Comparison of the changes in adenine nucleotides of rat liver mitochondria induced by tamoxifen and 4-hydroxytamoxifen

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Abstract

The antiestrogen tamoxifen (TAM) inhibits the growth of different estrogen receptor (ER)-negative cells. Recently, multiple effects of TAM on mitochondrial bioenergetic functions have been pointed to explain its ER-independent cell death mechanisms. We have shown that TAM and its major active metabolite 4-hydroxytamoxifen (OHTAM) induce depolarization of the mitochondrial membrane potential (ΔΨ) and uncouple the mitochondrial respiration, depressing the oxidative phosphorylation efficiency. To clarify the biochemical mechanisms underlying the changes in the regulation of ATP synthesis and yield, in this work we evaluated the alterations of mitochondrial adenine nucleotides induced by both drugs and ascertained whether such changes could reflect a specific inhibition of either the adenine nucleotide translocase (ANT) or the phosphate carrier, as well as the activation of ATP hydrolysis due to ΔΨ depolarization. We found that both antiestrogens caused a concentration-dependent decrease in mitochondrial ATP levels. Mitochondrial ADP and AMP were concomitantly increased with a subsequent decrease in the ATP/ADP or ATP/AMPratios. The total concentration of adenine nucleotides also changed. Additionally, both drugs decreased the ANT content of mitochondria, inhibited the phosphate carrier and induced ATP hydrolysis. However, the effects of TAM were more drastic than those induced by OHTAM. Therefore, the depletion of ATP might result from an activation of ATP catabolism, as well as from a decrease in the mitochondrial content of ANT and partial inhibition of the phosphate carrier. Our data may explain the ER-independent effects and cytotoxicity of both drugs and, in agreement with other previous studies, suggest that OHTAM is much less toxic to mitochondria than TAM.

Keywords: 4-Hydroxytamoxifen; Mitochondrial energetics; Mitochondrial adenine nucleotides; Phosphate carrier; Adenine nucleotide translocase; Tamoxifen

1. Introduction

The antiestrogen tamoxifen (TAM) is widely used in the treatment of breast cancer and has been recognized as a cancer chemopreventive agent in high-risk women (Neven and Vernaeve, 2000; Radmacher and Simon, 2000). It is extensively metabolized in humans being 4-hydroxytamoxifen (OHTAM) its major active metabolite, which has been found in high levels in both the plasma and the tumor tissues (Daniel et al., 1984; Lien et al., 1991). This metabolite has been reported to possess several hundred times more affinity toward estrogen receptor (ER) as compared with TAM (Borgna and Rochefort, 1981). Thus, OHTAM is accepted as a more active drug than its precursor in retarding the estrogen-dependent cell growth and proliferation of ER-positive mammary tumours (Jordan et al., 1988; Osborne et al., 1992). However, large clinical trials have indicated that a favourable response to adjuvant therapy with TAM is also observed in different cell types that lack the ER (Couldwell et al., 1993; Croxton et al., 1994; Charlier et al., 1995), thereby suggesting that, at least in part, its
antineoplastic action may not be related to the anti-
hormone properties of these drugs. Also, there are
clinically significant ER-independent TAM actions for
which the biochemical mechanisms still remain obscure.

Mitochondria, which dominate the intracellular scenery
of most animal cells as their total number per cell
usually far exceeds that of any organelle, are the site of
most ATP cell production (Pederson, 1999), that is
essential to maintain and regulate several physiological
functions (Gupta et al., 2001 and references therein).
Therefore, the amount of ATP available for the cell can
in turn be considered as one of the key events in pre-
serving cell integrity. Thus, chemicals causing mito-
chondrial damage and inducing depletion of ATP may
in turn cause cell dysfunction or even cell death.

Recently we have shown that TAM depresses the
oxidative phosphorylation efficiency and uncouples the
mitochondrial respiration (Cardoso et al., 2001) and
that OHTAM affects the phosphorylation capacity of
mitochondria in a slight way as compared with the
damaging effects of TAM (Cardoso et al., 2002). These
effects are essentially a consequence of the membrane
potential (ΔΨ) depolarization due to an increase in the
proton leak through the mitochondrial inner membrane
(Cardoso et al., 2002). Therefore, the cytotoxicity of
these drugs is associated with ATP depletion via
impairment of mitochondria function related to ΔΨ
and/or oxidative phosphorylation. However, although the ATPsynthase couples the transport of the protons
across the membrane to the synthesis of ATP inside the
mitochondrial matrix, the phosphate for the phosphor-
ylation is imported into the mitochondria by the phos-
phate carrier and the ATP is exported to the cytosol in
exchange for ADP by the adenine nucleotide translocase
(ANT).

Since the TAM action on mitochondrial phosphoryl-
ation is not completely understood and its OHTAM
metabolite has been shown to possess growth inhibitory
activity, the aim of this work was mainly to evaluate the
alterations of mitochondrial energetic charge induced
by both drugs and ascertain whether such changes could
reflect a specific inhibition of either the ANT or the
phosphate carrier, as well as the activation of ATP
hydrolysis.

2. Materials and methods

2.1. Chemicals

Bovine serum albumin (BSA), 4-(2-hydroxymethyl)-1-
-piperazineethanesulfonic acid (HEPES), ethylene gly-
col-bis-(β-amoethylether) N,N,N′,N′-tetraacetic acid
(EGTA), 5, 5′-dithio-bis(2-nitrobenzoic acid), N-ethyl-
maleimide, oligomycin, 4-Hydroxytamoxifen (OHTAM),
tamoxifen (TAM) and ATP, ADP and AMP for HPLC
analysis were obtained from Sigma Chemical Co (St.
Louis, MO). Carbonyl cyanide p-trifluoromethoxyphenyl-
hydrazone (FCCP) and sucrose were obtained from
Merck (Dramstadt, Germany). All other chemicals were
commercial products of the highest purity grade avail-
able 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 (Car-
doso et al., 2001), with slight modifications. Animals
were killed by cervical dislocation 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 5000 g for 10 min at 4 °C
and mitochondria were recovered from the supernatant
by centrifugation at 10000 g for 10 min. The mitochon-
drial pellet was resuspended twice in the washing med-
tum 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 respiration

The oxygen consumption 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 and TAM in ethanolic solutions (up
to 6 μl) were incubated for 3 min at 25 °C in 1 ml of the
standard respiratory medium consisting of 130 mM
succrose, 50 mM KCl, 2.5 mM MgCl2, 2.5 mM KH2PO4,
5 mM HEPES, 0.1 mM EGTA, pH 7.2, previously
supplemented with mitochondria (1 mg) and 4 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 cal-
culated assuming an oxygen concentration of 240 nmol
O2/ml in the experimental medium at 25 °C.

For the titration experiments with carboxyatractilo-
side (CATR) the respiration of isolated mitochondria
(0.5 mg/ml) was determined at 25 °C using also a Clark
oxygen electrode. The respiratory medium contained
200 mM sucrose, 5 mM succinate (sodium salt), 5 μM
rotenone, 5 mM KH2PO4 and 20 mM Tris–HCl (pH
7.4). The ANT content was determined by titrating the
rate of state 3 respiration with increasing concentra-
tions of CATR (Vignais, 1976). Mitochondria were pre-incu-
bated with CATR in the respiratory medium for 1 min
before ADP (0.1 mM) addition to initiate state 3
respiration. Plots of O\textsubscript{2} consumption versus CATR appeared biphasic, with an increasing inhibitory segment followed by a steady respiration indicating that state 3 respiration was completely inhibited. The mitochondrial content of ANT was determined by the amount of CATR required to completely inhibit state 3 respiration, assuming a 1:1 binding stoichiometry, which was expressed as CATR content per mg mitochondrial protein (Roussel et al., 2000; Zhou et al., 2001).

2.4. Enzymatic activities

ATP-synthase activity was determined by measuring the pH changes of the medium associated with ATP synthesis as reported by Moreno and Madeira (1991). The reactions were carried out at 25 \degree C in 2 ml of reaction medium containing 130 mM sucrose, 50 mM KCl, 2.5 mM MgCl\textsubscript{2}, 5 mM KH\textsubscript{2}PO\textsubscript{4}, 0.5 mM HEPES, 0.1 mM EGTA, pH 7.2, supplemented with 2 \muM rotenone, 150 \muM ADP and mitochondria (1 mg). The experiments were initiated by the addition of 5 mM succinate and TAM and OHTAM were added to induce hydrolysis of the ATP produced following ADP phosphorylation by the mitochondria. ATPase activity was estimated by monitoring the pH changes of the same medium associated with ATP hydrolysis (Madeira et al., 1974).

2.5. HPLC determination of adenylates

Adenine nucleotides (ATP, ADP and AMP) were extracted using an acidic extraction procedure and separated by reverse-phase HPLC as described by Stochi et al. (1985). Briefly, the acidic extraction was performed as follows: 1 mg mitochondria protein was incubated in 1 ml standard respiratory medium containing 130 mM sucrose, 50 mM KCl, 2.5 mM MgCl\textsubscript{2}, 2.5 mM KH\textsubscript{2}PO\textsubscript{4}, 5 mM HEPES, 0.1 mM EGTA, pH 7.2, supplemented with 4 \muM rotenone. TAM or OHTAM were added at the concentrations of 20, 40 and 60 nmol/mg protein. A control assay in the absence of any of those drugs, was also carried out. After the incubation period (3 min), 5 mM succinate was added to induce state 4 respiration and, 2 min later, 150 nmol ADP were added to the reaction vessel to induce state 3 respiration. After 1 min of reaction, 250 \mul of the suspension were added to 250 \mul ice–cooled HClO\textsubscript{4} 0.6 M (EDTA–Na\textsubscript{2} 25 mM). After vortexing, this mixture was centrifuged at 10000 rpm for 2 min at 4 \degree C in an Eppendorf table centrifuge (UNIEQUIP UEC 14/B) and the pellet was discarded. The supernatant was neutralized with KOH 3 M (Tris 1.5 M) and centrifuged again (10000 rpm, 2 min) to precipitate all the KClO\textsubscript{4} produced. The supernatant was then stored at −80 \degree C for further chromatographic analysis. All the extraction procedure was carried out at 0−4 \degree C to minimize degradation of nucleotides. The chromatographic apparatus was a Beckman System Gold, consisting of a binary pump (model 126) and a variable UV detector (model 166), controlled by a computer. The detection wavelength was 254 nm and the column was a Lichrospher 100 RP-18 (5 \muM) from Merck (Darmstadt, Germany). An isocratic elution with 100 mmol/l phosphate buffer (KH\textsubscript{2}PO\textsubscript{4}) (pH 6.5) and 1.2% methanol was performed with a flow rate of 1 ml/min. The time required for each analysis was about 5 min.

2.6. Mitochondrial swelling

Mitochondrial osmotic volume changes were measured by the apparent absorbance changes at 540 nm, with a Perkin-Elmer Lambda 6 UV/vis spectrophotometer computer controlled (Norwalk, USA). The reactions were carried out at 25 \degree C in 2.5 ml of the required isoosmotic media: 135 mM ammonium phosphate, 5 mM HEPES, 0.1 mM EDTA, 2 \muM rotenone (pH 7.2).

3. Results

To assess whether TAM or OHTAM affect the mitochondrial bioenergetic charge, we determined the ATP, ADP and AMP levels in rat liver mitochondria under the action of different concentrations of both drugs. As shown in Table 1, the pre-incubation of mitochondria with TAM or OHTAM for 3 min, before energization with succinate, causes a concentration-dependent decrease in ATP levels, with concomitant increases in levels of ADP and AMP in the mitochondrial population. The total concentration of adenine nucleotides also changes. Moreover, both drugs decrease the ATP/ADP and ATP/AMP concentration ratios as a consequence of

<table>
<thead>
<tr>
<th>Table 1</th>
<th>Changes in the concentrations of adenine nucleotides in rat liver mitochondria induced by different concentrations of TAM and OHTAM\textsuperscript{a}</th>
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<tr>
<td></td>
<td>ATP</td>
</tr>
<tr>
<td>Control</td>
<td>189.90±7.07</td>
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<tr>
<td>TAM 20</td>
<td>139.19±1.83</td>
</tr>
<tr>
<td>TAM 40</td>
<td>81.95±3.07</td>
</tr>
<tr>
<td>TAM 60</td>
<td>55.97±5.59</td>
</tr>
<tr>
<td>OHTAM 20</td>
<td>166.72±3.82</td>
</tr>
<tr>
<td>OHTAM 40</td>
<td>118.77±4.18</td>
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<tr>
<td>OHTAM 60</td>
<td>73.92±1.55</td>
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\textsuperscript{a} Mitochondria (1 mg) in 1 ml of the standard respiratory medium were previously incubated with different concentrations of TAM or OHTAM (20, 40 and 60 nmol/mg protein) for 3 min and adenine nucleotides concentrations were determined by HPLC, as described in Section 2. Each value (nmol/mg protein) represents the mean±S.D. of at least three independent determinations.
ATP levels depletion and increases in ADP and AMP levels. These effects are more pronounced with TAM as compared to OHTAM, according to other observed effects on mitochondrial bioenergetic functions (Cardoso et al., 2001, 2002).

Our recent reports pointing to a decrease in the oxidative phosphorylation capacity of mitochondria treated with either TAM or OHTAM, as a consequence of proton leak across the mitochondrial inner membrane (Cardoso et al., 2001, 2002), led us also to investigate whether the ANT and/or the phosphate carrier are involved in changes of oxidative phosphorylation efficiency induced by both drugs. Thus, to assess the involvement of ANT, titrations with CATR, an inhibitor of the mitochondrial ANT, were performed. Rat liver mitochondria were pre-incubated with TAM or OHTAM (20 and 40 nmol/mg protein) and the results are shown in Figs. 1 and 2. In both cases, either with TAM (Fig. 1A) or with OHTAM (Fig. 1B), the ANT sites were blocked by lower concentrations of CATR than in the control assay without drugs, suggesting that both drugs could affect the phosphorylation efficiency by interference with the ANT.

CATR reacts with the ANT in a 1:1 stoichiometry (Vignais, 1976; Klingenberg et al., 1975; Zhou et al., 2001). Accordingly, the ANT content in mitochondria can be estimated by titrating the active-state respiration with CATR as illustrated in Fig. 1. The mitochondrial content of active ANT was determined by the amount
of CATR required to reduce state 3 respiration to the state 4 respiratory rate, when CATR presumably saturates the translocase sites (Vignais, 1976), and the results are shown in Fig. 2A and B. TAM decreases the ANT content of mitochondria in a higher extension than OHTAM in relation to the control (Fig. 2), i.e., mitochondria incubated without the drugs. In fact, OHTAM and TAM at concentrations of 20 nmol/mg protein induce decrease in the ANT content to about 81 and 64% of the control, respectively. Moreover, while mitochondria pre-incubated with 40 nmol OHTAM/mg protein still exhibit 67% of the ANT content, they only exhibit 46% of this at the same concentration of TAM. Such decrease in ANT content correlates with both the decrease in phosphorylation efficiency of mitochondria (Cardoso et al., 2002) and the changes in the ATP levels induced by both drugs (Table 1). Therefore, the mechanisms underlying the induced ATP depletion can in theory be caused either by inhibition of ATP synthesis due to ΔΨ depolarization or by changes in mitochondrial ANT content. However, either the putative inhibition of the phosphate carrier or the degradation of ATP to ADP and AMP can not be excluded.

In order to investigate the effect of both antiestrogens on mitochondrial phosphate transport, the permeability of the inner mitochondrial membrane in isoosmotic medium of ammonium phosphate was analysed by monitoring the mitochondrial swelling at 540 nm (Fig. 3). After suspension in isoosmotic ammonium phosphate, mitochondria swell spontaneously since, accompanying Pi accumulation, driven by its concentration gradient, there is one proton entry, as dictated by transport through the Pi carrier and matrix acidification takes place causing accumulation of matrix NH$_4^+$ after diffusion of NH$_3$ across the inner membrane (Klingenberg, 1970; Klingenberg et al., 1974) (Fig. 3—line a). Therefore, after NH$_3$ permeation through the mitochondrial membrane, the resulting ammonium phosphate entry leads to the influx of water molecules with the consequent mitochondrial swelling. In agreement with this interpretation, any compound that affects the phosphate carrier induces changes in the rate of swelling of mitochondria in ammonium phosphate medium. In fact, in the presence of N-ethylmaleimide (Fig. 3—line f), an inhibitor of the phosphate carrier, the mitochondrial swelling in isoosmotic medium of ammonium phosphate is significantly decreased. In the presence of TAM (Fig. 3—lines d, e) and OHTAM (Fig. 3—lines b, c) a decrease occurs in mitochondrial swelling, in a concentration-dependent way, suggesting that both drugs cause a partial inhibition of the phosphate transport. This effect is higher for TAM according to its stronger inhibitory effect in phosphorylation efficiency, as compared with OHTAM (Cardoso et al., 2001, 2002). Thus, we can infer that the depression of ATP synthesis is partially related to the observed inhibition of mitochondrial phosphate transport. Therefore, we can conclude that the interference of TAM and OHTAM with oxidative phosphorylation results not only from the interference with ANT (Figs. 1 and 2) but

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**Fig. 3.** Effects of TAM and OHTAM on swelling of mitochondria, followed by monitoring the decrease in absorbance at 540 nm. Mitochondria (0.5 mg) were suspended in 2.5 ml of isoosmotic ammonium phosphate medium containing 5 mM HEPES, 0.1 mM EDTA, 135 mM ammonium phosphate, 2 μM rotenone, pH 7.2. The reactions were started at 25°C by the addition of mitochondria in the absence (a) and presence of 20 (b) and 40 nmol (c) OHTAM/mg protein or 20 (d) and 40 (e) nmol TAM/mg protein or 2 μmol N-ethylmaleimide/mg protein (f), added as indicated by the arrow. The traces are typical of four experiments with different mitochondrial preparations.

**Fig. 4.** Comparison of OHTAM and TAM effects on ATPase activity of intact mitochondria. The experiments were carried out at 25°C in 2 ml of reaction medium containing 130 mM sucrose, 50 mM KCl, 2.5 mM MgCl$_2$, 5 mM KH$_2$PO$_4$, 0.5 mM HEPES, 0.1 mM EGTA (pH 7.2), supplemented with 2 μM rotenone, 150 μM ADP and mitochondria protein (1 mg). The reactions were started by the addition of 5 mM succinate and were monitored by following the production of protons with a pH electrode. After phosphorylation of the added ADP, TAM (80 nmol/mg protein), OHTAM (80 nmol/mg protein), FCCP (4 nmol/mg protein) and Triton X-100 (0.1%) were added as indicated by the arrow. The traces are typical of three different experiments.
also from the partial inhibition of phosphate transport into the mitochondria (Fig. 3).

In order to verify whether the effect of TAM and OHTAM on mitochondrial ATP levels may also result from an increase in ATP catabolism to ADP and AMP, TAM and OHTAM were added to succinate-energized mitochondria after ADP phosphorylation and the H⁺ consumption and production were followed by using a pH electrode (Fig. 4). In fact, ADP phosphorylation by mitochondria after addition of succinate is accomplished by H⁺ removal from the medium associated with ATP synthesis (Fig. 4). The addition of TAM (80 nmol/mg protein) and OHTAM (80 nmol/mg protein), as indicated by the arrows (Fig. 4), stimulate ATP hydrolysis as proved by the release of H⁺ to the medium, suggesting that both drugs increase the ATPase activity of tightly coupled intact mitochondria. However, this effect induced by OHTAM is much smaller than those promoted by FCCP (4 nmol/mg protein) and TAM, which, like Triton (0.1%), induces disruption of the mitochondrial inner membrane integrity (Cardoso et al., 2001, 2002).

4. Discussion

The cytotoxic effects of TAM in estrogen-dependent breast cancer cells are mediated by high affinity binding to the ER (Coezy et al., 1982). Previous studies have reported that, in comparison with TAM, OHTAM has higher affinity and specificity for the ER (Borgna and Rochefort, 1981) and, besides being 100–200 times more potent (Osborne et al., 1992; Jordan et al., 1988), it is pharmacologically well tolerated (Sauvez et al., 1999). Although most studies of TAM have centred on its ability to compete with estrogens for binding to the ER and thereby antagonize their ability to induce conformational change in the ER and therefore activate transcription, several recent studies have described additional ER-independent activities of TAM (Perry et al., 1995; Fattman et al., 1998). In fact, TAM inhibits also the growth of other cell types including ER-negative breast cancer cells (Jordan, 1990), lung adenocarcinoma (Croxtall et al., 1994), prostate cancer (Bergan et al., 1999), ovarian carcinoma (Trope et al., 2000), virus (Laurence et al., 1990) and bacteria (Luxo et al., 1996). Furthermore, in spite of several often-conflicting studies showing that TAM and OHTAM can actively induce programmed cell death in cancer cells (Dietze et al., 2001; Mandlekar and Kong, 2001), it was reported that both drugs could induce ER-dependent and -independent programmed cell death (Obrero et al., 2002). However, the ER-independent action mechanisms of both drugs and their side effects have yet to be clarified.

Mitochondria of living cells participate in a number of different processes, among which is ATP synthesis by the oxidative phosphorylation system required to maintain the cell viability (Wallace and Starkov, 2000). ATP synthesis occurs in the inner mitochondrial membrane and requires the electron transport chain to generate the electrochemical proton gradient (ΔpH) consisting of a membrane potential (ΔΨ) and a pH gradient (ΔpH) (Brown, 1992). Moreover, this complex process also requires two key transport systems for the entry of ADP and inorganic phosphate into the matrix space and the ATP synthase complex. On the other hand, almost all intracellular ATP is generated in the mitochondria and about one-third of the cellular adenine nucleotides are located in these organelles (Pederson, 1999).

Recently, we have observed that TAM and, although in a lesser extension, OHTAM induce both ΔΨ depolarization, due to an increase in proton leak across the inner mitochondrial membrane, and uncoupling the mitochondrial respiratory chain, depressing the mitochondrial phosphorylative efficiency, as inferred from the changes in the phosphorylative cycle induced by ADP (Cardoso et al., 2002). However, it is noteworthy that the oxidative phosphorylation is seriously compromised at concentrations of TAM below those affecting the electron transfer along the respiratory chain. Therefore, in addition to the effects on redox chain and ΔΨ depolarization, these antiestrogens could affect the rate of ATP synthesis as a consequence of ATP hydrolysis stimulation and changes in either ANT or phosphate carrier that use the components of Δp.

HPLC analysis of adenine nucleotides revealed that indeed both antiestrogens decrease ATP levels, as well as ATP/ADP and ATP/AMP concentration ratios and increase ADP and AMP levels in the mitochondrial population (Table 1). However, the effects induced by TAM are more pronounced than those of OHTAM, in agreement with our previous results (Cardoso et al., 2002). In addition to the substantial alterations of mitochondrial energetic charge promoted by these drugs, the increase in the catabolism of ATP to ADP and AMP (Fig. 4) may be a key factor involved in the bioenergetic changes, rendering the mitochondria unable to fulfil the cell energy requirements. Thus, these effects reinforce that the hydrolysis of ATP (Fig. 4), in addition to the ΔΨ depolarization as consequence of permeability to protons due to destructive effects in the structural integrity of the mitochondrial inner membrane (Cardoso et al., 2001, 2002), may also account for the decrease in ATP levels induced by both drugs. However, this decrease can also result from the partial inhibition of both the ADP–ATP carrier (Figs. 1 and 2) and the phosphate transport into the mitochondria (Fig. 3). So, our data suggest that TAM and OHTAM, in addition to the respiratory chain inhibition and membrane proton leak with consequent depolarization of ΔΨ (Cardoso et al., 2002) disturb the mitochondrial energetic charge (Table 1), affect the mitochondrial...
transport of ADP (Figs. 1 and 2) and of Pi (Fig. 3), and stimulate the ATPase activity (Fig. 4). Taken together, these effects should be responsible for the observed decreased efficiency in the phosphorylative capacity of mitochondria induced by these drugs. It is noteworthy that in agreement with other previous studies, OHTAM is much less toxic to mitochondria than TAM.

In conclusion, the reduction in ATP levels induced by these drugs can dictate the future of the cell and the decision of the cell to undergo apoptosis and/or necrosis. Although these effects may not be specific for cancer cells, they may account for the antiproliferative ER-independent action mechanisms and for the cytotoxic effects of these drugs in different cell types.

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References


Madeira, V.M.C., Antunes-Madeira, M.C., Carvalho, A.P., 1974. Activation energies of the ATPase activity of sarcoplasmic reticulum. Biochemical and Biophysical Research Communications 58, 897–904.


