Comparing the level of bystander effect in a couple of tumor and normal cell lines

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ABSTRACT

Radiation-induced bystander effect refers to radiation responses which occur in non-irradiated cells. The purpose of this study was to compare the level of bystander effect in a couple of tumor and normal cell lines (QU-DB and MRC5). To induce bystander effect, cells were irradiated with 0.5, 2, and 4 Gy of 60Co gamma rays and their media were transferred to non-irradiated (bystander) cells of the same type. Cells containing micronuclei were counted in bystander subgroups, non-irradiated, and 0.5 Gy irradiated cells. Frequencies of cells containing micronuclei in QU-DB bystander subgroups were higher than in bystander subgroups of MRC5 cells (P < 0.001). The number of micronucleated cells counted in non-irradiated and 0.5 Gy irradiated QU-DB cells was also higher than the corresponding values for MRC5 cells (P < 0.001). Another difference between the two cell lines was that in QU-DB bystander cells, a dose-dependent increase in the number of micronucleated cells was observed as the dose increased, but at all doses the number of micronucleated cells in MRC5 bystander cells was constant. It is concluded that QU-DB cells are more susceptible than MRC5 cells to be affected by bystander effect, and in the two cell lines there is a positive correlation between DNA damages induced directly and those induced due to bystander effect.

Key words: Medium transfer, MRC5, QU-DB, radiation-induced bystander effect

Introduction

Radiation-induced bystander effect (RIBE) refers to radiation responses which occur in non-irradiated cells. It reverses the traditional paradigm that only direct irradiation causes biological effects in living cells. RIBE was observed in patients who underwent radiotherapy 50 years ago, but as it was not fully understood, did not draw much attention.\[1,2\] It was considered as a new phenomenon after Nagasawa and Little reported their observation in 1992. Nagasawa and Little irradiated less than 1% of CHO cells, but sister chromatid exchanges were observed in more than 30% of the cells. Since then, RIBE has been observed in many experiments. The subject has been reviewed in some articles\[1,6\] and some authors have described the applications of RIBE in radiotherapy.\[6,7,9\] RIBE is observed in both tumor and normal cells, as well as for both low- and high-dose radiation. RIBE in radiation therapy resembles a dual-edge sword. It is beneficial as it increases tumor cell killing, while it has adverse effects in normal cells either in near or far distances from the treatment volume. Our knowledge of the RIBE in both tumor and normal cells would help to have a quantitative assessment of the opposing roles of this phenomenon in radiotherapy. If the amount of damages induced by bystander effect (RIBE level) in tumor cells is more than that in normal cells, it may be used to enhance therapeutic ratio. Shao et al. have measured RIBE level for glioma and normal human skin cells. The frequency of micronucleated cells induced by bystander effect was much higher in gliomas relative to normal skin fibroblasts.\[10\] The authors predicted that intrinsic radiosensitivity may be important in defining the bystander response. In this study, we compared the level of micronuclei induced in another couple of tumor and normal cell lines (QU-DB and MRC5) as a result of RIBE to investigate which of them are more susceptible to be affected by RIBE. QU-DB is a continuous human cell line established from a patient with large cell anaplastic lung carcinoma\[11\] and MRC5 cell line was derived from normal lung tissue of a 14-week-old male fetus.
in 1966. MRC5 cells are capable of 42–46 population doublings before the onset of senescence.

Materials and Methods

Cell cultures
Cell lines were supplied by the Pasteur Institute in Tehran, Iran. Cells were cultured in RPMI-1640 medium (Biosera, England) supplemented with 10% fetal bovine serum (FBS; Biosera, England), 100 units/ml penicillin, and 100 µg/ml streptomycin (Sigma, USA), and maintained at 37°C in a humidified atmosphere with 5% CO₂. Sub-confluent cells were subcultured (5 x 10⁴ cells) in 25 cm² flasks (SPL, South Korea). Flasks containing individuals of each cell line were divided into two groups, target and bystander ones.

Irradiation
Exponentially growing QU-DB and MRC5 target cells were irradiated with 0.5, 2, and 4 Gy of gamma rays emitted from a ⁶⁰Co teletherapy unit (Theratron, Phoenix model, average dose-rate of 0.764 Gy/min) at room temperature. A group of cells were sham-irradiated and handled in parallel with irradiated cells. To deliver the exact doses, the flasks were placed on a water phantom (30 cm × 30 cm × 10 cm), with source to medium surface distance was 15 cm and source to medium surface distance was 80 cm. The proper time of irradiation was calculated by dividing the considered dose (0.5, 2, and 4 Gy) by dose-rate at 0.5 cm depth.

Medium transfer
After irradiation, the flasks were incubated for 24 hours, and then their media were harvested, filtered through 0.22 µm acetate cellulose filter (Orange Scientific, Belgium), and transferred to bystander flasks containing the same cell line. As a result, for each cell line, four bystander subgroups were formed (control, 0.5, 2, and 4 Gy bystander subgroups). The first bystander subgroup received medium from sham-irradiated cells and was used as control for the other bystander subgroups. At the same time, as transferring medium, cytochalasin B (Sigma Aldrich, USA) was added to a final concentration of 0.8 and 2 µg/ml to QU-DB and MRC5 flasks, respectively. The appropriate concentrations of cytochalasin B were determined before the main experiment. Different amounts of cytochalasin B were added to the flasks. Cells were placed in an incubator for 1.5 doubling time (24 hours for QU-DB and 45 hours for MRC5) and then they were fixed. The least concentration of cytochalasin B which induced more than 50% binucleated cells was determined as the proper concentration for each cell line.

Micronucleus assay
The cytokinesis-block micronucleus assay was used to score micronuclei as an end point of RIBE. For this purpose, 24 and 45 hours (1.5 doubling time) following addition of cytochalasin B, QU-DB and MRC5 cells were fixed respectively. QU-DB cells attached to the flasks were washed with phosphate-buffered saline (PBS) and then they were fixed with methanol:acetic acid taken at a ratio of 3:1 (Merck, Germany) for three times. Each time methanol:acetic acid was poured on the cells and they were incubated at 4°C for 20 min. After the last incubation, walls of the flasks were removed and cells left to dry. MRC5 cells were dried in air and then they were fixed only once with absolute methanol which was poured on the bottom of the flasks and let to dry in air. Fixed cells attached to the bottom of the flasks (as microscopic slides) were stained with 10% Geimsa (Merck, Germany) for 5–6 minutes. In each slide, 1000 binucleated cells were scored. The number of cells containing micronuclei (micronucleated cells) per 1000 binucleated cells (MC) was counted at 400× magnification. Micronucleus assay was performed for non-irradiated, 0.5 Gy irradiated, bystander, and control subgroups. The first group is representative of cells with spontaneously induced micronuclei.

Statistical analysis
All the data were distributed normally, therefore parametric tests were used to examine whether the differences between and among the data were statistically significant. One-way analysis of variance (one-way ANOVA) and Tukey’s multiple comparison test were performed to compare MC observed in bystander subgroups of individual cell line. Student's t-test was applied when only two sets of data were being compared.

Results
Direct irradiation induced micronuclei in both cell lines. We also examined the frequency of MC in non-irradiated flasks (spontaneous MC). The average number of spontaneous MC and MC induced by 0.5 Gy irradiation is shown in Figure 1. Student's t-test indicated a significant difference between frequencies of MC in 0.5 Gy irradiated cells and spontaneous MC for both cell lines (P < 0.001). There was also significant difference between spontaneous MC of QU-DB and MRC5 cells (P < 0.001) and between MC induced due to irradiation in these two cell types (P < 0.001).

The yield of MC induced in bystander subgroups is shown in Figure 2a. Medium transfer induced MC in bystander subgroups of both cell lines. Mean of MC induced in QU-DB and MRC5 bystander cells receiving medium from 0.5, 2, and 4 Gy irradiated cells was significantly different from MC values in cells which received medium from sham-irradiated cells (P < 0.005). While the statistical analysis revealed that there was significant difference between 4
and 0.5 Gy and also between 4 and 2 Gy QU-DB bystander subgroups \( (P < 0.001) \), there was no significant difference among the MRC5 bystander subgroups \( (P > 0.05) \). This observation indicated that RIBE in MRC5 cells was not dose dependent, unlike the QU-DB cells.

To compare the RIBE level in the two cell lines, the corresponding subgroups of QU-DB and MRC5 cells were compared by Student's \( t \)-test. The results indicated statistically significant differences between all couples (same subgroups of the two cell types) \( (P < 0.001) \).

Relative MC (RMC) is the ratio of MC induced in bystander subgroups to spontaneous MC of the same cell type. The values of RMC are represented in Figure 2b.

**Discussion**

Data represented in Figure 2a indicate that RIBE level in QU-DB cells is dose dependent. Statistical analysis indicated that there was significant difference between 0.5 and 4 Gy and also between 2 and 4 Gy QU-DB bystander subgroups. However, the number of MC induced in MRC5 bystander cells was equal at all doses \( (P > 0.05) \), and therefore their response was not dose dependent. The literature indicates that the type of relationship between dose and response to RIBE is dependent on cell type, but no one has evaluated the viewpoint that the association between dose and response to RIBE may be related to the cell being normal or tumoral. If the difference observed between these two cell types is demonstrated *in vivo* and also in other normal and tumor cell types, it may be related to fractionation regimens used in radiotherapy.

It is evident from Figure 1 that spontaneous MC and MC induced in QU-DB irradiated cells are higher than the corresponding values in MRC5 cells. To our knowledge, there are no published data concerning the radiosensitivity of QU-DB and MRC5 cells, but from this observation it may be interpreted that QU-DB cells are less capable of repairing DNA injuries as spontaneous MC is higher in this cell line, and these cells are more radiosensitive as they are more strongly affected by direct irradiation.

Figure 2a indicates that MC induced in QU-DB bystander cells is higher than MRC5 cells, similar to the results of direct irradiation. Therefore, it is concluded that QU-DB cells are more vulnerable than MRC5 cells to be affected by RIBE. It can also be concluded that there is a positive correlation between RIBE level and damages induced in QU-DB and MRC5 cells spontaneously or by direct irradiation.

MC values illustrated in Figure 2a are absolute quantities; however, it is appropriate to consider RMC, which represents RIBE level relative to spontaneous MC. RMC values were calculated by dividing MC of bystander subgroups by spontaneous MC in the same cell type [Figure 2b]. Substantial differences observed in absolute MC of the two cell lines were not noticeable in RMC values.
observation can be explained by the speculation that cells’ ability of repairing DNA damages determines both RIBE level and spontaneous MC, and consequently cannot affect the ratio of them; therefore, the ratios are not very different in the two cell lines. To explain higher RMC of MRC5 at 0.5 Gy, it may be predicted that at 0.5 Gy, without considering the final result or DNA damages induced in bystander cells, the level of RIBE signaling is higher in MRC5 cells. The fact that normal cells can communicate better than tumor cells supports this idea. This statement is related to a query by Mothersill et al. as to whether bystander signaling between normal and normal cells is different than between normal and malignant cells. Perhaps MRC5 cells can produce more bystander signals than irradiated QU-DB cells, but because of repairing damages in MRC5 bystander cells, the final result is less manifested. It might be better to follow an early indicator of DNA injuries such as gammaH2AX or molecules considered as bystander signaling carriers instead of micronuclei as an end point to test this hypothesis. GammaH2AX is an early marker of DNA double-strand break (DSB), and repair ability determines its value hours or days after irradiation. This end point has been used in some RIBE studies to investigate bystander kinetics. 

However, in view of radiation protection and radiation therapy which may be affected by RIBE, the above argument (the quantity of signaling or the kinetics of RIBE) is not as important as DNA damages fixed in bystander cells; and our results indicated that the level of fixed damages (micronuclei) induced by bystander effect was more pronounced in QU-DB cells which are more affected by direct irradiation. Intention of some investigations to find out whether there is a similar correlation between damages induced by bystander effects and direct irradiation in other tumor and normal cells is valuable as radiotherapy is still based on difference in radiation sensitivity of normal and tumor cells.

Besides the visible characteristic of RIBE remarked by the level of damages induced in DNA, the mechanism of induction of bystander effect in tumor and normal cells is also important as it may be different in tumor and normal cells. If this is true, it may be used to enhance therapeutic ratio. To achieve this goal, the different likely mechanisms of RIBE in tumor and normal cells will be applied to enhance RIBE in tumors and decrease it in normal tissues. Some differences have been observed in the factors responsible to transfer bystander signals in tumors versus normal cells. Burdak-Rothkamm et al. observed that 90 minutes after irradiation, both reactive oxygen spices (ROS) and cell membrane were still active in glioma cells to transfer bystander signals, whereas in normal human astrocytes only the cell membrane was active. In addition, differences have been noticed in the molecules responsible to transfer bystander signals in P53-mutated and wild-type cells. Nitric oxide (NO) production is downregulated (inhibited) by P53 proteins, thus it is abundant in P53-mutant or -deficient cells and acts as a RIBE signaling factor; however, some researchers have reported that in wild-type P53 cells, ROS substitutes the NO. It may be predicted that by scavenging ROS, it is possible to decrease or inhibit RIBE in normal (wild-type P53) cells while it may remain constant in P53-mutant tumor cells. Despite this conclusion, Harada et al. have reported that NO contributes to bystander signal formation in normal human fibroblasts. Konopacka and Rezszowska-Wolny observed that vitamins C and E do not protect human leukemia cells against death induced by direct X-irradiation or RIBE, whereas in other studies the protective effect of vitamins C and E against chromosomal damage, as well as induction of resistance to different death stimuli in normal cells, has been reported. Burdak-Rothkamm et al. detected gammaH2AX only in S phase bystander cells, and not in G1 or G2 phases of cell cycle. Therefore, it is reasonable to conclude that tumor cells, irrespective of their repair ability, are more vulnerable to bystander DSB because relatively larger fractions of these cells are in S phase.

Burdak-Rothkamm and Prise have reported some differences in the repair system of direct and bystander damages. They concluded that by manipulating repair system of “direct damages,” it will be possible to enhance radiosensitivity of tumors which are directly irradiated in patients who underwent radiotherapy, whereas radiosensitivity of normal tissues adjacent to tumors acting as bystander tissues remains constant. This procedure can be performed by disabling one or more molecules responsible to repair direct DNA injuries. For instance, inhibition of ataxia telangiectasia mutated (ATM) and ataxia telangiectasia and Rad3-related protein (ATR) molecules which participate in repairing only direct DNA damages, and not bystander ones, increases direct radiosensitivity while decreasing RIBE cell killing in bystander cells.

These facts illustrate that broadening of our knowledge in phenomenology and mechanistic differences of RIBE in tumor and normal cells is an important task and may be exploited in future radiotherapy.

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