

# The Advantage of FLASH Radiotherapy Confirmed in Mini-pig and Cat-cancer Patients

Marie-Catherine Vozenin<sup>1</sup>, Pauline De Fornel<sup>2</sup>, Kristoffer Petersson<sup>1,3</sup>, Vincent Favaudon<sup>4</sup>, Maud Jaccard<sup>1,3</sup>, Jean-François Germond<sup>3</sup>, Benoit Petit<sup>1</sup>, Marco Burki<sup>5</sup>, Gisèle Ferrand<sup>6</sup>, David Patin<sup>3</sup>, Hanan Bouchaab<sup>1</sup>, Mahmut Ozsahin<sup>1,6</sup>, François Bochud<sup>3</sup>, Claude Bailat<sup>3</sup>, Patrick Devauchelle<sup>2</sup>, and Jean Bourhis<sup>1,6</sup>



## Abstract

**Purpose:** Previous studies using FLASH radiotherapy (RT) in mice showed a marked increase of the differential effect between normal tissue and tumors. To stimulate clinical transfer, we evaluated whether this effect could also occur in higher mammals.

**Experimental Design:** Pig skin was used to investigate a potential difference in toxicity between irradiation delivered at an ultrahigh dose rate called "FLASH-RT" and irradiation delivered at a conventional dose rate called "Conv-RT." A clinical, phase I, single-dose escalation trial (25–41 Gy) was performed in 6 cat patients with locally advanced T2/T3N0M0 squamous cell carcinoma of the nasal planum to determine the maximal tolerated

dose and progression-free survival (PFS) of single-dose FLASH-RT.

**Results:** Using, respectively, depilation and fibronecrosis as acute and late endpoints, a protective effect of FLASH-RT was observed ( $\geq 20\%$  dose-equivalent difference vs. Conv-RT). Three cats experienced no acute toxicity, whereas 3 exhibited moderate/mild transient mucositis, and all cats had depilation. With a median follow-up of 13.5 months, the PFS at 16 months was 84%.

**Conclusions:** Our results confirmed the potential advantage of FLASH-RT and provide a strong rationale for further evaluating FLASH-RT in human patients.

*See related commentary by Harrington, p. 3*

## Introduction

Radiotherapy is an essential part of cancer treatment; more than half of patients with cancer are treated with radiotherapy, mostly with curative potential. In the last decade, major advances in high precision treatment delivery and multimodal imaging allowed radiotherapy to be better tolerated with an increased proportion of patients living longer, free of tumor, and with fewer side effects. Despite these progresses, radiation resistance remains an unmet clinical need, because an important proportion of solid tumors can be resistant to conventional radiotherapy. For patients harboring those tumors, a more effective radiotherapeutic strategy is needed. Radiation-induced side effects limit dose escalation and

compromise quality of life; therefore, a more powerful and better tolerated radiotherapy remains a major need to improve the ratio tolerance/efficacy of radiotherapy. With the objective of fostering innovation in radiotherapy, we have been the first to propose a completely novel modality of irradiation, named FLASH radiotherapy (FLASH-RT), that markedly increases the differential effect between tumors and normal tissues and is able to destroy tumors, while better protecting normal tissues and preventing side effects in experimental models (1, 2). We also developed accurate and traceable dosimetry methods to compare irradiation performed at dose rates ranging from conventional to FLASH (3, 4). Then, we started to delineate the physical parameters required for FLASH-induced protection of normal tissue and defined the threshold dose rate required to start observing some protective effect at 30 Gy/second, whereas this protective effect was more pronounced above 100 Gy/second (2).

FLASH-RT delivery uses irradiators with a high radiation output. We performed our studies using two experimental linear accelerators (linacs), specifically constructed to deliver pulsed electron beam irradiation at ultrahigh dose rates ( $\geq 100$  Gy/second), but also able to operate at conventional dose rates commonly used in radiotherapy (a few Gy/minute; ref. 5). Interestingly, the group of B. Loo at Stanford University (Palo Alto, CA) recently published a procedure on how to modify a clinical linac into an irradiator capable of delivering FLASH-RT, with dose rates exceeding 200 Gy/second (6) and produced preliminary data showing gut protection in mice (7). This does stimulate the research possibilities on FLASH-RT, and will help to investigate the radiobiological advantage of increasing the radiotherapy dose rate. An additional potential interest of these ultrahigh dose rates with treatment times generally shorter than

<sup>1</sup>Radio-Oncology Laboratory, Department of Radiation Oncology, Lausanne University Hospital, Lausanne, Switzerland. <sup>2</sup>Micen-vet, Créteil, France. <sup>3</sup>Institute of Radiation Physics (IRA), Lausanne University Hospital, Lausanne, Switzerland. <sup>4</sup>Institut Curie, INSERM U1021/CNRS UMR3347, Université Paris-Sud, Orsay, France. <sup>5</sup>Service de Chirurgie Expérimentale, Département de chirurgie et anesthésie, Lausanne University Hospital, Lausanne, Switzerland. <sup>6</sup>UNIL, Lausanne, Switzerland.

**Note:** Supplementary data for this article are available at Clinical Cancer Research Online (<http://clincancerres.aacrjournals.org/>).

M.-C. Vozenin, P. De Fornel, K. Petersson, P. Devauchelle, and J. Bourhis contributed equally to this article.

**Corresponding Author:** Marie-Catherine Vozenin, Laboratoire de Radio-Oncologie, Centre Hospitalier Universitaire Vaudois, Bugnon 46, Lausanne 1011, Switzerland. Phone: 417-9199-0204; Fax: 412-1692-5901; E-mail: marie-catherine.vozenin@chuv.ch

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0.1 second is the possibility to minimize treatment delivery uncertainties caused by intra-fraction motion.

Given both its potential radiobiological and delivery advantages, FLASH-RT could revolutionize the field of radiotherapy. To advance the clinical transfer of the modality, it was essential to evaluate whether the increased differential effect of FLASH-RT could also be observed in higher mammals. Consequently, we compared the effect of FLASH-RT with radiotherapy at conventional dose rate (Conv-RT) on the skin of a mini pig, which represents a well characterized radiobiological model, mimicking the reactions of human skin to radiotherapy and used high dose and single fraction as described previously (8–11). This experience confirmed our results obtained in mice, showing an important protective effect of FLASH-RT (at least 20% dose difference), compared with Conv-RT. These data prompted us to perform a clinical trial in domestic animals with spontaneous cancer. The model of locally advanced squamous cell carcinoma (SCC) of the nasal planum of cats was selected because the tumors are generally treated with radiotherapy but with a relatively poor tumor control rate (12). Surgery is mostly restricted to early-stage tumors (Tis and T1) because of the poor cosmetic and functional outcome in more advanced cases (13). Brachytherapy has been used with favorable results in Tis/T1 tumors (14, 15), whereas fractionated orthovoltage irradiation has also been used for relatively superficial invasive tumors (16, 17). Similarly, fractionated radiotherapy using electron (18, 19) or proton beams (20) at conventional dose rates showed 1-year progression-free survival (PFS) rates ranging from 50% to 60% in locally advanced cases. These treatments also induced substantial acute and late side effects, and required multiple fractions with multiple anesthesia. In contrast to these observations, in our study, an excellent normal tissue tolerance to the high single dose of FLASH-RT was found along with a high tumor control rate.

## Materials and Methods

The study was approved by the Swiss "Ethics Committee for Animal Experimentation," and performed within institutional guidelines.

### FLASH-RT linacs

Irradiation was performed using two prototype linacs of type Kinetron (4.5 MeV electrons) and Oriatron 6e (6 MeV electrons; PMB-Alcen; ref. 5). These machines are able to produce electron beams at a mean dose rate ranging from a few Gy/minute (i.e., similar to dose rates conventionally used for radiotherapy) to thousands of Gy/second (2). These wide ranges of dose rates are made possible by varying the linac gun-grid tension, the pulse-repetition frequency, pulse width, and the source-to-surface distance.

### Response of pig skin to FLASH-RT

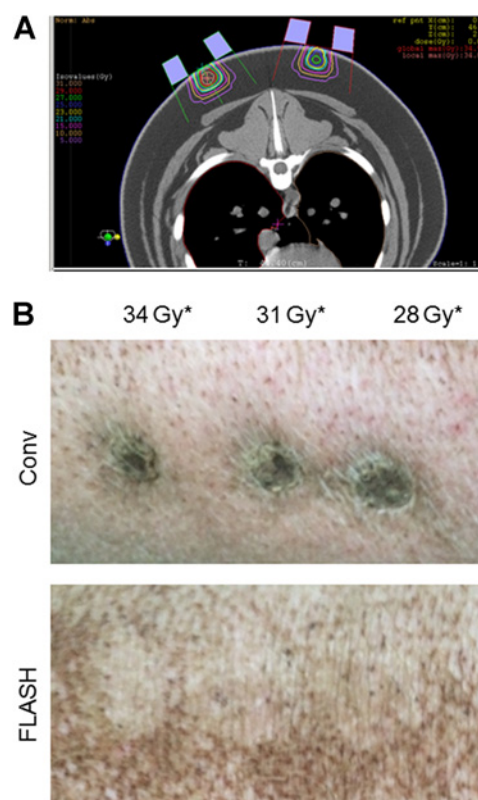
One female Göttingen mini pig (43 kg) entered the study, and was housed at the animal research facilities of the University of Lausanne (Lausanne, Switzerland). Irradiation took place under general anesthesia. The irradiation was performed on the back of the pig using a graphite applicator, with a 26-mm diameter circular aperture, in direct contact with the skin. The doses at 12 mm (5), which ranged from 22 to 34 Gy, were delivered in two ways. Some doses were delivered with a conventional radiotherapy dose rate ( $\approx 5$  Gy/minute), each dose to a different skin patch (one side of back), and then the same doses were delivered to other skin patches (the other side of back) with

FLASH-RT ( $\approx 300$  Gy/second). *In vivo* dosimetry was performed with film (Gafchromic EBT3) and alanine pellets (2, 3) positioned on the skin surface in the center of the beams. A reconstruction of the dose distribution (Fig. 1A) was performed using a CT scan of the pig and dose calculation in XiO (Elekta AB) treatment planning system.

The skin response to irradiation was carefully monitored weekly through visual examination, with any toxicity scored (Table 1) and photographed for 48 weeks postirradiation. At 36 weeks, skin biopsies were performed and analyzed by histology.

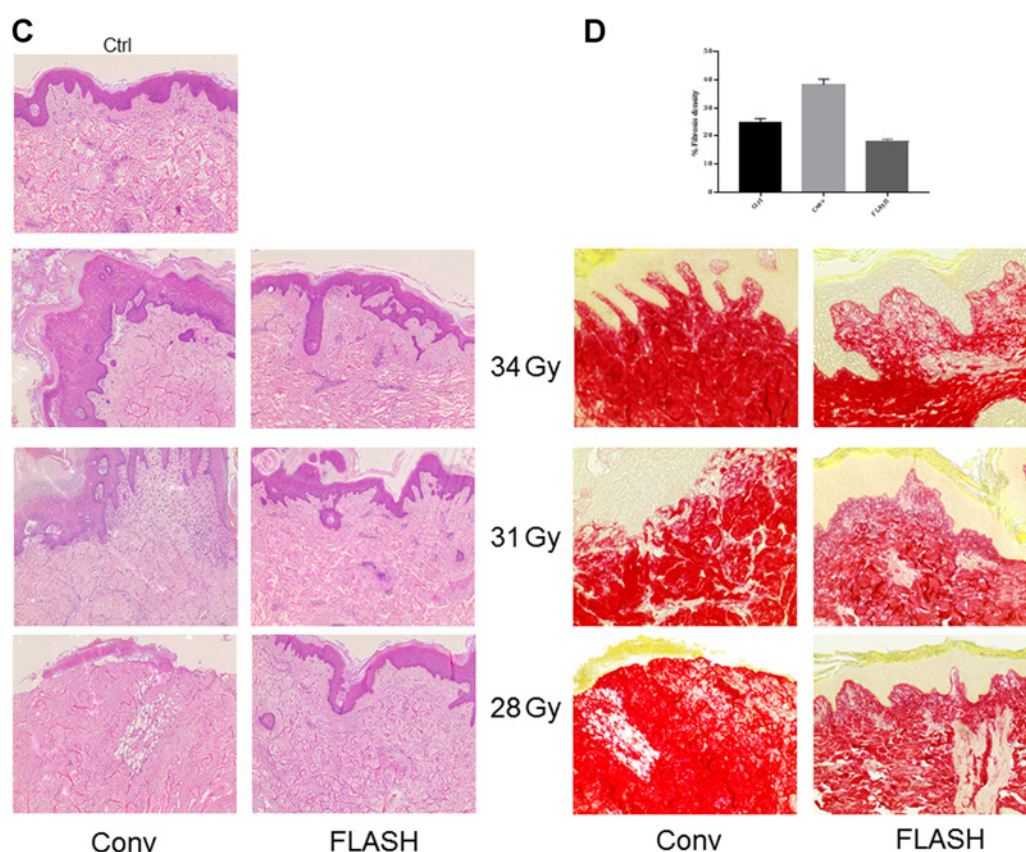
### Cat patient population

From September 2015 to October 2016, 6 previously untreated cats (5 males and 1 female) with histologically confirmed SCC of the nasal planum non-eligible for surgery were enrolled in the FLASH-RT trial. Written consent was obtained from each cat's owner before inclusion in the trial. Pretreatment evaluation included a complete physical examination, a complete blood and platelet count, serum chemistry profile, urine analysis, LDH, CT scan, and three-dimensional measurements of the primary tumor. All cat patients were clinically staged according to the WHO scoring system (Supplementary Table S1). Four patients had a stage T2 tumor and two had a stage T3 (Supplementary Table S2) with no evidence of nodal (N0) or metastatic tumor



**Figure 1.**

**A**, Dose distributions calculated in XiO for the mini pig irradiation; a transversal slice reconstructed from the CT-scan showed beam apertures and dose distributions. **B**, Thirty-six weeks postradiotherapy, macroscopic visualization showed severe fibronecrotic lesions in Conv-irradiated spots and the normal appearance of the skin in FLASH-irradiated spots (\*, maximal dose at 12 mm; ref. 5). (Continued on the following page.)



**Figure 1.**

(Continued.) **C**, H&E staining sections cut from 6-mm punch biopsies sampled in irradiated spots 36 weeks postradiotherapy (magnification  $\times 40$ ). Ctrl = normal pig skin sampled in nonirradiated area. Conv = punch biopsies taken from Conv-irradiated spots show necrotic area, hyperkeratosis, and dermis remodeling. FLASH = punch biopsies taken from FLASH-irradiated spots show normal histology with stratified epidermis, papillary dermis, and intact skin appendages. **D**, Sirius Red staining of sections cut from 6-mm punch biopsies sampled in irradiated spots 36 weeks postradiotherapy (magnification  $\times 40$ ) and densitometric analysis. Enhanced collagen deposition in Conv-RT biopsies as compared with FLASH-RT and control is measured. (Continued on the following page.)

spread (M0). Exclusion criteria included FELV/FIV serum positivity, treatment with corticoids, anti-Cox1/Cox2 drugs, or an anesthesia contraindication.

#### Cat patient treatments

The cats were given a single-fraction radiotherapy while under general anesthesia. The doses followed a dose-escalation scheme, and were prescribed to the surface of the nose with a margin of about 0.5 cm around the gross tumor mass (Table 1). The treatments were planned (set-up, treatment angle, field size, and with/without bolus) for optimally achievable dose distribution using XiO TPS. Graphite applicators with different rectangular apertures were used to limit the treatment field size (Supplementary Table S2), and the cats were in direct contact with the applicator during treatment. *In vivo* dosimetry was performed using either thermoluminescent dosimeters (TLD-100) or alanine pellets (3), with dosimeters positioned on the cats' noses in the center of the beams.

#### Trial design

This study was designed as a phase I dose-escalation trial of FLASH-RT. The first dose level was set at 25 Gy, which is a dose expected to be well tolerated according to the results observed

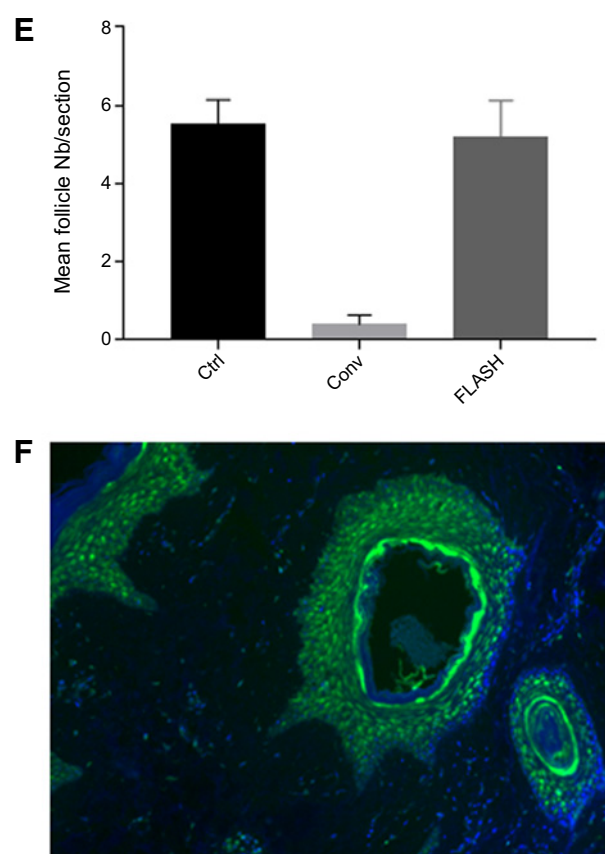
following the pig skin irradiation (see "Results" section). The escalation criteria were as follows: the first cat patient was included at the first dose level and if no dose limiting toxicity (DLT) was observed, the next cat patient would be included with an increased treatment dose. For safety purposes, a minimal interval of 4 weeks between two dose levels was recommended as per protocol along with no evidence of DLT during this follow-up period.

#### Clinical response and follow-up evaluations

The primary endpoint of this phase I study was to determine the MTD, which was defined as the highest dose level at which a cat-patient did not experience a DLT within the 4 weeks postradiotherapy. Normal tissue toxicity was assessed as the primary endpoint according to the VCOG-CTCAE v1.1 (21) implemented with the toxicity score (Table 1) used for the pig (22). A DLT was defined as any acute toxicity above or equal to A5 occurring in 4 weeks postradiotherapy in the treatment field, including stomatitis, mucositis, dysphagia, skin ulceration, or any grade equal or above A6 toxicity outside the radiotherapy field. In the case of DLT, dedicated analgesia and intensive supportive care were planned.

The secondary endpoint encompasses the late normal tissue tolerance in the radiotherapy field and PFS at 12 months. The



**Figure 1.**

(Continued.) **E**, Thirty-six weeks postradiotherapy, quantification of hair follicles per tissue sections has been performed on sections cut from 6-mm punch biopsies taken from nonirradiated, Conv-RT, and FLASH-RT patches (data are presented as number of hair follicle per tissue sections + SE). **F**, CD34<sup>+</sup> cells were detected by immunofluorescence (green intracellular staining) in the preserved hair follicles of FLASH-RT skin; a representative picture taken from 34 Gy FLASH-RT spot is shown.

following criteria were used for evaluating the antitumor efficacy: macroscopic complete response (CR) was defined as the disappearance of all clinical evidence of tumor on physical examination; partial response represents more than 50% decrease of tumor volume on physical examination; stable disease represents a tumor volume change observed on physical examination between a 50% decrease and a 25% increase. In case of doubtful clinical examination, a CT scan and/or biopsies were recommended whenever possible. The overall survival and metastatic dissemination were recorded by the veterinarian along with any acute or late toxicity. These endpoints were checked every day

during the first 15 days postradiotherapy, and at 21 and 28 days postradiotherapy and then every 3 months postradiotherapy. If applicable, any symptomatic medications used after radiotherapy were recorded.

## Results

### Reduced pig skin toxicity with FLASH-RT

The film and alanine pellet *in vivo* dosimetry verified the doses delivered to the pig skin surface (mean and maximum deviation of 0 and 3%, respectively). The reconstruction of the delivered dose distributions visualized the irradiated volumes (Fig. 1A). Acute toxicity was macroscopically evaluated and found to be limited to depilation, which occurred 3 weeks later for the skin patches exposed to FLASH-RT and was transient (lasting only 4 weeks) for doses  $\leq 31$  Gy FLASH-RT (Table 1A and B). Hair follicles were preserved within the skin patches exposed to FLASH-RT, whereas in the skin patches exposed to Conv-RT, the hair follicles appeared to be permanently destroyed and no hair regrowth was observed  $>6$  months (Supplementary Fig. S1). No other acute toxicity was observed (no erythema or moist desquamation). More interestingly, severe late skin fibronecrosis was observed only with Conv-RT and associated with skin contraction appeared 32 weeks posttreatment with 34 Gy and 36 weeks posttreatment with lower doses of 31 and 28 Gy (Table 1). Figure 1B shows macroscopic view of FLASH-RT and Conv-RT irradiated spots, 36 weeks postradiotherapy. At this time point, histologic analysis revealed that skin fibronecrosis in Conv-irradiated spots was associated with epithelial ulceration and hyperkeratosis located at the margin of the wound as well as inflammatory infiltration and severe dermal remodeling (Fig. 1C). In contrast, none of the FLASH-irradiated spots showed skin alteration and histologic analysis was comparable with the nonirradiated skin (Fig. 1C); the cutaneous tissue remained soft and healthy during the 48 weeks of follow-up, whereas some depilation occurred 36 weeks posttreatment and remained stable. Sirius Red staining and quantification confirmed fibrotic remodeling in Conv-irradiated samples (Fig. 1D). Quantification of hair follicle showed preservation of the follicles' number in biopsies taken from the FLASH-RT patches 36 weeks postradiotherapy, whereas they were destroyed in the biopsies taken from Conv-RT spots at the same time point (Fig. 1E). Using the same biopsies, immunofluorescence staining revealed preservation of CD34<sup>+</sup> cells defined as epidermal stem cell in the bulge of the hair follicle in FLASH-irradiated biopsies (Fig. 1F). All fibronecrotic scabs fell off around 42 weeks posttreatment and retractile scars remained stable over the time of follow up (48 weeks).

These results are consistent with our previous observations performed in mice and confirm that FLASH-RT is a nonfibrosing radiotherapy compared with Conv-RT. Furthermore, it shows a

**Table 1A.** Skin toxicity scoring (from ref. 22)

Acute/subacute toxicity	Description	Late toxicity	Description
A0	Depilation	L0	Depilation
A1	Erythema	L1	Atrophy
A2	Pigmentation	L2	Flaky appearance
A3	Dry desquamation	L3	Pigmentation
A4	Moist desquamation that heals	L4	Fibrosis
A5	Moist desquamation that partially heals	L5	Telangiectasia
A6	Moist desquamation that does not heal	L6	Necrosis

Abbreviations: A, acute toxicity; L, late toxicity.

**Table 1B.** Pig skin toxicity follow-up

RT	Dose (Gy)	7w	10w	14w	20w	24w	32w	36w	42w	48w
Conv	22	LO	LO	LO	LO	R	R	N/A	N/A	N/A
Conv	25	LO	LO	LO	LO	LO	LO	N/A	N/A	N/A
Conv	28	LO	LO	LO	LO	LO	LO + L4	LO + L4 + L6	LO + L4	LO + L4
Conv	31	LO	LO	LO	LO	LO	LO + L4	LO + L4 + L6	LO + L4	LO + L4
Conv	34	LO	LO	LO	LO	LO	LO + L1 + L4	LO + L4 + L6	LO + L4	LO + L4
FLASH	22	—	LO	R	R	R	R	N/A	N/A	N/A
FLASH	25	—	LO	R	R	R	R	N/A	N/A	N/A
FLASH	28	—	LO	R	R	R	R	LO	LO	LO
FLASH	31	—	LO	R	R	R	R	LO	LO	LO
FLASH	34	—	LO	LO	LO	LO	LO	LO	LO	LO

NOTE: Dashed line indicates the time of biopsy (at 36w).

Abbreviations: L, late toxicity; N/A, results are not yet available; R, regrowth of hair; RT, radiotherapy; w, week; —, no alteration of the skin.

dose modifying factor of at least 20% for single-fraction treatment in favor of FLASH-RT for the protection of normal tissue and prevention of fibrosis development as endpoints.

#### Cat patient trial: DLT and MTD

The TLD/alanine pellet *in vivo* dosimetry measured the absorbed doses delivered to the skin surface of the cats' noses. It showed a deviation between prescribed and measured doses of 6% for cat no. 1 (measured dose higher) and cat no. 4 (measured dose lower). For the remaining 4 cats, the deviations were <1%. The deviations between prescribed and measured dose values were compatible with the uncertainty of our dose determination with TLD, which is 8% ( $k = 2$ ). As per protocol, cat no. 1 received a single dose of 25 Gy with a  $25 \times 34$  mm field size. Moderate and transient A4 acute mucosal/skin toxicity was scored, including stomatitis of the upper oral cavity (Table 2A). Because DLT ( $\geq A5$  toxicity) was not reached, cat no. 2 with a deeply infiltrating tumor was included but only a moderate dose escalation was used with a dose of 27 Gy and a field size  $25 \times 34$  mm (Fig. 2A–D). Cat no. 2 displayed no detectable signs of "in-radiation field" acute toxicity (no mucositis and no skin erythema). Cat no. 3 had a large ulcerative tumor, requiring a field size of  $34 \times 45$  mm. A single dose of 28 Gy was applied, representing a moderate dose escalation. This cat (no. 3) displayed no detectable signs of acute toxicity (no mucositis and no skin erythema). Cat no. 4 had a bifocal deeply ulcerative tumor, requiring a treatment field size of

$18 \times 26$  mm. A dose escalation to 31 Gy was applied. This cat (no. 4) displayed no detectable signs of acute toxicity (no mucositis and no skin erythema). Cat no. 5 required an irradiation with the same field size ( $18 \times 26$  mm) and a dose escalation to 34 Gy was applied. This cat displayed transient grade A4 "in-radiation field" moist desquamation that healed. Finally, a considerable dose escalation to 41 Gy (field size of  $18 \times 26$  mm) was applied for cat no. 6, who had a deeply infiltrating and ulcerative tumor. Cat no. 6 developed a relatively strong "in-radiation field" mucositis with an exsudative component that was scored A4. Complete healing of the mucosa was obtained 5 weeks after irradiation. Although A4 toxicity observed at 41 Gy was transient and manageable, that is, not considered as a DLT, we did not continue to escalate the dose because CR was observed at all the lower dose levels. Regarding the other side effects observed during the acute phase, no smell and nutrition problems were observed for cats 2–6. Because of a stomatitis, cat no. 1 had difficulties swallowing during weeks 2–4 after irradiation. For all the cats, depilation in the irradiation field was observed, starting about 3 weeks after FLASH-RT (Table 2).

With a median follow-up of 18 months, all the cats revealed permanent depilation that was restricted to the field of irradiation. No other permanent, late toxicity, or "outside-radiation field" side effect has been observed. All 6 cat patients had an excellent functional result, with no limitation of food uptake or smelling capability.

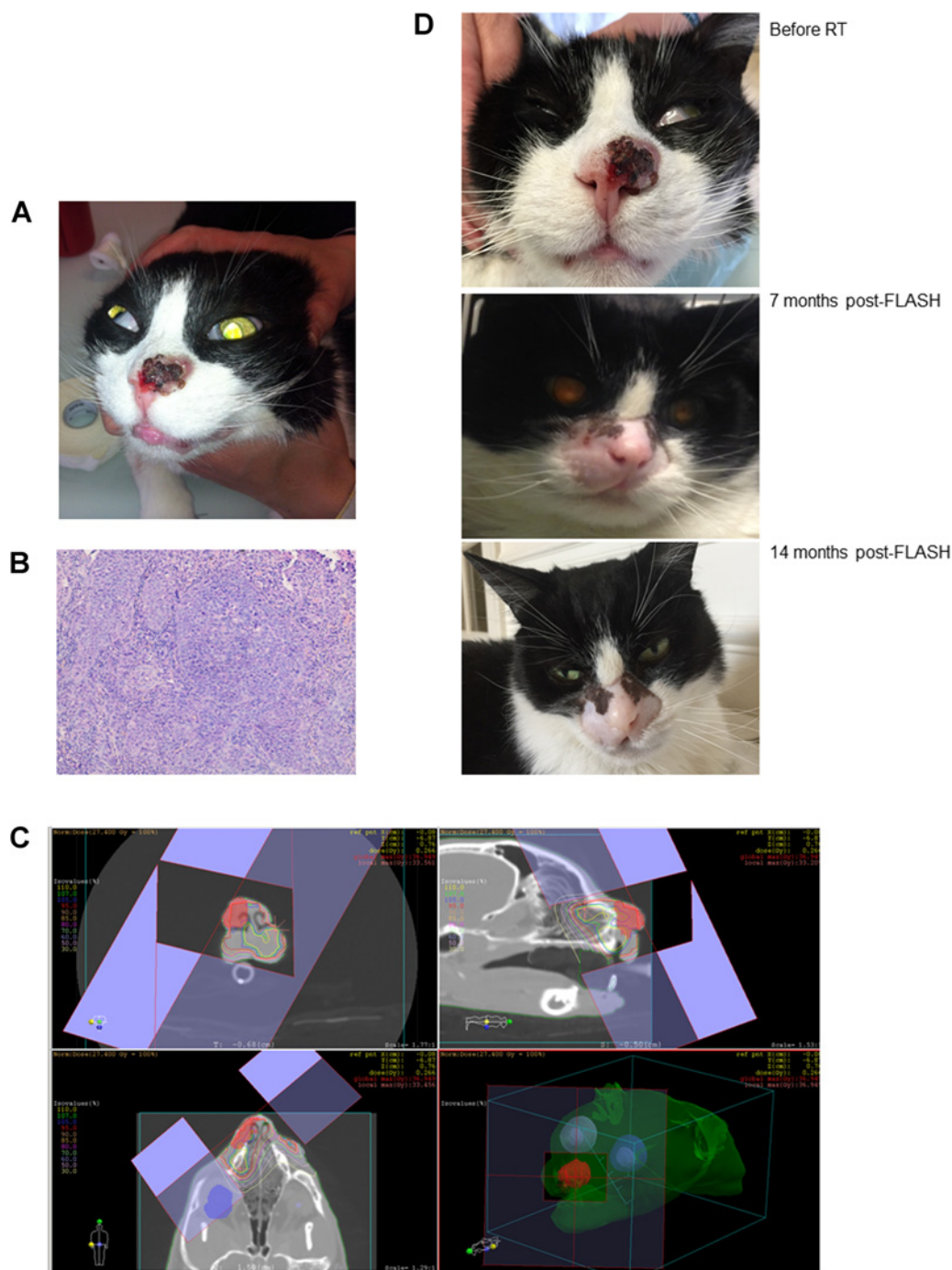
**Table 2.** Follow-up of cats

(A) Adverse events defined by dermatologic/skin scoring system						
Cat no.	Dose (Gy)	Acute toxicity <10 days	Acute toxicity <20 days	Late toxicity 3 months–6 months	Late toxicity 6 months–12 months	Late toxicity >12 months
1	25	A1 + A4	A1 + A4	LO	LO	LO
2	27	—	—	LO	LO	LO
3	28	—	—	LO	LO	N/A
4	31	—	—	LO	LO	LO
5	34	—	A4	LO	LO	LO
6	41	A1 + A4	A1 + A4	LO	LO	LO

(B) Tumor response (CR = macroscopic complete response)			
Cat no.	Dose (Gy)	Response at 3 months	Response at 16 months
1	25	CR	CR
2	27	CR	CR
3	28	CR	Dead
4	31	CR	CR
5	34	CR	CR
6	41	CR	CR

Abbreviations: A, acute toxicity; L, late toxicity; N/A, not available; —, no toxicity.



**Figure 2.** **A**, Picture of cat no. 2 at diagnosis. **B**, Histologic diagnosis of SCC (magnification  $\times 40$ ). **C**, Dose distributions calculated in XiO for the treatment of cat no. 2, with a transversal slice (left), a reconstructed sagittal slice from the CT, a coronal slice, and a 3D view showing the beam apertures, dose distributions, and applicator position directed toward the tumor. **D**, Macroscopic evolution of the lesion in cat no. 2 over time.

**Antitumor efficacy and overall survival**

All 6 cat patients had tumor assessments performed at inclusion and during follow-up. CR was observed for all cats (100%) at 6 months (Table 2B). Cat no. 3 exhibited a recurrence not confirmed by biopsy but with a clinical tumor progression leading

to euthanasia at 8 months. At 16 months, cats 1–6 were disease free and at 18 months, cats 1, 2, and 4 were disease free. For cat no. 4, a biopsy was performed 21 months following FLASH-RT, and showed a SCC corresponding to a local recurrence, the cat is still alive and was proposed for salvage surgery.

## Discussion

The markedly better normal tissue tolerance to FLASH-RT as compared with Conv-RT has been a serendipitous observation, obtained a few years ago in mice exposed to electron-beam irradiation delivered at ultrahigh dose rates. This enhanced tolerance of normal tissues appears at irradiation dose rates above 30–40 Gy/second, whereas on all the tumor models tested so far, FLASH-RT has been shown to be as effective for tumor control as Conv-RT (Montay-Gruel and Vozenin, in preparation; ref. 1). The mechanisms underlying this effect are under investigation, and not yet fully elucidated but the observed increase of the differential effect between tumors and normal tissues appeared to be very promising, and might significantly enhance the radiotherapy therapeutic index.

To prepare the transfer of FLASH-RT in human patients, we tested to which extent the effect obtained with FLASH-RT in mice could be a cross-species observation reproducible in higher mammals. For this purpose, we used two well-characterized models: mini pig and cat patients. Pig skin was chosen as a suitable model for various reasons: (i) it is known to be very close in structure to human skin (8); (ii) it is a well-characterized radiobiological model (8–11); (iii) the size of the pig enabled us to directly compare at the same time and same conditions response to FLASH-RT versus Conv-RT with several spots/dose levels; and (iv) our 6-MeV electron-beam linac is appropriate for skin irradiation. The results obtained when using radiation-induced depilation as acute/delayed endpoint and skin fibrosis as late endpoint were very consistent with our previous observations in mice. Our results show that FLASH-RT minimizes normal tissue damages at the skin level even when large high doses are applied. Using CD34 as a surrogate marker of the epidermal stem cells (23), our results suggest that FLASH-irradiated skin area retain the expression of CD34 supporting the idea that FLASH-RT has minimal impact on stem cell consistently with what was already reported for neural (2) and intestinal stem cells (7). We could estimate a dose-protective factor of at least 20% in favor of FLASH-RT as 28 Gy administered with Conv-RT–induced fibronecrosis, whereas 34 Gy administered FLASH-RT only induced depilation.

In parallel, the effect of FLASH-RT was explored in cat-patients, allowing us to evaluate normal tissue toxicity in the frame of a dose-escalation study. The cohort was too small to allow definitive conclusion for tumor control and, the anti-tumor effect of FLASH-RT was evaluated as a secondary endpoint. SCC of the nasal planum is an appropriate model for which surgery can be functionally devastating and for which Conv-RT is commonly used in clinical practice, albeit with relatively poor outcome and severe toxicity (16, 17, 24). An obvious interest of FLASH-RT is the possibility to use a single curative fraction, whereas Conv-RT requires relatively complex logistics with several fractions and multiple anesthetics that impair the cats' quality of life. The results obtained in our small cohort of T2/T3N0M0 cat patients showed that a MTD was not reached even at 41 Gy single dose. In fact, no DLT was scored and normal tissues exhibited a very good tolerance when irradiated with such a high single dose (25–41 Gy). Indeed, 3 of 6 cats exhibited no acute/late side effects except depilation, and the 3 others had only mild/moderate

and manageable acute mucositis/dermatitis and no late toxicity was seen. Interestingly, a durable tumor control was obtained in 5 of 6 cats at 16 months. Among the most advanced cases of this series, 2 cats (no. 2 and no. 3) had a T3 disease, and were treated with 27 Gy and 28 Gy, respectively. Two cats (no. 3 and no. 4) showed a recurrence suggesting that a dose above 31 Gy would be required for the control of such invasive and spread out tumors.

The good tumor-response rate that was observed is in agreement with previous (15–17) and recent publications (24) that used accelerated fractionation ( $10 \times 4.8$  Gy). However, the main improvement due to FLASH-RT was the good tolerance profile offering a further opportunity for dose escalation to enhance tumor control probability. This observation needs to be extended in a larger cohort of cats but it is in contrast with all previously mentioned studies where frequent grade 2 of 3 (21) toxicity (>30% of patients in ref. 24) was reported, along with PFS at one year of between 50% and 80% (16, 24). A second superiority of FLASH-RT for the treatment of domestic animals is its very convenient single-dose administration.

In conclusion, single-dose FLASH-RT shows promise as a new treatment option for cat patients with locally advanced SCC of the nasal planum. Our results in pig and cats provide a strong rationale for further evaluating FLASH-RT in human patients.

## Disclosure of Potential Conflicts of Interest

M. Jaccard is an employee of Geneva University hospital. J. Bourhis is a consultant/advisory board member for ASTRA, Bristol-Myers Squibb, Merck, and MSD. No potential conflicts of interest were disclosed by the other authors.

## Authors' Contributions

**Conception and design:** M.-C. Vozenin, K. Petersson, M. Jaccard, M. Ozsahin, F. Bochud, J. Bourhis

**Development of methodology:** M.-C. Vozenin, K. Petersson, V. Favaudon, M. Jaccard, G. Ferrand, M. Ozsahin, F. Bochud, C. Bailat, J. Bourhis

**Acquisition of data (provided animals, acquired and managed patients, provided facilities, etc.):** M.-C. Vozenin, P. De Fornel, K. Petersson, V. Favaudon, M. Jaccard, B. Petit, M. Burki, D. Patin, P. Devauchelle, J. Bourhis

**Analysis and interpretation of data (e.g., statistical analysis, biostatistics, computational analysis):** M.-C. Vozenin, K. Petersson, F. Bochud, C. Bailat, J. Bourhis

**Writing, review, and/or revision of the manuscript:** M.-C. Vozenin, P. De Fornel, K. Petersson, V. Favaudon, M. Jaccard, J.-F. Germond, D. Patin, M. Ozsahin, F. Bochud, P. Devauchelle, J. Bourhis

**Administrative, technical, or material support (i.e., reporting or organizing data, constructing databases):** M.-C. Vozenin, K. Petersson, B. Petit, M. Burki, H. Bouchaab, J. Bourhis

**Study supervision:** M.-C. Vozenin, F. Bochud, C. Bailat, J. Bourhis

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