Tetraconazole-induced Programmed Cell Death in 

*Schizosaccharomyces pombe*

**ABSTRACT**

Tetraconazole, a systemic triazole fungicide, shows potential toxic effects in agriculture and human health. Therefore, its cytotoxic effects and accompanying mechanisms should be unraveled. *S. pombe* (ED666) was used in this study, as a unicellular biology and toxicology model. Cells were grown on standard media and all treatments were done at 30 °C and shaking at 180 rpm 1-10 mg/L tetraconazole induced a dose-dependent cell death. Apoptosis was monitored by DAPI and AO/EB staining. Excessive ROS production and mitochondrial impairment were shown by DCFDA/NBT assays and Rhodamine 123 staining, which were supported by increased expressions of superoxide dismutases and glutathione peroxidase. Involvement of one of the potential apoptotic genes, *Cnx1*, in apoptosis was shown by increased transcription whereas two other potential genes, *Pca1* and *Aif1*, were not affected by tetraconazole treatment. In conclusion, tetraconazole-induced cytotoxicity and underlying mechanisms which were mediated via ROS damage and mitochondrial dysregulation (*Cnx1*-driven) were clarified in *S. pombe*.

**Keywords** - *Schizosaccharomyces Pombe*, Tetraconazole, Apoptosis, *Cnx1*, Oxidative Damage

**ÖZ**

Sistemik triazol bir funisit olan tetrakonazol tarımda ve insan sağlığından potansiyel toksik etkileri göstermektedir. Bu yüzden, sitotoksik etkileri ve eşlik eden mekanizmaları açığa çıkarmak önemlidir. Çalışmamızda, tek hücreli biyoloji ve toksikoloji modeli olarak *S. pombe* (ED666) kullanılmıştır. Hücreler standart medyumda büyütülmüş, muameleler 30 °C ve 180 rpm hızda çalkalamalı olarak yapılmıştır. 1-10 mg/L tetrakonazol doz-bağımlı hücre ölümüne sebep olmuştur. Apoptoz DAPI ve AO/EB boyamasya görüntülendiştir. Aşırı ROS üretimi ve mitokondriyel bozulma DCFDA/NBT deneyleri ve Rhodamin 123 boynamasıyla gösterilmiştir, bu sonuçlar da süperoksitdismutazlar ve glutatyonperoksidad ifade eden diğer bringenleri de gösterilmiştir. Potansiyel apoptotik genlerden biri olan *Cnx1*’in apoptozla ilişkisi transkripsiyonunda artırılmış, diğer iki potansiyel gen, *Pca1* ve *Aif1* tetrakonazozi etkilenmemiştir. Sonuç olarak, tetrakonazol kaynaklı apoptoz ile, ROS hasarı ve mitokondriyel düzentsizliğinin (*Cnx1*-oluşlu) aracılık etmiş olduğu mekanizmalar *S. pombe*’de açıklığa kavuşturulmuştur.

**Anahtar Kelimeler** - Schizosaccharomyces Pombe, Tetraconazole, Apoptoz, *Cnx1*, Oksidatif Hasar
I. INTRODUCTION

Azoles are antifungal agents showing variation in activity, toxicity, and interaction potential [1]. While some azoles can potentially inhibit oxidative drug-metabolizing enzyme systems [2–4], others were shown to interact with transporter proteins [5]. Amongazole antifungal agents, tetraconazole is a systemic triazole fungicide, which is formed by a chiral carbon atom and two enantiomers and is widely used as an eradicant and protectant against fungal infections [6, 7]. Although little is known on the potentially toxic effects of tetraconazole in human health, researchers have identified its inhibitory effects related to its high affinity for fungal demethylases, particularly for fungal P450 enzymes [8–10] having roles in ergosterol biosynthesis in cell membranes [11].

The median lethal dose (LD₅₀) for tetraconazole was calculated as 1,030 mg/kg in rats [12], whereas 7-pentoxyresorufin O-depenthylase values were dramatically elevated [13]. Hepatocyte necrosis and inflammation along with benign and malignant liver cell tumors, bodyweight loss, a decrease of plasma glucose, and increased levels of blood nitrogen and alkaline phosphatase were found in rats received 640-2500 ppm tetraconazole [14]. In addition, moderate-to-severe inhibition of cell viability and proliferation in Chinese hamster ovary cell line (CHO) was reported when the cells were treated with tetraconazole for 72 h (the calculated %C1/2-50% decrease in cell density- was 39.2 µM) and/or with other azoles such as tebuconazole (%C1/2: 98.1 µM) and propiconazole (%C1/2: 44.0 µM) [15]. Besides, tetraconazole was considered highly toxic to fish and other aquatic organisms in which 96 h the lethal concentration 50 (LC₅₀) values were calculated between 2.5-4.3 mg/L in fish. Given the significance of azoles in the pharma industry, chemical industry, and medicine, possible cytotoxic, genotoxic, inhibitor, carcinogenic, and/or anti-cancer potentials are to be investigated. The study for anti-cancer potentials mainly focused on azole and triazole derivatives, such as 1, 2, 4-triazoles [16], ribavirin [17], and DAN94 [18]. On the other hand, there are a lot of research focused on azoles-fungi interactions. Candida species, which are well-known pathogenic fungi class, showed a range of minimum inhibitory concentration (MIC: 0.03-128 µg/mL) when a variety of azoles (fluconazole, itraconazole, voriconazole, amphotericin B, tetraconazole and tebuconazole) were administered [19]. As genetic tools and its manipulation is limited in pathogenic fungi, Saccharomyces cerevisiae, which was known to have ScErg5 gene (the target of azoles), was used to monitor cellular metabolism of antifungal drugs, particularly azoles and derivatives [20]. 19-104 µM MICs were reported when general antifungal azoles, fluconazole and miconazole, were administered to S. cerevisiae [21]. However, a complete understanding of azoles, in particular, tetraconazole toxicity, and regulation of cell death in the fission yeast (Schizosaccharomyces pombe), which has many advantages and is more similar to higher eukaryotes in some issues, are currently unknown and waiting to be explained.

Analogous cell cycle control [22], mitochondrial biogenesis [23], and programmed cell death subroutines [24–26], in addition, a small genome that can be easily manipulated [27] present Schizosaccharomyces pombe as a well-established model cell for molecular toxicology, cell biology and biochemistry studies [28,29]. Besides, the Warburg effect, which refers to the reprogrammed energy metabolism in cancer cells, resembling the energy metabolism of highly proliferative yeast cells, constitutes a valuable opportunity in cancer research [30–32]. Consequently, the underlying mechanisms for cancer therapeutic candidates can be understood by using fission yeast as a eukaryotic unicellular cell death model [33–35].

There are several programmed cell death subroutines were found in the fission yeast: Apoptosis, necroptosis, and autophagy. Numerous research focused on fission yeast apoptosis and necrosis. However, cellular physiology of yeast apoptosis is not fully understood and waiting to be unraveled [36]. Whereas oxidative stress [25], mitochondrial impairment and lipotoxicity are well-known stimulations for the positive regulation of apoptosis in yeast, we do not know much about the dark side of the genetic regulation[37]. Indeed, many genes (caspases and endonucleases) having potential roles in apoptosis of Baker’s yeast (S. cerevisiae) were investigated [38] however, the orthologous genes (at least several of them) in the fission yeast (S. pombe) are waiting to be enlightened.

This study aims to evaluate the cytotoxicity of tetraconazole and its mechanism and to show the potential of S. pombe for cell biology, toxicology, and cancer research. Mortality, proliferation, and apoptosis along with candidate genes for apoptosis were assessed. Besides, oxidative stress, DNA damage, and mitochondrial damage were evaluated as accounting mechanisms. This study contributes to literature for tetraconazole toxicity and evaluation of its medicinal potential.
II. MATERIALS AND METHODS

A. Reagents

Media components, methylene blue, arsenic(III) oxide, hydrogen peroxide, DCFDA, Rhodamine 123, NBT, and DAPI were purchased from Sigma (Istanbul, Turkey). Tetraconazole (dichlorophenyl)-tetrafluoroethoxy)propyl]-1,2,4-triazole was a kind gift from E. Yoruk (Istanbul Yeni Yuzyl University) (Sigma, U.S.A.).

B. Yeast, Media, and Growth Conditions

*S. pombe* wild-type strain ED6666 was a kind gift from B. Palabiyik (Istanbul University). Yeast was grown in YEL media containing yeast extract and glucose on a shaker at 150 rpm. at 30 °C in all experiments. 1x10^6 cells/ml cultures were used for experiments.

C. Tetraconazole Exposure and Cytotoxicity

Yeast culture (OD_600 ~1, in YEL media) from overnight incubation (16 h) were counted using Carl-Zeiss, Axio Observer 3 microscope followed by adjusting to 1x10^6 cells/ml in conical flasks. Tetraconazole (TCN) solution was prepared in ethanol. Cells were exposed to different concentrations of tetraconazole (0-10 mg/L in ethanol) and solvent control for 24 h. Arsenic(III) oxide was positive control of apoptosis[39]. Hemocytometer was used for the evaluation of relative cell proliferation. Cells were prepared in PBS and were stained with methylene blue. Cells were observed under a light microscope from 200-250 cells in each biological replicate (n=5).

D. Detection of Apoptosis by AO/EB and DAPI staining

1 μg/ml DAPI was used to monitor the cell nucleus as previously reported[40]. Carl-Zeiss, Axio Observer 3 fluorescent microscope (63x objectives) was used to observe cells at 358 nm and 461 nm. Besides, yeast was stained using AO/EB as previously described [38,39] and examined under Carl-Zeiss, Axio Observer 3 using 63x objectives at 500 nm and 530 nm for AO (acridine orange), and 510 nm and 595 nm for EB (ethidium bromide).

E. ROS detection by DCFDA staining and NBT assay

ROS levels were measured using DCFDA as declared previously [25,43] and examined under Carl-Zeiss, Axio Observer 3 fluorescent microscope (63x objectives) at 495 nm and 529 nm. NBT reduction experiment was done as described previously [43]. NBT was used at a concentration of 0.1 % and cells were incubated for 1 hour at 1x10^6 cells/ml concentration. Cells were washed and fixed using absolute methanol and diluted methanol (70%). 2 M KOH and DMSO were used to solubilize the dry pellet. The optical density of the final solution was calculated by microplate spectrophotometer (Thermo Scientific, Multiskan Go) at 620 nm.

F. Detection of MTP by Rhodamine 123 assay

Rhodamine 123 was used to stain mitochondria as indicated previously[44]. The reaction was performed in sodium citrate buffer containing glucose and rhodamine. Incubation was 15 min at room temperature. Cells were visualized by Carl-Zeiss, Axio Observer 3 fluorescent microscope (63x objectives) at 505 nm and 534 nm.

G. RNA Extraction and RT-PCR

Alteration in mRNA levels for *Sod1, Sod2, GPx1*, and pro-apoptotic genes was given in ESI.[†] Oligonucleotides were designed using Primer3Plus and given in Table S1.[†]

H. Statistical analysis

Experimental data were shown as mean ± standard error of the mean (SEM). One-way ANOVA and Tukey’s tests (Graphpad, USA) were used to understand the statistical difference.
III. RESULTS

A. Tetraconazole-Induced Cytotoxicity is Dose-dependent

Tetraconazole was previously shown to induce cell death in fungi [45]. We hypothesized that tetraconazole-induced cell death in our model fungi S. pombe was related to the apoptotic activity of this triazole. To test this hypothesis, we first analyzed cell growth using a hemocytometer, which was significantly inhibited \( (p < 0.01) \) after tetraconazole exposure between 2-20 mg/L (Figure 1A), whereas mortality notably and gradually increased \( (p < 0.01) \), which was evaluated using methylene blue assay. Methylene blue permeates dead cell membranes and stains cytoplasm and nucleus, in contrast, live cells having intact cell membranes are capable of blocking permeation. The calculated the half-maximum inhibitory concentration\( (IC_{50}) \) value was 7.03 mg/L.

Figure 1. Cell proliferation, viability, apoptosis, and nuclear morphology after exposure to tetraconazole solutions for 24 h: A. Cell proliferation and mortality were assessed by hemocytometer and by methylene blue assay in comparison to ethanol (solvent) control (0 mg/L refers to solvent control). Solvent control includes 0.05% ethanol. B-F, Viable and dead cells were shown via fluorescent microscope after treatment with 1-10 mg/L tetraconazole and arsenic. Arsenic(III) was apoptosis-positive control. Arrows indicate apoptotic yeast. G. The graph shows the percentage of apoptosis in yeast treated with 0-10 mg/L tetraconazole and arsenic \((n=5)\). Asterisks were used to show significantly different values \((**p<0.01, *** p<0.001)\). H, DAPI assay was performed after treatment with 1-10 mg/L tetraconazole and arsenic. Arrows: Anucleation or fragmentation.

B. Cell Death is Dependent on Apoptosis

We observed apoptotic cells (Figure 1B-F) by using AO/EB assay showing both dead and live cell membranes along with cell nuclei. Ethidium bromide stains only the dead cell nucleus, whereas acridine orange penetrates both live and dead cells and stains the nucleus. While regular green nucleus is showing live cell, fragmented orange-bright or red nucleus is indicating apoptotic cell. Cells were exposed to tetraconazole at 1, 2, and 10 mg/L doses for 24h. Apoptotic cells were marked by arrows in Figure 1B-F. Number of apoptotic cells was significantly different in 2 mg/L and 10 mg/L tetraconazole groups\( (p<0.01 \) and \( p<0.001) \) as illustrated in Figure 1G. Apoptosis dramatically increased at 2 and 10 mg/L doses \((%20.21 \) and \%56.79\) compared to the control group.
The percentages of apoptotic cells were comparable with mortality rates (%21.57 and %59.60) at 2 and 10 mg/L doses.

In addition, apoptosis was confirmed by DAPI staining. After the exposure period, cells were fixed with formaldehyde. When cells were stained with DAPI, we observed nuclear fragmentation, condensed, crescent- and dot-shaped cell nuclei, which were known as typical apoptotic markers [46, 47], at 2-10 mg/L doses and in arsenic control (apoptosis control) cells (Figure 1H, h1: 1 mg/L; h2: 2 mg/L; h3: 10 mg/L; h4: 3 mM arsenic trioxide).

**C. Tetraconazole-Induced ROS Production Potentially Activated Apoptosis**

DCFDA immediately reacts with ROS, mostly superoxide anions, and transforms to its oxidized fluorescent form (DCF) after permeation to the membrane and localization to the cytoplasm. As demonstrated in Figure 2, green fluorescence gradually increased in the experimental group at all concentrations of tetraconazole and in hydrogen peroxide control. In addition, using NBT reduction assay, levels of reactive oxygen species were measured and calculated as percentages of the control group. NBT reduction reflects ROS generation [48]. As shown in Figure 2F, NBT reduction significantly increased (2-5-fold) at all dose groups (p < 0.05 for 1 mg/L, p < 0.01 for 2 mg/L and 10 mg/L tetraconazole concentrations, which was consistent with increase in mortality (%21.57 and %59.60; see Figure 1A) and apoptosis (%20.21 and %56.79; see Figure 1G). As illustrated in Figure 2G-I, Sod1 and Gpx1 mRNA levels increased 1.5-2-fold at all concentrations of tetraconazole, whereas Sod2 mRNA levels remained unchanged except at 2 mg/L tetraconazole concentration. Positive regulation of the antioxidant enzyme system generally indicates excessive reactive oxygen species production [49], which means that organism is to recover antioxidant enzymes exhausted by superoxide and other oxidizing molecules [50].

**Figure 2. ROS levels and antioxidant gene expression: A-E.** ROS generation in yeast treated with 1-10 mg/L tetraconazole and 1 mM hydrogen peroxide were visualized and calculated using a fluorescence microscope. F. ROS production in yeast treated with 0-10 mg/L tetraconazole and 1 mM hydrogen peroxide was also measured by NBT assay. ROS production was determined at 620 nm and given as a percentage of a control group. G-I. Sod1 (G), Sod2 (H), and Gpx1 (I) expressions were analyzed using RT-PCR. Significance is marked by asterisks (*p<0.05, **p<0.01; n=3).
D. Loss of MTP Elucidate the Effect of Tetraconazole on Mitochondria

Disruption of mitochondria is an early apoptotic marker, which can be measured as transmembrane potential (ΔΨm) [51]. Loss of MTP was monitored using mitochondrial fluorescence stain Rhodamine 123. Active mitochondrion sequesters fluorescence stain via the activity of intact membrane pumps and emits green fluorescence, whereas apoptotic mitochondrion cannot pump the dye to the matrix, therefore the dye cannot fluoresce. The difference in fluorescence intensity of experimental and control groups was demonstrated in Figure 3. A dramatic decrease in fluorescence intensity was calculated at all dose groups. Intensities were dramatically lowered 2-3-fold at 1-10 mg/L tetraconazole concentrations (p<0.001) as well as in arsenic control (see Figure 7F).

**Figure 3.** Mitochondrial transmembrane potential (MTP) was demonstrated by Rhodamine 123 staining: A-E. MTPs treated with 1-10 mg/L tetraconazole and arsenic were visualized and measured. F. Decrease of fluorescent intensity in the cells treated with tetraconazole (0-10 mg/L) was measured and expressed as a percentage of a control group. Significance was indicated by asterisks (***p<0.001; n=3).

E. Candidate apoptosis-related genes showed different expressions

*Pca1*, *Cnx1*, and *Aif1* were reported to regulate and progress the apoptotic process in *S. pombe* [53–55]. Although the function of *Pca1* is currently not clear, we showed 2-fold decrease (p<0.05) in *Pca1* expression at 10 mg/L tetraconazole group (p<0.05, Figure 4). Another pro-apoptotic gene, *Cnx1* (Calnexin-1), which was previously reported to regulate ER-stress-induced cell death [53], was found transcriptionally activated (1.5-2-fold, p<0.05) in response to tetraconazole exposure in this study. However, *Aif1*, a FAD-dependent flavoenzyme homologous to mammalian *Aif1*, which is known to activate chromatin condensation and other apoptosis-related processes after its release from mitochondria [56], remained unchanged (p>0.05) after tetraconazole exposure.
In *S. pombe*, programmed cell death subroutines occur in several ways including apoptosis, autophagy, and necrosis, which are highly responsive to chemical and physical stressors, such as drugs, plant-originated chemicals, heat, and starvation [57]. In our research, fission yeast responded to tetraconazole, anazole derivative that is used as an antifungal agent, and cell proliferation and viability gradually decreased in a dose-dependent manner. Although limited data is known on tetraconazole toxicity, generally azoles are defined as N-demethylase inhibitors that can subsequently inhibit cell membrane biosynthesis [11]. However, cell death, induced by tetraconazole in this study, was found mainly dependent on apoptosis, which was demonstrated with mitochondrial impairment, ROS accumulation, DNA fragmentation, and condensed cell nuclei, while many other azoles were found to induce cell death via blocking glycolysis, calcium influx and membrane biosynthesis [58].

Mitochondrial impairment and oxidative stress following the elevation of ROS production are known to be deleterious, which are having the potential to start cell death signaling [59]. An increase in oxidative molecules, such as hydrogen peroxide, hydroxyl radical, and superoxide anion was detected by NBT and DCFDA assays in our study. Besides, our results demonstrated a dose-dependent increase in oxidation of cellular compartments which was supported by elevations in expressions of cytosolic (*Sod1*) and mitochondrial superoxide dismutases (*Sod2*), and also in glutathione peroxidase. Similarly, a recent study reported that tetraconazole-induced oxidative stress, which was shown by MDA levels, superoxide dismutase, and catalase activities, led to chromosomal aberrations accompanying cell death in *Allium cepa* [60]. However, we found a negative correlation between ROS levels (oxidative stress) and mitochondrial inner membrane potential indicating oxidative damage on a mitochondrial membrane (see Fig. 2 and Fig. 3F). Previous studies elucidated the disruptive effects of oxidative stress on mitochondrial membrane permeability and mitochondrial DNA damage [61–63].

There are several candidate genes, which are suggested to be responsible for apoptosis in *S. pombe*. The fission yeast genome encodes only one caspase, which is known as *Pca1*. It is believed that *Pca1* plays a dual role in cell death decisions [64]. Although in our previous study, we showed elevated *Pca1* mRNA levels after chemical stress [42], however, in this study, tetraconazole treatment did not cause alteration in *Pca1* levels except 10 mg/L, which can be accepted as an overdose for tetraconazole treatment. The precise roles of *Pca1* and its ortholog *Yca1* (in *S. cerevisiae*), which were suggested to be metacaspases, in regulated cell death, are still waiting to be enlightened [65]. Similarly, *Aif1* mRNA levels were not altered in any tetraconazole dose group in this study. In baker’s and fission yeasts, involvement of *Aif1* in apoptosis activated by chemical stress and acidic pH was shown by a few articles [66–68]. But, in contrast, tetraconazole did not show a similar effect on *Aif1* transcription in this study. Finally, *Cnx1*, which was suggested to regulate ER stress-related proteotoxicity and apoptosis in *S. pombe* [53], was demonstrated to have a potential role in tetraconazole-induced cell death in our study. Transcriptional activation of *Cnx1* after tetraconazole treatment supports our hypothesis of its involvement in the fission yeast cell death signaling. However, some reports are suggesting that *Cnx1* is involved in both autophagy and apoptosis or manage crosstalk between them [69].
V. CONCLUSION

In conclusion, tetraconazole caused dose-dependent apoptotic cell death. Cytotoxicity was related to oxidative damage and disruption of mitochondria. This study, also, examined three S. pombe genes that were believed to positively regulate apoptosis, and, consequently, induce cell death. Moreover, this study warrants further study aiming at programmed (or regulated) cell death pathways, lipotoxicity, and autophagy, or, cellular aging using molecular genetics insights.

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