

## Background

Historically, collecting physiological recordings requires movement to be restricted such as MRI and EEG. In both cases the recording of interest can be confounded by a single minute movement of the head or an eye blink. Similarly, the effectiveness of data collected from a wrist-worn bio-sensor is sensitive to small environmental factors (Boucsein, 2012).

In this experiment we identify a particular artifact that would likely occur as a result of normal movement, to further remove this uninformative segment from the data. This is a novel method since most approaches to filtering data control for and attend to the informative data rather than the uninformative data lacking a physiological covariate (Taylor, *et al.*, 2015).

## Experiment Design

This experiment investigated two wrist worn bio-sensors: the Q-Sensor, made by Affectiva, and the E4, made by Empatica (Garbarino, *et al.*, 2014). A total of 38 participants were asked to wear either the Q sensor or the E4 sensor. A subgroup of 23 participants were asked to wear both the E4 and Q-Sensor.

Participants were asked to complete a survey regarding caffeine and food intake, as well as sleep quality. At least three minutes were allowed for the sensors to reach close to baseline levels of EDA. An experimenter then lifted each sensor from the wrist five times, indicated by an event marker. Participants were asked to repeat this process, lifting the sensor from their own wrist.

### Empatica E4 Specifications

**Battery life**  
 Streaming Mode: 20+hrs  
 Memory mode: 36+ hrs

**Data Management**  
 Flash memory  
 Bluetooth LE (Smart)

**Form Factor**  
 Small and comfortable  
 Case: 44 mm x 40mm, height 16 mm  
 Weight: 25 gr

**Event Mark Button**

**Certification**  
 CE certification  
 FCC certification

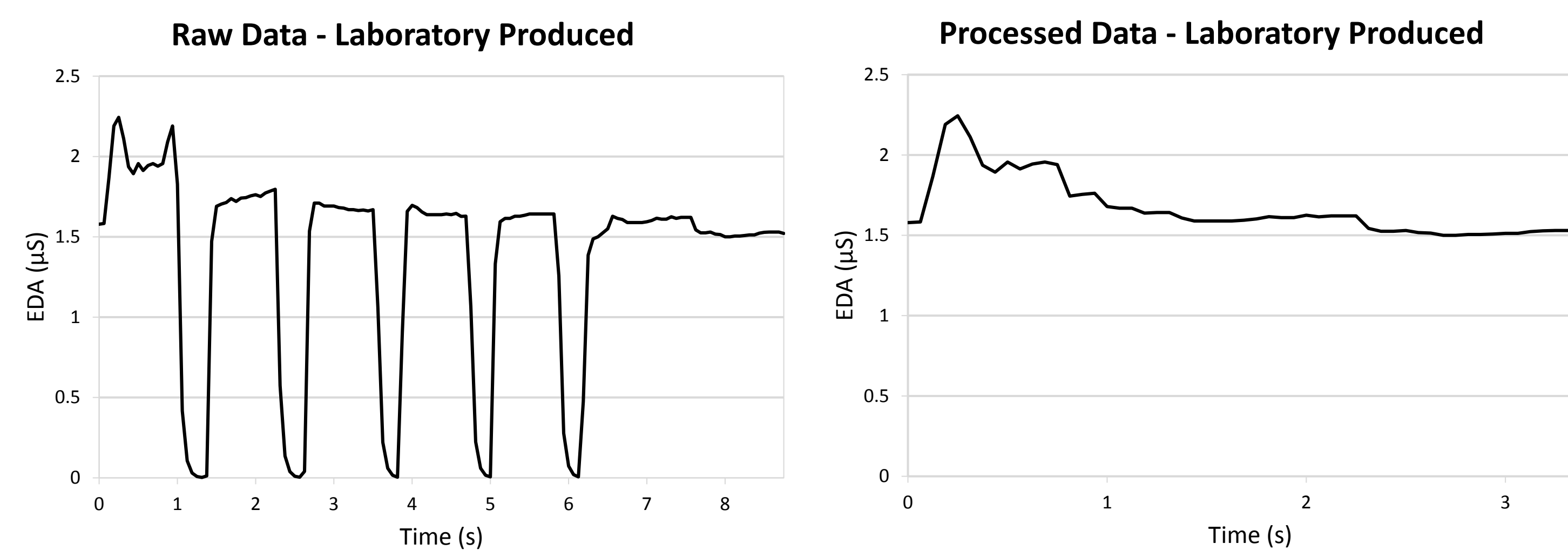
**Sensors**  
 Photoplethysmography (PPG)  
 Continuous Heart Rate (HRV, Stress, Relaxation)  
 3-axis Accelerometer  
 Movement, Activity  
 Temperature + Heat flux  
 Activity, Context  
 Electrodermal Activity (EDA)  
 Skin conductance (Arousal, Excitement)



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## Laboratory Generated Artifacts

We then compared each artifact to the normalized artifact by a modified Kolmogorov-Smirnov two sample test at an  $\alpha$  level of 0.05. Potential epochs were labelled as any two points having a slope  $< -2$ . After labelling, the lowest point within one second of the flag was queried, including each .5 or 2 second segment surrounding it. If resulting in a significant KS test, data was left untouched, otherwise removed. KS analysis flagged 100% of the controlled artifacts.



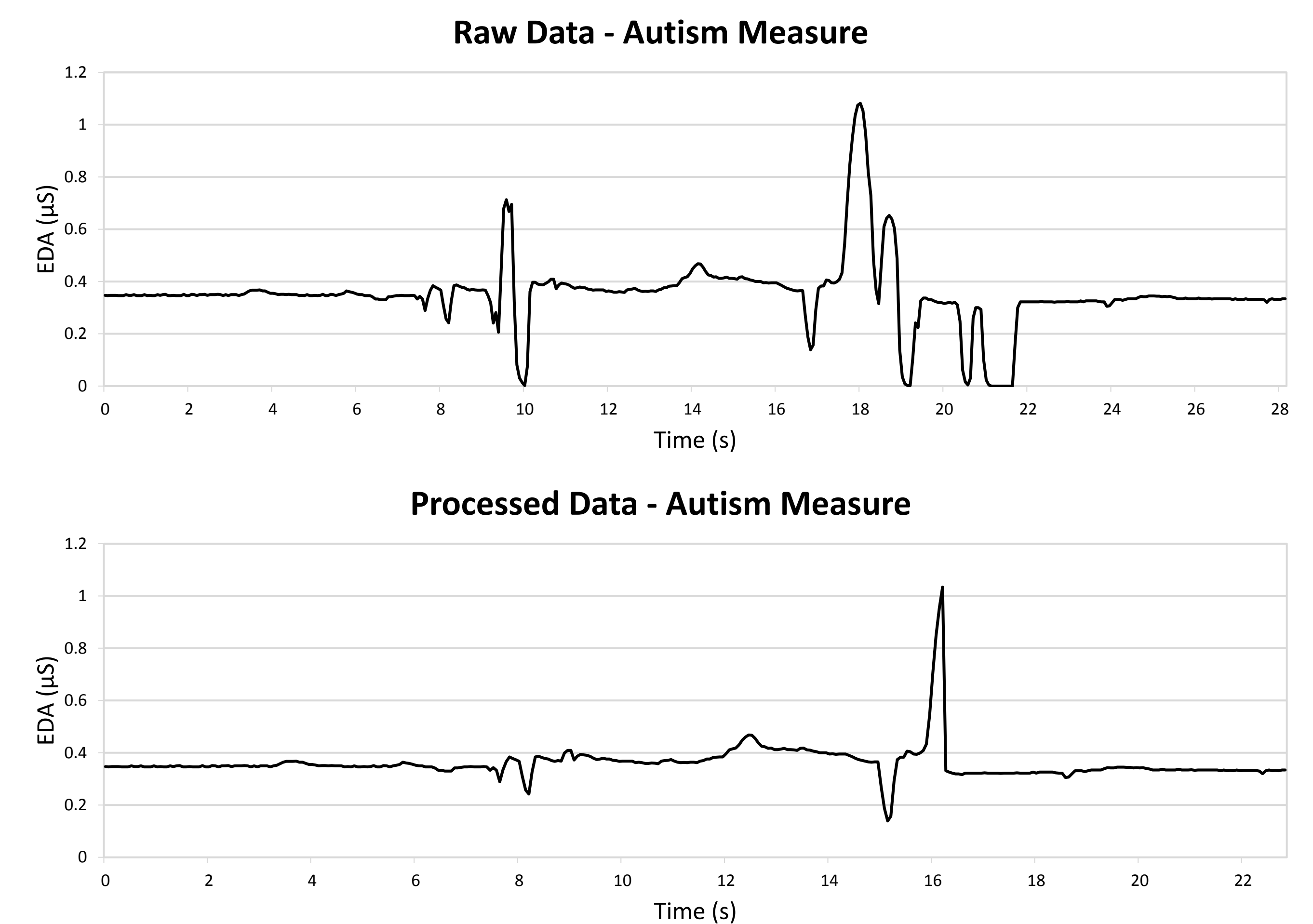
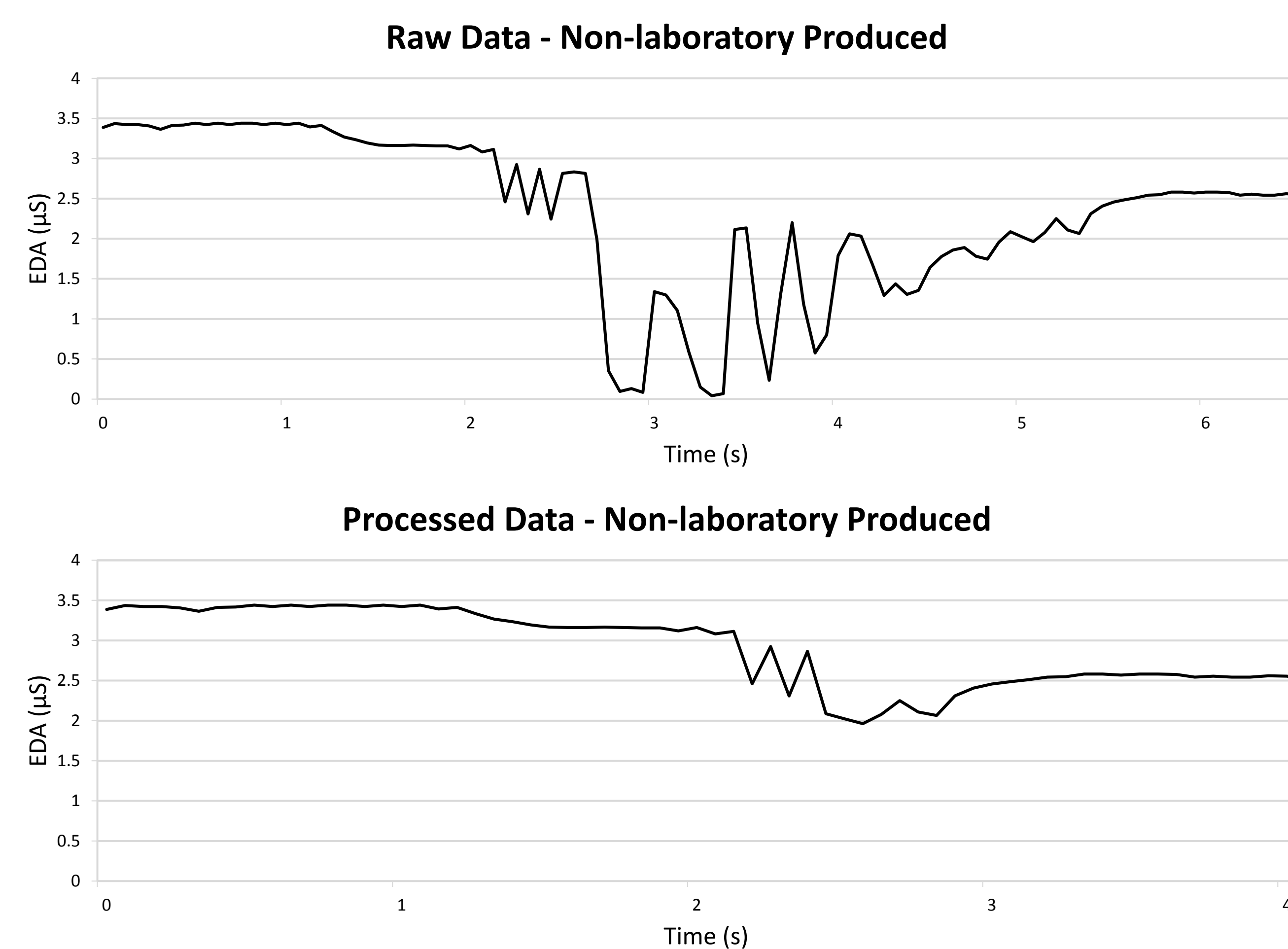
## Analysis

Each artifact was manually parsed from the data. Three participants with abnormally low EDA ( $< .1\mu\text{S}$ ) were excluded from the study. The controlled artifacts were observed to be, and subsequently defined by 155 one-second and 80 four-second artifacts for the Q and E4 sensor, respectively. The lowest value in the given epoch was identified. This value was surrounded by .5 seconds (Q Sensor) or 2 seconds (E4 Sensor) of data. Each epoch was averaged among participants to generate a normalized artifact.

## Conclusion

We have shown that comparisons of raw data to normalized artifacts can be valid measures to strongly reduce variability and can be done so automatically. This greatly reduces the impact of sensor disconnect on aspects of EDA such as baseline levels and arousal reactivity. Externally, this method could aid in analysis of both typically developing and autistic individuals. This helps reduce the effect of data artifacts in an already sensitive measure without unnecessarily discarding data.

## Externally Generated Artifacts



The same approach was applied to a noisy EDA sample collected in a non-laboratory setting. All artifacts specific to electrode disconnect were accurately identified, as determined by an expert labeler. Of the 6.5 second sample, 2.25 seconds were determined to fit the normalized artifact and removed.

Two-way t test yielded a significant increase in mean EDA of  $.577\mu\text{S}$  ( $t(162) = 5.2, p < .001$ ). Test for equal variances yielded a  $\Delta\text{SD}$  of  $-.49\mu\text{S}$  ( $F(1,170) = 16.02, p < .001$ ). We performed the same process on a noisy EDA sample collected from an individual diagnosed with autism. Two-way t test yielded no significant change in mean EDA ( $t(696) = -1.21, p = .227$ ), however Levene's test for equal variance did show a  $\Delta\text{SD}$  of  $-.079\mu\text{S}$  ( $F(1,815) = 21.94, p < .001$ ).

### References

- Taylor, S., Jaques, N., Chen, W., Fedor, S., Sano, A., & Picard, R. (2015, August). Automatic identification of artifacts in electrodermal activity data. In *Engineering in Medicine and Biology Society (EMBC), 2015 37th Annual International Conference of the IEEE* (pp. 1934-1937). IEEE.
- Boucsein, W. (2012). *Electrodermal activity*. Springer Science & Business Media.
- Garbarino, M., Lai, M., Bender, D., Picard, R. W., & Tognetti, S. (2014). Empatica E3 - A wearable wireless multi-sensor device for real-time computerized biofeedback and data acquisition. In *2014 EAI 4th International Conference on Wireless Mobile Communication and Healthcare (Mobihealth)* 39-42.

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