Physiological measures and self-report to evaluate neutral virtual reality worlds

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ABSTRACT

Using virtual reality technology for exposure therapy to treat patients with anxiety disorders is attracting considerable research attention. The ability to monitor patient anxiety level helps therapists to set appropriate anxiety arousing situations. Physiological measures have been put forward as objective indicators of anxiety levels. Because of individual variation, they need a baseline recording which is often conducted in neutral virtual worlds which do not include phobic stressors. Still because of the novelty of the virtual worlds, reports in the literature suggest that individuals already show some level of arousal when placed in these worlds. This paper presents two studies which look at the effect two different neutral virtual worlds can have on individuals. Findings suggest that a neutral world does not have to result in an increased level of arousal.

Keywords
Virtual Reality, Exposure Therapy, Physiological Measurements, Neutral Worlds, Fear of Flying

1. INTRODUCTION

Virtual Reality Exposure Therapy (VRET) is receiving considerable research attention for treatment of patients suffering from anxiety disorder, such as claustrophobia, fear of driving, acrophobia, spider phobia, social phobia, panic disorder with agoraphobia, post traumatic stress disorder, and fear of flying. VRET is based on the ideas of gradual exposure in vivo, considered the gold standard for treatment of phobias. Recent meta-studies (Gregg & Tarrier, 2007; Parsons & Rizzo, 2008; Powers & Emmelkamp, 2008) show that
exposure in VR is as effective as exposure in vivo. An important element of the therapy is that the exposure is done gradually to more anxiety arousing situations. Therapists are, therefore, continuously monitoring the anxiety level of a patient. This can be done using Subjective rating of Anxiety (SUD), behavioural observations or physiological measures. The latter has the advantage of being more objective, and can be used directly by a computer to assist a therapist in a multi-patient VRET setting (Paping, Brinkman, & van der Mast, 2010). Physiological measures need however a baseline measurement because of individual variation. One often used procedure is to obtain a physiological baseline recording when the patient is placed in a neutral VR world, i.e. a VR world which should not include phobia related stressors. Even if this world has no phobia related stressor, it is not clear whether experience of being placed in a Virtual Environment (VE) causes some level of anxiety. Some authors (Wiederhold & Wiederhold, 2005) have suggested that the majority of non-phobic individuals do get some level of arousal when placed in a VE. For example Jang et al (Jang, et al., 2002) report a study with non-phobic individuals and observed that participants were initially aroused in the VR exposure, but returned to a normal base line after approximately 7 minutes. In another study, Wiederhold et al. (Wiederhold, Davis, & Wiederhold, 1998) also report that non-phobics, when placed in a VE, initially show some level of anxiety. They argued that the VE is a new and novel stimulus and therefore causing this effect. Extending on this line of reasoning, this paper explores whether the design of the neutral world can also contribute to this effect. Or in other words, would it be possible to design a truly neutral world. As reported in this paper, we were confronted with this question after results of our first study suggested that both phobic and non-phobic participants showed higher heart rates during exposure in a neutral virtual world than both in the VE with phobic stressors and in the recovery phase after the VR exposure. Furthermore, both phobics and non-phobic individuals experienced moderate to severe nausea in the neutral VR condition. This called into question the neutrality of the neutral virtual world and led our research into the creation of neutral virtual worlds.

The paper starts with briefly discussing key concepts such as VR systems, presence and problems experienced by patients. After this, the first study is presented in which both non-phobic and phobic individuals are placed in a neutral VR world, a virtual airplane, and a recovery phase. The second study starts with a discussion of the design of a new neutral VR world. This virtual world aims to be an almost identical representation of the room the individual is sitting in. Results are presented from data collected in four conditions: the real world room, the new neutral VR world, virtual airplane, and recovery phase. The paper concludes by discussing the findings which suggests that it might be possible to design a truly neutral world. Also no support was found for a possible transfer of habitation from the physical room to the neutral virtual room.

2. BACKGROUND

The sense of being a part of the VE even when a person is physically situated in a totally different real world is considered a key element of VRET. This concept of presence is related to four components: technological devices; user-computer interactions; main task and the user (IJsselsteijn, de Ridder, Freeman, & Avons, 2000;
Witmer, Jerome, & Singer, 2005). In the application field of VRET, the main technical devices used are a head mounted display (HMD) and a computer automatic environment (CAVE). The CAVE has a relatively higher immersion level with stereoscopic images on four to six sides around the user while the HMD has only one stereoscopic image in front of the user. In a study on the effects of VRET in patients with acrophobia using CAVE and HMD, it is reported that VRET was superior to no-treatment on anxiety, behavioral avoidance and attitudes towards heights. Although the therapy given in the CAVE resulted in higher level of presence than the therapy given through HMD, no differences in effect were found between them and the results remained stable during the following six months (Krijn, Emmelkamp, Biemond, et al., 2004). This therefore seems to suggest that only a certain level of presence is needed for treatment to be effective. Even with devices as a HMD or a CAVE patients can still experience low level of presence causing them to drop out of the treatment (Krijn, Emmelkamp, Olafsson, & Biemond, 2004). This underlines that presence is also determined by individual factors (Ling, Nefs, Brinkman, Heynderickx, & Qu, 2010) such as vision ability, cognitive processing ability (T. W. Schubert, 2009) of the VE, and personality (Wallach, 2010). It has also been suggested that imaginative power influences presence (Huang, Himle, & Alessi, 2000; Regenbrecht, Schubert, & Friedmann, 1998). Being able to visualize more vividly could intensify the experience of the VE. Cybersickness is another potential intervening factor. Cybersickness is a form of motion sickness that occurs as a result of exposure into VE and can range from a slight headache to an emetic response (Stanney, Mourant, & Kennedy, 1998). Although physiological measurements can be used for determining anxiety during VRET, the side effect of cybersickness can also arouse physiological changes in people (Min, Chung, Min, & Sakamoto, 2004). Both cybersickness and presence therefore seem important factors that might explain, besides the initially suggested habitation, physiological effects in neutral virtual worlds.

3. STUDY 1

3.1 Method

The first study was initially set out to study physiological response of both phobic and non-phobic individuals in a VE with phobic stressors. Both groups were exposed to three conditions: a neutral virtual world, a virtual flight, and a recovery phase. Both the effects for two groups and conditions on physiological recordings and self-reported anxiety were analyzed.

3.1.1 VR system

The VRET Delft 2007 system is described in detail elsewhere (Aslan, 2007; Brinkman, van der Mast, Sandino, Gunawan, & Emmelkamp, 2010; Gunawan, Mast, Neerincx, Emmelkamp, & Krijn, 2004; Schuemie, 2003). Briefly, the HMD used was the stereoscopic Cybermind Visette Pro with a resolution of 640x480 per display and a 60 Hz refresh rate. An Ascension Flock of Birds was used as the tracking tool. Two personal computers (PC) were used in the system, the therapist computer where the therapist controls the therapy session and a
patient PC which gets input from the HMD and therapist computer. Both the neutral virtual world and the flight world were created with WorldUp R4 by Sense8.

Figure 1: Left, the neutral courtyard, right virtual flight world.

### 3.1.2 Participants

Participants for study 1 were aviophobics that applied for therapy at the VALK foundation, and non-paid volunteers without fear of flying who acted as a control sample. The VALK foundation is a mental health clinic that specializes in aviation related anxiety. During the recruitment period 46 phobic clients who applied for treatment received written information regarding the VR study at their home address two weeks before their first visit. Out of this group, 40 phobics were willing to participate. One client was excluded because of the use of cardioactive medication (β-blockers). This left 39 phobic clients (15 men) with an average age of 44.5 ($SD = 12.4$), who fulfilled the DSM-IV criteria for specific situational phobia furnishing usable data. In the same period 22 non-paid volunteers without fear of flying and an average age of 48.3 ($SD = 11.4$) successfully completed a part of the same protocol. Volunteers were recruited through the social network of the research institution’s staff. One of them received a positive diagnosis for aviophobia during the intake and was excluded. Another control subject’s questionnaire data rendered unusable, her physiological data was included for analyses. The 21 non-phobic healthy subjects had flown at least several times; most of them had flown within 18 months of the experiment. None of the control subjects was ever treated for fear of flying. Before start of the experiment, informed consent was obtained from all participants. The research protocol has been approved by the local medical ethics committee.

### 3.1.3 Measures

For the physiological recordings, the three target variables were Heart Rate (HR), Pre Ejection Period (PEP) and Respiratory Sinus Arrhythmia (RSA). PEP is considered a measure of (activating) sympathetic cardiac control (Sherwood, et al., 1990) whereas RSA is a measure of (calming) parasympathetic control (Berntson, et al., 1994). Scoring of these variables from thorax impedance and the ECG is described in detail elsewhere (Goedhart, Kupper, Willemsen, Boomsma, & de Geus, 2006; Goedhart, Van der Sluis, Houtveen, Willemsen, & De Geus, 2007). Briefly, from the ECG (sampling rate 1000 Hz) the HR was obtained from the time
between two adjacent R waves. PEP was defined from the ECG and ICG as the time interval from the Q-wave onset, the onset of the electromechanical systole, to the B-point (from the ICG), which signals opening of the aortic valves (Sherwood, et al., 1990; Willemsen, DeGeus, Klaver, VanDoornen, & Carroll, 1996). RSA was obtained from the ECG and respiration signals by subtracting the shortest IBI during HR acceleration in the inspirational phase from the longest IBI during deceleration in the expirational phase (i.e. the peak-through method) (Grossman, 1990). When no phase-related acceleration or deceleration was found, the breath was assigned a RSA score of zero. Automatic scoring of PEP and RSA was checked by visual inspection of the impedance and respiratory signal from the entire recording. Our focus on cardiac parameters reflects two major considerations: measurements needed to be as non-invasive as possible and they needed to respond to changes in psychological state over a time scale of a few minutes. The PEP and RSA measures are uniquely qualified to meet both demands (Goedhart, et al., 2006; Willemsen, et al., 1996). Using a visual display of the output of an inbuilt vertical accelerometer, we identified artefact free periods in each condition that lasted at least 5 minutes each.

All questionnaires were administered in the Dutch language, they were:

- The Subjective Units of Discomfort (SUD) scale was used to examine to what extent participants were feeling anxious at several moments. They had to indicate their perceived anxiety on a scale from 1 (“totally relaxed”) to 10 (“extremely anxious”) (Wolpe, 1973).

- The Visual Analogue Flight Anxiety Scale (VAFAS) was used to examine to what extent participants were anxious about flying. The one-tailed scale ranges from 0 (No flight anxiety) to 10 (Terrified or extreme flight anxiety) (Van Gerwen, Spinhoven, Van Dyck, & Diekstra, 1999).

- The Igroup Presence Questionnaire (IPQ) was used to measure the feeling of being in the VE. Each of the 14 items is rated on a 7-point Likert-type scale, ranging from -3 (totally disagree) to 3 (totally agree). The IPQ scale consists of 3 subscales: spatial presence, involvement and realness. The psychometric properties proved to be good to excellent (T. Schubert, Friedmann, & Regenbrecht, 2001). In the present study only the total score on the IPQ was used. The internal consistency in the present study was good, Cronbach’s Alpha .95

3.1.4 Procedure

All measurements took place at the VALK facility. Upon arrival participants were informed about the procedure. For the aviophobics it was emphasized that participating was voluntary and neither participation nor refusal to participate impacted on the quality of treatment. After informed consent was given, six electrodes were attached and connected to the Vrije Universiteit Ambulatory Monitoring System (VU AMS) which records the thorax impedance and the ECG in freely moving individuals (Goedhart, et al., 2006; Goedhart, et al., 2007; Houtven, Groot, & de Geus, 2006; Riese, et al., 2003; Willemsen, et al., 1996).
Participants were then seated upright in a normal chair and partook in three experimental conditions, always in the same fixed order. Participants first received a 7 minutes VR exposure in a neutral VE after which they were asked to fill out the IPQ. The neutral VE (Schuemie, 2003) consisted of a courtyard in which participants moved around under therapist control, i.e. locomotion is not controlled by the participants. The locomotion was standardized and automated. Participants completed two rounds along the outer perimeter of the courtyard (figure 1, left). This condition was followed by 7 minutes VR flight simulation in a real airplane seat. Participants were seated upright and followed a standardized program consisting of taxi-out, take-off, a short cruise flight, descent, approach and landing. Subsequently, participants were given 7 minutes of recovery time while seated in the airplane seat. Subjective units of distress (SUD) were measured at four discrete moments: before the start of the experiment, directly after both VR presentations and at the end of the recovery period. The VAFAS was administered before start of the experiment.

### 3.2 Results

Comparison of phobic and non-phobic control participants on sociodemographic characteristics and the VAFAS scale were performed with one-way ANOVA. Table 1 shows the main characteristics for the group of phobic participants and the control group.

**Table 1: Number of participants, gender, age, BMI and VAFAS score in study 1**

<table>
<thead>
<tr>
<th></th>
<th>Phobics</th>
<th>Non-phobics</th>
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</thead>
<tbody>
<tr>
<td></td>
<td>Mean (SD or %)</td>
<td>Mean (SD or %)</td>
</tr>
<tr>
<td>Number of participants</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Total</td>
<td>39</td>
<td>21</td>
</tr>
<tr>
<td>Men</td>
<td>15 (38%)</td>
<td>11 (52%)</td>
</tr>
<tr>
<td>Women</td>
<td>24 (62%)</td>
<td>10 (48%)</td>
</tr>
<tr>
<td>Age (years)</td>
<td>44.5 (12.4)</td>
<td>48.3 (11.4)</td>
</tr>
<tr>
<td>BMI</td>
<td>24.6 (3.8)</td>
<td>23.5 (2.4)</td>
</tr>
<tr>
<td>VAFAS</td>
<td>8.0 (1.4)*</td>
<td>0.6 (0.7)</td>
</tr>
</tbody>
</table>

* Phobics differ from non-phobics at $p < .001$

An ANOVA was conducted on the SUD scores collected in three conditions (neutral VR world, virtual flight, and recovery) from the two participant groups (phobic, and control). Significant condition ($F(1.73,91.54) = 3.81, p = .031$) and group ($F(1,53) = 21.68, p < .001$) effects were found for self-reported distress. Phobics had higher levels throughout, while on average participants reported less fear during recovery compared to the
virtual flight. Follow-up analyses for both groups separately showed significant differences in reported anxiety between the recovery condition and both the neutral VR world ($t(1, 37) = 2.51, p = .017$) and the virtual flight ($t(1,33) = 3.09, p = .004$) for the flight phobics, while no significant differences between conditions were seen for the control group (Figure 2).

Figure 2: SUD scores for phobic and control participants.

An ANOVA with the same independent variable was also conducted on the physiological data. Of the physiological variables, RSA had to be log (ln) transformed to obtain normal distributions. For HR a significant main effect was found for condition ($F(1.78, 101.62) = 16.94, p < .001$). Both control participants and phobic participants had higher heart rates during the neutral VR world than in any other condition (Figure 3). In contrast to the main effect of condition for HR, no significant effects of condition were found in RSA and PEP data. In fact, there was no significant main or interaction effect in RSA at all. A significant main group effect did emerge in PEP data, phobic participants had significantly shorter PEP values than control subjects, indicating higher cardiac sympathetic control ($F(1,58) = 5.83, p = .019$).

A one-way ANOVA was conducted for the IPQ. Non-phobic subjects scored significantly higher on the total IPQ scale ($F(1,57)=10.42, p=0.002$) including its subscales Spatial Presence (SP: $F(1,57)=11.45, p < 0.05$) Involvement (INV: $F(1,57)=5.24, p < 0.05$) and Realism (Real: $F(1,57)=4.08, p < 0.05$). IPQ scores were
relatively high compared with other studies\(^1\). No significant correlations between IPQ scores and SUD scores were found.

Almost all participants complained either during or directly after the neutral VE exposure about dizziness and nausea. This was corroborated by an elevated HR during this supposedly non-provocative neutral condition. No difference in anxiety between the neutral VR world and the flight world was reported by the flight phobics. All other measures did not differentiate between conditions. This led us to the conclusion that the neutral VR world probably was not truly neutral after all.

Figure 3: Average HR for phobic and non-phobic combined

4. STUDY 2

4.1 Method

Novelty of a new environment or cybersickness might have caused the higher level of arousal in the neutral VR world in study 1. This would suggest that arousal level could be reduced by removing the novelty and therapist controlled locomotion element from the neutral VR world. The aim of the second study was therefore to examine whether a new neutral world would still result in an elevated level of arousal. In addition the study was also set out to study the suggested novelty effect or possible transfer of habituation by changing the order in which physiological recording was collected (the actual room first, or virtual room first).

\(^1\) For comparison data see www.igroup.org
4.1.1 VR system and new neutral world

To make a possible transfer of habitation possible a virtual world was created which was a close replication of the actual room the individual was situated in, causing participants to see the same environment when they would put on or take off the HMD. Participants were seated in front of a television (Figure 4, right) showing a documentary about wildlife. In the new neutral VR world (Figure 4, left) participants were seated in front of the same television set which shows the same documentary. Looking around with or without the HMD would give the same view of the room. The new neutral VR world ran on the same hardware as the VR flight and the old neutral VR world but used different software with exception of the Windows XP operating system. The Vizard Virtual Reality Toolkit, Vizard 3.0 was used to create an executable that provided head tracking and the image for the HMD. The model of the room was created with Autodesk Maya 2008 and textures were edited with Adobe Photoshop CS2. The model consisted of the room in which a table, television set, room dividers and a metal rail were modelled in detail. There were 18 textures made with several different sizes ranging from 2048 x 2048 pixels (the wall closest to the patient) to 64 x 512 pixels (a table leg). The world was displayed with a resolution of 640 x 480 to match the resolution of the HMD used in VR flight. All textures where file textures. The textures were made out of photographs taken from the location where the patient would sit. Every visible face got its own unique texture. No dynamic lights or computer generated shadows were used. Distortion was removed from the images and the colour balance of several images was altered. Some objects were edited out of textures, like the table that was removed from the photograph that formed the texture for the wall behind the television set, which was facing the patient. Shadows belonging to objects that were removed from the scene were edited out like the radio and chair in front of the table. Some shadows had to be drawn in by hand like the shadow of the table on the part of the wall behind the table. A video could be displayed on the television set triggered by a keyboard button press. The video used was ripped from DVD and recompressed with a resolution of 720 x 576 at 25 frames/second. The video format and codec used was VC-1, WMV3 (Windows) and the audio format and codec used was WMA2, 161 (Windows). The video was edited to a duration of 6 minutes and 29 seconds. A DVD with the exact same edited video was made so that the video could be played on the DVD player in the actual room.

Figure 4: Left, new neutral virtual world, Right, picture of the actual room
4.1.2 Participants

44 People participated, 32 students who earned credits by participating and 12 non-paid volunteers recruited by means of word-to-mouth. All participants received an email with information regarding the study before start of the experiment. One participant was excluded because of the use of cardioactive medication. Another two participants were partly excluded from analyses because of equipment failure during physiological recordings; their questionnaire data was included for analysis. Average age of all 43 (16 men) participants was 25.6 (SD = 8.0), the youngest being 18 years old, the oldest 51.

4.1.3 Measures

In addition to the three physiological measurements and questionnaires used in the previous study the following two questionnaires were added:

- The Simulator Sickness Questionnaire (SSQ) was used to examine to what extent participants experienced symptoms associated with simulator sickness caused by the VR exposure. The SSQ consists of a checklist of 27 symptoms, each of which is rated in terms of degree of severity (none, slight, moderate, severe). It is normally administered twice, before and after a VR exposure (Kennedy, 1993). The instrument provides three subscales (Nausea, Oculomotor and Disorientation) and a composite Total Severity Score, which is used in the present study. The instrument’s psychometric properties are good (Johnson, 2005). The internal consistency in the present study was good, Cronbach’s Alpha .78.

- The Vividness of Visual Imagery Questionnaire (VVIQ) was used to examine the ability to form mental pictures (Marks, 1973). The vividness of the image is rated along a 5-point Likert-type scale, from 1 (perfectly clear and as vivid as normal vision) to 5 (no image at all, you only “know” that you are thinking of an object). All items for images obtained are first answered with eyes open, secondly with eyes closed. Note that a low VVI score means vivid imagery and a high score means vague imagery. The psychometric properties proved to be good to excellent (Campos & Perez-Fabello, 2009). In the present study the average score on the VVIQ was used. The internal consistency in the present study was good, Cronbach’s Alpha .95.

4.1.4 Procedure

Participants started with filling out the VAFAS and the SSQ (pre-exposure). After attachment of the electrodes of the VU AMS participants were seated upright in a normal seat. Participants randomized started either with the new neutral VR world, or the neutral real world. Participants were asked to complete the IPQ and SSQ-post-exposure directly after the neutral VR world. These two conditions were followed by 7 minutes VR flight simulation seated in a real airplane chair. Participants were seated upright and followed a standardized program consisting of taxi-out, take-off, a short cruise flight, descent, approach and landing. Subsequently, participants were given 7 minutes of recovery time while seated in the airplane seat. SUD score were recorded at five discrete moments: before the start of the experiment, directly after both neutral worlds,
after the virtual flight and at the end of the recovery period. Before start of the experiment informed consent was obtained from all participants. The research protocol had been approved by the local medical ethics committee.

### 4.2 Results

As was done with study 1, a series of ANOVAs was conducted to study the effect of two independent variables: condition (real world, new neutral VR world, virtual flight, recovery), and group (first real world then new neutral VR world, or first new neutral VR world and then real world).

<table>
<thead>
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<th>Number of participants, gender, age, BMI and VAFAS score in study 2</th>
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<tbody>
<tr>
<td><strong>Mean (SD or %)</strong></td>
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<tr>
<td>Number of participants</td>
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<tr>
<td>• Total</td>
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<tr>
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<tr>
<td>• Women</td>
</tr>
<tr>
<td>Age (years)</td>
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<tr>
<td>BMI</td>
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<tr>
<td>VAFAS</td>
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A significant main effect was found for condition ($F(2.46, 100.9) = 3.29, p =.032$) in the SUD scores. Participants reported lower levels of anxiety during the real world than during any other condition. Follow-up analyses for both groups separately showed a significant difference in reported anxiety between the real world and both the new neutral VR world ($t(1, 20) = -2.32, p =.031$) and the virtual flight ($t(1,20) = -2.35, p =.029$) for the participants who saw the real world first (Figure 5). Interestingly no significant differences between conditions were found when the new neutral VR world was presented first.
The analysis of heart rate found a significant condition by group interaction \((F(2.19, 85.32) = 5.48, p = .005)\), together with a main effect of condition \((F(2.19, 85.32) = 10.12, p < .001)\). Follow-up tests revealed that HR during virtual flight was significantly lower than HR in any other condition (all \(p < .001\)), while the interaction with group was driven by an increase of HR during the real world condition for the participants who saw the real world first (Figure 6).

In contrast to the condition by group interaction for HR, no significant condition by group interactions were found for RSA and PEP. Significant main condition effects did emerge for overall RSA and PEP levels. Participants had significantly longer RSA values during virtual flight compared to all other conditions, indicating higher parasympathetic control during virtual flight \((F(2.72, 106.04) = 9.06, p < .001)\), and significant longer PEP values during virtual flight compared to the new neutral VR world and the recovery condition \((F(3, 37) = 5.12, p = .003)\), indicating less cardiac sympathetic control during virtual flight.
On average participants had a significant decrease in SSQ from pre- to post-presentation measurement ($t(1, 41)=2.65$, $p=.011$). These changes from pre-to post scores on the SSQ were significantly correlated to the SUD values from the Virtual Flight such that decreased simulator sickness was accompanied by a lower anxiety score during the flight condition ($r = -.437$, $p = .003$). SSQ-post scores were significantly correlated with SUD-Flight ($r = .508$, $p = .001$) and SUD-Recovery ($r=.522$, $p<.001$). Participants with lower post presentation simulator sickness scores report lower anxiety during virtual flight and the recovery condition, while participants with higher post presentation SSQ values report more anxiety in both conditions.

No significant correlations between IPQ scores, SUD scores and VVIQ were found. A significant negative correlation was found between IPQ and SSQ-Post ($r = -.325$, $p =.033$) and a significant positive correlation was found between IPQ scores and SSQ pre-post ($r=.391$, $p=.009$). On average participants with a higher presence score had a lower post-presentation simulator sickness score than participants with lower IPQ score, while participants with a higher presence score show a stronger decrease in simulator sickness compared to participants with lower IPQ scores.
5. CROSS COMPARISON

Cross comparison was performed on control participants from study 1 with all participants from study 2. Table 3 shows the main characteristics for both groups. ANOVAs were conducted with condition (Neutral VR world, Flight world, Recovery) and group (participants study 1, participants study 2) as independent variables. Age was added as a covariate.

Table 3: Number of participants, gender, age, BMI and VAFAS score in study 1 and 2

<table>
<thead>
<tr>
<th></th>
<th>Participants Exp.1</th>
<th>Participants Exp.2</th>
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<tbody>
<tr>
<td>Number of participants</td>
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<td></td>
</tr>
<tr>
<td>• Total</td>
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<td>43</td>
</tr>
<tr>
<td>• Men</td>
<td>11 (52%)</td>
<td>16 (37%)</td>
</tr>
<tr>
<td>• women</td>
<td>10 (48%)</td>
<td>27 (63%)</td>
</tr>
<tr>
<td>Age (years)</td>
<td>48.3 (11.4)</td>
<td>25.6 (8.0)*</td>
</tr>
<tr>
<td>BMI</td>
<td>23.5 (2.4)</td>
<td>22.6 (2.8)</td>
</tr>
<tr>
<td>VAFAS</td>
<td>0.6 (0.7)</td>
<td>0.8 (1.2)</td>
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* Participants study 1 differ from Participants study 2 at $p < .001$

A significant group by condition interaction was found for self-reported distress ($F(1.71, 104.1) = 6.76, p = .003$), together with main effects of group ($F(1.61) = 5.10, p = .028$) and condition ($F(1.71,104.1) = 9.19, p < .001$). In figure 7 it can be seen that the control participants in study 1 had higher levels of distress throughout, while in contrast with the participants from study 2, their reported distress was highest during the neutral VR condition. In parallel with self-reported distress a significant group by condition interaction was found for heart rate ($F(4.57,101.21) = 4.57, p = .017$), with a main effect for condition, not for group ($F(1.72,101.21) = 4.21, p = .023$). Control participants in study 1 had significantly higher HR during the neutral VR condition than during the virtual flight and recovery condition, while participants from study 2 showed no significant difference in HR between the neutral and recovery condition.

Analogue with both subjective distress and heart rate a significant group by condition interaction ($F(2,118) = 4.52, p = .013$) showed for the parasympathetic measure RSA, together with a main effect of condition ($F(2,118) = 5.87, p = .004$). Post-hoc analyses revealed that participants in the second study had nearly the same RSA values during the neutral VR world and the recovery condition, while RSA values during virtual flight were significantly longer than RSA values during the neutral VR world ($t(1,40)= 4.55, p<.000$) and
recovery condition \((t(1,40)=2.50, p=.016)\). Differences between conditions did not reach significance for participants from the first study (Figure 8).

Figure 7: SUD scores for participants from both studies

Figure 8: Average RSA for participants from both studies
For the sympathetic measure PEP only a significant group by condition interaction was found \((F(1.57,92.68) = 3.57, p = .042)\). Again participants in the second study had no significant differences between the neutral VR world and the recovery period, with significant longer PEP values during the virtual flight. Participants in study 1 showed no significant difference in PEP between all three conditions.

A one-way ANOVA was conducted for the IPQ. No significant differences between both groups were found.

**6. CONCLUSION AND DISCUSSION**

In the second study all physiological measures differentiated between the flight condition on one side and the VR neutral world and recovery condition on the other side, while no physiological difference was apparent between the VR neutral world and recovery. Even self-reported distress showed no significant differences between conditions when the new VR neutral world was presented first. This seems to refute the idea that a virtual world by definition will generate arousal and anxiety (Wiederhold & Wiederhold, 2005). The second study found only an interaction effect between the condition and the groups in HR. Still, follow-up analyses only found a significant decrease, instead of an increase, in HR when recordings were first collected in actual room and then in neutral VR room. This observation is therefore contra to the idea that novelty of VE would always cause arousal. Also the follow-up analyses did not find a significant difference in HR of participants from the group in which the recording took place in the opposite order (first neutral VR world, second actual room). A significant decrease would have provided support for the hypothesis of transfer of habituation from one environment to another. The lack of interaction effects in the other physiological measures makes this hypothesis again less likely. Thus suggesting that to obtain neutral physiological measurements the VE does not have to be a replication of the actual room the individual is situated in. Cross comparison of control participants of study 1 and all participants from study 2 strengthen our findings. Both subjective measures of anxiety as well as all physiological measures of arousal indicated equal anxiety and arousal in the neutral VR world and the recovery condition in the second study, while participants in study 1 had elevated values for SUD and HR during the supposedly neutral VR condition when compared to the virtual flight and recovery condition. The study also found presence and cybersickness to be negatively related. Although only a certain level of presence is needed for treatment to be effective (Krijn, Emmelkamp, Biemond, et al., 2004), maximizing presence might reduce simulator sickness and thereby minimize drop out. No relationship was found between imaginative power, anxiety and presence. Our data therefore did not corroborate the idea that imaginative power influences presence (Huang, et al., 2000; Regenbrecht, et al., 1998). This might be caused by relatively higher IPQ scores compared to other studies. It is reported that imagination has an important effect on presence when the VR is limited. However, if the VR is vivid enough, the participants do not need to use their imagination to create a convincible virtual environment (Wallach, 2010). The average lower heart rate in the virtual flight condition compared to the neutral and recovery conditions could reflect different types of coping mechanisms. Our data are strongly reminiscent of similar data in dental phobics exposed to a
stressful video showing surgical operations (Bosch, et al., 2001) as well as non-phobics exposed to neutral-and flight-videos (Busscher, Van Gerwen, Spinhoven & De Geus, 2010). Exposure to phobic stimuli is a complex stressor in that it can invoke both flight-flight responses, characterized by increased sympathetic and reciprocal decreased parasympathetic activity, and a passive coping response (freeze), characterized by increased sympathetic activity paired to increased parasympathetic activity. A principal finding in study 1 is that phobics were more anxious during the entire experiment than non-phobics, as expressed in significantly higher SUDs and sympathetic activation (PEP). This contributes to the validity of VR as a useful tool in exposure based therapy.

7. REFERENCES


