



DPK2025 - Bergfest in Berlin

Uncovering daily symptoms: **long-term digital phenotyping and EMA** in patients undergoing cognitive behavioral therapy for internalizing disorders (SP6)

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FOR SP6/ PREACT-digital

Can we depict symptoms in everyday life and predict therapy non-response using digital phenotyping and EMA?
→ focus on emotion regulation



FOR SP6/ PRACT-digital

Can we depict symptoms in everyday life and predict therapy non-response using digital phenotyping and EMA?

- ⇒ Digital Phenotyping and EMA in Mental Health
- ⇒ PRACT-digital study (SP6)
- ⇒ DP in Psychotherapy? Feasibility and Engagement
- ⇒ Ongoing study





Background: DP & EMA in Mental Health

Background

Collect data in the patients' **everyday life**

Digital Phenotyping (DP)

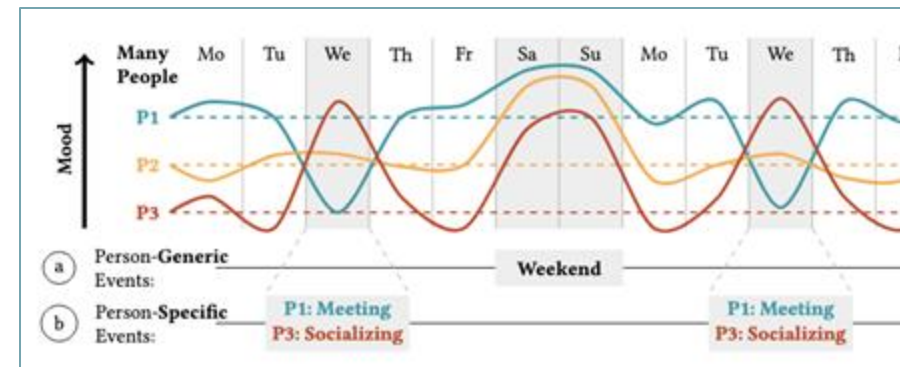
- **passive**
- via wearables (i.e. Smartwatch) or smartphones



- low burden
- allows long-term continuous, real time data collection

EMA (*Ecologically momentary assessment*)

- **active**, i.e. user input required
- short, repeated self-reports on the smartphone
- high temporal resolution, no retrospective bias
- catches fluctuations in real time (i.e. mood)



Background

Uncover **daily symptoms** in internalizing disorders using **DP** and **EMA**

Internalizing Disorders

- most common: diagnosis of depression (i.e. RADAR study)
- i.e. time at home predicts change in PHQ-8 scores (Zhang et al., 2022)
- Large heterogeneity in study designs and outcomes; many shortcomings
- **DP** in its **infancy**!

Psychotherapy

- no studies predicting **psychotherapy outcomes** using DP!
- De Angel et al (2023): feasibility of DP in patients doing psychotherapy
 - good adherence; declines when therapy starts; but small sample
- need for more studies on DP during psychotherapy in internalizing disorders!



PREACT-digital

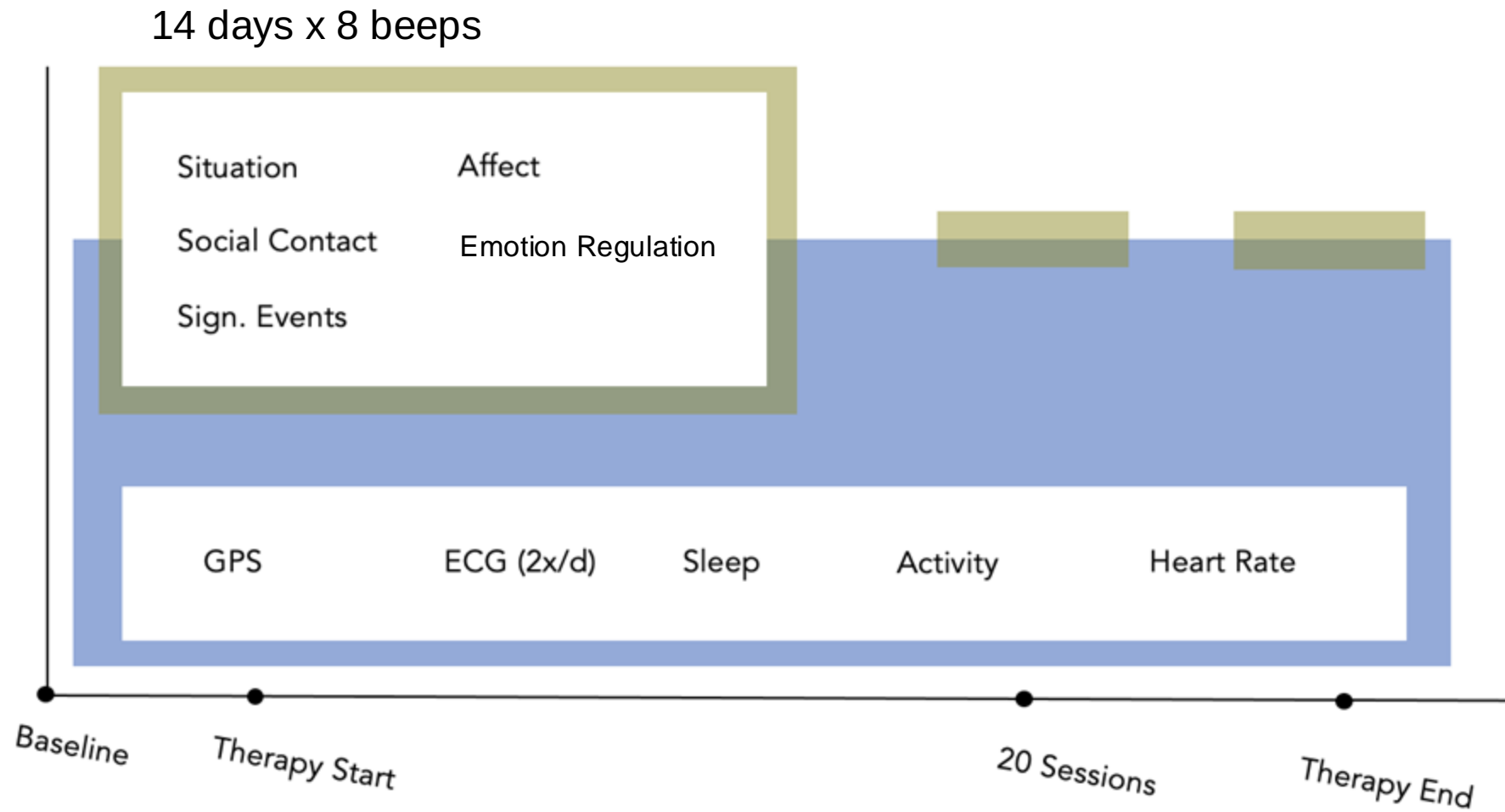
Study procedures

TIKI App

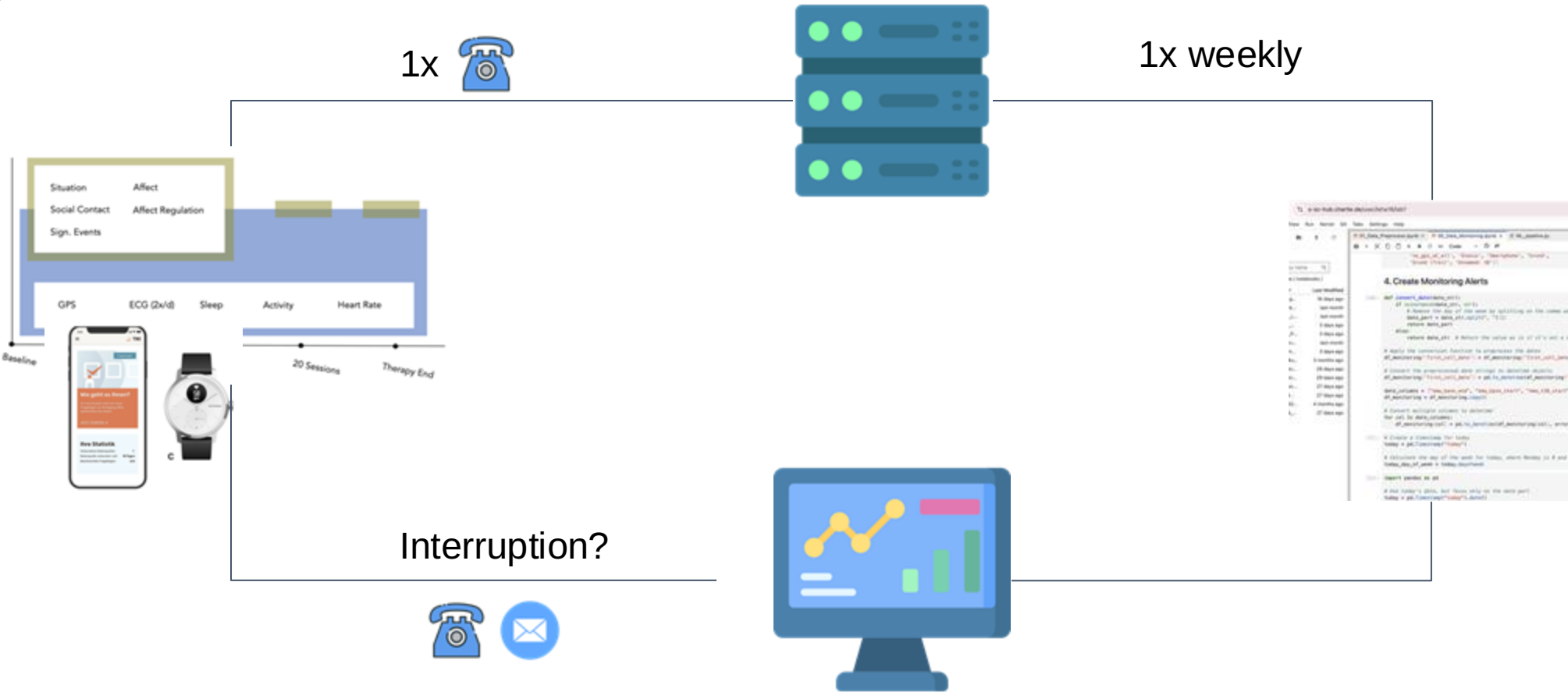


Withings
Scanwatch

Study procedures



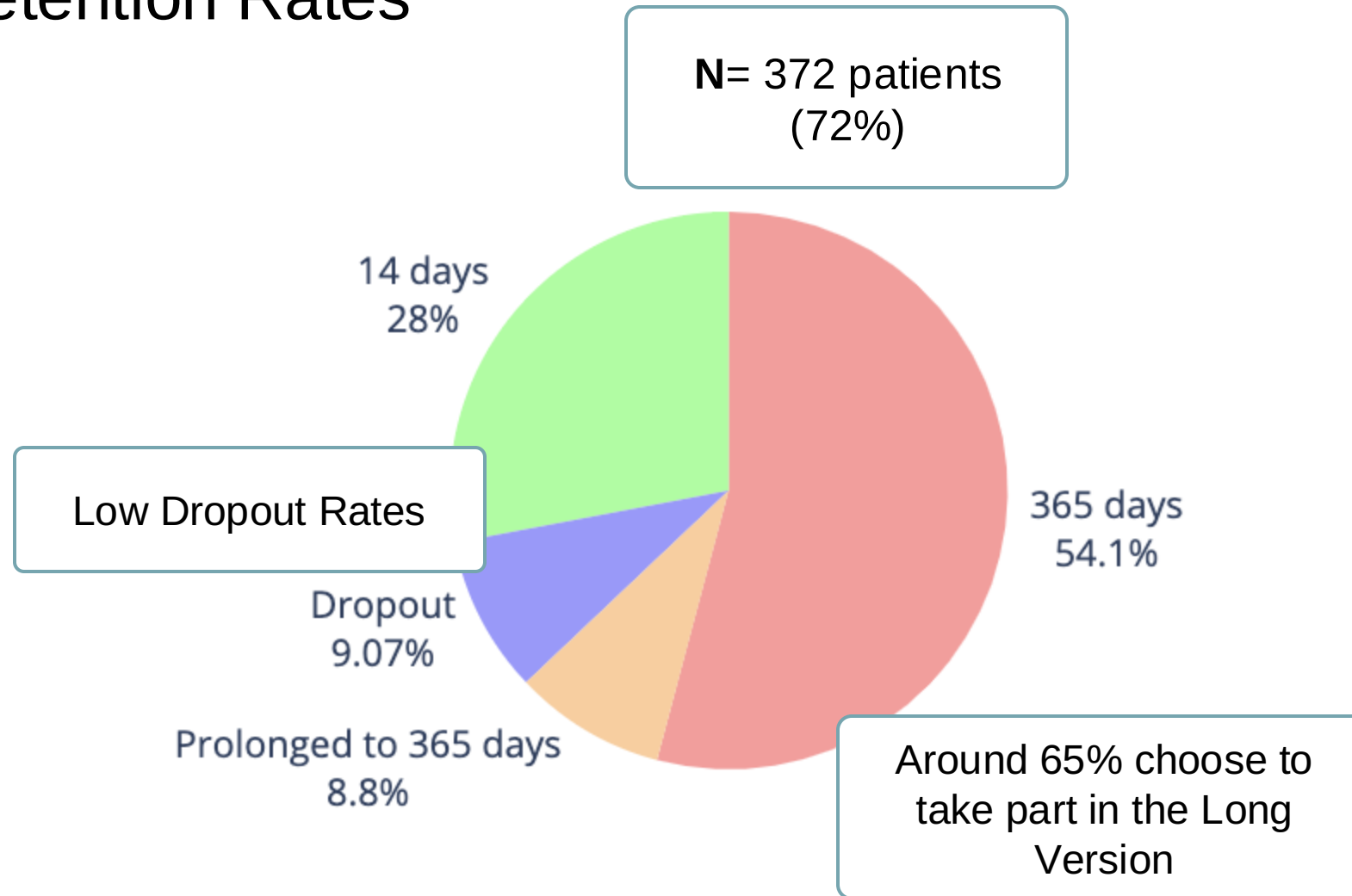
Study





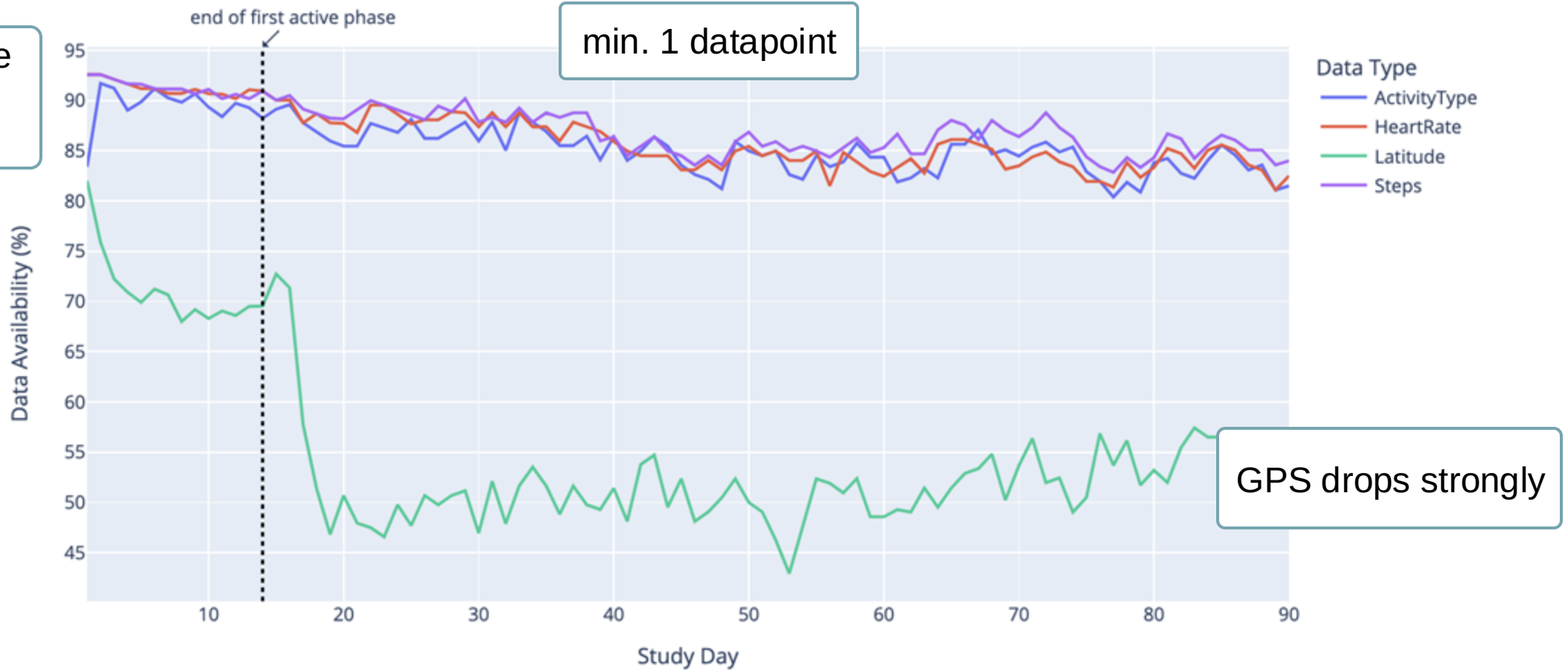
Feasibility & Adherence

Retention Rates

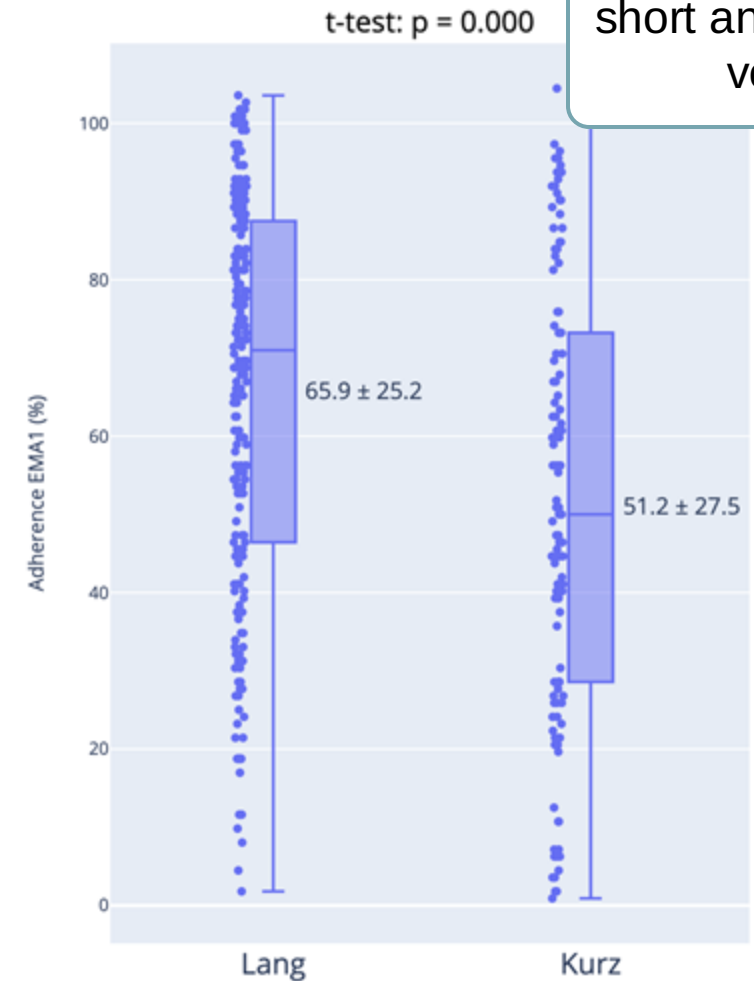
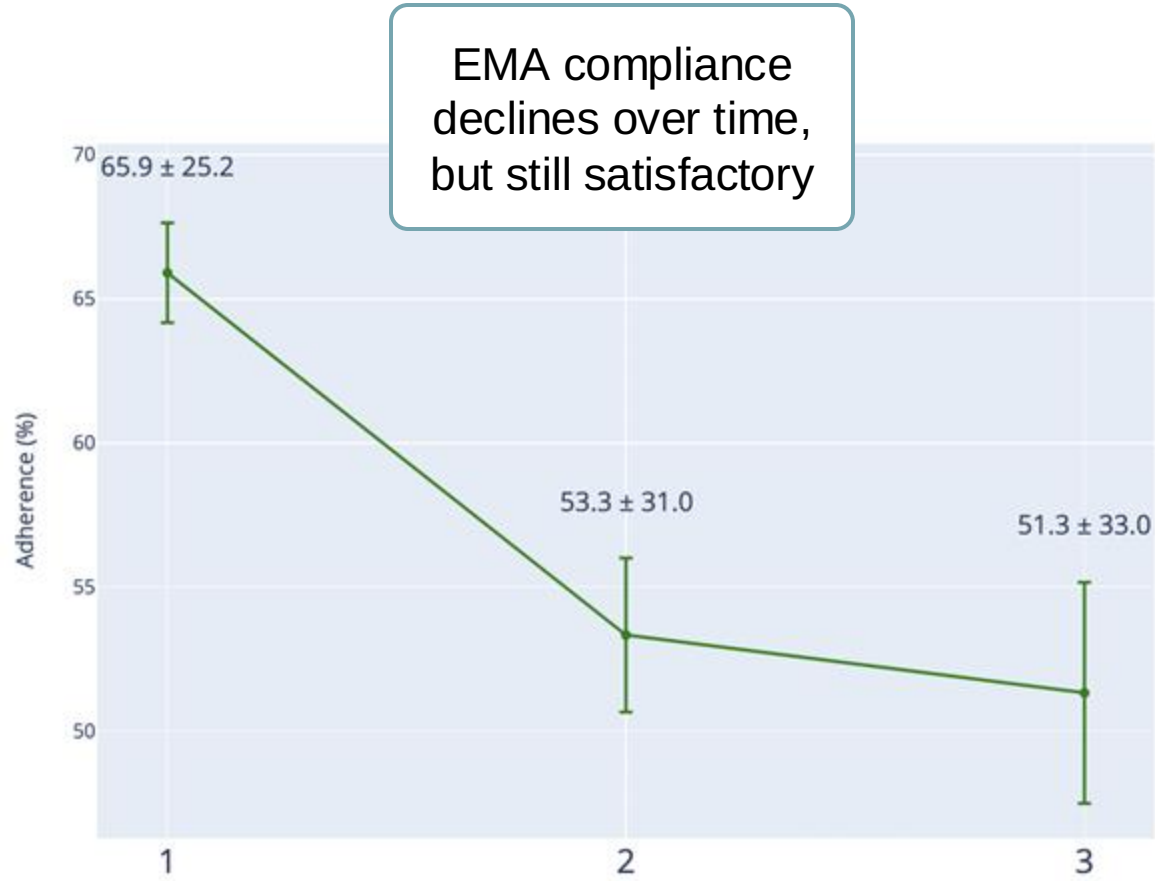


Adherence: DP

slight decrease
for wearable
data



Engagement: EMA



Discussion Feasibility

Conclusion

- lessons learned: choose study-app with better maintenance-service
- but: Long-term DP and EMA are feasible in clinical populations, also during psychotherapy!

Passive

↑ High long-term adherence
for Smartwatch-derived data;
adherence comparable to
previous studies

↓ Lower adherence for GPS
data due to technical
constraints

Active

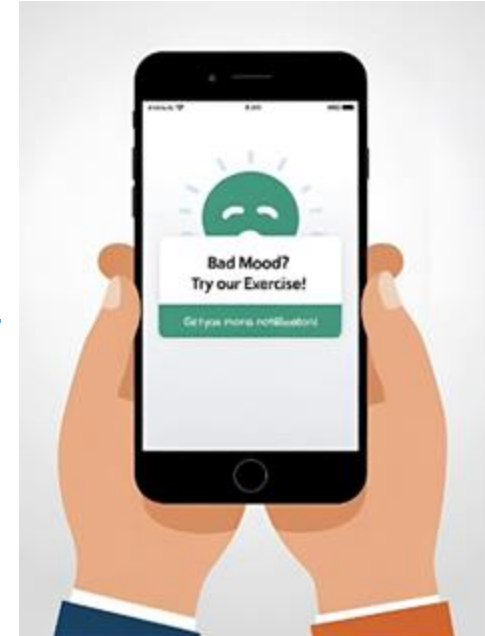
→ Sufficient adherence across
study phases

→ difference between short vs.
long version = therapy
motivation?

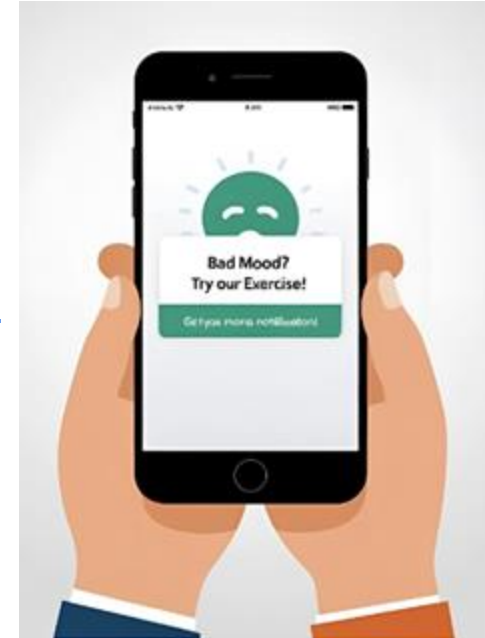


Ongoing study

Towards JITAI



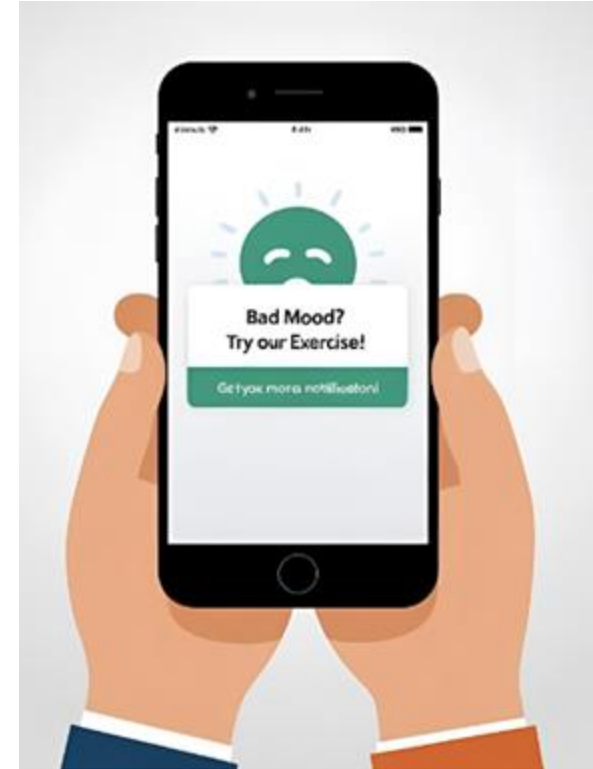
Towards JITAI



Towards JITAI

Can we predict **short-term** fluctuations in negative affect with **entirely passive data**?

→ compare **subject-independent** with **subject-dependent** machine learning (“personalized ML”)



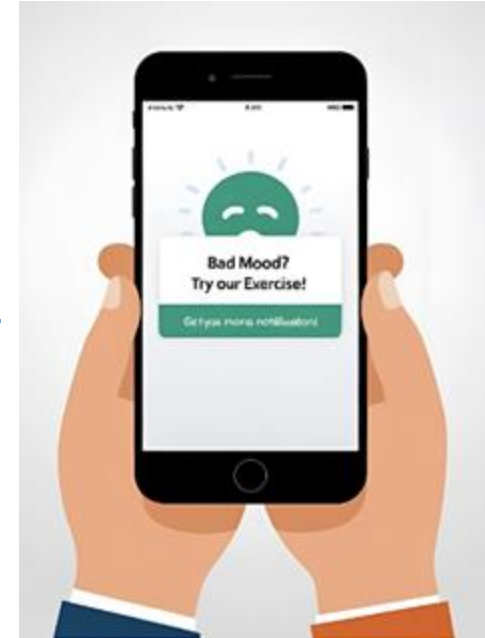
Towards JITAI



2 hours,
passive data



personalized ML

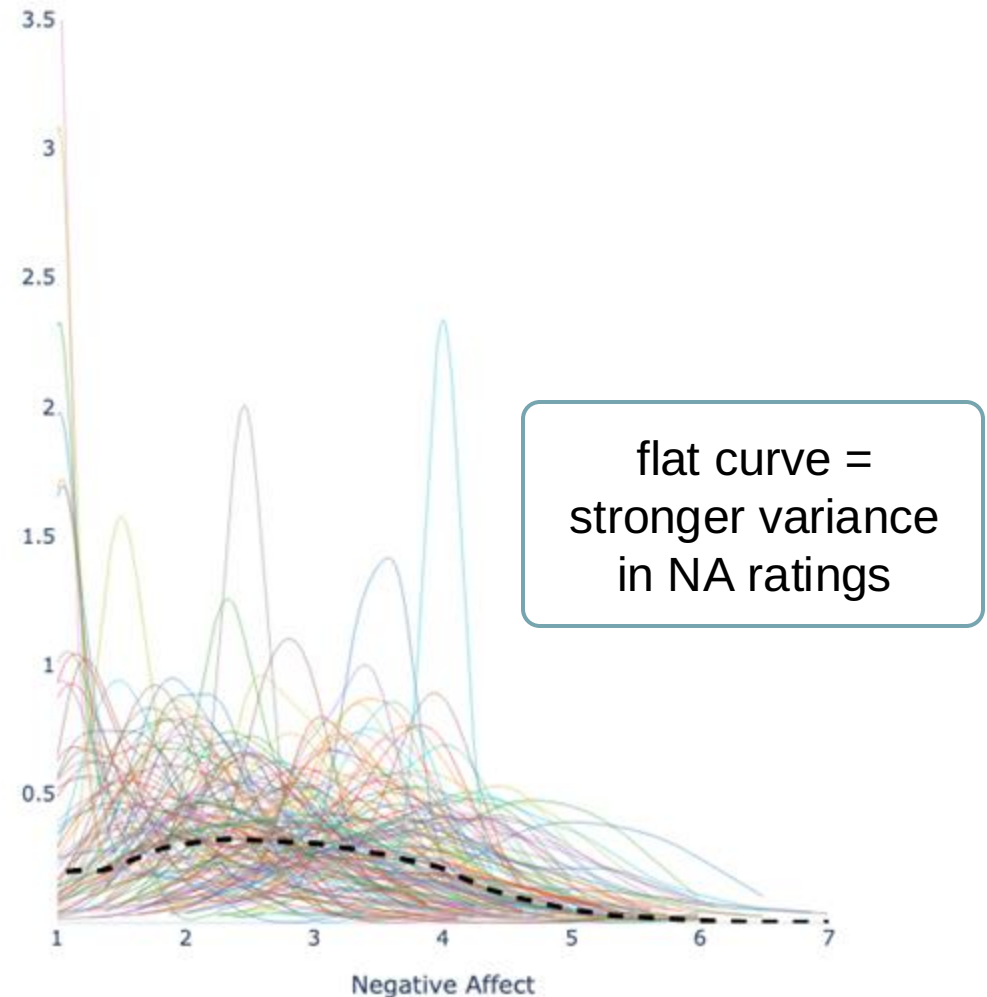


predict short-term
negative affect

Short-term prediction of negative affect

Can we predict **short-term** fluctuations in negative affect with **entirely passive data**?

→ high variation in EMA-derived negative affect ratings within and between participants



Short-term prediction of negative affect

Can we predict **short-term** fluctuations in negative affect with **entirely passive data**?

→ high variation in EMA-derived negative affect ratings within and between participants

→ **personalized ML** models perform best; however, best models achieve similar errors as the average NA per person.

→ passive data alone do not contain enough signal!

Model	MAE ^a	R ² ^b	RMSE ^c
Global Intercept	.878	-.007	1.053
Per Person Intercept	.631	.522	.818
LR	..964	.010	1.177
RF	.959	.015	1.174
FFNN	.984	-.015	1.192
LR + PS	.944	.051	1.153
RF + PS	.912	.108	1.117
FFNN + PS	.968	.011	1.177
MERF	.609	.525	.816
MERF + PS	.614	.522	.818
FFNN + Embedding	.608	.532	.809

Thank you!

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Study Protocol SP6



Preregistration



Literature

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