Effects of *Pluchea lanceolata* Root Extract on Cisplatin--induced Nausea and Vomiting in Rat Pica Model

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Received June 12, 2012; Revised September 27, 2012; Accepted November 8, 2012

This paper is available online at http://ijpt.tums.ac.ir

**ABSTRACT**

Cisplatin is an effective chemotherapeutics against a wide range of cancers. However, it causes significant nausea and vomiting which limit its usefulness. In the present study, the effects of methanolic root extract of *Pluchea lanceolata* (DC.) C. B. Clarke, asteraceae (MPL) was investigated against cisplatin-induced nausea using a rat pica model. In rat pica model, rats react to cisplatin (emetic/nausea stimuli), with altered feeding habits, manifested by increased consumption of kaolin. The pica in rats was measured to quantify cisplatin-induced nausea, and to evaluate the protective effect of pretreatment with MPL given orally. Cisplatin at 3 mg/kg (i.p.) induced significant pica indicated by reduced food intake and increased kaolin consumption, suggesting the presence of nausea/emesis. Cisplatin-induced pica decreased significantly when animals were pretreated with MPL at doses of 400 mg/kg p.o. (p < 0.05). MPL pretreatment decreased cisplatin-induced kaolin intake in the rat model of simulated nausea, suggesting that MPL and/or its active constituent(s) may play a therapeutic role as protective against chemotherapy-induced emesis.

**Keywords:** Cisplatin, Pica, Pluchea lanceolata, Asteraceae
of life. The pathophysiology of these symptoms has been partly attributed to oxidant injury to the gastrointestinal epithelium [8,9]. The mucosal injury results in leakage of voucherm (PP-569) was deposited in the Department of Pharmaceutical Science, Guru RamDas University, Hisar, India. Since 1999, the plant material was further size reduced and oxidant injury to the gut may be the primary event. A stored until further use in an air tight container. The defatted pretreatment with an antioxidant should ameliorate these symptoms. Despite advances in antiemetic therapy, nausea and vomiting remain among the most feared adverse events associated with chemotherapy. Herbal medicines may represent an alternative new class of low-cost antiemetics. To get a reddish brown powder with 6.2% yield, hereafter referred to as MPL (Methanolic extract of Pluchea lanceolata).

**MATERIALS AND METHODS**

**Drugs and Chemicals**

Cisplatin injection (Cipla, Ltd., India), Kaolin and Methanol (SD Fine-Chem Ltd, India) and all other chemicals were of analytical grades.

**Animals**

Male Wistar strain rats (150-250 g, 3-4 months of age) were procured from the disease-free small animal house of CCS Haryana Agriculture University, Hisar, Haryana, India. The animals were housed at 24 ± 1°C temperature, 45 ± 5% humidity, 12-h light-dark cycle, and left to acclimatize for 1 week before the experiments. Rats were allowed free access to water, standard laboratory rat chow and kaolin, placed in separated containers, continuously available throughout the experiment. Experiments were carried out between 09:00 and 17:00 h. The experimental protocol was approved by the Institutional Animal Ethics Committee, GJUS&T, Hisar, Haryana and the care of the laboratory animals was taken as per the guidelines of CPCSEA, Ministry of Forests and Environment, Government of India.

**Preparation of extracts of Pluchea lanceolata**

The shade dried roots of the plant *Pluchea lanceolata* (DC.) C. B. Clarke, asteraceae, was collected from waste land of Dist. Hisar and Sirsa, Haryana, India, in October 2009 and authenticated by Rawat, Dept. of Botany, GJUS&T, Hisar. The plant material was further size reduced and oxidant injury to the gut may be the primary event. A stored until further use in an air tight container. The defatted pretreatment with an antioxidant should ameliorate these symptoms. Despite advances in antiemetic therapy, nausea and vomiting remain among the most feared adverse events associated with chemotherapy. Herbal medicines may represent an alternative new class of low-cost antiemetics. To get a reddish brown powder with 6.2% yield, hereafter referred to as MPL (Methanolic extract of *Pluchea lanceolata*).

**Experimental design**

The rats were randomly assigned to six groups of six animals each. Group I and II treated with vehicle (distilled water) was kept as normal and control group respectively. Group III and IV were administered with MPL (200 and 400 mg/kg body wt; p.o.) for 7 days. Group V and VI were also administered with MPL (200 and 400 mg/kg body wt; p.o.) for 7 days. Group II, III and IV were injected with a single dose of cisplatin (03 mg/kg body weight; i.p.) on day 4, to induce the pica behavior. On each experimental day (next five consecutive days), kaolin intake (g), food intake (g), and body weight (g) were measured. To measure kaolin and food intake, the remaining kaolin and food from the day prior was collected including that spilled outside the containers. The collected kaolin and food were dried for 72 h to obtain dry weight (g).

**Statistical analysis**

The statistical significance of differences among values of individual parameters was evaluated by using the Student’s *t* test. All the values are expressed as mean ± SD. The significance was set at *p* < 0.05.

**RESULTS**

Kaolin intake (pica) was measured in rats of various groups under study. Fig 1 demonstrates that MPL from waste land of Dist. Hisar and Sirsa, Haryana pretreatment significantly reduced kaolin intake induced by cisplatin. Cisplatin induced a significant increase in kaolin consumption in the animals of group II at 24, 48,
72, 96 and 120 h compared to normal animals of group I
168 (p < 0.05). The MPL (200 mg/kg) pretreatment
significantly decreases the kaolin intake compared to
the group II at 24, 48, 72 and 96 h (p < 0.05). Kaolin intake at 24 h (4.1±0.27 g) was significantly lower in
MPL (200 mg/kg) pretreated animals than the animals
of group II (6.9± 0.43 g). However, kaolin intake was
still higher than normal baseline intake at 0 h (0.3 ±0.02
g). Pretreatment with MPL (400 mg/kg) significantly
reduced kaolin intake compared to group II at 24, 48,
and 72, 96 and 120 h (p < 0.05). Moreover, the kaolin
treatment with MPL (400 mg/kg) significantly
reduced kaolin intake compared to group II at 24, 48,
and 72, 96 and 120 h (p < 0.05). Moreover, the kaolin
consumption was near to the baseline intake at 0 h. This suggests that MPL at 400 mg/kg reduced the pica for cisplatin-treated rats. Additionally, the antioxidant activity of MPL may be one of the mechanisms by which MPL attenuates cisplatin-induced nausea/emesis. The present study demonstrated that a single dose of cisplatin (3 mg/kg; i.p.) induced an alteration in food habit, indicated by increased kaolin consumption and reduced food intake. The mechanism of cisplatin-induced nausea/vomiting is possibly mediated via cytotoxic damage to the enterochromaffin cells in the small intestine by ROS release and treatment with an antioxidant should reduce these side effects. Based on the results, it can be concluded that MPL may be a potential candidate for the treatment of cisplatin-induced nausea/vomiting in rats.

**Figure 1.** Effect of cisplatin (3 mg/kg) and cisplatin plus MPL (200 and 400 mg/kg) on kaolin intake. Values are expressed as mean ± SD. *p < 0.05 with respect to normal, *p < 0.05 with respect to control.

**Figure 2.** Effect of cisplatin (3 mg/kg) and cisplatin plus MPL (200 and 400 mg/kg) on reduced food intake (% baseline) induced by cisplatin in rats. Values are expressed as mean ± SD. *p < 0.05 with respect to normal, *p < 0.05 with respect to control.

**Discussion**

The present study inferred that methanolic extract from *Pluchea lanceolata* attenuated kaolin intake (pica) and cisplatin-induced nausea/vomiting in rats. Additionally, the antioxidant activity of MPL may be one of the mechanisms by which MPL attenuates cisplatin-induced nausea/emesis. The mechanism of cisplatin-induced nausea/vomiting is possibly mediated via cytotoxic damage to the enterochromaffin cells in the small intestine by ROS release and treatment with an antioxidant should reduce these side effects. Based on the results, it can be concluded that MPL may be a potential candidate for the treatment of cisplatin-induced nausea/vomiting in rats.
these facts, the present investigation was done to evaluate the efficacy of Pluchea lanceolata, in cisplatin-induced pica. In vitro antioxidant activity of methanolic root extract of Pluchea lanceolata was already determined by DPPH free radical scavenging assay and hydrogen peroxide scavenging activity [26,27]. The results showed that MPL at dose of 200 mg/kg and 400 mg/kg reduced cisplatin-induced pica. This suggests that cisplatin-induced pica (nausea) could be treated with MPL. Although low doses of MPL caused reduced pica in cisplatin-treated rats, the improvement was still less as compared to normal kaolin intake. These findings support the notion that herbal medications, such as MPL, could be an effective and inexpensive alternative for preventing chemotherapy-induced emesis without troublesome side effects. Further, earlier studies also showed that herbal antioxidants may have a role in attenuating cisplatin-induced nausea and vomiting [28]. However, it is important to examine the interaction between the herbal extract and cisplatin, which could either hamper or augment the anticancer actions of cisplatin. As cisplatin act by oxidative stress in tumor cells and treatment with antioxidants could detoxify ROS, the herb may prevent oxidative injury to tumor cells and sensitize the tumor cells to the anticancer effects of chemotherapy [29].

We conclude that herbal antioxidants potentially represent a new class of low-cost antiemetic agents for the treatment of chemotherapy-induced nausea/vomiting. Additional studies are required to further investigate the antiemetic actions of such herbal medications and the effects of interaction with the chemotherapeutic agents.

REFERENCES

Pluchea Lanceolata and cisplatin-induced nausea/vomiting

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Published online: January 31, 2013