Collagen fiber analysis in melanoma: a new candidate to precise lesion border

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Abstract

In melanoma, collagen has been described as a potential new prognostic biomarker. In this way, our purpose was to evaluate the quantity, uniformity and organization of peri- and intratumoral collagen, and to correlate these parameters with clinicopathological features. For this purpose, a total of 22 specimens of cutaneous melanoma were submitted to second-harmonic generation microscopy and image evaluation. Our results suggest that the collagen fiber parameters, mainly organization, are potential candidates to determine with higher precision the border between neoplasia and non-neoplastic dermis.

Key words: Melanoma; Second-harmonic generation image; Collagen; Diagnosis

Introduction

Non linear optics, including two-photon excited fluorescence (TPEF) and second-harmonic generation (SHG), has provided a breakthrough in collagen research in many diseases. In melanoma, collagen has been described as a potential new prognostic biomarker, since it can influence tumor growth, invasion and metastasis. In this perspective, our purpose was to evaluate the quantity, uniformity and organization of peri- and intratumoral collagen, and to correlate these parameters with clinicopathological features.

Materials and Methods

22 samples of paraffin embedded human melanoma were sectioned (4 μ m) and stained with hematoxylin and eosin. Slides were analyzed in an Inverted Zeiss LSM 780-NLO confocal system (Carl Zeiss AG, Germany). Three SHG images from intratumoral (close to melanoma cells) and three from peritumoral (skin distant from melanoma) collagen fibers were captured with an excitation wavelength of 800 nm, a pulse length of approximately 100 fs, and emission filter of SP405 nm (Semrock, Rochester, NY, USA) (**Figure 1**). Images were analyzed in 3 randomly chosen regions for each image, using the ImageJ software (NIH, available at <http://rsb.info.nih.gov/ij/>). Collagen fiber quantity, uniformity and organization of peri- and intratumoral areas were compared using t-test, and the sensitivity and specificity for collagen parameters to distinguish peri- and intratumoral areas were assessed with ROC-curve, using RStudio (CRAN, available at <https://www.rstudio.com/>). Significance was set at $p < 0.05$.

Results and Discussion

The median age of patients was 58 years (31 – 81 years). The tumor was Clark stage I in 09 (40.9%) patients, II in 03 (13.6%), III in 06 (27.3%), IV in 02 (9.1%) and V in 02 (9.1%). The median Breslow was 1,07 mm (0.0 – 6.3 mm). Quantitative analyses of the SHG images demonstrated that intratumoral collagen fibers showed high values of collagen parameters (**Table 1**) in relation to peritumoral collagen fibers. ROC curve was constructed for all parameters to verify the potential of differentiation between peri- and intratumoral collagen: quantity showed 52.% sensitivity and 90.5% specificity; uniformity, 61.9% and 85.7%, and organization 42.9% and 100%.

Figure 1: Representation of autofluorescence tissue sections (A and B) and SHG images (C and D), in peri- (A and C) and intratumoral (B and D) areas.

Table 1. Results for peri- and intratumoral collagen fiber parameters.

	Peritumoral	Intratumoral	p-value
Quantity	14.26 $\pm$ 0.62	14.76 $\pm$ 0.63	<0.001
Uniformity	9.41 $\pm$ 0.74	10.05 $\pm$ 0.77	<0.001
Organization	0.08 $\pm$ 0.04	0.14 $\pm$ 0.06	<0.001

Conclusions

Our results suggest that the collagen fiber parameters, mainly organization, are candidates to determine lesion borders with higher precision.

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