

# EXHALED BREATH MEASUREMENT FOR BIOLOGICAL MONITORING OF ORGANIC COMPOUNDS FROM ENVIRONMENTAL EXPOSURES

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## INTRODUCTION

The organic constituents of exhaled human breath are representative of blood concentrations through gas exchange in the blood-breath interface in the lungs. The presence of specific compounds can be an indicator of recent exposure or biological response of the subject. The analysis of breath is preferred to direct measurement from blood samples because breath collection is a non-invasive procedure and the measurement of gas-phase analytes is much simpler in a gas matrix than in a complex biological tissue like blood. Between the several types of exhaled breath samples used, ambient air (and exhaled breath) sampling permits valid estimation of partial compound pressure in arterial blood; additionally, this technique can be used as an indicator of the partial solvent pressure of mixed venous blood if the samples are suitably obtained.

In this study we present the results of biological monitoring, by means of exhaled breath, of n-hexane, toluene and styrene in several industry workers, and of isoflurane (2-chloro,2,2,2-trifluoroethyl difluoromethyl ether) in anaesthetists, surgeons, and nurses in operating theatres.

## EXPERIMENTAL

### Biological sampling

A portable system for end-exhaled breath sampling shown in figure 1 was used. The system is based in a Haldane-Priestley tube modified to concentrate aliquots of end-exhaled air from one or more exhalations into a 60/80-m cartridge, has been previously designed and validated in our laboratory [1]. Activated charcoal for solvent desorption and Chromosorb 106 or Tenax for thermal desorption were used as sorbents. The samples were taken at the end of shift.



Fig. 1. Schematic diagram of the system of sampler.

1. End-exhaled breath
2. Haldane-Priestley tube
3. Sorbent cartridge
4. Sampling pump
5. Sampling valve
6. Sampling container
7. Sampling pump
8. Sampling valve



Fig. 2. Diffusive sampler for thermal desorption (ATD-50) and diffusive sampler for solvent desorption (SM-2000).

### Ambient air sampling

For each occupationally exposed subject, an evaluation of environmental level of these compounds was carried out during the shift, by means of personal diffusive samplers attached to the clothing within the breathing zone during the exposure. Diffusive samplers for thermal and solvent desorption (figure 2), were used. Experimental conditions for biological and environmental sampling are shown in table I.

### Analysis

For thermal desorption, direct analysis by means of ATD-50 system gas chromatograph, equipped with a 25-m FFAP capillary column and FID, was used.

For solvent desorption, after desorption with  $CS_2$ , an aliquot was injected into gas chromatograph equipped with a 5-m column packed with 10% FFAP on Chromosorb W-AW.

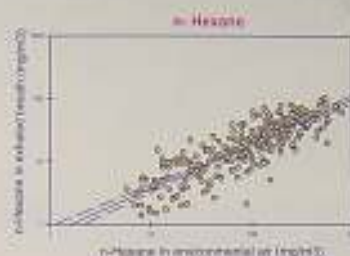


Figure 3

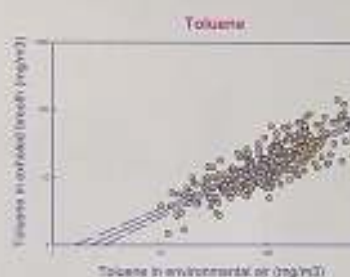


Figure 4

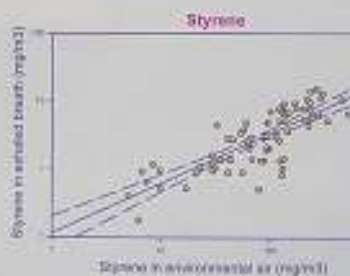


Figure 5

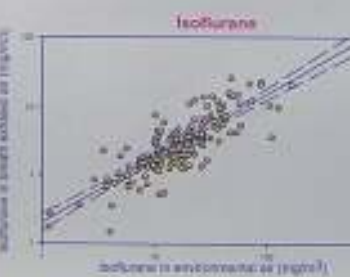


Figure 6

TABLE II  
Results of analyses in environmental air and exhaled breath and parameters of the regression equation  $\log Y = a + b \log X$

Compound	CONCENTRATION (ng/ml)		REGRESSION PARAMETERS				
	n	range	a	b	r	s	
n-Hexane	234	10 (10.0)	28 (137.1)	0.92	0.70	0.88	0.001
Toluene	205	171 (873.1)	15 (254.7)	0.90	0.70	0.81	0.000
Styrene	80	77 (329.8)	4.7 (24.02)	0.86	0.70	0.86	0.001
Isoflurane	118	37 (137.7)	47 (263.7)	0.88	0.86	0.81	0.001

TABLE I  
Experimental conditions for biological and environmental sampling

Compound	SAMPLING	SAMPLING	BIOLOGICAL		ENVIRONMENTAL	
			DESORPTION	DESORPTION	DESORPTION	DESORPTION
n-Hexane	1.0h	Active Charcoal	$CS_2$	SM-2000	$CS_2$	2
Toluene	1.0h	Active Charcoal	$CS_2$	SM-2000	$CS_2$	2
Styrene	1.0h	Tenax	Thermal	ATD-50	$CS_2$	1.4
Isoflurane	1.0h	Chromosorb 106	Thermal	ATD-50	Thermal	1

## RESULTS AND DISCUSSION

Figures 3 to 6 show the correlation found between environmental and biological results. We have studied 234 subjects exposed to n-hexane, 205 to toluene, 80 to styrene and 178 to isoflurane. The parameters of the regression equation are shown in table II. From the good correlations between exhalatory concentrations and personal air samples, the concentration in ambient air can be predicted from the exhaled air concentration and vice versa. Thus, Threshold Limit Values in ambient air have their equivalent concentration in the exhaled air.

In conclusion our results show that the proposed method may be useful for the biological monitoring of environmental exposures to volatile organic compounds.

## REFERENCES

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