Introduction

T-2 toxin, a fungal secondary metabolite, is one of the type A Trichothecenes [1,2]. Ingestion by humans or livestock of cereals contaminated by T-2 toxin can cause adverse reactions, such as vomit, diarrhea, and even death [3,4]. Alimentary toxic aleukia (ATA), mainly due to ingestion of cereal containing large amounts of T-2 toxin has been reported to cause the death of a large number of people [5]. Injection of large dose of T-2 toxin to rat caused cardiomyopathy, which was similar to the symptom of ATA [6,7]. In view of the great harm to the health of human and livestock, the toxicological effects of T-2 toxin was reported in the Joint Food and Agriculture Organization/World Health Organization (FAO/WHO) Expert Committee on Food Additives [4,6–8].

It was reported that T-2 toxin could affect protein synthesis by its affinity with trans-peptidase, one of the important subunits in ribosome, and the biosynthesis of DNA and RNA were also inhibited by T-2 toxin [9,10]. It was also found that T-2 toxin could interfere with the cytomembrane phosphorylation and cause lipid peroxidation in liver [11]. Islam et al. (1998) reported that the effect of T-2 toxin on mice thymocytes was apoptosis [12]. Shinozuka et al. (1997) also found that the T-2 toxin induced lymphocyte death was by apoptosis [13]. It was confirmed by in situ hybridization that the apoptotic process was accompanied by DNA damage [14]. Wang et al. (2012) further reported that JAK/STAT (Janus kinase/signal transducers and activators of transcription) might play an important role in the trichothecenes induced apoptosis [15]. T-2 toxin mainly acted on the metabolically active organs, such as spleen, thymus, marrow, stem cells, and so on [4,16,17]. The apoptotic process is initiated by a series of oxidative stress, and subsequently cells enter the mitochondrial death pathway [18,19]. It was reported that T-2 toxin not only could down-regulate intracellular reduced GSH (Glutathione), but also could up-regulate intracellular total ROS (Reactive oxygen species) [5], which showed intrinsic link between T-2 toxin-induced apoptosis and oxidative stress. But the mechanism of how oxidative stress induces apoptosis is still unclear.

JunD, a member of the AP-1 family of transcription factors, regulates genes involved in antioxidant defense. Gerald et al. (2004) found that JunD could reduce angiogenesis in tumor by reducing ROS, and demonstrated that JunD involved in regulation of antioxidant defense [20]. Toullec et al. (2010) took advantage of JunD deletion cell strain to examine the role of ROS in tumor development, and uncovered the role of JunD in the suppression of the migratory properties of stromal fibroblasts, which in turn potentiate tumor dissemination [21]. However, there are still not
any reports on the effect of JunD in the process of apoptosis induced by T-2 toxin.

In view of the harmful effects of T-2 toxin, this study focused on the mechanism of T-2 toxin-induced apoptosis, and on the role of oxidative stress, especially the function of JunD, in T-2 toxin induced apoptotic process.

Materials and Methods

Ethics Statement

All animal work was performed according to relevant national and international guidelines. All animal experiments were complied with the rules by the Animal Ethics Committee of the Fujian Agriculture and Forestry University.

Materials

T-2 toxin was purchased from Sigma Corporation (USA), and caspase-3 colorimetric assay kit and MTT (3-(4, 5-Dimethylthiazol-2-yl)-2, 5-diphenyltetrazolium) assay kit were from Nanjing Keygen Biotech Co. Ltd (China). Lipofectamine 2000, anti-caspase-3 antibody, anti-caspase-8 antibody, anti-caspase-9 antibody, anti-JunD antibody, GSH assay kit, and MDA (Malondialdehyde) assay kit were from Beyotime Institute of Biotechnology (China). The other chemical reagents used were of analytical grade. E. coli BL21 (DE3) and DH5α, and interference vector pSliencer4.1 were preserved in our lab. Cell lines (Hela, Bel-7402, and Chang liver cells) were purchased from a typical cell culture collection Committee of the Chinese Academy of Sciences Library, and cultured in RPMI medium 1640 supplemented with 10% FBS (fetal bovine serum, Biotechnology Ltd. Co., Shanghai, China) [22].

Inhibition Effect of T-2 Toxin on Cells

Cells (Hela, Bel-7402, and Chang liver) in logarithmic growth phase were transferred into 96-well plate (10⁵ cells per well, the cell density in the following experiments was the same), and were cultivated overnight. Then, 100 μL of T-2 toxin of various concentration (2000, 1000, 500, 250, 125, 62, and 30 ng/mL) was added respectively. DMSO (Dimethyl sulfoxide), the solvent for T-2 toxin, was added as control. After the cells were incubated with T-2 toxin at a concentration of LC50 (50% lethal concentration) was added, and the cells were cultivated for 0, 4, 8, 16, 24 h respectively. After the incubation, cells were washed twice by PBS, the cells were mixed with protein removal reagent (S solution) at an amount of three times volume of cell pellet, and fully shocked by vortex. Following twice rapid freezing in liquid nitrogen and thawing in 37°C water bath, the samples were keep on ice for 5 min and centrifuged at 10000 g for 10 min under 4°C. At last, the volume of endogenic GSH was detected following the instruction of GSH extract and assay kit, and the volume of endogenic MDA was detected following the instruction of MDA extract and assay kit.

Effect of NAC on Cells’ Viability and GSH Level under T-2 Toxin Stress

Cells in logarithmic growth phase were transferred into 96-well plate, and were cultivated overnight. In every well 100 μL 5 μmol/L NAC (N-acetyl cysteine) was added and incubated for 4 h. After the cells were washed twice by PBS, every well was treated with 100 μL T-2 toxin at a concentration of LC50, and cultivated for 6, 12, 18, 24 h respectively. Following steps referred to the methods mentioned in MTT assay kit and GSH detection and assay kit.

Detection of Caspase-3, 8, 9, p53, and JunD by Western-blot Analysis

Cellular cultivation and T-2 toxin treatment was the same to that mentioned above. The cells in 96-well plate were cultivated for 8, 16, 24 h respectively. After the cells were collected and washed twice by PBS, cells were lysed for 3 min in 100 μL lysis buffer, then were centrifuged at 10000 g for 10 min, and the supernatant was kept on ice for further Western-blot analysis. Samples were used for SDS/PAGE on 12% gels by discontinuous buffer system at 15 mA. Proteins from the gels were transferred to Nitrocellulose (NC) membranes for 1 h at 60 V in transfer buffer (48 mM Tris, 39 mM glycine and 20% methanol) at 4°C. The membranes were incubated with corresponding antibody at a dilution of 1:1000 in TNT buffer (1.211 g Tris, 8.77 g NaCl and 500 mL tween-20 in 1 L TNT, pH 7.0) containing 5% skim milk for 1 h at room temperature on a gentle shaker. The membranes were rinsed three times for 10 min with TNT buffer and incubated with goat anti-mouse HRP-conjugated IgG at a dilution of 1:4000 in TNT buffer containing 5% skim milk for 1 h at room temperature.

Detection of the GSH and MDA under T-2 Toxin Stress

Cells in logarithmic growth phase were transferred into 96-well plate, and were cultivated overnight. In every well 1.6 mL T-2 toxin at a concentration of LC50 (50% lethal concentration) was added, and the cells were cultivated for 0, 4, 8, 16, 24 h respectively. After the incubation, cells were washed twice by PBS, the cells were mixed with protein removal reagent (S solution) at an amount of three times volume of cell pellet, and fully shocked by vortex. Following twice rapid freezing in liquid nitrogen and thawing in 37°C water bath, the samples were keep on ice for 5 min and centrifuged at 10000 g for 10 min under 4°C. At last, the volume of endogenic GSH was detected following the instruction of GSH extract and assay kit, and the volume of endogenic MDA was detected following the instruction of MDA extract and assay kit.

Figure 1. The inhibition effect of T-2 toxin on three cell strains. A. Dose-dependent inhibition ration of T-2 toxin at 24 h. B. Time-dependent effect of T-2 toxin to cells. Data was presented as mean ± SD. *p<0.05, **p<0.01, ***p<0.001. doi:10.1371/journal.pone.0083105.g001
temperature. The membranes were developed with substrate (ECL, Electrochemiluminescence) until optimum color developed [23].

Effect of Over-expressed JunD on Cells' Viability under T-2 Toxin Stress

The recombinant plasmid (JunD-PCDNA3.0, constructed previously in our lab) was extracted from E. coli DH5a, and transfected into cells by liposome (Lipofectamine 2000). After screening by G418, the JunD expression level was detected by Western-blot analysis with anti-JunD antibody as the first antibody [23]. The survival rate was calculated according to that mentioned in MTT assay.

Statistical Analysis

The data from this experiment was analyzed by statistic software (SPSS 13.0), and all data was presented on the form of mean ± standard deviation (x ± s). The comparison between each group was based on single-factor analysis of variance. The comparison between parallel groups was analyzed with LSD test. The assessment of statistical significance of differences was carried out with one way ANOVA in Microsoft Excel. P<0.05 means that the differences are statistically significant.

Results

Dose and Time Dependent Cytotoxicity of T-2 Toxin to Cells

MTT assay was used to detect the inhibition effect of T-2 toxin on cells. It could be concluded from Figure 1A that T-2 toxin was toxic to three lines of cells (Hela, Bel-7402, and Chang liver cell), and the virulence of T-2 toxin was different towards different cells. With the increase of the toxin dose, the cell mortality was increased, too. According to the method provided above, LC50 of T-2 toxin at 24 h to three cell lines (Hela, Bel-7402, and Chang liver cells) was found to be 357, 63, and 412 ng/mL respectively. The concentrations of T-2 toxin used in subsequent experiments were the LC50 of T-2 toxin presented above. Time effect was also observed when cells were stressed under LC50 of T-2 toxin. It could be found from the Figure 1B that as the time of T-2 stress extended, the survival rates of all three cell lines were decreasing obviously. It indicated that the effect of T-2 toxin on cells survival rate was not only dose-dependent, but also time-dependent.

The Effect of T-2 Toxin on Endogenous GSH and MDA

Endogenic GSH is an important antioxidant in cells, and its level impacts on cell anti-oxidative capacity directly. The decrease in GSH level would initiate intracellular oxidative stress. It could be found from Figure 2A that with the elongation of T-2 toxin treatment time, endogenic GSH in these cells was gradually reduced, and the endogenous GSH in test groups was less than
half of that in the control group at 24 h. This indicated that the level of endogenous GSH would decrease when the cells were induced by T-2 toxin, and certain degree of oxidative stress took place in the cells. This was a hint for us to explore the level of MDA, a product of lipid peroxidation. The level of MDA inside cells usually increases significantly, when the cells are under oxidative stress. As shown in Figure 2B, all three cell lines produced different levels of MDA under T-2 toxin stress, and the level of MDA was gradually increased with the elongation of the treatment time. These results and previous reports showed that there was a link between the T-2 toxin stress and intracellular oxidative stress.

The Protective Effect of Antioxidant NAC to the Cells against T-2 Toxin Stress

In the study, antioxidant NAC was employed to test if it had protective effect on cells stressed by T-2 toxin. It could be found from Figure 3A that the survival rate of NAC pretreated Hela cells increased compared to the control group, which meant that the antioxidant NAC could relief the virulence of T-2 toxin, and alleviate T-2 toxin induced cell death. Endogenous GSH was detected, and the result from Figure 3B showed that there was more GSH in NAC pretreated Hela cells. But no obvious protective effect of NAC to the Chang liver and Bel-7402 cell lines under T-2 toxin stress was observed (data no shown). These results suggested that the pernicious effect of T-2 toxin on cells was at least partially caused by oxidative stress, and the protective effect of NAC to Hela cells against T-2 toxin stress was through protecting GSH.

The Effect of T-2 Toxin Stress on Caspase-3

Caspase 3, an important factor in apoptosis, is activated in both death ligand and mitochondrial pathways. In cell, Caspase 3 is an enzyme precursor that is activated only when cell initiates an apoptotic process. In the study, the hydrolase activity of Caspase 3 was detected to reflect the activity level of Caspase-3 in cells (Hela, Bel-7402, and Chang liver) under T-2 toxin stress. It could be observed from Figure 4A that the activity of Caspase-3 in three cell lines under T-2 toxin stress was increased 2–7 times compared to control group at 24 h. These results reflected that Caspase-3 played an important role in the process, and preliminary suggested that the process induced by T-2 toxin in cells was apoptosis.

Figure 4. Activation of Caspase-3 induced by T-2 toxin. A. Detection of Caspase-3 hydrolase activity under T-2 toxin stress. B. Western blot result of Caspase-3 activated fragments in Hela cells when treated with T-2 at the concentration of LC50 for 8, 16, and 24 h respectively. C. Western blot result of Caspase-3 activated fragments in Chang liver cells. D. Caspase-3 activity in Bel-7402 cell. Data was presented as mean ± SD. *p<0.05, **p<0.01, ***p<0.001.
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Figure 5. Western-blot analysis on activated Caspase-8 and Caspase-9 level. A. Activated Caspase-8 and Caspase-9 level in Hela cells when treated with T-2 toxin at the concentration of LC50 for 8, 16, and 24 h respectively. B. Activated Caspase-8 and Caspase-9 level in Chang liver cells were treated with T-2 toxin. C. Bel-7402 cells were treated with T-2 toxin.
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The hydrolyzed or activated fragment of Caspase-3 was further analyzed by Western-blot analysis. The result showed that the level of activated hydrolysis fragment increased in all three cell lines (Figure 4 C–D). These results illuminated the effect of T-2 toxin to cells was apoptosis rather than necrosis.

T-2 Toxin Induced Apoptosis by Mitochondrial Pathway

Members of the Caspase family are critical in the process of apoptosis. Caspase-8 mainly takes part in the death receptor pathway, and Caspase-9 is primarily working in mitochondrial pathway. The results of our Western-blot analysis (Figure 5) in present study showed that Caspase-8 was hydrolyzed and activated at about 16 h, and Caspase-9 was activated at about 8 h (Figure 5A and B). The same situation happened to the line of Bel-7402, and Caspase-8 was hydrolyzed and activated relatively later than that of Caspase-9. Caspase-8 was activated at about 24 h, but Caspase-9 was about 16 h (Figure 5C). These results indicated that the apoptosis induced by T-2 toxin was probably through mitochondrial pathway, which coincided with the previous reports [24,25].

The Effect of T-2 Toxin on the Level of p53

The role of p53 in T-2 toxin induced apoptosis has been controversial. Some reports declared that the level of p53 did not change in the T-2 toxin mediated apoptosis [26], other reports found that p53 level was up-regulated in the process [25]. So it is necessary to further clarify the role of p53 played in the apoptotic

Figure 6. Levels of p53 under T-2 toxin stress. A. Levels of p53 in Hela cells when treated with T-2 at the concentration of LC50 for 8, 16, and 24 h respectively. B. Levels of p53 in Chang liver cells. C. Bel-7402 cells were treated with T-2 Toxin.

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Figure 7. The effect of up-regulated JunD on cells. A. The expression levels of JunD induced by T-2 toxin. B. The expression of JunD after transfection of JunD over-expression vector in three cell strains. wt: wild type. +: cell strains transfected with over-expression vector. C. The effect of over-expressed JunD on cell survival rate. T-2: wild-type cells under T-2 toxin stress. JunD (+/+)/: JunD over-expressed cell lines under T-2 toxin stress.

Data was presented as mean ± SD. *p<0.05, **p<0.01, ***p<0.001.

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process. We found that the protein level of p53 in these three cell lines (Hela, Chang liver, and Bel-7402) was up-regulated under T-2 toxin stress (Figure 6 A–C) suggesting that p53 took part in the apoptotic process induced by T-2 toxin.

The Effect of Up-regulated JunD on Cells under T-2 Toxin Stress

JunD plays a role in intracellular antioxidant system. In the present study, we explored the relationship between JunD and apoptosis induced by T-2 toxin. By Western-blot analysis, we found that the protein levels of intracellular JunD were obviously suppressed under T-2 toxin stress in all three cell lines we used (Figure 7A). The down-regulation of JunD results in the suppression of intracellular antioxidant system, which makes the cells susceptible to oxidative stress.

These three cell lines (Hela, Chang liver, and Bel-7402) were all transfected with an over-expression vectors of JunD. As shown in Figure 7B, the expression levels of JunD were obviously up-regulated in all three cell lines after transfection. It could also be seen from Figure 7C that the survival rate of cell lines transfected with JunD over-expression vector was significantly increased under T-2 toxin stress compared to wild-type cells. These results showed that JunD played a critical role in the apoptotic process, and its over-expression could effectively prevent the cells from damaging effects by T-2 toxin.

Discussion

It had been found in many reports that intracellular reduced GSH was down-regulated by T-2 toxin, and at the same time, lipid oxidation occurred and total intracellular ROS was up-regulated [5,27]. Reduced GSH can effectively remove intracellular free radicals, and the decline in GSH expression level would lead to unbalance of intracellular antioxidant system. The up-regulated ROS and MDA reflected that cells had been coerced by oxidative stress. In the current study, it was clear that all three cell lines (Hela, Bel-7402, and Chang liver) could be induced into the process of apoptosis by T-2 toxin, and the death effect of the toxin against cells was time-dependent and dose-dependent. Oxidative stress might be the main factor which leads to apoptosis under T-2 toxin stress. In the experiment in which antioxidant NAC was used to supplement, the survival rate of cell lines was all improved, which might suggest that there are other factors that could complement the function of JunD. Thus, we found it is important to clarify the role of p53 in T-2 toxin induced apoptosis. It was confirmed in the present study that p53 was up-regulated, which also suggested that the pathway of T-2 induced apoptosis was through the mitochondrial pathway [35].

Author Contributions

Conceived and designed the experiments: DBY ZHZ YLH. Wrote the paper: SHW ZHZ.

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