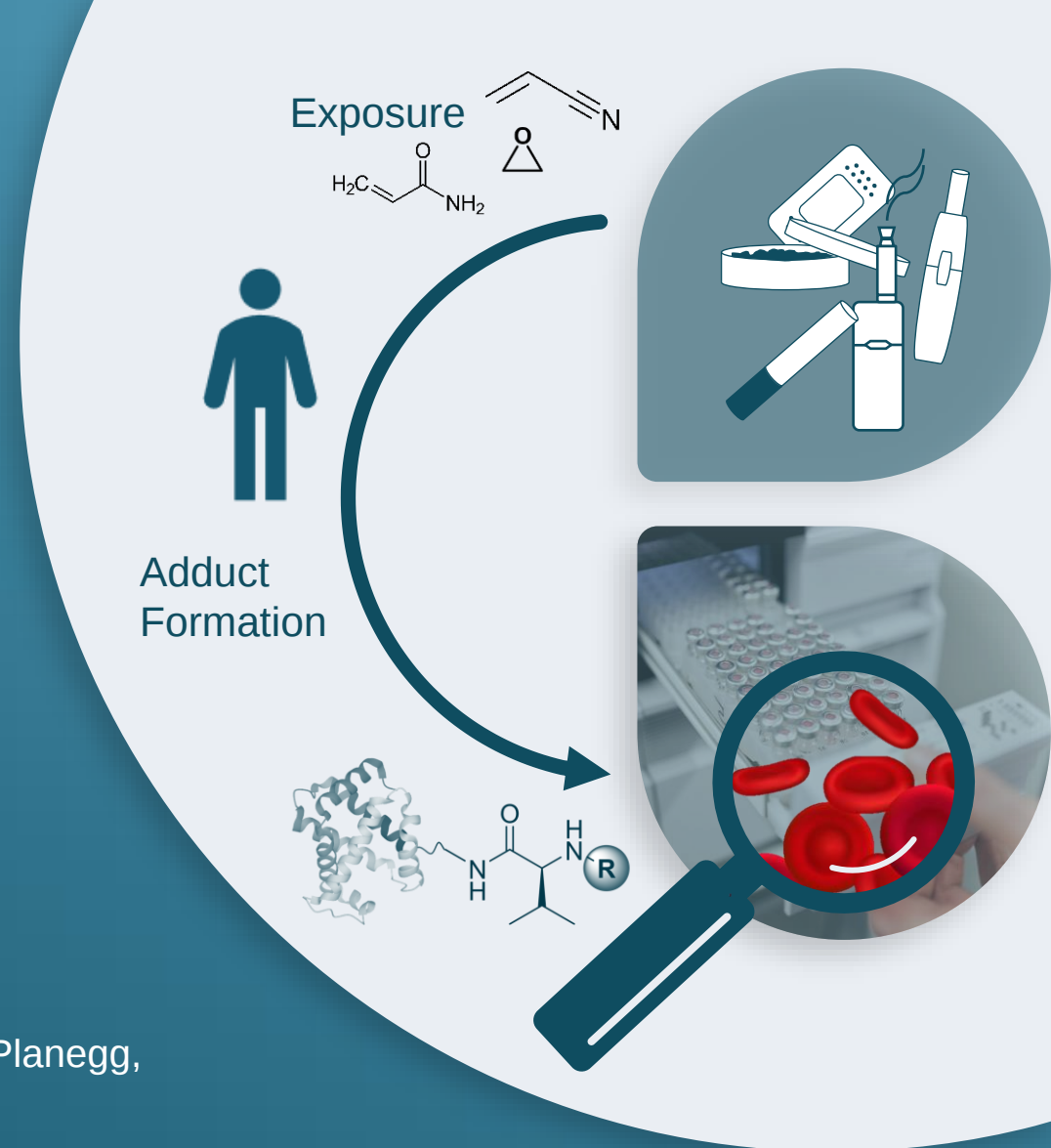


Identification of use-specific hemoglobin adduct patterns for different tobacco/nicotine product user groups by non-targeted GC-MS/MS analysis

M. Scherer, F. Pilz, T. Burkhardt, G. Scherer, N. Pluym

Analytisch-Biologisches Forschungslabor GmbH, Semmelweisstr. 5, 82152 Planegg, Germany

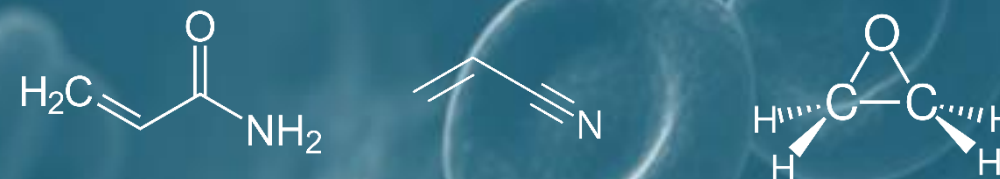


Introduction

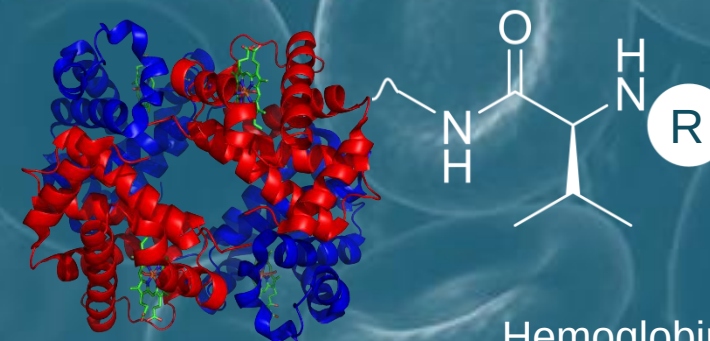
- **Biomarkers of exposure** commonly monitored in urine
→ short half-life
- **Hemoglobin (Hb)**: Life-time of ~ 120 days
→ Formation of adducts with electrophilic compounds
- **Electrophilic compounds**: formed *in vivo* by the metabolism of molecules derived from endogenous (e.g., oxidative stress) and exogenous (e.g., diet, smoking) sources
 - Modification of nucleophilic sites on DNA and functional proteins (e.g., HSA and Hb)
 - Increased cancer risk
- Chemically stable adducts accumulate during exposure

Potential **biomarker** for monitoring
long-term *in vivo* exposure

Electrophilic compounds



Formation of hemoglobin adducts



Hemoglobin with N-terminal adduct

Introduction

- **Biomarkers of exposure** commonly monitored in urine
→ short half-life
- **Hemoglobin (Hb)**: Life-time of ~ 120 days
→ Formation of adducts with electrophilic compounds
- **Electrophilic compounds**: formed *in vivo* by the metabolism of molecules derived from endogenous (e.g., oxidative stress) and exogenous (e.g., diet, smoking) sources
 - Modification of nucleophilic sites on DNA and functional proteins (e.g., HSA and Hb)
 - Increased cancer risk
- Chemically stable adducts accumulate during exposure

Potential **biomarker** for monitoring
long-term *in vivo* exposure

REQUIREMENT

Identification of product use-specific
hemoglobin-adducts



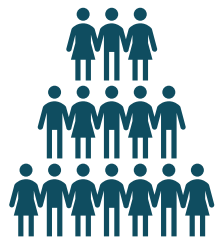
Risk evaluation



Compliance

Clinical Study – Study Design

Clinical Study with 5 different nicotine product user groups and a non-user group (control)

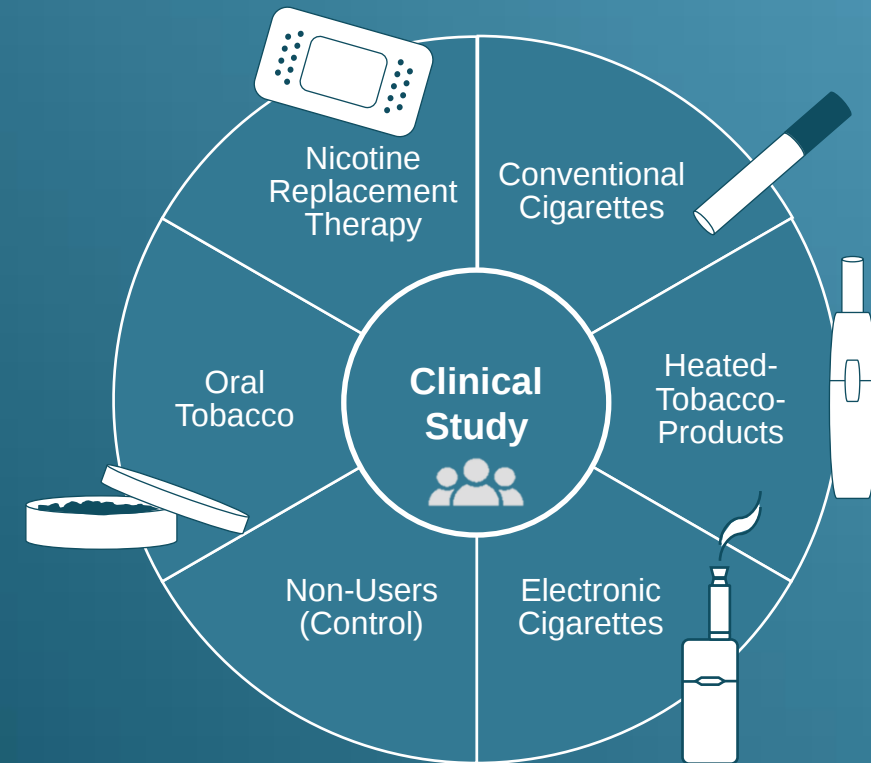


60 healthy volunteers

- Males and Females
- Aged 19 – 65 years
- BMI: 18 – 33 kg / m²

GENERAL STUDY OBJECTIVE

Identify biomarkers or biomarker patterns capable to discriminate between different nicotine product user groups and non-users.



10 subjects per group

Day 1

Day 2

Day 3

Analytical Strategy

Bioanalytical methods

Holistic



Introduction

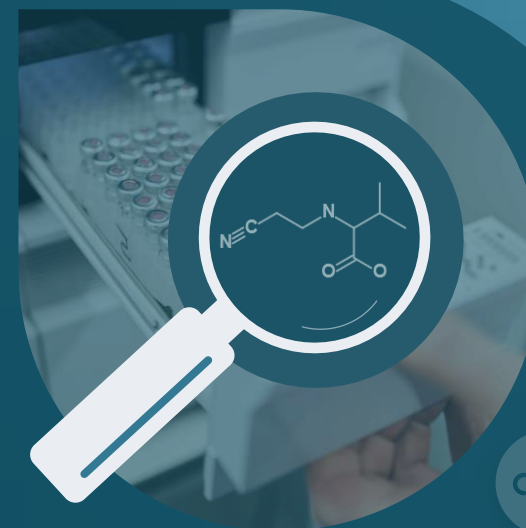
Study Design & Method

Results & Discussion

Conclusion

We have a problem !

Screening
Hemoglobin adducts
for specific product categories



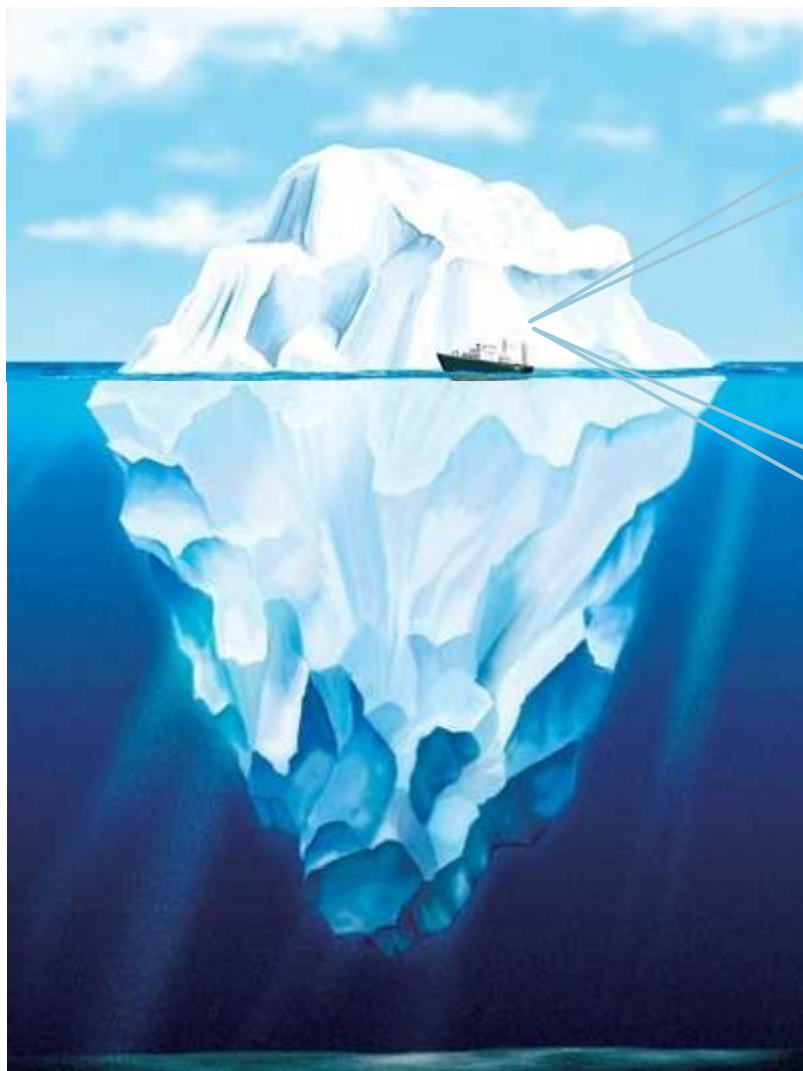
abf

Analytical Strategy

Bioanalytical methods

Targeted

Holistic



Introduction

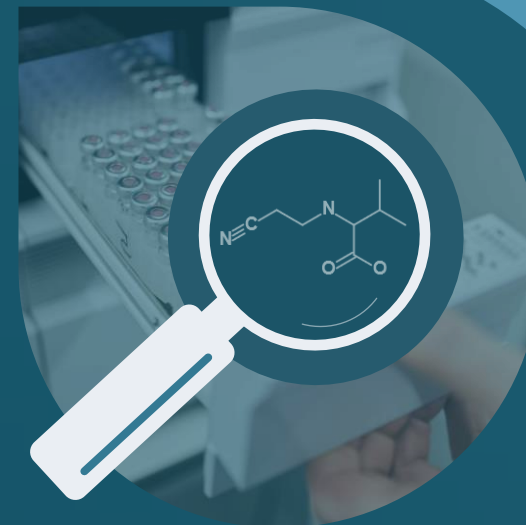
Study Design & Method

Results & Discussion

Conclusion

We have a problem !

Screening
Hemoglobin adducts
for specific product categories



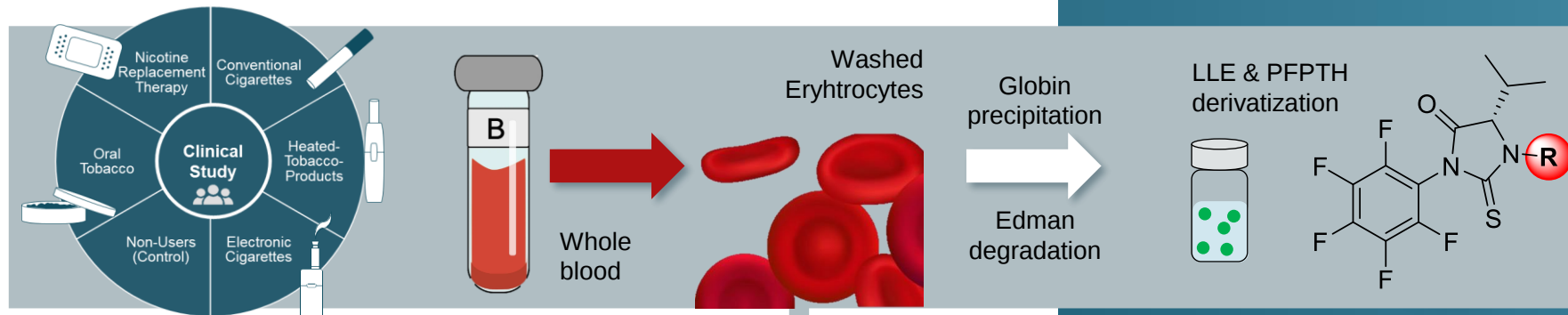
Can we
identify the
root cause?

Targeted quantitative profiles



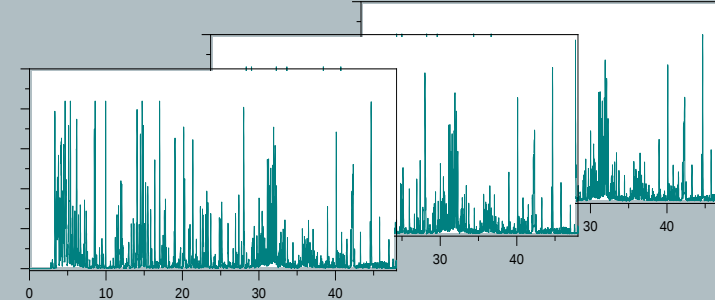
abf

Non-targeted screening – Workflow

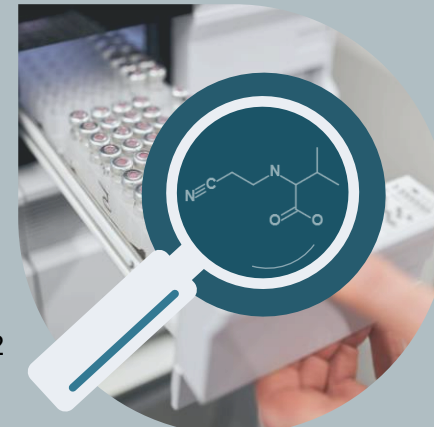
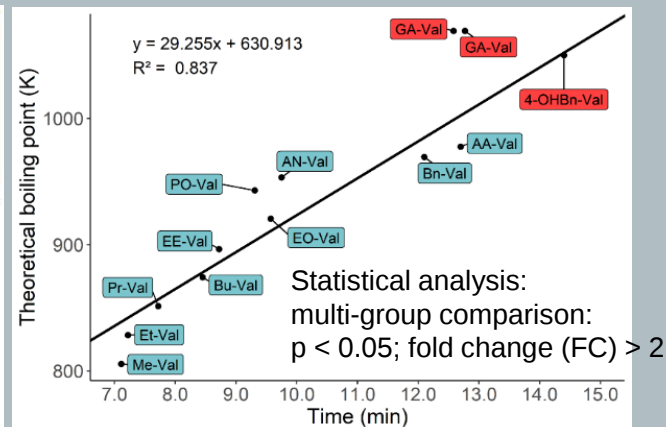
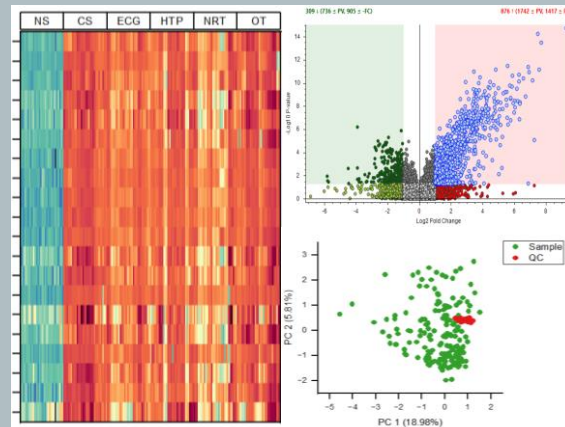


Sampling &
Sample Preparation

GC-MS/MS
Shimadzu TQ8050



Sample Analysis



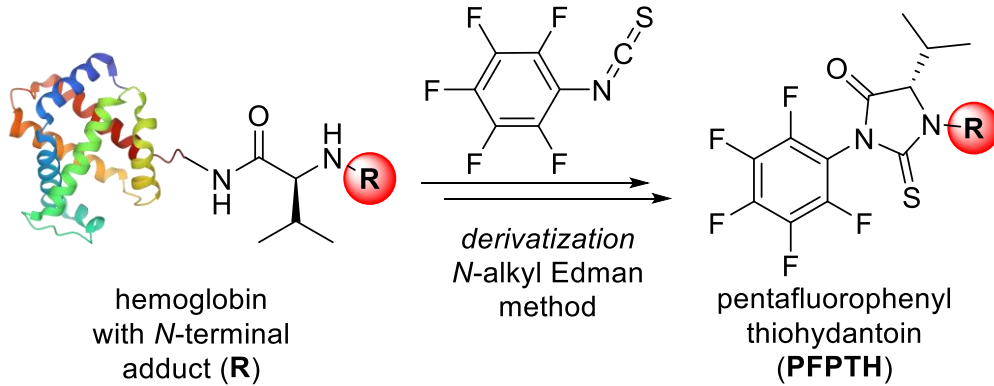
Data Processing
Statistical Analysis
Compound Identification

Non-Target Screening

Derivatization of Hb with pentafluorophenyl isothiocyanate (PFPITC)

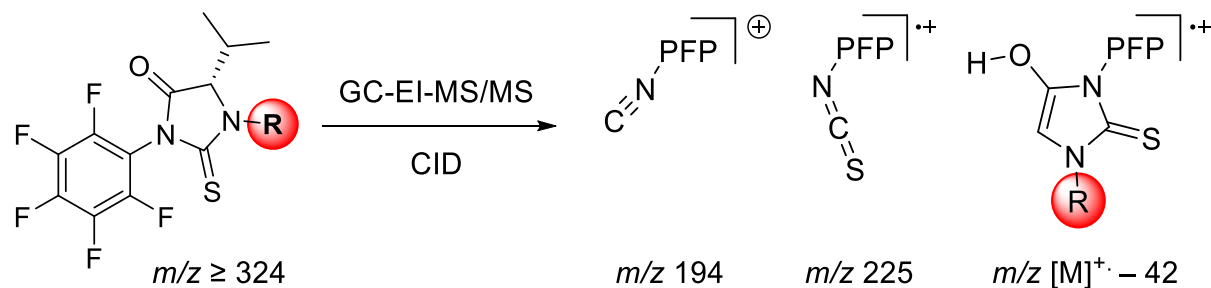


Formation of small molecules (PFPTH) for analysis

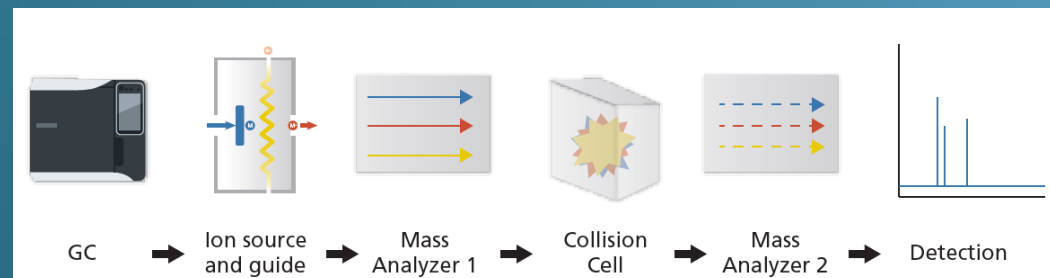


Non-Target Screening

- PFPTH derivatives of N-terminal Hb adducts show a similar fragmentation pattern in MS/MS



- MRM screening approach uses the structural relation and similar core structure of PFPTHs
- Screening range in Q1: m/z 338 – 487 (multiple injections)

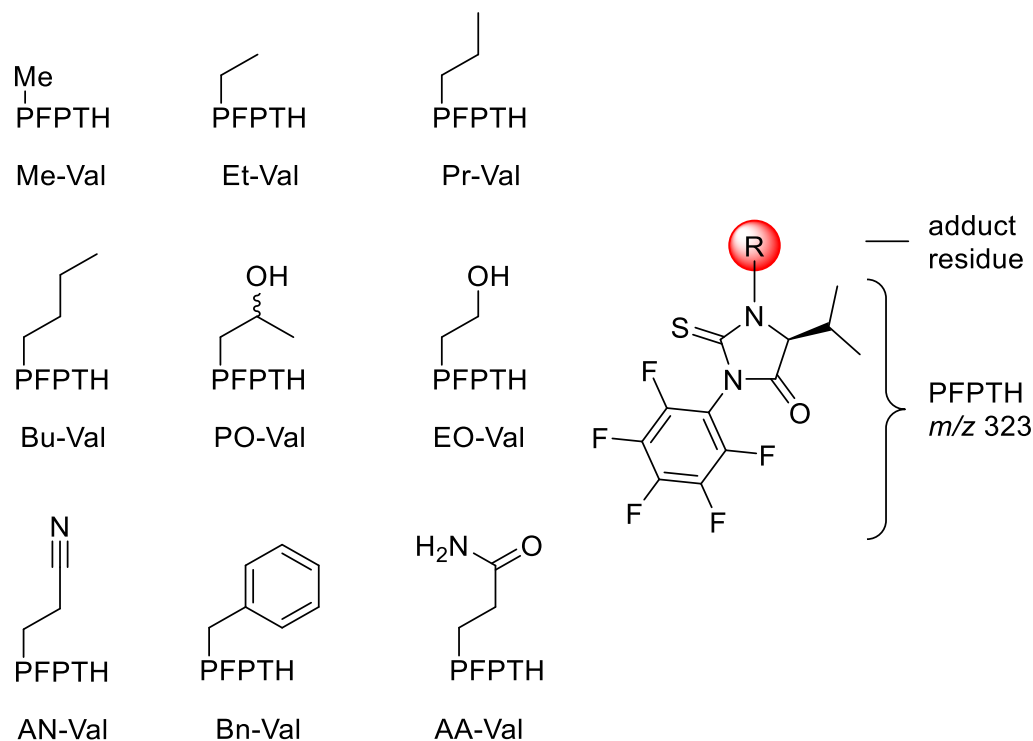


MRM method used to screen for N-terminal Hb adducts

Q1: Parent ion		Q3: Fragments	
$[M]^+$	$[M]^+ - 42$	225	194
338	296	225	194
339	297	225	194
340	298	225	194
⋮	⋮	⋮	⋮
487	445	225	194

Method Validation

9 compounds evaluated:



→ Cover the expected retention time (RT), polarity and mass range

Results

Repeatability (CV, N = 5)

4.6 – 13.1 %

Precision Intraday (CV, N = 3) Interday (CV, N = 4)

1.4 – 19.0 %

6.6 – 19.0 %

Accuracy Intraday (N = 3) Interday (N = 4)

82.6 – 107.9 %

89.2 – 117.8 %

Carry over

0.0 – 0.3 %

Linearity (R², Range: 15, 60, 120 – 3600 pmol/g Gb)

≥ 0.9903

Post-preparative stability (room temperature, 5 d)

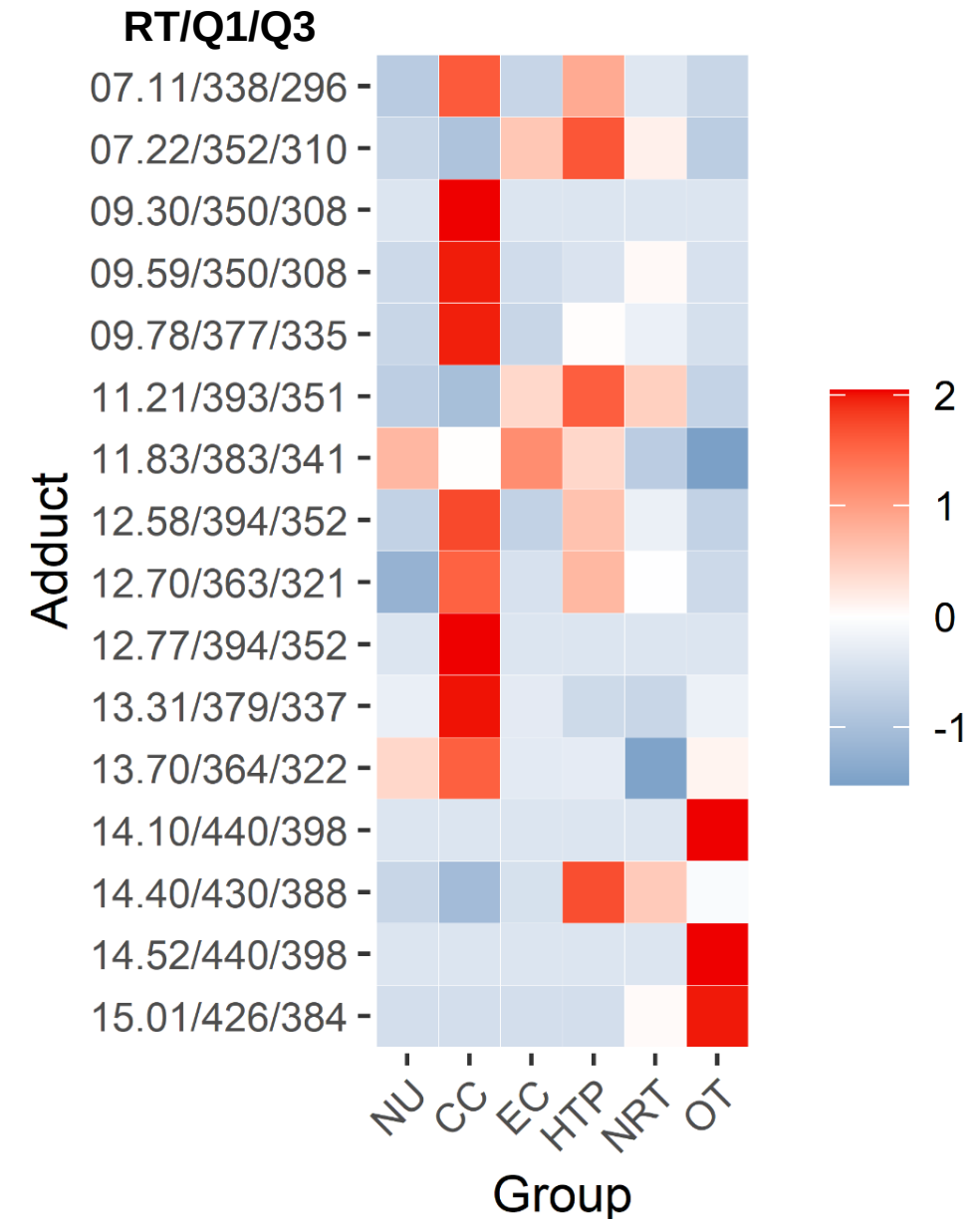
76.8 – 123.6 %

Non-Target Screening

- 13 adducts showed significant differences among groups ($p < 0.05$)
- 3 more adducts determined by fragmentation pattern ($p > 0.05$)

Identification based on...

- Adductome data**
 - Parent ions
 - Retention time
 - Isotopic distribution
- Literature review and database search** on exposure from tobacco/nicotine products and previously reported adducts



Identification Strategy

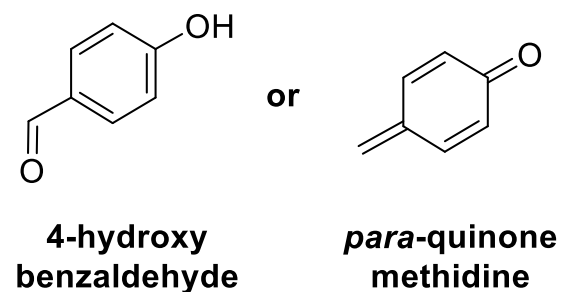
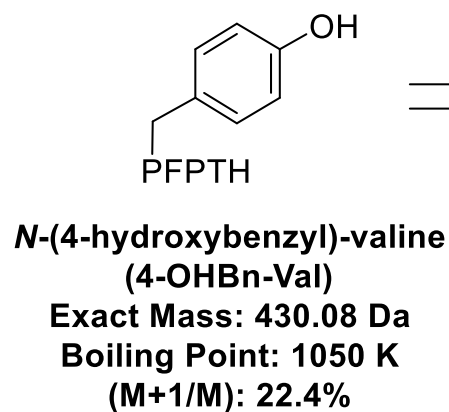
Adduct at 14.40 min

RT (min)	Parent Ion (m/z)	Relative Intensities of Mass Transitions (%) ^a		
		194	225	precursor ion -42
14.40	430	39.6 (55.7)	35.9 (26.6)	100.0
	431	14.9 (12.6)	-	22.4 (20.6)

^aRSDs in percent in brackets.

Potential Adduct

Potential Precursor

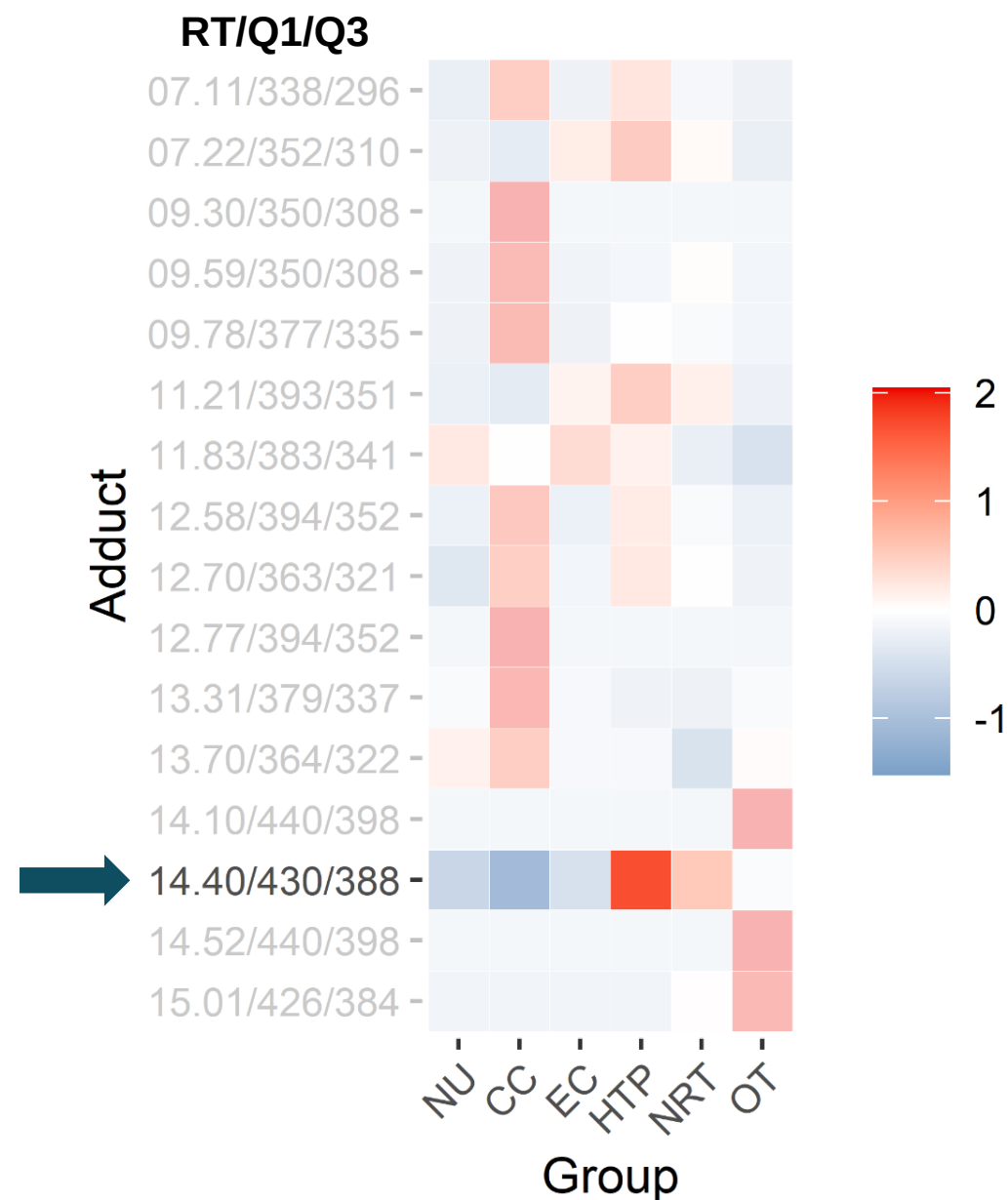


Introduction

Study Design & Method

Results & Discussion

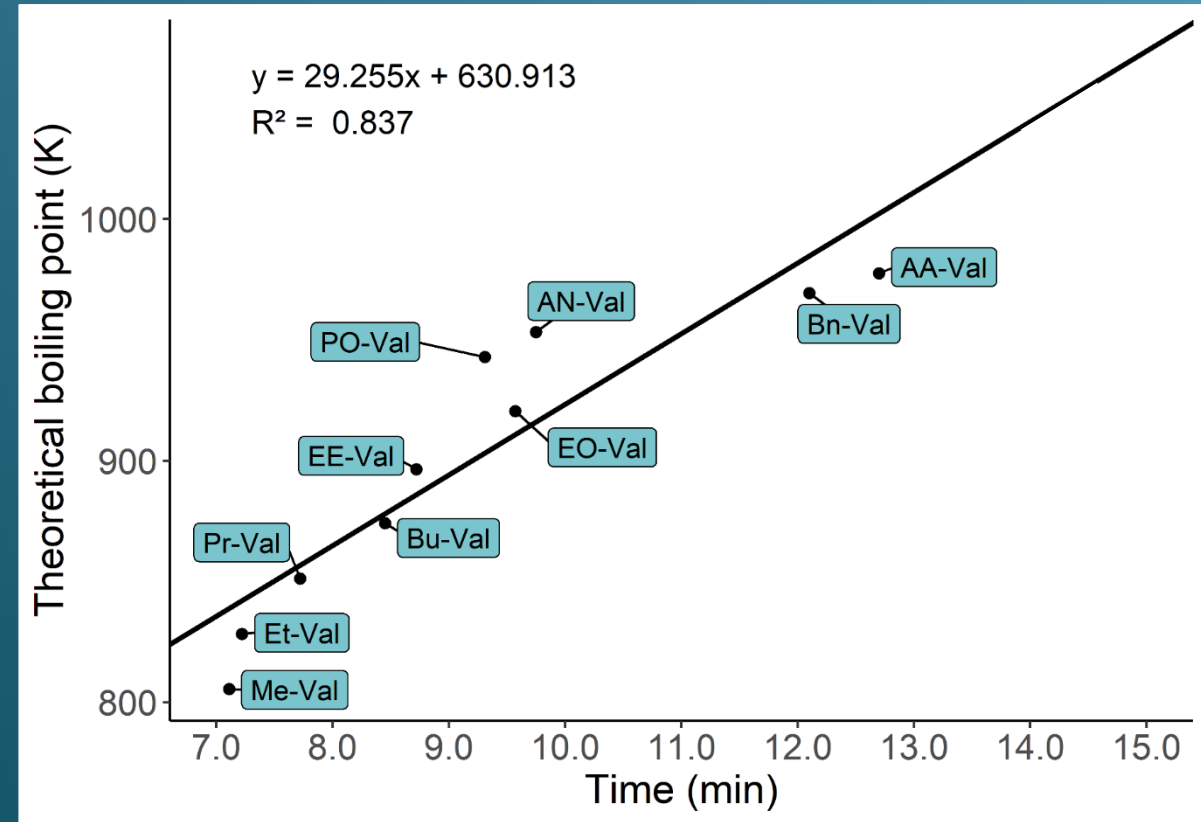
Conclusion



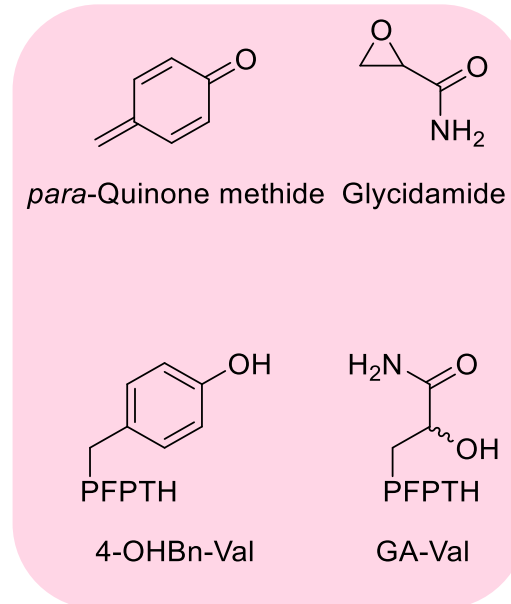
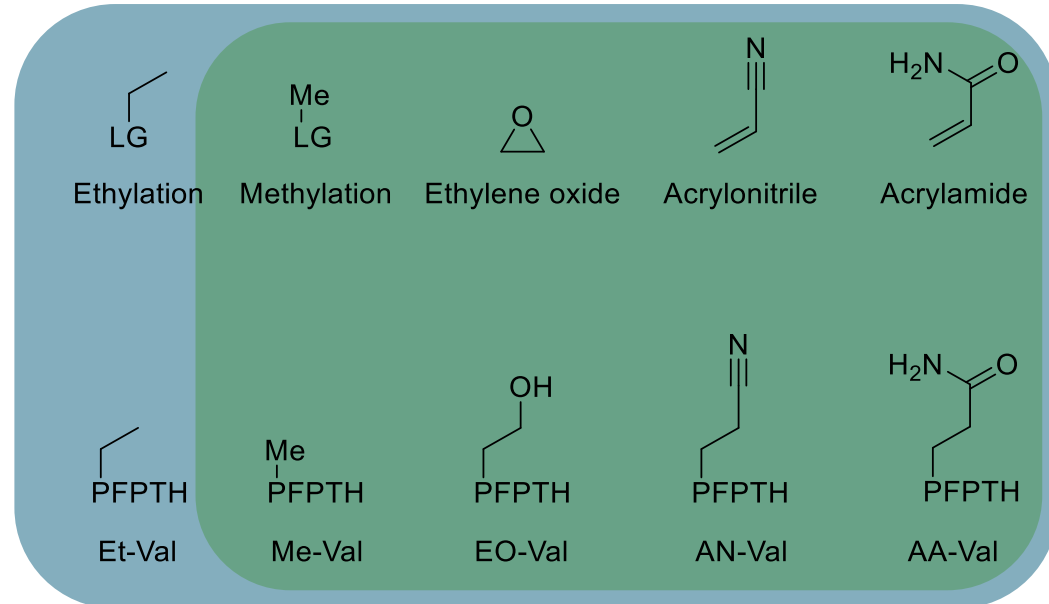
Identification Strategy

Boiling Point Model

Hb adducts used to generate a linear model to predict boiling points of unknowns based on the observed retention times (RTs)

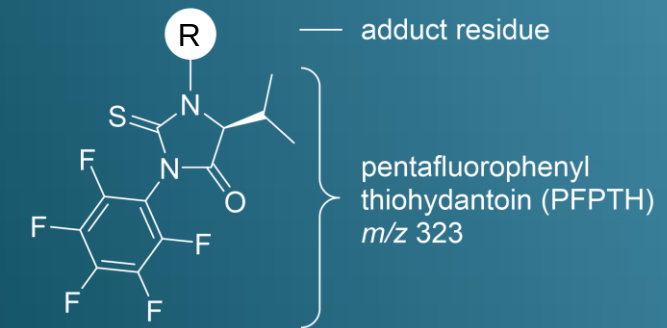


Identified Hb Adducts



Precursor

Hb Adduct

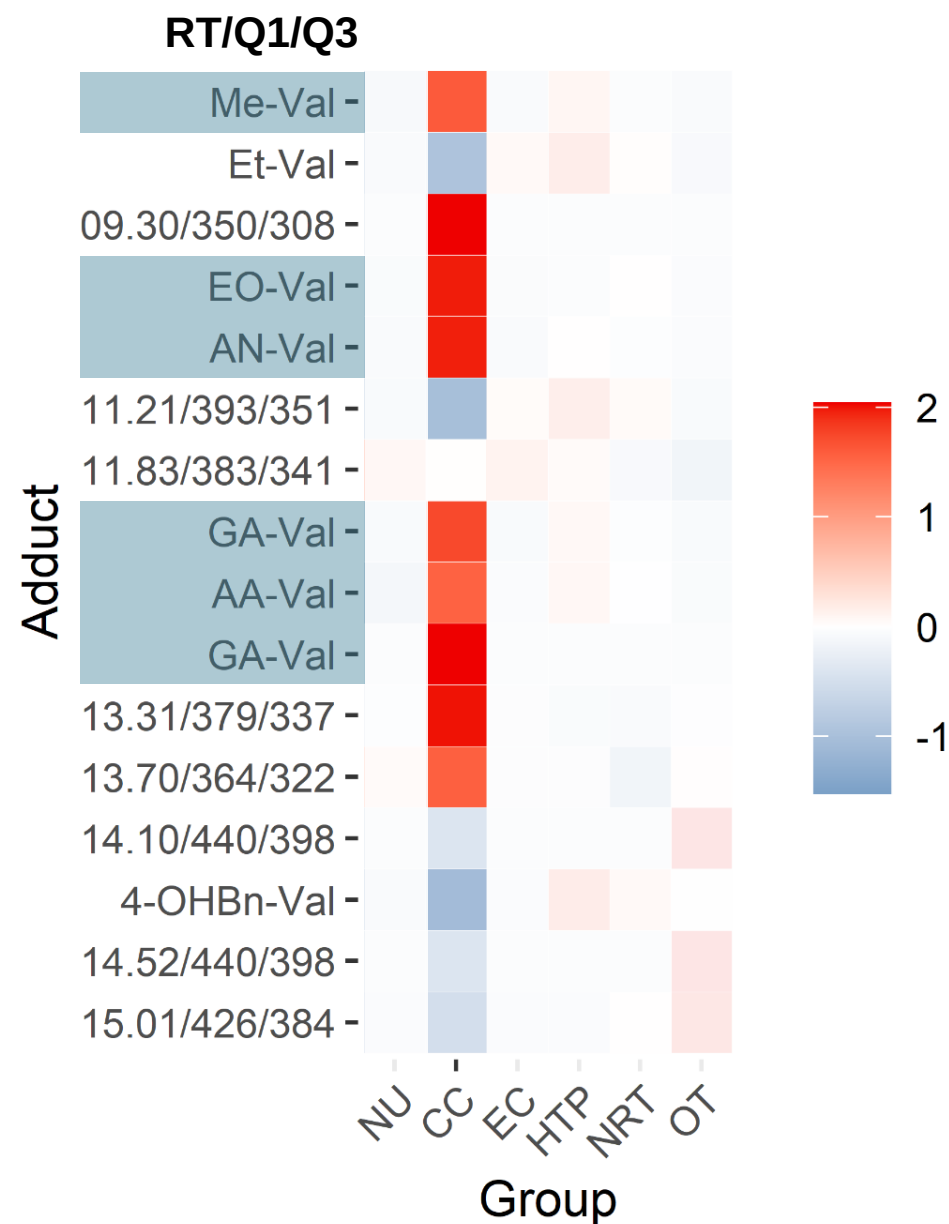


LG - leaving group

Identified Hb Adducts

Cigarette Smokers (CC):

High levels of Me-Val, EO-Val, AN-Val, AA-Val and its metabolite GA-Val



Identified Hb Adducts

Cigarette Smokers (CC):

High levels of Me-Val, EO-Val, AN-Val, AA-Val and its metabolite GA-Val

Users of Heated Tobacco Products (HTP):

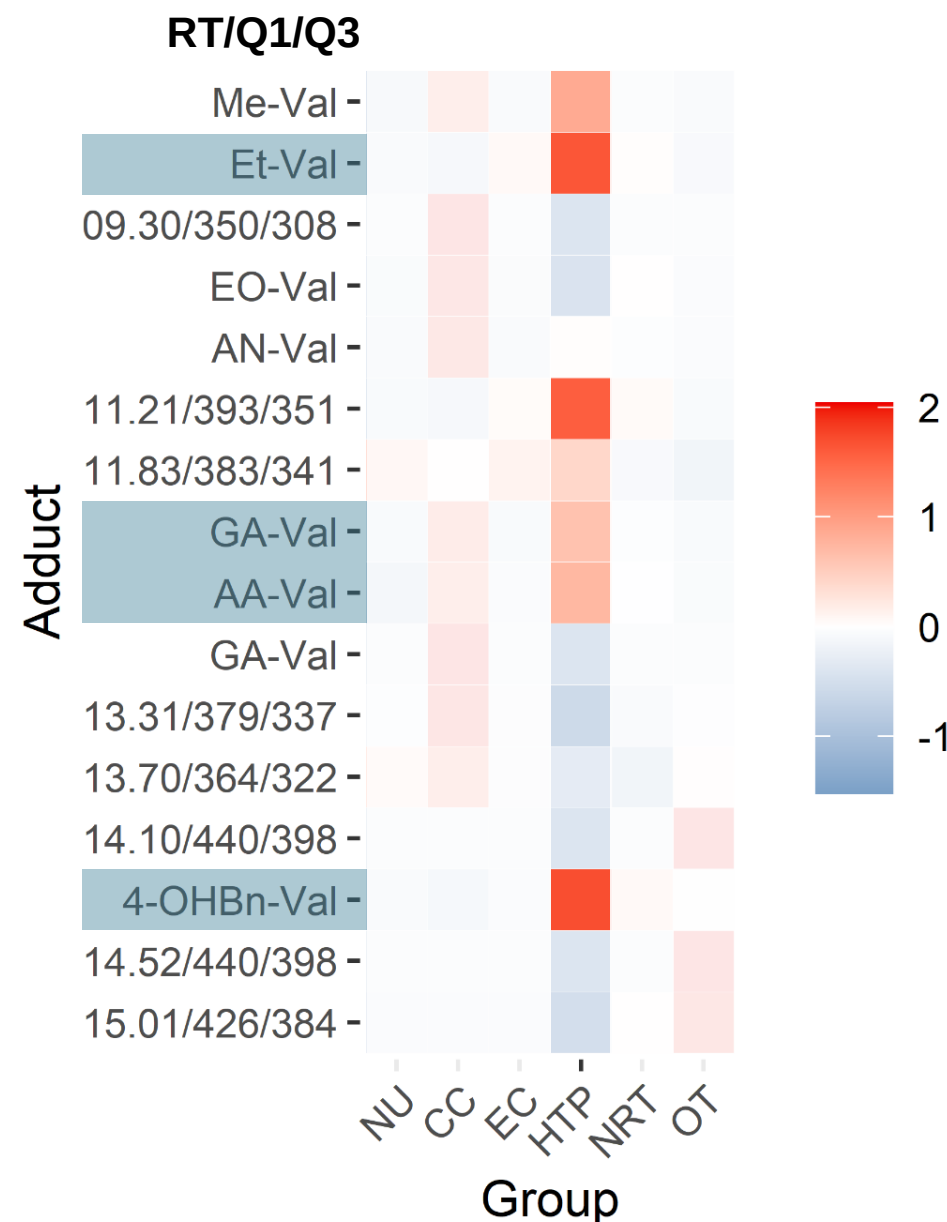
Significantly elevated levels of Et-Val, AA-Val and GA-Val as well as 4-OHBn-Val

Introduction

Study Design &
Method

Results & Discussion

Conclusion



Identified Hb Adducts

Cigarette Smokers (CC):

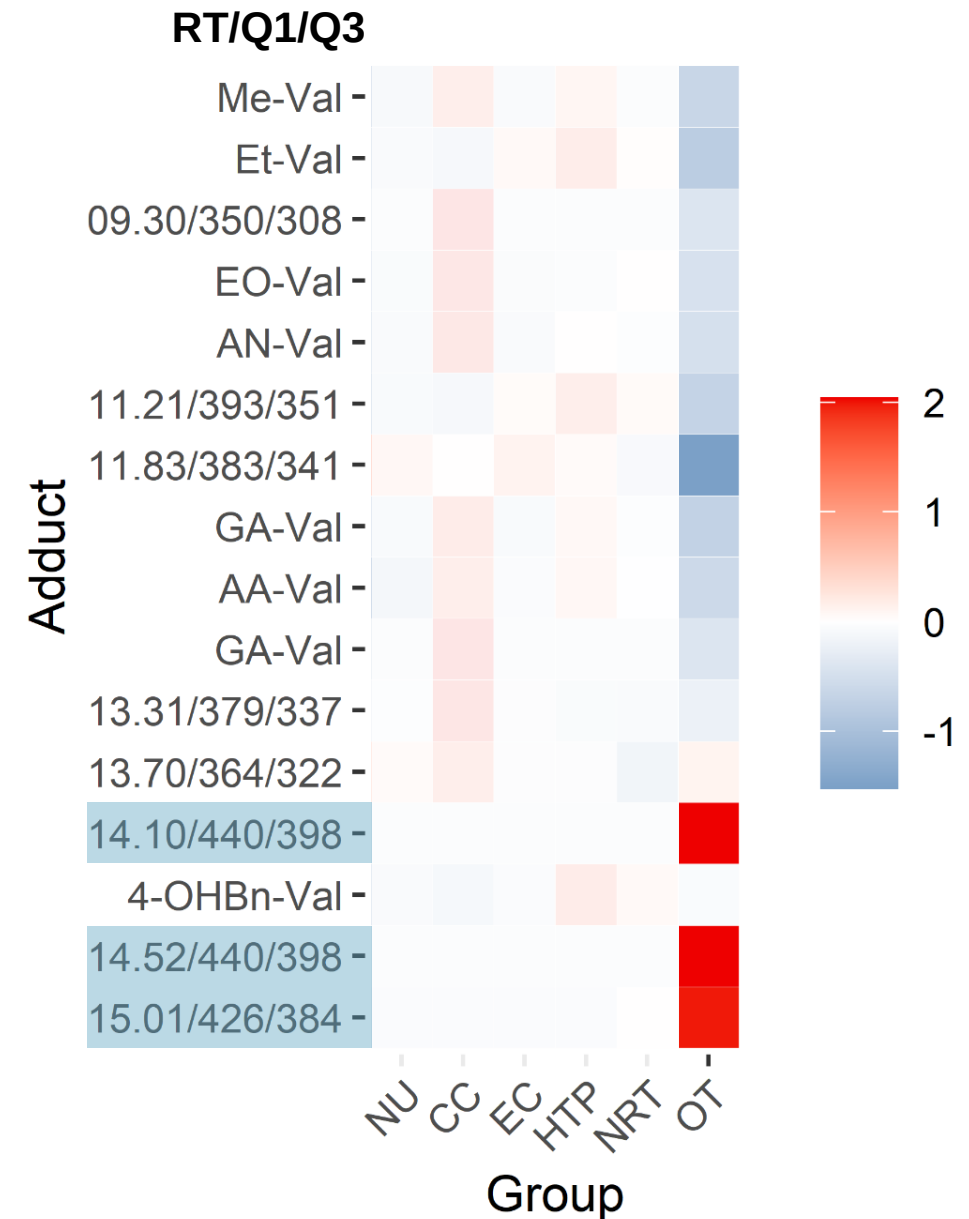
High levels of Me-Val, EO-Val, AN-Val, AA-Val and its metabolite GA-Val

Users of Heated Tobacco Products (HTP):

Significantly elevated levels of Et-Val, AA-Val and GA-Val as well as 4-OHBn-Val

Users of Oral Tobacco (OT):

Elevated levels of unknown adducts at high RTs



Identified Hb Adducts

Cigarette Smokers (CC):

High levels of Me-Val, EO-Val, AN-Val, AA-Val and its metabolite GA-Val

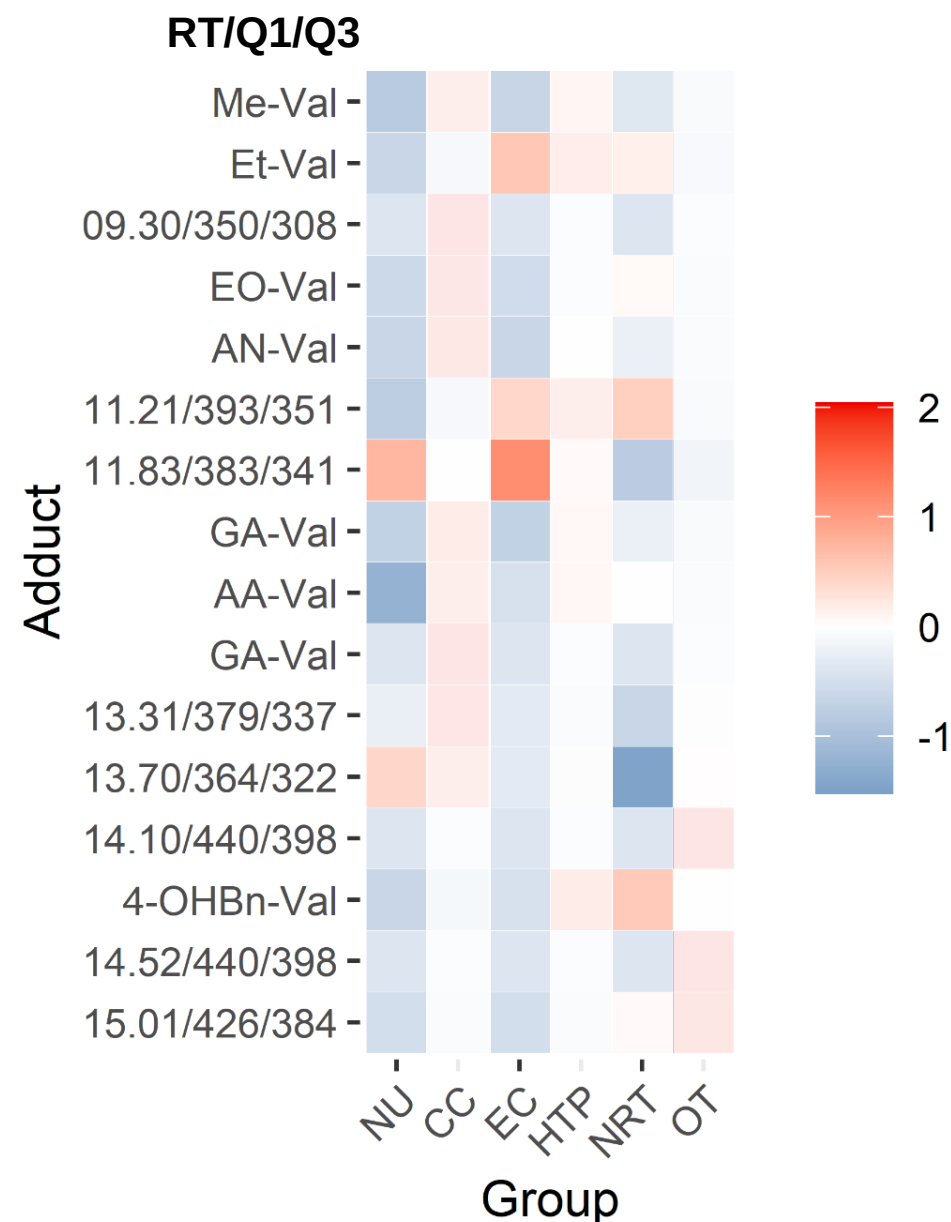
Users of Heated Tobacco Products (HTP):

Significantly elevated levels of Et-Val, AA-Val and GA-Val as well as 4-OHBn-Val

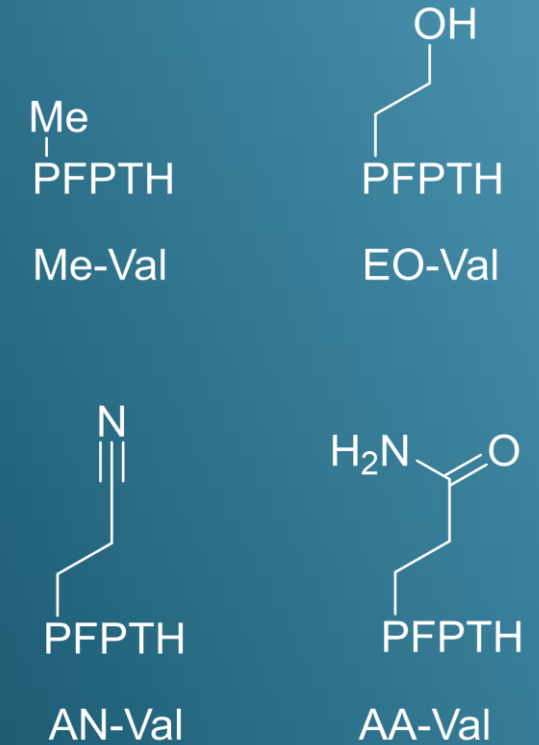
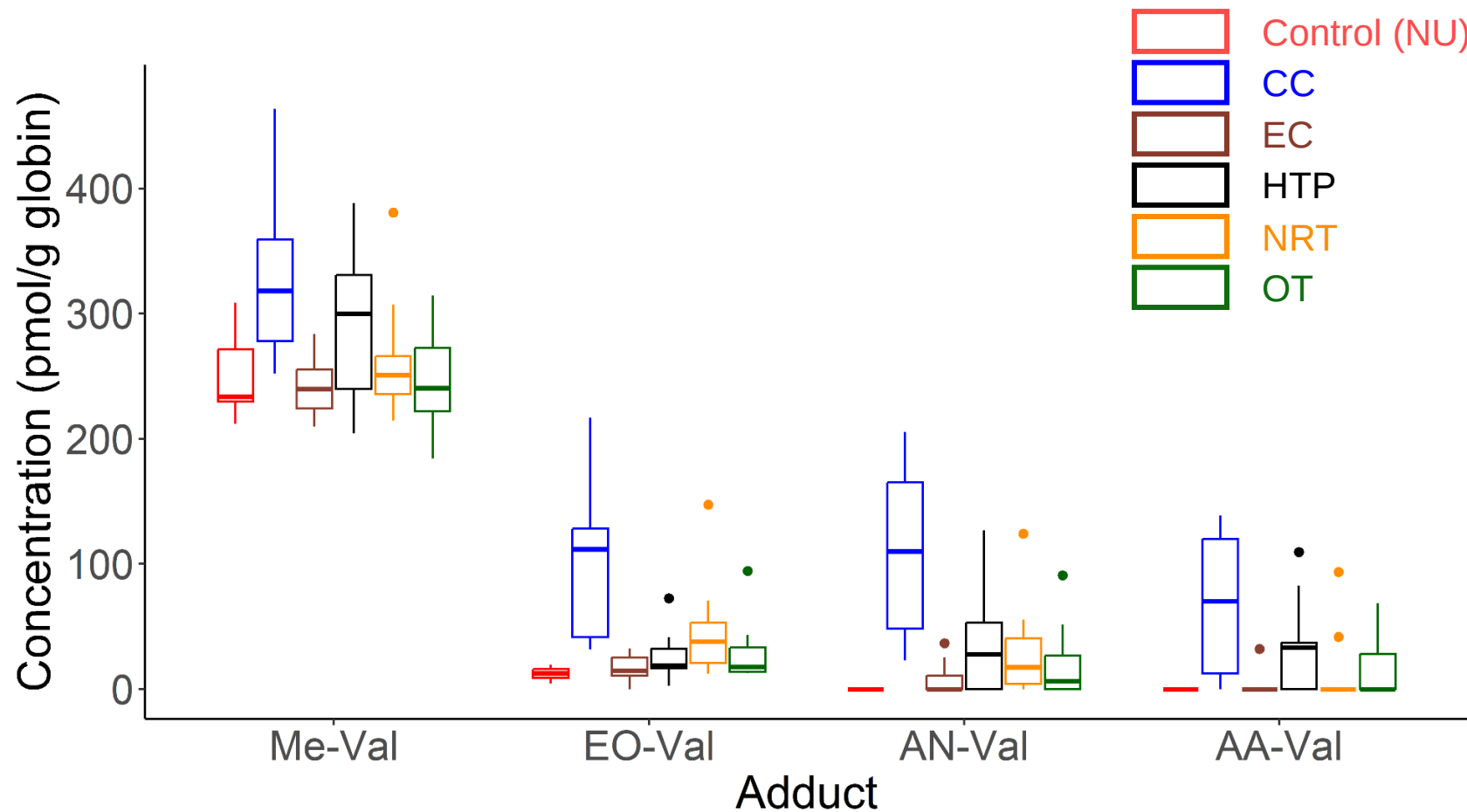
Users of Oral Tobacco (OT):

Elevated levels of unknown adducts at high RTs

The adduct profile of e-cigarette (EC), nicotine replacement therapy (NRT) users, and the control group (NU) is not significantly different

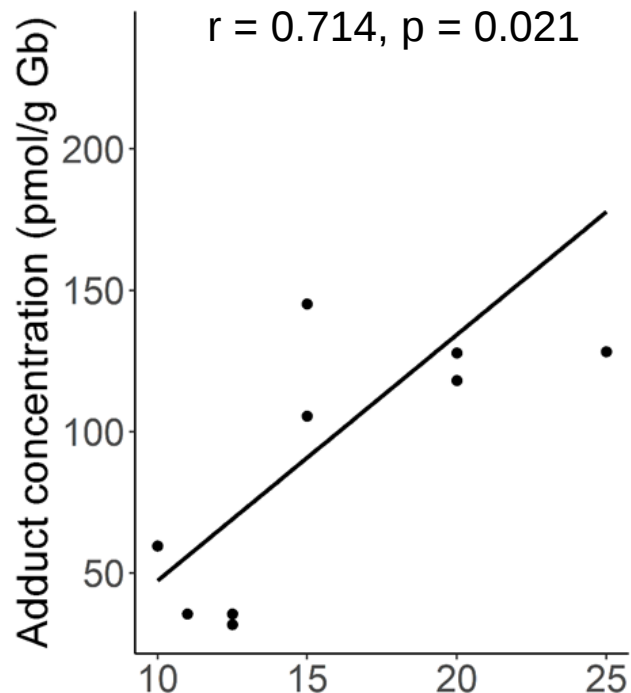


Quantification of Me-Val, EO-Val, AN-Val and AA-Val

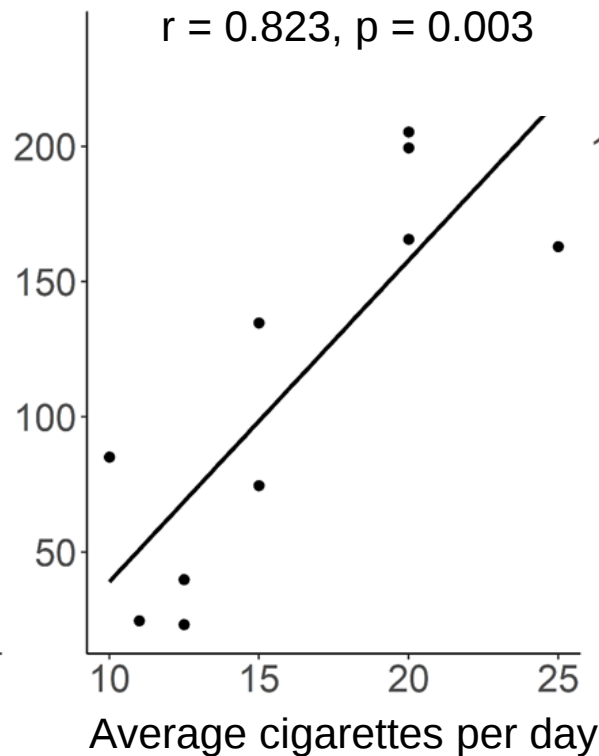


Correlation between Adduct Levels and Cigarette Consumption in Smokers

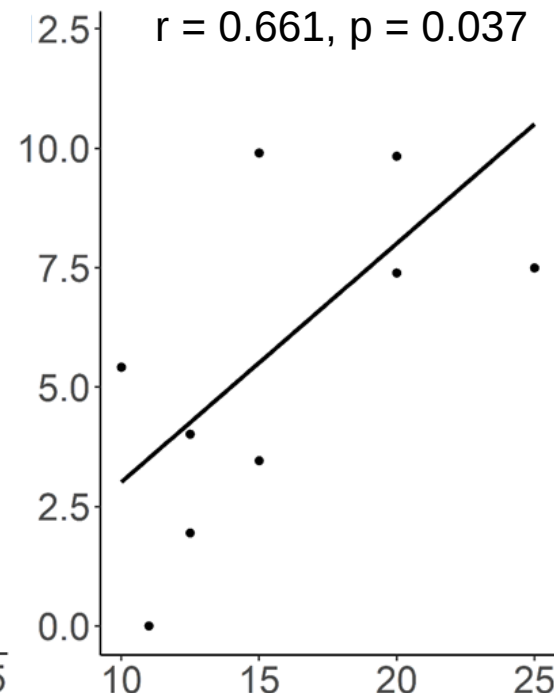
EO-Val



AN-Val



GA-Val (sum)



For Smokers (CC)

Significant moderate to strong **correlation** between selected adducts and cigarette consumption ($p < 0.05$)

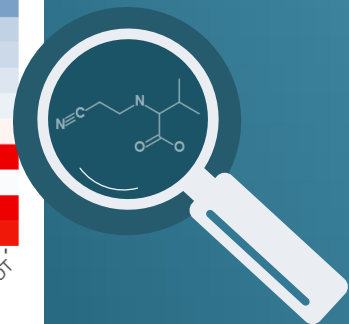
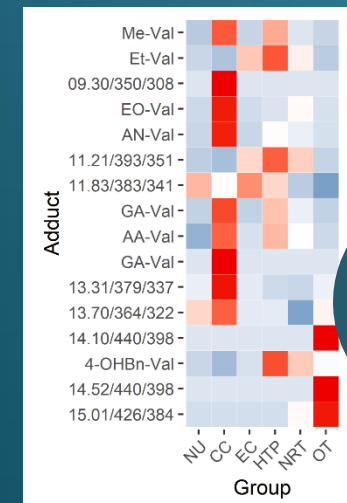
Conclusion

Method Development & Validation

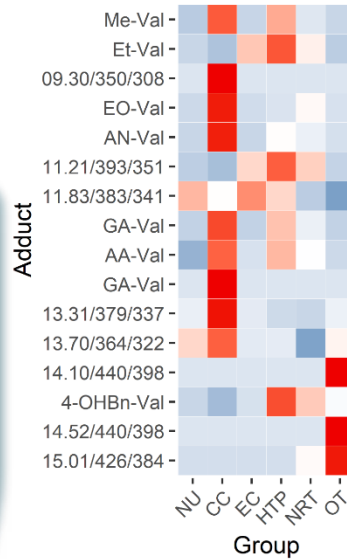
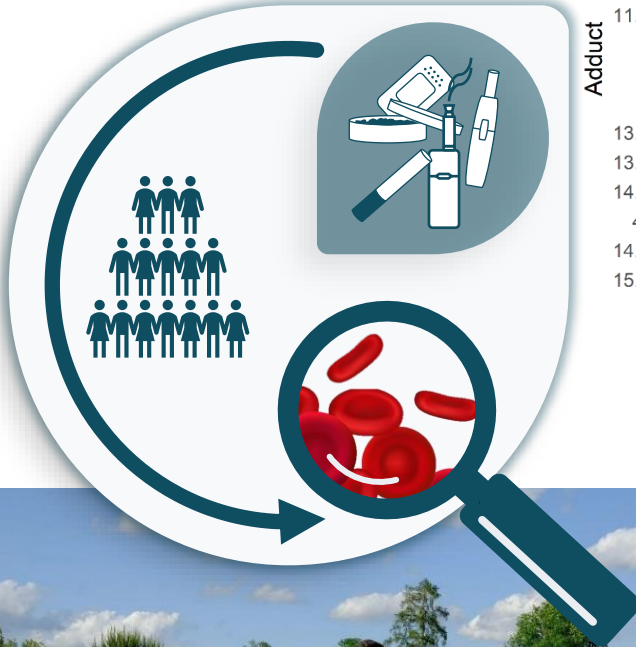
- Successful **development** of a **non-targeted GC-MS/MS method** for the analysis, evaluation and identification of hemoglobin adducts
- Successful **method validation** with 9 representative standards
- Development of a targeted method Hb-adduct method for quantification

Application of the Method & Identification

- Detection of **13 adducts** showing **significant differences** between the different nicotine user groups
- **Identification** of features using modified dipeptides and incubation experiments
- **Smokers (CC)** exhibited the **highest exposure to electrophiles** and showed good correlation with cigarette consumption for EO-Val, AN-Val and GA-Val
- Users of **HTPs** show elevated levels of Et-Val, AA-Val, GA-Val and 4-OHBn-Val



Acknowledgment



ABF Team
Fabian Pilz
Dr. N. Pluym
Dr. Therese Burkhardt
Prof. Dr. Gerhard Scherer



Funding:

This study was funded with a grant from the Foundation for a Smoke-Free World. The Foundation for a Smoke-Free World is an independent, US nonprofit 501(c)(3) grantmaking organization with the purpose of improving global health by ending smoking in this generation. Through September 2023, the Foundation received charitable gifts from PMI Global Services Inc. ("PMI"). Independent from PMI since its founding in 2017, the Foundation operates in a manner that ensures its independence from PMI and any commercial entity. The Foundation seeks to collaborate with associations and institutions to accelerate its investments in life-saving research projects.