

Identifying Cybersickness through Heart Rate Variability alterations

Augusto Garcia-Agundez ¹, Christian Reuter ², Polona Caserman ¹, Robert Konrad ¹ and Stefan Göbel ¹

¹ KOM – Multimedia Communications Lab, Technische Universität Darmstadt, Darmstadt, Germany

Abstract - Numerous users of Head Mounted Displays develop symptoms similar to motion sickness. This phenomenon is known as cybersickness. The current standard method to evaluate it is the Simulator Sickness Questionnaire (SSQ). Additionally, detection by methods such as biosignals is possible, but it requires sophisticated equipment and expertise. In order to detect early cybersickness symptoms, the availability of real-time detection by using simple equipment, such as a 2-lead ECG, would be highly useful. This contribution aims to analyze the feasibility of detecting cybersickness with a simple 2 lead ECG. A total of 13 participants played the VR game QuakeVR wearing an Oculus Rift DK2 for 15 minutes, during which a 2-lead ECG was recorded. Additionally, pre- and post-SSQ questionnaires were given. Four of 13 participants had to end the experiment prematurely due to cybersickness. The difference in SSQ scores is statistically significant (p<.01), as is the ECG (p=.02), in these participants. This study shows the utility of a simple 2-lead ECG to detect cybersickness. These findings raise the possibility of real-time monitoring and prediction of cybersickness with simple devices and open the question of whether photoplethysmography could be used with the same purpose.

Index Terms - Virtual Reality, Electrocardiography, Cybersickness, Heart Rate Variability

I. Introduction

Cybersickness is a term used to refer to the cluster of symptoms that users experience during, or after, Virtual Reality (VR) exposure (McCauley and Sharkey 1992). It is acknowledged as a physiological response to an unusual stimulus, similar to motion sickness or seasickness (Bouchard, Robillard, and Renaud 2007). The reported incidence of cybersickness amongst users of VR is varied, but at least, 60% of participants in a first VR experience will suffer its symptoms to some degree, and although most users adapt to the environment after few immersions, approximately 5 percent will never do so. The degree of intensity depends on the nature of the VR environment, and previous works have shown the incidence of mild cybersickness ranges between 60% and 90%, with 5-30% of participants having to discontinue research

E-mail: augusto.garcia@kom.tu-darmstadt.de

evaluations due to strong symptoms (Johnson 2005; Regan 1995).

The amount of time required to cause significant sickness depends on the environment, but generally, symptoms can already be expected 5 to 15 minutes after starting (DiZio and Lackner 1997; DiZio and Lackner 2000; Lampton et al. 1994; So and Lo 1999), in some cases even under less than 5 minutes (Dennison, Wisti, and D'Zmura 2016). Although these symptoms usually disappear 15 minutes after the experience is concluded, users can remain sensitized for hours, and re-immersion causes cybersickness to reappear strongly and abruptly, although this is not always the case (DiZio and Lackner 2000). Cybersickness is regarded as a real problem with medical and safety implications, which will surely hamper its integration into the many fields it could be useful for, such as education, entertainment, engineering, and video games (Kennedy and Lilienthal 1995).

Head-mounted devices such as the Oculus Rift have already been proven to cause vection and sickness (Palmisano, Mursic, and Kim 2017), but the way through which vection acts is unknown, and symptoms vary greatly from user to user. Currently, the standard method to evaluate cybersickness is the Simulator Sickness Questionnaire (SSQ) (Kennedy et al. 1993). This questionnaire, derived from a prior motion sickness questionnaire, classifies symptoms into three blocks: Nausea, Oculomotor and Disorientation, and then scores each symptom from 1 (none) to 4 (severe). An SSQ Score of 10 or higher is considered to be substantial, and a score higher than 20 is considered to be due to bad VR design (Stanney, Kennedy, and Drexler 1997).

At the moment, no specific questionnaire exists to address cybersickness. However, studies have been done (Bruck and Watters 2011) on the viability of the Simulator Sickness Questionnaire (SSQ) as a correct tool to assess cybersickness. These studies suggest that, in the specific environment of virtual reality, symptoms can be structured in 4 factors that account for 78% of the alterations in physiological behavior: General cybersickness (cardiac activity, fatigue, increased nausea and salivation, burping, anxiety), vision factor (eyestrain and headache), arousal (respiration alterations) and fatigue (blurred vision and dizziness), where general cybersickness alone accounts for 38.75% of biosignal variation. Nevertheless, the design of

² Deck13 Interactive GmbH, Franfurt am Main, Germany

a questionnaire specific to cybersickness remains an issue, especially when considering that it has been proven that the pathophysiology of cybersickness and simulator sickness is not identical. Curiously enough, in (Wilson 1996) and (Young, Adelstein, and Ellis 2007), the possibility whether taking an SSQ/MSQ before and after, or just after the experiment, has a measurable effect on the amount of sickness reported, was studied. Results suggest that indeed reports of motion sickness after immersion are greater when a preemptive questionnaire (such as MSSQ) is given. That is, giving a pre-test SSQ substantially increases the post-test SSQ score. This stresses the importance shown by previous research on attempting to move to more objective measurement methods, such as postural sway (Chardonnet, Mirzaei, and Merienne 2017) or biosignals.

Previous studies have reported for cybersickness to increase the levels of cortisol in saliva (Kennedy, Drexler, and Kennedy 2010), cause tachygastria, correlate with sweating and respiratory rate variations (Johnson 2005) heart rate variability (Gavgani, Hodgson, and Nalivaiko 2017; Malińska et al. 2015; Nakagawa 2015; Rieder, Kristensen, and Pinho 2011) and galvanic skin response (GSR) (Gavgani et al. 2017). Other studies state that the onset of cybersickness is induced by specific factors (or triggers), but the increase in cybersickness depends on individual differences in autonomic regulation, which makes it difficult to detect it (Kiryu and Iijima 2014). A summary of the results from studies attempting to correlate vital parameters and cybersickness (Dennison, Wisti, and D'Zmura 2016; Gavgani, Hodgson, and Nalivaiko 2017: Gavgani et al. 2017; Kim et al. 2005; Nam et al. 2001; Roberts and Gallimore 2005) is presented in Table 1.

It thus seems the best results are obtained when evaluating ECG, EEG, alterations in the respiratory rate, and GSR. It is generally accepted that sweating is the firstmost response to cybersickness, and the autonomic response is mostly secondary, although present (Gavgani et al. 2017). However, ECG still presents several advantages: It is much more simple to capture, and it is possible to obtain the same parameters from simpler methods, such as Photoplethysmography (PPG) (Garcia-Agundez, Dutz, and Goebel 2017)

The aim of this contribution is to analyze the feasibility of detecting cybersickness with a simple 2 lead ECG and thus potentially with other solutions such as Smartphoneor wearable PPG, by finding possible correlations between SSQ scores and relevant ECG parameters.

II. METHODS

Participants

A total of n=13 healthy users (2 females), median age 22, participated in our evaluation.

Materials

An Oculus Rift Developer Kit 2 was used connected to a pc with an Intel Core i7 6700k CPU, Nvidia GTX 980

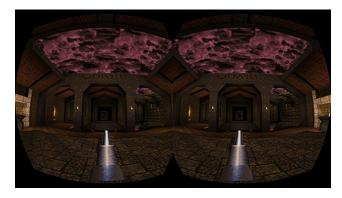


Figure 1: Illustration of the Virtual Environment (phoboslab.org)

graphics card and 16 GB of RAM, with which we ensured the maximum framerate (75 fps) and resolution (960x1080 per eye) could be achieved. Electrocardiography was performed using a 2-lead USBAmp Biosignal Amplifier by g.Tec (www.gtec.at). The signal was captured on Matlab (www.mathworks.com), sampled at 256 Hz and passed through a 10th order Butterworth filter to remove the effect of movement and sweating. A simple maxima detection algorithm was then applied to detect N-N intervals. N-N intervals were then normalized using a baseline taken at rest during two minutes prior to the evaluation start. Finally the average value of all N-N intervals (AVGNN) and the standard deviation of all N-N intervals (STDNN) was measured.

Procedure

Users were asked to play a VR shooter (Quakespasm Rift, a VR modification of Quake compatible with the Oculus Rift Developer Kit 2, available at www.phoboslab.org, see Figure 1) for at least 15 minutes, or until the user decided to stop, while sitting. In the game, they had to navigate through three levels and shoot at enemies. No instructions were provided regarding navigation, to make it as natural as possible, regarding the first person shooter scenario. This duration should suffice according to previous research (DiZio and Lackner 1997; DiZio and Lackner 2000; Lampton et al. 1994; So and Lo 1999). The shooter was chosen given the increased amount of rapid lateral movements and combined head rotations along several axes, which makes the event of cybersickness more likely (Bonato, Bubka, and Palmisano 2009). The ECG electrodes were attached before users filled a pre-immersion SSQ. This was done purposely to capture the baseline data. Users were asked to report the evaluators in case signs of discomfort started showing up. In case this happened, the evaluation was interrupted. After the evaluation users filled a post-immersion SSQ as well.

Ethical approval was obtained from the Ethics Committee of the TU Darmstadt. Prior to taking part in the evaluation, users were asked to fill an informed consent.

III. RESULTS & DISCUSSION

The goal of this work was to analyze if ECG data from users who experience cybersickness during a VR game differs from users who do not. This effect was analyzed by performing a statistical analysis of normalized and non-normalized ECG data, extracting the average duration of N-N intervals (AVGNN) and standard deviation of N-N intervals (STDNN), and comparing these values to the resulting scores of the SSQ in both global and specific areas (nausea, oculomotor, disorientation).

Observations during the evaluation: Interruptions and SSQ Scores

Four out of 13 users (Users 5 to 8) had to interrupt the experiment prematurely due to severe cybersickness. The time of their interruptions is presented on Table 3. About half of users (6 out of 13) reported significant symptoms of cybersickness. Across all users, the highest symptom increases were reported for eyestrain, sweating, nausea and vertigo. There was no report on fatigue, and only minor reports on headache and blurred vision occurred. These observations are based on the results presented in Table 2.

Effect of Cybersickness on ECG data and data quality

Complete results for SSQ scores and ECG data are presented in Table 3. In contrast to other studies, where tachycardia was detected in users (Gavgani et al. 2017), users in our evaluation who suffered more cybersickness presented lower heart rates, with heart rates as low as 54 beats per minute at some points. Due to sweating and extreme movement artifacts, the ECGs of users 3 and 13 had to be discarded.

SSQ Scores and ECG Data

On Average, the experiment increased the SSQ total score by 8 points. This increase was greater in the nausea symptom group, but significant for all three groups. The normalized AVGNN and STDNN remained mostly the same, but it is easy to appreciate greater changes in users who experienced severe cybersickness and had to abandon (Users 5 to 8). This phenomenon is further studied in section 3.5. These observations are based on the results presented in Table 3.

Linear regression between SSQ scores and ECG data

In general, it seems there is a higher correlation between STDNN and SSQ scores than between AVGNN and SSQ. In particular, the regression coefficients between STDNN and oculomotor (r=0.47, presented in Figure 2) or disorientation (r=0.38) are higher. The oculomotor value is slightly better

than the best one obtained so far (r=.43) on (Kim et al. 2005) for the heart period, and is of particular interest, considering previous works also report higher correlations between oculomotor symtpoms and HR, even though disorientation symptoms are believed to be predominant in Cybersickness (Stanney, Kennedy, and Drexler 1997). These results are similar for both normalized and non-normalized ECG data. All linear regression coefficients are presented in Table 4.

T-tests between SSQ scores and ECG data

Across all users, differences between pre- and post SSQ scores are statistically significant in all test sections (nausea, oculomotor, disorientation and total score, p<0.03). Differences in ECG between users who stop prematurely and those who did not are statistically significant as well (p<0.05) except for normalized AVGNN values. This suggests indeed STDNN is a better parameter to evaluate the possibility of cybersickness occurring. In particular, the difference in total SSQ score between users who interrupted the evaluation and those who did not is very significant (p<0.0001). Results of all t-tests are presented in Table 5.

IV. CONCLUSION

The aim of this study was to provide insight into whether it is possible to detect Cybersickness by focusing on changes in AVGNN and STDNN in comparison with SSQ scores using a simple 2-lead ECG. We found significant differences in AVGNN and STDNN of users who report severe cybersickness in the SSQ, as well as a good correlation between Oculomotor SSQ scores and STDNN. We can thus conclude that there is a correlation between ECG parameters and certain SSQ scores.

Since our contribution provides additional evidence on the possible use of ECG to detect cybersickness, we conclude using simple devices to measure biosignals instead, and not in addition to, a pre- and post-immersion SSQ questionnaire may be indeed feasible. We acknowledge that GSR is more reliable, albeit requiring specific measurement hardware. On the other hand, AVGNN and STDNN are simpler to measure, since a simple smartphone using PPG can be used for this purpose. Although it is also possible to measure GSR using a smartphone, it always requires the use of an external device, while PPG is feasible by only using the smartphone itself (Garcia-Agundez, Dutz, and Goebel 2017).

In the future, we plan to evaluate the collected vital data and develop a machine learning algorithm to evaluate the possibility of automatically detecting the onset of cybersickness, and whether there is a consistent correlation between in-game events (such as the number of collisions on-screen, or the speed and acceleration of the user) and the user's physiological response. In addition, we will analyze whether PPG data correlate with ECG data during cybersickness as we demonstrated previously at rest (Garcia-Agundez, Dutz, and Goebel 2017) with the ultimate goal of detecting cybersickness with a simple smartphone.

Acknowledgements

All devices employed during this study were acquired with funds from the Hochschulpakt 2020 program of the German Federal Ministry of Education and Research (BMBF). This project (HA project no. 480/15-22) is funded in the framework of Hessen ModellProjekte, financed with funds of LOEWE – Landes-Offensive zur Entwicklung Wissenschaftlich-ökonomischer Exzellenz, Förderlinie 3: KMU-Verbundvorhaben (State Offensive for the Development of Scientific and Economic Excellence).

Author disclosure statement

Authors report no conflict of interest.

REFERENCES

- [Bonato, Bubka, and Palmisano 2009] Bonato, F.; Bubka, A.; and Palmisano, S. 2009. Combined pitch and roll and cybersickness in a virtual environment. *Aviation, space, and environmental medicine* 80(11):941–945.
- [Bouchard, Robillard, and Renaud 2007] Bouchard, S.; Robillard, G.; and Renaud, P. 2007. Revising the factor structure of the simulator sickness questionnaire. *Annual review of cybertherapy and telemedicine* 5(Summer):128–137.
- [Bruck and Watters 2011] Bruck, S., and Watters, P. A. 2011. The factor structure of cybersickness. *Displays* 32(4):153–158.
- [Chardonnet, Mirzaei, and Merienne 2017] Chardonnet, J.-R.; Mirzaei, M. A.; and Merienne, F. 2017. Features of the postural sway signal as indicators to estimate and predict visually induced motion sickness in virtual reality. *International Journal of Human–Computer Interaction* 33(10):771–785.
- [Dennison, Wisti, and D'Zmura 2016] Dennison, M. S.; Wisti, A. Z.; and D'Zmura, M. 2016. Use of physiological signals to predict cybersickness. *Displays* 44:42–52.
- [DiZio and Lackner 1997] DiZio, P., and Lackner, J. R. 1997. Circumventing side effects of immersive virtual environments. In *HCI* (2), 893–896.
- [DiZio and Lackner 2000] DiZio, P., and Lackner, J. R. 2000. Motion sickness side effects and aftereffects of immersive virtual environments created with helmetmounted visual displays. Technical report, BRANDEIS UNIV WALTHAM MA ASHTON GRAYBIEL SPA-TIAL ORIENTATION LAB.
- [Garcia-Agundez, Dutz, and Goebel 2017] Garcia-Agundez, A.; Dutz, T.; and Goebel, S. 2017. Adapting smartphone-based photoplethysmograpy to suboptimal scenarios. *Physiological measurement* 38(2):219.
- [Gavgani et al. 2017] Gavgani, A. M.; Nesbitt, K. V.; Blackmore, K. L.; and Nalivaiko, E. 2017. Profiling subjective symptoms and autonomic changes associated with cybersickness. *Autonomic Neuroscience* 203:41–50.

- [Gavgani, Hodgson, and Nalivaiko 2017] Gavgani, A. M.; Hodgson, D. M.; and Nalivaiko, E. 2017. Effects of visual flow direction on signs and symptoms of cybersickness. *PloS one* 12(8):e0182790.
- [Johnson 2005] Johnson, D. M. 2005. Introduction to and review of simulator sickness research. Technical report, ARMY RESEARCH INST FIELD UNIT FORT RUCKER AL.
- [Kennedy and Lilienthal 1995] Kennedy, R. S., and Lilienthal, M. G. 1995. Implications of balance disturbances following exposure to virtual reality systems. In *Virtual Reality Annual International Symposium*, 1995. Proceedings., 35–39. IEEE.
- [Kennedy et al. 1993] Kennedy, R. S.; Lane, N. E.; Berbaum, K. S.; and Lilienthal, M. G. 1993. Simulator sickness questionnaire: An enhanced method for quantifying simulator sickness. *The international journal of aviation psychology* 3(3):203–220.
- [Kennedy, Drexler, and Kennedy 2010] Kennedy, R. S.; Drexler, J.; and Kennedy, R. C. 2010. Research in visually induced motion sickness. *Applied ergonomics* 41(4):494–503.
- [Kim et al. 2005] Kim, Y. Y.; Kim, H. J.; Kim, E. N.; Ko, H. D.; and Kim, H. T. 2005. Characteristic changes in the physiological components of cybersickness. *Psychophysiology* 42(5):616–625.
- [Kiryu and Iijima 2014] Kiryu, T., and Iijima, A. 2014. A multi-timescale autonomic regulation model for interpreting visually induced motion sickness. In Consumer Electronics (GCCE), 2014 IEEE 3rd Global Conference on, 254–257. IEEE.
- [Lampton et al. 1994] Lampton, D. R.; Kolasinski, E. M.; Knerr, B. W.; Bliss, J. P.; Bailey, J. H.; and Witmer, B. G. 1994. Side effects and aftereffects of immersion in virtual environments. In *Proceedings of the Human Factors and Ergonomics Society Annual Meeting*, volume 38, 1154–1157. SAGE Publications Sage CA: Los Angeles, CA.
- [Malińska et al. 2015] Malińska, M.; Zużewicz, K.; Bugajska, J.; and Grabowski, A. 2015. Heart rate variability (hrv) during virtual reality immersion. *International Journal of Occupational Safety and Ergonomics* 21(1):47–54.
- [McCauley and Sharkey 1992] McCauley, M. E., and Sharkey, T. J. 1992. Cybersickness: Perception of self-motion in virtual environments. *Presence: Teleoperators & Virtual Environments* 1(3):311–318.
- [Nakagawa 2015] Nakagawa, C. 2015. Toward the detection of the onset of virtual reality sickness by autonomic indices. In *Consumer Electronics (GCCE)*, 2015 IEEE 4th Global Conference on, 662–663. IEEE.
- [Nam et al. 2001] Nam, Y.; Kim, Y.; Kim, H.; Ko, H.; and Park, K. 2001. Automatic detection of nausea using biosignals during immersion in a virtual reality environment. In *Engineering in Medicine and Biology Society, 2001. Proceedings of the 23rd Annual International Conference of the IEEE*, volume 2, 2013–2015. IEEE.

- [Palmisano, Mursic, and Kim 2017] Palmisano, S.; Mursic, R.; and Kim, J. 2017. Vection and cybersickness generated by head-and-display motion in the oculus rift. *Displays* 46:1–8.
- [Regan 1995] Regan, E. 1995. Some evidence of adaptation to immersion in virtual reality. *Displays* 16(3):135–139.
- [Rieder, Kristensen, and Pinho 2011] Rieder, R.; Kristensen, C. H.; and Pinho, M. S. 2011. Identifying relationships between physiological measures and evaluation metrics for 3d interaction techniques. In *IFIP Conference on Human-Computer Interaction*, 662–679. Springer.
- [Roberts and Gallimore 2005] Roberts, W. K., and Gallimore, J. J. 2005. A physiological model of cybersickness during virtual environment interaction. In *Proceedings of the Human Factors and Ergonomics Society Annual Meeting*, volume 49, 2230–2234. SAGE Publications Sage CA: Los Angeles, CA.
- [So and Lo 1999] So, R. H., and Lo, W. 1999. Cybersickness: an experimental study to isolate the effects of rotational scene oscillations. In *Virtual Reality*, 1999. Proceedings., IEEE, 237–241. IEEE.
- [Stanney, Kennedy, and Drexler 1997] Stanney, K. M.; Kennedy, R. S.; and Drexler, J. M. 1997. Cybersickness is not simulator sickness. In *Proceedings of the Human Factors and Ergonomics Society annual meeting*, volume 41, 1138–1142. SAGE Publications Sage CA: Los Angeles, CA.
- [Wilson 1996] Wilson, J. R. 1996. Effects of participating in virtual environments review of current knowledge. *Safety science* 23(1):39–51.
- [Young, Adelstein, and Ellis 2007] Young, S. D.; Adelstein, B. D.; and Ellis, S. R. 2007. Demand characteristics in assessing motion sickness in a virtual environment: Or does taking a motion sickness questionnaire make you sick? *IEEE Transactions on Visualization & Computer Graphics* (3):422–428.



Augusto Garcia-Agundez (augusto.garcia@kom.tu-darmstadt.de) is a PhD student in the Serious Games group of the TU Darmstadt. He specializes in Games for Health and Rehabilitation, particularly the measurement of the effect of Serious Games.



Robert Konrad (Robert.konrad@kom.tu-darmstadt.de) is a PhD student in the Serious Games group of the TU Darmstadt, and creator and developer of Kha (kha.tech). His main research topic is GPU Programming and Cross-Platform development.



Dr. Christian Reuter (christian.reuter@kom.tu-darmstadt.de) is a game developer at Deck13 GmbH. Previously, he wrote his PhD in the Serious Games research group at the TU Darmstadt, working on topics such as VR, Multiplayer Design Patterns, Automated Balancing Analysis and Deadlock Detection and Rapid Development.



Dr. Stefan Göbel (stefan.goebel@kom.tu-darmstadt.de) is an assistant professor and academic councilor, lecturer and head of the Serious Games group at the Multimedia Communications Lab of the TU Darmstadt. His research topics cover standardization, authoring, personalized and adaptive games and effect measurement of serious games. He also provides lectures and practical courses in Serious Games and Game Technologies.



Polona Caserman (polona.caserman@kom.tu-darmstadt.de) is a PhD student in the Serious Games group of the TU Darmstadt. Her research areas are VR/AR body tracking, particularly accuracy issues and latency reduction.

Table 1: Results of studies vinculating cybersickness and biosignals

Biosignal	Correlated with VR im- mersion (Nam et al. 2001)	Correlated with cyber- sickness (Kim et al. 2005)	Correlated with cybersickness (Roberts and Gallimore 2005)	Correlated with cy- bersickness (Gavgani, Hodgson, and Nalivaiko 2017)	Correlated with cybersickness (Gavgani et al. 2017)	Correlated with SSQ Scores (Den- nison, Wisti, and D'Zmura 2016)
Heart Rate	_	_	_	No	No	r = 0.09
Heart Rate Variability	Yes $(p < 0.001)$	r = 0.43	_	Yes $(p < 0.05)$	_	r = 0.11
Pulse Volume	Yes $(p < 0.001)$	_	_	_	_	_
Respiratory Rate	No $(p = 0.15)$	r = 0.39	_	_	No	r = -0.31
Respiratory Arrythmia	Yes (p < 0.001)	r = 0.33	_	_	_	_
Skin Tempera- ture	Yes $(p < 0.001)$	_	_	_	_	_
GSR Fore- head:	_	_	_	Yes $(p < 0.01)$	Yes $(p < 0.001)$	r = 0.20
GSR Finger:	Yes $(p < 0.001)$	_	_	Yes $(p < 0.01)$	Yes $(p < 0.001)$	_
Gastric Rhythm	Yes $(p < 0.001)$	r = 0.36	Yes $(p = 0.04)$	_		_
Eyeblink Rate	Yes $(p = 0.01)$	r = 0.30	_	_	_	r = 0.50
EEG Delta F3	Yes $(p < 0.001)$	r = 0.30	_	_	_	_
EEG Delta T3	Yes $(p < 0.001)$	r = 0.31	_	_	_	_
EEG Delta O1	_	r = 0.34	_	_	_	_
EEG Beta F3	Yes $(p = 0.02)$	r = -0.38	_	_	_	_
EEG Beta T3	Yes $(p < 0.001)$	r = -0.32	_	_	_	_
EEG Beta T4	_	r = -0.30	_	_	_	_
EEG Beta O1	_	r = -0.39	_	_	_	_
EEG Gamma F3	_	r = -0.32	_	_	_	_
EEG Theta P3	_	r = 0.39	_	_	_	_

Table 2: Evaluation symptom results

SSQ Score Change
7
-7
2
12
1
2
11
11
2
-2
2
3
7
10
8
2

Table 3: SSQ and ECG results

Массилот	Table 3: SSQ and ECG results MeasuremenTotal Im- SSQ SSQ SSQ ECG ECG ECG ECG						ECG		
method	mersion	SSQ	SSQ	SSQ	SSQ	ECG	ECG	ECG	ECG
memod	Time (s)								
User	Time (s)	Nausea	Oculomote	or Disorienta	ntioTotal	AVGNN	STDNN	AVGNN	STDNN
Osci		rausea	Oculomot	or Disorient	idolotai	msec	msec	(Normal-	(Normal-
						Insec	msee	ized)	ized)
1	919.44	-3.74	-3.74	-3.74	-3.74	696.049	52.268	1.014	0.076
2	1426.7	3.74	0	0	1.25	760.181	65.445	0.914	0.079
3	1581.38	3.74	11.22	11.22	8.73	_	_	_	_
4	1237.91	14.96	-3.74	11.22	7.48	680.01	65.029	1.023	0.098
5 (Aban-	687.05	41.14	11.22	37.4	29.92	703.75	66.635	1.052	0.1
doned)									
6 (Aban-	1017.1	22.44	11.22	7.48	13.71	768.112	64.467	1.123	0.094
doned)									
7 (Aban-	685.83	14.96	18.7	22.44	18.7	711.095	94.394	1.124	0.149
doned)									
8 (Aban-	1055.06	29.92	11.22	11.22	17.45	895.182	119.756	0.976	0.131
doned)									
9	1152.72	22.44	7.48	3.74	11.22	596.989	48.38	0.914	0.074
10	1064.49	22.44	3.74	3.74	9.97	651.544	47.749	1.025	0.075
11	1102.48	-7.48	7.48	3.74	1.25	646.893	70.302	1.063	0.116
12	1200.24	0	-3.74	0	-1.25	672.112	75.837	1.13	0.128
13	1154.04	-7.48	0	0	-2.49		_	_	_
Average	1098.8	11.49	5.08	7.48	8.01	707.447	70.024	1.033	0.102

Table 4: Linear Regression Results

Linear Regression	AVGNN	STDNN	AVGNN (Normal-	STDNN (Normal-
Coefficients			ized)	ized)
Nausea SSQ	r = 0.29	r = 0.18	r = -0.09	r = -0.01
Oculomotor SSQ	r = 0.27	r = 0.46	r = 0.24	r = 0.47
Disorientation SSQ	r = 0.15	r = 0.31	r = 0.27	r = 0.38
Average SSQ	r = 0.27	r = 0.33	r = 0.12	r = 0.26

Table 5: T-test Results

	Significant
Nausea, pre- vs. Post scores	Yes (p=0.01)
Oculomotor, pre- vs. Post scores	Yes (p=0.02)
Disorientation, pre- vs. Post scores	Yes (p=0.03)
Total, pre- vs. Post scores	Yes (p<0.01)
Total SSQ score, stops prematurely vs doesn't	Yes (p<0.001)
AVGNN, stops prematurely vs doesn't	Yes (p=0.04)
STDNN, stops prematurely vs doesn't	Yes (p=0.05)
AVGNN, same data, normalized	No (p=0.07)
STDNN, same data, normalized	Yes (p=0.02)

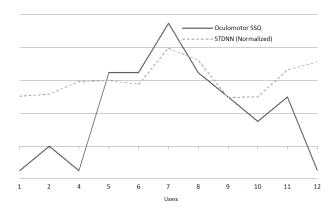


Figure 2: Comparison between Oculomotor SSQ scores and normalized STDNN (multiplied by 100 for scale), excluding users without ECG Data