

PsPM course session 3: SCR

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Clinical Research Priority Program "Synapse & Trauma" & Department of Psychiatry, Psychotherapy, and Psychosomatics, University of Zurich*

16.04.2020

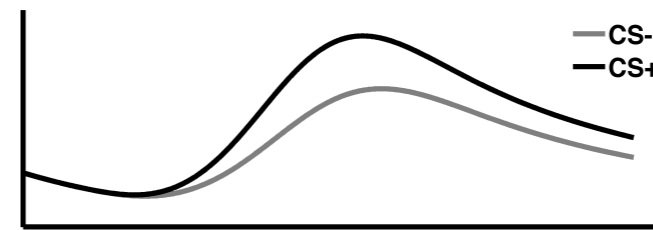
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@bachlab_cog*



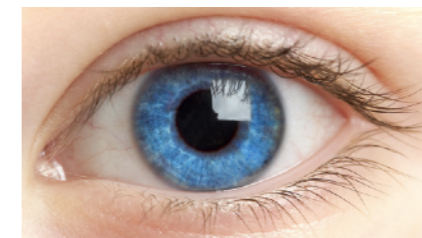
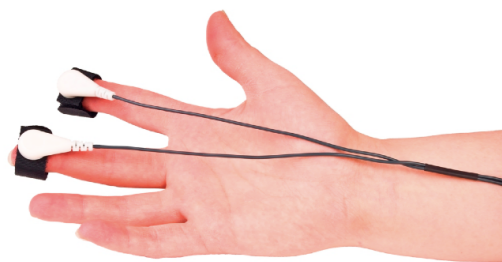
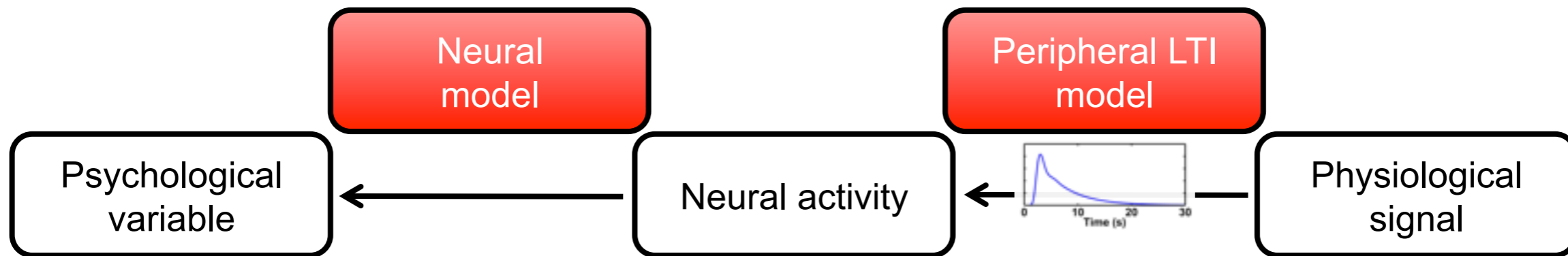
The "best possible" approximation to the true psychological variable.

Memory difference between CS+/CS-?

Lecture 7: 14.05.2020



Lecture 2: 09.04.2020



Lecture 3: 16.04.2020

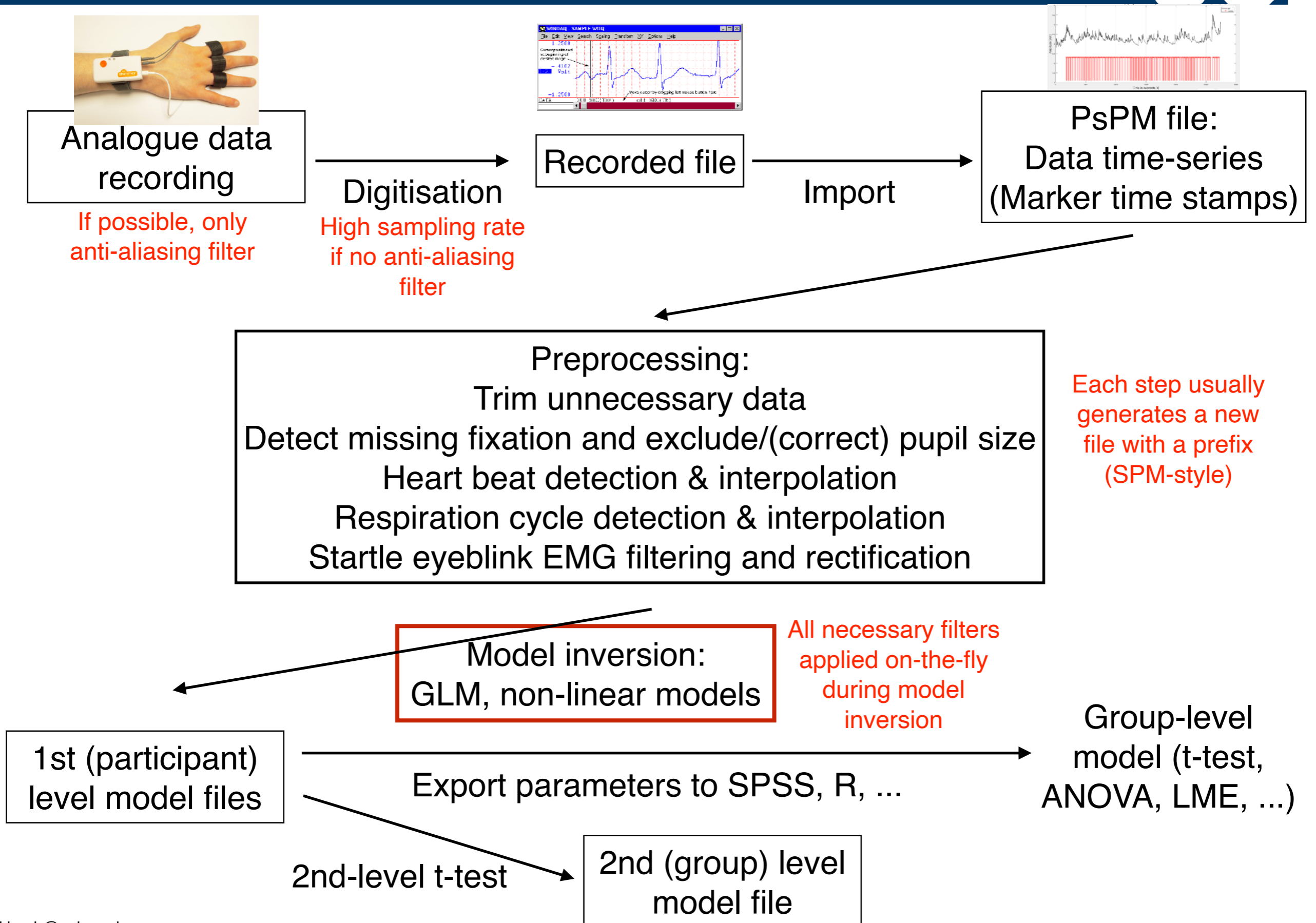
Lecture 5: 30.04.2020

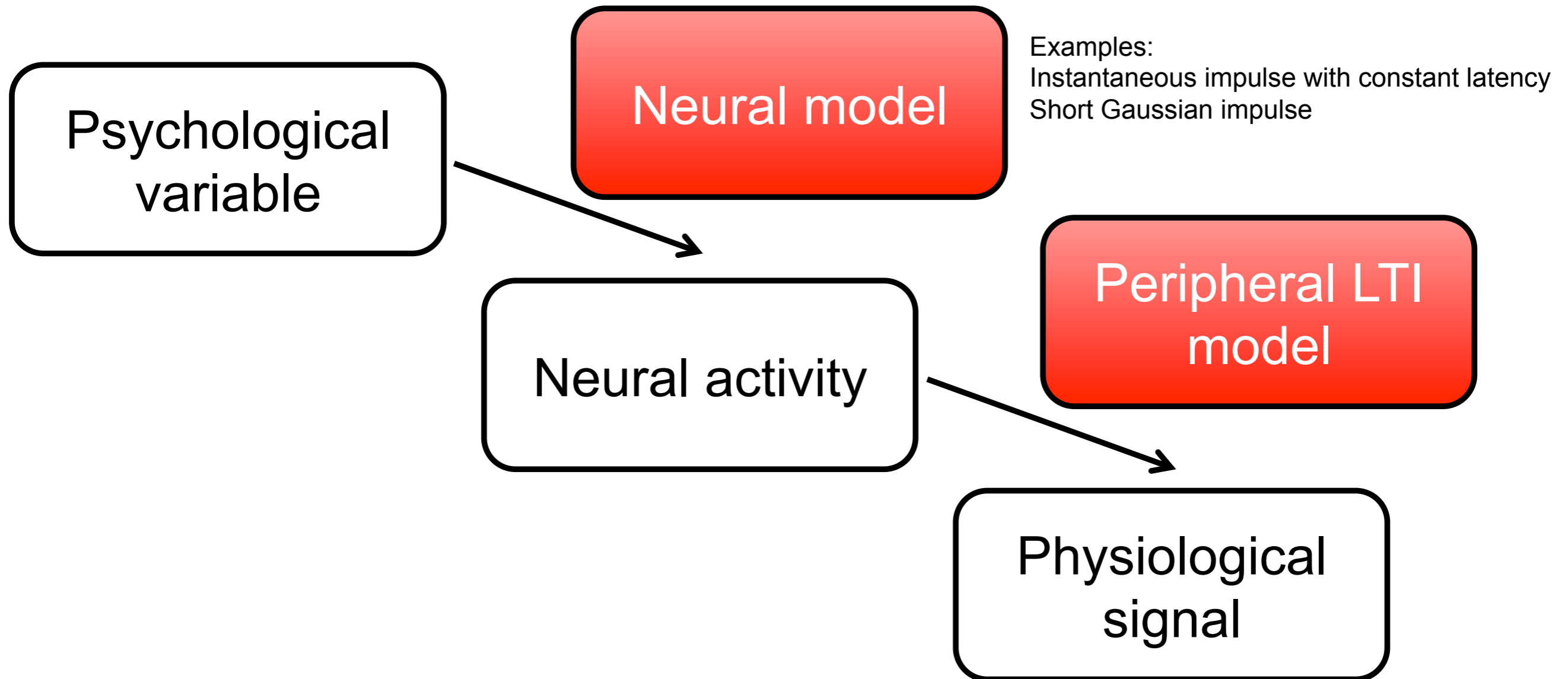
Lecture 6: 07.05.2020

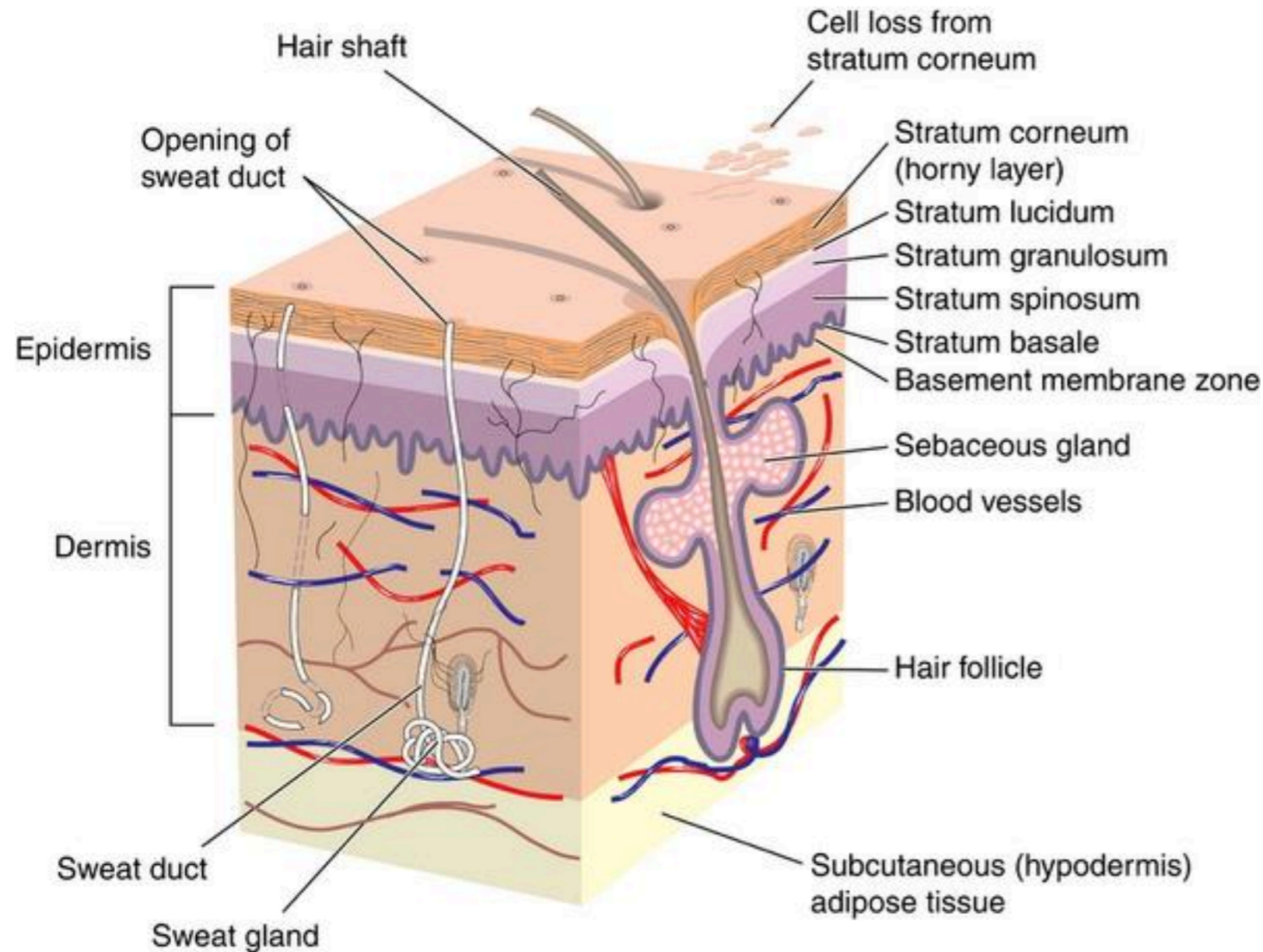
Lecture 4: 23.04.2020

Lecture 6: 07.05.2020

PsPM pipeline overview





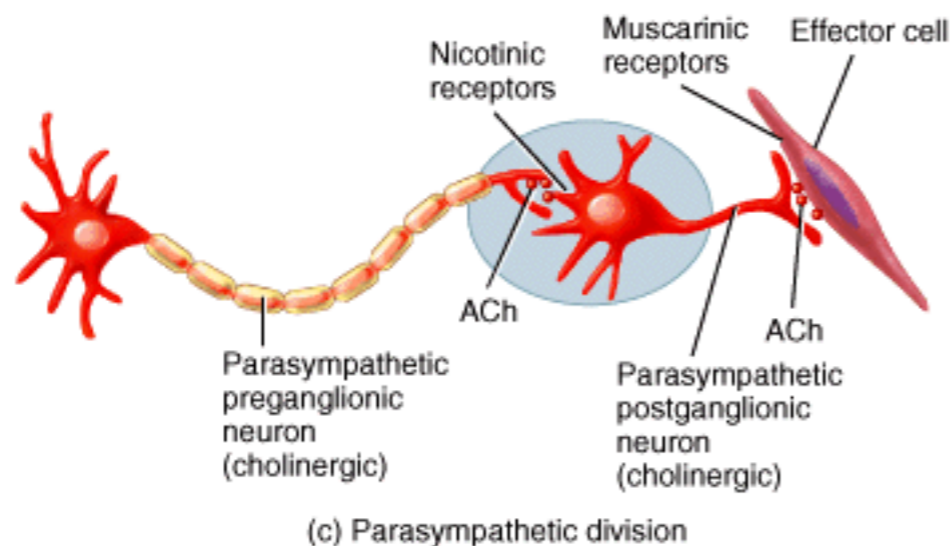
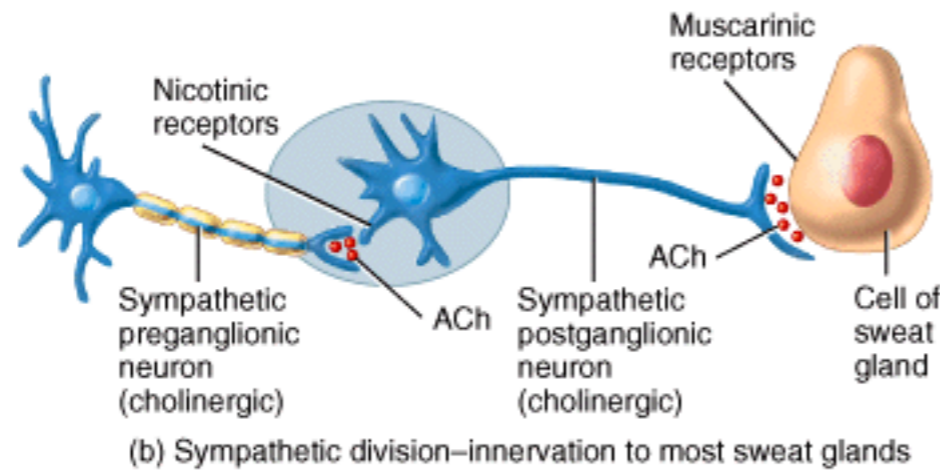
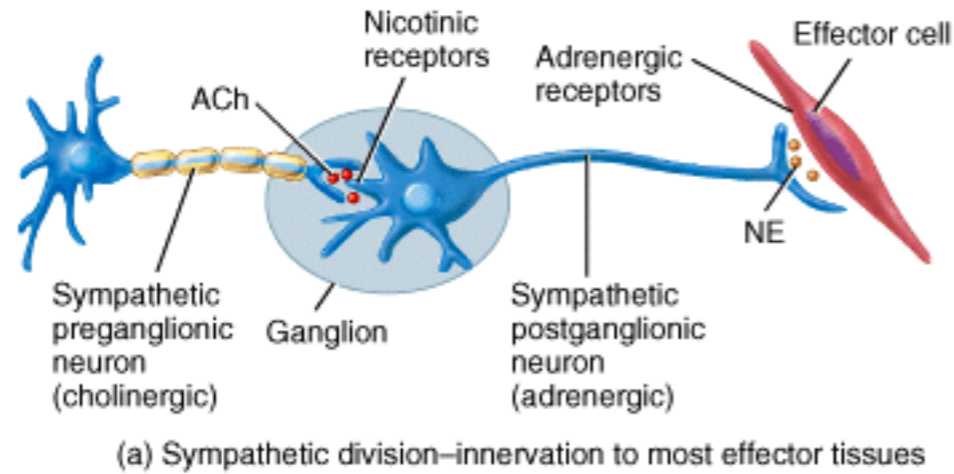


Constant voltage (DC) coupler:

$$G = \frac{1}{R} = \frac{I}{U}$$

Applies constant voltage (e.g. 0.5 V), quantifies current via a measured voltage proportional to the current

Measured voltage depends on sensitivity, needs to be transformed to mcS for comparability between subjects & studies



Thermoregulation

"Emotional sweating": mainly measured on palmar & plantar skin with high sweat gland density, but no positive evidence for regional specificity

"Gustatory sweating"

Orienting response

- Novelty
- Salience

Fear conditioning

- Debate as to how many components: 1-3 depending on ITI, evidence is weak
- Amygdala-dependent threat learning or (hippocampus-dependent?) contingency
- awareness? [1]

Motor action

- Often strongest determinant of SCR

Operant conditioning

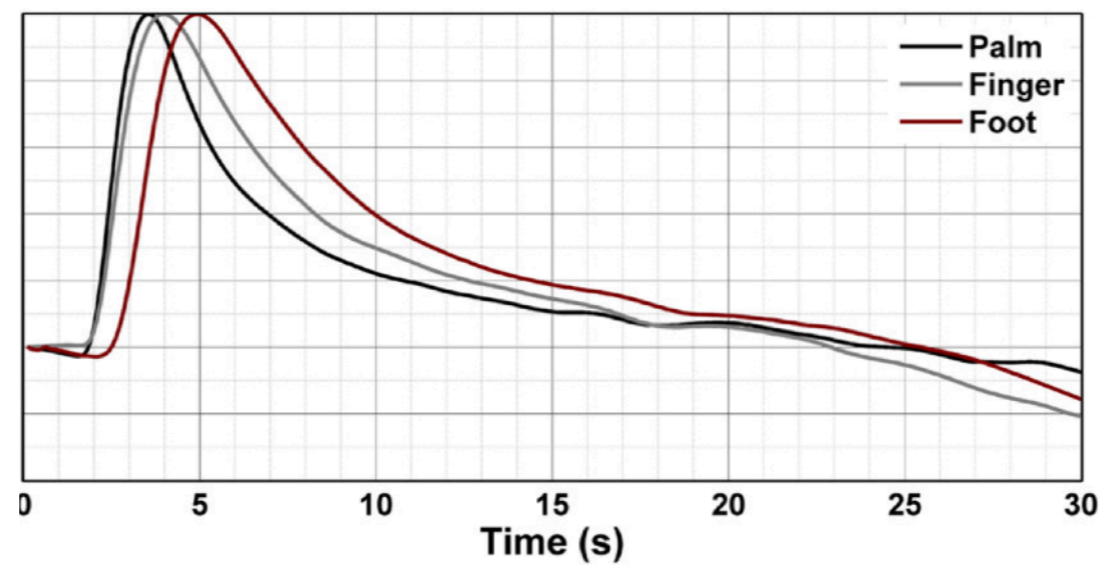
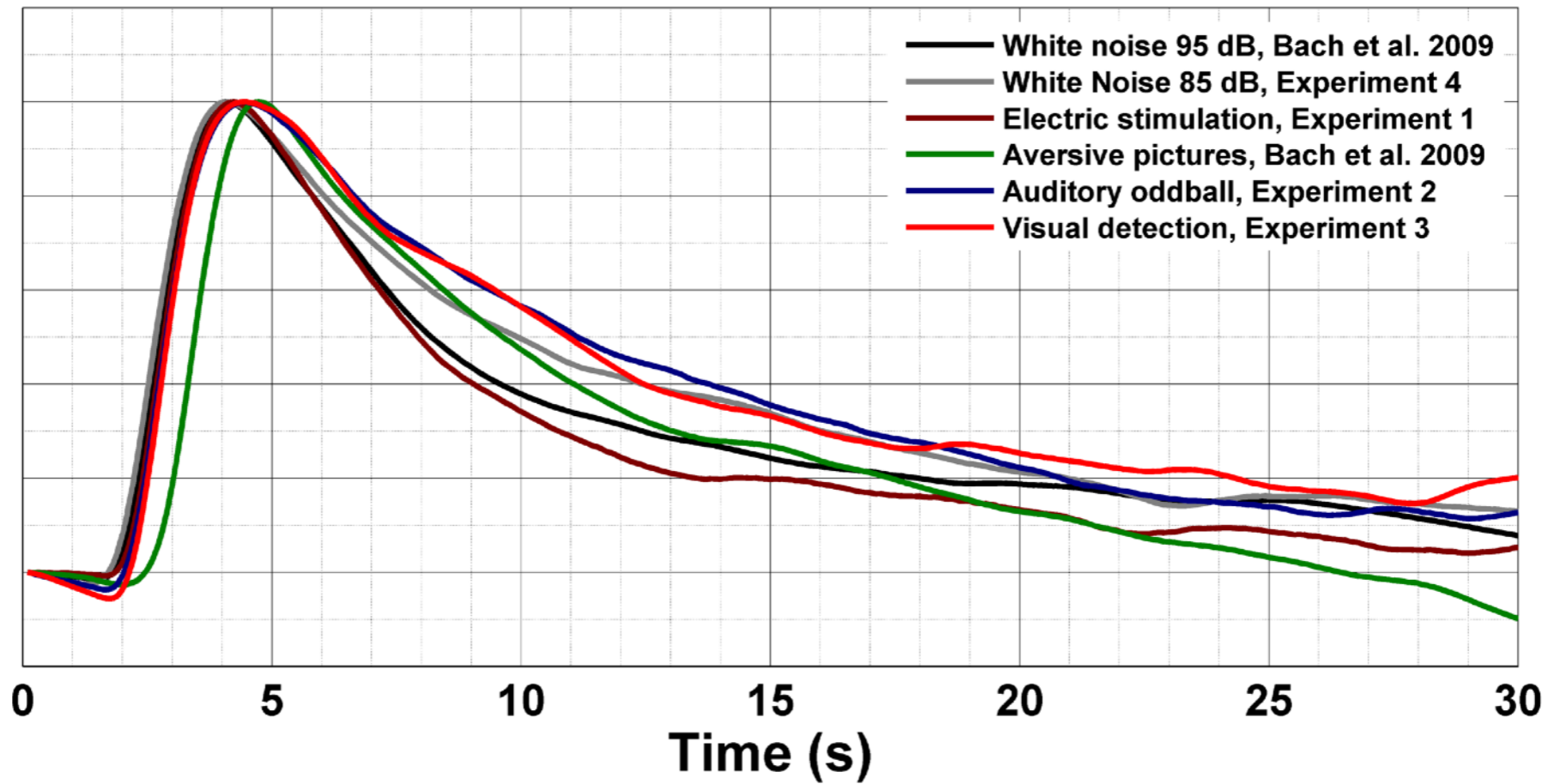
- SCR can be reinforced

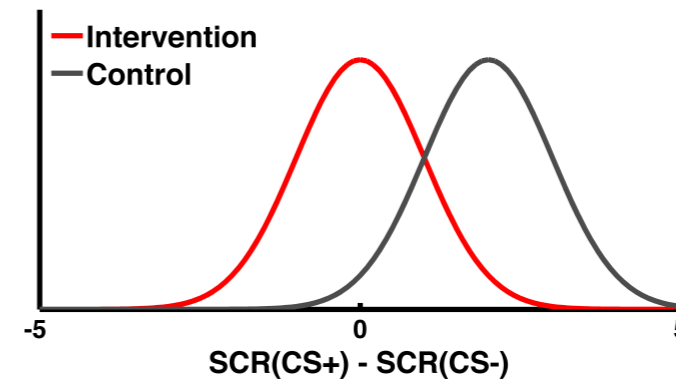
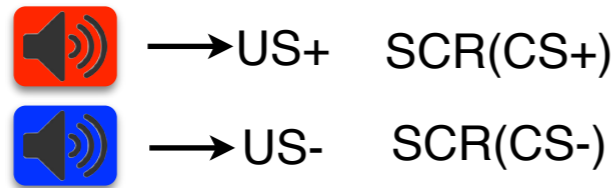
Information processing

- Cognitive load, decision-making, memory

Tonic arousal

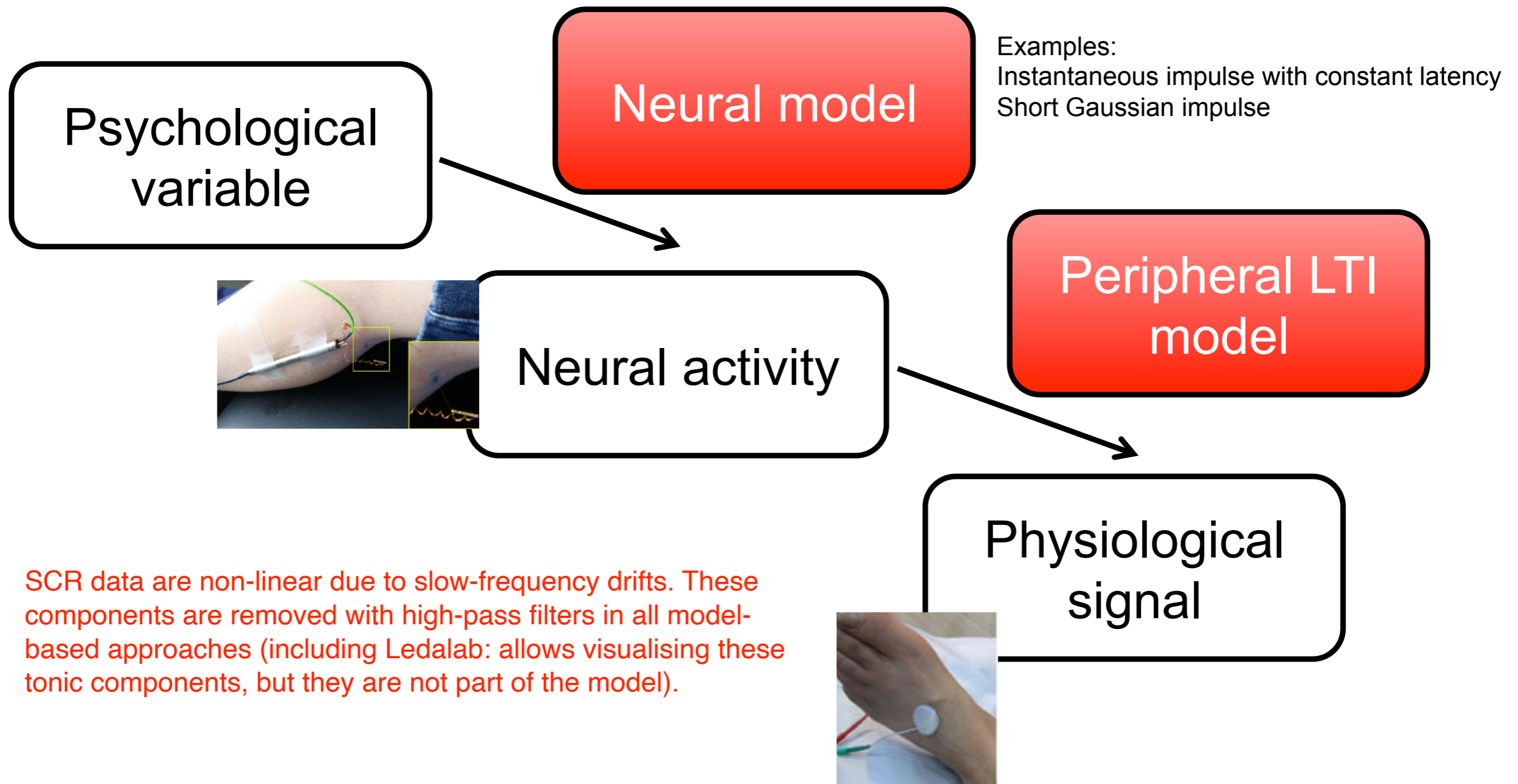
- Anxiety, stress





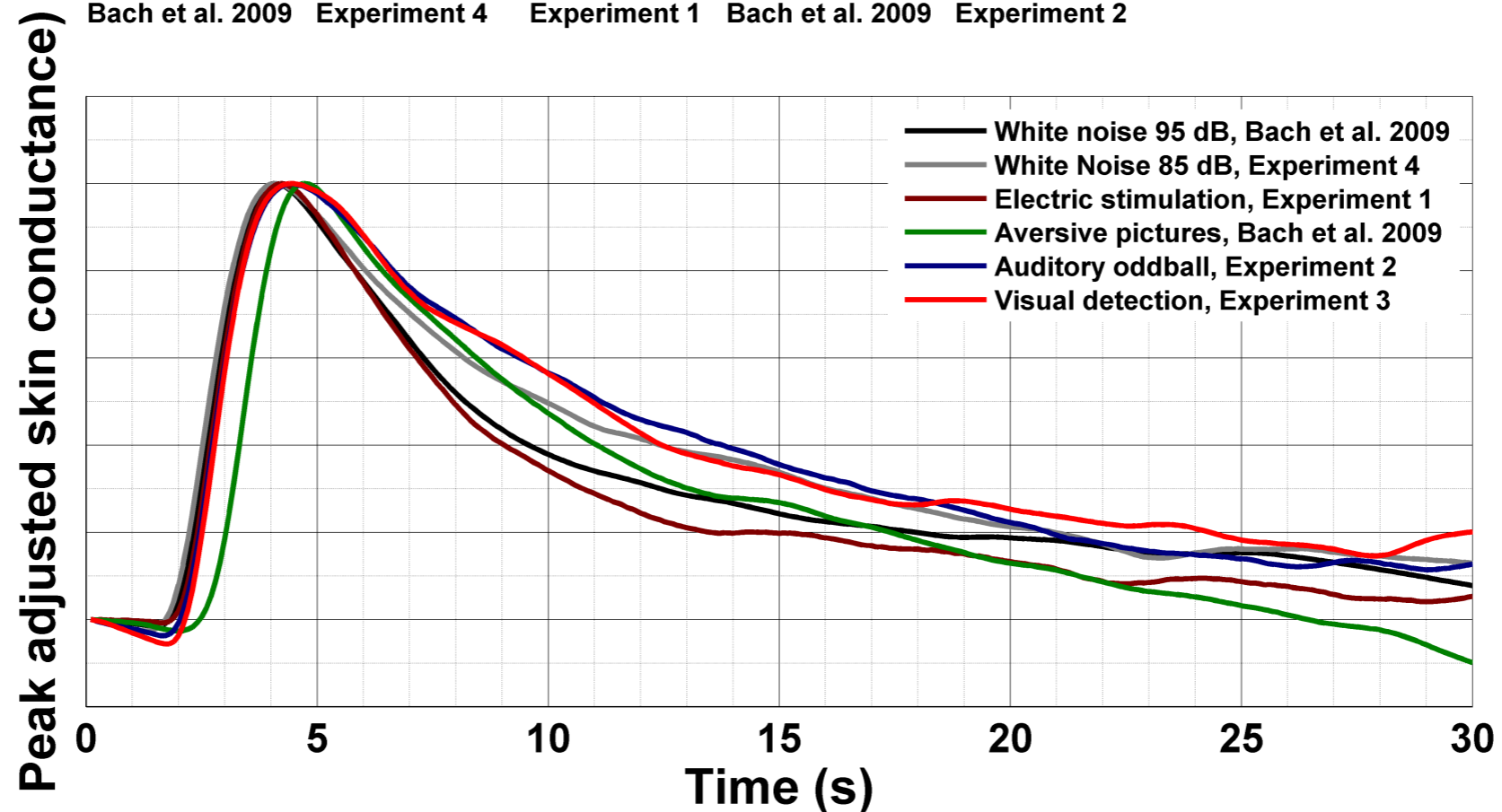
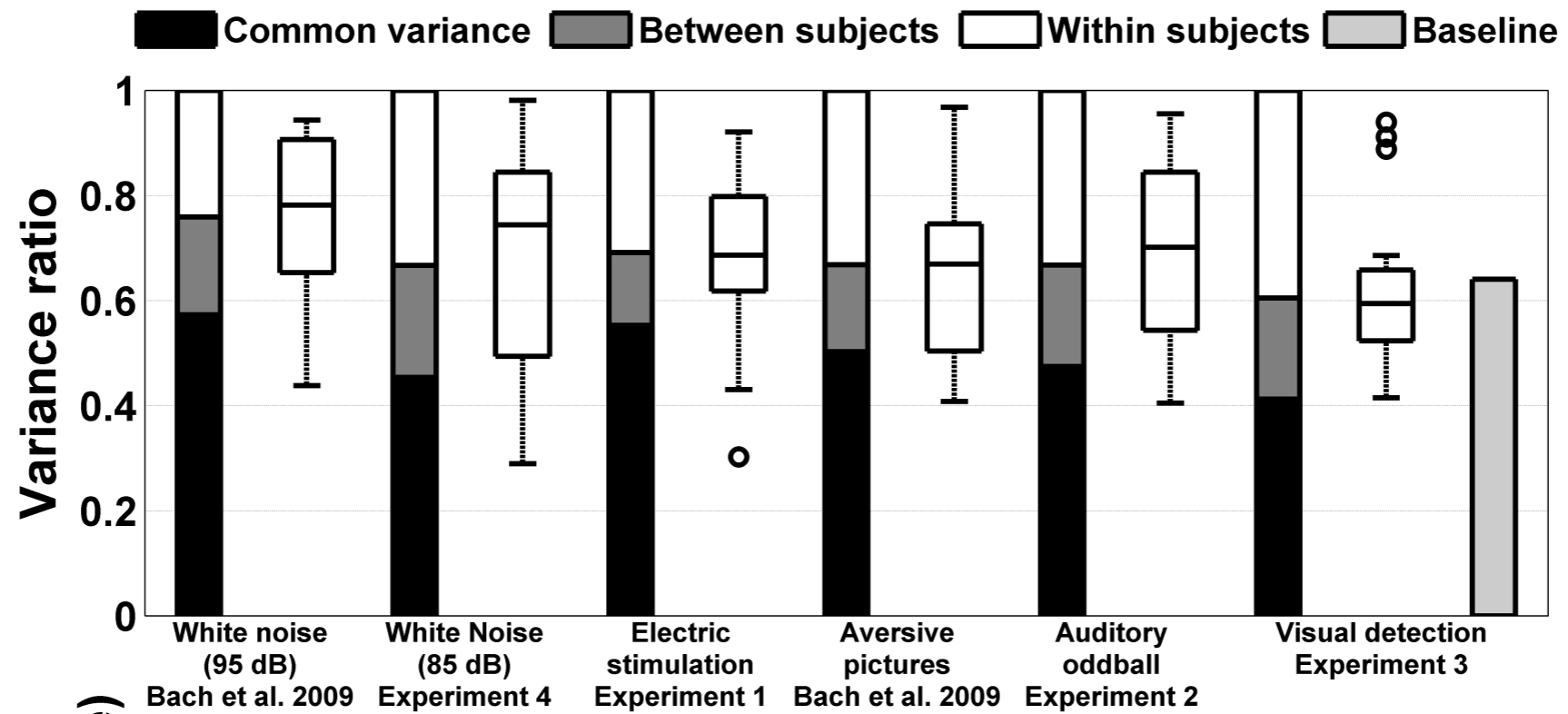
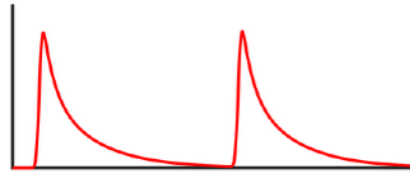
	Measure	d	N*
	SCR peak scoring	0.44	514
	SCR model-based	0.75	174
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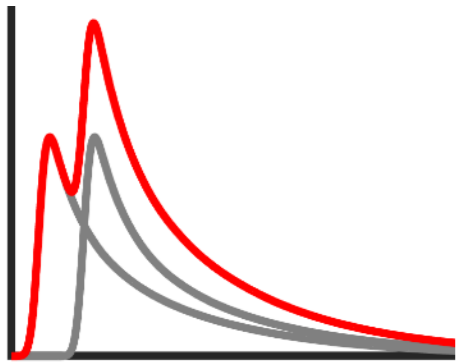
*: Sample size required to achieve 80% power at $\alpha = .05$ in a one-tailed test, if intervention reduces fear memory at least 50% and has no variability (best-case scenario)



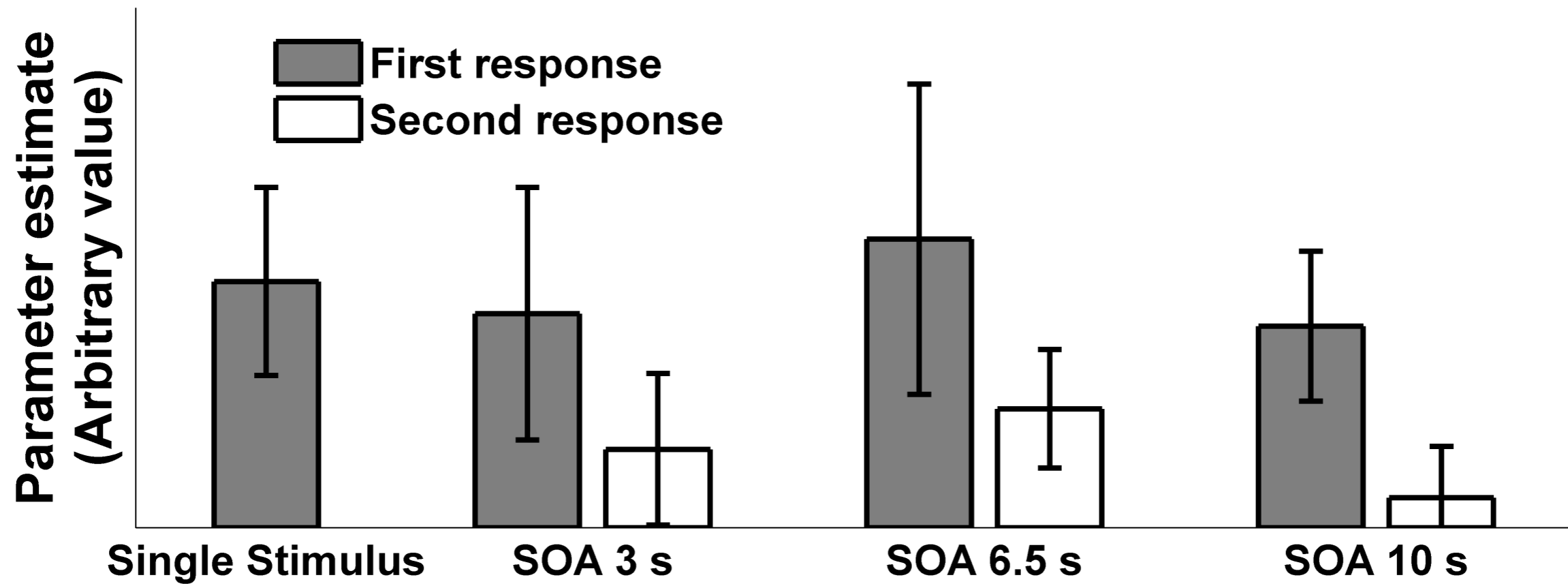
SCR data are non-linear due to slow-frequency drifts. These components are removed with high-pass filters in all model-based approaches (including Ledalab: allows visualising these tonic components, but they are not part of the model).

Indirect tests: time invariance

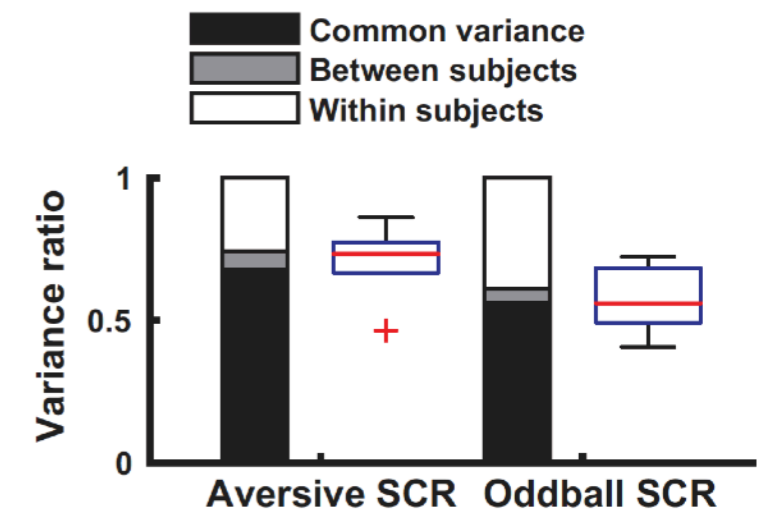
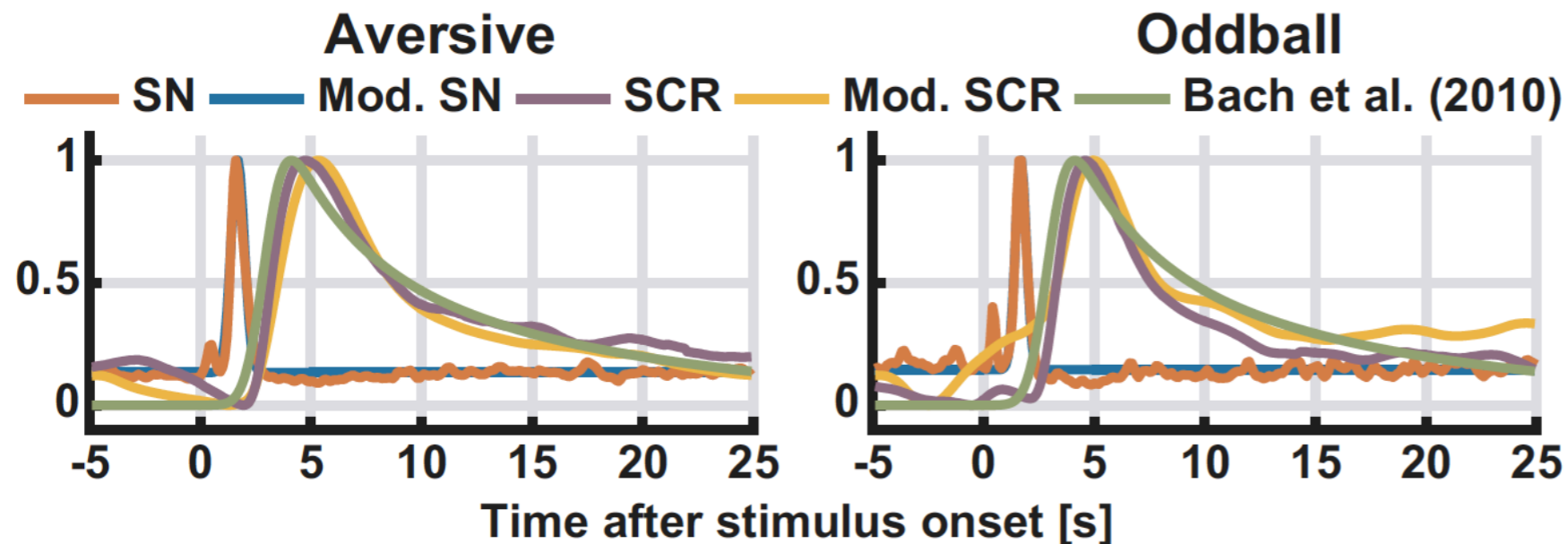




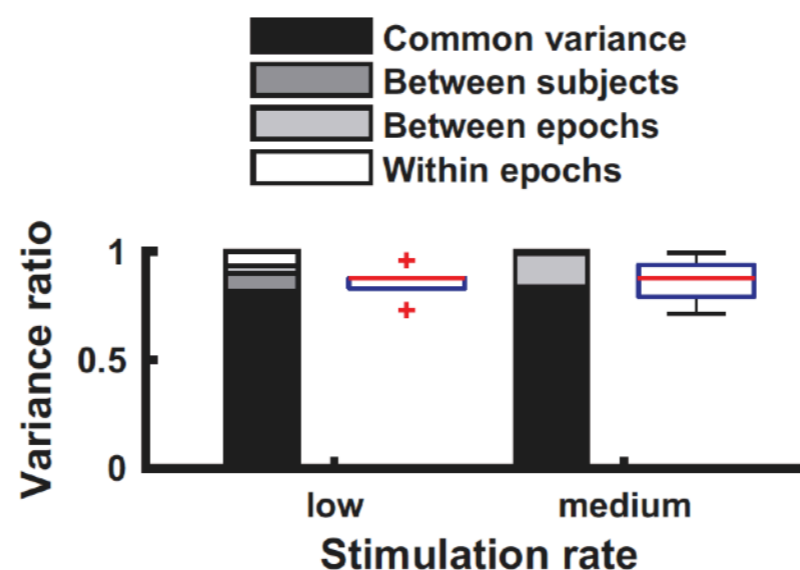
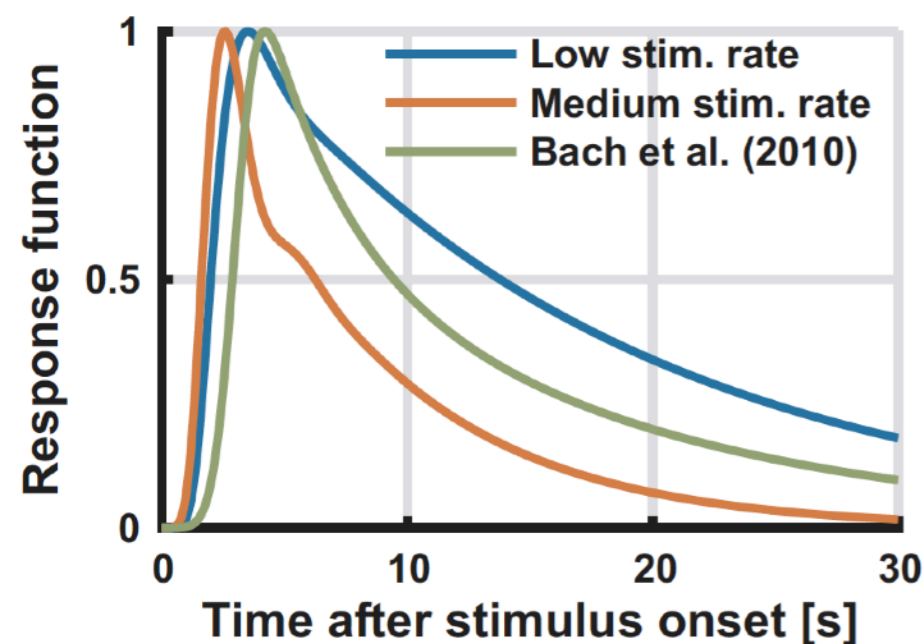
Estimated response amplitude



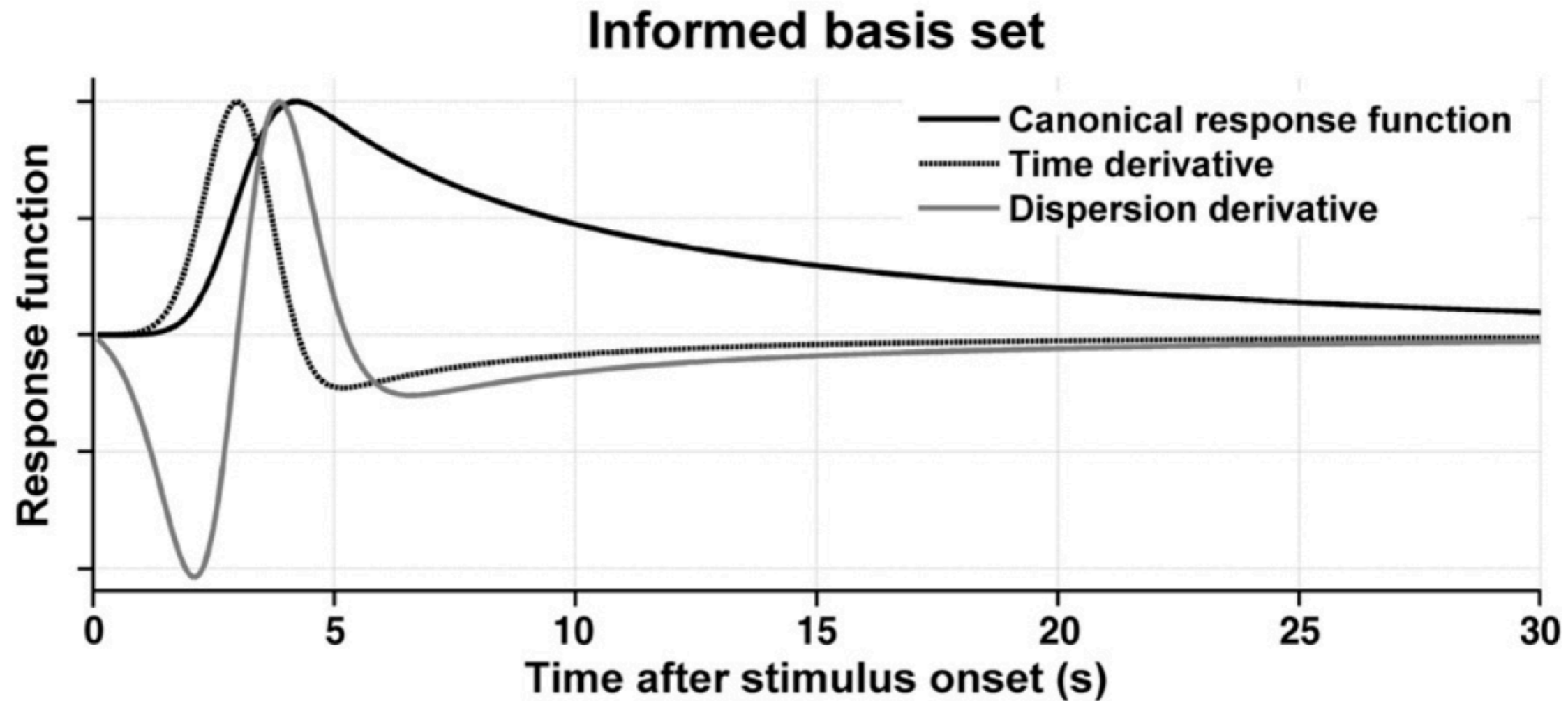
Intraneural recordings



Intraneural stimulation (under peripheral anesthesia): results for frequencies below 0.6 Hz



High trial-by-trial variability in gain factor, but no evidence for peripheral habituation

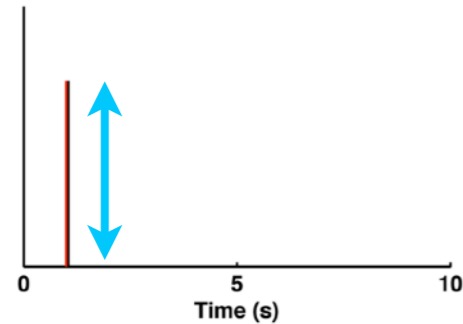


$$h(t) \propto \int_0^t N(t-\tau) (E_1(\tau) + E_2(\tau)) d\tau, \quad t \geq 0,$$

$$N(t) = \frac{1}{\sqrt{2\pi}\sigma} e^{-(t-t_0)^2/2\sigma^2},$$

$$E_i(t) = e^{-\lambda_i t},$$

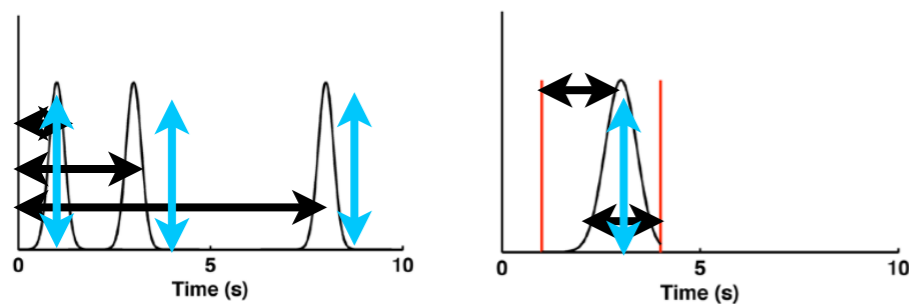
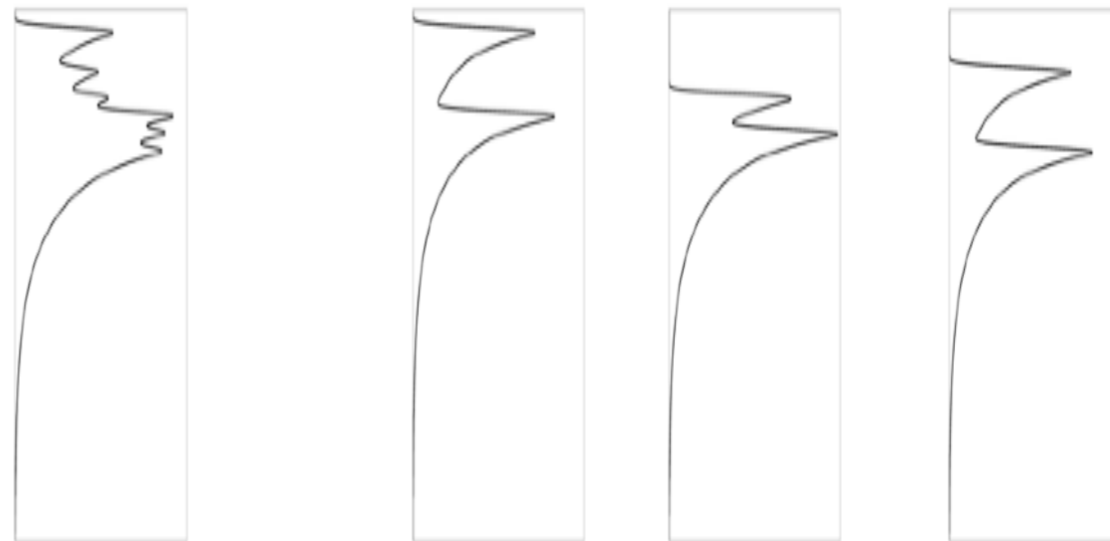
$$\ddot{y} + \vartheta_1 \ddot{y} + \vartheta_2 \dot{y} + \vartheta_3 y - u(t - \vartheta_4) = 0$$



General Linear Model

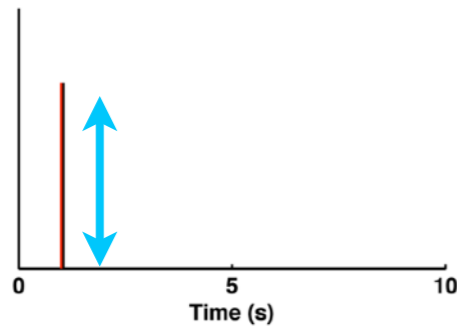
$$\hat{\beta} = \arg \max L(SCR|\beta)$$

$$SCR = \beta_0 + \beta_1 X_1 + \beta_2 X_2 + \beta_3 X_3 + \varepsilon$$



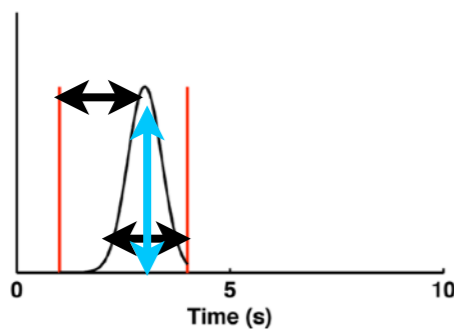
Non-linear methods:

- maximum-a-posteriori estimation using Variational Bayes ("DCM")
- for SF also: approximation using Matching Pursuit (100 x faster)



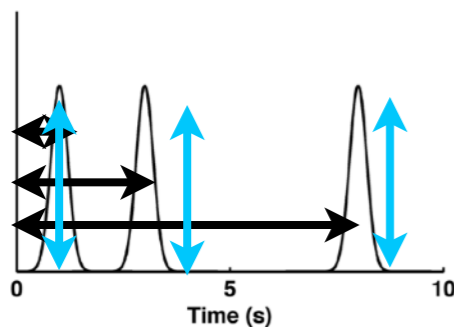
$$SN = \begin{cases} a & t = 0 \\ 0 & \text{otherwise} \end{cases}$$

evoked SCR:
constant-latency input



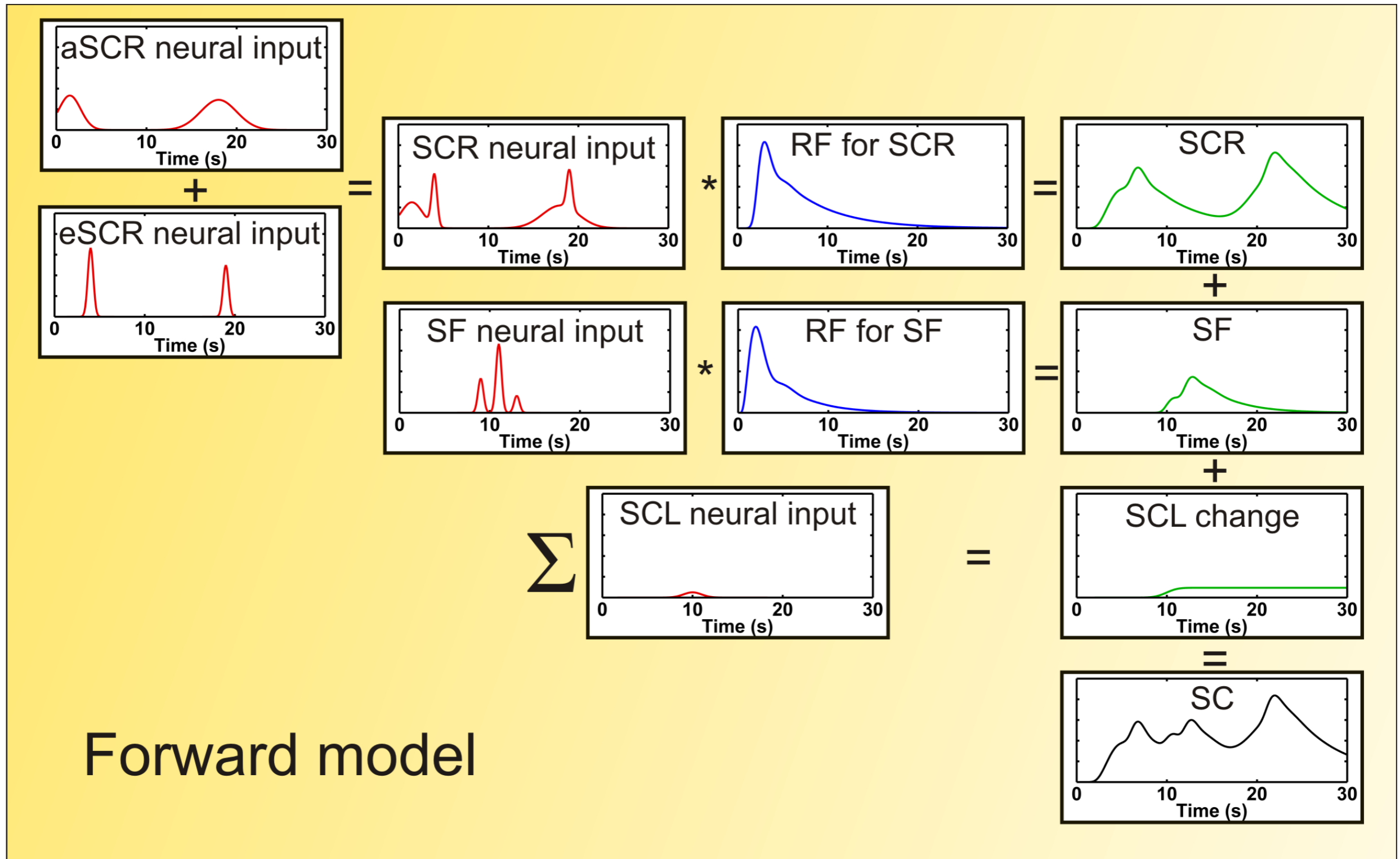
$$SN = a \exp \frac{-(t - \mu)^2}{2\sigma^2}, \quad 0 < \mu < \mu_{max}, \quad \sigma_{min} < \sigma < \sigma_{max}$$

event-related SCR:
constrained-latency input



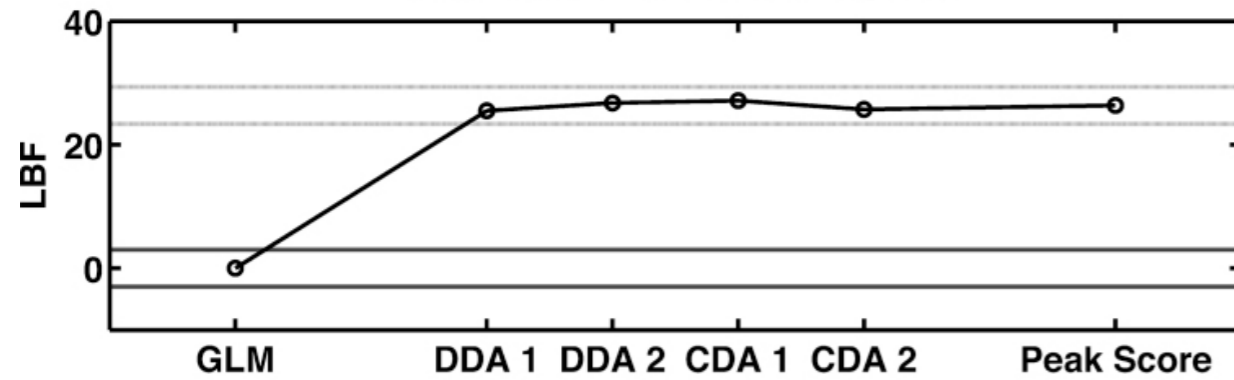
$$SN = a \exp \frac{-(t - \mu)^2}{2\sigma^2}, \quad \sigma = \sigma_0$$

spontaneous
fluctuations (SF):
input at unknown time points

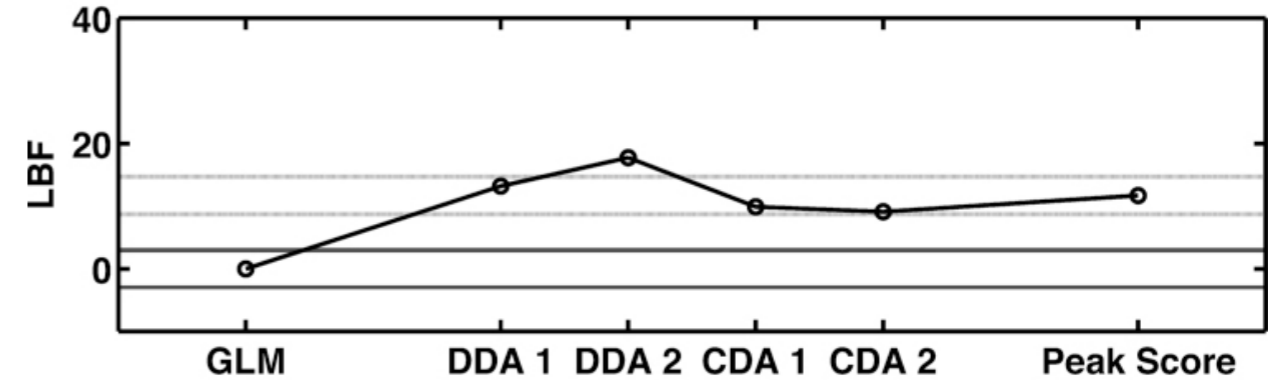


Forward model

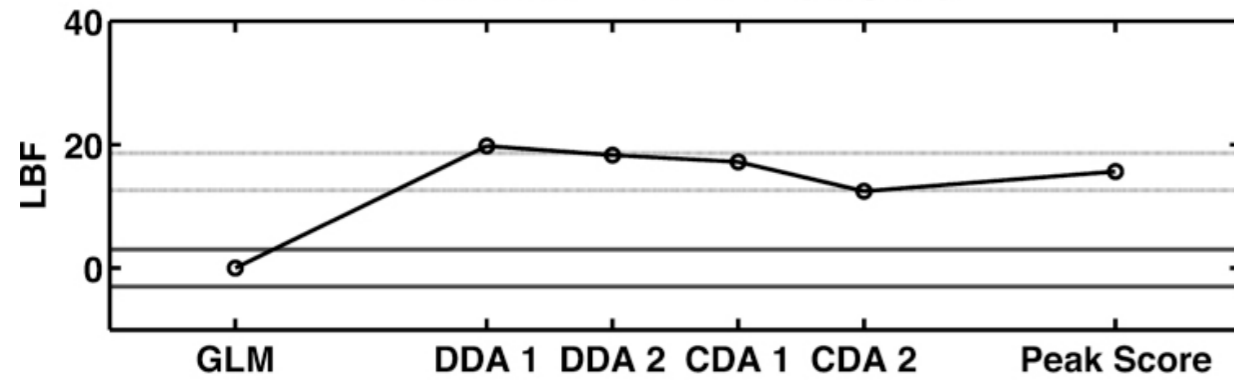
Aversive > neutral (Exp. 1)



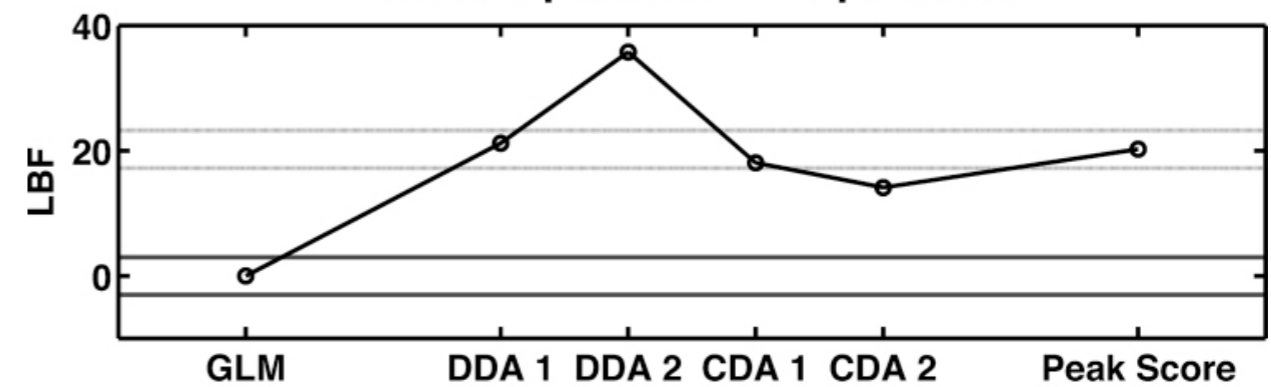
Fearful > angry



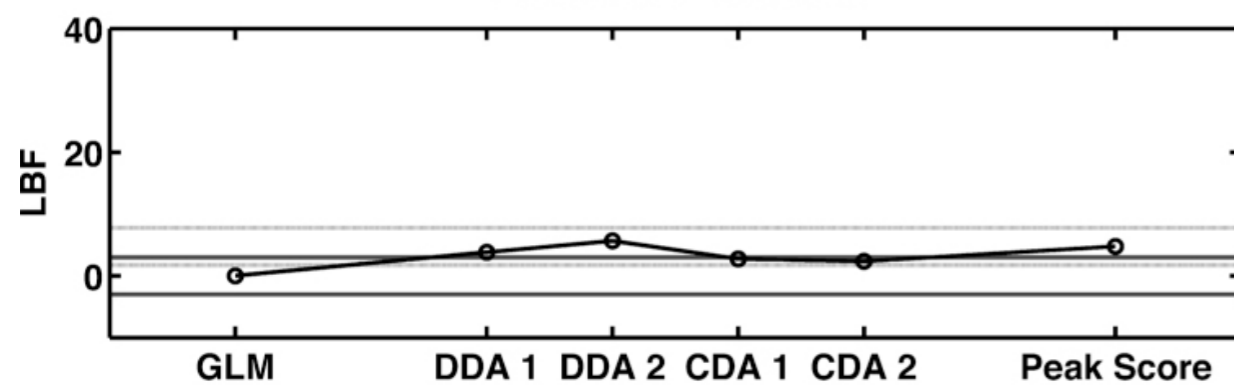
Aversive > neutral (Exp. 2)



Neutral pictures > no pictures



Positive > neutral

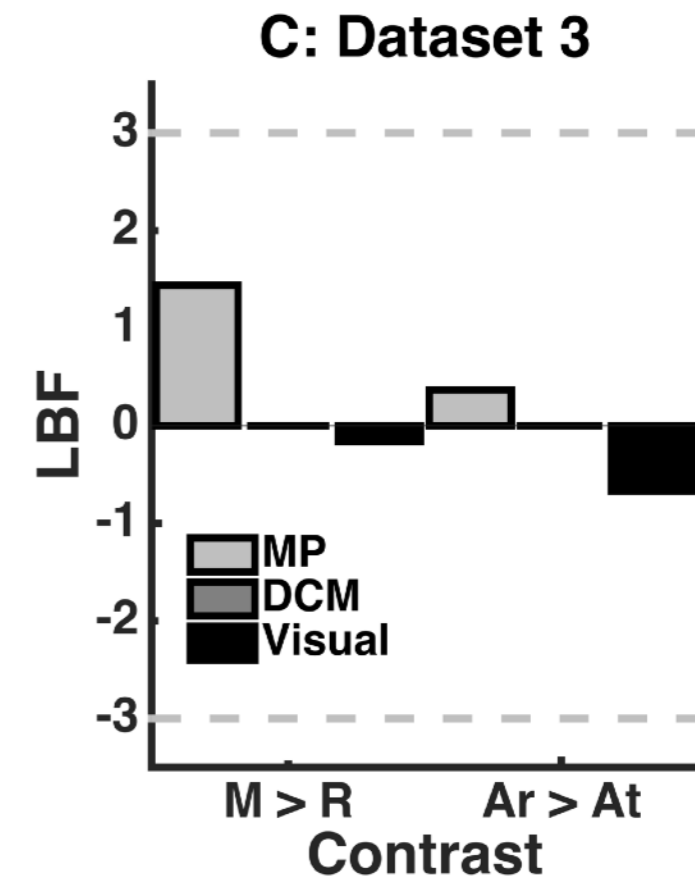
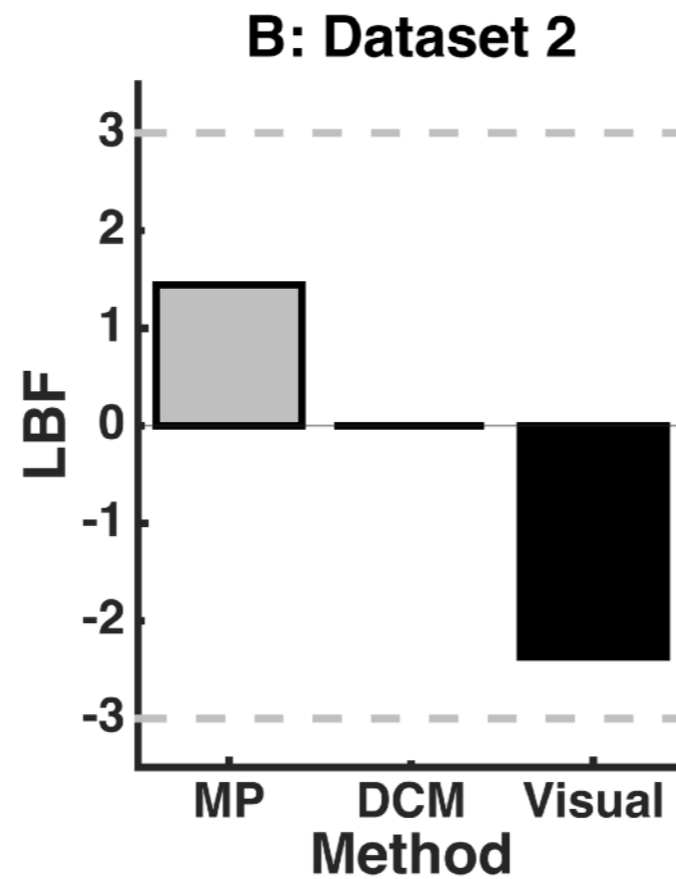
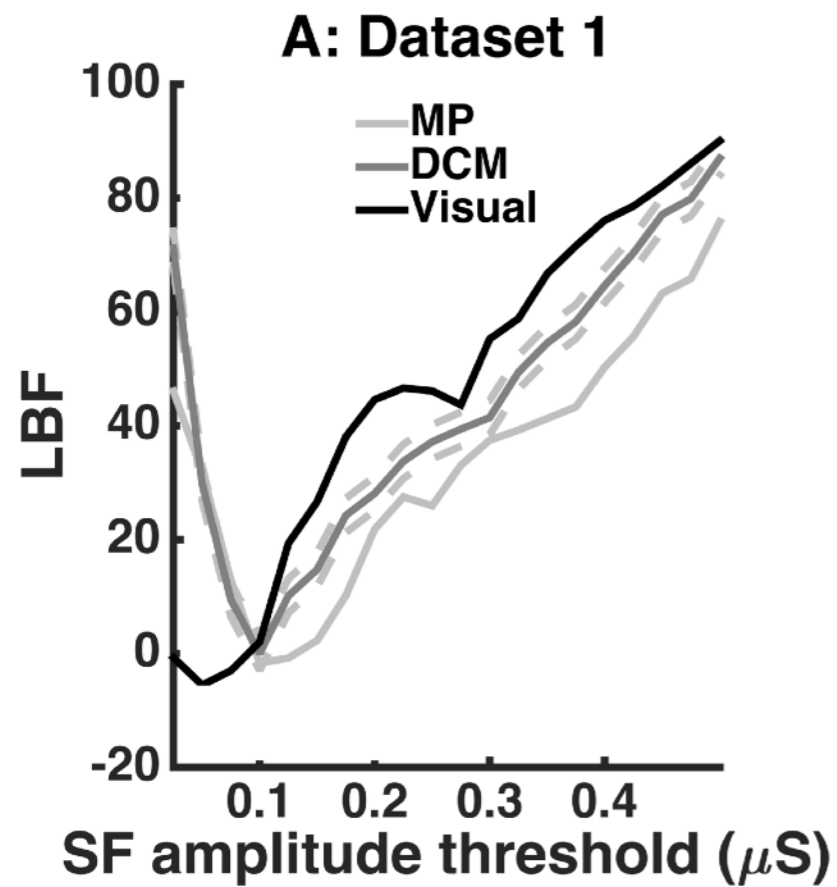


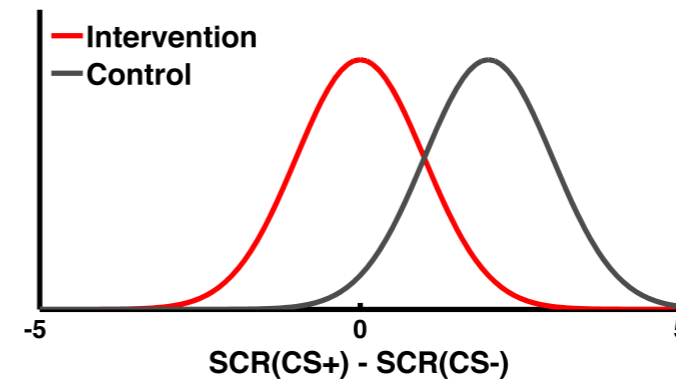
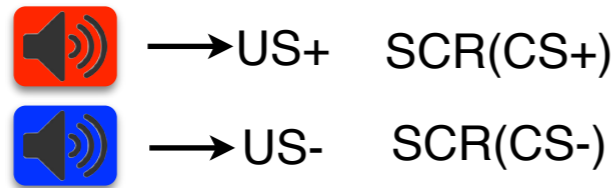
Model evidence (LBF)

- smaller is better
- decisiveness threshold 3

	Experiment 1		
	CS+ > CS-		Comparison with default DCM: LBF (smaller is better)
	<i>t</i> (19)	<i>p</i>	
Default DCM	3.88	0.001	
Peak	1.99	0.062	18
CDA ('AmpSum')	2.39	0.027	15
CDA ('SCR')	2.56	0.019	13
DCM (z-standard)	4.31	<0.001	-2
Peak (z-standard)	2.12	0.047	17
CDA ('AmpSum') (z-standard)	2.59	0.018	13
CDA ('SCR') (z-standard)	2.92	0.009	10

	Experiment 2		
	CS+ > CS-		Comparison with default DCM: LBF (smaller is better)
	<i>t</i> (29)	<i>p</i>	
Default DCM	3.55	0.001	
Peak	2.32	0.027	11
CDA ('AmpSum')	2.77	0.010	08
CDA ('SCR')	2.64	0.013	09
DCM (z-standard)	3.64	0.001	-1
Peak (z-standard)	2.64	0.013	09
CDA ('AmpSum') (z-standard)	2.95	0.006	06
CDA ('SCR') (z-standard)	2.94	0.006	06





	Measure	d	N*
	SCR peak scoring	0.44	514
	SCR model-based	0.75	174
	HPR model-based	0.97	108
	RAR model-based	0.61	268
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*: Sample size required to achieve 80% power at $\alpha = .05$ in a one-tailed test, if intervention reduces fear memory at least 50% and has no variability (best-case scenario)

Filtering (done on-the-fly)

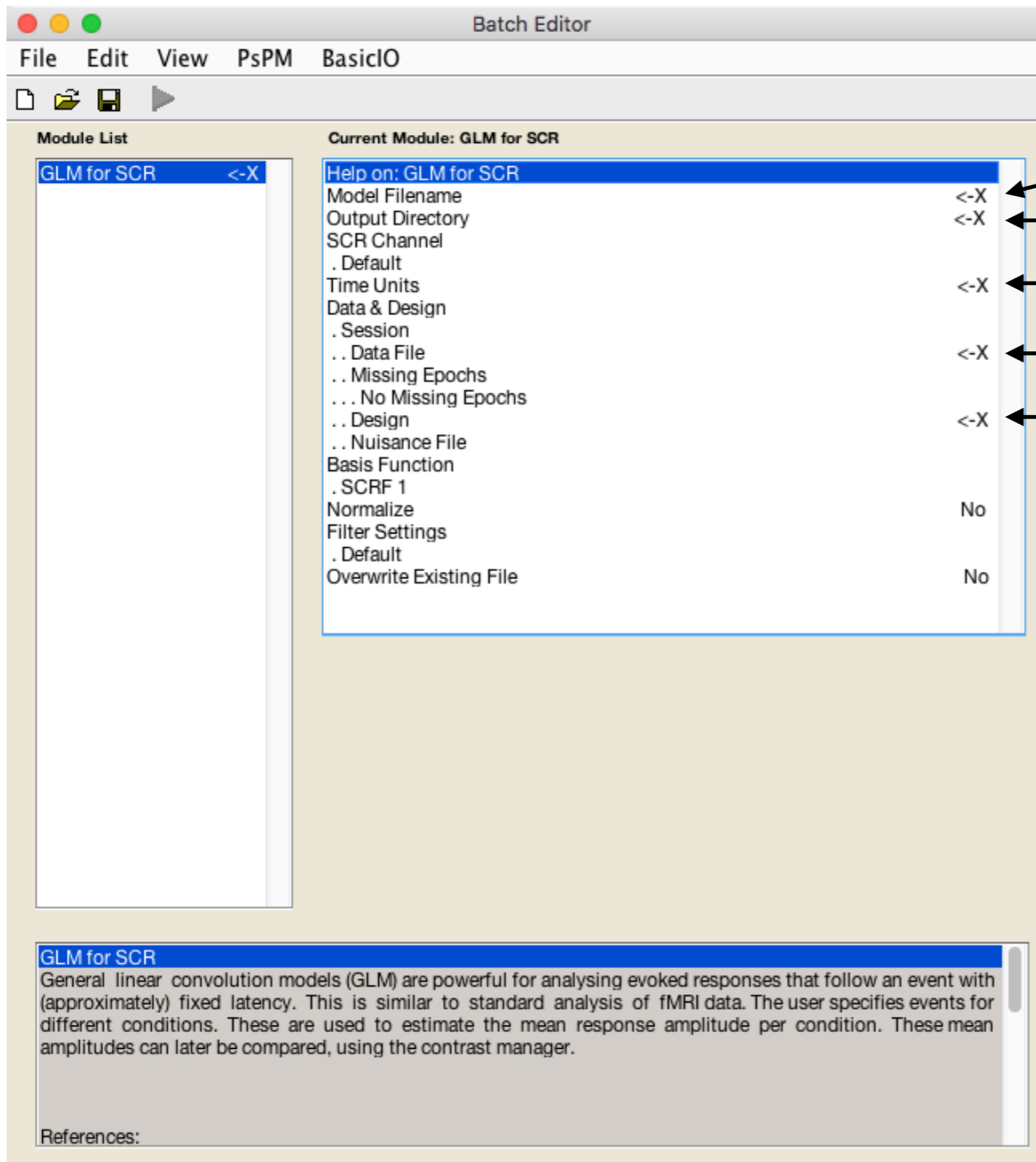
- Low-pass 5 Hz, downsampling to 10 Hz
- Optimal high-pass filters (empirically determined): 0.05 Hz unidirectional (GLM), 0.0159 Hz bidirectional (non-linear model), 0.0159 Hz unidirectional (SF model, not optimised)

Response function

- Canonical SCRF, applies to all manipulations/conditions
- Interpret parameter estimates for individual conditions and trials
- GLM: constrained RF with canonical RF + time derivative is best [1]
- non-linear model: no benefit from IRF if estimated from short ITI data (7-11 s, Staib et al. 2015) but could be beneficial if estimated from external data set with long ITI

Conclusion & open questions

- GLM: fairly mature model, optimised preprocessing and RF
- DCM:
 - best neural forward model for fear conditioning with different SOAs
 - global optimisation as a faster and more precise inversion algorithm



Name for 1st level model file ...

... and directory

Seconds, samples, markers?

Data file (1 per session)

Timings (specify in GUI or 1 file per session)

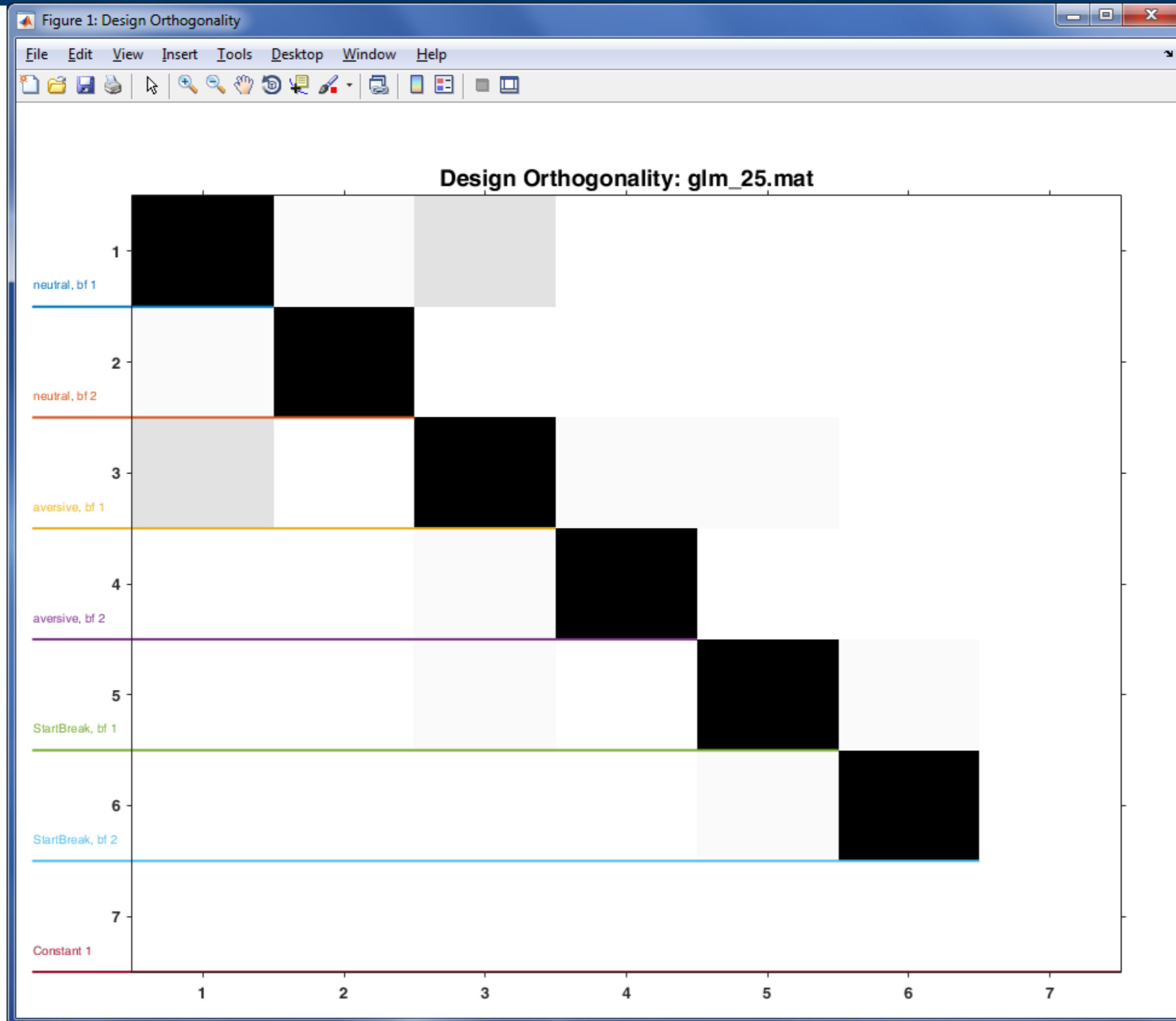
SPM style timing files:

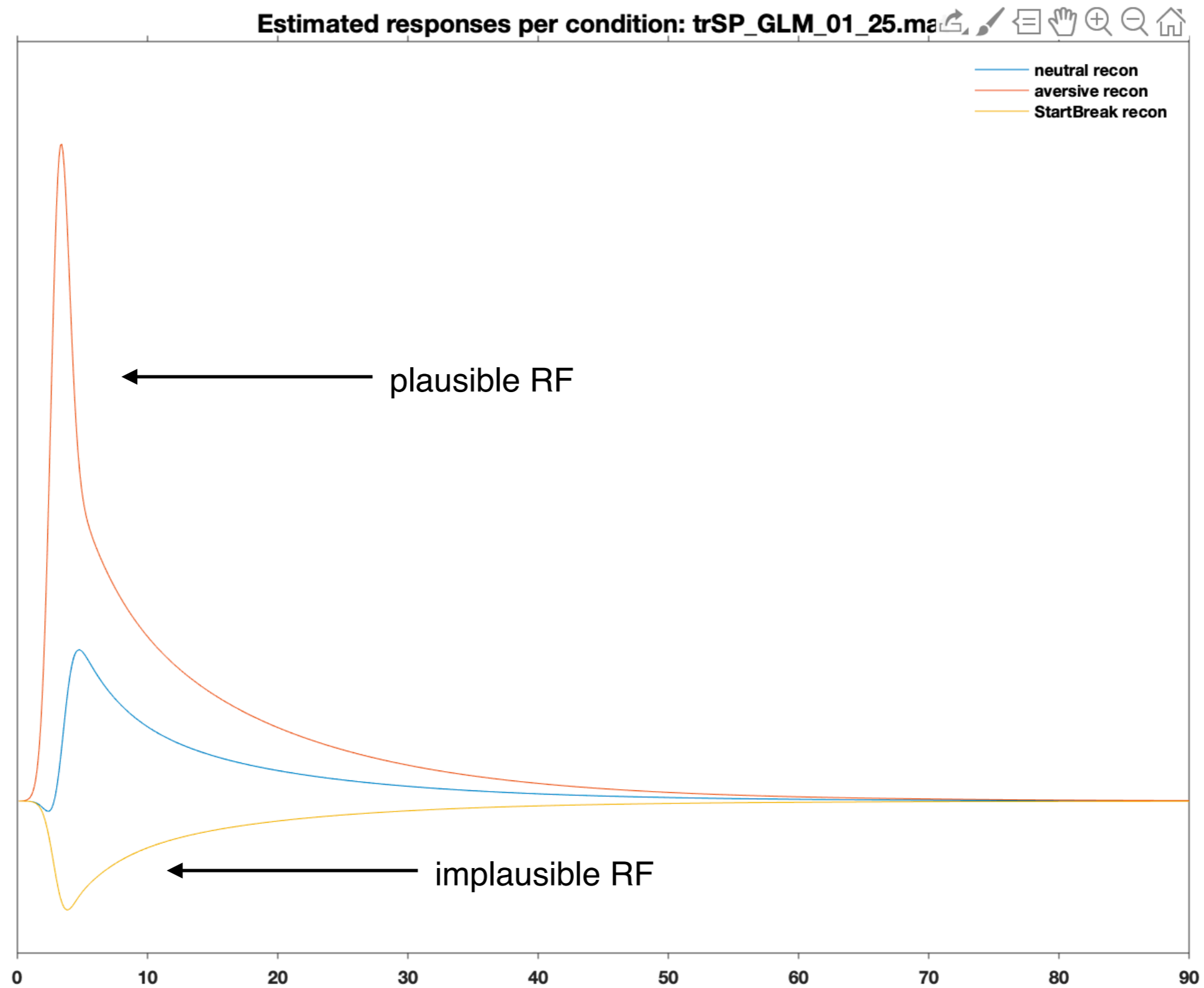
```
names = {'Neutral', 'Aversive'};  
onsets = {[1, 5], [2, 3, 4]};
```

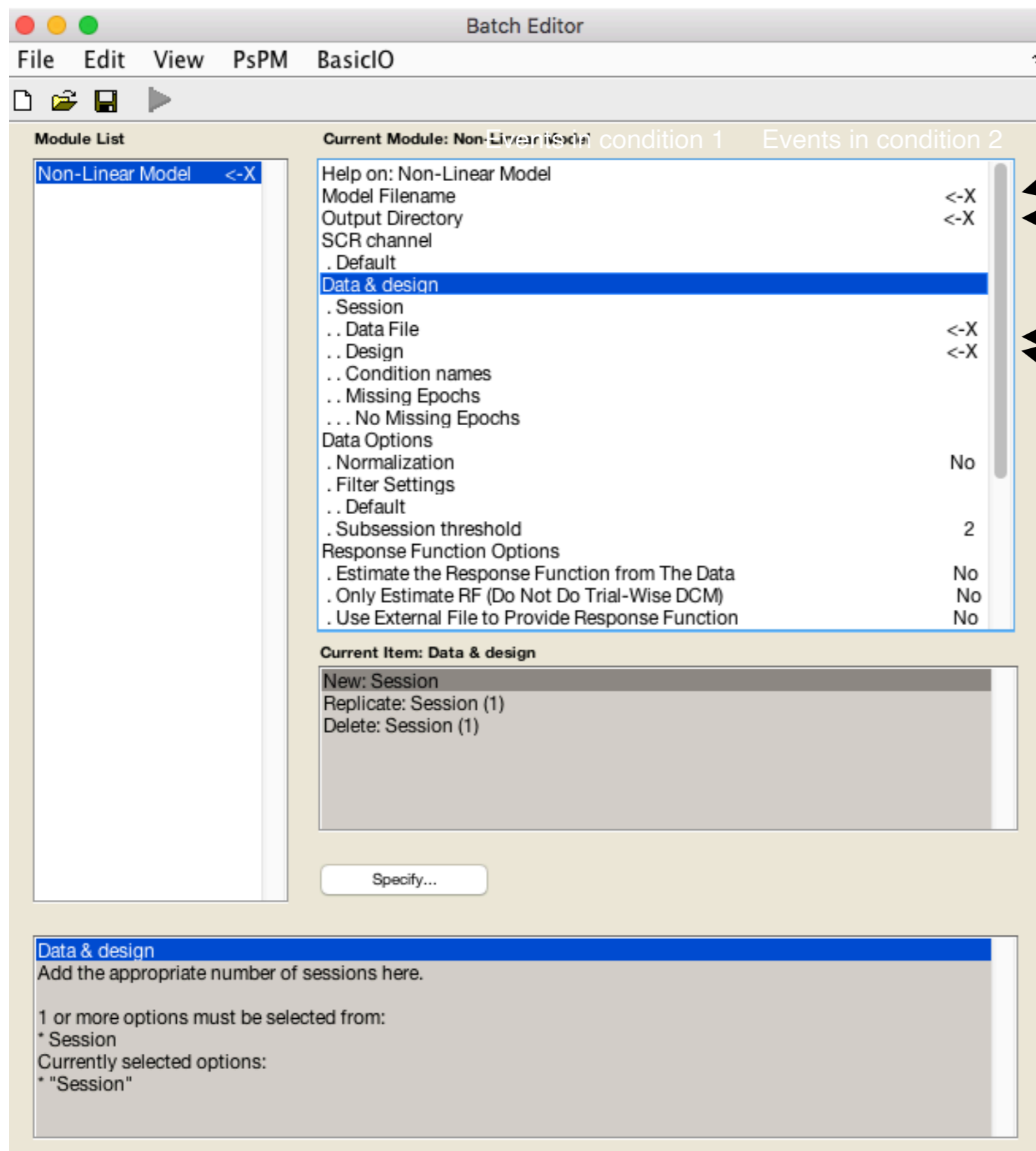
```
pmod(1).name{1} = 'Arousal_rating';  
pmod(1).param{1} = [0.1 1.3];
```

```
pmod(2).name{1} = 'Arousal_rating';  
pmod(2).param{1} = [4.7, 8.3, 7.0];
```

Design orthogonality







Name for 1st level model file ...

... and directory

Timing always in seconds!

Data file (1 per session)

Timings (specify in GUI or 1 file per session)

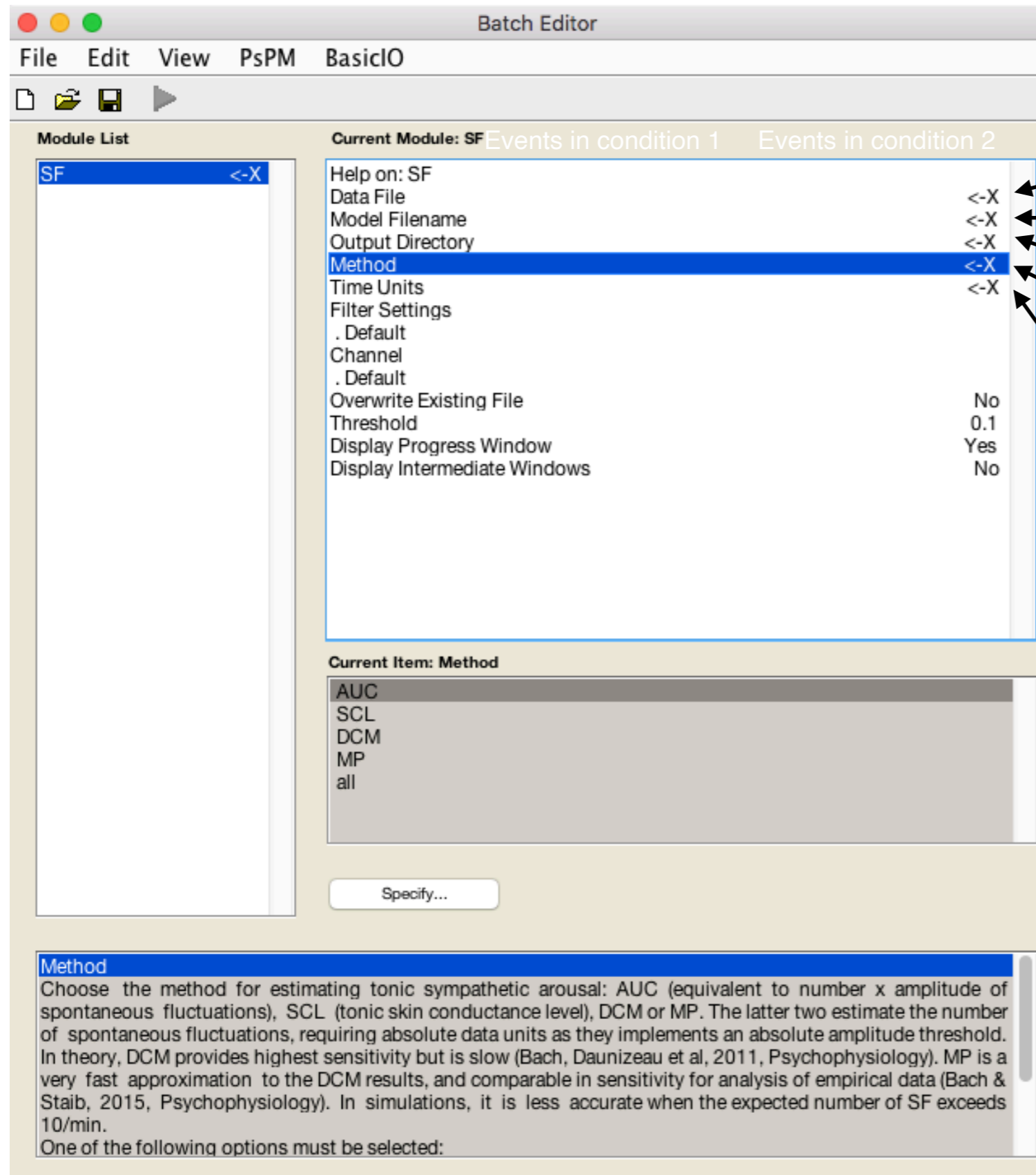
Specific conventions, different from GLM timing files.

Flexible responses (constrained latency & dispersion):

```
events{1} = [1 4;
            10 14];
```

Fixed responses (constant latency & dispersion):

```
events{2} = [5;
            15];
```



Data file (1 per session)

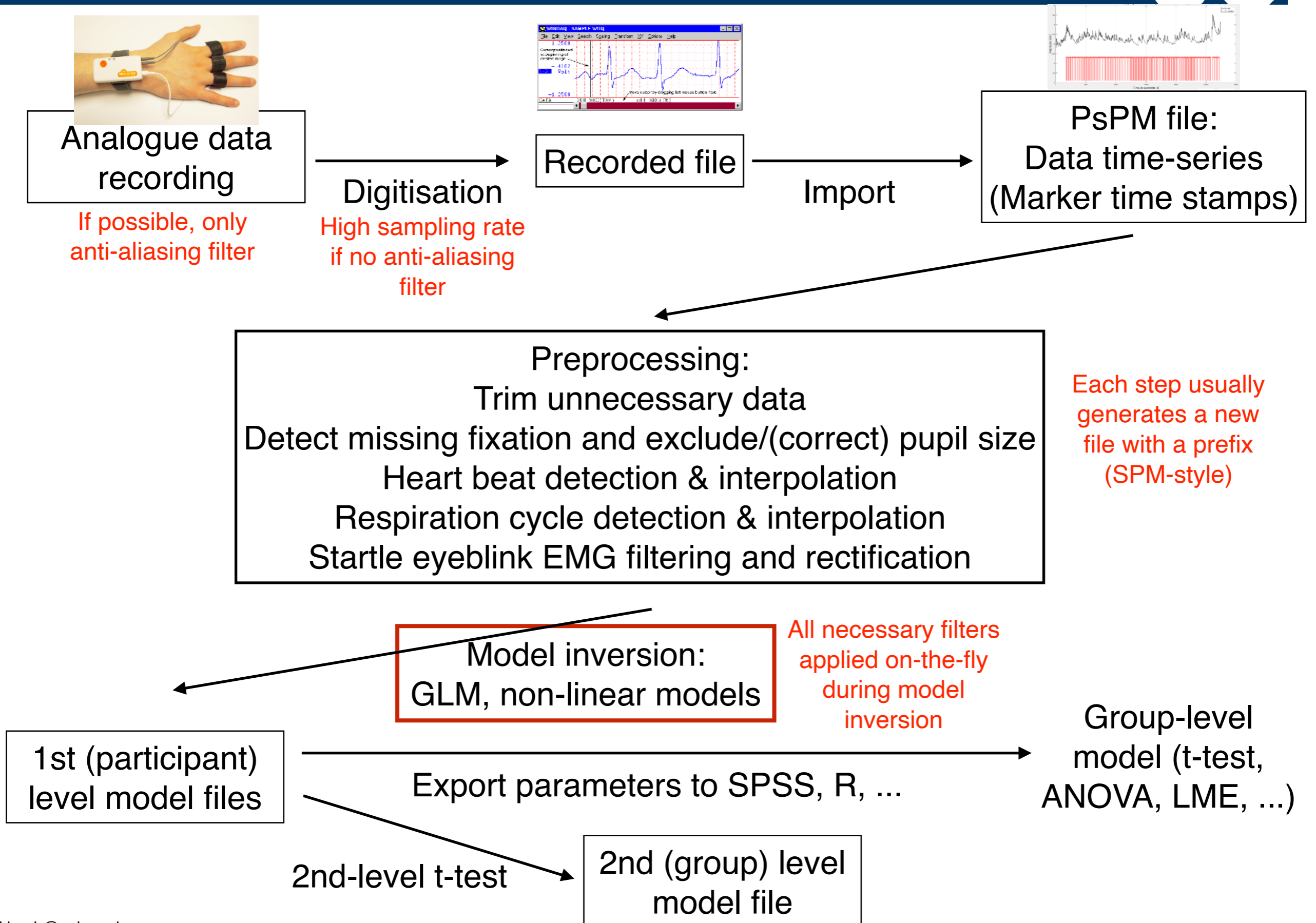
Name for 1st level model file ...

... and directory

AUC (mean-baseline), SCL (mean), DCM (number of responses), MP (number of responses), all

Timings: seconds, samples, markers (provide epochs) or entire file

PsPM pipeline overview: today's tutorial



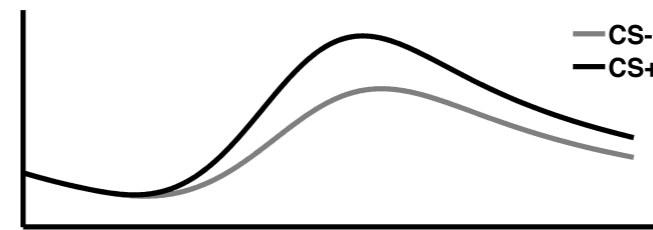
Next lecture: Pupil responses (Christoph Korn)



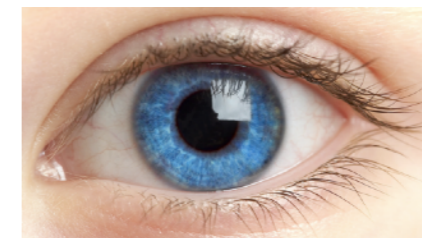
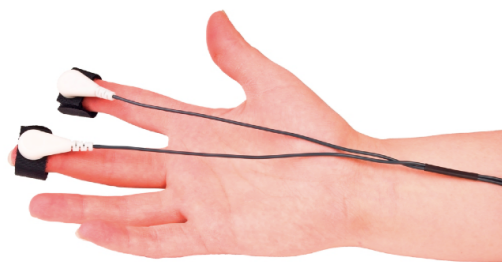
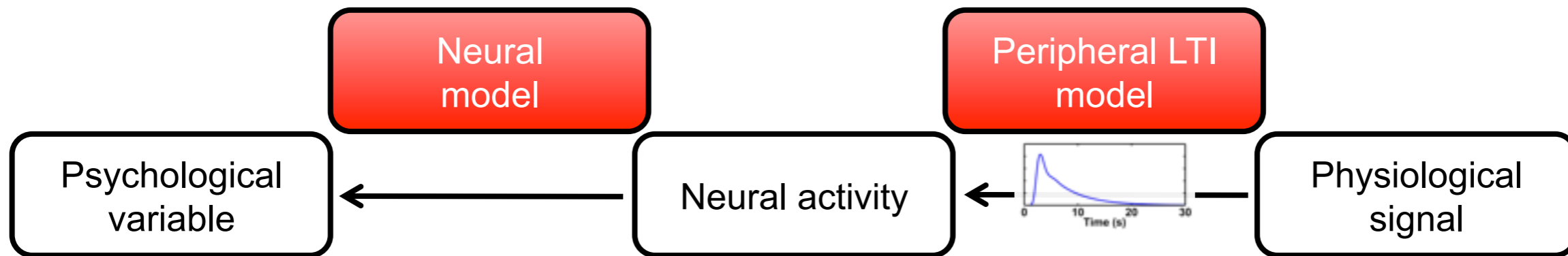
The "best possible" approximation to the true psychological variable.

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Thank you!

Funders



SWISS NATIONAL SCIENCE FOUNDATION



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