

Why do scientists care about sweat sensors?



Research hotspot: extract physiological information from biological fluid

- Sweat VS plasma: **non-invasive**
- Sweat VS urine, tear : **elegant**
- Fast → **high frequency, large-scale**



Relative research focus on:

- **Collection** methods (amount)
- **Analysis** methods (sensitivity, accuracy)
- Extract **insights** from sweat

For example

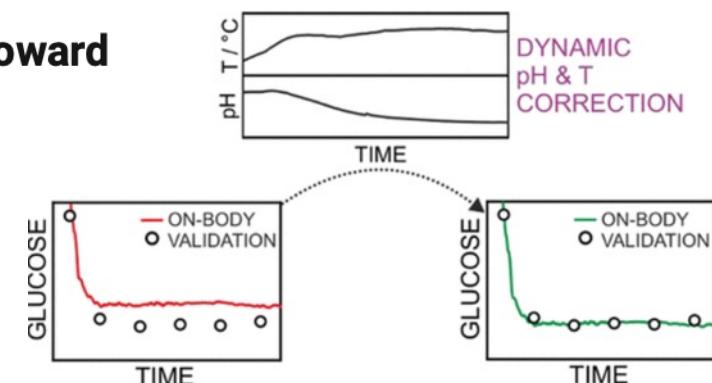
ARTICLE | June 26, 2020

Epidermal Patch with Glucose Biosensor: pH and Temperature Correction toward More Accurate Sweat Analysis during Sport Practice

Alexander Wiorek, Marc Parrilla, María Cuartero, and Gastón A. Crespo*

Citations

168





nature communications

Article | [Open access](#) | Published: 13 October 2021

Finger sweat analysis enables short interval metabolic biomonitoring in humans

[Julia Brunmair](#), [Mathias Gotsmy](#), [Laura Niederstaetter](#), [Benjamin Neuditschko](#), [Andrea Bileck](#), [Astrid Slany](#), [Max Lennart Feuerstein](#), [Clemens Langbauer](#), [Lukas Janker](#), [Jürgen Zanghellini](#), [Samuel M. Meier-Menches](#) & [Christopher Gerner](#) 

[Nature Communications](#) **12**, Article number: 5993 (2021) | [Cite this article](#)

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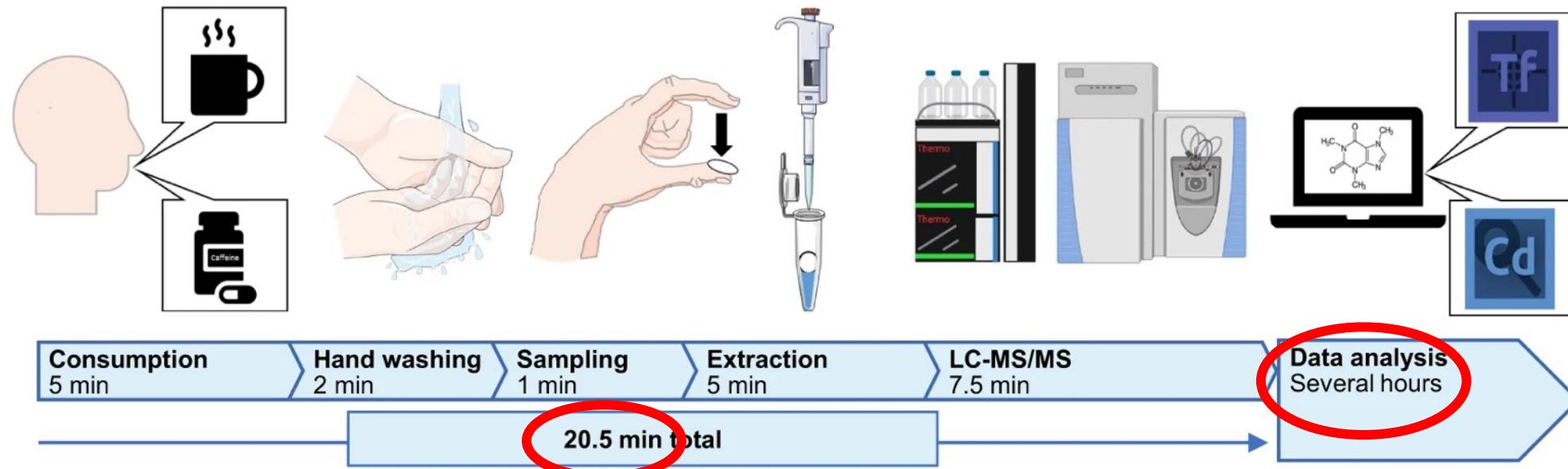
12212859 Sijie Li

The pipeline to collect sweat



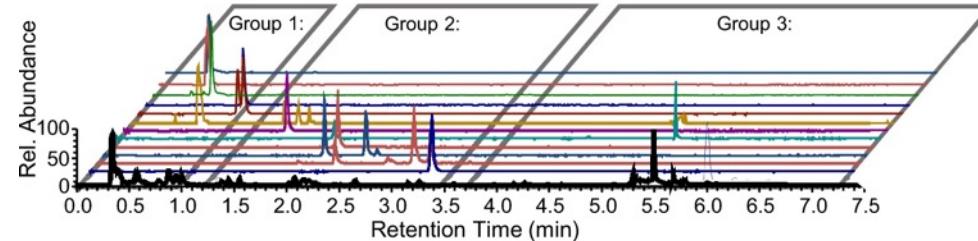
南方科技大学
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Department of Biomedical Engineering

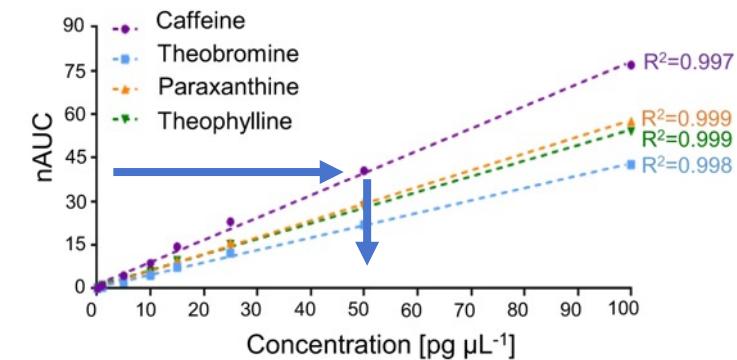


$$2 \text{ min} \times 2 \text{ cm}^2 \times 50\text{--}500 \text{ nL min}^{-1} \text{ cm}^{-2} = 200\text{--}2000 \text{nL sweat}$$

About MS



AUC = Area Under Curve → concentration

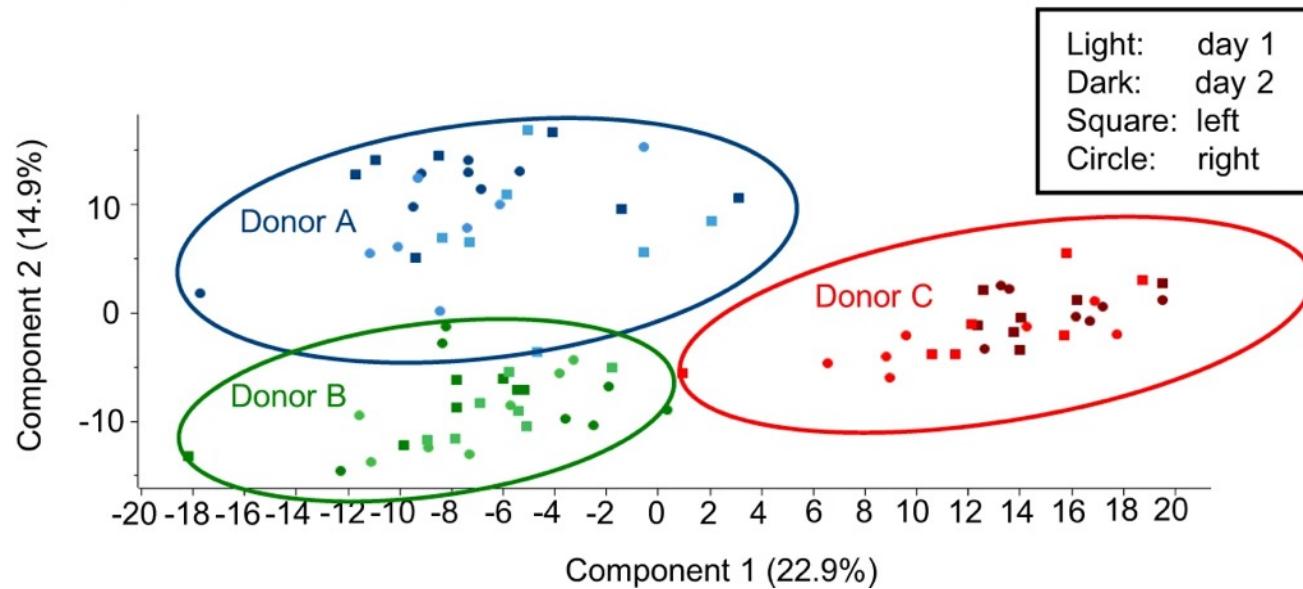


The pipeline works

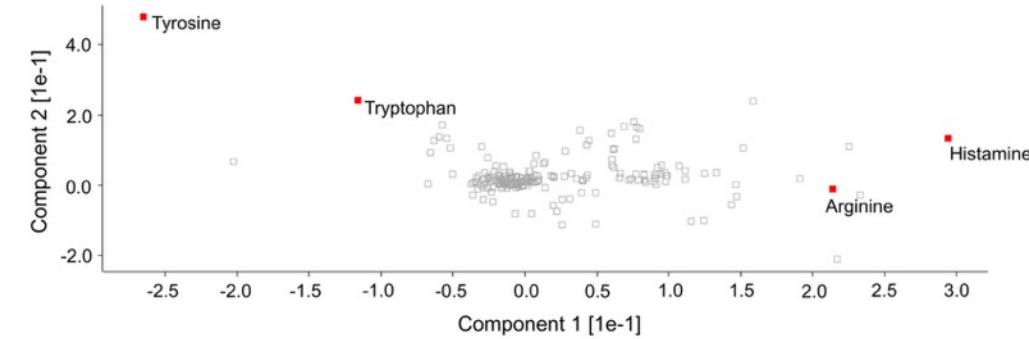


Identified and verified **250 metabolites**

- ✓ Previous known + unknown
- ✓ Endogenous + exogenous



Study A: observational

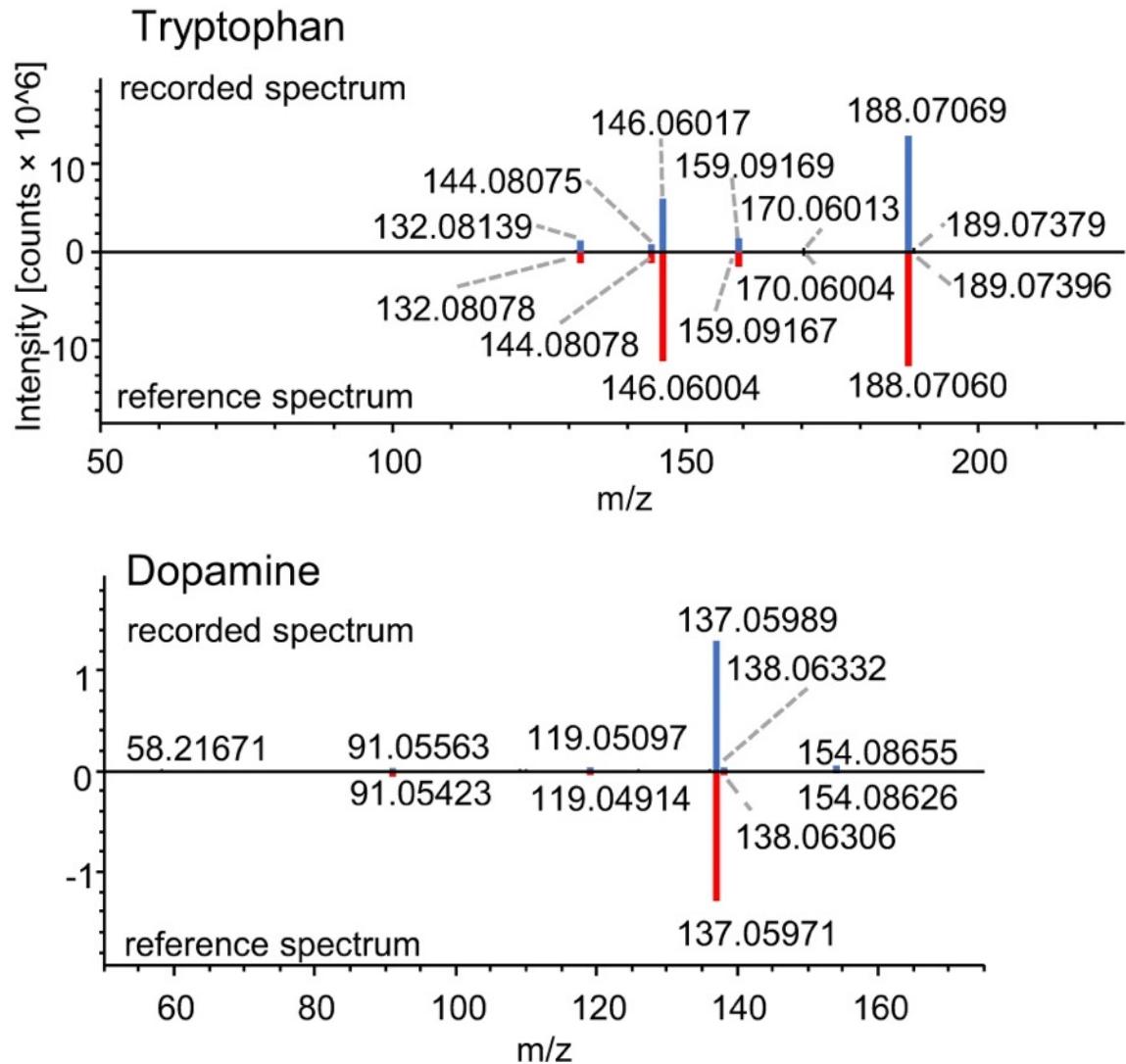
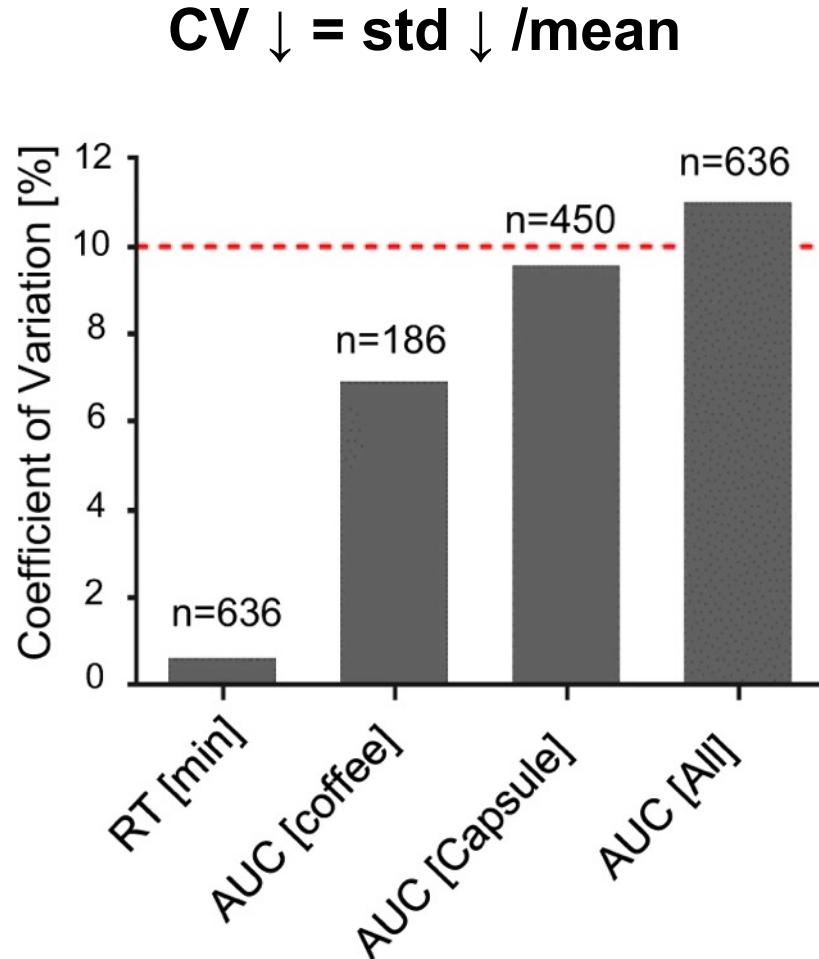


PC loading

Some small discoveries

- Sample clustered according to individuals
- No obvious differences between left and right hands
- PCs strongly determined by several endogenous metabolites

The pipeline is robust

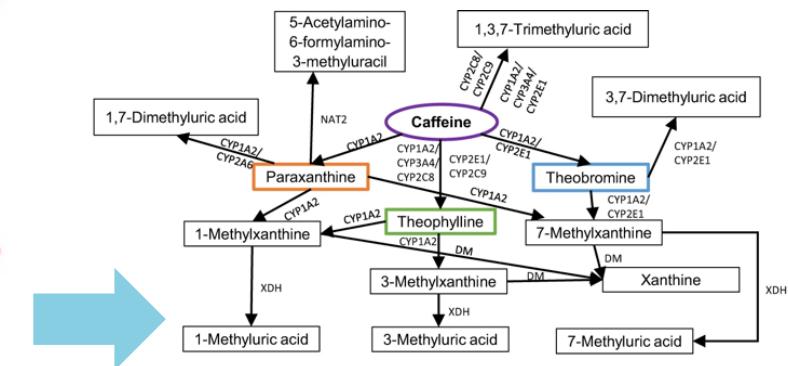
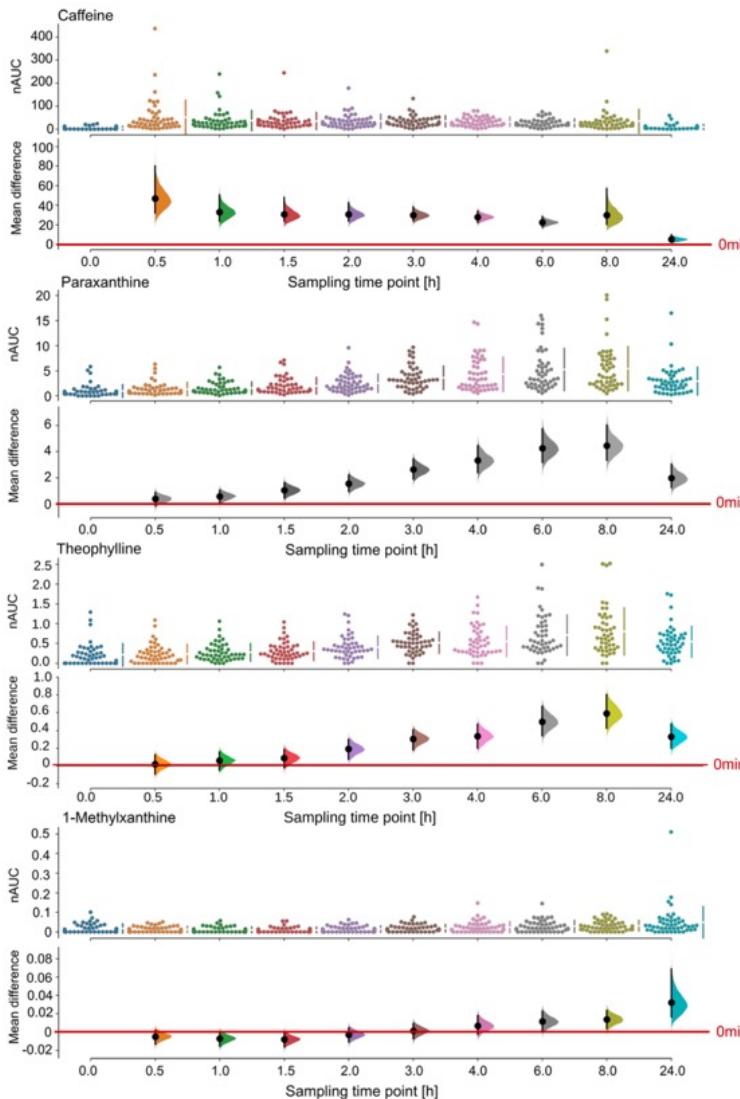
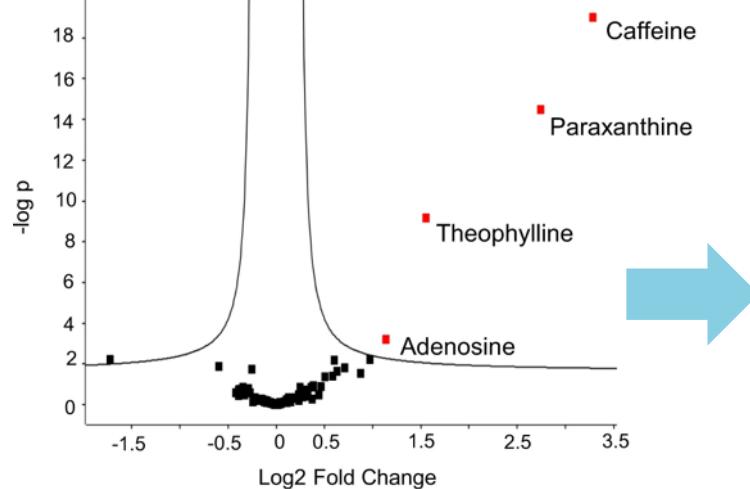


Finger sweat reflects caffeine metabolism

🤔 Why caffeine: excellent oral bioavailability

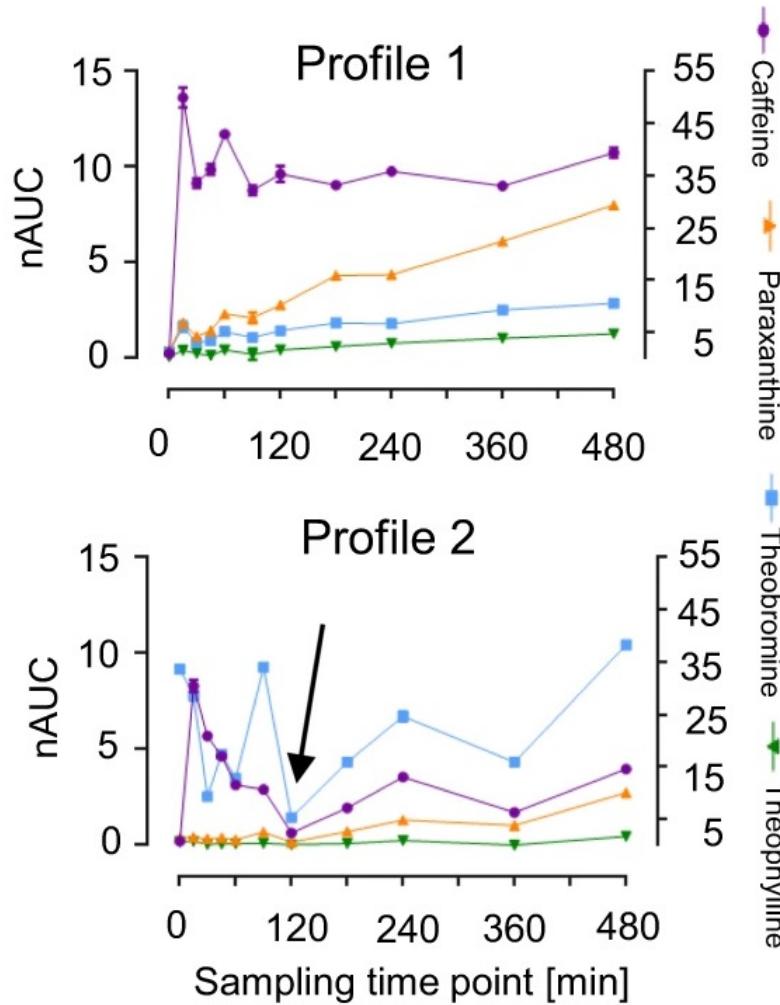
Study B: espresso or control group
Study C: Caffeine capsule

4hr later

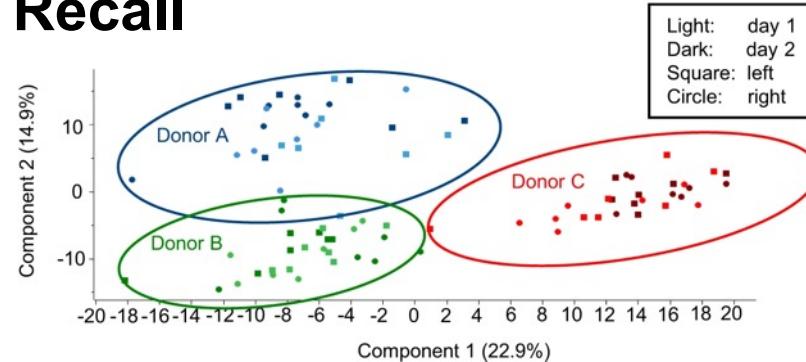


Align with what's already known

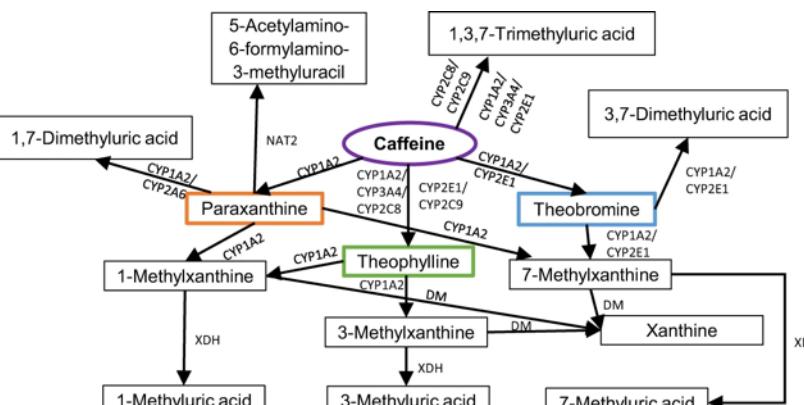
Finger sweat elucidates individual metabolic traits



Recall



Hepatic enzymes participate



caffeine elimination → a proxy for liver function

Study C: Caffeine capsule

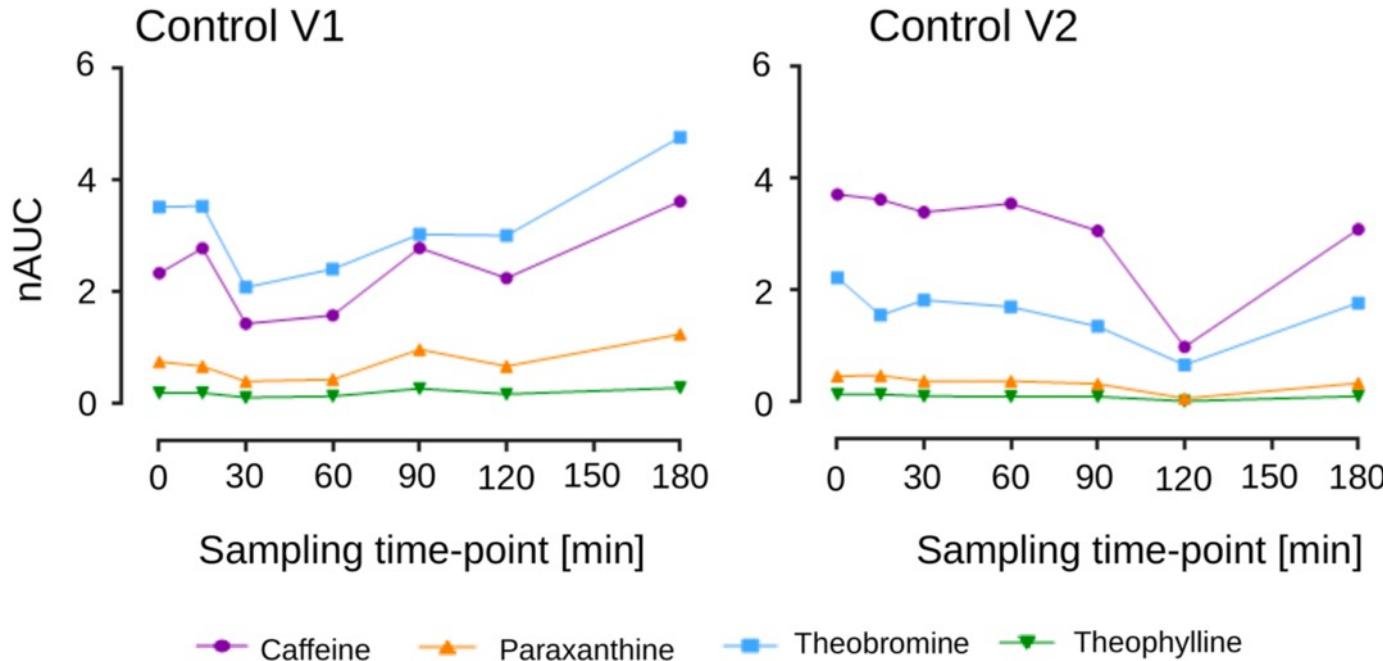
May tell:
Cirrhotic or normal livers

But for accurate diagnosis,
QUANTIFICATION
is required

Challenge: fluctuations in the rate of sweat



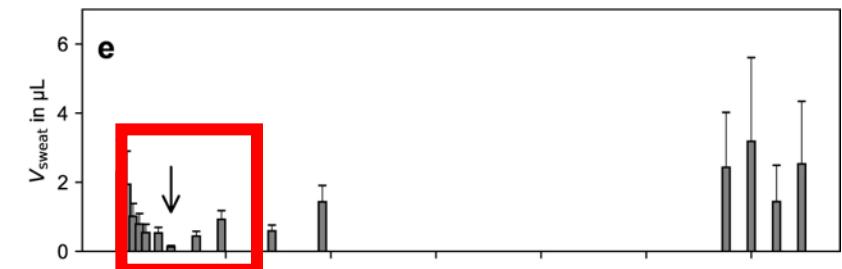
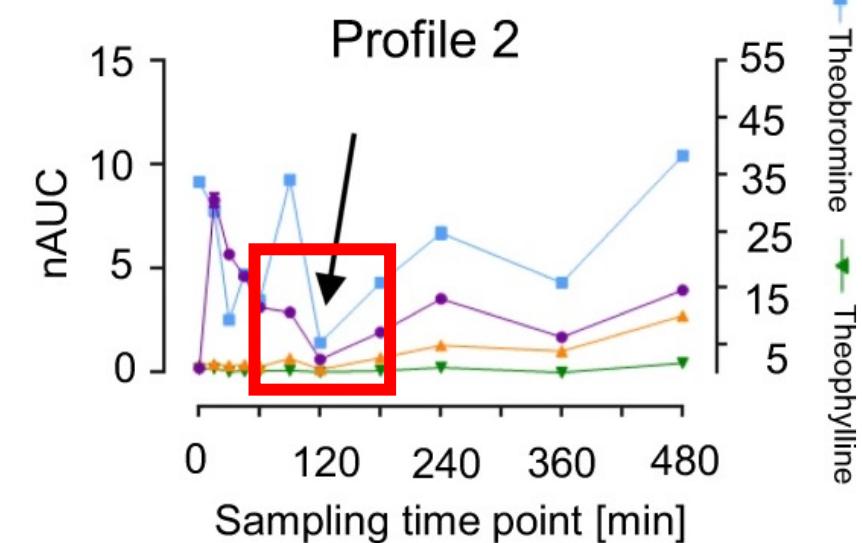
Even control group, nAUC vary with time



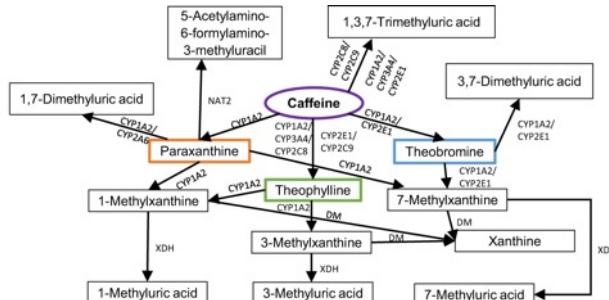
Fluctuate!

$$C \times V_{\text{sweat}} = \text{Mass}$$

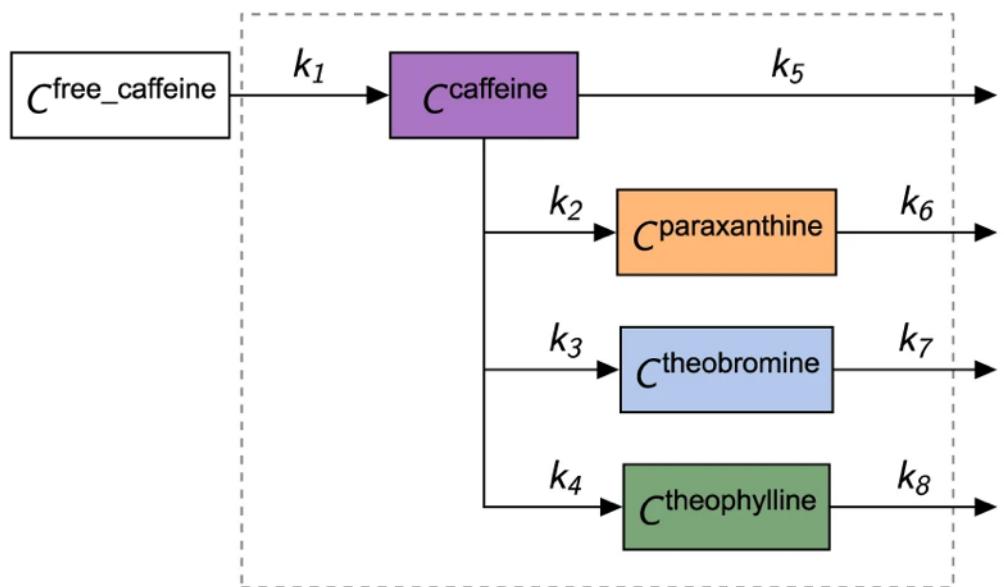
$$\text{Mass} / V_{\text{sample}} = C_{\text{sample}} \rightarrow \text{nAUC}$$



Kinetics model is built to obtain V_{sweat}



Abstract relations

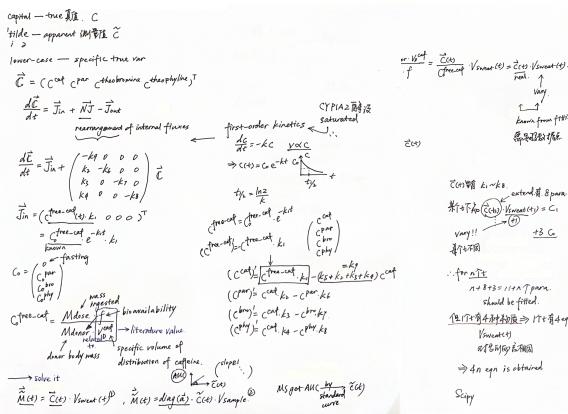


Assumptions

- ## ➤ first-order kinetics

$$\frac{dC}{dt} = -kC$$

After some calculation..



- list and solve the differential equations for the 4 metabolites in sweats
 - get the relation
 $C = f(t, k_{1-8})$

Key points

$$\text{Fit} \quad \tilde{\mathbf{M}}(t) = \mathbf{C}(t)V_{\text{sweat}}(t),$$

Known $\tilde{\mathbf{M}}(t) = \text{diag}(\mathbf{a})\tilde{\mathbf{C}}(t)V_{\text{sample}}$.

$$C_0^{\text{free-caf}} = \frac{M_{\text{dose}}}{M_{\text{donor}}} \frac{f}{v_D^{\text{caf}}},$$

Boundary Conditions fasting

$$\mathbf{C}_0 = \begin{pmatrix} 0 \\ C_0^{\text{par}} \\ C_0^{\text{bro}} \\ C_0^{\text{phy}} \end{pmatrix}$$

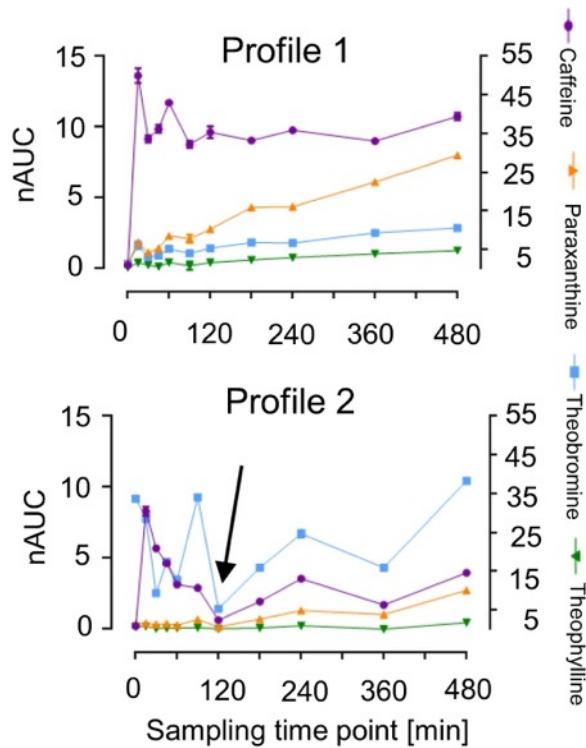
★ Get k_{1-8} , $V_{sweat}(t)$ (for each time point)

Evaluate the kinetics model



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Fit well

