Depletion of new neurons by image guided irradiation

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Ionizing radiation continues to be a relevant tool in both imaging and the treatment of cancer. Experimental uses of focal irradiation have recently been expanded to studies of new neurons in the adult brain. Such studies have shown cognitive deficits following radiation treatment and raised caution as to possible unintentional effects that may occur in humans. Conflicting outcomes of the effects of irradiation on adult neurogenesis suggest that the effects are either transient or permanent. In this study, we used an irradiation apparatus employed in the treatment of human tumors to assess radiation effects on rat neurogenesis. For subjects we used adult male rats (Sprague-Dawley) under anesthesia. The irradiation beam was directed at the hippocampus, a center for learning and memory, and the site of neurogenic activity in adult brain. The irradiation was applied at a dose-rate 0.6 Gy/min for total single-fraction, doses ranging from 0.5 to 10.0 Gy. The animals were returned to home cages and recovered with no sign of any side effects. The neurogenesis was measured either 1 week or 6 weeks after the irradiation. At 1 week, the number of neuronal progenitors was reduced in a dose-dependent manner with the 50% reduction at 0.78 Gy. The dose–response curve was well fitted by a double exponential suggesting two processes. Examination of the tissue with quantitative immunohistochemistry revealed a dominant low-dose effect on neuronal progenitors resulting in 80% suppression of neurogenesis. This effect was partially reversible, possibly due to compensatory proliferation of the remaining precursors. At higher doses (>5 Gy) there was additional, nearly complete block of neurogenesis without compensatory proliferation. We conclude that notwithstanding the usefulness of irradiation for experimental purposes, the exposure of human subjects to doses often used in radiotherapy treatment could be damaging and cause cognitive impairments.

Keywords: adult neurogenesis, dentate gyrus, radiation, radiotherapy, hippocampus, imaging

INTRODUCTION

Ionizing radiation has emerged as a standard method in animal studies on functional effects of adult neurogenesis (Wojtowicz, 2006). It is the only method that is currently available for relatively uninvasive deletion of new neurons in selected brain regions in a dose-dependent manner. It can be used on any species and its effects are readily translated to humans (Monje et al., 2002; Monje and Palmer, 2003). With the advent of improved focusing methods combined with brain imaging the irradiation beam can be applied to brain targets as small as a few millimeters in diameter. In this study, we employed such an approach to construct a dose–response curve showing effects of irradiation on the population of neuronal progenitors in the hippocampus, a brain structure involved in spatial and episodic memory (Cohen and Eichenbaum, 1993). Previous studies generally used one or two doses and focused on the mechanisms of cell depletion. Several studies have established that within hours of the treatment many of the proliferating cells die through apoptosis (Peissner et al., 1999; Mizumatsu et al., 2003). This is consistent with the established vulnerability of the mitotically active cells to radiation. However, there are indications of other damaging processes as well. Inflammation with the resulting release of toxic cytokines (Monje et al., 2003), destruction of blood capillaries (Monje et al., 2003) and killing of quiescent stem cells (Encinas et al., 2008) have been reported. There are conflicting reports on irreversible and reversible nature of these effects (Tada et al., 2000; Ben Abdallah et al., 2007). This is an important issue since behavioral tests require a stable baseline of neurogenesis at a given level, presumably lower than in the intact brain. If the effects of irradiation are in fact reversible the factors that can speed up or enhance its recovery are of importance in clinical practice where irradiation is used to treat malignant tumors often with unavoidable collateral damage on the surrounding tissue. If the effects are irreversible, the cause of the permanent damage should be understood since it could influence the mechanism of the remaining neurogenesis and the properties of the surviving neurons. This study shows that dose-dependence is bimodal and can be accounted for by low-dose reversible and high-dose irreversible effects of irradiation.

METHODS

SUBJECTS

Thirty 12-week-old male S-D rats (Charles River) were used in this study.

The animals were allowed to acclimate in the facility for 1 week after the delivery. All procedures were approved by the University of Toronto and University Hospital Network ethics committees.

IRRADIATION

Just prior to irradiation the animals were anesthetized with IP injection of ketamine/xylazine at 85/5 mg/kg. Animals were placed one by one on the treatment couch of the irradiation unit (Elekta Synergy)...
and aligned at the intersection of isocenter marking lasers just in front of the ears (Figure 1). The shielded door to the treatment room was shut and the animal imaged using the integrated cone-beam CT system (imaging dose of <0.03 Gy) to obtain an outline of the skull together with the body landmarks such as oral cavity, ear canals, etc. The graphical representation of the isocenter of the treatment beam was aligned with the approximate location of the hippocampus as determined from a pre-existing MRI and CT scan from a generic rat. The position of the rat with respect to the actual isocenter was then automatically adjusted using the robotic table of the treatment unit. The dosimetry with the resulting fields as seen from the side and from the front was done using Pinnacle software (Figure 2). Thus the 98, 80, 60%, isodose level were seen superimposed on the image of the skull aligned with the approximate location of the hippocampus. The dose-rate of irradiation was 0.6 Gy/min and the total dose was obtained by irradiating equally from each side (left and right lateral beams). Doses were applied over the range from 0.5 to 10 Gy with three animals per dose. The treatment unit is referenced to the National Research Council of Canada (NRCC) radiation dosimetry standards through the

![FIGURE 1](image1.png)  
**FIGURE 1** (A) Elekta synergy CT system used for imaging and irradiation in this study. Same system is also used for human radiotherapy. Inset. Red arrow points at the approximate location of the hippocampus at the intersection of the two guiding laser beams. (B) Screen-print of the targeting and dosimetry software allowing visualization of the rat’s skull (white outline). The intersection of two yellow beams indicates the center point of the irradiation (isocenter).

![FIGURE 2](image2.png)  
**FIGURE 2** (A) Dosimetry data show approximate irradiated area on the animal’s skin. (B) Sagittal CT scan with 98% dose (red), 80% dose (green), and 60% dose (blue) outlines. Brain outline is shown in purple and the hippocampus is sketched in the location determined from a brain atlas. Red circle and blue square are part of the targeting software with cross-hairs visible only with zoom-in at the isocenter. (C) Frontal view with the same isodose outlines.
American Association of Physicists in Medicine TG-51 dosimetry protocol. Monitor units used for the irradiation were derived from the clinical dosimetry database of the Princess Margaret Hospital’s Radiation Medicine Program. Sham controls were anesthetized but not imaged or irradiated. Following treatment, the animals were transferred to home cages and kept for 1 week or 6 weeks in standard conditions. There was no effect of the treatment on animal’s behavior and no side effects such as hair loss or weight loss. The weights of the control animals \( n = 3 \) were 394.7 before, and 425.7 g 1 week after treatment. The weights of the 10 Gy (the highest dose) irradiated animals \( n = 3 \) were 392.7 g before treatment and 425.7 g 1 week after treatment.

**BrdU Administration**

5-Bromo-2-deoxyuridine (BrdU; Sigma, St. Louis, MO, USA) was dissolved in phosphate-buffered saline (PBS) containing 10 mM NaOH for a final concentration of BrdU 20 mg/ml. The solution was injected i.p. at a dose of 200 mg/kg of body weight at 9 AM and again at 9 PM on the 14th day after irradiation. The dose of BrdU was chosen to allow for maximal labeling of cells born during a 24-h period according the previously described procedures (McDonald and Wojtowicz, 2005).

**Perfusion and Tissue Preparation**

Most animals were perfused at 1 week after irradiation with three animals per irradiation dose. This time point was chosen on the basis of published data showing a peak of proliferation following cell birth (McDonald and Wojtowicz, 2005). Several additional animals were perfused at 6 weeks, allowing measurement of cell survival and examination of long-term effects of irradiation on cell differentiation and maturation. Following intracardiac perfusion with PBS and 4% paraformaldehyde, the right hippocampus was sectioned into three parts – dorsal, medial, and ventral – and post-fixed overnight in 4% paraformaldehyde at 4°C. Serial 30 μm sections of all segments were cut with a vibratome resulting in 200–250 sections per animal. One in 10 transverse sections were sampled using a systematic random sampling procedure along the dorso-ventral extent of the hippocampus.

**Immunohistochemistry**

Procedures for detection of calbindin (CaBP), doublecortin (DCX), Ki67, and BrdU were the same as described in several research and methods articles recently published by us (Kee et al., 2002; McDonald and Wojtowicz, 2005; Wojtowicz and Kee, 2006). Primary antibodies, dilutions, and incubation times were as follows: CaBP (Chemicon AB1778), 1:200, 72 h at 4°C; DCX (Santa Cruz; C-18, sc-8066), 1:200, 24 h at 4°C; Ki67 (Vector labs. VP-K451), 1:200, 18 h at room temp.; BrdU (Cedarlane, OBT0030), 1:200, 24 h at 4°C). The secondary antibodies were tagged with Alexa 594 (red) or 488 (green) fluorophores (Molecular Probes). Selected sections were stained for a microglia marker ED1 as described previously (Tan et al., 2010).

**Data Analysis**

Cells were counted in each of the sampled sections (20–25 sections per animal for each marker at each time point). In all experiments with 1-week survival, the cell counts were expressed per dentate gyrus (unilateral) by counting all labeled cells in the subgranular zone of the dentate gyrus in sampled sections and multiplying the average by the total number of sections per animal. In experiments with 6-weeks survival, the cells were counted per section, since we wished to compare three regions (dorsal, middle, and ventral) of the dentate gyrus with regard to effects of radiation. In these experiments, BrdU/CaBP double-labeled cells were examined under a confocal microscope (Leica, TC, SSL).

**RESULTS**

Post-irradiation status of neurogenesis was assessed 1 week after treatment. This time point was chosen because previous BrdU tracking experiments revealed maximal density of the immature and proliferating cells at that time (McDonald and Wojtowicz, 2005). In addition, brain trauma, such as ischemia, produces maximal rebound neurogenesis 1 week after treatment (Tada et al., 2000; Dash et al., 2001; Sharp et al., 2002). Thus, the 1-week time point is an important milestone in normal and pathological cell development.

To examine possible effects of this procedure on neurogenesis we examined a wide range of doses, beginning with very low levels and spanning the previously used higher doses. As expected we observed strong inhibitory effect of high doses, but surprisingly even a low dose of 1 Gy had clear, pronounced effect (Figure 3). The irradiation produced statistically significant reductions in the total number of DCX+ cells per dentate gyrus at all doses (ANOVA, with pairwise comparisons between the control and the irradiated groups, \( P = 0.001 \)). We showed this decrease using another marker of neurogenesis, PSA-NCAM, that labels young neurons in the similar age range, but has very different chemical characteristics and entirely different cellular distribution (Seki and Arai, 1993). The controls and the 5 Gy-irradiated groups showed similar cell counts using DCX and PSA-NCAM labeling, 21.334 ± 1082, 2.150 ± 56 for DCX and 21.308 ± 654, 1.839 ± 652 for PSA-NCAM, respectively. This confirmed that irradiation had a robust effect on cell number and was not marker-specific.

Densities of the proliferating Ki67+ cells in the same animals were not reduced at any of the doses. In fact, the proliferation appeared to be enhanced at the intermediate doses of 2, 3, and 4 Gy (ANOVA, \( F = 3.978, P = 0.016 \)) compared to controls. These results suggest that reduced proliferation is either not the cause of lowered neurogenesis or that a delayed compensatory increase in proliferation occurred as a result of the initial insult. The observed increase in proliferation cannot instantly compensate for the underlying decrease in neurogenesis but could account for its recovery within a few weeks.

We next analyzed the dose–response relationship by fitting the DCX data to a double exponential curve with four parameters corresponding to the maxima and the rate constants of the two exponential components (Figure 4). A good fit to the data suggests two processes, one with a maximum of 80% suppression and sharp rising rate constant with a half dose of 0.78 Gy, and the other with
We investigated the possibility that increased proliferation at moderate doses of irradiation can compensate for the loss of DCX+ cells. In this experiment we exposed the animals to 3 Gy, which at 1 week causes suppression of neurogenesis by nearly 90% but increases proliferation by over two-fold. We hypothesized that with time the increased proliferation of the precursors should compensate for the initial cell loss. The results are consistent with this hypothesis. The number of Ki67+ cells was significantly higher in the irradiated animals at 1 week post-irradiation. Two-way ANOVA, taking time (1 or 6 weeks) and treatment (control or irradiation) as variables, shows a significant effect of time ($F_{1,11} = 39.26, P < 0.001$) and treatment ($F_{1,11} = 18.06, P = 0.003$) and significant interactions ($F_{1,11} = 8.86, P = 0.018$). In particular, with regard to the above hypothesis, the treatment had a significant effect at 1 week ($P < 0.0001$) but not at 6 weeks ($P = 0.39$). In contrast, the number of DCX+ cells was significantly reduced at 1-week time point. Two-way ANOVA shows a significant effect of time ($F_{1,11} = 32.63, P < 0.001$) and treatment ($F_{1,11} = 128.73, P < 0.001$) and significant interactions ($F_{1,11} = 76.63, P < 0.001$). In particular, the treatment had a significant effect at 1 week ($P < 0.0001$) but not at 6 weeks ($P = 0.10$). These data are presented graphically using the relative values (irradiated/control) to show the recovery of the effects at 6 weeks. The absolute values for cell numbers are given in the legend of Figure 5.

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In the next experiment we showed that the remaining proliferation after irradiation resulted in the production of mature neurons, albeit at lower rates. Two groups of animals, 3 Gy irradiated and sham controls, were injected with BrdU at 2 weeks and killed at 6 weeks after irradiation, thus allowing 4 weeks for neuronal maturation of the BrdU-tagged neurons. The surviving neurons were co-labeled for BrdU and calbindin (CaBP), a neuronal marker that is expressed during neuronal maturation (McDonald and Wojtowicz, 2005). In this case we show separately the total number of BrdU+ and BrdU/CaBP+ cells and we include the distribution among the three hippocampal regions (Figure 6).

**DISCUSSION**

High energy radiation surrounds us in the environment, it is also used in high doses in ionizing X-ray procedures, various imaging devices such as CT scans, and in radiotherapy for treatment of cancer or benign conditions (Koh et al., 2006). One specialized application of the ionizing radiation is to inhibit growth of endogenous stem cells in adult brain in order to assess their function in normal physiology and in potential regeneration of the brain tissue (Saxe et al., 2006; Wojtowicz, 2006). The present study examined the effects of irradiation on the hippocampus, a key brain structure involved in learning and memory. There is now convincing evidence that inhibition of adult neurogenesis in the dentate gyrus, the integral part of the hippocampus, affects some types of learning and memory in experimental animals. In particular, the memory that relates past events to context is very sensitive to reduced neurogenesis (Winocur et al., 2006; Wojtowicz et al., 2008; Hernandez-Rabaza et al., 2009). With continuing use of the irradiation in animal experiments on neurogenesis it is important to understand the mechanisms and dose dependence of the treatment. It is also useful to assess the extent of cell depletion along the dorso-ventral axis of the hippocampus. Dorsal and ventral regions are unevenly involved in various behaviors and recent evidence suggests that neurogenesis is also polarized (Snyder et al., 2008). Our results show that the targeting procedure ensures uniform effects of radiation in all regions of the hippocampus along its dorso-ventral axis (Figure 6).

There exists conflicting evidence with regard to the duration of radiation effects. In most studies the effects were irreversible (Wojtowicz, 2006), but in others, either spontaneous or experimentally induced recovery has been observed (Monje et al., 2003; Rola et al., 2004; Ben Abdallah et al., 2007). Our new data reconcile these discrepancies by showing that the effects of irradiation are bimodal. There appear to be a low-dose effect of irradiation that is associated with enhanced compensatory proliferation of precursors and, a high dose effect that lacks such compensation. Most of previous reports of the irreversible effects on neurogenesis used the doses in the higher range, above 5 Gy. The lower doses, below 5 Gy may be reversible by allowing sufficient time for recovery after treatment. However, it is also apparent that full recovery is unlikely since according to the exponential model shown in Figure 4A the high dose effect begins from 0 and grows steadily with dosage. This finding appears ominous for possible side effects of radiotherapy that uses large doses (>50 Gy) in fractions of 1-2 Gy (Laack and Brown, 2004; Barani et al., 2007). It can also be expected that even some forms of imaging such as CT scans, that require significant exposure to radiation, may affect neurogenesis. It should be noted that the irreversible effects are not necessarily due to persistent inflammation since our irradiation procedure did not produce any detectable inflammation (Figure 4B). Unlike in our previous studies where we used pentobarbital anesthesia (Wojtowicz, 2006), the present study utilized a more common ketamine/xylazine anesthesia. Thus, the proposed protective action of the pentobarbital does not seem to be necessary to prevent the inflammation at least in the dose range utilized here.

The apparent increase in proliferation at lower doses (Figure 3) may seem paradoxical but in fact is in agreement with the well established compensatory or rebound response of proliferation to various types of brain injury including ischemia or mechanical trauma (Dash et al., 2001; Kee et al., 2001; Sharp et al., 2002). There is no doubt that proliferation is reduced within 24 h of the irradiation but the rebound at 1 week suggests a homeostatic mechanism that compensates for
the initial damage. This partial recovery seems to take the course of de novo renewal of the young cell population. Accordingly, the DCX+ cells observed at 1 week of recovery are very similar to 1-week-old neurons in their normal development (McDonald and Wojtowicz, 2005). Such cells are characterized by clustering and possess short, stubby dendrites extending horizontally, in parallel to the hilar/GCL border (McDonald and Wojtowicz, 2005; Seki et al., 2007). More developed neurons with long dendrites are missing. In contrast, at 6 weeks the cells appear to include a normal range of shapes and sizes typical for the young neuronal population. At higher doses there is no rebound and ultimately the damage is permanent.

In summary, our results demonstrate the usefulness of the irradiation approach to studies of adult neurogenesis. With the targeted application of the radiation, the full dose was centered between regions or treatment (two-way ANOVA, $P = 0.645$) indicating that the irradiation is equally effective in all three regions and there is no effect on cell maturation. (B) Panels show examples of images stained for BrdU (left panel) or BrdU and CaBP (right panel). BrdU is shown in green and CaBP is shown in red. Calibration bars are 200 and 15 μm in left and right panels, respectively.
in the 5 mm × 5 mm × 5 mm voxel of the brain tissue that comprised both hippocampi. There was an inevitable “spill-over” effect that included the surrounding brain tissue but the 80–60% isodose levels declined sharply within the approximate volume of 10 mm × 10 mm × 10 mm. The effects were dose-dependent and easily verifiable with post hoc immunohistochemistry. The effects of irradiation were uniform along the dorso-ventral axis of the hippocampus with equal effect on the number of BrdU+ cells in the dorsal, medial, and ventral regions. This result serves to verify the accuracy of our irradiation targeting procedure. The equal proportions of the CaBP/BrdU-labeled cells further show that the cells spared by irradiation develop normally according the same time line as that observed in control animals (Figure 6).

These results may be used in assessing possible collateral damage caused by radiotherapy and other types of treatment or imaging procedures involving radiation. It would appear that doses as small as 100 cGy (0.1 Gy) could cause noticeable damage in the neurogenic regions of the hippocampus. This in turn could cause deficits in learning and memory as well as possible changes in sensitivity of the subjects to antidepressants. Some aspects of depression, schizophrenia, and opioid-dependence have been linked to adult neurogenesis (Eisch et al., 2000; Santarelli et al., 2003; Reif et al., 2006; Canales, 2007). Reduced neurogenesis is expected to negatively affect the symptoms and treatment of these conditions. A proof of principle evidence has recently emerged from a study Dr. C. Menard and colleagues who reported memory deficits in patients undergoing radiation therapy at doses as low as 0.75 Gy (Bernard et al., 2010).

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