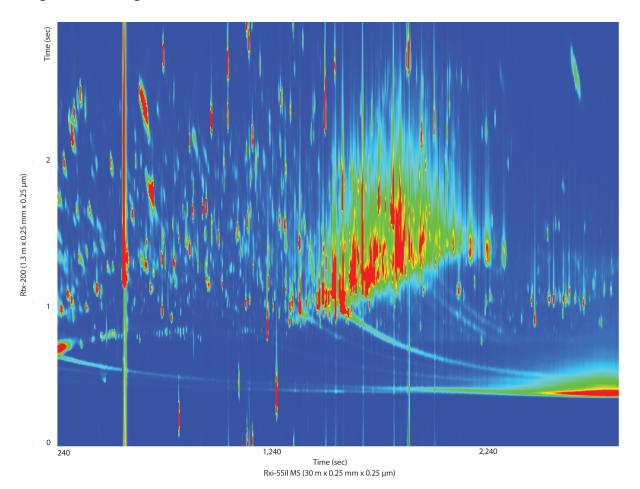
## GCxGC-TOFMS Analysis of Unfortified Tobacco Extract After QuEChERS Extraction and dSPE Cleanup With 25 mg PSA and 7.5 mg GCB



GC\_FF1242

Notes

Column Rxi-5Sil MS 30 m, 0.25 mm ID, 0.25 µm (cat.# 13623) Rtx-200 1.3 m, 0.25 mm ID, 0.25  $\mu$ m (cat.# 15124) Standard/Sample

Tobacco extract with dSPE cleanup Injection Inj. Vol.: 1.0 µL splitless (hold 1.0 min)

Liner: Premium 4 mm single taper w/wool (cat.# 23303)

Ini. Temp.: 250°C Oven

Oven Temp.: Rxi-5Sil MS: 90 °C (hold 1.0 min) to 310 °C at 5 °C/min (hold 2.0 min) Rtx-200: 95 °C (hold 1.0 min) to 315 °C at 5 °C/min (hold 2.0 min) He, corrected constant flow (2 mL/min) **Carrier Gas** 

Modulation Modulator Temp. Offset: +20 °C Second Dimension Separation Time: 3 sec Hot Pulse Time: 0.9 sec Cool Time between Stages: 0.6 sec Detector MS Mode:

Transfer Line Temp.: 300°C Analyzer Type: TOF 225°C Source Temp.: Electron Energy: 70 eV -20 mu/100 u Mass Defect:

Ionization Mode:

**Acquisition Range:** Spectral Acquisition Rate: Instrument

Sample Preparation

LECO Pegasus 4D GCxGC-TOFMS

QuEChERS Extraction:

45 to 550 amu

100 spectra/sec

A 2 g sample of tobacco was weighed into a 50 mL polypropylene centrifuge tube (cat.# 26239). After the addition of 10 mL of organic-free water to the sample, 100 µL of QuEChERS internal standard mix for GC-MS analysis (cat.# 33267) was added to each sample. Next, 10 mL of acetonitrile was added and the samples were vortexed for 30 min. Immediately after vortexing, pre-packaged QuEChERS European EN 15662 method formulation extraction salts (cat.# 26236) were added to each centrifuge tube. The tubes were immediately shaken for 1 min and then centrifuged for 5 min at 3000 g.

Sample Cleanup:

A 1 mL aliquot of the extract was fortified with 5 µL of an anthracene standard (cat.# 33264) and added to a dSPE tube containing 150 mg MgSO<sub>4</sub>, 25 mg PSA and 7.5 mg GCB (cat.# 26218). The tubes were gently shaken for 2 min and then centrifuged for 5 min using a Q-sep 3000 centrifuge (cat.# 26230). A 0.5 mL portion of the supernatant extract was removed and placed into an autosampler vial and 5  $\mu$ L of a 5% formic acid solution in acetonitrile was added to each sample prior to analysis. Rtx-200 (cat.# 15124) is a 2 m column. A 1.3 m section was cut off and used

as the second dimension column.



