Cytotoxic effects of aripiprazole on MKN45 and NIH3T3 cell lines and genotoxic effects on human peripheral blood lymphocytes

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INTRODUCTION

Cancer is now believed to rank among the three major causes of death and among them gastric cancer is known as the third leading cause of cancer-related death worldwide; however, the incidence of gastric cancer varies greatly and this number can change across populations¹,². Although there are various treatment options to treat localized gastric cancer, ranging from minimally invasive EMR to aggressive extended lymphadenectomy and perioperative em

ABSTRACT – Background – Gastric cancer is known as the fourth most common cancer. Current treatments for cancer have damaged the sensitive tissues of the healthy body, and in many cases, cancer will be recurrent. Therefore, need for treatments that are more effective is well felt. Researchers have recently shifted their attention towards antipsychotic dopamine antagonists to treat cancer. The anticancer activities of aripiprazole remain unknown.

Objective – This study aimed to evaluate the efficacy and safety of aripiprazole on gastric cancer and normal cell lines. Methods – In this regard, the cytotoxicity and genotoxicity of aripiprazole were investigated in MKN45 and NIH3T3 cell lines by methyl tetrazolium assay and on peripheral blood lymphocytes by micronucleus assay. For this purpose, cells were cultured in 96 wells plate. Stock solutions of aripiprazole and cisplatin were prepared. After cell incubation with different concentrations of aripiprazole (1, 10, 25, 50, 100 and 200 μL), methyl tetrazolium solution was added. For micronucleus assay fresh blood was added to RPMI culture medium 1640 supplemented, and different concentrations of aripiprazole (50, 100 and 200 μL) were added. Results – The finding of present study showed that the IC₅₀ of aripiprazole in the cancer cell line (21.36 μg/mL) was lower than that in the normal cell line (54.17 μg/mL). Moreover, the micronucleus assay showed that the frequency of micronuclei of aripiprazole at concentrations below 200 μM was much less than cisplatin. Conclusion – Aripiprazole can be a good cytotoxic compound and good candidate for further studies of cancer therapy.

essential. Actually, a favorable profile of safety and tolerability and good clinical effectiveness are among the advantages of aripiprazole\(^{18,19}\). However, the anticancer effects of aripiprazole on gastric cancer MKN45 cell line, and the possible genotoxicity of it in human cells has not been confirmed yet. This study aimed to evaluate the efficacy and safety of aripiprazole on gastric cancer and normal cell lines. In this regard, the cytotoxicity and genotoxicity of aripiprazole were investigated in MKN45 and NIH3T3 cell lines by MTT assay and on peripheral blood lymphocytes by MN assay.

**METHODS**

**Cell culture**

NIH3T3 and MKN45 Cell lines (Pasteur Institute, Tehran, Iran) were cultured in Dulbecco’s Modified Eagle Medium (DMEM) (GIBCO, Berin, Germany) with 10% fetal bovine serum (Gibco-BRL, Germany) and 100 μg/mL streptomycin (Gibco-BRL, Germany) and 100 IU/mL penicillin (Gibco-BRL, Germany). Cell cultures were adjusted to allow for exponential growth.

**MTT assay**

NIH3T3 and MKN45 Cell lines (10\(^4\) cells) were cultured in 200 μL DMEM/F12 medium containing 10% bovine serum in 96 wells plate and incubated at 37°C for 24h. Stock solutions of aripiprazole and cisplatin were prepared in 1% DMSO and phosphate buffered saline (PBS), respectively. Twenty-μL of MTT solution (5 mg/mL) was added to each well following 48h incubation with different concentrations of aripiprazole (1, 10, 25, 50, 100 and 200 μL). The optical density (OD) of the MTT reaction was measured on a microplate ELISA reader at 570 nm. All experiments were repeated two times and each treatment was run in triplicate. The percentage of cell viability was calculated using the equation: (mean (OD) of treated cells/mean OD of control cells (1% DMSO)) \(\times 100\).

**Micronucleus assay (CBMN assay)**

Fresh blood was collected from 10 healthy, no smoking, no alcoholic, male donors aged between 25-35 years by venepuncture in heparinized falcons. 0.5 mL of whole blood was added to 4.5 mL of Roswell Park Memorial Institute (RPMI) culture medium 1640 supplemented with 10% fetal bovine serum containing L-glutamine, antibiotics, and phytohemagglutinin (PHA), and different doses of duloxetine (1, 10, 25, 50, 100 and 200 μL). The binucleated lymphocytes were harvested 28h after adding Cyt-B (Sigma, Missouri, USA); they were treated by hypotonic KCl (0.075M) to red blood cell (RBC) lysis. Then fixative solution (methanol: acetic acid =6:1) was added to the cells prior to slide preparation and staining. For slide preparation, 2-3 drops of cell suspension were thrown on a clean slide. The slides were stained with Giemsa solution (4%) using a light microscope to estimate mitotic index (the cells with two or more nuclei per 1000 observed cells) and micronuclei frequency (the number of micronuclei in 1000 binucleated cells) are lymphocytes that were once divided by mitosis. The experiment was performed two times. Mitotic Index has a direct relation with cells’ proliferative activity.

**Statistical analysis**

One way analysis of variance and Tukey’s honestly significant differences (HSD) test were used for multiple comparisons of data. A P value less than 0.05 was considered as significant. The IC\(_{50}\) (half maximal inhibitory concentration) values were calculated by PRISM software using nonlinear regression. Standard deviations represent average results of double experiments. The IC\(_{50}\) values were compared using the Student’s t-test measuring the effectiveness of a substance to cause cell death or inhibit cell growth. Therefore, the lower amount of IC\(_{50}\) represents a higher toxicity of a compound, which leads to death or inhibition of cell growth.

**RESULTS**

**MTT assay**

The IC\(_{50}\) of aripiprazole on MKN45 and NIH3T3 cell lines were examined using MTT assay. The IC\(_{50}\) of aripiprazole on MKN45 cell line was 21.36 μg/mL and on NIH3T3 cell line was 54.17 μg/mL. The lower IC\(_{50}\) value is representative of the higher ability of a cytotoxic compound to cause cell death or inhibit cell growth\(^{20}\). The results of the MKN45 cancer cell line (TABLE 1 and FIGURE 1) showed that aripiprazole compared to the negative control group had more cytotoxic effects in every concentration. Compared to the positive control group; cisplatin (the common anticancer drug) it had no significant difference at 100 and 200 μM, but at lower concentrations: 50, 25, 10 and 1 μM, it showed less cytotoxic effects \((P>0.05)\). Furthermore, the results of the NIH3T3 cell line (TABLE 1 and FIGURE 2) showed that aripiprazole in comparison with the negative control group had more cytotoxic effects at 10, 25, 50, 100, 200 μM and no significant difference at 1 μM. Moreover, in comparison with the positive control group (cisplatin), it showed less cytotoxic effects in every concentration but 100 μM \((P<0.05)\).

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<th>TABLE 1. Cell viability assay.</th>
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<td>Control</td>
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<td>Effect of aripiprazole on NIH3T3 cell viability</td>
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<td>Micronuclei frequency in different concentrations of aripiprazole</td>
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The genotoxic effects of aripiprazole were studied based on the number of MN produced in peripheral blood lymphocytes following treatment with different concentrations of aripiprazole (TABLE 1 and FIGURE 3). Results showed that the MN number was relatively increased based on the increase in the aripiprazole concentration. Fifty-μM concentration of aripiprazole did not produce any significant difference in MN number relative to the control group. While, 100 and 200 μM concentrations of aripiprazole significantly increased the number of MN. Comparison of different concentrations of aripiprazole and cisplatin showed that treatment of lymphocytes with 50 and 100 μM of aripiprazole significantly increased the number of MN. Two hundred-μM concentration of aripiprazole did not produce any significant difference in MN number relative to the cisplatin (P<0.05).

**DISCUSSION**

In this study, the cytotoxic effect of Duloxetine on MKN45 gastric cancer cell line was investigated and compared to NIH3T3 normal cell line by MTT assay. Moreover, the genetic damage caused by this drug was also evaluated using MN assay. The IC$_{50}$ of aripiprazole on MKN45 cell line and NIH3T3 cell line was calculated 21.36 μg/mL and 54.17 μg/mL respectively. However, the IC$_{50}$ of cisplatin on MKN45 was 12.49 μg/mL and on NIH3T3 cell line was 24.9 μg/mL. The lower IC$_{50}$ value is representative of the higher ability of a cytotoxic compound to cause cell death or inhibit cell growth. Taken together, it seems that this drug can be mentioned as a good cytotoxic compound. As reported here, the cytotoxicity of aripiprazole on MKN45 cancer cell line is consistent with some other researches(21). Anti-cancer effects of aripiprazole on various malignant tumor cells and its molecular mechanism were examined by using cell proliferation assay, xenograft mouse model, immunoblotting analysis, migration assay, luciferase reporter gene assay, kinase assay, and overexpression strategy. Treatment with aripiprazole induced cytotoxicity in U251 glioma cells, MKN-1 gastric adenosquamous carcinoma cells, and CT26 colon carcinoma cells(25). Moreover, the effects of aripiprazole alone and in combination with chemotherapeutic agents in order to test growth ability, the ability to form stem cells/differentiation/chemical resistance, the markers of cancer stem cells, cancer cells and normalized cells was investigated. According to their study, the growth of cancer cells and cancer stem cells was inhibited by aripiprazole in non-toxic concentrations of normal cells. Generally, it could be concluded that aripiprazole could be a good candidate as an anticancer stem cell(26).

Additionally, we used lymphocytes in our in vitro studies to assess the potential genotoxic effect of aripiprazole. Results from the micronucleus assay confirmed the ability of aripiprazole to induce the formation of micronuclei. The induction of micronuclei is commonly used to evaluate the chromosomal damage. The cellular and tissue toxicity was observed in the increased therapeutic concentrations of aripiprazole. Aripiprazole and its metabolites can bind DNA, causing damage that can result in chromosome breaks, micronucleus formation, and cell death. As evident, the frequency of micronuclei was increased with aripiprazole in comparison with the negative control group but in comparison with the positive control group, it caused less DNA damage at 100 and 50 μM concentrations. The genetic damage caused by aripiprazole at concentrations below 200 μM was much less than cisplatin, indicating that it could be studied more widely due to its low genotoxicity. The results of our findings in this regard coincided with the work of other researchers(22,23). The mutagenic and antimutagenic effects of some quinoline- and isoquinoline-sulfonamide analogs of aripiprazole were evaluated and their...
results showed that newly synthesized azinesulfonamide analogs of aripiprazole might be considered as genotoxicity safe as they do not display mutagenic activity on the tester strains\(^2,2\). In addition, genetic toxicity of 545 medications including aripiprazole were studied by using information from 1999 to 2008. The data also contained anti-cancer and antiviral drugs, nucleosides (all with known mechanistic genotoxicity), steroids with class-specific genotoxicity and biologicals or peptide-based drugs. Their results discussed the link between postive genetic toxicity findings, rodent carcinogenesis, and silica predictions and found that there was supporting evidence for the idea that just the presence of an N-dialkyl group or piperidine aryl ketone may somehow be associated with genotoxicity\(^2,2\).

CONCLUSION

Aripiprazole can be a great cytotoxic compound and beneficial candidate for further studies of cancer therapy. This study in comparison to other studies showed that aripiprazole can be considered as one of the candidates for the synthesis of a safe anticancer drug. Further studies are needed.

ACKNOWLEDGEMENT

This research was supported/partially supported by Sana institute of high education. We thank our colleagues from [mehdi abbas roshan] who provided insight and expertise that greatly assisted the research, although they may not agree with all of the interpretations/conclusions of this paper.

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