Morinda citrifolia (Noni) Alters Oxidative Stress Marker and Antioxidant Activity in Cervical Cancer Cell Lines

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Abstract

Background: Cervical cancer, the second most common cancer in women, has a high mortality rate. Cisplatin, an antitumor agent, is generally used for its treatment. However, the administration of cisplatin is associated with side effects and intrinsic resistance. Morinda citrifolia (Noni), a natural plant product, has been shown to have antioxidant activities in vitro and in vivo. Materials and Methods: Both HeLa and SiHa cervical cancer cell lines were treated with 10% Noni, 10 mg/dl cisplatin, or a combination of both 10% Noni and 10 mg/dl cisplatin for 24 hours. Post culturing, the cells were pelleted and stored at -70°C for malondialdehyde and catalase assays. Results: On treatment with Noni, CP, and their combination, the level of MDA decreased by 0.76 fold, 0.49 fold, and 0.68 fold respectively in HeLa cells; and by 0.93 fold, 0.67 fold, and 0.79 fold respectively in SiHa cells, as compared to their controls; whereas catalase activity increased by 1.61 fold, 0.54 fold, and 2.35 fold, respectively in HeLa cells; and by 0.98 fold, 0.39 fold, and 1.85 fold respectively in SiHa cells. Conclusions: A decrease in level of lipid peroxidation and an increase in catalase activity were observed with Noni by itself and the effect ameliorated changes observed with cisplatin when given in combination.

Keywords: Morinda citrifolia (Noni) - cisplatin - malondialdehyde - catalase - HeLa - SiHa
prevent angiogenesis and metastatic spread, suggesting a potential role for antioxidants as adjuvants in cancer therapy (Borek, 2004b). Even though some studies have examined the antioxidant effects of Noni on cancer and have elucidated some of the mechanisms involved, there is still very little information available on the usefulness of Noni in the field of cancer, especially in terms of its antioxidant effects in cervical cancer. Hence, the aim of this study was to study the antioxidant properties of Noni by itself and in combination with Cisplatin (a commonly used chemotherapeutic agent for treatment of cervical cancer) in human cervical cancer HeLa and SiHa cell lines.

Materials and Methods

Cell line (s)

Two cell lines, i.e. HeLa (HPV18+) and SiHa (HPV16+) cervical cancer cell lines were used in this study. These cell lines were cultured in DMEM (Sigma), supplemented with 10% fetal bovine serum (FBS), 100 U/ml Penicillin, and 100 μg/ml Streptomycin. Cultures were maintained at 37°C in 5% CO₂ and 95% humidified atmosphere.

Cell culture and treatments

Noni juice was obtained from Health India Laboratories (a unit of Noni BioTech Pvt. Ltd., Chennai, India). Cisplatin was purchased from Sigma, USA. The cells were treated with different concentrations of CP and Noni and incubated at 37°C for 24 hrs.

Both the cervical cancer HeLa and SiHa cell lines were treated with 10% Noni (v/v), 10 μg/ml Cisplatin, and combination of both 10% Noni (v/v), 10 μg/ml Cisplatin for 24 hours at 37°C in 5%CO₂ and 95% humidified atmosphere. Post culturing, the cells were pelleted and stored at -70°C for various assays (Gupta et al., 2013).

Measurement of malondialdehyde (MDA) level

The cells were collected and re-suspended in 1 ml of DMEM culture medium. The samples were then sonicated and used for MDA measurement as described in the protocol. The Thiobarbituric Acid Reactive Substances (TBARS) assay was performed using protocol provided in the TBARS Assay Kit, Cayman Chemical Company.

Assay of catalase activity

The cells were collected by centrifugation and the cell pellet was then sonicated on ice in 1-2 ml of cold buffer (i.e. 50 mM potassium phosphate, pH 7.0, containing 1 mM EDTA). After sonication, the product was centrifuged at 10,000× g for 15 min at 40°C. The supernatant was removed for assay and stored on ice. The assay was performed with Catalase Assay Kit, Cayman Chemical Company.

Statistical analysis

Results of each experiment represent the mean±standard deviation (SD) of three independent experiments carried out in triplicate. All the data were analysed using Student’s t-test. Statistical analysis showing a value of p<0.05 was considered significant.

Figure 1. Effect of Noni (10%), CP (10 μg/ml), and their Combination on A) MDA Level and B) Catalase Activity. The results shown in the bar diagram are means±SD of three individual experiments. Statistical analysis showing a value of p<0.05 was considered significant

Results

Effect of noni and cisplatin on the level of malondialdehyde

Increased levels of Malondialdehyde (MDA) indicate upsurge in lipid peroxidation which is a consequence of increased free radical generation. It causes profound alterations in the function of the cell membrane and structural organization of DNA leading to mutations. Therefore, it can be stated that lipid peroxidation may possibly be involved in cervical carcinogenesis. On treatment with Noni, CP, and their combination, the level of MDA decreased by 0.76 fold, 0.49 fold, and 0.68 fold respectively in HeLa cells; and by 0.93 fold, 0.67 fold, and 0.79 fold respectively in SiHa cells, as compared to their controls (Figure 1A).

Effect of noni and cisplatin on catalase activity

Catalase is an ubiquitous antioxidant enzyme that is present in most aerobic cells. Catalase is involved in the detoxification of hydrogen peroxide, a reactive oxygen species (ROS), which is a toxic product of both normal aerobic metabolism and pathogenic ROS production. On treatment with Noni, CP, and their combination, Catalase activity was found to be increased by 1.61 folds, 0.54 fold, and 0.79 fold respectively in HeLa cells; and by 0.93 fold, 0.67 fold, and 0.79 fold respectively in SiHa cells, as compared to their controls (Figure 1B).

Discussion

Search for new chemopreventive and antioxidant agents that are more effective but less toxic has kindled great interest in phytochemicals. Noni, fruit juice derived from the plant Morinda citrifolia, is one such compound which was used in this study. Noni is a herbal remedy with promising antioxidant properties (Wang et al., 2009; Serafini et al., 2011).

We studied the antioxidant properties of Noni by itself, Cisplatin by itself, and their combination, on two cervical cancer cell lines (HeLa and SiHa). These cell lines were chosen for the study because they harbor the human papilloma virus (HPV) type 18 (HeLa) and type 16 (SiHa) genotypes. These HPVs have been shown in multi-institutional studies as etiological agents in the development of cervical cancer and these genotypes account for >70% of all HPV DNA positive invasive cervical cancers (Gupta et al., 2013).
Reactive oxygen species (ROS) such as hydrogen peroxide, superoxide anions, and hydroxyl radicals promote apoptosis. However, the hydroxyl radicals are capable of abstracting a hydrogen atom from polyunsaturated fatty acids in membrane lipids to initiate lipid peroxidation. These free radicals can evoke extensive tissue damage, reacting with macromolecules, such as membrane lipids, proteins, and nucleic acids (Gupta et al., 2012). Thus, an alteration in enzymatic antioxidant status with increase in lipid peroxidation indicates that the enzymes play an important role in combating free radical-induced oxidative stress. Hence, in this study, we measured the ability of Noni/Cisplatin to inhibit lipid peroxidation and also studied the activity of antioxidant enzymes.

The level of Malondialdehyde (MDA), a marker for lipid peroxidation, was found to be maximum in controls as compared to the treated cells (p<0.001), in both HeLa and SiHa cells, therefore reflecting insufficient antioxidant potential in both the cervical cancer cells. The level of MDA decreased significantly on treatment with Noni/Cisplatin alone and with their combination, in both HeLa and SiHa cells; with Noni showing greater decrease in lipid peroxidation. This suggests that antioxidants present in Noni, mainly the lipid soluble ones, might exert a profound effect on inhibition of lipid peroxidation and free radical generation. However, the exact mechanism is not clear.

Catalase is a free radical scavenging enzyme, present in all oxygen metabolizing cells and provides defense against potentially damaging entities of hydrogen peroxide. Our study showed increased level of Catalase on various treatments, indicating their protective role. The activity of Catalase increased significantly with Noni alone (p<0.05), but on Cisplatin treatment, it was found to decrease as compared to Noni alone (p<0.01). This implies that Noni is a better enhancer of Catalase activity. The activity of Catalase increased significantly with their combination as compared to control (p<0.01) and as compared to either Noni alone (p<0.001) or Cisplatin alone (p<0.001). This could be due to the ability of Noni to scavenge free radicals more effectively, magnifying the $\text{H}_2\text{O}_2$ level when treated in the combination, and hence, magnified Catalase activity could be a part of the defense system specific to the epithelial cells, designed to eliminate $\text{H}_2\text{O}_2$ produced. The increase in Catalase activity could also be due to decrease in lipid peroxidation with Noni/Cisplatin alone and with their combination, thus enhancing Catalase activity.

Based on the results obtained, it is highly possible that several compounds of different polarity may contribute to the antioxidant activity of M. citrifolia fruit juice (Noni). Part of the antioxidant activity may be due to lipid soluble polyphenols, anthraquinones, $\alpha$-tocopherol, and $\beta$-carotene present in Noni, and many natural phytochemicals have been demonstrated to promote antioxidant activity in cancer cells, including phenolics, alkaloids, dammacanthal, and flavonoids (Alshatwi et al., 2011; Wong and Kadir, 2012). It has also been reported that most natural antioxidant compounds often work synergistically with each other to produce a broad spectrum of effects against free radical attack. Our data also demonstrate the presence of possible interrelationship and crosstalk between Noni and Cisplatin which may be related, at least, partly to the antioxidant status in cervical cancer cell lines.

Taken together, this study showed that Noni/Cisplatin by themselves and their combination were able to decrease lipid peroxidation and enhance Catalase activity in both the cervical cancer HeLa and SiHa cell lines. Hence, Noni offers potential to be used as a chemo adjuvant, as an antioxidant, especially for the treatment of cervical cancer. Although further studies are needed to conclude that Noni, as antioxidants, do not conflict with the use of chemotherapy in the treatment of a cervical cancer and that, it may significantly mitigate the adverse effects of that treatment.

References
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