A Network Approach to Mapping Mouse Brain-wide Mitochondrial Respiratory Chain Capacity in Relation to Behavior

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Abstract

The brain and behavior are under energetic constraints, limited by mitochondrial energy production capacity. However, the mitochondria-behavior relationship has not been systematically studied on a brain-wide scale. Here we examine the association between multiple features of mitochondrial respiratory chain capacity and stress-related behaviors in mice with diverse behavioral phenotypes. Miniaturized assays of mitochondrial respiratory chain enzyme activities and mitochondrial DNA (mtDNA) content were deployed on 571 samples across 17 brain regions, defining specific patterns of mito-behavior associations that vary across brain regions and behaviors. Furthermore, multi-slice network analysis applied to our brain-wide mitochondrial dataset identified three large-scale networks of brain regions with shared mitochondrial signatures. A major network composed of cortico-striatal regions exhibited the strongest mitochondria-behavior correlations, suggesting that this mito-based network is functionally significant. Mito-based networks also overlap with regional gene expression and structural connectivity, thereby providing convergent multimodal evidence of mitochondrial respiratory chain functional organization anchored in gene, brain and behavior.

*Keywords:* mitochondria, behavior, stress, connectivity, network analysis, mouse


**Introduction**

The shaping of behaviors by life experiences is driven by energetically demanding circuitry across the brain\(^1\). The brain’s enormous energetic demand is mainly met by ATP produced through oxidative phosphorylation (OxPhos), subserved by the combined activities of respiratory chain (RC) enzymes within mitochondria\(^2\). As a result, mitochondria influence multiple aspects of brain development and function ranging from dendritic and axonal branching\(^3,4\), remodeling of gene expression\(^5\), regulation of neurotransmitter release and excitability in mature synapses\(^6-8\), neurogenesis\(^9\), and inflammation\(^10\). Thus, the mounting molecular and functional evidence that the brain is under energetic constraints\(^11,12\) suggests that if we want to understand the basis of brain function and behavior, we must understand key aspects of brain mitochondrial biology.

Mitochondria are small, dynamic, multifunctional organelles with their own genome\(^13\), but not all mitochondria are created equal. Mitochondria serving different cellular demands (i.e., in different cell types) have different relative molecular compositions, morphologies, and functional phenotypes\(^14-19\). Therefore, developing a comprehensive understanding of the association between mitochondrial biology and animal behavior calls for assessments of multiple functional and molecular mitochondrial features, across multiple brain regions. By analyzing mitochondrial features across multiple brain regions simultaneously, we can also potentially uncover unknown brain “mitochondrial circuits”. This idea aligns with the evolving understanding of large-scale brain circuitry and metabolism\(^11,20\), and of the network distribution of neural activity achieved using brain-wide, high spatial resolution methods (e.g., MRI-based functional and structural connectivity maps)\(^21,22\). Although a similar degree of resolution for mitochondrial phenotyping is not feasible with current technologies, miniaturization of biochemical and molecular mitochondrial assays open new possibilities to systematically map mitochondria-to-behavior associations across multiple cortical and sub-cortical brain regions in the same animal.
Both mitochondrial biology and animal behaviors measured on standardized tests exhibit *naturally occurring* and *acquired* (i.e., experience-dependent) variation, providing an opportunity to map their associations. For example, behaviorally, exposure to social stress such as chronic social defeat predictably increases potential threat vigilance and social avoidance\(^{23}\), but animals differ in their vulnerability/resilience to these stress-induced behavior changes\(^{24,25}\). Experimental challenges aimed at modeling neuroendocrine disturbances resulting from chronic stress, such as chronic exposure to corticosterone, also induce avoidance behaviors associated with recalibrations in specific brain circuitry\(^{26}\), gene expression\(^{27,28}\), and anatomical plasticity (i.e., atrophy) in stress-sensitive brain regions like the hippocampus\(^{29}\). In relation to mitochondria, a separate body of literature similarly documents naturally-occurring mitochondrial variation\(^{30-32}\), as well as stress-induced functional mitochondrial recalibrations that occur within days to weeks of stress exposure\(^{33-35}\) (meta-analysis in \(^{36}\)), potentially linking mitochondrial biology to behavior. More direct causal experiments show that mitochondrial RC enzyme activities directly influence the brain and specific behavioral domains including working memory\(^{37,38}\), social dominance\(^{31,39,40}\) and anxiety-related behavior\(^4\). Targeted mitochondrial defects even cause mood disorder-like phenotypes in animals\(^{41}\), positioning mitochondria as upstream modulators of brain function and behavior. Moreover, mitochondrial RC defects are likely implicated in the etiology of psychiatric, neurological and degenerative disorders in humans\(^{42}\), and *in vivo* brain metabolic imaging studies show that energy metabolism in specific brain areas (e.g., nucleus accumbens, NAc) predict cognitive performance and anxiety\(^{43-45}\), making these biological questions also potentially relevant to human mental health.

Although the importance of mitochondria for brain structure and function is unequivocal, we lack an understanding of potential differences in mitochondria across different brain regions. This calls for more systematic mapping of region-specific brain mitochondrial biology in relation to behavior. To address the hypothesis that mitochondrial phenotypes in specific brain regions
are associated with behaviors, we have further miniaturized existing biochemical and molecular assays of mitochondrial OxPhos enzyme activities for sub-milligram tissue samples and deployed them across 17 cortical and sub-cortical brain regions in mice with a wide range of behavioral phenotypes, quantified through four behavioral tests. Using network-based connectivity analysis, we also sought to explore the distribution of mitochondrial phenotypes across brain regions, finding evidence that mouse brain mitochondria specialize as distinct functional networks linked to behavior. Further, we use gene co-expression and structural/anatomical connectivity data, which anchor the newly observed mitochondrial circuits into other modalities, and provide a foundation for future mechanistic studies to elucidate the specific energetic contributions to behavioral variation.

Results

Miniaturization of mitochondrial assays for sub-milligram resolution

Although enzymatic activity assays directly reflect the capacity of RC complexes and therefore energy production capacity, previously available assays suffer from low throughput that preclude a multi-region, brain-wide analysis in dozens of animals. Building from our efforts to miniaturize and scale the throughput of mitochondrial RC activity assays in immune cells\textsuperscript{30,32}, here we miniaturized and optimized spectrophotometric assays in 96-well plate format for brain tissue, validated against the standard cuvette-based reaction (Extended Data Fig. 1, and Methods). Using this optimized platform, we can quantify the enzymatic activities of RC complex I (CI, NADH-ubiquinone oxidoreductase), complex II (CII, succinate-ubiquinone oxidoreductase), complex IV (CIV, cytochrome c oxidase), and citrate synthase (CS, a Krebs cycle enzyme and marker of mitochondrial content) in two 1mm-diameter, 200\textmu m deep tissue punches, establishing in our hands the lowest detection limit for mouse brain tissue. This represents <1mg of tissue (estimated 0.33mg), over an order of magnitude more sensitive than currently available methods.
In this pipeline, the same biological sample used for enzymatic activities is also used to quantify mtDNA abundance, both by i) the classical metric relative to the nuclear genome – mtDNA/nDNA ratio, termed mtDNA copy number (mtDNAcn)\textsuperscript{46}, and ii) per tissue volume (mtDNA copies per μm\textsuperscript{3} or per mg), termed mtDNA density. qPCR data of nuclear genome abundance across the brain, independent of the mtDNA, showed that cellular density varies by up to ~8.5-fold between brain regions (highest: cerebellum, lowest: visual cortex). Unlike in peripheral tissues, this remarkably large variation in neuronal and glial cell somata density significantly skews mtDNAcn estimates. Extended Data Fig. 2 shows the inter-correlations of mtDNAcn and mtDNA density in relation to enzyme activities. Compared to mtDNAcn, which is confounded by the presence or absence of somata/nuclear genome, mtDNA density was more consistently associated with RC enzymatic mitochondrial phenotypes across all brain regions, and therefore likely represents a more generalizable estimate of mitochondrial genome abundance across mouse brain regions.

We subsequently combined these five primary mitochondrial measures into a mitochondrial health index (MHI) by dividing RC activities (CI+CI\texttext{I+}CV) by mitochondrial content (CS+mtDNA density), thus creating an index of energy production capacity on a per-mitochondrion basis\textsuperscript{30} (see Methods for details). Although each of the six resulting features are partially correlated with other features, the proportion of shared variance between individual features across brain regions is 31-64%, indicating that they each contribute some non-redundant information about mitochondrial phenotypes, which can be deployed at scale.

Our study design first aimed to profile animal-to-animal differences in mitochondrial phenotypes across a broad set of brain regions known to be associated with anxiety-related behavior, social behaviors, cognition, or mitochondrial disorders, and to relate these measures to each animal’s behavioral phenotypes. In total, we enzymatically and molecularly phenotyped 571 samples covering 17 cortical, sub-cortical and brainstem brain regions isolated by bilateral
punches at defined stereotaxic coordinates (Table 1, Extended Data Fig. 3). To eventually compare the specificity of our findings related to brain and behavior, 5 peripheral tissues were also collected and analyzed from each animal, which we expected to show somewhat related but more modest associations with behavioral outcomes.

Protein levels and enzymatic activity

We initially explored if RC protein abundance is a viable surrogate for mitochondrial RC activity\textsuperscript{31}, which could theoretically allow high spatial resolution imaging of the entire brain. We focused on the cerebellum due to its well-defined cellular composition and laminar organization, where the Purkinje cell layer is flanked by molecular and granular layers\textsuperscript{14} (Extended Data Fig. 4a). Compared to protein abundance, enzymatic activity ultimately determines mitochondrial RC function and energy production capacity and consequently should be regarded as the most representative measure of mitochondrial phenotypes. In consecutive cerebellar slices, we compared RC complex II enzymatic activity measured spectrophotometrically, to the protein abundance of a complex II subunit, SDHA (succinate dehydrogenase, subunit A), for which a validated high-affinity antibody allows its quantification by microscopy (Extended Data Fig. 4b-c). Across the three cerebellar layers, enzyme activity did not correlate with protein abundance assessed by immunohistochemistry and densitometry (proportion of shared variance, $r^2=0.02$-0.07), indicating that protein abundance and enzymatic activity are not equivalent (Extended Data Fig. 4d). The reasons for this finding could include the action of post-translational modifications, stoichiometry of the four SDH subunits, or the biochemical context that drive biochemical activity independent of protein content (e.g.,\textsuperscript{47}). Therefore, we focus all downstream analyses on direct measures of mitochondrial RC enzymatic activity and mtDNA density.

Diversity of mitochondrial RC activities between animals

We next examined mitochondria-behavior associations in a cohort of mice exhibiting naturally-occurring behavioral and mitochondrial variation. Our goal was to identify robust and
generalizable associations between mitochondrial phenotypes and behavior, rather than potential correlations that would exist among only subgroups of animals (naïve or stressed). Therefore, to further extend the spectrum of mito-behavior variation using well-characterized rodent stress models, subgroups of the cohort were either chronically administered corticosterone (CORT)\textsuperscript{48} for three months, or exposed to 10 days of chronic social defeat stress (CSDS)\textsuperscript{49} (see Extended Data Fig. 3a). To create additional diversity among groups, half of the CSDS mice were allowed to recover for two months, as some stress-induced behavioral and mitochondrial changes may change again when the stressor is removed. Based on previous work\textsuperscript{24,50}, we also expected naturally-occurring behavioral and brain molecular phenotype differences between animals that are resilient or susceptible to CSDS (based on the social interaction test; see Methods)\textsuperscript{51}. The naturally-occurring variation in mitochondria and behavior plus the effects of various exposures provides a strong test of robustness for our hypothesis, which was that across a diverse population of mice, mitochondrial phenotypes in specific brain regions are consistently associated with behaviors.

Across our cohort of inbred male mice with a range of exposures, a wide spectrum of mitochondrial phenotypes was observed. The average variation for all measures (4 individual RC enzymatic activities and mtDNA density) across all animals and all 17 brain regions was a C.V. of 36\% (coefficient of variation = standard deviation / mean). For any given brain region, the absolute variation in mitochondrial phenotypes between mice reached up to 2.9-fold between the animal with the lowest and the animal with the highest activities. This means that for a given brain region, there are large mouse-to-mouse differences in mitochondrial content and RC activities, even among inbred naïve mice not exposed to stressors (Extended Data Fig. 5). Peripheral tissues showed an average animal-to-animal C.V. of 25\%, about a third less variation than for brain regions, indicating that the brain may exhibit particularly large inter-individual differences.
To verify that the interventions used to enhance naturally-occurring variation in behavior and mitochondria were effective, we first compared mitochondrial features in mice exposed to CORT and CSDS relative to naïve mice. The effects of CORT and CSDS on mitochondrial phenotypes were quantified as standardized effect sizes (Hedges g) and shown in Figure 1a-c (detailed in Extended Data Fig. 6). Both interventions altered mitochondrial RC complexes, CS enzymatic activities, mtDNA density, and MHI in a region-specific manner. Although not statistically significant, CORT-treated mice tended to have higher mitochondrial activities than non-stressed animals in ~60% of brain regions, whereas CSDS animals trended towards lower activities in ~82% of brain regions, which was statistically significant for CI, CIV, and MHI measures (Figure 1d), suggesting opposing effects of these two different stress models on brain mitochondria. This could be due to either the nature of the stressor, or their durations. The amygdala (Amyg) showed the greatest CORT-induced increase in CII enzymatic activity (+49%, p<0.05, unadjusted p value), whereas the periaqueductal gray (PAG) showed the largest CSDS-induced decrease in CI activity (-42%, p<0.05, unadjusted).

To determine if brain regions were co-regulated in their stress-induced mitochondrial recalibrations, we employed a topological data analysis (TDA)-based Mapper approach. Mapper is a variant of nonlinear dimensionality reduction methods (manifold learning) that produces a graph or network embedding of the high-dimensional data (a.k.a. shape graph) while recovering projection loss using an additional partial clustering step. When applied to the six mitochondrial features (i.e., four enzyme activities, mtDNA density, and the MHI) separately for CORT and CSDS (both measured as delta of stress vs the naïve group average), the shape graphs revealed differences in regional mitochondrial recalibrations across the two groups (Figure 1e-f). This was quantified using a graph-theoretical measure of participation coefficient (PC), where higher values of PC indicate uniformly distributed connectivity across regions (integrated network) and lower values indicate a segregated connectivity pattern. CORT-
induced mitochondrial recalibrations were relatively more region-specific or segregated, whereas CSDS caused a more coherent or integrated mitochondrial response across all brain regions (Figure 1g).

Although we were not powered to make statistical comparisons between recovery-susceptible and -resilient mice based on the social interaction test for CSDS mice, we did observe relatively large differences in effect sizes (g > 0.8) in mitochondrial recalibrations between the groups that could be explored in the future work (Extended Data Fig. 7). Additionally, compared to non-recovered CSDS mice, animals who were allowed to recover from CSDS for 57 exhibited mitochondrial phenotype changes only marginally different from control mice, and in a limited number of brain regions (Extended Data Fig. 8). This result suggests, along with the main group differences, that the brain mitochondrial recalibrations are dynamic over time scales ranging from days to weeks.

Overall, these univariate and TDA-based results established the existence of naturally-occurring and acquired variation in brain mitochondrial phenotypes between animals, providing a strong basis to test the existence of conserved associations with behaviors.

Diversity of anxiety and depressive like behaviors

Similar to the spectrum of mitochondrial phenotypes across mice, as expected from previous work, animals also naturally exhibited large variation in their behavioral phenotypes. Behavioral tests included potential threat (“anxiety”) monitored by the open field test (OFT) and elevated plus maze (EPM), hyponeophagia monitored by novelty suppressed feeding test (NSF), and approach-avoidance conflict using the social interaction test (SI). Because specific behavioral tests are generally administered in conjunction with specific interventions (CORT, CSDS), some animals were only tested on some and not all behaviors. Both CORT and CSDS produced the expected elevation in anxiety-related behavior compared to naïve mice (Extended Data Fig. 9). Results from the behavioral tests either correlated moderately with each other
(EPM and OFT, r=0.60), were not correlated (OFT and NSF, r=-0.02), or were negatively correlated (OFT and SI, r=-0.46). This indicated that each test captures different aspects of behavior, thereby providing a basis to examine how different aspects of behavior might relate to region-specific brain mitochondrial phenotypes.

**Brain MHI correlates with specific behaviors**

We next evaluated the extent to which mitochondrial phenotypes in different brain regions was associated with behavior across all animals (Figure 2a). Behavioral scores were transformed so that higher scores on each test indicate higher avoidance/anxiety-like behaviors (as in 24) (See Methods, Extended Data Fig. 9). Behavioral scores were then correlated with the 6 measures of mitochondrial phenotypes, where higher values indicate higher mitochondrial content or RC functioning. A frequency distribution of the effect sizes (Spearman r) for all mitochondrial-behavior pairs revealed a significant non-zero correlation between MHI in the 17 brain regions and behavior on OFT (p<0.01), EPM (p<0.0001), and SI (p<0.0001), but not for NSF (Figure 2b, Gaussian frequency distributions for the other mitochondrial features are shown in Extended Data Fig. 10). Time to feed on the NSF was capped at 600 seconds, so the correlations are less precise than for other tests.

To better understand which brain regions were driving the direction and magnitude in the distributions, we then examined the patterns of correlations for all 17 brain regions, for all 6 mitochondrial features, across the 4 behavioral tests (Figure 2c). In the majority of brain regions, higher mitochondrial metrics were correlated with higher avoidance scores based on OFT (average r=0.12) and EPM (average r=34), although these overall correlations did not reach statistical significance. The strongest correlations were of 0.51 for CII in the primary motor cortex (M1) and OFT (p=0.025, linear regression, unadjusted), and 0.92 for MHI in the nucleus accumbens (NAc) and EPM (p=0.0005, unadjusted). Previous research in rodents showed that mitochondrial RC function in brain regions such as the NAc is linked to complex behaviors such
as social dominance and anxiety\textsuperscript{4,31}. For NSF there was more heterogeneity between brain regions, with a similar number of positive and negative correlations. Finally, for SI, we observed the opposite relationship; animals with higher mitochondrial content and RC activities generally displayed greater sociability (lower social avoidance) (average \( r=-0.21 \), strongest for MHI in the substantia nigra (SN) and SI, \( r=-0.78, p=0.0035, \) unadjusted). Interestingly, of the six mitochondrial features, the strongest mito-behavior correlations (for 3 out of 4 behaviors) were for MHI, which may reflect the superiority of MHI as an integrative measure of mitochondrial energy production capacity over individual enzymatic and molecular features, as previously observed\textsuperscript{30,55,56}.

As expected, the average correlation between RC activities and behaviors was significantly more consistent for brain mitochondria than for mitochondria in peripheral tissues (Figure 2d). For example, whereas mitochondrial phenotypes in several brain regions correlate with anxiety-related behavior on the EPM, mitochondrial measures in the muscles, heart, liver, or adrenal glands of the same animals on average, did not correlate with behavior. This finding aligns with recent work in mice showing that mitochondrial phenotypes exhibit strong segregation between different cell types and tissues\textsuperscript{57}, and has two implications. First, it reinforces the specificity of these mito-behavior findings for the brain. Second, it implies that mitochondria across the brain and other tissues within an individual mouse are not equivalent, and likely differentially regulated\textsuperscript{57}. This naturally raised the question whether specific brain regions within an animal could also exhibit independently regulated mitochondrial properties, and whether brain regions could be functionally organized into separate networks based on their mitochondrial properties.

\textit{Mitochondrial phenotype-based organization of the brain}

To address this question, we first asked whether mitochondrial features in each brain region/tissues were statistically independent or correlated with other tissues. Using the same
rationale that underlie functional connectivity analysis in fMRI data, we first generated a correlation-based similarity matrix of all mitochondrial measures across brain regions and non-brain tissues (Figure 3a). Within the 17 brain regions, mitochondrial features were generally positively correlated (average $r=0.22$, $p<10^{-100}$, two-sample t-test), with a few exceptions. Thus, within the brain, higher mitochondrial activities in one region generally also implies higher activities in other regions ($p<10^{-82}$, one-sample t-test).

Consistent with the co-regulation of RC enzymes within the mitochondrion, a modular structure was also apparent, indicating that mitochondrial features within each region were more similar than with other regions ($p<0.0001$, permutation test). Figure 3b shows the average correlations of each mitochondrial feature with other features among all brain regions – RC enzymes were most strongly co-regulated (average $rs=0.7-0.8$), followed by mtDNA density ($r=0.6$), and MHI ($r=0.5$), again suggesting that MHI captures different information than individual measures. To compute how similar each region is to other regions, we computed nodal degree (i.e., average inter-regional correlation, similar to the concept of global connectivity) for all regions. Nodal degree was highest in the cerebellum (average $r=0.34$) and lowest in the brainstem vestibular nucleus ($r=0.05$) (Figure 3c), suggesting that the region-to-region similarity in mitochondrial phenotypes is not randomly distributed across the mouse brain.

As suggested above from the divergence between brain regions and peripheral tissues in the mito-behavior correlation patterns, the mitochondrial features in peripheral tissues were not correlated with brain mitochondria (average $r=0.02$) or with other peripheral tissues mitochondria (average $r=-0.03$) (Figure 3d). The lack of association between brain and non-brain tissues of the same animals suggest that MHI (as well as content and RC activities, data not shown) is not a ubiquitous animal-level feature, but rather relatively independently defined in the brain, and further specified among each brain region.
Given the overall positive connectivity across the brain and recent evidence of circuit-level metabolic coupling across large neural networks in the Drosophila brain\textsuperscript{12}, we examined whether certain regions exhibited particularly strong mitochondrial feature co-regulation. Functional brain networks in the living brain are defined by synchronous activity patterns (e.g., default-mode network, fronto-parietal network). Similarly, we reasoned that connectivity patterns of mitochondrial features (based on single measurements) could reflect networks of brain regions with similar bioenergetic properties. To examine this hypothesis with a framework agnostic to anatomical categorization and inclusive of all mitochondrial features, we performed multi-slice community detection analysis\textsuperscript{61}, with mitochondrial features represented in six separate layers (Figure 3e). Categorical multi-slice community detection allow to detect cohesive groups of brain regions, or communities, that: i) are more similar to each other than they are to the rest of the regions, and ii) have cohesion converging across the six layers of mitochondrial features. Representing all 17 brain regions, and across 6 mitochondrial features, we found 3 separate communities, or networks: 1) \textit{Cortico-striatal network}: CPu, visual, motor, mOFC, mPFC and NAc; 2) \textit{Salience/Spatial navigation network}: cerebellum, vestibular nuclei, VTA, thalamus, hippocampus (CA3), and dentate gyrus (dorsal and ventral); and 3) \textit{Threat response network}: amygdala, hypothalamus, substantia nigra, and periaqueductal grey (Figure 3f).

To examine whether this functional organization of the brain revealed using mitochondrial phenotypes was also evident cross-modally, we used two Allen Brain Atlas datasets of brain-wide gene co-expression\textsuperscript{62}, and EYFP-labeled axonal projections that define the structural connectome\textsuperscript{63}, developed in the same mouse strain. Specifically, we examined whether gene expression correlations and structural connectivity within brain regions that are functionally grouped together as a network, based on mitochondrial features, is higher than expected by chance. We used two independent graph theoretical metrics: strength fraction
(S.F.)∞4 and quality of modularity (Q_mod)∞5, to examine whether similar communities exist in gene and structural connectivity data as in our mitochondrial phenotype data. Using permutation testing, we randomly shuffled community structure of brain regions 10,000 times to determine whether the community structure derived from mitochondrial features is also evident in gene co-expression and structural connectivity data. Both strength fraction and quality modularity statistics indicated that mitochondria-derived networks also have higher similarity in gene co-expression (S.F. p=0.020; Q_mod p=0.008) and structural connectome data (S.F. p=0.029; Q_mod p=0.015) than expected by chance (Extended Data Fig. 11). Hence, this provides convergent multimodal evidence of mitochondrial phenotypic organization overlapping with gene expression and structural connectivity.

**Network-level mitochondria-behavior correlation**

Finally, to examine the potential significance and added value of this effective brain-wide mitochondrial connectivity in relation to animal behavior, we used the mitochondria-derived network results to partition the brain into three networks, and then analyzed the ability of each network to linearly predict behaviors. The mitochondrial-behavior correlations varied both in strength and direction between the three networks, and the mitochondrial phenotypes among the three networks had largely divergent associations with behaviors. The average mitochondrial activity in the cortico-striatal network consistently showed the strongest average correlations with behaviors measured in the OFT, EPM, and SI tests (r=0.20, 0.54, -0.41, respectively). For the EPM, mitochondrial features in the cortico-stratal network (1) accounted for up to 33% of the animal-to-animal variance in behavior (EPM). In contrast, the salience/spatial navigation network (2) exhibited the weakest average correlations for the same three behaviors (r=-0.03, 0.18, -0.14), accounting for <5% of behavioral variance (Figure 3g). Together, these results suggest the presence, specificity, and functional significance of brain-wide mitochondrial networks, embedded within existing neural circuitry in the mouse brain.
Discussion

Using a high-throughput approach to functionally phenotype hundreds of brain samples from a heterogenous mouse cohort, we have defined brain-wide associations between mitochondrial phenotypes and behaviors. Combined with previous findings\textsuperscript{31,35}, the diverging mito-behavior associations between brain regions, and between brain and non-brain tissues, brought to light the possibility that different brain regions might exhibit different mitochondrial phenotypes. In particular, the network characteristics of mitochondrial phenotypes across the brain provided evidence for the modular distribution of mitochondria across cortical and subcortical regions, as well as their relevance to animal behaviors. Based on these data, we conclude that mouse brain mitochondria may exist as behaviorally-relevant networks overlapping with, but distinct from, other modalities including gene expression and structural connectivity.

Utilizing CORT and CSDS interventions to induce behavioral variation in our animal cohort also allowed us to compare the effects of these two interventions on mitochondrial phenotypes. Some brain regions were found to respond, in some cases, in opposite directions, particularly after exposure to CORT. In contrast to CORT, the recalibrations of brain mitochondria to CSDS was more uniform, with the majority of brain regions exhibiting a coordinated reduction in most mitochondrial features. This difference in mitochondrial recalibrations between both stress models may be driven by several factors. This includes the stressor duration, although the temporal dynamics over which stress-induced mitochondrial recalibrations take place remain poorly defined and will require further focused attention. Differences in the effects of CORT vs CSDS on mitochondria also could be related to their neuroendocrine underpinnings (single hormone for CORT vs multiple physiological neuroendocrine recalibrations for CSDS), regional differences in glucocorticoid and
mineralocorticoid receptor density, or other factors generally relevant to interpreting chronic stress rodent models.

One valuable aspect of our study is that it highlights the specificity of earlier findings indicating a connection between mitochondrial RC function in the NAc and anxiety behaviors\(^4,31\). In our study, the strongest correlation between EPM-based anxiety-like behavior and MHI was in the NAc, confirming the strong association between NAc mitochondrial energy production capacity and anxious behavior. We also extend this finding to show that mitochondrial phenotypes are linked to behaviors across not only isolated regions, but likely distributed among brain networks. Similar to the conceptual shift from regional and cellular perspectives towards distributed brain networks, circuits, and neuronal ensembles\(^66,67\), our findings therefore advance the notion that mitochondria may modulate brain function and behavior through distributed mito-networks. This notion is consistent with recent evidence of metabolic coupling with distributed brain-wide patterns of neural activity linked to behavior in Drosophila\(^12\).

While we cannot directly explain why distinct mitochondrial phenotypes appear to exist across brain regions, this may be driven by three main factors. First, mitochondria could respond to differences in neuronal circuit functioning, in agreement with observed coupling of neuronal and metabolic activities\(^12\), such that the cellular infrastructure of neural circuits that fire together not only wire together, but also generate similar mitochondrial phenotypes (i.e., mitotypes). A second possibility is that brain regions which are regularly co-activated, i.e., within the same functional networks, harbor similar levels of receptors for neuroendocrine factors (stress and sex steroids) known to influence mitochondrial biogenesis and/or functional specialization\(^68,69\). A third possibility would involve differences in metabolites and substrates that regulate mitochondrial RC activities, arising similarly among co-activated brain regions. These and other potential factors underlying the modularity of mitochondria within the mouse brain remain to be investigated.
From a mitochondrial biology perspective, our data highlights a potentially important distinction between mitochondrial content (abundance of mitochondria) and RC activity normalized to mitochondrial content, such as the MHI (respiratory chain activity per mitochondrion). Frozen tissue measurements naturally reflect the maximal functional capacity of the mitochondria RC, rather than their actual in vivo rates that are driven by neural activity and metabolic demands. Interestingly, the behavioral correlations with mitochondrial content features (mtDNA density and CS activity) for social avoidance behavior were similar to one another, and differed somewhat from RC enzyme activities. Experimentally, the specificity of these findings contrasting content from RC enzyme activities highlights the value of parallel assessments of multiple mitochondrial features reflecting unique mitochondrial phenotypes (i.e., mitochondria are not all created equal). In neuroimaging terms, the composite MHI can be understood as the mitochondrial analog to fMRI-based multi-variate pattern analysis (MVPA)\textsuperscript{70,71}, where multiple features (mitochondrial enzymes for MHI, voxels for MVPA) are combined to create a more stable and statistically accurate metric of the desired outcome (mitochondrial health for MHI, or brain activation for MVPA). Moreover, mtDNAcn has previously been assessed across multiple brain regions\textsuperscript{72}, but our results go beyond these observations in showing that quantifying mtDNA content on a per-cell basis (mtDNA:nDNA ratio) is heavily skewed by cellularity variations across brain regions and not directly related to RC energy production capacity. As such, the mtDNA:nDNA ratio (mtDNAcn) is driven by how many cell bodies are present in the tissue, and correlates poorly with either mitochondrial content or RC activity. Therefore, our data reinforce the notion that mtDNAcn on its own is not a valid measure of mitochondrial phenotypes\textsuperscript{73,74}. In the mouse brain, our findings suggest that mtDNA density per unit of volume (rather than per cell) is a more biologically meaningful mitochondrial feature when comparing brain regions that differ in cellularity.
By examining mitochondrial features across the mouse brain, we discovered a moderate level of global functional connectivity across mitochondria in most brain regions, but not among peripheral tissues (Figure 3). By functional connectivity, we do not imply that mitochondria are directly connected to each other in the same way that neurons project and chemically (de)activate each other; but rather that they share functional properties. If mitochondrial phenotypes are directly determined by genetic and physiological factors, the logical expectation is that all mitochondria within different organs of the same organism should exhibit a high degree of coherence (i.e., correlated with each other). In other words, the animal with the highest mitochondrial content or RC activities in the one brain region should also be the animal with the highest activities in other brain regions and tissues. Our results strongly disprove this point. Similarly, previous work on multiple human tissues showed that mtDNAcn was not significantly correlated between organs, including across three brain regions\textsuperscript{75}. Here we extend these data to 17 brain regions and 5 non-brain tissues, demonstrating that identified correlations are relatively modest (the strongest is $r=0.31$ for the cerebellum, representing less than 10% of shared variance). Notably, while on average we observed some degree of mitochondrial phenotype coregulation across the brain, some brain regions exhibited no consistent correlation with other regions. The most parsimonious explanation for this result is that individual animals differentially recruit different circuitry, which secondarily shape their mitochondria and drive animal-specific patterns of regional mitochondrial variation. Considering theories that stipulate that the brain strives for maximal energetic efficiency\textsuperscript{1} and that there may be limits to cellular energy conversion rates\textsuperscript{76}, we also cannot exclude the possibility that a limited quantity of resources (i.e., mitochondrial content or energy conversion) is available within the brain or the whole organism, which are distributed unequally in an activity-dependent manner among different organ systems, functional networks, and individual brain regions. Thus, we speculate that different animals may achieve an optimal balance of systemic and neural functions through
specific combinations of mitochondrial activity in different brain regions, a possibility that remains to be tested.

In trying to better understand why certain regions displayed stronger brain-wide connectivity than others, we identified mitochondria-based communities, or networks of brain regions. Interestingly, these mitochondria-derived networks share general anatomical features with established large-scale networks. For example, the identified cortico-striatal network 1 includes the CPu, NAc, mOFC, mPFC, and motor and visual cortex, which are implicated in decision making and executing actions. Network 2 (Cereb, VN, VTA, Thal, CA3, DGv and DGd) is the most heterogenous but comprised of regions that are known to be connected and involved in salience and spatial navigation. Lastly, network 3 includes limbic and limbic-associated regions (Amyg, Hypoth, PAG and SN) involved in threat responses. In comparing these mitochondria-derived networks to gene expression and structural connectome data we found that the communities significantly overlap across modalities. However, they are not identical, suggesting that mitochondrial properties are not the simple product of either gene expression nor structural connectivity. Finally, each network’s integrated mitochondrial phenotype exhibited different associations with behavioral responses. Unsurprisingly, the cortico-striatal network 1 explained the greatest proportion of variance in behavior, accounting for up to one third of the variance in anxiety-related behavior among animals. This quantitative observation increases the likelihood that the identified large-scale mitochondrial networks are functionally relevant. This network perspective may provide a basis to further delineate and develop accurate models brain-wide mitochondria organization and specialization.

Until now, the methods used to reveal neurobiological and metabolic networks in mammals have typically been through indirect functional and structural connectivity analysis. Here we have developed a scalable approach to examine mitochondrial phenotypes across a large number of brain regions in mice with a range of behavioral phenotypes. We showed that
mitochondrial phenotype connectivity was non-random, and linked with gene co-expression and structural connectivity, thereby providing converging evidence of mitochondria-based networks across modalities. This study synergizes with recent work providing the technical and empirical foundation to bring mitochondrial biology into brain-wide, network-based models of neural systems in mammals. Applied to mitochondria, network-based analytics should contribute to develop more accurate maps, and eventually an understanding of what drives mitochondrial and metabolic properties across interacting neural circuits. Developing a spatially-resolved understanding of brain mitochondrial biology will help to resolve the energetic contraints on brain function and behavior.

**Limitations.** Notable limitations of this study include the lack of cell type specificity. Neurons operate in a metabolic partnership with astrocytes and glial cells, and different cell types exhibit different molecular mitochondrial phenotypes (e.g.,) that cannot be disentangled in tissue homogenates. While neither enzymatic or functional mitochondrial profiling at the single brain cell is currently technically feasible, it remains possible that mitochondrial phenotypes between brain regions are influenced by differences in cell type proportions. Moreover, our molecular and biochemical mitochondrial phenotypes do not reflect other factors that can influence the efficiency or activity of the mitochondrial RC or OxPhos system in vivo, such as variations in cofactor abundance and RC structural assembly (e.g., supercomplexes). Because functional assays require harvesting brain tissue, it also was not feasible to ascertain within a given animal how stable (trait) or dynamic (state) the brain biochemical mitochondrial phenotypes are. If mitochondrial phenotypes were more dynamic than expected, our estimated proportion of behavioral variance attributable to mitochondrial biology could be substantially underestimated.
Methods

Animals

This study was carried out in accordance with NIH Guidelines, and was approved by the Institutional Animal Care and Use Committee (IACUC) at New York State Psychiatric Institute. Adult (52 weeks old) C57BL/6J male mice were obtained from Jackson Laboratories (n=29). Three month-old CD1 retired breeder male mice were obtained from Charles River, and were used as the aggressors in the social defeat model.

Chronic Social Defeat Stress (CSDS)

Aggressors Prescreening: All CD1 mice used in the experiment were pre-screened for aggressive behaviors as previously described\(^5^0\). During a three-day screening procedure, a novel C57BL/6J mouse was placed in the cage of the CD1 mouse for 3 min. C57BL/6J screener mice were not further used in the study. The latency of the CD1 mouse to attack the C57BL/6J screener mouse was recorded. CD1 mice that attacked in less than 1 minute on at least the last two consecutive screening days were considered to be aggressive.

Experimental Groups: Thirteen C57BL/6J mice underwent social defeat stress, while 6 remained in their cages, to serve as the naïve group. Animals were randomly assigned to these groups. One mouse subjected to social defeat died of unknown causes during the duration of the experiment. Of the remaining 12 social defeat mice, half (n=6) were sacrificed two days after the completion of the stressor, and they are referred to as the ‘stressed’ CSDS group. Three naïve mice were sacrificed at the same time. The other half of the CSDS mice (n=6) along with 3 naïve mice, were allowed an 8.5 weeks (59 days) stress recovery period prior to sacrifice. This group is referred to as the ‘recovered’ group. Animals were selected for the recovery group from the CSDS group so that there was an even ratio of susceptible to resilient mice in the ‘stressed’ and ‘recovered’ groups.
**Social Defeat Paradigm:** Adult experimental male C57BL/6J mice (n = 13) were exposed to a CSDS paradigm daily, for 10 days. The experimental C57BL/6J mice were placed in the cage of a new CD1 aggressor mouse for 5 minutes every day, for 10 consecutive days. After the 5 minutes of physical defeat, the C57BL/6J mice were housed in the same cage as the CD1 aggressor, with a perforated plexiglass divider to separate them for 24 hours. After the 10 days of defeats, experimental C57BL/6J mice and CD1 mice were singly housed. Adult naïve male C57BL/6J mice (n = 6) were housed 2 mice per cage separated by a perforated plexiglass divider. To control for the effects of experimental handling, each naïve mouse was paired with a new naïve mouse every day for 10 days.

**Corticosterone administration (CORT)**

Corticosterone (Sigma Aldrich, St Louis, MO) was dissolved in vehicle (0.45% β-cyclodextrin) at a concentration of 35 µg/ml, equivalent to administration of ~5 mg/kg/day per mouse\(^48\). C57BL/6J mice (n=5) were group-housed and administered corticosterone in their drinking water. Naïve mice (n=5) were group-housed and received only vehicle. Animals were randomly allocated to the two groups. Water bottles were prepared twice a week, and the mice never had access to other water. Behavioral testing began on day 56 of CORT administration. Experimental and naïve mice were sacrificed on day 63 following the completion of behavioral testing. The duration of administration was 9 consecutive weeks. The study design with the number of animals allocated to each condition are listed in **Supplemental Table 1** and in **Extended Data Figure 2**.

**Behavioral tests**

**Social Interaction (SI) Test:** Social avoidance was measured 24 hrs after the completion of the last day of defeat (day 11). During the first trial, experimental mice were allowed to
explore an open field arena (40 cm × 40 cm × 40 cm) containing an empty wire enclosure for 2.5 minutes. During the second trial, a CD1 mouse was placed into the wire enclosure, and the C57BL/J6 mouse was reintroduced for 2.5 minutes. Time spent in the SI zone and time spent in the corner zones during the first and second trial were recorded. SI ratios were calculated as time spent in the SI zone during the second trial divided by time spent in the interaction zone during the first trial. Mice with SI ratios of <1 were considered ‘susceptible’ (n = 6), while mice with SI ratios >1 were considered ‘resilient’ (n = 7). Corner zone ratios were calculated as time spent in the corner zones during the second trial divided by time spent in the corner zones during the first trial.

Open Field Test (OFT)- both groups: OFT were run 24 hrs after the SI tests for CSDS mice, and on day 56 of CORT administration. Each mouse was placed in an open field arena (40 cm × 40 cm × 40 cm) for 10 minutes. A camera on a tripod stand was set up above the arena to record the activity, and the video was later analyzed using Ethovision XT (Noldus). The percent of time spent in the center of the open field (20 cm x 20 cm) and the percent distance traveled in the center of the open field were analyzed.

Elevated Plus Maze (EPM): EPM tests were done the day after the OFT for CORT mice (day 57 of treatment). Each mouse was placed in an elevated plus maze for 10 minutes. A camera on a tripod stand was set up above the arena to record the activity, and the video was later analyzed using Ethovision XT (Noldus). The percent of time spent in the open arms was analyzed.

Novelty Suppressed Feeding (NSF): NSF was performed after EPM for CORT mice (day 59 of treatment), and as previously described. Briefly, NSF testing apparatus consisted of a plastic box (50 x 50 x 20 cm) with 2 cm of wood chip bedding. The center of the arena was brightly lit (1200 lux). Mice were food restricted for 15 h during the dark phase prior to testing. At the time of testing, a single pellet of food (regular chow) was placed on a white paper platform.
positioned in the center of the box. Each animal was placed in a corner of the box, and a stopwatch was immediately started. The latency of the mice to begin eating was recorded during a 10-min test. Immediately after mice took a bite from the food pellet, the pellet was removed from the arena. Mice were then placed back in their home cage and latency to eat and the amount of food consumed in 5 min were measured (home cage consumption). NSF latency was capped at 10 min, with animals that did not consume any portion of the pellet receiving a score of 600 sec.

Behavioral Z-Scores: Social avoidance scores were determined by averaging the z scores for 4 measures as previously described\textsuperscript{24}; SI ratio, corner ratio, time spent in SI zone, and time spent in corner zones. The z scores for SI ratio and time spent in SI zone were multiplied by -1, so that higher scores across all 4 measures indicated higher avoidance. Similarly, the OFT score was determined by averaging the z scores for the 2 measures. The z scores were multiplied by -1, so that higher OFT score indicate higher avoidance of the brightly lit center of the OF. EPM scores were also multiplied by -1, so that higher EPM scores indicate higher avoidance of the open arms of the EPM. High NSF scores already indicate higher avoidance, so they were not inverted. Therefore, across all 4 tests, higher scores indicate higher anxiety-like/avoidant behavior.

Tissue Collection

Animals were sacrificed by rapid decapitation to maintain mitochondrial integrity in brain tissue. Brains were rapidly flash frozen in ice-cold isopentane, stored at -80°C, and later transferred to -170°C (liquid nitrogen vapor) until mitochondrial measures were performed. Brains were transferred back to -80°C during the week prior to being sectioned and were then transferred to -30°C the night before sectioning. Brains were sectioned coronally on a Leica Model CM3050 S cryostat. The cryostat internal temperature and blade temperature were set to
-22°C during sectioning. Brains were mounted onto a specimen disk using optimal cutting temperature (OCT) compound. The brain was sectioned coronally, alternating between two 200μm thick slices and then two 20μm, and were deposited onto microscope slides. Brain sections were kept at -80°C until brain region-specific tissue sample collection. One of the recovered CDSD mouse brain contained blood and could not be reliably sectioned, so this animal’s brain tissue was excluded from analysis. A second CSDS mouse brain and a naïve brain both cracked during slicing, so some of the brain regions (n= 7 for CSDS mouse, n=8 for naïve) could not be obtained from those brains, but the regions that were obtained were included in analysis.

Tissue biopsy punches on frozen brain sections: The scalable Allen Mouse Brain volumetric atlas 2012 was used to determine the location of each brain region of interest, and their distance from bregma. The atlas’ Nissl-stained images were used as a reference for estimating each section’s distance from bregma, which determined which brain slices would be used for punch biopsy collection of each region of interest (Extended Data Fig. 3b). Brain slices were punched using 1.00mm diameter Robbins True-Cut Disposable Biopsy. Two bilateral punches were collected for each brain region over dry ice. For brain regions in the midline, punches were taken from two consecutive slices. All tissue punch locations were approximated by using landmarks on the slice and by comparing to the atlas. It is important to note that the 1mm puncher was larger than the actual brain region in some instances, and so parts of neighboring brain regions may have been included in the punches. Therefore, the punching technique may include some error. Punches were stored at -80°C until they were ready to be used for enzymatic activity assays. Intact consecutive 200μm cerebellar slices were stored at -80°C for immunohistochemical staining.

The tissue punches were too light to be accurately weighed, so we had to estimate the weight based on the reported mouse brain density of 1.04g/cm³. The punches were 0.5mm
(diameter) by 200μm (height), so using the equations \( v = \pi r^2 \times h \) and \( d = \frac{m}{v} \) we obtain an estimated mass of 0.163mg per punch, thus 2 punches were approximated to weigh 0.327mg.

**Mitochondrial Measurements**

*Tissue Preparation:* The punches from each brain region were homogenized in 0.2mL of homogenization buffer (1mM EDTA and 50mM Triethanolamine) (2x 1mm tissue punch/ 0.2mL of homogenization buffer), with 2 tungsten beads to disrupt the tissues’ cells and release the mitochondria. Tissues were homogenized using a Tissue Lyser (Qiagen cat# 85300), which was run at 30 cycles/sec for 1 minute. The tissues were then incubated in ice for 5 minutes, and were then re-homogenized for 1 minute. Tissues were vortexed to ensure homogeneity. Peripheral tissues were cut over dry ice, weighed, and were then homogenized 1:180 (weight:volume, mg: μL), except for heart samples that were further diluted to 1:720 to be in the dynamic range of the assays.

**Enzymatic activities**

Enzymatic activities were quantified spectrophotometrically for Citrate Synthase (CS), complex I (Cl, NADH-ubiquinone oxidoreductase), Succinate Dehydrogenase (CII, succinate-ubiquinone oxidoreductase, also known as SDH), Cytochrome C Oxidase (CIV, COX) and were expressed per mg of tissue, as described previously\(^{32}\), with some modifications as described below in full details. All miniaturized assay measurements were performed in 96-well plates and enzymatic activity assays recorded on a Spectramax M2 (Spectramax Pro 6, Molecular Devices). Linear slopes reflecting changes in absorbance of the reporter dye were exported to Microsoft excel and converted into enzymatic activities using the molar extinction coefficient and dilution factor for each assay. The assays were optimized to determine the minimal amount of tissue required to obtain reliable results assessed by the C.V. between duplicates. The assays were then further optimized to determine the minimal amount of brain tissue homogenate.
required for each individual assay. Assay validation involved regressing increasing tissue amounts (number of punches) with observed activities, which confirmed that an increase or decrease in tissue used produced proportional changes in total activity.

To validate the miniaturized biochemical enzymatic assays, we performed each assay in both the miniaturized 200ul format in 96-well plates, and the traditional 1ml cuvette format, using increasing tissue homogenate volumes (4, 8, 12, 16, 20ul) from the same brain region (cerebellum) of a wild-type control mouse. The same homogenized tissue (1:200 weight:volume, mg:μL) was used for CS, CI, CII, and CIV spectrophotometric assays from which the respective enzyme activities were quantified using the reagents and procedures described below. Both the miniaturized and standard-size cuvette assays showed high agreement ($r^2 = 0.81-0.96; ps = 0.0032-0.037$) (Extended data Fig. 1). For 3 out of 4 assays, compared to the traditional 1ml cuvettes, the 96-well plate exhibited substantially less variation between consecutive dilutions (CS, 8% in the 96-well plate vs. 20% in the cuvette; CI, 15% vs 13%; SDH, 4% vs 9%; COX, 5% vs 9%).

Samples were run in duplicates for each enzyme, along with a nonspecific activity control, and every plate had a positive control (heart homogenate). The 96-well plates were designed so that each brain region/tissue from all animals were run on a single plate, which prevents potential batch variation for comparisons between the animals. However, due to the size of the plates, no more than 2 types of tissues could be run together on a single plate. This limits the accuracy of the comparisons of enzymatic activities between tissues, but maximizes the accuracy of comparisons between animals for each tissue. Reference positive controls on each plate were used to control for potential batches/plates effects. None of the enzymatic activities of some samples (n=8 out of 579, resulting in n=571) could be measured for technical reasons and are therefore not included in the analyses.
Citrate synthase (CS) enzymatic activity was determined by measuring the increase in absorbance of DTNB at 412nm at -30°C in 200μL of a reaction buffer (200 mM Tris, pH 7.4) containing acetyl-CoA 10 mM, 10 mM 5,5'- dithiobis- (2-nitrobenzoic acid) (DTNB), 2 mM oxaloacetic acid, and 10% w/v Triton-x-100. The rate of conversion of DTNB into NTB\(^2^-\) ions indicates the enzymatic activity and is used as a marker of mitochondrial content. Oxaloacetate is removed from the assay mix as a way to measure non-specific activity. The final CS activity was determined by integrating OD412 change over 150-400 seconds and then subtracting the non-specific activity. 10μL of homogenate was used to measure CS activity.

Complex I (CI, NADH-ubiquinone oxidoreductase) activity was determined by measuring the decrease in absorbance of DCIP. The rate of absorbance of DCIP is measured at 600nm at 30°C, in 200μL of a reaction buffer (potassium phosphate 100mM, pH 7.4) containing 550mg/ml bovine serum albumin (BSA), 50mM potassium cyanide (KCN), 20 mM decylubiquinone, and 0.4mM antimycin A. 10μL of homogenate was used to measure CI activity. Antimycin A and KCN are used to inhibit electron flow through complexes III and IV. The negative control condition includes rotenone (200mm) and piericidin A (0.2mM), which selectively inhibit NADH-ubiquinone oxidoreductase. The final CI activity was determined by integrating OD600 change over 150-500 seconds, and by subtracting the rate of NADH oxidation in the presence of rotenone and piericidin A from the total decrease in absorbance.

Complex II (CII, succinate-ubiquinone oxidoreductase, also known as SDH, succinate dehydrogenase) activity was determined by measuring the decrease in absorbance of DCIP. The rate of absorbance of DCIP was measured at 600nm at 30°C, in 200μL of a reaction buffer (potassium phosphate 100mM, pH 7.4) containing 50mg/mL bovine serum albumin (BSA), 500μM rotenone, 500mM succinate-tris, 50mM potassium cyanide (KCN), 20 mM decylubiquinone, 20mM DCIP, 50mM ATP, 0.4mM antimycin A. 15μL of homogenate was used to measure CII activity. The negative control condition includes sodium-malonate, which inhibits
succinate-ubiquinone oxidoreductase. The final CII activity was determined integrating OD600 change over 300-800 seconds, and by subtracting the absorbance in the presence of malonate (500mM) from the total decrease in absorbance.

Complex IV (CIV, also cytochrome c oxidase) activity was determined by measuring the decrease in absorbance of cytochrome c. The rate of conversion of cytochrome c from a reduced to oxidized state was measured at 550nm at 30°C, in 200μL of reaction buffer (100mM potassium phosphate, pH 7.5) containing 10% w/v n-dodecylmaltoside and 120μM of purified reduced cytochrome c. 6μL of homogenate was used to measure COX activity. The negative control condition omits tissue homogenate determine the auto-oxidation of reduced cytochrome c. The final CIV activity was determined by integrating OD550 change over 150-500 seconds, and by subtracting the non-specific activity from the total decrease in absorbance.

Mitochondrial enzymatic activities were determined by averaging the duplicates. The technical variation of the duplicates was measured with and a 10% cutoff. The specific activity of each sample was calculated as the total activity minus non-specific activity (negative control), times the normalization factor. Plates were normalized by their positive controls. Because of some variation in positive controls, each plate’s positive control was z-scored to the average of the positive control activity per assay. All of the activities on the plate were then multiplied by their normalization factor which is determined by 1/z-scored positive control.

**Mitochondrial DNA (mtDNA) quantification**

mtDNA density and mtDNA copy number (mtDNAcn) were measured as previously described 88, with minor modifications. The homogenate used for the enzymatic activity measures was lysed at a 1:10 dilution in lysis buffer (100 mM Tris HCl pH 8.5, 0.5% Tween 20, and 200 g/mL proteinase K) for 10 hours at 55°C, 10 minutes at 95°C, and were kept at 4°C until used for qPCR. qPCR reactions were measured in triplicates in 384 well qPCR plates using a liquid handling station (ep-Motion5073, Eppendorf), with 12μL of master mix (TaqMan
Universal Master mix fast, Life Technologies #4444964) and 8μL of lysate. Each plate contained triplicates of a positive control (heart) and of a negative control (lysate without homogenate). qPCR reaction with Taqman chemistry was used to simultaneously quantify mitochondrial and nuclear amplicons in the same reactions: Cytochrome c oxidase subunit 1 (COX1, mtDNA) and β-2 microglobulin (B2M, nDNA). The Master Mix included 300 nM of primers and 100 nM probe: COX1-Fwd: ACCACCATCATTTCTCCTTC, COX1-Rev: CTCTGCGATGGGCTAGATT, COX1-Probe: HEX/AAGCAGGAG/ZEN/CAGGAACAGGATGAA/3IABkFQ. mB2M-Fwd: GAGAATGGGAAGCCGAACATA, mB2M-Rev: CCGTTCCAGCATTTGGATT, B2M-Probe: FAM/CGTAACACA/ZEN/GTTCACACCCGCTC/3IABkFQ.

The plate was quickly centrifuged and cycled in a QuantStudio 7 flex instrument (Applied Biosystems Cat# 448570) at 50°C for 2 min, 95°C for 20 sec, 95°C for 1 min, 60°C for 20 sec for 40x cycles. To ensure comparable Ct values across plates and assays, thresholds for fluorescence detection for both mitochondrial and nuclear amplicons were set to 0.08. Triplicates for each sample were averaged for mtDNA and nDNA, and an exclusion cutoff of Cts >33 was applied. For samples with triplicates C.V.s > 0.02, the triplicates were checked, and outlier values removed where appropriate, and the remaining duplicates were used. The final cutoff was C.V. > 0.1 (10%); and any samples with a C.V. > 0.1 were discarded. The mtDNAcn was derived from the ΔCt calculated by subtracting the average mtDNA Ct from the average nDNA Ct. mtDNAcn was calculated by 2^{ΔCt} x 2. For measures of mtDNA density, the Ct value was linearized as 2^{Ct} / (1/10^{-12}) to derive relative mtDNA abundance per unit of tissue.

In tissues of similar cellular density (number of cell nuclei per area or mass of tissue), mtDNAcn (mtDNA:nDNA) provides an accurate reflection of mtDNA genome density per cell. However, different brain regions vary widely in their cellularity (up to 8.5-fold), mostly because some defined areas such as the granular layer of the cerebellum are populated with numerous small cell bodies, whereas other regions such as the molecular layer of the DG are completely
acellular and devoid of cell bodies/nuclei. Nevertheless, acellular tissue compartments filled with dentrites can be rich in mitochondria and mtDNA, and therefore the number of mtDNA copies per unit of tissue (μm³ or mg of tissue) is a more generalizable and accurate estimate of mtDNA density between brain regions.

**Mitochondrial Health Index**

The mitochondrial health index (MHI) integrates the 5 primary mitochondrial features, yielding an overall score of mitochondrial respiratory chain activity on a per mitochondrion basis\(^30\). The simple equation uses the activities of Complexes I, II, and IV as a numerator, divided by two indirect markers of mitochondrial content, citrate synthase activity and mtDNA density: \(\text{MHI} = \frac{(\text{CI}+\text{CII}+\text{CIV})}{(\text{CS}+\text{mtDNA density}+1)} \times 100\). A value of 1 is added as a third factor on the denominator to balance the equation. Values for each of the 5 features are mean centered (value of an animal relative to all other animals) such than an animal with average activity for all features will have an MHI of 100 \([(1+1+1) / (1+1+1) \times 100 = 100\].

**Immunohistochemical staining**

Immunofluorescent staining was used to quantify local Succinate dehydrogenase complex, subunit A (SDHA) expression in the cerebellum. Slides with 200μm coronal sections bearing the cerebellum were taken from storage at -80°C and allowed to warm up to room temperature for one hour. The sections were obtained during tissue collection from the same brains as those used for measuring enzymatic activites, but were stored at -80°C and were not biopsied (Extended Data Fig. 4b). The sections were then fixed in freshly prepared 4 % PFA for 5 minutes and immediately washed twice in a 0.1% PBS-Tween 20 (PBS-Tw) solution. They then underwent dehydration through iced cold 70% methanol for 10 min, 95% methanol for 10 min, 100% methanol for 20 min, 95% methanol for 10 min and 70% methanol for 10 min followed by two washes in PBS-Tw. In a PBS-Tw bath, sections were carefully unmounted from the slides using a razor blade and artist’s paintbrush. One section containing an appropriate
The cerebellum region was selected from each animal and individually placed in 10% Normal Donkey Serum (NDS) overnight at 4 °C. They were then incubated in a 1:100 dilution of Rabbit anti-SDHA 1° antibody in PBS overnight at 4°C on a gentle shaker. The following day, they were washed repeatedly in PBS-Tw. Sections were then incubated in a 1:500 dilution of donkey anti-Rabbit 2° conjugated with AlexaFluor 546 in NDS for two hours in a dark box at room temperature. They were then washed in PBS, incubated in 0.625 µg/mL 4′,6-diamidino-2-phenylindole (DAPI) and washed once more. Finally, sections were mounted on Superfrost slides with Prolong Diamond mounting media and a No. 1.5 coverslip.

**Confocal imaging**

The immunofluorescent staining was imaged on a Leica SP8 confocal microscope equipped with Lightning super resolution software. The regions of interest were located using a Leica 20x 0.75 NA objective to determine areas of suitable staining surrounding the purkinje cell layer in the cerebellum. High-resolution images of the area were taken with a Leica 63x 1.40 NA oil objective. For highest resolution, the images were formatted at 2496 x 2496 pixels, with a zoom of 2.50x, line average of 8, pinhole size of 0.50 AU. The final pixel size was 29.58nm x 29.58nm and a z-stack with a step size 0.15 µm was taken through the stained area. A 405nm laser was used for exciting the DAPI channel with a power of 8.0% and gain of 60%. A 561nm laser was used for the AlexaFluor 546 channel with a power of 4.0% and a gain of 20%.

**SDHA staining analysis**

Final images were deconvolved in the Lightning software using the default settings. Images were analyzed in ImageJ version 1.52p. Fifty consecutive slices from each stack were selected for a section thickness of 7.5µm, which roughly covered the depth of penetration of the antibodies. The 16bit images were first thresholded using the Triangle algorithm within the integrated Auto-thresholding plugin to create a binary image. The particle analyzer was then used to quantify the percent of the area stained for each slice. The percent area stained from
each slice was summated to calculate the percent of volume stained. For compartmentalized
data, the molecular and granular layers were separated using a rectangular selection of
approximately 50px width through each stack. For the purkinje layer, a rectangle was drawn that
narrowly captured the stained area of each slice. The volumetric staining for each region of
interest was calculated from the percent of area stained quantified using the particle analyzer.

**TDA-based mapper analysis**

After creating a delta of mitochondrial features between stressed (CORT or CSDS) and
naïve mice, the data matrix for each group was processed through the TDA-based Mapper
pipeline. The input data matrix contained 102 concatenated rows for brain regions (17 brain
regions x 6 mitochondrial features) and 5 or 6 columns for individual mice, based on the number
of animals per group. Missing values, if any, in the input data matrix were interpolated within
group using a linear interpolation. The TDA-based Mapper analysis pipeline consists of four
steps. First, Mapper involves embedding the high-dimensional input data into a lower dimension
$d$, using a filter function $f$. For ease of visualization, we chose $d = 2$. The choice of filter
function dictates what properties of the data are to be preserved in the lower dimensional space.
For example, linear filter functions like classical principal component analysis (PCA) could be
used to preserve the global variance of the data points in the high dimensional space. However,
a large number of studies using animal models and computational research suggest that inter-
regional interactions in the brain are multivariate and nonlinear. Thus, to better capture the
intrinsic geometry of the data, a nonlinear filter function based on neighborhood embedding was
used. The second step of Mapper performs overlapping n-dimensional binning to allow for
compression and reducing the effect of noisy data points. Here, we divided the data into lower
dimensional space into 64 bins with 70% overlap. Similar results were observed for different
number of bins (e.g., 49, and 81). Third, partial clustering within each bin is performed, where
the original high dimensional information is used for coalescing (or separating) data points into
nodes in the low-dimensional space. Partial clustering allows to recover the loss of information incurred due to dimensional reduction in step one\textsuperscript{92}. Lastly, to generate a graphical representation of the shape of the data, nodes from different bins are connected if any data points are shared between them.

The Mapper-generated graphs can be annotated (or colored) using meta-information that was not used to construct the graphs. Here, we annotated these graphs using region-labels to examine whether mitochondrial features were similarity expressed across all regions or whether regional specificity was observed across the two groups. To quantify the extent of segregation (high regional specificity) or integration (low regional specificity) across the six mitochondrial features, we used a graph theoretical measurement of participation coefficient\textsuperscript{54}. Participation coefficient $P_i$ of a node $i$ is defined as:

$$P_i = 1 - \sum_{s=1}^{N_M} \left( \frac{k_{is}}{k_i} \right)^2$$

where $k_{is}$ is the number of links of node $i$ to nodes in community $s$, $k_i$ is the total degree of node $i$ and $N_M$ is the number of communities. The $P_i$ of a node $i$ is close to 1 if its links are uniformly distributed among all communities of the graph (and hence integrated) and it is close to 0 if its links are mostly within its own community (and hence segregated).

**Multi-slice community detection**

One of the most commonly studied mesoscale aspect of a graph is modularity, where highly modular graphs consist of cohesive groups of nodes (or communities) that are more strongly connected to each other than they are to the rest of the network\textsuperscript{93}. Examination of modularity has been recently extended to multi-slice networks that are defined by coupling multiple adjacency matrices across time or modality\textsuperscript{61}. Here, we used six slices, each derived from one of the mitochondrial features, where each slice represented weighted adjacency matrix
of pair-wise Pearson’s correlation between brain regions. We use categorical multi-slice community detection algorithm with the presence of all-to-all identity arcs between slices. The generalized modularity function for multi-slice community detection is given by

\[
Q_{\text{multislice}} = \frac{1}{2\mu} \sum_{ij} \left[ \left( A_{ij} - \frac{k_i k_j}{2m_s} \right) \delta_{sr} + \delta_{ij} C_{jsr} \right] \delta(g_{is}, g_{jr})
\]

Where \( A_{ij} \) represents weighted adjacencies between nodes \( i \) and \( j \) for each slice \( s \), with interslice couplings \( C_{jsr} \) that connect node \( j \) in slice \( r \) to itself in slice \( s \). \( \gamma_s \) represents resolution parameter in each slice; higher value of \( \gamma_s \) (e.g., > 1) detects smaller modules, while lower values (e.g., between 0 and 1) detects bigger modules. In line with previous work, here, we set the resolution parameter across slices to be unity, i.e., \( \gamma_s = 1 \). For simplicity, and as done previously, the interslice coupling parameters were also kept same across slices. Here, we used \( C_{jsr} = 0.1 \). Perturbation of interslice coupling values around 0.1 produced similar results.

To estimate the stability of identified communities across the six mitochondrial slices, under the assumption that stable communities could represent convergence across mitochondrial features, we estimated module allegiance matrix. The module allegiance matrix summarizes the stability of community structure across slices, such that each entry \( i, j \) of the matrix corresponds to the percentage of slices in which regions \( i \) and \( j \) belong to the same community. Finally, an iterative consensus community detection algorithm was run on the module allegiance matrix to define the large-scale networks that are convergent across slices defined by mitochondrial features. The iterative community detection was run multiple times (1,000), followed by consensus clustering to get stable results for identifying large-scale networks of brain regions.

**Comparison of mito-based networks with transcriptomic- and structural connectivity-based networks**
To compare our mitochondrial communities against other modalities, we examined the organization of gene co-expression in the mouse brain as well as mouse structural connectome data. We utilized the Allen Brain Atlas’ ISH (in situ hybridization) feature, which maps each gene onto a reference standard coordinate atlas image, providing a spatial estimate of transcript levels representing gene expression. Specifically, we used the Anatomic Gene Expression Atlas (AGEA) (https://mouse.brain-map.org/agea), which is a three-dimensional male adult C57BL/6J mouse brain atlas based on the ISH gene expression images. The AGEA feature allows for selecting both a ‘seed voxel’ and a target ‘selected voxel’ from exact brain coordinates, yielding the transcriptome-wide correlation between the seed and target regions. The correlation is a measure of the average co-expression between the two voxels. The co-expression values were obtained for all possible pair of brain regions (17x17 matrix), yielding a gene co-expression correlation matrix to which the structure of our mitochondrial networks could be compared.

The structural connectome data were obtained from The Allen Mouse Brain Connectivity Atlas, which is a mesoscale connectome of the adult mouse brain. We utilized the normalized projection strength values for the all brain regions of interest. Because this atlas does not distinguish between dorsal and ventral dentate gyrus, data was obtained for 16 brain regions. For three brain regions where the Atlas provides connectivity values for multiple subregions (OFC, VN, Cerebellum), the connectivity values were averaged across the subregions to yield a global measure for each region, thus matching the dimensionality of our mitochondrial dataset.

We used non-parametric permutation statistics with 10,000 permutations to examine whether the mitochondrial features derived networks were also more densely connected than expected by chance in other modalities (transcriptomic and structural connectivity data). To measure the degree of within-network connectedness we used two established metrics: modularity index (Q_mod; ) and strength fraction (S.F.; ). As presented in Extended Data
**Fig. 11**, for each modality, and across the two metrics, the networks derived from mitochondrial features were more tightly knit than expected by chance (all $ps<0.05$). Thus providing a convergent multimodal evidence of mitochondrial phenotypic organization overlapping with gene expression and the structural connectome.

**Statistical analysis**

Standardized effect sizes (Hedge’s $g$) were computed to quantify the effect of stress conditions on mitochondrial features. Significant effect sizes were determined by the 95% confidence intervals. Two-way ANOVAs were used to compare the effects on a variable between groups. Frequency distributions of the effect sizes for the two stressors and of the mitochondrial-behavior correlations were fitted with Gaussian curves and analyzed by one sample t tests compared to the null distribution. A survival curve using Mantel-Cox log-rank test was generated for novelty suppressed feeding test latencies because the test is capped at 600 sec. Behavioral scores between groups were analyzed using Tukey’s multiple comparison ordinary one-way ANOVA. Correlations between behavioral scores and mitochondrial activities were estimated using Spearman’s $r$ ($r$) to guard against inflation. Correlations between tissues’ mitochondrial activities were measured as Pearson’s $r$, and hierarchical clustering was performed on the data using Euclidian distance with Ward’s clustering algorithm. T-tests were used to compare groups’ correlations. Permutation testing was used to analyze mitochondrial features between versus within brain regions and to analyze the mitochondrial networks against gene co-expression and structural connectome data. To assess the group (CORT vs CSDS) differences in the regional response to stressors are statistically robust, a phase randomized (PR) null approach (Theiler et al. 1992) was used, where we generated 1000 iterations of null data separately for CORT and CSDS groups and ran our TDA-based Mapper approach on each null dataset. PR null preserved the covariance across mice but shuffled any relation between
regional responses. Non-parametric permutation statistics were later estimated for each group. Statistical analyses were performed with Prism 9 (GraphPad) and Metaboanalyst version 4.38.

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**Author contributions**

M.P., C.A., D.D. conceived the project. A.R., P.R., C.A. collected the data. A.J. performed validation experiments. A.R., P.R., A.W.L., M.S. analyzed the data. A.R., M.S., P.R., A.J. generated the figures. A.R., M.S., C.S., E.V.M., D.D., C.A., M.P. interpreted the data. A.R. and M.P. drafted the manuscript. All authors reviewed and edited the final version of the manuscript.

**Competing Interests**

The authors declare no competing interest related to this work. M.P. has consulted and received funding from Epirium Bio. C.A. receives research funding from Sunovion Pharmaceuticals.
Data Availability

The datasets generated during and/or analyzed during the current study are available from the corresponding author upon request.

Code Availability

For TDA Mapper analysis, the generic toolbox is available at https://braindynamicslab.github.io/dyneusr/. The software used for multi-slice network analysis can be accessed at http://netwiki.amath.unc.edu/GenLouvain/GenLouvain. Custom scripts generated for this paper will be made available at https://github.com/braindynamicslab/tda-mito-networks upon publication.
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Figure legends

Figure 1. Behavioral and neuroendocrine stressors enhance the diversity of mitochondrial phenotypes across brain regions. (a) Effect of CORT and CSDS on mitochondrial features across brain regions and peripheral tissues relative to naïve mice. Effect sizes are quantified as Hedge’s g, with significant effect sizes (95% confidence interval) labeled with the fold difference. Unadjusted p value from two-way ANOVA, n=27 mice, 5-6 per group. (b) Pairwise comparisons between each brain regions’ responses to the stressors (Hedge’s g) from (A) as compared to each other region, colored by p-value. (c) Gaussian fit for the frequency distribution of the effect sizes in A on all 6 mitochondrial features in all 17 brain regions (n=102 pairs); one-sample t test (two tailed) against null hypothesis g=0. (d) Number of brain regions in which mitochondrial features are either above or below the control group average in CORT and CSDS mice relative to naïve mice; p values from binomial test (two-tailed), *p<0.05, ***p<0.001. (e) Exemplar representations of Mapper input and output. Nodes in the Mapper output represent regional mitochondrial features (rows from input matrix) that are highly similar across mice. Thus, the brain regions that undergo similar stress-induced recalibrations in specific mitochondrial features cluster together in single nodes (most similar) or interconnected nodes (moderately similar), whereas regions that undergo divergent recalibrations are not connected. The pie-chart-based annotation of graph nodes allowed us to examine the degree of co-regulation of mito-features across brain regions. (f) Topological data analysis (TDA) based Mapper approach to determine if brain regions were co-regulated in their stress-induced mitochondrial recalibrations for the two groups. (g) Graph-theoretical measure of participation coefficient (PC), measuring the degree of segregation/integration among brain regions in each graph, indicating that CORT-induced mitochondrial recalibrations were more region-specific (or segregated) whereas CSDS caused a more integrated response. A phase randomized null approach was used to estimate the robustness of these results, such that null data were generated where covariance across mice (columns in the input matrix) was preserved while the regional relations (rows) were disrupted. Non-parametric permutation statistics revealed CORT group has a significantly lower participation coefficient (PC) as compared to the Mapper graphs generated from the null models (p=0.037). A similar analysis for the CSDS group revealed a trend towards significance for higher PC as compared to the null models (p=0.069).

Figure 2. Association patterns for brain-wide mitochondrial phenotypes and mouse behavior. (a) Mitochondrial phenotyping and behavioral profiling of inter-individual variation in a heterogenous population of mice; OFT, open field test; EPM, elevated plus maze; NSF, novelty suppressed feeding; SI, social interaction test. (b) Gaussian fits of the frequency distributions for all correlations (n=102) between the composite mitochondrial health index (MHI) and each behavioral test; one-sample t test (two tailed) against null hypothesis r=0 (other mitochondrial features are shown in Extended Data Fig. 8). (c) Individual correlations for the 17 brain region, across the 6 mitochondrial features, for each behavioral score, quantified as Spearman’s r. OFT and EPM behavioral scores were inverted so that higher scores on all four tests indicate higher anxiety (see Extended Data Fig. 8 for additional details). The strongest correlations for each behavioral test are denoted by yellow boxes, with the scatterplots shown below. An adjusted p
value of <0.002 was applied (false-discovery rate 1%). All tests have been adjusted so that a higher score indicates higher anxiety-like behavior (see Methods for details). (d) Average correlation of each behavior for the brain (B) and tissue (T) mitochondrial features; two-way ANOVA. n=10-27 mice per behavioral test.

**Figure 3.** Mitochondrial phenotype-based connectivity across anatomical brain regions. (a) Connectivity matrix of mitochondrial features across brain regions, using all 6 mitochondrial features across the animal cohort (n=27 mice), quantified as Pearson’s r. The matrix is ordered by hierarchical clustering (Euclidian distance, Ward’s clustering). (b) Cross-correlation of each mitochondrial feature to the other 5 measures within each brain region (n=17 brain regions). (c) Global connectivity based on the average correlation for each brain region with all other regions. (d) Average correlation of mitochondrial features between brain region’s, between peripheral tissues, and between brain regions and tissues; p<0.0001, Ordinary one-way ANOVA with multiple comparisons. (e) Multi-slice community detection analysis on mitochondrial measures across the 17 brain regions, with mitochondrial features represented in six separate layers, resulting in (f) three distinct communities or brain networks. Modular structure confirmed by permutation test, p<0.0001). (g) Average mito-behavior correlation by module for each behavioral test (top). The bottom panel shows networks with each region color-coded by its average correlation with behaviors; for OFT and EPM (left) and SI (right). Modularity metrics anchored in whole-brain transcriptome and the structural connectome data are shown in Extended Data Fig. 10.

**EXTENDED DATA FIGURE LEGENDS**

**Extended Data Fig. 1.** Validation of miniaturized 96-well plate enzymatic activity assays. Comparison between miniaturized 96-well plate and traditional 1ml cuvette for enzymatic activities of (a) citrate synthase (CS), (b) NADH-ubiquinone oxidoreductase (Complex I), (c) succinate-ubiquinone oxidoreductase (Complex II), and (d) cytochrome c oxidase (Complex IV).

**Extended Data Fig. 2.** Comparison of mtDNA copy number versus mtDNA density. (a) Diagram to illustrate the difference between measures of mtDNA copy number (mtDNAcn=mtDNA/nDNA ratio) and mtDNA density (mtDNA copies per mg of tissue). (b) Correlation between the average mtDNAcn and average mtDNA density per brain region. (c) Correlations between mtDNAcn (left) and mtDNA density (right) with the other five mitochondrial measures in all animals for each brain region and tissue, measured by Spearman’s r. The strongest positive and negative brain correlations for each are highlighted with yellow boxes and plotted in (d).

**Extended Data Fig. 3.** (a) Experimental design. (b) 17 brain regions of interest, labeled by name and distance from bregma, with red circles indicating bilateral punch locations. Images acquired from the Allan Mouse Brain Atlas (Dong, H. W. The Allen reference atlas: A digital color brain atlas of the C57Bl/6J male mouse. John Wiley & Sons Inc. (2008). Abbreviated brain
region names are as follows (orders by Bregma coordinates): mOFC, mPFC, CPu, NAc, M1, Hypoth, Thal, DGd, Amyg, CA3, VTA, V1, SN, DGv, PAG, Cereb, VN.

Extended Data Fig. 4. Cerebellar staining for mitochondrial protein quantification. (a) Coronal 200μm-thick cerebellar slice stained for DAPI (blue, cell nuclei), and the mitochondrial respiratory chain Complex II subunit SDHA (yellow). (a’) higher magnification of A, and (a”) example high-magnification of SDHA signal along the purkinje cell layer, highlighting three regions of interests used for quantification of SDHA density among the cerebellar molecular, purkinje, and granular layers. (b) Diagram illustrating that spectrophotometric measurements of CII activity and immunofluorescence confocal microscopy measurements for protein abundance were measured in contiguous 200μm-thick cryosections from the same animals. (c) Example raw spectrophotometric trace for CII activity in the cerebellum, integrating OD600 change over 300-800 seconds. (d) Correlations between layer-specific SDHA density (% SDHA-positive pixels per volume) and the biochemical activity of Complex II, demonstrating that protein abundance is not an appropriate surrogate for biochemical activity.

Extended Data Fig. 5. Animal-to-animal differences in mitochondrial features for each brain region and peripheral tissue. (a) Diagram illustrating the range of mitochondrial content and function across animals. (b) Example C.V. (coefficient of variation) calculation for CS activity in the substantia nigra (SN). Each datapoint is an animal (n=27 mice), illustrating the variation in mitochondrial features between animals. (c) Individual C.V.s per mitochondrial feature for each brain region and peripheral tissue. (d) C.V.s for brain regions by mitochondrial feature for naïve animals only. Each datapoint is the average C.V. for a brain region or tissue (n=22). (e) C.V.s for brain regions versus tissues by mitochondrial feature for all animals (naïve, CORT, CSDS), two-way ANOVA.

Extended Data Fig. 6. Effect size of stressors on brain regions’ and other tissues’ mitochondrial content and functioning. (a) Effect of CORT treatment and (b) effect of CSDS treatment on brain regions and tissues, quantified by standardized Hedges g, compared to naïve mice. (c) Example plot if both stressors had the same effect on a brain region, with quartiles labeled and (d) plot of the effect sizes of both stressors on each brain region’s MHI.

Extended Data Fig. 7. Exploratory analysis comparing susceptible and resilient subgroups of CSDS mice. Effect sizes for mitochondrial outcomes shown separately by brain region for animals classified as susceptible or resilient to CSDS (Methods). Effect sizes are Hedge’s g, with significant effect sizes (95% confidence interval) labeled with the % change compared to naïve (non-stressed) mice.

Extended Data Fig. 8. Effect of recovery from social defeat stress on mitochondrial features across brain regions. (a) Average effect of 2 months recovery (Rec) from CSDS stress across the 6 mitochondrial features for each brain region, as compared to non-recovered CSDS mice. Effects sizes are quantified by Hedges g, with 95% confidence intervals. MHI in recovered mice was higher than non-recovered CSDS in some regions (Caudoputamen, Hypothalamus-CA3), and lower in others (Hypothalamus, Medial orbitofrontal cortex, Vestibular
nucleus). (b) Average effect sizes of 6 mitochondrial features of non-recovered CSDS mice and recovered mice compared to naïve mice in three selected brain regions from A (red boxes). (c) Frequency distribution of the effect size of CORT, CSDS, and recovery as compared to naïve mice on all 6 mitochondrial measures in all 17 brain regions, gaussian-fitted curve, indicating that recovered mice have lower MHI than naïve mice and marginally higher MHI than CSDS (**p<0.0001, one-sample t test (two tailed)). (d) TDA analysis using Mapper of mitochondrial measures in naïve mice, CORT, CSDS, and CSDS-recovered mice, revealing the recovered group to be more similar to the non-recovered CSDS group than to naïve mice.

Extended Data Fig. 9. Behavioral test results (a) To account for both the % time and % distance traveled in the center, the two measures were z-scored and averaged, creating a single Open Field Test (OFT) score. Z-scores were inverted so that a higher OFT score represents less time and distance in the center, indicating higher anxiety. Open field was run for all groups. (b) Elevated plus maze (EPM), measuring percent of time spent in open arms/total time, with the score being multiplied by -1 so that higher scores represent less time spent in the open arms, which indicates higher anxiety. EPM was run only on CORT mice. (c) Novelty suppressed feeding (NSF) test used to measure the latency to feed in a novel environment, with the test being capped at 600 sec. Cum. summ=cumulative survival, meaning percent of mice that have not eaten; Survival curve: Mantel-Cox log-rank test. NSF was run only on CORT mice. (d) Social Interaction test (SI), represented by a Social Avoidance score for all CSDS mice, separated by susceptible (sus.) and resilient (res.), and recovered (rec). Social avoidance was measured as the average of 4 scores; z scored social interaction ratio, z scored time spent in interaction zone, z scored time spent in corner, and z scored corners ratio. Social interaction ratio and time spent in SI zone were inverted so that higher values for all four measures indicate higher avoidance. p<0.05, p<0.01, Tukey’s multiple comparison ordinary one-way ANOVA.

Extended Data Fig. 10. Frequency distribution of all correlations of mitochondrial-behavior pairs by behavioral test. Gaussian fits, p values from one-sample t test (two tailed) against the null hypothesis (r=0).

Extended Data Fig. 11. To examine whether the functional organization of the brain revealed using mitochondrial features is also evident cross-modally, we utilized non-parametric permutation statistics with 10,000 permutations to compare the topology of our mito-derived networks with the expected topology based on the transcriptomic and structural connectomes in the Allen Brain Atlas (see Online Methods for details). Specifically, we examined i) whether the tightly knit networks (or communities) observed in the mitochondrial features are more densely connected than expected by chance (a-b); and ii) whether the networks derived from mitochondrial features were also more densely connected than expected by chance in other modalities, including gene-coexpression data (c-d) and EYFP-based structural connectivity (e-f). To measure the degree of within-network connectedness we used two established metrics: modularity index (Q_mod; Newman 2006) and strength fraction (S.F.; Richardi et al. 2015). The histograms depict distribution of within-network connectedness generated using 10,000 permutations (or data shuffling). The real value of within-network connectedness is shown using red dots across all plots. For each modality, and across the two metrics, the networks derived
from mitochondrial features were more tightly knit than expected by chance. Hence, providing a convergent multimodal evidence of mitochondrial functional organization overlapping with gene expression and the structural connectome.
Table 1. Expanded abbreviations and Bregma coordinates. All regions were taken bilaterally except those marked with *, which were obtained by collecting tissue from two consecutive slices.

| Abbreviation | Region name                      | Bregma / Anatomical location |
|--------------|----------------------------------|------------------------------|
| mOFC         | Medial orbitofrontal cortex      | 2.42                         |
| mPFC         | Medial prefrontal cortex         | 1.22                         |
| CPu          | Caudoputamen                     | 1.22                         |
| NAc          | Nucleus accumbens                | 1.22                         |
| M1           | Primary motor cortex             | 1.22                         |
| Hypoth       | Hypothalamus                     | -0.48*                       |
| Thal         | Thalamus                         | -0.68*                       |
| DGd          | Dorsal dentate gyrus             | -1.78                        |
| Amyg         | Amygdala                         | -1.78                        |
| CA3          | CA3 region                       | -2.58                        |
| VTA          | Ventral tegmental area           | -2.58                        |
| V1           | Primary visual cortex            | -2.58                        |
| SN           | Substantia nigra                 | -3.08                        |
| DGv          | Ventral dentate gyrus            | -3.38                        |
| PAG          | Periaqueductal grey              | -3.78*                       |
| Cereb        | Cerebellum                       | -6.38                        |
| VN           | Vestibular nuclei                | -6.38                        |
| AG           | Adrenal glands                   | Both glands pooled           |
| Liver        | Liver                            | Inferior portion of left lobe|
| Heart        | Heart                            | Inner wall of left ventricle |
| Soleus       | Red oxidative skeletal muscle    | Tendon-to-tendon, right leg  |
| WG           | White glycolytic skeletal muscle | Superficial medial head, right leg |
**Supplemental Table 1.** Study design with the number of animals allocated to each condition

| Experimental group | CSDS | CSDS-recovered | CORT | Naïve (CSDS-matched) | Naïve (CSDS Recovered) | Naïve (CORT) |
|--------------------|------|----------------|------|----------------------|------------------------|--------------|
| Number of animals  | 6    | 6              | 5    | 3                    | 3                      | 5            |
| Day at sacrifice   | 14   | 71             | 63   | 14                   | 71                     | 63           |

(total n=28).
**Fig. 1**

### a. Mitochondrial recalibrations (stress-control)

| Region | CORT (2 months) | CSDS (10 days) |
|--------|-----------------|-----------------|
| Amyg   |                 |                 |
| CPu    |                 |                 |
| SN     |                 |                 |
| NAc    |                 |                 |
| V1     |                 |                 |
| CA3    |                 |                 |
| mPFC   |                 |                 |
| M1     |                 |                 |
| Hypoth |                 |                 |
| Cereb  |                 |                 |
| mOFC   |                 |                 |
| PAG    |                 |                 |
| DGd    |                 |                 |
| VTA    |                 |                 |
| Thal   |                 |                 |
| DGv    |                 |                 |
| VN     |                 |                 |
| Heart  |                 |                 |
| Liver  |                 |                 |
| Soleus |                 |                 |
| AG     |                 |                 |
| WG     |                 |                 |

### b. Statistical differences between regions

- Amyg: 42% 48%
- CPu: 31%
- SN: 31%
- NAc: 31%
- V1: 42%
- CA3: 31%
- mPFC: 42%
- M1: 31%
- Hypoth: 42%
- Cereb: 31%
- mOFC: 42%
- PAG: 31%
- DGd: 42%
- VTA: 31%
- Thal: 42%
- DGv: 31%
- VN: 42%

### c. Global effects of stressors

- Number of brain-mito features combinations
- Global effects of stressors
- Participation Coefficient
- Degree of segregation vs integration

### d. Direction of mitochondrial responses to stressors

- Hedge's g
- Effect sizes (g)

### e. Mapper input:

- Mouse: M, M', M''
- Stressor: St, St', St''

### f. Regional responses to stressors — Topological Data Analysis (TDA)

- Amyg
- CPu
- SN
- NAc
- V1
- CA3
- mPFC
- M1
- Hypoth
- Cereb
- mOFC
- PAG
- DGd
- VTA
- Thal
- DGv
- VN
- Heart
- Liver
- Soleus

### g. Degree of segregation vs integration

- CORT
- CSDS

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**Note:** The image contains complex illustrations and tables that represent mitochondrial recalibrations, statistical differences between regions, global effects of stressors, and regional responses to stressors using Topological Data Analysis (TDA). The data is presented in a visual format to aid in understanding the underlying concepts and results.
Fig. 2

(a) Heterogenous animal population

(b) MHI-behavior correlations

(c) Spearman r

(d) Average correlation
Connectivity matrix based on mitochondrial features

- Brain regions (17):
  - Cereb
  - NAc
  - CA3
  - PAG
  - DGd
  - DGv
  - mPFC
  - CPu
  - M1
  - VN
  - Thal
  - VTA
  - mOFC
  - Hypoth
  - Amyg
  - V1
  - SN
  - CA3
  - Soleus
  - AG
  - Liver
  - Heart
  - WG

- Peripheral tissues (5):
  - Cereb
  - NAc
  - PAG
  - DGd
  - DGv
  - mPFC
  - CPu
  - M1
  - VN
  - Thal
  - VTA
  - mOFC
  - Hypoth
  - Amyg
  - V1
  - SN
  - CA3
  - Soleus
  - AG
  - Liver
  - Heart
  - WG

- Connectivity by mito feature:
  - CS
  - CI
  - CII
  - CIV
  - mtDNA
  - MHI

- Global connectivity by brain region

- Average Inter-regional Correlation

- Average Correlation with other mito-measured Inter-mito feature correlations

- Brain vs tissues

- Average inter-tissue correlation

- Regional co-variation for each mito feature across 17 brain regions

- 6 mito features

- Mitochondria-driven community detection among brain regions

- Categorical multi-slice community detection

- Module allegiance

- Mito-behavior correlations by network

- Effect size (r)

- Anxiety-related behaviors correlations (OFT & EPM)

- Social avoidance correlations (SI)

- Fig. 3