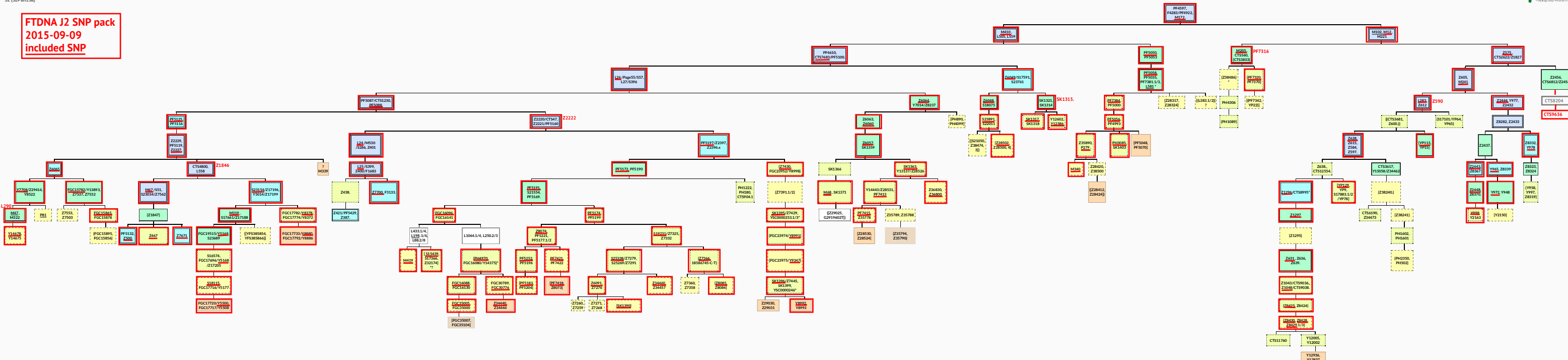


FTDNA J2 SNP pack 2015-09-09 included SNP

J2 (J2Pan15a)



Phylogeny and Haplogroup SNPs (two most prominent/good-coverage + historically informative/recurrent) based on J2-M172_spreadsheet (anonymous SNP-names not always listed). Usually listed SNPs are checked for genome-seq uniqueness with BLAT 1Mb and YFull SNP search (see Spreadsheet>Phylo-SNPs for details). Recurrent SNPs are considered to define clades when no alternative is available and they are found in more than two samples. Recurrency is shown: SNR (found once in a sample outside of Hg J), SNP.1 or SNP.2 or SNP.x (not more than twice in Hgs), SNP.1/3 (3 times recurrent).

Legend: [under research] [tentative split] [Single/Private] [placement vs parallel branches unavailable] [* upstream placement uncertain]

Priority for Haplogroups: Backbone, Skeleton, Body, Informative, Expected Informative, Low Diversity, Clon/Geographical, Diverse, Research, Sparse, 700-1400 years, Surname

Terminal samples / Listing of SNP count: FGC (Full Genomes Core), BiGy (FTDNA), Geographic Proj., Single SNP etc., M413 (1000 Genomes Project, Magoon et al 2013), PGP (Personal Genome Project), GoNL (Y-CCF 2012), Chromo2 (2014), SK14 (SkStoneking, Lippold et al 2014), PH14 (Hallast et al 2014), EB15 (Estonian Biocentre CG, Karmin et al 2015), PF15 & PF16 (Sundstrom-Forsell et al 2015, 6 2015), YFS (YFull Single Best+Acceptable).