Mitochondria Express α7 Nicotinic Acetylcholine Receptors to Regulate Ca\textsuperscript{2+} Accumulation and Cytochrome c Release: Study on Isolated Mitochondria

Galyna Gergalova\textsuperscript{1}, Olena Lykhmus\textsuperscript{1}, Olena Kalashnyk\textsuperscript{1}, Lyudmyla Koval\textsuperscript{1}, Volodymyr Chernyshov\textsuperscript{1}, Elena Kryukova\textsuperscript{2}, Victor Tsetlin\textsuperscript{2}, Sergiy Komisarenko\textsuperscript{1}, Maryna Skok\textsuperscript{1}\textsuperscript{*}

\textsuperscript{1}Department of Molecular Immunology, Palladin Institute of Biochemistry, Kyiv, Ukraine, \textsuperscript{2}Department of Molecular Bases of Neurosignaling, Shemyakin-Ovchinnikov Institute of Bioorganic Chemistry, Russian Academy of Sciences, Moscow, Russia

Abstract

Nicotinic acetylcholine receptors (nAChRs) are ligand-gated ion channels that mediate synaptic transmission in the muscle and autonomic ganglia and regulate transmitter release in the brain. The nAChRs composed of α7 subunits are also expressed in non-excitable cells to regulate cell survival and proliferation. Up to now, functional α7 nAChRs were found exclusively on the cell plasma membrane. Here we show that they are expressed in mitochondria and regulate early pro-apoptotic events like cytochrome c release. The binding of α7-specific antibody with mouse liver mitochondria was revealed by electron microscopy. Outer membranes of mitochondria from the wild-type and β2−/− but not α7−/− mice bound α7 nAChR-specific antibody and toxins: FITC-labeled α-cobratoxin or Alexa 555-labeled α-bungarotoxin. α7 nAChR agonists (1 μM acetylcholine, 10 μM choline or 30 nM PNU-282987) impaired intramitochondrial Ca\textsuperscript{2+} accumulation and significantly decreased cytochrome c release stimulated with either 90 μM CaCl\textsubscript{2} or 0.5 mM H\textsubscript{2}O\textsubscript{2}. α7-specific antagonist methyllycaconitine (50 nM) did not affect Ca\textsuperscript{2+} accumulation in mitochondria but attenuated the effects of agonists on cytochrome c release. Inhibitor of voltage-dependent anion channel (VDAC) 4,4'-diisothio-cyano-2,2'-stilbene disulfonic acid (0.5 μM) decreased cytochrome c release stimulated with apoptogens similarly to α7 nAChR agonists, and VDAC was co-captured with the α7 nAChR from mitochondria outer membrane preparation in both direct and reverse sandwich ELISA. It is concluded that α7 nAChRs are expressed in mitochondria outer membrane to regulate the VDAC-mediated Ca\textsuperscript{2+} transport and mitochondrial permeability transition.

Introduction

Nicotinic acetylcholine receptors (nAChRs) are pentameric ligand-gated ion channels that were initially explored in muscle and autonomic ganglia and shown to mediate fast synaptic transmission [1]. In the brain, they regulate glutamate-, GABA-and dopamine-mediated transmission and are involved in the establishment of nicotine dependence [2]. Studies of the last decade documented the presence of nAChRs and nAChR-like receptors in many non-excitable cells of mammals, as well as in invertebrates, plants and even bacteria, where their functions are related to the general vital properties of living cells like proliferation, survival, adhesion and motility [3–5]. It is becoming increasingly clear that nAChRs have appeared in evolution long before the development of the nervous system and that they are multifunctional receptors employing different kinds of signaling in the cells of various origin.

Structurally, the nAChRs are composed of combinations of ten alpha (α1 to α10) and four beta (β1 to β4) subunits, muscle-type receptors contain also γ, δ or ε subunits. They can be assembled as heteropentamers (eg. (α1)\textsubscript{α} (β1)(β2)\textsubscript{β}) and, correspondingly, differ in their cation selectivity, kinetics of the ion channel opening and desensitization [1].

The homopentameric nAChRs composed of α7 subunits (α7 nAChRs) are of special interest, because they belong to the most ancient branch of this receptor’s family and were shown to be expressed in both neurons and non-excitable cells to mediate pro-proliferative, survival and anti-inflammatory signaling [6–9]. Previously we found that the absence of these receptors in α7−/− mice resulted in poorer survival of B lymphocyte precursors within the bone marrow [7]. Activation of nAChRs stimulated the growth of cancer cells and suppressed apoptosis [10], and the nAChR agonist nicotine could abolish the chemotherapy-induced apoptosis [11]. However, up to now, the pro-survival signaling was attributed exclusively to α7 nAChRs exposed on the cell plasma membrane. We posed a question: whether functional α7 nAChRs can be found in intracellular organelles, in particularly, in mitochondria, which are involved in inducing intracellular apoptotic pathway? Here we show that α7 nAChRs are expressed in the outer mitochondria membrane to regulate Ca\textsuperscript{2+} accumulation and cytochrome c release stimulated with apoptogens like high Ca\textsuperscript{2+} dose or H\textsubscript{2}O\textsubscript{2}.}
Results

The presence of α7 nAChRs in mitochondria

Mitochondria isolated from the liver of C57Bl/6 mice were treated with the antibody against the whole extracellular domain (1–208) of α7 nAChR subunit, followed by the 10 nm colloidal gold conjugated secondary antibody, and examined by electron microscopy. As shown in Fig. 1, the binding of α7(1–208)-specific antibody was detected on mitochondria bodies. However, positive staining was quite rare, probably, due to the mode of sample processing (cutting) for electron microscopy.

As the next step, we studied the presence of nAChRs in detergent lysates of mitochondria isolated from the liver of either wild-type or α2−/− mice. For this purpose, two types of sandwich assays were developed (Fig. 2, A, D). The αChR contained within the mitochondria preparation was captured with the antibody against α7(1–208) and was further revealed with either fluorescein isothiocyanate-labeled α-cobratoxin (CTX-FITC) or α7(179–190)-specific antibody. CTX is a long-chain α-neurotoxin from Naja kaouthia cobra venom; a specific ligand for the muscle-type, α7 and α9(α10) nAChRs of mammals [12]. Antibody against the extracellular epitope (179–190) of α7 nAChR subunit was generated by us previously [13] and was proven to be α7-specific in numerous experimental systems and assays including ELISA, Western blot and flow cytometry [7,14]. As shown in Fig. 2, B and E, the binding of both toxin and antibody was observed with the mitochondria of the wild-type but not α2−/− mice in two independent assays. When the wild-type mitochondria were fractionated into the outer (OM) and inner (IM) membranes, the binding of both CTX-FITC and α7(179–190)–specific antibody was found with the OM, but not IM preparation (that explained the rare positive staining of cut mitochondria in electron microscopy, Fig. 1). We further compared the OM preparations of the wild-type, α7−/− and α2−/− mice in the antibody- and toxin-based sandwich assays; in this case we used Alexa Fluor 555-labeled α-bungarotoxin previously shown to bind specifically α7 nAChRs in both α7-transfected model cells and in dorsal-root ganglia naturally expressing this nAChR subtype [15]. As shown in Fig. 2, C and F, positive signal was found in the preparations of the wild-type and α2−/−, but not α7−/− mitochondria. This data clearly indicated that α7 nAChR was present in the mitochondria outer membrane.

Figure 1. Electron microscopy images of mouse liver mitochondria. Isolated mitochondria were stained with rabbit α7(1–208)-specific antibody followed by 10 nm colloidal gold-conjugated anti-rabbit IgG. A – secondary antibody only (control); B – α7(1–208)-specific antibody plus secondary antibody; arrows (1) indicate the sites of positive staining.

Regulation of Ca2+ accumulation in mitochondria by α7 nAChR ligands

To reveal possible functions of α7 nAChRs in mitochondria, we took into account that this nAChR subtype is highly permeable to Ca2+ [1], whereas mitochondria are well-known intracellular Ca2+ depots. To find out whether mitochondrial α7 nAChRs are involved in Ca2+ transport we studied intramitochondrial Ca2+ accumulation in the presence or absence of α7-specific ligands. To exclude any contamination with the whole cells or plasma membranes, the isolated mitochondria were allocated by flow cytometry according to their size and granularity (Fig. 3, A). 95% of particles within the gated population incorporated mitochondria-specific fluorescent dye acridine orange 10-nonyl bromide [16] indicating that it contained pure mitochondria (Fig. 3, B). Experiments were performed in live mitochondria maintaining their membrane potential monitored with the potential-sensitive fluorescent dye tetramethyl rhodamine methyl ester [17]. Addition of 90 µM CaCl2 to the mitochondria loaded with Ca2+-sensitive fluorescent dye Fluo 3-AM evoked the fluorescent signal within the first two minutes; the signal was maintained during at least 10 min more and was completely abolished by the addition of 1 mM EGTA (Fig. 3, C) or 0.1 µM carbonyl cyanide-3-chlorophenylhydrazide (CCCP, data not shown). When CaCl2 application was preceded with that of α7 nAChR agonists choline, acetylcholine or PNU-282987, but not a specific competitive antagonist methyllycaconitine (MLA), Ca2+ accumulation was inhibited by about 20% (Fig. 3, D). Similar effect was exerted by the VDAC inhibitor 4,4’-disothiocyano-2,2’-stibene disulfonic acid (DIDS). VDAC is also located in mitochondria outer membrane and facilitates Ca2+ entry from the cytosol into the intermembrane mitochondria space [18]. Similar effects of DIDS and α7 nAChR agonists on Ca2+ accumulation in mitochondria pushed us to search a further physical and functional connection between VDAC and α7 nAChR.

Connection of α7 nAChR to VDAC in the outer mitochondria membrane

To elucidate if there is an interaction between α7 nAChR and VDAC in the outer mitochondria membrane we developed another set of sandwich immunoenzyme assays presented in Fig. 4 A, C. These assays are analogues of immunoprecipitation where the complex of two interacting proteins is captured from the mixture with the antibody against one component and is revealed in Western blot with the antibody against another component. The advantage of sandwich assay is that both steps are performed in the same media (immunoplate) and that the second antibody binding is evaluated photometrically as in conventional ELISA. Previously, we used such an approach to reveal the interaction of α7 nAChR with CD40 and of αβ2 nAChRs with IgM in B lymphocytes [19]. In the first assay, the α7 nAChR was captured from the OM preparation with anti-α7(1–208) and was revealed with either anti-α7(179–190) or anti-VDAC (Fig. 4, A). As shown in Fig. 4, B, the OM preparation captured with anti-α7(1–208) was revealed with both anti-VDAC and anti-α7(179–190). In the “reverse” assay (Fig. 4, C), the complex was captured with either anti-VDAC or the antibody against mitochondria outer membrane translocase (anti-TOM22) and was revealed with anti-α7(1–208). In this case, the nAChR-specific antibody recognized the complex captured with anti-VDAC and not with anti-TOM22 (Fig. 4, D). These data clearly demonstrated that α7 nAChR interacts with the VDAC in the outer mitochondria membrane.

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Regulation of cytochrome c release from mitochondria by α7 nAChR ligands

VDAC is a key element in mitochondria permeability transition pore (MPTP) formation accompanied with cytochrome c (cyt c) release [20] which is the initial step of mitochondria-driven apoptosis. To test if mitochondrial α7 nAChRs are involved in apoptosis-related processes, we studied the effects of α7 nAChR ligands on the cyt c release from purified mitochondria stimulated with either high dose of CaCl2 or H2O2. Preliminary data demonstrated that cytochrome c release stimulated with 0.5 mM H2O2 was inhibited with α7 nAChR agonists and DIDS [21]. Here we show that 90 μM CaCl2 stimulated extensive cyt c release from live mitochondria similarly to 0.5 mM H2O2; both were significantly inhibited with 0.5 μM DIDS or α7 nAChR agonists (choline, acetylcholine or PNU-282987, Fig. 5). MLA did not affect cyt c release itself but prevented inhibitory effects of all agonists. This data clearly indicated that α7 nAChRs were involved in regulating MPTP formation, and their activation with agonists produced an effect similar to inhibition of VDAC by DIDS.

Discussion

The presence of nicotinic receptors in mitochondria has initially been discussed in connection with the neuroprotective role of nicotine. Then it was shown that nicotine affected mitochondria respiratory chain independently on nAChRs [22]. However, in other studies, the decrease of mitochondria membrane potential caused by ethanol was prevented with specific α7 nAChR agonist 2,4-dimethoxybenziliden anabasein and this effect was blocked with MLA [23]. In the latter work, α7 nAChR agonists also attenuated cytochrome c release stimulated by ethanol in rat hippocampal neuronal cultures that is in good agreement with our data. In our earlier published experiments, 1 μM nicotine prevented Ca2+ accumulation in isolated mitochondria similarly to 1 μM choline, and mitochondria from mice injected with α7(1–208)-specific antibody possessed lower membrane potential than those from mice injected with non-specific IgG, suggesting the involvement of α7 nAChRs [24]. The binding of α7-specific antibody with mitochondria of rat hippocampus was demonstrated by electron microscopy; however, this was not confirmed by Western blot analysis [25] and, therefore, did not allow the authors to state the expression of α7 nAChRs in mitochondria. Our data clearly demonstrate that nAChRs of α7 subtype are present in the outer membranes of mitochondria isolated from the mouse liver. This was shown by electron microscopy with α7(1–208)-specific antibody and by sandwich ELISA with α7(179–190)-specific antibody. Since the specificity of many antibodies against nAChRs was put in doubt [26–27], we confirmed our results by independent binding of either α7-cobratoxin or α7-bungarotoxin fluorescent derivatives and used mitochondria from the wild-type, α7−/− or β2−/− mice to prove the specificity of binding (Fig. 2 B, C). Mouse liver is a recognized source for mitochondria isolation [28]; it is yet to be established if mitochondria from other...
tissues and species contain similar or different nAChR quantity and/or subtype.

To reveal the functions of α7 nAChRs in mitochondria, we took into account that nAChRs expressed in non-excitable cells trigger intracellular signaling and affect the activity of adjacent receptors by either ion-dependent or independent mechanism [29–31]. Previously we found that α7 nAChR expressed in mouse B lymphocytes was coupled with CD40 and regulated CD40-mediated B lymphocyte activation [19]. Similar approach applied to mitochondria indicates that mitochondrial α7 nAChR is coupled to VDAC that may underlie similar effects of α7 agonists and DIDS on Ca2+ accumulation and cyt c release (Fig. 3, D and 5).

VDAC, positioned on the interface between mitochondria and the cytosol, is responsible for the fluxes of various metabolites across mitochondria outer membrane [32–33]. It is suggested to increase the local Ca2+ concentration in the intermembrane mitochondria space thus facilitating the Ca2+ uniporter activity [34]. It has been also recognized as a key protein in MPTP formation and induction of mitochondria-mediated apoptosis [35]. VDAC is easily converted from anion-selective to cation-permeable state by environmental conditions [36] and is oligomerized to be involved in MPTP [18]. Similar effects of α7 nAChR agonists and of DIDS, which prevents VDAC’s oligomerization [18], allow suggesting that mitochondrial α7 nAChR signaling affects the neighboring VDAC thus preventing its involvement in MPTP. Interestingly, the doses of choline (10 μM) and acetylcholine (1 μM) affecting both Ca2+ accumulation and cyt c release were much lower than those reported to open the α7 nAChR ion channel (EC50 1.6 mM for choline and 79–316 μM for acetylcholine); that of synthetic agonist PNU-282987 (30 nM) was closer to but still less than its reported functional potency (128 nM) [37]. Since the effects of all tested agonists on cyt c release were inhibited with the competitive inhibitor MLA, it may be suggested that mitochondrial α7 nAChRs are much more sensitive to natural agonists than those expressed in the plasma membrane, possibly, due to specific lipid surrounding of mitochondria outer membrane [38]. The lipids were shown to influence the ability of the nicotinic acetylcholine receptor to gating in response to neurotransmitter binding [39]. In addition, the exact subunit composition of α7-containing mitochondrial nAChRs is still to be elucidated, since heteromeric α7β2 nAChRs were shown to possess different pharmacological sensitivity compared to α7 homopentamers [40].

Recognizing the α7 nAChR presence in mitochondria poses a question about its natural ligand(s) and physiological significance. Choline is actively transported into the cell from extracellular space and is present in the cytosol at 30 to 50 μM [41]. According to our data, it is sufficient to activate mitochondrial α7 nAChRs and to keep mitochondria “resistant” to apoptogenic agents. However, the half-life time of intracellular choline is very short [42], therefore, the mitochondria protection depends on the

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**Figure 3. The effects of α7-specific ligands and DIDS on Ca2+ accumulation in mitochondria studied by flow cytometry.**

A – purified liver mitochondria gated by size (Forward scatter) and granularity (Side scatter) in flow cytometry. B – binding of 0.1 μM acidine orange 10-nonyl bromide (NAO) to the gated mitochondria population; Control – the non-stained mitochondria. C – Ca2+ accumulation in mitochondria loaded with Fluo 3-AM. D – Ca2+ accumulated in mitochondria during 2 min after 1 min pretreatment with DIDS, acetylcholine, choline, PNU-282987 or MLA; data are shown as normalized mean fluorescence values of 5 independent experiments for each ligand; * – P<0.05 compared to the fluorescence in the absence of α7 nAChR agonists or DIDS.

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balance of its intake and degradation. Recent evidence demonstrates the increased susceptibility of mitochondria to calcium-induced permeability transition upon choline deficiency [43] that supports our idea. In contrast, intracellular release of choline upon ischemia [42] obviously has the protective anti-apoptotic effect. In addition, according to proteomic studies, mitochondria contain choline acetyltransferase [44] and, therefore, are able to locally synthesize acetylcholine.

In summary, our results demonstrate that α7 nAChRs are expressed on mitochondria outer membrane and regulate Ca\(^{2+}\) accumulation and cyt c release, the initial step of apoptosis induction. This means that, in addition to established anti-apoptotic signaling pathways mediated by plasma membrane α7 nAChRs [8–9], there is an endogenous, previously unrecognized cholinergic mechanism to control mitochondria functions and their apoptotic susceptibility. Probably, it belongs to the most

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**Figure 4. Connection of α7 nAChR and VDAC studied by sandwich assays.** The schemes (A, C) and results of direct (B, n = 3) and reverse (D, n = 3) sandwich assays demonstrating the connection of α7 nAChR and VDAC in the outer membranes (OM) of the wild-type (WT) mitochondria. Anti-TOM22 – antibody against mitochondrial outer membrane translocase. doi:10.1371/journal.pone.0031361.g004

**Figure 5. Cytochrome c release into mitochondria supernatants.** Cyt c released from mitochondria in 2 min after addition of either 0.5 mM H\(_2\)O\(_2\) or 90 μM CaCl\(_2\) in the presence or absence of DIDS or α7 nAChR ligands. Each column corresponds to M±SE of three independent measurements. Mit – mitochondria without apoptogens; Contr – mitochondria treated with H\(_2\)O\(_2\) or CaCl\(_2\) only. doi:10.1371/journal.pone.0031361.g005
ancient survival mechanisms inherited by mitochondria from their hypothetic prokaryotic ancestor [45]. This finding offers a novel view on the mitochondria protection in apoptosis and opens the way for its pharmacological regulation.

Materials and Methods

Ethics Statement
We used age-matched male wild-type and mutant (lacking either α7 or β2 nicotinic receptor subunit [46–47]) mice with common C57BL/6J background. The mice were kept in the animal facilities of Pasteur Institute, Paris and Palladin Institute of Biochemistry, Kyiv. They were housed in a quiet, temperature-controlled room (22–23°C) and were provided with water and dry food pellets ad libitum. Before removing the liver mice were sacrificed by cervical dislocation. All procedures conformed to the guidelines of the Centre National de la Recherche Scientifique or IACUC of Palladin Institute. Before starting the experiments, the protocols were approved by the Animal Care and Use Committee of Palladin Institute of Biochemistry (Protocol 1/7-421).

Reagents
All reagents were of chemical grade and were purchased from Sigma-Aldrich unless specially indicated. Alexa Fluor 555 α-bungarotoxin was purchased from Invitrogen (USA). CTX-FITC and antibodies against α7(1–208) or α7(179–190) were obtained by us previously [13,48–49].

Mitochondria purification and fractionation
Mitochondria isolation from the mouse liver and fractionation into inner and outer membranes was performed by differential ultracentrifugation according to standard procedure described [50]. The separation medium contained 10 mM HEPES, 1 mM EGTA and 250 mM sucrose, pH 7.4, 4°C. For the membrane preparation, the isolated mitochondria were resuspended in 10 ml 10 mM KH2PO4, left on ice for 5 min and spun at 8000 g for 10 min. The pellet was resuspended in 10 ml 125 mM KCl, 10 mM Tris-MOPS (pH 7.4, 4°C) and centrifuged again. The supernatant was withdrawn, while the pellet was resuspended in the mixture of 10 mM KH2PO4 (10 ml) and 1.8 M sucrose, 2 mM ATP, 2 mM MgSO4 (3.5 ml). The sample was sonicated at 4 W for 20 min, laid on the top of 15 ml 1.18 M sucrose and spun for 2 h at 90 000 g. The inner membranes were collected as the brown pellet in the tube bottom, while the outer membranes were precipitated in the interphase. To prepare detergent lysates, the mitochondria suspension or membrane fractions were frozen at −70°C, thawed and treated with the lysing buffer (0.01 M Tris –HCl, pH 8.0; 0.14 M NaCl; 0.025% NaN3; 1% Tween 20) and protease inhibitors cocktail for 2 h on ice upon intensive stirring. The resulting lysate was cleared by centrifugation (20 min at 20 000 g) and dialysed against PBS containing 0.025% NaN3 and protease inhibitors. The protein concentration was established by Bradford assay.

Electron microscopy
The purified mitochondria were treated in suspension with α7(1–208)-specific rabbit antibody (0.05 mg/ml) for 30 min at RT, washed by centrifugation and subsequently incubated with 10 nm colloidal gold-labeled goat anti-rabbit immunogoldlubins for additional 30 min. Control preparation was treated with the second antibody only. The washed mitochondria were pelleted and fixed with 2.5% glutaraldehyde in 0.1 M sodium cacodylate buffer pH 7.4 during 3 h at RT. The samples were additionally fixed with 1% OsO4 in 0.05 M cacodylate buffer during 2 h and were dehydrated in 30%, 50%, 70%, 90% and 100% acetone subsequently. The obtained preparations were polymerized in epoxy resin Apon-Argylt (AGAR, UK). Ultrathin lesions (70–100 nm) were prepared with Ultratome LKB-V (LKB, Sweden) and were stained with 1% uranyl acetate (60 min) and the lead dye (2 min). The preparations were analyzed under electron microscope Hitachi, Japan with 20,000 amplification.

Flow cytometry studies
The purified mitochondria (200 μg of protein per ml in the standard sample) were resuspended in the incubation medium containing 10 mM HEPES, 125 mM KCl, 25 mM NaCl, pH 7.4. For Ca2+-related studies the incubation medium was supplemented with 5 mM sodium succinate and 0.1 mM Pi(K), pH 7.4. The purity of gated mitochondria in flow cytometry was assessed using 0.1 μM acridine orange 10-nonyl bromide (NAO) added immediately before flow cytometry examination. For Ca2+ uptake studies, mitochondria were pre-incubated with 1 μM Fluo 3-AM for 30 min in the dark. CaCl2, DIDS and α7 nAChR ligands were added as described in the Fig. 3 and its legend. The studies were performed with the COULTER EPICS-XL™ fluorescent flow cytometer (Beckman Coulter, USA) at room temperature.

Sandwich assays
The 96-well plates (Nunc MaxiSorp, Denmark) were coated with rabbit α7(1–208)-specific antibody and were subsequently blocked with 1% BSA/PBS. The detergent lysates of mitochondria or their membranes were applied into the coated wells, 600 μg/ml (for the whole lysate) or 100 μg/ml (for membranes). After 2 h of incubation at 37°C the plates were washed with water. In ELISA version, the bound antigen was revealed with bioinylated rabbit α7(179–190)-specific antibody (Santa Cruz Biotechnology, USA) applied for additional 2 h and followed by Extravidin-peroxidase conjugate and o-phenylenediamine-containing substrate solution. The absorbance at 490 nm was read by the StatFax-2100 Microplate reader (Awareness Technology, USA). In the combined antibody-toxin version, the bound antigen was detected with either CTX-FITC (10–20 μg/ml) or Alexa Fluor 555-labeled α-bungarotoxin in 30 nM final concentration applied for 1 h (CTX) or overnight (α-bungarotoxin) at RT and washed out with PBS. The fluorescence of dry plates was then read with FLx800 Multi-detection microplate reader (BioTek, USA) or fluorimeter FFM-01 (Kortek, Russia) using excitation/emission wavelength of 485/530 nm or 546/607, respectively.

Cyt c release studies
The purified mitochondria (120 μg of protein per ml) were incubated with either 90 μM CaCl2 or 0.5 mM H2O2 in the presence or absence of DIDS or nAChR ligands for 2 min at room temperature and were immediately pelleted by centrifugation (10 min., 7000 g at 4°C). The supernatants were collected and tested by sandwich assay. The plates were coated with ammonium sulfate-precipitated fraction of bovine cyt c-specific rabbit antiserum and blocked with 1% BSA. Mitochondria supernatants were applied at optimal dilutions established in preliminary experiments. Calibration curve was built using bovine cyt c. The bound cyt c was revealed with biotinylated immunoglobulins from cyt c-specific rabbit serum followed by Extravidin-peroxidase conjugate and o-phenylenediamine-containing substrate solution. This assay became possible because of multiple epitopes recognized on cyt c molecule, so that antibodies raised against bovine cyt c could recognize its mouse analogue.
Statistical analysis
Each experiment was reproduced independently minimum three times with 3 to 6 repeats for each point. Statistical analysis was performed according to Student’s test using OriginPro 8.6 software. The data are presented as mean and SEM. The difference was considered significant at P<0.05 (*).

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Author Contributions
Conceived and designed the experiments: MS GG OL VT EK. Performed the experiments: GG OL OK LC VC EK. Analyzed the data: GO OL MS VT SR EK VC. Contributed reagents/materials/analysis tools: VT. Wrote the paper: MS GG VT.

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