Article

Nrf2-Knockout Protects from Intestinal Injuries in C57BL/6J Mice Following Abdominal Irradiation with γ Rays

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Abstract: Radiation-induced intestinal injuries (RIII) commonly occur in patients who suffer from pelvic or abdominal cancer. Nuclear factor-erythroid 2-related factor 2 (Nrf2) is a key transcriptional regulator of antioxidant, and the radioprotective role of Nrf2 is found in bone marrow, lung, and intestine, etc. Here, we investigated the effect of Nrf2 knockout on radiation-induced intestinal injuries using Nrf2 knockout (Nrf2−/−) mice and wild-type (Nrf2+/+) C57BL/6J mice following 13 Gy abdominal irradiation (ABI). It was found that Nrf2 knockout promoted the survival of irradiated mice, protected the crypt-villus structure of the small intestine, and elevated peripheral blood lymphocyte count and thymus coefficients. The DNA damage of peripheral blood lymphocytes and the apoptosis of intestinal epithelial cells (IECs) of irradiated Nrf2−/− mice were decreased. Furthermore, compared with that of Nrf2+/+ mice, Nrf2 knockout increased the number of Lgr5+ intestinal stem cells (ISCs) and their daughter cells including Ki67+ transient amplifying cells, Villin+ enterocytes, and lysozyme+ Paneth cells. Nuclear factor-κB (NF-κB) was accumulated in the crypt base nuclei of the small intestine, and the mRNA expression of NF-κB target genes Bcl-2, uPA, and Xiap of the small intestine from irradiated Nrf2−/− mice were increased. Collectively, Nrf2 knockout has the protective effect on small intestine damage following abdominal irradiation by prompting the proliferation and differentiation of Lgr5+ intestinal stem cells and activation of NF-κB.

Keywords: Nrf2; intestinal injury; abdominal irradiation; intestinal stem cells; NF-κB

1. Introduction

The small intestine is one of the most sensitive organs of the body to ionizing radiation (IR). The radiotherapy for abdominal or pelvic cavity tumors may destroy intestinal absorption and barrier function and cause radiation-induced intestinal injuries (RIII) [1]. The major symptoms of radiation-induced intestinal injuries include diarrhea, nausea, bleeding, abdominal or rectal pain, and electrolyte loss, etc. These adverse side effects of radiotherapy limit the effective radiation dosage applied to eradicate tumors. Therefore, radiation-induced intestinal injuries seriously interfere with the treatment of patients with abdominal or pelvic cavity tumors and reduce the patients’ quality of life.

Nuclear factor-erythroid 2-related factor 2 (Nrf2), the redox sensitive transcriptional activating factor, is a member of the cap “n” collar (CnC) family that is composed of the transcription factors SKN-1, NRF1, NRF2, NFE2, NRF3, CncC, BACH1, and BACH2. These factors are characterized by a leucine zipper protein-protein dimerization domain, as well as CnC and basic domains that confer DNA
binding activity [2]. Nrf2 heterodimerization with small musculoaponeurotic fibrosarcoma protein (sMaf) binds to cis antioxidant response elements (AREs) to govern the expression of Nrf2 target genes [3], which are involved in a wide range of antioxidant response and xenobiotic metabolism [4].

It is generally accepted that exposure to IR can produce radical oxygen species (ROS), including hydroxyl radicals, hydrogen peroxide and superoxide, which contribute to cell and tissue injury. Hydroxyl radicals can result in single strand breaks, double strand breaks, protein DNA crosslinks, as well as other types of DNA damage. These DNA damages caused by hydroxyl radicals, and hydrogen peroxide and superoxide as a secondary consequence of IR, can induce cell apoptosis [5]. Under oxidative stress, Nrf2 upregulates the antioxidant enzymes and detoxifying proteins. These proteins can scavenge radical oxygen species to reduce DNA damage and cell apoptosis following IR [6]. Therefore, Nrf2 has a radioprotective effect on cell and tissue damage induced by IR.

It was reported that the activation of Nrf2 mitigated radiation-induced lung injury [7,8], hematopoietic system injury [9–11], and bone loss [12]. Some studies showed that Nrf2 activation alleviated radiation-induced intestinal damage in mice post total body irradiation (TBI) [13–16]. Conversely, Berhane et al., reported that Nrf2 knockout increased the radioresistance of bone marrow stromal and hematopoietic progenitor cell lines derived from Nrf2−/− homozygous deletion recombinant-negative mice [17]. Until now, the role of Nrf2 on radiation-induced intestinal injuries remains poorly understood. Here, we explored the effect of Nrf2 on intestinal injuries to elucidate whether Nrf2 plays a radioprotective role in radiation-induced intestinal injuries. In this study, a mouse model of radiation-induced intestinal injuries in Nrf2 knockout (Nrf2−/−) mice and wild-type (Nrf2+/+) C57BL/6J mice was built by exposure to 13 Gy abdominal irradiation (ABI). The 30-day survival rate of mice, the crypt-villus structure of the small intestine, the apoptosis of intestinal epithelial cells (IECs), and the proliferation and differentiation of Lgr5+ intestinal stem cells (ISCs) of the small intestine were respectively assessed.

2. Results

2.1. Nrf2 Knockout Improved the Survival of Mice after Abdominal Irradiation

To investigate whether Nrf2 has radioprotective effect in vivo, we first analysed the 30-day survival of Nrf2−/− mice and Nrf2+/+ mice following 13 Gy abdominal irradiation. As shown in Figure 1A, all Nrf2−/− mice survived after exposure to abdominal irradiation. In contrast, 80% of Nrf2+/+ mice survived 10–30 days after abdominal irradiation, which was significantly lower than that of the irradiated Nrf2−/− mice (p < 0.05). At the same time, the body weight of survival mice was monitored for 30 days after abdominal irradiation. The body weight of survival mice was decreased four days after abdominal irradiation, subsequently, the body weight gradually increased in the following days. The speed of body weight recovery in Nrf2−/− mice was significantly faster than that in Nrf2+/+ mice (p < 0.05) (Figure 1B). These results suggested that Nrf2 knockout improved the survival of mice treated with abdominal irradiation and their body weight recovered rapidly.

In addition, the 30-day survival of Nrf2−/− mice and Nrf2+/+ mice exposed to 7.2 Gy total body irradiation was observed. As shown in Figure 1C, the survival of Nrf2−/− mice was higher than that of Nrf2+/+ mice within 15 days following 7.2 Gy total body irradiation. However, 7.2 Gy total body irradiation resulted in 14% survival in Nrf2−/− mice by day 30. In contrast, the survival rate of Nrf2+/+ mice at day 30 was 43%, which was significantly higher than that of the Nrf2−/− mice (p < 0.05). The data suggested that the effect of abdominal irradiation and total body irradiation on the survival of Nrf2−/− mice were totally different, that was, Nrf2 knockout protected from intestinal injuries induced by abdominal irradiation, but did not protect from tissue damage induced by total body irradiation.
were calculated. The spleen coefficient decreased rapidly from the irradiated mice one day after abdominal irradiation (Figure 2B). Nrf2−/− coefficient of mice after abdominal irradiation, the value remained at a stable level in the following days. The thymus coefficient of mice 2–5 days after abdominal irradiation was higher than that of Nrf2+/+ mice 2–5 days after abdominal irradiation (Figure 2A). While the thymus coefficient of Nrf2+/+ mice was decreased rapidly two days after abdominal irradiation, the value remained at a stable level in the following days. However, there was no distinct difference in the spleen coefficient of Nrf2−/− mice and Nrf2+/+ mice (Figure 2A). While the thymus coefficient of Nrf2+/+ mice was decreased rapidly two days after abdominal irradiation, the value remained at a stable level in the following days. The thymus coefficient of Nrf2−/− mice was higher than that of Nrf2+/+ mice 2–5 days after abdominal irradiation (p < 0.01), suggesting that Nrf2 knockout promoted the thymus immune function of mice exposed to abdominal irradiation (Figure 2B).

2.2. Nrf2 Knockout Increased the Immune Organ Coefficient of Mice after Abdominal Irradiation

To investigate the immune function of Nrf2−/− mice and Nrf2+/+ mice post-abdominal irradiation, the spleen coefficient and thymus coefficient of mice at 1, 2, 3.5, 4 and 5 days after abdominal irradiation were calculated. The spleen coefficient decreased rapidly from the irradiated mice one day after exposure to abdominal irradiation, subsequently, the coefficient gradually increased in the following days. However, there was no distinct difference in the spleen coefficient of Nrf2−/− mice and Nrf2+/+ mice (Figure 2A). While the thymus coefficient of Nrf2+/+ mice was decreased rapidly two days after abdominal irradiation, the value remained at a stable level in the following days. The thymus coefficient of Nrf2−/− mice was higher than that of Nrf2+/+ mice 2–5 days after abdominal irradiation (p < 0.01), suggesting that Nrf2 knockout promoted the thymus immune function of mice exposed to abdominal irradiation (Figure 2B).

Figure 2. The effect of Nrf2 knockout on the immune organ coefficients of mice after exposure to abdominal irradiation. (A) The spleens were collected and weighted after Nrf2+/+ mice and Nrf2−/− mice were euthanized 1, 2, 3.5, 4, and 5 days post-abdominal irradiation. The spleen coefficient was calculated by spleen weight (mg)/mice weight (g). The data were expressed as mean ± SD (n = 4 mice/group); (B) the thymus were collected and weighted after Nrf2+/+ mice and Nrf2−/− mice were euthanized 1, 2, 3.5, 4, and 5 days post-abdominal irradiation. The thymus coefficient was calculated by thymus weight (mg)/mice weight (g). The data were expressed as mean ± SD (n = 4 mice/group). *** p < 0.001, Nrf2−/− group versus Nrf2+/+ group.
2.3. Nrf2 Knockout Mitigated Peripheral Blood Lymphocyte Count Decreases of Mice after Abdominal Irradiation

To evaluate the blood system of Nrf2−/− mice and Nrf2+/+ mice post-abdominal irradiation, the numbers of circulating white blood cell (WBC), blood platelets (PLT), and lymphocytes from the peripheral blood at 1, 3, and 5 days after abdominal irradiation were measured. The number of WBC decreased rapidly from the mice one day after abdominal irradiation and remained at a stable level in the following days. WBC counts showed no significant difference between Nrf2−/− mice and Nrf2+/+ mice (Figure 3A), and there was no marked difference in PLT counts between Nrf2−/− mice and Nrf2+/+ mice (Figure 3B). The lymphocyte count of Nrf2−/− mice decreased rapidly three days post-abdominal irradiation, whereas, the lymphocyte count of Nrf2−/− mice was little changed after abdominal irradiation (Figure 3C). The lymphocyte count of Nrf2−/− mice was higher than that of Nrf2+/+ mice 3–5 days after abdominal irradiation (p < 0.001), suggesting that Nrf2 knockout mitigated peripheral blood lymphocyte count decreases of mice following abdominal irradiation.

Figure 3. The effect of Nrf2 knockout on various blood cells of mice after exposure to abdominal irradiation. Whole blood samples were obtained from the orbital sinus of Nrf2+/+ mice and Nrf2−/− mice to count various cells at 1, 3, and 5 days post-abdominal irradiation. Then, (A) white blood cell (WBC), (B) blood platelets (PLT), and (C) lymphocytes were counted using a MEK-7222k hemocytometer. *** p < 0.001, Nrf2−/− group versus Nrf2+/+ group.

2.4. Nrf2 Knockout Attenuated DNA Damage to Peripheral Blood Lymphocytes of Mice After Abdominal Irradiation

We further evaluated the effect of Nrf2 on the DNA damage to peripheral blood lymphocytes from Nrf2−/− mice and Nrf2+/+ mice by the comet assay. The DNA percentage in the comet tail (Tail DNA %), tail moment (TM), and olive tail moment (OTM) of lymphocytes represent the degree of DNA damage. As demonstrated in Figure 4, compared with those of Nrf2+/+ mice, there was an obvious decrease in Tail DNA%, TM and OTM of Nrf2−/− mice at 4 h, 2 days, and 5 days after abdominal irradiation, respectively (p < 0.001). The results indicated that Nrf2 knockout reduced abdominal irradiation-induced DNA damage to lymphocytes.
Figure 4. The effect of Nrf2 knockout on DNA damage of mice after exposure to abdominal irradiation. Blood samples were obtained from the orbital sinus of Nrf2+/+ mice and Nrf2−/− mice to isolate lymphocytes at 4 hours, 2 days, and 5 days post-abdominal irradiation. Then, the alkali comet assay was adopted to determine DNA damage to lymphocytes from Nrf2+/+ mice and Nrf2−/− mice. (A) Representative images of the DNA damage to peripheral blood lymphocytes. The comet tail length represents DNA damage. Scale bar, 50 μm; (B) graph displaying the DNA percentage in the comet tail (Tail DNA%) of lymphocytes; (C) graph showing the tail moment (TM) of lymphocytes; and (D) graph showing the olive tail moment (OTM) of lymphocytes. The bar graph represents mean ± SD obtained by analysing 200 cells/mouse (n = 4 mice/group). *** p < 0.001, Nrf2−/− group versus Nrf2+/+ group.

2.5. Nrf2 Knockout Alleviated the Damage to Crypt-Villus of the Small Intestine from Mice after Abdominal Irradiation

To determine the effect of Nrf2 knockout on radiation-induced intestinal injuries, the crypt-villus structure in the small intestine of mice was examined at 3.5 and five days post-abdominal irradiation by haematoxylin and eosin (H and E) staining. As shown in Figure 5, compared with that of the control mice, the crypt-villus structure in the small intestines of Nrf2−/− mice was well preserved at 3.5 and 5 days after abdominal irradiation. In contrast, the intestinal villi in Nrf2+/+ mice were obviously stunted and blunt at 3.5 days after abdominal irradiation, and the number of intestinal crypts continued to decline at five days after abdominal irradiation. The results suggested that Nrf2 knockout reduced abdominal irradiation-induced damage to the crypt-villus of the small intestine.
Abdominal Irradiation

Nrf2 knockout from abdominal irradiation. Compared with that of Nrf2+/+ mice, the rate of apoptosis of the small intestine was significantly decreased (p < 0.001).

**Figure 5.** The effect of Nrf2 knockout on the crypt-villus structure of the small intestine after exposure to abdominal irradiation. Representative haematoxylin and eosin (H and E)-stained sections of the small intestines. Small intestines from Nrf2+/+ mice and Nrf2−/− mice were harvested for pathological examination after mice were euthanized 3.5 and 5 days post-abdominal irradiation. Scale bar, 100 μm.

2.6. Nrf2 Knockout Decreased the Rate of Apoptosis of the Small Intestine from Mice after Abdominal Irradiation

IR induces apoptosis of intestinal epithelial cells, which is p53 dependent [18]. In the study, the rate of apoptosis of the small intestine was tested at 6 h after abdominal irradiation by terminal deoxynucleotidyl transferase dUTP nick end labeling (TUNEL) assay. As shown in Figure 6, the substantial numbers of apoptotic cells occurred in the intestinal villi of Nrf2+/+ mice exposed to abdominal irradiation. Compared with that of Nrf2+/+ mice, the rate of apoptosis of the small intestine from Nrf2−/− mice were significantly decreased (p < 0.001).

**Figure 6.** The effect of Nrf2 knockout on the apoptosis of the small intestines after exposure to abdominal irradiation. The small intestines of Nrf2+/+ mice and Nrf2−/− mice were harvested at 6 h post-abdominal irradiation for TUNEL assay. (A) Representative TUNEL staining images of the small intestines. Scale bar, 50 μm; (B) TUNEL-positive cells in a single field of view were quantified. The data were shown as mean ± SD. *** p < 0.001, Nrf2−/− group versus Nrf2+/+ group.
2.7. Nrf2 Knockout Resulted in an Increase in Lgr5+ Intestinal Stem Cells and Their Progeny of the Small Intestine from Mice after Abdominal Irradiation

Lgr5+ intestinal stem cells are indispensable for intestinal regeneration following radiation [19]. To investigate the proliferation and differentiation ability of Lgr5+ intestinal stem cells, the Lgr5+-, Ki67+-, Vil1in+-, and lysozyme+-stained sections, representing Lgr5+ intestinal stem cells, transiently-amplifying cells, villus cells, and Paneth cells respectively, were observed at 3.5 and five days after abdominal irradiation. As shown in Figure 7A, the number of Lgr5+ intestinal stem cells in Nrf2−/− mice at 3.5 and five days post abdominal irradiation was significantly higher than that of Nrf2+/+ mice. Similarly, the numbers of Ki67+, villin+, and lysozyme+ cells of Nrf2−/− mice were also markedly higher than that of Nrf2+/+ mice at 3.5 and five days post abdominal irradiation (Figure 7B–D). The data indicated that Nrf2 knockout accelerated the regenerative response of radiation-induced intestinal injuries by promoting the proliferation and differentiation of Lgr5+ intestinal stem cells in the small intestine.

![Figure 7](image-url)

Figure 7. The effect of Nrf2 knockout on the proliferation and differentiation ability of Lgr5+ of small intestine after exposure to abdominal irradiation. Small intestines of Nrf2+/+ mice and Nrf2−/− mice were harvested at 3.5 and 5 days post-abdominal irradiation for immunohistochemical analysis. Representative immunohistochemical images for (A) Lgr5-, (B) Ki67-, (C) lysozyme-, and (D) Villin-stained sections of the small intestine. Scale bar, 100 μm.
2.8. Nrf2 Knockout Enhanced the Activation of NF-κB of the Small Intestine from Mice after Abdominal Irradiation

To explore nuclear factor-κB (NF-κB) role in the protection effect of Nrf2 knockout on radiation-induced intestinal injuries, the expression of NF-κB p65 of the small intestine at 1 h, 3 h, and 6 h after abdominal irradiation was examined by immunohistochemistry analysis. As demonstrated in Figure 8A, NF-κB p65 was accumulated in the nuclei of the intestinal crypt base. Compared with that of Nrf2+/+ mice, the accumulation of NF-κB p65 of Nrf2−/− mice occurred at 3 h after abdominal irradiation, and was increased at 6 h after abdominal irradiation.

Figure 8. The effect of Nrf2 knockout on activation of NF-κB of the small intestine after exposure to abdominal irradiation. (A) Immunohistochemical analysis for NF-κB p65-stained sections of the small intestine. Small intestines of Nrf2+/+ mice and Nrf2−/− mice were harvested at 1, 3, and 6 h post-abdominal irradiation for immunohistochemical analysis. Scale bar, 100 μm; (B–D) The expression of NF-κB p65 in the cytoplasmic and nuclear lysates. Small intestines of mice were harvested at 1, 3, and 6 h post-abdominal irradiation. The crypt cells were isolated from the small intestines to extract the cytoplasmic and nuclear lysates respectively. The cytoplasmic protein was identified by an anti-GAPDH antibody, and the nuclear protein was identified by an anti-LaminB1 antibody; (E) The mRNA levels of NF-κB target genes Bcl-2, uPA, and Xiap. Small intestines of mice were harvested at one day post-abdominal irradiation and total RNA was extracted to performed RT-PCR reaction. The data were expressed as the mean ± SD. ** *p < 0.01, Nrf2−/− group versus the Nrf2+/+ group.

When it is activated under stress situations, NF-κB p65 immigrates from the cytoplasm into the nuclei of cells. The location of NF-κB p65 expression in the intestinal crypts cells at 6 h post-abdominal irradiation was verified by Western blot analysis. As shown in Figure 8D, the level of NF-κB p65
expression in the nuclei of the crypts cells from Nrf2−/− mice was higher than that of Nrf2+/+ mice. In addition, the mRNA expression of NF-κB target genes Bcl-2, uPA, and Xiap in the small intestine after abdominal irradiation were examined. As shown in Figure 8E, the mRNA expression of Bcl-2, uPA, and Xiap in Nrf2−/− mice was significantly higher than that of Nrf2+/+ mice at one day after abdominal irradiation (p < 0.01). The NF-κB target genes Bcl-2, uPA, and Xiap have the properties of anti-apoptosis and pro-survival. Taken together, these results indicated that the activation of NF-κB and upregulation of NF-κB target genes in Nrf2−/− mice may be involved in the event that Nrf2 knockout alleviated radiation-induced intestinal injuries.

3. Discussion

Radiation-induced intestinal injuries, the side effects of abdominal or pelvic cavity tumour radiotherapy, severely affect the curative effect. In the study, Nrf2−/− mice following abdominal irradiation were used to investigate the effect of Nrf2 on radiation-induced intestinal injuries. It was reported that radiation-induced intestinal injuries occurred in a dose-dependent manner. Exposure of 12 Gy or lower-dose abdominal irradiation for the gastrointestinal tract was considered nonlethal to mice [20,21] and exposure of 14 Gy abdominal irradiation may be the maximum dose to allow for full intestinal recovery [22]. In our study, the dose of 13 Gy abdominal irradiation was adopted for the experimental mice. We found that Nrf2 knockout promoted the survival rate and the body weight retention of mice after exposure to abdominal irradiation, indicating that Nrf2 knockout had a protective effect on radiation-induced intestinal injuries.

Under physiological conditions, epithelial homeostasis is maintained by proliferative cells within crypts [23]. However, injuries to proliferative cells commonly occur in irradiated intestine. Various degrees of villus blunting and fusion may occur, with attenuation and hypertrophy of the villus epithelial cells [24] and severe loss of crypts [25], resulting in the destruction of epithelial homeostasis and epithelial integrity. An incomplete epithelium cannot easily maintain the absorptive and defensive functions of the intestine [26]. Thus, a rapid decline of body weight, and even death, were observed in the irradiated mice. Our findings demonstrated that the crypt-villus structure of the small intestines from Nrf2−/− mice was well preserved after abdominal irradiation and fewer apoptotic cells occurred in the small intestines, indicating that Nrf2 knockout alleviated the degree of villus blunting and fusion and the loss of crypts, and preserved the epithelial integrity of mice post-abdominal irradiation. Consequently, it could explain why the survival rate and the body weight in Nrf2−/− mice after abdominal irradiation was increased. In addition, our results indicated that Nrf2 knockout attenuated peripheral blood lymphocyte count decreases and DNA damage, as well as increased the thymus coefficient after abdominal irradiation. The boost in the blood system and the immune function may be the cause of the increase of the survival rate and the body weight in Nrf2−/− mice after abdominal irradiation.

In the present study, we found the radioprotective effect of Nrf2 knockout on radiation-induced intestinal injuries. To explore the potential molecular mechanism, Lgr5+ intestinal stem cells and their progeny in the small intestine were detected. The columnar cells of the intestinal crypt base expressing high levels of Wnt target gene Lgr5 are defined as Lgr5+ intestinal stem cells, which are indispensable for intestinal regeneration [19]. Lgr5+ intestinal stem cells can divide into differentiated intestinal cell types, which migrate either upward into the villus to differentiate into functional cells, such as enterocytes, or downward into the crypt base to differentiate into transiently-amplifying cells [27,28]. Our findings demonstrated that the number of Lgr5+ intestinal stem cells in Nrf2−/− mice after abdominal irradiation was increased and the increased Lgr5+ intestinal stem cells differentiated into more transiently-amplifying cells, Paneth cells, and villus cells. Therefore, Nrf2 knockout has the protective effect on radiation-induced intestinal injuries by promoting the proliferation and differentiation of Lgr5+ intestinal stem cells.

NF-κB is a dimeric DNA binding protein, which is a member of the NF-κB/Rel family. It was reported that Nrf2 absence was susceptibility to activation of NF-κB in vivo and in vitro [29–33], and
NF-κB activation selectively protects the small intestine from radiation-induced intestinal injuries [34]. Consistently, the nuclear accumulation, the protein level of NF-κB p65 and NF-κB target genes in the intestinal crypt base of Nrf2−/− mice were significantly higher than that of Nrf2+/+ mice. Our findings demonstrated that Nrf2 knockout enhanced NF-κB activity of the intestinal crypt base in Nrf2−/− mice. The intestinal crypt base is the place that intestinal stem cells reside. As intestinal stem cells display high NF-κB activity [35] and NF-κB activation can make transformed cells grow and survive [36], it is plausible that NF-κB may promote the proliferation and differentiation of Lgr5+ intestinal stem cells. Therefore, NF-κB activation in Nrf2−/− mice may be participated in the regeneration of radiation-induced intestinal injuries and may be one reason of the protective effect of Nrf2 knockout on radiation-induced intestinal injuries.

However, it was reported that Nrf2 activation alleviated radiation-induced intestinal injuries of mice following total body irradiation [13–16]. We found that the effect of exposure to abdominal irradiation or total body irradiation on Nrf2−/− mice was completely opposite, that was, the survival of Nrf2−/− mice after abdominal irradiation was higher than that of Nrf2+/+ mice, but the survival of Nrf2−/− mice after total body irradiation was lower than that of Nrf2+/+ mice. It is known that abdominal irradiation can only lead to radiation-induced intestinal injuries, and total body irradiation can result in not only radiation-induced intestinal injuries, but also hematopoietic system injury. We speculate that the reason of the protective effect of Nrf2 knockout on radiation-induced intestinal injuries may be attributed to different methods of radiation exposure. It has been reported that the roles and functions of Nrf2 in intestinal stem cells and hematopoietic stem cells (HSCs) are diverse. Hochmuth et al., found that CncC, which is a Drosophila homolog of Nrf2, was constitutively active in intestinal stem cells of Drosophila and maintained intestinal stem cells quiescent with low radical oxygen species concentration [37]. However, Nrf2 deficiency attenuated the reconstitution capacity of hematopoietic stem cells after the transplantation, indicating that Nrf2 was required for the hematopoietic stem cells’ maintenance [6,38]. The different role of Nrf2 in intestinal stem cells and hematopoietic stem cells may explain why Nrf2 knockout has the protective effect on radiation-induced intestinal injuries in this study, while Nrf2 alleviates radiation-induced hematopoietic system damage. It is important to note that non-Nrf2 knockout mice were adopted to investigate the role of Nrf2 in radiation-induced intestinal injuries in previous studies, and Nrf2 knockout mice were used to explore the effect of Nrf2 knockout in the present study. The effect of Nrf2 knockout mice on radiation-induced intestinal injuries was markedly different from that of non-Nrf2 knockout mice, which may be one of reasons that our results were inconsistent with that of previous reports. However, it is necessary to further verify whether Nrf2 knockout truly has the protective effect on radiation-induced intestinal injuries following abdominal irradiation, but not total body irradiation. In addition, the associated mechanisms by which Nrf2 knockout facilitated the protective effect on radiation-induced intestinal injuries have yet to be elucidated thoroughly. Further study is needed to understand the contribution of radical oxygen species to radiation-induced intestinal injuries and explore how Nrf2 knockout enhances the NF-κB activity.

4. Materials and Methods

4.1. Mice

Wild-type (Nrf2+/+) C57BL/6j mice were purchased from Beijing HFK Bioscience Co. Ltd. (Beijing, China). Nrf2 knockout (Nrf2−/−) C57BL/6j mice were generously provided by Thomas W. Kensler of the University of Pittsburgh and backcrossed into the C57BL/6j background in our laboratory. All mice were housed in a temperature-controlled, specific pathogen free environment with a 12-h light/dark cycle, and fed standard chow and water. All experimental procedures and protocols were approved by the Animal Care and Ethics Committee of the Institute of Radiation Medicine (approval number: 170512, 5 August 2016), and performed according to the principles of the Institutional Animal Care and Ethics Committee guidelines.
4.2. Irradiation of Mice

Mice were anaesthetized with an intraperitoneal injection of 3.5% chloral hydrate (0.1 mL/10 g body weight) and placed on a platform to receive 13 Gy abdominal irradiation. The irradiated field was from the xiphoid process to the pubic symphysis. Mice were exposed to ionizing irradiation in a 137 Cesium γ-ray irradiator (Atomic Energy of Canada, Chalk River, ON, Canada) at a rate of 1 Gy/min.

4.3. Analysis of the 30-Day Survival Rate

Mice were monitored for up to 30 days after 13 Gy abdominal irradiation or 7.2 Gy total body irradiation. The survival curves and the body weight curves were plotted by the Kaplan–Meier method and the ratio of body weight to initial weight, respectively.

4.4. Organ Coefficient Calculation

The spleen and thymus were collected and weighted after mice were euthanized. The organ coefficients were calculated by dividing individual organ weight (mg) with their body weight (g). The spleen coefficient = spleen weight (mg)/mice weight (g), the thymus coefficient = thymus weight (mg)/mice weight (g).

4.5. Peripheral Blood Cells Counts

Whole blood samples were obtained from the orbital sinus of mice. The numbers of various blood cell types, white blood cells (WBC), platelets (PLT), and lymphocytes were counted using a MEK-7222k hemocytometer (Nihonkohden Corp, Tokyo, Japan).

4.6. Analysis of DNA Damage to Peripheral Blood Lymphocytes

The comet assay, also called single-cell gel electrophoresis (SCGE), was performed under alkaline conditions as described by Banath et al. [39]. Whole blood was obtained from the orbital sinus of mice. Lymphocytes were harvested from peripheral blood using lymphocyte separation medium, mixed with low-melting agarose and fixed onto frosted slides pre-coated with normal melting agarose. The slides were immersed in cold, freshly made alkalineysis solution at 4 °C for 2.5 h and then rinsed with PBS. Slides were placed in a horizontal electrophoresis device pre-filled with cold alkaline electrophoresis buffer for 20 min at 4 °C, followed by electrophoresis for 20 min (30 V, 300 mA). The slides were fixed in neutralization buffer, stained with ethidiumbromide (2.5 µg/mL), and visualized by a ECLIPSE 90i fluorescence microscopy (Nikon, Tokyo, Japan). Two hundred cells were analysed using the Comet Assay Software Project (CASP, Wroclaw University, Poland) [40], and the DNA percentage in the comet tail (Tail DNA%), tail moment (TM), and olive tail moment (OTM) were recorded to describe DNA damage to the lymphocytes.

4.7. Pathological Analysis

Dissected small intestines were harvested after mice were euthanized and fixed in neutral formalin for 24 h prior to being embedded in paraffin. Paraffin-embedded small intestines were cut into 5 µm sections at the maximum cross-section. Sections were stained with haematoxylin/eosin and alcian blue.

4.8. Immunohistochemical Analysis

Paraffin-embedded sections were dewaxed and rehydrated. The sections were then immersed in Tris/EDTA (pH 9.0) and incubated for 15 min at 98 °C for antigen retrieval. The sections were incubated with serum for 30 min at room temperature to block non-specific antigen-binding sites. Anti-Ki67 antibody, anti-NF-κB p65 antibody, anti-Lgr5 antibody (Abcam, Cambridge, MA, USA), anti-lysozyme antibody, and anti-Villin antibody (Proteintech, Chicago, IL, USA) was diluted as recommended by the manufacturer, added to the sections, and incubated overnight at 4 °C. Unbound antibodies were washed away with PBS. Sections were incubated in 0.3% H2O2 for 15 min to block
endogenous peroxidase. Secondary antibody was added and incubated at 37 °C for 2 h. DAB was added to detect positive cells.

4.9. Apoptosis Assay

The apoptosis assay was tested using one step TUNEL apoptosis assay kit (Beyotime, Shanghai, China). Paraffin-embedded sections were dewaxed and rehydrated. The sections were then treated with proteinase K for 5 min in a 37 °C water bath and incubated with TUNEL detection liquid for 1 h at 37 °C. After washes, sections were counterstained with DAPI. These slides were observed under an ECLIPSE 90i fluorescence microscope (Nikon). TUNEL-positive cells in a field of view were quantified.

4.10. Isolation of Crypt Single Cells

The method of isolating the intestinal crypts of mice was described by Mahe et al. [41]. Briefly, the small intestines of mice were sliced and opened longitudinally. After washing with ice-cold PBS, the small intestines were cut into small pieces and then incubated in ice-cold PBS containing 2 mM EDTA for 30 min. After rinsing twice with ice-cold PBS, the fragments were resuspended in ice-cold PBS containing 54.9 mM D-sorbitol and 43.4 mM sucrose, and filtered with a 70-µm filter. After centrifuging, the pellet was incubated in pancreatin until it was digested to single cells, and filtered with a 40 µm cell strainer.

4.11. Preparation of Cytoplasmic and Nuclear Extracts

Cytoplasmic and nuclear extracts were prepared from crypts single cells using NE-PER™ nuclear and cytoplasmic extraction reagents (ThermoFisher, Rockford, IL, USA) according to the manufacturer’s instructions. The single cells pellet was suspended with ice-cold cytoplasmic extraction reagent I for 10 min and then added ice-cold cytoplasmic extraction reagent II. After centrifuging, the supernatant (cytoplasmic extract) was collected. The insoluble fraction was suspended with nuclear extraction reagent. The supernatant (nuclear extract) was collected following centrifuging.

4.12. Western Blot Analysis

The concentrations of protein lysates were quantified using a BCA protein kit (Beyotime). Samples containing equal amounts of protein (20 µg) were mixed with loading buffer containing 5% 2-mercaptoethanol, heated for 10 min at 99 °C, and loaded onto a 10% SDS-PAGE gel. After electrophoresing, the proteins from the gel were then transferred onto polyvinylidene difluoride membranes. These membranes were blocked with 5% milk and 0.1% Tween 20 in Tris-buffered saline, and incubated overnight at 4 °C with anti-NF-κB p65 (Abcam), anti-LaminB1, anti-GAPDH and anti-β-Actin (Proteintech). Then, the appropriate horseradish peroxidase-conjugated secondary antibody was added on the membranes at room temperature. Finally, the proteins were detected with chemiluminescent substrate.

4.13. RNA Isolation and Real-Time Polymerase Chain Reaction

Total RNA was extracted from the small intestines using TRIzol Reagent according to the manufacturer’s instructions (Invitrogen, Carlsbad, CA, USA). A reverse transcription reaction was performed with 10 µg aliquot of each RNA sample using oligo-dT random primers and reverse transcriptase (Takara, Dalian, China), and real-time PCR was performed using a SYBR Premix Ex TaqTM II (Takara). Primer sequences were as follows:

Bcl-2: 5′-AGGATTGTGGCCTTTTGA-3′ (forward) and 5′-CAGATGCCGCCTCACGACT-3′ (reverse); XIAP: 5′-TGGAACCTCACTGTGGAA-3′ (forward) and 5′-CTGCCCCACCTTGATAC-3′ (reverse); uPA: 5′-CGCACACTGCTTCATTCAAC-3′ (forward) and 5′-GCTGCTCCACCTCAA-3′ (reverse).
4.14. Statistical Analysis

The data were analysed using Student’s t-test and expressed as the mean ± SD. The Kaplan–Meier method was used to analyse animal survival curves. A p-value of <0.05 was taken to be significant.

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Conflicts of Interest: The authors declare no conflict of interest.

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