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Use of Protoporphyrin IX Fluorescence to Determine Clinical Target Volume for Non-Melanotic Skin Cancers Treated with Primary Radiotherapy

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Background

Squamous cell (SCC) and basal cell (BCC) of the skin continue to be the most commonly diagnosed cancers worldwide. ¹ Radiotherapy provides local control of 90% and 80% for small and large BCCs respectively. ^{2,3}

The challenge with radiotherapy lies in determining treatment volumes, especially for poorly defined tumors.

Photodynamic therapy (PDT), which involves preferential uptake of a photo-sensitizer that also fluoresces, provides a unique method that may increase accuracy for tumor delineation. ^{4,5}

Two previous surgical series (Brandt, Tierney) demonstrated that protoporphyrin IX (PpIX) fluorescence strongly correlates with the surgical extent of disease. ^{4,6}

This study prospectively investigated PpIX fluorescence as a means of determining the clinical target volume (CTV) necessary to encompass disease for non-melanotic skin cancers.

Materials and Methods

33 biopsy proven SCCs or BCCs. 7 were controls (well demarcated), 26 were poorly demarcated (experimental group).

Gross lesion size recorded (GTV)

ALA, MAL applied, left on 2 hours

Lesion fluorescence recorded (CTV)
Compared to: 5mm expansion if well demarcated, or 10mm if poorly demarcated

Lesion treated to PTV (CTV + 2-3mm), 50Gy in 20 fractions

Results

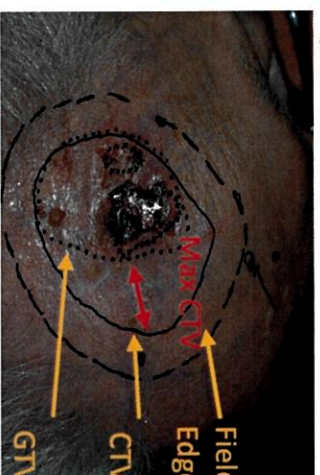
Table 1: Average PpIX CTV margin

Patients	Control	Study
7	7	26
Standard CTV	5mm	10mm
	p=0.50	p=0.01

Max PpIX CTV	9mm	14mm
Patients	Nose 17	Other 16
PpIX CTV	14mm	14mm
	p=0.11	p=0.04

Multi-focal	94%	44%
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Figure 3: treatment plan



Follow-up

Average follow-up was 14 months. 1 local recurrence, was excised Nov. 2012. Patient died 11 months post-excision. 1 distant recurrence – axillary metastases 1 month after completion. Died 7 months after therapy.

Conclusions

PpIX photo-delineation suggests that 10 mm margins may inadequately cover areas of subclinical disease in poorly demarcated skin tumors. Long follow-up may be necessary to determine if this increase in CTV is clinically significant. PpIX photo-delineation also demonstrated that most nasal lesions are multifocal and thus, treatment of the entire nose should be considered.

References

- American Joint Committee on Cancer. Chapter 29. Cutaneous squamous cell carcinoma and other cutaneous carcinomas. AJCC cancer staging handbook, 7th Ed. P3539376.
- Venets ML. The important role of radiotherapy in patients with non-melanoma skin cancer and other cutaneous entities. J Med Imaging Radi Oncol 2008; 52: 278-96.
- Venets ML, Ang KK. Chapter 39. Cutaneous Malignancies. Clinical Radiation Oncology, 3rd Edition. Gunderson & Tepper. Elsevier, 2012. Philadelphia, pp756-67.
- Brandt MG, Moore CC, Jordan K. Randomized control trial of fluorescence-guided surgical excision of nonmelanotic cutaneous malignancies. J Otolaryngol 2007; 36(3): 148-155.
- Lee V, Baron ED. Photodynamic therapy: current evidence and applications in dermatology. Sem Cutan Med Surg 2011; 36: 199-209.
- Tierney E, Petersen J, Hanke CW. Photodynamic diagnosis of tumor margins using methyl aminolevulinate before Mohs micrographic surgery. J Am Acad Dermatol 2011; 64: 913-8.



Figure 1: lesion under white light

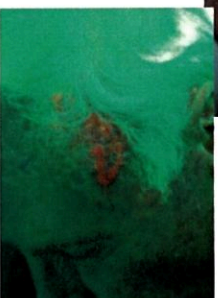


Figure 2: lesion after PpIX fluorescence