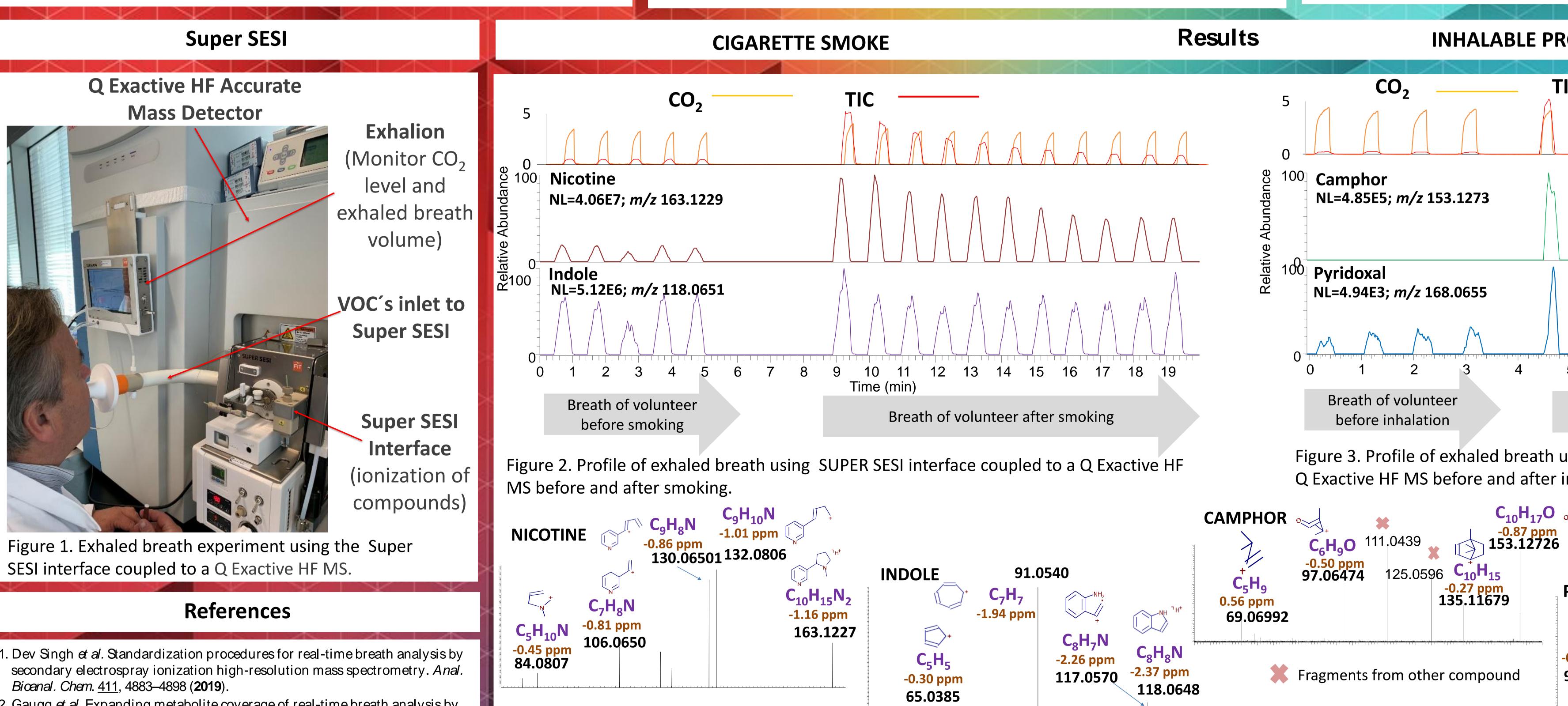


Introduction and Objectives

VOCs in breath are produced either by various biochemical processes within the body or as a result of external factors such as environmental exposure, lifestyle, diet, and/or therapeutic interventions. Real-time breath analysis is an advantageous analytical approach by which information about physiological changes over a short period of time can be obtained. Real-time analysis of human exhaled breath enables rapid monitoring of exposure-driven absorption of exogenous VOCs from the lungs into the bloodstream.

The aim of this study was to detect, confirm and monitor absorption of exogenous compounds originating from cigarette smoke and various inhalable products from the lungs into the bloodstream.



- 2. Gaugg *et al.* Expanding metabolite coverage of real-time breath analysis by coupling a universal secondary electrospray ionization source and high resolution mass spectrometry--a pilot study on tobacco smokers. J. Breath Res. <u>10,</u> 1, 016010 (**2016**).
- 3. Zivkovic Semren et al. Monitoring of metabolite kinetics of tobacco users by real-time exhaled breath analysis. Application note (2020)

ASMS Conference | June 1 – 12, 2020

Monitoring of exogenous compound kinetics in exhaled breath

Methods

Human exhaled breath samples were analyzed with an Exhalion Super SESI coupled to a Q Exactive HF mass spectrometer. The system measures CO₂ levels (%), pressure drop (mbar), exhalation flow rate (L/min), and total exhaled volume (L) in real-time. **Compounds present in exhaled breath are ionized by the Super SESI** interface and detected by high-resolution MS.

Human volunteers exhaled before and after exposure to specific intervention, at a rate of one exhalation per minute. MS acquisition was performed in full-scan positive ionization mode by scanning m/z 50–600 at a resolution of 240,000. Putative compound identification was supported by the mass accuracy of the instrument (5 ppm tolerance) and further confirmed by tandem MS experiments using high-energy collisional dissociation (HCD).

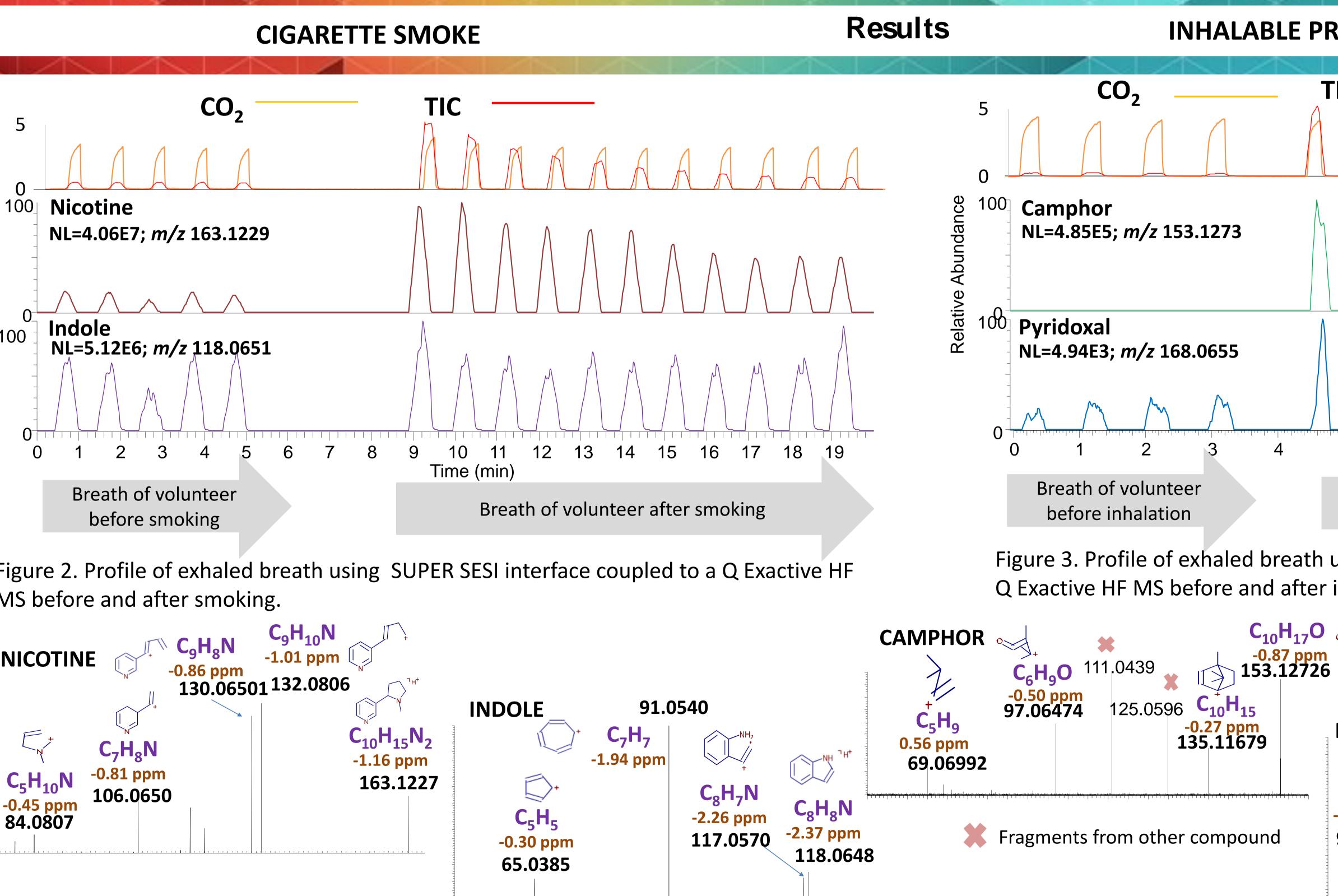


Figure 4. Experimental data of fragmentation of *m/z* 163.1229 (nicotine), *m/z* 118.0651 (indole), *m/z* 153.1273 (camphor), *m/z* 168.0655 dissociation (HCD).

T. Zivkovic Semren * ¹, C. Laszlo¹, M. Gomez², G. Vidal-de-Miguel², J. Hoeng¹, M. Peitsch¹, N. Ivanov¹, P.A. Guy¹ ¹*Philip Morris International,* ² *Fossil Ion Technology*

- from the lungs into the bloodstream.
- profile depending on the type of exposure.
- time exhaled breath samples.



Conclusions

 Exhalion Super SESI coupled to a Q Exactive HF MS system allows rapid monitoring of the absorption of exogenous compounds originating from cigarette smoke and/or aerosol from various inhalable products

 \checkmark Nicotine, one the main compound inhaled from smoking, showed a well-defined washing pattern in the lungs, where the intensity increased right after smoking and slowly decreased afterwards.

✓ Indole, known as an endogenous metabolite, showed a relatively flat

 Camphor, and pyridoxal—which were confirmed in a tested inhalable products—showed a similar washing pattern as nicotine.

 \checkmark These results demonstrate the benefits of this device to study real-

ODUCTS						
IC						1
						<u></u>
	·····	·····			,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,	
5	6 Time (r	7 nin)	8	9	10	11
Breath of volunteer after inhalation						
using SUPER SESI interface coupled to a nhalation. f^{H^+} $C_8H_8NO_2$ -0.97 ppm						
PYRIDOXAL C.H.N		C ₇ -0.	H ₈ NO 41 ppm 2.0599		50.0548 +0 C ₈ -0	^{¬н+} H ₁₀ NO ₃ .99 ppm 58.0653
(pyridoxal) from high-energy collisional						

Competing Financial Interest – The research described in this poster was sponsored by Philip Morris International