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1 **Abstract**

2 Touch has been shown to regulate emotions, stress responses, and physical pain. However, its
3 impact on cognitive functions, such as inhibitory control, remains relatively understudied. In this
4 experiment, we explored the effects of low-force, slow-moving touch—designed to optimally activate
5 unmyelinated cutaneous low-threshold mechanoreceptor C-tactile (CT) afferents in human hairy skin—
6 on inhibitory control and its psychophysiological correlates using the Stroop Task, a classic paradigm
7 commonly employed to assess inhibitory control capacity. The Stroop Task was repeated twice before
8 and once after receiving either gentle touch or no-touch. Participants were assigned to two groups: the
9 touch group (n = 36), which received low-force, slow-moving touch on their forearms at a stroking
10 velocity of ~3 cm/sec, and the no-touch group (n = 36), which did not receive any touch stimulation.
11 Changes in autonomic nervous system activity were also assessed by measuring heart rate variability
12 (HRV) and skin conductance levels before and during cognitive performance. Compared to the no-touch
13 group, participants who received gentle, low-force, slow-moving touch demonstrated faster responses
14 and higher HRV during the Stroop Task. Additionally, within the touch group, individuals with higher
15 HRV exhibited even quicker performance on the cognitive task. While we cannot draw definitive
16 conclusions regarding the CT velocity-specific effect, these results provide preliminary evidence that
17 low-force, slow-moving touch may influence cognitive processes involved in the inhibitory control of
18 goal-irrelevant stimuli.

19 **Keywords:** gentle, low-force, slow-moving touch; autonomic nervous system; HRV; inhibitory control;
20 Stroop Task

21

22 **1. Introduction**

23 Touch plays a crucial role in fostering social interactions (Suvilehto et al., 2023), bonding and

24 attachment (Duhn, 2010; Jablonski, 2021), and human development (Cascio et al., 2019). The
25 identification of a system of unmyelinated cutaneous low-threshold mechanoreceptor (LTMR) C-fibres
26 in human hairy skin has redefined the traditional understanding of touch as being solely discriminative
27 in nature. These C-tactile (CT) afferents, characterised by a preference for low-force, skin temperature,
28 caress-like stroking touch of between 1 and 10 cm/s (Ackerley et al. 2014a, 2014b; Löken et al. 2009),
29 are not well-suited for precise tactile discrimination (see McGlone, Wessberg, & Olausson, 2014 for an
30 extensive review). Psychophysical studies consistently show that participants find this stimulus more
31 pleasant compared to touch delivered at slower or faster velocities (Ackerley et al., 2014b; Essick et al.,
32 1999; Löken et al., 2009). According to the affective touch hypothesis (Morrison & Croy, 2021), these
33 CT afferents have been found to play a key role in conveying touch's pleasant and rewarding properties
34 (Morrison et al., 2010; Löken et al., 2009; Vallbo et al., 1999). It also reduces negative emotions (e.g.,
35 social exclusion; Oya & Tanaka, 2023; von Mohr et al., 2017), buffers physical pain (Gursul et al., 2018;
36 von Mohr et al., 2018), and increases body awareness (Crucianelli et al., 2018; Cazzato et al., 2021;
37 Jenkinson et al., 2020).

38 From a physiological perspective, CT-targeted touch has been shown to regulate stress responses
39 (Kidd et al., 2023; Morrison, 2016; Walker et al., 2022) and autonomic nervous functions (Püschel et
40 al., 2022; Manzotti et al., 2023; Tricoli et al., 2017). For instance, maternal stroking touch has been
41 found to increase heart rate variability (HRV) (Manzotti et al., 2023; Van Puyvelde et al., 2019). HRV,
42 i.e., the beat-to-beat changes in heart rate, is an indirect, well-validated vagal tone index (Laborde et al.,
43 2017). Higher levels of resting HRV, indicating increased activity of the parasympathetic nervous
44 system (Berntson et al., 1997; Kop et al., 2011), are linked to improved emotional and behavioural
45 regulation (Balzarotti et al., 2017; Cai et al., 2019; Mather & Thayer, 2018), as well as enhanced overall
46 mental and physical wellbeing (Cai et al., 2019; Kemp & Quintana, 2013; Sloan et al., 2017). Changes
47 in HRV are thought to be pivotal in maternal-infant physiological and behavioural regulation and

48 resilience (Poehlmann et al., 2011; Porter, 2003; Suga et al., 2019). On the other hand, low levels of
49 resting HRV have been associated with a range of mental health conditions, including anxiety (e.g.,
50 Chalmers et al., 2014; Thayer et al., 1996; Kemp et al., 2014), panic disorder (e.g., McCraty et al., 2001),
51 post-traumatic stress disorder (Cohen et al., 1998), depression (e.g., Hartmann et al., 2019; Nahshoni et
52 al., 2004), and suicide ideation and behaviour (Adolph et al., 2018).

53 While most existing research has predominantly focused on affective touch as a source of affect
54 regulation (Fotopoulou et al., 2022; Silvestri et al., 2024), less attention has been given to its potential
55 effects on cognitive processes, exploring the bottom-up influence of touch on top-down mechanisms.
56 According to the “embodied cognition” framework (Gallese & Ebisch, 2013; Wilson & Golonka, 2013),
57 bodily experiences—particularly tactile sensations—play a crucial role in shaping and influencing our
58 cognitive functions. As such, touch is not merely a passive experience but an active process that
59 integrates with and affects cognitive mechanisms.

60 To date, only a few studies have focussed on how interpersonal touch affects the neurocognitive
61 processes that underlie flexible goal-directed behaviour involved in cognitive control (Dydenkova et al.,
62 2024; Saunders et al., 2018). In particular, the study by Saunders and colleagues (2018) recruited
63 romantic partners, with the active partner performing a speeded inhibitory control task modified version
64 of a Go-no-Go Task while either holding (touch condition) or not holding their partner’s hand (no-touch
65 condition), whilst Electroencephalography (EEG) activity was also recorded throughout. The results
66 demonstrated that touch (handholding) enhanced cognitive control, as evidenced by reduced error rate
67 on the task and increased error-related negativity amplitudes, which reflect the neural response to
68 recognising mistakes and potentially triggering cognitive control mechanisms to correct or adjust
69 behaviour. Additionally, holding the partner’s hand elicited positive emotional responses, including
70 increased happiness, suggesting that interpersonal touch can enhance cognitive control through
71 modulation of emotional and neural mechanisms. A possible explanation for these findings is that human

72 proximity can enhance personal efficacy (Coan & Sbarra, 2015), helping individuals reduce their
73 tendency to ignore or minimise negative feedback signals (e.g., error monitoring), which may, in turn,
74 lead to exert inhibitory control over interference. While the study's findings suggest a potential link
75 between touch and the cognitive/neural monitoring processes underlying flexible goal-directed
76 behaviour, several issues might limit the conclusions of this investigation. The (handholding)
77 interpersonal touch manipulation used in the study by Saunders and colleagues (2018) could not
78 disentangle the specific effects of social (interpersonal proximity and interaction) versus affective
79 (pleasant) touch on cognitive control. Additionally, it cannot determine whether changes in autonomic
80 nervous system (ANS) activity may mediate psychophysiological regulation of inhibitory control. In
81 light of this, we adopted a touch condition involving gentle, low-force, slow-moving touch to the skin
82 specifically designed to activate CT afferents, which are thought to regulate stress response in rats
83 (Walker et al. 2022) and in certain individuals (Kidd et al., 2023; Morrison, 2016) as well as a more
84 controlled method (Löken et al., 2009; Wijaya et al., 2020). Furthermore, to mitigate potential order
85 effects associated with a within-subject design (as used by Saunders and colleagues, 2018), we chose to
86 employ a between-subjects design to compare low-force, slow-moving touch with no-touch conditions.
87 Importantly, our study aimed to explore whether and how interpersonal touch enhances cognitive control
88 via emotional regulation. Specifically, we sought to account for the potential role of vagal activity in
89 supporting response inhibition, as highlighted in prior research (e.g., Thayer & Lane, 2006). As an
90 important hallmark of executive functions, primarily regulated by the prefrontal regions of the brain,
91 inhibitory control refers to the capacity to suppress automatic responses and irrelevant information (Bari
92 & Robbins, 2013; Cristofori et al., 2019; Grafman, 2002). According to the Neurovisceral Integration
93 Model (NIM; Thayer et al., 2009a; Thayer & Lane, 2000), prefrontal cortex engagement during
94 inhibitory control is crucially associated with vagally-mediated high-HRV (parasympathetic activity) and
95 reduced sympathetic activation. Research highlights the significance of high-frequency (HF) HRV as an

96 index of parasympathetic activity in assessing the autonomic regulation linked to demanding tasks (Forte
97 et al., 2019; Forte & Casagrande, 2025). HF-HRV is particularly valuable because it is sensitive to short-
98 term fluctuations in autonomic tone, making it highly responsive to potentially stressful stimuli that
99 require rapid autonomic adjustments (Thayer & Lane, 2000). Conversely, heightened sympathetic
100 activation, as indicated by galvanic skin response (Kim et al., 2023), appears to result from lower
101 prefrontal cortex activation and impaired cognitive control mechanisms (Boberg et al., 2022; Clark et al.,
102 2018). This leads to disinhibition and altered cognitive performance (Thayer & Lane, 2000). Hence, the
103 ANS activity, as indexed by increased vagal tone, is proposed to reflect attentional regulation and overall
104 adaptive and flexible behavioural strategies in response to high-cognitive tasks or demands (Colzato et
105 al., 2017; Grol & De Raedt, 2020; Hovland et al., 2012; Park & Thayer et al., 2014; Thayer & Lane,
106 2000). These findings are further supported by studies showing that autonomic reactivity, particularly as
107 indicated by changes in HF-HRV in healthy adults, is heightened during demanding tasks measuring
108 inhibition (e.g., Stroop Task; Stroop, 1935) or executive functioning, thus confirming a strong connection
109 between ANS function and cognitive performance (Forte et al., 2019; Forte & Casagrande, 2025; Huang
110 et al., 2021; Renaud & Blondin, 1997; Thayer et al., 2009). Therefore, an outstanding research question
111 is whether the ability to inhibit a response can be influenced by manipulating the ANS activity through
112 gentle, low-force, slow-moving touch. Most touch-based interventions have been found to benefit mental
113 and physical health (Alp et al., 2021; McGlone et al., 2024). However, the specific impact of gentle, low-
114 force, slow-moving touch on autonomic regulation during cognitive inhibition is poorly understood.

115 This study investigated whether gentle, low-force, slow-moving touch, specifically through
116 stimulation designed to activate CT-targeted touch preferentially, could enhance inhibitory control
117 during a Stroop task. The Stroop Task is a standard test that measures participants' abilities to suppress
118 cognitive interference and to examine the efficiency of attentional control, processing speed, and overall
119 executive processing abilities. During the Stroop Task, the capacity to overcome reaction conflict caused

120 by the intentional suppression of irrelevant and incompatible information may elicit physiological stress
121 that can involve the sympathetic nervous system (responsible for fight or flight response) and the
122 parasympathetic nervous system (responsible for recovery and rest) (Hoshikawa & Yamamoto, 1997;
123 Mathewson et al., 2010; Vazan et al., 2017; Waxenbaum et al., 2023).

124 Importantly, in this study, participants completed the Stroop Task whilst indexes of the
125 sympathetic and parasympathetic activity, including Electrodermal Activity (EDA) and
126 Electrocardiogram (ECG), were collected to measure Skin Conductance Level (SCL) and HRV for HF-
127 HRV power, respectively. Physiological indexes were obtained before and after receiving gentle, low-
128 force, slow-moving touch or without receiving any touch at all. We expected that participants who
129 received gentle, low-force, slow-moving touch stimulation would perform better on the Stroop Task than
130 those who did not receive any touch stimulation (Saunders et al., 2018). Accordingly, touch stimulation
131 might modulate participants' physiological states (Mazza et al., 2023; Pawling et al., 2024; Triscoli et
132 al., 2017), aiding in the implementation of flexible and adaptive control over conflicting information
133 during prefrontal task performance (Thayer et al., 2009a). In agreement with the NIM model (Thayer &
134 Lane, 2000, 2009a,2009b), and following touch stimulation, we also anticipated increased HF-HRV
135 levels (parasympathetic activity) during Stroop Task performance compared to SCL (sympathetic
136 activity).

137 **2. Methods**

138 **2.1 Participants**

139 The sample size calculation was determined using G*Power 3.0.10 (Faul et al, 2007) based on the
140 outcome measures of RTs and accuracy. Calculations indicated a minimum of 27 participants per group
141 (touch vs no-touch) and Time (pre- vs post-manipulation) for a small effect size ($f^2 = 0.25$), with 95%
142 power and an α level set at 0.05, using a mixed design. A total of 72 participants took part in this study,
143 with 36 adults (23 females, mean age = 42.78yrs, $SD = 21.90$) assigned to the touch group and 36 adults

144 to the no-touch group (22 females, mean age = 45.03yrs, $SD = 21.65$). Participants were recruited from
145 external sources, including poster advertisements in public places, social media, and personal contacts
146 of the researcher, as well as internally through the Liverpool John Moores University (LJMU)
147 Psychology SONA system. Participants were free of neurological diseases and psychiatric disorders,
148 skin conditions or nerve impairment, and visual-perception disorders (e.g., colour blindness). The study
149 was carried out in accordance with the Helsinki Declaration of ethical standards. The study protocol was
150 approved by the LJMU's University Research Ethics Committee (UREC, 22/PSY/019). All participants
151 gave their written informed consent to take part in the study. Participants were rewarded with a £5
152 shopping voucher or SONA credits if they were LJMU students.

153 **2.2 General procedure**

154 A schematic representation of the general procedure is presented in Figure 1.
155 On the day of testing, participants gave written consent and were asked to fill out a questionnaire
156 concerning demographic details (i.e., gender, age, education), the Positive and Negative Affect Schedule
157 (PANAS; Watson, 1988) for rating positive and negative emotions, and the Depression anxiety stress
158 Scale-21 (DASS-21; Lovibond & Lovibond, 1995) to provide a measure of anxiety, depression, and
159 stress levels. Then, all participants were asked to perform the Stroop Task at Time 1 (T1). At Time 2
160 (T2), the touch group received gentle, low-force, slow-moving touch stimulation delivered at a velocity
161 of ~3 cm/sec—a speed typically perceived as pleasant and optimal for activating the CT system (Löken
162 et al., 2009)—before performing the Stroop Task for the second time. The interval between the two
163 times was about 7 minutes, consistently maintained across groups and participants. After receiving
164 manual stroking through a cosmetic soft brush applied over their ventral forearm, participants were
165 required to report their pleasantness on a Visual Analogue Scale (VAS, e.g., Bellard et al., 2023;
166 Sacchetti et al, 2021). Participants assigned to the no-touch group underwent the same procedure except
167 for the touch stimulation. Participants in the no-touch group were instructed to remain quietly without

168 being engaged in stimulating activities to prevent any sensory/affective input that could potentially
169 influence the Stroop Task performance for a time equal to that of the participants receiving touch
170 stimulation. In this case, the experimenter maintained a non-intrusive presence, staying two metres away
171 from the participant to minimise engagement and prevent heightened arousal. In the touch condition that
172 closely mirrored this setup, participants were invited to remain still, calm, and away from the tactile
173 stimulation. We implemented a standardised interaction script for the experimenter during the touch
174 stimulation. This script reduced variability in non-verbal cues, such as body language and tone of voice,
175 ensuring that every participant experienced the same level of engagement. Additionally, both groups
176 were exposed to the same ambient lighting and room temperature settings to avoid sensory differences
177 that could influence arousal levels. All participants were randomly assigned to either the touch or no-
178 touch condition to ensure that any physiological and cognitive differences observed were attributable to
179 touch rather than pre-existing differences between participants. Participants were informed in the
180 participant information sheet that they might receive touch during the experiment, although the timing
181 was not specified. On the testing day, participants were informed about their group allocation (whether
182 they would receive touch or not) after the first Stroop Task (T1) and just before they performed the task
183 again (T2) to minimise biases and anticipatory effects that might arise from knowing about the touch
184 stimulation.

185 EDA and ECG signals were measured throughout the experiment to evaluate sympathetic and
186 parasympathetic activity, respectively. During this time, participants were instructed to maintain regular
187 breathing and minimise body movements during the physiological recording before performing the task.
188 At the end of the experiment, they were asked to fill out the PANAS a second time. Overall, the testing
189 procedure lasted approximately 45 minutes.

190

191 -----Please Insert Figure 1 about here -----

192 **2.3 Material and measures**

193 **2.3.1 Stroop Task**

194 The colour word Stroop Task (Stroop, 1935) was performed using Millisecond software (Inquisit
195 6; Draine, 1999; <https://www.millisecond.com>). This task measures the ability to inhibit automatic
196 responses by requiring participants to ignore the meaning of a word and focus on naming the colour of
197 the word's ink. In this study, participants were asked to type specific keys corresponding to the colour of
198 the word displayed on the screen [i.e., D = red, F = green, J = blue, and K = yellow]) as quickly and
199 accurately as possible. Each word was displayed until one of the four keys was pressed. The task included
200 84 trials [4 colours × 3 stimuli (congruent, incongruent, control) × 7 repetitions]. This resulted in 28
201 congruent trials (word and colour match), 28 incongruent trials (word and colour do not match), and 28
202 control trials (coloured rectangles), randomly presented (Parkin et al., 2017). Prior to the start of the task,
203 participants were trained with a short practice consisting of 12 practice trials (4 for each trial type). If the
204 response was correct during the experiment, the subsequent trial started immediately. A red X was flashed
205 on the screen if an incorrect response was made. Accuracy was determined by the percentage of correct
206 responses ($\text{Tot correct}/\text{Ntrial}$) with a score of 1 for correct and 0 for incorrect answers. RTs were recorded
207 by measuring the time lapse between the presentation of the stimulus and the participant's response on
208 the keyboard. We calculated the mean latency of congruent or incongruent trials (in milliseconds) to
209 assess RTs for our analyses. Data from the practice and control trials were not included in accuracy and
210 RTs performance counts.

211

212 **2.3.2 Touch stimulation**

213 Participants received manual gentle strokes on the ventral forearm using a soft brush (No7 cosmetic
214 brush, Boots UK; Cazzato et al., 2021; Pawling et al., 2024; Sacchetti et al., 2021) for two minutes (Della

215 Longa et al., 2021; Ree et al., 2019) before performing the Stroop Task a second time. This interval
216 length is sufficient for obtaining accurate measures of physiological arousal (Della Longa et al., 2021;
217 Munoz et al., 2015). The brush was employed for tactile stimulation as materials perceived as soft are
218 typically rated as pleasant (Tarvainen et al., 2014; Wijaya et al., 2020). Following the procedure adopted
219 in a study previously published by our research group, each stroking was applied at a velocity of ~3 cm/s
220 on the ventral forearm (Sacchetti et al., 2021). The rationale for this choice was that this velocity
221 preferentially activates CT afferents, a type of nerve fibre that typically responds to gentle, slow stroking
222 touch (Löken et al., 2009; Olausson et al., 2010; Vallbo et al., 1999), triggers pleasant feelings (Löken
223 et al., 2009; Tricoli et al., 2017) and buffers stress (Morrison, 2016). Accordingly, we delivered 12
224 strokes, each separated by a 6-second interval, in a single session to account for CT-afferents' tendency
225 to fatigue after repeated exposure to tactile stimuli (Schirmer & McGlone, 2022; Vallbo et al., 1999).
226 Strokes were delivered at a constant pressure of 22 gr/cm² on about 9 cm long by a (female) research
227 assistant trained to deliver the strokes on a scale to replicate the same movements on participants'
228 forearms during the experiment. A visual metronome was programmed on PsychoPy (Peirce, 2007) to
229 guide the research assistant in delivering the strokes. During the touch manipulation, participants were
230 asked to look at a blank screen presented on the computer in front of them. After touch manipulation, a
231 VAS was used to evaluate the pleasantness of touch. The VAS consisted of a horizontal line measuring
232 20 cm. Participants were instructed to make a mark on the line using a pen, indicating the level of
233 pleasantness experienced during the touch. The scale ranged from -10 to +10, representing unpleasant,
234 neutral, and pleasant touch.

235

236 **2.3.3 Physiological arousal**

237 A Biopac System, Inc., MP36 was utilized to record electrocardiogram (ECG) signals from which
238 High-Frequency Heart Rate Variability (HF-HRV; variation in time between each heartbeat for high

239 power frequency) was taken. HF-HRV (HRV in the 0.15-0.4 Hz band range) was used for assessing
240 vagal tone as an index of the parasympathetic nervous system activity (Laborde et al., 2017; Shaffer et
241 al., 2014).

242 During the experiment, three sensors were applied to the torso to reproduce Einthoven's triangle
243 (i.e., one electrode on each shoulder and one on the left hip). Then, these were connected to The Biopac
244 Student Lab Pro 3.7 software. The software was programmed to filter real-time data using a band-pass of
245 0–35 Hz and .5-35 Hz, respectively. The sampling rate for data acquisition was set at 2000Hz. The
246 recordings were interspersed with 30s breaks. To facilitate data recording, we configured a graphical
247 template in the Biopac Student Lab software allowing us to manually add markers for precise
248 visualisation of time intervals within the software's dialogue box (e.g., beginning and end of resting
249 state; start and end for HRV during Stroop task, etc.).

250 ECG signals were first visually inspected to remove artifacts and subsequently imported into
251 Kubios HRV software (Tarvainen et al., 2014) to obtain the frequency domain measure of the High-
252 Frequency band (i.e., 0.15-0.4 Hz). The software retrieves the interbeat (or RR) intervals from the
253 original ECG signal and applies the smoothness prior's method to remove the low-frequencybaseline
254 trend component. The normalised HF-HRV units were acquired through frequency domain estimation
255 employing powerspectrum density. This estimation method involved Welch's periodogram method,
256 which leverages the fast Fourier transformation.

257 ECG signals were captured in conjunction with electrodermal activity (EDA) signals, as shown in
258 previous studies investigating the link between touch and ANS activity (Chatel-Goldman et al., 2014;
259 Sacchetti et al., 2021). EDA signals, which refer to the electrical activity of the skin resulting from
260 variations in sweating, were used for calculating Skin Conductance Level (SCL), a measure of the tonic
261 arousal regulated by the sympathetic nervous system (SNS; Dawson et al., 2007; see Braithwaite et al.,
262 2015, a guide for analysing SCL). When the sympathetic system is activated, the electrical activity of the

263 skin results in increased sweating and, thus, increased SCL (Gordan et al., 2015).

264 While arousal levels were recorded throughout the experiment, our analysis focused on changes
265 in HRV and SCL during two distinct phases: resting (pre task) and during task performance. These
266 phases were analysed at two different time points, i.e., time 1 (T1) and time 2 (T2). Therefore, the study
267 design resulted in a total of four recordings for each participant, as follows:

- 268 • Pre-Task at T1: before participants performed the Stroop Task during the first session;
- 269 • During task at T1: during Stroop Task performance in the first session;
- 270 • Pre-Task at T2: prior to touch stimulation (touch group) or task performance in the second session;
- 271 • During task at T2: during Stroop Task performance in the second session.

272 Notably, for resting state measurement, participants were instructed to remain still and relaxed with their
273 eyes open for 3 minutes, a sufficient time interval length for obtaining accurate measures of
274 physiological arousal (Della Longa et. al., 2021; Munoz et al., 2015; Ree et al., 2019). The rationale for
275 recording physiological arousal before the task was to ensure that any differences observed during the
276 tasks were not influenced by pre-existing group differences in the arousal levels (Liang et al., 2009;
277 Pendleton et al., 2016). Moreover, real-time assessments of the HF-HRV/SCL levels during the task
278 contributed to examining specific changes in arousal linked to task engagement (Culver et al., 2012;
279 Liang et al., 2009; Pendleton et al., 2016), particularly in relation to touch stimulation.

280 **2.3.4 Self-report questionnaires**

281 **2.3.4.1 The Depression, Anxiety, and Stress Scale (DASS-21)**

282 DASS-21 (Lovibond & Lovibond, 1995) is a self-report scale of mood that consists of 21 items
283 divided into three subscales assessing depression (e.g., lack of interest/involvement in activities,
284 anhedonia, etc), anxiety (e.g., restlessness, and physiological arousal associated with anxiety), and stress
285 (e.g., being easily upset/agitated, irritable/over-reactive, etc). Participants are asked to rate the presence
286 and intensity of their symptoms over the past week on a 4-point Likert scale. Each item can be rated

287 from “0” which indicates the symptoms were not experienced at all to “4” which indicates that the
288 symptoms were experienced most of the time.

289

290 **2.3.4.2 Positive and Negative Affect Schedule (PANAS)**

291 The Positive and Negative Affect Schedule (PANAS; Watson et al., 1988) was used to evaluate
292 positive and negative emotions before and after completing the Stroop Task. Participants were asked to
293 respond to a 20-item self-report using a 5-point scale with 10 items assessing positive affect and 10 items
294 assessing negative affect. Each item can be rated from “1” (very slightly or not at all) to “5” (extremely).
295 Scores ranged from 10 to 50 on each scale, with higher scores on the positive affect scale indicating a
296 more pronounced positive mood (e.g., “enthusiast”) whereas items with higher scores on the negative
297 affect scale indicate a more pronounced negative mood (e.g., “nervous”).

298 **2.4 Data handling**

299 Statistical analyses were conducted using IBM SPSS 26 (SPSS Inc., Chicago, IL). A series of
300 independent sample t-tests were performed to determine whether there were any baseline statistically
301 significant differences in the demographics (e.g., age and education), DASS-21 subscales, PANAS
302 scores, and HF-HRV/SCL levels between the two groups (touch vs. no-touch group). For the analysis
303 of Stroop Task performance, we calculated the mean of response times (RTs) in msec and the % of
304 correct responses for assessing the accuracy for each word category (congruent and incongruent). To
305 assess changes in Stroop Task performance, two separate mixed-design two-way ANOVAs were
306 performed, with Group [touch vs. no-touch] as a between-subjects factor, and Congruency [congruent
307 vs. incongruent words] as a within-subjects factor, using RTs or Accuracy as a dependent variable.

308 Then, we ran two one-way ANOVAs using Group [touch vs no-touch] as a between-subjects factor
309 and HF-HRV or SCL as a dependent variable to assess changes in the parasympathetic and sympathetic
310 activity respectively. Prior to these analyses, we calculated the difference (Δ) in mean scores between

311 T1 and T2 for HF-HRV and SCL measurements. For both HF-HRV and SCL measures, we considered
312 two temporal windows, i.e., recordings before and during the Stroop Task.

313 To account for a potential trade-off between accuracy and speed, we calculated an inverse
314 efficiency score (IES) by taking the ratio of the percentage of correct responses (expressed as a decimal)
315 to the mean latency for both congruent and incongruent trials. This calculation was carried out separately
316 for T1 and T2, for each group. We conducted a 2 Group [touch vs no-touch] \times 2 Time [T1 vs T2] mixed
317 ANOVA to assess changes in the IES.

318 An additional 3-way mixed design ANOVA was performed with Group [touch vs no-touch] as a
319 between-subjects factor, and Time [T1 vs T2] and Valence [positive vs negative emotions] as within-
320 subjects factors to assess changes in emotions based on the PANAS questionnaire scores from T1 to T2.

321 A series of Pearson correlations were performed to explore the relationship between physiological
322 arousal (SCL and HF-HRV) and cognitive outcomes (RTs and Accuracy) obtained from the Stroop Test
323 within each touch/no-touch group. For our analyses, we calculated the Δ difference in mean scores
324 between T1 and T2 for RTs and Accuracy (for congruent and incongruent words). Similarly, to establish
325 the change indices for arousal levels, we calculated the change (Δ) in mean scores for HF-HRV and SCL
326 between T1 and T2 across two phases: resting state (before the task) and during task performance. After
327 obtaining the Δ change index values for all variables, we proceeded to examine the correlations.

328 Before performing the ANOVAs, all dependent variables were tested for homogeneity of variance
329 and sphericity assumptions. To follow-up all significant interactions, we conducted a series of
330 independent sample t-tests to examine differences between the touch and no touch groups. P-values were
331 corrected using the Bonferroni method to account for multiple comparisons (Rogers & Weiss, 2009). A
332 significance threshold of $p < .05$ was set for all effects. Effect sizes were estimated using partial eta
333 square (η^2_p) and Cohen's d.

334

335 3. Results

336 3.1 Descriptive statistics

337 Overall, participants in the touch group reported the touch stimulation as relatively pleasant (Mean
338 = 12.95cm; SD = 3.5). Baseline descriptive statistics for demographics, mood (DASS-21), emotions
339 (PANAS), and physiological measures (HF-HRV and SCL) for each group (touch vs no-touch) are
340 reported in Table 1. Overall, we observed no significant differences when comparing baseline
341 measurements between the two groups. Therefore, the two groups were comparable in all measures.

342
343 ----- Please insert Table 1 about here -----

344

345 3.2 PANAS analysis

346 The 3-way mixed ANOVA on mean scores obtained at the PANAS for positive and negative
347 emotions revealed a significant main effect of Valence [$F(1, 70) = 363.61, p < .001, \eta^2p = .84$], which
348 was corroborated by a significant interaction of Time \times Valence [$F(1, 70) = 21.15, p < .001, \eta^2p = .09$].
349 In both groups, post-hoc tests revealed that positive emotions significantly increased, $t(71) = 3.02, p =$
350 $.004, d = .35$, whereas negative emotions decreased at T2, $t(71) = 3.08, p = .003, d = .36$. However, there
351 was no variation in PANAS scores across the touch and no-touch groups from T1 to T2, suggesting that
352 positive and negative emotions did not differ between the two groups before and after completing the
353 Stroop Task.

354

355 3.3 Stroop Task outcomes

356 3.3.1 Response times (RTs)

357 Findings revealed significant main effects of Group [$F(1, 70) = 11.09, p < .001, \eta^2p = .14$] and
358 Congruency [$F(1, 70) = 11.09, p < .001, \eta^2p = .14$]. These effects were further qualified by a significant

359 Group \times Congruency interaction [$F(1, 70) = 8.39, p = .005, \eta^2_p = .11$]. As shown in Figure 2, an
360 independent sample t-test revealed a greater reduction in RTs in the touch group for congruent trials
361 (Mean = 284.60 msec, SD = 69.22) compared to the no-touch group (Mean = 98.14 msec, SD = 153.94),
362 $t(70) = 6.63, p < .001$, Cohen's $d = 1.03$. Similarly, a greater reduction in RTs was observed in the touch
363 group for the incongruent trials (Mean = 168.22 msec, SD = 113.70) compared to the no-touch group
364 (Mean = 90.04 msec, SD = 103.69), $t(70) = 3.05, p = .002$, Cohen's $d = .72$.

365
366 ----- Please insert Figure 2 about here -----
367

368 To summarise, these findings indicate that the group receiving gentle, low-force, slow-moving
369 touch exhibited faster processing in both congruent and incongruent trials, compared to the no-touch
370 group.

371 3.3.2 Accuracy

372 The analyses did not yield a significant main effect of Congruency [$F(1, 70) = .98, p = .33, \eta^2_p =$
373 $.02$]. Similarly, there were no significant effect of Group [$F(1, 70) = 3.88, p = .05, \eta^2_p = 0.05$] or the
374 Group \times Congruency interaction [$F(1, 70) = 3.39, p = .07, \eta^2_p = 0.05$].

375 3.3.3 Inverse efficiency score (IES)

376 The results showed a significant effect of Time [$F(1, 70) = 297.90, p < .001, \eta^2_p = .81$], and a
377 significant Time \times Group interaction [$F(70) = 51.55, p < .001, \eta^2_p = .42$], as shown in Fig. 3.

378
379 ----- Please insert Figure 3 about here -----
380

381 T-test results revealed no significant difference between the touch and no-touch groups at T1

382 (touch group: Mean = 18.02, SD = 2.48; no-touch group: Mean = 17.54, SD = 2.93), $t(70) = 0.76$, $p =$
383 .45, Cohen's $d = .18$. However, at T2, there was a significant difference between the two groups (touch
384 group: Mean = 14.87, SD = 2.18; no-touch group: Mean = 16.24, SD = 3.06), $t(70) = 2.18$, $p = .03$,
385 Cohen's $d = 0.51$. In line with our main results, results suggest that the touch group showed better
386 performance at T2, with faster responses while maintaining high accuracy, as indicated by the
387 significantly lower IES at T2.

388 **3.4 High-Frequency Heart Rate Variability (HF-HRV) outcomes**

389 HRV during task

390 When looking at the HF-HRV during the task, results revealed a significant main effect of Group
391 [$F(1, 70) = 48.55$, $p < .001$, $\eta^2p = .41$], indicating a difference in HF-HRV levels between groups. As
392 shown in Figure 4, an independent sample t-test revealed a significant difference in the change of HF-
393 HRV between groups, $t(70) = -6.96$, $p < .001$, Cohen's $d = 1.64$. Specifically, HF-HRV was significantly
394 greater in the touch group (Mean = -6.13, SD = 3.92) than in the no-touch group (Mean = -1.22, SD =
395 1.56).

396
397 ----- Please insert Figure 4 about here -----

398
399 Overall, these results showed a greater increase in HF-HRV in the touch group compared to the
400 no-touch group.

401 **3.5 Skin conductance level (SCL) outcomes**

402 We did not observe any significant main effect of Group [$F(1, 70) = 1.72$, $p = .20$, $\eta^2p = .03$] for
403 SCL during the Stroop Task performance.

404

405 **3.6 Correlations analyses: physiological arousal and cognitive outcomes**

406 Correlational analyses between measures of physiological arousal (HF-HRV and SCL pre and
407 during task) and Stroop outcomes (RTs and Accuracy) were performed for each group. In the touch
408 group, we observed a significant and negative association between HF-HRV during task and RTs for
409 incongruent words ($r = -.36$, $p = .02$) but not for congruent words ($p = .36$). However, no significant
410 correlations were found between physiological measures during task and accuracy (all $ps \geq .40$).
411 Moreover, when looking at the no-touch group, we did not observe any significant association between
412 physiological measures during the task and Stroop outcomes (all $ps > .33$).

413 Lastly, no significant correlations were found between HF-HRV or SCL pre-task and cognitive
414 outcomes within each group. Specifically, in the touch group, the p-values ranged from above 0.40 to
415 0.80. Similarly, the no-touch group also exhibited no significant correlations, with p-values ranging
416 between 0.40 and 0.90. These results suggest that the physiological state at rest did not relate to cognitive
417 performance.

418

419 **4 Discussion**

420

421 CT afferents contribute to affective touch processing and the regulation of social behaviours
422 (Hazard et al. 2022), including modulating stress response and resilience (Walker et al. 2022). This study
423 explored the effects of touch, specifically gentle, low-force, slow-moving touch, designed to optimally
424 activate CT afferents on physiological arousal and cognitive performance, with a particular emphasis on
425 inhibitory control of goal-irrelevant stimuli. We hypothesised that touch stimulation would positively
426 influence participants' physiological states, enhancing their ability to manage conflicting information
427 during a cognitive task. Although Saunders and colleagues (2018) were the first to examine the impact
428 of touch (i.e., handholding with a romantic partner) on cognitive functioning (i.e., error monitoring), to

429 our knowledge, this study is the first to explore the beneficial effects of gentle, low-force, slow-moving
430 touch on inhibitory control ability through the modulation of psychophysiological reactivity. Our
431 findings suggest that participants receiving gentle, low-force, slow-moving touch exhibited increased
432 physiological arousal, as evidenced by higher HF-HRV, and reduced RTs during the Stroop Task,
433 compared to those who did not receive touch. These results may point to a potential link between gentle,
434 low-force, slow-moving touch and cognitive performance, particularly in a task involving inhibitory
435 control. However, further research is necessary to fully elucidate the nature of this relationship and
436 determine the specific underlying mechanisms involved.

437 It is important to note that a practice effect was observed in both groups, with a greater reduction
438 RTs in the touch group, suggesting that touch may play an active role in cognitive processing, potentially
439 extending its influence beyond mere repeated exposure. These findings seem to align with the "embodied
440 cognition" framework, which proposes that sensory experiences, including tactile sensations, could play
441 a significant role in bolstering cognitive processes (Gallese & Ebisch, 2013; Wilson & Golonka, 2013).

442 The mechanism for the increased cognitive performance, as indicated by reduced RTs, may also
443 be grounded in the homeostatic and allostatic regulation properties of affective touch (Fotopoulou et al.,
444 2022). It is possible that in our study, the touch manipulation could have facilitated an increase in internal
445 control (e.g., heightened body awareness; "homeostatic mechanism"), which might have contributed to
446 the regulation of affective and physiological states ("allostatic mechanism") (Burleson & Quigley, 2021;
447 Fotopoulou & Tsakiris, 2017; Fotopoulou et al., 2022). This effect may be amplified when touch involves
448 the activation of CT afferents, as is the case with affective/pleasant stimulation (e.g., Ree et al., 2019;
449 Silvestri et al., 2024; Van Puyvelde et al., 2019). Affective regulation is crucial in achieving optimal
450 goal-directed behaviour (Cardinale et al., 2019; Rónai et al., 2024). It can be speculated that integrating
451 sensory information from gentle, low-force, slow-moving touch with higher-level cognitive control
452 activity might have enabled participants to regulate task-induced negative emotions (Ellingsen et al.,

453 2016; McRae et al., 2012), which could have helped them to cope with inhibitory control mechanisms
454 (Gliga et al., 2019; McCabe et al., 2008). Although speculative, this interpretation resonates with the
455 findings of the Saunders et al. study (2018), which suggest that touch between romantic partners can
456 increase self-reported positive emotions and buffer against the threat of negative information, possibly
457 making people more open to negative signals or processing negative, affectively charged events (e.g.,
458 impulses and mistakes) during a conflict task performance. Furthermore, touch is known to have
459 significant implications for the regulation of the Hypothalamic-Pituitary-Adrenal (HPA) axis (Yachi et
460 al., 2018), a critical system involved in stress management (Smith et al., 2006). Although this study did
461 not explicitly test this hypothesis, it is possible that the type of touch used in our study may have
462 stimulated the release of oxytocin (Portnova et al., 2020; Walker et al., 2017), a hormone associated with
463 stress reduction (Lee et al., 2009). This release might contribute to lower cortisol levels by influencing
464 the hippocampus and other brain regions that regulate the HPA axis (Matsushita et al., 2019).
465 Consequently, gentle, low-force, slow-moving touch may promote a more adaptive stress response,
466 facilitating a timely deactivation of the HPA axis and supporting overall physiological homeostasis (Kidd
467 et al., 2023; Lupien et al., 2009; McEwen, 2007).

468 Another potential mechanism to support the findings observed here may be attributed to changes
469 in physiological arousal following touch manipulation. Participants in the touch group exhibited a more
470 pronounced increase in HF-HRV compared to the no-touch group. According to the NIM (Thayer &
471 Lane, 2000, 2009a), this effect might reflect a boost of flexible and adaptive responses to increasingly
472 cognitive demand. One crucial component of this flexibility could be inhibitory control, which involves
473 a series of feedback loops between frontal brain areas in the central nervous system, and the ANS, which
474 in turn regulates heart rate, as indexed by HRV (Thayer & Friedman, 2002; Thayer, 2006). It is
475 reasonable to suggest that enhanced physiological reactivity, supported by increased HRV levels—
476 potentially fostered by gentle, low-force, slow-moving touch (Tricoli et al., 2017; Van Puyvelde et al.,

477 2019)—might have contributed to participants' quicker reactions during the Stroop Task performance
478 (Pallak et al., 1975). These mechanisms could include increased allocation of anticipatory attentional
479 resources (Bastiaansen & Brunia, 2001; Weiss et al., 2018), cognitive control over conflicting and
480 irrelevant information (Banich et al., 2019), and error monitoring (Saunders et al., 2018). Supporting
481 this idea, neuroimaging studies revealed that, in particular being gently stroked, activates a brain network
482 including, e.g., the orbitofrontal, insular, and cingulate cortices, all of which are involved in
483 interoception, autonomic regulation, and high-level cognitive processes (e.g., Craig, 2002, 2008;
484 Fotopoulou et al., 2022; Gordon et al., 2013; McCabe et al., 2008; McGlone et al., 2012).

485 It should be noted that even the no-touch group showed an improvement in HF-HRV levels. We
486 speculate that participants' expectations regarding the upcoming tasks may have heightened their arousal
487 levels in preparation for the next phase of the experiment (Knutson & Greer, 2008). Another possible
488 explanation is that repeated exposure to the tactile stimulus may have led to sensitization, where initial
489 arousal during the first Stroop task primes the nervous system for increased arousal in later sessions
490 (Stevens & Bruck, 2019). Presumably according to the NIM (Thayer & Lane, 2000), an increase in
491 parasympathetic activity is typically expected to enhance executive functioning, even in the no-touch
492 group. However, the lack of a significant correlation between Stroop performance and HF-HRV suggests
493 a more complex relationship between physiological measures and cognitive outcomes, particularly in
494 the context of CT-targeted touch. In the absence of touch stimulation, this relationship could be weaker.

495 Partially consistent with NIM (Thayer & Lane, 2000), the changes observed in physiological
496 responses during the Stroop Task may have been driven by parasympathetic activity, as indicated by
497 significant changes in HF-HRV levels. Accordingly, we did not observe any significant difference in
498 sympathetic activity as measured by SCL levels across the two groups. One possible explanation for the
499 divergence between SCL and HRV effects is that during cognitive challenges, individuals might
500 experience increased sympathetic activation that does not correspond to changes in SCL. This could be

501 due to a “feedback loop” from cognitive engagement that enhances parasympathetic activity (i.e.,
502 increased HRV) while inhibiting sympathetic activation (i.e., lower SCL) (Knight et al., 2021). This
503 concept further highlights that dimensions of arousal may not be uniform and affect all physiological
504 parameters (like SCL and HRV) (Dickman, 2002).

505 Nevertheless, our findings revealed that in the touch group, higher HF-HRV levels were linked to
506 faster reaction times compared to the no-touch group, but no changes were observed in levels of SCL.
507 This finding could be consistent with a relationship between parasympathetic activity and cognitive
508 performance (Lazaridi et al., 2022; Nicolini et al., 2024), particularly under increased cognitive
509 demands, as evidenced by the highest HRV levels observed during incongruent trials (Solhjoo et al.,
510 2019). These findings may imply that HRV could serve as an indicator of an adaptive stress response
511 (Thayer et al., 2012), where greater mental effort may contribute to improved performance, especially
512 in more complex tasks (Solhjoo et al., 2019). Indirect support for this idea comes from findings that CT
513 mediated touch may have a regulatory effect on the parasympathetic nervous system (i.e., as reflected
514 in increased HRV), as observed in previous research (Manzotti et al., 2023; Triscoli et al., 2017; Van
515 Puyvelde et al., 2019).

516 **4.1 Limitations**

517 Although our findings seem to suggest that gentle, low-force, slow-moving touch may enhance
518 cognitive performance through physiological regulation, the absence of a group receiving CT-targeted
519 touch at suboptimal velocities (e.g., faster speeds outside the optimal CT range, such as 30 cm/s; Sacchetti
520 et al., 2021, or static touch; Ali et al., 2023) limits our ability to draw definitive conclusions about the
521 specific velocity effects of CT-targeted touch. Including such control groups in future research could
522 help disentangle the unique contributions of CT-targeted touch from general tactile stimulation,
523 providing a clearer understanding of its specific influence on cognitive processes. Furthermore, the
524 current study did not determine whether the effects observed are specific to CT-targeted touch or could

525 be attributed to any form of tactile stimulation such as tapping and light finger touch (non-affective touch;
526 Della Longa et al., 2023; Lee et al., 2018a). Future research could explore this distinction to better isolate
527 the potential contributions of CT-targeted touch to the observed effects. It is also important to highlight
528 that gentle skin stroking activates various classes of C-fiber low-threshold mechanoreceptors (CLTM),
529 including A β field low-threshold mechanoreceptors which are highly sensitive to gentle stroking but
530 unresponsive to other types of stimuli like hair deflection (Walker et al., 2022; Watkins et al., 2021; Bai
531 et al., 2015). Future studies should further investigate the sensory role of these mechanoreceptors,
532 particularly in distinguishing their contributions to affective touch vs discriminative touch.

533 In this study, other touch properties (such as duration and manual stimulation) may have played a
534 significant role in the interaction between autonomic regulation and task performance. Therefore, future
535 research might consider investigating the impact of various CT-touch characteristics (e.g., velocity,
536 temperature, skin locations; Ackerley et al., 2014a, 2014b), or non-CT touch characteristics, on both
537 physiological and cognitive outcomes. Furthermore, we do not exclude the potential beneficial effects
538 of different tactile texture stimuli (e.g., satin; haptic glove; Etzi et al., 2018; Terrile et al., 2021), as well
539 as sensorial activities (e.g., light, aroma; Chamine & Oken, 2015; Siraji et al., 2023) could influence
540 physiological patterns and cognitive processes related to inhibitory control. Including control conditions
541 would enhance the validity of our findings, allowing to determine whether the observed effects are
542 indeed attributable to the specific tactile or sensory modalities being tested.

543 It is also important to acknowledge that improvements in cognitive performance may stem from
544 attentiveness or motivation related to social facilitation, such as the awareness or presence of other
545 individuals (Belletier et al., 2019). To minimise contextual variability, our experimental setup
546 consistently included both the researcher and assistant researcher across all participants. However, our
547 effort to keep the experimenter's presence non-intrusive or at a distance from the participant in the control
548 condition may have unintentionally drawn attention to proximity as a potential confounding factor.

549 Future studies could incorporate more rigorous control over proximity, such as setting fixed distances
550 between the participant and the experimenter or using a transparent partition to control for the visual and
551 spatial presence of the experimenter, thereby reducing its influence on the physiological-cognitive
552 outcomes. Furthermore, we recommend an experimental design that incorporates additional conditions
553 to isolate the effects of touch from social presence, such as using a Rotary Tactile Stimulator (RTS). The
554 RTS allows for the delivery of precise, controlled force and velocity, potentially reducing variability
555 introduced by human touch and establishing a control condition in which participants receive identical
556 tactile stimuli without the influence of social context (Lee et al., 2018b). Our study utilised the HF
557 parameter to measure parasympathetic activity in the autonomic nervous system through high-frequency
558 bands of HRV. Future research should consider incorporating a variety of HRV measures, such as time-
559 domain indices or additional metrics, (e.g., SDNN Index, RMSSD, NN50, and pNN50, see Shaffer &
560 Ginsberg, 2017 for an overview) to provide a more comprehensive analysis of the cardiac vagal tone
561 related to sensory-cognitive stimulation.

562 During the experiment, gentle, low-force, slow-moving touch at 3cm/sec was delivered for two
563 minutes, which is generally sufficient to elicit physiological activation (Della Longa et al., 2021; Ree et
564 al., 2019). Despite this, it is important to note that too high or too low arousal levels elicited can have a
565 detrimental effect on cognitive performance (Storbeck et al., 2008; Yerkes & Dodson, 1908). Thus,
566 further research could explore the optimal duration of touch stimulation to achieve the desired level of
567 arousal without negatively impacting cognitive performance.

568 Lastly, we assessed emotional distress using the DASS-21 questionnaire and affect through PANAS
569 before and after the experimental manipulation. These measures provided insight into participants'
570 emotional states, which helped understand potential confounding factors, such as whether any observed
571 changes in physiological responses or cognitive performance could be due to pre-existing emotional
572 states. While PANAS has been previously used in studies related to affective touch (Mammarella et al.,

573 2012; Sailer et al., 2024), we are aware that these questionnaires may not directly capture the influence
574 of social or environmental factors.

575 **5 Conclusion**

576 While we cannot make definitive claims about the specific role of CT-targeted touch in enhancing
577 inhibitory control through physiological regulation, our study provides preliminary evidence of a
578 potential connection between gentle, low-force, slow-moving touch and autonomic regulation, as
579 indicated by increased HF-HRV. This was accompanied by enhanced processing speed during the Stroop
580 Task. Future research employing more rigorous control conditions is necessary to further clarify the role
581 of CT-targeted touch in shaping physiological and cognitive outcomes.

582 **CRedit authorship contribution statement**

583 Loredana Frau: Writing – original draft, Writing – review & editing, Conceptualization, Resources,
584 Methodology, Investigation, Validation, Software, Project administration, Data curation, Formal
585 analysis. Davide Bruno: Review & editing, Validation, Supervision. Francis McGlone: Review &
586 editing. Valentina Cazzato: Writing – Review & editing, Validation, Supervision.

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590 **Data Availability**

591 All the relevant data are freely available from OSF at the following weblink:
592 https://osf.io/wbgjv/?view_only=35b5166bea004ce9974476c099dc317f. The current study and its
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