In Vitro Investigation of the Effects of Imidacloprid on AChE, LDH, and GSH Levels in the L-929 Fibroblast Cell Line

L929 Fibroblast Hücre Hattında İmidoklorit Etkisinin In Vitro Araştırılması

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ABSTRACT

Objectives: There are several types of pesticides to control pests and several new types coming into use that could be less toxic compared to the old ones. Pesticide-induced oxidative stress, which is one of the main mechanisms of toxicity, is the research area focused most on over the last decade. There are several different studies in the literature on whether pesticide exposure induces oxidative stress parameter-mediated toxicity. Pesticide-induced oxidative stress level depends on the biochemical features of mammalian systems. Imidacloprid is a neonicotinoid pesticide in wide use that is considered safe; however, it has been reported in different studies that it may cause changes in oxidative stress parameters.

Materials and Methods: We investigated the dose- and time-dependent effects of imidacloprid on acetylcholinesterase (AChE), lactate dehydrogenase (LDH), and glutathione (GSH) levels in the L-929 fibroblast cell line. The effects of 1-500 µg imidacloprid dose range on AChE, GSH, and LDH were investigated.

Results: LDH levels were significantly increased dose dependently in the 250 and 500 ng imidacloprid groups compared to the control group. GSH levels nonsignificantly decreased dose dependently and GSH levels were lower in the 500 ng imidacloprid group compared to the control group. There were no significant differences between the groups in AChE levels.

Conclusion: These results indicated that high doses of imidacloprid may induce oxidative stress in fibroblast cells.

Key words: Imidacloprid, L-929 cell line, oxidative stress, AChE

ÖZ

Amaç: Haşereleri kontrol altında almak için çeşitli pestisit türleri ve eskilere kıyasla daha az toksik olan yeni tür pestisitler kullanma giriyor. Toksitsitenin ana mekanizmalarından biri olan pestisit kaynaklı oksidatif stres, son on yılda en çok odaklanılan araştırma alanıdır. Literatürde pestisit maruziyetinin oksidatif stres parametresi aracılı toksisiteyi indüklemesine ilişkin farklı çalışmalar mevcuttur. Pestisit kaynaklı oksidatif stres seviyesi, memelilerin biyokimyasal özelliklerine bağlıdır. İmidakloprid, güvenli olduğu düşünülen, yaygın olarak kullanılan bir neonicotinoid pesticidir; ancak oksidatif stres parametrelerinde değişikliklere neden olabileceği farklı çalışmalarda bildirilmiştir.

Gereç ve Yöntemler: Bu çalışmada doza ve zamana bağımlı olarak imidaklopridin L-929 fibroblast hücrelerinde AChE, lactate dehydrogenase (LDH), ve glutathione (GSH) düzeyleri üzerine etkisini inceledik. 1-500 µg imidakloprid dozlarının AChE, GSH, ve LDH düzeyleri üzerine etkileri araştırıldı.

Bulgular: LDH seviyeleri doza bağımlı olarak 250 ve 500 ng imidakloprid doz gruplarında kontrol grubundan anlamlı olarak arttıldı. GSH seviyeleri dozdan bağımsız olarak 500 ng imidakloprid dozunda kontrol grubuna kıyasla anlamlı olarak azaldığı tespit edildi. Asetilkolinesteraz seviyeleri arasında anlamlı bir fark gözlenmedi.

Sonuç: Bu sonuçlara göre yüksek doz imidakloprid maruziyeti fibroblast hücrelerinde oksidatif stres parametrelerini uyarabileceği gözlenmiştir.

Anahtar kelimeler: İmidakloprid, L-929 hücre hattı, oksidatif stres, AChE

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INTRODUCTION

Pesticides are mixtures of substances used to prevent, control, or reduce harmful organisms. Pesticides include active substances and filling material that facilitate usage and increase the effect of their active substances. Commercially available drugs are manufactured and launched to market in mixed form with filling material. In Turkey, 2,500 tons of pesticides are used every year.1-3

It has been speculated that neonicotinoid pesticides exhibit much lower toxic effects on mammals than on insects and therefore can replace organophosphates and carbamate insecticides. Neonicotinoid pesticides are a high risk for humans as they are often used not only in agricultural applications but also in the removal of domestic pests. Neonicotinoid pesticides show their selective toxic effects on insects via nicotinic acetylcholine receptors. Imidacloprid, which is a member of a new neuroactive neonicotinoid insecticide class, is a commonly used insecticide around the world as well as in Turkey. It has been reported that 120 countries have commonly used imidacloprid since 1990, when it was first introduced commercially. It has been reported that imidacloprid’s oral LD50 values for rats were 380-650 mg/kg body weight and for mice 130-170 mg/kg body weight. Under aerobic conditions, imidacloprid is an environmentally persistent chemical with a 3-year half-life, which increases its risk.4,7

Pesticide-induced oxidative stress has been the focus of toxicological research over the last decade as a possible mechanism of toxicity. Several studies have been conducted to determine whether oxidative stress in humans or animals is caused by various agents in this group and is related to their toxic effects.6,9 It has been reported in different studies that imidacloprid induced oxidative stress parameters’ imbalance in different organisms in vitro and in vivo;10-17 however, there are no studies on imidacloprid’s oxidative imbalance effects on fibroblast tissue in the literature.

In the present study we aimed to investigate the in vitro effects of imidacloprid on acetylcholinesterase (AChE), lactate dehydrogenase (LDH), and glutathione (GSH) levels in the L929 fibroblast cell line.

MATERIALS AND METHODS

Chemicals

Dulbecco’s Modified Eagle’s Medium (DMEM), fetal bovine serum (FBS), phosphate-buffered saline (PBS), penicillin streptomycin solution, and trypsin-EDTA solution were purchased from Sigma-Aldrich Co. (St. Louis, MO, USA). The AChE assay kit (Fluorometric Red, ab138873), LDH assay kit (Fluorometric, ab197000), and GSH assay kit (Fluorometric, ab65322) were purchased from Abcam (Cambridge, UK). The L929 cell line (CRL-6364) was purchased from the American Type Culture Collection (ATCC, VA, USA). The imidacloprid-based herbicide (GORTCA FS 600) was purchased from Safa Tarim Ltd. (Turkey) and contained pure imidacloprid (CAS number: 138261-41-3, product code: 0210-207).

Cell culture and treatments

All experiments on L929 were performed within 20 passages. The cells were grown with DMEM containing 10% FBS and 1% penicillin-streptomycin-amphotericin B in a humidified incubator supplied with 5% of CO₂ at 37 °C. Before the treatments were conducted, the cells were cultured for 24 h to ensure attachment.

The L-929 cells were incubated with imidacloprid dissolved in 100% of dimethyl sulfoxide (DMSO) and diluted with medium to the desired concentrations as 500 µg, 250 µg, 125 µg, 50 µg, 25 µg, 5 µg, and 1 µg. Studies reported that cytotoxicity is observed at concentrations higher than 500 µg/mL. In the present study a high concentration of imidacloprid was determined as 500.18 Vehicle control cells received equal volumes of DMSO (0.5%) as the treatment for 24 h. LDH, GSH, and AChE assays were performed after 24 h of exposure.

Lactate dehydrogenase and glutathione parameters

LDH is a cytosolic enzyme and the measurement of its leakage to the extracellular matrix due to cell membrane damage is an indicator of cell membrane integrity loss and oxidative stress.19

For this purpose, the L-929 cells were plated in 96-well plates at 1x10⁴ cells/well and grown for 24 h. Following the cell treatments, the LDH release in supernatants due to membrane damage was quantified using a rat LDH ELISA kit (Cat. number: E-EL-R2547) in accordance with the manufacturer’s protocol. To each well was added 100 µL of standard or sample, followed by incubation for 90 min at 37 °C. After the liquid was removed, 100 µL of Biotinylated detection Ab was added, followed by incubation for 1 h at 37 °C. After aspiration and washing 5 times, 90 µL of substrate reagent was added, followed by incubation for 15 min at 37 °C. Under aerobic conditions, imidacloprid is an environmentally persistent chemical with a 3-year half-life, which increases its risk.4,7

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GSH is an important molecule for the cellular antioxidant system and under oxidative stress conditions GSH level decreases. The L-929 cells were cultured in 25-cm² flasks to determine GSH levels. Following exposure of cells to imidacloprid, 1x10⁶ cells were harvested in 1 mL of PBS and were homogenized by sonication, and the GSH content of the L-929 cells was determined using the GSH ELISA kit (Cat. number: E-EL-0026) to assay human GSH following the manufacturer’s procedure and referred to as µmol/g protein. To each well was added 50 µL of standard or sample and then immediately 50 µL of Biotinylated detection Ab was added, followed by incubation for 45 min at 37 °C. After aspiration and washing 3 times, 100 µL of horseradish peroxidase (HRP) conjugate was added, followed by incubation for 15 min at 37 °C. After more aspiration and washing 5 times, 90 µL of substrate reagent was added, followed by incubation for 15 min at 37 °C. When 50 µL of stop solution was added, it was read at 450 nm using a micro-plate reader (Biotek Epoch, USA).
Acetylcholinesterase assay

AChE is strikingly distributed in the cell-substrate interface of radiated and migrating fibroblasts (morphoregulation by AChE in fibroblasts and astrocytes) and it helps us to understand how the nervous system works. The L-929 cells were cultured in 25-cm² flasks to determine AChE levels. Following exposure of cells to imidacloprid, 1x10⁶ cells were harvested in 1 mL of PBS and were homogenized by sonication, and the AChE content of L-929 cells was determined using the AChE assay kit (colorimetric, Cat. number: ab138871). To each well of AChE standard, blank control, and test samples was added 50 µL of AChE reaction mixture to make the total AChE assay volume 100 µL/well, followed by incubation for 30 min at room temperature. Increased fluorescence monitored absorbance optical density at 410 nm with a micro-plate reader (Biotek Epoch, USA).

Statistical analysis

All the experiments were performed as three replicates and the results were presented as the mean ± standard deviation. The statistical comparisons were evaluated using One-Way ANOVA followed by Tukey’s test for post hoc analysis, and the statistical significance was set at p<0.05 (SPSS, version 21.0, USA).

RESULTS

The LDH results of imidacloprid exposure on L929 cells are shown in Figure 1. We observed that LDH levels increased dose dependently, and 250 and 500 ng imidacloprid increased LDH levels significantly compared to the control group. GSH results of imidacloprid exposure on L929 cells are shown in Figure 2. GSH levels nonsignificantly decreased dose dependently and GSH levels decreased in the 500 ng imidacloprid group compared to the control group. AChE results of imidacloprid exposure on L929 cells are shown in Figure 3. There were no significant differences between the groups, but the AChE level nonsignificantly decreased in the 125 ng group.

DISCUSSION

Neonicotinoids are pesticides popular worldwide whose use has increased since 2000. Neonicotinoids exert their effects on insects by cellular nAChR mechanism. In recent years, neonicotinoids have taken the place of pyrethroid, organophosphorus, and carbamate insecticides. It is thought that they are a safe pesticide group for nontarget species; however, there are several recent studies in the literature about different toxic effects of neonicotinoids on nontarget organisms.

The stable cytoplasmic enzyme LDH is an important biomarker for oxidative stress, apoptosis, necrosis, and other forms of cellular damage, expressed in all cells and rapidly released when the plasma membrane is damaged. In a study conducted in 2018, Kumar et al. showed that LDH level increased with cellular damage, in accordance with our study. Abu Zeid et al. reported that imidacloprid exposure in the Rock pigeon (Columba livia domestica) resulted in a significant increase in plasma LDH levels in high dose (6 mg/kg) and median dose (3 mg/kg) imidacloprid groups; however, there were no significant increase in the low dose (2 mg/kg) group compared to the control group. Plasma AChE’s enzyme activities in all imidacloprid dose groups significantly increased compared to the control group. Lonare et al. demonstrated that, 45 and 90 mg/kg body weight oral exposure of imidacloprid for 28 days significantly decreased AChE level in rats' erythrocytes. In addition, the brain AChE activity in rats in the imidacloprid treatment groups was decreased in a dose dependant manner compared to the control group. They also demonstrated that LDH levels increased and
GSH levels decreased significantly imidacloprid treated groups in different tissues of rats. Imidacloprid exposure increased the GSH and AChE activities in Chinese rare minnows’ brain. This result indicates that imidacloprid has no effect on Chinese rare minnows. It has been demonstrated that imidacloprid has neurotoxic effects through AChE inhibition and induces oxidative stress in rainbow trout brain tissue. Moreover, imidacloprid exposure significantly decreased AChE levels in the plasma and brain of 10 and 20 mg/kg treated female rats. In another study 24-h 20 mg/kg imidacloprid administration decreased AChE activity approximately 22% in the brain and 28% in red blood cells.

CONCLUSION

Studies about imidacloprid’s effects on AChE, LDH, and GSH express controversial results, which may be related to different study conditions such as species variety, exposure time variety, in vitro and in vivo conditions, and different doses. There are studies that include oxidative stress inducing effects of imidacloprid with different species in the literature; however, the underlying mechanism is not elucidated yet. Further studies are needed to clarify the toxic effects of neonicotinoids, especially imidacloprid, in different species and in different tissues.

Conflicts of interest: No conflict of interest was declared by the authors. The authors alone are responsible for the content and writing of the paper.

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