

Applications note

MSTips No.398 GC-QMS Application

Analysis of di(2-ethylhexyl)phthalate, dichloroacetonitrile and chloral hydrate by GC-MS method using nitrogen carrier gas

Product used: Mass spectrometer (MS)

1. Overview

Helium (He), which is widely used as a carrier gas for GC, can have problems such as temporary price increases and unstable supply conditions due to various reasons, and when supply delays occur, it is necessary to consider using another type of carrier gas as an alternative. Hydrogen and nitrogen are the main alternative gases being considered, with nitrogen gas being relatively easier to introduce into GC-MS when safety is a priority. In this report, di(2-ethylhexyl)phthalate, dichloroacetonitrile, and chloral hydrate, which are water quality control target items in water quality testing, were measured by the same column using nitrogen carrier gas. (For He, see MSTips No.325.) As a result, good linearity of the calibration curve and reproducibility at the lower limit of quantification were obtained for all compounds, and are presented in this report.



JMS-Q1600GC UltraQuad™ SQ-Zeta

1.1. Measurement Condition

The measurements were performed using a gas chromatograph quadrupole mass spectrometer "JMS-Q1600GC UltraQuad™ SQ-Zeta". The measurement conditions are shown in Table 1. As mentioned above, the column used was the same, and each compound was measured by changing the GC conditions. The monitor ions for SIM described in the Japanese "Notification Law" were used.

Parameter Di(2-ethylhexyl)phthalate Dichloroacetonitrile & Chloral hydrate 50°C(2min)→20°C/min→180°C(0min)→ 35°C(3.5min)→15°C/min→100°C Oven temp. 5°C/min→260°C(10.5min)→10°C/min→ $(0min)\rightarrow 20^{\circ}C/min\rightarrow 250^{\circ}C(3min),$ 280°C(5min), Total 42min Total 18.3min GC Column flow (Nitrogen) 1mL/min 1mL/min Splitless, Purge time 1min Injection mode Splitless, Purge time 0.5min Injection volume 1uL GL Sciences Inc. InertCap 1MS, 30m x 0.25mm id, 1µm film thickness Column 250°C Inlet temp. Interface temp. 250°C lon source temp. 250°C MS Ionization EI(20eV, 50µA) Acquisition mode SIM

Table. 1 Measurement Condition of Each Compound

2. Di(2-ethylhexyl)phthalate

2.1. Measurement

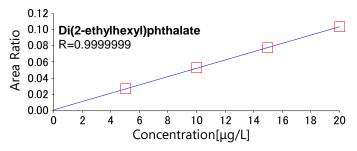
The test solution was diluted with n-hexane to achieve concentrations of 5, 10, 15, and 20 µg/L of di(2-ethylhexyl)phthalate in the test water before pretreatment. Phenanthrene-D10 was added as an internal standard at a concentration of 250 µg/L.

2.2 Result

The calibration curve for di(2-ethylhexyl)phthalate is shown in Figure 1. The coefficient of correlation for the relative calibration curve was more than 0.999. SIM chromatogram and the coefficient of variation of the quantitation value for the 5 μ g/L sample measured continuously at n=5 are shown in Figure 2 and Table 2, respectively. The coefficient of variation of the quantification value at 5 μ g/L, which is less than 1/10 of the target value of 80 μ g/L, was within 5%, a good result.

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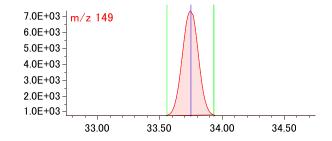


Figure 1. Calibration curve of Di(2-ethylhexyl)phthalate Figure 2. SIM chromatogram of Di(2-ethylhexyl)phthalate at 5µg/L

Table 2. Coefficient variation (C.V.) of Di(2-ethylhexyl)phthalate at 5µg/L

	C.V. %					
#1	#2	#3	#4	#5	C.V. 70	
5.16	5.19	5.19	5.18	5.18	0.2	

3. Dichloroacetonitrile and Chloral hydrate

3.1. Measurement

The test solution was prepared by dilution with MTBE to achieve concentrations of 1, 3, 5, and 15 μ g/L of dichloroacetonitril e and chloral hydrate in the test water before pretreatment. 1,2,3-trichloropropane was added as an internal standard at a concentration of 125 μ g/L.

3.2. Result

The calibration curves for dichloroacetonitrile and chloral hydrate are shown in Figure 3. The correlation coefficients for the relative calibration curves were more than 0.999. The SIM chromatograms and the coefficient of variation values of the quantification values for the continuous measurement of 1 μ g/L samples at n=5 are shown in Figure 4 and Table 3, respectively. The coefficient of variation of the quantification value at 1 μ g/L, which is less than 1/10 of the target value (dichloroacetonitrile: 10 μ g/L, chloral hydrate: 20 μ g/L), was within 5%, a good result.

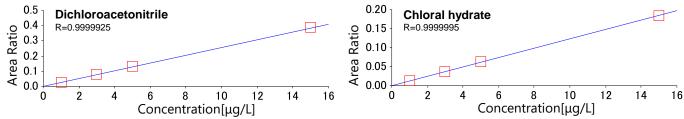


Figure 3. Calibration curve of dichloroacetonitrile & chloral hydrate

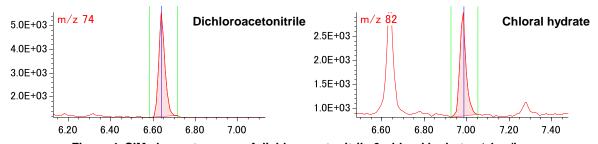


Figure 4. SIM chromatograms of dichloroacetonitrile & chloral hydrate at $1\mu g/L$

Table 3. Coefficient variation of dichloroacetonitrile & chloral hydrate at 1µg/L

Compound name	Q	C.V.%				
Compound name	#1	#2	#3	#4	#5	C.V./0
Dichloroacetonitrile	0.93	0.92	0.95	0.93	0.91	1.9
Chloralhydrate	0.99	1.03	1.01	1.00	1.00	1.7

4. Summary

Di(2-ethylhexyl) phthalate, dichloroacetonitrile, and chloral hydrate were measured by the same column using a nitrogen carrier gas. The linearity of the calibration curve and the reproducibility at concentrations less than 1/10 of the target value were good, confirming that the measurement by the same column using a nitrogen carrier gas can be used adequately.

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