Mitochondrial DNA polymorphisms specifically modify cerebral \(\beta\)-amyloid proteostasis

Katja Scheffler · Markus Krohn · Tina Dunkelmann · Jan Stenzel · Bruno Miroux · Saleh Ibrahim · Oliver von Bohlen und Halbach · Hans-Jochen Heinze · Lary C. Walker · Jörg A. Gsponer · Jens Pahnke

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Abstract Several lines of evidence link mutations and deletions in mitochondrial DNA (mtDNA) and its maternal inheritance to neurodegenerative diseases in the elderly. Age-related mutations of mtDNA modulate the tricarboxylic cycle enzyme activity, mitochondrial oxidative phosphorylation capacity and oxidative stress response. To investigate the functional relevance of specific mtDNA polymorphisms of inbred mouse strains in the proteostasis regulation of the brain, we established novel mitochondrial congenic mouse lines of Alzheimer’s disease (AD). We crossed females from inbred strains (FVB/N, AKR/J, NOD/LtJ) with C57BL/6 males for at least ten generations to gain specific mitochondrial conplastic strains with pure C57BL/6 nuclear backgrounds. We show that specific mtDNA polymorphisms originating from the inbred strains differentially influence mitochondrial energy metabolism, ATP production and ATP-driven microglial activity, resulting in alterations of cerebral \(\beta\)-amyloid (A\(\beta\)) accumulation. Our findings demonstrate that mtDNA-related increases in ATP levels and subsequently in microglial activity are directly linked to decreased A\(\beta\) accumulation in vivo, implicating reduced mitochondrial function in microglia as a causative factor in the development of age-related cerebral proteopathies such as AD.

Keywords Abeta · Mitochondrial congenic mice · Conplastic · Alzheimer’s disease · Amyloid-beta · Microglia · Mitochondria

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K. Scheffler · M. Krohn · T. Dunkelmann · J. Stenzel · H.-J. Heinze · J. Pahnke
Department of Neurology, Neurodegeneration Research Laboratory (NRL), Universities of Rostock, Rostock, Germany

M. Krohn · J. Pahnke
German Centre for Neurodegenerative Diseases (DZNE), Magdeburg, Germany

B. Miroux
Laboratoire de Biologie Physico-Chimique des Protéines Membranaires, UMR 7099, Institut de Biologie Physico-Chimique, Paris, France

S. Ibrahim
Department of Dermatology, University of Lübeck, Lübeck, Germany

O. von Bohlen und Halbach
Institute of Anatomy, University of Greifswald, Greifswald, Germany

H.-J. Heinze · J. Pahnke
Leibniz Institute for Neurobiology, Magdeburg, Germany

L. C. Walker
Yerkes National Primate Research Center and Department of Neurology, Emory University, Atlanta, GA, USA

J. A. Gsponer
Center for High-Throughput Biology, University of British Columbia, Vancouver, Canada

Present Address:
K. Scheffler
Department of Biochemistry, University of Oslo, Oslo, Norway

J. Pahnke
Department of Neurology, Neurodegeneration Research Laboratory (NRL), Universities of Magdeburg, Leipziger Str. 44, H15, 39120 Magdeburg, Germany
e-mail: jens.pahnke@gmail.com
URL: www.NRL.ovgu.de
Introduction

mtDNA mutations become increasingly common throughout the body with advancing age [20] and are thought to be associated with several age-related neurodegenerative disorders [35]. However, there is little compelling evidence linking these mutations to specific pathological changes [6, 22]. Because engineering and integrating specific mutated mtDNA in mice is technically challenging, clinically relevant mouse models with pathological mtDNA mutations are rare [36]. Therefore, naturally occurring mtDNA polymorphisms in common inbred strains hold considerable promise for analyzing the pathogenic effects of polymorphic mtDNA. mtDNA variants interfere with cognitive abilities [30], and differentially modulate mitochondrial oxidative phosphorylation (OXPHOS) and the generation of reactive oxygen species (ROS) [26]. The age-related decline of respiratory chain function in mtDNA-mutant mice was found to be more strongly affected by mtDNA point mutations than by deletions [11]. Numerous findings also implicate mitochondrial dysfunction and oxidative damage in the pathogenesis of neurodegenerative diseases, such as Alzheimer’s disease (AD) [24]. For instance, depleting cells of endogenous mtDNA in vitro and repopulating them with mitochondria from AD patients resulted in respiratory chain deficiency and β-amyloid (Aβ) accumulation [18, 33]. Sequencing of mtDNA from the blood or brain tissue of AD patients however yielded contradictory results [12, 34]. Hence, the hypothesized connection between specific mtDNA polymorphisms and defects in protein maintenance (protein homeostasis or proteostasis), specifically the cerebral accumulation of Aβ, remains uncertain [9, 10].

Materials and methods

Animal models and generation of congenic mouse lines

(1) APP-transgenic mice [29] (C57BL/6 gDNA and mtDNA background) were obtained from R. Radde and M. Jucker (University of Tübingen, Hertie Institute for Clinical Brain Research, Tübingen, Germany) and further maintained with C57BL/6 females for more than six generations (APP-B6). C57BL/6 female mice (B6) were purchased from Jackson Laboratory (Bar Harbor, ME, USA). (2) APP-FVB mice were generated by crossing APP-transgenic mice to FVB/N females for more than ten generations. FVB/N female mice were purchased from Jackson Laboratory. (3) Mouse inbred strains (FVB/N, AKR/J, NOD/LtJ) used to generate mtDNA conplastic strains were purchased from Jackson Laboratory. The mtDNA of each strain had been sequenced previously [38]. C57BL/6 (B6) was chosen as the gDNA and mtDNA reference strain because the APP-transgenic mouse model on the same background is well characterized [29, 31]. Based on the exclusive inheritance of maternal mtDNA [1], we mated female mice manifesting the desired mtDNA variants with male B6 mice for more than ten generations to eliminate the nuclear DNA of the inbred strains AKR/J, FVB/N, and NOD/LtJ, respectively. APP-B6 males (Sect. 1 above) were then crossed to mtDNA strains to produce APPxmtAKR, APPxmtFVB and APPxmtNOD mice, respectively. (4) Uncoupling protein 2-knockout mice (Ucp20/0, C57BL/6 gDNA and mtDNA background) were provided by Bruno Miroux. APP-B6 mice were crossed to Ucp20/0 mice to produce APP/+xUCP20/0 mice.

All mice were housed in a climate-controlled environment on a 12 h light/dark cycle with free access to rodent food (SNIFF, Germany) and water. All procedures were conducted in accordance with animal protocols approved by the University of Rostock and according to the state law of the government of Mecklenburg-Western Pomerania.

Tissue preparation

Mice were killed by cervical dislocation and transectually perfused with PBS. The brain was removed, and one hemisphere was fixed in buffered, 4 % paraformaldehyde (PFA) for paraffin-embedding and immunohistochemistry, while the other hemisphere was snap-frozen in liquid nitrogen and stored at −80 °C for biochemical analysis.

Enzyme-linked immunosorbent assay (ELISA)

ELISA kits (TK42HS—high sensitivity from The Genetics Company, Schlieren, Switzerland) were used for the quantification of Aβ42 in whole brain hemispheres from which the cerebellum and brain stem had been removed at the level of the midbrain. Hemispheres were homogenized using PreCellys24 (12 s, 6,500 rpm). After addition of carbonate buffer (pH 8.0), homogenates were mixed using PreCellys (5 s, 5,000 rpm) and centrifuged for 90 min (4 °C) at 24,000g to separate insoluble from soluble Aβ species. The resulting supernatant (buffer-soluble fraction) was mixed with 8 M guanidine hydrochloride at a ratio of 1:1.6. To extract aggregated Aβ species, the pellet was dissolved in 8 volumes of 5 M guanidine hydrochloride, shaken at room temperature for 3 h and centrifuged at 24,000g for 20 min at 4 °C. The resulting supernatant represents the guanidine-soluble fraction. Protein contents of all samples were measured in triplicate using a Nanodrop1000 (Thermo Fisher Scientific, USA). ELISAs were performed according to the manufacturer’s instructions using appropriate dilutions.
TCA and OXPHOS activity measurements in isolated mitochondria

Mitochondria from freshly prepared brain hemispheres were isolated as described elsewhere [38]. The mitochondria were adjusted to the same protein concentration as determined by BCA assay (Pierce, a division of Thermo Fisher Scientific, USA). The different enzymatic activities were assessed using the PARADIGM spectrophotometer (Beckman Coulter, USA). Pyruvate DH activity was measured using a PDH enzyme activity microplate assay kit (MitoScience, USA). Malate DH activity was measured by monitoring the absorbance at 340 nm in a reaction mixture containing 50 mM potassium phosphate buffer (pH 7.4), 0.4 mM NADH, 2 mM oxaloacetate and 10 μM rotenone. Complex I, III and ATPase activities were measured as described elsewhere [2, 38]. The ATPase activity represents complex V activity. Complex II and IV activities were measured using the respective enzyme activity microplate assay kits (MitoScience, USA).

Assessment of ATP and H2O2 levels

ATP and H2O2 levels were measured in freshly prepared total brain homogenates. Assays were performed with either a luminescence (Infinite200Pro, Tecan, Switzerland) or spectrophotometric plate reader (PARADIGM, Beckman Coulter, USA). Tissue ATP levels were determined by a luciferin/luciferase ATP Bioluminescent Assay kit (Sigma-Aldrich, Germany). The Amplex Red hydrogen peroxide assay kit (Invitrogen, USA) was used to detect H2O2 release. Sample preparation and assay procedures were performed according to the manufacturers’ protocols. Due to the high activities of ATPases in fresh samples, tissue was immediately snap frozen in liquid nitrogen. All steps for sample preparation were carried out at 4 °C in order to avoid the recovery of ATPase activity and subsequently degradation of ATP in the tissue.

Catalase and β-secretase activity measurements

Enzyme activities were measured in total brain homogenates. The different enzymatic activities were assessed using the PARADIGM spectrophotometer (Beckman Coulter, USA). Catalase activity was measured by using an OxiSelect catalase activity assay kit (Cell Biolabs, USA). The activity of β-secretase was determined with a β-secretase activity assay kit (BioVision, USA). Sample preparation and assay procedures were performed according to the manufacturers’ protocols.

Western blots

For Western blotting, tissue homogenates were prepared as described by Lesné et al. [23]. Total protein concentrations of the extracts were determined using a BCA assay (Pierce, division of Thermo Fisher Scientific, USA). After electrophoresis of 10 μg total protein per lane, proteins were blotted onto PVDF membranes. After blocking in 5 % dry milk in TBST buffer (50 mM Tris pH 7.4, 150 mM NaCl, 0.1 % Tween20) for 1 h at room temperature, blots were probed for ADAM10 (1:1,000, Calbiochem, USA), BACE1 (1:1,000, Abcam, UK), IDE (1:500, Abcam, UK), ApoE (1:2,000, Abcam, UK), ABCA1 (1:1,000, Novus Biologicals, USA), total OXPHOS antibody cocktail (1:1,000, MitoSciences, USA) or β-actin (1:40,000, Sigma-Aldrich, Germany) overnight at 4 °C. As detection antibodies, anti-mouse-HRP and anti-rabbit-HRP, respectively, were used. The Amersham ECL Plus Detection kit (GE Healthcare, UK) and a Roper CoolSnap HQ2 Camera (Roper, Germany) were used for visualization.

Immunohistochemistry

Brain hemispheres were post-fixed for at least 24 h in buffered, 4 % paraformaldehyde. Paraffin-embedded, 4-μm thick coronal sections were stained using a BondMax™ (Leica Microsystems GmbH, Germany) automated immunostaining system. Sections were pretreated with 98 % formic acid for 5 min and immunostained for Aβ using the anti-human Aβ clone 6F3D (1:200, 30 min, DAKO, Germany) and the Bond™ Polymer Refine Detection kit (Leica Microsystems GmbH, Germany). For double-stained slides, microglia were subsequently immunostained on the same sections using anti-Iba1 (1:1,000, 30 min, Wako GmbH, Germany) and the Bond™ Polymer AP-Red Detection kit (Leica Microsystems GmbH, Germany) [31]. Whole tissue sections were fully digitized at a resolution of 230 nm using Mirax Desk/Midi slide scanners (Zeiss, Germany) and then analyzed semi-automatically using the AxioVision software package (Zeiss Microimaging GmbH, Germany) as previously described in [31].

Primary microglia preparation and Aβ-phagocytosis assays

Highly purified primary microglial cultures were prepared as described by Horvath et al. [15] with slight modifications. In brief, cerebral cortices of 2- to 3-day-old mice were minced and digested with 1 x Trypsin/EDTA for 15 min at 37 °C. Pelleted tissue was triturated several times on ice in Dulbecco’s modified Eagle’s medium supplemented with 10 % FBS, 1 % GlutaMax, 1 % penicillin/streptomycin and 2,000 units of DNase. Supernatant containing glial cells was collected, triturated again until...
the tissue clumps disappeared and finally centrifuged at 340g for 15 min. The cell pellet was re-suspended in media and plated at 2 × 10^6 cells per 75 cm² flask. Cultures were maintained at 37 °C and 5 % CO₂. Media were changed every 3–4 days. After 20 days in culture, the flask was shaken at 250 rpm for 45 min, and the primary microglia-enriched supernatant was removed and centrifuged at 340g for 15 min. To assess the phagocytosis of Aβ, the cell pellet was resuspended in media and plated at 30,000 primary microglia per well onto a 96-well plate. Peptide phagocytosis was quantified by measuring the uptake of FITC-labelled Aβ42 (rPeptide, USA). Aβ aggregates (500 nM) were incubated with the microglia for 6 h. Then the medium was removed and the remaining cells were washed twice with PBS and trypsinized. The cells were centrifuged at 340g for 15 min and resuspended in PBS. Intracellular fluorescence (excitation 485 nm and emission 535 nm) was measured with the PARADIGM spectrophotometer (Beckman Coulter, USA).

Immunofluorescence

Primary microglia were seeded at 200,000 cells per 6-well plate onto poly-d-lysine-coated cover slips. After 24 h incubation with 500 nM FITC-labeled Aβ42, cells were fixed with 2 % paraformaldehyde for 5 min and then ice-cold methanol for an additional 5 min. Primary microglia were immunolabelled using an antibody against Iba1 (1:1,000, Wako GmbH, Germany) for 1 h. Detection was performed with goat anti-rabbit Cy3 (1:1,000, Dianova, Germany). Cells were counterstained and mounted using VECTASHIELD with DAPI (Biozol, Germany). Visualization was achieved with a Zeiss Laser Scanning Microscope (LSM700, Zeiss, Germany).

Results

Inbred AD mouse strains show distinct differences in Aβ load

Support for a link between DNA polymorphisms and altered protein homeostasis comes from the initial evaluation of the influence of the C57BL/6J (B6) and FVB/N murine strain backgrounds on cerebral Aβ accumulation in APP-transgenic mice. B6 and FVB/N mice differ substantially in nuclear and mitochondrial DNA. ELISA quantifications revealed significantly lower Aβ levels in APP-FVB mice compared to APP-B6 mice at 200 days of age (Fig. 1). This indicates a nuclear and/or mitochondrial DNA variations may cause these differences while expressing the identical APP transgenes.

Conplastic mice show distinct biochemical differences

To determine whether the observed differences in Aβ levels are indeed caused by mtDNA differences between B6 and FVB/N, we established new mtDNA congenic AD mouse lines by using common inbred strains (FVB/N, AKR/J, NOD/LtJ) with naturally occurring mtDNA polymorphisms leading to coding and non-coding nucleotide exchanges (Table 1). As mitochondria are inherited maternally, we backcrossed female mice of these strains for at least ten generations with B6 males to gain solely mitochondrial differences on an otherwise pure B6 background (B6xmtAKR/J, B6xmtFVB/N, and B6xmtNOD/LtJ). The offsprings after the tenth generation were regarded as congeneric/conplastic strains that carried the nuclear genome of the recipient strain and the mitochondrial genome from the donor strains. The resulting congeneric mtDNA mice were then crossed to APP-B6 males to evaluate intracellular Aβ proteostasis in the context of variant mtDNA (conplastic AD mice: APP-B6xmtFVB/N [mtFVB], APP-B6xmtAKR/J [mtAKR], APP-B6xmtNOD/LtJ [mtNOD]).

Sequencing the mtDNA of these common inbred strains revealed four genetic variants (Table 1): (1) a poly A insertion in the D-loop of mtRNAArg in all three mtDNA congenic mouse strains previously described to alter not only hearing ability in mice, but also to differentially affect ROS production in transmitochondrial cybrids [17, 26]; (2) a coding substitution in the ATP synthase subunit 8 (Atp8) gene in the mtFVB DNA, altering mitochondrial function [38]; (3) a coding substitution in the cytochrome c oxidase subunit 3 (Cox3) gene in mtNOD mice, leading to decreased complex IV activity (Supplementary Fig. 2); (4) a non-
coding substitution in the NADH dehydrogenase subunit 3 (Nd3) gene in all three mtDNA congeneric mouse strains. The relevance of non-coding single nucleotide exchanges in the NADH dehydrogenase gene is as yet unknown; however, transcriptional regulation of NADH dehydrogenase subunits has been discussed in the context of proteostasis defects [5].

To evaluate the impact that the mtDNA variants have on cellular metabolism, we first analyzed the enzymatic activities of the components of the respiratory chain complex, the tricarboxylic acid cycle (TCA) and the resulting cellular ATP levels in the brain. In all mtDNA mouse strains, we found significant differences in TCA enzymes and OXPHOS complex activities (Supplementary Figs. 1, 2) that ultimately resulted in significantly increased ATP levels in mtNOD mice (2) that ultimately resulted in significantly increased ATP levels in mtNOD mice (+75 %), and non-significantly in mtAKR mice (+35 %) and in mtFVB mice (+52 %, Fig. 3a). Because ROS production differs in relation to murine mitochondrial DNA haplotypes [26], we then measured hydrogen peroxide (H2O2) levels and catalase activity in the conplastic AD mice. H2O2 levels were reduced only in the mtFVB strain (−48 %), whereas catalase activity was unchanged in all mtDNA strains and B6 controls (Supplementary Fig. 3). Overall these results demonstrate that single nucleotide differences in the mtDNA can cause substantial metabolic changes.

Conplastic mice exhibit differences in Aβ load

To reveal which impact these metabolic changes have on cerebral proteostasis, we quantified Aβ in the conplastic AD strains. Immunohistochemical analysis showed that Aβ-immunoreactive senile plaques were reduced both in size and number in all mtDNA conplastic strains compared to the APP-B6 controls (Fig. 2a–g). Quantification of cortical plaques in progressively older age groups of mice showed that mtNOD mice had a lower areal density of plaques evident as early as at 75 days of age, whereas plaque load in the other groups (mtAKR, mtFVB) was similar to controls until 175 days of age. At 200 days of age, plaque load was significantly lower in all mtDNA-variant mice (Fig. 2h). These findings were confirmed by ELISA measurements in which Aβ levels were significantly lower in the mtDNA variant mice (−59 % mtAKR, −65 % mtFVB, −82 % mtNOD) (Fig. 2i). Again, lower Aβ levels were apparent in the mtNOD mice as early as 75 days of age (Fig. 2i and Supplementary Fig. 4). To exclude that the differences in Aβ deposition are due to alterations in APP processing pathways, we measured the levels of several proteins/proteases involved in the production, degradation or trafficking of Aβ. The expression of α-secretase (ADAM10), β-secretase (BACE1), ABCA1, ApoE or IDE did not differ as a function of mitochondrial status. In addition, β-secretase activity did not differ significantly among the mouse strains as determined at 200 days of age (Supplementary Fig. 5). These results reveal that other mechanisms are likely involved in the changed Aβ proteostasis.

Microglia function is regulated by conplastic mitochondria

Activated microglial cells rely strongly on mitochondrial function; they are a prominent feature of Aβ plaques and are known to be involved in the phagocytosis of Aβ [14, 28]. Thus, we were interested whether the mtDNA polymorphisms result in variations of the cerebral proteostasis due to changes in microglial activity. We found severe microgliosis in mtDNA-variant mice at 200 days of age, most prominently in the mtNOD mice (Fig. 3c–f). All conplastic strains exhibited significantly greater numbers of Aβ plaques that were heavily covered (>50 %) by microglia at day 100 (+128 % mtAKR, +153 % mtFVB and +186 % mtNOD), whereas only mtNOD mice maintained a significantly elevated microglial response at 200 days of age (+93 %, the endpoint of our analysis) (Fig. 3b). To assess the potential association between microglial mtDNA polymorphisms and the number and size of Aβ plaques, we studied the phagocytosis of Aβ by microglia in vitro. Immunofluorescence analysis of Iba1-positive microglia isolated from mtDNA mice and B6 controls incubated with fluorescein-coupled Aβ42 revealed an intravesicular localization of Aβ aggregates (Fig. 3g). Quantification of Aβ phagocytosis in mtNOD mice, which show the lowest

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**Table 1 mtDNA sequence variation among the four inbred mouse strains**

<table>
<thead>
<tr>
<th>Position (bp)</th>
<th>C57BL/6J</th>
<th>AKR/J</th>
<th>FVB/N</th>
<th>NOD/LtJ</th>
<th>Gene/RNA</th>
<th>Amino acid substitution</th>
</tr>
</thead>
<tbody>
<tr>
<td>7778</td>
<td>G</td>
<td>G</td>
<td>T</td>
<td>G</td>
<td>mt-Atp8</td>
<td>Asp → Tyr</td>
</tr>
<tr>
<td>9348</td>
<td>R</td>
<td>G</td>
<td>G</td>
<td>A</td>
<td>mt-Cox3</td>
<td>Val → Ile</td>
</tr>
<tr>
<td>9461</td>
<td>T</td>
<td>C</td>
<td>C</td>
<td>C</td>
<td>mt-Nd3</td>
<td>None</td>
</tr>
<tr>
<td>9828</td>
<td>A</td>
<td>AA</td>
<td>AA</td>
<td>AAA</td>
<td>mtRNA17s</td>
<td></td>
</tr>
</tbody>
</table>

C57BL/6J was set as the reference strain. Shown are nucleotide positions of mtDNA-encoded genes at which differences occur (indicated in bold letters).
cerebral Aβ load, demonstrated significantly higher uptake levels in microglia compared to B6 controls (Fig. 3h). These findings suggest that ATP-triggered microglial activity plays an important role in Aβ homeostasis.

High ATP levels reduce Aβ load

The most striking effects in Aβ reduction are found in mtNOD mice compared to mtAKR, mtFVB and B6 control...
mice. Interestingly, these mice also exhibit the highest ATP levels. It is known that release of ATP triggers microglial response, and this might therefore explain amplified microglial activity toward aggregated Aβ [7, 8]. To support the ATP dependency of this phenomenon, we additionally established an mtDNA-independent AD model lacking the nuclear genome-encoded ATP-bypass enzyme uncoupling protein 2 (APP × UCP2). Higher levels of ATP have...
been observed in various cells types of UCP2-deficient mice [19, 39]. As expected, the APP-B6 × UCP20/0 mice also exhibit much higher ATP levels. Most importantly, these mice have significantly lower Aβ levels in the brain (Fig. 4), which suggests an inverse correlation of increased ATP and reduced Aβ levels.

**Discussion**

In this study, we analyzed conplastic mouse models with identical nuclear but divergent mitochondrial genomes, and thereby provide the first in vivo evidence that specific mtDNA variants in common inbred mouse strains can substantially influence the cerebral proteostasis network, indicating a potential role of mtDNA mutations in the ontogeny of neurodegenerative diseases. In particular, we found distinct biochemical alterations resulting in higher ATP levels, lower plaque number and size, lower Aβ levels and a pronounced microglial response to Aβ. Energetic differences influence the phagocytic activity of microglia toward Aβ, which is necessary for plaque removal; hence, the microglial activation state may be an important regulator of the progression of cerebral Aβ amyloidosis. In line with this, ablation of the majority of parenchymal microglia in APP-transgenic mice on a B6 genomic background that miss the energetic phagocytic microglial activation [13]. Senescence leads to the accumulation of random mitochondrial changes in cells and tissues and to an age-related switch in microglial function from phagocytic to more cytotoxic in vivo [16, 31]. Increased heteroplasmy of random mitochondrial polymorphisms accumulating with age could engender gross functional and regulational deficiencies in specific mitochondrial pathways, thus contributing to functional changes of ATP-consuming mechanisms, e.g., Aβ export from the brain [3, 21, 25], fostering sporadic forms of AD. Consequently, no specific, inherited mtDNA variations have yet been discovered as linked to AD, although several studies have sought to ascertain the presence of mtDNA polymorphisms and maternal inheritance in AD patients [4, 5, 27, 34, 37]. The FVB/N and AKR/J strains have a double and the NOD/LtJ a treble A-repeat. Mutations in mtRNAArg result in an altered conformation and OXPHOS deficiencies in patients [32]. Moreover, alterations of regulatory mechanisms, such as a reduction in the amount of transcripts of NADH dehydrogenase subunits, have been proposed as potential risk factors for AD [5].

Together, ATP level alterations due to functional variations in OXPHOS and/or TCA cycle variations that are caused by accumulating mtDNA mutations lead to beneficial or detrimental effects for the brain’s proteostasis...
networks and may thus play an important role in the development or even prevention of age-related neurodegenerative diseases.

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Conflict of interest The authors declare that they have no competing financial interests.

References


Electronic Supplementary Information

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Supplementary Information – Table of content

Supplementary Figures .........................................................pages 2-6
Supplementary Figure 1. Activities of TCA enzymes are influenced by mtDNA variations.

Shown are the relative enzymatic activities of key TCA enzymes (pyruvate dehydrogenase [PDH] and malate dehydrogenase [Malate DH]) of either the nontransgenic (nTg) background strains (left side) or the APP-expressing mouse models (right side) at the age of 200 days. The nontransgenic C57BL/6J strain served as reference and was set at 100%. Asterisks indicate significant differences relative to either the C57BL/6J or the APP-B6 control strain.

Data are presented as means ± SEM (n=3-6 per group), *p<0.05.
Supplementary Figure 2. Activities of mitochondrial electron transport chain (mtETC) enzymes are influenced by mtDNA variations.

Shown are the relative enzymatic activities of mtETC enzymes of either the nontransgenic (nTg) background strains (left side) or the APP-expressing mouse models (right side) at the age of 200 days. The nontransgenic C57BL/6J strain served as reference and was set at 100%. Asterisks indicate significant differences relative to either the C57BL/6J or the APP-B6 control strain. (Inset) Immunoblot showing unaltered expression levels of mETC complexes in APP-expressing mitochondrial inbred strains.

Data are presented as means ± SEM (n=4-6 per group), *p<0.05.
Supplementary Figure 3. ROS and defence activity in the mitochondrial congeneric mouse strains.

Relative H$_2$O$_2$ content is only significantly reduced in APP-B6xmt$^FVB$ mice at 200 days of age, but catalase activity is unaltered.

Data are presented as means ± SEM (n=4 per group), *p<0.05.
Supplementary Figure 4. Age-dependent measurements of Aβ42 levels in the buffer-soluble Aβ fraction in mtDNA-variant mouse strains.

The APP-B6xmt<sup>NOD</sup> mice show an early and persistent deficit in buffer-soluble Aβ42 levels relative to the APP-B6 control mice, whereas (with the exception of the APP-B6xmt<sup>AKR</sup> group at 150 days) similar amounts of Aβ42 were found in the APP-B6xmt<sup>AKR</sup> and the APP-B6xmt<sup>FVB</sup> mice compared to the APP-B6 controls.

Data are presented as means ± SEM (n = 6-10 per group), *p<0.05.
Supplementary Figure 5. Unaltered APP processing and degradation pathways in mtDNA mouse strains.

(a) Immunoblot analyses of proteins involved in the production, trafficking, or degradation of APP in 200-day old congenic mouse strains. β-actin serves as a loading control. Representative results of three independent experiments are shown. (b) Measurement of β-secretase activity in 200-day old APP-expressing mtDNA mouse strains reveals no differences.

Data are presented as means ± SEM (n=6 per group), *p<0.05.