The mitochondrial permeability transition phenomenon elucidated by cryo-EM reveals the genuine impact of calcium overload on mitochondrial structure and function

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Mitochondria have a remarkable ability to uptake and store massive amounts of calcium. However, the consequences of massive calcium accumulation remain enigmatic. In the present study, we analyzed a series of time-course experiments to identify the sequence of events that occur in a population of guinea pig cardiac mitochondria exposed to excessive calcium overload that cause mitochondrial permeability transition (MPT). By analyzing coincident structural and functional data, we determined that excessive calcium overload is associated with large calcium phosphate granules and inner membrane fragmentation, which explains the extent of mitochondrial dysfunction. This data also reveals a novel mechanism for cyclosporin A, an inhibitor of MPT, in which it preserves cristae despite the presence of massive calcium phosphate granules in the matrix. Overall, these findings establish a mechanism of calcium-induced mitochondrial dysfunction and the impact of calcium regulation on mitochondrial structure and function.

Mitochondria regulate cell fate through a variety of means1–5. Their extensive networks and dynamic architecture facilitate metabolic signaling to ensure proper cellular function and survival. Mitochondria achieve this by integrating intracellular cues and physiological stimuli to regulate ATP production, metabolite oxidation, calcium signaling, phospholipid and steroid hormone biosynthesis, and mitochondrial fission and fusion processes6–13. As such, mitochondria must operate under a range of physiological conditions including transient changes in energy demand, oxidative stress, and moderate calcium overload. For example, in highly metabolic organs such as the heart, brain, and kidney, their response to these conditions is crucial for cell survival14. However, in pathological conditions, such as during an ischemia/reperfusion event, mitochondria undergo a phenomenon known as the mitochondrial permeability transition (MPT). MPT is a gateway mechanism for cell death and involves the opening of a non-selective pore that allows small molecules and metabolites up to 1.5 kDa in size to freely diffuse across the inner mitochondrial membrane15,16. When the pore is open, the membrane potential is dissipated, there is a loss of respiratory control, ATP is hydrolyzed, and osmotic swelling occurs5. The swelling causes inner membrane unfolding, outer membrane rupture, and eventually release of apoptogenic molecules, including cytochrome c (cyt. c) that ends in cell death.

While the consequences of MPT are well appreciated, the molecular composition of the pore is currently unknown. The MPT phenomenon was first observed nearly seven decades ago when early studies in the mid-1950s to early 1960s demonstrated massive mitochondrial swelling under certain conditions17–21. These conditions involved calcium overload, high inorganic phosphate concentrations, fatty acids, oxidative stress, and...
protective therapies. However, our new data add exciting therapeutic targets for mitochondrial overload. This mechanism is not mutually exclusive with the current dogma as it integrates many past findings in a concise, overarching theoretical framework. Instead of focusing on the pore, we sought to investigate the consequence of excessive calcium overload on a population of isolated mitochondria by analyzing cryo-electron microscopy (cryo-EM) time-course data. This powerful imaging technique was coupled with high-resolution respirometry and spectrofluorimetry to structurally analyze the effect of calcium overload on mitochondrial function. We identified a novel mechanism that links calcium phosphate granule formation to cristae structural changes, inner membrane fragmentation, and ultimately mitochondrial permeabilization.

Figure 1. EGTA rescues mitochondrial function at low-to-moderate calcium loads but not high loads. (A) Representative traces of ADP-stimulated respiration from calcium loaded mitochondria following the addition of EGTA. Mitochondria (0.1 mg/mL) were energized with 5 mM sodium pyruvate and 1 mM L-malate and exposed to various calcium boluses (0, 12.5, 25, and 50 µM). Five mins after calcium uptake, 1 mM EGTA was added to chelate all calcium in the system. Following an additional 5 min, a bolus of 500 µM ADP was added to induce maximal ADP-stimulated respiration. (B) ADP-stimulated mitochondrial respiration is recovered after EGTA addition for all but the 50 µM calcium bolus. (C) However, the outer membrane integrity is maintained at the highest calcium dose as shown after the addition of a 10 µM cyt. c. Data are presented as mean ± standard deviation for a sample size of n = 4. Statistical comparisons are made with respect to 0 µM calcium. *Represents a p value < 0.01.

Results

Respiratory inhibition by calcium overload is reversible in low-to-moderate calcium load. While mitochondrial calcium concentrations lower than 100 nmol calcium/mg mitochondria support ATP production, levels above 500 nmol/mg mitochondria depress oxidative phosphorylation. In one of these studies, it was proposed that calcium phosphate precipitates form in the mitochondrial matrix at high calcium loads and reduce ATP production rates by either impeding metabolite transport and diffusion or destabilizing cristae, the functional units of mitochondria. However, the lasting effects of significant calcium accumulation were not explored in either of these studies. To test this, we monitored mitochondrial respiration rates following the addition of the calcium chelator EGTA under various calcium boluses in the range of 0–500 nmol/mg as shown in Fig. 1.

The results in Fig. 1 show that the inhibitory effect of calcium overload is reversible for all but high calcium loads. As expected, the respiratory rates before calcium addition were equal across conditions. After the calcium-dependent transient increase in respiration, respiration remains elevated due to the activation of calcium-sensitive matrix dehydrogenases and sodium/calcium cycling. When 1 mM EGTA was added, the ADP-driven respiratory rates were similar across all conditions except for the highest dose tested.

To rule out that the observed decrease in mitochondrial respiration for the 50 µM CaCl₂ bolus is due to mPT activation, we assessed outer membrane integrity by exposing mitochondria to an exogenous bolus of 10 µM cytochrome c (Fig. 1C). As shown, mitochondrial respiration did not go above 5% relative to the baseline for either condition confirming that the outer membrane was mostly intact at the highest calcium dose and disputes the notion that the observed detrimental effects on respiration are caused by mPT activation. These results suggest
that when calcium overload exceeds a certain threshold, mitochondrial oxidative phosphorylation is irreversibly inhibited. This effect does not involve mitochondrial calpains34 and may involve some sort of structural change that lowers ATP production rates. Thus, the effect of calcium overload lies on a spectrum whereby higher levels of calcium result in detrimental changes in mitochondrial bioenergetic pathways.

CsA preserves the mitochondrial function under high calcium loads. We then measured mitochondrial respiratory rates during excessive calcium overload by adding a 75 μM or 100 μM calcium bolus in the presence or absence of CsA, a known PTP inhibitor (Fig. 2A,B). In agreement with results from Fig. 1, increasing the extent of calcium overload impairs oxidative metabolism. However, the depressive effects of calcium on ADP-stimulated respiration are much more severe at these higher doses. The respiratory rate after ADP addition drops below 50 nmol O2/mg/min after the 75 μM CaCl2 bolus and drops below 20 nmol O2/mg/min for the 100 μM CaCl2 bolus. When CsA was present, this calcium-dependent inhibitory effect is partially mitigated with rates reaching nearly 320 nmol O2/mg/min after the 75 μM bolus and 280 nmol O2/mg/min for the 100 μM bolus. Therefore, as others have found, CsA partially preserves mitochondrial function in the face of overwhelming calcium overload31,38–41. This effect is typically attributed to the ability of CsA to inhibit PTP opening. Our structural data shown in the following sections suggest the existence of a novel protective effect of CsA.

In addition to the respirometry studies, mitochondrial absorbance data obtained in parallel (Fig. 2C) shows that only the 100 μM calcium bolus elicited large amplitude swelling, a classic indicator of mitochondrial permeability transition42. In contrast, the addition of a 75 μM calcium chloride bolus induced an increase in absorbance due to the formation of calcium phosphate granules scattering light at this wavelength43. The gradual decrease in absorbance that follows is attributed to mitochondria fragmenting over time in response to the calcium insult. For both CsA-treated groups, the calcium-dependent increase in absorbance was sustained followed by a much...
effectively take up and store the 100 µM CaCl2 bolus at all when CsA was absent. This level of calcium overload and the process repeats until the entire population is compromised. In contrast, mitochondria were not able to take up even more calcium16,43,47. This results in additional mitochondria losing their ability to retain calcium, which mitochondria with lower calcium tolerances release their calcium loads and force other mitochondria to differ.

Results are similar to findings from a recent study that looked at the effects of the mitochondria-targeting peptide SS-31 on reducing infarct size of reperfused ischemic hearts46. Our interpretation of the absorbance data is supported by the calcium uptake data shown in Fig. 2D. These data also demonstrate the profound beneficial effects of CsA on mitochondrial calcium sequestration. When CsA was absent, mitochondria were not able to maintain calcium homeostasis and calcium was released into the buffer. For the 75 µM CaCl2 challenge, this release was gradual and suggests there is a snowball-like effect in which mitochondria with lower calcium tolerances release their calcium loads and force other mitochondria to take up even more calcium16,43,47. This results in additional mitochondria losing their ability to retain calcium, and the process repeats until the entire population is compromised. In contrast, mitochondria were not able to effectively take up and store the 100 µM CaCl2 bolus at all when CsA was absent. This level of calcium overload is sufficient to rapidly compromise the entire population in short order.

Elucidating the effects on calcium overload on mitochondrial ultrastructure. To capture mitochondria undergoing MPT during calcium overload, we used the sampling scheme shown in Fig. 3. These samples were drawn from a cuvette of isolated mitochondria at the indicated time points and subsequently vitrified in liquid ethane and imaged using cryo-EM. A total of 1345 cryo-EM images were analyzed and organized by sample time-point; before adding calcium (t0), approximately 1.5 min after calcium addition (t1), 4 min after calcium addition (t2), and 10 min after calcium addition (t3). The effect of calcium in the presence or absence of CsA was quantified for each time-point. In the absence of CsA, mitochondrial respiration decreases dramatically as a function of time and the effect is exacerbated at greater calcium loads. In the presence of CsA, mitochondrial respiration was maintained. Data are presented as mean ± standard deviation for n = 3–5 biological replicates.

Mitochondria in stage 1 have intact inner and outer membranes and are typically round (Fig. 4A and Supplementary Fig. S1). Cristae structures in this set of images are hard to distinguish; however, some are identifiable. Before the addition of a calcium bolus mitochondria are smaller with some calcium granules due to contaminating calcium (~4 µM) from buffer solutions as shown by the insets in Fig. 4B–D. The number of calcium phosphate granules is relatively low with sizes averaging less than 100 nm in size. After the addition of 75 µM CaCl2, mitochondria begin to fragment and lose bioenergetic competency. The beginning of this process is characterized by stage 2 (Fig. 4 and Supplementary Fig. S2). In this stage, regions of localized outer membrane rupture are observed and are always accompanied by the appearance of calcium phosphate granules. While the size of granules within a mitochondrion does not vary significantly, differences between mitochondria are common and noticeable (Supplementary Fig. S3). During the transition from stage 2 to stage 3, outer membrane definition is lost, and the inner membrane is released. The inner membrane also begins to fragment in this stage. In some instances, calcium phosphate granules are still present indicating that the inner membrane is still energized. However, there are also images of this stage showing granules in the middle of dissolution (Supplementary Fig. S4), suggesting this stage is when depolarization begins. Unexpectedly, the core of the granules appears to dissolve first. In stage 4, the outer membrane is almost entirely gone, and the inner membrane is extensively fragmented. Stage 5 is characterized by the complete fragmentation of the mitochondrial inner membrane and is the dominant stage at the 15 min time point. In this stage, mitochondria are deenergized and contain no calcium phosphate granules.
There were no large differences in mitochondrial sizes between the time points (ranging 500–3000 nm), but there were some clear differences in the size and number of granules (Fig. 4B–D). As mitochondrial transition from stage 1 to stage 3, the increase in absorbance shown in Fig. 2C is caused by the increases in numbers and sizes of calcium phosphate granules. In fact, the number of calcium phosphate complexes per mitochondrion reached a maximum by stage 3 and decreased in the following stage as shown in Fig. 4C. The decreased in size and abundance by stage 4 is due to more complete mitochondrial permeabilization and fragmentation. Hence, for the first time to our knowledge, the MPT phenomenon now has direct visual confirmation of the processes proposed to occur. However, our results elucidate a mechanism that pinpoints cristae remodeling and inner membrane fragmentation as the key determinant of mitochondrial dysfunction as discussed further below.

CsA preserves the inner membrane, promotes the formation of granules of greater size, and increases the abundance. Next, we repeated the calcium overload imaging experiments in the presence of CsA to understand how mitochondrial respiration and calcium handling were preserved from an ultrastructural perspective (Fig. 5 and Supplementary Fig. S5). Like control mitochondria, CsA-treated samples were grouped into 4 classes based on morphology (Supplementary Table 2). However, the classes are not related to a sequence of events like the stages, rather they are descriptive. Many of the images showed normal-looking mitochondria with well-defined inner and outer membranes. These are class 1 mitochondria. Some of these mitochondria contained granules caused by the uptake of low levels of contaminant calcium. In addition, some mitochondria had a condensed inner membrane that was sometimes localized to one side of the mitochondrion. These electron-dense regions are presumably areas of high cristae density. Interestingly, some images showed mitochondria with the outer membrane ruptured with the inner membrane partially or more completely ejected from the mitochondrion. These mitochondria are classified as class 2 mitochondria. In other images, mitochondria were clustered together and are defined as class 3 mitochondria. Lastly, after the treatment, images revealed mitochondria with no outer membrane, large calcium phosphate granules, and the inner membrane spread across the carbon grid. These mitochondria are classified as class 4 mitochondria. Because of morphological changes induced by CsA and calcium addition, the sizes of these mitochondria are larger than mitochondria in the other classes. In addition, mitochondria in this class had granules of heterogeneous sizes between mitochondria but rarely within a single mitochondrion (Supplementary Fig. S6). Despite these radical changes in ultrastructure, the mitochondria remain functionally competent as shown in Fig. 2. The best explanation for this observation is that the cristae junctions and inner membrane integrity are preserved by the CsA treatment.

After calcium addition, the abundance and size of granules per mitochondrion and the mitochondrial size increased in the presence of CsA compared to the control group (Fig. 5). Before the addition of calcium, the average control mitochondrial size was 1320 ± 550 nm, the average granule size of 68 ± 14 nm, and the average number of granules per mitochondrion was 5.1 ± 2.3. These results show that CsA does not influence any of these parameters before the large calcium bolus was administered. However, after calcium addition, there are noticeable differences between control and CsA-treated mitochondria. The mitochondrial size for control averages 1470 ± 530 nm with a granule size of 90 ± 22 nm and an abundance of 18.0 ± 4.3 per mitochondrion. Whereas CsA-treated mitochondria size average was 1630 ± 400 nm with a granule size of 102 ± 36 nm and an abundance of 26.0 ± 5.1 per mitochondrion. Of note, only a small number of control mitochondria survived by the last time point (Fig. 5C).

Mitochondrial membrane fragmentation occurs more rapidly at greater calcium loads but is mitigated by CsA. Seeking to understand the observed large amplitude swelling for the control group after the addition of a 100 μM CaCl₂ bolus, images of control mitochondria were collected after a 100 μM CaCl₂ addition. Most of the images displayed outer membrane rupture at multiple regions suggesting a rapid expansion of the inner membrane compared to the 75 μM CaCl₂ bolus (Fig. 6 and Supplementary Fig. S7 and S8). Thus, at this high of a calcium bolus, the morphological changes were caused by what appears to be bona-fide permeability transition pore opening. As expected, CsA prevented this rapid expansion and led to the formation of numerous and large calcium phosphate granules. Without CsA, the size and abundance of the granules were noticeably decreased (Fig. 6B). While there were no differences in mitochondrion sizes between treatments shortly after calcium addition (1320 ± 370 nm vs 1490 ± 380), the average control mitochondrial size decreased to 1160 ± 440 nm in the last time point (Fig. 6C). In contrast, the average CsA-treated mitochondrial size increased to 1710 ± 440 nm. The average number of granules in control mitochondria as a function of time was reduced from 9.6 ± 3.1 to 6.6 ± 2.6. The average size of these granules marginally increased from 84 ± 32 to 90 ± 37 nm. Whereas the average number of granules in CsA-treated mitochondria increased from 46.1 ± 6.8 to 107 ± 10 with average sizes increasing from 121 ± 21 to 132 ± 28 nm. These values are greater compared to the values measured after a 75 μM CaCl₂ bolus was given. This is consistent with CsA increasing calcium accumulation and preserving mitochondrial function even at these high calcium loads. However, the oxygen consumption rate after the 100 μM CaCl₂ bolus was significantly lowered compared to the 75 μM CaCl₂ bolus (Fig. 2A,B). Hence, we conclude that calcium induces irreversible effects on mitochondrial function and CsA, although not entirely protective, delays complete loss of function, and allows more calcium uptake.

Calcium phosphate granules are composed of smaller structural units. Calcium phosphate complexes are considered the main component of the mitochondrial calcium sequestration system. Pioneering studies by Posner and others suggested that amorphous calcium phosphate consists of many smaller spherical elements with a chemical composition of Ca₅(PO₄)₆·H₂O. These elementary units were named Posner clusters with a diameter ranging from 0.7 to 1.0 nm. In the present study, we lack the image resolution to resolve indi-
that show CsA increases the calcium retention capacity by nearly 3-fold. While an attractive hypothesis, this idea is based on studies that show that it only increases the calcium threshold required to open the pore. This fact that both images in Fig. 7B show intact outer membranes, combined with the results from Fig. 1C, further supports that the absence of granules in uncoupled mitochondria is not due to bona fide transition but rather the effect of a collapsed membrane potential and subsequent calcium efflux. Absence of granules in mitochondria under extreme high calcium conditions.

Membrane potential is required to form calcium phosphate granules. Calcium uptake by the mitochondria is highly dependent on the mitochondrial energetic status to regulate ATP production. While the intracellular calcium content is maintained in the 100 nM range, mitochondria gradually accumulates and buffers calcium within the mitochondrial matrix using the membrane potential generated across the inner mitochondrial membrane. This results in calcium phosphate granules formation. To further show this dependency, we used cryo-EM to visualize mitochondria treated with and without the protonophoric uncoupler carbonyl cyanide-p-trifluoromethoxyphenylhydrazone (FCCP) in the presence of 50 µM CaCl₂ (Fig. 7B). While the addition of a 50 µM CaCl₂ generated granules, uncoupled mitochondria did not contain calcium phosphate granules in their matrices. The data from histograms were obtained from only mitochondria containing calcium phosphate granules. Arrows point to calcium phosphate granules.

Absence of granules in mitochondria under extreme high calcium conditions. While most of the isolated mitochondria contained calcium phosphate granules following the calcium addition, a few did not. There are two possible explanations for this phenomenon; either this group of mitochondria is (1) de-energized preventing calcium uptake or (2) they lack mitochondrial calcium uniporters (MCU). Based on the following statistical arguments, the latter is a more likely explanation. Assuming there are 40 MCUs per mitochondrion with an estimated standard deviation of 20, the probability of randomly selecting a mitochondrion without an MCU channel is 2.3%. This corresponds to 11 mitochondria in our total set of 502 images. In agreement with this estimation, our data show that 17 mitochondria do not possess granules after either calcium bolus was given which corresponds to 3.4% of the number of mitochondria imaged. This percentage is independent of treatment with 3.5% of control mitochondria and 3.6% of CsA treated mitochondria without any calcium phosphate granules. These results also match the respirometry data given in Fig. 2 whereby even after a large bolus of CaCl₂, some mitochondria are bioenergetically competent and synthesize ATP after the ADP bolus. Assuming that the boluses of calcium were enough to elicit MPT in the mitochondria with an MCU channel, the measured ADP-stimulated respiratory rate increase must be due to mitochondrial lacking an MCU. In line with this observation, the maximum ADP-stimulated respiratory rate for each calcium treatment relative to the maximum rate without calcium (as shown in Fig. 1), is 10.9% ± 5.2 and 4.3% ± 1.1 for the 75 µM and 100 µM CaCl₂ bolus, respectively. These values are strikingly close to the value estimated from the imaging data.

Discussion

The current leading hypothesis of calcium-induced mitochondrial dysfunction involves the peptidyl-prolyl cis–trans isomerase, cyclophilin D (CypD), interacting with as yet to be identified inner membrane proteins to form the permeability transition pore. When open, the pore results in sustained membrane depolarization, large amplitude swelling, calcium release, and loss of mitochondrial respiratory control. CsA can bind to CypD and sequester it so that its interaction with its target is prevented. However, CsA is not fully protective. It is believed that it only increases the calcium threshold required to open the pore. This idea is based on studies that show CsA increases the calcium retention capacity by nearly 3-fold. While an attractive hypothesis, this model has problems that are easier to explain using a different mechanism. As an alternative, we propose a novel...
mechanism of action whereby CsA enables robust calcium accumulation in the context of promoting calcium uptake and calcium phosphate granule formation. This mechanism involves the interaction between putative CsA-regulated proteins, and cristae structural proteins to preserve the inner membrane intactness. While the calcium phosphate granules may induce changes in morphology by mechanically disrupting membranes, it is plausible that free calcium interacts with proteins regulating inner membrane and cristae maintenance (namely the optic atrophic factor 1 and the mitochondrial contact site and cristae organizing system; known as OPA1 and MICOS) or additional regulators of this system. For instance, the stress-sensing overlapping activity with m-AAA protease 1 (OMA1) is a zinc metallopeptidase found in the inner mitochondrial membrane regulating mitochondrial dynamics through OPA1 processing59–61. OMA1 is activated under stress conditions including membrane potential dissipation, decreased ATP levels, and oxidative stress, among other insults61. Upon activation, OMA1 mediates the proteolytic processing of OPA1 inducing cristae remodeling and cyt. c release59,62–65. This mechanism can explain the morphological and functional changes included by calcium overload that we observed in our cryo-EM images and bioenergetics data.

Indeed, we demonstrated that calcium overload impairs mitochondrial ATP production at greater calcium loads and depleting mitochondria of calcium did not fully restore function—indicating an irreversible component. These data revealed an underappreciated energetic consequence of calcium overload on the mitochondrial function that supports a direct role of the mitochondrial calcium buffering system. In cardiac tissue, the steady-state cycling of calcium across plasma membranes maintains cytosolic calcium levels at ~ 100 nM during diastole; however, the peak calcium concentration in the cytosol during systole can rise to the low micromolar range9,33,66. Whether the mitochondria can respond to these transient changes to meet metabolic demand is a subject of debate (reviewed in9) that revolves around the mitochondrial calcium uniporter (MCU) being unable to approach maximum flux rates in the transient rise of cytosolic calcium due to its low affinity for calcium67. Alternative hypotheses regarding calcium microdomains have been proposed in an attempt to argue in favor of significant mitochondrial calcium uptake during systole68–72; however, direct imaging studies do not support

Figure 5. CsA disrupts OMM morphology, causes IMM release, and enhances the number and size of granules. Representative images before the addition of a 75 µM calcium bolus (t0) in the presence of 1 µM CsA. Mitochondria were energized with 5 mM sodium pyruvate and 1 mM L-malate. (A) CsA induced morphological changes to mitochondria that can be grouped into 4 classes as described in Supplementary Table 2. (B) The mitochondrial size, calcium phosphate granules size and number per mitochondrion were quantified for each time-point (t0–t3) before and after the addition of a 75 µM calcium bolus in the presence or absence of CsA. (B) There are no differences in the mitochondrial size of control to CsA-treated mitochondria before the addition of calcium. (C) After the calcium addition, the number and size of the granules increased in CsA-treated mitochondria were much larger than in the control mitochondria. Scale bars are 250 nm. n represents the number of images analyzed by the time point for control and CsA treated conditions. Gray zones represent the overlapping points between histograms.
A recent study by Wescott et al. found that physiological cytosolic calcium transients cause a gradual, step-wise increase in matrix calcium concentration per beat rather than large transient peaks. They also showed that at high pacing rates, the matrix calcium concentration did not change any further. Further studies are required to determine whether these results are due to equal influx and efflux of calcium per cycle or due to calcium buffering. At this point, it is possible that calcium buffering in the form of calcium phosphate granule formation becomes relevant. In a separate study, calcium phosphate granules were observed in the matrix near cristae junctions in a variety of different eukaryotic cells under physiological conditions. Given the relevance of calcium in bioenergetics, the presence of these calcium deposits may exert some degree of control over mitochondrial signaling and metabolism.

In Fig. 8, we present a model that accounts for various characteristics of membrane fragmentation before the MPT onset. This model integrates findings from our cryo-EM analysis with mitochondrial function and recapitulates the effects of calcium on the mitochondrial structure. Based on our findings, we believe that changes in...
mitochondrial ultrastructure can explain the loss of function in calcium overload as well as the protective effects of CsA. Our results suggest that mitochondrial outer membrane rupture and inner membrane fragmentation are caused by calcium overload whereas the formation of granules is a consequence of calcium uptake and accumulation. In the present study, the detrimental effects of calcium overload on mitochondrial function are mitigated when CsA is present. Regardless of the calcium bolus, the number and size of granules in CsA-treated
mitochondria increased, suggesting that CsA increases the mitochondrial calcium buffering capacity, thus explaining why CsA allows robust calcium uptake and increases the threshold for permeability transition pore activation.

To interpret these results, we sought confirmation of our findings from work by others. A study by Pinton’s group studied the effect of calcium overload on mitochondria in HeLa cells. Exposing HeLa cells to the ionophore ionomycin resulted in mitochondrial network fragmentation. However, in the presence of CsA, the mitochondrial network condensed and maintained its integrity after ionomycin treatment. Another study looking at mitochondrial swelling using light transmittance in a single mitochondrion showed that calcium induces mitochondrial swelling in a concentration-dependent fashion. CsA decreased this effect in a calcium-dependent manner, which led the authors to conclude that either CsA induces mitochondria shrinkage or calcium accumulation induces light scattering. We show that CsA increases the absorbance in a calcium-dependent manner and induces changes in mitochondria ultrastructure including condensed inner membranes and loss of outer membrane. Therefore, our results are consistent with these studies but quantitatively describe the ultrastructural changes associated with calcium overload and how these changes are linked to mitochondrial function.

A major challenge in this study is the lack of cristae structural definition in our set of images. As dynamic structures, cristae are the functional units of mitochondria that lock cyt. c in the cristae lumen and provide sufficient membrane surface area to sustain oxidative phosphorylation at high rates. Under certain conditions when the cristae junctional width is enlarged, cyt. c escapes the lumen and causes loss of mitochondria function and cell death. While the expected outcomes during calcium overload were addressed, intricate details of the cristae structure including junction width, length, density, and shape must be incorporated to better understand the implications of cristae remodeling as key mediators in mitochondrial function. While energy-dispersive X-ray spectroscopy (EDX) was not included in our studies, others have determined these granules are majorly composed of calcium and phosphorous. Earlier studies looking at calcium phosphate granule composition relied on staining, fixing, or dehydrating samples, introducing artifacts which makes them less reliable. More recently, changes in mitochondrial structure were analyzed using high-pressure techniques and freeze-substitution to minimize sample structural distortion resulting from fixation or dehydration. However, details such as granule space distribution and structure are not as well defined with this method relative to the latest advanced cryo-EM techniques. Hence, visualization of the mitochondria in 3D by cryo-electron tomography (cryo-ET) would be an avenue for future studies to address. Nonetheless, our finding that CsA preserves the inner membrane integrity suggests that cristae remodeling and cyt. c release from the cristae lumen is likely avoided. This poses a new approach by which therapies targeting cristae remodeling can be identified to prevent pathological mitochondrial dysfunction leading to tissue injury.

Materials and methods

Ethical approval. This work conformed to the National Institutes of Health’s Guide for the Care and Use of Laboratory Animals and was approved by Michigan State University’s Institutional Animal Care and Use Committee.

Mitochondria isolation and protein quantification. Cardiac mitochondria were isolated from guinea pig hearts using differential centrifugation as described in Wollenman et al. Briefly, Hartley albino guinea pigs weighing 350–450 g (4–6 weeks) were injected with heparin (500 units/mL) into the intraperitoneal cavity to prevent blood clotting during the cardiac mitochondrial isolation. Before heart removal, the animals were deeply anesthetized with 4–5% isoflurane. Prior to decapitation by guillotine, a noxious stimulus (paw pinch and eyelid reflex) confirmed the animals were fully sedated. After decapitation, a thoracotomy was performed. The heart was then perfused with cold cardioplegia solution and homogenized as described previously. Mitochondrial protein content was quantified using the BIO-RAV Bovine Serum Albumin (BSA) Standard Set Kit and the BCA assay. The mitochondrial suspension was diluted to a working concentration of 40 mg/mL and kept on ice for the duration of the experiment (4–8 h). Substrate stock solutions were neutralized to pH 7.0.

Mitochondrial quality control. The mitochondrial quality was determined using an Oxygraph 2 k (Oroboros Instruments Corp., Innsbruck, Austria) under constant stirring. The O2k chambers were loaded with 2 mL respiratory buffer containing 130 mM KCl, 5 mM K2HPO4, 20 mM MOPS, and 1 mM MgCl2, 1 mM EGTA, 0.1% (w/v) BSA at a pH of 7.1 and 37 °C. All subsequent experiments were done using this buffer and temperature. At 0 min, 5 mM sodium pyruvate and 1 mM L-malate were added followed by 0.1 mg/mL mitochondria. Here we defined leak state as the rate of oxygen consumption by mitochondria only in the presence of substrates. At 5 min a bolus of ADP (500 µM) was added to induce maximal ADP-stimulated respiration. Quality was assessed by computing the respiratory control ratio (maximal ADP-stimulated rate divided by the leak rate). Only mitochondria with an RCR value greater than or equal to 16 were used in the experiments.

Calcium contamination and buffer calcium measurements. The amount of contaminating calcium present in the respiratory buffer was 4.0 µM ± 0.43 µM which comes from reagent impurities. This was measured using a perfectION calcium selective electrode (Mettler Toledo, Columbus, OH). Results were further confirmed using 1 µM calcium fluorescent indicator calcium green 5N (503 nm excitation and 531 nm emission) using an Olis DM245 spectrofluorimeter (Olis, Inc., Bogart, GA, USA).

Calcium effects on respiration and oxidative phosphorylation. Calcium effects on mitochondrial leak and ADP-stimulated respiration were determined by quantifying changes in leak and ADP-stimulated respiration rates after a calcium challenge in the presence or absence of cyclosporin A (CsA). At 0 min, 5 mM

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Calcium effects on the outer membrane integrity. The effect of calcium on the outer membrane integrity was determined by quantifying changes in ADP-stimulated respiration rates after a calcium challenge by the addition of cytochrome c. At 0 min, 5 mM sodium pyruvate, 1 mM L-malate, and 0.1 mg/mL mitochondria were injected into each 2 mL chamber containing respiratory buffer. At 5 min, a calcium bolus of 50 µM CaCl₂ or ddH₂O was injected. At 5 min, 1 mM EGTA was added onto each chamber. At 10 min, 500 µM ADP was added to induce maximal ADP-stimulated respiration. The respiratory buffer contained no EGTA for these studies.

Mitochondrial swelling assay. Mitochondrial swelling was quantified by measuring absorbance at 540 nm using an Olis DM245 spectrophotometer with a dual-beam absorbance module. At 0 min, 5 mM pyruvate and 1 mM L-malate was added to a polystyrene cuvette with respiration buffer containing ±1 µM CsA followed by the addition of 0.1 mg/mL mitochondria. At 5 min, a 75 or 100 µM CaCl₂ bolus was added and the absorbance was recorded for a total of 15 min. The minimum absorbance signal was determined by adding the uncoupler FCCP (1 µM) and the channel forming peptide Alamethicin (10 µg/mg). To normalize the raw traces, we used the minimum absorbance value followed by the absorbance just before the addition of a calcium bolus.

Cryo-EM sample vitrification and imaging. Isolated mitochondria were suspended at a concentration of 0.1 mg/mL in 2 mL respiration buffer with 5 mM sodium pyruvate and 1 mM L-malate. At the collection times indicated, 5 µL samples were pipetted from the mitochondrial suspension and deposited on Quantifoil R2/2 Holey Carbon grids that had been plasma-cleaned for 20 s using a Fischione Instruments model 1020 plasma cleaner. Grids were blotted to thin the water layer, and subsequently plunged into liquid ethane at room temperature using a manual plunge-freezing device (Michigan State University Physics Machine Shop). Grids were then transferred and stored in liquid nitrogen until imaging. Data for the 75 µM CaCl₂ experiments were collected in the cryo-EM facility at the University of Pittsburgh School of Medicine using a FEI Polara G2 cryo-electron microscope with a field emission gun operating at 300 kV at nominal magnification of 9400× with a post-column magnification of 1.4× to obtain a ~ 12–10 Å/pixel resolution. Images were recorded on a FEI Falcon 3 direct electron-detecting camera at an electron dose of 0.2 e⁻/Å². Data for the 100 µM CaCl₂ experiments were collected in the cryo-EM facility at the University of Pittsburgh School of Medicine using a FEI TF20 cryo-electron microscope with a field emission gun operating at 200 kV at nominal magnification of 3000× on a TVIPS XF416 CMOS camera with a post-column magnification of 1.4× to obtain a ~ 10–8 Å/pixel resolution. Data for the ±1 µM FCCP condition at 50 µM CaCl₂ were collected in the cryo-EM facility on Michigan State University using a Talos Arctica outfitted with a Falcon 3 direct electron-detecting camera. The field emission gun was operating at 200 kV using a magnification of 22,000× at an electron dose of 2.03 e⁻/Å² to obtain a 4.69 Å/pixel resolution. Images were recorded on a FEI Polara G2 cryo-electron microscope with a field emission gun operating at 200 kV at nominal magnification of 9400× with a post-column magnification of 1.4× to obtain a ~12–10 Å/pixel resolution. Data for the ±1 µM FCCP condition at 50 µM CaCl₂ were collected in the cryo-EM facility on Michigan State University using a Talos Arctica outfitted with a Falcon 3 direct electron-detecting camera. The field emission gun was operating at 200 kV using a magnification of 22,000× at an electron dose of 2.03 e⁻/Å² to obtain a 4.69 Å/pixel resolution. Grids were blotted for 5 s before vitrification in liquid ethane using a Vitrobot Mark IV System from Thermo Fisher Scientific. At these magnifications, the electron dose (e⁻/Å²) is low enough to avoid significant sample destruction.

Calcium phosphate granules, posner’s clusters, and mitochondrial structure quantification. The program EMAN2 was used to quantify the total number of granules for each mitochondrion under each condition from TEM images. A total of 1345 individual mitochondrial images were acquired in the presence and absence of CsA for two calcium treatments. For the 75 µM CaCl₂ treatment, there were 235 images of control mitochondria and 645 images of CsA-treated mitochondria. For the 100 µM  CaCl₂ treatment, there were 235 images of control mitochondria and 645 images of CsA-treated mitochondria. Mitochondrial and phosphate granule diameters were computed from three averages of two diagonal and one horizontal diameter measurement. Pixel resolution was converted to nanometers based on the magnification level. The fractional area that the calcium phosphate granules occupy per mitochondrion was calculated by multiplying the number of granules within a mitochondrion times the sum of all the granule areas divided by the area of the mitochondrion (Ngranules * Agranules / A mitochondria ). The calcium phosphate nanoclusters (n = 227) were determined by measuring the electron-dense regions located within the granules using ImageJ (NIH, Bethesda, MD, USA).
Statistics. The Shapiro–Wilks test was used to confirm data normality. All data were analyzed and plotted using either MATLAB 2019a or MATLAB 2019b (Mathworks, Inc., Natick, MA, USA). The data in Figs. 1, 2 and 3 (n = 3–4) and stats presented for the calcium phosphate nanoclusters are presented as mean ± standard deviation. Mitochondrial images with calcium phosphate granules were only included for the histogram analysis (n value in the figures). An unpaired Student’s t test was used to compare the CaA treatment with the control group. An n-way ANOVA was run to determine significant effects between treatments at various calcium loads and different time-points. A p value < 0.05 was assumed to be statistically significant.

Reagents. All reagents were purchased from Sigma-Aldrich unless otherwise stated. Calcium Green-5 N hexapotassium salt was purchased from Thermo Fisher Scientific.

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Author contributions

J.O.S.R. and J.N.B. conceived the approach, J.O.S.R. prepared the specimens, J.O.S.R. performed experiments, J.R.S., I.F.C., K.N.P. performed cryo-EM tomography, J.O.S.R., E.V.P., J.R.S., K.N.P., and J.N.B. developed methods, J.O.S.R. and J.N.B. analyzed the data, J.O.S.R. and J.N.B. wrote the manuscript. J.O.S.R., J.R.S., I.F.C., E.V.P., K.N.P., and J.N.B. edited and reviewed the manuscript. J.O.S.R. drew the schematic representation image.

Competing interests

The authors declare no competing interests.

Additional information

Supplementary Information

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