGCTTTATGTACTATGTAC | | |
| m5 | L-16210 | CCCCATGCTTACAAGCAAGT | 62°C | 178 |
| | H-16347 | ATGGGGACGAGAAGGGATTTG | | |
| m17 | L-16211 | CTCCCCATGCTTACAAGCAGTA | 60°C | 100 |
| | H-16266 | AGGTTTGTTGGTATCCTAGTGG | | |
| m22 | L- 16226 | GCAAGTACAGCAATCAACCCTC | TDP62-57-52 | 109 |
| | H- 16290 | GTACTATGTACTGTTAAGGGTGG | (5,5,25) | |
| m7 | L-16288 | CACTAGGATACCAACAAACC | 62°C | 161 |
| | H-16409a | GCGGGATATTGATTTCACGG | | |
| m25 | L- 16314 | CCACCCTTAACAGTACATAGTAC | TDP62-57-52 | 140 |
| | H-16409b | GTGCGGGATATTGATTTCACGG | (5,5,25) | |

*TDP – Touchdown protocol

Table 2. Mitochondrial primers used for PCR amplification

and aliquots of the extracted DNA. An alternative reaction mix, consisting of 100 mM dNTPs (25 mM each dNTP) and 0.125–0.5 µl of Herculase® II Fusion DNA polymerase solution with the 1X Stratagene buffer, was used for the aDNA samples that were not successfully amplified using the AmpliTaq Gold reaction mix. The amplification cycling conditions were: 94°C for 5 minutes, then 35 cycles of 94°C for 30 seconds, 52–63°C for 30 seconds, and 72°C for 30 seconds, followed by 72°C for 10 minutes as a final extension, using a GeneAmp® PCR System 9700 Thermocycler (Applied Biosystems).

Before proceeding with the cloning steps, excision and purification of the gel bands of specific sizes were executed using the Qiaquick® Gel Extraction Kit according to the manufacturer's

instructions. Cloning of the PCR products was carried out using the TOPO TA Cloning® Kit and One Shot MAX Efficiency DH5α-T1 chemically competent *E. coli* cells (Invitrogen), following the standard protocols. Successful ligation and transformation were detected by the blue-white selection. Colony PCR was performed in a reaction mix of total volume 50µl, using the universal M13 primers, forward 5'-GTAAAACGACGGCCAG-3' and reverse 5'-CAGGAAACAGCTATGAC-3' (Eurofins MWG Operon). The cycling protocol was 94°C for 10 minutes, and then 25 cycles of 94°C for 1 minute, 55°C for 1 minute, 72°C for 1 minute, with a final extension at 72°C for 10 minutes.

For Sanger sequencing, ExoSAP-IT® (USB) was used for DNA purification. Cyclic sequencing for the colony PCR was carried out using a reaction mix containing 10 pmol of each M13 primer and 1X BigDye® Terminator v3.1 Cyclic Sequencing Kit (Applied Biosystems). The cycling conditions were 96°C for 1 minute, then 25 cycles of 96°C for 10 seconds, 50°C for 5 seconds and 60°C for 4 minutes. Sodium dodecyl sulphate (Phytotechnology laboratories) was added to each PCR product, at a final concentration of 0.2% for further incubation at 98°C for 5 minutes, then 25°C for 10 minutes and Centrisep columns were used for the filtration of the product prior to sequencing. A Genetic Analyzer (ABI Prism3130, Applied Biosystems) was used for the electrophoresis of purified amplicons, according to standard protocols.

Diagnostic mutations in the mitochondrial sequences were determined using the majority rule. Sequence alterations that occurred with a frequency of more than 75% were defined as diagnostic mutations. Positions thought to be prone to post-mortem damage and artefacts were omitted from subsequent analyses (Gilbert et al. 2003, 32–47; Pusch and Bachmann 2004, 957–964; Gilbert et al. 2007, 1–10).

Bioinformatic analysis:

The Whit Athey's haplogroup predictor (Athey 2006, 34–39) [http://www.hprg.com/hapest5] was used to calculate percentages of the suggested Y-chromosomal haplotypes of the male mummies, (KV55/Akhenaten, Tutankhamun, Amenhotep III, Yuya, TT320-Cairo CG 61065) and to define the haplogroups of the ancient and modern control individuals.

Matrilineal inheritance of mtDNA and characteristic single nucleotide polymorphisms (SNPs) of the royal family under investigation were determined using Lasergene (SeqMan and EditSeq) version 8.0 (DNAstar, Madison, Wisconsin). Mitochondrial DNA haplogroups were determined using the van Oven and Kayser phylogenetic tree (http://www.phylotree.org – mtDNA tree 2014; van Oven and Kayser 2009, E386–E394) and the Genebase mtDNA Haplogroup Reference Guide (www.genebase.com).

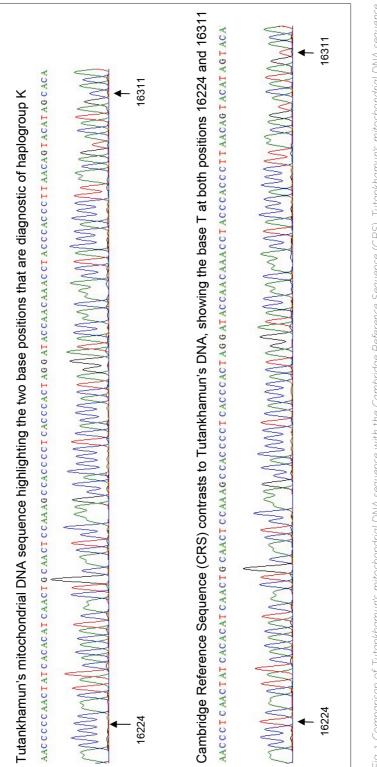


Fig. 1. Comparison of Tutankhamun's mitochondrial DNA sequence with the Cambridge Reference Sequence (CRS). Tutankhamun's mitochondrial DNA sequence highlighting the two base positions that are diagnostic to haplogroup K. Cambridge Reference Sequence (CRS) contrasts to Tutankhamun's DNA, showing the base T at both positions 16224 and 16311

| | VIIImmies | 2 |
|---|---------------|---|
| 2 | Α | |
| | UDSOUI | ,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,, |
| • | | |
| | Y-chromosome | |
| • | | |
| | ROV2 | |
| | 5 | 5 |

| Sample No. | DYS | SYO | DΥS | DYS DYS DYS | DYS | Sλα | DYS | DYS | DYS | DYS | DYS DYS I | DYS | DYS YGATA | DΥS | DΥS | DYS | Predicted |
|---------------|-----|------|-----|-------------|-----|-----|--------|-----|-----|-----|-----------|-----|-----------------|-----|-----|-----|-----------|
| 0 | 456 | 3891 | 390 | 38911 | 458 | 19 | 385a,b | 393 | 391 | 439 | 635 | 392 | H4 | 437 | 438 | 448 | haplotype |
| Yuya | 15 | 14 | _ | 29 | | | | | 1 | 1 | 20 | | 11 . | | - | 24 | G2a* |
| Amenhotep III | 15 | 13 | 24 | 30 | 16 | _ | -,14 | 13ª | 11 | | | | 11 ^a | 14 | | | R1b** |
| KV55 | 15 | 13 | 24 | 30 | 16 | 14 | 11,14 | 13ª | 11 | 10 | 23 | 13 | 11 ^a | 14 | 12 | 19 | R1b* |
| (Akhenaten) | | | | | | | | | | | | | | | | | |
| Tutankhamun | 15 | 13 | 24 | _ | 16 | | 11,- | 13ª | 11 | | 23 | | 11 ^a | 14 | | - | R1b** |

b- Reference Mummies

| Sample No. | DYS | DYS | DYS | DYS | DΥS | DYS | DYS | DYS | DYS | DYS | DYS | DYS | YGATA | DYS | DYS | DYS | Predicted |
|---------------------------|-----|------|-----|-------|-----|-----|------------|---------|-----|-----|-----|-----|-------|-----|-----|-----|------------------------|
| | 456 | 3891 | 390 | 38911 | 458 | 19 | 385a,b 393 | 393 | 391 | 439 | 635 | 392 | H4 | 437 | 438 | 448 | haplotype ¹ |
| TT320- | 15 | 12 | 19 | _1 | 18 | 13 | | 0a B | | 8 | 20 | | Da | 15 | | | *. |
| CCG61065 | | | | | | | | | | | | | | | | | |
| Ramesses III ^b | 13 | 13 | 21 | 33 | | 19 | 20 | 00 | | | | 17 | 13 | 14 | 10 | 20 | E1b1a* |
| Unknown ManE ^b | 13 | 13 | 21 | 33 | | 19 | 20 | 00 | | | | 17 | 13 | 14 | 10 | 20 | E1b1a* |

c- DNA Controls

| Sample No. | DYS | DYS | DYS | DYS | DYS | DYS | DYS | DYS | DYS | DYS | DYS | | DYS YGATA | DYS | DYS | DYS | Predicted |
|------------|-----|------|-----|-------|-----|-----|--------|-----|-----|-----|-----|-----|-----------|-----|-----|-----|------------------------|
| | 456 | 3891 | 390 | 389II | 458 | 19 | 385a,b | 393 | 391 | 439 | 635 | 392 | H4 | 437 | 438 | 448 | haplotype ¹ |
| DNA 007 | 15 | 13 | 24 | 29 | 17 | 15 | 11,14 | 13 | 11 | 12 | 24 | 13 | 13 | 15 | 12 | 19 | R1b*** |
| Staff 1 | 16 | 13 | 24 | 29 | 15 | 14 | 11,11 | 12 | 11 | 12 | 23 | 13 | 12 | 15 | 12 | 19 | R1b*** |
| Staff 2 | 14 | 11 | 24 | 27 | 19 | 13 | 13,17 | 12 | 10 | 12 | 25 | 11 | 11 | 14 | 11 | 19 | Q*** |
| Staff 3 | 16 | 13 | 24 | 29 | 15 | 14 | 11,11 | 12 | 11 | 12 | 23 | 13 | 12 | 15 | 12 | 19 | R1b*** |
| | | | | | | | | | | | | | | | | | |

Note: All female mummies were also tested with Y-chromosomal markers and did not yield any positive results. ^a Data already published in Hawass *et al.* (2010)

99.9%, eqp. 99.6%), the G2a hg (meditp: 94.9%, eqp. 89.2%), the L hg (meditp: 92.9%, eqp. 98.4%) and E1b1a hg (meditp: 86.7%, eqp. 98.3%). ** Since the weights of the various alleles and loci in haplogroup prediction are apparently unequal, the Amenhotep III-KV55-Tutankhamun male lineage haplotype was predicted based on the complete KV55 data rather than the partial data of Tutankhamun and Amenhotep III which gave different probabilities. Amenhotep III partial profile gave (meditp: R1a: 55.5%, R1b: 31.5% and E1b1b: 12.5%; eqp: R1a: 84.1%, R1b: 9.3% and E1b1b: 5.4%), while that of Tutankhamun gave (meditp: R1b: 71.7%, R1a: 28% and E1b1b: 0.0%; eqp: R1a: 82.9% and E1b1b: 0.3% and E1b1b: 5.4%). *The haplogroup (hg) prediction was carried out through the website: www.hprg.com/hapest5/, by matching the concluded ancient Y-STR profiles versus those of the Mediterranean populations (meditp) as well as of the equal priors (eqp) selection (Athey 2006, 34-39). The predicted hg was suggested based on the probability that is more than 85%. The actual probabilities for the detected 4 ancient haplotypes were for the R1b hg (meditp) selection. All probabilities exceeded 95%.

Table 3. Y-chromosome data of male mummies and DNA controls

Results

Y-chromosomal data:

The Y-chromosomal data of the 7 male mummies are collectively shown in Table 3. Those subjects included 4 mummies from the Tutankhamun family group, and the father-son pair from another dynasty. One complete (KV55-Akhenaten) and 6 partial profiles (Amenhotep III, Tutankhamun, Yuya, TT320, Ramesses III, Unknown Man E) were obtained for the tested male subjects. On the other hand, all female mummies repeatedly tested negative.

Using the Whit Athey haplogroup predictor, the most probable haplogroups obtained for the Tutankhamun family members were G2a for Yuya and R1b for KV55. Due to the incomplete profiles observed for Tutankhamun and Amenhotep III, analysis led to different probability figures, despite their concordant allele results. The haplogroup for these two mummies was thus predicted based on the full KV55 data, particularly since the relationships were confirmed through a previous study (Hawass et al. 2010, 638–647). The mummies that did not belong to the Tutankhamun lineage showed other haplotype probabilities (L and E1b1a). It should be noted that the identified R1b haplotype in Tutankhamun, KV55 and Amenhotep III was different from those detected in the kit control DNA 007 as well as in the staff members (1 and 3) as the profiles differed in several loci.

Mitochondrial data:

The mtDNA of 9 individuals was analyzed; 7 belonged to the Tutankhamun family (4 males, 3 females) and 2 females who were not (Table 4). Using PCR primers specifically designed to amplify short mtDNA fragments, partial analysis of the HVRI (16056–16409) of the D loop control region was done. Diagnostic mutations in the sequences from the tested mummies were determined using the majority rule. The total number of sequences obtained to characterize the HVRI region was 631. Considering the alignment data from Thuya (generation I in the family), 87 sequences were generated in total using 6 bone biopsies and 9 different primer combinations. Position 16224 was covered by 36 sequences from 10 different PCR products using 4 primer combinations. Of these 36 sequences, 32 showed a C and 4 showed a T residue. According to the majority rule and in the absence of contaminant sequences, this defines C to be the authentic base at mtDNA position 16224. Moreover, position 16311 was covered by 38 sequences in total and derived from 9 PCR products using 3 different combinations; 35 showed a C at this position, while 2 showed a T and 1 showed an A.

Likewise for Tutankhamun (generation IV), 79 sequences were obtained in total using 4 bone biopsies and 8 different primer combinations. Position 16224 was covered by 51 sequences in total from 10 different PCR products using 5 different combinations. Out of 51 sequences, 49 showed a C, while 2 showed a T. Position 16311 was covered by 51 sequences in total from 10 different PCR products using 3 different primer combinations. Out of these 51 sequences, 50 showed a C

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

a- Royal Mitochondrial Lineage Mummies

b- Control Mummies

| 0 00110 | ormunnies | | | | |
|---------------|--|--------------------------|---------------|----------|----------------|
| Mummy | HVRI Diagnostic Mutation*, | Predicted | Sequences | Min/ Max | Coverage |
| (tomb) | (Frequency, No. of Trials [†]) | Haplogroups [#] | per Alignment | Coverage | |
| Yuya | 16224 (80% - 2 exps) | K | 64 | 2/58 | Full |
| (KV46) | 16311 (89% - 3 exps) | | | | |
| Amenhotep III | 16311 (100% - 1 exp) | H2b | 80 | 17/47 | Partial 16118- |
| (KV35) | M ~ 10 | | | | 16409 |
| Sitra-In | None | K excluded | 18 | - | Partial 16118- |
| (KV60) | | | | | 16232 |
| Hatshepsut | None | K excluded | 20 | - | Partial 16118- |
| (KV60) | | | | | 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

while only 1 showed a T. Using the same rule, this also defines C to be the authentic base at positions 16224 and 16311, respectively.

The two diagnostic mutations observed in Tutankhamun's DNA sequences (Fig. 1) are indicative of the mitochondrial haplogroup K according to the van Oven and Kayser (2009, E386–E394) phylogenetic tree (http://www.phylotree.org) and the Genebase mtDNA Haplogroup Reference Guide (www.genebase.com). The haplogroup data for the other individuals is listed in Table 4. Based on the occurrence of the two diagnostic mutations 16224 and 16311, six members of the

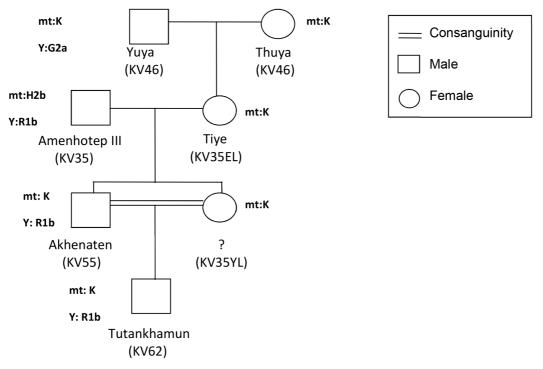


Fig. 2. Family pedigree depicting mitochondrial and Y chromosome haplogroups

Tutankhamun family group were characterized as haplogroup K, while one was characterized with haplogroup H2b. For the two KV60 reference mummies, a small PCR product encompassing 16118 to 16232 was amplified and sequenced but it did not show the 16224 diagnostic mutation associated with haplogroup K.

A pedigree representing the generated mitochondrial and Y-chromosome haplogroups of the investigated Tutankhamun family members is shown in Fig. 2.

Discussion

Implications from Y-chromosomal DNA analysis:

Haplogroup R1b

The proposed haplogroup for three of the investigated mummies, namely Tutankhamun, KV55 (Akhenaten) and Amenhotep III, was R1b. The R1b haplogroup is defined by the mutation M343 and thought to originate in the western Asian region (Wood et al. 2005, 867–876; Cruciani et al. 2010, 800–807; Myres et al. 2011, 95–101; Klyosov 2012, 87–105). R1b is not abundant in contemporary

Africa (Cruciani et al. 2010, 800–807), yet it is geographically widespread among the European populations making up to 50–60% of the genetic pool of the modern Europeans (Arredi et al. 2004, 338–345; Cinnioğlu et al. 2004, 127–148; Karafet et al. 2008, 830–838; Myres et al. 2011, 95–101; Klyosov 2012, 87–105). The introduction of this Y-chromosomal signature may be due to old introgression of lineages to the European gene pool from the Near East during the Neolithic spread of farming (Myres et al. 2011, 95–101). The phylogeographic analysis of the R1b haplogroup distribution provided strong support to the back-migration theory from Asia to Africa (Cruciani et al. 2010, 800–807). The R1b group was estimated to have an age of ca. 16,000 years before present (YBP) (Klyosov 2012, 87–105), and its suggested expansion time nearly approximates that of the G2a haplogroup (Cinnioğlu et al. 2004, 127–148). Less than 7% of the contemporary Egyptians share the R1b Y-chromosomal fingerprint (Cruciani et al. 2010, 800–807; el-Sibai et al. 2009, 568–581).

Haplogroup G2a

The great-grandfather of Tutankhamun, Yuya, carries a Y-chromosomal signature that could be assigned to the haplogroup G2a. Haplogroup G is an F-affiliated clade (Luis et al. 2004, 532–544; Wood et al. 2005, 867–876), and it is defined by the mutation M201 (Cinnioğlu et al. 2004, 127–148; Luis et al. 2004, 532–544; Wood et al. 2005, 867–876; Karafet et al. 2008, 830–838). This clade is not globally abundant, and its prevalence is mainly in the Middle East (highest in Druze), the Mediterranean basin and Caucasus Mountains where it exhibits its maximum frequency (Cinnioğlu et al. 2004, 127–148; Karafet et al. 2008, 830–838; Balanovsky et al. 2011, 18255–18259; Lacan et al. 2011, 2905–2920). The pattern of this haplogroup distribution in the Caucasus suggests a Near Eastern origin (Cinnioğlu et al. 2004, 127–148; Balanovsky et al. 2011, 18255–18259). The genetic share of the F-affiliated groups (G, H, I, J) is around 40% of the modern Egyptians, with G-M201 representing approximately 9% of the population (Luis et al. 2004, 532–544).

Haplogroup L

One of the male control mummies, an unidentified 18th Dynasty royal, previously thought to be Thutmose I (TT320-Cairo CG 61065), was proposed to belong to haplogroup L. This haplogroup is mainly defined by the M20 mutation, with additional markers to further confirm the clade definition (Sengupta et al. 2006, 202–221). It is found at low frequencies in some parts of the Middle East and Mediterranean Europe, and is mainly observed in southern, western and central Asia (highest in Indian subcontinent) (Kivisild et al. 2003, 313–332; Cordaux et al. 2004, 231–235; Sengupta et al. 2006, 202–221; Thanseem et al. 2006, 42; Karafet et al. 2008, 830–838). In Y-chromosome studies on modern Egyptians, no haplogroup L findings were reported (Arredi et al. 2004, 338–345; Luis et al. 2004, 532–544; el-Sibai et al. 2009, 568–581).

Haplogroup E1b1a

The father-son pair, Ramesses III and Unknown Man E/Pentawere, was found to have the haplogroup E1b1a (Hawass et al. 2012, e8268) that shows its highest frequencies in modern populations from

West Africa (~80%) and Central Africa (~60%). It is less than 10% among North Africans and nearly absent in East Africa, which is where it was hypothesized to have originated (Trombetta et al. 2011, e16073). Alternatively, an earlier origin of around 20,000–30,000 YBP, was hypothesized in West Africa due to its very high diversity and frequency in that location (Rosa et al. 2007, 124). Although the frequency of haplogroup E among contemporary Egyptians is 39.5%, that of the E1b1 subclade was only 1.4% (Cruciani et al. 2004, 1014–1022; Luis et al. 2004, 532–544; el-Sibai et al. 2009, 568–581).

Implications from mitochondrial DNA:

Haplogroup K

Haplogroup K most likely belonging to a Near Eastern lineage (Brandstätter et al. 2008, 191; Fernández et al. 2014, e1004401), is a subcluster of the macrohaplogroup U (Richards et al. 1998, 241–260; Kivisild et al. 1999, 1331–1334; Macaulay et al. 1999, 232–249; Maca-Meyer et al. 2001, 13). The age estimate of the K cluster is ~15,500–25,500 YBP (Richards et al. 2000, 1251–1276). The K haplogroup has been found to be present in 4.7% of contemporary Egyptian populations (Saunier et al. 2009, e97–e103).

Haplogroup H

The haplogroup of one of the studied mummies (Amenhotep III) was proposed as being H2b. Haplogroup H displays a sequence in the hypervariable region I (HVRI) identical to that of the Cambridge Reference Sequence (CRS) (Torroni et al. 1996, 1835–1850; Macaulay et al. 1999, 232–249; Richards et al. 2000, 1251–1276; van Oven and Kayser 2009, e386–e394). It is believed that the HVRI 16311 transition is an emerging subclade of the H-CRS and named as H2b (Roostalu et al. 2007, 436–448; Brandstätter et al. 2008, 191). The age estimate of cluster H in Europe can be dated to 19,200–21,400 YBP; however, it is older in the Near East and can be dated back to 23,200–28,400 YBP. Haplogroup H is substantially prevalent in Europe as it is in the Near East but with a lower frequency (Torroni et al. 1996, 1835–1850; Richards et al. 2000, 1251–1276; Tambets et al. 2000, 219–235; Malyarchuk et al. 2008, 1651–1658). This indicates that, although this cluster might have its roots in the Middle East, the significant evolution of this mitochondrial signature occurred in Europe (Richards et al. 1996, 185–203), probably in the late Upper Paleolithic expansions (14 500 YBP) (Richards et al. 1998, 241–260; Richards et al. 2000, 1251–1276). The frequency of haplogroup H was, respectively, 3.6 and 5.94% among modern Egyptians (Saunier et al. 2009, e97–e103; Elmadawy et al. 2013, 338–341).

Concluding remarks:

This study reconfirms and complements the previously published family pedigree of the King Tutankhamun (Hawass et al. 2010, 638–647). It also presents the plausible mitochondrial and Y-chromosomal signatures of the family members. The data suggest geographical and temporal

matching of the origins of Y-chromosomal and the mitochondrial signatures in the family to possibly Near Eastern lineages, with time coverage believed to be between ~14,500 and 28,400 YBP. A recent study by Schuenemann et al. (2017, 15694) has also shown the propensity of Near Eastern lineages among their ancient Egyptian sample coming from Faiyum, although these were from later ancient periods. A distinction between the geographical origins of the investigated late 18th Dynasty family and the succeeding 20th Dynasty pharaohs (Ramesses III and Unknown Man E/Pentawere) was also shown. The latter was estimated to be of African origin, not a Middle or Near Eastern one.

Due to its geographical location, Egypt could have over time created a melting pot of various gene pools. The genetic variation outside Africa reflects only a subset of the African gene pool (Underhill and Kivisild 2007, 539–564). Studies on today's Egyptian population samples have resulted in haplogroups that are predominantly European or west Eurasian (Stevanovitch et al. 2003, 23–39; Saunier et al. 2009, e97–e103; Elmadawy et al. 2013, 338–341; Fadhlaoui-Zid et al. 2013, e80293). It was also reported that the modern Egyptian haplogroups in a study were more closely associated with non-African populations than with the West Africans (Pagani et al. 2015, 986–991). This accords with our findings that the same trend was indicated through the analysis of 18th Dynasty family members.

Conflict of Interest (COI):

The authors declare no conflict of interest to the submitted work.

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