Anti-Proliferative Activity of \textit{Nigella sativa} Chloroform Extract on 7,12-Dimethylbenz[a]anthracene Induced Female Rats Spleenocyte

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Abstract

Previous study reported that \textit{Nigella sativa} has in vitro and in vivo cancer activity. This study was conducted to observe the effect of chloroform extract of \textit{Nigella sativa} seed (NCE) on 7,12-Dimethylbenz[a]anthracene (DMBA)-induced female rats’ spleenocyte. The experiment consisted of five groups: corn oil solvent control group, DMBA group, DMBA+250 mg/kgBW NCE, DMBA+500 mg/kgBW NCE and DMBA+750 mg/kgBW NCE. Extract was dissolved in corn oil and oral administered daily for 2 weeks before and during the DMBA induction. Observation of cell proliferation was performed using haematoxylin and eosin (H&E) and AgNOR stainings. H&E staining showed decreased necrosis activity extract groups compared to DMBA group. From AgNOR staining results, mean AgNOR (mAgNOR) of extract groups was less in number compared to DMBA group. The mAgNOR in corn oil solvent control group, DMBA group, DMBA+250 mg/kgBW NCE, DMBA+500 mg/kgBW NCE and DMBA+750 mg/kgBW NCE were 1.22, 1.91, 1.29, 1.36 and 1.33, respectively. Our current results showed that NCE reduces the proliferation of DMBA-induced rat spleenocytes. Thus, NCE has potency to be developed as a chemopreventive agent.

Keywords: \textit{Nigella sativa}, spleen, DMBA, anti-proliferative

INTRODUCTION

Malignant lymphocytes can grow in various lymphoid organs in the body. One of the lymphoid organs is spleen which is the largest lymphoid organ in the organism. One of the carcinogenic substances that caused cancer is 7,12-Dimethylbenz[a]anthracene (DMBA). DMBA is metabolized by cytochrome P450 enzymes. DMBA will react with DNA to initiate cancer. DMBA caused normal cells in the spleen to change into lymphoblastic cell (Gao, et al., 2007).

Chemotherapy and radiotherapy have been developed to treat cancer. These treatment methods cause a lot of side effects, including hair loss, feeling or being sick, feeling tired, and having low levels of blood cells (ACS, 2010). Therefore, we need to search for new ingredients which can be used safely by patients to inhibit cancer proliferation. Herbal plants have been studied to prevent and to treat diseases. One of the herbal plants is black cumin (\textit{Nigella sativa}).

Compounds of \textit{N.sativa} which have anti-proliferative activity are palmitic acid, stearic acid, oleic acid and linoleic acid (Syazana, et al., 2011). The gas chromatography mass spectrometry (GCMS) showed that chloroform extract of \textit{N.sativa} contained 95.5% fatty acids which the main component fats are linoleic acid (55.6%), oleic acid (23.4%) and palmitic acid (12.5%) (Ekowati, et al., 2011).

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Based on previous studies, chloroform extract of *N. sativa* (NCE) has *in vitro* anti-cancer activity on T47D and MCF-7 breast cancer cells, HeLa cells, and WiDr colon cancer cells. This study was conducted to examine the effect of NCE in inhibiting the proliferation of malignant lymphocytes in spleen induced by DMBA.

**MATERIALS AND METHODS**

**Materials**

*N. sativa* seed was oven-dried at temperature of 50 ± 1 °C. The powder was extracted by maceration technique using petroleum ether for 1x24 hours and continued using chloroform for 3x24 hours. DMBA was obtained from Sigma (Saint Louis, MO). DMBA and *N. Sativa* chloroform extract (NCE) were dissolved in corn oil.

**Animals**

*Sprague Dawley* female rats weighed 75±15 g. were obtained from Laboratory of Pharmacology and Toxicology, Faculty of Pharmacy, Universitas Gadjah Mada, Yogyakarta, Indonesia. The rats were quarantined for at least one week.

Sixty rats were divided into five groups (12 rats per group). The experimental design is summarized in Fig.1. For solvent control (group 1), rats were administered with corn oil. Meanwhile, for treatment groups (groups 2, 3, 4 and 5), rats were induced by oral administration of 20 mg/kgBW DMBA (Sigma), twice a week, for 5 weeks, then were treated with oral administration of 0 (Group 2), 250 (Group 3), 500 (Group 4) and 750 (Group 5) mg/KgBW NCE, once a day for 6 weeks, started from 1 week before DMBA. BWs were recorded weekly.

**Haematoxylin & Eosin Staining**

At the end of the experiment (16 weeks) all rats were sacrificed by decapitation. Spleens were then removed and fixed in 10% buffered formalin. After 12-24 h of fixation, tissues were embedded in paraffin, sliced with thickness of 3-5 μM, mounted on glass slides, processed further for H&E staining.

**AgNOR Staining**

AgNOR staining was performed according to the modified method (Pich, *et al.*, 2003; Bankafalvi, *et al.*, 2003). The staining solution was prepared by mixing one part of 2% gelatin in 1% formic acid with two parts of 50% aqueous silver nitrate (AgNO₃). All sections were sliced with thickness of 3 μm from routinely processed paraffin blocks. Sections were immersed in sodium citrate buffer (pH 6.0) and were incubated for 20 minutes in autoclave (120°C, 1.1-1.2 Bar).

Sections were then covered with the staining solution at room temperature in the dark for 15-20 minutes. The specimens were then washed with 5% sodium thiosulfate and distilled deionized water, dehydrated through graded ethanol to xylene, and mounted in synthetic medium. AgNORs, which appeared as dots both outside and within the nucleoli, were counted according to the description of previous report (Rizali and Auerkari, 2003).

A minimum of one hundred nuclei per specimen were observed randomly in three different views. mAgNOR is the mean of number of black dots observed in a cell, computed from total amount of blackdots (minimal 100 cells) divided with amount of cells. All specimens were observed on a binocular microscope (Olympus® DP12 microscope digital camera system, NY) with an immersion oil lens at magnification of 1000 x.

**Statistical Analysis**

A statistically significant BW difference was evaluated by ANOVA, continued with HSD (*p*<0.05) between groups was considered statistically significant using SPSS (SPSS version17.0; SPSS Inc.).

**RESULTS AND DISCUSSION**

**Effect of DMBA on Female Rats BW**

There was no direct evidence of toxicity due to NCE treatment. BW changes of animals treated with corn oil, DMBA, and DMBA+NCE were shown in (Table 1). In the beginning of the study, BW of corn oil group was different from other groups (*p*<0.05). Final BW showed that DMBA group was not significant different from DMBA+NCE groups (*p*<0.05) (Table 1).
Table I. Effect of DMBA, NCE & Corn Oil on Rats-BW

| Group                        | BW Before Treatment (g) | BW After Treatment (g) |
|------------------------------|-------------------------|------------------------|
| DMBA                         | 90.000 ± 15.374         | 121.429 ± 68.947       |
| DMBA + 250 mg/Kg BW NCE      | 90.000 ± 20.449         | 112.000 ± 58.981       |
| DMBA + 500 mg/Kg BW NCE      | 78.333 ± 17.494         | 121.429 ± 63.883       |
| DMBA + 750 mg/Kg BW NCE      | 64.167 ± 15.050*        | 82.000 ± 46.212*       |
| Corn oil                     | 54.167 ± 9.003*         | 90.000 ± 35.707*       |

Values are expressed as mean±SD; n=12, * Statistical significant in comparison to DMBA group p < 0.05
Histopathological Studies of Spleen

Histopathology profiles of spleen treated with DMBA and DMBA+NCE were depicted in Fig. 2. In general, H&E staining showed no morphological differences between DMBA and DMBA+NCE-treated groups. There were only some spleens in DMBA and and DMBA+NCE groups showed necrosis cells (Fig. 2).

Figure 2. Histological evaluation of spleen tissues. (A) Corn oil, (B) DMBA control group, (C) DMBA+250 mg/KgBW NCE; (D) DMBA+500 mg/KgBW NCE, (E) DMBA+750 mg/KgBW NCE. Black arrow shows necrotic cell with magnification 1000x

Anti-Proliferative Activity of NCE

AgNOR staining on spleen’s solvent (Fig.3) showed that DMBA group has numerous blackdots than other groups. mAgNOR scores (Table II) of corn oil and DMBA are 1.22 ± 0.12 and 1.92 ± 0.55 respectively. Treatments of 250 mg/KgBW, 500 mg/KgBW, and 750 mg/KgBW NCE exhibited a significant anti-proliferative activity of DMBA (Table II) with mAgNOR scores of 1.29 ± 0.39, 1.36 ± 0.08 and 1.33 ± 0.11 (p<0.05). In general, treatment of NCE inhibited proliferation of DMBA-induced spleenocytes.
DMBA is one of Polycyclic Aromatic Hydrocarbons carcinogenic compounds (PAHs) metabolized by cytochrome P450 (Rajapaksa, et al., 2007). DMBA is a substrate of the enzyme cytochrome P450 (CYP), CYP1A1, and CYP1B1 (Shimada and Guengerich, 2006). In the phase I metabolism, DMBA is changed to 8.9; 5.6; 3,4-epoxide DMBA by CYP1A1 and 3,4-epoxide DMBA by CYP1B1 (Hermawan, et al., 2011). DMBA enhances the production of reactive oxygen species (ROS). DMBA can also decrease the body's antioxidant level leading to radicals attack that cannot be neutralized by the body. It will cause cell damage and cell necrosis (Patri, et al., 2009). DMBA also increase intracellular Ca2+ which would interfere membrane permeability, increase the influx of extracellular fluid into cells, and reduce cell viability (Golstein and Kroemer, 2006). Therefore, membrane permeability will be impaired and increased intracellular fluid. Cell lysis will then be followed by the release of the contents of the cytoplasm. N. sativa is a plant that has antioxidant activity ability to protect the body from oxidative stress (Gilani, et al., 2004). It can be seen in treatment groups, which have less necrotic cells compared to DMBA groups.

Cellular proliferation requires an orderly progression through the cell cycle, primarily driven by protein complexes composed of cyclins and cyclin-dependent kinases (CDKs). Progression through the G1-S transition requires the activity of at-least two different types of kinases, cyclin D-Cdk4/6 and cyclin E/A-Cdk2. N.sativa decreases the expression of cyclin D1 (Aggarwal, et al., 2008) and inhibits cell proliferation, leading to cell cycle arrest (Ilairaja, et al., 2010). Treatment groups
showed less black dots number than control (DMBA group).

Tumor cells evolve a variety of strategies to limit or circumvent apoptosis. The most common hallmark is the loss of p53 tumor suppressor function. Tumors may achieve similar ends by increasing expression of anti-apoptotic regulators (Bcl-2, Bcl-xL) or survival signals (Igf1/2), by downregulating proapoptotic factors (Bax, Bim, Puma), or by short-circuiting the extrinsic ligand-induced death pathway (Hanahan and Weinberg, 2011). *N. sativa* stimulates apoptosis by increasing p53 expression (Yazan, et al., 2009; Ilaiyaraja and Khanum, 2010) and inhibiting anti-apoptotic proteins (Ivankovic, et al., 2006).

*N. sativa* stimulates the expression of GST (Ilaiyaraja and Khanum, 2010) which will conjugate DMBA compounds to prevent DMBA from binding with DNA, RNA, or protein (Murray, et al., 2006). GST is a phase II metabolic enzyme that detoxify carcinogens and facilitate their excretion by promoting the conjugation of electrophilic compounds with glutathione. GST deactivates and protects the surrounding tissues from mutagenesis and carcinogenesis.

Our current results showed that oral administered of NCE decreased necrosis and proliferative activity of DMBA-induced rat spleenocytes. These results indicated that NCE has a potency to be developed as a chemopreventive agent.

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