



# **B-Adrenoceptor-blocking drugs**

## Analysis of esmolol and flumolol in whole blood

### Application Note

Clinical Research

#### **Authors**

Agilent Technologies, Inc.

#### **Introduction**

According to K. Kylberg-Hanssen et al, sodium dodecyl sulfate (SDS) is an effective esterase inhibitor in the determination of labile  $\beta$ -blockers: esmolol and flumolol. The limit of detection (coefficient of variation: 10 - 15%) is 5 nmol/L of whole blood.



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## Conditions

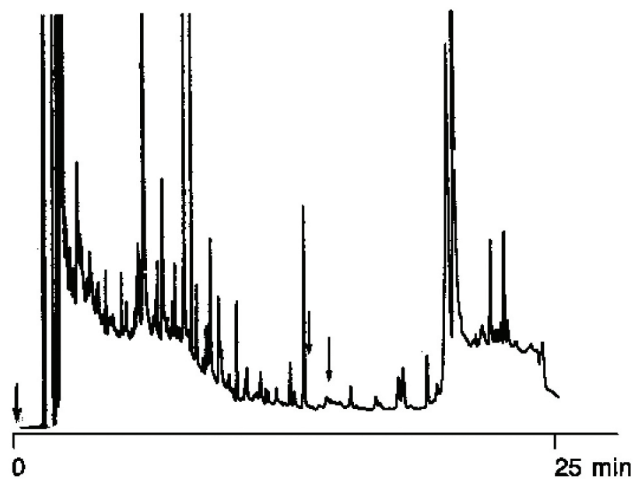
Technique : GC-capillary  
Column : Agilent CP-Sil 8 CB, 0.32 mm x 25 m fused silica  
WCOT CP-Sil 8 CB (0.12  $\mu$ m) (Part no. CP7741)  
Temperature : esmolol determination:  
100 °C (1 min)  $\rightarrow$  185 °C 10 °C/min;  
185 °C (15 min)  $\rightarrow$  250 °C, 30 °C/min;  
250 °C (7 min)  
flumolol determination:  
100 °C (1 min)  $\rightarrow$  200 °C, 15 °C/min;  
200 °C (10 min)  $\rightarrow$  250 °C, 30 °C/min;  
250 °C (7 min)  
Carrier Gas : He, 60 kPa  
Injector : Splitter, 20 mL/min  
T = 270 °C  
Detector : ECD  
T = 350 °C

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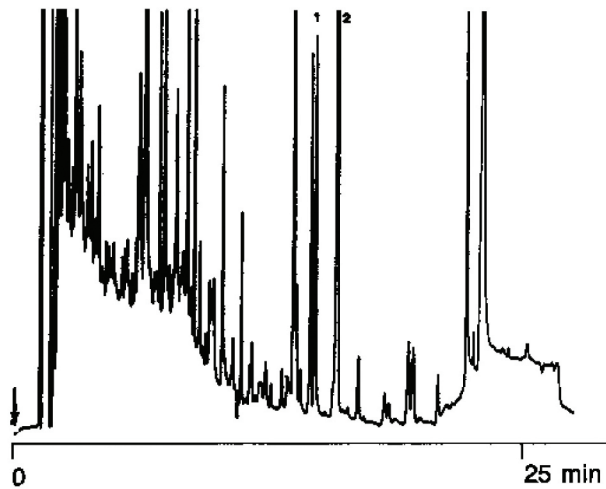
## Peak identification

1. flumolol (100 nmol/L)
2. Int. Standard (H 163/37) (500 nmol/L)

*Blood sample*



*Blood sample with added flumolol*



## Analytical procedure

Collect 5 mL of blood in a tube containing 0.5 mL SDS solution (200 g/L). After mixing transfer 1.0 g of the sample to a 15 mL centrifuge tube. Add 100  $\mu$ L of the internal standard solution (5  $\mu$ mol in phosphate buffer, pH = 6.0, 0.1 mol/L). Add 4 mL of toluene and 0.5 mL of phosphate buffer (pH = 11.8, 0.2 mol/L). Shake (15 min) and centrifuge. Immerse the tube in an acetone/solid CO<sub>2</sub> mixture to freeze the aqueous layer.

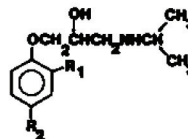
Pour the organic layer into a 5 mL centrifuge tube and evaporate under a stream of N<sub>2</sub> at 35 °C. Redissolve the residue in 100  $\mu$ L of ethylacetate and 25  $\mu$ L of pentafluoropropionic anhydride. After reacting for 20 min at room temperature, evaporate under nitrogen at 35 °C. Dissolve the residue in toluene (500  $\mu$ L) and inject 1.5  $\mu$ L.

## Acknowledgement

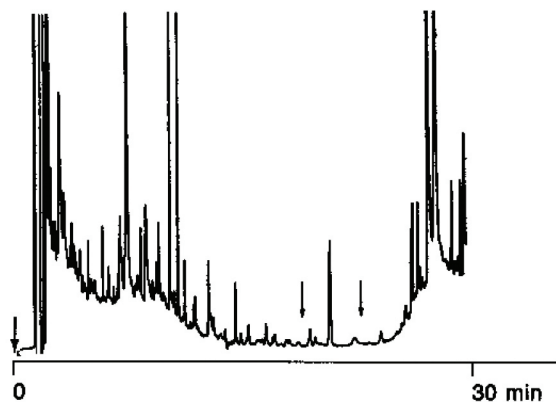
Agilent Technologies thanks Kerstin Kylberg-Hanssen, Department of Bioanalytical Chemistry, AB Hässle, S-431 83 Mölndal, Sweden for all technical information and the chromatogram.

## Peak identification

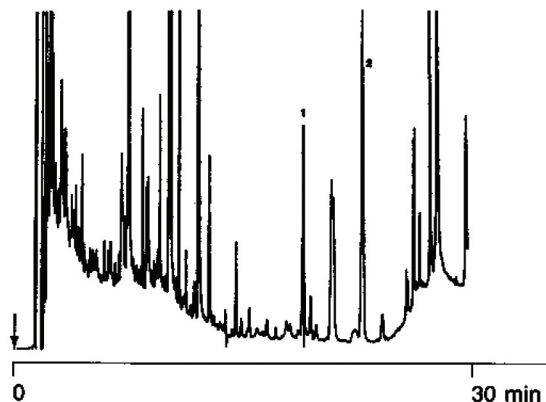
1. esmolol (300 nmol/L)
2. Int. Standard (H 163/37) (500 nmol/L)



*Blood sample*



*Blood sample with added esmolol*



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