

Nota de aplicación

Application of UPC² in Extractables Analysis

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Abstract

In this application, four different types of packaging material were extracted, including a high density polypropylene pill bottle (HDPE), a low density polypropylene bottle (LDPE), an ethylene vinyl-acetate plasma bag (EVA), and a polyvinyl chloride blister pack (PVC). The extracts were screened for 14 common polymer additives.

Benefits

- UPC² provides a technique for analysis of non-volatile and semi-volatile extractables, as well as polar and non-polar compounds
- Provides a turnkey single instrument approach for extractable and leachable studies
- Rapid analysis of container closure systems used for pharmaceutical, food, and clinical products

Introduction

Extractables from packaging materials are a concern to manufacturers and suppliers of containers for the heavily regulated pharmaceutical and food industries.¹⁻³ Due to these regulations, packaging material manufacturers are motivated to control and monitor their product to ensure that no potential risk exists from extractable and leachable material. Similarly, the manufacturers of supplies for industrial processes, such as plastic vessels and filters, are required to demonstrate that their products do not add any leachables in the production process.

The initial investigation, called a controlled extraction study, qualitatively and quantitatively investigates the nature of extractable profiles from critical container closure system components. It is performed early in device and packaging development. The testing involves solvent extraction techniques encompassing a range of polarity, solvent compatibility studies, and multiple analytical techniques. One of the limitations encountered in these studies involved matching the solvent extracts with the appropriate analytical technique. For example, non-polar solvent extracts can be directly injected into a gas chromatography (GC) system but must be evaporated and reconstituted with a solvent compatible with a liquid chromatography (LC) system. Likewise, water extracts must be back-extracted into a non-polar solvent for analysis by GC. UltraPerformance Convergence Chromatography (UPC²), built on the principles of supercritical fluid chromatography (SFC), allows different types of extraction solvents to be injected for separation on one system for analysis, thereby saving time and reducing sample preparation efforts.

In this application, four different types of packaging material were extracted, including a high density

polypropylene pill bottle (HDPE), a low density polypropylene bottle (LDPE), an ethylene vinyl-acetate plasma bag (EVA), and a polyvinyl chloride blister pack (PVC). The extracts were screened for 14 common polymer additives. Hexane, isopropanol (IPA), and water were used as the extraction solvents. GC-MS was used to analyze hexane and IPA extracts, the ACQUITY UPLC System was used to analyze water and IPA extracts, and the ACQUITY UPC² System was used to analyze all three solvent extracts. The UPC² analysis was compared to the GC and UPLC chromatographic profiles.

Experimental

Sample description

Samples were prepared by microwave extraction. The samples of HDPE, LDPE, EVA, and PVC (2 g) were extracted in 10 mL of isopropanol or hexane for 3 h at 50 °C. Water extracts were prepared by placing 2 g of sample into 20 mL headspace vials with 10 mL of water, and keeping them in a conventional oven for 72 h at 50 °C.

GC-MS conditions

Column:	HP-5MS 30 m x 0.32 mm, 1.0 μm film
Carrier gas:	He at 2 mL/min
Temperature program:	35 °C for 5 min, 20 °C/min to 320 °C, hold 20.75 min
Injection port:	300 °C
Injection type:	1 μL splitless, 1 min purge
Makeup gas:	N ₂ at 400 mL/min
Transfer line:	350 °C

Scan range:	100 to 1500 <i>m/z</i>
Run time:	40 min
Data management:	MassLynx v4.1 Software
UPC ² Conditions	
System:	ACQUITY UPC ²
Detection:	Photodiode Array (PDA) Detector and SQD Mass Spectrometer
Column:	ACQUITY UPC ² BEH 2-EP 3.0 x 100 mm, 1.7 µm
Mobile phase A:	CO ₂
Mobile phase B:	1:1 methanol/acetonitrile
Flow rate:	2.0 mL/min
Gradient:	1% B for 1 min, to 20% over 2.5 min, hold for 30 s, re- equilibrate back to 1%
Column temp.:	65 °C
APBR:	1800 psi
Injection volume:	1.0 µL
Run time:	5.1 min

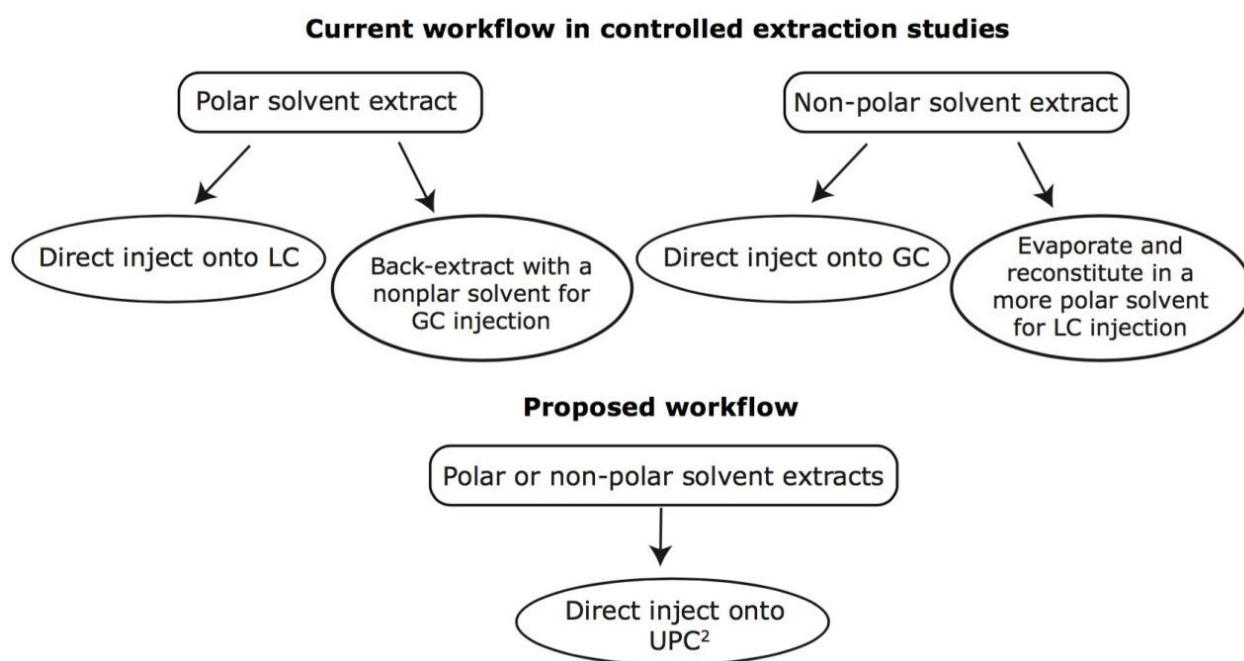
UPC² Conditions

Wavelength:	220 nm
MS scan range:	200 to 1200 <i>m/z</i>
Capillary:	3 kV
Cone:	25 V
MS make-up flow:	0.1% formic acid in methanol, 0.2 mL/min
Data management:	Empower 3 Software

UPLC conditions

System:	ACQUITY UPLC
Column:	ACQUITY UPLC BEH Phenyl 2.1 x 100 mm, 1.7 μ m
Mobile phase A:	0.1% formic acid in water
Mobile phase B:	0.1% formic acid in acetonitrile
Flow rate:	0.9 mL/min
Gradient:	50% B to 90% over 10 min, re- equilibrate back to 50% B
Column temp.:	50 °C
Injection volume:	2 μ L

Run time:	12 min
Wavelength:	220 nm
MS scan range:	200 to 1500 <i>m/z</i>
Cone:	30 V
Capillary:	3 kV
Data Management:	Empower 2 Software



Results and Discussion

The structures for polymer additives screened in this method are shown in Figure 1. They cover different classes of additives, such as plasticizers, antioxidants, and UV-absorbers.

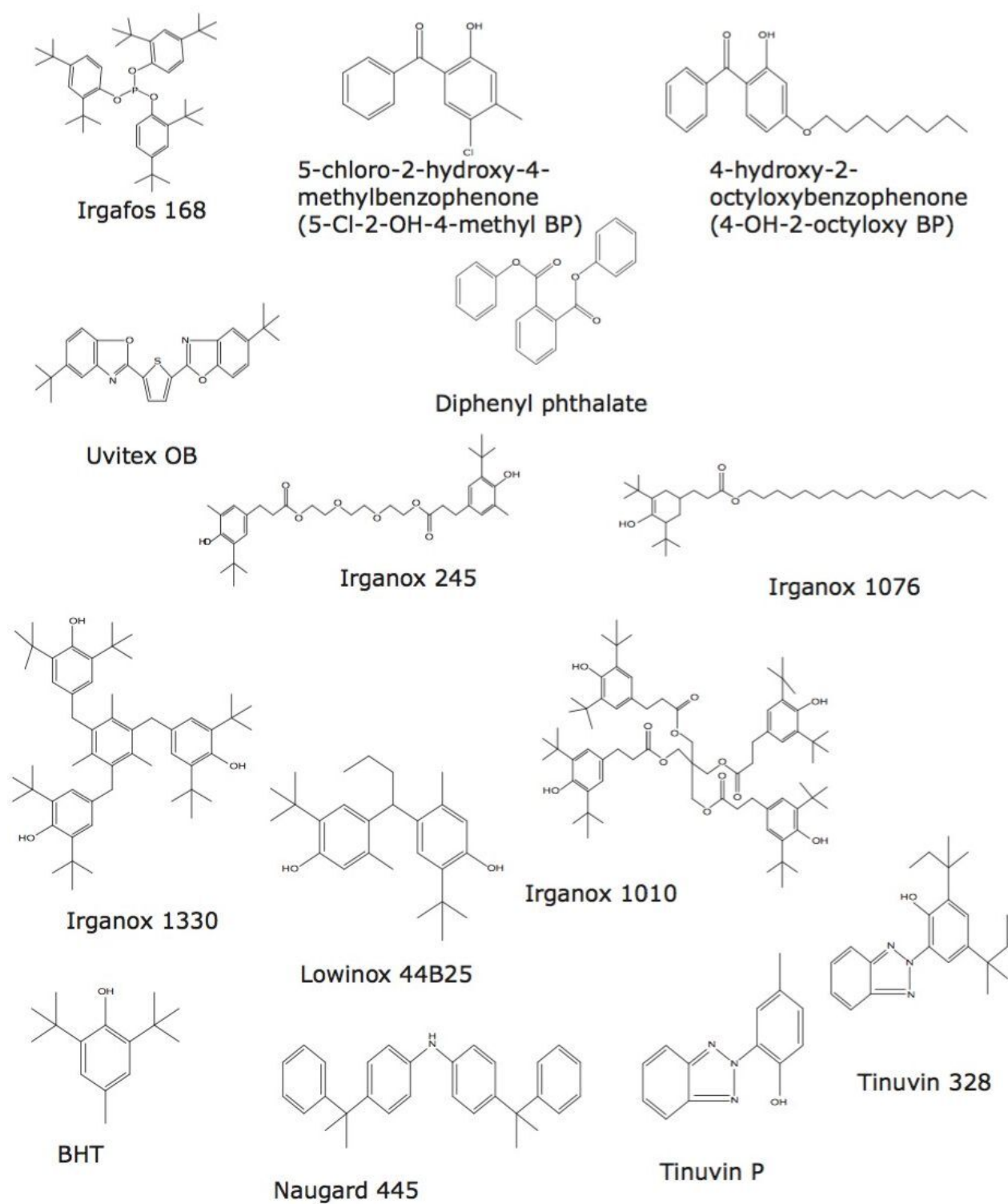


Figure 1. Polymer additives and their structures.

Comparing the separation of the standards by each analytical technique, as shown in Figure 2, UPLC and

UPC² were applicable to all 14 compounds chosen. The elution order was different for both methods due to orthogonal selectivity. The ACQUITY UPC² System provided a shorter run time compared to the ACQUITY UPLC System. It was observed that the thermal instability of some analytes, such as Irganox 1010 and Irganox 245, prevented successful chromatographic separation by GC-MS. Late eluters from Irgafos 168 to Uvitex OB produced wide peaks in GC-MS, possibly due to secondary interactions with the stationary phase or on-column degradation. The compounds selected for this screening were more compatible with liquid chromatography or convergence chromatography than with gas chromatography analysis.

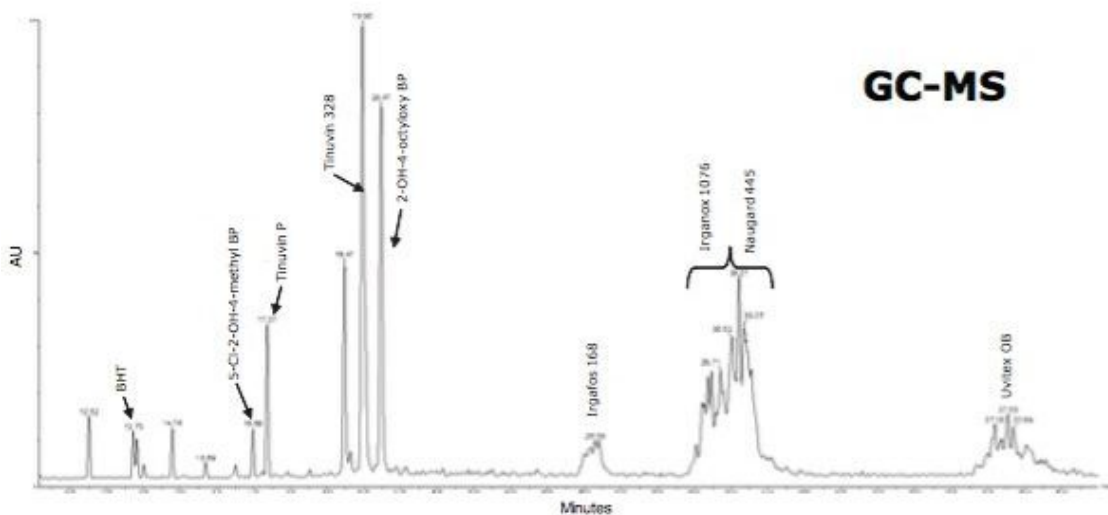
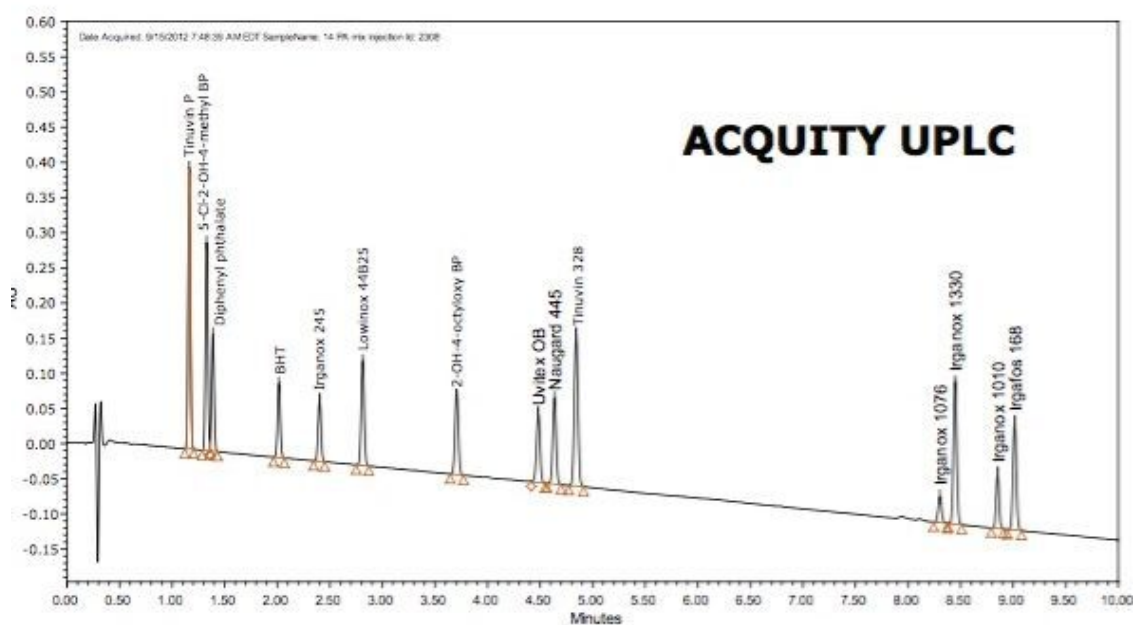
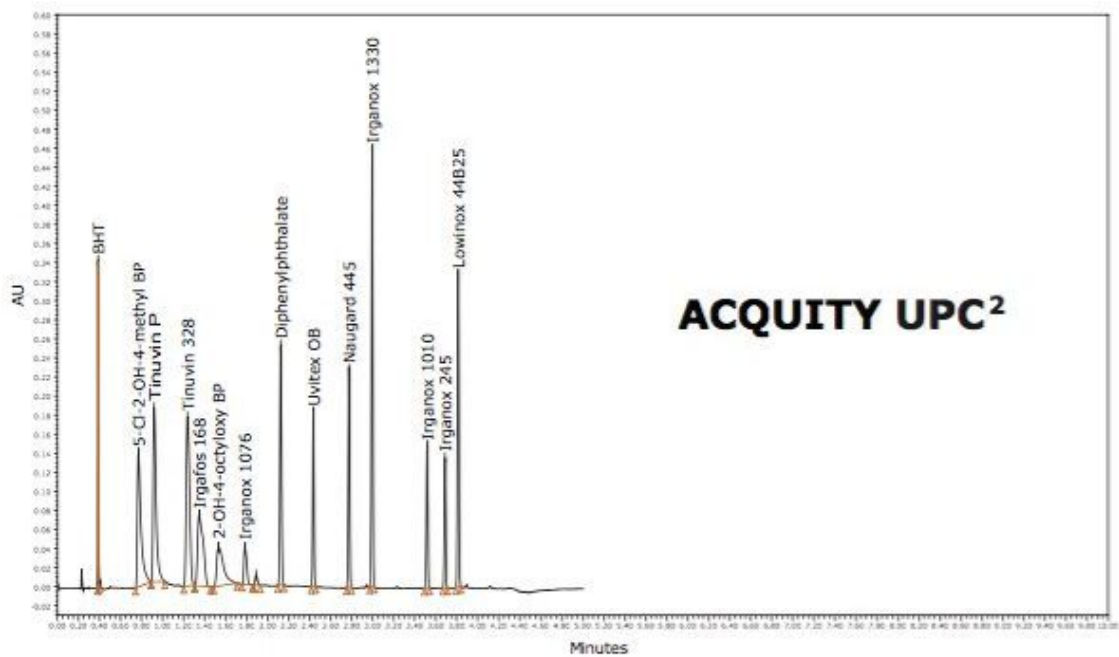


Figure 2. Chromatograms for standards separation.

systems did not have any peaks present (data not shown). This was expected, since water is the most common solvent present in the environment. Manufacturers avoid formulating their products to be susceptible to water solubility.

In the other two extracts, hexane and IPA, LDPE had the most extractables present, as seen in Figure 3. IPA extracts analyzed by UPLC (data not shown) produced less intense peaks than UPC². Prior to UPLC analysis, the hexane extracts were reduced to dryness, re-dissolved in solvent, and analyzed by UPLC (data not shown). Both the ACQUITY UPLC and ACQUITY UPC² systems showed the same set of extractable compounds present in the samples.

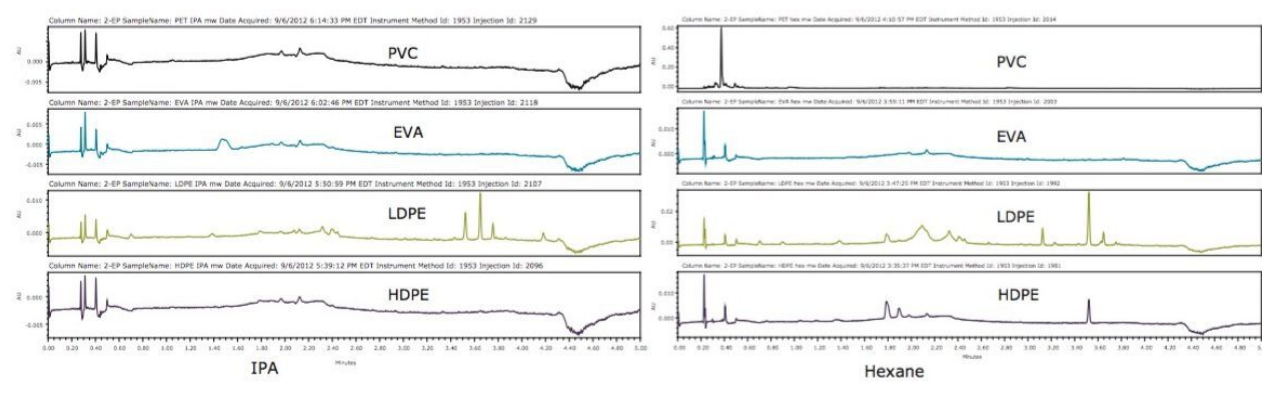


Figure 3. ACQUITY UPC² System chromatograms for IPA and hexane sample extracts.

Noisy baselines were observed with the GC-MS analysis. When utilizing this technique, extracted ion chromatograms of known polymers had to be performed, thus making it difficult to screen for unknown extractables in packaging products, as shown in Figure 4. A sample pre-concentration step could have improved the intensity of the detected peaks.

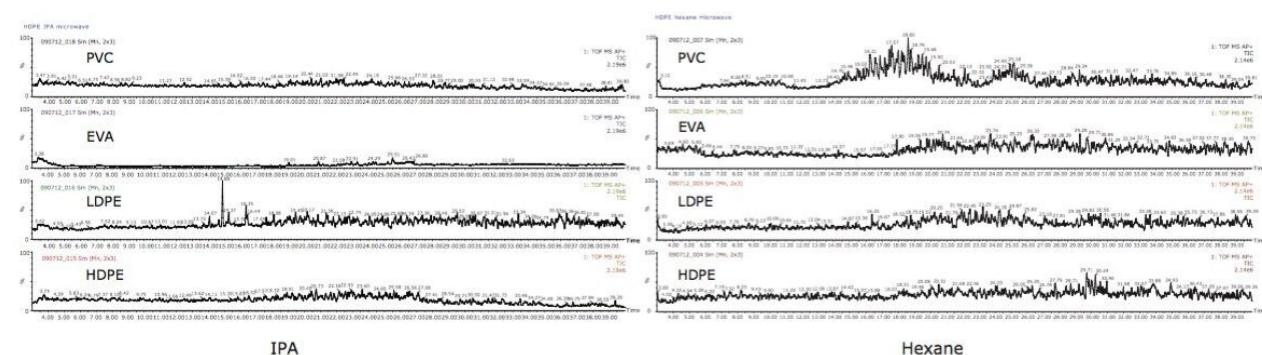


Figure 4. GC-MS chromatograms for IPA and hexane extracts.

Three known polymer additives were identified in LDPE samples by ACQUITY UPC², including Irganox 1010, Irganox 1076, and Irgafos 168, as shown in Figure 5. These are commonly used antioxidants that improve the stability of polymers. The identity of each extractable was confirmed by injection of authentic standards, comparison of the retention time, and MS data. An example for Irganox 1076 is shown in Figures 6 and 7. Each of these additives was detected in either hexane or isopropanol extracts of LDPE.

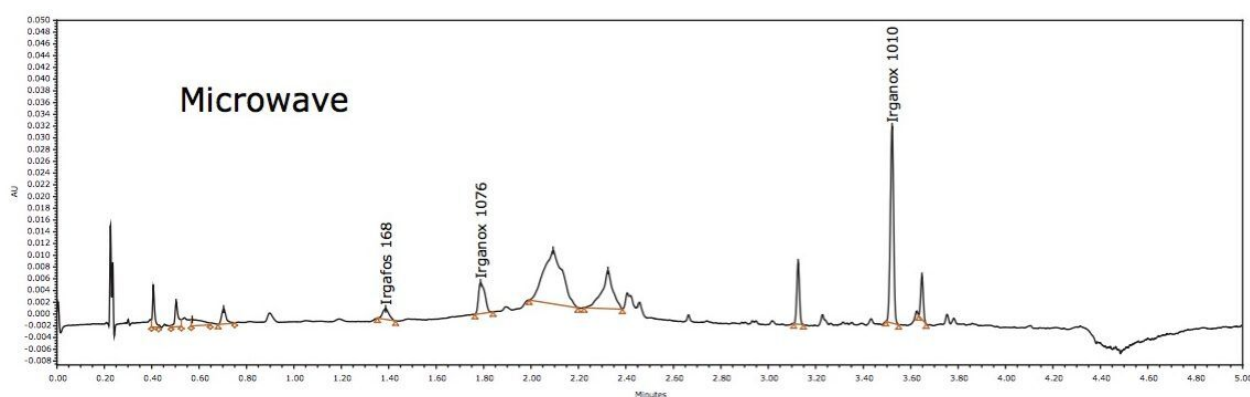


Figure 5. Identified extractables in LDPE hexane extract using ACQUITY UPC².

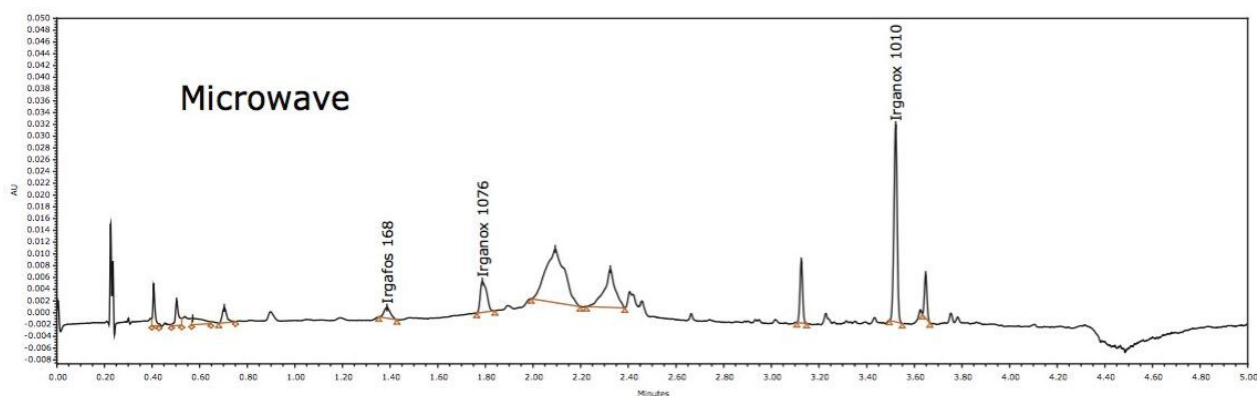


Figure 6. Irganox 1076 in LDPE hexane extract by UPC².

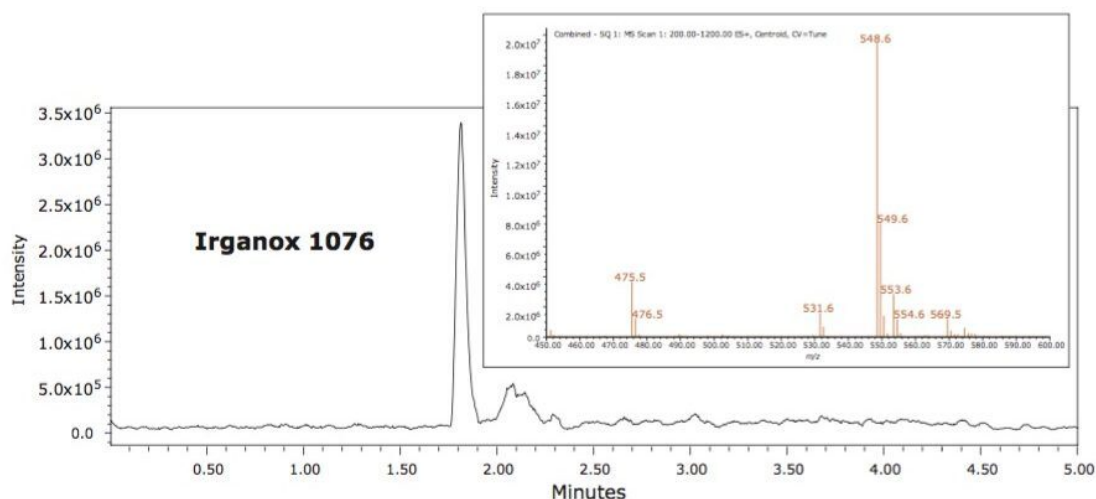


Figure 7. Irganox 1076 standard by UPC².

In GC-MS analysis, the presence of Irgafos 168 and Irganox 1076 was also confirmed using standard retention time and mass spectra.

Conclusion

In this application, a single technique was found to be compatible for all extracts of different packaging material. This capability allowed for a streamlined, simplified sample preparation workflow with better asset utilization, since all of the solvent extracts can be directly injected onto the ACQUITY UPC² System. Using other separation techniques, such as LC and GC, some extracts are not compatible requiring additional processing steps prior to analysis.

UPC² offered better information for non-volatile and thermally labile compounds than GC due to lower analysis temperatures. The UPC² analysis provided a two-fold improvement in run time compared to UPLC, and an eight-fold improvement in run time compared to GC.

The ease-of-use coupled with the MS detector provided quick polymer identification for known entities in the sample extracts.

References

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3. Norwood DL, Fenge Q. Strategies for the analysis of pharmaceutical excipients and their trace level impurities. *Am Pharm Rev*. 2004; 7(5): 92,94, 96-99.

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720004490, November 2012