



UNIVERSITY  
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# Ecological momentary assessment (EMA): examining behaviours and processes in daily life

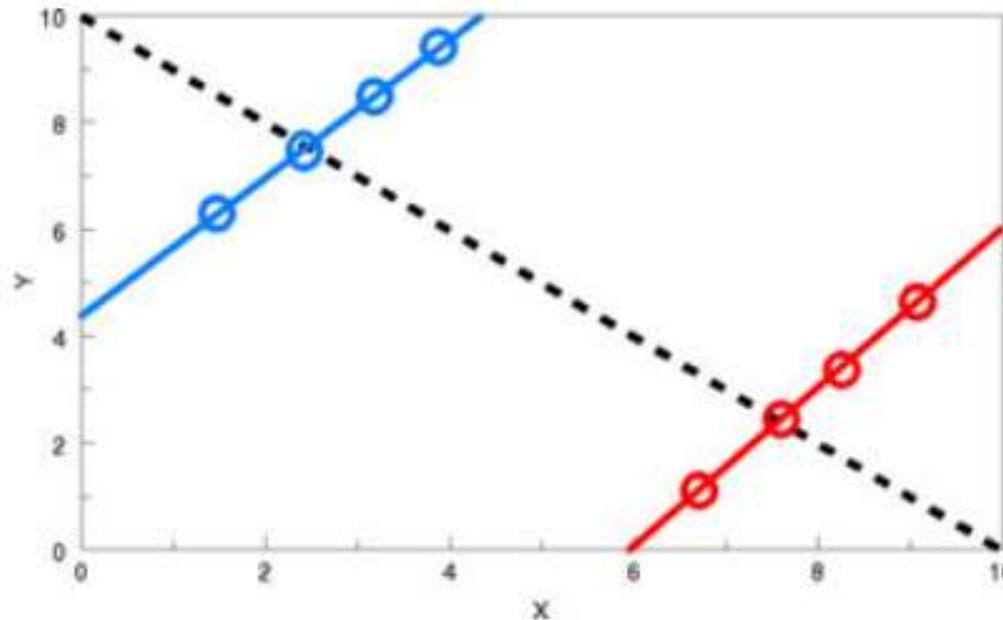
Dan Powell, University of Aberdeen

12<sup>th</sup> October, 2018

# Why use EMA? Paradigmatic Rationale

**THINKING WITHIN-PERSON**

# Thinking Within-Person versus Between-Person

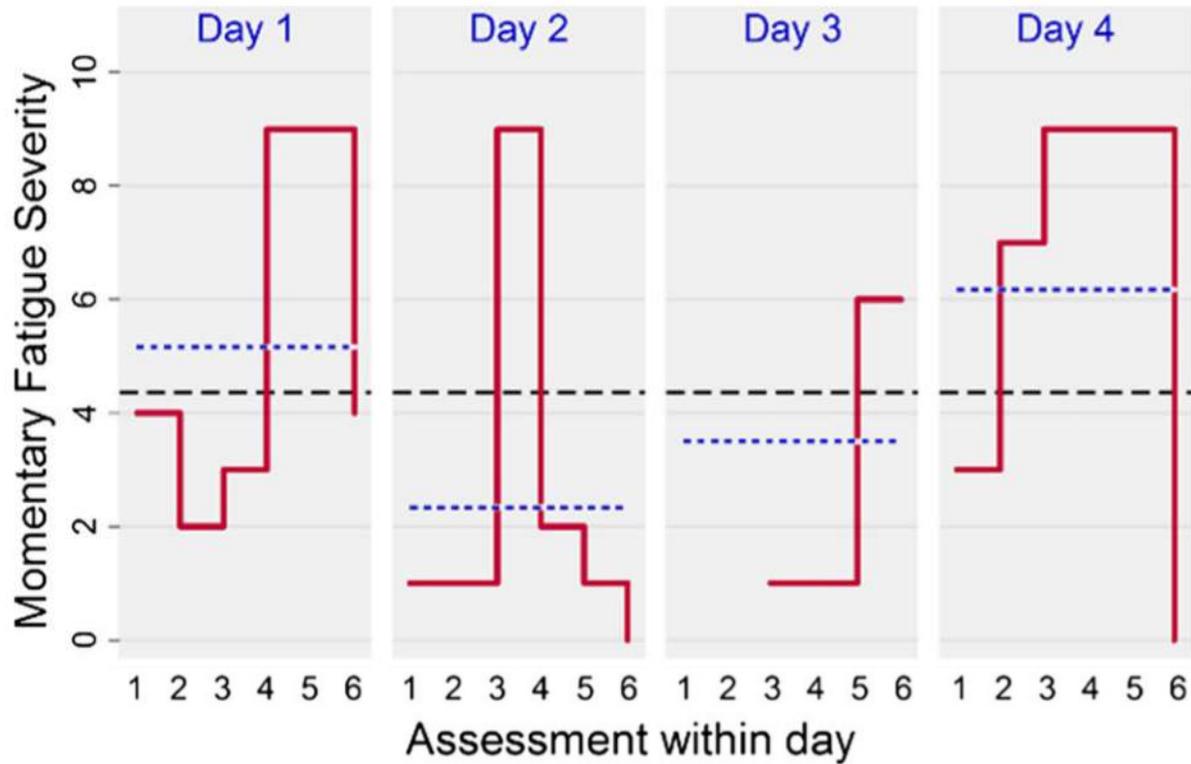


**Figure 1.** Illustration of a possible result of testing the relationship between variables  $x$  and  $y$  where the between-result (dotted line) is negative, but the result for individuals studied over days or locations (illustrated by continuous lines for two people) is positive.

Johnston & Johnston (2013) *British Journal of Health Psychology*.

### Participant A

Mean = 4.4; Med = 4.0; MSSD = 19.1; PAC = 0.38; Proportion  $\geq 5 = 0.41$



$$x_{it} = M_i + e_{it}$$

$x_{it}$  = score for individual  $i$  at time  $t$ .

$M_i$  = mean score for individual  $i$

$e_{it}$  = deviation from mean for individual  $i$  at time  $t$

Powell, Lioffi, Schlotz, & Moss-Morris (2017). Tracking daily fatigue fluctuations in multiple sclerosis: ecological momentary assessment provides unique insights. *Journal of Behavioral Medicine*, 40(5), 772-783.

# Practicalities: centring your predictors

		Person mean of all Score_S for individual	Overall mean of all PersonMean_S in sample	Score_S – PersonMean_S	PersonMean_S – GrandMean_S
ID	Score_S	PersonMean_S	GrandMean_S	Within_S	Between_S
1	7	4	5	3	-1
1	3	4	5	-1	-1
1	1	4	5	-3	-1
1	5	4	5	1	-1
2	9	6	5	3	1
2	2	6	5	-4	1
2	6	6	5	0	1
2	7	6	5	1	1

See Curran & Bauer (2011)

@danpowell83



# SNAPSHOT project

**S**NAcking, **P**hysical activity, **S**elf-regulation, and  
**H**eartrate **O**ver **T**ime



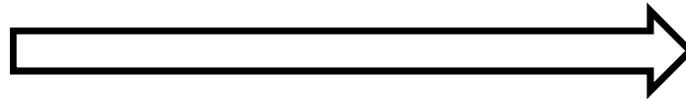
# SNAPSHOT project

**SNA**cking, **P**hysical activity, **S**elf-regulation, and  
**H**eartrate **O**ver **T**ime

# Health behaviours and self-control

Unhealthy behaviours often have significant long-term costs but immediate benefits

(Hall & Fong, 2007; 2015)



**TODAY**

**6 MONTHS LATER**

# Executive function & health behaviour

Exerting self-control over behaviour in the face of temptation places heavy demand on top-down cognitive processes known as the *executive functions*

Three core facets:

- Inhibitory Control
- Working Memory
- Set Shifting



# Behavioural Theories and EF



Several theories of health behaviour and conceptual models of self-regulation posit EF as an important determinant of health-relevant behaviour

For example:

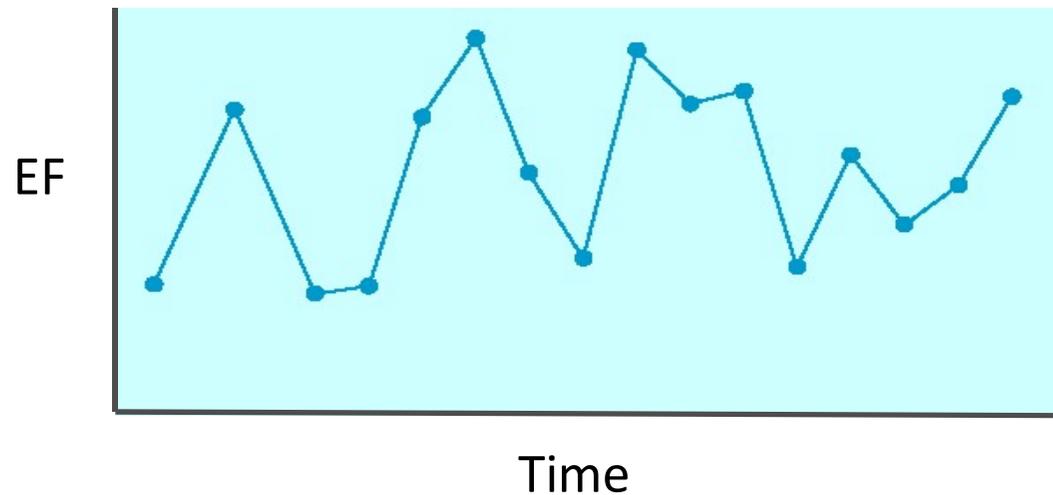
- Temporal Self-Regulation Theory (Hall & Fong, 2007, 2013)
- Reflective Impulsive Theory (Strack & Deutsch, 2004; Strack et al., 2014)
- Models of self-control (Baumeister & Vohs, 2007; Inzlicht & Schmeichel, 2012)

# Which EF facet?

## Inhibitory control seems most relevant to unhealthy eating and obesity

- Poor performance on cognitive tests assessing inhibitory control (e.g. Go/No-Go) associated with weaker control of food intake in the lab; particularly of high-fat foods (Allom & Mullan, 2014; Hall, 2012; Hall, Lowe & Vincent, 2014; Limbers & Young, 2015)
- Obese adults and children show marked inhibitory deficits relative to controls (Lavagnino et al., 2016) – impaired inhibition a “critical feature” of obesity
- Other facets (e.g. planning skills, updating) related with initiation of healthy eating behaviours such as fruits and vegetables (Allom & Mullan, 2014; Limbers & Young, 2015)

# Studies are focussed on individual differences

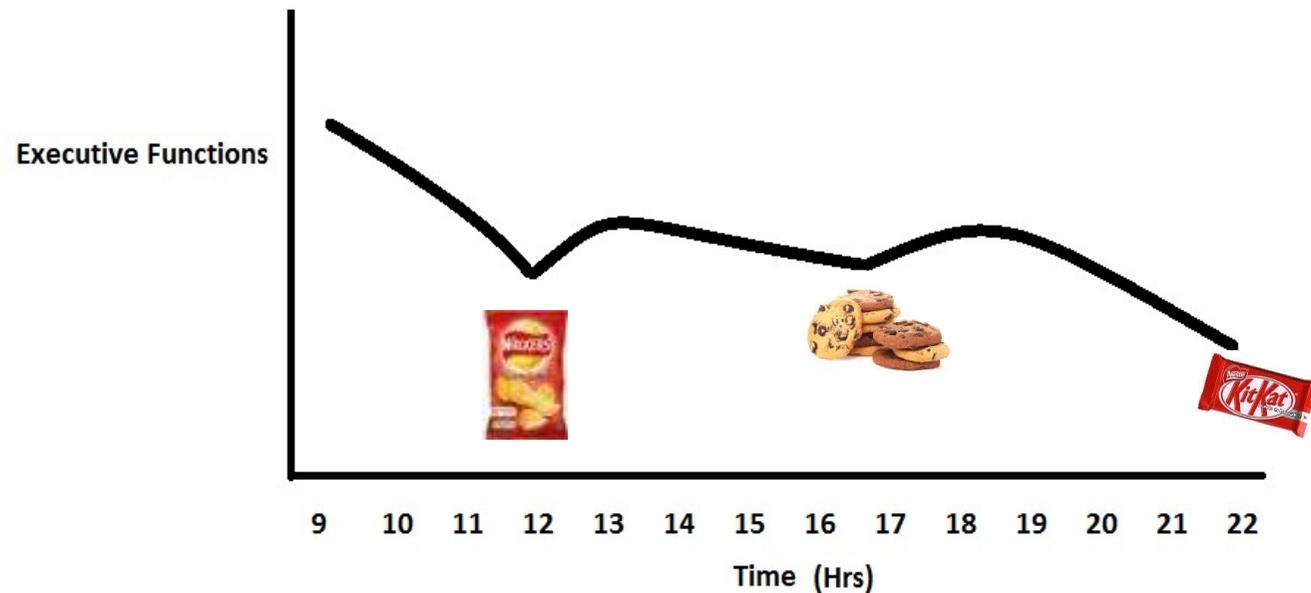


.....but all models imply within-person variability!

EF at any given time important to whether impulsive action occurs or not

# Main Hypothesis

Within-person: when EF is poorer than usual, initiation and consumption of energy-dense snacks will be higher than usual



# Methodological Challenge



How to assess objective fluctuations in inhibitory control efficiency “in the wild”?

# Methodological Challenge: Solution



In collaboration with CamnTech, we developed a Go/No-Go Test to deploy via PRO-Diary devices

# Go/No-Go Test

Testing inhibitory control efficiency

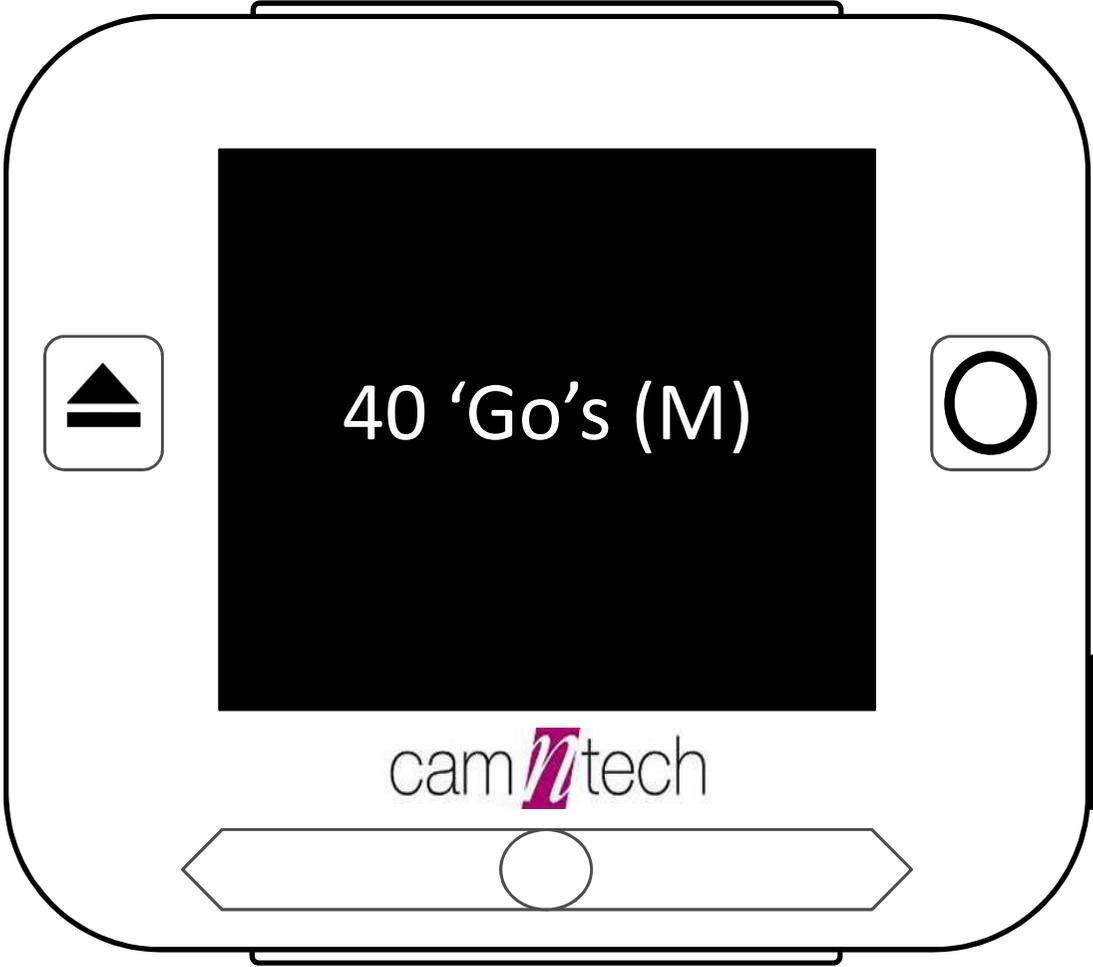
# Go/No-Go task



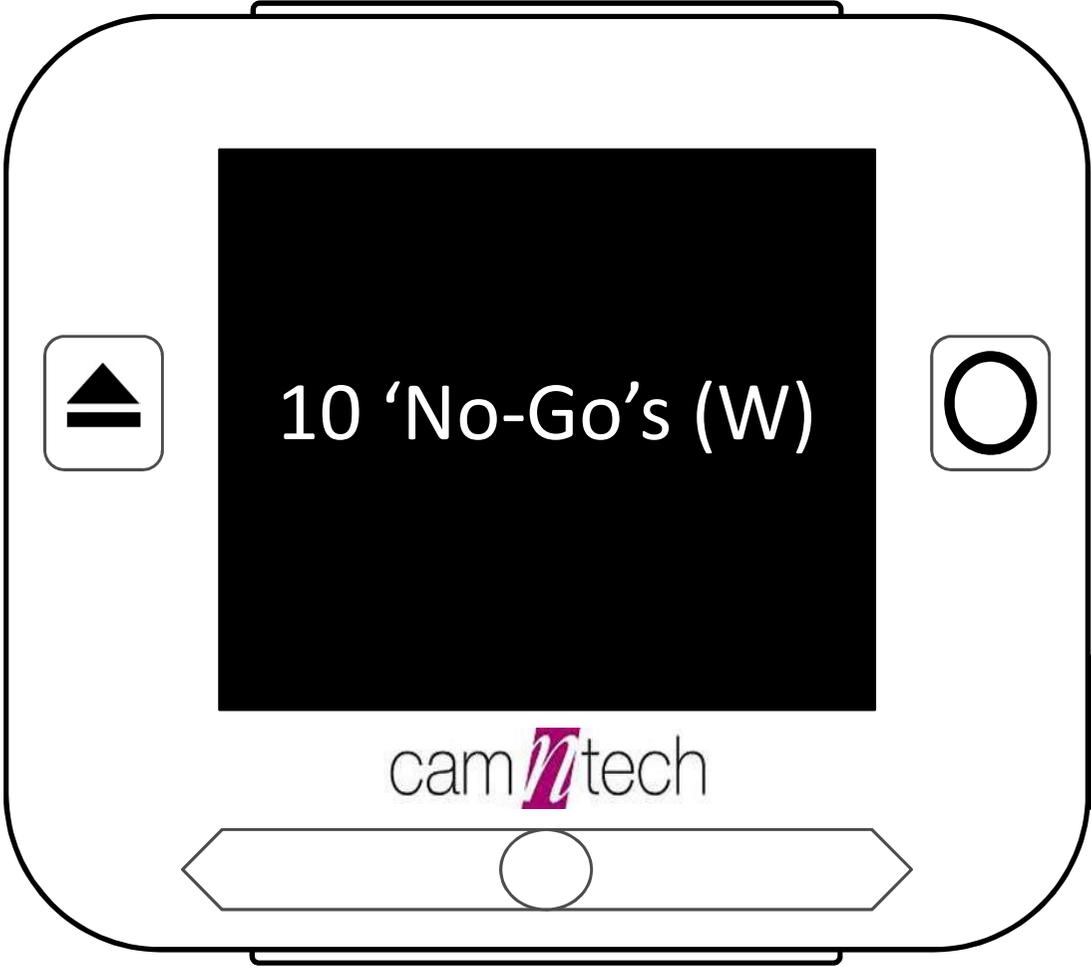
# Go/No-Go task



# Go/No-Go task



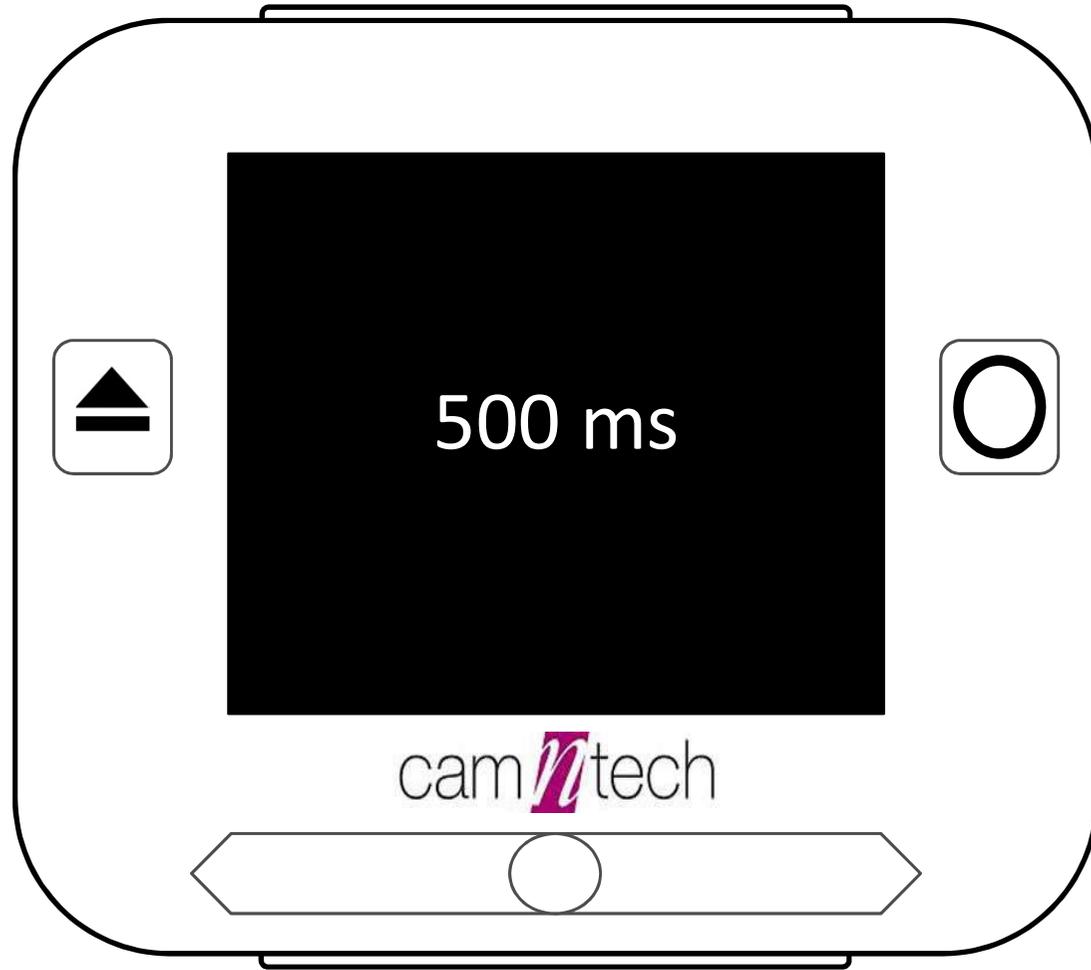
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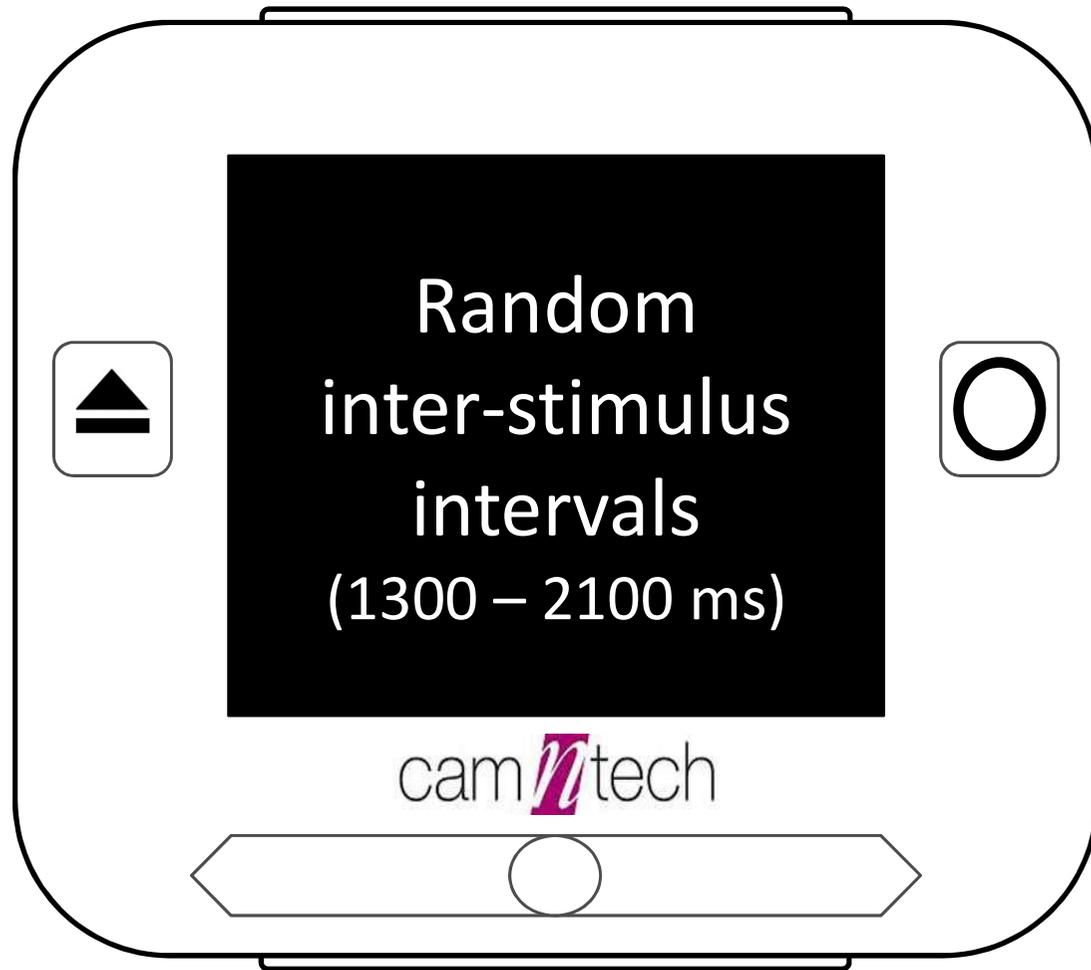
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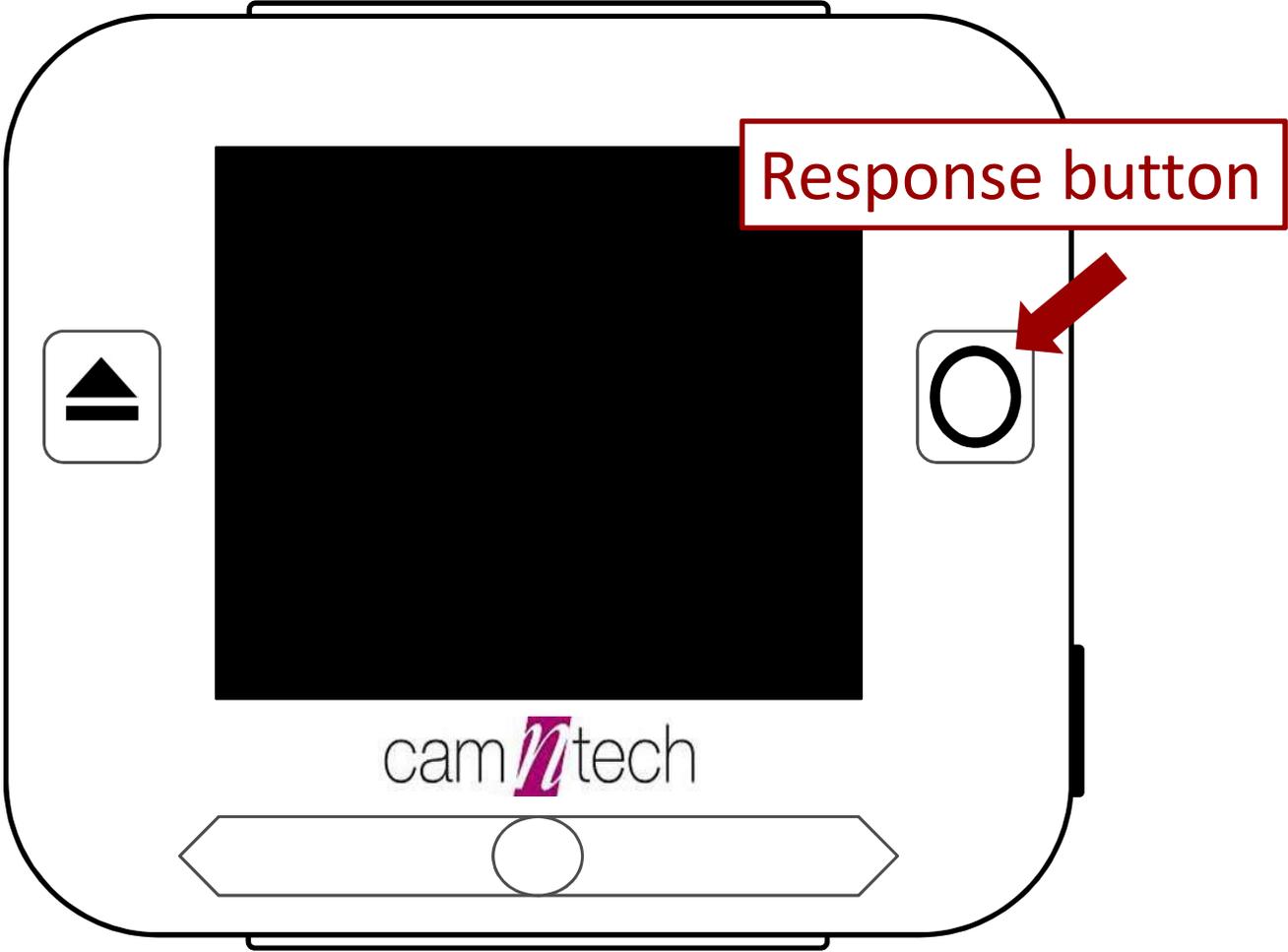
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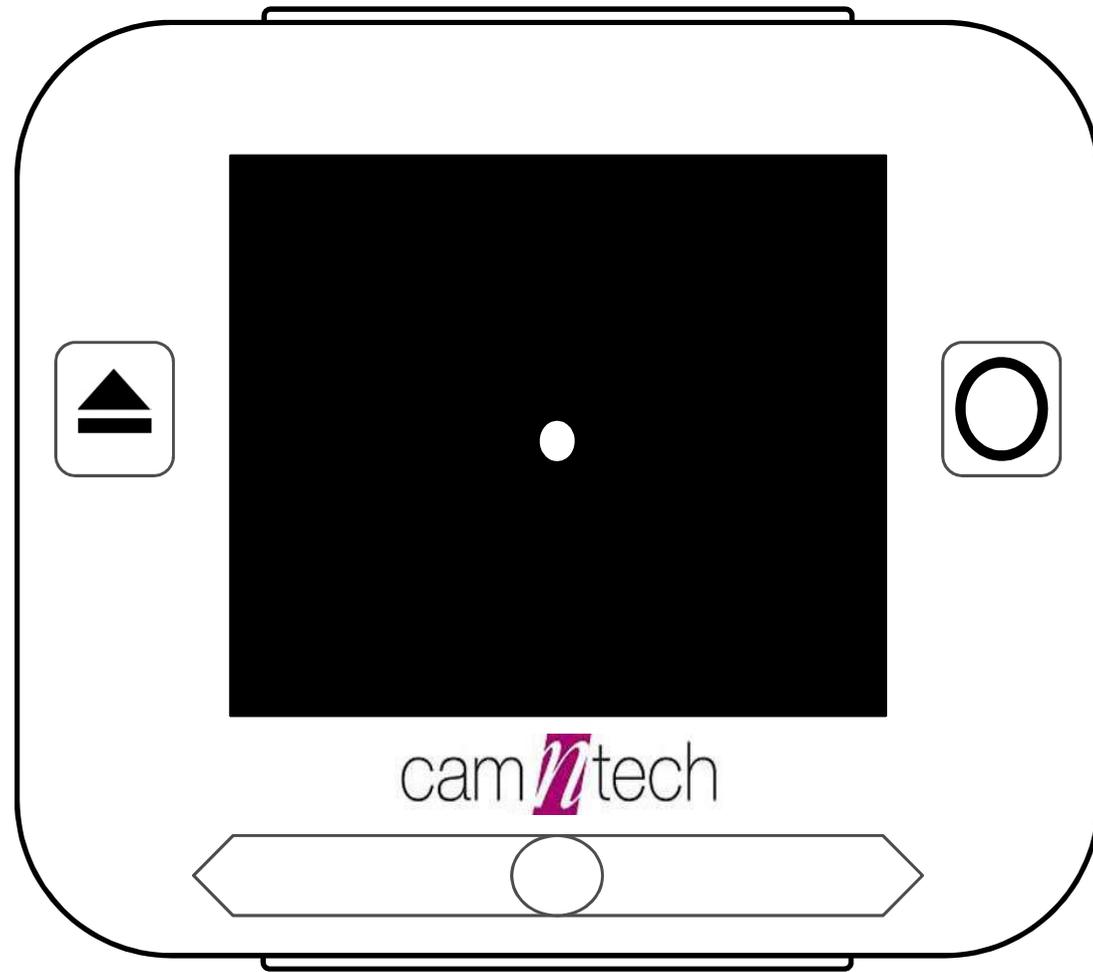
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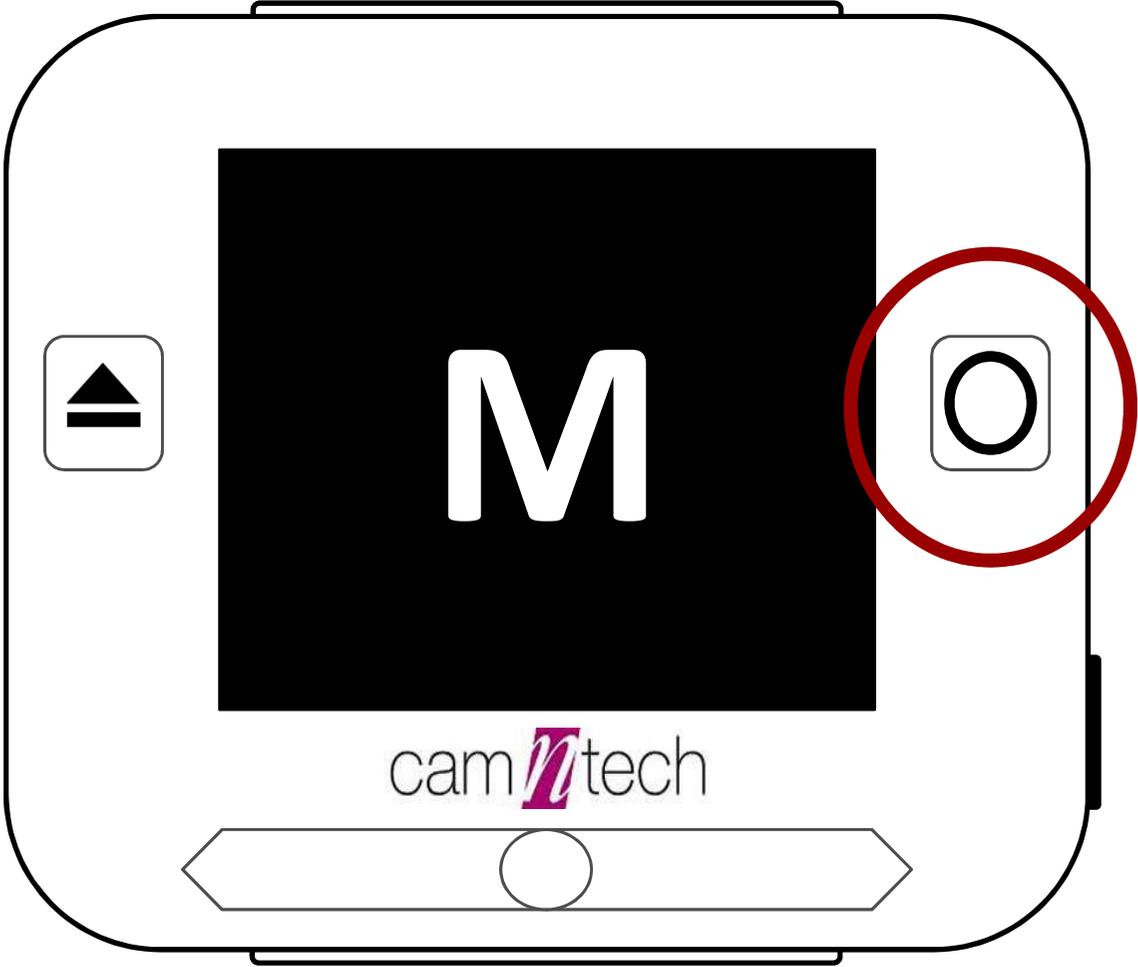
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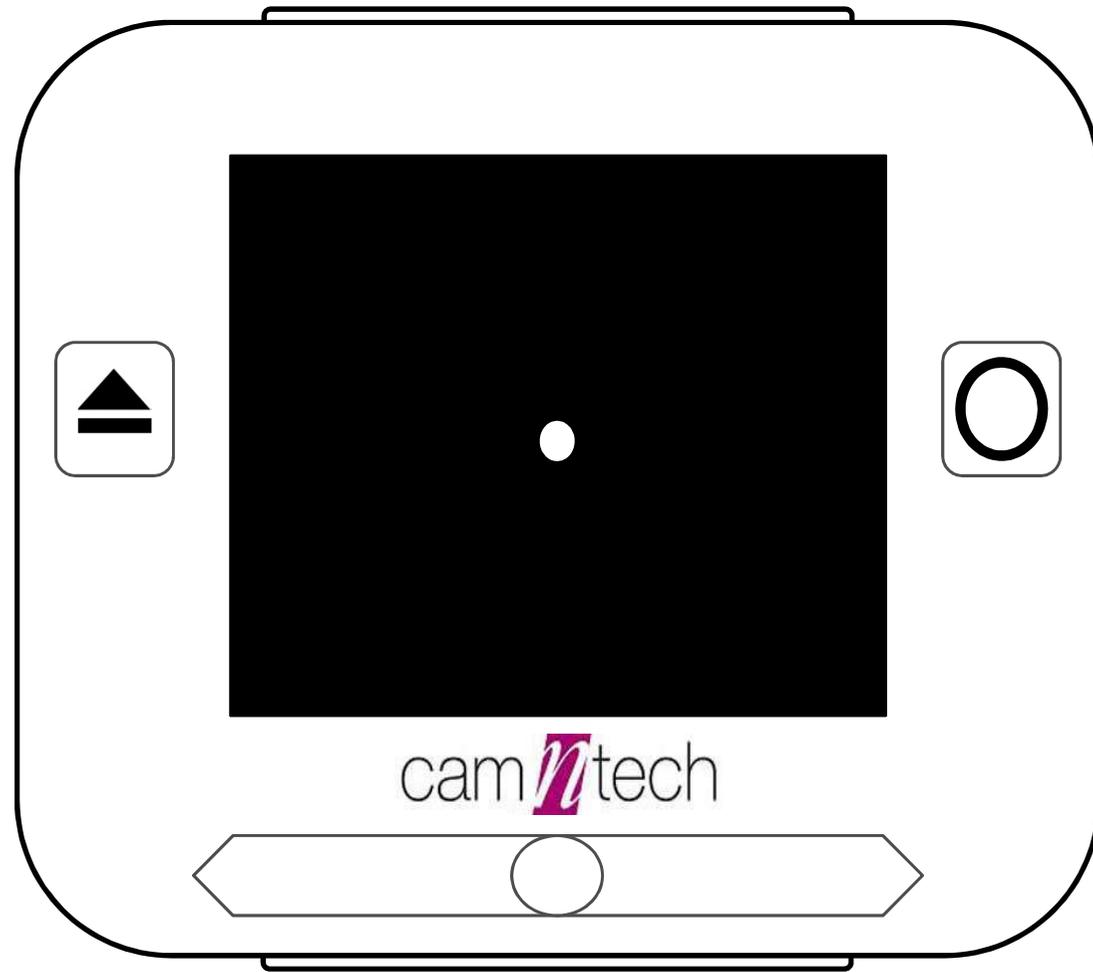
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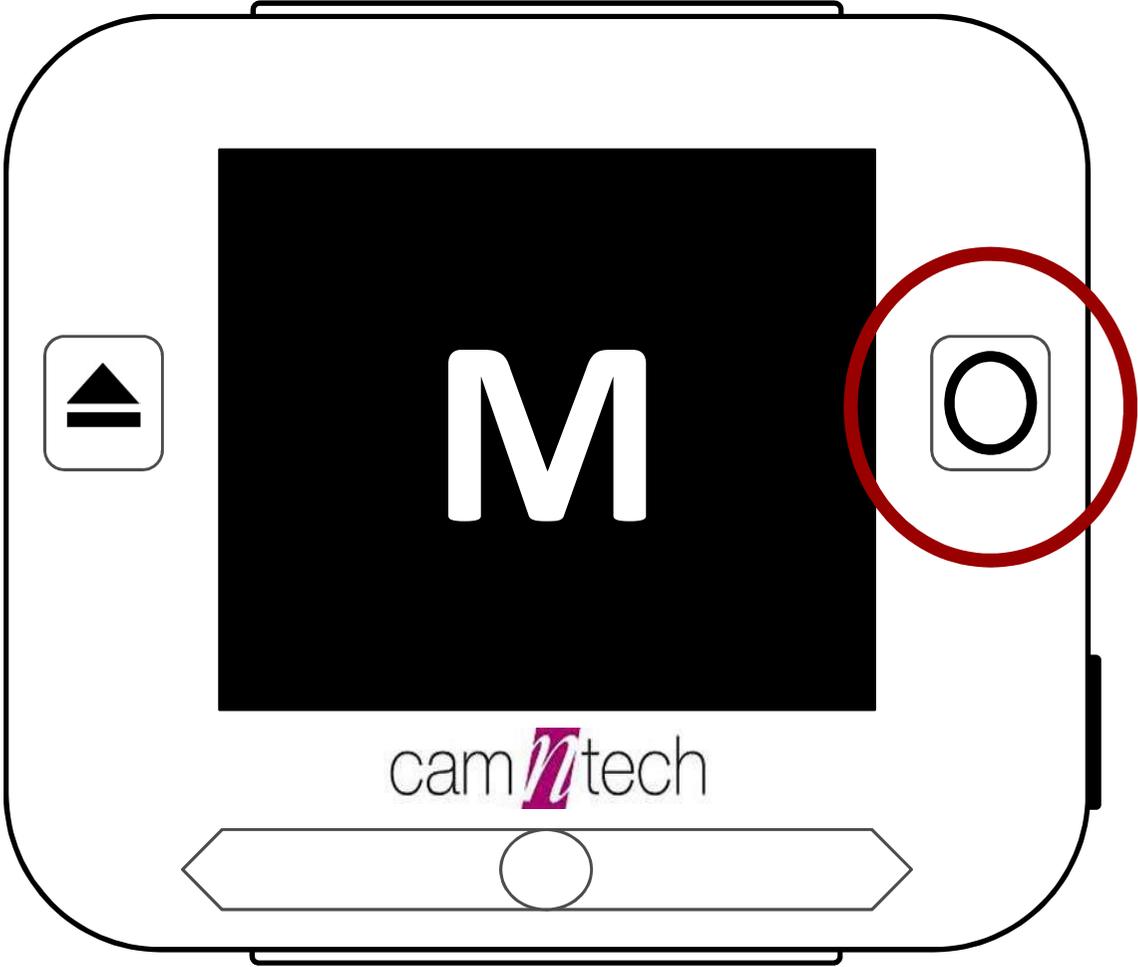
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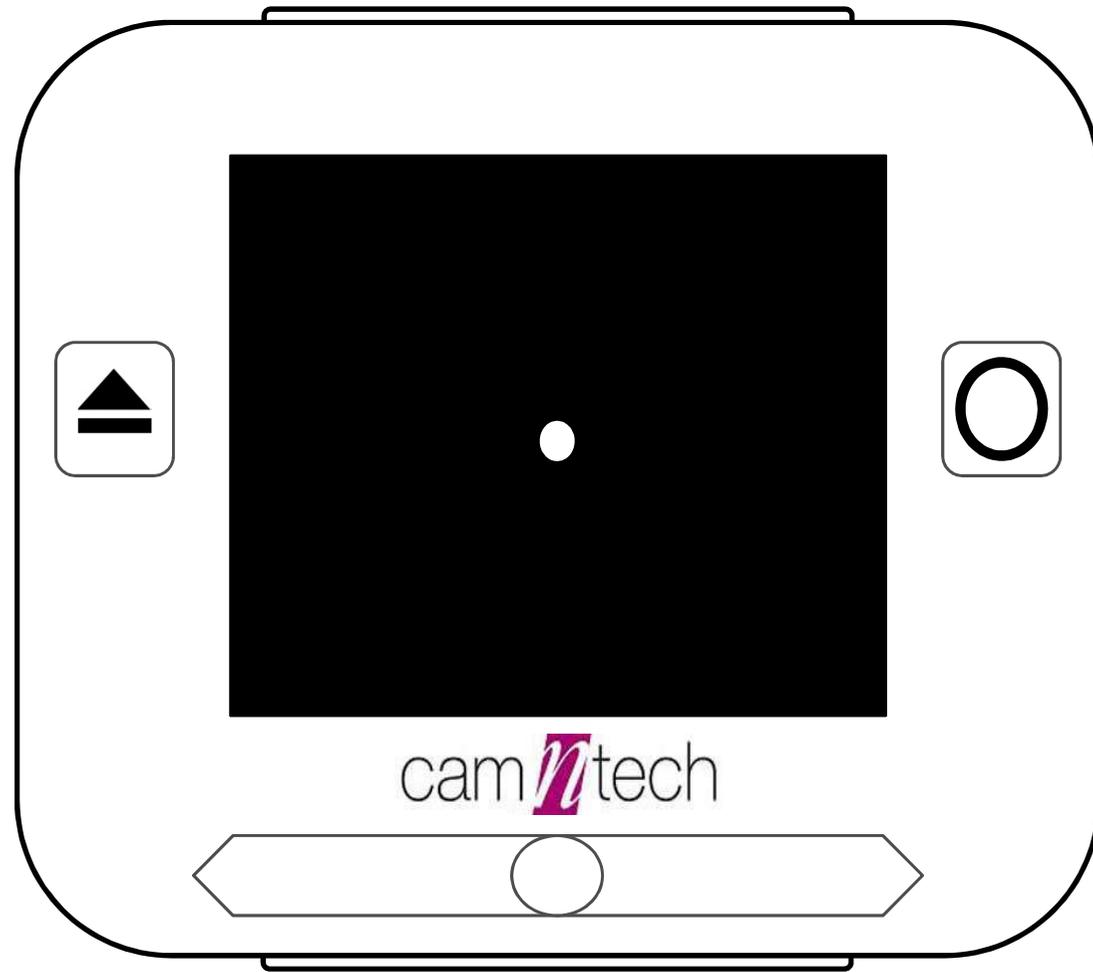
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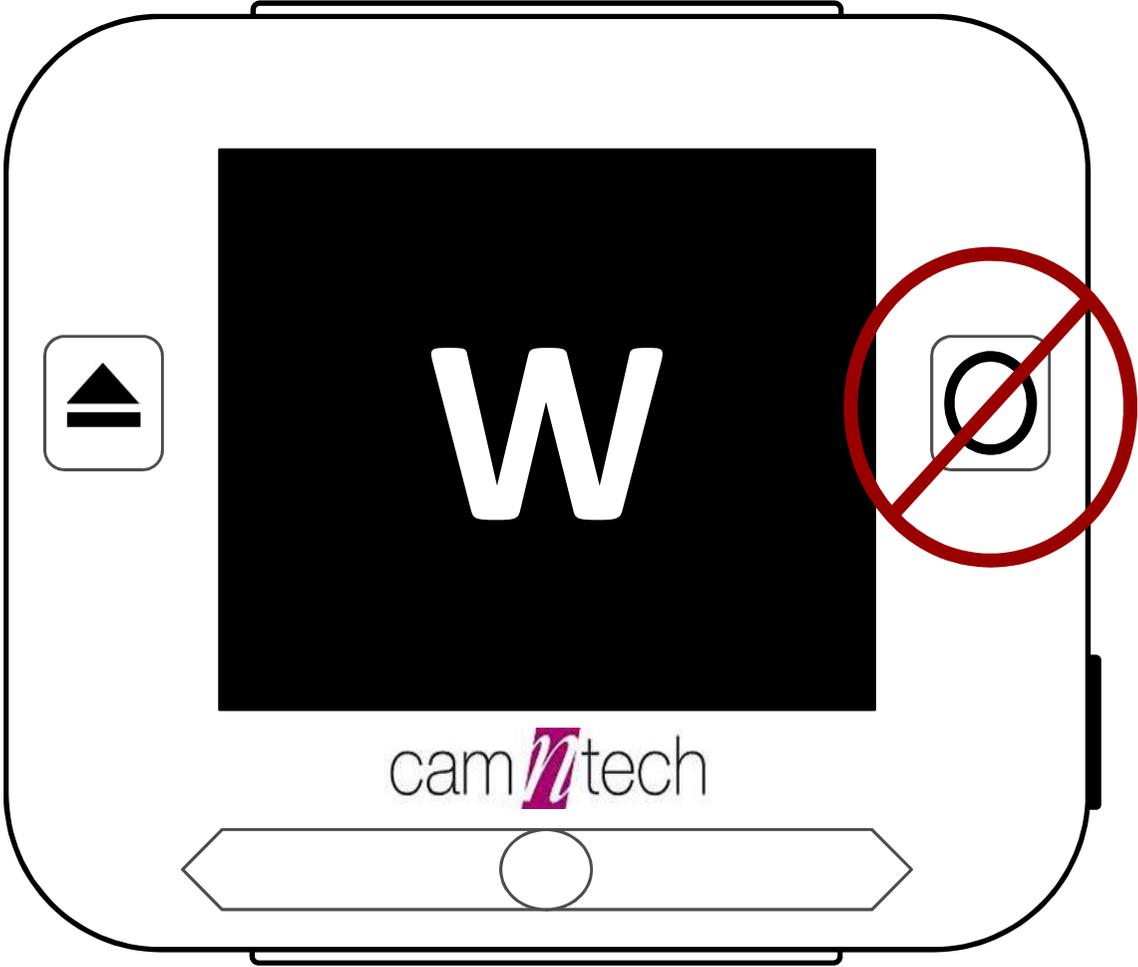
# Go/No-Go task



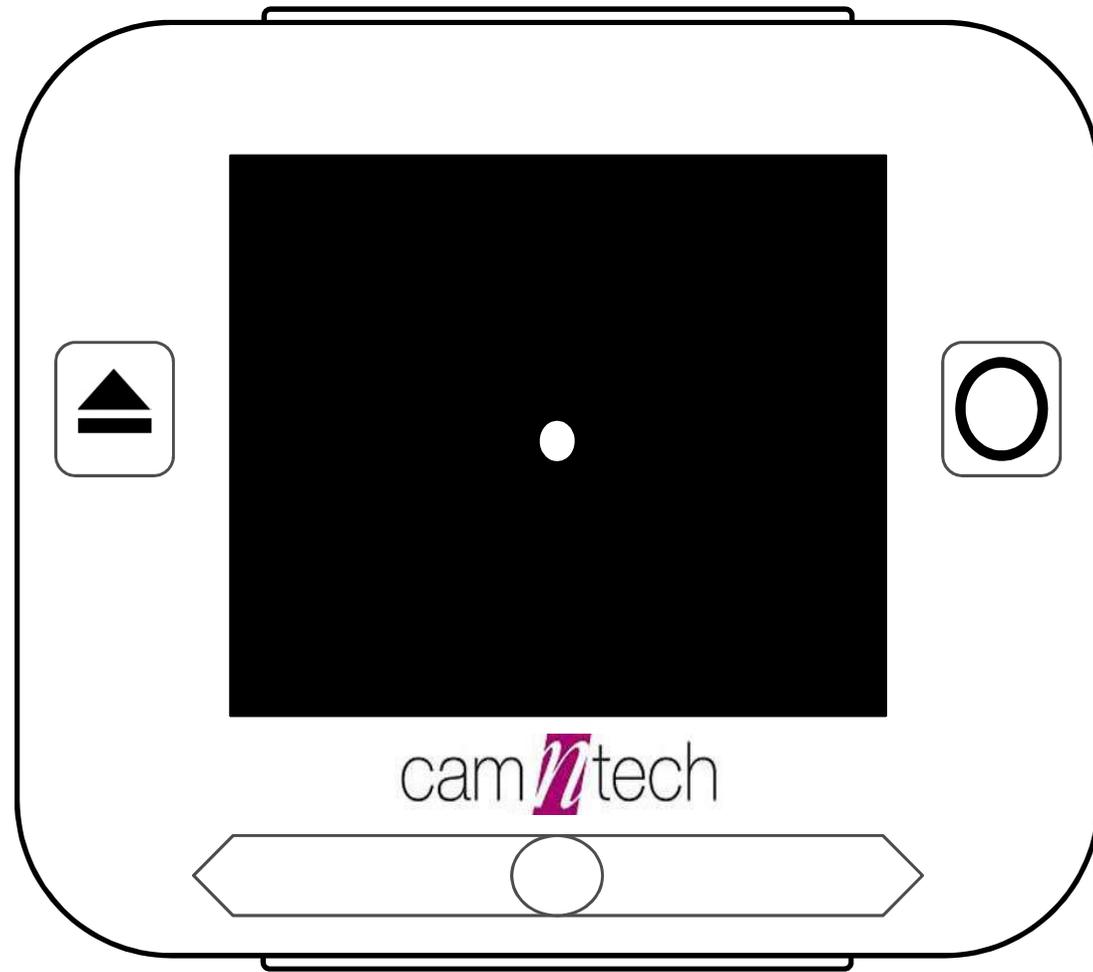
# Go/No-Go task



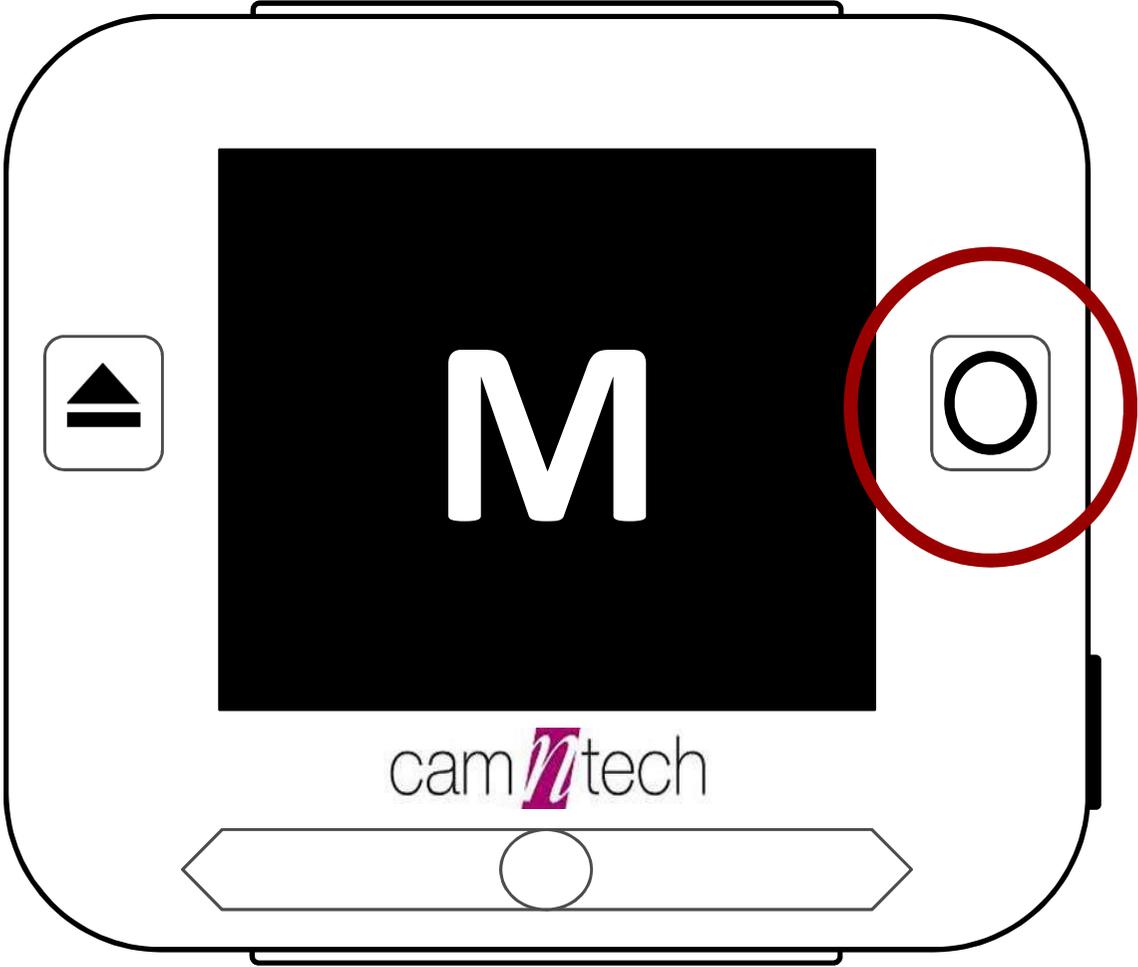
# Go/No-Go task



# Go/No-Go task



# Go/No-Go task



# Go/No-Go task – assessing inhibitory control



Performance Indicator:

Reaction time for correct responses (ms)

Slower reaction times = poorer real-time inhibitory control

Correct responses (%)?

# SNAPSHOT Design

- 7 consecutive days of EMA
- Fixed time-based design - hourly from 7am – 10pm
- Real-time Go/No-Go test assessments
- Short recall of snacking episodes



# Go/No-Go Test Validation

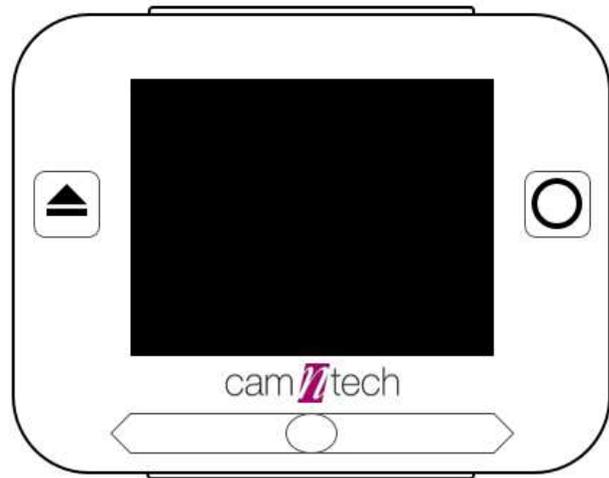


Person-mean Go/No-Go reaction times correlated (in expected direction) with several measures taken at baseline (in the lab) that purport to measure inhibitory control

- Attention-Switching Task: Congruency Cost ( $r = .380, p = .002$ )
- Stop-Signal Task: Stop-Signal Delay ( $r = -.538, p < .001$ ) and reaction time to Go Trials ( $r = .560, p < .001$ ), but not Stop-Signal RT ( $r = .165, p = .20$ )
- Behaviour Rating Inventory of Executive Function - Behavioural Regulation Index ( $r = .262, p = .037$ )

Intra-class correlation coefficient (ICC) of EMA Go/No-Go test = .55

# Snacking self-reports



Over the last hour.....

Have you eaten (non-core foods)...

- Chocolates/Sweets?
- Biscuits/Cakes/Pastries?
- Crisps/Savoury Snacks?
- Savoury pies/pastries?
- Takeaway/fast food?
- Soft drinks?

If Yes: 'Small', 'Typical' or 'Large/Multiple'

Operationalised as a count variable = snack consumption

# Participants (n = 68)

	Count or mean (SD)
Gender, Female	49
Current employment <sup>a</sup>	
Paid	39
Student	18
Retired	4
Housewife/husband	2
Household income <sup>b</sup>	
£0 – £20,000	20
£21,000 - £40,000	20
£41,000 – £60,000	7
£61,000 – £80,000	5
£81,000 – £100,000	4
> £101,000	5
Age (years) <sup>c</sup>	38.58 (15.54)
BMI <sup>d</sup>	25.67 (4.83)
Subjective Social Status <sup>e</sup>	6.42 (1.40)
Years in formal education <sup>f</sup>	16.94 (3.10)

<sup>a</sup> One response missing; <sup>b</sup> Three responses missing;

<sup>c</sup> Range = 18 – 70 years; <sup>d</sup> Range = 17.54 – 39.56;

<sup>e</sup> Range = 3.00 – 9.00; <sup>f</sup> Range = 10.00 – 23.00.

# Results: Missing data



All participants completed 7 days of EMA

## Go/No-Go test

74.2% (4664/6284) of requested Go/No-Go tests initiated;

Of these, 670 incomplete or noncompliant (replaced with a missing value)\*

## Real-time snacking

78.2% (4912/6284) of diary reports completed

*Negative Binomial Generalized Linear Mixed Model of Snacking Consumption*

Effect	$\gamma$	SE	p	Exp ( $\gamma$ ) <sup>a</sup>	95% confidence interval	
					Lower	Upper
Fixed effects						
Intercept	-2.00	.14	<.001	.14	.10	.18
Time of day (centered at 11:50 a.m.)	.08	.02	<.001	1.08	1.05	1.12
Go/No-Go RT (between-person) <sup>b</sup>	.02	.18	.965	1.02	.71	1.46
Go/No-Go RT (within-person) <sup>b</sup>	.23	.09	.002	1.26	1.06	1.49
Random effects <sup>c</sup>						
Intercept	.77	.22	<.001		.45	1.33
Time of day	.004	.003	.095		.001	.02
Covariance: Intercept and time of day	-.05	.02	.022		-.09	-.01
Residual <sup>d</sup>						
AR1 diagonal	1.24	.03	<.001		1.18	1.31
AR1 $\rho$	-.03	.02	.105		-.07	.007

*Note.* Probability distribution: Negative Binomial; link function: log.

<sup>a</sup> Exp ( $\gamma$ ) is interpreted as a percentage increase (values > 1) or decrease (values < 1) in the consumption rate for a 1-unit increase in the predictor. <sup>b</sup> higher RTs indicate less inhibitory control. RT was transformed such that 1-unit equated to 100 ms (i.e. one-tenth of a second). <sup>c</sup> random effect covariance structure: unstructured. <sup>d</sup> residual covariance structure: first-order autoregressive (AR1).

# Results

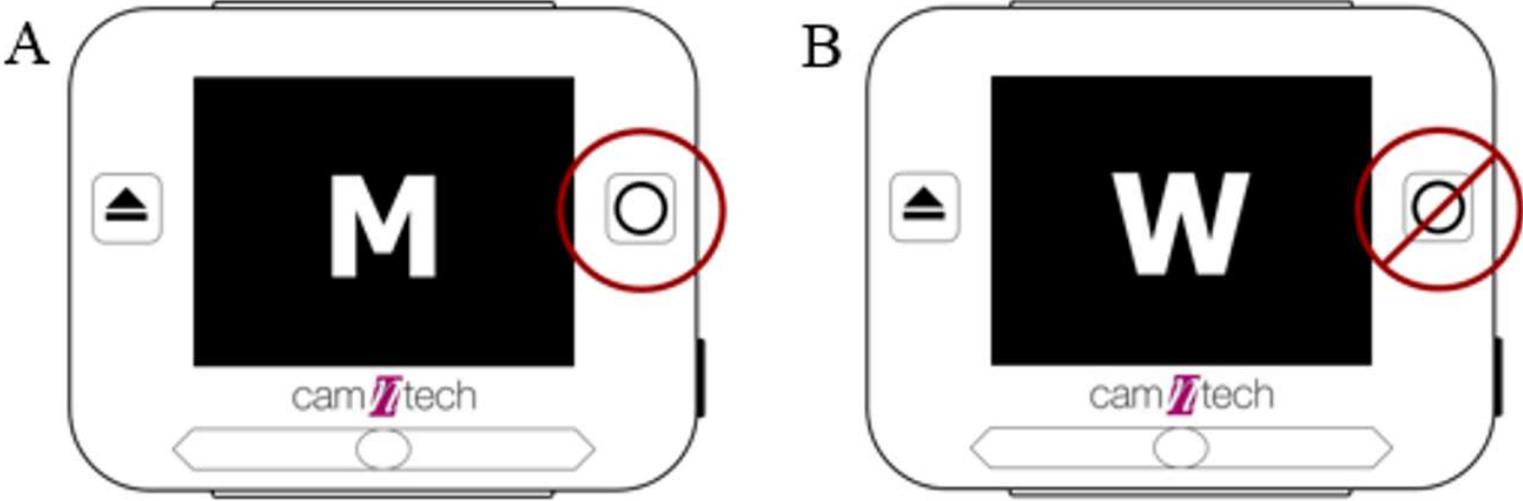
**Within-person:**  $\text{Exp}(\gamma) = 1.26$ ,  $p = .002$ ,  $\text{CI}_{95}: 1.06, 1.49$

100 ms slower RT (than usual) = 25.67% higher consumption in subsequent hour

**Between-person:**  $\text{Exp}(\gamma) = 1.02$ ,  $p = .965$ ,  $\text{CI}_{95}: 0.71, 1.46$

Results robust to analyses adjusting for BMI, alcohol intake, outliers.

# Go/No-Go tests: spotting non-compliance



*Figure S1.1.* Representation of the PRO-Diary watch-faces for ‘Go’ trials (A) and ‘No-Go’ trials (B). Participants were asked to respond as quickly as possible to ‘Go’ trials using the response button on the right.

# Go/No-Go tests: “Multi-tappers”

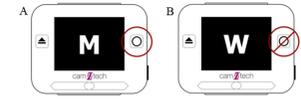


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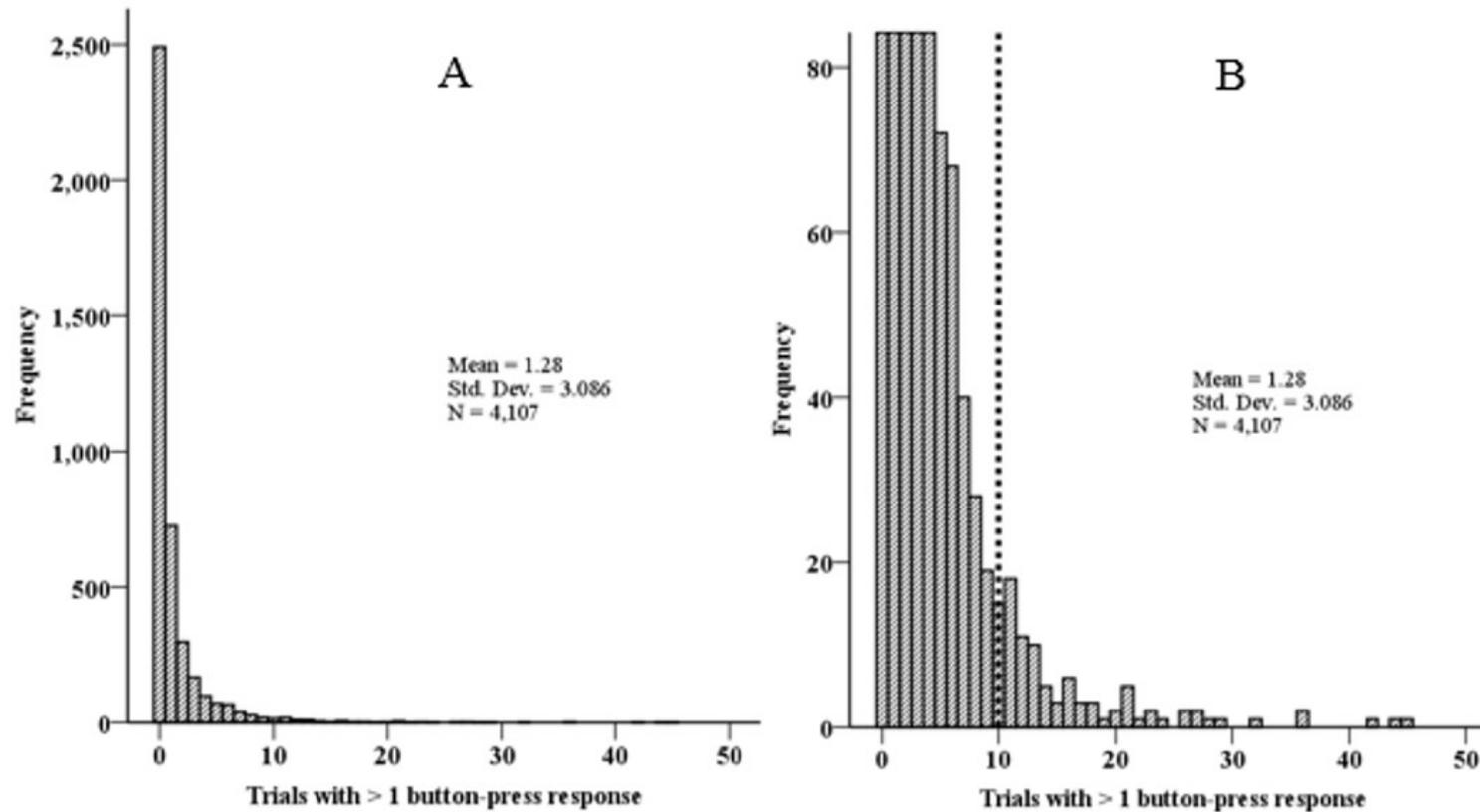


Figure S1.3. Frequency of tests by the number of constituent ‘Go’ trials with a button-press response (A) and focussing on the lower frequencies of the same histogram (B). The cut-off decision is denoted by the dashed line.

# Go/No-Go: “Tap-and-goers”

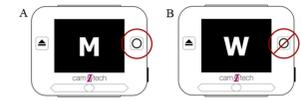


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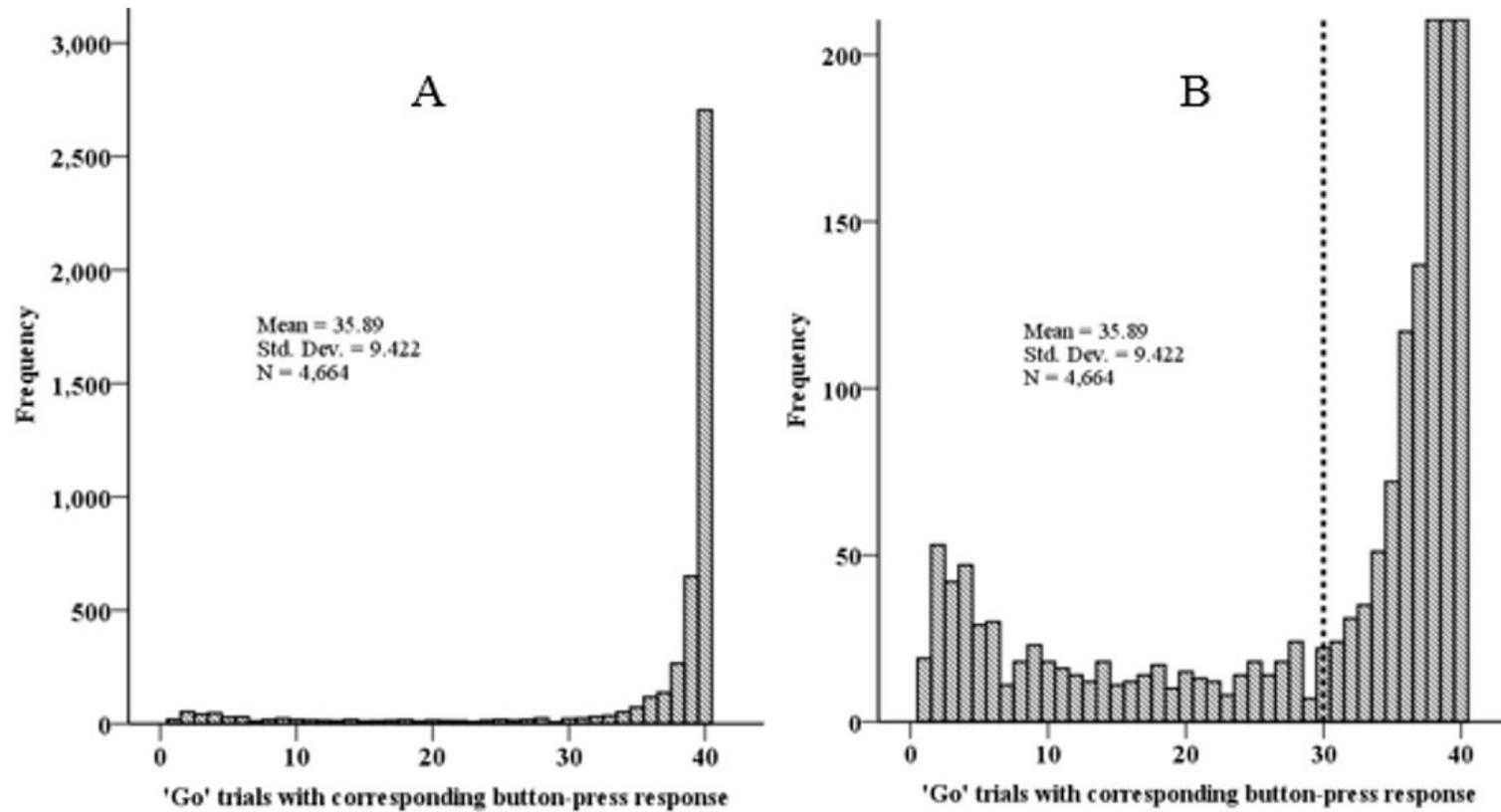


Figure S1.2. Frequency of tests by the number of constituent 'Go' trials with a button-press response (A) and focussing on the lower frequencies of the same histogram (B). The cut-off decision is denoted by the dashed line.

# Solution to non-compliance in data

- Realise that full compliance at all times is an unrealistic expectation, especially when burden is quite high
- Know what to look for – “cheat” in a pilot run yourself. How does this look in the data? Create syntax/code to identify
- Set pre-specified rules for where a test result is insufficiently or inappropriately completed
- Check for any predictors of invalid responses in your data. Include these predictors in the model (as not missing completely at random)

# Summary

- Inhibitory control fluctuates significantly within-persons
- Consistent with theory, the relationship between inhibitory control and snacking appears best understood as a within-person process.
- Mediating effect on intention-behaviour relation untested.
  
- Go/No-Go test can detect changes in inhibitory control efficiency in an EMA study
- Too simple? A more selective inhibitory mechanism based on behavioural goals may be more realistic – deciding which cues to suppress and which to enact
- Careful data cleaning is needed. Know what to look for....!



# MS Cortisol & Fatigue Study

# Fatigue in MS

- 60 – 85% prevalence (Lerdal et al., 2003; Minden et al., 2006)
  - Often described as the most disabling and distressing symptom
  - Cortisol is the “end product” of the HPA axis
  - Cortisol is an important regulator of the immune system (downregulation)
  - HPA axis hyperactive in MS? (Gold et al., 2010; Kern et al., 2011)
  - Cortisol important in energy metabolism
  - Little examination in daily life using EMA
-

# Main Hypothesis

Fatigue severity associated with an attenuated cortisol awakening response (CAR)

- Between individuals
- Within individuals (i.e. day-to-day)

# MS Fatigue and Cortisol - Recruitment



## People with relapsing-remitting MS (RRMS)

### Exclusion criteria:

- Within 3 months of a clinical relapse
- (Self-reported) inability to walk 300m with/without use of a walking aid
- Diagnosed physical or psychiatric comorbidity
- HADS score  $\geq 8$  (indicative of moderate depression)
- Current antidepressant use
- Pregnancy
- Caregiving
- Shift-working

# MS Fatigue and Cortisol Design

- Case-Control
- 4 consecutive days of EMA
- 4 x CAR; 4 x diurnal slopes
- CAR: fixed event-based design (awakening, +30min, +45min)
- Diurnal slope: variable time-based design (10am – 8pm)
- Baseline fatigue severity scale (Chalder Fatigue Scale)
- Momentary fatigue severity ratings
  - How fatigued (tiredness, weariness, problems thinking clearly) do you feel right now?*

# Salivary cortisol in daily life



- Enables repeated assessments
- Non-invasive, relatively easy to administer

Markers of HPA axis activity:

- (1) Cortisol awakening response
  - (2) Diurnal slope
  - (3) Stress reactivity
-

# Salivary cortisol in daily life

- Enables repeated assessments
- Non-invasive, relatively easy to administer

Markers of HPA axis activity:

(1) Cortisol awakening response

(2) Diurnal slope

(3) Stress reactivity

Sampling:

T0      Awakening

T30      Awakening + 30mins

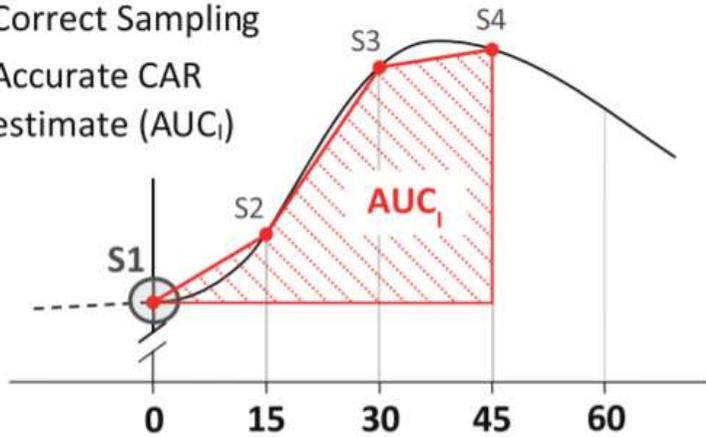
T45      Awakening + 45 mins

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# Methodological Challenge

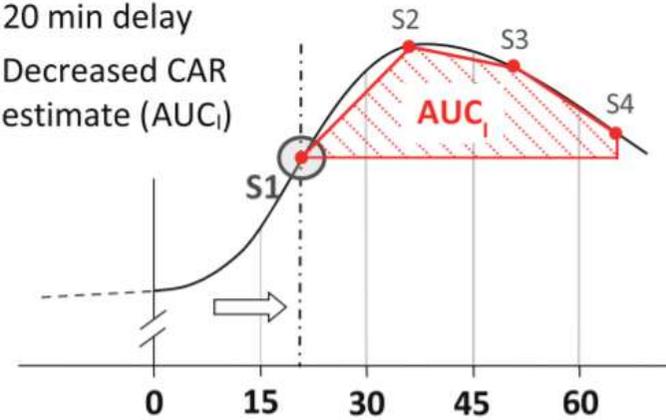
a) Correct Sampling

→ Accurate CAR estimate ( $AUC_i$ )



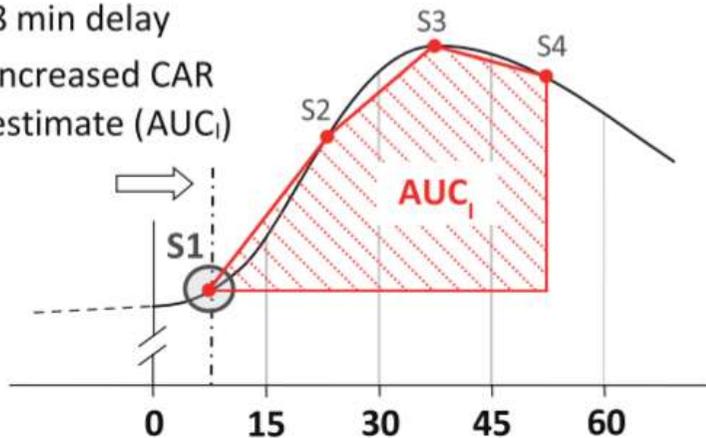
c) 20 min delay

→ Decreased CAR estimate ( $AUC_i$ )



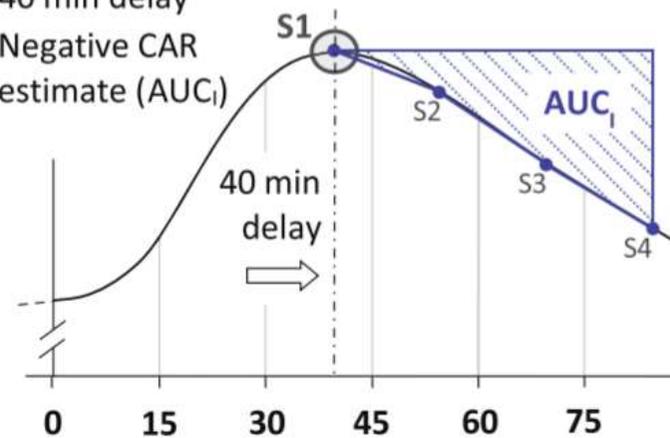
b) 8 min delay

→ Increased CAR estimate ( $AUC_i$ )



d) 40 min delay

→ Negative CAR estimate ( $AUC_i$ )



# Methodological Challenge



How to ensure participants collect samples promptly when asked?

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# Methodological Solution



Stetler, Dickerson, & Miller (2004)

- Use the device as an alarm clock
- Present time-limited code
- Require code be transferred to label
- Exclude any delayed samples, or samples with incorrect code

## Remaining weaknesses

- Spontaneous awakenings

# Participants (n = 76)

	RRMS	Control
N	38	38
Age	41.89 (7.53)	40.34 (8.16)
Gender	31f/7m	31f/7m
EDSS	4.35 (1.40)	
Years since diagnosis	6.03 (5.18)	
HADS-D	4.00 (2.29)	2.08 (2.27)
HADS-A	7.50 (3.90)	4.82 (3.12)
Fatigue Scale	17.58 (7.09)	11.55 (2.87)
FS-Phys	11.18 (4.89)	7.26 (2.34)
FS-Ment	6.39 (2.66)	4.29 (0.96)
Sleep Quality (Mean)	6.07 (1.57)	6.22 (1.97)
Sleep Hrs (Mean)	7.83 (1.00)	7.63 (0.89)

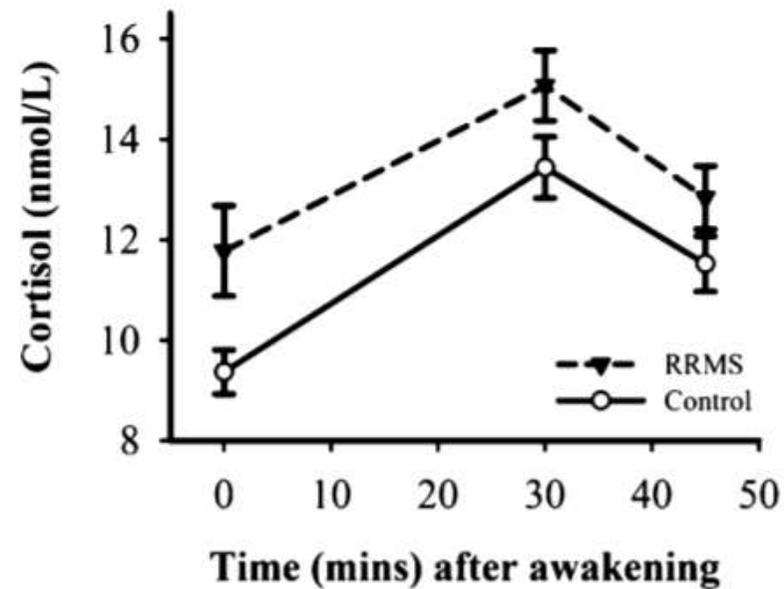
# Results: Missing data



All participants completed 4 days of EMA

## Salivary cortisol

One participant did not provide any CAR samples, leaving 75 participants with data  
42 from 300 (14%) remaining CARs had missing or **delayed** samples



**Figure 1** Cortisol awakening response represented by the mean of the within-subject means for samples at 0 (S1), 30 (S2), and 45 min (S3) post-awakening. Error bars represent the standard error of the mean.

# Results

Fixed effect	RRMS					Control				
	Coef.	SE	t	p	95% CI	Coef.	SE	t	p	95% CI
Models for CAR AUCi										
FS Score	0.157	0.064	2.442	.017	[0.029, 0.285]	0.087	0.160	0.543	.588	[-0.231, 0.405]
FS Physical	0.216	0.093	2.316	.023	[0.031, 0.400]	0.094	0.194	0.482	.631	[-0.292, 0.480]
FS Mental	0.391	0.175	2.235	.028	[0.044, 0.739]	0.219	0.502	0.324	.663	[-0.776, 1.214]
Models for (ln) S1 cortisol										
FS Score	-0.022	0.008	-2.781	.007	[-0.038, -0.006]	-0.008	0.019	-0.387	.700	[-0.046, 0.031]
FS Physical	-0.028	0.012	-2.454	.016	[-0.051, -0.005]	-0.009	0.024	-0.393	.695	[-0.058, 0.039]
FS Mental	-0.060	0.021	-2.865	.005	[-0.102, -0.018]	-0.011	0.059	-0.191	.849	[-0.129, 0.106]

Analyses robust to sensitivity analyses with “responders”-only

Powell, Moss-Morris, Liossi, & Schlotz, W. (2015)

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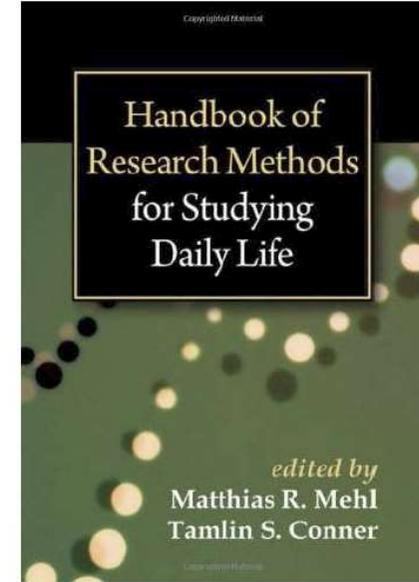
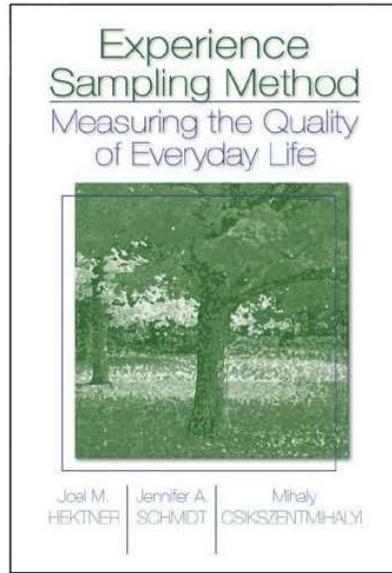
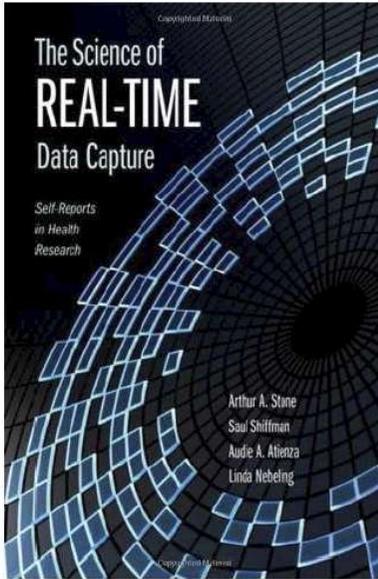
# Summary

- Fatigue in MS associated with lower CAR AUCi and higher S1 cortisol
- No relation between fatigue in controls with any CAR marker
- Timing is crucial in salivary cortisol studies
- In 2013-14, 7.9% of CAR studies had objective control of awakening time (Stalder et al., 2016)
- In 2013-14, 18.6% of CAR studies had objective control of sampling time (Stalder et al., 2016)
- In 2013-14, a diary log method in 65.5% of studies (Stalder et al., 2016)
- CAR on a single day is determined mostly by situational factors (Hellhammer et al., 2007) yet in 2013-14, 31.7% of studies observed CAR on 1 day alone and only 4.8% over 4-5 days (Stalder et al., 2016)

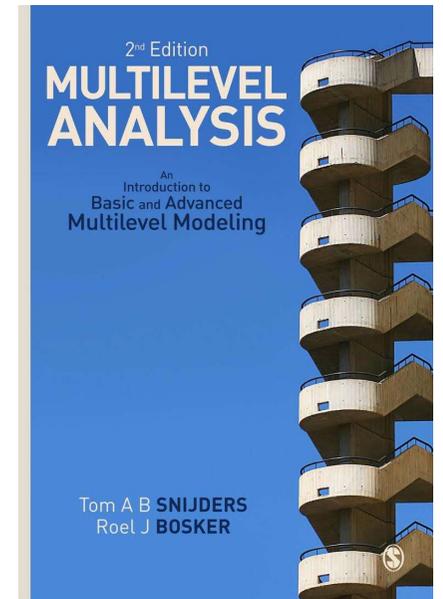
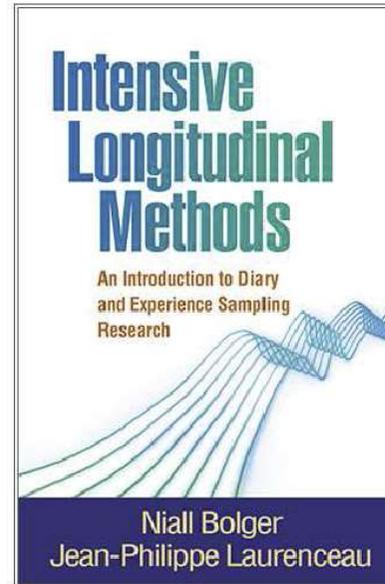
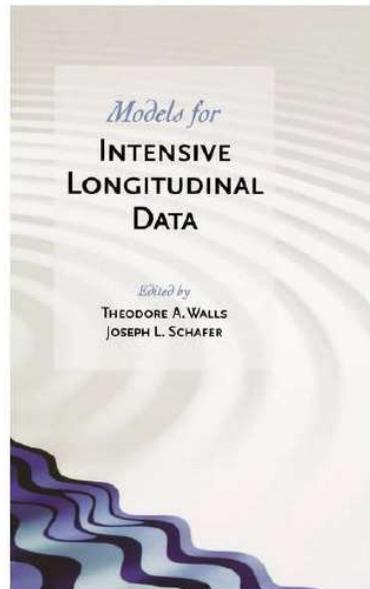
# Overall EMA Summary and Tips



- EMA methods present unique opportunities but also challenges
- Have a theory of change and design your study accordingly
- Pilot, pilot, pilot
- Account for time in your models – avoid spurious relationships
- Deal with missing data appropriately. MLMs do not automatically resolve this issue
- You will have missings, but also ‘partial missings’ and ‘invalid completes’. Decide how you will (i) discourage these; (ii) identify these and (iii) what you’ll do about them.
- Do not fall into the trap of thinking more-objective measures are immune from challenges. They aren’t.
  
- Lots of useful, accessible texts on both the method and its analysis



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Data Analysis  
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# Thanks. Any questions?

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