

# UC Davis

## UC Davis Previously Published Works

### Title

Emerging Translational Opportunities in Comparative Oncology With Companion Canine Cancers: Radiation Oncology

### Permalink

<https://escholarship.org/uc/item/4t16q23h>

### Authors

Nolan, Michael W

Kent, Michael S

Boss, Mary-Keara

### Publication Date

2019

### DOI

10.3389/fonc.2019.01291

### Copyright Information

This work is made available under the terms of a Creative Commons Attribution License, available at <https://creativecommons.org/licenses/by/4.0/>

Peer reviewed



# Emerging Translational Opportunities in Comparative Oncology With Companion Canine Cancers: Radiation Oncology

Michael W. Nolan<sup>1,2,3\*</sup>, Michael S. Kent<sup>4</sup> and Mary-Keara Boss<sup>5</sup>

<sup>1</sup> Department of Clinical Sciences, North Carolina State University, Raleigh, NC, United States, <sup>2</sup> Comparative Medicine Institute, North Carolina State University, Raleigh, NC, United States, <sup>3</sup> Duke Cancer Institute, Duke University, Durham, NC, United States, <sup>4</sup> Department of Surgical and Radiological Sciences, School of Veterinary Medicine, University of California, Davis, Davis, CA, United States, <sup>5</sup> Department of Environmental and Radiological Health Sciences, Flint Animal Cancer Center, Colorado State University, Fort Collins, CO, United States

## OPEN ACCESS

### Edited by:

Mark W. Dewhirst,  
Duke University, United States

### Reviewed by:

Maria Shoshan,  
Karolinska Institutet (KI), Sweden  
Karishma Rajani,  
Mayo Clinic, United States  
Mark A. Ritter,  
University of Wisconsin-Madison,  
United States

### \*Correspondence:

Michael W. Nolan  
mwolan@ncsu.edu

### Specialty section:

This article was submitted to  
Cancer Molecular Targets and  
Therapeutics,  
a section of the journal  
Frontiers in Oncology

**Received:** 27 August 2019

**Accepted:** 07 November 2019

**Published:** 22 November 2019

### Citation:

Nolan MW, Kent MS and Boss M-K  
(2019) Emerging Translational  
Opportunities in Comparative  
Oncology With Companion Canine  
Cancers: Radiation Oncology.  
*Front. Oncol.* 9:1291.  
doi: 10.3389/fonc.2019.01291

It is estimated that more than 6 million pet dogs are diagnosed with cancer annually in the USA. Both primary care and specialist veterinarians are frequently called upon to provide clinical care that improves the quality and/or quantity of life for affected animals. Because these cancers develop spontaneously in animals that often share the same environment as their owners, have intact immune systems and are of similar size to humans, and because the diagnostic tests and treatments for these cancers are similar to those used for management of human cancers, canine cancer provides an opportunity for research that simultaneously helps improve both canine and human health care. This is especially true in the field of radiation oncology, for which there is a rich and continually evolving history of learning from the careful study of pet dogs undergoing various forms of radiotherapy. The purpose of this review article is to inform readers of the potential utility and limitations of using dogs in that manner; the peer-reviewed literature will be critically reviewed, and current research efforts will be discussed. The article concludes with a look toward promising future directions and applications of this pet dog “model.”

**Keywords:** radiation oncology, radiobiology, canine comparative radiation oncology, medical physics, animal models of cancer, imaging, theranostics, translational research

## INTRODUCTION

Radiotherapy (RT) is most frequently applied to pet animals with naturally-occurring cancer as a means for improving animal health. However, there is also a long and rich history of studying radiation responses in the tumors and normal tissues of dogs, with a goal of informing therapeutic development in a manner that directly benefits both canine and human cancer patients. Such research forms the basis for the field of canine comparative radiation oncology and radiobiology. The purpose of this review article is to inform readers of the potential utility and limitations of using dogs in that manner; the peer-reviewed literature will be critically reviewed, and current research efforts will be discussed. The article will conclude with a look toward promising future directions and applications of this pet dog “model”; but we will begin with a brief overview of current practices in veterinary radiation oncology, which will enable readers to gain an appreciation for how the field

of canine comparative radiation oncology has developed, and how canine cancer studies can be efficiently, effectively and ethically conducted.

The American Veterinary Medical Association estimated that 38% of households in the United States owned a pet dog in 2018, and the National Cancer Institute's Comparative Oncology Program reports that nationwide ~6 million new canine cancer diagnoses are made annually. RT is an important component of cancer care for dogs; external beam RT is the most common form of treatment. But despite the high rate of canine cancers, and the efficacy of RT, it is actually quite uncommon for dogs to be treated as such; this is largely attributable to limited accessibility. Financial cost is another important barrier to pet owners who might otherwise want to pursue advanced cancer treatments. The economics of veterinary RT are not well-documented. The cost of care varies with geographic location and type of center (university run academic veterinary teaching hospitals vs. private practice specialty clinics), but anecdotally, a course of palliative-intent RT in the US currently costs between ~\$1,000 and \$3,500, and the cost for a definitive-intent course of therapy is often between \$5,000 and \$12,000. This is challenging because fewer than 10% of households carry pet health insurance, and so for most families, veterinary care represents an out-of-pocket expense. Another important factor that limits accessibility is the distance that must be traveled to pursue care; a 2010 report identified 66 veterinary-specific external beam RT centers in the United States (1). While that number is low relative to the number of canine cancer cases, geographic access is improving; indeed, there were only 42 US-based veterinary RT centers in 2001 (2).

Dogs develop a wide range of neoplasms that are treated with RT. Some of the most common indications with translational relevance to humans include soft tissue sarcoma, extremity osteosarcoma, glioma, and genitourinary carcinomas (both muscle-invasive urothelial carcinomas of the urinary bladder, and prostatic carcinomas). Many of these canine cancers share striking similarities with the equivalent human diseases, not only clinically and histologically, but also at a molecular and genomic level. For example, the most commonly altered gene in both canine and human osteosarcoma is TP53; the genomic imbalance in the two species is similar, and expression of several genes (including PTEN and RUNX2) are correlated with ploidy (3–5). Likewise, in addition to certain similarities between human and canine glioma, the process of PDGF-induced gliomagenesis seems to be well-conserved across mammalian species (6, 7).

A majority of canine radiotherapy patients are treated using conventional C-arm linear accelerators. Due to the need for general anesthesia, full-course definitive-intent, treatment protocols tend to be hypofractionated relative to protocols in common use for humans; often a total of 16–20 daily (Monday through Friday) fractions are administered, with fractional doses used for dogs typically ranging from 2.5 to 4 Gy. As with physician-based oncology, there has also been a tremendous shift in recent years toward much higher doses per fraction, with a rapid increase in access to, and application of, stereotactic radiosurgery (SRS) and stereotactic body RT (SBRT) for various malignancies (1).

## HISTORICAL USES OF DOGS IN TRANSLATIONAL RADIATION RESEARCH

### Normal Tissue Toxicity

In an online query of the US National Library of Medicine's PubMed database for the terms "canine" and "radiation," the earliest return was a 1922 article that was written by Stafford Warren and George Whipple; it was published in the *Journal of Experimental Medicine*, and described small intestinal radiosensitivity in dogs (8). This early use of dogs for radiobiology research may have been driven at least in part by the ease of housing and handling dogs, as well as access to hospital-grade irradiation equipment (X- and gamma-ray) for which it would have been straightforward to perform partial or total body irradiations. The similar anatomy of dogs and humans would also have been favorable, thus allowing experiments to be performed with similar radiation field sizes, and targets, as compared with other research species of the times, including various large animal agricultural species (e.g., swine, cattle) and small fish (9–11). Fortunately, it was later learned that DNA repair mechanisms are highly conserved between mammalian species, and that there is high homology between key DNA damage response genes in humans and dogs (12–14).

Dogs have been used to model radiation injury in a wide range of normal tissues. And indeed, just as veterinarians have borrowed from physicians to inform the practice of veterinary radiation oncology, lessons learned from canine radiobiology research have also had important influences to optimize human cancer care. Through the 1980's and 1990's significant efforts at both Colorado State University and the National Cancer Institute (NCI) were directed toward defining the tolerance of canine tissues to intra-operative RT (IORT); similar studies were performed with multi-fraction irradiation protocols to estimate alpha-to-beta ratios for various normal tissues. These studies evaluated both conventional pathologic endpoints, and a wide range of clinically-relevant functional endpoints. For example, in a canine study, the volume of lung irradiated was found to be a critical limiting factor in thoracic RT (15). The assumption had already been made that a dose-volume relationship existed for lung tissue (16); the TD<sub>50/5</sub> (the radiation dose that causes toxicity in 50% of patients, 5 years after irradiation) was estimated to be 65 Gy when 33% of the total lung volume was irradiated. However, there were no cases of severe symptomatic pneumonitis in dogs for whom 33% of lung received up to 72 Gy; this suggests that when patients have otherwise healthy lungs (with normal compensatory function) the dose tolerance limits for lung may actually be higher than those originally proposed by Emami et al. (16).

It is beyond the scope of this manuscript to review all normal tissue injury studies for which dogs were utilized; however, another noteworthy endeavor was the use dogs to investigate tolerance of peripheral nerves to IORT. In the 1990's, investigators at NCI demonstrated a progressive sciatic neuropathy in American Foxhounds: with 3.5 years of follow-up, doses of up to 20 Gy were not found to cause clinically significant neuropathies, but 5 years after irradiation, doses of 15 Gy or more of IORT were injurious (17, 18). Similar findings

were reported in beagles, whose nerve function was evaluated via histology, neurologic examination, and electrophysiology (19). These experiments confirmed that nerves are dose-limiting for IORT, and that the neurovasculature plays a critical role in nerve injury. These studies also yielded insight regarding combinatorial therapies; while combining IORT with external beam RT resulted in a similar incidence and latency of neuropathy vs. IORT alone, the combination of IORT with hyperthermia led to increased incidence and decreased time to onset of nerve damage. More recently, hounds were used to demonstrate that both altered function of the internal pudendal artery, and pudendal nerve, accompany erectile dysfunction that follows treatment of prostate cancer with SBRT (20). That study identified potential strategies for mitigating radiation-induced sexual dysfunction in men. It also provided data to support the clinical observation that high-dose SBRT may lead to severe colorectal injury, for which latency can be increased, and incidence reduced via either reduction of the radiation dose, or increasing the duration of the interfraction interval (21, 22).

While some laboratory groups have studied both purpose-bred and tumor-bearing pet dogs, the latter have largely been used to study radiation effects on tumors, rather than normal tissue. Rainer Storb's team at the Fred Hutchinson Cancer Research Center has taken this approach for many years, demonstrating dose, fractionation and dose rate effects of total body irradiation in research colony animals, and providing proof-of-principle for clinical applications in the management of lymphomatous diseases in pet dogs (23–36). With rising clinical interest in bone marrow transplantation (including total body irradiation for marrow ablation) in pet dogs with high grade (non-Hodgkin's) multicentric lymphomas, opportunities exist to test novel radiomitigators (37). Certainly, infrastructure exists to enable canine clinical trials to study radiation-modifying drugs and devices. A classic example is a 1986 paper describing the role of canine comparative radiation oncology research in early development of the FDA-approved radioprotector amifostine (WR-2721); a bi-institutional canine clinical trial was performed, in which 73 pet dogs with spontaneously occurring soft tissue sarcomas were randomized (over a period of 3 years) to two dose response assays to receive irradiation alone, or with the radioprotector WR-2721 (38). With the dose and schedule used (40 mg/kg given intravenously, 15 min before each of 10 radiation fractions), WR-2721 provided no protection against acute skin reactions, little to no protection against late complications, and there was a suggestion that the drug protocol provided protection of the tumors at the low end of the radiation dose range. A contemporary example of how a similar study design can be deployed is provided by ongoing research by two authors of this review; both Nolan and Boss are involved in an NIH-sponsored bi-institutional canine clinical trial (5R01CA232148-02) which is currently enrolling 104 dogs with spontaneously occurring soft tissue sarcomas (over a 5 year period) to determine the safety of, and radiation enhancement provided by ultrasound-guided oxygen release from microbubbles. Conduct of these large-scale studies has historically been overseen and facilitated by clinical study coordinators and study teams at the local institutions. Today, large-scale multi-center trials can be coordinated by the Comparative Oncology Program (COP),

which is a core resource of the Center for Cancer Research at the National Cancer Institute. The COP centrally manages a network of 20 academic comparative oncology centers; these centers comprise the Comparative Oncology Trials Consortium (COTC). Similarly, and with support from the V-Foundation, a second trials network is currently being planned. As envisioned, that group, called the Canine Oncology Research Consortium (CORC), will include multiple academic partners, with each partner defined as an academic veterinary center paired with an NCI-designated Comprehensive Cancer Center. Both the COP/COTC and CORC exemplify the expanded support that is now available for efficient conduct of canine comparative oncology clinical trials.

Another contemporary example of how normal tissue toxicity data can be directly gathered from pet dogs is provided by an author of this review; while not yet published, Nolan et al., have presented an abstract which describes how pain can be measured and modeled in dogs in pet dogs that develop acute radiodermatitis while undergoing post-operative RT for incompletely excised extremity soft tissue sarcomas (39). Their work not only indicated that mechanical quantitative sensory testing can be used as a reliable tool for preclinical evaluation of novel analgesic strategies, but also provides evidence that localized RT-induced pain is accompanied by widespread somatosensory sensitization. That set of experiments provides a template for how pet dog studies might fit into the traditional paradigm of therapeutic development: (1) hypothesis generating observations can first be made in people or dogs undergoing cancer therapy; (2) mechanistic studies can then be performed in more conventional preclinical models; (3) novel therapies that arise from the preclinical work can then be efficiently testing in dogs, to provide proof-of-concept for safety and clinical efficacy, before advancing to early phase human clinical trials.

## Tumor Microenvironment

Tumor hypoxia is associated with relative radioresistance and aggressive biological behavior. Canine comparative oncology research has contributed meaningfully to the advancement of radiobiology research through tumor oxygenation studies. To validate canine cancer as a translationally relevant model of tumor hypoxia, Cline et al. detected the *in vivo* binding of a 2-nitroimidazole hypoxia (CCI-103F) marker in histochemical sections of canine tumors (40). The binding pattern was consistent with the expected location of hypoxic cells in tissues for which oxygen concentration gradients have been established by diffusion. The hypoxic fractions appeared in regions adjacent to necrosis, but also in regions free of necrosis. In addition to interest in hypoxic cells, populations of both non-cycling quiescent cells and rapidly-cycling proliferating cells can also influence tumor radioresponses. Zeman et al. investigated the relationships between hypoxia and proliferative status semi-quantitatively via immunohistochemical analysis of CCI-103F and proliferating cell nuclear antigen (PCNA), respectively, in canine tumor samples (41). Tumors with both high and low hypoxic and proliferative area fractions were identified; the hypoxic and proliferative cell populations overlapped to varying extents.

Direct, real-time quantification of tissue oxygenation was enabled by emergence of the Eppendorf method of direct oxygen partial pressure measurements. This technique, which involves intratumoral placement of polarographic oxygen needle electrodes, opened the door for comparative veterinary trials characterizing the tumor microenvironmental effects of hypoxia in spontaneous canine tumors; it also allowed trials designed to investigate the impact of tumor oxygenation on treatment outcomes. Achermann et al. evaluated the oxygenation of canine soft tissue sarcomas via the Eppendorf method and determined that 44% of tumors had oxygenation measurements consistent with hypoxia (42). Soon after, trials were performed in dogs undergoing fractionated RT. Polarographic needle electrodes and OxyLite fluorescence probes were used to document the presence and changes of hypoxia during fractionated RT; 58% of the dog tumors in one study were hypoxic prior to treatment (43). The  $pO_2$  of initially hypoxic tumors remained unchanged during fractionated RT, whereas the  $pO_2$  decreased in initially normoxic tumors. Brurberg et al. evaluated  $pO_2$  fluctuations in spontaneous canine tumors prior to and during RT (44). It was found that overall oxygenation status differed substantially among the tumors, and RT had no consistent effect on overall oxygenation status. Fluctuations in  $pO_2$  were detected in both unirradiated and irradiated tumors, and those fluctuations were independent of the baseline tumor oxygenation status. This study was important as it demonstrated for the first time in canine cancer the dynamic changes in tumor oxygenation in spontaneous tumors over an extended time period. The influence of tumor oxygenation status on the response to RT was first described for spontaneous canine tumors by Bley et al. (45). Pretreatment oxygen level measurements in spontaneous canine tumors were correlated with local tumor response after RT; after curative-intent full-course irradiation, hypoxic tumors had a significantly shorter median progression-free interval and a shorter overall survival time compared to better oxygenated tumors.

Comparative canine oncology trials were instrumental to understanding how hyperthermia can be combined with RT to improve tumor control. A number of positive randomized studies in dogs provided initial evidence supporting the therapeutic benefit of such combinatorial therapy (46–48). In canine soft tissue sarcomas (STS), Vujaskovic et al. identified changes in tumor oxygenation, extracellular pH, and blood flow after hyperthermia (49). They also found that hyperthermia has biphasic effects on tumor physiologic parameters: lower temperatures tend to favor improved perfusion and oxygenation, whereas higher temperatures are more likely to cause vascular damage, leading to greater hypoxia.

## EMERGING USES OF DOGS IN TRANSLATIONAL RADIATION RESEARCH

### Imaging/Theranostics

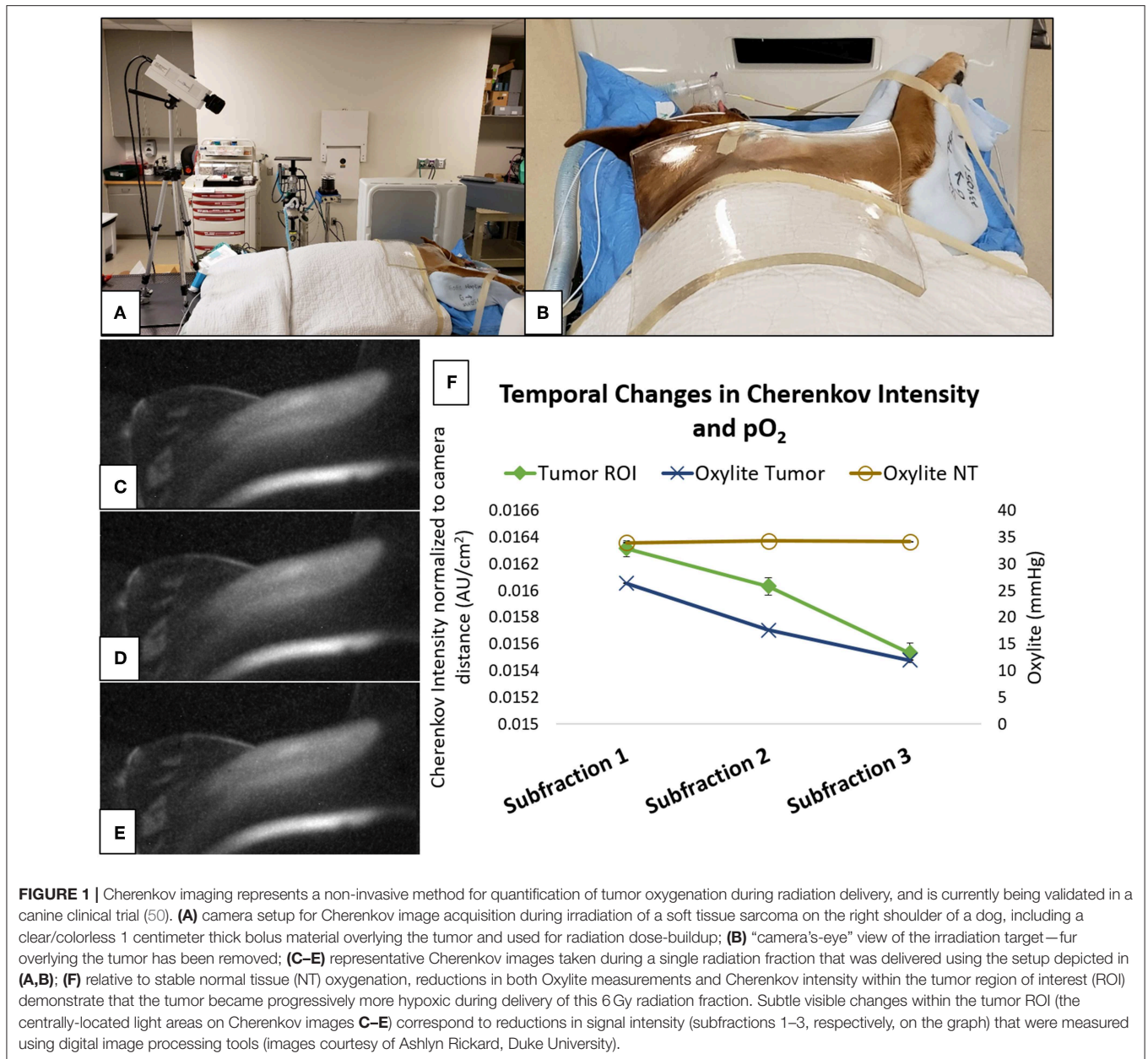
Canine comparative oncology studies that incorporate functional imaging technologies have been used to characterize the tumor microenvironment, improve target delineation, optimize

biological dose delivery, and correlate imaging characteristics with clinical outcomes. Building upon the early oxygenation and radioresponse research which relied on tissue sampling or direct insertion of electrodes for measurements, functional imaging studies provide opportunities for serial, non-invasive, quantitative or semi-quantitative analyses of the tumor microenvironment without tissue disruption (**Figure 1**).

Various positron emission tomography (PET) radiotracers have been used in comparative oncology studies to characterize the tumor microenvironment. The glucose analog 2-deoxy-2- $[^{18}F]$ -Fluoro-D-glucose (FDG), a marker of glucose uptake, is the most commonly used PET tracer in clinical oncology, and canine cancer patients were among the earliest to be imaged with FDG PET (51). FDG PET/CT has become increasingly available in veterinary medicine (52), and it has been used to characterize tumor biology and treatment responses (53–57).

As tumor hypoxia is associated with both radioresistance and tumor aggressiveness, PET-based approaches have been developed for measuring tumor hypoxia (**Figure 2**). Bruehlmeier et al. were the first to examine tumor hypoxia in canine STS using  $[^{18}F]$ -fluoromisonidazole ( $[^{18}F]$ -FMISO); FMISO tumor oxygenation measurements correlated well with Eppendorf electrode measurements (58). However, when evaluating tumor hypoxia via  $[^{18}F]$ -FMISO in cats with fibrosarcomas, the polarographic  $pO_2$  measurements did not confirm PET results; this lack of concordance was attributed to extensive tumor necrosis, and heterogeneous patterns of hypoxia (59). An alternative hypoxia imaging tracer is  $^{64}Cu$ -ATSM. Hansen et al. performed a study to compare uptake characteristics of pimonidazole immunohistochemistry (IHC) to  $^{64}Cu$ -ATSM autoradiography, and to PET uptake levels of  $^{64}Cu$ -ATSM and  $[^{18}F]$ -FDG in spontaneous canine sarcomas and carcinomas (60). Tumors with high levels of pimonidazole staining displayed high uptake of  $[^{18}F]$ -FDG and  $^{64}Cu$ -ATSM; the regional distribution of  $^{64}Cu$ -ATSM and pimonidazole correlated with each other in heterogeneous tumor regions. The potential of using  $^{64}Cu$ -ATSM to characterize the tumor microenvironment longitudinally over time was evaluated in canine tumors by measuring tumor uptake and distribution characteristics between consecutive PET scans (61). In this study,  $^{64}Cu$ -ATSM uptake was also compared to uptake and spatial distributions of  $[^{18}F]$ -FDG and dynamic contrast enhanced perfusion CT perfusion maps.  $^{64}Cu$ -ATSM uptake was positively correlated to FDG, signal was relatively stable between PET scans, and temporal changes were observed in hypo-perfused regions.  $^{64}Cu$ -ATSM PET/CT scan has also been used to detect hypoxia in feline head and neck squamous cell carcinoma (HNSCC), with PET/CT results verified by pimonidazole IHC and  $O_2$  detection probes (62).

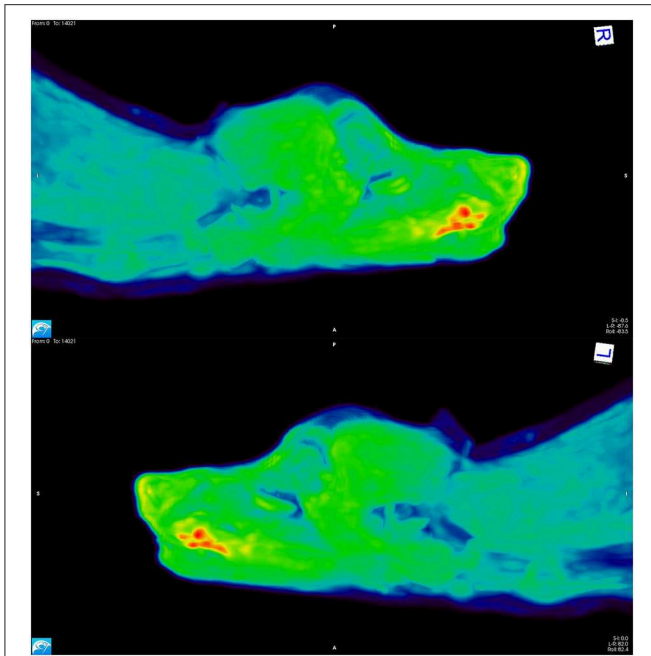
Comparative canine oncology trials have supported investigations into the kinetics of established and novel PET/CT approaches, as well as characterization of the spatial distribution of these biological markers (63–65). Bradshaw et al. concurrently evaluated the predictive value of numerous quantitative imaging biomarkers derived from multitracer PET imaging in tumors before and during RT in dogs with sinonasal tumors (66). The strongest predictors of poor outcome were derived from fluorothymidine (FLT) imaging, a marker of proliferation. The



combination of high mid-treatment standardized uptake values ( $SUV_{max}$ ) and large decreases in FLT signal from pretreatment to mid-treatment was associated with worse clinical outcome. In this study, neither FDG PET nor Cu-ATSM PET were predictive of outcome. PET/CT has also been utilized in comparative oncology studies to investigate the potential for radiation “dose painting,” which aims to improve therapeutic outcomes by increasing radiation dose in tumor regions that are identified as being at risk of relative radioresistance based upon imaging features (67, 68). Early results have been mixed.

Dynamic contrast-enhanced magnetic resonance imaging (DCE-MRI) can be used to assess tumor physiology by exploiting abnormal tumor microvasculature (Figure 3). This

enables quantitative assessment of tissue vessel density, integrity, and permeability (69). In a canine STS study, DCE-MRI was performed before and following the first hyperthermia treatment, and parameters associated with increased tumor perfusion were predictive for overall and metastasis-free survival (70). This was the first time that DCE-MRI was shown to be predictive of clinical outcome for STS. A subsequent study performed an integrative analysis of gene expression and diffusion weighted imaging (DWI) parameters, pre- and post-treatment, in dogs with STS treated with thermoradiotherapy. DWI is an MRI-based technique which quantifies the diffusion of water to characterize tumor tissue (71). Significant correlations were identified between gene expression and DWI. An unsupervised



**FIGURE 2** | [ $^{18}\text{F}$ ]-FAZA PET scan images from a dog with a squamous cell carcinoma on the ventral aspect of the tongue. The areas depicted in red have high uptake of the tracer, indicating hypoxia; the yellow and green regions have intermediate uptake, and the blue areas have low tracer uptake, thus indicating the presence of well-oxygenated tissue.

analysis of the gene sets revealed two clusters: (1) tumors with an increase in tissue water content (corresponding to an increased ADC on the DWI) after treatment showed increases in genes associated with tissue remodeling (e.g., IL1 $\beta$ , IL6, IL8, IL10) and inflammation (e.g., MMP1, TGF $\beta$ ); (2) tumors with less change in the ADC had more signs of more mature vasculature (i.e., higher gene expression of CD31 and vWF). These observations demonstrated how one can link changes in tumor physiology to changes in gene expression. This work demonstrates how early changes in functional imaging parameters might be used to aid in prognostication. Furthermore, the authors provided a blueprint for how such data can be manipulated to identify potential new drug targets.

## DNA Damage Responses

The DNA damage response (DDR) is a complex network of pathways that responds to both endogenous and exogenous DNA damage. DDR deficiencies can be targeted for cancer therapy. For example, the DDR can be inhibited as a radiosensitization strategy. Additionally, because DNA damage drives chronic inflammation and various molecular and cellular pathways of the DDR activate immune signaling (72) and since checkpoint inhibitors may work best in the setting of tumors with high mutation load, DDR inhibition during RT can generate and preserve treatment-induced DNA damage which sensitizes the tumor to subsequent checkpoint blockade (73). There are many DDR components that can be inhibited to sensitize cancer cells to

DNA damaging agents, including PARP, DNA-PK $_{cs}$ , ATM, ATR, Chk1, and Chk2.

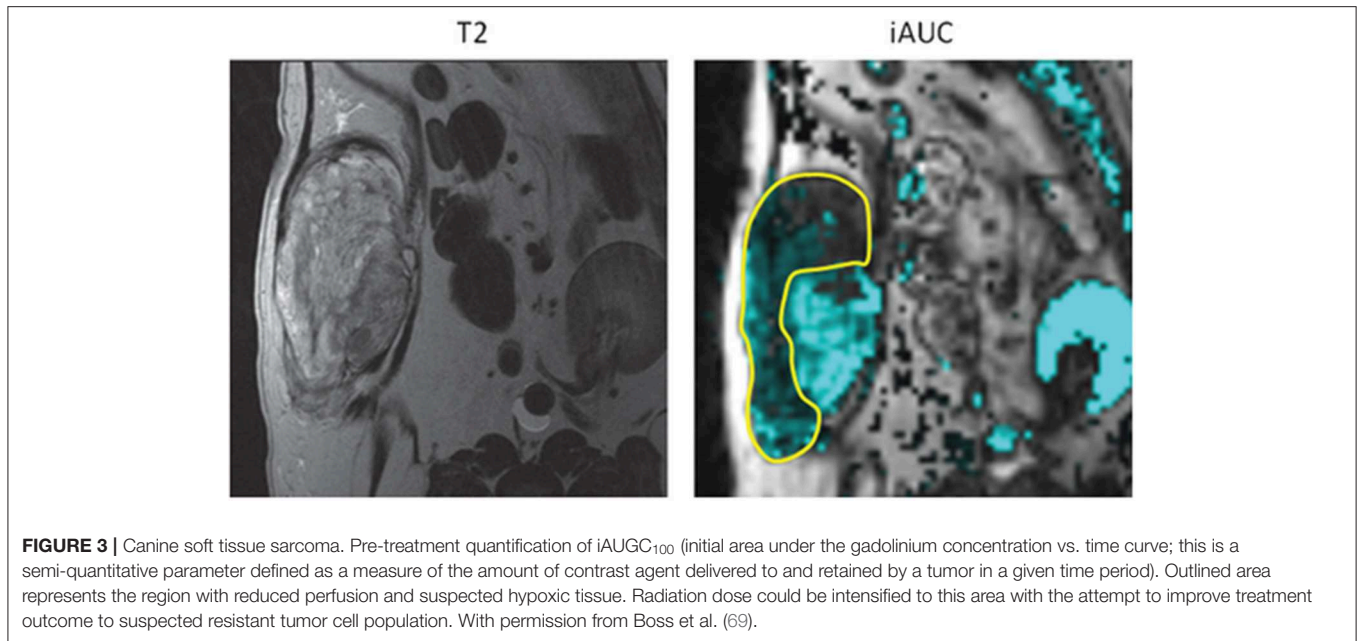
Many novel agents that show great promise in mice ultimately fail to prove efficacious in humans. Despite improved rodent models, methodologies, and study designs, many factors still contribute to these failures (74). With specific regard to DDR modifiers, a recent review suggested that better predictive biomarkers are needed to identify patients that would benefit from treatment, and that better therapeutic response biomarkers are also needed to quantify the pharmacodynamic impact and clinical gains that are achieved in humans (75). Successful and efficient translation of DDR inhibitors will rely upon preclinical studies that are designed to evaluate appropriate endpoints that can also be measured in human trials; when possible, they should also determine whether synthetic lethality contributes to efficacy of the drug as a chemo-radiosensitizer.

Dogs have previously been proposed as a model for studying the DDR (76). Indeed, the intrinsic radiosensitivity of various canine tumors and tumors cell lines is increasingly well-understood (77, 78), and methods for measuring the canine DDR have also been developed, including validation of the comet assay and immunohistochemistry for phosphorylated H2AX (79).

Canine extremity osteosarcoma (OS) may be particularly useful in the development of novel DDR inhibitors that can be combined with RT—ATR, ATM, and DNA-PK inhibitors in particular. As with pediatric OS, tumor resection plus chemotherapy is the standard treatment for canine OS. However, because RT can also provide significant analgesia and local tumor control, and because tumor necrosis is a validated surrogate for local control of canine OS, an intriguing study design would be to treat OS-bearing dogs with chemoRT, with or without a novel DDR inhibitor, and subsequently pursue tumor removal to provide standard-of-care treatment to the dog, and to provide investigators access to the resected tumor, which would thus allow robust measurement of treatment effects. Furthermore, approximately half of canine OS tumors have aberrant p53 function, making it possible to compare the effects of chemo-radiosensitizing effects of ATR or DNA-PK inhibition in normal tissues and tumors that may or may not benefit from synthetic lethality (80, 81). Thus, by studying canine OS, investigators would be able to learn about both mechanisms of interaction and clinically relevant biomarkers that directly translate to a variety of human cancers in a manner that is essentially agnostic of tumor histology.

## Immuno-Radiotherapy

Advanced metastatic disease is the most common cause of death in human patients and is a significant cause of death in dogs as well (82). Cancer immunotherapy is not a new concept, but interest re-emerged when in 2010 a phase 3 clinical trial in people with metastatic melanoma showed a survival advantage for those treated with ipilimumab, a monoclonal antibody that targets CTLA-4 (83). Since then, the use of immunotherapy in human clinical trials has expanded rapidly, with some amazing successes. The problem remains that the majority of patients do not respond favorably; furthermore, unacceptable and sometimes fatal toxicities have occurred (84, 85).



The idea that RT could induce a tumor response distant from the irradiated field has been around since the 1950s when the so-called “abscopal effect” was first proposed (86). In the context of RT, the abscopal effect refers to regression of metastatic lesions that are distant from the primary tumor and irradiated field. These are rare events; one study found 46 cases reported in the literature between 1969 and 2014 (87). That the abscopal effect occurs secondary to an immune response was first demonstrated in a 2004 mouse study which characterized radiation-induced abscopal effects as being a T-cell dependent event (88). Since that time it has become obvious that the mechanism is actually more complex, and involves a multi-faceted immune response (89).

While local irradiation can result in immunogenic cell kill by releasing neoantigens and creating local inflammation, irradiation can also attract immunosuppressive cells into the tumor microenvironment. This includes myeloid-derived suppressor cells, M2 tumor-associated macrophages and T regulatory cells; this cellular response is associated with release of cytokines (e.g., TGF- $\beta$  and IL-10) which causes local immunosuppression (90). Given the clinical rarity of measurably and clinically beneficial RT-induced immune responses, and because of the potential for post-irradiation immunosuppression, it is logical that patients may benefit from a combination of RT plus immunotherapeutics that can more reliably induce beneficial systemic immune responses (91).

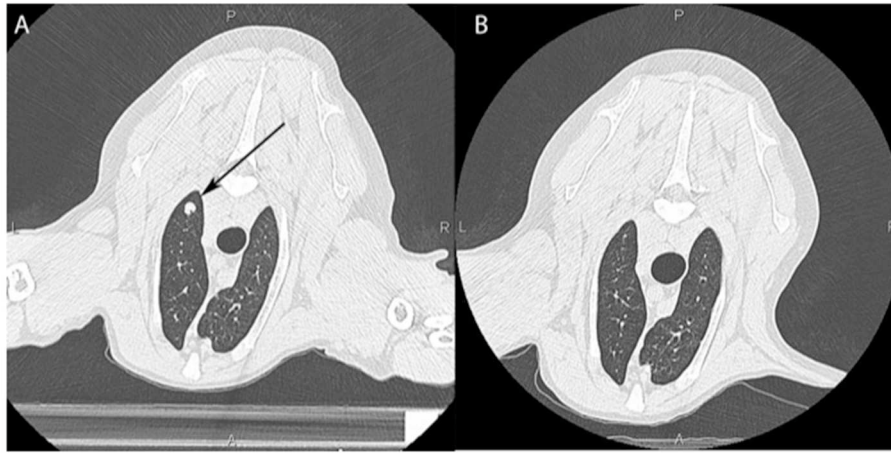
The dog immune system has been fairly well-characterized and shows great homology to humans (92). Because of these similarities, pet dogs with cancer can provide a useful model when looking to translate potential new radioimmunotherapies from mouse studies to human clinical trials.

Several small clinical trials using radio-immunotherapy protocols in dogs have been published. One recent example involved testing a novel immunotherapy combination in

dogs with metastatic melanomas and sarcomas. Immediately after each fraction of primary tumor irradiation, intratumoral injections of canine CpG oligodeoxynucleotides [CpG ODNs; immune stimulatory toll-like receptor 9 (TLR9) agonists] were done; dogs were also orally dosed with 1-methyl-tryptophan [an indolamine-2,3 dioxygenase (IDO) inhibitor] (93). The idea was that localized tumor irradiation would induce immunogenic cell death, the CpG ODNs would stimulate an immune response, and the IDO checkpoint inhibitor would counter tumor-induced immunosuppression. The dog trial was paired with mouse studies which revealed a rebound immunosuppression after mice were treated with radiation or CpG ODNs alone. While all dogs showed a local response to irradiation, there were also abscopal responses in their metastatic lesions with one dog having a complete response, two having partial responses, one with stable disease and one with progressive disease (**Figure 4**). There was no toxicity beyond what was expected for a palliative course of localized RT. Interestingly, both circulating and tumoral T regulatory cells were decreased in the dogs who responded and increased in the dog with progressive disease—suggesting this as a potential biomarker. Due to the promising preliminary results and the lack of toxicity in dogs a clinical trial using this same strategy has now opened for people with advanced metastatic cancers (94).

In another study using an adoptive immunotherapy approach combined with RT, dog natural killer cells were isolated from peripheral blood (95). Those cells could be expanded and activated, and NK cells were capable of killing osteosarcoma tumor cells *in vitro*. Furthermore, cytotoxicity was improved when tumor cells were pretreated with ionizing radiation. This was repeated *in vivo* using canine patient-derived xenograft (PDX) tumors; adoptively-transferred canine NK cells delayed the growth of tumors in mice, and focal irradiation increased





**FIGURE 4 |** Computed tomographic image of a dog's lungs showing a metastatic lesion (arrow) taken prior to local irradiation of an oral tumor (**A**) and 3 months post treatment (**B**) showing an abscopal effect with near complete resolution of the mass.

NK cell homing to these sarcoma xenografts. As part of the same study, a proof of concept clinical trial was carried out in 10 dogs with osteosarcoma. Treatment consisted of a course of palliative-intent RT followed by two intra-lesional injections of autologous activated canine NK cells. The NK cells were isolated, expanded, and activated *ex vivo* and supplemented with rhIL-2. The study demonstrated that NK cells persisted at the local tumor site for at least 1 week after injection. There was acceptable toxicity and in one of the cases there was resolution of a suspected metastatic lung nodule suggesting a possible abscopal effect.

Despite these early successes, there are several challenges in carrying out immune-RT trials in dogs, including: (1) limited availability of validated reagents, canine-specific monoclonal antibodies, and canine interleukins; and (2) limited access to properly staffed and equipped veterinary radiation oncology and RT centers (92). Work is ongoing to better characterize the canine immune system, and while many canine-specific antibodies are still not commercially available, this is improving. The majority of available checkpoint inhibitors are humanized monoclonal antibodies for which commercially available caninized versions do not exist. Furthermore, while it is possible to use recombinant human interleukins in dogs (they have been shown to be effective), there is concern that with prolonged use dogs could develop neutralizing antibodies and/or adverse immune reactions to the human-derived products. Thus, it will be difficult to fully realize the potential of this model until more immunotherapies become available for safe use in dogs.

## FUTURE DIRECTIONS

The existing experimental animal models (e.g., rodents, zebrafish) are, and will remain, vitally important for *in vivo* radiation oncology research. Yet, there are opportunities in using pet dogs that have not been fully explored; pet dogs with spontaneously arising tumors can better inform the use of RT for clinical management of various human cancers. All of the topics

discussed above will continue to spawn new research. Some of the additional areas where companion animal studies could make an impact in human medicine include radiobiology studies looking at different dosing and fractionation schemes, *in vivo* dosimetry, and testing of newly developed radiation sensitizing and radiation mitigating agents.

Small animal irradiators used in rodent research are increasingly sophisticated, but remain unable to recapitulate medical linear accelerators in terms of beam energy and dose rate (96, 97). Because companion animal oncology patients are routinely treated using standard clinical linear accelerators, they are also readily available for studies that seek to better understand the radiobiological effects of such factors as spatial fractionation (e.g., GRID and Lattice RT) and dose rate (e.g., FLASH-RT) (98, 99). For example, canine tumors are particularly well-suited to understanding the underlying biology of FLASH-RT. This is because the apparent selective normal tissue sparing (vs. tumor sparing) achieved by FLASH-RT is likely dependent upon  $pO_2$ ; and as discussed above, because the size and growth rate of canine tumors is more similar to human cancers than experimentally-induced tumors in rodents, the oxygen (hypoxia) profiles of canine tumors also tend to be similar to those of human tumors. FLASH-RT also provides a particularly good example of how studies in companion animal species can be used as an intermediate step in scaling technologies from geometries that work for rodents, to those that would work well in the context of human tumors (100). In much the same way that treatment of canine intranasal tumors was used in the early development of helical Tomotherapy, these types of companion animal studies provide an excellent vehicle for testing the safety of, and establishing feasible clinical workflows for, novel treatment devices and approaches (101).

One area where dog studies may be particularly useful is understanding the interplay between radiation dose and immune responses. Some *in vivo* work suggests that the ideal fractional

dose for radioimmunotherapy may be 6–8 Gy per fraction, rather than the more extreme doses that are commonly used in modern SRS/SBRT (89, 102). Interestingly, a recent study showed an inverse relationship between radiation dose and survival in humans having undergone RT for stage III non-small cell lung cancer; worsening outcomes with increasing radiation dose was counterintuitive but may have been attributable to delivery of higher radiation doses to tumor-infiltrating immune cells. Hypofractionation is commonly used in veterinary medicine both for palliation, and in SRS/SBRT; this practice pattern makes it relatively straightforward to interpose scientific research into clinical practice in order to use dogs as a model for studying the biology of hypofractionation.

Research in the field of therapeutic radiation physics could also benefit from a canine comparative radiation oncology approach. Because of similarities between human and canine anatomy and treatment approach, dogs can be a valuable model for validation of new techniques and methods; this use of canine comparative radiation oncology research has been exemplified for *in vivo* dosimetry (103).

There are also several limitations of the canine “model” that must be addressed. First, access to clinical outcomes data, and canine cancer tissues is limited; at the present time, there are no canine cancer registries in the United States, there are few well-validated canine cancer cell lines, and there are very few well-curated canine cancer tissue banks exist. Second, while dogs and their cancers more closely recapitulate the geometry of human cancers than do tumors of rodents, it should be noted that many pet dogs weigh far less than half the average human adult. Third, while there are many similarities between various human and canine malignancies, important differences also exist. For example, while human prostate cancer most often arises from the glandular acini, carcinomas of the canine prostate seem to most often be of urothelial origin. And although IDH1 and

IDH2 mutations are frequent in human gliomas, they are not a consistent or common feature of the canine condition (104). The value of any model system is maximized by understanding its strengths and limitations; thus, moving forward, it will be of utmost importance to focus significant energy on describing the biology of canine cancers, and making rigorous comparisons with the analogous human conditions.

Though the field of canine comparative radiation oncology research is still in its infancy, canine studies have already helped advance the understanding of human tumor biology and treatment. Through continued efforts to improve our understanding of canine tumor biology, and careful application of the canine comparative oncology model, we expect that man’s best friend will be key to reducing the global burden of cancer and improving cancer care.

## AUTHOR CONTRIBUTIONS

MN, MK, and M-KB each contributed to the drafting and editing of this manuscript, and each also approved the final version.

## FUNDING

This work was supported in part by NIH R01 CA204723-03, and a seed grant from the Consortium for Canine Comparative Oncology (a joint initiative of the NC State University College of Veterinary Medicine, and Duke University’s Duke Cancer Institute).

## ACKNOWLEDGMENTS

The authors thank Ms. Ashlyn Rickard (Duke University) for her assistance with figure preparation.

## REFERENCES

- Farrelly J, McEntee MC. A survey of veterinary radiation facilities in 2010. *Vet Radiol Ultrasound*. (2014) 55:638–43. doi: 10.1111/vru.12161
- McEntee MC. A survey of veterinary radiation facilities in the United States during 2001. *Vet Radiol Ultrasound*. (2004) 45:476–9. doi: 10.1111/j.1740-8261.2004.04082.x
- Angstadt AY, Motsinger-Reif A, Thomas R, Kisseberth WC, Guillermo Couto C, Duval DL, et al. Characterization of canine osteosarcoma by array comparative genomic hybridization and RT-qPCR: signatures of genomic imbalance in canine osteosarcoma parallel the human counterpart. *Genes Chromosomes Cancer*. (2011) 50:859–74. doi: 10.1002/gcc.20908
- Angstadt AY, Thayanithy V, Subramanian S, Modiano JF, Breen M. A genome-wide approach to comparative oncology: high-resolution oligonucleotide aCGH of canine and human osteosarcoma pinpoints shared microalterations. *Cancer Genet*. (2012) 205:572–87. doi: 10.1016/j.cancergen.2012.09.005
- Sakthikumar S, Elvers I, Kim J, Arendt ML, Thomas R, Turner-Maier J, et al. SETD2 is recurrently mutated in whole-exome sequenced canine osteosarcoma. *Cancer Res*. (2018) 78:3421–31. doi: 10.1158/0008-5472.CAN-17-3558
- Connolly NP, Shetty AC, Stokum JA, Hoeschele I, Siegel MB, Miller CR, et al. Cross-species transcriptional analysis reveals conserved and host-specific neoplastic processes in mammalian glioma. *Sci Rep*. (2018) 8:1180. doi: 10.1038/s41598-018-19451-6
- Koehler JW, Miller AD, Miller CR, Porter B, Aldape K, Beck J, et al. A revised diagnostic classification of canine glioma: towards validation of the canine glioma patient as a naturally occurring preclinical model for human glioma. *J Neuroopathol Exp Neurol*. (2018) 77:1039–54. doi: 10.1093/jnen/nly085
- Warren SL, Whipple GH. Roentgen ray intoxication: I. Unit dose over thorax negative over abdomen lethal Epithelium of small intestine sensitive to X-rays. *J Exp Med*. (1922) 35:187–202. doi: 10.1084/jem.35.2.187
- Trum BF, Shively JN, Kuhn USG, Carll WT. Radiation injury and recovery in swine. *Radiat Res*. (1959) 11:326–42. doi: 10.2307/3570681
- Brown DG, Thomas RE, Jones LP, Cross FH, Sasmore DP. Lethal dose studies with cattle exposed to whole-body Co<sup>60</sup> gamma radiation. *Radiat Res*. (1961) 15:675–83. doi: 10.2307/3571149
- Shechmeister IL, Watson LJ, Cole VW, Jackson LL. The effect of x-irradiation on goldfish. I The effect of x-irradiation on survival and susceptibility of the goldfish, *Carassius auratus*, to infection by *Aeromonas salmonicida* and *Gyrodactylus* spp. *Radiat Res*. (1962) 16:89–97. doi: 10.2307/3571133
- Gentilini F, Turba ME, Forni M, Cinotti S. Complete sequencing of full-length canine ataxia telangiectasia mutated mRNA and characterization of its putative promoter. *Vet Immunol Immunopathol*. (2009) 128:437–40. doi: 10.1016/j.vetimm.2008.12.006
- Koike M, Yutoku Y, Koike A. Cloning, localization and focus formation at DNA damage sites of canine Ku70. *J Vet Med Sci*. (2017) 79:554–61. doi: 10.1292/jvms.16-0649

14. Koike M, Yutoku Y, Koike A. Cloning, localization and focus formation at DNA damage sites of canine XLF. *J Vet Med Sci.* (2017) 79:22–8. doi: 10.1292/jvms.16-0440
15. Poulson JM, Vujaskovic Z, Gillette SM, Chaney EL, Gillette EL. Volume and dose-response effects for severe symptomatic pneumonitis after fractionated irradiation of canine lung. *Int J Radiat Biol.* (2000) 76:463–8. doi: 10.1080/095530000138457
16. Emami B, Lyman J, Brown A, Coia L, Goitein M, Munzenrider JE, et al. Tolerance of normal tissue to therapeutic irradiation. *Int J Radiat Oncol Biol Phys.* (1991) 21:109–22. doi: 10.1016/0360-3016(91)90011-R
17. Kinsella TJ, DeLuca AM, Barnes M, Anderson W, Terrill R, Sindelar WF. Threshold dose for peripheral neuropathy following intraoperative radiotherapy (IORT) in a large animal model. *Int J Radiat Oncol Biol Phys.* (1991) 20:697–701. doi: 10.1016/0360-3016(91)90011-R
18. Johnstone PAS, Deluca AM, Bacher JD, Hampshire VA, Terrill RE, Anderson WJ, et al. Clinical toxicity of peripheral nerve to intraoperative radiotherapy in a canine model. *Int J Radiat Oncol Biol Phys.* (1995) 32:1031–4. doi: 10.1016/0360-3016(95)00028-W
19. LeCouteur RA, Gillette EL, Powers BE, Child G, McChesney SL, Ingram JT. Peripheral neuropathies following experimental intraoperative radiation therapy (IORT). *Int J Radiat Oncol Biol Phys.* (1989) 17:583–90. doi: 10.1016/0360-3016(89)90110-7
20. Nolan MW, Marolf AJ, Ehrhart EJ, Rao S, Kraft SL, Engel S, et al. Pudendal nerve and internal pudendal artery damage may contribute to radiation-induced erectile dysfunction. *Int J Radiat Oncol Biol Phys.* (2015) 91:796–806. doi: 10.1016/j.ijrobp.2014.12.025
21. King CR, Brooks JD, Gill H, Presti JC Jr. Long-term outcomes from a prospective trial of stereotactic body radiotherapy for low-risk prostate cancer. *Int J Radiat Oncol Biol Phys.* (2012) 82:877–82. doi: 10.1016/j.ijrobp.2010.11.054
22. Kim DW, Cho LC, Straka C, Christie A, Lotan Y, Pistenmaa D, et al. Predictors of rectal tolerance observed in a dose-escalated phase 1–2 trial of stereotactic body radiation therapy for prostate cancer. *Int J Radiat Oncol Biol Phys.* (2014) 89:509–17. doi: 10.1016/j.ijrobp.2014.03.012
23. Storb R, Epstein RB, Ragde H, Bryant J, Thomas ED. Marrow engraftment by allogeneic leukocytes in lethally irradiated dogs. *Blood.* (1967) 30:805–11. doi: 10.1182/blood.V30.6.805.805
24. Weiden PL, Storb R, Lerner KG, Kao GF, Graham TC, Thomas ED. Treatment of canine malignancies by 1200 R total body irradiation and autologous marrow grafts. *Exp Hematol.* (1975) 3:124–34.
25. Deeg HJ, Meyers JD, Storb R, Graham TC, Weiden PL. Effect of trimethoprim-sulfamethoxazole on hematological recovery after total body irradiation and autologous marrow infusion in dogs. *Transplantation.* (1979) 28:243–6. doi: 10.1097/00007890-197909000-00017
26. Kolb H, Sale GE, Lerner KG, Storb R, Thomas ED. Pathology of acute graft-versus-host disease in the dog. An autopsy study of ninety-five dogs. *Am J Pathol.* (1979) 96:581–94.
27. Weiden PL, Storb R, Deeg HJ, Graham TC, Thomas ED. Prolonged disease-free survival in dogs with lymphoma after total-body irradiation and autologous marrow transplantation consolidation of combination-chemotherapy-induced remissions. *Blood.* (1979) 54:1039–49. doi: 10.1182/blood.V54.5.1039.1039
28. Deeg HJ, Storb R, Prentice R, Fritz TE, Weiden PL, Sale GE, et al. Increased cancer risk in canine radiation chimeras. *Blood.* (1980) 55:233–9. doi: 10.1182/blood.V55.2.233.233
29. Deeg HJ, Prentice R, Fritz TE, Sale GE, Lombard LS, Thomas ED, et al. Increased incidence of malignant tumors in dogs after total body irradiation and marrow transplantation. *Int J Radiat Oncol Biol Phys.* (1983) 9:1505–11. doi: 10.1016/0360-3016(83)90325-5
30. Shulman HM, Luk K, Deeg HJ, Shuman WB, Storb R. Induction of hepatic veno-occlusive disease in dogs. *Am J Pathol.* (1987) 126:114–25.
31. Deeg HJ, Storb R, Longton G, Graham TC, Shulman HM, Appelbaum F, et al. Single dose or fractionated total body irradiation and autologous marrow transplantation in dogs: effects of exposure rate, fraction size, and fractionation interval on acute and delayed toxicity. *Int J Radiat Oncol Biol Phys.* (1988) 15:647–53. doi: 10.1016/0360-3016(88)90307-0
32. Storb R, Raff RF, Graham T, Appelbaum FR, Deeg HJ, Schuening FG, et al. Marrow toxicity of fractionated vs. single dose total body irradiation is identical in a canine model. *Int J Radiat Oncol Biol Phys.* (1993) 26:275–83. doi: 10.1016/0360-3016(93)90207-C
33. Storb R, Raff RF, Appelbaum FR, Deeg HJ, Graham TC, Schuening FG, et al. Fractionated versus single-dose total body irradiation at low and high dose rates to condition canine littermates for DLA-identical marrow grafts. *Blood.* (1994) 83:3384–9. doi: 10.1182/blood.V83.11.3384.3384
34. Storb R, Raff R, Deeg HJ, Graham T, Appelbaum FR, Schuening FG, et al. Dose rate-dependent sparing of the gastrointestinal tract by fractionated total body irradiation in dogs given marrow autografts. *Int J Radiat Oncol Biol Phys.* (1998) 40:961–6. doi: 10.1016/S0360-3016(97)00913-9
35. Storb R, Raff RF, Graham T, Appelbaum FR, Deeg HJ, Schuening FG, et al. Dose rate-dependent marrow toxicity of TBI in dogs and marrow sparing effect at high dose rate by dose fractionation. *Biol Blood Marrow Transplant.* (1999) 5:155–61. doi: 10.1053/bbmt.1999.v5.pm10392961
36. Graves SS, Storer BE, Butts TM, Storb R. Comparing high and low total body irradiation dose rates for minimum-intensity conditioning of dogs for dog leukocyte antigen-identical bone marrow grafts. *Biol Blood Marrow Transplant.* (2013) 19:1650–4. doi: 10.1016/j.bbmt.2013.08.007
37. Gieger TL, Nolan MW, Roback DM, Suter SE. Implementation of total body photon irradiation as part of an institutional bone marrow transplant program for the treatment of canine lymphoma and leukemias. *Vet Radiol Ultrasound.* (2019) 60:586–93. doi: 10.1111/vru.12776
38. McChesney SL, Gillette EL, Dewhirst MW, Withrow SJ. Influence of WR 2721 on radiation response of canine soft tissue sarcomas. *Int J Radiat Oncol Biol Phys.* (1986) 12:1957–63. doi: 10.1016/0360-3016(86)90132-X
39. Kelsey KL, Lascelles BDX, Enomoto M, Gieger TL, Nolan MW. Widespread central sensitization is associated with acute radiation dermatitis in pet dogs undergoing radiotherapy for spontaneous tumors. In: *63rd Annual Meeting of the Radiation Research Society.* Cancun (2017).
40. Cline J, Thrall D, Page R, Franko A, Raleigh J. Immunohistochemical detection of a hypoxia marker in spontaneous canine tumours. *Br J Cancer.* (1990) 62:925. doi: 10.1038/bjc.1990.411
41. Zeman EM, Calkins DR, Cline JM, Thrall DE, Raleigh JA. The relationship between proliferative and oxygenation status in spontaneous canine tumors. *Int J Radiat Oncol Biol Phys.* (1993) 27:891–8. doi: 10.1016/0360-3016(93)90465-8
42. Achermann R, Ohlerth S, Fidel J, Gardelle O, Gassmann M, Roos M, et al. Ultrasound guided, pre-radiation oxygen measurements using polarographic oxygen needle electrodes in spontaneous canine soft tissue sarcomas. *In Vivo.* (2002) 16:431–7.
43. Achermann RE, Ohlerth SM, Bley CR, Gassmann M, Inteworn N, Roos M, et al. Oxygenation of spontaneous canine tumors during fractionated radiation therapy. *Strahlenther Onkol.* (2004) 180:297–305. doi: 10.1007/s00066-004-1193-6
44. Brurberg KG, Skogmo HK, Graff BA, Olsen DR, Rofstad EK. Fluctuations in pO<sub>2</sub> in poorly and well-oxygenated spontaneous canine tumors before and during fractionated radiation therapy. *Radiother Oncol.* (2005) 77:220–6. doi: 10.1016/j.radonc.2005.09.009
45. Bley CR, Ohlerth S, Roos M, Wergin M, Achermann R, Kaser-Hotz B. Influence of pretreatment polarographically measured oxygenation levels in spontaneous canine tumors treated with radiation therapy. *Strahlenther Onkol.* (2006) 182:518–24. doi: 10.1007/s00066-006-1519-7
46. Dewhirst MW, Sim DA, Sapareto S, Connor WG. Importance of minimum tumor temperature in determining early and long-term responses of spontaneous canine and feline tumors to heat and radiation. *Cancer Res.* (1984) 44:43–50.
47. Gillette EL, McChesney SL, Dewhirst MW, Scott RJ. Response of canine oral carcinomas to heat and radiation. *Int J Radiat Oncol Biol Phys.* (1987) 13:1861–7. doi: 10.1016/0360-3016(87)90353-1
48. Gillette SM, Dewhirst M, Gillette E, Thrall D, Page R, Powers B, et al. Response of canine soft tissue sarcomas to radiation or radiation plus hyperthermia: a randomized phase II study. *Int J Hyperthermia.* (1992) 8:309–20. doi: 10.3109/02656739209021786
49. Vujaskovic Z, Poulson JM, Gaskin AA, Thrall DE, Page RL, Charles HC, et al. Temperature-dependent changes in physiologic parameters of spontaneous canine soft tissue sarcomas after combined radiotherapy and hyperthermia treatment. *Int J Radiat Oncol Biol Phys.* (2000) 46:179–85. doi: 10.1016/S0360-3016(99)00362-4

50. Zhang X, Lam SK, Palmer G, Das S, Oldham M, Dewhirst M. Noninvasive measurement of tissue blood oxygenation with Cerenkov imaging during therapeutic radiation delivery. *Opt Lett.* (2017) 42:3101–4. doi: 10.1364/OL.42.003101
51. Paul R, Johansson R, Kiuru A, Illukka T, Talvio T, Roeda D, et al. Imaging of canine cancers with 18F-2-fluoro-2-deoxy-D-glucose (FDG) suggests further applications for cancer imaging in man. *Nucl Med Commun.* (1984) 5:641–6. doi: 10.1097/00006231-198410000-00006
52. Hansen AE, McEvoy F, Engelholm SA, Law I, Kristensen AT. FDG PET/CT imaging in canine cancer patients. *Vet Radiol Ultrasound.* (2011) 52:201–6. doi: 10.1111/j.1740-8261.2010.01757.x
53. LeBlanc AK, Miller AN, Galyon GD, Moyers TD, Long MJ, Stuckey AC, et al. Preliminary evaluation of serial 18fdg-pet/ct to assess response to toceranib phosphate therapy in canine cancer. *Vet Radiol Ultrasound.* (2012) 53:348–57. doi: 10.1111/j.1740-8261.2012.01925.x
54. Yoshikawa H, Randall EK, Kraft SL, LaRue SM. Comparison between 2-18f-fluoro-2-deoxy-d-glucose positron emission tomography and contrast-enhanced computed tomography for measuring gross tumor volume in cats with oral squamous cell carcinoma. *Vet Radiol Ultrasound.* (2013) 54:307–13. doi: 10.1111/vru.12016
55. Randall E, Kraft S, Yoshikawa H, LaRue S. Evaluation of 18F-FDG PET/CT as a diagnostic imaging and staging tool for feline oral squamous cell carcinoma. *Vet Comp Oncol.* (2016) 14:28–38. doi: 10.1111/vco.12047
56. Griffin LR, Thamm DH, Selmic LE, Ehrhart E, Randall E. Pilot study utilizing Fluorine-18 fluorodeoxyglucose-positron emission tomography/computed tomography for glycolytic phenotyping of canine mast cell tumors. *Vet Radiol Ultrasound.* (2018) 59:461–8. doi: 10.1111/vru.12612
57. Griffin LR, Thamm DH, Brody A, Selmic LE. Prognostic value of fluorine18 fluorodeoxyglucose positron emission tomography/computed tomography in dogs with appendicular osteosarcoma. *J Vet Int Med.* (2019) 33:820–6. doi: 10.1111/jvim.15453
58. Bruehlmeier M, Kaser-Hotz B, Achermann R, Bley CR, Wergin M, Schubiger PA, et al. Measurement of tumor hypoxia in spontaneous canine sarcomas. *Vet Radiol Ultrasound.* (2005) 46:348–54. doi: 10.1111/j.1740-8261.2005.00065.x
59. Allemann K, Wyss M, Wergin M, Ohlerth S, Rohrer-Bley C, Evans S, et al. Measurements of hypoxia ([18F]-FMISO,[18F]-EF5) with positron emission tomography (PET) and perfusion using PET ([15O]-H2O) and power Doppler ultrasonography in feline fibrosarcomas. *Vet Comp Oncol.* (2005) 3:211–21. doi: 10.1111/j.1476-5810.2005.00081.x
60. Hansen AE, Kristensen AT, Jørgensen JT, McEvoy FJ, Busk M, van der Kogel AJ, et al. 64 Cu-ATSM and 18 FDG PET uptake and 64 Cu-ATSM autoradiography in spontaneous canine tumors: comparison with pimonidazole hypoxia immunohistochemistry. *Radiat Oncol.* (2012) 7:89. doi: 10.1186/1748-717X-7-89
61. Hansen AE, Kristensen AT, Law I, McEvoy FJ, Kjær A, Engelholm SA. Multimodality functional imaging of spontaneous canine tumors using 64Cu-ATSM and 18FDG PET/CT and dynamic contrast enhanced perfusion CT. *Radiother Oncol.* (2012) 102:424–8. doi: 10.1016/j.radonc.2011.10.021
62. Ballegeer EA, Madrill NJ, Berger KL, Agnew DW, McNeil EA. Evaluation of hypoxia in a feline model of head and neck cancer using 64 Cu-ATSM positron emission tomography/computed tomography. *BMC Cancer.* (2013) 13:218. doi: 10.1186/1471-2407-13-218
63. Black NF, McJames S, Rust TC, Kadrmas DJ. Evaluation of rapid dual-tracer 62Cu-PTSM+ 62Cu-ATSM PET in dogs with spontaneously occurring tumors. *Phys Med Biol.* (2007) 53:217. doi: 10.1088/0031-9155/53/1/015
64. Bradshaw TJ, Bowen SR, Jallow N, Forrest LJ, Jeraj R. Heterogeneity in intratumor correlations of 18F-FDG, 18F-FLT, and 61Cu-ATSM PET in canine sinonasal tumors. *J Nucl Med.* (2013) 54:1931–7. doi: 10.2967/jnumed.113.121921
65. Simoncic U, Jeraj R. Heterogeneity in stabilization phenomena in FLT PET images of canines. *Phys Med Biol.* (2014) 59:7937. doi: 10.1088/0031-9155/59/24/7937
66. Bradshaw TJ, Bowen SR, Deveau MA, Kubicek L, White P, Bentzen SM, et al. Molecular imaging biomarkers of resistance to radiation therapy for spontaneous nasal tumors in canines. *Int J Radiat Oncol Biol Phys.* (2015) 91:787–95. doi: 10.1016/j.ijrobp.2014.12.011
67. Clausen MM, Hansen AE, Af Rosenschold PM, Kjær A, Kristensen AT, McEvoy FJ, et al. Dose escalation to high-risk sub-volumes based on non-invasive imaging of hypoxia and glycolytic activity in canine solid tumors: a feasibility study. *Radiat Oncol.* (2013) 8:262. doi: 10.1186/1748-717X-8-262
68. Bradshaw T, Fu R, Bowen S, Zhu J, Forrest L, Jeraj R. Predicting location of recurrence using FDG, FLT, and Cu-ATSM PET in canine sinonasal tumors treated with radiotherapy. *Phys Med Biol.* (2015) 60:5211. doi: 10.1088/0031-9155/60/13/5211
69. Boss MK, Muradyan N, Thrall DE. DCE-MRI: a review and applications in veterinary oncology. *Vet Comp Oncol.* (2013) 11:87–100. doi: 10.1111/j.1476-5829.2011.00305.x
70. Viglianti BL, Lora-Michiels M, Poulson JM, Lan L, Yu D, Sanders L, et al. Dynamic contrast-enhanced magnetic resonance imaging as a predictor of clinical outcome in canine spontaneous soft tissue sarcomas treated with thermoradiotherapy. *Clin Cancer Res.* (2009) 15:4993–5001. doi: 10.1158/1078-0432.CCR-08-2222
71. Chi JT, Thrall DE, Jiang C, Snyder S, Fels D, Landon C, et al. Comparison of genomics and functional imaging from canine sarcomas treated with thermoradiotherapy predicts therapeutic response and identifies combination therapeutics. *Clin Cancer Res.* (2011) 17:2549–60. doi: 10.1158/1078-0432.CCR-10-2583
72. Nakad R, Schumacher B. DNA damage response and immune defense: links and mechanisms. *Front Genet.* (2016) 7:147. doi: 10.3389/fgene.2016.00147
73. Liontos M, Anastasiou I, Bamias A, Dimopoulos M-A. DNA damage, tumor mutational load and their impact on immune responses against cancer. *Ann Transl Med.* (2016) 4:264. doi: 10.21037/atm.2016.07.11
74. Coleman CN, Higgins GS, Brown JM, Baumann M, Kirsch DG, Willers H, et al. Improving the predictive value of preclinical studies in support of radiotherapy clinical trials. *Clin Cancer Res.* (2016) 22:3138–47. doi: 10.1158/1078-0432.CCR-16-0069
75. Weber AM, Ryan AJ. ATM and ATR as therapeutic targets in cancer. *Pharmacol Ther.* (2015) 149:124–38. doi: 10.1016/j.pharmthera.2014.12.001
76. Grosse N, van Loon B, Rohrer Bley C. DNA damage response and DNA repair - dog as a model? *BMC Cancer.* (2014) 14:203. doi: 10.1186/1471-2407-14-203
77. Maeda J, Froning CE, Brents CA, Rose BJ, Thamm DH, Kato TA. Intrinsic radiosensitivity and cellular characterization of 27 canine cancer cell lines. *PLoS ONE.* (2016) 11:e156689. doi: 10.1371/journal.pone.0156689
78. Yoshikawa H, Sunada S, Hirakawa H, Fujimori A, Elmegerhi S, Leary D, et al. Radiobiological characterization of canine malignant melanoma cell lines with different types of ionizing radiation and efficacy evaluation with cytotoxic agents. *Int J Mol Sci.* (2019) 20:841. doi: 10.3390/ijms20040841
79. Schulz N, Chaachouay H, Nytko KJ, Weyland MS, Roos M, Fuchsli RM, et al. Dynamic *in vivo* profiling of DNA damage and repair after radiotherapy using canine patients as a model. *Int J Mol Sci.* (2017) 18:16. doi: 10.3390/ijms18061176
80. Loukopoulos P, Thornton JR, Robinson WF. Clinical and pathologic relevance of p53 index in canine osseous tumors. *Vet Pathol.* (2003) 40:237–48. doi: 10.1354/vp.40-3-237
81. Kirpensteijn J, Kik M, Teske E, Rutteman GR. TP53 gene mutations in canine osteosarcoma. *Vet Surg.* (2008) 37:454–60. doi: 10.1111/j.1532-950X.2008.00407.x
82. Seyfried TN, Huysentruyt LC. On the origin of cancer metastasis. *Crit Rev Oncog.* (2013) 18:43–73. doi: 10.1615/CritRevOncog.v18.i1-2.40
83. Hodi FS, O'Day SJ, McDermott DF, Weber RW, Sosman JA, Haanen JB, et al. Improved survival with ipilimumab in patients with metastatic melanoma. *N Engl J Med.* (2010) 363:711–23. doi: 10.1056/NEJMoa1003466
84. Puzanov I, Diab A, Abdallah K, Bingham CO III, Brogdon C, Dadu R, et al. Managing toxicities associated with immune checkpoint inhibitors: consensus recommendations from the Society for Immunotherapy of Cancer (SITC) Toxicity Management Working Group. *J Immunother Cancer.* (2017) 5:95. doi: 10.1186/s40425-017-0300-z
85. Ventola CL. Cancer immunotherapy, part 3: challenges and future trends. *Pharm Ther.* (2017) 42:514–21.
86. Mole RH. Whole body irradiation: radiobiology or medicine? *Br J Radiol.* (1953) 26:234–41. doi: 10.1259/0007-1285-26-305-234

87. Abuodeh Y, Venkat P, Kim S. Systematic review of case reports on the abscopal effect. *Curr Probl Cancer*. (2016) 40:25–37. doi: 10.1016/j.crrprobcancer.2015.10.001
88. Demaria S, Ng B, Devitt ML, Babb JS, Kawashima N, Liebes L, et al. Ionizing radiation inhibition of distant untreated tumors (abscopal effect) is immune mediated. *Int J Radiat Oncol Biol Phys*. (2004) 58:862–70. doi: 10.1016/j.ijrobp.2003.09.012
89. Siva S, MacManus MP, Martin RF, Martin OA. Abscopal effects of radiation therapy: a clinical review for the radiobiologist. *Cancer Lett*. (2015) 356:82–90. doi: 10.1016/j.canlet.2013.09.018
90. Shevtsov M, Sato H, Multhoff G, Shibata A. Novel approaches to improve the efficacy of immuno-radiotherapy. *Front Oncol*. (2019) 9:156. doi: 10.3389/fonc.2019.00156
91. Baird JR, Monjazeb AM, Shah O, McGee H, Murphy WJ, Crittenden MR, et al. Stimulating innate immunity to enhance radiation therapy-induced tumor control. *Int J Radiat Oncol Biol Phys*. (2017) 99:362–73. doi: 10.1016/j.ijrobp.2017.04.014
92. Park JS, Withers SS, Modiano JF, Kent MS, Chen M, Luna JI, et al. Canine cancer immunotherapy studies: linking mouse and human. *J Immunother Cancer*. (2016) 4:97. doi: 10.1186/s40425-016-0200-7
93. Monjazeb AM, Kent MS, Grossenbacher SK, Mall C, Zamora AE, Mirsoian A, et al. Blocking indolamine-2,3-dioxygenase rebound immune suppression boosts antitumor effects of radio-immunotherapy in murine models and spontaneous canine malignancies. *Clin Cancer Res*. (2016) 22:4328–40. doi: 10.1158/1078-0432.CCR-15-3026
94. Monjazeb AM. *A Phase I/II Trial of an Indolamine 2,3 Dioxygenase Blockade, Intralesional Toll-receptor 9 Agonist, and Radiotherapy in Patients with Advanced Solid Tumors and Lymphomas*. (2019). Available online at: <https://ccresources.ucdmc.ucdavis.edu/csr/clinicaltrialdisplay.csr?studynumdisplayed=UCDCC%23271&orgid=51&searchtext=-1&agegroup=-1&cancercategory=-1&piid=1551&piname=Arta%20Monjazeb&studynumber=-1&phaseid=-1> (accessed July 31, 2019).
95. Canter RJ, Grossenbacher SK, Foltz JA, Sturgill IR, Park JS, Luna JI, et al. Radiotherapy enhances natural killer cell cytotoxicity and localization in pre-clinical canine sarcomas and first-in-dog clinical trial. *J Immunother Cancer*. (2017) 5:98. doi: 10.1186/s40425-017-0305-7
96. Sharma S, Narayanasamy G, Przybyla B, Webber J, Boerma M, Clarkson R, et al. Advanced small animal conformal radiation therapy device. *Technol Cancer Res Treat*. (2017) 16:45–56. doi: 10.1177/1533034615626011
97. Verhaegen F, Dubois L, Gianolini S, Hill MA, Karger CP, Lauber K, et al. ESTRO ACROP: technology for precision small animal radiotherapy research: optimal use and challenges. *Radiother Oncol*. (2018) 126:471–8. doi: 10.1016/j.radonc.2017.11.016
98. Nolan MW, Gieger TL, Karakashian AA, Nikolova-Karakashian MN, Posner LP, Roback DM, et al. Outcomes of spatially fractionated radiotherapy (GRID) for bulky soft tissue sarcomas in a large animal model. *Technol Cancer Res Treat*. (2017) 16:357–65. doi: 10.1177/1533034617690980
99. Harrington KJ. Ultrahigh dose-rate radiotherapy: next steps for FLASH-RT. *Clin Cancer Res*. (2019) 25:3–5. doi: 10.1158/1078-0432.CCR-18-1796
100. Vozenin MC, De Fornel P, Petersson K, Favaudon V, Jaccard M, Germond JF, et al. The advantage of FLASH radiotherapy confirmed in mini-pig and cat-cancer patients. *Clin Cancer Res*. (2019) 25:35–42. doi: 10.1158/1078-0432.CCR-17-3375
101. Forrest LJ, Mackie TR, Ruchala K, Turek M, Kapatoes J, Jaradat H, et al. The utility of megavoltage computed tomography images from a helical tomotherapy system for setup verification purposes. *Int J Radiat Oncol Biol Phys*. (2004) 60:1639–44. doi: 10.1016/j.ijrobp.2004.08.016
102. Deloch L, Derer A, Hartmann J, Frey B, Fietkau R, Gaipl US. Modern radiotherapy concepts and the impact of radiation on immune activation. *Front Oncol*. (2016) 6:141. doi: 10.3389/fonc.2016.00141
103. Hsieh ES, Hansen KS, Kent MS, Saini S, Dieterich S. Can a commercially available EPID dosimetry system detect small daily patient setup errors for cranial IMRT/SRS? *Pract Radiat Oncol*. (2017) 7:e283–90. doi: 10.1016/j.prro.2016.12.005
104. Reitman ZJ, Olby NJ, Mariani CL, Thomas R, Breen M, Bigner DD, et al. IDH1 and IDH2 hotspot mutations are not found in canine glioma. *Int J Cancer*. (2010) 127:245–6. doi: 10.1002/ijc.25017

**Conflict of Interest:** The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

Copyright © 2019 Nolan, Kent and Boss. This is an open-access article distributed under the terms of the Creative Commons Attribution License (CC BY). The use, distribution or reproduction in other forums is permitted, provided the original author(s) and the copyright owner(s) are credited and that the original publication in this journal is cited, in accordance with accepted academic practice. No use, distribution or reproduction is permitted which does not comply with these terms.