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# 1 Attention Orientation to Pleasantness and Depressive 2 Symptomatology Predict Autonomic Reactivity

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## 14 *Abstract*

15  
16 Depression is characterised by attentional bias to emotional information and  
17 dysregulated autonomic reactivity. Despite its relevance to understanding depressive  
18 mechanisms, the association between attentional bias and autonomic reactivity to emotional  
19 information remains poorly characterised. This study compared behavioural and autonomic  
20 responses to emotional images in 32 participants in whom subclinical depressive  
21 symptomatology was quantified using the Beck Depression Inventory. Pairs of emotional and  
22 neutral images (unpleasant-neutral, U-N; pleasant-neutral, P-N; neutral-neutral, N-N) were  
23 presented while attentional indices (eye movements) and autonomic activity (skin conductance  
24 responses, SCRs; heart rate, HR) were recorded. Results showed that all recorded ocular  
25 parameters indicated a preferential orientation and maintenance of attention to emotional  
26 images. SCRs were associated with a valence effect on fixation latency: lower fixation latency  
27 to pleasant stimuli leads to lower SCRs whereas the opposite was observed for unpleasant  
28 stimuli. Finally, stepwise linear regression analysis revealed that latency of fixation to pleasant  
29 images and scores of depression predicted SCRs of participants. Thus, our research reveals an  
30 association between autonomic reactivity and attentional bias to pleasant information, on the  
31 one hand, and depressive symptomatology on the other. Present findings therefore suggest that  
32 depressive individuals may benefit from attention training towards pleasant information in  
33 association with autonomic biofeedback procedures.

34 **Keywords:** Emotion, Depression, Attention bias, Eye movements, Autonomic responses, skin  
35 conductance.

## Introduction

37 Healthy individuals mobilise privileged attentional resources toward emotional  
38 information (Vuilleumier, 2015) and express associated autonomic and behavioural responses  
39 (Damasio, 2000). However, abnormalities in the amount of attention dedicated to emotional  
40 information are implicated in the etiology and maintenance of depressive symptomatology  
41 (Gotlib and Joormann, 2010). For mood-congruent information, depressed individuals are  
42 characterised by biases at all stages of attentional processing (Ingram et al., 1998). Meta-  
43 analysis reveals that, in depression, attentional bias toward negative information is observed  
44 for verbal and non-verbal stimuli, in both clinical and subclinical populations (Peckham et al.,  
45 2010). It is also reported that, in depression, less attention is dedicated to pleasant information  
46 (Duque & Vázquez, 2015). This may reflect an absence of the ‘protective bias’ toward positive  
47 stimuli that is usually observed in healthy or non-dysphoric individuals (Shane & Peterson,  
48 2007). Furthermore, attentional bias to pleasant information is found to correlate negatively  
49 with the onset of depressive symptomatology, and is associated with greater trait resilience.  
50 Hence, attentional bias to pleasant information can be considered as an index of adaptive  
51 emotion regulation (Thoern et al., 2016). Conversely, reduced attentional bias to pleasant  
52 material increases vulnerability to stress-related psychopathology (Fox et al., 2010). In brief,  
53 individuals with depressive symptomatology show greater orientation and maintenance of  
54 attention towards unpleasant stimuli and reduced orientation towards pleasant stimuli (Gotlib  
55 and Joormann, 2010).

56 The dot probe task is one established approach for measuring the time course of  
57 attentional processes in depression (MacLeod et al., 1986). Here, the participant views a pair  
58 of stimuli, usually one emotional and one neutral image, which are immediately followed by  
59 a stimulus (‘probe’), which appears at the location of one of two images. The participant is  
60 instructed to make a reaction time response to the probe. Emotional attentional bias to it  
61 expresses faster responses on trials when the probe is presented in the location of the emotional  
62 compared to the neutral image. This task and variations of it has become a gold standard for  
63 investigating attentional bias and its time course. However, the use of reaction times does not  
64 allow direct measurement of the attention span and has recently been criticised regarding its  
65 psychometric properties (Chapman et al., 2019). As an alternative, measures of eye  
66 movements can provide a more direct index of attentional deployment. In depressed  
67 individuals, they showed an increased attention to negative stimuli and decreased attention to  
68 positive stimulation when compared to nondepressed individuals (Armstrong & Olatunji,

69 2012). Eye movements can reflect both orienting (e.g. initial orientation and latency to first  
70 fixation) and maintenance (e.g. number of fixations, or total duration of fixation; Duque &  
71 Vazquez, 2015) components of attention, as well as attentional reorientation to stimuli.  
72 Indeed, unlike reaction times, eye movements allow the continuous measurement of  
73 attentional processes and can thus better characterise attention biases to emotional stimuli. A  
74 few studies in the literature have investigated the psychometric properties of eye movement  
75 indices and most of them used different stimuli, paradigm or sample characteristics (Waechter  
76 et al., 2014; Lazarov et al., 2016, Skinner et al., 2018, Sears et al., 2019). Although there are  
77 still mixed results for the early attention cues (Skinner et al., 2018), the results for indices of  
78 maintenance of attention such as total fixation time and number of fixations appear  
79 encouraging concerning psychometrics properties (Sears et al., 2019). Consequently, some  
80 authors suggest that eye-tracking measures of attentional bias may have better overall  
81 psychometric properties as compared than traditional RT measures of attentional bias for  
82 children and adults (Chong & Meyer, 2020).

83 Depressive symptoms are associated with altered patterns of autonomic activity, which  
84 has been related to a disengagement from emotional information (Bylsma et al., 2008).  
85 Increases in electrodermal activity (Branković, 2008), skin temperature and respiratory  
86 frequency (Wenzler et al., 2017) and decreases in heart rate variability (HRV; Kemp et al.,  
87 2010) are reported. However, across studies of depression, there is heterogeneity in autonomic  
88 reactivity to emotive stimuli, which remains to be clarified. We propose this heterogeneity  
89 may reflect individual differences in attentional focus (De Zorzi et al., 2021). For example,  
90 depressed individuals show heightened electrodermal reactivity only for stimuli in direct  
91 attentional focus (i.e., central vision) and not for stimulation presented in peripheral vision  
92 (De Zorzi et al. 2020). More generally, attentional processes are linked to the modulation of  
93 autonomic activity. Thus, electrodermal activity, notably the amplitude of sympathetic skin  
94 conductance responses (SCRs), is related to attention-orientation behaviours and reflects  
95 focused attention to new stimuli and their salience (Boucsein, 2012). Similarly, HRV,  
96 reflecting both parasympathetic and sympathetic influences on heart rate, is also considered  
97 as an objective indicator of attentional processes. In this context, superior selective or  
98 sustained attention is associated with increased HRV, particularly in the dominant high-  
99 frequency range (HF-HRV, indexing vagal parasympathetic autonomic activity) (Suess &  
100 Porges, 1994). Interoceptive feedback of autonomic bodily signals also influences emotional  
101 and attentional processes (Critchley & Harrison, 2013). Altered sensitivity to bodily arousal

102 is observed in depression (Paulus & Stein, 2010). As depressive individuals suffer from  
103 attentional disturbance when emotional information is involved, the use of a task allowing to  
104 measure attentional processes appears relevant in the study of their autonomic reactivity to  
105 emotion.

106 The present study aimed to characterise the association of attentional bias and autonomic  
107 reactivity to emotional information in individuals expressing different levels of depression.  
108 To this end, in an original paradigm, we presented pairs of emotional and neutral images, at  
109 near eccentricities within left (-12°) and right (+12°) visual fields. Eye movements were  
110 recorded to enable the tracking of attentional deployment toward one or other images, while  
111 we simultaneously measured autonomic variables (SCR and HR). Accordingly, we  
112 hypothesised that: 1) Attention will be preferentially directed to emotional images in all  
113 participants; 2) attentional bias to emotional information will be associated with autonomic  
114 reactivity (SCR and HR responses); 3) depressive symptoms will be associated with greater  
115 attention towards unpleasant stimuli and reduced orientation towards pleasant stimuli on the  
116 one hand, and with autonomic reactivity on the other hand.

## 117 **Method**

### 118 *Participants*

119 Thirty-four healthy unmedicated participants were recruited through an online  
120 questionnaire. All were French speakers, right-handed and had normal or corrected-to-normal  
121 vision. Individuals with a history of neurological disorders or regular and/or recent illicit drug  
122 consumption were not included. Two participants were excluded due to recording problems,  
123 giving a sample of 32 participants (24 females and 8 males; **Table 1**). Each participant provided  
124 an informed consent statement and received a 20 € compensation for his or her participation.  
125 This study was approved by the Ethics Committee of Université de Lille [Référence: 2019-352-  
126 S73], and conducted in accordance with the Declaration of Helsinki at Faculté de Medecine,  
127 Pôle Recherche, Université de Lille, France.

128 **[Insert Table 1 about here]**

### 130 *Stimuli and Apparatus*

131 The stimuli used were pairs of images of emotional or neutral scenes selected from the  
132 International Affective Pictures System (IAPS; Lang, Bradley, & Cuthbert, 2008), which

133 provides standardised *a priori* values for each image on valence and arousal dimensions. One  
134 value is provided for men, and another for women. Given the recognised differences in  
135 gender-based emotional assessments (Bradley, Codispoti, Cuthbert, & Lang, 2001), we  
136 performed two image selections adapted to each gender, but resulting in equivalent valence  
137 and arousal values. Ninety-six images were selected and used to build three kinds of pairs: 16  
138 unpleasant-neutral pairs (|UN|), 16 pleasant-neutral pairs (|PN|) and 16 neutral-neutral pairs  
139 (|NN|). In order to control the salience of the two images that made up each pair, the difference  
140 of valence and arousal between the two images constituting each pair were calculated. These  
141 within-pairs differences significantly differed between the three kinds of pairs on their *a priori*  
142 valence values (*women*: |U - N| = 2.56, |P - N| = 2.47, |N - N| = 0.54,  $F_{1,15} = 290.196$ ;  $p <$   
143  $0.001$ ; *men*: |U - N| = 2.65, |P - N| = 2.28, |N - N| = 0.40,  $F_{1,15} = 640.406$ ;  $p <$  0.001), and  
144 on their arousal *a priori* values (*women* : |U - N| = 2.74, |P - N| = 2.80, |N - N| = 0.50,  $F_{1,15}$   
145 = 263.916;  $p <$  0.001 ; and *men*: |U - N| = 2.48, |P - N| = 2.67, |N - N| = 0.54,  $F_{1,15} =$   
146 232.177;  $p <$  0.001) with higher within-pairs difference for |UN| and |PN| pairs than |NN| ones,  
147 but there were no differences between |UN| and |PN| pairs. For each pair of images, no within-  
148 pairs differences of valence or arousal was observed between the selections for men and  
149 women (all  $F_s <$  0.275 and  $p_s >$  0.609). For each image, the angular size ( $12^\circ \times 8^\circ$ ), the energy  
150 across spatial frequencies (Delplanque, N'diaye, Scherer, & Grandjean, 2007) and the main  
151 physical properties were extracted (*ImageJ v1.50 software*), including the luminance and  
152 contrasts for the greyscale version and the RGB (red, green and blue) layers. No significant  
153 differences were observed between the three sets of images for both genders (all  $p_s >$  .20).  
154 Thus, the image pairs differed only in terms of their emotional dimensions (see table in  
155 **Appendix Table S1**).

156 Participants were seated at a fixed viewing distance of 60 cm from the projection screen  
157 (30 inches, 256 x 160 ppi, DELL 3007WFP HC), which was connected to a computer (DELL  
158 Optiplex 9020, Windows 7 Professional) that managed the presentation of the pairs of images.  
159 The images were displayed on a black background and each pair of images was presented  
160 pseudo-randomly, based on a Latin squares design, at near visual eccentricities ( $-12^\circ$ ,  $+12^\circ$ ).  
161 The presentation of the 48 trials lasted approximately 15 minutes.

162

163 *Recordings*

164 Anxiety state, trait and depressive scores were measured using French language versions  
165 of the State-Trait Anxiety Inventory (STAI-A & B; Spielberger, 1983) and Beck Depressive  
166 Inventory (BDI-II; Beck, Steer, & Brown, 1996) respectively.

167 Regarding behavioural data, the eye movements were recorded using an eye tracker  
168 (SMI RED-m Eye Tracking System) connected to the projection computer with *SMI iView*  
169 *RED-m 2.11* software for an acquisition at 120 Hz. The skin conductance (SC) and  
170 electrocardiogram (ECG) were recorded during two minutes of baseline, during the task and  
171 over 2-minute recovery periods, using a BIOPAC MP35 system connected to a second  
172 computer (running *BIOPAC Student Pro 3.7* software) for an acquisition at 200 Hz. SC was  
173 recorded using bipolar Ag/AgCl surface electrodes (BIOPAC EL507) pre-gelled with an  
174 isotonic electrolyte (0.05 molar NaCl) and attached to the palmar side of the middle phalanges  
175 of the index and middle fingers of the participant's non-dominant hand. SC was measured  
176 with a gain of 5  $\mu\text{S}/\text{V}$  and a 10 Hz low-pass filter. The ECG was recorded using a DI modified  
177 bypass placing the Ag/AgCl pre-gelled (BIOPAC EL503, 7% NaCl) surface electrodes on the  
178 participant's left and right wrists and with a band-pass filter set between 0.5 and 66.5 Hz. At  
179 the end of the experiment, the participant was required to review each of the images and to  
180 rate them individually for valence and arousal values using two nine-point SAM scales (Self-  
181 Assessment Manikin; Bradley & Lang, 1994), ranging from 1, very unpleasant, to 9, very  
182 pleasant, and from 1, very calm, to 9, very arousing. Ratings were recorded with *OpenSesame*  
183 (Mathôt et al., 2012).

184

185 *Procedure*

186 The experimental procedure was divided into three steps. First, the SCR and ECG  
187 electrodes were attached and the participant was acclimatised to the experimental environment.  
188 The participant completed a psychometric measure of anxiety state (State Anxiety Inventory,  
189 STAI-A; Spielberger et al., 1993), then the task was again explained orally in full.

190 Next, autonomic responses were recorded over a task-free 2-minute baseline period.  
191 This period was then followed by the main task, with recording of behavioural (eye-tracking)  
192 and autonomic responses together. The participant saw 48 pairs of images; 16 pairs of  
193 unpleasant-neutral |UN| images, 16 pairs of pleasant-neutral |PN| images and 16 pairs of neutral-



194 neutral |NN| images presented in a pseudo-random order. Each trial had the following sequence:  
195 First, a central fixation cross was projected for a duration of 0.5 s then this was replaced by the  
196 probe; a digit number (between 1 and 9) replacing the fixing cross for 1 s. The participant was  
197 instructed to say this number as quickly as possible. As shown by Duque and Vazquez (2015),  
198 this procedure ensured that the participant watched the centre of the screen before the  
199 presentation of the stimuli. The procedure also helped maintain the participants' attention  
200 during the experimental phase. Following the presentation of the number, a pair of images was  
201 presented simultaneously, at  $-12^\circ$  and  $+12^\circ$  on the projection screen for 3.5 s, followed by a  
202 black screen for a random duration between 9 to 13 s. Participants were invited to view the  
203 images naturally without any further requirements. The inter-stimulus interval (ISI) varied from  
204 10.5 to 14.5 s; an ideal interval to avoid habituation inherent to autonomic responses,  
205 particularly electrodermal activity. After the task, we recorded autonomic activity during a 2-  
206 minute recovery period.

207 Finally, to validate our selection of images, the participant was asked to rate the valence  
208 and arousal dimensions using the two 9-point SAM scales. These subjective ratings correspond  
209 to *a posteriori* values of images.

210

## 211 *Data and Statistical Analyses*

212 All trials containing eye movements before the onset of the image were rejected to  
213 exclude trials for which gaze is already directed to a image's side. Across all participants and  
214 conditions, 5.14% of the trials were rejected.

### 215 *Eye Movements*

216 Attentional deployment was assessed based on eye movements  
217 (*OpenGazeAndMouseAnalyzer*; Voßkühler et al., 2008). For initial fixation, the percentage of  
218 first fixation for each type of image (unpleasant, pleasant and neutral) in pairs (|UN| and |PN|)  
219 was calculated for each participant. In addition, total fixation duration, number and latency of  
220 fixations were recorded.

### 221 *Conductance and Cardiac Activities*

222 SC and ECG variations initially sampled at 200 Hz, were down-sampled at 10 Hz using  
223 *LabChart7*. For the SC variations to the presentation of pairs of images, phasic waveforms were  
224 derived from the tonic signals with an offline 0.05 Hz high-pass filter using *AcqKnowledge 4.1*  
225 software. SCRs were analysed by computing the integrals of SC amplitude variations over time

226 for each condition and participant. One participant with no SCR was excluded from the analysis.  
227 For the ECG, the instantaneous heart rate in beats per minute (BPM) was calculated from the  
228 R wave intervals and smoothed using the triangular Bartlett window with a 1 s width using  
229 *LabChart7*. SC variations in response to stimulation were obtained by subtracting the average  
230 over a 3 s pre-stimulus period from the 10 s post-stimulus period data. For each condition and  
231 participant, after the baseline correction (-3 to 0 s), we averaged the epochs (-3 to 10 s), time-  
232 locked to the stimulus onset. Finally, we analysed the heart rate variability (HRV) during the 2  
233 minutes of baseline and recovery periods. HRV quantification was computed with an in-house  
234 customised program *MATLAB* program referring to HRV guidelines (Berntson et al., 1997).  
235 The R-R intervals were detrended with a smoothness-prior method in order to remove the slow  
236 (< 0.04 Hz) non-stationary trends from the HRV signal. For the frequency domain method, a  
237 power spectrum density analysis was performed for the RR interval series using fast Fourier  
238 transform method with the low frequency (LF) band set at 0.04-0.15 Hz and a high frequency  
239 (HF) band set at 0.15-0.4 Hz. The LF/HF ratio was also computed. For the time domain method,  
240 we computed the mean heart rate (HR), the standard deviation of HR (namely the variability of  
241 the HR) as well as the root mean square of successive RR intervals differences (RMSSD).

## 242 *Statistical Analyses*

243 Regarding emotion and in accordance with its dimensional theory (Lang et al., 1993), we  
244 tested for two emotional effects: 1) A valence effect (Unpleasant vs. Pleasant), being modelled  
245 by a first degree polynomial contrast (*Linear Contrast*,  $LC = |PN| - |UN|$  ; and 2) an arousal  
246 effect (Emotion vs. Neutral) being modelled by a second degree polynomial contrast (*Quadratic*  
247 *Contrast*,  $QC = (|UN| + |PN|) / 2 - |NN|$ ). These contrasts were assessed with a repeated  
248 measure analysis of variance (ANOVA), applied to the individual subjective image  
249 assessments, eye movements, electrodermal and cardiac measurements with emotion (type of  
250 pairs:  $|UN|$ ,  $|NN|$ ,  $|PN|$ ) as intra-subject factors.

251 The analysis of the factors associated with the autonomic variables was performed by  
252 calculating the Pearson correlation coefficient given linear relationships were expected between  
253 variables and after inspection of scatterplots. Partial correlations were then assessed controlling  
254 for age and gender. The search for predictors of autonomic variables was performed with  
255 stepwise linear regression analyses. The multivariate model includes variables for which  
256 associations were observed between ocular and autonomic parameters. Thus, the model was  
257 constructed by including variables associated with SCRs, but also including covariates such as  
258 the age of participants, and their state and trait anxiety (STAI-A and STAI-B scores), regardless

259 of their degree of significance in the univariate analyses. The model selection was based on  
260 considerations of the corrected Akaike information criterion (AICc). The validity of the  
261 multivariate model was established by a study of the residuals.

## 262 **Results**

263 Concerning psychometry, STAI-B (trait anxiety) scores correlated with BDI (depression)  
264 ( $r_{33} = 0.810$ ;  $p < 0.001$ ) and STAI-A (state anxiety) scores ( $r_{33} = 0.631$ ;  $p < 0.001$ ). BDI scores  
265 correlated with STAI-A scores ( $r_{33} = 0.651$ ;  $p < 0.001$ ).

### 266 *Eye Movements and Emotional Arousal*

267 Analysis of the contrasts revealed an emotional arousal effect on the initial fixation (QC:  
268  $F_{(1,31)} = 7.144$ ;  $p = 0.012$   $\eta^2 = 0.187$ ), first fixation latency (QC:  $F_{(1,31)} = 75.624$ ;  $p < 0.001$ ;  $\eta^2$   
269  $= 0.709$ ), fixation duration (QC:  $F_{(1,31)} = 107.966$ ;  $p < 0.001$ ;  $\eta^2 = 0.777$ ) and number of fixation  
270 (QC:  $F_{(1,31)} = 125,181$ ;  $p < 0.001$ ;  $\eta^2 = 0.802$ ; **Figure 1**). However, no differences were observed  
271 between unpleasant and pleasant for these parameters (initial fixation: LC:  $F_{(1,31)} = 2.827$ ;  $p =$   
272  $0.103$   $\eta^2 = 0.084$ ; first fixation latency : LC:  $F_{(1,31)} = 0.040$ ;  $p = 0.843$ ;  $\eta^2 = 0.001$ ; fixation  
273 duration : LC:  $F_{(1,31)} = 0.92$ ;  $p = 0.764$ ;  $\eta^2 = 0.003$ ; number of fixation : LC:  $F_{(1,31)} = 0.049$ ;  $p$   
274  $= 0.826$   $\eta^2 = 0.002$ ). In sum, eye movements were initially oriented and engaged by emotional  
275 images.

276 **[Insert FIGURE 1 about here]**

277

### 278 *Psychometry, Eye Movements and Autonomic Activity*

279 Mean SCRs to pairs of images correlated with depression ( $r_{32} = 0.475$ ;  $p = 0.005$ ). Thus,  
280 higher SCRs to images were associated with higher depression scores even when age or sex  
281 were controlled (respectively  $r_{30} = 0.476$ ;  $p = 0.006$  and  $r_{30} = 0.495$ ;  $p = 0.004$ ). HF-HRV at  
282 baseline correlated with the duration of fixation on pleasant images. Thus, higher HF-HRV at  
283 baseline was associated with a longer duration of fixation on pleasant images during the task  
284 ( $r_{31} = 0.374$ ;  $p = 0.035$ ). This association was still observed when age was controlled ( $r_{29} =$   
285  $0.375$ ;  $p = 0.037$ ) and was still marginally significant when controlled for sex ( $r_{29} = 0.320$ ;  $p =$   
286  $0.079$ ). During the task, a correlation was observed between ocular parameters and autonomic  
287 reactivity: the difference of fixation latency between pleasant and unpleasant images (valence  
288 effect) correlated with mean SCRs ( $r_{30} = -0.381$ ;  $p = 0.034$ ) even when age or sex were

289 controlled (respectively  $r_{28} = -0.380$ ;  $p = 0.038$  and  $r_{28} = -0.438$ ;  $p = 0.015$ ). No other  
290 associations were found between ocular and autonomic variables ( $r_s > 0.281$ ;  $p_s > 0.120$ ) nor  
291 between ocular variables and depression or anxiety ( $r_s < 0.249$ ;  $p_s > 0.170$ ).

### 292 *Depression, Eye Movements and Autonomic Reactivity*

293 As SCRs correlated with depression scores and with the valence effect on the fixation  
294 latency, the stepwise linear regression model was performed by including depression (BDI  
295 scores) and the latency of fixing pleasant and unpleasant images, the age of participants, and  
296 their state and trait anxiety (STAI-A and STAI-B scores).

297 On the basis of the corrected Akaike information criterion (AICc), the selected model  
298 for predicting SCRs was found to explain 21% of the variance in the integral of SCR ( $F_{3,29} =$   
299  $3.60$ ,  $p = 0.027$ ) and include the depression scores (BDI), the latency to fix pleasant images and  
300 the trait-anxiety scores (STAI-B) as predictors. The measures found did not point to the  
301 existence of significant collinearity between the predictors, minimal tolerance = 0.429 and  
302 maximal variance inflation factor (VIF) = 2.33 (**Figure 2 A1**).

303 **[Insert FIGURE 2 about here]**

304 In the model, depression scores ( $t = 2.22$ ,  $p = 0.035$ ) and the latency to fix pleasant  
305 images ( $t = 2.33$ ,  $p = 0.028$ ) significantly contributed to a better prediction of the SCRs while  
306 the trait-anxiety scores ( $t = -1.92$ ,  $p = 0.066$ ) did not significantly contribute to the predictive  
307 power of the model. Hence, greater SCRs were associated with higher depression, with a  
308 coefficient of 1.30, and higher latency to fix pleasant images, with a coefficient of 0.05.  
309 Interestingly, contrary to what has been observed for depression scores and the latency to fix  
310 pleasant images, the coefficient describing the direction of the relation between STAI-B scores  
311 and SCRs was negative.

312 In sum, the best model to predict SCRs integrate latency of fixation to pleasant images,  
313 depression and anxious scores such higher depression level and lower orientation to pleasant  
314 stimuli were expected to predict higher SCRs, after controlling for the other variables.

## 315 **Discussion**

316 The aim of this study was to investigate potential links between attentional bias and  
317 autonomic reactivity to emotional information and the implication of depressive  
318 symptomatology on these potential associations. Firstly, all recorded parameters of ocular

319 behaviour indicated a preferential orientation of attention to emotional images. Secondly,  
320 higher orientation and maintenance of attention towards pleasant images were associated with  
321 lower values of autonomic arousal during baseline (HRV) and during the task (SC). Thirdly,  
322 the best model to predict SCRs of participants includes latency of fixation to pleasant images,  
323 and scores of depression and anxiety.

324 The first result, showing a preferential orientation and maintenance of attention toward  
325 emotional contents, reflected by the initial fixation and the shorter latency to fixate upon  
326 emotional images, extends data from previous eye-tracking studies demonstrating that  
327 emotional stimuli benefit from enhanced perceptual processing, with more fixations, especially  
328 during the first saccades (Niu et al., 2012). In terms of unpleasant and pleasant value of images,  
329 no attentional preference was observed for one over the other, suggesting that attentional bias  
330 depends only on the arousal dimension of emotion.

331 Depressive symptomatology was not correlated with any ocular parameter which is  
332 consistent with few studies that failed to report bias for emotion in people with depression or  
333 dysphoria (Koster et al., 2006; Elgersma et al., 2018). Moreover, data from the literature report  
334 an attentional bias in depression towards “mood-congruent” or “dysphoric” information, using  
335 discrete emotional stimuli, and mainly for attention maintenance indicators (Armstrong &  
336 Olatunji, 2012). The present study did not use mood-congruent stimuli but natural scene based  
337 on dimensional theory of emotion, which could explain that such correlation was not observed.  
338 Besides, in the present study, the images’ selection was carried out in order to ensure  
339 homogenised arousal differences between the two images constituting the pairs. Thereby,  
340 attention may have been captured by the emotional image regardless of symptomatology for all  
341 participants.

342 The second result supports specific links between attentional bias to pleasant  
343 information and the expression of autonomic arousal both at baseline and during the task.  
344 During the baseline, greater HF-HRV was associated with longer fixation on pleasant images.  
345 Hence, higher parasympathetic influence, and thus increased HF-HRV, appears linked to  
346 maintenance of attention toward pleasant information. The polyvagal theory (Porges, 2007)  
347 proposes that baseline cardiac measures of parasympathetic activity can index the capacity to  
348 adapt to the environment. More precisely, parasympathetic HRV activity at rest and reactivity  
349 are associated with adaptive expression of emotion and self-regulatory skills. Therefore, our  
350 observed association between baseline autonomic activity and preferential attention to pleasant  
351 information reveals a positive impact of a more parasympathetic psychophysiological state on

352 an upcoming emotional task in the domain of emotional regulation (Beauchaine, 2001).  
353 Individuals with higher parasympathetic activation at baseline may be better able to employ the  
354 best strategies to respond to stressful emotional challenges by focusing on pleasant information.  
355 The present results also reinforce the neurovisceral integration model (Thayer and Lane, 2000),  
356 which postulates that cardiac activity, through HRV, is informative about the integrity of brain  
357 networks supporting interaction between emotion and cognition. However, this association  
358 became marginally significant when sex is controlled. Indeed, sex differences on HRV have  
359 been reported with higher HRV in women characterised by a relative dominance of vagal  
360 activity (Koenig and Thayer, 2016). Due to the sample size, it seems difficult to conclude on  
361 sex differences for HRV and it is therefore advisable to remain cautious about this result.  
362 Additional data on the association between attentional and autonomic measurement in relation  
363 to sex are necessary and may point out different associations for these parameters for men and  
364 women.

365         During the task, a valence effect on the fixation latency was associated with autonomic  
366 reactivity. Individuals who fixated upon pleasant stimuli more quickly showed lower SC to the  
367 pairs of images whereas the reverse was observed for fixation on unpleasant stimuli. This  
368 positivity bias, described in the attentional literature (Troller-Renfree et al., 2017) bears witness  
369 to a regulatory interaction between the capture of information and autonomic adaptation.

370         The third result showed that the best model to predict SCRs integrate latency of fixation  
371 to pleasant information with depression and trait-anxiety scores. At psychometric level, higher  
372 depression predicted greater SCRs in response to pair of images while trait anxiety did not  
373 significantly contribute to the prediction of SCRs. At attentional level, measured by images'  
374 fixation latency, a lower attentional orientation towards the pleasant images was associated with  
375 greater SCRs. This association between pleasant orientation and autonomic reactivity is  
376 particularly interesting since depressed individuals appear to lack positive attentional bias  
377 (Duque & Vazquez, 2015) and are characterised by autonomic dysfunctions sometimes reported  
378 as increased autonomic activity (Branković, 2008; Wenzler et al., 2017). Consequently,  
379 attempts to reinforce such bias could help to attenuate autonomic activation and potentially  
380 serve as a protective homeostatic adaptation or a coping strategy. In the same vein, similar  
381 autonomic hyporeactivity to emotional challenges has already been reported, and interpreted as  
382 the expression of a coping strategy engaged to improve performance of a behavioural task  
383 (Naveteur et al., 2005). In this context, an intervention to attenuate autonomic reactivity could  
384 increase capacity to orient attention towards pleasant information which may be beneficial

385 especially for depressed individuals. Therefore, these results suggest that the pleasantness bias  
386 could constitute a cognitive marker of behavioural and autonomic adaptations to emotion.

387 Finally, this study has several methodological strengths. First, we carefully considered  
388 influences of physical saliency of images on visual search and attention (Lucas & Vuilleumier,  
389 2008) and gender differences in emotional assessment and reactivity (Bradley et al., 2001).  
390 Thus, we carried out a rigorous selection of stimuli for each type of pair of images (see  
391 **Appendix 1; Table S1**). Second, this study was enriched by taking into account both facets of  
392 the autonomic nervous system, sympathetic and parasympathetic, which potentially have  
393 distinct contributions to attentional and emotional processes. Third, the integration of  
394 behavioural and autonomic parameters, encompassing their reciprocal influences, allowed us  
395 to clarify the relationship between attentional bias and autonomic reactivity and the relation  
396 with depressive symptomatology. The study also had some limitations, notably the sex-ratio of  
397 participants. Indeed, even if images selection was homogenised between women and men, sex  
398 of participants seems to influence some of the association observed (e.g. HRV) and the sample  
399 size for men do not allow to examine potential sex differences properly on these associations.  
400 Therefore, additional researches are needed to determine potential implication of this variable  
401 on interaction between attention and autonomic expression. Moreover, the choice of a non-  
402 clinical population constrains the scope of our results to moderate levels of depression. Indeed,  
403 participants are not clinically depressed in our sample. Nonetheless, even when considering  
404 lower depression levels, prominent effects were observed. Moreover, findings remain relevant  
405 since attentional bias is considered to be implicated in the development of clinical depression  
406 (Gotlib and Joorman, 2010).

407 In conclusion, our characterisation of the association between attentional processes and  
408 specific patterns of autonomic reactivity extends a growing literature, and points in the direction  
409 of reciprocal influences of both systems that can lead to the personalisation of clinical  
410 remediation and rehabilitation procedures, including attentional training and/or biofeedback  
411 therapies. Furthermore, the data obtained in this work argue in favour of evidence-based  
412 interventions using attentional training towards pleasant information. Consequently,  
413 considering the lack of orientation to pleasant information and autonomic dysfunction in  
414 depression, two strategies can emerge from these results. On the one hand, in depressed  
415 individuals who lack strong attentional bias toward pleasantness, attentional training towards  
416 emotionally positive information may be beneficial in reducing autonomic hyperarousal and  
417 reactivity to emotional information. On the other hand, in depressed individuals who show  
418 autonomic hyperarousal, biofeedback procedures based on HRV may be therapeutically

419 beneficial, in part by fostering enhanced attentional bias toward pleasant information. Finally,  
420 our research supports an approach integrating both cognition and physiology to better  
421 understand their interdependence in healthy and pathological expressions of emotion.

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## 427 **Disclosure statement**

428 No potential conflicts of interest were reported by the author(s).

429

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437

438



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547

548 *Table and Figures*

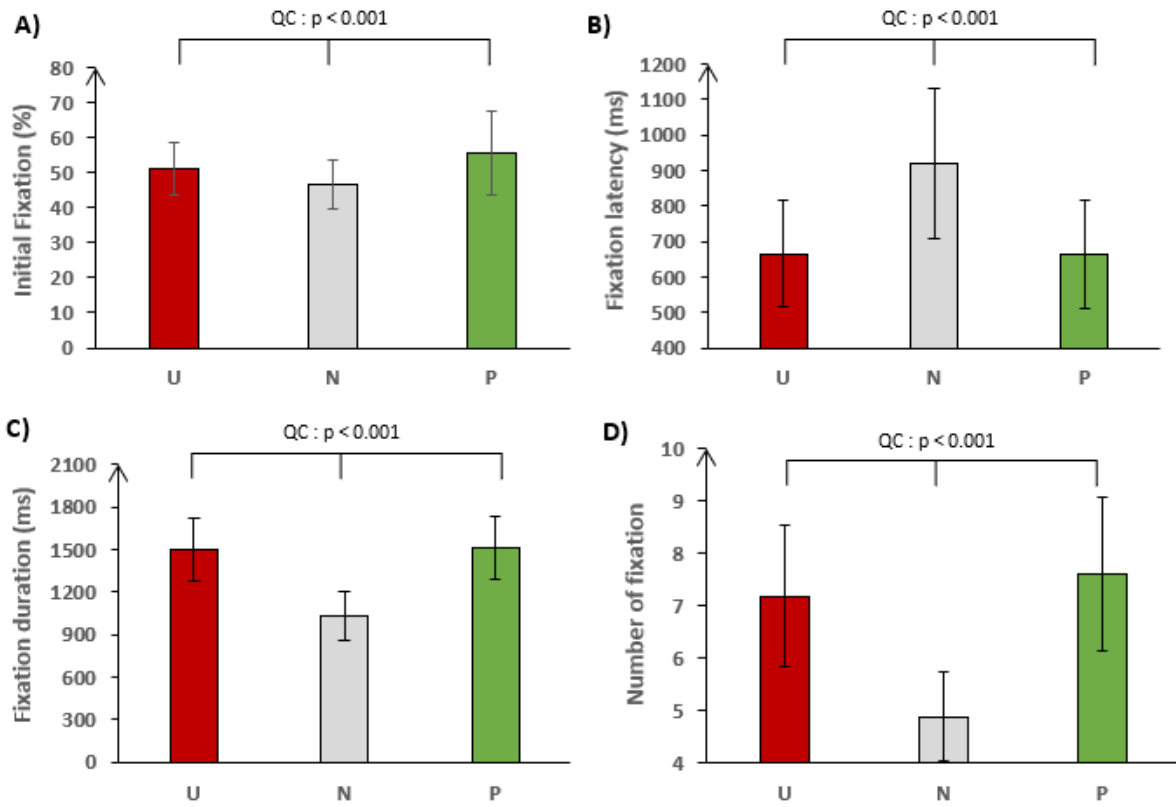
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Sample (N = 32)			
	Mean	SD	(Min – Max)
Age	21	2	(18 – 24)
STAI-A	30.4	9.7	(20 – 58)
STAI-B	47.1	12.8	(25 – 71)
BDI-II	13.3	10.4	(1 – 41)

550

**Table 1. Demographic and psychometric characteristics of participants.**

551



552 **Figure 1. Initial fixation and fixation latency (A-B) and total fixation duration and number of fixations (C-D) to image**  
 553 **emotion in |UN| and |PN| pairs. U: Unpleasant; N: Neutral; P: Pleasant; QC: Quadratic contrast.**

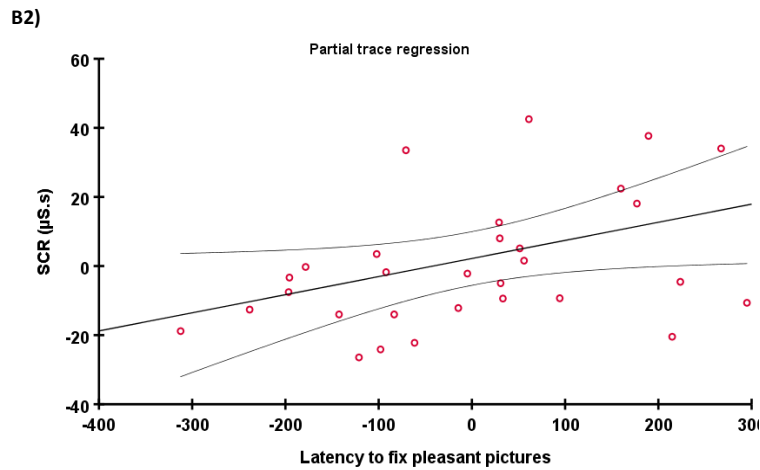
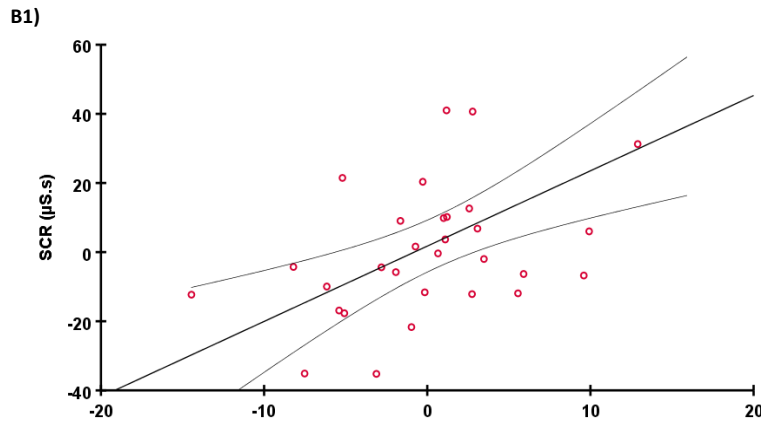
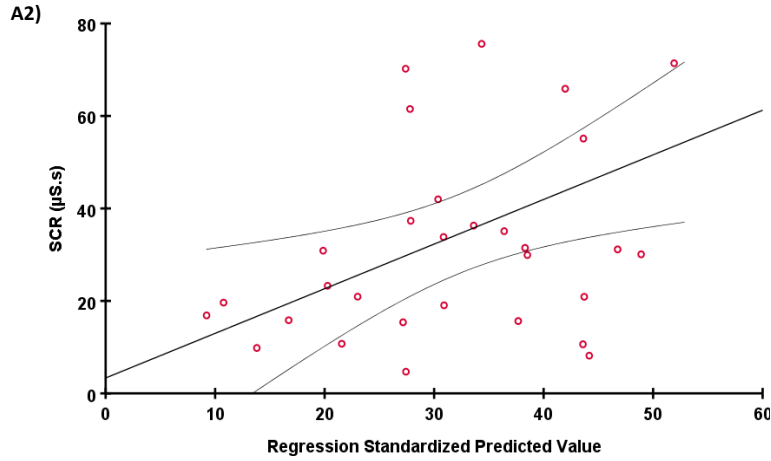
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A1)

Predictors	Slope	$\beta$	t	Sig.	VIF
STAI-B	-0.842	-0.484	-1.922	0.066	2.329
BDI	1.302	0.559	2.220	0.035	2.332
Latency to fix pleasant	0.050	0.384	2.325	0.028	1.003

One participant was excluded (N = 30) due to a Cook's distance value greater than 0.2.



**Figure 2. A. Multiple linear regression model to predict SCRs.** VIF: maximal variance inflation factor, index allowing to verify the premise of multicollinearity (A1) and representation of the model (A2) B. Partial trace regression with intervals of confidence for BDI scores (B1) and for latency of fixation to pleasant images (B2).

	Women				Men				W/M
	U-N	N-N	P-N	p	U-N	N-N	P-N	p	p
<b>Valence</b>	2,563 (0,829)	0,548 (0,407)	2,473 (0,668)	> 0,001	2,652 (0,621)	0,401 (0,241)	2,281 (0,663)	> 0,001	0.502
<b>Activation</b>	2,746 (0,949)	0,501 (0,344)	2,805 (0,984)	> 0,001	2,485 (0,937)	0,547 (0,349)	2,678 (1,066)	> 0,001	0.502
<b>Luminance</b>	42,973 (37,520)	37,069 (25,382)	41,300 (34,078)	0,871	41,292 (24,261)	41,985 (30,566)	45,121 (27,293)	0,916	0.704
<b>Contrast</b>	19,411 (13,826)	21,620 (12,766)	18,570 (12,383)	0,792	15,798 (11,064)	17,726 (9,295)	16,286 (14,325)	0,890	0.200
<b>Luminance (R)</b>	42,846 (35,427)	36,542 (22,893)	46,769 (35,522)	0,659	45,321 (35,155)	45,965 (29,572)	48,051 (32,176)	0,969	0.504
<b>Contrast (R)</b>	19,717 (13,838)	22,692 (15,125)	18,869 (12,797)	0,720	20,565 (14,880)	16,888 (10,719)	17,884 (11,968)	0,698	0.468
<b>Luminance (G)</b>	41,892 (38,487)	39,115 (32,486)	43,611 (34,817)	0,936	44,000 (26,756)	44,386 (31,426)	47,708 (31,503)	0,929	0.569
<b>Contrast (G)</b>	18,969 (14,722)	21,646 (15,968)	19,000 (12,785)	0,837	16,549 (13,788)	16,536 (12,298)	16,997 (14,822)	0,994	0.273
<b>Luminance (B)</b>	54,553 (41,430)	44,421 (30,608)	53,355 (41,504)	0,716	47,938 (18,047)	52,475 (37,773)	48,761 (28,520)	0,897	0.880
<b>Contrast (B)</b>	23,999 (14,607)	27,288 (17,579)	23,142 (18,017)	0,763	17,734 (14,854)	21,057 (17,558)	26,196 (21,305)	0,419	0.380
<b>Low frequencies (Grey)</b>	6,775 (6,253)	9,781 (11,364)	6,164 (3,779)	0,381	6,148 (9,917)	8,0253 (7,804)	6,3499 (3,820)	0,747	0.643
<b>High frequencies (Grey)</b>	1,152 (1,387)	1,479 (2,568)	0,925 (0,812)	0,669	1,380 (2,560)	1,479 (1,762)	0,975 (0,804)	0,719	0.801
<b>Low frequencies (R)</b>	6,149 (4,876)	8,315 (9,291)	6,220 (3,798)	0,562	7,133 (8,703)	5,921 (6,927)	4,930 (3,348)	0,651	0.504
<b>High frequencies (R)</b>	0,999 (0,997)	1,698 (3,140)	0,9172 (0,764)	0,468	1,363 (2,586)	1,705 (2,357)	0,944 (0,727)	0,583	0.748
<b>Low frequencies (G)</b>	6,020 (5,721)	9,132 (11,022)	5,576 (3,305)	0,345	5,193 (8,357)	7,280 (8,227)	5,560 (3,812)	0,678	0.546
<b>High frequencies (G)</b>	1,048 (1,133)	1,583 (2,716)	0,929 (0,756)	0,537	1,319 (2,591)	1,579 (1,767)	0,861 (0,751)	0,548	0.858
<b>Low frequencies (B)</b>	7,178 (5,619)	10,506 (14,983)	4,958 (4,328)	0,267	6,911 (10,878)	7,728 (10,712)	6,085 (5,361)	0,884	0.741
<b>High frequencies (B)</b>	1,110 (1,309)	1,286 (1,790)	0,949 (0,935)	0,792	1,053 (1,653)	1,479 (1,523)	0,994 (1,037)	0,582	0.834

569

570

**Table S1. Physical properties from the pairs of images selected.** Mean value and standard deviation of

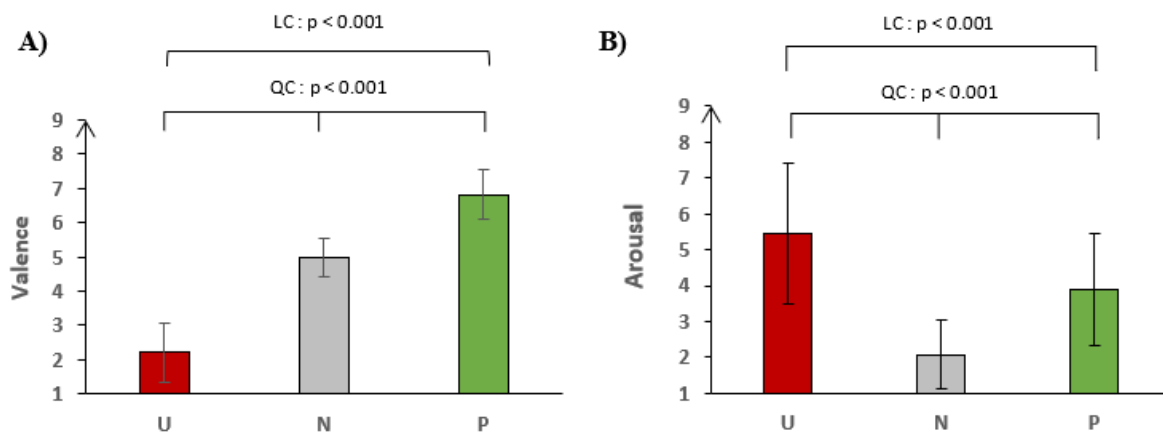
571 differences between the two images composing each pair concerning valence, activation and physical properties for women

572 and men and for unpleasant-neutral (U-N), neutral-neutral (N-N) and pleasant-neutral (P-N) conditions. R = red, B = blue, G

573 = green, p = p value of multivariate analyses for each sex, W/M = comparisons between women and men.

574 *Appendix 2: Subjective assessment of images*

575 As expected, the valence assessment by the participants differed according to the  
576 emotional category ( $F_{1,41,46.67} = 322.085$ ;  $p < 0.01$ ;  $\eta^2 = 0.907$ ;  $U = 2.21$ ,  $N = 4.98$ ,  $P = 6.82$ ).  
577 Thus, participants rated unpleasant images with a lower valence than pleasant ones (LC:  $F_{1,33} =$   
578  $387.70$ ;  $p < 0.001$ ;  $\eta^2 = 0.922$ ) but the valence gap with neutral images was more important for  
579 unpleasant images (QC:  $F_{1,33} = 24.113$ ;  $p < 0.001$ ;  $\eta^2 = 0.422$ ; **Figure S1.A**). The arousal  
580 assessment by the participants also differed according to the emotional category ( $F_{1,67,55.34} =$   
581  $109.110$ ;  $p < 0.001$ ;  $\eta^2 = 0.768$ ;  $U = 5.45$ ,  $N = 2.07$ ,  $P = 3.89$ ). Participants rated emotional  
582 images with a greater arousal than neutral ones (QC:  $F_{1,33} = 217.006$ ;  $p < 0.001$ ;  $\eta^2 = 0.868$ ) but  
583 they evaluated unpleasant images as more arousing than pleasant ones (LC:  $F_{1,33} = 38.370$ ;  $p$   
584  $< 0.001$ ;  $\eta^2 = 0.868$  **Figure S1.B**).



585 **Figure S1. Subjective assessment of valence (A) and arousal (B).** U: Unpleasant; N: Neutral; P: Pleasant; LC: Linear  
586 contrast; QC: Quadratic contrast.