Effect of treatment Osteosarcoma (Mg-63) cells with Vitamin C, Green tea and their combination (An-in vitro study)

Hiam R. Hussien*, Amr H.M. El Bolok, Sherif F. Elgayar and Maii I. A. Sholqamy

Introduction

We deal with cancer as it is one of the major causes of death all over the world [1]. Osteosarcoma is the most common primary pediatric bone malignancy, derived from primitive bone-forming (osteoid producing) mesenchymal cells [2]. Until now the real cause of bone cancer is unknown, this may due to different genetic types with lack of molecular alterations [3], [4]. For several years ago, natural plants and their extracts have been used for various treatment purposes such as cancer [5]. In ancient years, various studies were confirmed that different concentrations of vitamin C inhibited the tumor growth and caused cancer death [6]. Different compounds are considered to be anticancer, polyphenols as gallacetehins, flavonoids and tannis are type of those compounds [7]. Green tea are added to a person’s nutrition for health improvement, Gallacetehins are present in green Tea, it is thought including polyphenols in a person’s diet can improve health and reduce risk of cancers by being natural antioxidants [8].

Aim

The aim of the current study is to investigate the effect of vitamin C, green tea and their combination on Mg-63 cell line.

Materials and Methods

(A)Cell line:-

Mg-63 osteosarcoma cell line was obtained from Cell Culture Department (NAWA SIENTIFIC) in the form of frozenvial.in vitro cultivation of human tumor tissues (Osteosarcoma).

(B)Drugs:-

Mg-63 Cells treated with vitamin C, green tea extracts and their combination with IC50 concentration.

1. Cytotoxicity Assay principal

SRB assay: -

It is SRB Sulforhodamine B colorimetric assay for cytotoxicity screening, used One of the most widely used methods for in vitro cytotoxicity screening. The assay relies on the ability of SRB to bind to protein components of cells that have been fixed to tissue-culture plates by trichloroacetic acid (TCA). SRB is a bright-pink amino-xanthene dye with two sulfonic groups that bind to basic amino-acid residues under mild acidic conditions, and dissociate under basic conditions. Cells were maintained in DMEM media supplemented with100mg/mL of streptomycin,100units/mL of penicillinand10%ofheat-inactivated fetal bovine serum in humidified, 5% (v/v) CO2atmosphere at 37° [9],[10].
Microscopic examination

Hematoxylin and eosin staining of MG-63 cell line :-

Hematoxylin and Eosin stain was performed according to Alshehri et al., (2019) where IC50 treated and untreated cells were harvested by cold centrifugation (Jouan-Ki-22,France) residual adherent cells were trypsenized and processed in the same way . [11] Deposited cells were re-suspended in 0.5 ml of phosphate buffer saline (PBS) Fifty microliters of treated and untreated cells were dispersed and spread in circular way on clean slides (3 for each treatment). Slides were air-dried, methanol fixed and rehydrated in descending concentrations of alcohol (100%, 90%, 75% and 50%). Slides were washed in distilled water for 5 min [11]. The slides were immersed in filtered hematoxylin (HE) stain for 3 min and washed with distilled water twice. Slides were immersed in filtered eosin stain for 5 seconds and washed with distilled water. Dried slides were immersed in xylene, mounted with Canada balsam then coverslips were placed and left to dry. Five microscopic fields of each slide were photomicrographed using the power of 400×. This was done using a digital camera (Canon, Japan), which was mounted on a light microscope. Images were transferred to the computer system for analysis. Field selection was based on the presence of the highest number of apoptotic cells. The photomicrographs were qualitatively evaluated for the presence of morphological criteria of apoptosis.

(3) Autophagy assay.

Autophagic cell death is assessed using acridine orange lysosomal stain coupled with flowcytometric analysis. After treatment with test compounds for 24/48or72 cells (105 cells) are collected by trypsinization and washed twice with ice-cold PBS (pH 7.4). Cells are stained with acridine orange (10 µM) and incubated in dark at 37°C for 30 minutes. After staining, cells are injected via ACEA Novocyte™ flowcytometer (and acridine orange fluorescent signals is analyzed using FL1 signal detector (λex/em 488/530 nm). For each sample, 12,000 events are acquired and net fluorescent intensities (NFI) are quantified using ACEA NovoExpress™ [12].

Results:

1. Cytotoxicity results:-

SRB Assay was employed. Data obtained revealed cytotoxic effect of Mg-63 cells with vitamin C, green tea and combination. Tab.(1)

2. Microscopic examination: -

Effects of different concentrations of vitamin, green tea and their combination by microscopic evaluations. Figs. (1),(2),(3),(4),(5).

3. Autophagy results:-

Autophagic cell death assessment in Mg-63 cells treated vitamin C, green tea and their combination with (IC50 µM) using the acridine orange lysosomal stain coupled with the flow cytomteric analysis. The green curve for the negative control (untreated), and the red one for the treated cells. Vitamin C caused autophagic cell death followed by combination then least effect was green tea. Fig.(6) Tab. (2)

(1)Fig1

Figure 1: A photomicrograph showing malignant cells showing hyperchromatism and cellular and nuclear pleomorphism (H and E, Original magnification 100X oil).

(2) Fig2

Figure 2: A photomicrograph showing sheets of malignant cells. Shrunken apoptotic cells with irregular cellular and nuclear membranes (Yellow arrows) (H and E, Original magnification 100X Oil).

(3)Vitamin C

Figure 3: A photomicrograph showing malignant cells. Few apoptotic cells could be detected showing cellular shrinkage (Yellow arrow) and peripheral condensation of chromatin (Red arrow) (H and E, Original magnification 100X Oil).
(4) Green tea

![Figure 4: A photomicrograph showing sheets of malignant cells. Shrunken apoptotic cells (Yellow arrows) with peripheral condensation of chromatin (Red arrows) could also be detected. An apoptotic body could also be seen (Green arrow) (H and E, Original magnification 100X Oil).](image)

(5) Combination:

![Figure 5: A photomicrograph showing sheets of malignant epithelial cells. Few apoptotic cells could be detected with irregular cellular and nuclear membranes (red arrows) (H and E, Original magnification 100X Oil).](image)

3. Autophagy results:

Autophagic cell death assessment in Mg-63 cells treated with vitamin C, green tea and their combination with (IC50 μM) using the acridine orange lysosomal stain coupled with the flow cytometric analysis. The green curve for the negative control (untreated), and the red one for the treated cells. Vitamin C caused autophagic cell death followed by combination then least effect was green tea.

Table (2)

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Discussion:

The aim of the present study is to investigate the cytotoxic effect of vitamin C, green tea and their combination on Mg-63 osteosarcoma cell line. We confirmed that all three drugs have cytotoxic effect but with different degree, resulting in cell cycle arrest in with inhibition of mitosis followed by inducing different types of cell death such as apoptosis, necrosis and autophagy but with varying degree.

Our data explained clearly the effect of in vitro cytotoxic effects of vitamin C green tea and their combination on Mg-63 osteosarcoma cell line after 72 hours. Photomicrography and cytological evaluation were done to confirm and clarify SRB results. Careful examination revealed that, the major criteria of apoptosis and necrosis were showed among different concentrations of the three drugs.

Microscopic examinations were done, in morphological analysis, untreated control cells showed a relatively some morphological changes in the form of nuclear and cellular pleomorphism due to the malignant nature of the cells. Only few control cells showed early morphological characteristics of apoptosis such as cellular and nuclear chromatin condensation and apoptotic bodies. Fig. (1), (2)

Cells treated with vitamin C of showed sheets of malignant cells. Few apoptotic cells could also be detected. A photomicrograph showing malignant cells. Few apoptotic cells could be detected showing cellular shrinkage and peripheral condensation of chromatin. Figs. (3)
Cells treated with green Tea reflected sheets of malignant cells. Apoptotic shrunken cells could be detected. The cells show peripheral condensation of chromatin and irregular cell and nuclear membranes. A photomicrograph showing sheets of malignant cells. Shrunken apoptotic cells with peripheral condensation of chromatin and apoptotic bodies were detected. Fig.(4)

Cells treated with combination explained sheets of malignant cells. Few apoptotic cells could be detected in this group. A photomicrograph showing sheets of malignant cells. Few apoptotic cells could be detected with irregular cellular and nuclear membranes. Fig.(5)

These indicated that these drugs induced in vitro cell death by three different mechanisms of autophagy apoptosis and necrosis. All of these results are in agreement with the following :- (O'Leary BR et al.,2020 ; Ma E.,et al.,2014 and Venturelli S.,) [13],[14],[15] confirmed that vitamin C possess anti-cancer effect and they explained their confirmations by examining the effect of vitamin C on various cancer cell lines of different type of tumor.

( Belin et al., 2009; Chen et al., 2005, 2008; Valenti et al., 2014), [16],[17],[18], studied the antitumor effect vitamin C on another osteosarcoma cell line (G292) as the following:- that vitamin C caused inhibition of cell growth of different osteosarcoma cell line such as G292 this inhibition is dose dependent of vitamin C resulting in decrease growth of tumor cells in low concentrations of vitamin C and apoptosis of tumor cells in high concentrations., this also confirmed by Wang J. et al., 2018, Ni J et al., 2018 and Luo kw. et al.,2017,[19],[20],[21].


Furthermore, (Liu L et al.,2015 and Tsao A S.),[22],[23] demonstrated that in these GTPs affect cell cycle regulation and caspase-3 that has essential role in apoptosis. Observing the results of our study, we concluded that autophagy cell death was induced with treatment of Mg-63 cells with vitamin C using acridine orange in relation to control group. Fig.(6B) also we noticed that Mg-63 cells treated with green tea produce autophagy but with lower percentage than that of vitamin C. Fig.(6C) when we used Mg-63cell cells treated with combination , autophagy cell death was observed but not more than Mg-63 cells treated with vitamin C and greater than green tea. Fig.(6D).

These results are in agreement with (Valenti MT, et al.2014) [18] reported that vitamin C used in treatment of osteosarcoma were able to produce apoptosis and autophagic cell death in vitro.

Wang, X et 2019 [19] assured that polyphenols by being the main component of GTPs contain special properties as they cause autophagic cell death in cancer death.

Conclusions

From the results of the current study we found that vitamin C, green tea and their combination have cytotoxic effect on Mg-63 cell line but with different levels, also induced apoptosis, necrosis and Autophagy.

List of Abbreviations

OS: Osteosarcoma.

GT: green tea.

IC50: Half-maximal inhibitory concentration.

SRB: Sulforhodamine B assay.

TCA: trichloroacetic acid.

DMEM: Modified Eagle Medium.

PI: Propidium iodide.

DNA: Deoxyribonucleic acid.

RNA: Ribonucleic acid.

DEMSO: Dimethyl Sulfoxide.

PBS: Phosphate Buffer Saline.

NFI: Net fluorescent intensities.

HE: Hematoxylin eosin.

GTPs: Green tea polyphenols.

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Sincerely,

Dr. Hiam Rifaat Hussein.

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