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Univariate and Multivariate Spectrophotometric Methods for Simultaneous Determination of Avobenzone and Octinoxate in Pure Form and in Cosmetic Formulations: A Comparative Study.

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Abstract

Simple, economic and precise spectrophotometric and chemometric techniques were used to determine UV filters namely; avobenzone (AV) and octinoxate (OCT) simultaneously in pure form and in cosmetic formulations in concentration range $(2 - 10 \ \mu g.mL^{-1})$ for both drugs. The spectrophotometric technique includes five different methods; Method (A) is first derivative (D¹) spectrophotometry at 380.6 nm for AV and 276.2 nm for OCT, Method (B) is first derivative of ratio spectra (DR¹) at 352.8 nm for AV and 312.2 nm for OCT, Method (C) is ratio difference spectrophotometry (RD) at 356 nm and nm 347.2 nm for AV and at 311.6 nm and 281 nm for OCT, Method (D) is mean centering spectrophotometry (MCR) at 356 nm for AV and 301.8 nm for OCT and method (E) is modified vierordt's method which involves absorbance measurement at 358 nm for AV and 309.2 nm for OCT and determination of the concentration of x and y from the two simultaneous equations. The chemometric technique includes multivariate calibration methods; partial least squares (PLS) and principle component regression (PCR) using the absorption spectra. The proposed methods were applied for determination of (AV) and (OCT) simultaneously in pure form and in cosmetic formulations. These methods were validated according to ICH guidelines.

Keywords: Avobenzone; Octinoxate; UV filters; Spectrophotometry; Chemometry.

1-Introduction:

Ultraviolet (UV) **light** differs in its effect according to its wavelength; UVA (wavelengths 320 to 400 nm) contributes to the long-term harmful effects of photoageing and cancers and can be subdivided into UVA-I (340 to 400 nm) and UVA-II (320 to 340 nm), UVB (wavelengths 290 to 320 nm) contributes to malignant changes and UVC (wavelengths 200 to 290 nm) are highly cytotoxic to human skin but the ozone layer protect earth's surface from its

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