

Phenols

Analysis of urinary phenols as acetate esters

Application Note

BioPharma

Authors

Agilent Technologies, Inc.

Introduction

Gas chromatography with an Agilent CP-Sil 8 CB column separates 11 urinary phenols as acetate esters in 15 minutes.

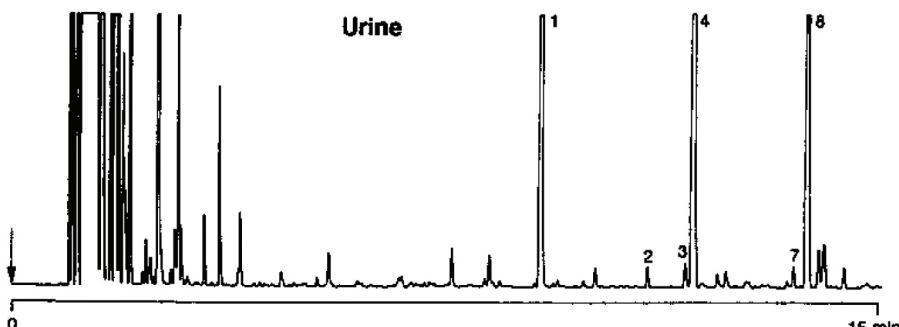


Agilent Technologies

Conditions

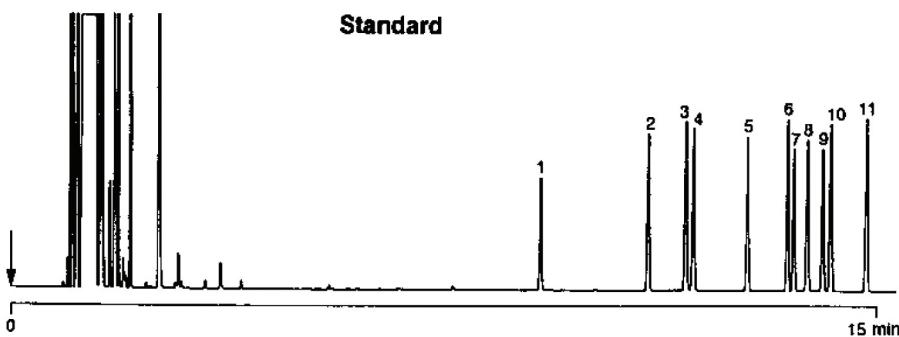
Technique : GC-capillary
Column : Agilent CP-Sil 8 CB, 0.32 mm x 25 m fused silica
Temperature : 50 °C - 140 °C, 6 °C/min
Carrier Gas : H₂, 45 kPa (0.45 bar, 6.4 psi)
Injector : Splitter, 50 mL/min
T = 250 °C
Detector : FID
T = 300 °C
Sample Size : 2 µL
Concentration Range : 50 ppm each component
Solvent Sample : hexane

Courtesy : Dr. C. Weber,
National Institute of
Occupational Health,
Budapest, Hungary



Peak identification as acetate esters

1. phenol
2. o-cresol
3. m-cresol
4. p-cresol
5. 2,6-dimethylphenol (2,6 Xylenol)
6. 2,5-dimethylphenol (2,5 Xylenol)
7. 2,4-dimethylphenol (2,4 Xylenol)
8. 3-ethyl phenol (I.S.)
9. 3,5-dimethylphenol (3,5 Xylenol)
10. 2,3-dimethylphenol (2,3 Xylenol)
11. 3,4-dimethylphenol (3,4 Xylenol)



Sample preparation

The urinary phenols are conjugated, thus they were first hydrolysed enzymatically, steam-distilled, buffered with H₃BO₃/NaOH and acetylated with acetic acid anhydride. The calibration standard contains the following phenols at the 50 ppm concentration level each.

www.agilent.com/chem

This information is subject to change without notice.

© Agilent Technologies, Inc. 2011

Printed in the USA

31 October, 2011

First published prior to 11 May, 2010

A00711



Agilent Technologies