

Application News

Quantitative Analysis of Tocopherols in Vegetable Oils

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User Benefits

- ◆ Tocopherols (α -, β -, γ -, and δ - tocopherol) can be simultaneously analyzed in about four minutes.
- ◆ Concentrations of tocopherols in vegetable oils can be determined.
- ◆ Limited consumption of organic solvents reduces operating costs.

Introduction

Tocopherols are a class of structurally related compounds that partly constitute vitamin E. They are important natural fat-soluble compounds with outstanding antioxidant properties, which assist with neurotransmission, keep the muscles working well, prevent blood clots, and boost immune system performance. Plant-based oils, nuts, seeds, fruits, and vegetables are rich in vitamin E.

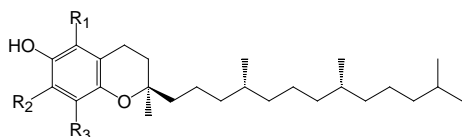
CO₂, which is used in supercritical fluid chromatography (SFC), is mainly used to analyze compounds subject to normal phase separation in liquid chromatography (NPLC). In addition, supercritical fluid is unique in terms of diffusivity and viscosity, with diffusivity about 100 times that of liquids, and viscosity approximately 1/10 that of liquids.

Tocopherols consist of four analogous forms, alpha-, beta-, gamma-, and delta-tocopherol, with slight structural differences. Quantitative tocopherol analysis can be performed using NPLC, but transferring from LC to SFC reduces the consumption of organic solvents. Furthermore, analysis time can be decreased while maintaining identical peak resolution by increasing the flowrate.

This article introduces examples of the simultaneous quantitative analysis of tocopherols in vegetable oils using the Nexera UC.

Analysis of a Mixed Standard Solution of Tocopherols

Fig. 1 shows the structural formula for tocopherol. There are four forms of tocopherol, α , β , γ , and δ , which differ depending on the number and position of methyl groups in the chromanol ring. Fig. 2 shows the chromatogram of a mixed standard solution of tocopherols (containing 50 mg/L of each standard in *n*-hexane) and Table 1 lists the analytical conditions.



| | R ₁ | R ₂ | R ₃ |
|----------|-----------------|-----------------|-----------------|
| α | CH ₃ | CH ₃ | CH ₃ |
| β | CH ₃ | H | CH ₃ |
| γ | H | CH ₃ | CH ₃ |
| δ | H | H | CH ₃ |

Fig. 1 Chemical Structure of Tocopherol

Table 1 Analytical Conditions

| | |
|------------------|--|
| System | : Nexera UC |
| Column | : Shim-pack™ UC-NH ₂ * ¹ (250 mm x 4.6 mm I.D., 5 μ m) |
| Mobile Phase | : A) CO ₂ B) Methanol |
| Time Program | : B conc. 3 % (0 - 4 min) → 50 % (4 - 5 min) → 50 % (5 - 7 min) → 3 % (7.01 - 9 min) |
| Flowrate | : 5.0 mL/min |
| Column Temp. | : 40 °C |
| Injection Volume | : 20 μ L |
| BPR Pressure | : 15 MPa |
| BPR Temperature | : 50 °C |
| Vial | : SHIMADZU LabTotal for LC 1.5 mL, Glass* ² |
| Detection | : 295 nm (SPD-M40 with a high-pressure flow cell) |

*1 P/N: 227-30423-04 *2 P/N: 227-34001-01

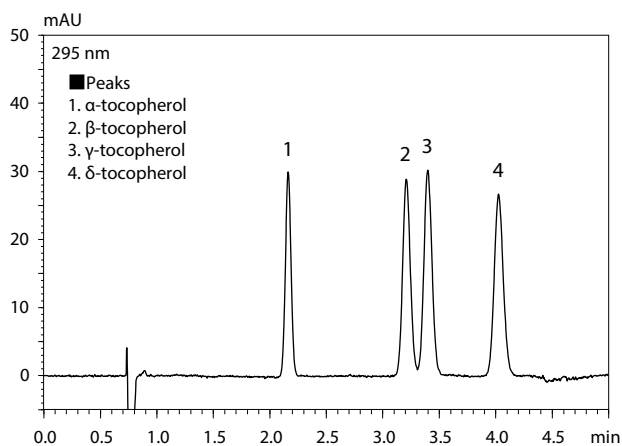


Fig. 2 Chromatogram of a Mixed Standard Solution of Tocopherols at 50 mg/L

Calibration Curves

Fig. 3 shows the calibration curves. For each of the tocopherols, a calibration curve was created with the range shown in Table 2. Table 2 shows the calibration ranges adjusted based on the actual content of the respective forms and the linearity of the respective calibration curves. All calibration curves yielded favorable linearity, with coefficients of determination of 0.9995 or higher.

Table 2 Linear range and coefficient of determination (r²)

| Compound | Calibration range (mg/L) | r ² |
|----------------------|--------------------------|----------------|
| α -tocopherol | 10 - 400 | 0.9995 |
| β -tocopherol | 1 - 20 | 0.9995 |
| γ -tocopherol | 10 - 400 | 0.9996 |
| δ -tocopherol | 2 - 100 | 0.9995 |

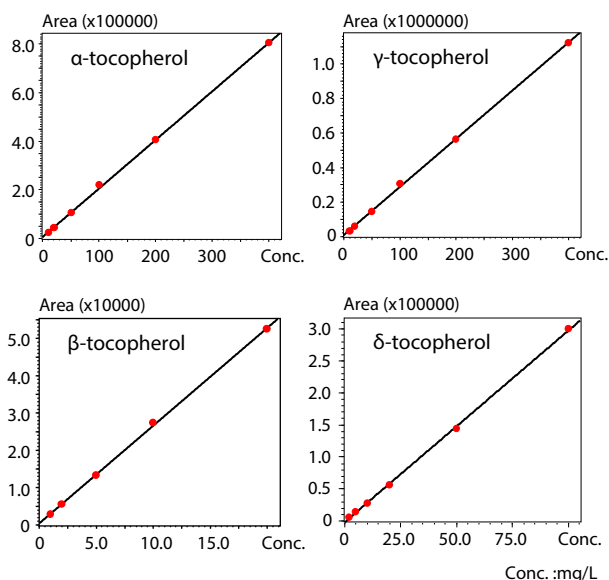


Fig. 3 Calibration Curves

■ Analysis of Vegetable Oils

Quantitative analyses of tocopherols were performed using commercially available sunflower oil, olive oil, palm oil, and sesame oil.

500 μL of each oil was diluted with *n*-hexane to prepare the respective oil samples. The prepared samples were subjected to vortex mixing before SFC analysis.

Table 3 shows the concentrations of tocopherols in the vegetable oils calculated at a specific gravity of 0.9 g/cm³. The chromatograms of the vegetable oils are shown in Figs. 4, 5, 6, and 7.

Table 3 Concentrations of Tocopherols in the Vegetable Oils

| Compound | Concentration (mg/100 g) | | | |
|--------------|--------------------------|-----------|----------|------------|
| | Sunflower oil | Olive oil | Palm oil | Sesame oil |
| α-tocopherol | 53.4 | 13.1 | 8.2 | 9.6 |
| β-tocopherol | 1.3 | N.D. | N.D. | 1.2 |
| γ-tocopherol | N.D. | N.D. | 4.9 | 32.6 |
| δ-tocopherol | N.D. | N.D. | 7.6 | 13.3 |

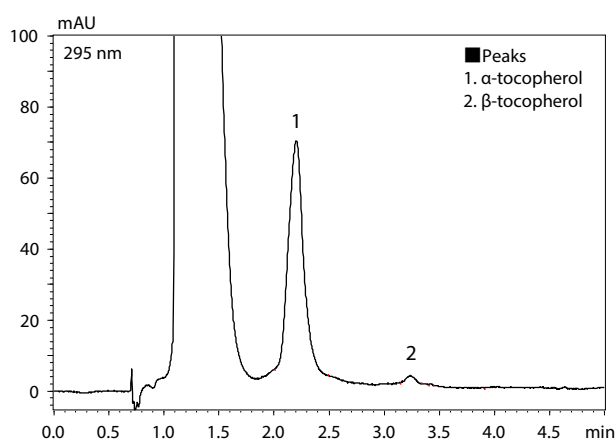


Fig. 4 Chromatogram of Sunflower Oil

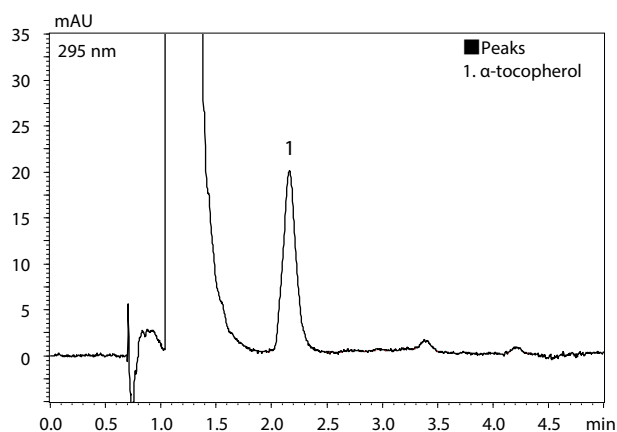


Fig. 5 Chromatogram of Olive Oil

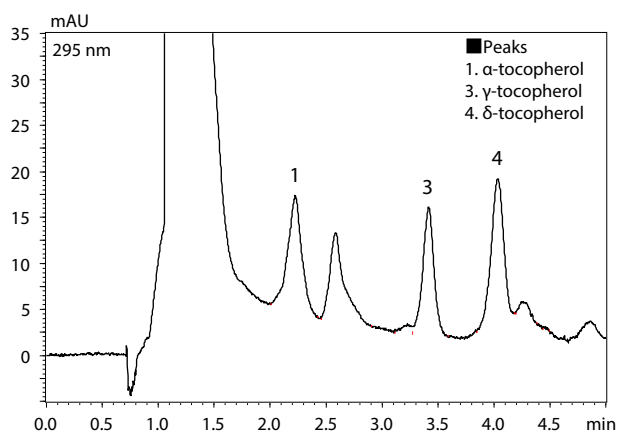


Fig. 6 Chromatogram of Palm Oil

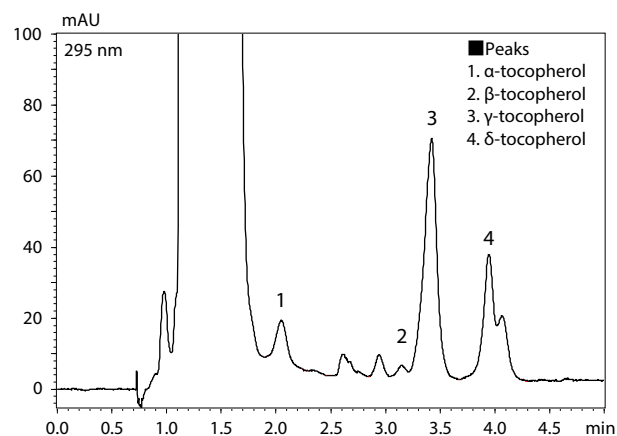


Fig. 7 Chromatogram of Sesame Oil

■ Conclusion

This article introduced examples of the quantitative analysis of tocopherols in vegetable oils using the Nexera UC. The elution of compounds takes at least ten minutes in tocopherol analysis using NPLC, but only four minutes in SFC analysis.

Furthermore, the CO₂ used for SFC analysis is less expensive than the *n*-hexane and other organic solvents used in NPLC, and waste disposal is also cheaper.

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