(BACK)

mthap version 0.19b (2015-05-11); haplogroup data version PhyloTree Build 17 (2016-02-18) +mods raw data source SNP raw v4 Full 20170514175358.txt (14MB)

Found 3270 markers at 3268 positions covering 19.7% of mtDNA.

Markers found (shown as differences to rCRS):

HVR2: 152C 263G

CR: 750G 1438G 4769G 8860G

HVR1:

IMPORTANT NOTE: The above marker list is almost certainly incomplete due to limitations of genotyping technology and is not comparable to mtDNA sequencing results. It should not be used with services or tools that expect sequencing results, such as mitosearch.

Best mtDNA Haplogroup Matches:

1) H(T152C)

Defining Markers for haplogroup H(T152C):

HVR2: 152C 263G

CR: 750G 1438G 4769G 8860G 15326G

HVR1:

Marker path from rCRS to haplogroup H(T152C):

 $\mathbf{H2a2a1(rCRS)} \Rightarrow 263\mathbf{G} \Rightarrow \mathbf{H2a2a} \Rightarrow 8860\mathbf{G} \ 15326\mathbf{G} \Rightarrow \mathbf{H2a2} \Rightarrow 750\mathbf{G} \Rightarrow \mathbf{H2a} \Rightarrow 4769\mathbf{G} \Rightarrow \mathbf{H2} \Rightarrow 1438\mathbf{G} \Rightarrow \mathbf{H} \Rightarrow 152\mathbf{C} \Rightarrow \mathbf{H(T152C)}$

Imperfect Match. Your results contained differences with this haplogroup:

Matches(6): 152C 263G 750G 1438G 4769G 8860G

Untested(1): 15326

2) H1(T152C)

Defining Markers for haplogroup H1(T152C):

HVR2: 152C 263G

CR: 750G 1438G 3010A 4769G 8860G 15326G

HVR1:

Marker path from rCRS to haplogroup H1(T152C):

 $\mathbf{H2a2a1(rCRS)} \Rightarrow 263G \Rightarrow \mathbf{H2a2a} \Rightarrow 8860G \ 15326G \Rightarrow \mathbf{H2a2} \Rightarrow 750G \Rightarrow \mathbf{H2a} \Rightarrow 4769G \Rightarrow \mathbf{H2} \Rightarrow 1438G \Rightarrow \mathbf{H} \Rightarrow \mathbf{3010A} \Rightarrow \mathbf{H1} \Rightarrow 152C \Rightarrow \mathbf{H1(T152C)}$

Imperfect Match. Your results contained differences with this haplogroup:

Matches(6): 152C 263G 750G 1438G 4769G 8860G

No-Calls(1): 3010A Untested(1): 15326

3) H

Defining Markers for haplogroup H:

HVR2: 263G

CR: 750G 1438G 4769G 8860G 15326G

HVR1:

Marker path from rCRS to haplogroup H (plus extra markers):

 $\mathbf{H2a2a1(rCRS)} \Rightarrow 263G \Rightarrow \mathbf{H2a2a} \Rightarrow 8860G \ 15326G \Rightarrow \mathbf{H2a2} \Rightarrow 750G \Rightarrow \mathbf{H2a} \Rightarrow 4769G \Rightarrow \mathbf{H2} \Rightarrow 1438G \Rightarrow \mathbf{H} \Rightarrow \mathbf{152C}$

Imperfect Match. Your results contained differences with this haplogroup:

Matches(5): 263G 750G 1438G 4769G 8860G

Extras(1): 152C Untested(1): 15326

3) H9

Defining Markers for haplogroup H9:

HVR2: 152C 263G

CR: 750G 1438G 4769G 8860G 13020C 15326G

HVR1:

Marker path from rCRS to haplogroup H9:

 $\mathbf{H2a2a1}(\mathbf{rCRS}) \Rightarrow 263G \Rightarrow \mathbf{H2a2a} \Rightarrow 8860G \ 15326G \Rightarrow \mathbf{H2a2} \Rightarrow 750G \Rightarrow \mathbf{H2a} \Rightarrow 4769G \Rightarrow \mathbf{H2} \Rightarrow 1438G \Rightarrow \mathbf{H} \Rightarrow 152C \Rightarrow \mathbf{H}(\mathbf{T152C}) \Rightarrow \mathbf{13020C} \Rightarrow \mathbf{H9}$

Imperfect Match. Your results contained differences with this haplogroup:

Matches(6): 152C 263G 750G 1438G 4769G 8860G

Mismatches(1): 13020T Untested(1): 15326

3) H52

Defining Markers for haplogroup H52:

HVR2: 152C 263G

CR: 750G 1438G 4769G 8860G 14220G 15326G

HVR1:

Marker path from rCRS to haplogroup H52:

 $\mathbf{H2a2a1(rCRS)} \Rightarrow 263G \Rightarrow \mathbf{H2a2a} \Rightarrow 8860G \ 15326G \Rightarrow \mathbf{H2a2} \Rightarrow 750G \Rightarrow \mathbf{H2a} \Rightarrow 4769G \Rightarrow \mathbf{H2} \Rightarrow 1438G \Rightarrow \mathbf{H} \Rightarrow 152C \Rightarrow \mathbf{H(T152C)} \Rightarrow \mathbf{14220G} \Rightarrow \mathbf{H52} \Rightarrow \mathbf{H53} \Rightarrow \mathbf{H53} \Rightarrow \mathbf{H53} \Rightarrow \mathbf{H53} \Rightarrow \mathbf{H54} \Rightarrow \mathbf{H54} \Rightarrow \mathbf{H55} \Rightarrow \mathbf{$

Imperfect Match. Your results contained differences with this haplogroup:

Matches(6): 152C 263G 750G 1438G 4769G 8860G

Mismatches(1): 14220A Untested(1): 15326

3) H69

Defining Markers for haplogroup H69:

HVR2: 152C 263G

CR: 750G 1438G 4646C 4769G 8860G 15326G

HVR1:

Marker path from rCRS to haplogroup H69:

 $\mathbf{H2a2a1}(\mathbf{rCRS}) \Rightarrow 263G \Rightarrow \mathbf{H2a2a} \Rightarrow 8860G \ 15326G \Rightarrow \mathbf{H2a2} \Rightarrow 750G \Rightarrow \mathbf{H2a} \Rightarrow 4769G \Rightarrow \mathbf{H2} \Rightarrow 1438G \Rightarrow \mathbf{H} \Rightarrow 152C \Rightarrow \mathbf{H}(\mathbf{T152C}) \Rightarrow 4646C \Rightarrow \mathbf{H69}$

Imperfect Match. Your results contained differences with this haplogroup:

Matches(6): 152C 263G 750G 1438G 4769G 8860G

Mismatches(1): 4646T Untested(1): 15326

3) H16(T152C)

Defining Markers for haplogroup H16(T152C):

HVR2: 152C 263G

CR: 750G 1438G 4769G 8860G 10394T 15326G

HVR1:

Marker path from rCRS to haplogroup H16(T152C):

 $\mathbf{H2a2a1(rCRS)} \Rightarrow 263G \Rightarrow \mathbf{H2a2a} \Rightarrow 8860G \ 15326G \Rightarrow \mathbf{H2a2} \Rightarrow 750G \Rightarrow \mathbf{H2a} \Rightarrow 4769G \Rightarrow \mathbf{H2} \Rightarrow 1438G \Rightarrow \mathbf{H} \Rightarrow \mathbf{10394T} \Rightarrow \mathbf{H16} \Rightarrow 152C \Rightarrow \mathbf{H16(T152C)}$

Imperfect Match. Your results contained differences with this haplogroup:

Matches(6): 152C 263G 750G 1438G 4769G 8860G

Mismatches(1): 10394C Untested(1): 15326

3) H3(T152C)

Defining Markers for haplogroup H3(T152C):

HVR2: 152C 263G

CR: 750G 1438G 4769G 6776C 8860G 15326G

HVR1:

Marker path from rCRS to haplogroup H3(T152C):

 $\mathbf{H2a2a1(rCRS)} \Rightarrow 263\mathbf{G} \Rightarrow \mathbf{H2a2a} \Rightarrow 8860\mathbf{G} \ 15326\mathbf{G} \Rightarrow \mathbf{H2a2} \Rightarrow 750\mathbf{G} \Rightarrow \mathbf{H2a} \Rightarrow 4769\mathbf{G} \Rightarrow \mathbf{H2} \Rightarrow 1438\mathbf{G} \Rightarrow \mathbf{H} \Rightarrow \mathbf{6776C} \Rightarrow \mathbf{H3} \Rightarrow 152\mathbf{C} \Rightarrow \mathbf{H3}(\mathbf{T152C})$

Imperfect Match. Your results contained differences with this haplogroup:

Matches(6): 152C 263G 750G 1438G 4769G 8860G

Mismatches(1): 6776T Untested(1): 15326

3) H46

Defining Markers for haplogroup H46:

HVR2: 152C 263G

CR: 750G 1438G 2772T 4769G 8860G 15326G

HVR1:

Marker path from rCRS to haplogroup H46:

 $\mathbf{H2a2a1(rCRS)} \Rightarrow 263G \Rightarrow \mathbf{H2a2a} \Rightarrow 8860G \ 15326G \Rightarrow \mathbf{H2a2} \Rightarrow 750G \Rightarrow \mathbf{H2a} \Rightarrow 4769G \Rightarrow \mathbf{H2} \Rightarrow 1438G \Rightarrow \mathbf{H} \Rightarrow 152C \Rightarrow \mathbf{H(T152C)} \Rightarrow \mathbf{2772T} \Rightarrow \mathbf{H46C} \Rightarrow \mathbf{$

Imperfect Match. Your results contained differences with this haplogroup:

Matches(6): 152C 263G 750G 1438G 4769G 8860G

Mismatches(1): 2772C Untested(1): 15326

Need help?

First, please check the Frequently Asked Questions for guidance on how to read this report. If you still have questions, there is a discussion about mthap on eng.molgen.org. You can also email your questions to me at james.lick@jameslick.com. So that I can best help you, please include a copy of the complete mthap report and/or your mtDNA data file in your email.