A Novel MitoTimer Reporter Gene for Mitochondrial Content, Structure, Stress, and Damage in Vivo*

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Background: Mitochondrial health is difficult to assess in vivo.

Results: We have generated a reporter gene, MitoTimer, which targets mitochondria, and fluoresces green and shifts to red when oxidized, for assessment of mitochondrial content, structure, stress, and damage under physiological and pathological conditions.

Conclusion: MitoTimer is useful for assessment of mitochondrial health in vivo.

Significance: MitoTimer could advance mitochondrial research in multiple disciplines.

Mitochondrial dysfunction plays important roles in many diseases, but there is no satisfactory method to assess mitochondrial health in vivo. Here, we engineered a MitoTimer reporter gene from the existing Timer reporter gene. MitoTimer encodes a mitochondria-targeted green fluorescent protein when newly synthesized, which shifts irreversibly to red fluorescence when oxidized. Confocal microscopy confirmed targeting of the MitoTimer protein to mitochondria in cultured cells, Caenorhabditis elegans touch receptor neurons, Drosophila melanogaster heart and indirect flight muscle, and mouse skeletal muscle. A ratiometric algorithm revealed that conditions that cause mitochondrial stress led to a significant shift toward red fluorescence as well as accumulation of pure red fluorescent puncta of damaged mitochondria targeted for mitophagy. Long term voluntary exercise resulted in a significant fluorescence shift toward green, in mice and D. melanogaster, as well as significantly improved structure and increased content in mouse FDB muscle. In contrast, high-fat feeding in mice resulted in a significant shift toward red fluorescence and accumulation of pure red puncta in skeletal muscle, which were completely ameliorated by voluntary wheel running. Hence, MitoTimer allows for robust analysis of multiple parameters of mitochondrial health under both physiological and pathological conditions and will be highly useful for future research of mitochondrial health in multiple disciplines in vivo.

Mitochondrial dysfunction, e.g. reduced respiratory function and increased production of reactive oxygen species (ROS), 3

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3 The abbreviations used are: ROS, reactive oxygen species; Cox4, cytochrome c oxidase subunit IV; FDB, flexor digitorum bevis; HFD, high-fat diet; OCR, oxygen consumption rate; DsRed, Discosoma red fluorescent protein.
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oxidase subunit VIII gene to the N terminus of the coding region of Timer, under control of the constitutive CMV promoter. Timer encodes a DsRed mutant (DsRed1-E5) that fluoresces like green fluorescence protein when newly synthesized, and shifts the fluorescent spectrum irreversibly to red (18) following a form of oxidation (dehydrogenization) of the Tyr-67 residue (19, 20). This physical property in a biological environment makes it suitable for reporting the age of proteins and cells (18), and thus was named “Timer.” Recently, Ferree et al. (21) and Hernandez et al. (22) have used inducible MitoTimer reporter to show its usefulness in cell culture to report changes in mitochondrial turnover and transport. Because the fluorescence spectrum shifts upon oxidation, we predicted that MitoTimer driven by a constitutively active promoter would report cumulative redox history of the labeled mitochondria. We also provide additional value by using MitoTimer in vivo in transgenic Caenorhabditis elegans and Drosophila melanogaster as well as following somatic gene transfer in adult mouse skeletal muscle.

Here we have validated the reporter properties and confirmed that MitoTimer could be used to report mitochondrial content, structure, stress, and damage in vivo under physiological and pathological conditions. We also used MitoTimer to address a biological question of whether exercise training was protective against HFD-induced mitochondrial damage in mouse skeletal muscle. Our findings provide morphological evidence demonstrating HFD-induced mitochondrial defects in vivo using two independent methods (transmission electron microscopy and MitoTimer fluorescence), and the benefits of exercise training for mitochondrial health. This reporter gene can be potentially used in multiple disease models, and its applications, beyond this study, will likely lead to advancement in multiple disciplines of mitochondrial health research.

EXPERIMENTAL PROCEDURES

Plasmid DNA Construct—We constructed pMitoTimer by inserting the BamHI-NotI fragments of pTimer-1 (Clontech) into pDsRed2-Mito (Clontech) vector digested with BamHI and NotI using T4 ligase (New England Biolabs) following DNA isolation by using the Qiaxil II gel extraction kit (Qiagen). The mitochondrial targeting sequence of the human cytochrome c oxidase subunit VIII gene from pDsRed2-Mito was then fused to the mutant DsRed1-E5 on the N terminus. Bacterial colonies were screened using a Qiaprep Spin Miniprep Kit (Qiagen) with the mutant DsRed1-E5 on the N terminus. Bacterial colonies were screened using a Qiaprep Spin Miniprep Kit (Qiagen) with the mutant DsRed1-E5 on the N terminus.

Drug Treatment of Transfected Cells—pMitoTimer-transfected C2C12 myoblasts were treated with 10 μM rotenone (electron transport chain complex I inhibitor; Sigma), 100 nM antimycin A (complex III inhibitor; Sigma), or 100 μM paraquat (mitochondrial ROS producer primarily through complex I) in normal growth medium for 6 h. In experiments using diethylthiainetamine-nitric oxide, cells were pre-treated with 200 μM diethylthiainetamine-NO for 1 h prior to co-treatment with either antimycin A or dimethyl sulfoxide for 6 h. Cells were fixed and mounted immediately following drug treatment for confocal microscopy analysis. To assess the effect of the drug treatment to mitochondrial function, C2C12 myoblasts were treated with the same drugs for 6 h prior to measurement of the oxygen consumption rate (OCR) using a Seahorse XF-24 Flux Analyzer (Seahorse Biosciences, Billerica, MA). Briefly, prior to the assay, the media was changed to 5 mM HEPES-buffered Seahorse medium (Invitrogen) (pH 7.4) and equilibrated for 30 min at 37°C. Oligomycin (Oligo, 1 μM), BAM15 (23) (2 μM), and rotenone and antimycin A (1 and 10 μM, respectively) were injected sequentially during the assay, whereas OCR was measured at 3-min intervals. OCR was normalized to α-Tubulin protein abundance in harvested cells following completion of the assay. Basal mitochondrial OCR, ATP-linked OCR, and reserve capacity were calculated.

D. melanogaster Strains and Drug Treatment—Wild type (w1118) and Mef2-Gal4 fly lines were obtained from the Bloomington Drosophila Stock Center (Indiana University, IN). Flies were housed in a humidified, temperature-controlled incubator at 25°C on a 12:12-h light-dark cycle. To generate the MitoTimer transgenic fly line, the coding region of MitoTimer was subcloned into pUAS vector. A UAS-MitoTimer transgenic fly line was established commercially (Genetic Services, Inc.). UAS-MitoTimer flies were crossed with Mef2-Gal4 to obtain Mef2-Gal4>UAS-MitoTimer flies. Adult female flies were collected under light CO2 anesthesia and transferred to fresh medium every 2–3 days (20 per vial). To induce oxidative stress in Drosophila, we prepared the drugs as following. Antimycin A was dissolved in 100% ethanol at 1 mM and then added to 5% sucrose at 0.1 mM. Paraquat was dissolved in water at 100 mM and added to 5% sucrose at 10 mM. These two drugs were administered to adult flies (20 days old) 24 h after 6 h of fasting as described by Lee et al. (24). Rotenone was first dissolved in dimethyl sulfoxide at 4 mM, added to freshly made fly food at
60 °C at 0.4 mm, and used to feed the flies for 4 days after 6 h of fasting (25).

**Exercise Training and Negative Geotaxis Assays in Flies—**Exercise training of UAS-MitoTimer-Mef2-Gal4 flies were performed according to a previously reported method (26). Negative geotaxis was assessed in Rapid Negative Geotaxis (RING) assays in groups of 100 flies as described (27). Flies were tested 5 times per week for 5 weeks to assess the decline in negative geotaxis speed with age. Climbing endurance was measured using the fatigue assay as previously described (28).

**Imaging Acquisition of Adult Fly Heart—**Adult fly hearts were dissected according to an established method (29) and fixed in 4% paraformaldehyde for 20 min. Confocal Z-stack images were obtained within abdominal segment A1 with a set of fixed acquisition parameters. One set of images per heart, both green (excitation/emission 488/518 nm) and red (excitation/emission 543/572 nm) channels, were selected for analysis that was 6–8 nm from the top section.

**C. elegans Experiments—**

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  - **mec-3** MitoTimer was constructed by digesting pPD57.56 with BamHI and Nhel and inserting a PCR fragment that contained the DsRed-2 mitochondrial portion of pTTimer-1 also cut with BamHI and Nhel. The sequence-confirmed plasmid was injected into wild type C. elegans to generate strain ZB4034. MitoTimer was expressed in 10 neurons including the touch receptor neurons. We imaged using a Zeiss Axiovert 200M upright microscope with deconvolution.

**Mouse Model—**

- **Male C57BL/6J mice (7–8 weeks old)** were obtained commercially (Jackson Laboratory) and housed in temperature-controlled (21 °C) quarters with a 12:12-h light-dark cycle and water and chow (Purina) provided *ad libitum.* All experimental protocols were approved by the University of Virginia, Institutional Animal Care and Use Committee. MitoTimer and pLamp1-YFP transfection by somatic gene transfer in the adult mouse FDB muscle was performed as previously published (30). Briefly, the mice were anesthetized (isoflurane) and the base of each hind limb foot, where the flexor digitorum brevis (FDB) muscle is located, was injected with 10 μl of hyaluronidase (0.36 mg/ml; Sigma) subcutaneously. One h later, the mice were anesthetized a second time and the FDB injected with 20 μg of pMitoTimer in saline. In the case of co-transfection, pMitoTimer and pLamp1-YFP were mixed in a 1:1 ratio, and 20 μg of plasmid DNA was injected. Following another 10 min, a pair of gold-plated acupuncture needles were inserted under the skin at the heel of the foot and the base of the toes parallel to each other and perpendicular to the long axis of the foot. Ten pulses with 20 ms duration at 1 Hz and 75 V/cm were applied to each foot.

**Diet and Exercise Interventions—**Following somatic gene transfer, the mice were allowed to recover for 10 days and were then allocated to one of the following groups (n = 6; sedentary normal chow (NC-Sed), sedentary with 60% high-fat diet (HFD-Sed); Research Diets Inc.), voluntary wheel running with NC (NC-Ex), or voluntary wheel running with 60% HFD (HFD-Ex) for 3 weeks. In a separate experiment, the mice (n = 6) were allocated to either sedentary (Sed) or voluntary wheel running (Ex) for 6 weeks. All mice were housed individually in cages with running wheels, and the running wheels were locked for the mice allocated to the sedentary groups. At the end of the studies, all mice were humanely sacrificed under anesthesia, and FDB muscles were carefully excised and immediately fixed in 4% paraformaldehyde for 20 min.

**Immunofluorescence—**For detection of Cox4, pMitoTimer-transfected FDB muscles were processed for paraffin embedding as described (31). Briefly, harvested muscles were placed in 10% formalin at room temperature overnight. Samples were transferred to 70% ethanol before being paraffin-embedded and sectioned by a microtome. The sections were incubated in a dry 37 °C incubator overnight before being deparaffinized and rehydrated in a series of xylene and ethanol washes. Muscle sections (5 μm) were then washed for 10 min in Tris-buffered saline (TBS) with 0.1% Tween 20, followed by incubating with blocking buffer (10% normal goat serum in TBS) for 2 h at room temperature. Sections were incubated with mouse anti-Cox4 antibody (Invitrogen A21348, 1:100) in TBS overnight at 4 °C, washed for 10 min in 0.1% Tween 20 TBS, and incubated for 1 h at room temperature with goat anti-mouse Cy5 antibody (Jackson 115-175-146, 1:50) followed by mounting with Pro-Long Gold Antifade Reagent (Invitrogen) for confocal microscopy.

For detections of LC3, pMitoTimer-transfected FDB muscles were immediately fixed in 4% paraformaldehyde for 20 min and incubated overnight in PBS with 0.2% saponin at 4 °C. Muscles were then incubated in blocking solution (PBS with 0.5% BSA, 0.2% saponin, 3% normal goat serum, and 3% normal rat serum) for 1 h at room temperature, followed by overnight incubation with rabbit anti-LC3 (Novus Biologics, NB100-2220; 1:300) in blocking solution at 4 °C. Following 5 × 20-min washes in PBS with 0.2% saponin, muscles were incubated with goat anti-rabbit Cy5 antibody (Jackson 111-175-144, 1:200) for 2 h at room temperature and washes were repeated. Muscles were then whole mounted and imaged by confocal microscopy as described.

**Confocal Microscopy—**Immediately following fixation (described above) the muscles were whole mounted on gelatin-coated glass slides with 50% glycerol in PBS with a coverslip. MitoTimer images were acquired at ×100 magnification under a confocal microscope (Olympus Fluoview FV1000) using the green (excitation/emission 488/518 nm) and red (excitation/emission 543/572 nm) channels with identical predetermined acquisition parameters for all samples to ensure no saturation of the signals and similar intensity of the green and red channels in control samples. Lamp1-YFP images were acquired using the yellow channel (excitation/emission 543/572 nm), whereas LC3 and Cox4 immunofluorescence (Cy5) was detected in the far-red spectrum (excitation/emission 635/672 nm).

**Ratiometric Analysis—**MitoTimer signals were analyzed using our custom-designed Matlab-based algorithm. Briefly, positive pixels were thresholded at a value of 1.5 times the mean gray value of the above background pixels for each channel. This made the threshold robust against differences in fiber size between images. Saturated pixels (gray level = 255) were removed from analysis. In addition, the pixels with a red to green ratio of >2.5 were removed for the ratiometric analysis. The ratio of red to green was calculated as a mean value of red to green ratio for all the remaining positive pixels in the image (Fig. 2c). The number of red dots in the images was
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calculated by counting areas with clusters of ≥5 pixels with high fluorescence signal (≥175) and a red to green ratio of >2.5. The red dot counting was done on images including saturated pixels (Fig. 2d).

Transmission Electron Microscopy—Transmission electron microscopy was performed as previously described (32).

Statistical Analysis—The data were analyzed using a t test or one-way analysis of variance with Student’s Newman-Kuels post hoc where appropriate. Data are presented as mean ± S.E. and statistical significance was set at p < 0.05.

RESULTS

MitoTimer Protein Targets to Mitochondria and Displays Heterogeneous Green and Red Fluorescence—We engineered the MitoTimer reporter gene to target the oxidation-sensitive Timer protein to mitochondria. To confirm the subcellular location, we transiently transfected C2C12 myoblasts with plasmid pMitoTimer (Fig. 1a) and compared the expression pattern with those of pGFP3, pDsRed-Mito, and pTimer-1 (Fig. 1b). Cells transfected with pMitoTimer showed typical mitochondrial network structures (Fig. 1a) identical to those transfected with pDsRed-Mito (Fig. 1b). Cells transfected with pMitoTimer also showed heterogeneous green and red fluorescence (Fig. 1a, panel labeled Merge), similar to that of pTimer-1 (Fig. 1b). Mitochondrial targeting of the protein was further confirmed by Western blot of the mitochondrial and cytosolic fractions of C2C12 myoblasts transiently transfected with pMitoTimer compared with cells transfected with an empty vector, pCI-neo (Fig. 1c). MitoTimer protein was highly enriched in the mitochondrial fraction as probed by anti-DsRed antibodies (Fig. 1c). To validate these findings in vivo, we generated transgenic C. elegans with touch neuron-specific expression under control of the neuron-specific promoter mec-4 and observed green and red mitochondria in the soma and distributed through the sensory neuron process (Fig. 1d), which is similar to mitochondrial targeting standard GFP as reported previously (33). We also generated UAS-MitoTimer transgenic flies and crossed with Mef2-Gal4 transgenic flies to generate Mef2-Gal4>UAS-MitoTimer flies with muscle expression (34). Confocal microscopy showed mitochondrial structure identical to that obtained in Mef2-Gal4>UAS-Mito-GFP flies not only in adult heart tube (Fig. 1, e and f), but also in other muscles, such as indirect flight muscle and larval body wall muscle (Fig. 1, g and h). There were variable green and red fluorescence in different regions of the heart tube. In addition, we detected pure red fluorescent puncta, which were often disconnected from, but in close proximity to, the mitochondrial reticulum (Fig. 1e).

To further validate the use of the MitoTimer reporter gene in vivo, we performed electric pulse-mediated gene transfer in adult mouse FDB muscle followed by whole mount confocal microscopy. FDB muscle fibers transfected with pMitoTimer showed a striking sarcomeric striation pattern at low magnification (Fig. 1i) identical to that in FDB muscle transfected with pDsRed-Mito (Fig. 1j). High magnification images showed labeling of inter myofibrillar mitochondria on each side of the Z-line as well as subsarcolemmal mitochondria with streaking patterns (Fig. 1i). These patterns have also been shown by the mitochondria-targeted reporter gene mt-cpYFP (35). Mitochondria targeting of the MitoTimer protein was further confirmed by co-localization of fluorescence with immunofluorescence staining of Cox4 (Fig. 1k). Similar to the fly heart tube, we also detected pure red puncta in mouse skeletal muscle transfected with pMitoTimer and confirmed that they were often positive for Cox4, suggesting that they are of mitochondrial origin and are likely degenerative mitochondria (Fig. 1k). In addition we found that some red puncta and yellow fluorescent subsarcolemmal mitochondria co-localized with LC3 immunofluorescence, a marker for autophagosome (Fig. 1l), as well as a YFP signal from pLamp1-YFP transfection (Fig. 1m), a marker of lysosome. The findings strongly suggest that red puncta were mitochondria targeted for removal from the cell via autophagy.

Quantification of Fluorescence Spectrum Shift and Pure Red Puncta—We developed a Matlab-based automated image analysis platform for quantification of fluorescence of each channel that was acquired with the same parameters of laser intensity and scanning speed for all samples under the confocal microscope. In the case of transfected FDB muscles, this Matlab-based computer algorithm segmented the muscle fiber and identified MitoTimer positive pixels within the fiber using a threshold value of 1.5 times the mean gray value above background pixels for each channel (Fig. 2, a and b). This processing controlled for differences in fiber number and size between images. The shift of the fluorescence spectrum was calculated by taking the ratio of red to green fluorescence intensity of each of the positive pixels (Fig. 2c). The algorithm also quantified the number of pure red puncta (Fig. 2d), which were defined as more than 5 pixels in contact with each other with a high fluorescence signal (>175) and a red to green ratio of >2.5. With this automated image analysis algorithm we were able to quantitatively characterize the fluorescent properties of MitoTimer signals in various model systems quickly and reproducibly. The algorithm can process 100 pairs of images in ~30 s.

MitoTimer Protein Changes Fluorescence from Green to Red under Conditions of Mitochondrial Stress—To determine whether mitochondrial stress promotes a shift of the fluorescent spectrum from green to red, we treated pMitoTimer-transfected C2C12 myoblasts with rotenone (10 μM), antimycin A (100 nM), or paraquat (100 μM) for 6 h. These drugs are known to induce mitochondrial ROS production from complex I or complex III of the electron transport chain. A ratiometric algorithm showed that treatment with any of these drugs resulted in a significant (p < 0.05) shift of mitochondrial fluorescence toward red (Fig. 3, a and d). The drug treatments significantly reduced mitochondrial respiratory function as measured by a mitochondrial stress test using a Seahorse oxygen flux analyzer. Specifically, these three drug treatments all resulted in reduced basal mitochondrial respiration and reserve capacity along with a trend of reduced ATP-linked respiration (Fig. 3, b and c). To further confirm that the change in MitoTimer fluorescence in cells is due to ROS production we showed that pre-treatment with diethylenetriamine-NO (200 μM) was sufficient to prevent the antimycin A-induced fluorescence shift (Fig. 3, e and f). When adult Mef2-Gal4>UAS-MitoTimer transgenic flies (20 days of age) were treated with these drugs supplemented in feed, there was also a significant shift toward
red fluorescence in the heart tube concurrent with increases of pure red puncta (Fig. 3, g–i). Similar changes were observed in aged (70 days old) compared with young (7 days old) flies (Fig. 3, j–l). These findings support the use of the MitoTimer reporter gene to report mitochondrial stress and damage in vitro and in vivo.

MitoTimer Reports the Beneficial Impact of Exercise on Mitochondria in Vivo—It is known that endurance exercise training increases mitochondrial quantity and improves mitochondrial quality in skeletal muscle (36). To determine whether the MitoTimer reporter gene could be used for this purpose, we performed somatic gene transfer of pMitoTimer in mouse FDB
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FIGURE 2. Computational analysis of MitoTimer fluorescence. A Matlab-based computer algorithm was designed for identification of MitoTimer positive pixels; quantification of MitoTimer fluorescence spectrum shift between green and red, and accumulation of pure red puncta; a, a representative confocal image (merged image of green and red) of a mouse FDB muscle fiber; b, the processed confocal image showing background (black), fiber (gray), and MitoTimer (white) pixels; c, corresponding heat map of the red:green ratio of MitoTimer positive pixels; d, pure red puncta identified by the MitoTimer image analysis algorithm.

Muscles followed by voluntary wheel running for 6 weeks. Whole mount confocal microscopy showed that exercise training promoted a significant increase in mitochondrial content per fiber area (Fig. 4, a and b), as well as a shift of the fluorescence toward green (Fig. 4, a and c), indicative of reduced mitochondrial stress. There was no significant change in the number of pure red puncta (Fig. 4d). Under higher magnification, we also observed improved mitochondrial alignment and uniformity of mitochondrial pairs along the Z-line in FDB fibers from exercised mice compared with those of sedentary mice (Fig. 4a). We then employed exercise training in flies using a unique training device known as the power tower (26). Exercise training in male flies resulted in improved exercise capacity (Fig. 4e) associated with a significant shift of the MitoTimer fluorescence toward green (Fig. 4, f and g) with no significant changes in pure red puncta in the heart tube (Fig. 4, f and h).

MitoTimer Reports Mitochondrial Stress and Damage under Physiological and Pathological Conditions in Vivo—To determine the utility of the MitoTimer reporter gene in a mammalian model for mitochondrial stress and damage under pathological conditions, we subjected mice transfected with pMitoTimer in the FDB muscles to a 60% HFD for up to 10 weeks. Confocal microscopy of whole mounted fixed muscles showed a considerable shift in fluorescence toward red in mice on the HFD along with a substantial increase of pure red puncta compared with mice on normal chow diet (Fig. 5a). These changes appear to be time-dependent as shown by increasing severity in a time course at 1, 3, 6, and 10 weeks (Fig. 5a). Transmission electron microscopy confirmed that the HFD resulted in accumulation of damaged mitochondria with various degrees of enlargement and vacuolization in skeletal muscles (Fig. 5b). We then set to determine whether exercise training could prevent these changes and found that the shift of the fluorescence toward red with increased pure red puncta following 3 weeks of HFD were completely prevented by voluntary wheel running (Fig. 5, c–e).

Taken together, we have shown that the MitoTimer reporter gene encodes an oxidation-sensitive protein that targets to mitochondria and reports in vitro and in vivo mitochondrial health under physiological and pathological conditions.

DISCUSSION

Mitochondrial dysfunction is one of the most important features of many prevalent diseases (1–9). For example, reduced mitochondrial content and electron transport chain activity in skeletal muscles have been reported both in obese, diabetic patients (37, 38) and in lean, insulin-resistant offspring of type 2 diabetic patients (2, 39), suggesting its importance in the pathogenesis. Accumulation of damaged mitochondria, due to lipotoxicity, in cardiac myocytes has been reported in a rat genetic model of obesity (4). Mitochondrial DNA damage and dysfunction are associated with heart failure induced by myocardial infarction (40). Abnormal regulation of genes essential for mitochondrial function is sufficient to induce heart failure in animals (5). Finally, there is accumulating evidence of loss of the mitochondrial function in the central nervous system in neurodegenerative diseases (3, 6, 41). It is clear that mitochondrial damage and dysfunction is detrimental, if not causal, to the aforementioned disease conditions. More importantly, mitochondrial abnormalities promote excess production of reactive oxygen species in a vicious cycle leading to further mitochondrial damage (42). In light of the importance of mitochondria in health and disease, there is an urgent need to develop appropriate animal models to improve our understanding of mitochondrial pathology and develop interventions to improve mitochondrial function.

This study has taken critical steps to advance our ability to measure mitochondrial stress and damage through validation of MitoTimer under physiological and pathological conditions in vitro and in vivo. We believe that the MitoTimer reporter gene will be highly useful for future in vivo research because it allows for robust measurement of multiple mitochondrial parameters. The most significant changes that we observed with the MitoTimer reporter gene in whole animals were the overall change of fluorescence spectrum and the abundance of pure red puncta. These two parameters may reflect two different features of mitochondrial health. Within a given region of mitochondrial network, the MitoTimer fluorescence spectrum is likely to be determined by a balance between the import of the newly synthesized protein and its oxidation. Because MitoTimer protein in this study is expressed under control of constitutively active promoters, an overall increase of red:green ratio of the mitochondrial network under the conditions of mitochondrial stress are likely due to increased oxidation of the protein rather than altered transcription/translation. This could be caused by increased reactive oxygen species produc-
FIGURE 3. MitoTimer shifts fluorescence spectrum in response to mitochondrial stress in C2C12 myoblasts and Drosophila heart tube. Confocal microscopy analysis of MitoTimer reporter gene expression was performed for transiently transfected C2C12 myoblasts and transgenic Drosophila heart tube following treatment or feeding, respectively, with drugs that are known to cause mitochondrial stress. a, representative merged images of pMitoTimer-transfected C2C12 myoblasts (scale bar = 10 μm). b, OCR normalized to α-tubulin during a Seahorse mitochondrial stress test. Baseline OCR was measured twice prior to addition of oligomycin (A) to determine ATP-dependent OCR followed by addition of the mitochondrial uncoupler BAM15 (B) for measurement of maximal capacity and finally addition of antimycin A and rotenone (C) for non-mitochondrial OCR. c, quantification of basal mitochondrial, ATP linked, and reserve capacity OCR normalized to α-Tubulin protein during a Seahorse mitochondrial stress test. d, quantification of the red:green fluorescence intensity. e, representative merged images of pMitoTimer-transfected C2C12 myoblasts (scale bar = 10 μm); and f, quantification of red:green fluorescence intensities following treatment with dimethyl sulfoxide or diethylenetriamine-NO (D-NO, 200 μM) for 1 h followed by either dimethyl sulfoxide (Con), antimycin A (Ant A, 100 μM), or D-NO alone for 6 h. g, representative merged images of Drosophila heart tube MitoTimer signals (scale bar = 10 μm). h, quantification of the red:green fluorescence intensity. i, quantification of number of pure red puncta in the A1 heart tube segment of Drosophila following treatment with rotenone (0.4 mM in food for 4 days, n = 39), paraquat (10 mM in 5% sucrose for 24 h, n = 21), or antimycin A (0.1 mM in 5% sucrose for 24 h, n = 25). Controls were either normal food for 4 days (n = 24) or sucrose for 24 h (n = 23). k, quantification of red:green fluorescence intensity. l, quantification of pure red puncta from 7-day-old flies (n = 21) compared with 70-day-old flies (n = 20). Data represent mean ± S.E.; *, **, and *** denote p < 0.05, 0.01, and 0.001, respectively.
tion and/or reduced antioxidant defense, *i.e.* oxidative stress. Because MitoTimer is translated in the cytosol and transported to mitochondria, increased oxidative stress in the cytosol, decreased import (of newly synthesized green MitoTimer), and/or reduced degradation (of old red MitoTimer) could all potentially lead to the finding.

In adult fly heart tubes and adult mouse FDB muscles, we readily detected pure red puncta that is approximately the size of enlarged mitochondria. We suspected that these pure red puncta were severely damaged mitochondria, as they were often disconnected from, but close to, normal healthy mitochondria, consistent with a phenomenon of mitochondrial fission after damage. Furthermore, we found that the pure red puncta were similar in size and location to those of enlarged, partially vacuolated mitochondria with remnant cristae structures detected by transmission electron microscopy of muscle.
sections from mice on a HFD. The direct proof of this speculation came as evidence of co-localization of Cox4 immunofluorescence with the pure red puncta. Some pure red puncta did not show detectable Cox4 immunofluorescence possibly due to complete degradation of electron transport chain proteins. It is likely that severely damaged mitochondria no longer import new MitoTimer protein, thus appearing as pure red puncta once the remaining protein is oxidized. Another possibility is that some of the autolysosome containing degenerating mitochondria were fused with lysosome to form autolysosome and display only red fluorescence due to acidic pH as demonstrated similarly by pmCherry-GFP-LC3B previously (43). In support of this notion, we found that some yellow and red puncta were positive for LC3 and Lamp1, suggesting that they were part of autophagosome and autolysosome, respectively, targeted for degradation and removal from the cell.

Morphological analysis of the total fluorescent signal of MitoTimer allows for estimation of mitochondrial content and examination of structure. As shown in Fig. 4b, the % area occupied by the mitochondrial fluorescence signal of myofibers increased significantly in response to voluntary wheel running exercise training. Furthermore, high magnification of the MitoTimer images showed a typical structure of paired mitochondria on either side of the Z-line of the sarcomere. Exercise training not only increased the density of mitochondria, but also improved the size uniformity of the mitochondrial pair. Taken together, we have shown that the MitoTimer reporter could be used for robust, simultaneous analysis of multiple parameters, including mitochondrial content, structure, stress, and damage in vivo. These unique features of MitoTimer will prove to be valuable for mitochondrial research.

While the current studies were underway, Hernandez et al. (22) and Ferree et al. (21) published findings using a similar structural design of MitoTimer with the reporter gene under control of a doxycycline-inducible promoter for assessment of mitochondrial turnover in cultured cells. Hernandez et al. (22) provided evidence in HEK293 cells that MitoTimer targeted to the mitochondrial matrix and transitioned to red fluorescence...
over time demonstrating the usefulness in assessment of mitochondrial turnover and protein import. Ferree et al. (21) expanded on these findings in other cell types to show that mitochondrial dynamics and motility could also alter Mitotimer fluorescence and would thus be a useful reporter under conditions where mitochondria turnover and transport is affected. These findings set an important stage for this novel reporter gene being used for assessment of mitochondrial dynamics and health. In the current study, our application of Mitotimer in various animal models ranging from worms, flies to mice, allows analysis of mitochondrial content, structure, stress, and damage. This robust in vivo analysis of mitochondrial health adds significant value to the application of this novel reporter gene.

In summary, we took advantage of cultured cells, transgenic flies, transgenic worms, and somatic gene transfer in adult mice to show that the Mitotimer reporter gene in combination with computer-assisted imaging analysis is an excellent tool for robust, simultaneous measurements of mitochondrial content, structure, stress, and damage in vitro and in vivo. We anticipate that a combination of this technology with other cell and animal models will aid innovative research and facilitate the development of effective interventions to prevent and treat many of the prevalent diseases related to abnormal mitochondrial structure and function in various tissue/organ systems.

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