

Research Article

Simple Determination of 17 α -Ethinylestradiol in Hair Restorer by High-Performance Liquid Chromatography Coupled with Fluorescence Detection After Pre-Column Derivatization 4-(*N*-Chloroformylmethyl-*N*-Methylamino)-7-Nitro-2,1,3-Benzoxadiazole

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Abstract

The concentration of 17 α -Ethinylestradiol (EE2) in hair restorer was analyzed by High-Performance Liquid Chromatography (HPLC) coupled with fluorescence detection (excitation wavelength, 470 nm; emission wavelength, 540 nm) after pre-column derivatization with 4-(*N*-chloroformylmethyl-*N*-methylamino)-7-nitro-2,1,3-benzoxadiazole (NBD-COCl). Derivatization with NBD-COCl was performed in borate buffer (pH 9.0) at room temperature for 3 min. The HPLC column was 150 mm \times 3.0 mm i.d., containing 5 μ m particles of C₂₂ packing material, and the mobile phase was prepared by addition of acetonitrile (580 mL) to 420 mL of Milli-Q water containing trifluoroacetic acid (0.1 v/v%). The sample was eluted from the column at room temperature at a flow rate of 0.5 mL/min, and the retention time of NBD-CO-EE2 was 10.7 min. The calibration plot was linear in the range of 0.05 to 1 μ g/mL with an *r*² value of 0.9952, and the lower limit of detection was 0.010 μ g/mL (at a signal-to-noise ratio of 3:1, absolute amount of 0.050 ng/20 μ L injection). The coefficient of variation was less than 9.5%. None of three tested estrogens (estriol, β -estradiol, estrone) interfered with the NBD-CO-EE2 peak. It was found that the content of EE2 in hair restorer was 14.3 \pm 0.9 μ g/mL (range, 13.3 to 15.9 μ g/mL). Recovery in addition-recovery tests was within the range of 70.5 to 98.4%.

Keywords: 17 α -Ethinylestradiol; High-performance liquid chromatography; Fluorescence; 4-(*N*-Chloroformylmethyl-*N*-methylamino)-7-nitro-2,1,3-benzoxadiazole; Derivatization; Hair restorer

Abbreviations

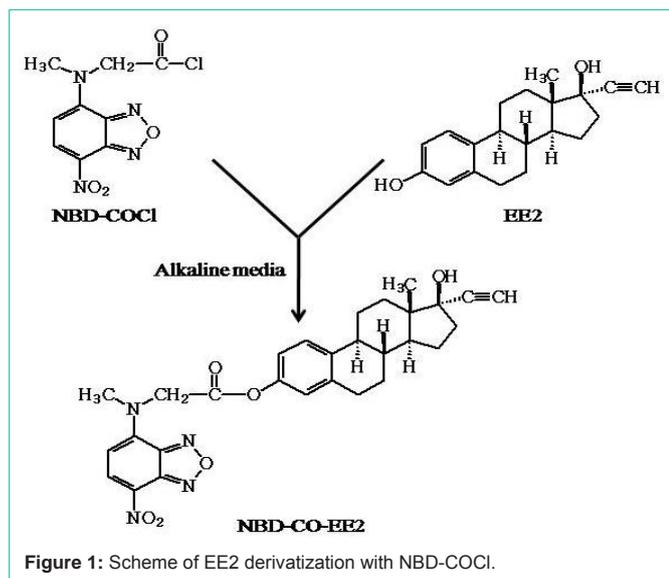
EE2: 17 α -Ethinylestradiol; HPLC: High-Performance Liquid Chromatography; FL: Fluorescence Detection; NBD-COCl: 4-(*N*-Chloroformylmethyl-*N*-methylamino)-7-nitro-2,1,3-benzoxadiazole; GC: Gas Chromatography; MS: Mass Spectrometry; E3: Estriol; E2: β -Estradiol; E1: Estrone

Introduction

17 α -Ethinylestradiol (EE2, 17 α -ethynyl-1,3,5(10)-estratriene-3,17 β -diol) is a synthetic female hormone that is sometimes used in hair restorer and other cosmetics to promote hair growth, skin suppleness, and so on. But, it can have serious side effects. For example, it was reported that a 36-month-old girl presented with vaginal bleeding and uterus enlargement after she had played with her mother's combs and empty vials that had contained hair lotion in which EE2 was present at the concentration of 0.5% [1]. Komori *et al.* suggested that long-term application of cosmetic cream containing EE2 to the face and body three times a day for 75 years might have been related to development of breast cancer and endometrial hyperplasia in a 93-year-old woman [2]. Since EE2 at extremely low levels may induce gynaecological side effects, a simple and sensitive

analytical method for EE2 determination is needed for routine quality control to ensure the safety of personal care products.

Various methods for EE2 determination have been reported, including High-Performance Liquid Chromatography (HPLC) or Gas Chromatography (GC) coupled with Mass Spectrometry (MS) [3-9]. The systems using MS are very sensitive, and have been employed for determination of EE2, other female hormones, and endocrine disrupters in river, sea, and waste water. However, MS is relatively expensive and complicated. In addition, some of the reported methods needed large volumes of sample (more than 1 L) [4,6]. An enzyme-linked immunosorbent assay for quantification of EE2 was also developed [10]. This method is simple and sensitive, but suffers from cross-reactivity, large standard deviation, and the lack of commercially available antiserum. Recently, micro fluidic immunoassay methodology for analysis of EE2 was established for sensitive determination of EE2 in river water samples [11]. In addition, sensitive methods for EE2 analysis by HPLC with Fluorescence detection (FL) have been developed [12-14], including a method employing dansyl chloride derivatization for EE2 analysis in pharmaceutical tablets [14].



However, for determination of EE2 in routine quality control of cosmetic products, we required a rapid, simple, convenient and inexpensive method. It was considered that an HPLC-FL method would be most appropriate, since the equipment is inexpensive, widely available and easy to use, and derivatization with a fluorescent labeling agent generally offers good selectivity and sensitivity. For derivatization, it was focused on 4-(*N*-chloroformylmethyl-*N*-methylamino)-7-nitro-2,1,3-benzoxadiazole (NBD-COCl), because it has already been used as a fluorescent labeling agent of primary and secondary amino groups and phenolic hydroxyl group for HPLC-FL [15-17], and it should also react with the phenolic hydroxyl group of EE2. In this paper, a simple HPLC-FL method for determination of EE2 in hair restorer after pre-column derivatization with NBD-COCl is presented. The derivatization scheme is illustrated in Figure 1.

Material and Methods

Reagents

EE2, estriol (E3), β -Estradiol (E2), and Estrone (E1) were purchased from Sigma-Aldrich Co. (St. Louis, MO, U.S.A.). NBD-COCl was obtained from Tokyo Chemical Industry Co., Ltd. (Tokyo, Japan). Hair restorer was purchased from a market in Kanazawa city, Ishikawa Pref., Japan. Although EE2 was stated to be present on the container label, the concentration was not given. Other general reagents were obtained from Wako Pure Chemical Industries (Osaka, Japan).

Equipment

The HPLC system comprised a model L-6200 pump (Hitachi, Tokyo, Japan), a Rheodyne injection valve (Cotati, CA, USA) with a 20- μ L loop and a model RF-10A fluorometer (Shimadzu, Kyoto, Japan) operating at an excitation wavelength of 470 nm and an emission wavelength of 540 nm. The HPLC column (C₂₂-MS-II, Nacalai tesque, Kyoto, Japan) was 150 mm \times 3.0 mm i.d., containing 5 μ m particles of C₂₂ packing material. Quantification of peaks was performed using a Chromatopac Model C-R3A integrator (Shimadzu). The mobile phase was prepared by the addition of acetonitrile (580 mL) to 420 mL of Milli-Q water containing trifluoroacetic acid (0.1 v/v%). The

samples were eluted from the column at room temperature at a flow rate of 0.5 mL/min.

Derivatization

Ultrapure water was from a Milli-Q water purification system (Simplicity[®] UV, Millipore Corporation, Bedford, MA, USA). A standard solution of EE2 (2 mg) in methanol (50 mL) was prepared and stocked at 4°C. Working standard solutions (0, 0.05, 0.1, 0.25, 0.5, and 1 μ g/mL) were prepared by dilution with 10% methanol. Borate buffer (0.1 M) was adjusted to pH 9.0 by the addition of NaOH. Borate buffer (50 μ L) was added to a diluted standard sample (50 μ L), then NBD-COCl solution in acetonitrile (50 μ L, 3 mg/mL) was added. The mixture was vortexed and allowed to react for 3 min at room temperature, and then ice-cold HCl solution (50 μ L, 0.1 M) was added to stop the reaction. An aliquot of 20 μ L was injected into the HPLC system.

Sample preparation and addition-recovery tests

An aliquot of hair restorer (200 μ L) was diluted to 20 mL with 10% methanol, derivatized, and analyzed as described above. Addition-recovery tests were carried out to assess the accuracy of the method by spiking hair restorer (200 μ L) with EE2 (1.50 or 3.00 μ g), and diluting it in the same manner. An aliquot of 50 μ L was analyzed and the EE2 concentration in the sample was determined. Recovery was calculated as follows.

$$\text{Recovery (\%)} = \frac{(\text{Total amount after spiking}) - (\text{Original amount})}{(\text{Spiked amount})} \times 100$$

Results and Discussion

For the time course study, the reaction time was set at 1, 2, 3, 5, 7, or 10 min at room temperature. EE2 (50 μ L, 1 μ g/mL), borate buffer (50 μ L, pH 9.0), and NBD-COCl (50 μ L, 3 mg/mL) were mixed as described in Derivatization. The derivatization of EE2 reached a maximum at 3 min (Figure 2). Next, pH dependency (pH 8.0 to 10.0) was examined at the derivatization time of 3 min at room temperature. The peak area of NBD-CO-EE2 was maximal at pH 9.0, and it was considerably reduced at pH 8.0 (Figure 3). In addition, the temperature dependency was examined at 3 min and pH 9.0. Peak areas at 4°C (on ice), room temperature, 35°C, 45°C, and 55°C were

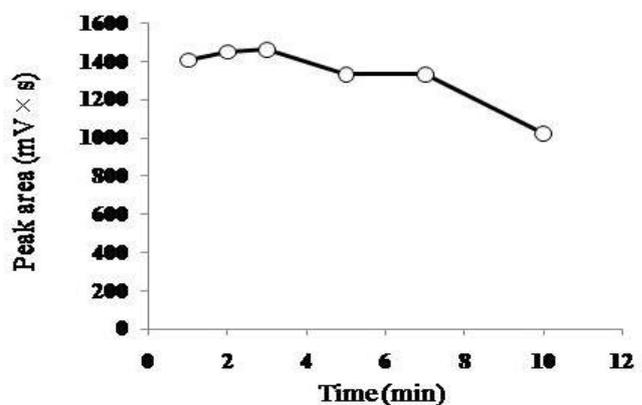


Figure 2: Time course of formation of NBD-CO derivative of EE2. A standard sample (1 μ g/mL, 5 ng/20 μ L of injection) was reacted with NBD-COCl in borate buffer at pH 9.0 at room temperature. Data are expressed as mean values of two experiments.

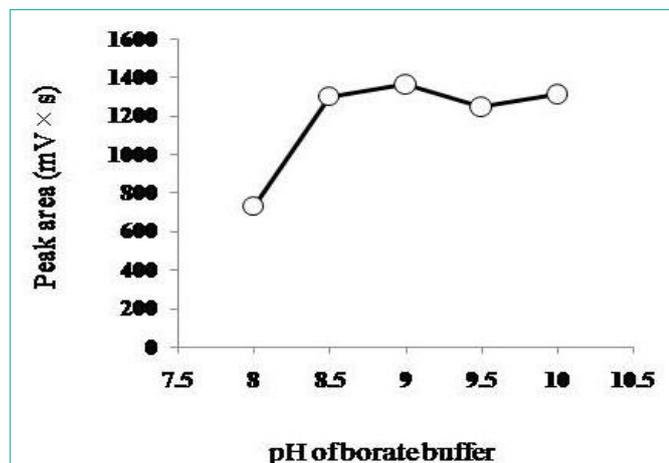


Figure 3: pH dependency of the formation of NBD-CO derivative of EE2. A standard sample (1 $\mu\text{g/mL}$, 5 ng/20 μL of injection) was reacted with NBD-COCl for 3 min in various borate buffers at room temperature. Data are expressed as mean values of two experiments.

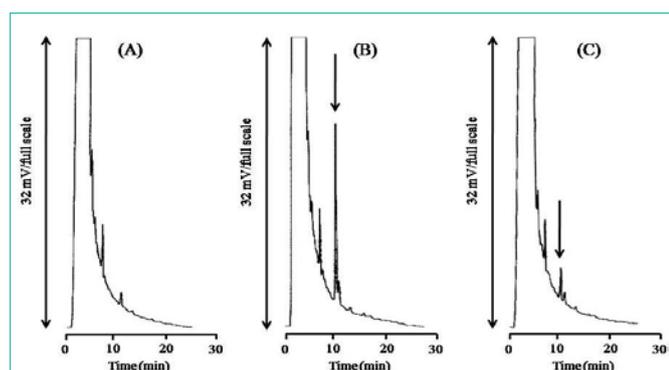


Figure 4A and 4B: Typical chromatograms of blank (A) and standard sample (B) after derivatization with NBD-COCl. A standard sample (0.5 $\mu\text{g/mL}$, 2.5 ng/20 μL of injection volume) was reacted with NBD-COCl for 3 min at pH 9.0 at room temperature. Retention time: 10.7 min, NBD-CO-EE2 (arrowed peak).

Figure 4C: Typical chromatogram of hair restorer sample (C) after derivatization with NBD-COCl. The test sample was reacted with NBD-COCl for 3 min at pH 9.0 at room temperature. NBD-CO-EE2 peak (arrowed peak) was detected at 10.7 min.

1082, 1459, 1360, 1103, and 998 $\text{mV} \times \text{s}$, respectively (mean values of two experiments). The peak area of NBD-CO-EE2 was maximum when the derivatization was performed at room temperature. Thus, the derivatization time of 3 min at pH 9.0 and room temperature was selected. Additionally, in preliminary test, peak area of NBD-CO-EE2 using C_{22} -MS-II column was tended to slightly increase compared with C_{18} -MS-II column (150 $\text{mm} \times 3.0$ mm i.d., containing 5 μm particles of C_{18} packing material, Nacalai tesque), and separated from the later blank noise peak at 11.3 min (data not shown). Therefore, C_{22} -MS-II column was chosen as an analytical column.

Figure 4 shows typical chromatograms obtained from (A) blank and (B) standard sample (0.5 $\mu\text{g/mL}$). The retention time of NBD-CO-EE2 was 10.7 min.

A standard curve was constructed by plotting integrated peak area vs. concentration of EE2. The plot was linear ($y = 1531x - 51.33$) in the range of 0.05 to 1 $\mu\text{g/mL}$ with an r^2 value of 0.9952. The values

Table 1: Sensitivity of various methods for determination of EE2.

Method	Limit of detection	Reference
HPLC-MS	1.4 ng/L ^{**}	[3]
HPLC-MS	5×10^{-3} ng [*] ; 0.45 ng/L ^{**}	[4]
Ultra HPLC-MS	1.2 ng/L ^{**}	[5]
HPLC-MS	6 ng/L ^{**}	[6]
HPLC-MS	10.5 ng/L ^{**}	[7]
GC-MS	5.0 ng/L ^{**}	[8]
GC-MS	1 ng/L ^{**}	[9]
Enzyme-linked immunosorbent assay	14 ng/L ^{**}	[10]
Microfluidic immunoassay	0.006 ng/L ^{**}	[11]
HPLC-FL	0.0566 ng [*] ; 283 ng/L ^{**}	[12]
HPLC-FL	6.5×10^2 ng/L ^{**}	[13]
HPLC-FL	0.04 ng [*]	[14]
HPLC-FL	0.050 ng [*] ; 1.0×10^4 ng/L ^{**}	This paper

The limit of detection was expressed as absolute amount (ng)^{*} and/or concentration (ng/L)^{**}.

Table 2: Intra- and Inter-day assay reproducibility for determination of EE2.

	Concentration ($\mu\text{g/mL}$)	Measured ($\mu\text{g/mL}$, Mean \pm S.D., n=5)	C.V. (%)	Recovery (%)
Intra-day	0.05	0.0452 ± 0.0038	8.4	90.4
	0.25	0.232 ± 0.013	5.6	92.8
	1	0.980 ± 0.044	4.5	98
Inter-day	0.05	0.0440 ± 0.0042	9.5	88
	0.25	0.228 ± 0.015	6.6	91.2
	1	0.964 ± 0.056	5.8	96.4

of the lower limits of quantification and detection were 0.033 $\mu\text{g/mL}$ (absolute amount of 0.17 ng/20 μL injection, signal-to-noise ratio of 10:1) and 0.010 $\mu\text{g/mL}$ (absolute amount of 0.050 ng/20 μL injection, signal-to-noise ratio of 3:1), respectively. As shown in Table 1, the sensitivity of the author's method is considerably inferior to those of previous HPLC- or GC-MS methods [3-9], two immunoassays [10,11], and the HPLC-FL method of Matsumoto *et al.* [13], but is similar to those of previously reported HPLC-FL methods [12,14].

Precision and accuracy for intra-day and inter-day assays of EE2 are shown in Table 2. In the intra-day assay, the range of standard deviation was within 4.5 to 8.4% of the mean, and recoveries were within the range of 90.4 to 98.0%. In the inter-day assay, the range of standard deviation was within 5.8 to 9.5% of the mean, and recoveries were within the range of 88.0 to 96.4%. The accuracy and precision was satisfactory in the tested concentration. However, the data showed the high C.V. value and low recovery on the assays at 0.05 $\mu\text{g/mL}$. It was considered that a blank peak at 11.3 min slightly would interfere with NBD-CO-EE2 peak.

Interference with the detection of NBD-CO-EE2 by E3, E2, and E1 (each 1 $\mu\text{g/mL}$) was investigated. Derivatization was performed as described in Derivatization. The retention times of NBD-CO-E2 and NBD-CO-E1 were 8.3 and 11.8 min, respectively. On the other hand, the NBD-CO-E3 peak was not observed, because it overlapped with a large blank peak. None of the three tested estrogens interfered with the NBD-CO-EE2 peak.

Table 3: Level of EE2 in hair restorer and recovery of spiked EE2.

Assay	Content ($\mu\text{g/mL}$)	Recovery (%)	
		Added (1.50 μg)	Added (3.00 μg)
Day 1	15.9	74.1	89.8
Day 2	13.6	87.7	98.4
Day 3	14.3	84.6	94.8
Day 4	14.2	70.5	93.7
Day 5	14.6	75.7	90.2
Day 6	14.3	76.4	93.5
Ave. \pm S.D. (R.S.D.)	14.3 \pm 0.9 (6.3%, $n=6$)	85.8 \pm 9.4% (11.0%, $n=12$)	

Figure 4C shows a typical chromatogram obtained from a sample of hair restorer. The peak of NBD-CO-EE2 was detected at 10.7 min, and the concentration of EE2 in the sample was determined as follows.

The described method was used to determine EE2 in the restorer and in samples spiked with standard compound. As shown in Table 3, the concentration of EE2 in the restorer was found to be 14.3 \pm 0.9 $\mu\text{g/mL}$ (range, 13.3 to 15.9 $\mu\text{g/mL}$). Recovery of spiked EE2 from the restorer was 85.8 \pm 9.4% (range, 70.5 to 98.4%). The restorer contained EE2 at the level of 1.72 \pm 0.11 mg per package. The present system is able to detect EE2 in hair restorer when the content is more than 108 μg per package (more than 0.9 $\mu\text{g/mL}$).

Conclusion

An HPLC-FL method for determination of EE2 in hair restorer by using NBD-COCl as a fluorescent labeling reagent has been developed. A tested hair restorer was found to contain EE2 at the mean concentration of 14.3 $\mu\text{g/mL}$ using the present system. This method is simple, convenient and inexpensive, and should be suitable for routine quality assessment of cosmetics.

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