

Establishing the Impact of Vascular Damage on Tumor Response to High-Dose Radiation Therapy

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Abstract

Approximately half of all patients with cancer receive radiotherapy, which is conventionally delivered in relatively small doses (1.8–2 Gy) per daily fraction over one to two months. Stereotactic body radiation therapy (SBRT), in which a high daily radiation dose is delivered in 1 to 5 fractions, has improved local control rates for several cancers. However, despite the widespread adoption of SBRT in the clinic, controversy surrounds the mechanism by which SBRT enhances local control. Some studies suggest that high doses of radiation (≥ 10 Gy) trigger tumor endothelial cell death, resulting in indirect killing of tumor cells through

nutrient depletion. On the other hand, mathematical models predict that the high radiation dose per fraction used in SBRT increases direct tumor cell killing, suggesting that disruption of the tumor vasculature is not a critical mediator of tumor cure. Here, we review the application of genetically engineered mouse models to radiosensitize tumor cells or endothelial cells to dissect the role of these cellular targets in mediating the response of primary tumors to high-dose radiotherapy *in vivo*. These studies demonstrate a role for endothelial cell death in mediating tumor growth delay, but not local control following SBRT.

Introduction

Stereotactic body radiation therapy (SBRT) is now routinely used in the clinic to deliver large doses of daily radiation in a small number of fractions to a very precise target volume (1). This type of radiation delivery was first developed for the treatment of brain tumors in the early 1950's (2) and was not shown to be feasible in the setting of extracranial tumors until the 1990's (3, 4). With advances in imaging and medical physics, patients with non-small cell lung cancer with inoperable tumors were among the first patients to be treated with SBRT (5). Remarkably, local control rates reached approximately 80%–90% across several clinical trials (6–9), significantly higher than historical rates achieved with conventional, low dose per fraction radiotherapy. Therefore, SBRT is now being employed for the treatment of a wide variety of cancers, including non-small cell lung cancer, hepatocellular carcinoma, and oligometastatic disease at a variety of sites, with impressive rates of local control (10–15). Despite the efficacy of SBRT in the clinic, controversy surrounds the mechanism by which high-dose radiotherapy leads to tumor eradication. Two competing models on the mechanistic basis for improved efficacy with SBRT predominate: (i) functional impairment of tumor vasculature results in indirect cell death by killing tumor cells that would otherwise not have died from radiation or (ii) higher radiation dose per fraction increases the biologically effective dose, which leads to more direct tumor cell

death. By understanding the mechanism of improved local control by SBRT, new therapeutic approaches with targeted agents can be designed to enhance the therapeutic ratio.

Indirect Tumor Cell Killing Hypothesis

Data supporting the first model were recently reviewed by Song and colleagues (16). They argue that radiation-induced tumor cell death alone is insufficient to explain the increased rates of local control achieved with SBRT as radiation doses are not large enough to kill every tumor cell (17–23). For example, *in vitro* survival studies using human tumor cell lines suggest that single radiation doses between 22 and 36 Gy are required to eradicate the number of cells estimated to be present in a 3-cm tumor (24). Although single doses of 30 Gy and higher are used in the clinic to achieve local control (25, 26), the *in vitro* survival studies found that the curative radiation dose increased by a factor of approximately three when 20% of the tumor cells were assumed to be hypoxic as hypoxic tumor cells are more resistant to radiation (24). As the calculated radiation dose for *in vitro* cell killing exceeded the SBRT dose used in the clinic, the investigators concluded that the efficacy of SBRT required indirect in addition to direct tumor cell death (27). Further supporting this model, the number of surviving tumor cells in a fibrosarcoma xenograft several days after high-dose radiotherapy was significantly less than the number of cells immediately following radiation exposure, suggesting that indirect killing mediates a second wave of tumor cell death after irradiation (21).

Early studies using tumor allografts revealed that low doses of radiation (<5 Gy) caused only a temporary impairment in vascular function, while irradiation with 10 Gy triggered a persistent decrease in circulating blood volume in tumors (28, 29). A connection between such vascular dysfunction and indirect tumor cell death has been well described (30–33). Garcia-Barros and colleagues assessed endothelial cell death in xenograft tumors along a range of radiation doses, but reported endothelial cell death only at doses greater than 8 Gy (34), suggesting that

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radiation doses associated with conventional radiotherapy are not capable of killing vascular endothelial cells. In this study, tumors transplanted into mice lacking acid sphingomyelinase were resistant to endothelial cell apoptosis and had a shorter growth delay following high-dose radiotherapy when compared with tumors implanted into wild-type mice (34). The authors concluded that a decrease in endothelial cell death caused resistance to SBRT, although this finding was later challenged (35, 36). Recently, these investigators further characterized the response of transplanted tumors in acid sphingomyelinase-knockout mice and proposed a new mechanism, which did not rely on endothelial cell death, to explain the indirect cell death associated with SBRT. They argued that high single-dose radiotherapy triggers ischemia/reperfusion injury that interferes with homologous recombination, thereby preventing the repair of radiation damage in tumor cells (37). Regardless of the mechanism of indirect cell death, these studies indicate that high dose radiotherapy (≥ 10 Gy) can cause indirect tumor cell death and suggest that this "new biology" could explain increased efficacy of SBRT.

Proponents of the indirect cell death hypothesis contend that the linear quadratic model, which is commonly used to model radiation-induced cell death, does not accurately estimate tumor cell killing at high doses per fraction. It is argued that this mathematical model overestimates the amount of cell death caused directly by radiation-induced DNA damage because it does not account for indirect cell death mediated by the immune system or vascular dysfunction (38). Incorporation of a secondary tumor cell killing mechanism into the model results in a downward bend of the dose-response curve at radiation doses greater than 10 Gy (27). Thus, the adapted model predicts that SBRT causes more cell death than would be estimated by the standard linear quadratic model, which may explain the increased rates of local control with SBRT.

Direct Tumor Cell Killing Hypothesis

The second model rejects the need for a "new biology" to explain the efficacy of SBRT and argues that improved outcomes after SBRT simply reflect increased direct tumor cell death from the higher radiation dose (39, 40). Experimental evidence supporting this model comes from Budach and colleagues, who investigated the radiation dose required to cure 50% of tumors implanted into either nude or SCID mice. Although SCID mice are 3-fold more radiosensitive than nude mice due to a mutation in *DNA-dependent protein kinase* that impairs DNA repair, no difference was detected in the single dose of radiation needed to achieve local control in a panel of human tumors (41). These data indicate that the radiosensitivity of tumor stromal cells, such as endothelial cells, is not a critical regulator of tumor cure. This study supports the direct cell killing hypothesis, which asserts that stromal cell death is not a critical mediator of local control. Importantly, the direct tumor cell killing hypothesis does not imply that stromal cells lack any role in tumor response to radiotherapy. Indeed, using doses of radiation that were not sufficient for tumor cure, Budach and colleagues observed increased growth delay of some xenografts in radiosensitive SCID hosts (41). Other studies combining molecularly targeted drugs with radiotherapy have also demonstrated that an increase in tumor growth delay does not always translate to an enhanced rate of local control with a curative radiation dose (42, 43). These

studies highlight the importance of characterizing the impact of stromal cell death directly on local control.

Furthermore, local control for patients with non-small cell lung cancer treated with conventional radiotherapy or SBRT is enhanced with an increase in the biologically effective dose (BED; ref. 44). BED provides a mathematical framework to adjust for dose fractionation to compare different radiotherapy prescriptions (45). Correcting the human lung cancer dataset for BED resulted in comparable rates of local control between the conventional radiotherapy and SBRT arms (39). These data suggest that the higher BED associated with SBRT may completely explain the increased rates of local control, thus eliminating the need for indirect tumor cell killing to explain the efficacy of SBRT.

Proponents of the direct tumor cell killing model also argue that the linear-quadratic model accurately predicts radiation-induced cell killing at high, single fraction doses (46). Data collected in irradiated rat spinal cord, mouse skin, and mouse small intestine *in vivo* fit the linear-quadratic model over a wide range of doses (47–49). Likewise, local control data in patients with primary lung cancer or brain metastases irradiated with SBRT were better fit by the linear-quadratic model than any adapted model that factored in indirect tumor cell death at high doses (50). Therefore, these results suggest that the linear-quadratic model should not be replaced with an adapted model that incorporates increased cell death at high doses and they argue against the need for a "new biology" to explain the efficacy of SBRT.

Using Primary Mouse Models to Dissect the Cellular Target of Radiotherapy

To address the controversy regarding the cellular target that mediates the efficacy of SBRT, we utilized genetically engineered mouse models (GEMM) of cancer. Although transplanted tumor models are relatively fast and inexpensive for radiotherapy studies, experiments with xenografts are often unable to predict clinical outcomes (51–53). GEMMs, by contrast, require additional time and expense to develop but enable the study of autochthonous tumors that arise in their native microenvironment with an intact immune system (51). These primary tumors may more accurately model the role of stromal cells in human cancer compared with transplanted tumors (54, 55). Indeed, patterns of tumor vascularization and hypoxia differ significantly between spontaneous and transplanted models (51, 56, 57). Therefore, GEMMs may better recapitulate the response of human cancer to treatment in the clinic (58–60). Like all model systems, GEMMs have limitations, including the limited number of genetic alterations and decreased heterogeneity. Nevertheless, the intact stromal cell compartment and similarity with human cancer are critical considerations when investigating the impact of the tumor microenvironment on response to radiotherapy.

Dual recombinase technology

The utilization of the Cre/loxP system to mediate site-specific recombination in the 1980's (61, 62) revolutionized the field of cancer biology by enabling the generation of sophisticated GEMMs of cancer. Cre is a bacteriophage P1-derived enzyme that recombines a pair of DNA sequences termed loxP sites. These sites are often engineered into genomic DNA to flank a target or "floxed" gene so that the flanked region will be deleted in the presence of Cre. In a similar manner, the Flp-FRT system, adopted

from *Saccharomyces cerevisiae* in the mid-1990's, employs the recombinase activity of flippase (Flp) to delete genomic targets flanked by Flp recombinase target (FRT) sites, referred to as "FRTed" regions (63). The recombinase activity of Cre and Flp can be controlled by a variety of mechanisms, including viral delivery and expression under the control of a cell-type-specific promoter (64). Thus, Cre and Flp can temporally and spatially restrict recombination to study tumor development and response to radiotherapy (65). By combining Cre and Flp in a GEMM, the sophisticated interactions between tumor cells and the supporting stroma can be dissected by using one recombinase for tumor initiation and the second recombinase to genetically manipulate a specific stromal compartment (66, 67). Recently, we applied this dual recombinase technology to investigate the role of endothelial cells in mediating the response of primary tumors to SBRT (68–70).

radiosensitization of specific cell types within a tumor

As DNA damage is a key cause of cell lethality following exposure to ionizing radiation (71), blocking DNA repair mechanisms may increase the number of unresolved DNA double-stranded breaks thus enhancing radiation-induced cell death. Ataxia telangiectasia mutated (ATM) is a protein kinase that regulates homologous recombination and cell-cycle arrest through the phosphorylation of a large number of downstream targets, including p53, MRE11, RAD50, BRCA1, and CHK2 (72–75). In addition, patients with inherited homozygous mutations in *ATM*, human cell lines lacking functional ATM, and *Atm*-knockout mice are hypersensitive to radiation (76–78). Therefore, genetic deletion of *Atm* in either primary tumor or endothelial cells can be utilized to radiosensitize specific cell populations to define the roles of these cell types during tumor response to radiotherapy. To specifically assess the impact of vascular damage on tumor response to radiation, we employed dual recombinase technology to delete floxed alleles of *Atm* specifically in endothelial cells. In this model, viral delivery of Flp recombinase initiated tumorigenesis by deleting FRTed alleles of the tumor suppressor *p53* and drove expression of the mutated oncogene *Kras*^{G12D} by deleting an upstream FRTed STOP cassette at the endogenous promoter (79). Cre recombinase was not used to initiate the primary tumor in this system, but instead was expressed under the control of the *VE-Cadherin* promoter to direct Cre expression to endothelial cells to delete floxed *Atm* alleles specifically in the vasculature without affecting *Atm* gene expression in the primary tumor cells (Table 1). To specifically radiosensitize tumor cells, we used Cre-loxP technology to simultaneously initiate tumorigenesis and modulate expression of *Atm* exclusively within the tumor cells. Cre expression in tumor initiating cells activated expression of oncogenic *Kras*^{G12D} by deleting a floxed STOP cassette and deleted floxed alleles of the tumor suppressor *p53* in addition to *Atm*. Because *Atm* was deleted in the tumor-initiating cells, which gave rise to the primary tumor,

this genetic approach specifically enhanced the radiosensitivity of the tumor cells (Table 1).

Primary sarcomas

A GEMM of primary soft tissue sarcoma (79–81) was the first setting in which we employed dual recombinase technology to examine the impact of endothelial or tumor cell radiosensitization on tumor response to SBRT. In this primary mouse model, an SBRT dose of 20 Gy caused vascular injury as measured by dual energy micro-CT and histology (82). To characterize whether endothelial cells are critical targets of SBRT, we used adenoviral delivery of Flp to initiate sarcomagenesis in *FRT-STOP-FRT (FSF)-Kras*^{G12D}; *p53*^{FRT/FRT}; *VE-Cadherin-Cre*; *Atm*^{FL/+} (*KPVA*^{FL/+}) or *Atm*^{FL/FL} (*KPVA*^{FL/FL}) mice (69). Thus, primary sarcomas expressing oncogenic *Kras*^{G12D} with both alleles of *p53* deleted were compared, with one cohort lacking expression of both alleles of *Atm* in endothelial cells (*KPVA*^{FL/FL}) and the other retaining expression of one wild-type allele of *Atm* in endothelial cells (*KPVA*^{FL/+}). As expected, loss of ATM signaling in the vasculature enhanced endothelial cell death 24 hours post irradiation with 20 Gy. Consistent with results in transplanted tumor models, radiosensitization of the endothelial cell compartment of primary sarcomas resulted in vascular dysfunction, as indicated by a decrease in perfusion after radiation exposure. Furthermore, an increase in the total amount of cell death in *KPVA*^{FL/FL} tumors suggested that functional changes to the tumor vasculature triggered indirect killing of adjacent tumor cells.

To evaluate the impact of enhanced vascular dysfunction after SBRT, tumor-bearing mice were treated with a single 20 Gy dose of focal irradiation (69). Growth delay to a volume tripling endpoint in tumors lacking expression of *Atm* in the vasculature was significantly longer, when compared with control tumors that retained expression of one allele of *Atm*. These results in a primary sarcoma model support the indirect tumor cell hypothesis by demonstrating that an increase in the number of dying tumor cells after SBRT leads to an increase in growth delay. However, these data are not sufficient to conclude that the increased indirect tumor cell death caused by vascular dysfunction contributes to the efficacy of SBRT to achieve local control, which is a more relevant clinical endpoint. To assess the role of endothelial cell radiosensitivity on primary tumor eradication by SBRT, sarcomas in *KPVA*^{FL/+} and *KPVA*^{FL/FL} mice were treated with a curative single dose of 50 Gy (68). Although 50 Gy was sufficient to cure approximately 10% of the sarcomas, mice bearing tumors with enhanced vascular radiosensitivity did not achieve a higher rate of local control. Why was the rate of local failure the same in tumors with enhanced indirect tumor cell killing? One potential explanation is that different tumor cells have different susceptibilities to indirect cell killing. Just as hypoxic tumor cells are resistant to radiotherapy, cells that have adapted to survive in the hypoxic microenvironment far from tumor vasculature may also be resistant to indirect cell death caused by vascular dysfunction. In this

Table 1. Genetic changes within primary tumor cells and tumor vasculature

	VE-Cadherin-Cre; <i>Atm</i> ^{FL/FL}		Viral Cre + <i>Atm</i> ^{FL/FL}	
	Tumor cell	Endothelial cell	Tumor cell	Endothelial cell
No Cre recombinase	Wild-type	N/A	N/A	Wild-type
Cre recombinase	No effect	<i>Atm</i> deleted	<i>Atm</i> deleted	No effect
Resulting radiosensitivity	Wild-type	Radiosensitive	Radiosensitive	Wild-type

Abbreviation: N/A, not applicable.

scenario, the radiation dose required to kill every hypoxic tumor cell with the capacity to cause local recurrence would not be affected by increased indirect tumor cell death adjacent to blood vessels. Regardless of the explanation, these findings support a role for endothelial cell death in sarcoma growth delay following SBRT, but not in local control following high single-dose irradiation.

As a positive control for the ability to modulate rates of tumor eradication with SBRT, we also deleted *Atm* specifically in tumor parenchymal cells. *Pax7-CreER; LoxP-STOP-LoxP (LSL)-Kras^{G12D}; p53^{FL/FL}; Atm^{FL/+} (P7KPA^{FL/+})* and *Atm^{FL/FL} (P7KPA^{FL/FL})* mice were injected into the gastrocnemius muscle with 4-hydroxy-tamoxifen to activate Cre recombinase to initiate sarcomagenesis and delete *Atm* within the same cell population (68). Deletion of both floxed alleles of *Atm* (*P7KPA^{FL/FL}*) within the tumor cells of the primary sarcoma resulted in enhanced radiosensitivity. This radiosensitivity translated to a significantly improved tumor response to 50 Gy compared with tumors with deletion of one floxed allele of *Atm* (*P7KPA^{FL/+}*), as measured by the time to tumor volume tripling and rate of local control. Collectively, these findings in a primary sarcoma mouse model demonstrate that tumor cell death, rather than endothelial cell death, is a critical mediator of achieving local control following SBRT, which supports the direct cell killing hypothesis.

Primary lung cancer

In the clinic, SBRT is routinely used to treat inoperable non-small cell lung cancer because of the high rate of local control (4). To study the role of endothelial cell death in SBRT for lung cancer, we utilized a sophisticated GEMM of non-small cell lung cancer to radiosensitize the tumor vasculature (79, 83). An adenovirus expressing Flp recombinase was administered intranasally to *KPVA^{FL/+}* and *KPVA^{FL/FL}* mice to initiate tumorigenesis in the lung epithelium, while the Cre-driver, *VE-Cadherin*, mediated recombination of *Atm* in the vasculature (70). Upon detection of lung tumors by micro-CT imaging, mice were treated with a single 15 Gy dose of whole-thorax irradiation and individual tumors were monitored every 2 weeks by micro-CT to evaluate tumor growth. As expected, an increase in tumor endothelial cell death as well as total cell death was observed in *KPVA^{FL/FL}* mice 24 hours post radiation exposure, which supports the occurrence of indirect tumor cell death. Despite the enhanced radiosensitivity of the tumor vasculature and indirect tumor cell death, no significant difference in tumor growth delay was detected 2–6 weeks after 15 Gy in tumors lacking *Atm* expression in the vasculature as compared with tumors retaining one wild-type allele of *Atm*. A small, although not statistically significant, decrease in tumor volume was detected in *KPVA^{FL/FL}* tumors 8 weeks after irradiation. Overall, this study suggested that endothelial cell death has only a modest effect on the response of primary lung tumors to SBRT.

To investigate the impact of tumor parenchymal cell radiosensitization on the lung tumor response to SBRT, lung tumors were initiated in *LSL-Kras^{G12D}; p53^{FL/FL}; Atm^{FL/+} (KPA^{FL/+})* and *Atm^{FL/FL} (KPA^{FL/FL})* mice (70). In this model, inhalation of a lentivirus expressing Cre facilitated the recombination of *Atm* specifically in lung tumor-initiating cells. Disruption of ATM signaling enhanced lung cancer cell radiosensitivity in a colony survival assay *in vitro*. Radiosensitization of lung tumor cells in *KPA^{FL/FL}* mice translated to a significant decrease in tumor volume 6–8 weeks after 15 Gy irradiation to the whole thorax.

Taken together, these results support a model in which tumor cells play a larger role than endothelial cells in regulating the response of primary lung cancers to SBRT and provide further evidence for the direct tumor cell killing hypothesis to explain the efficacy of SBRT.

Discussion

The large dose per fraction (≥ 10 Gy) radiotherapy schedules used to treat cancer with SBRT have increased local control rates for several diseases when compared with 2 Gy daily fractions delivered with conventional radiotherapy (10, 11, 84–86). However, biologists and physicists continue to debate the mechanism by which SBRT improves tumor response to radiotherapy. Many investigators have shown that radiation can induce proliferative defects in endothelial cells, thereby triggering endothelial cell death, increased vascular permeability, and indirect tumor cell death (34, 69, 70, 87–90). The controversy surrounding the mechanism of SBRT revolves around whether such functional changes to the vasculature and the accompanying indirect tumor cell death are sufficient to enhance tumor eradication. While some investigators argue that indirect cell killing caused by vascular impairment can regulate tumor cure in response to radiation (34, 91, 92), others have employed mathematical modeling to counter that the level of endothelial cell death does not accurately predict clinical outcomes and therefore increased dose per fraction simply kills more tumor cells directly (41, 93, 94). These divergent views may be reconciled by strictly limiting the conclusions of transplanted tumor models to the endpoints studied: indirect tumor cell death does occur as a consequence of vascular injury following single high-dose radiotherapy, and this increases tumor growth delay but not local control. Indeed, others have previously demonstrated that prolonged tumor growth delay using a radiotherapy regimen with a targeted agent does not always translate into increased tumor cure when a curative dose of radiotherapy is delivered (42, 43). This conclusion is consistent with the results of Budach and colleagues studying tumors transplanted into nude and SCID mice (41) as well as our results using GEMMs of sarcoma (68, 69) and lung cancer (70) following SBRT. Furthermore, this discrepancy between growth delay endpoints and tumor cure may explain why some clinical trials of radiotherapy and targeted agents do not recapitulate preclinical data (95).

Tumor cells as critical mediators of single high-dose radiotherapy

By genetically manipulating the radiosensitivity of either tumor cells with Cre-loxP technology or endothelial cells with dual recombinase technology, we generated data to address the controversy of the critical cellular target of SBRT. Importantly, SBRT triggered an increase in fractional blood volume and vascular permeability in primary soft tissue sarcomas in a dose-dependent manner (82, 96). Radiosensitization of the endothelial cells through the deletion of *Atm* further disrupted AngioSense accumulation and blood flow into sarcomas, indicating an impairment of vascular function (69). Importantly, radiation-induced endothelial cell death in both the primary sarcoma and lung tumor models triggered indirect cell death of neighboring tumor cells (69, 70). Thus, primary tumors lacking *Atm* expression specifically in the vasculature represent powerful tools to assess whether vascular impairment can regulate growth delay and local

tumor control following radiotherapy. Despite the observed increase in endothelial cell death and indirect tumor cell death in these GEMMs following irradiation with 15–20 Gy, tumors lacking functional ATM in the vasculature displayed only a modest increase in growth delay (69, 70). Remarkably, endothelial cell radiosensitization and the resulting vascular dysfunction did not enhance local control in sarcomas treated with a curative radiation dose (69). While additional experiments with dual recombinase technology are needed to determine whether these results extend beyond soft tissue sarcoma and lung cancer to other primary cancer mouse models, the available data in these two autochthonous tumor models suggest that the indirect tumor cell killing hypothesis may play a role in

extending tumor growth delay and palliation with radiotherapy, but is unlikely to contribute to the impressive local control that can be achieved with SBRT (Fig. 1).

In contrast, radiosensitization of the tumor cell population in the GEMMs significantly prolonged growth delay of lung tumors (70) and increased the incidence of sarcoma eradication (69). These results support the direct tumor cell killing hypothesis, in which an increase in direct tumor cell death can promote local control following high dose per fraction radiotherapy (Fig. 1). Taken together, these results suggest that a "new biology" mediated via endothelial cell death is not required to explain the increased rate of local control observed with SBRT.

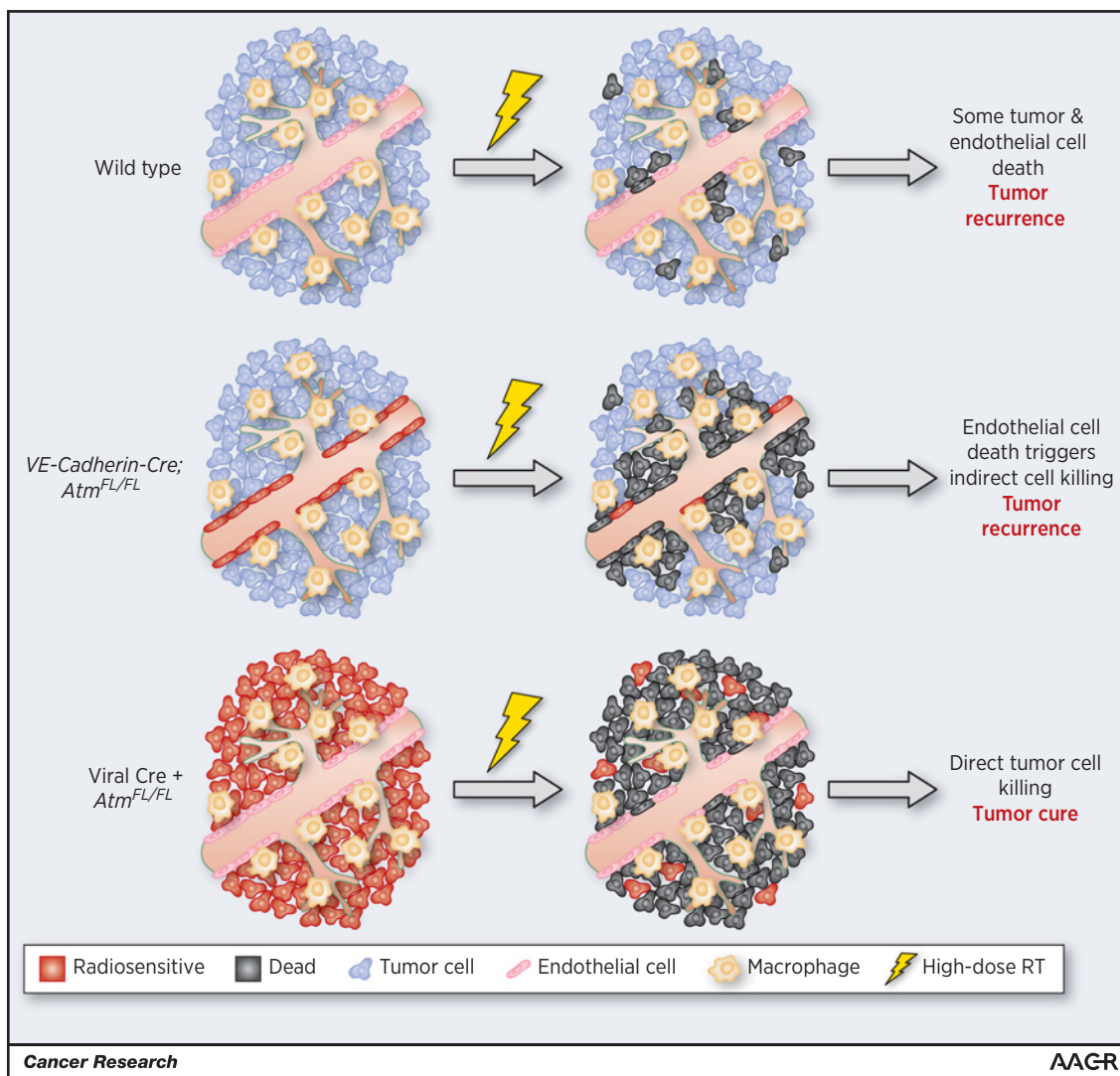


Figure 1.

Enhancing tumor parenchymal cell radiosensitivity preferentially promotes tumor cure. In wild-type tumors that maintain functional ATM signaling in all cellular compartments, high-dose radiotherapy (RT) induces both endothelial and tumor cell death. Deletion of both alleles of *Atm* in the tumor vasculature using dual recombinase technology increases the amount of endothelial and total cell death by approximately two- to threefold. Despite the resulting vascular dysfunction and indirect tumor cell killing, the tumors still recur following high-dose radiotherapy. In contrast, radiosensitizing tumor parenchymal cells through the deletion of *Atm* enhances both radiation-induced growth delay and tumor cure.

Stromal cells as potential mediators of SBRT

Although endothelial cell death may not regulate tumor eradication by SBRT, this does not rule out the possibility of other stromal cell populations regulating tumor response and cure following SBRT. For example, Brown and colleagues have reported that macrophages and endothelial progenitor cells can replenish tumor endothelium after single high-dose radiotherapy to impact the response of xenografts (97, 98). Similarly, T cells within the immune system can respond to SBRT to promote or impede transplanted tumor response to radiotherapy (99–101). In the future, dual recombinase technology can be applied to these cell populations to investigate their role in primary tumor response to SBRT.

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Disclosure of Potential Conflicts of Interest

D.G. Kirsch reports receiving a commercial research grant from XRAD Therapeutics, Merck, and Eli Lilly & Company, reports receiving other commercial research support from Lumicell Inc., has ownership interest (including patents) in Lumicell Inc. and XRAD Therapeutics, and has a paid consultant/advisory board relationship with Sarcoma Alliance for Research through Collaboration. No potential conflicts of interest were disclosed by the other author.

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