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Were inefficient mitochondrial haplogroups selected during radiations of modern humans? – a test using modular kinetic analysis of coupling in mitochondria from cybrid cell lines.

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Short title: General test of the effects of mitochondrial DNA variants

Key words: mitochondrial DNA, human evolution, coupling efficiency, oxidative phosphorylation, cybrid, modular kinetic analysis

Abbreviations: ρ^0 , mtDNA-less; $\Delta\psi$, mitochondrial membrane potential; TPMP, triphenylmethylphosphonium; FCCP, carbonyl cyanide *p*-trifluoromethoxyphenylhydrazone.

SYNOPSIS

We introduce a general test of the bioenergetic importance of mitochondrial DNA variants: modular kinetic analysis of oxidative phosphorylation in mitochondria from cybrid cells with constant nuclear DNA but different mitochondrial DNA. We apply this test to the hypothesis [Ruiz-Pesini, Mishmar, Brandon, Procaccio and Wallace (2004) *Science* **303**, 223-226] that particular mitochondrial DNA haplogroups (specific combinations of polymorphisms) that cause lowered coupling efficiency, leading to generation of less ATP and more heat, were positively selected during radiations of modern humans into colder climates. Contrary to the predictions of this hypothesis, mitochondria from Arctic haplogroups had similar or even greater coupling efficiency than mitochondria from tropical haplogroups.

INTRODUCTION

Many mutations in mitochondrial DNA (mtDNA) lead to severe pathologies, such as LHON (Leber hereditary optic neuropathy) and MELAS (mitochondrial myopathy, encephalopathy, lactic acidosis and stroke-like episodes), although the penetrance of the pathological phenotype is often puzzling [1, 2]. The bioenergetic consequences of normal mtDNA variation are less well understood, although there are indications that such variation may affect bioenergetic function [3, 4], disease susceptibility [5, 6] and longevity [7]. Such analyses are difficult, however, because mitochondrial functions are mostly controlled by genes encoded in nuclear DNA, and it can be very difficult to be sure that small effects attributed to mtDNA variations are not confounded by different nuclear DNA backgrounds. In the present paper we use cybrids with different mtDNA complements but the same nuclear DNA background to overcome this problem [8]. We then assay bioenergetic function using the modular kinetic approach [9] to determine the effect of mtDNA sequence on mitochondrial bioenergetics. This novel application of modular kinetic analysis to cybrid mitochondria provides a powerful new tool, of general applicability, to analyse bioenergetic effects of differences in mtDNA in healthy human populations or in mitochondrial diseases.

Analysis of human mtDNA variation has identified specific combinations of polymorphisms that have been used systematically to classify mtDNAs into haplogroups, and to study the origin, radiations and evolution of human populations [10, 11]. More recently, it has been proposed that these polymorphisms interacted with environmental factors, leading to positive selection of adaptive variants with decreased mitochondrial coupling efficiency and increased heat production that were beneficial for humans moving out of Africa into colder Eurasia and through the Arctic to the Americas [12, 13]. Specifically, haplogroups A, C, D and X, which are dominant in North-eastern Eurasian and native American populations, are predicted to have lower oxidative phosphorylation coupling efficiency than the other haplogroups, particularly L, which is dominant in African populations and B, which is present in central Asian and American populations and is thought not to have been cold-selected. These haplogroups have specific polymorphisms in the mitochondrial genes *MTATP6*, *MTCYB*, *MTND2* and *MTND4*, which code for components of the protein complexes of oxidative phosphorylation, so might cause slip or leak reactions and affect coupling efficiency. The present study tests this hypothesis experimentally, to clarify whether and how oxidative

phosphorylation coupling efficiency depends on genetic variants of mtDNA generated after modern humans migrated out of Africa.

EXPERIMENTAL

Subjects

Ethics approval was obtained from the Cambridge Research Ethics Committee. We recruited healthy volunteers by advertisement and obtained written informed consent from all participants. DNA was extracted from buccal swab samples by QIAamp DNA Mini Kit (Qiagen, Hilden, Germany). Mitochondrial haplogroups were determined by PCR-RFLP (restriction fragment length polymorphism) analysis according to criteria described in [11]. Entire mtDNAs were amplified in several overlapping fragments by PCR [14]. Each fragment was digested by restriction enzymes, and resolved on agarose gels. 15 ml of blood were taken from each of six volunteers who had Arctic (A, C and D) or tropical (L1, L2 and L3) haplogroups, for platelet preparation and cybrid construction.

Cells

A549.B2 ρ^0 (mtDNA-less) cells derived from human lung carcinoma A549 were cultivated in Dulbecco's modified Eagle's medium (DMEM) containing 4.5 g/l glucose and 110 $\mu\text{g/ml}$ sodium pyruvate, supplemented with 10% foetal bovine serum (FBS) and 50 $\mu\text{g/ml}$ uridine. Platelets from the blood samples of volunteers were fused with A549.B2 ρ^0 cells as described elsewhere [15]. The resulting cybrid cell lines were routinely maintained in DMEM with 10% dialysed FBS. DNA was extracted from cultured cybrid cells as described previously [16] to confirm mitochondrial haplogroups by PCR-RFLP analysis.

Cybrid B2.3243 cells, which have the same A549 nuclear background, but harbour a pathogenic mutation at np 3243 in the tRNA-leu(UUR) gene (*MTTL1*) [17, 18] were kindly donated by Ian Holt (MRC Dunn Human Nutrition Unit). The mutation was heteroplasmic: clone Z had a high proportion of wild type and clone X had a high proportion of mutant.

Mitochondria

Human cell mitochondria were prepared from cultured cybrids cells grown in DMEM with 10% FBS. Cells were allowed to grow up to approximately 90% confluence on 3 or 4

trays (500 cm², Nunclon™). Following two washes with ice-cold STE buffer (comprising 0.25 M sucrose, 5 mM Tris-HCl (pH 7.4) and 2 mM EGTA), cells were scraped off the trays and harvested in STE by centrifugation at 480 g (10 min, 4°C). Cell pellets were resuspended in 5 ml of STE supplemented with a protease-inhibitor cocktail (Calbiochem #539131) and homogenized using a glass homogenizer. After 8-fold dilution in STE and removal of cell debris (3 min centrifugation at 1100 g followed by filtration through muslin), mitochondria were pelleted by centrifugation at 12000 g (11 min, 4°C). This procedure gave good quality mitochondria (respiratory control ratio with α -ketoglutarate + malate approximately 11) and yielded typically 5-8 mg of mitochondrial protein per 500 cm² tray. Protein concentration was determined by the biuret method with BSA as a standard.

Respiratory flux and membrane potential

Oxygen consumption was measured at 37°C using a Clark-type electrode (Rank Brothers, Cambridge, UK) calibrated with air-saturated medium comprising 0.115 M KCl, 10 mM KH₂PO₄, 3 mM Hepes (pH 7.2), 1 mM EGTA, 2 mM MgCl₂ and 0.3% (w/v) defatted BSA, assumed to contain 406 nmol atomic oxygen · ml⁻¹ [19]. No correction was made for consumption of oxygen by the electrode, so mitochondrial oxygen consumption at low rates will have been slightly overestimated. Simultaneously with respiratory activity, $\Delta\psi$ (mitochondrial membrane potential) was measured using an electrode sensitive to the lipophilic cation TPMP⁺ (triphenylmethylphosphonium) [20]. Mitochondria were incubated at 1.0 mg · ml⁻¹ (for succinate respiration) or 1.5 mg · ml⁻¹ (for α -ketoglutarate + malate respiration) in the presence of 80 ng · ml⁻¹ nigericin to collapse Δ pH and 4 μ M rotenone to inhibit complex I. The TPMP⁺-sensitive electrode was calibrated with sequential additions of TPMP⁺ up to 2 μ M, then 4 mM succinate or 3.2 mM α -ketoglutarate + 0.8 mM malate (with rotenone omitted) was added to initiate respiration. Experiments were terminated with 1.6 μ M FCCP (carbonyl cyanide *p*-trifluoromethoxyphenylhydrazone), allowing correction for any small baseline drift. $\Delta\psi$ values were calculated from the distribution of TPMP⁺ across the mitochondrial inner membrane using a binding correction factor of 0.35 mg of protein · μ l⁻¹. Respiratory control ratios (the State 3 respiration rate with 0.8 mM ADP divided by the State 4 rate with oligomycin) with α -ketoglutarate + malate as substrate were determined in the absence of nigericin.

Modular kinetics

Mitochondrial oxidative phosphorylation can be divided conceptually into three modules. These are (i) the reactions that produce $\Delta\psi$, consisting of the substrate translocases, dehydrogenases and other enzymes and the components of the respiratory chain, here called 'substrate oxidation', (ii) the reactions that consume $\Delta\psi$ and synthesize, export and dephosphorylate ATP, consisting of the ATP synthase, the phosphate and adenine nucleotide translocases and any ATPases that may be present, called the 'phosphorylating system', and (iii) the reactions that consume $\Delta\psi$ without ATP synthesis, called the 'proton leak' [21]. Oxygen consumption and $\Delta\psi$ were measured simultaneously using mitochondria incubated with 80 ng · ml⁻¹ nigericin and 4 μ M rotenone. Respiration was initiated by 4 mM succinate (or 3.2 mM α -ketoglutarate + 0.8 mM malate, with rotenone omitted). The kinetic behaviour of a ' $\Delta\psi$ -producer' can be established by specific modulation of a ' $\Delta\psi$ -consumer' and reciprocally, the kinetics of a consumer are revealed upon modulation of a producer [9]. To measure the kinetic response of proton leak to $\Delta\psi$, the state 4 (non-phosphorylation) respiration of mitochondria in the presence of oligomycin (0.8 μ g · ml⁻¹; to prevent any residual ATP synthesis), which was used solely to drive the proton leak, was titrated with malonate (up to 0.5 mM). In a similar way, the state 4 respiration was titrated by FCCP (up to 0.8 μ M) for measurement of the kinetic response of substrate oxidation to $\Delta\psi$. State 3 (maximal rate of ATP synthesis) was obtained using an ADP regenerating system (100 μ M ATP, 20 mM glucose, and 10 units · ml⁻¹ hexokinase (from baker's yeast; Sigma)). Titration of state 3 respiration with malonate (up to 1.15 mM) allowed measurement of the kinetics of the $\Delta\psi$ -consumers (the sum of the phosphorylating system and proton leak).

Catalase assay

Cells were harvested by trypsinisation, washed with ice-cold PBS, and suspended in 50 mM potassium phosphate buffer (pH 7.0) containing 0.5% Triton X-100 and 0.1mM EDTA. The cells were disrupted by brief sonication, and the supernatant was obtained by centrifugation. Catalase activity of the supernatant was measured at 37°C in triplicate as described elsewhere [22]. One unit of catalase is defined as the activity required to decompose 1 μ mol of hydrogen peroxide per min. Specific activity is expressed as units per mg protein, estimated using a BCATM protein assay kit (Pierce, Rockford, IL, USA) with BSA as a standard.

Statistics

Values are given as mean $\pm$ SEM, with n being the number of independent mitochondrial preparations (Fig. 1) or cell clones (nine for Fig. 2 and Fig. 4; three for Fig. 3). To test for significant differences in modular kinetics between haplogroups, we interpolated the mean respiration rates of mitochondria from the individual preparations or clones at the lowest individual state 4 potential (for proton leak) or state 3 potential (for the phosphorylating system) or at any potential between the highest and lowest membrane potential (for substrate oxidation). The significance of differences between means was assessed by unpaired Student's t -test using Microsoft Excel X; p values < 0.05 were taken to be significant. There were no significant differences between Arctic and tropical haplogroups for any of the data in Fig. 2 or Fig. 4.

RESULTS

Generation of cybrid cell lines

To overcome effects of different nuclear backgrounds, cybrids were used to analyse the functional effects of mtDNA. Cybrid cell lines are constructed by repopulation of mtDNA-less (ρ^0) cells with exogenous mitochondria [8]. A549.B2 ρ^0 cells derived from human lung carcinoma A549 were used in the present study. To obtain different populations of mitochondria, we recruited healthy human volunteers, determined their mitochondrial haplogroups and selected six with Arctic (A, C or D) or tropical (L1, L2 or L3) haplogroups. Platelets, which have no nuclei, were isolated from blood samples and fused with the ρ^0 cells. The resultant cybrids had mitochondrial DNA from the different donors, but their nuclear DNA was identical. After many generations of cybrid growth (diluting platelet-derived nuclear-encoded subunits), all nuclear-encoded mitochondrial protein subunits will be specified by the host cell, but all mitochondrial-encoded subunits will be specified by the donor mtDNA. Using cybrids made it possible to investigate the effects of mtDNA variation in cells with identical nuclear DNA.

Modular kinetic analysis of oxidative phosphorylation – validation of the method

Rather than arbitrarily measuring individual reactions to look for differences caused by mitochondrial DNA variants, we applied a systems approach: top-down elasticity analysis

[9]. This is better described as modular kinetic analysis, since it analyses the kinetics of the whole of oxidative phosphorylation divided into a small number of modules (in this case three: substrate oxidation, proton leak and the phosphorylating system), connected by their common substrate or product, mitochondrial membrane potential, $\Delta\psi$. Thus the analysis picks up any change in oxidative phosphorylation that is functionally important, and is unresponsive to any changes that have no functional consequences. Comparison of the kinetic responses of each of the three modules to $\Delta\psi$ obtained using mitochondria isolated from different cybrids reveals any effects of mitochondrial haplogroup on the kinetics of oxidative phosphorylation. The coupling efficiency of oxidative phosphorylation (the percentage of respiration rate at a given $\Delta\psi$ that is used for ATP synthesis) can then be calculated from the kinetic curves. Note that any slip reactions will appear as proton leak in this analysis [23].

To validate this experimental system, we examined two models: a known chemical inhibitor and a known pathogenic mutation. We first measured the kinetics of oxidative phosphorylation in the absence and presence of a low concentration of the complex III inhibitor, myxothiazol. Fig. 1A shows that only the kinetics of substrate oxidation were affected, as predicted, and that a change in kinetics giving rate changes of 10% was readily observed. In the second model system, we measured the kinetics of oxidative phosphorylation in mitochondria isolated from cybrids harbouring a pathogenic mutation at np 3243 in the tRNA-leu(UUR) gene (*MTTL1*) [17, 18]. This mutation decreases expression of all mitochondrially-encoded proteins, so is expected to decrease the activity of all the respiratory chain complexes (except complex II, whose subunits are all nuclear-encoded) and the ATP synthase. Fig. 1B shows that our assay system reported the expected difference in the kinetics of substrate oxidation. The kinetics of the phosphorylation system were not significantly different, despite the anticipated decrease in ATP synthase activity, presumably because of low control of the ATP synthase over the phosphorylation rate. As expected, the kinetics of proton leak were not changed. Thus our test system was able to identify small changes, less than 10%, in the kinetics of different modules of oxidative phosphorylation if they were present.

Modular kinetic analysis of oxidative phosphorylation – test of the hypothesis.

The electrochemical proton gradient across the mitochondrial inner membrane, formed as electrons are passed down the respiratory chain, is the primary energy source for cellular ATP synthesis. However, not all of the energy available in the redox reactions is

coupled to ATP synthesis. Some is consumed by proton slip reactions (fewer protons are pumped out of the matrix during electron flow) or proton leak reactions (protons are pumped normally but leak back through proton conductance pathways that bypass the ATP synthase). As a result of these slip and leak reactions, heat is generated instead of ATP. These non-productive proton slip and leak pathways are physiologically important. They account for a surprisingly high proportion of cellular and organismal respiration rate, estimated to be up to 20% [24, 25], so differences in their capacities could plausibly play an adaptive role that could be selected during ancient human migrations.

To test the hypothesis, we measured the kinetics and coupling efficiency of oxidative phosphorylation using isolated mitochondria from cybrid cells harbouring Arctic or tropical haplogroups. Fig. 2A shows the averaged kinetics of the three modules of oxidative phosphorylation using succinate grouped by Arctic (closed symbols) or tropical (open symbols) mtDNA haplogroup (the results for the six individual cybrid lines are shown in Fig. 3).

There was a small difference, not statistically significant, between the Arctic and tropical haplogroups in the rate of the substrate oxidation module at any given $\Delta\psi$ (Fig. 2A, circles). This small difference was caused entirely by one of the three tropical haplogroup cybrids, L3, which had a lower maximum respiration rate than the other tropical haplogroup lines (Fig. 3A). We also measured the kinetics of substrate oxidation using α -ketoglutarate + malate as substrate instead of succinate to check for any effects of differences in the mitochondrial complex I genes. Once again, there was a small, but not significant, difference in the kinetics of substrate oxidation between the Arctic and tropical haplogroups (Fig. 2C) that was caused by lower rates only in the L3 cybrids (Fig. 3D). As there was otherwise no difference between Arctic and tropical haplogroups in the kinetics of substrate oxidation with either substrate, we assume this was an individual difference in the L3 line possibly caused by somatic mutations in mtDNA. We conclude that there is no general difference between Arctic and tropical mtDNA haplogroups in the overall kinetics of substrate oxidation.

Contrary to the hypothesis, the rate of the proton leak of mitochondria with the Arctic haplogroups was, if anything, slightly lower on average than that of the tropical ones at any value of $\Delta\psi$ (Fig. 2A, squares) (differences not statistically significant). For the individual haplogroups, the rate for tropical haplogroup L3 was relatively high and the rates for Arctic haplogroups C and D were relatively low (Fig. 3B), although any differences were small. This shows that mtDNA variations between the Arctic and tropical haplogroups had at most a

modest effect on mitochondrial proton leak, but in the wrong direction to support the hypothesis.

Fig. 2B shows that on average, there was no difference in the rate of the phosphorylating system at any given value of $\Delta\psi$ between the Arctic and tropical haplogroups. The kinetic curves of the six individual haplogroups almost overlapped (Fig. 3C). We conclude that there is no difference between Arctic and tropical mtDNA haplogroups in the overall kinetics of the phosphorylating system.

From the kinetic curves shown in Figs 2 and 3, we can analyse the coupling efficiency of oxidative phosphorylation. The classic measure is the respiratory control ratio: respiration at maximal rate of ATP synthesis (state 3) divided by respiration with no ATP synthesis (state 4) (Fig. 4A). We also independently measured the respiratory control ratio using α -ketoglutarate + malate as substrates (Fig. 4D and 4E). Fig. 4A and 4E show that there were small differences in average respiratory control ratio; mitochondria from the Arctic haplogroups were slightly better coupled than those from the tropical haplogroups. However, these differences were not statistically significant, and they were in the wrong direction to support the hypothesis even if haplogroup L3, which has lower respiration rate, was omitted (data not shown).

A more precise measure of coupling efficiency is given by the percentage of respiration that is coupled to ATP synthesis, calculated as shown in Fig. 2A at any chosen membrane potential. If this percentage is lower, then a greater proportion of the redox energy is diverted to heat production. Fig. 4B shows that at the value of $\Delta\psi$ achieved in state 3 (maximal rate of ATP synthesis), mitochondria from the Arctic haplogroups were slightly better coupled than those from the tropical haplogroups. Again, this difference was not statistically significant, and it was in the wrong direction to support the hypothesis. Mitochondria in cells normally operate at less than maximal rates. Fig. 4C examines the coupling efficiency across the whole range of rates of ATP synthesis and values of $\Delta\psi$ from state 3 to state 4 (where ATP synthesis is minimal). As expected [26], coupling efficiency decreased as mitochondria moved from state 3 towards state 4. Once again, there was a small difference, but not statistically significant, between the Arctic and tropical haplogroups, and again it was in the wrong direction to support the hypothesis.

Catalase activity

Differences in respiration rates of mouse cybrid cells harbouring different mtDNA sequences can be masked by compensatory changes in mitochondrial density due to a retrograde response that correlates with three-fold changes in catalase activity [27]. However, such compensatory changes would not mask changes in the modular kinetics of oxidative phosphorylation in mitochondria isolated from cybrids, and did not prevent the expected results in our pathogenic mutant model (Fig 1B). Nonetheless, to test whether catalase-associated compensation occurred in our human cybrids, we measured catalase activity to see if depended on mitochondrial haplogroup. Catalase specific activity did not differ between arctic (15.4 ± 1.0 units/mg protein, $n = 9$) and tropical (14.6 ± 1.1 units/mg protein, $n = 9$) cybrid cells ($p = 0.6$), suggesting that this retrograde response was not a factor in our experiments.

Discussion

We conclude that the hypothesis that mitochondrial haplogroups that lower the coupling efficiency of oxidative phosphorylation were positively selected during radiations of modern humans [12, 13] is not supported by our experimental test using modular kinetic analysis of cybrid mitochondria. Recent mitochondrial genetic studies have also cast doubt on the hypothesis [28, 29].

The strengths of our approach to test this hypothesis are (i) for the first time, we used a direct empirical measurement, rather than inference from genetic and physiological traits, (ii) we separated the effects of mitochondrial haplogroup from any confounding effects of nuclear DNA background and (iii) we used a simple in vitro system with the powerful modular kinetic approach where the relevant variables could be tightly controlled and manipulated. However, our approach also has some weaknesses. (i) A small effect of mitochondrial haplogroups operating over many generations might still be too small to be detected by biochemical experiments. We estimate that our experiments had the statistical power to identify a 7.3% difference in coupling efficiency between the two haplogroups. (ii) We measured the coupling efficiency of oxidative phosphorylation using isolated mitochondria in simple defined media, but effects of mitochondrial haplogroups might only emerge at cellular or higher levels (for example, they might involve mitochondrial-cytoplasmic feedback loops or retrograde signalling to nuclear genes, or might be tissue-

specific). (iii) Interaction between nuclear and mitochondrial genomes may affect oxidative phosphorylation [30], so the effect of mitochondrial haplogroup might depend on nuclear background. A549, the parental line of the A549.B2 ρ^0 cells used here, has mtDNA of European haplogroup H, which is proposed to be relatively cold-adapted [12, 13]. Perhaps mtDNA of tropical haplogroups does not complement A549 nuclear DNA optimally and causes slightly lower coupling efficiency, masking evolutionarily relevant efficiency changes in human populations.

The novel method introduced in this study, modular kinetic analysis using cybrids, has provided a powerful empirical test of a specific hypothesis of altered mitochondrial coupling efficiency in human populations carrying different mitochondrial haplogroups. However, it is also a more general method that can be used to investigate the bioenergetic consequences of other differences in mitochondrial haplogroups and haplotypes in humans or other species. As shown in Fig. 1B, it is also a powerful method to investigate in a precise and controlled way the bioenergetic phenotypes of hereditary or sporadic pathological mitochondrial DNA mutations that are thought to cause many mitochondrial diseases.

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Figure legends

Fig. 1. Modular kinetic analysis – validation of the method.

Modular kinetic analysis of the kinetic responses of the three modules of oxidative phosphorylation (substrate oxidation, circles; proton leak, squares; $\Delta\psi$ -consumers (sum of phosphorylating system and proton leak), triangles) to membrane potential, $\Delta\psi$, in mitochondria isolated from cybrids, using 4 mM succinate as substrate. (A) Kinetic responses in mitochondria isolated from A, D, L1 and L2 cybrids in the absence (filled symbols) and presence of 25 nM of the complex III inhibitor, myxothiazol (open symbols). Data are means $\pm$ SEM ($n = 4$ independent mitochondrial preparations). The kinetics of substrate oxidation were significantly different at membrane potentials above 98 mV. (B) Kinetic responses in mitochondria isolated from cybrids harbouring a pathogenic mutation at np 3243 in the tRNA-leu(UUR) gene (*MTTL1*). Filled symbols, clone Z (high proportion of wild type); open symbols, clone X (high proportion of mutant). Data are means $\pm$ SEM ($n = 3$ independent mitochondrial preparations). The kinetics of substrate oxidation were significantly different at all membrane potentials.

Fig. 2. Modular kinetic analysis – averaged values.

Modular kinetic analysis of the kinetic responses of the three modules of oxidative phosphorylation to membrane potential, $\Delta\psi$, in mitochondria isolated from cybrids. Filled symbols, Arctic haplogroups (A, C and D); open symbols, tropical haplogroups (L1, L2 and L3). (A) Succinate as substrate. Comparison of the kinetic responses of substrate oxidation (circles; $\Delta\psi$ titrated with uncoupler, FCCP), proton leak (squares; $\Delta\psi$ titrated with inhibitor, malonate), and $\Delta\psi$ -consumers (sum of phosphorylating system and proton leak, triangles; $\Delta\psi$ titrated with malonate). Circled are state 3 (condition with maximal ATP synthesis) and state 4 (condition with no ATP synthesis). The coupling efficiency of oxidative phosphorylation (percentage of respiration used for ATP synthesis at a given $\Delta\psi$) was calculated from the kinetic curves: the inset bar graph shows the example of Arctic haplogroups, at the $\Delta\psi$ of state 3. (B) Kinetic response of the phosphorylating system to $\Delta\psi$, calculated from (A) by subtracting respiration driving proton leak from respiration driving the $\Delta\psi$ -consumers at each $\Delta\psi$. (C) Kinetic response of substrate oxidation using 3.2 mM α -ketoglutarate + 0.8 mM

malate as substrate, with rotenone omitted. Data are means $\pm$ SEM ($n = 9$ cell clones - three different clones for each of the three constituent haplogroups; 3 or 4 (**A**, **B** and **C**) or 1 (**D**) independent mitochondrial preparations per clone).

Fig. 3. Modular kinetic analysis – full dataset.

Modular kinetic analysis in mitochondria isolated from cybrids, using succinate as substrate, of (**A**) substrate oxidation, (**B**) proton leak and (**C**) the phosphorylating system. (**D**) Modular kinetic analysis of substrate oxidation using α -ketoglutarate + malate as substrate. Filled symbols, Arctic haplogroups (circles, haplogroup A; squares, haplogroup C; triangles, haplogroup D); open symbols, tropical haplogroups (circles, haplogroup L1; squares, haplogroup L2; triangles, haplogroup L3). Data are means $\pm$ SEM ($n = 3$ cell clones; 3 or 4 (**A**, **B** and **C**) or 1 (**D**) repeats using independently prepared mitochondria for each clone). The kinetic response of the phosphorylating system to $\Delta\psi$ was calculated by subtracting respiration driving proton leak from respiration driving the $\Delta\psi$ -consumers at each value of $\Delta\psi$.

Fig. 4. Coupling efficiency of oxidative phosphorylation in mitochondria isolated from cybrids.

Filled bars, Arctic haplogroups (A, C and D); open bars, tropical haplogroups (L1, L2 and L3). (**A**) Respiratory control ratio using succinate as substrate (State 3 respiration rate/State 4 respiration rate). (**B**) Coupling efficiency using succinate as substrate at the $\Delta\psi$ of state 3. (**C**) Coupling efficiency using succinate as substrate at selected $\Delta\psi$ values between state 3 and state 4. Values at 145 mV or less were significantly different from values at 155 mV. (**D**) Respiration rates with α -ketoglutarate + malate as substrate. (**E**) Respiratory control ratio using α -ketoglutarate + malate as substrate. Data are means $\pm$ SEM ($n = 9$ cell clones - three different clones for each of the three constituent haplogroups; 3 or 4 (**A**, **B** and **C**) or 1 (**D** and **E**) independent mitochondrial preparations per clone). Values in (**A**, **B** and **C**) were calculated from data in Fig. 2A.

Figure 1

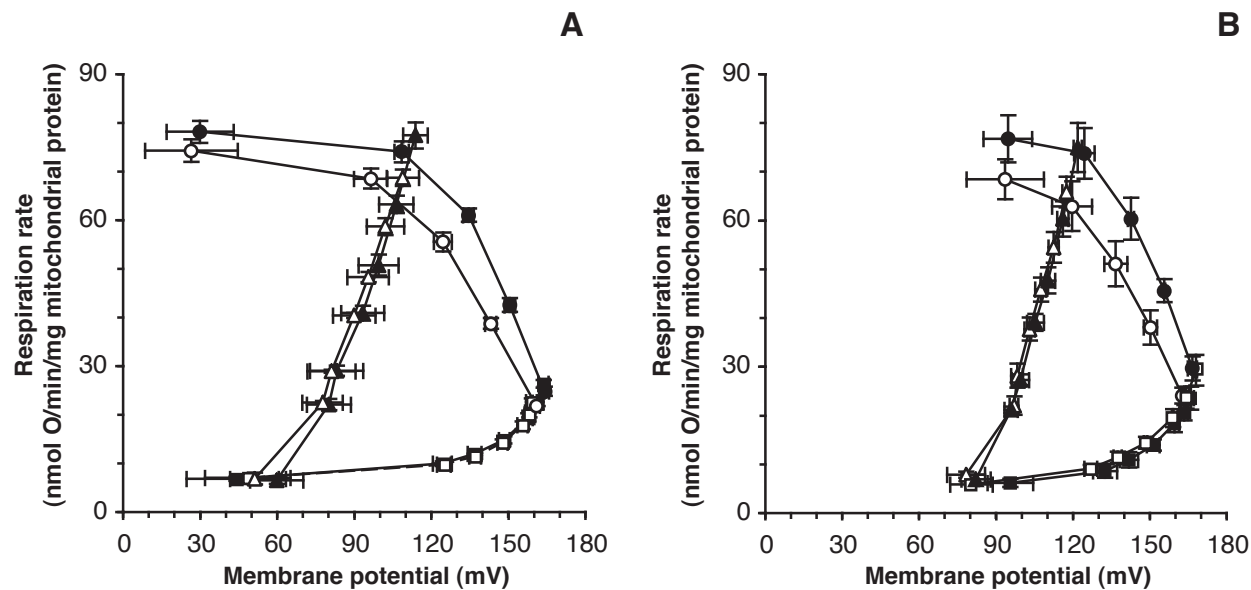


Figure 2

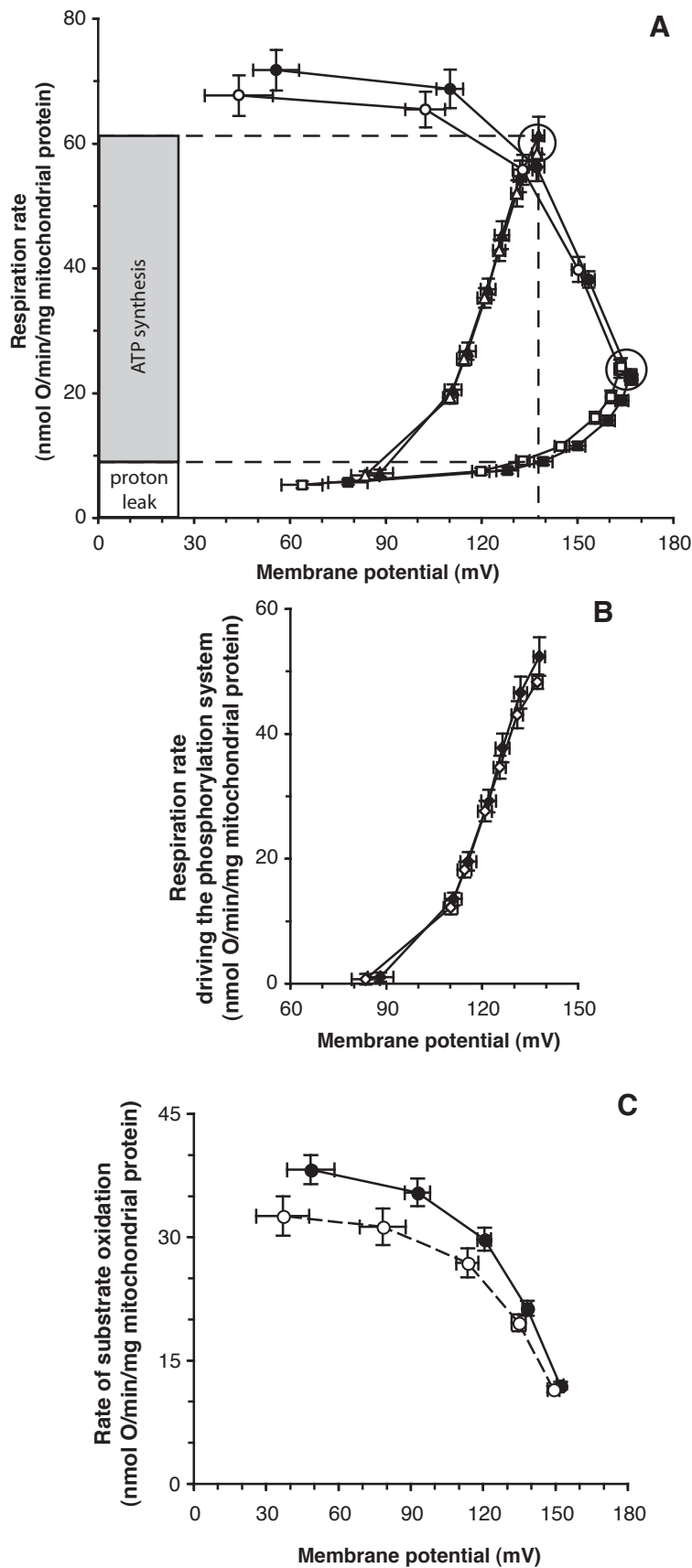


Figure 3

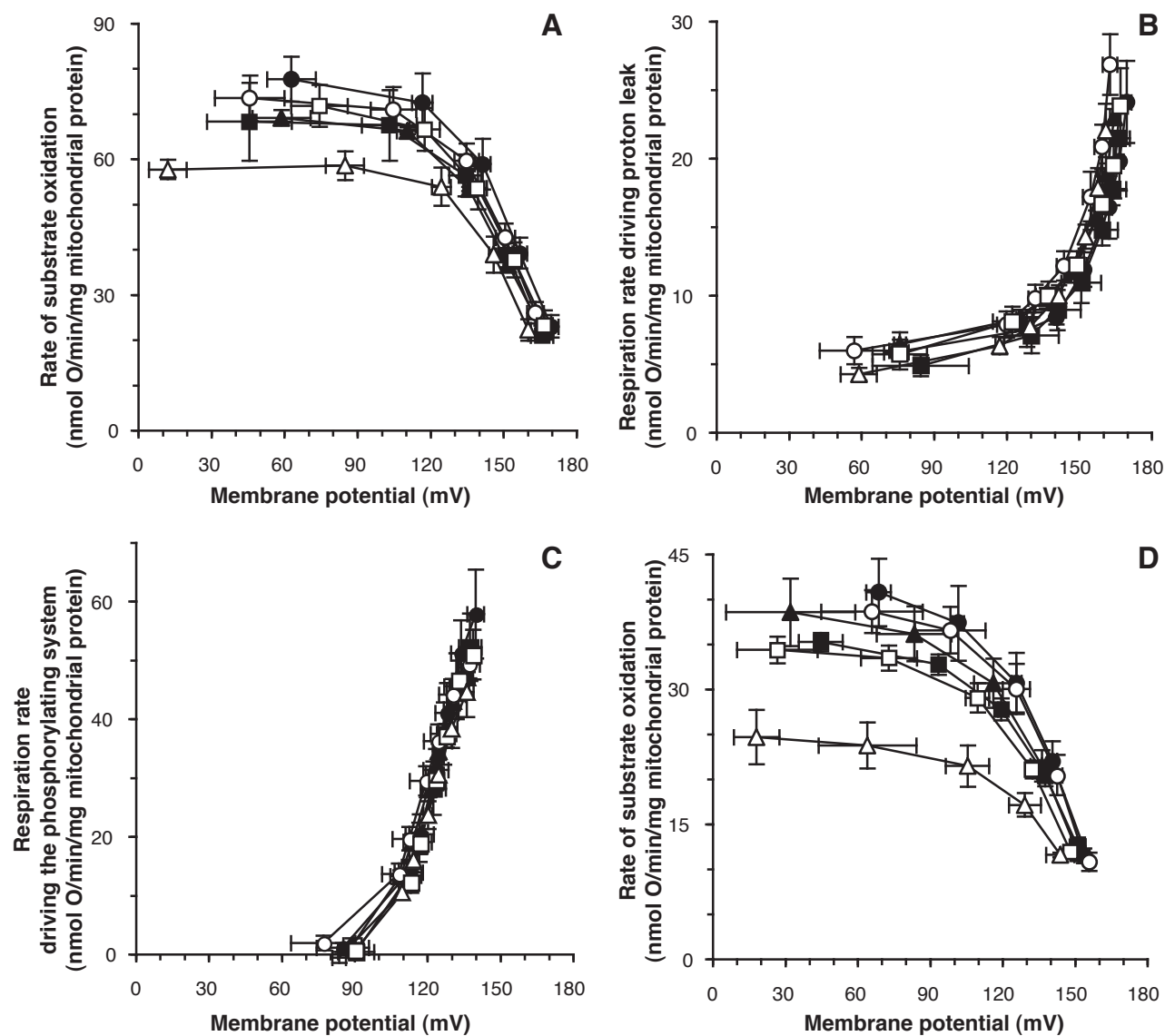


Figure 4

