1	The neural correlates of economic value and valuation context: An event-related
2	potentials study
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26 Abstract

27 The value of environmental cues and internal states is continuously evaluated by the human brain and it is this subjective value that largely guides decision-making. The present 28 29 study aimed to investigate the initial value attribution process, specifically the spatio-30 temporal activation patterns associated with values and valuation context using 31 electroencephalographic event-related potentials (ERPs). Participants completed a stimulus 32 rating task in which everyday household items marketed up to a price of £4 were evaluated 33 with respect to their desirability or material properties. The subjective values of items were 34 evaluated as willingness-to-pay (WTP) in a Becker-DeGroot-Marschak auction. Based on the 35 individual's subjective WTP values, the stimuli were divided into high and low value items. 36 Source dipole modelling was applied to estimate the cortical sources underlying ERP 37 components modulated by subjective values (high vs. low WTP) and the evaluation condition 38 (value-relevant vs. value-irrelevant judgments).

39 Low WTP items and value-relevant judgements both led to a more pronounced N2 40 visual evoked potential at right frontal scalp electrodes. Source activity in right anterior insula and left orbitofrontal cortex was larger for low vs. high WTP at around 200 ms. At a similar 41 42 latency, source activity in right anterior insula and right parahippocampal gyrus was larger for value-relevant vs. value irrelevant judgements. A stronger response for low- than high-43 44 value items in anterior insula and orbitofrontal cortex appears to reflect aversion to low-45 valued item acquisition which, in an auction experiment, would be perceived as a relative loss. This initial low-value bias occurs automatically irrespective of the valuation context. 46

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50 New & Noteworthy

51	We demonstrate the spatio-temporal characteristics of the brain valuation process using
52	event-related potentials and willingness-to-pay as a measure of subjective value. The N2
53	component resolves values of objects with a bias toward low-value items. The value-related
54	changes of N2 component are part of an automatic valuation process.
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75 **1. Introduction**

Economic values of stimuli are continuously and automatically encoded in the human brain. Previous brain imaging studies show that valuation occurs predominantly in the orbitofrontal cortex (OFC), ventromedial prefrontal cortex (vmPFC) and the ventral striatum (Bartra et al. 2013; Clithero and Rangel 2014; Padoa-Schioppa 2007; Raghuraman and Padoa-Schioppa 2014).

81 Value attribution is one of the first stages of any value based decision (Rangel et al. 82 2008). Previous studies investigated the modulation of event-related potential (ERP) 83 components by hedonic aspects of visual stimuli (for a review, see Hajcak et al. 2012). For 84 example, a negativity bias reflecting preferential processing of unpleasant stimuli may result in greater ERP responses (Delplanque et al. 2006; Huang and Luo 2006; Smith et al. 2003). 85 86 Some studies identified the role of the late positive potential in the encoding of emotional 87 stimulus valence (Foti et al. 2009; Macnamara et al. 2009; Moser et al. 2006), however, the late positive potential also varies as a function of motivational significance (i.e., salience; 88 89 Weinberg and Hajcak 2010). Although the subjective pleasantness of a stimulus may contribute to the value of perceived goods, economic value is not identical to emotional 90 91 valence.

92 Electrophysiological studies have highlighted that value-related signals appear as 93 early as 200 ms post-stimulus presentation in binary decision tasks where a choice between 94 two options is required (Larsen and O'Doherty 2014; Tzovara et al. 2015). Differences in 95 ERPs were also observed across multiple time windows ranging from 150 to 800 ms (Harris 96 et al. 2011). However, ERPs were not investigated in relation to behavioural measures 97 concerning economic value directly. Other investigations of the value-encoding phase were focussed within specific brain regions (Hunt et al. 2012). A common finding in previous ERP 98 99 studies investigating the representation of value-based preferences in binary reaction time

100 tasks was a progression of activations from the occipito-temporal cortical regions to frontal 101 and prefrontal sites over the course of the ERP (Harris et al. 2011; Larsen and O'Doherty 102 2014). However, the involvement of a reaction time response in experiments investigating the 103 representation of value also adds a motor readiness component to ERPs which may interact 104 with activations related to the automatic valuation process occurring in absence of decision 105 making (Gluth et al. 2013; Polania et al. 2014). Further, binary decision making as compared 106 to reporting hedonic ratings has been found to involve different brain regions, such as 107 anterior cingulate cortex (Rolls et al. 2009).

108 Several ERP components relevant to value-based decision making have been revealed 109 in previous literature. The event-related negativity (ERN) and feedback-related negativity 110 (FRN) are two ERP components that, due to their nature, allow us to investigate decision 111 making processes (Walsh and Anderson 2012). These two components are elicited by 112 feedback following decision tasks and are relevant to reward-prediction errors (Gehring et al. 113 2012; Nieuwenhuis et al. 2004; Yu and Huang 2013). Additionally, the P300 ERP component 114 is often implicated in which the P300 encodes outcome valence (San Martin 2012; Yeung and 115 Sanfey 2004). It is generally found that these ERP components are specific to outcome 116 processing, though it has been revealed that the eliciting stimuli can modulate the ERP magnitude at the outcome stage (Yeung and Cohen 2006). 117

A common method for estimating the economic value of goods is via auction tasks such as the Becker-DeGroot-Marschak (BDM) mechanism (Becker et al. 1964). The BDM mechanism is an incentive compatible method for estimating a subject's willingness-to-pay (WTP) for goods and prospects (Wilkinson and Klaes 2012). Previous functional magnetic resonance imaging (fMRI) studies have established that the brain valuation system activates during the BDM mechanism (Chib et al. 2009; Plassmann et al. 2010; 2007).

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124 The context in which economic decisions are made can also influence the neural 125 activations within the brain valuation system. For example, neural responses within valuation regions can be modulated during an auction task in which bids may be forced (Plassmann et 126 127 al. 2010; 2007), passive viewing tasks (Levy et al. 2011) and tasks in which value is 128 irrelevant (Grueschow et al. 2015; Polania et al. 2014) or where outcomes are uncertain 129 (Payzan-LeNestour et al. 2013). Activation of the brain valuation system during tasks in 130 which it was not required demonstrates the automaticity of valuation processes (Lebreton et 131 al. 2009).

The aim of the present study was to investigate the spatio-temporal aspects of brain economic evaluation of everyday household items during a task in which value was either task-relevant or irrelevant. Subjects viewed each item but were not requested to make a speeded response, rather, they rated the likeability or the material features of the item. A BDM auction experiment was used to evaluate WTP in a separate session, and the WTP values were correlated with ERPs and subjective ratings.

138 **2. Methods**

139 2.1. Participants

Twenty-five healthy participants (14 females) with a mean age of 24 ± 4.67 (mean ±
SD) years took part in the study. The experimental procedures were approved by the
Research Ethics Committee of the University of Liverpool. All participants gave written
informed consent in accordance with the declaration of Helsinki. Participants were
reimbursed for their time and travel expenses.

145 2.2. Procedure

146 All experimental procedures were carried out in a dimly lit, sound attenuated room.

147 Participants sat in front of a 19-inch LCD monitor. The study was carried out in two sessions

148 approximately 2-5 days apart. During the first session, participants completed the auction

149task. During the second session, participants completed the rating task. The stimuli comprised15090 everyday household items varying in value from £0.75 to £4.00 with a mean value of151£2.52 \pm £1.01 (mean \pm SD) obtained from a shopping catalogue. Food items were excluded to152avoid confounds arising from difference in the appetitive value of stimuli between session 1153and 2 of the study. Stimuli were presented in random order. Presentation of stimuli was154controlled using Cogent 2000 (UCL, London, UK) in Matlab 7.8 (Mathworks, Inc., USA).155Experimental protocols and stimulus timings are illustrated in Figure 1.

156 2.3. Auction task

157 The protocol for the auction task was adapted from previous studies (Plassmann et al.
158 2007, 2010) and employed the BDM mechanism (Becker et al. 1964; Wilkinson and Klaes
159 2012). Each stimulus was presented once resulting in a total of 90 auctions.

160 Each auction consisted of a fixation cross followed by an evaluation stage, a bidding 161 period and then feedback. During the evaluation stage, participants appraised the stimulus 162 that was presented on-screen. The bidding period required the participants to bid on the item. 163 Here, participants were asked to bid between £0 and £4 in increments of £0.50 giving a total of nine options. During the feedback stage, participants were notified as to whether or not the 164 item was won. The outcome of an auction was dependent on the bid and a randomly 165 generated number, in which the item was purchased when $b \ge r$, where b represents the bid 166 167 and r represents the randomly generated number for that auction. At the end of the 168 experiment, three auctions resulting in a purchase were selected at random. For each auction 169 selected, a price equal to r was subtracted from an initial endowment of $\pounds 12$. Therefore, the 170 actual endowment could vary between $\pounds 0$ and $\pounds 12$. The participant could pick up the items 171 won within a few days of completion of the full experiment.

172 *2.4. Rating task*

173 Approximately 2-5 days following completion of the auction task, participants 174 returned to take part in session 2. EEG was recorded continuously using the 128-channel Geodesics EGI system (Electrical Geodesics, Inc., Eugene, Oregon, USA) with the sponge-175 176 based HydroCel Sensor Net. The sensor net was aligned with respect to three anatomical 177 landmarks (two pre-auricular points and the nasion). Electrode-to-skin impedances were kept 178 below $50k\Omega$ and at equal levels across all electrodes as recommended for the system (Ferree 179 et al. 2001; Luu et al. 2003; Picton et al. 2000). The sampling rate was 1000 Hz and Cz was 180 used as the initial reference. Data was filtered online using a 0.1-200 Hz bandpass filter. 181 After fitting the EEG cap, participants completed a computerized rating task. Each 182 trial began with a fixation cross followed by an instruction stage, evaluation period and then 183 rating. During the instruction stage, participants were presented with either the word 184 'DESIRABILITY' or 'MATERIAL', which served to cue the participant to the required type 185 of evaluation. The evaluation stage began with the presentation of one of the visual stimuli, 186 followed by the presentation of a visual analogue scale (VAS) for the rating stage. In the 187 value-relevant condition, the participant would have to rate the desirability of the preceding stimulus (anchors: "neutral"-"very desirable"), while in the value-irrelevant condition, the 188 189 participant would rate the proportion of the preceding stimulus composed of a certain 190 material (for example, "none"-"plastic"). Here, the proportion of the scale that is shaded 191 indicated the percentage of plastic composition. Desirability and material estimation trials 192 were randomly intermixed within blocks.

Investigating the neural basis of subjective value is complicated by the multiple nonspecific neural processes elicited during experimental paradigms used to reveal subjective value. During the rating task, the only difference between these two conditions was the calculation of subjective value for the trials in which desirability was rated. Any differences in ERPs between these two trials can, therefore, be attributed to computation required to

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report subjective value. Of course, automatic processes involved in valuation would still be
present. Each stimulus was presented in both conditions, yielding a total of 180 trials, split
into three blocks.

201 2.5. Median split of WTP values

202 The stimulus set was divided into high and low WTP items using a median split of 203 subjective values. In the case of items with identical value on both sides of the split, the items 204 with that value were removed in such a manner that there was no overlap in value between 205 the two sides and there was an equal number of stimuli in each category. For an unequal 206 number of stimuli of identical value on each side of the split, stimuli of that value were 207 removed randomly from the side with more. This produced two categories of stimuli (high 208 and low value) of equal size for each participant, with a mean of 38.48 ± 5.02 (mean \pm SD) 209 items remaining in each condition.

210 2.6. ERP analysis

EEG data were pre-processed using BESA v. 6.0 program (MEGIS GmbH, Munich, Germany). Oculographic artefacts and electrocardiographic artefacts were removed using principle component analysis based on averaged eye-blinks and artefact topographies (Berg and Scherg 1994). Data were also visually inspected for the presence of atypical electrode artefacts due to muscle movement. Data were filtered from 1-45 Hz and epochs contaminated with artefacts were excluded manually.

ERPs in response to stimulus presentation were computed separately for each level within conditions (High Value Item & Desirability Rating; High Value Item & Material Estimation; Low Value Item & Desirability Rating; Low Value Item & Material Estimation) by averaging respective epochs in the intervals ranging from 300 ms before image onset to 1000 ms following image onset. Epochs were baseline corrected using a time window of -300

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- to 0 ms relative to stimulus onset. The mean number of accepted trials in each condition
- (following the median split and artefact rejection) was 32.4 ± 5.8 (mean \pm SD).
- 224 2.7. Source dipole reconstruction

225 Grand average potentials were computed by combining all conditions. The grand 226 average waveform was used to define a source dipole model in BESA v. 6.0 program. Using 227 a sequential strategy (Hoechstetter et al. 2001; Stancak et al. 2002), Equivalent Current 228 Dipoles (ECDs) were fitted to describe the 3-dimensional source currents in the regions 229 contributing predominantly to the data (Scherg and Von Cramon 1986). Six ECDs were 230 consecutively seeded until the source mode explained 91.6% of the variance. This amount of 231 explained variance is comparable to previous ERP source dipole localisation studies 232 (Hämäläinen et al. 1993; Schlereth et al. 2003; Stancak et al. 2012; Vrana et al. 2005), and 233 suggests that the six-dipole model explained all major ERP components. Classical LORETA 234 analysis recursively applied (CLARA) method, which is an iterative application of the LORETA algorithm (Pascual-Marqui et al. 1994), was used as an independent source 235 236 localisation method to confirm the locations of the ECDs (Wright et al. 2015). The 237 orientations of ECDs were fitted with the constraint of fixed dipole locations and determined 238 at the maximum of the source strength. A 4-shell ellipsoid head volume conductor model was employed, using the following conductivities (S/m = Siemens per meter): brain = 0.33 S/m; 239 240 scalp = 0.33 S/m; bone = 0.0042 S/m; cerebrospinal fluid = 1 S/m. 241 Source waveforms for each condition were exported and analysed using the EEGLab toolbox (Delorme and Makeig 2004). Due to the large number of statistical tests that this 242 243 requires, P values were corrected using permutation-based repeated-measures ANOVA 244 utilising 5000 permutations (Maris and Oostenveld 2007). For each latency identified, mean

activation over a 10 ms period was calculated, centred on the peak of the observed effect and

for each participant. The data were exported to SPSS Statistics version 22.0 (IBM Corp,
2013) for further analysis.

It is important to note the limitations of source analysis techniques due to the inverse 248 249 problem manifesting in the possibility to generate a number of plausible source dipole models (Michel and Murray 2012). Therefore, a priori information, such as constraining the source 250 251 dipole locations to the cortical mantle, has been implemented in source dipole localisation methods to reduce the number of possible solutions (Michel et al. 2004). To build a plausible 252 253 source dipole model, we applied two different source dipole modelling methods. Firstly, the 254 sequential method consisting of fitting equivalent current dipoles sequentially, and secondly, 255 a distributed source dipole modelling method (CLARA). Both methods yielded highly 256 convergent source dipole models which mitigates but does not completely overcome the 257 limitations associated with the large number of potential source dipole solutions given the 258 mathematical features of the inverse problem

259 **3. Results**

260 3.1. Behavioural data

261 The high value items had a mean WTP of 2.1 ± 0.87 (mean \pm SD) and desirability rating of 50.4 \pm 29.7, whereas the low value items had a mean WTP of 0.66 \pm 0.62 and 262 desirability rating of 27 ± 25.3 . To ensure this finding was not confounded by individual 263 264 differences, a regression model for each participant was created with WTP as a predictor and desirability as a dependent variable. This produced a mean unstandardized coefficient of 15.5 265 266 \pm 9.37; a one-sample t-test revealed this to be significantly different from zero, t(24) = 8.27, P < .001. A mean adjusted R² of 0.23 \pm 0 .17 (mean \pm SD) was also found across subjects. 267 Therefore, desirability of objects was linearly related to WTP (see Figure 2). 268

269 *3.2. Source dipole model*

270 Figure 3 illustrates the ERPs at each electrode site in response to stimulus 271 presentation across all conditions in the form of a butterfly plot; ERP components and their 272 corresponding latencies and topographies are labelled. Four distinct ERP components were 273 observed across the epoch beginning with the visually evoked P1 component peaking at 99 ms, a component related to the early processing of visual stimuli (Hopf et al. 2002) and 274 275 characterised by the strong positivity over the central occipital electrodes with reversed 276 polarity over the frontal electrodes. A P2 component peaked at 209 ms with bilateral 277 positivity over the occipital electrodes but with negativity restricted over a frontal region on 278 the right side of the head (Freunberger et al. 2007; Luck 2005). Although clearly overlapping 279 with the P2, the N2 component peaking at 243 ms can be differentiated by the additional 280 negativity over a frontal region (Folstein and Van Petten 2008). The P3 component (Polich 281 2007) emerges at approximately 316 ms in a parietal region on the right side of the scalp, 282 before reaching a positive maximum at 354 ms over the midline frontal electrodes. 283 Figure 4A shows the source waveforms and the appropriate topographic maps for 284 different ECDs and Figure 4B illustrates the spatial localisation of the ECDs. ECD 1 was located in the right lingual gyrus (Brodmann area 18; approximate Talairach coordinates: x =285 18 mm, y = -59 mm, z = 9 mm) with a peak latency at 95 ms and again at 121ms. ECD 2 286 showed similar characteristics being located in the left lingual gyrus (Brodmann area 18; x = -287 17 mm, y = -59 mm, z = 9 mm) with a peak latency at both 100 ms and 215 ms. Both ECD 1 288 289 and 2 showed a positive maximum over the medial occipital electrodes and a negative 290 potential over a frontal region of the scalp. The latency and the topographical pattern indicate 291 that these two sources were equivalent to the visual P1 component. ECD 3 was located in the

right anterior insula cortex (Brodmann area 13; x = 32 mm, y = 15 mm, z = 0 mm), peaking at

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293 233 ms and showing maximum negativity over a frontal region on the right side of the scalp.

294 This spatial map corresponds to the frontal portion of the N2 component. ECD 4 was located

295 in the left orbitofrontal cortex (Brodmann area 11; x = -26 mm, y = 34 mm, z = -2 mm) 296 showing a small peak at 230 ms. ECD 4 projected positivity over a frontal region localised marginally on the left side. However, this was masked by the N2 component. ECD 5 was 297 298 located in the right parahippocampal gyrus (Brodmann area 28; x = 19 mm, y = -17 mm, z = -17299 21 mm), showing two peak latencies of 215 ms and 316 ms corresponding to both the P2 and 300 the early P3 component. ECD 5 accounted for positivity over a posterior region, localised primarily on the right side of the scalp. ECD 6 was fitted in the posterior cingulate cortex 301 302 (bordering closely with the anterior cingulate cortex; Brodmann area 31; x = 3 mm, y = -18303 mm, z = 42 mm). The source peaked at 248 ms and 431 ms with negativity being distributed 304 across a frontal region of the scalp at 248 ms (contributing to the N2 component at the vertex) 305 and positivity at 431 ms. The final source dipole model accounted for 91.6% of the total 306 variance. CLARA method was used to verify the origins of the fitted ECDs. A mean 307 discrepancy of approximately 15 mm was found between the location of each ECD and the 308 maxima of the nearest cluster.

309 3.3. Effects of rating task and WTP

310 To test the effect of rating task and value on ERPs, a two-way ANOVA for repeated 311 measures was carried out over the latency interval ranging from -200 ms to 450 ms using 312 permutation analysis (Maris and Oostenveld 2007) with 5000 permutations. The F value 313 waveforms were masked inclusively to highlight significant latencies that extended beyond 314 three standard deviations of the source's mean baseline amplitude. Figure 4 shows the 315 topographies at the peak significance of each observed main effect with the corresponding 316 source waveform. Activity over a 10 ms interval centred on the peak significance for each 317 effect (indicated by the shaded region on the source waveform) was exported for further analysis. Table 2A and 2B summarise the mean amplitude and test statistics for each 318

condition over the stated time interval for the main effects of rating task (desirability vs.
material) and value (high vs. low); significant interactions are highlighted in Table 2C.

321 Figure 5A indicates three significant main effects of rating task on the activity from 322 ECD 2, 3 and 5. The waveforms for these ECDs all demonstrate larger activation for 323 desirability ratings than for material estimation ratings. Figure 5B illustrates the two 324 significant main effects of value on the activity from ECD 3 and 4, each displaying greater 325 activation for low value items. Despite the main effect of value at 233ms in ECD 4, it is 326 important to note the difficulty in discerning the differences on scalp topographies due to the 327 dominance of the negativity originating from ECD 3 which peaked at approximately the same 328 time. Only one significant interaction between rating task and value was observed (ECD 6) 329 which is visualised in Figure 5C. During the value rating condition, source activation for a 330 desirability rating of a high value item was higher than in other conditions. Pairwise 331 comparisons indicate that this activation was significantly stronger than during the material 332 estimation and high value condition, t(24) = 2.23, P = 0.035, and also the desirability rating 333 and low value condition, t(24) = 2.1, P = 0.046, but not the material estimation and low value condition, t(24) = 0.65, P = 0.524. No other significant differences were found (P > 0.05). 334

335 A possible explanation for this interaction could be a result of task-switching. For 336 example, upon presentation of a high value item, participants would need to suppress their 337 response if the task required material estimation with a low composition of the given 338 material, with the same going for a low value item in the material estimation task in which 339 composition was high. To test this, a regression model was produced for each subject with desirability as the independent variable and material composition as the dependent variable. 340 This produced a mean unstandardized coefficient of -0.063 which was not significantly 341 different from zero, t(24) = -1.51, P = .145, thus suggesting that task-switching does not 342 adequately explain the interaction effect in PCC. 343

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344 **4. Discussion**

345 This study explored the cortical representation of value by comparing items associated with high or low WTP, and recorded ERPs during passive viewing of items in two 346 347 different valuation contexts, allowing us to disentangle the automatic and the elaborate and 348 conscious valuation processes. Results showed increased cortical activity following the 349 presentation of low value stimuli at the latency of approximately 200 ms, corresponding to 350 the N2 and P2 components of ERPs. Although multiple sources contributed to ERP data at 351 this latency, the economic value of items only modulated the activation in the right AIC and 352 the left OFC. The effects of valuation context were seen in the left LG, right AIC and right 353 PHG.

354 Modulation of source activity within the right AIC peaked at 200 ms, and activity was 355 the strongest for rating of low value items. Although overlapping with the P2 component, 356 source dipole orientation and topographical differences in the negativity over the forehead 357 indicated that the N2 component that demonstrated an effect of value was distinct from the 358 P2 component. The N2 potential was previously reported as being related to aspects of 359 attentional selection (Codispoti et al. 2006; Näätänen and Picton 1986; Patel and Azzam 360 2005), or emotional content of visual stimuli (Olofsson and Polich 2007). The anterior N2 component has been related more specifically to novelty detection and cognitive control 361 362 (Folstein and Van Petten 2008). The present study shows that the right AIC, a region known 363 to be involved together with the OFC and amygdala in loss aversion (Canessa et al. 2017; Canessa et al. 2013; Markett et al. 2016; Tom et al. 2007), contributed to effects of economic 364 value on the amplitude of the N2 component. Therefore, it is possible that the bias towards 365 366 low value items reflects a loss averse response as low value items could represent possible sources of financial loss. However, without more experimental control, it is difficult to 367 368 speculate on the underlying cognitive processes.

369 The low-value bias seen in the N2 component might have been boosted in the present 370 study by the relatively limited range of value among the items on offer. Bartra et al. (2013) report a quadratic pattern within the AIC showing increased BOLD signal in response to 371 372 extreme outcomes, positive or negative, and decreased BOLD for neutral stimuli. With a 373 relatively small range of values in the current study (£0 - £4), the low value items may well 374 have been negatively encoded (high arousal). In contrast, the high-value items may not have 375 passed a threshold in order to be perceived as truly rewarding thus eliciting no arousal 376 response.

377 A similar low-value bias was also seen in left OFC at a latency of 233 ms; despite 378 falling within the N2 component latency, this effect was characterised by increased positivity 379 over the left frontal region but masked by the negativity of the N2. The modulation of source 380 activity for this ECD by stimulus value exhibits an automatic valuation, independent of the 381 valuation context. Modulation of BOLD signal by subjective value has been observed 382 frequently, often within the OFC (Clithero and Rangel 2014). Interestingly, this modulation 383 has been observed for various paradigms utilising several measures of value such as 384 hedonicity ratings (Grabenhorst and Rolls 2009; Lebreton et al. 2009), binary choice tasks 385 (FitzGerald et al. 2009) and importantly, BDM auctions (Plassmann et al. 2010; 2007). The same modulation is also found for multiple reward types and across multiple stages of the 386 387 decision making process (for a review, see Peters and Buchel 2010). Further to this, animal 388 research utilising electrophysiological methods have highlighted the encoding of subjective 389 value within the OFC (Padoa-Schioppa 2013; Padoa-Schioppa and Assad 2006). Similar 390 conclusions have been drawn regarding the vmPFC (Bartra et al. 2013; Clithero and Rangel 391 2014), however given the limitations to spatial resolution that EEG presents, the current 392 findings may not differentiate the activation of the OFC from the neighbouring vmPFC. The 393 emergence of value-based signals in electrophysiological animal research has been observed

394 in OFC at latencies as early as 150 ms (Padoa-Schioppa 2013). Thus, formation of subjective 395 value occurs automatically at an early stage and aids subsequent decision, regardless of whether this signal is an accurate depiction of the ultimate value assigned to the stimulus 396 397 after further deliberation. However, given the task order in the current study, it is important to 398 recognise the potential role that memory may have played in producing this automatic 399 valuation. The auction task always preceded the stimulus rating task resulting in participants 400 having already reported their valuation of each of the stimuli. Although the time between the 401 two tasks was between two and five days, it is possible that the valuation of the stimuli prior 402 to the EEG task may have contributed to this finding due to memories originating from the 403 auction task.

404 The cortical activity in the 200 ms latency range was also modulated by the valuation 405 context. Given that the only computational difference between the two rating tasks is the 406 presence of valuation, any differences in ERPs between the two contexts likely represent the 407 cortical responses associated with attribution of value. The first modulation by the context 408 was observed within the latency of the P2 component at 177 ms; the source activity in the LG 409 was stronger when subjects focused on desirability of items, rather than the material 410 compositions. It has been suggested that the P2 is involved in working memory processes 411 (Finnigan et al. 2011; Lefebvre et al. 2005; Taylor et al. 1990; Wolach and Pratt 2001), visual 412 feature recognition (Hillyard and Münte 1984), and attention allocation (Martin-Loeches et 413 al. 1997). Federmeier and Kutas (2002) reported context-dependent modulations of the P2 in 414 the left hemisphere which finding accords the present study. 415 An effect of the valuation context was also observed in the P2 component at a slightly

An effect of the valuation context was also observed in the P2 component at a slightly later latency of 209 ms. This modulation was related to an increase in source activity in right PHG when evaluating the desirability of items compared to evaluating materials. Given the role of the PHG in memory processes (Aminoff et al. 2013), it is likely here that focusing on the desirability of a stimulus has elicited working memory processes to a greater extent, or required a greater magnitude of attentional allocation. This may be due to the more complex analysis required to reach a decision about value rather than a more simple perceptual evaluation. Assuming value-based decisions require an in depth analysis of the stimuli, in contrast to the perceptual decision requiring estimation of a single material, this modulation may simply be a result of visual feature recognition regarding multiple aspects of the stimuli (Hillyard and Munte 1984).

Finally, the right AIC also showed an increased source activity for the rating of 426 427 desirability resulting in greater negativity over the right forehead. Augmentation of anterior N2 components have been attributed to attentional processes (Codispoti et al. 2006; Näätänen 428 429 and Picton 1986; Patel and Azzam 2005) and it seems the differing computational demands 430 of the value-based and perceptual decisions augmented the observed N2 in the current study. 431 The additional requirement of value-computation for the value-based decision could be the contributing factor to this increased amplitude. Indeed, Näätänen and Picton (1986) highlight 432 433 that the N2 component can be modulated by conscious processing of stimuli, and thus, this 434 processing may well be value specific.

A final modulation of ERPs by the valuation context was observed at approximately 429 ms in PCC. The source activity in PCC, manifested as the negativity potential at vertex electrodes, was prominent for the rating of desirability of high value items, indicating this activation to be specific to highly valued stimuli in an economically relevant context. However, this finding should be interpreted with caution due to the lack of statistically significant differences between the desirability rating of high value items condition, and the material estimation of low value items at the same latency.

442 To conclude, we show that the subjective value of simple household items, measured 443 as WTP in an auction experiment, manifests in ERPs in the latency window and electrodes

444	corresponding to the N2 component. The value-related cortical response, purportedly
445	originating in right AIC and left OFC, is enhanced for low-value items possibly by eliciting
446	loss aversion. The low-value bias in these cortical regions occurred across two different
447	valuation contexts suggesting that this response is a part of an automatic valuation process. In
448	contrast to the subjective value, the valuation context modulates the P2 and N2 components
449	with stronger cortical responses in left LG, right AIC and right PHG occurring whilst subjects
450	focused on desirability than on material aspects of items.
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Figure legends.

471	Figure 1 Experimental protocol. A. Timeline of auction task. A fixation cross was presented
472	at the beginning of each trial for 2 s. Following offset of the fixation cross, an image was
473	presented for 3 s followed by the bidding options for 4 s. A total of nine options were
474	available between £0 and £4 in increments of £0.50. Following the selection of a bid,
475	feedback was presented for 1 s to indicate the outcome of that auction. B. Timeline of EEG
476	task. A fixation cross was presented at the beginning of each trial for 3 s. Next, an instruction
477	was presented for 2 s to indicate the demands of the trial, followed by an image for 3 s.
478	Following image offset, a VAS was presented for 4 s to allow either a desirability rating or
479	material estimation depending on the preceding instruction.
400	
480	Figure 2 Regression lines for each subject predicting desirability from WTP. Grand average
481	regression line is shown in black.
482	Figure 3 Butterfly plot of grand average ERPs in response to stimulus presentation. Distinct
483	ERP components are highlighted with arrows (99, 209, 243, 316 and 354 ms). The
484	topographic map for each ERP component is also displayed.
185	Figure 4 Source dipole model of EDPs. A. Source dipole waveforms in six ECDs. Peak
405	rigure 4 Source upple model of ERIS. A. Source upple wavelorms in six ECDS. I eak
486	latencies and the topographic maps for each of the ECDs are shown. B. Locations and
487	orientations of the six ECDs in the schematic glass brain.
488	Figure 5 Effects of subjective value and context on source dipole waveforms. Each line
489	represents the source dipole waveform for each condition ($D =$ desirability rating; $M =$
490	material estimation; $H = high$ value items; $L = low$ value items; desirability of high [HD] and
491	low [LD] value items; material estimation of high [HM] and low [LM] value items). The

492 shaded grey region on the source dipole waveforms indicates a 10 ms latency period in which

493	a main effect or interaction was revealed, centred on the peak significance. Topographic
494	maps for each condition are displayed. A. ECDs demonstrating a main effect of rating task
495	(ECD 2, 3 and 5). B. ECDs demonstrating a main effect of value (ECD 3 and 4). C. ECD
496	demonstrating significant interaction between rating task and value (ECD 6).
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512 REFERENCES

- 513 Aminoff EM, Kveraga K, and Bar M. The role of the parahippocampal cortex in cognition.
- 514 Trends Cogn Sci 17: 379-390, 2013.
- 515 Bartra O, McGuire JT, and Kable JW. The valuation system: a coordinate-based meta-
- analysis of BOLD fMRI experiments examining neural correlates of subjective value.
- 517 *Neuroimage* 76: 412-427, 2013.
- 518 Becker G, Degroot M, and Marschak J. Measuring utility by a single-response sequential
- 519 method. *Behav Sci* 9: 226-232, 1964.
- 520 Berg P, and Scherg M. A multiple source approach to the correction of eye artifacts.
- 521 *Electroencephalogr Clin Neurophysiol* 90: 229-241, 1994.
- 522 Canessa N, Crespi C, Baud-Bovy G, Dodich A, Falini A, Antonellis G, and Cappa SF.
- Neural markers of loss aversion in resting-state brain activity. *Neuroimage* 146: 257-265,
 2017.
- 525 Canessa N, Crespi C, Motterlini M, Baud-Bovy G, Chierchia G, Pantaleo G, Tettamanti
- 526 M, and Cappa SF. The functional and structural neural basis of individual differences in loss
- 527 aversion. J Neurosci 33: 14307-14317, 2013.
- 528 Chib VS, Rangel A, Shimojo S, and O'Doherty JP. Evidence for a common representation
- 529 of decision values for dissimilar goods in human ventromedial prefrontal cortex. J Neurosci
- 530 29: 12315-12320, 2009.
- 531 Clithero JA, and Rangel A. Informatic parcellation of the network involved in the
- 532 computation of subjective value. Soc Cogn Affect Neurosci 9: 1289-1302, 2014.
- 533 Codispoti M, Ferrari V, Junghofer M, and Schupp HT. The categorization of natural
- scenes: brain attention networks revealed by dense sensor ERPs. *Neuroimage* 32: 583-591,
- 535 2006.

536	Delorme A, and Makeig S. EEGLAB: an open source toolbox for analysis of single-trial
537	EEG dynamics including independent component analysis. J Neurosci Methods 134: 9-21,
538	2004.
539	Delplanque S, Silvert L, Hot P, Rigoulot S, and Sequeira H. Arousal and valence effects
540	on event-related P3a and P3b during emotional categorization. Int J Psychophysiol 60: 315-
541	322, 2006.
542	Federmeier K, and Kutas M. Picture the difference: electrophysiological investigations of
543	picture processing in the two cerebral hemispheres. Neuropsychologia 40: 730-747, 2002.
544	Ferree TC, Luu P, Russell GS, and Tucker DM. Scalp electrode impedance, infection risk,
545	and EEG data quality. Clinicial Neurophysiology 112: 536-544, 2001.
546	Finnigan S, O'Connell RG, Cummins TD, Broughton M, and Robertson IH. ERP
547	measures indicate both attention and working memory encoding decrements in aging.
548	Psychophysiology 48: 601-611, 2011.
549	FitzGerald TH, Seymour B, and Dolan RJ. The role of human orbitofrontal cortex in value
550	comparison for incommensurable objects. J Neurosci 29: 8388-8395, 2009.
551	Folstein JR, and Van Petten C. Influence of cognitive control and mismatch on the N2
552	component of the ERP: A review. Psychophysiology 45: 152-170, 2008.
553	Foti D, Hajcak G, and Dien J. Differentiating neural responses to emotional pictures:
554	Evidence from temporal-spatial PCA. Psychophysiology 46: 521-530, 2009.
555	Freunberger R, Klimesch W, Doppelmayr M, and Holler Y. Visual P2 component is
556	related to theta phase-locking. Neurosci Lett 426: 181-186, 2007.
557	Gehring W, Liu Y, Orr JM, and Carp J. The error-related negativity (ERN/Ne). In: The
558	Oxford Handbook of Event-Related Potential Components, edited by Luck SJ, and
559	Kappenman ES2012.

- 560 Gluth S, Rieskamp J, and Buchel C. Classic EEG motor potentials track the emergence of
- value-based decisions. *Neuroimage* 79: 394-403, 2013.
- 562 Grabenhorst F, and Rolls ET. Different representations of relative and absolute subjective
- value in the human brain. *Neuroimage* 48: 258-268, 2009.
- 564 Grueschow M, Polania R, Hare TA, and Ruff CC. Automatic versus Choice-Dependent
- 565 Value Representations in the Human Brain. *Neuron* 85: 874-885, 2015.
- 566 Hajcak G, Weinberg A, MacNamara A, and Foti D. ERPs and the study of emotion. In:
- 567 The Oxford Handbook of Event-Related Potential Components, edited by Luck SJ, and
- 568 Kappenman ES. New York: Oxford University Press, 2012, p. 441-472.
- 569 Hämäläinen M, Hari R, Ilmoniemi RJ, Knuutila J, and Lounasmaa OV.
- 570 Magnetoencephalography—theory, instrumentation, and applications to noninvasive studies
- 571 of the working human brain. *Reviews of Modern Physics* 65: 413-497, 1993.
- 572 Harris A, Adolphs R, Camerer C, and Rangel A. Dynamic construction of stimulus values
- 573 in the ventromedial prefrontal cortex. *PLoS One* 6: e21074, 2011.
- 574 Hillyard SA, and Munte TF. Selective attention to color and location: an analysis with
- 575 event-related brain potentials. *Percept Psychophys* 36: 185-198, 1984.
- 576 Hoechstetter K, Rupp A, Stancak A, Meinck HM, Stippich C, Berg P, and Scherg M.
- 577 Interaction of tactile input in the human primary and secondary somatosensory cortex--a
- 578 magnetoencephalographic study. *Neuroimage* 14: 759-767, 2001.
- 579 Hopf J, Vogel E, Woodman G, Heinze H, and Luck SJ. Localizing visual discrimination
- 580 processes in time and space. *J Neurophysiol* 88: 2088-2095, 2002.
- 581 Huang YX, and Luo YJ. Temporal course of emotional negativity bias: an ERP study.
- 582 Neurosci Lett 398: 91-96, 2006.

583 Hunt LT, Kolling N, Soltani A, Woolrich MW, Rushworth MF, and Behrens TE.

- Mechanisms underlying cortical activity during value-guided choice. *Nat Neurosci* 15: 470476, \$471-473, 2012.
- 586 Larsen T, and O'Doherty JP. Uncovering the spatio-temporal dynamics of value-based
- 587 decision-making in the human brain: a combined fMRI-EEG study. *Philos Trans R Soc Lond*
- 588 *B Biol Sci* 369: 2014.
- 589 Lebreton M, Jorge S, Michel V, Thirion B, and Pessiglione M. An automatic valuation
- 590 system in the human brain: evidence from functional neuroimaging. *Neuron* 64: 431-439,
- 591 2009.
- 592 Lefebvre CD, Marchand Y, Eskes GA, and Connolly JF. Assessment of working memory
- bilities using an event-related brain potential (ERP)-compatible digit span backward task.
- *Clin Neurophysiol* 116: 1665-1680, 2005.
- 595 Levy I, Lazzaro SC, Rutledge RB, and Glimcher PW. Choice from non-choice: predicting
- 596 consumer preferences from blood oxygenation level-dependent signals obtained during
- 597 passive viewing. J Neurosci 31: 118-125, 2011.
- 598 Luck SJ. An introduction to the event-related potential technique. Cambridge, Mass: MIT
 599 Press, 2005.
- 600 Luu P, Tucker DM, Derryberry D, Reed M, and Poulsen C. Electrophysiological
- 601 responses to errors and feedback. *Psychological Science* 14: 47-53, 2003.
- 602 Macnamara A, Foti D, and Hajcak G. Tell me about it: neural activity elicited by
- 603 emotional pictures and preceding descriptions. *Emotion* 9: 531-543, 2009.
- 604 Maris E, and Oostenveld R. Nonparametric statistical testing of EEG- and MEG-data. J
- 605 Neurosci Methods 164: 177-190, 2007.

- 606 Markett S, Heeren G, Montag C, Weber B, and Reuter M. Loss aversion is associated
- with bilateral insula volume. A voxel based morphometry study. *Neurosci Lett* 619: 172-176,
 2016.
- 609 Martin-Loeches M, Schweinberger SR, and Sommer W. The phonological loop model of
- 610 working memory: an ERP study of irrelevant speech and phonological similarity effects.
- 611 *Mem Cognit* 25: 471-483, 1997.
- 612 Michel CM, and Murray MM. Towards the utilization of EEG as a brain imaging tool.
- 613 *Neuroimage* 61: 371-385, 2012.
- 614 Michel CM, Murray MM, Lantz G, Gonzalez S, Spinelli L, and Grave de Peralta R.
- EEG source imaging. *Clin Neurophysiol* 115: 2195-2222, 2004.
- 616 Moser JS, Hajcak G, Bukay E, and Simons RF. Intentional modulation of emotional
- 617 responding to unpleasant pictures: an ERP study. *Psychophysiology* 43: 292-296, 2006.
- 618 Näätänen R, and Picton TW. N2 and automatic versus controlled processes.
- 619 *Electroencephalogr Clin Neurophysiol Suppl* 38: 169-186, 1986.
- 620 Nieuwenhuis S, Holroyd CB, Mol N, and Coles MG. Reinforcement-related brain
- 621 potentials from medial frontal cortex: origins and functional significance. *Neurosci Biobehav*
- 622 *Rev* 28: 441-448, 2004.
- 623 Olofsson K, and Polich J. Affective visual event-related potentials: arousal, repetition, and
- 624 time-on-task. *Biol Psychol* 75: 101-108, 2007.
- 625 Padoa-Schioppa C. Neuronal origins of choice variability in economic decisions. *Neuron*
- 626 80: 1322-1336, 2013.
- 627 Padoa-Schioppa C. Orbitofrontal cortex and the computation of economic value. Ann NY
- 628 Acad Sci 1121: 232-253, 2007.
- 629 Padoa-Schioppa C, and Assad JA. Neurons in the orbitofrontal cortex encode economic
- 630 value. *Nature* 441: 223-226, 2006.

- 631 Pascual-Marqui RD, Michel CM, and Lehmann D. Low resolution electromagnetic
- tomography: a new method for localizing electrical activity in the brain. *Int J Psychophysiol*18: 49-65, 1994.
- 634 Patel SH, and Azzam PN. Characterization of N200 and P300: selected studies of the event-
- related potential. *International Journal of Medical Sciences* 2: 147-154, 2005.
- 636 Payzan-LeNestour E, Dunne S, Bossaerts P, and O'Doherty JP. The neural representation
- of unexpected uncertainty during value-based decision making. *Neuron* 79: 191-201, 2013.
- 638 Peters J, and Buchel C. Neural representations of subjective reward value. *Behav Brain Res*639 213: 135-141, 2010.
- 640 Picton TW, Bentin S, Berg P, Donchin E, Hillyard SA, Johnson J, Miller GA, Ritter W,
- 641 Ruchkin DS, Rugg MD, and Taylor MJ. Guidelines for using human event-related
- potentials to study cognition: Recording standards and publication criteria. *Psychophysiology*37: 127-152, 2000.
- 644 Plassmann H, O'Doherty JP, and Rangel A. Appetitive and aversive goal values are
- 645 encoded in the medial orbitofrontal cortex at the time of decision making. *J Neurosci* 30:
- 646 10799-10808, 2010.
- 647 Plassmann H, O'Doherty JP, and Rangel A. Orbitofrontal cortex encodes willingness to
- 648 pay in everyday economic transactions. *J Neurosci* 27: 9984-9988, 2007.
- 649 Polania R, Krajbich I, Grueschow M, and Ruff CC. Neural oscillations and
- 650 synchronization differentially support evidence accumulation in perceptual and value-based
- 651 decision making. *Neuron* 82: 709-720, 2014.
- 652 **Polich J.** Updating P300: an integrative theory of P3a and P3b. *Clin Neurophysiol* 118: 2128-
- 653 2148, 2007.
- 654 Raghuraman AP, and Padoa-Schioppa C. Integration of multiple determinants in the
- neuronal computation of economic values. J Neurosci 34: 11583-11603, 2014.

- 656 Rolls ET, Grabenhorst F, and Parris BA. Neural systems underlying decisions about
- 657 affective odors. J Cogn Neurosci 22: 1069-1082, 2009.
- 658 San Martin R. Event-related potential studies of outcome processing and feedback-guided
- 659 learning. Front Hum Neurosci 6: 304, 2012.
- 660 Scherg M, and Von Cramon D. Evoked dipole source potentials of the human auditory
- 661 cortex. *Electroencephalogr Clin Neurophysiol* 65: 344-360, 1986.
- 662 Schlereth T, Baumgärtner U, Magerl W, Stoeter P, and Treede R-D. Left-hemisphere
- 663 dominance in early nociceptive processing in the human parasylvian cortex. *Neuroimage* 20:
- 664 441-454, 2003.
- 665 Smith NK, Cacioppo JT, Larsen JT, and Chartrand TL. May I have your attention,
- 666 please: electrocortical responses to positive and negative stimuli. Neuropsychologia 41: 171-
- 667 183, 2003.
- 668 Stancak A, Hoechstetter K, Tintera J, Vrana J, Rachmanova R, Kralik J, and Scherg
- 669 M. Source activity in the human secondary somatosensory cortex depends on the size of
- 670 corpus callosum. *Brain Res* 936: 47-57, 2002.
- 671 Stancak A, Johnstone J, and Fallon N. Effects of motor response expectancy on cortical
- 672 processing of noxious laser stimuli. *Behav Brain Res* 227: 215-223, 2012.
- 673 Taylor MJ, Smith ML, and Iron KS. Event-related potential evidence of sex-differences in
- 674 verbal and nonverbal memory tasks. *Neuropsychologia* 28: 691-705, 1990.
- 675 Tom SM, Fox CR, Trepel C, and Poldrack RA. The neural basis of loss aversion in
- decision-making under risk. *Science* 315: 515-518, 2007.
- 677 Tzovara A, Chavarriaga R, and De Lucia M. Quantifying the time for accurate EEG
- decoding of single value-based decisions. J Neurosci Methods 250: 114-125, 2015.

- 679 Vrana J, Polacek H, and Stancak A. Somatosensory-evoked potentials are influenced
- 680 differently by isometric muscle contraction of stimulated and non-stimulated hand in humans.
- 681 Neurosci Lett 386: 170-175, 2005.
- 682 Walsh MM, and Anderson JR. Learning from experience: event-related potential correlates
- 683 of reward processing, neural adaptation, and behavioral choice. *Neurosci Biobehav Rev* 36:

684 1870-1884, 2012.

- 685 Weinberg A, and Hajcak G. Beyond good and evil: the time-course of neural activity
- elicited by specific picture content. *Emotion* 10: 767-782, 2010.
- 687 Wilkinson N, and Klaes M. An Introduction to Behavioral Economics. Houndsmills,
- 688 Basingstoke: Palgrave Macmillan, 2012.
- 689 Wolach I, and Pratt H. The mode of short-term memory encoding as indicated by event-
- related potentials in a memory scanning task with distractions. *Clin Neurophysiol* 112: 186-197, 2001.
- 692 Wright H, Li X, Fallon NB, Giesbrecht T, Thomas A, Harrold JA, Halford JC, and
- 693 Stancak A. Heightened eating drive and visual food stimuli attenuate central nociceptive
- 694 processing. J Neurophysiol 113: 1323-1333, 2015.
- 695 Yeung N, and Cohen JD. The impact of cognitive deficits on conflict monitoring.
- 696 *Psychological Science* 17: 164-171, 2006.
- 697 Yeung N, and Sanfey AG. Independent coding of reward magnitude and valence in the
- 698 human brain. J Neurosci 24: 6258-6264, 2004.
- 699 Yu R, and Huang Y. A nickel ain't worth a dime anymore: the illusion of money and the
- rapid encoding of its true value. *PLoS One* 8: e55025, 2013.







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Downloaded from www.physiology.org/journal/jn by \${individualUser.givenNames} \${individualUser.surname} (138.253.205.169) on February 23, 2018. Copyright © 2018 American Physiological Society. All rights reserved. A. Mean source amplitude ± SDs for both desirability and material estimation conditions over the stated time interval for each significant latency and the corresponding ECD.
 F and P values for the relevant ANOVA are also displayed.

ECD	Time Interval	Desirability	Material	F ₍₂₄₎	Р
	[ms]				
ECD2	172 – 182	14.2 ± 23.2	9.32 ± 22.3	9.93	0.004
ECD3	201 - 211	19.28 ± 14.51	12.04 ± 12.2	17.6	< 0.001
ECD5	204 - 214	37.26 ± 20.81	27.49 ± 20.09	8.34	0.08

B. Mean source amplitude ± SDs for both high and low value conditions over the stated time interval for each significant latency and the corresponding ECD. F and P values for the relevant ANOVA are also displayed.

ECD	Time Interval	High Value	Low Value	F ₍₂₄₎	Р
	[ms]				
ECD3	195 – 205	10.07 ± 12.85	17.81 ± 15.31	9.19	0.006
ECD4	228 - 238	5.09 ± 8.07	9.36 ± 8.89	12.57	0.002

C. Mean source amplitude for desirability ratings of high value (HD) and low value (LD) items and for material estimation ratings of high value (HM) and low value (LM) items for each significant latency and the corresponding ECD over the stated time interval. F and P values for the relevant ANOVA are also displayed.

ECD	Time Interval	HD	HM	LD	LM	F ₍₂₄₎	Р
	[ms]						
ECD6	424 - 434	7.44 ± 12.76	1.85 ± 10.13	1.47 ± 10.32	5.05 ± 15.2	8.25	0.008