

# Exploration of New Low-Pressure GC Columns for Food and Environmental Emerging Contaminants

Jessi Collier<sup>1</sup>, Jana Hepner<sup>1</sup>, Whitney Dudek-Salisbury<sup>1</sup>, Kristi Sellers<sup>1</sup>, Jaap de Zeeuw<sup>1</sup>, Chris English<sup>1</sup>

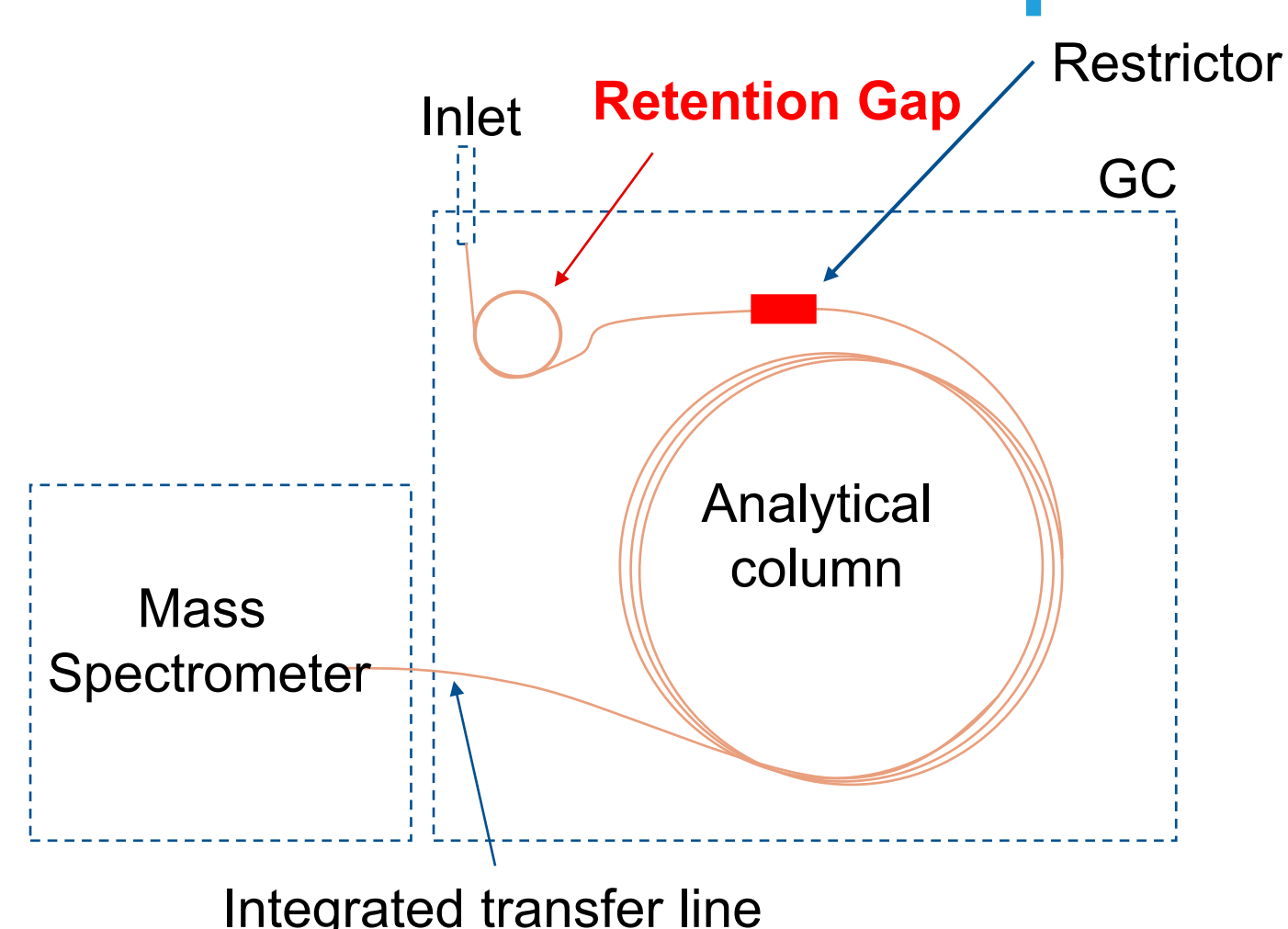
<sup>1</sup>Restek Corporation, 110 Benner Circle; Bellefonte, PA 16823, USA; [jessi.collier@restek.com](mailto:jessi.collier@restek.com)

## Introduction

The Low-Pressure GC (LPGC technique) has been successfully used in the past for pesticide residues' analysis. However, the technique is very versatile, and it allows for other applications, especially if different column phases are used. So far, the majority of the applications have been using the "5"-type phase (95% dimethylpolysiloxane, 5% diphenyl polymer). To expand on the previous applications, four additional column phases were selected (cyanopropylphenyl dimethylpolysiloxane; 50% dimethylsiloxane, 50% diphenyl; 65% dimethylsiloxane, 35% diphenyl; and trifluoropropylmethyl polysiloxane phases) to analyze various food and environmental contaminants, such as nitrosamines, alkylfurans, phthalates, arylamines and fluorotelomer alcohols.

The LPGC techniques provided significant reduction in run times (up to 3.3x faster runs) and helium consumption reduction (up to 81% less helium used), while keeping an acceptable resolution.

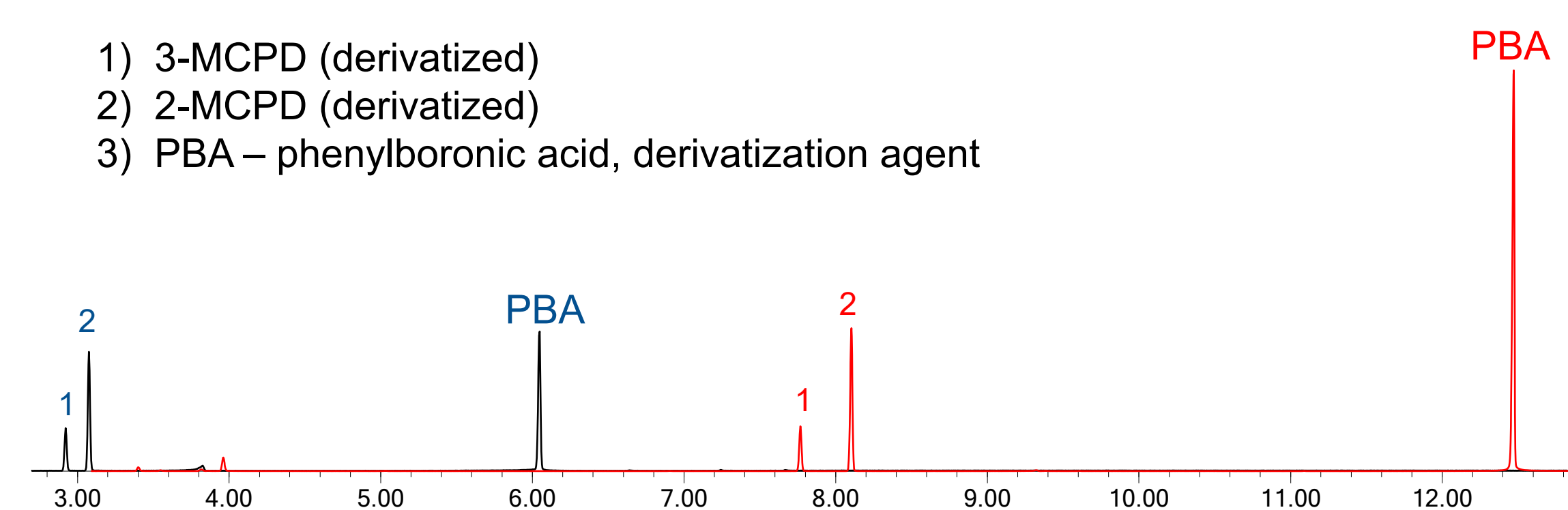
## LPGC setup



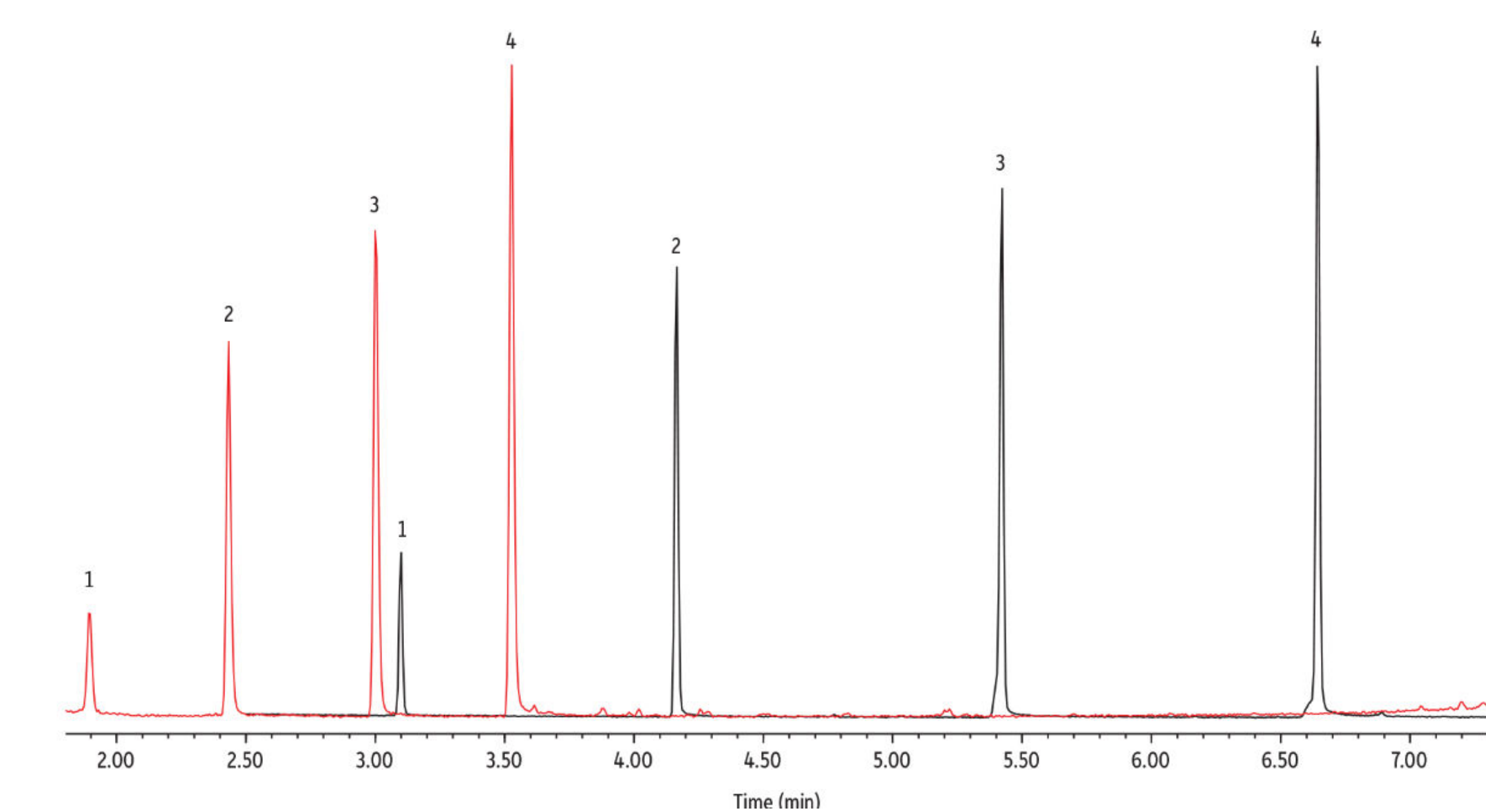
Column set is delivered pre-connected in the box  
Only extra consumable needed is 0.8 mm vespel/graphite ferrule for MS transfer line

## LPGC Rxi-17Sil MS Application - MCPDs

- 1) 3-MCPD (derivatized)
- 2) 2-MCPD (derivatized)
- 3) PBA – phenylboronic acid, derivatization agent



## LPGC Rtx-200 – Fluorotelomer Alcohols (FTOHs)



- 1) 4:2 FTOH (2-perfluorobutyl alcohol)
- 2) 6:2 FTOH (2-perfluorohexyl alcohol)
- 3) 8:2 FTOH (2-perfluorooctyl alcohol)
- 4) 10:2 FTOH (2-perfluorodecyl alcohol)

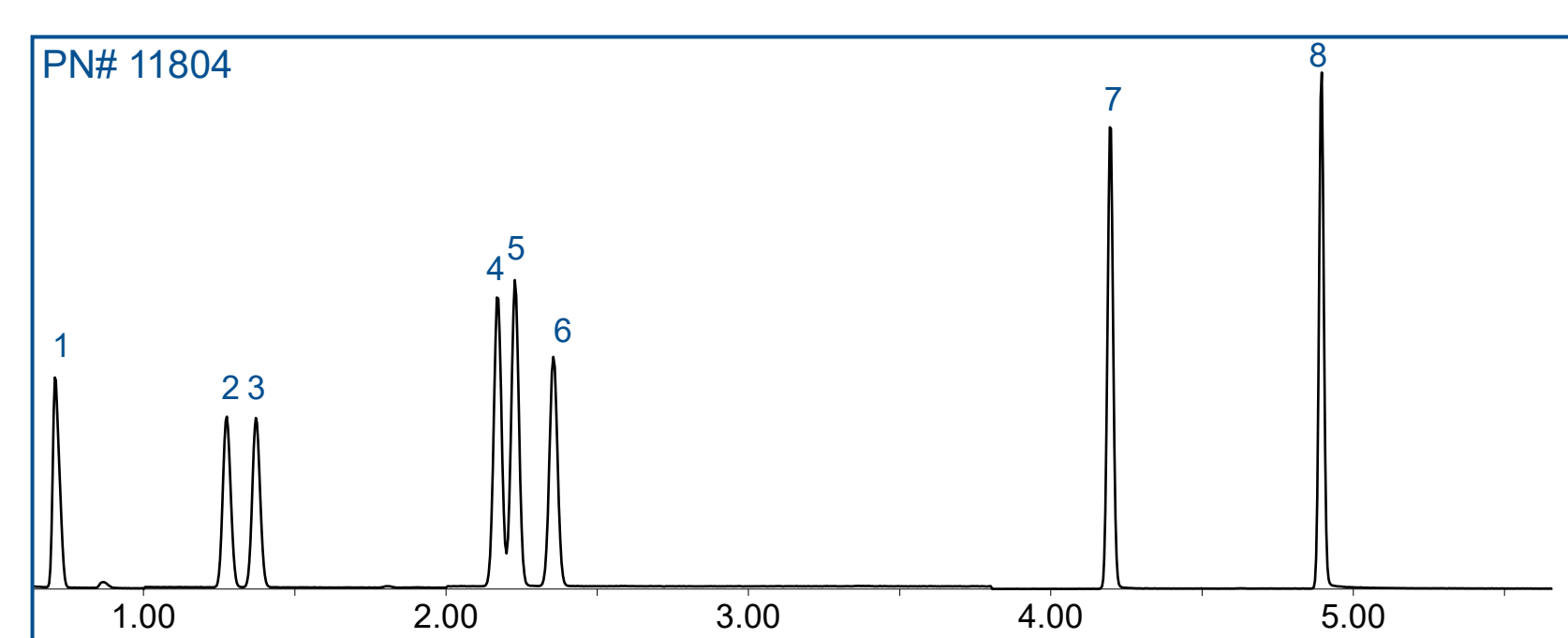
## Conclusions

- **LPGC offering has been expanded**
  - Analysis up to 3.3x faster
  - Limited by the temperature program
  - Up to 81% less helium used
- **No need to change instrumentation**
  - Now compatible with all instruments

## Advantages of LPGC

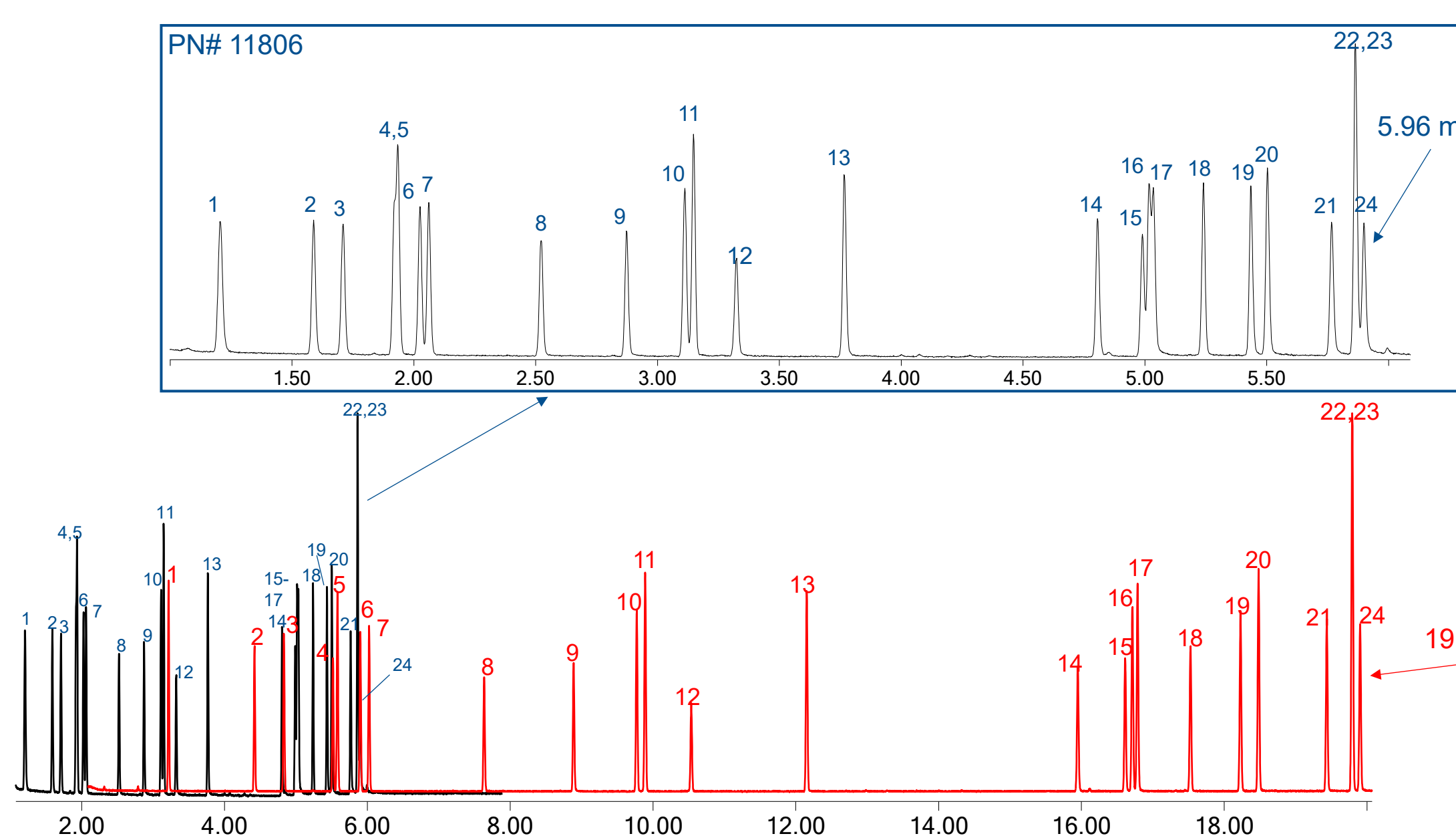
- **Fast analysis with short 0.53 mm or 0.32 mm capillaries**
  - Short analysis times
  - Increased sensitivity
  - Potentially higher capacity
- **Peak width enough for any type of MS**
- **Lower elution temperatures**
- **Elution at 50-80°C lower temperatures**
  - Lower bleed
  - Standard injection techniques, high volume injections
- **All stationary phase chemistries**

## LPGC Rxi-624Sil MS Application #2 - Alkylfurans



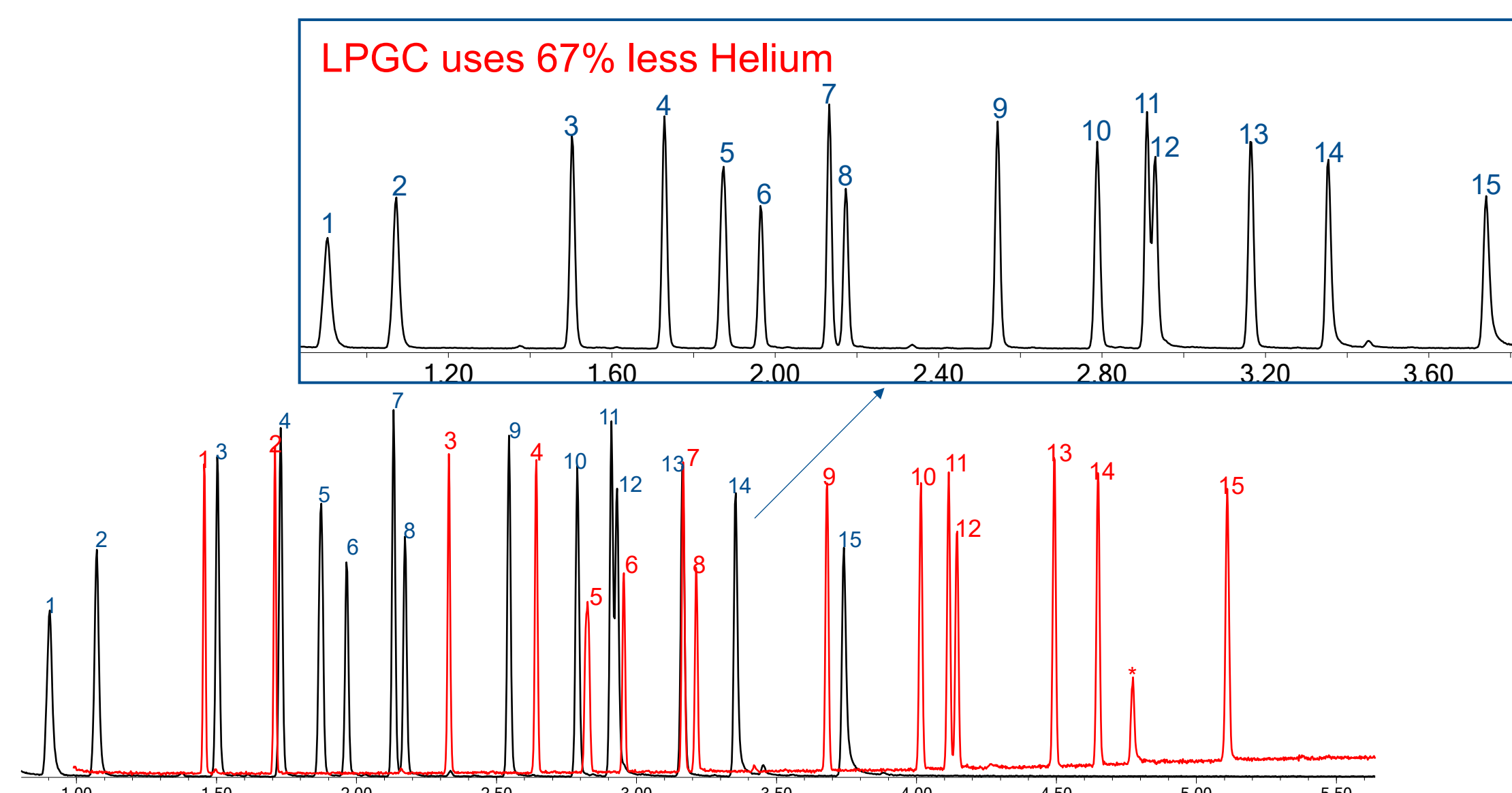
- 1) Furan
- 2) 2-methylfuran
- 3) 3-methylfuran
- 4) 2-ethylfuran
- 5) 2,5-dimethylfuran
- 6) 2,3-dimethylfuran
- 7) 2-butylfuran
- 8) 2-pentylfuran

## LPGC Rxi-35Sil MS Application #1 - Arylamines



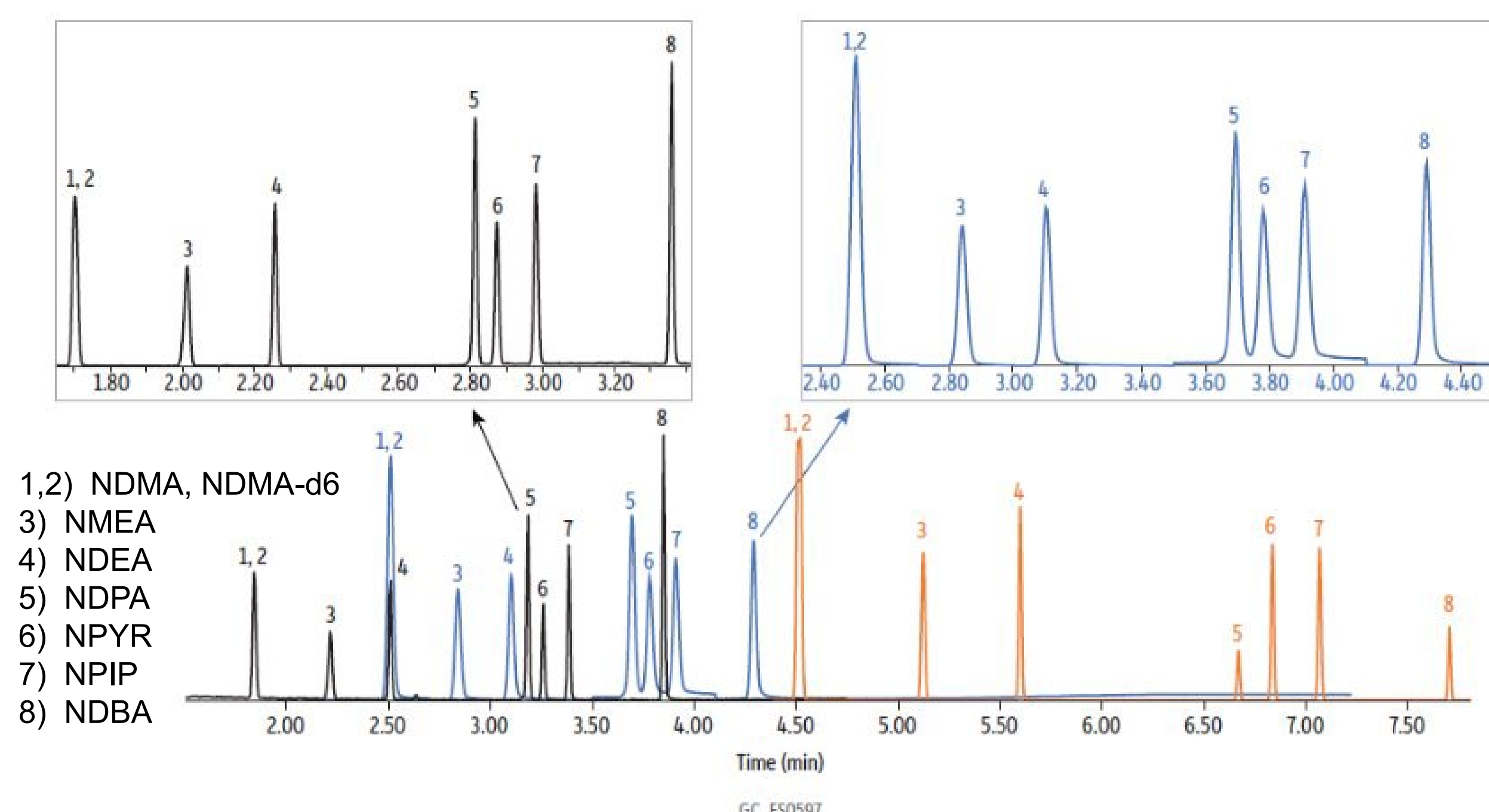
- 1) o-Toluidine
- 2) o-Anisidine
- 3) 4-Chloroaniline
- 4) p-Cresidine
- 5) 2,4,5-Trimethylaniline
- 6) 3-Chloro-o-toluidine
- 7) 4-Chloro-o-toluidine
- 8) 2,4-Diaminotoluene
- 9) 2,4-Diaminoanisole
- 10) 2-Naphthylamine
- 11) 2-Aminobiphenyl
- 12) 2-Amino-4-nitrotoluene
- 13) 4-Aminobiphenyl
- 14) p-Aminoazobenzene
- 15) 4,4'-Oxydianiline
- 16) 4,4'-Diaminodiphenylmethane
- 17) Benzidine
- 18) o-Aminoazotoluene
- 19) 3,3'-Dimethyl-4,4'-diaminodiphenylmethane
- 20) 3,3'-Dimethylbenzidine
- 21) 4,4'-Thiodianiline
- 22) 3,3'-Dichlorobenzidine
- 23) 4,4'-Methylenebis(2-chloroaniline)
- 24) 3,3'-Dimethoxybenzidine

## LPGC Rxi-35Sil MS Application #2 - Phthalates



- 1) Dimethyl phthalate
- 2) Diethyl phthalate
- 3) Diisobutyl phthalate
- 4) Di-n-butyl phthalate
- 5) Bis(2-methoxyethyl) phthalate
- 6) Bis[4-methyl-2-pentyl] phthalate isomers
- 7) Di-n-pentyl phthalate
- 8) Bis(2-ethoxyethyl) phthalate
- 9) Di-n-hexyl phthalate
- 10) Butyl benzyl phthalate
- 11) Bis(2-ethylhexyl) phthalate
- 12) Bis(2-butoxyethyl) phthalate
- 13) Dicyclohexyl phthalate
- 14) Di-n-octyl phthalate
- 15) Dinonyl phthalate

## LPGC Rxi-624Sil MS Application #1 - Nitrosamines



## Limitations of LPGC Strategies for Addressing Shortcomings

- **Loss of theoretical plates (compared to conventional column)**
  - Can be mitigated by selective detection by MS
- **Greater potential for leaks**
  - Pre-connected set
- **More complicated to cut analytical column**
  - Less need to cut column
- **Need for MS instrument**
- **Fast oven heating needed**
  - 120 V instruments might need an accelerator

## Can I use LPGC for my application?

- **Is MS a suitable detector?**
  - Vacuum is needed
- **What kind of column is the conventional method using?**
  - In general, most conventional methods with 30 m column can be translated to LPGC method
- **Are there isobars to resolve?**
  - What level of resolution is needed?
  - What is the resolution in the conventional method?
  - If it is above 2.0, usually we can get peaks resolved

## Large Volume Injections

- GC oven set to temperature below the boiling point of the solvent
- Fast injection with liquid band formation
- Liquid sample is deposited on glass wool
- Pressure surge from evaporating solvent "pumps" sample into retention gap
- Most of solvent goes into retention gap

- Lower detection limits
- Saves time in sample prep
- Eliminate need for expensive PTV
- Large volume splitless injection needs:
  - A splitless injection device
  - A liner with glass wool
  - A retention gap that has to be coupled

## Peak shapes at 5-25 µL injections

### Methyl Pirimiphos

