Effect of withaferin, a radiosensitizer, on the erythrocyte antioxid-ants in carcinoma of uterine cervix

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Abstract

Withaferin, an active component obtained from the root extracts of Withania somnifera (Ashwaganda), showed antitumor and radiosensitising effects in animals. A similar approach in human cancer patients could probably increase the therapeutic outcome. Antioxidants are good markers of free radical induced tissue damage or in other words, radiosensitisation of tumor cells. Therefore an assay of erythrocyte antioxidant levels namely GSH, GSH-PX, SOD and G6PD were performed in cancers of uterine cervix, before, in the midst of, and post radiation and compared with the erythrocyte levels of the same in normal controls. Although a high level of GSH was observed in the baseline samples, this level was consistent after treatment. A significant decrease in GSHPX values following radiation was observed. SOD and G6PD levels remained non significant. Therefore establishing the role of withaferin as a radiosensitiser seems to be ambiguous.

Introduction

Cancer of uterine cervix is one of the leading causes of cancer death among women worldwide[1]. Early stages of this disease can be treated by surgery or with radiation. Failure of radiotherapy in the local control of solid tumors is often attributed to the presence of radioresistant hypoxic cells. Therapeutic outcome can be improved by using the chemicals(radiosensitizers) that increase the radiosensitivity of tumor cells, so that a higher tumor killing is achieved at conventional doses of RT. Furans, thiophenes, imidazoles, pyrazoles, pyrroles and thiazoles have been synthesised and tested as hypoxic
cell radiosensitizers[2]. Withaferin A, the active component obtained from the alcoholic extracts of the dried roots of the plant Withania somnifera showed significant antitumor and radiosensitising effects in experimental tumors induced in mice without any noticeable systemic toxicity [3,4]. As radiation kills tumor cells by generating free radicals and since withaferin sensitizes the tumor tissue to radiation we sought to study the influence of the latter on the erythrocyte antioxidant levels.

**Materials and Methods**

20 cases of carcinoma of uterine cervix (stage 111 B ) were considered for the study. All patients were treated with radiation at Kasturba Medical college, Hospital, Mangalore, India.

**Inclusion criteria**

All patients selected were aged between 30 and 70 years. All cancer patients were selected based on the karnofsky’s performance scale KPS>70% (10).[Cares for self but unable to carry out normal activity: shows some signs or symptoms of the disease]

The patients had no previous history of treatment and received radiotherapy at a dose of 60 Gy in 30 fractions over 6 weeks.

**Exclusion criteria**

All patients were subjected to thorough clinical examination and those with severe systemic illness like diabetes mellitus, coronary artery disease and tuberculosis were excluded.

All patients with carcinoma of cervix were treated with withaferin prior to radiotherapy, at a dose of 400mg/m², 2 hrs prior to each sitting. The Institutional Ethical Committee had approved the drug trials.

Age and sex matched healthy non hospitalized controls (n=25) were considered for the comparative study with the patients.

NADPH, Riboflavin, Lmethionine and Glutathione standard were obtained from SRL company limited. Glucose 6 phosphate was purchased from Loba chem.

Cyanomethemoglobin standard was bought from Ranbaxy. DTNB was obtained from SISCO, NBT from S.D. Fine chem. Ltd. Cumene hydroperoxide from Fluka, Ag L Buchio, Switzerland and Glutathione reductase (E.C.1.6.4.2.) Type111 from Bakers yeast from Sigma chemicals, U.S.A.

Heparinised vacuotainers were purchased from Babul Biomedicals Pvt. Ltd, Ahmedabad.

5 ml of venous blood was collected from patients in three stages:
a. 0 days of radiation (Baseline sample)
b. 15 days of radiation (1 follow up sample)
c. 30 days of radiation (11 follow up sample all patients were not available)

Likewise 5 ml of blood was collected from controls.

Blood samples were kept in an upright position at room temperature for one hour. Once the plasma separated erythrocytes were collected and the hemolysate was prepared by the reported protocol.

This hemolysate was used for estimation of different parameters. Glutathione (GSH) was assayed by the method of Beutler et al [5]. Glutathione peroxidase (GSH-PX) by the method of Paglia and Valentine [6,7]. Glucose 6 phosphate dehydrogenase (G6PD) was estimated by monitoring the increase in absorbance at 340 nm [8]. Superoxide dismutase (SOD) was assayed by the method of Beauchamp and Fridovich [9,10]. Haemoglobin (Hb) concentration in the RBC was determined by cyanomethemoglobin method [11]. The activities of GSH was expressed as nmol/g Hb and the activities of other antioxidants were expressed in terms of units/g Hb.

**Statistical analysis**

Kruskal Waalis test was used for comparison between independent groups. Wilcoxon’s rank sign test was used for comparing the follow up cases.

**Results**

Erythrocyte GSH increased in cancer of uterine cervix as compared to controls p<0.01. Following treatment, a decrease is observed which is not statistically significant but significantly high as compared to controls p<0.05. There were no significant changes when GSH-PX values of patients were compared with controls. However, the values were significantly low in the second follow up as compared to first follow up p<0.05. SOD and G6PD values did not vary between follow up samples or between the two groups.

**Table 1: Erythrocyte parameters in controls and patients treated with radiation and withaferin**

<table>
<thead>
<tr>
<th></th>
<th>Controls</th>
<th>Patients</th>
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<tbody>
<tr>
<td></td>
<td>n =25</td>
<td>Baseline n =20</td>
</tr>
<tr>
<td>GSH (nmol/g Hb)</td>
<td>58.5± 6.20</td>
<td>133.3± 18.20*</td>
</tr>
<tr>
<td>GSH-PX (nmol of NADP reduced mm/g Hb)</td>
<td>14.4± 1.46</td>
<td>14.0± 2.26</td>
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<tr>
<td>G6PD (nmol of NADPH oxidized mm/g Hb)</td>
<td>2.7± 0.30</td>
<td>2.9± 0.26</td>
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<tr>
<td>SOD (U/g Hb)</td>
<td>630± 0.9± 386.80</td>
<td>668± 43± 503.50</td>
</tr>
</tbody>
</table>

(For larger image, click [here](#))
Discussion

Certain works have reported low levels of GSH in cancer of uterine cervix and cervical intraepithelial neoplasia [12,13]. A fall in GSH concentration in blood plasma in cancer of uterine cervix after one fraction of RT was also observed which correlated positively with tumor response as reported by Jadhav et al [14]. On the contrary, elevated levels of GSH in cancer tissues (oral cancer, lung squamous cell carcinoma, cervical and other squamous cell carcinoma) has been observed by Wong et al [15] which has been attributed to abnormal proliferative activities in cancer tissues. Our work has observed an elevation of GSH in the blood which indicates that the abnormal proliferative activities reflects in the blood as well. However, treatment with radiation and withaferin have not resulted in alteration in the values of GSH.

GSHPX values have decreased in cancer tissues [16] and in the blood of cancer patients [17]. A fall in erythrocyte GSHPX levels was reported in oesophageal cancer [18] advanced gastrointestinal cancer, breast cancer [19], colon cancer [20] and lung cancer [21]. Although we report no change in the baseline samples of cancer patients as compared to controls, RT has led to a fall in the GSHPX levels which could not be reverted by Withaferin. G6PD activities have neither altered in baseline samples nor in the post treated samples as compared to controls which would mean that the reducing equivalents for the generation of reduced glutathione is provided by some other source. SOD levels remained nonsignificant in the pretreated and post treated samples versus controls.

An increase in the antioxidant enzymes namely SOD, GSHPX and catalase was reported in the rat brain following withaferin treatment [22]. No such effects were observed with respect to withaferin in the present work. Therefore, the role of withaferin in sensitizing the cancer tissue to radiation remains obscure.

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