4-Hydroxytamoxifen induces slight uncoupling of mitochondrial oxidative phosphorylation system in relation to the deleterious effects of tamoxifen

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Abstract

The use of tamoxifen (TAM) has been questioned on the chemotherapy and chemoprevention of breast cancer due to several estrogen receptor-independent cytotoxic effects. As an alternative, its more active metabolite 4-hydroxytamoxifen (OHTAM) has been proposed with presumed lower side effects. In this work, the potential OHTAM toxicity on rat liver mitochondrial bioenergetics in relation to the multiple deleterious effects of TAM was evaluated. OHTAM, at concentrations lower than those putatively reached in tissues following the administration of TAM, does not induce significant perturbations on the respiratory control ratio (RCR), ADP/O, transmembrane potential (∆Ψ), phosphorylative capacity and membrane integrity of mitochondria. However, at high concentrations, OHTAM depresses the ∆Ψ, RCR and ADP/O, affecting the phosphorylation efficiency, as also inferred from the ∆Ψ fluctuations and pH changes associated with ADP phosphorylation. Moreover, OHTAM, at concentrations that stimulate the rate of state 4 respiration in parallel to the decrease in the ∆Ψ and phosphorylation rate, causes mitochondrial swelling and stimulates both ATPase and citrate synthase activities. However, the OHTAM-observed effects, at high concentrations, are not significant relatively to the damaging effects promoted by TAM and suggest alterations to mitochondrial functions due to proton leak across the mitochondrial inner membrane. © 2002 Elsevier Science Ireland Ltd. All rights reserved.

Keywords: 4-Hydroxytamoxifen; Breast cancer; Mitochondrial bioenergetics; Mitochondrial transmembrane potential; Oxidative phosphorylation efficiency; Proton leak; Membrane permeabilization

Abbreviations: ADP/O ratio, number of ADP molecules added to the medium per oxygen atom consumed during phosphorylation; ANT, adenine nucleotide translocator; BSA, bovine serum albumin; ER, estrogen receptor; EGTA, ethylene glycol-bis (β-aminoethyl ether) NNN’-tetraacetic acid; FCCP, carbonyl cyanide p-trifluoromethoxyphenylhydrazone; HEPES, 4-(2-hydroxymethyl)-1-piperazineethanesulfonic acid; Mops, 3-(N-Morpholino)propanesulfonic acid; OHTAM, 4-hydroxytamoxifen; RCR, respiratory control ratio (= state 3/state 4); SR, sarcoplasmic reticulum; TAM, Tamoxifen; Tris, Tris (hydroxymethyl)-aminomethan; TPP⁺, tetraphenylphosphonium; ∆Ψ, Mitochondrial membrane potential; ∆ρ, proton motive force.

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1. Introduction

Tamoxifen (TAM) is an important adjunct chemotherapeutic agent for treating breast cancer (Early Breast Cancer Trialists’ Collaborative Group, 1998). This non-steroidal antiestrogen has also been reported as an effective agent in the chemoprevention of healthy women at high risk of the disease (Ault and Brandbury, 1998). However, TAM causes serious side effects, such as hepatocellular damage and agranulocytosis (Ching et al., 1992), increased risk of uterine and endometrial cancers (Ault and Brandbury 1998; Early Breast Cancer Trialists’ Collaborative Group, 1998), hemolytic anemia (Ching et al., 1992; Suwalsky et al., 1998), inhibition of Ca²⁺ uptake by sarcoplasmic reticulum (SR) (Custódio et al., 1996; Kargacin et al., 2000), and erythrocyte hemolysis (Cruz Silva et al., 2000). Therefore, an alternative breast cancer treatment with presumed lower side effects, such as the percutaneous administration of 4-hydroxytamoxifen (OHTAM), the major active metabolite of TAM and responsible for its antiestrogenic activity, has already been performed (Mauvais-Jarvis et al., 1986).

Previous studies have reported that, in comparison with TAM, OHTAM has higher affinity and specificity for the estrogen receptor (ER) (Borgna and Rochefort, 1981) and, besides being 100- to 200- times more potent (Osborne et al., 1992; Jordan et al., 1988), it is pharmacologically well tolerated (Sauvèz et al., 1999) and less toxic to erythrocytes (Cruz Silva et al., 2001) and to the SR calcium pump (Custódio et al., 1996). Furthermore, in contrast to TAM, OHTAM fails to induce either mitochondrial depolarization or caspase activation and, subsequently, does not induce apoptosis in p53(−) human mammary epithelial cells [p53(−)HMEC] (Dietze et al., 2001). Despite these and other studies, including those reporting the differences between the OHTAM percutaneous and oral administration (Pujol et al., 1995; Sauvèz et al., 1999), the toxicity mechanisms of OHTAM are not yet clarified. Therefore, toxicological studies are required to assure a safe and appropriate administration of OHTAM on breast cancer therapy and to achieve a better understanding of the molecular pharmacology of this drug.

Mitochondria are important subcellular sites recognized as energy generators for the cell, as well as control points of cell replication, differentiation, ion regulation and other active transport processes of intermediary metabolism (Wallace et al., 1997; Wallace and Starkov, 2000). Moreover, recent studies have demonstrated that mitochondria play a key role in cell signaling and in the apoptotic cell death (Loeffler and Kroemer, 2000). Consequently, changes in the structural and functional characteristics of mitochondria provide a number of primary targets for xenobiotic-induced toxicity and cell death (Wallace and Starkov, 2000). Therefore, in addition to the different effects reported for TAM and its metabolite OHTAM on mitochondria structure and apoptosis of p53(−)HMEC cells (Dietze et al., 2001), the evaluation of OHTAM effects on mitochondrial bioenergetic functions are important to a better understanding of its molecular action mechanisms and potential mitochondrial toxicity in relation to the damaging effects of TAM (Cardoso et al., 2001).

2. Materials and methods

2.1. Chemicals

OHTAM, bovine serum albumin (BSA), 4-(2-hydroxymethyl)-1-piperazinethanesulfonic acid (HEPES), ethylene glycol-bis (β-aminoethyl ether) NNN’N-tetraacetic acid (EGTA), 3-[N-Morpholino]propanesulfonic acid (Mops), 5,5’-dithiobis(2-nitrobenzoic acid), oxaloacetic acid, acetyl-coA, nigericin and oligomycin were purchased from Sigma Chemical Co (St. Louis, MO). Tris (hydroxymethyl)-aminomethan (Tris), tetraphenylphosphonium (TPP⁺), carbonyl cyanide p-trifluoromethoxyphenylhydrazone (FCCP) and sucrose were obtained from Merck (Dramstadt, Germany). All other chemicals were commercial products of the highest purity grade available and solutions were prepared in deionized ultra pure water.
2.2. Preparation of rat liver mitochondria

Mitochondria were isolated from the liver of overnight fasted Wistar rats (250–300 g) of either sex by differential centrifugation as described elsewhere (Custódio et al., 1998), with slight modifications. Animals were killed by decapitation and the liver was quickly homogenized. The homogenate was prepared in 250 mM sucrose, 10 mM HEPES (pH 7.4), 1 mM EGTA and 0.1% (w/v) free fatty acid BSA. The liver homogenate was centrifuged at 500 × g for 10 min at 4 °C and mitochondria were recovered from the supernatant by centrifugation at 10 000 × g for 10 min. The mitochondrial pellet was resuspended twice in the washing medium containing 250 mM sucrose, 10 mM HEPES, pH 7.2. Mitochondrial protein was determined by the biuret method using BSA as the standard (Gornall et al., 1949).

2.3. Mitochondrial membrane potential measurements

Mitochondrial membrane potential (ΔΨ) was estimated by calculating transmembrane distribution of TPP⁺ with a TPP⁺-selective electrode prepared according to Kamo et al. (1979), using an Ag/AgCl-saturated electrode as reference (model MI 402; Microelectrodes, Inc., Bedford, NH). ΔΨ is expressed in mV and TPP⁺ uptake was measured from the decreased TPP⁺ concentration in the medium sensed by the electrode. The differences between the selective and the reference electrodes were measured with an electrometer and were recorded continuously (Kamo et al., 1979; Moreno and Madeira, 1991). Reactions were carried out in a closed chamber at 25 °C, with magnetic stirring in 1 ml of standard respiratory medium (130 mM sucrose, 50 mM KCl, 2.5 mM MgCl₂, 2.5 mM KH₂PO₄, 5 mM HEPES and 0.1 mM EGTA) supplemented with 2 μM rotenone and 4 μM TPP⁺. OHTAM was added from ethanolic solutions to mitochondria (1 mg) and pre-incubated for 3 min before the addition of 5 mM succinate to ensure the complete incorporation of the compound into the membrane due to its lipophilic characteristics (Custódio et al., 1991). After reaching a steady-state distribution of TPP⁺, ADP (150 nmol/mg protein) was added. Preliminary calibrations run in the presence of OHTAM excluded any direct interference of the drug with the electrode signal.

2.4. Mitochondrial respiration

The oxygen consumption (respiration rate) of isolated mitochondria was measured polarographically using a Clark-type oxygen electrode (YSI model 5331, Yellow Spring Inst.), in a 1 ml thermostated water-jacketed closed chamber with magnetic stirring and connected to a suitable recorder. OHTAM in ethanolic solutions (up to 3 μl) was incubated for 3 min at 25 °C in 1 ml of the standard respiratory medium previously supplemented with mitochondria (1 mg) and 2 μM rotenone. State 4 respiration was initiated with 5 mM succinate and ADP (150 nmol/mg protein) was added to establish state 3 respiration. Respiration rates were calculated assuming an oxygen concentration of 240 nmol O₂/ml in the experimental medium at 25 °C. Respiratory control ratio (RCR = state 3/state 4), respiratory rates, and ADP/O ratios (number of ADP molecules added to the medium per oxygen atom consumed during phosphorylation) were calculated according to Chance and Williams (1956).

2.5. Simultaneous measurements of respiration rate and ΔΨ

In order to determine the effects of OHTAM on mitochondrial function and to discriminate its site of action on mitochondria we adopted the approach commonly used for top–down metabolic control analysis (Brand, 1990; Hafner et al., 1990; Lionetti et al., 1998). With this approach, oxidative phosphorylation is conceptually divided into three subsystems, i.e., mitochondrial respiratory chain, phosphorylation system and the proton leak. To assess the effects of OHTAM on the proton motive force (Δp) producing system (the respiratory chain complexes and the substrate transporters), mitochondrial respiration and ΔΨ were measured simultaneously at 25 °C in a 1-ml incubation chamber fitted with a Clark-type oxygen electrode and a TPP⁺ sensitive electrode.
Mitochondria (1 mg) were suspended in the reaction medium containing 130 mM sucrose, 100 mM KCl, 2.5 mM MgCl₂, 2.5 mM KH₂PO₄, 5 mM HEPES and 0.1 mM EGTA, supplemented with 2 μM rotenone, 4 μM TPP⁺, 1 μg oligomycin (to eliminate phosphorylation) and 50 ng nigericin, which was used to collapse the pH gradient across the mitochondrial inner membrane (Murphy and Brand, 1987; Hafner and Brand, 1991), making the ΔΨ the sole component of Δp. Mitochondria were incubated under these conditions for 3 min in the presence of OHTAM. Respiration was initiated by the addition of 5 mM succinate and after a steady-state TPP⁺/C₂⁷ distribution the titration was performed by successive additions of FCCP (10 nM), according to previously described procedures (Brand, 1990; Lionetti et al., 1998). The same protocol was followed to assess the effects of OHTAM on proton leak (i.e. the passive permeability of the inner mitochondrial membrane to protons and any cation cycling reactions), but the titration was done with successive additions of malonate (0.5 mM). Titration experiments with OHTAM were conducted under similar conditions in the absence of either FCCP or malonate.

2.6. Measurement of mitochondrial swelling

Mitochondrial osmotic volume changes were followed by monitoring the decrease in absorbance (light-scattering) at 540 nm (Custo´dio et al., 1998) with a Perkin Elmer, Lambda 6 UV/VIS spectrophotometer. The reactions were performed with 1 mg mitochondrial protein in 2 ml of a reaction medium containing 200 mM sucrose, 10 mM Tris-Mops (pH 7.4), 1 mM KH₂PO₄ and 10 μM EGTA. OHTAM and TAM were added to mitochondrial suspensions in the absence of respiratory substrates and preincubated for 3 min at 30 °C.

2.7. Enzymatic activities

ATPase activity was estimated by monitoring the pH changes of the medium associated with ATP hydrolysis (Madeira et al., 1974). The experiments were carried out at 25 °C in 2 ml of the standard respiratory medium lightly buffered containing 0.5 mM HEPES, 130 mM sucrose, 50 mM KCl, 2.5 mM MgCl₂, 5 mM KH₂PO₄, 0.1 mM EGTA, pH 7.2 and supplemented with 2 μM rotenone. OHTAM and TAM were incubated with mitochondria (1 mg) for 3 min before starting the reactions with Mg-ATP as described elsewhere (Ferreira et al., 1997). Oligomycin (1 μg/mg protein) was added 1.5 min after starting the reactions.

ATP-synthase activity was determined by measuring the pH changes of the medium associated with ATP synthesis as reported by Moreno and Madeira (1991), at 25 °C, in 2 ml of reaction medium containing 130 mM sucrose, 50 mM KCl, 2.5 mM MgCl₂, 5 mM KH₂PO₄, 0.5 mM HEPES, 0.1 mM EGTA, pH 7.2, supplemented with 2 μM rotenone, 100 μM ADP and mitochondria (1 mg). The experiments were initiated by the addition of 5 mM succinate. TAM and OHTAM, in ethanolic solutions, were added and allowed to incubate with mitochondria for 3 min before starting the reactions.

Citrate synthase activity was monitored by the increase in absorbance at 412 nm as previously described (Shepherd and Garland, 1969; Trounce et al., 1996), using a Perkin Elmer, Lambda 6 UV/VIS spectrophotometer. The reactions were performed with 100 μg of mitochondrial protein suspended in 2 ml of the reaction medium containing 100 mM Tris (pH 8), 200 μM 5,5’-dithio-bis(2-nitrobenzoic acid) and 200 μM acetylCoA. The assays were started by the addition of 100 μM oxaloacetic acid after incubation of mitochondria in the absence and presence of either 0.1% Triton X-100, TAM or OHTAM for 3 min at 30 °C.

3. Results

To assess whether OHTAM affects mitochondrial bioenergetic functions, we evaluated the respiration parameters, both the RCR and ADP/O ratio (Fig. 1), and the ΔΨ (Fig. 2) of mitochondria energized with succinate. OHTAM pre-incubated with mitochondria progressively decreases RCR and ADP/O ratios as a function of concentration (Fig. 1). However, both ratios are not significantly affected by OHTAM as opposed to the high decrease observed at much lower concen-
Addition- ally, OHTAM at concentrations up to 70 nmol/mg protein does not yet induce complete uncoupling of mitochondria (Figs. 1 and 2). In contrast, TAM induces such drastic effect at 40 nmol TAM/mg protein (Cardoso et al., 2001), suggesting that OHTAM is much less toxic to mitochondrial membrane than TAM.

The effects of several concentrations of OHTAM on ΔΨ fluctuations associated with either the mitochondrial respiration or the phosphorylation cycle induced by ADP are shown in Fig. 2. Under succinate energization, mitochondria build up a potential of about −220 mV (negative inside). Upon ADP addition (150 nmol/mg protein), to initiate state 3 respiration (synthesis of ATP), the potential drops immediately to −190 mV, since the phosphorylation system consumes ΔΨ to transport and to phosphorylate ADP. After a short lag phase, when the phosphorylation cycle is completed, the transmembrane potential returns to nearly its initial value. OHTAM, at concentrations lower than those putatively reached in tissues following the administration of TAM (McCague et al., 1990; Jordan, 1990; Lien et al., 1991), slightly depresses the ΔΨ and the depolarization after ADP addition. However, at high concentrations (> 40 nmol/mg protein), the lag phase preceding repolarization displays a marked increase and the rate of repolarization progressively decreases (Fig. 2), suggesting that OHTAM affects the efficiency of the mitochondrial phosphorylative system. These effects become more pronounced at higher OHTAM concentrations whereas TAM, at much lower concentrations, induces a drastic depolarization of ΔΨ and total uncoupling of mitochondria, making them unable to phosphorylate the added ADP (Cardoso et al., 2001). This is evidenced in Fig. 3 that shows the effects of both drugs on pH changes associated with ATP synthesis. Thus, the consequences of OHTAM on the phosphorylation cycle induced by ADP, as shown in Fig. 2, are consistent with those on ADP/O ratio (Fig. 1) and with the promoted decrease in the rate of ATP synthesis by intact mitochondria (Fig. 3). In fact, increasing concentrations of OHTAM depress the rate of ATP synthesis, increasing the time required to phosphorylate the ADP (200 nmol/mg protein) (Fig. 3). However, it is noteworthy that TAM, at a concentration of 50 nmol/mg protein, completely inhibits the phosphorylation of the added ADP, in contrast to OHTAM that at the same concentration only decreases the rate of phosphorylation (Fig. 3).

A simultaneous follow-up of OHTAM effects on both ΔΨ and state 4 respiration (Fig. 4) shows that sequential additions of this drug to mitochondria causes a small decrease in ΔΨ and a parallel stimulation of state 4 respiration to compensate the induced depolarization of ΔΨ. Subsequent additions of OHTAM lead to a lower stimulation of respiration rate, concomitantly with a significant depolarization of ΔΨ (results not shown).

In agreement with the depolarization of ΔΨ (Fig. 2) and the stimulation of state 4 (Fig. 4) caused by a proton leak through the mitochondrial inner membrane, OHTAM increases the ATPase activity of tightly coupled intact mitochondria (Fig. 5). However, these effects are smaller than those promoted by TAM, which induces disruption of the mitochondrial inner membrane integ-
rity, as evidenced in Fig. 5 and further in Fig. 8 and Fig. 9.

To clarify the mechanisms responsible for the stimulation of state 4 respiration and ΔΨ depolarization, the effects of OHTAM on mitochondria were evaluated according to the approach developed by Brand (1990) (Fig. 6). This approach has been pointed to be useful to determine how compounds of pharmacological interest affect the mitochondria (Fusi et al., 1992). Nonphosphorylating mitochondria (in the presence of oligomycin) were titrated with FCCP to promote an increase in respiratory rate and a decrease in ΔΨ. The plot of ΔΨ versus respiration rate describes the kinetic response of the Δρ generators to ΔΨ (Brand, 1990) (Fig. 6). If a compound inhibits the respiratory chain, this curve will be displaced downward and to the left (Brand, 1990). Fig. 6 shows ΔΨ changes versus the respiration rate obtained in the absence (control) and in the presence of several concentrations of OHTAM (20, 40 and 60 nmol OHTAM/mg protein). At any given value of ΔΨ, OHTAM induces a slight decrease in the respiration rate. The displacement of the curve induced by OHTAM indicates that one or more components of the Δρ generating system are slightly inhibited by this drug, in contrast with the severe effects induced by TAM on the respiratory chain (Cardoso et al., 2001).

The putative proton leaks induced by OHTAM through the mitochondrial inner membrane were investigated in nonphosphorylating mitochondria titrated with malonate, a respiratory inhibitor, and ΔΨ was plotted versus respiration rate (Fig. 7). In the steady state, the proton efflux must equal the proton leak, assuming that no slip in the proton pump occurs (Murphy, 1989) and any secondary effects on the leak due to alterations in the Δρ value were eliminated. When a compound increases the proton leak across the mitochondrial

![Figure 2](image-url)
inner membrane, the curve will be displaced downward and to the right (Brand, 1990). Accordingly, OHTAM increases the proton leak across the mitochondrial inner membrane (Fig. 7) since at any given DC value the respiration rate is greater in the presence of 20 (b), 40 (c) and 50 (d) nmol OHTAM/mg protein or 50 nmol TAM/mg protein (e). The reactions were started by adding 5 mM succinate and were monitored by following the consumption of protons with a pH electrode. The traces are typical of four different experiments.

Fig. 3. Effects of OHTAM and TAM on pH changes associated with ATP synthesis. The ATP synthesis was evaluated at 25 °C in 2 ml of reaction medium containing 130 mM sucrose, 50 mM KCl, 2.5 mM MgCl₂, 5 mM KH₂PO₄, 0.5 mM HEPES, 0.1 mM EGTA (pH 7.2), supplemented with 2 μM rotenone and 200 nmol ADP. Mitochondria (1 mg) were incubated for 3 min in the absence (a) and presence of 20 (b), 40 (c) and 50 (d) nmol OHTAM/mg protein or 50 nmol TAM/mg protein (e). The reactions were started by adding 5 mM succinate and were monitored by following the consumption of protons with a pH electrode. The traces are typical of four different experiments.

Fig. 4. Effect of sequential additions of OHTAM on ΔΨ (○) and state 4 respiration (●) in non-phosphorylating mitochondria. Mitochondria (1 mg) were suspended in 1 ml of the standard respiratory medium supplemented with nigericin (50 ng/mg protein), oligomycin (1 μg/ml), 2 μM rotenone and 4 μM TPP⁺ at 25 °C and were energized with 5 mM succinate. Mitochondrial respiration rates and ΔΨ were evaluated simultaneously using a Clark-type electrode and a TPP⁺ selective electrode, respectively, placed in the same reaction chamber. After steady-state distribution of TPP⁺, the mitochondrial suspension was titrated with sequential additions of 10 nmol OHTAM/mg protein. The results are means ± SD of three separate experiments with different mitochondrial preparations. Actually, in the presence of 20 nmol TAM/mg protein, the respiratory rate of mitochondria to sustain −190 mV is about 32 nmol O₂/min/mg protein, a value about 3 times higher than that observed at the same concentration of OHTAM and similar to that induced by 60 nmol OHTAM/mg protein (Fig. 7).

In order to clarify whether the proton permeabilization is a consequence of either mitochondrial membrane disruption or proton shuttling, the effects of OHTAM on the integrity of the mitochondrial membrane of intact non-energized mitochondria were evaluated. OHTAM causes a
small dose-dependent decrease in the light scattering of non-energized mitochondria in suspension (Fig. 8), reflecting a slight direct effect of this drug on mitochondrial membrane integrity. Such effect becomes more evident at OHTAM proportions above 40 nmol/mg mitochondrial protein, suggesting that this drug only at high concentrations may interfere with the integrity of mitochondrial membranes, perturbing their permeability. However, these effects are not significant relatively to those induced by TAM, which at 50 nmol/mg protein causes an extensive swelling of mitochondria. The assessment of mitochondrial integrity was further confirmed by using the accessibility of citrate synthase from matrix to oxaloacetic acid through the mitochondrial membrane (Fig. 9). According to that observed in mitochondrial swelling studies (Fig. 8), increasing concentrations of OHTAM induce relative slight effects on mitochondrial integrity, since proportions of 10, 25 and 50 nmol/mg protein only affect 13, 17 and 40% of mitochondria, respectively. In contrast, TAM at concentrations of 25 and 50 nmol/mg protein disrupts 66 and 80% of the mitochondrial population, respectively, pointing out that this anticancer drug induces a much stronger disruptive effect on the mitochondrial membrane structure (Fig. 9).

The concentrations of OHTAM usually reached in blood plasma after therapeutic administration are about 3–5 ng/ml (Lien et al., 1987; McCague et al., 1990), although recently it has been reported that OHTAM attained in rat serum a C_max of 385 ± 132 ng/ml (0.9 μM) after s.c. administration of a second dose of TAM at 1 mg/kg (Mandlekar et al., 2000). Owing to its high hydrophobic character and partition into biological membranes (Custódio et al., 1991), the levels of OHTAM in humans are 10- to 60-fold higher in tissues than in plasma (Lien et al., 1991). Therefore, the therapeutics with OHTAM may cause tissue concentra-

**Fig. 5.** Comparison of OHTAM and TAM effects on ATPase of intact mitochondria. The ATPase activity was evaluated at 25 °C in 2 ml of reaction medium containing 130 mM sucrose, 50 mM KCl, 2.5 mM MgCl₂, 5 mM KH₂PO₄, 0.5 mM HEPES, 0.1 mM EGTA (pH 7.2) and supplemented with 2 μM rotenone. After pre-incubation of mitochondria (1 mg) for 3 min with different concentrations of OHTAM or TAM (nmol/mg protein), as indicated by the values adjacent to the traces, the reactions were started by adding 2 mM Mg-ATP and were monitored by following the production of protons with a pH electrode. Oligomycin (1 μg/mg protein) was added 1.5 min after Mg-ATP addition. The traces are typical of three different experiments.

**Fig. 6.** Effect of OHTAM on the kinetic response of Δρ generators to Δp. Nonphosphorylating mitochondria (1 mg) suspended in 1 ml of the standard respiratory medium supplemented with nigericin (50 ng/mg protein), oligomycin (1 μg/ml), 2 μM rotenone and 4 μM TPP⁺ were incubated at 25 °C for 3 min in the absence (○) and presence of 20 (□), 40 (■) and 60 (●) nmol OHTAM/mg protein. Mitochondrial respiration and Δp were measured simultaneously with a Clark-type O₂ electrode and a TPP⁺ sensitive electrode, respectively, and were placed in the same closed reaction chamber. The reactions were started with 5 mM succinate and after a steady-state distribution of TPP⁺, mitochondrial suspensions were titrated by sequential additions of FCCP (10 nM). The results are typical of six different experiments performed with three different mitochondrial preparations.
tions similar to those observed for TAM (≈ 50 μM) since the partitioning of OHTAM into membranes is significantly higher than that of TAM (Custódio et al., 1991). However, it is worthy of notice that, in this study we have also used higher concentrations only to compare the effects of this metabolite with those previously reported for TAM (Cardoso et al., 2001).

4. Discussion

Mitochondria serve as sensors of the xenobiotic-induced toxicity that ultimately leads to cellular disassembly (Wallace and Starkov, 2000). Recently, we have reported that TAM-induced alteration of mitochondrial bioenergetic capacity may be involved in its antiproliferative ER-independent activity and side effects on different cells types (Cardoso et al., 2001).

In the present study, we evaluated the effects of OHTAM, recognized to be more active and possessing several hundred times more affinity to the ER than TAM, on mitochondrial bioenergetic functions. As reported for TAM, OHTAM depresses the oxidative phosphorylation capacity of liver mitochondria in a concentration-dependent way, as deduced from a decrease of respiratory coefficient and of the ADP/O ratio (Fig. 1), but TAM has expressed much more potent effects (Cardoso et al., 2001). Moreover, OHTAM slightly depresses the mitochondrial ΔΨ (Fig. 2) and the state 3 respiration (Figs. 1 and 2) but induces stimulation of state 4 respiration (Fig. 4) as a function of concentration that also reflect a partial impairment of the phosphorylation efficiency of mitochondria. However, as opposed to TAM that promotes a strong inhibition of the oxidative phosphorylation of added ADP, as shown in Fig. 3 and also according to our previous results (Cardoso et al., 2001), the phosphorylation efficiency of mitochondria is not significantly affected by OHTAM (Fig. 1). This is also inferred from the fluctuations associated with the ADP-induced phosphorylative cycle (Fig. 2) and from the effects on pH changes associated with ATP synthesis (Fig. 3).
The OHTAM-induced increase in the respiratory rate in state 4 that parallels the partial collapse of the DC (Fig. 4) and the decrease of the ADP phosphorylation indicate an uncoupling effect of this drug on the oxidative phosphorylation system. The observed partial uncoupling effect observed is probably due to a slight permeabilization of the mitochondrial inner membrane to protons as confirmed by the Brand approach (Fig. 7). This effect could be related with a protonophoretic action and/or a nonspecific membrane permeabilization induced by this tertiary-amine antiestrogen (Murphy, 1989). In fact, a proton shuttle mechanism has been proposed to explain the ability of some tertiary-amine local anesthetics and other amine compounds to uncouple oxidation from phosphorylation (Garlid and Nakashima, 1983). Similarly, this mechanism could also explain the uncoupling effect of OHTAM, since the amine group of this drug can bind and translocate protons across the mitochondrial inner membrane due to its high lipophilic character and strong partition into biomembranes (Custódio et al., 1991). Moreover, the charged protonated form of OHTAM, preferentially located in the outer regions of the bilayer (Custódio et al., 1993), could be membrane-permeable and determine the increase in proton leak, acting in low concentrations like a classic protonophore. In addition to its uncoupling effect, the ionizable group of OHTAM, by changing the membrane surface potential, may induce a nonspecific inhibition of the phosphorylation system and a specific inhibition of either the adenine nucleotide translocator or the phosphate transporter. Further studies, currently in progress, are obviously required to assess these hypotheses focused on the membrane transporters. Both suggestions could explain the decrease in the oxidative phosphorylation capacity and the partial collapse of the ΔΨ due to proton leak that parallels the stimulation of state 4 respiration rate (Fig. 4), as previously reported for several lipophilic compounds possessing base- or acid-dissociative groups (Garlid and Nakashima, 1983; Fromenty et al., 1990; Keller et al., 1992). However, it cannot be excluded that the uncoupling action of OHTAM, at high concentrations, might be due to a nonspecific increase in the membrane permeability as a consequence of the mitochondrial membrane disruption.

To assess the relationship between the proton leak and ultrastructural changes induced by OHTAM, the stimulated ATPase activity was measured in intact mitochondria pre-incubated with this drug (Fig. 5). Furthermore, the OHTAM effects on mitochondrial swelling (Fig. 8) and on the citrate synthase activity of intact mitochondria (Fig. 9) were also evaluated. OHTAM at high concentrations stimulates ATPase activity of intact mitochondria, but to a smaller degree than FCCP or TAM (Fig. 5) which induces membrane disruption as previously described (Chen et al., 1999; Cruz Silva et al., 2000; Cardoso et al., 2001). Moreover, the decrease in light scattering of non-energized mitochondria (Fig. 8) and the stimulation of citrate synthase (Fig. 9) induced by OHTAM, together with the previous data demonstrating the release of Ca^{2+} by SR (Custódio et al.,

![Fig. 9. Comparison of OHTAM and TAM effects on citrate synthase activity of intact mitochondria. The citrate synthase activity was measured in 2 ml of reaction medium containing 100 mM Tris (pH 8) and supplemented with 200 μM 5,5’-dithiobis(2-nitrobenzoic acid) and 200 μM acetylCoA. After incubation of mitochondria (100 μg) at 30 °C for 3 min in the absence (a) and presence of either 10 (b), 25 (c), 50 (d) nmol OHTAM/ mg protein, or 25 (e) and 50 (f) nmol TAM/mg protein or 0.1% Triton (g), the reactions were started by adding 100 μM oxaloacetic acid. The citrate synthase activity expressed in μmol/min/mg protein, as indicated by the values adjacent to the traces, was followed by the increase in absorbance at 412 nm as function of time. The recordings are representative of four different mitochondrial preparations.](image-url)
and of K⁺ from erythrocytes (Cruz Silva et al., 2001), point to the membrane permeabilization ability of this drug. Therefore, at low concentrations OHTAM may promote membrane permeabilization to protons by acting as a proton shuttle, whereas TAM has showed to induce a much more potent proton leak (Fig. 7). At high concentrations, OHTAM partially uncouples oxidative phosphorylation due to a slight permeabilization related with the disruption of the structural integrity of the mitochondrial inner membrane. Besides the increase in the membrane permeability to protons, the inhibition of respiration induced by this drug at high concentrations (Fig. 6) may contribute to the understanding of the mechanism underlying its depressive effect on ΔΨ.

In conclusion, our results suggest that OHTAM slightly affects the mitochondrial bioenergetic functions, inducing a partial uncoupling of the oxidative phosphorylation system, in contrast with the deleterious effects of TAM in rat liver mitochondria (Cardoso et al., 2001). The low depressive effect of OHTAM on ΔΨ may activate the apoptotic machinery of cytochrome c release, leading to induction of caspase-dependent apoptosis. Moreover, these data indicate that OHTAM is much less toxic to mitochondria than TAM, in agreement with the different effects of both drugs in human erythrocytes (Cruz Silva et al., 2000, 2001) and with the different apoptotic effects in p53(−) human mammary epithelial cells (Dietze et al., 2001).

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References


