The assessment of cytotoxic T cell and natural killer cells activity in residents of high and ordinary background radiation areas of Ramsar-Iran

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ABSTRACT

The effective radiation dose of human from natural sources is about 2.4 mSv/y and the dose limit for radiation workers is 20 mSv/y. Ramsar, a city in Iran, has been the subject of concern in the last forty years for a high level of radiation measured in some spots as high as 260 mSv/y. Carcinogenesis is one of the most studied effects of radiation especially in high doses. Recent studies showed that the high level of natural radiation received by inhabitants of this area, paradoxically don’t have significant health effect. Natural killer (NK) cells and cytotoxic T cells are the most important cells in tumor immune surveillance and CD107a is a widely expressed intracellular protein located in the lysosomal/endosomal membrane. CD107a transiently located on the cell membrane can be used as a marker of CD8+ T cell degranulation following stimulation. It is also expressed, to a lower extent, on activated NK cells. In this study, 60 healthy people were selected randomly and their consent obtained and confounding factors such as sex, age, life-styles was matched then the count of activated NK and CD8+ cells was compared in high and normal background radiation areas inhabitants of Ramsar. After filling the questionnaire and measurement of background radiation, blood samples of 30 healthy people from each region were analyzed immediately by means of flowcytometry. The leukocytes and their subsets were not significantly different between two groups and the count of active cells was higher in control group. The result shows that the changes in immune system occur due to radiation and maybe it is as a result of higher radiosensitivity of activated cells.

Key words: CD107a, cytotoxic T cell, high background radiation area, natural killer cell, Ramsar

Introduction

Treg (T) cells can be broadly classified as cytotoxic CD8+ T cells, which can directly kill an antigen-expressing cell or cytokine-secret ing CD4+ T cells. The CD4+ T-cell (T helper) response can elicit both immune stimulatory or immune inhibitory effects. T helper type 1 (Th1) lymphocytes (LYM) secrete interleukin (IL)-2, interferon (INF)-γ, and lymphotoxin-a and stimulate type 1 immunity, which is characterized by intense phagocytic activity. But, Th2 cells secrete IL-4, IL-5, IL-9, IL-10, and IL-13 and stimulate type 2 immunity, which is characterized by high antibody titers. Type 1 and type 2 immunity are not strictly synonymous with cell-mediated and humoral immunity, because Th1 cells also stimulate moderate levels of antibody production, whereas Th2 cells actively suppress phagocytosis.[1] Specific CD4+ T-helper cell phenotypes are crucial for the expansion and persistence of tissue-destructive CD8+ T cells.[2] Natural killer (NK) cells are a subset that plays a central role in the innate immune response to tumors and infections.[3] NK cells are an important component of the innate immune responses they are able to lyse tumor cells or virally infected cells without prior antigen sensitization.[3] Cytotoxic T cells destroy virally infected cells and tumor cells, and are also implicated in transplant rejection. These cells are also known as CD8+ T cells since they express the CD8 glycoprotein at their surface. These cells recognize
their targets by binding to antigen associated with major histocompatibility complex (MHC) class I, which is present on the surface of nearly every cell of the body. Through IL-10, adenosine and other molecules secreted by regulatory T cells, the CD8 + cells can be inactivated to an anergic state, which prevent autoimmune diseases such as experimental autoimmune encephalomyelitis.[2]

There are a few areas in the world that have high natural background radiation. Among these areas, Ramsar, a coastal city in Iran, has the highest dose level.[5,6] This high dose radiation is mainly from Radium-226 and its progenies coming from hot springs eventually spreading in the environment.[7]

The effective dose received by the inhabitants of Ramsar, especially in Talesh Mahalleh, is even many times greater than the dose limit for radiation workers (260 mSv/y versus 20 mSv/y).[8] According to international commission on radiological protection recommendation, the maximum effective dose to public at large should be < 1 mSv/y.[9] Many studies about radiation doses and health indices of inhabitants of Ramsar are conducted. In the study of their health status and frequency of cancer by Monfared et al., they have reported that although the radiation doses received by these individuals is many times greater than residents of normal background radiation areas (NBRAs) and the cytogenetic studies showed that there is higher chromosomal aberration in the LYM of the residents of high background radiation area (HBRA) of Ramsar,[10] the frequency of cancer in these people, was not higher than other people living in NBRAs of Ramsar.[11‑14] According to immune surveillance theory, one of the important roles of immune system is to identify and to kill tumor cells. CD8 + T cells and NK cells lyse tumor cells after activation.[15]

Lysosomal-associated membrane protein-1 (LAMP-1 or CD107a) has been described as a marker of CD8 + T cell degranulation following stimulation and Recently, Galit Alter et al., described CD107a as marker of NK cell functional activity using multi-parameter flowcytometry.[3] CD107a is significantly upregulated on the surface of NK cells following stimulation with MHC devoid targets. Additionally, CD107a expression correlates with both cytokine-secretion and NK cell-mediated lysis of target cells.[3]

In this study, we have counted activated CD8 + T cells and NK cells and our hypothesis was that HBRA residents exhibit a similar immunological change observed in acutely and heavily exposed people. The higher number of activated cells in HBRA residents could indicate that high level of radiation dose received by these individuals could induce tumor cells and activated NK and CD8 + cells, eradicate these tumors. Thus it can be as evidence for the relationship between activation of immune system and cancer prevention, and immunological implication in radiation carcinogenesis.

**Materials and Methods**

Thirty healthy residents of HBRA and 30 healthy residents of NBRA were selected randomly and a questionnaire were filled by interview. The age and sex was matched and pregnant women, recently operated, and individuals suffering from an illness or taking medications were excluded from the study.

The people who were living for at least 10 years in this area considered as study population. Radiation dose rates of their homes were measured by calibrated dosimeter (Gratez A, Germany). To estimate the annual effective dose that is the total dose resulting from exposures of ionizing radiation over a year (µSv/y), the dose rate (nSv/h) was multiplied by occupancy factor. The occupancy factor was the proportion of exposure time multiplied by 4380 (365 days × 24 h).

Two milliliter fresh peripheral blood of them was taken in sterile conditions. The samples carried to laboratory (Tooba Lab., Sari, Iran) in K3EDTA tubes and the cell count was performed automatic hematology analyzer (sysmex kx-21, Japan).

Twenty microliter of each sample was stained with 10 µL of CD107a-PE (Exbio, Czech Republic) and 10 µL CD45-PE (Partec, Germany) monoclonal antibody for 15 min in dark. Then we added buffer 1 and 2 according to catalog for detection of all CD45 + cells (Partec, Germany). This method is simplified volumetric flowcytometry that allows feasible and accurate determination of CD45 + cells that don’t need to lyse and washing buff er. Afterward, the samples were shaken and stimulated CD107a + cells were counted by flowcytometer (Partec Pas, Germany). This study was approved by the accredited Ethics Committee of Babol University of Medical sciences, Babol, Iran.

The statistical analysis of results was done by SPSS 16. The results of HBRA and NBRA inhabitants were compared by independent sample t test and P < 0.05 assumed significant.

**Results**

The radiation dose rate at HBRA was significantly more than NBRA [Table 1]. Annual effective doses of HBRA group were between 4511 and 57000 and its mean was 9770 μSv/y. On the other hand NBRA group received 570-3020 μSv/y and the mean of their effective dose was 1207 μSv/y.

The difference in counts of white blood cells (WBC), content of the mixture (MXD), neutrophils and LYM between two groups was not significant [Table 2].

The difference between MXD, neutrophils and LYM percentages of two groups was not significant [Table 3].
The number of CD107a+ cells in NBRA group was higher in comparison with HBRA inhabitants [Figure 1].

**Discussions**

There are a lot of studies that have evaluated the radiation effects on the immune system of animals.\[16,17\] Studies showed that following a single dose of 100 and 200 mGy, increase in cytotoxic activity of NK cells is observable 24 until 72 h later.\[18-21\]

Fuggetta et al. have shown impaired NK activity in vitro after gamma irradiation (20 Gy).\[22\] INF-\(\beta\) (200 IU/mL) was able to completely reverse this inhibitory radiation induced effect, but was not able to modify the number of CD16+ and CD56+ cells that died by apoptosis after irradiation. Concerning in vivo radiation exposure, no changes in NK cell activity were found in 1341 atomic bombing survivors residing in Hiroshima.\[23\] Koike et al. evaluated the immune status in children with goiter living in highly contaminated area of Gomel. They found increased serum levels of Immunoglobulin G (IgG), IgM and IgE and depressed NK cell activity.\[24\] Although IFN-\(\gamma\) and IL-2 enhanced the cytotoxicity of these NK cells, the response to IFN-\(\gamma\) was still below control values.

Attar et al. have studied the immune responses to infection and inflammation of the exposed residents of Ramsar.\[25\] Their results showed that the total serum antioxidant level in the exposed people was significantly lower than control and the exposed individuals also had higher LYM-induced IL-4 and IL-10 production, and lower IL-2 and IFN-\(\gamma\) production. In addition, neutrophil Nitroblue Tetrazolium, phagocytosis, and locomotion were higher in the exposed group. In contrast, LYM proliferation in response to phytohemagglutinin was unaffected. The authors conclude that the immune system of individuals exposed to high dose ionizing radiation has adapted to its environment by shifting from a Type 1 to a Type 2 response to promote anti-inflammation. This may be because inflammatory Type 1 responses generate more free radicals than Type 2 responses, in addition to the free radicals generated as a result of high environmental radiation. Thus, the serum total antioxidant level in the exposed residents was lower than the unexposed group.

Zakeri and Kariminia have evaluated hormone levels associated with immune responses of inhabitants in HBRAs of Ramsar and their study demonstrated a decreased level of dehydroepiandrosterone (DHEA) and increased level of cortisol in the elderly HBRA group which might be possibly associated with higher cumulative doses. There was a higher incidence of hypothyroidism in the HBRA group, mostly in elder group. A significant increase of CD69 expression on CD+ cells that died by apoptosis after gamma irradiation \[22\] INF-\(\gamma\) and IL-2 enhanced the cytotoxicity of these NK cells, the response to IFN-\(\gamma\) was still below control values.

Former study reported a significant increase in CD4+ and CD8+ in HBRA group. But the absolute counts of CD4+ and CD8+ and CD4+/CD8+ ratio was not significantly different between HBRA and NBRA residents of Ramsar.\[27\]

**Table 1: Radiation dose rate of habitats of individuals (nSv/h), their ages (year) and the habitance duration of them (years), difference between doses is significant**

| Habitat        | Dose rate of habitat (nSv/h) | Age (year) | Habitance duration (year) |
|----------------|------------------------------|------------|----------------------------|
| Talesh mahalleh | 1115±238                     | 48±3       | 32±4                       |
| Chaparsar (control) | 140±14             | 46±3       | 27±4                       |

**Table 2: White blood cells, mixture, Neutrophils, lymphocytes counts (per µL) in two regions. None of the differences are statistically significant**

| Habitat            | WBC (per µL) | LYM (per µL) | NEUT (per µL) | MXD (per µL) |
|--------------------|--------------|--------------|---------------|--------------|
| Talesh mahalleh    | 6753±253     | 2329±163     | 3860±308      | 440±48       |
| Chaparsar (control)| 7300±299     | 2496±130     | 4274±201      | 766±198      |

**Table 3: The difference between mixture, neutrophils and lymphocytes percentages of two groups. None of the differences are statistically significant**

| Habitat       | LYM (%)   | NEUT (%) | MXD (%) |
|---------------|-----------|----------|---------|
| Talesh mahalleh | 35.84±2.05 | 56.29±2.96 | 6.62±0.85 |
| Chaparsar (control) | 35.25±1.21 | 57.85±1.54 | 9.47±1.95 |

**MXD: Content of the mixture including monocytes, basophils and eosinophils, LYM: Lymphocytes, NEUT: Neutrophils**

Figure 1: The number of CD107a+ cells per µL (P < 0.005)
Mechanisms for putative tumor suppression or other mechanisms defending against tumor induction and development include the DNA repair, apoptosis, terminal differentiation, phenotypic suppression and radioprotectors should be considered. Altogether, these mechanisms will reduce the probability that a specifically damaged target cell will progress to frank malignancy.[20]

We recommend the measurements of the proportions or counts of the other T-cell subsets including T helper 1 (Th1), Th2, Th17 and regulatory T cells to determine other aspects of the immune involvement. The ratios of Th1/Th2 and Th17/Tregs that could reflect altered states of immune responses against foreign antigens or tumor antigens and the proportions or counts of peripheral leukocytes including monocytes and polymorph nuclear leukocytes, which also modify or even suppress antitumor immunity also, are important.

Conclusion

Our results showed that although the effective radiation doses received by inhabitants of HBRA of Ramsar were significantly higher than normal, the count of CD107a+ cells is lower than NBRA inhabitants. This maybe as the result of higher radiosensitivity of active cells and shows that these cells are present and immune surveillance are active there. Other measured parameters, WBC and MXD, neutrophil, LYM counts and percentages were not significantly different between two groups.

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