Wavelength characteristics for UVA1 phototherapy with suppressed immediate pigment darkening

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Introduction

UVA1 phototherapy selectively uses the longer UVA1 wavelengths (340-400 nm), and does not include the shorter UVA2 wavelengths (320-340 nm) or UVB wavelengths (290-320 nm) that cause an erythema reaction. Several studies report the effectiveness of UVA1 phototherapy for various diseases such as atopic dermatitis, T-cell lymphoma, and systemic sclerosis. Previously we reported wavelength characteristics of UVA1 phototherapy

Irradiation condition

Table Intensity and irradiation time

	Dose [J/cm ²]	Irradiation distance [mm]	Intensity [mW/cm ²]	Irradiation time [mm:ss]
Sham irradiation	0	-	-	00:00
Filter (-)	15	30	111.4	02:15
A filter (+)	15	30	85.8	02:55
B filter (+)	15	30	97.2	02:34





While UVA1 phototherapy has a high therapeutic effect, it also causes immediate pigment darkening (IPD) as a deleterious effect. IPD is a dull grayish-brown pigment enhancement observed during or immediately after UVA irradiation. IPD begins to disappear 5 to 10 min after irradiation and generally disappears within several hours. Irradiation with UVA at levels several times the threshold level at which IPD develops leads to a persistent pigment darkening that does not disappear even after 24 h and may continue for several weeks. Therefore, IPD may interfere with treatment completion. The peak wavelength of the IPD action spectrum is around 340 nm¹). In an effort to decrease IPD, we investigated the wavelength characteristics of UVA1 phototherapy.



Fig.1 Relative IPD spectral distribution and transmittance of the filter.



Materials&Methods

Normal human epidermal melanocytes (HEM) were irradiated with UVA1 using a short wavelength cut filter or no filter. After irradiation, the cells were incubated for 24 h and tyrosinase mRNA expression was measured by real-time polymerase chain reaction. In addition, the cells were incubated for 8 days and melanin darkening was evaluated by measuring the absorbance of 405 nm using a plate reader.





Fig.2 Absorbance of 405 nm (melanin darkening) and tyrosinase mRNA expression.

Summary

UVA1 irradiation increased the tyrosinase mRNA levels and absorbance – that is, promoted both melanin production and darkening, which was suppressed by use of the filter.





These results suggest that the use of a cut filter may suppress IPD during UVA1 phototherapy.



1) Irwin C, et al., An ultraviolet radiation action spectrum for immediate pigment darkening, Photochem Photobiol. 1993 Mar;57(3):504-7.