Arteether Exerts Antitumor Activity and Reduces CD4⁺CD25⁺FOXP3⁺ T-reg Cells in Vivo

Maryam Azimi Mohamadabadi¹, Zuhair Mohammad Hassan¹*, Ahmad Zavaran Hosseini¹, Mehrdad Gholamzad¹, Shekoofe Noori², Mehdi Mahdavi³, Hamidreza Maroof⁴

¹Department of Immunology, School of Medical Sciences, Tarbiat Modares University, ²Department of Biochemistry, School of Medical Sciences, Shahid Beheshti University, ³Department of Virology, Pasteur Institute of Iran, Tehran, ⁴Department of Microbiology, Zanjan Branch, Islamic Azad University, Zanjan, Iran

ABSTRACT

Background: Chemo-immunotherapy is one of the new achievements for treatment of cancer, by which the success of anti-cancer therapy can be increased. In vitro studies have been shown that Arteether (ARE) induces apoptosis in tumor cells, but not in normal cells. Objective: To investigate the cytotoxic and immunomodulatory properties of Arteether in-vivo and in-vitro. Methods: In this study, we used MTT assay for evaluation of cytotoxicity of Arteether on tumor cell line and Peripheral Blood Mononuclear Cells (PBMCs) from healthy individuals. Balb/c mice were subcutaneously transplanted with tumor tissue taken from Spontaneous Mouse Mammary Tumor (SMMT) bearing female mice. Arteether was administered to breast tumor-bearing Balb/c mice at a dose of 6 mg/kg/day intraperitoneally. Tumor sizes, lymphocyte proliferation, cytokines production, and the percentage of splenic T-reg cells were measured. Results: We observed that ARE could reduce the cell growth of 4T1 cell line in a dose-dependent manner but it had no cytotoxic effect on the growth of peripheral blood lymphocytes. ARE administered intraperitoneally to tumor-bearing Balb/c mice could reduce the tumor growth rate and splenic T-reg cells. No difference in the IFN-γ, IL-10 and IL-4 production was observed between tumor antigen-stimulated splenocytes of mice treated with ARE and control mice. Conclusion: These results underscore antitumor properties of Arteether that may aid in development of more effective antitumor agents.


Keywords: Arteether, Breast Cancer, Immunotherapy

*Corresponding author: Dr. Zuhair Mohammad Hassan, Department of Immunology, School of Medical Sciences, Tarbiat Modares University, Tehran, Iran, Tel: (+) 98 21 82883565, Fax: (+) 98 21 82884555, e-mail: hasan_zm@modares.ac.ir
INTRODUCTION

Among all types of cancer, breast cancer is one of the most important causes of women’s death (1); the increasing speed of cancer research has failed to keep pace with breast cancer growth, leaving many people suffering from it. Hence, cancer control is a major health care priority. There are several cancer therapy methods like surgery, chemotherapy, and radiotherapy, to name a few, but each of them has a specific shortcoming; for example, metastasis impedes surgical success. In addition, chemotherapy not only has a non-specific mode of killing dividing cells, but also is toxic to normal cells (2). It is therefore important to develop new methods for cancer therapy that targets tumor cells in a specific manner. Artemisia annua in Chinese traditional medicine is used for extracting Artemisinin (ART), which its anti malarial activity was published in early 1980s. Due to its molecular structure, Artemisinin has low solubility in water and oil, and much research has focused in modifying its structure in order to enhance its solubility. Many structural derivatives of Artemisinin have been synthesized since 1980. Most acknowledged derivatives including Arteether are very active against acute malaria and possess very low side effects. Arteether (ARE) is an ethyl ether derivative of dihydroartemisinin, a metabolite of Artemisinin, which is characterized by a rapid oral absorption due to its high hydrophobic properties (3-6). Alcohol soluble derivatives of Artemisinin also show effects against malignant cells. Although their mode of action in these cells is not clear; recent studies have been focusing on the antitumor activity of Artemisinin's derivatives, especially Arteether; which show an extreme anticancer activity in vitro and in vivo.

ART and ARE are sesquiterpene lactones with a highly active endoperoxide bridge. They can produce great amounts of reactive oxygen species in the presence of iron (7-9), which is distinct from other anti-malarial drugs, a property most interesting to cancer therapists. To facilitate the rapid growth, tumor cells need to increase the influx of iron; these cells have higher amounts of transferrin receptors and intracellular iron, hence increasing sensitivity of tumor cells to Arteether's killing mechanism as opposed to normal cells (9).

In general, tumor cells utilize the regulatory mechanism of T-reg cells to evade the immune system. That is why many studies have focused on methods that can reduce the number of T-reg cells in the tumor (10). Artemisins and its derivatives also posses immunomodulatory effects in a dose- and time-dependent manner; although their stimulatory effect on T-lymphocyte-mediated immune responses has also been reported in a study by Yang et al. (11-15).

In this study, we investigated the cytotoxic and immunomodulatory properties of Arteether in vivo.

MATERIALS AND METHODS

Mice. A group of inbred female BALB/c mice, at 4-6 weeks of age, were purchased from Pasteur Institute of Iran. Animal care and treatment were conducted in compliance with the guideline of Animal Care and Research Committee of Tarbiat Modares University, which is in conformity with the Guide for the Care and Use of Laboratory Animals (DHEW Publication No. (NIH) 85-23, Revised 1985, Office of Science and Health Reports, DRR/NIH, Bethesda, MD 20205) THE.
Arteether. Arteether powder was purchased from Exim-Pharm International Co., India. It was freshly prepared for each administration by dissolving the powder in absolute ethanol (Merck Co.), and further dilution in PBS (40:60). The solution was checked for endotoxin and mycoplasma contamination in our lab and Pasteur Institute of Iran, respectively. Solubilized Arteether was then filter sterilized to avoid any contamination.

Cell culture. Mice breast cancer cell line (4T1) was purchased from Pasteur Institute of Iran. The cells were cultured in RPMI 1640 medium (Gibco), supplemented with 10% FCS, 100 μg/ml streptomycin, and 100 IU/ml penicillin. All cells were grown in a humidified atmosphere containing 5% CO2 at 37°C. Normal lymphocytes were collected from healthy individuals in the Medical School clinic.

Cell Viability Analysis. Cell viability was evaluated by MTT viability assay. Briefly, growing cells were re-cultured (10000 cells/well) in 96-well tissue culture microplates (Nunc, Denmark). Three doses of ARE was added to the respective wells. After 48 and 72 hours, 20 μl MTT (Sigma-Aldrich) was added to each well with a final concentration of 5 mg/ml. After 4 h incubation, the medium containing MTT was discarded and 100 μl DMSO (Sigma-Aldrich) was added. MTT crystals were completely solubilized with vibration for 10 min. Absorbance was measured at 450 nm using ELISA reader and viability was expressed as stimulation index (SI), using the following formula:

\[ SI = \frac{OD \text{ of ARE+ cells}}{OD \text{ of ARE- cells}} \]

In Vivo Breast Cancer Model. Spontaneous mouse mammary tumor (SMMT) is an invasive ductal carcinoma, which develops spontaneously in female BALB/c mouse after transplantation with tumor cells. SMMT is regarded to be similar to the invasive ductal carcinoma observed in human samples (16).

In our study, mice were subcutaneously transplanted with 0.5 mm³ of tumor tissue taken from SMMT-bearing BALB/c mice. Tumors were allowed to reach an approximate size of 500 mm³. Ten tumor-bearing mice were randomly divided into two groups, each consisting of five mice. The first group was treated with 0.1 ml of Arteether at a dose of 6 mg/kg/day via the intraperitoneal (IP) route. The controls received ARE diluents in the same volume and via the same route. The treatments were applied for thirteen consecutive days. The tumor volumes were measured every other day using digital vernier callipers (Mitutoyo, Japan). The volumes were calculated with the following formula (17):

\[ V = \frac{1}{2} (LW^2) \]


Antigen Preparation. A volume of approximately 3000 mm³ tumor tissue was extracted from the breast cancer-bearing BALB/c mouse. The tumor suspension was prepared using five rounds of freezing/thawing and was passed through a 150 μm stainless steel mesh. The suspension was sonicated in a power of 4W for 30 seconds, followed by a 20-second pause for five times. To inactivate serine proteases, 1 mM of phenyl methyl sulfonyl fluoride (PMSF) was added to the cell lysate. The extract was filtered through a 0.22 μm filter and its concentration was determined using the Bradford method. Finally, it was stored at -20°C until use.

Separation of Splenic Mononuclear cells (MNC). The mice were sacrificed on day 13; spleens were resected under sterile conditions and were suspended in PBS. The splenic cell suspension was RBC-lysed with a solution of 0.75% NH4Cl and Tris buffer (0.02%) (pH=7.4). The cells were washed and the single-cell suspension was prepared...
in RPMI 1640 containing stable glutamine (Cytogen) and 10% heat-inactivated fetal calf serum (Gibco, UK). To define the viability and density of cells in the suspension, Trypan blue dye exclusion method was used. The cells were counted using a hemocytometer by light microscopy method. The viability of splenocytes was generally above 95%. After additional washing, the suspension was adjusted to $4 \times 10^6$ cells per milliliters in RPMI 1640 supplemented with 10% FCS, 100 µg/ml streptomycin, and 100 IU/ml penicillin (complete RPMI), and kept at 4°C.

**Lymphocyte Proliferation Assay.** Splenic MNCs were cultured in RPMI 1640 (Gibco, UK) (adjusted to $2 \times 10^6$ cells/ml in RPMI supplement). Briefly, 100 µl of cell suspensions were transferred to 96-well flat-bottom microplates (Nunc, Denmark). 6 wells were considered for each sample. 20 µl of tumor antigen was added to three wells of each sample. Plates were incubated at 37°C (separate wells were cultured with untreated normal splenocytes and incubated with PHA as positive control). Proliferation assay was done using Roche Cell Proliferation ELISA, BrdU kit according to the manufacturer's protocols. The absorbance of the samples was measured at 450 nm using ELISA reader. Results are presented as stimulation index (SI), using the following formula:

$$SI = \frac{OD \text{ of tumor Ag}^+ \text{ splenocytes}}{OD \text{ of tumor Ag}^- \text{ splenocytes}}$$

**Splenocyte Cytokine Production.** The isolated spleen MNCs were cultured in 24-well plates (Nunc, Denmark) in a final concentration of $2 \times 10^6$ cells/ml. To stimulate the cell, 20 µl of purified tumor antigen was added to each well, the supernatants were collected after 72 h incubation at 37°C and 5% CO2 (11), and were kept frozen at -70°C until used. IFN-γ, IL-4 and IL-10 concentrations were measured using R&D DuoSet ELISA Development kit according to the manufacturers’ protocol. Each sample was analyzed in triplicate.

**Three-Color Immunostaining and Flow Cytometry Analysis.** The MNCs, purified from mice spleens, were immunostained with FITC anti-mouse CD4, PE anti-mouse CD25, and subsequently PE-Cy5 anti-mouse Foxp3, according to the instructions given in the eBioscience mouse regulatory T cell staining kit. The samples were analyzed by FACS Calibur flow cytometer (BD, USA) and the results were analyzed using WinMDI software.

**Statistical Analysis.** In this study, each experiment was performed in duplicate or triplicate, using either one-way analysis of variance (ANOVA) or the Mann-Whitney non-parametric test, to determine the statistical significance ($p<0.05$) between the test and control groups. The data were analyzed using SPSS software version 16. The results are expressed as the mean ± standard error (mean ± SE).

**RESULTS**

**Effect of Arteether on Cell Growth.** To evaluated the effect of ARE on the growth of tumor cells and peripheral blood lymphocytes, three doses of ARE (10-20 µM) was applied in vitro. The result showed that ARE could reduced the cell growth of 4T1 cell line in a dose-dependent manner but it has no cytotoxic effects on the growth of peripheral blood lymphocytes (Figure 1).
Figure 1. Cytotoxicity effect of arteether against 4T1 cell line. Cells were incubated with increasing concentration (10-20 μM) for 48 and 72h. Graph shows the significant difference between ARE-treated 4T1 cell line and ARE-treated normal cells (p<0.05).

**Effect of Arteether on Lymphocyte Proliferation Assay.** In order to assess the effect of ARE on specific Lymphocyte proliferation, ten mice were used and the protocol in figure 2 was applied. Our results indicated a slightly increase in SI at the dose of 6 mg/kg but no significant difference was observed (p>0.05).

Figure 2. The results of lymphocyte proliferation assay: ARE-treated group showed no significant differences in comparison with control (p>0.05). Positive group (PHA) showed a significant difference (p<0.05). Tip=arteether-treated group (Test, IntraPeritoneally), Cip=control group (Control IntraPeritoneally).
The results of proliferation assay were evaluated in triplicate after stimulation with a specific tumor antigen. **Effect of Arteether on Lymphocyte Cytokine Production.** The concentration of IFN-γ and IL-4, typical cytokines produced by Th1 and Th2 cells, in both treated and untreated tumor-bearing mice were evaluated by ELISA technique. The result have shown no significant difference (p>0.05) between the two groups.

**Figure 3.** The results of cytokine ELISA assay: the levels of IFN-γ and IL-4 cytokines produced from stimulated splenocytes by specific tumor antigen are shown. Result showed no statistically significant difference between the ARE-treated group and untreated group (p>0.05).

The concentration of IL-10 as an important mediator of regulatory T cells was also detected in a same manner, our results indicated that ARE could decrease the level of IL-10 in the treated group, but this difference was not statistically significant (p>0.05). The results are shown in Figures 3 and 4, respectively.

**Figure 4.** The results of cytokine ELISA assay: the level of IL-10 cytokine produced from stimulated splenocytes by specific tumor antigen is shown. Result showed no statistically significant difference between the ARE-treated group and untreated group (p>0.05).
Effect of Arteether on Splenic CD4+CD25+FoxP3+ T cells. The flowcytometry technique was used to define the percentage of splenic CD4+CD25+Foxp3+ T cells (T-reg) in tumor-bearing mice. As shown in Figure 5, the results indicated a significant difference in the percentage of splenic T-reg cell between the ARE-treated group and the untreated group; i.e., the percentage of splenic T-reg in the treated group was lower than the untreated group (p=0.049).

**Figure 5.** The frequency of T regulatory cells measured by flow cytometry assay, A. graph showing the percentage of splenic CD4+CD25+FoxP3+ T cells (T-reg) in ARE-treated group and control. Result showed a statistically significant difference between the percentage of splenic T-reg cells in ARE-treated group and untreated group (p<0.05). B: Dot plots representing the percentages of CD4 and Foxp3 expressing cells on the CD25 gated population from treated group (right) and non treated group (left).
Effect of Arteether on Tumor Volume. To define the effect of ARE on the tumor volume, 10 mice were used. ARE at a dose of 6 mg/kg/day was intraperitoneally injected for 13 consecutive days. The changes in the tumor volume during the experiment period are shown in figure 6. Our results indicated that there was a significant difference in the tumor volume between the treated and untreated groups, detected from day 6 of measurement (p<0.05). The results showed that ARE administration could decrease tumor growth in the treated group.

Figure 6. The results of tumor size measurement: graph showing the Mean ± SE of tumor volume in 2 groups of mice (n=5 for each group) in 13 consecutive day. 6 mg/Kg/day of ARE and ARE diluents (PBS:ethanol 60:40) were intraperitoneally injected to ARE- treated group and control, respectively. A statistically significant difference was detected between the ARE-treated group and control from 6th day of measurement (p<0.05)

DISCUSSION

During the past decade, medical researchers have focused on herbal medicine, especially Artemisinin and its derivative Arteether, which its anti malarial activity dates back to 1596. Li Shizen, a famous researcher used the plant Quighaosu, for treating fever and chill. In addition to their anti-malarial activity, research has shown that these extracts exert a profound cytotoxicity towards tumor cells (18-22). In the present investigation we explored the cytotoxic and immunomodulatory activities of ARE in vivo (in a mouse model of breast cancer). At first, to define the effect of arteether on tumor cell growth in vitro, MTT assay was applied. Our result showed a significant and selective cytotoxicity of ARE against 4T1 cell line. This result was not unexpected: our previous studies also showed that artemisinin and its derivatives have a selective
cytotoxicity toward tumor cells owing to their novel killing mechanism. Based on our previous findings and other published studies (9,23-25), the dose of 6 mg/Kg was identified as the optimal immunostimulatory dose, and was used for further tests. The recent studies show that ART and its derivatives have a different dose-dependent pattern on the production of hemagglutination antibody (14,15), which was confirmed by our previous results. The same result has also been established for other antitumor agents (26).

Lymphocyte proliferation assay showed no statistically significant difference in the ARE-treated group in comparison with the untreated group; a similar result has previously been reported in a study of Artemisinin and its derivatives, suggesting that they had no effect on lymphocyte proliferation (11). As shown in Figure 5 and Figure 6, the levels of IFN-γ, IL-4 and IL-10 cytokines were not statistically significant, while we indicated a significant increase in the level of IFN-γ after the treatment with Artemisinin. It may be that the dose-dependent manner of ARE and other derivatives or increasing in hydrophobicity can reduce the intracellular communication of ARE. In another study, we noticed that an Artemisinin derivative, Artemetheter, had no effect on the pattern of cytokine production (27), while other investigations indicated that these derivatives have different immunomodulatory or immunosuppressive effects on the immune system based in their structure and dose (13,14). Nevertheless, further investigation is needed in order to determine an effective and immunostimulator dose of ARE.

We indicated that ARE can decrease the tumor growth rate in tumor transplanted mice. This result is not unexpected; our previous studies also showed that the dose-dependent use of ART derivatives is more effective in reducing the tumor mass than ART (11,27). We also showed that ARE could decrease the total number of splenic T-reg cells in the treated group in comparison with the untreated group. Although the mechanism by which ARE reduces T-reg cells has yet to be understood, the decrease in T-reg cells may be because of the tumor mass reduction. However, other studies reported that ART derivatives can reduce the number of T-reg cells through inhibiting the production of NO, a inducer metabolite for T-reg expansion, from macrophage cells (28,29).

In conclusion, our results showed that ARE is more effective in slowing down the tumor growth than to be an immunomodulator. Decrease in the level of IFN-γ, IL-4, IL-10 and T-reg cell expansion were also noted; and these abilities might be ascribed to the hydrophobic properties of ARE. Our findings in this study are consistent with our previous findings about Artemether, a methyl ether derivative of ART (27).

ACKNOWLEDGEMENTS

In present study financial support was provided by Tarbiat Modares University and we also thank Dr. Ladan Langroudi for technical assistance.

REFERENCES

Anti-tumor properties of Arteether


