Evaluation of the synergistic effect of danazol and radiation exposure on some biochemical functions in female albino rats.

Ramadan, F. L.
Radiation Biology Dep., National Center for Radiation Research and Technology (NCRRT), Atomic Energy Authority (AEA), Egypt.

Abstract

Background: Radiation generates a variety of free radicals during the exposure of biological tissues through radiolysis of water. These free radicals are highly reactive and cause oxidative damage to biological molecules. The present study was carried out to investigate the synergistic effect of danazol and radiation exposure on certain functional disorders in female rats.

Material and methods: Mature female rats weighing about 120-150g were divided into four groups. Group 1: control animals, group 2: animals orally administrated with danazol at a daily dose of 1.8 mg/100 g b.wt/day for four weeks., group 3: animals exposed to whole body gamma irradiation (5 Gy), group 4: animals orally administrated with danazol two weeks before and two weeks after irradiation.

Results: Blood and liver samples were obtained two week post irradiation. Erythrocyte counts, haemoglobin concentration (Hb), hematocrite percentage (Ht) and leucocyte counts (WBCs) were determined in blood. Total cholesterol (TC), triglycerides (TG), high density lipoprotein-cholesterol (HDL-c), low density lipoprotein-cholesterol (LDL-c), glucose as well as gamma glutamyle transferase (γ-GT), transmaminase aspartat (AST), alanine (ALT) and estradiol serum hormone level were assessed. Peroxidative hepatic damage was investigated by assessing thiobarbituric acid reactive substance (TBARS) and total protein content in liver tissues. The data obtained revealed that exposure of rats to gamma radiation and / or danazol treatment or dual treatment caused a significant increase in ALT, AST, glucose, γ-GT, TC, TG, LDL-c and liver TABRS. While a significant decrease were recorded in RBCs Hb, WBCs and HT.

On the other hand, serum HDL, estradiol and liver total protein in group treated with danazol declined compared to control group.

Conclusion: The results are of great importance from the stand point of radiation protection and drug safety.

Introduction

Ionizing radiation has been considered a source of naturally occurring physical damage to living organisms. At present, various man-mad therapeutic diagnostic or occupational sources of exposure to ionizing radiation are of far greater importance (Leadon, 1996). Radiation interacts with matter by direct and indirect processes to form ion pairs, some of which may be free radicals that lead to molecular damage translated to biochemical damage that may be then amplified and expressed as biological injury (Abou-Safi et al., 2005).

Danazol (DA-Na-Zole) is a synthetic steroid derived from 17-alpha ethinyltestosterone and it is used for curing number of different medical problems (Cai, et al., 1999). These included pain or infertility due to endometriosis and hereditary angioedema which causes swelling of sexual organs (Momeda, 2001). Its effectiveness is due to a reversible hypoestrogenic and hyperandrogenic state which lead to atrophy of the ectopic endometrial tissues (Tamaya, 2001). Radiation therapy is widely used because these is often no other
choice for treatment endometriosis in infertility female. Revelance, the steadily are gowing number of female workers integrating their activities with such kind of radiation exposure and / or therapy.

In this respect, our study was carried out to study the synergistic effect of danazol and radiation exposure induced certain functional disorders in female albino rats.

**Material and Methods**

Mature female albino rats weighing 120-150g were obtained from the animal farm of the Egyptian Organization for Vaccine and Biological products. Rate were housed in cages and maintained under standard condition of ventilation, temperature and humidity. Food and water were available ad-libitum.

**Radiation facility:**
Irradiation was performed by Cesium-137 biological irradiation Gamma Cell-40 belonging to the NCRRT, Nase City, Egypt. Irradiation dose (5Gy) was applied as a shot dose at a dose rate of 1Gy/1.2 min.

**Material:**
Danazol was purchased from SAND1g-Synthelabo, Canada. It was dissolved in saline solution and orally administrated to rats at a dose of 1.8 mg/100 kg b.wt/day for four weeks.

**Experimental design:**
Animals were divided into 4 groups each of 5 rats:
1-Control group: Animals did not receive any treatment.
2-Danazol group: Animal orally administrated with danazol at a dose of 1.8 mg/100g b.wt/day all over the experiment (four weeks).
3-Irradiated group: Animals irradiated at a dose of 5 Gy.
4-Danazol and irradiated group: Animals treated with danazol two week pre and post irradiation (5 Gy).

**Biochemical Studies**
Five rats from each group were anaesthetized with ether and blood collected by heart puncture. Part of the blood was collected on EDTA for assay of RBc,s Hb and WBC,s Ht and Dacie and Lewis (1991), blood haemoglobin was assayed as cyanmethaemoglobin using Spectrum Diagnostic kit according to Teitz, (1994). The rest of blood was centrifuged to separate serum to determine glucose level according to Trinder, (1969) using Scicodiagnosis kit. Aminotransaminases activity levels were determined as described by Reitman and Frankel (1957) by kit of plasmate laboratory LTd. Activity level of gamma glutamy transferase (γ-GT) was measured colorimetrically according to the procedure of Szasz and Persijn (1974). Also, estradiol was determined by using Radioimmunoassay RIA technique (Diagnostic products corporation, Los Angeles, USA). Total cholesterol, high density lipoprotein cholesterol, low density lipoprotein cholesterol and triaglycerides were assayed in serum according to the method of Allan et al., (1974), Demacer et al., (1980), Friedewald et al. (1972) and Fossati and Prencipe (1982) respectively. Liver were excised and weighed then homogenized to determine total protein according to Lowry et al. (1951). Lipid peroxidation in liver tissue was ascertained by the formation of TBARS and measured as described by Yoshioka et al. (1979). Statistical analysis of data were performed using student t-test (Snedector and Cochron 1989).

**Results**
In the present study, experimental animals received danazol for 4 weeks at a dose equivalent to 1.8 mg/100 g b.wt/day and / or exposed to gamma radiation (5 Gy) exerted a significant decrease in RBc,s Hb, WBC,s Ht and HDL-c (P ≤ 0.001) as compared to control group table (1 and 2). Table (3) indicated that whole body gamma irradiation (5 Gy) induced a significant increase in serum estradiol levels whereas treatment with danazol induced a significant decrease in the serum estradiol (P < 0.001) as compared with control. Meanwhile, the tested treatment combination with danazol and radiation exhibited a tendency to decrease serum
estradiol as compared to irradiated group. On the contrary, danazol administration and/or exposure to gamma radiation showed a significant elevation in serum TC, TG, LDL-c, glucose level, AST, ALT and \( \gamma \)-GT activity (\( P < 0.001 \)) in comparison with those of control group (table 2, 3). Data presented in table (4) showed a significant decline in total protein (\( P < 0.001 \)) paralled with a significant elevation of TBARS in experimental groups when compared to with control group.

### Table (1): Effect of danazol administration to irradiated female rats on RBc (10\(^6\)/mm\(^3\)), Hb (g/dI), WBe’s (10\(^3\)/mm\(^3\)) and HT (%).

<table>
<thead>
<tr>
<th>Groups</th>
<th>RBc,( \times 10^6 )/mm(^3)</th>
<th>Hb (g/dI)</th>
<th>WBe,( \times 10^3 )/mm(^3)</th>
<th>HT (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>5.644 ± 0.241</td>
<td>12.56 ± 0.903</td>
<td>6.17 ± 0.324</td>
<td>34.43 ± 0.673</td>
</tr>
<tr>
<td>Danazol</td>
<td>3.560 ± 0.377***</td>
<td>9.00 ± 0.631**</td>
<td>4.06 ± 0.172***</td>
<td>30.04 ± 0.696***</td>
</tr>
<tr>
<td>( \gamma )-irradiation</td>
<td>3.062 ± 0.343***</td>
<td>8.06 ± 0.617***</td>
<td>3.36 ± 0.192***</td>
<td>29.86 ± 0.399***</td>
</tr>
<tr>
<td>Danazol + ( \gamma )-radiation</td>
<td>2.928 ± 0.146***</td>
<td>7.36 ± 0.640***</td>
<td>2.95 ± 0.212***</td>
<td>27.298 ± 0.645***</td>
</tr>
</tbody>
</table>

Each value represents the mean of 5 observations ± S.E.

* \( P < 0.05 \), ** \( P < 0.01 \), *** \( P < 0.001 \).

### Table (2): Effect of danazol administration to irradiated female rats on serum of (TC), (TG), HDLC and LDL-c (mg/dI).

<table>
<thead>
<tr>
<th>Groups</th>
<th>TC (mg/dI)</th>
<th>TG (mg/dI)</th>
<th>HDL-c (mg/dI)</th>
<th>LDL-c (mg/dI)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>76.800 ± 3.513</td>
<td>82.800 ± 6.127</td>
<td>44.308 ± 0.489</td>
<td>30.088 ± 0.035</td>
</tr>
<tr>
<td>Danazol</td>
<td>98.801 ± 0.583***</td>
<td>106.300 ± 6.066***</td>
<td>41.628 ± 0.417***</td>
<td>42.056 ± 1.105***</td>
</tr>
<tr>
<td>( \gamma )-irradiation</td>
<td>116.801 ± 5.314***</td>
<td>135.400 ± 9.389***</td>
<td>36.638 ± 1.379***</td>
<td>46.916 ± 1.164***</td>
</tr>
<tr>
<td>Danazol + ( \gamma )-radiation</td>
<td>146.5 ± 3.449***</td>
<td>151.600 ± 8.280***</td>
<td>32.860 ± 1.287***</td>
<td>51.07 ± 1.125***</td>
</tr>
</tbody>
</table>

### Table (3): Effect of danazol administration to irradiated female rats on serum glucose, AST, ALT, activity of \( \gamma \)-GT and estradiol level.

<table>
<thead>
<tr>
<th>Groups</th>
<th>Glucose (g/dI)</th>
<th>AST (U/ml)</th>
<th>ALT (u/ml)</th>
<th>( \gamma )-GT (U/L)</th>
<th>Estradiol (P g/ml)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>106.28±3.512</td>
<td>63.88±0.671</td>
<td>31.58±0.565</td>
<td>20.270±0.765</td>
<td>19.198±0.058</td>
</tr>
<tr>
<td>Danazol</td>
<td>139.11±8.429**</td>
<td>69.37±0.666***</td>
<td>37.38±0.704***</td>
<td>29.648±1.284***</td>
<td>16.450±0.182***</td>
</tr>
<tr>
<td>( \gamma )-irradiation</td>
<td>188.88±3.187***</td>
<td>71.22±0.665***</td>
<td>39.65±0.692***</td>
<td>46.270±1.753***</td>
<td>42.560±0.510***</td>
</tr>
<tr>
<td>Danazol + ( \gamma )-radiation</td>
<td>192.49±4.045***</td>
<td>75.82±0.652***</td>
<td>41.76±0.559***</td>
<td>49.710±0.271***</td>
<td>30.530±0.268***</td>
</tr>
</tbody>
</table>

Leganed as Table (1).
Table (4): Effect of danazol administration to irradiated female rats on thiobarbituric acid reactive substance (TBARS), total protein content in liver tissue.

<table>
<thead>
<tr>
<th>Groups</th>
<th>Parameter</th>
<th>TBARS (m mol/g tissue)</th>
<th>Total protein (g/g tissue)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td></td>
<td>371 ± 21.51</td>
<td>0.51 ± 0.020</td>
</tr>
<tr>
<td>Danazol</td>
<td></td>
<td>425 ± 1.78**</td>
<td>0.42 ± 0.028**</td>
</tr>
<tr>
<td>γ-irradiation</td>
<td></td>
<td>492 ± 8.746***</td>
<td>0.40 ± 0.017**</td>
</tr>
<tr>
<td>Danazol + γ-radiation</td>
<td></td>
<td>509 ± 5.586***</td>
<td>0.36 ± 0.016***</td>
</tr>
</tbody>
</table>

Leganed as Table (1).

Discussion

In the present work, the results revealed that whole body gamma irradiation (5 Gy) or / and danazol administration at a dose of 1.8 mg/100 g b.wt/day produced haematological disorders. This was evident from the significant decrease in RBC’s, remarkable fall in Hb concentrations, significant drop in HT percentage and significant decrease in total WBC’s. The decrease in erythrocyte counts could be due to a drop in their production, increased destruction and / or haemorrhage due to the failure of thrombopoiesis. Our results are in complete harmony with those of (Hussein et al., 2007, Ashry, 2003 and Lee and Ducoff 1994). Radiation damage of haemoglobin is considered as one of the most important mechanisms triggering radiation sickness (Kurbanov et al., 1995). The decrease in Hb content (HT %) due to the radiation induced changes in erythrocyte membrane emphasizes the formation of free radicals. The effect of free radicals on erythrocyte membrane may contribute to the eventual leak of haemoglobin out of the cells (Hussein et al., 2007). Irradiation of rats caused retardation in incorporation of iron and decrease in haemoglobin binding to erythrocyte membrane. This finding agree with that observed by Tanikawa et al. (1990).

Leucocytes are highly radiosensitive to radiation and are considered as a sensitive biological indicator for the diagnosis of radiation injuries (Moss et al., 1979). In the present study, the reduction in total leucocytes count (leucopenia) after irradiation of animals could be attributed to mitotic inhibition of the bone marrow precursors (Hassan et al., 1996). On the other hand, danazol caused impairment in immune system which may has resulted in a defect in the ability to remove debris in the peritoneum and lacke of serum-soluble human leukocyte antigen which lead to hemolytic anemia (Moen et al., 1984). The present results demonstrated that exposure of animals to gamma radiation and / or danazol treatment showed a significant elevation in serum cholesterol, triglyrcide, low density lipoprotein and a decrease in high density lipoprotein. These data are in accordance with previous results of Ragab and Ashry 2004, and Abou Safi et al., 2005 who observed that the elevation in serum lipid fractions might result from ionizing radiation ability to accelerate other pathways of cholesterol formation like increasing its rate of biosynthesis in the liver and other tissues, or destruction of cell membrane by radiation and also to disturbance LDL cholesterol receptors, leading to hypercholesterolemia which affect particularly on polyunsaturated fatty acids and increase lipid peroxidation Kolomigseva, 1986 and Karbownik and Reiter 2000. On the other hand, danazol administration indicated that VLDL was catabolized and converted to LDL while the formation of HDL was impaired due to low activity of apoprotein A and B (Fukuda and Tamura 1987). The recorded consequently elevated level of triglycerides correlates previous finding of Osman, (1996) and Abou-Safi et al. (1999), who observed that after irradiation insulin level increased and synthesis of triglycerides was increased in both adipose tissues and liver which was accompanied by an acceleration of fatty acids mobilization from fat dopots to blood.
Moreover, Ellefson and Caraway (1976) stated that hyperglyceridemia may be caused by metabolic disorder as a result of endocrine dysfunction and increased level of glucose. The recorded hyperglycemia in our results could be attributed to endocrine abnormalities induced by irradiation that promote the secretion of peptide which has relation to carbohydrate metabolism by increasing glyconeogenesis in liver (Darwish et al., 2007).

Results of the present investigation showed that irradiation (5 Gy) of female rats or treatment with danazol induced disturbance in the secretory activity of the ovaries. Such effect was manifested by the significant increase in serum estradiol level after exposure of rats to gamma radiation. These observation were similar to those described by Banetskay and Amvrosev (1995) who indicated that marked stimulation of the ovulation was detected at once after irradiation, which can be connected with changes of the hormonal balance in animal body and also increased number of atretic follicles as a result of production of free radicals which induce cellular changes (Hassan and Abou-Safi 1998). Therefore, the lower penetration of hormones in a tissue target leads to its accumulation in blood (Ramadan and Rezk 2004). The decrease in serum steroid hormones (estradiol) after treatment of rats with danazol could attributed to danazol effect on ovarian and anterior pituitary gland which lead to an increase in androgenic activity (Fedele et al., 1993 and Liu et al., 1998). Furthermore, Meldrum et al. (1983) observed that sex hormone binding globulin was markedly suppressed throughout danazol treatment resulting in elevation of testosterone and decrease estrogen level. Our results manifested a significant increase in ALT, AST and GGT activities after exposure of rats to gamma radiation or danazol administration. GGT activity is considered to be one of the best indicators of liver damage since it is embedded in the hepatocytes plasma membrane and the libration of this enzyme to sera levels damage to the cell and then injury of the liver, which lead to increase in lipid peroxides (Muriel et al., 1992 and Gharib, 2007). The present data further demonstrated a significant increase in serum transaminase activity in all experimental groups. This conclusion agree with that found by (Abdel-Gawad and Ahmed 2005). The increased levels of AST and ALT could be referred to the drastic dysfunction of the liver cells induced by radiation interaction with cellular membranes and also related to extensive breakdown of liver parenchyma (Kafafy 2000). However, the increase in TBARS might be explained on the basis that exposure of female rats to ionizing radiation increased the amount of free radicals in the body especially potent hydroxyl radicals (OH) attack the polyunsaturated fatty acids in the phospholipids portion of cell membranes initiating the lipid peroxidation chain reaction (Darwish et al., 2007). Moreover, also danazol caused cytotoxicity for cell which causes elevation of lipid peroxidation (Stevenson et al., 2000). The registered decrease in total protein content of liver tissue could be similar to other findings of (Azab, 1996 and Abou Safi et al., 2005) who reported that radiation could induce variety of membrane changes including lipid peroxidation and amino acid damage in membrane proteins.

On the other hand, danazol treatment interfered with the function of liver which lead to severe anemia and it caused hepatic failure (Holt and Keller 1984). In addition, the treatment of animals with danazol was related to the decline in Ca++ ion concentration which lead to decrease total protein (Kriplani et al., 1998).

According to the present results it could be concluded that radiation exposure or / danazol intake could exert a deleterious biological effect. Therefore, it is recommend that radiation occupational workers especially females have to be careful toward danazol intake.

References


26. Kurbanov F F, Mamedov T C, Abdullaev......


تقييم التأثيرات المؤازرة للدانزول والتعرض الإشعاعي على بعض الوظائف الحيوية
في إناث الجرذان البيضاء
فاتمة لطفى رمضان
قسم بحوث البيولوجيا الإشعاعية - المركز القومي لبحوث وتكنولوجيا الإشعاع
هيئة الطاقة الذرية - جمهورية مصر العربية

يًولَد الإشعاع أنواعًا مختلفة من الشوارد الحرة أثناء تعرض الأنسجة حيث تسمح هذه الشوارد النشطة في تدمير الخلايا. استهدف هذا البحث دراسة التأثير المؤازر للتأثيرات البيولوجية الناتجة عن الإشعاع والمعاملة بالدانزول على إناث الجرذان البيضاء. وقد قسمت إناث الجرذان إلى أربعة مجموعات، المجموعة الأولى استخدمت كمجموعة ضابطة، المجموعة الثانية تُجريع بالدانزول لمدة أربعة أسابيع بمقدار (1.8 ملجم لكل 100 جرام من وزن الجسم يوميًا)، والمجموعة الثالثة تم تعرضها كلية للإشعاع الجامي بجرعة كلية مقدارها 5 جرائم والمجموعة الرابعة تُجريع بمساحة جزءية من جرعة الدانزول لمدة أسبوعين قبل التعرض للجرعة الإشعاعية (5 جرائم) ولمدة أسبوعين بعد التعرض للإشعاع. ينهاية التجربة تم سحب عينات من الدم ونسيج الكبد بعد أسبوعين من التعرض للإشعاع.

أظهرت نتائج الدراسة أن التعرض للإشعاع أو المعاملة بالدانزول أو المعاملة المزدوجة بالإشعاع والدانزول قد تسبب في زيادة مستويات الجلوكوز ونشاط إنزيمات ترانس أمينيز الألدهنا، ترانس أمينيز الأسيرتيت، الكولسترول الكلي، الكولسترول منخفض الكثافة، الدهون الثلاثية، إنزيم الجاماجتاميلي وترانسفيراز في السيرم وكذلك المواد المتفاعلة مع حمض الثيوريثيوريك في نسيج الكبد بينما سبب التعرض للإشعاع أو المعاملة بالدانزول أو الاثنين معاً إلى انخفاض معنويًا في عدد كرات الدم الحمراء والبيضاء، الهيموجلوبين، الهرمون الكولسترول العالي للكثافة في السيرم وأيضاً هرمون الأستراديلل للمجموعة المعالمة بالدانزول بالإضافة إلى حدوث انخفاضاً معنويًا في محتوى البروتين الكلى في أنسجة الكبد.

وتستخلص من هذه النتائج على ضرورة إستعمال الدانزول بحذر للإناث العاملين في مجال الإشعاع.