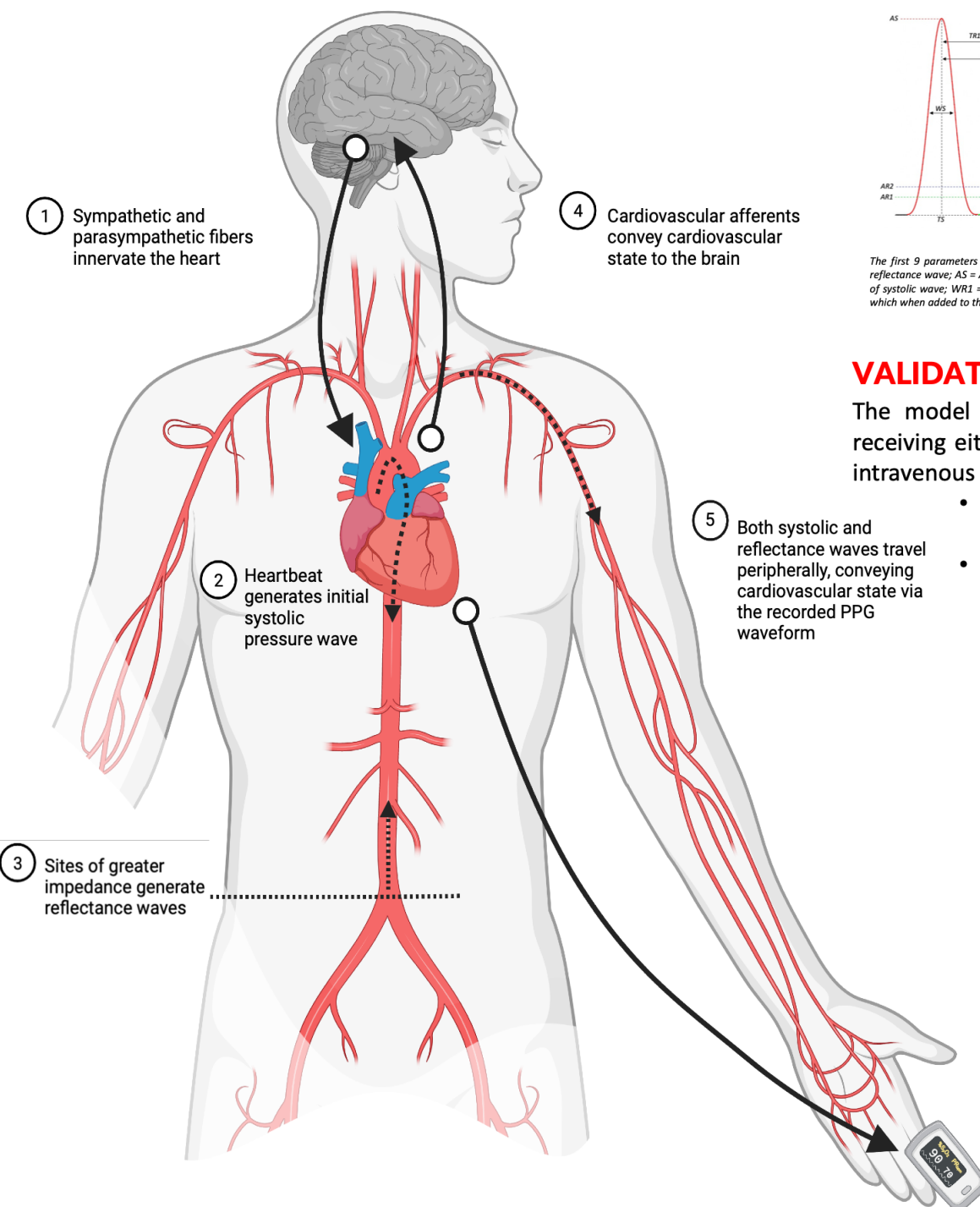


Photoplethysmography: a window onto psychological processes?

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INTRODUCTION

- The photoplethysmography (PPG) signal is often used for **basic assessment** of heart rate and heart rate variability. The signal's waveform, however, is **rich in cardiovascular information**. It is shaped by both **psychological** and **physiological** factors that influence **the entire arterial tree**.
- By harnessing the waveform, neuroscience studies could **better characterize** psychophysiological states within and between individuals. We developed an **automated pipeline** and model for systematic analysis of PPG waveforms.

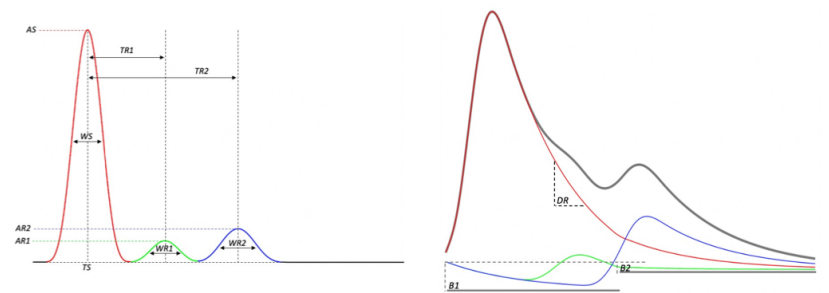


DISCUSSION

- PulseWaveform offers a new level of nuance in non-invasive cardiovascular measurement, and a new means to characterize brain-body interactions. This is directly relevant to disorders of perturbed interoception (e.g. panic disorder) and those with established vascular changes (e.g. schizophrenia or depression).
- With the rise in popularity of wearable devices, the potential for PPG to aid in the study and monitoring of mental health conditions is significant.

THE 'PulseWaveform' PACKAGE

- PulseWaveform is a PPG analysis pipeline that isolates waveforms to extract features reflective of cardiovascular state.
- As well as traditional measures of morphology (fiducial points), PulseWaveform enables the decomposition of waveforms into three underlying pressure waves – one systolic and two reflectance.
- Decomposition is achieved using the Hybrid Excess and Decay (HED) Model, which models waveforms as three component waves and an exponential decay:

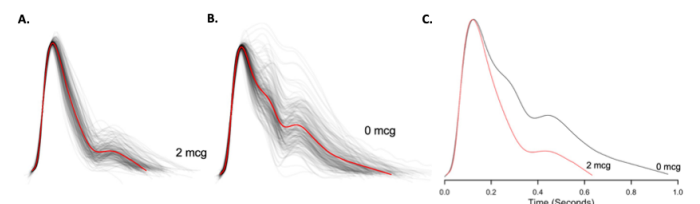


The first 9 parameters (left) are the excess element. TS = Timing of systolic wave; TR1 = Timing of 1st reflectance wave; TR2 = Timing of 2nd reflectance wave; AS = Amplitude of systolic wave; AR1 = Amplitude of 1st reflectance wave; AR2 = Amplitude of 2nd reflectance wave; WS = Width of systolic wave; WR1 = Width of 1st reflectance wave; WR2 = Width of 2nd reflectance wave. 2b: The final 3 parameters are the decay element, which when added to the excess yield the final modelled waveform (right). DR = Decay rate; B1 = 1st baseline; B2 = 2nd baseline.

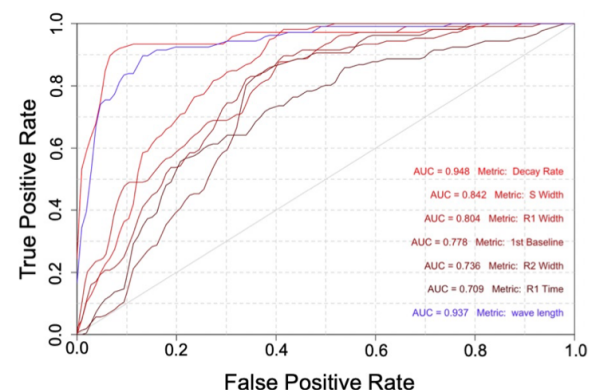
VALIDATION OF THE HED MODEL

The model was applied to PPG data of participants (n = 105) receiving either saline or the adrenaline-like drug isoproterenol by intravenous infusion:

- Waveforms were modelled with high precision (median NRMSE = 0.9).
- The change in waveform shape induced by isoproterenol was best classified by a model parameter (Decay rate; AUC = 0.948), which numerically outperformed all traditional measures (heart rate; AUC = 0.931).



A: Averaged waveforms (black) during pharmacological manipulation (one per subject/time series) aligned by systolic peak. The mean of these (red) is overlaid. B: Averaged waveforms during infusion of saline (one per subject/time series) aligned by systolic peak. The mean (red) is overlaid. C: The mean waves derived from A and B, overlaid to display the morphological change between saline and 2 mcg isoproterenol infusion.



ROC curve of top performing (highest AUC) model parameters (with best performing morphological feature, wavelength, overlaid in blue) in classifying between 0 mcg and 2 mcg morphologies.

