The Effects of Ara-C, Simvastatin and combo therapy on energy metabolism of HL-60 promyelocytic leukemia cell lines

[H.60 lösömi hücrelerinde Ara-C ve Simvastatinin tek tek ve kombine kullanımlarının enerji metabolizması üzerindeki etkileri]


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ABSTRACT

Objective: Cancer cells choose their metabolic pathway depending on the oxygen content and substrate concentration of the medium. A wide spectrum of therapeutic agents regulating the energy metabolism of cancer cells are in still in use. Cytosine arabinoside (Ara-C) is a pyrimidine analogue used in the treatment of acute myeloid leukemia (AML) and simvastatin is an inhibitor of HMG-CoA (3-hydroxy-3-methyl-glutaryl-CoA) reductase, which regulates cholesterol biosynthesis. Thus, this study aimed to assess the energy metabolism of HL-60 promyelocytic leukemia cells and healthy white blood cells, additionally to determine the effects of simvastatin and Ara-C, alone or in combination on the energy metabolism of these cells.

Materials and Methods: Healthy white blood cells, untreated and treated HL-60 promyelocytic leukemia cell lines were incubated for 4 hours with radiolabelled glucose. Following incubation, lactate, which is one of the end products of the carbohydrate catabolism, and radiolabelled CO2 produced by cells were collected and measured by the liquid scintillation device. In addition, glycogen consumption per hour was determined in each group.

Results and Conclusion: We found that untreated HL-60 promyelocytic cells use anaerobic glycolysis pathway whereas healthy white blood cells use aerobic glycolysis for energy gain. It was concluded that combined use of Ara-C and Simvastatin might lead to significant increase in the rate of aerobic glycolysis of HL-60 promyelocytic cells and the metabolism of these leukemia cells become more similar to the metabolism of healthy white blood cells which they originate from.

Key Words: Warburg effect, HL-60 cell lines, energy metabolism, simvastatin, Ara-C.

Conflict of Interest: Authors declare no conflict of interest.

ÖZET


Gereç ve Yöntemler: Bu amaçla, ilaç kullanılmayan tüm HL-60 akut promyelositik lösemi hücre hataları,uida radyoaktif glukoz ile 4 saat inkübasıона bırakıldı. Inkübasyonun sonrasi, karbonhidrat katabolizmasının son ürünlerinin kolleksiyonunun yapılışı, hücreler tarafından üretilen radyoaktif işaretli CO2 toplanarak ölçülmiştir. Ayrıca, araçsal, simvastatin uygulamasının hücrelerin enerji metabolizmalarını etkilediğini bulmak amaçlandı.

Introduction

Many structural and metabolic changes take place during carcinogenic transformation of a normal cell under the influence of chemical agents, radiation and/or microbial carcinogens [1]. Metabolic changes are principally seen at the pathways cells use for energy gain. It was first reported by Warburg in 1930, at a publication named “Tumor Metabolism”, that the rate of anaerobic glycolysis is high in cancer cells [2]. Other studies performed in the following years also reported varying degrees of anaerobic glycolysis in cells undergoing cancer transformation, thus this condition was named as “the Warburg Effect” [3-5]. The choice of metabolic pathway depends mainly on the oxygen and nutrient concentration of the medium. When the oxygen concentration is insufficient, anaerobic glycolytic pathway, which has less energy yield than aerobic glycolysis, is activated so that cells can continue to survive under low oxygen concentrations. Depending on the low oxygen concentration, the choice of anaerobic pathway for ATP production is called the “Pasteur Effect” [6-8]. Apart from the “Pasteur Effect”, the nutrient concentration of the medium also affects the choice of the metabolic pathway even though the oxygen concentration is sufficient [9-12]. This characteristic is named as the “Crabtree Effect” leading these cells choose the anaerobic pathway under high nutrient concentrations and the aerobic glycolytic pathway under low nutrient concentrations [13-14]. Tumor cells usually choose anaerobic glycolytic pathway for energy gain as it is shorter, with the aim yielding their high metabolism while rapidly consuming the sources of the organism. However, when the nutrient concentration of the medium decreases the tumor cells choose the more energy yielding aerobic glycolytic way in order to survive [15-17]. Thus, knowing whether the tumor cell displays the Pasteur or Crabtree Effects is important in terms of the success of anti-cancer therapy.

Acute myeloid leukemia (AML) is one of the most common types of leukemia in adults. In younger patients, the prognosis is better than for their older counterparts. Cytosine arabinoside (Ara-C) is a deoxycytidine (dCyd) analog that remains one of the most effective drugs used in the treatment of acute leukemia as well as other hematopoietic malignancies. More recent work has attempted to improve the patient outcomes by adding targeted therapies to the current induction chemotherapy [18]. Simvastatin (Sim) is one of the statins, which are commonly used for treating hypercholesterolemia but also have some unexpected benefits such as reducing the overall cancer incidence and inducing apoptosis [19].

In this study, the effects of simvastatin, Ara-C, and combination of Simvastatin + Ara-C therapy over the energy metabolism of HL-60 acute promyelocytic leukemia cells and untreated control white blood cells were investigated.

Materials and Methods

Chemicals and Biomaterials:

D-(6-C14) Glucose (Amersham) was used for radiolabeling, hexokinase and glucose-6-phosphate dehydrogenase enzymes were obtained from Boehringer. All other chemicals used were analytically purified. HL-60 acute promyelocytic leukemia cells were from the Karolinska Institute, Hematology Department, Huddinge, Sweden and they were maintained at 37°C, at the RPMI 1640 medium supplemented with 10 % fetal calf serum, 2 mM L-Glutamine, 100 µg/mL streptomycin and 100 u/mL penicillin, in a medium containing 5 % CO₂.

MTT assay

The potential growth-inhibitory and cytotoxic effects of Ara-C (0.001-10 µM) and Simvastatin (0.0001-100 µM) in HL-60 cell line were evaluated using the the MTT (3-[4,5-dimethylthiazol-2-yl][2,5-diphenyl-tetrazolium bromide) assay according to the study of Mosmann et al [20]. Cells, at a concentration of 3×10⁶ cells/ml, were exposed to a concentration range of drugs and incubated at 37°C for 72 hr. Cell viability was expressed as IC₅₀, which was defined as the concentration of drug causing 50% growth inhibition.

Radioactive incubation and analysis of end products:

HL-60 cells were seeded at 1´10⁵ cells/mL in growth medium in 96-well microtiter plates (Costar Co., Cambridge, MA) and were divided into 4 groups. Control cells remained untreated, whereas Simvastatin (0.0001-100 µM) and Ara-C (0.001-10 µM) were added to treatment groups. The combo treatment regimen including both Simvastatin (0.1µM) and Ara-C (0.01µM), the doses equivalent to IC₅₀ values, was also tested on the cells. All the cells were incubated at 37°C in an atmosphere of 5% CO₂ for 5 days which was the commonly used incubation time reported in similar studies.

White blood cell isolation was performed on the blood samples of healthy donors and was used for the aerobic control group. Then, all cells were incubated with radioactive glucose [D-(6-C14) Glucose], at 37°C for 4 hours. The radiolabeled glucose was taken by the cells during the incubation and catabolized by the glycolytic pathway, finally converted to end products like lactate, acetate, and propionate. Following the incubation, the radioactive CO₂ left in the equipment was hold in the scintillation vials via the nitrogen. Then, the rest of the content was centrifuged at 3000g for 10 minutes. The supernatant and the cell pellet were collected into different tubes. Lactate, as the end product of anaerobic glycolysis was measured in the supernatants collected in the scintillation vials by anion exchange chromatography. The data was calculated with Microsoft excel hardware with standard graphics. The protein content of the cell pellets was measured by the modified Lowry
The glycogen amount in the cell pellet was measured with the enzymatic method of Hassid and Abraham [22].

**Statistical Analysis**

Mann-Whitney U-test was used to compare groups. P values less than 0.05 were considered significant.

**Results**

**The effects of Ara-C and combination therapy on cell proliferation**

The effect of different concentrations of Sim, Ara-C and combination therapy was determined in HL-60 cell line by MTT assay. Ara-C (0.001-10 μM) and Sim (0.0001 to 100 μM) alone inhibited cell proliferation in a concentration-dependent fashion in the range of concentrations used. IC$_{50}$ value for Simvastatin was found as 0.1μM and 0.01μM for Ara-C. Co-incubation of Sim (0.1 μM) with Ara-C (0.01 μM) resulted in a marked and significant inhibition of proliferation of HL-60 cells compared with Sim and Ara-C alone (p<0.001 for each) as determined by MTT assay and trypan blue dye exclusion assay run based on previous related studies [23, 24].

**The effects on energy metabolism**

We first compared the CO$_2$ and lactate production and glycogen consumption of healthy white blood cells (WBC) and the untreated HL-60 cells, and we observed statistically significant difference between their energy metabolisms. The results showed that the healthy white blood cells use the aerobic glycolytic pathway whereas the untreated HL-60 promyelocytic cell cultures principally use the anaerobic glycolytic pathway for energy gain. When HL-60 leukemia cells treated with Simvastatin or Ara-C or Sim + Ara-C were compared with the untreated HL-60 leukemia cells or healthy white blood cell cultures, the difference in CO$_2$ and lactate production, in addition to glycogen consumption was also statistically significant (Table 1).

![Figure 1: The amounts of CO$_2$ and lactate production (pmol glucose / h / μg protein) and glycogen consumption (nmol glucose / h / μg protein) of HL-60 leukemia cells after treatments for 5 days.](image)

Table 1. The amounts of glycogen consumptions and end-products of untreated control cells and treatment groups (n=7).

<table>
<thead>
<tr>
<th>Treatment</th>
<th>Glycogen consumption (nmol glucose / h / μg protein)</th>
<th>CO$_2$ Production (pmol glucose /h / μg protein)</th>
<th>Lactate production (pmol glucose /h / μg protein)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Leukocyte culture (WBC)</td>
<td>1026 ± 27</td>
<td>3522 ± 276</td>
<td>463 ± 57</td>
</tr>
<tr>
<td>HL-60 Control</td>
<td>2870 ± 152</td>
<td>1530 ± 379</td>
<td>43205 ± 6761</td>
</tr>
<tr>
<td>Ara-C</td>
<td>1854 ± 154</td>
<td>858 ± 276</td>
<td>1790 ± 112</td>
</tr>
<tr>
<td>Simvastatin</td>
<td>1796 ± 168</td>
<td>1094 ± 632</td>
<td>1715 ± 124</td>
</tr>
<tr>
<td>Ara-C + Simvastatin</td>
<td>1572 ± 81</td>
<td>1526 ± 268</td>
<td>1461 ± 101</td>
</tr>
</tbody>
</table>
Discussion

Increased anaerobic glycolysis rate of cancer cells was first reported by Warburg in 1930 [2]. Then new studies were planned to investigate the pathways for obtaining ATP during carcinogenesis, with the aim of developing more effective anti-cancer drugs. The results showed that cancer cells use different metabolic pathways for energy gain when compared with the normal cells they are stemmed from. Furthermore, as Warburg suggested, high anaerobic glycolysis rate can not be generalized for all cancer cells, in some cancer types like Morris Hepatoma, the glycolysis rate is normal and the mitotic division rate is low [25,26]. However, most cancer types principally use anaerobic glycolysis with varying degrees and the mitotic division rate is very high. Under normal conditions, the choice of metabolic pathway is related with the oxygen and nutrient concentration of the medium. When the ambient oxygen concentration is decreased and the glycogen and lipid depots of the organism are also rapidly consumed, cells which have undergone cancer transformation chose anaerobic glycolysis instead of aerobic glycolysis, which is known as the Pasteur Effect [27-33].

The Crabtree Effect which is not seen at normal cells but observed at cancer cells may suggest that the substrate concentration of the environment plays a principal role in the selection of metabolic pathway. As long as the substrate concentration is rich and sufficient, cancer cells use anaerobic glycolysis for obtaining ATP as seen frequently depending on the marked overexpression of type II hexokinase (HKII). In anaerobic glycolysis process, one mole of glucose is degraded into lactate and the net energy gain is 2 moles of ATP. Furthermore, the increase in lactate concentration causes a decrease in pH of the medium. Thus, in aerobic glycolysis, ATP gain is 36-38 moles for every mole of glucose, and the substrate consumption is 19 times higher than seen in physiological conditions. When the nutrient concentration of the medium is extremely decreased, the cell that have undergone tumor transformation chose aerobic glycolysis and start to use the nutrient content of the medium more effectively [34-39]. The change in the aerobic or anaerobic pathway limits the effect of chemotherapy which depends on the energy metabolism of cancer cells.

In recent years, increasing evidence suggests that statins have antiproliferative, antiangiogenic, and antitumoral properties in addition to their well-described antiatherosclerotic, antithrombotic, and anti-inflammatory properties. Statins have been shown to inhibit the growth of a variety of tumor cell types, including prostate, gastric, and pancreatic carcinoma, as well as colon adenocarcinoma, neuroblastoma, glioblastoma, mesothelioma, melanoma, and acute myeloid leukemia cells. Statins have also been shown to activate caspase proteases involved in programmed cell death, additionally upregulate proapoptotic protein expression and decrease antiapoptotic protein expression. Statins also impair the metastatic potential of tumor cells by inhibiting cell migration, attachment to the extracellular matrix, and invasion of the basement membrane. Interestingly, normal cells appear to be more resistant to the antiproliferative effects of statins relative to tumor cells [40]. Therefore, statins may thus represent a novel therapeutic approach for cancer prevention and treatment when combined with other cytotoxic agents.

In this study, we aimed to determine how statins effect the energy metabolism of HL60 cells when used single or in combination with Ara-C therapy. We observed that the healthy white blood cells produced CO2 at a rate of 88.38 % and the rate of lactate production was 11.62 %, showing that they principally use aerobic glycolysis. On the contrary, when the end products of the untreated HL-60 promyelocytic cells was taken into consideration, the rate of lactate production was 96.58 % whereas CO2 production was 3.42 % showing that they principally use anaerobic glycolysis as shown in Table 1.

When HL-60 cells were incubated with only Ara-C, lactate production was 67.6 % and CO2 production was 32.4 %; when only Simvastatin was given to the cells these rates were 61.06 % and 38.94 %, respectively. In case of a combination therapy (Ara-C + Sim), lactate production rate decreased to 47 % and CO2 amount increased to 51 % showing that combo therapy lowered the rate of anaerobic glycolysis. When CO2 and lactate production in addition to glycogen consumption of treatment groups were compared with healthy white blood cell cultures and untreated HL-60 leukemia cells, the difference was statistically significant in both groups (p<0.001), (p=0.002). The present data showed that combined drug use increased the rate of aerobic glycolysis, additionally the metabolism of these cells become more similar to the metabolism of healthy white blood cells they originate from.

When CO2 and lactate production in addition to glycogen consumption of Ara-C and Sim + Ara-C groups were compared, there was a statistically significant difference in between these three indicators (p=0.018, p=0.003, p=0.002, respectively). The difference between Simvastatin group and combo therapy groups also had statistical significances (p=0.0018, p=0.004, p=0.007, respectively).

When the effects of these drugs on energy metabolism was compared, it was observed that single use of Simvastatin and Ara-C did not show any statistically significant difference either in glycogen consumption or lactate production (p>0.05), Thus, Simvastatin treatment given to HL-60 promyelocytic cells significantly lead to aerobic glycolysis, especially in terms of CO2 production. Especially, when compared with Ara-C treatment, a statistically significant difference was observed favoring Simvastatin in terms of CO2 production (p=0.002).

Our results also showed that combination of Sim and
Ara-C produced an antigrowth interaction which was more active than the individual compounds on inhibiting cell growth. According to these results, it is possible to conclude that combined drug use increase the rate of aerobic glycolysis of HL-60 cells, and the metabolism of these cells become more similar to the metabolism of healthy white blood cells they originate from. Based on these observations, we can conclude that Simvastatin enhances the therapeutic efficacy of Ara-C, therefore Simvastatin and Ara-C may be suggested as the promising agents which can be used together in routine chemotherapy protocols for hematological malignancies, however different clinical studies are required to assess the promising results of these agents.

The authors declare no conflict of interest.

References


