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Repeated Menthol spray application enhances exercise capacity in the heat

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56 **Abstract**

57

58 **Purpose.** Exercise performance is impaired in the heat and a contributing factor to
59 this decrement is thermal discomfort. Menthol-spraying of skin is one means of
60 alleviating thermal discomfort but has yet to be shown to be ergogenic using single
61 spray applications. We examined whether repeated menthol-spraying could relieve
62 thermal discomfort, reduce perception of exertion and improve exercise performance
63 in hot (35°C), dry (22% RH) conditions; we hypothesised it would. **Method.** Eight
64 trained cyclists completed two separate conditions of fixed intensity (FI) cycling
65 (50% P_{Max}) for 45-minutes before a test to exhaustion (TTE; 70% P_{Max}) with 100 mL
66 of menthol-spray (0.20% menthol) or control-spray applied to the torso after 20 and
67 40-minutes. Perceptual (thermal sensation (TS), thermal comfort (TC), RPE)
68 performance (TTE duration), thermal variables (skin temperature (T_{skin}), rectal
69 temperature (T_{rec}), cardiac frequency (fc)) and sweating were measured. Data were
70 compared using ANOVA to 0.05 alpha level. **Results.** Menthol-spray improved TS
71 (*'cold'* sensation *cf* *'warm/hot'* after first spraying; p=.008) but only descriptively
72 altered TC (*'comfortable'* *cf* *'uncomfortable'*; p=.173). Sweat production (994 (380)
73 mL *cf* 1180 (380); p=.020) mL and rate (827 (327)mL·hr⁻¹ *cf* 941 (319)mL·hr⁻¹;
74 p=.048) lowered. TTE performance improved (4.6 (1.74) *cf* 2.4 (1.55) minutes
75 (p=.004). Menthol-spray effects diminished despite repeated applications indicating
76 increased contribution of visceral thermoreceptors to thermal perception. **Conclusion.**
77 Repeated menthol-spray improves exercise capacity but alters thermoregulation
78 potentially conflicting behavioural and thermoregulatory drivers; care should be taken
79 with its use. Carrying and deploying menthol-spray would impose a logistical burden
80 which needs consideration against performance benefit.

81

82 **Keywords.** TRPM8 receptors, thermoregulation, sweating, thermal perception.

83

84

85 **Introduction**

86
87 Exercise performance is impaired in hot conditions with fatigue occurring
88 prematurely compared to cool environments¹. The aetiology of this fatigue is complex
89 and multifaceted but is in part attributable to increased thermal sensations (i.e. feeling
90 hot) and thermal discomfort². Accordingly, any intervention that offsets these
91 disturbances in thermal perception may prove to be ergogenic and influence exercise
92 behaviour³. One such intervention with the potential to do so is the topical application
93 of menthol to the skin. This has been found to change the action potential of the
94 Transient Receptor Potential Melastatin 8 (TRPM8) subfamily of thermoreceptors
95 thereby inducing cool sensations^{4,5}. Although, menthol is also known to activate TRP
96 vanilloid (TRPV) and ankyrin (TRPA) receptors⁶ above temperatures of 37 °C
97 thereby inducing warm sensations⁷. Accordingly in exercise and environmental
98 scenarios where skin temperatures do not exceed 37 °C (i.e. the majority of scenarios)
99 the chemical stimulation of the skin by menthol appears to be a viable means of
100 improving thermal perception and potentially exercise performance.

101
102 Only one study to date has revealed an ergogenic benefit following the topical
103 application of an 8% menthol gel applied to the face during self-paced exercise
104 performed at a fixed perception of exertion². Menthol application induced an
105 approximate 18% increase in total work during the study where thermal stress was
106 applied through a water-perfused suit². Thermal perception was shown to be a
107 relatively independent behavioural regulatory influence on exercise termination as
108 shorter exercise duration was observed with the induction of hot sensations by
109 capsaicin application to the skin². However, in studies performed using ecologically
110 valid laboratory protocols^{3,8,9} an ergogenic effect has proved illusive leading to
111 suggestions that menthol-spraying may only improve thermal perception but not
112 performance¹⁰. Menthol applied to the skin at concentrations (0.05 to 0.20 % L-
113 Menthol in solution), similar to that of commercially available products (Physicool™,
114 London, U.K), has been reliably shown to induce improvements in thermal sensation
115 and comfort, during fixed intensity¹¹ and self-paced exercise^{3,8,9} in the heat. However,
116 it has also been shown to induce heat gain responses (i.e. vasoconstriction¹¹) and alter
117 sweating responses¹²; in the latter case at higher concentrations (i.e. 4.6%¹²).
118 Therefore, it is also plausible that menthol application could increase the risk of heat-
119 illness and place behavioural and thermoregulatory drivers in conflict.

120
121 Nevertheless, there are iterations on the timing of menthol application that have not
122 been explored experimentally which may mean concluding a lack of ergogenic effect
123 is premature. To date, we have explored whether relieving thermal discomfort and
124 improving thermal sensation is performance enhancing prior to and during the early
125 minutes of a 40 km cycling time trial; it was not⁸. We have examined whether
126 inducing hot and uncomfortable sensations using a heat pre-load followed by menthol
127 application would result in improved performance of a shorter duration exercise of 5
128 km running but it did not⁹. Most recently we examined whether applying menthol
129 towards the end of an exercise task (i.e. at 10 km of a 16.1 km cycling time trial; TT)
130 would result in benefits to TT completion time³. Once again we saw no improvement
131 although menthol-spray application did result in lowered RPE in addition to benefits
132 to thermal perception. Each of these studies, and others where perceptual
133 manipulation was the primary goal¹¹ involved *single* applications of menthol-spray. It
134 has yet to be investigated whether *repeated* menthol application can act as an

135 ergogenic aid. Theoretically, in prior studies the acute bouts of thermal discomfort
136 relief through menthol-spray application may have been insufficient to perturb the
137 behavioural thermoregulatory drivers towards altering exercise performance. Whereas
138 *repeated* application may provide a greater driver to change this. Moreover, the nature
139 of the exercise task may also be important. Menthol is evidently more likely to
140 influence an exercise task where tolerance is the critical factor¹⁰ (e.g. test to
141 exhaustion; TTE) rather than the spontaneous variation in power output (e.g. TT)
142 which have consistently failed to be responsive to menthol in three of our previous
143 studies^{3,8,9}. Accordingly, the present study sought to examine this possibility.

144

145 We hypothesised that menthol application, applied every 20-minutes during exercise
146 in the heat¹¹, would enhance exercise performance in a subsequent TTE where heat
147 tolerance is the main limiting factor to performance (H₁). We also hypothesised that
148 menthol-spray application would enhance thermal perception by inducing cool
149 thermal sensations and relieving thermal discomfort which may result in reduced
150 perception of exertion in contrast to a control-spray condition (H₂).

151

152 **Method**

153

154 ***Experimental Design***

155 The local ethics committee approved the study which used a within participant,
156 repeated measures design in which participants completed three exercise conditions.
157 The first condition took place in a temperate environment and was to establish their
158 maximal power output (P_{Max}) for use during the subsequent two conditions which
159 took place in a hot environment. Conditions two and three were counter-balanced
160 where the participants' t-shirt was repeatedly sprayed (i.e. every 20-minutes) with a
161 menthol-spray or a control-spray. Tests took place at the same time of day (\pm 1 hour)
162 with a minimum of 48 hours between tests.

163

164 ***Participants***

165 Eight trained cyclists (mean \pm SD: age 22 \pm 2 yrs; height 1.84 \pm 0.1 m; body surface
166 area¹³ 2.05 \pm 0.1 m² P_{Max} 362.5 \pm 35.4 W) volunteered and provided written informed
167 consent. Participants were considered trained if they achieved a minimum P_{Max} of
168 \geq 350 watts¹⁴. Participants abstained from alcohol, caffeine consumption and
169 strenuous exercise 24 hours prior to each test and were non-smokers.

170

171 ***Procedures***

172 *Condition One - P_{Max} Test:* Participants arrived at the laboratory wearing cycle
173 clothing. They were instrumented with a heart rate monitor (FT1, Polar Electro Oy,
174 Kempele, Finland) and entered the environmental chamber held at a temperature of
175 16°C. Participants were made comfortable on the cycle ergometer (Velotron,
176 Racermate, Seattle, USA); bike positioning was replicated for subsequent visits.
177 Participants completed a standardised 5-minute warm up at 150 W and a cadence of
178 70 rev·min⁻¹ followed by stretching. They remounted the ergometer and
179 recommenced cycling at the same power output and cadence as the warm-up. The
180 required power output was increased by 50 W every 2-minutes until volitional
181 exhaustion or when the prescribed cadence could not be maintained for 15-seconds
182 and having achieved a heart rate within 10 b.p.m⁻¹ of age predicted maximum.
183 Participants were instructed prior that they should make a maximal effort during the
184 test.

185 *Conditions Two and Three - Repeated Spray Applications:* Participants arrived in a
186 hydrated state; i.e. having consumed 500 mL of water the preceding night and 500
187 mL in the two hours prior to arrival at the laboratory. Participants were allowed to
188 drink tepid tap water during the trials. Participants first voided and naked body mass
189 was measured in private (Seca, Model 705 2321009, Vogel & Halke, Hamburg,
190 Germany). They then donned their cycling shorts and were instrumented with a
191 calibrated, insulated rectal thermistor (Grant Instruments Ltd, Cambridge [Shepreth],
192 U.K) inserted (in private) 12-15 cm beyond the anal sphincter. They were also
193 instrumented with skin thermistors (Grant Instruments Ltd, Cambridge [Shepreth],
194 U.K) placed at eight different body sites¹⁵ on the left side of the body secured by
195 breathable tape (TransporeTM,1527-1, 3M Health Care, MN, USA). A heart rate
196 monitor was also worn to measure cardiac frequency (fc). Rectal temperature (T_{rec})
197 and skin temperature (T_{skin}) were logged automatically every 5-seconds using a
198 remote data logger (Squirrel 2020 series, Grant Instruments Ltd, Cambridge
199 [Shepreth], U.K). Following instrumentation participants completed dressing by
200 wearing socks, shoes and a close-fitting long sleeve t-shirt (100% polyester; Campri
201 Sports Baselayer, Sportsdirect, Shirebrook, U.K). Identical clothing was worn in each
202 condition that involved repeated spraying.

203
204 Participants then entered an environmental chamber set to 35°C and 20% relative
205 humidity (RH). Environmental conditions were measured by a wet-bulb, globe,
206 temperature (WBGT) station (1000 series, Squirrel Data Logger, Grant Instruments
207 Ltd, Cambridge [Shepreth], U.K). One minute prior to the start of exercise, all data
208 logging systems were activated and synchronised. Prior to the commencement in
209 exercise participants provided a resting capillary sample of blood for measurement of
210 blood lactate concentration (B_{lac}). Participants also reported their resting thermal
211 comfort (TC^{16}) and thermal sensation (TS^{16}). Participants then mounted the cycle
212 ergometer and completed the same standardised warm up as prior to the P_{Max} , and
213 then commenced fixed intensity (FI) cycling at 50% P_{Max} for 45-minutes. Participants
214 cycled in front of a fan positioned 80 cm from the velotron (Wahl, Model ZX220,
215 Wahl, Sterling, IL, USA) and pointed at the participants' torso. The wind speed
216 produced by the fan was verified at a fixed position by an anemometer (LM-8000
217 Anemometer, Digital Instruments, New York, USA; this approximated between 1.6
218 and 2.1 $m \cdot s^{-1}$).

219
220 Perceptual responses including RPE¹⁷, TC and TS were obtained initially every 10-
221 minutes of the FI period, until (i.e before) the first spray application at 20-minutes.
222 They were recorded every 5-minutes thereafter; RPE was not collected at 30-minutes.
223 After 20 and 40-minutes of exercise participants' jerseys were sprayed evenly with
224 100 mL of either the control-spray or the menthol-spray which was heated in a water
225 bath to match environmental temperature³. Spray volume was measured on each
226 occasion using calibrated, digital, weighing scales (Sartorius Mechatronics UK Ltd,
227 TE6100, Surrey, U.K; 1 g resolution). Intervals between sprays were 20-minutes on
228 the basis that the menthol-spray perceptual response has been shown to decay
229 thereafter¹¹. Sprays were produced by an independent chemical consultant (Chemical
230 Associates, Rosemead, Frodsham, United Kingdom). The control-spray contained 3%
231 surfactants mixed in water, while the menthol-spray contained a concentration of 0.20
232 wt/wt L-menthol in 3% surfactants plus water.

233

234 Upon completion of the FI period participants provided another capillary blood
235 sample and immediately commenced a test to exhaustion (TTE) at 70 % P_{Max}.
236 Participants received no feedback of exercise time elapsed or encouragement during
237 the TTE. Upon TTE cessation (i.e. volitional exhaustion) the participant exited the
238 chamber and were weighed naked and, in conjunction with measured fluid intake,
239 sweat production and sweat rate were calculated. Performance times were not
240 revealed until the post-experiment debrief.

241

242 *Statistical Analysis*

243 Mean (SD) were calculated for perceptual (TS, TC, RPE), performance, (B_{lac}, TTE
244 duration), thermal (T_{skin}, T_{rec} and *fc*) spray variables (temperature and volume),
245 environmental conditions and sweat production including rate. The normality of
246 distribution was verified using a Kolmogorov-Smirnov test. Data were
247 compared using a repeated measures analysis of variance (ANOVA) at rest and fixed
248 points during the FI period including TTE end point for the two hot trials (9 x 2
249 ANOVA) for perceptual (no RPE measure at rest and 30-minute point) and thermal
250 variables. Sphericity was checked using Mauchly's test and, where necessary, a
251 Greenhouse-Geisser adjustment was applied. The direction of statistically significant
252 effects were determined using Fisher's (LSD) *post-hoc* pair-wise comparisons. Partial
253 eta squared (η^2) are reported as estimates of effect size. Environmental conditions,
254 spray temperature, volume, TTE duration, fluid consumed, sweat data and terminal
255 B_{lac} were compared using paired samples t-test. The 95% confidence interval (CI) was
256 calculated for the TTE data. Data are otherwise presented as mean (SD). An alpha
257 level of 0.05 was used for all statistical tests which were conducted using SPSS (SPSS
258 v 21, IBM, Chicago, Illinois, USA) and Prism (Graphpad, Prism v 6, San Diego,
259 USA).

260

261 **Results**

262

263 *Environmental Conditions*

264 Ambient temperature averaged 35.0 (1.3) °C and 34.6 (1.2) °C in the control-spray
265 and menthol-spray conditions respectively and did not differ ($t = .846$, $p = .213$). RH
266 averaged 21.8 (0.90) % and 22.2 (1.0) % and did not differ ($t = -1.06$, $p = .162$).

267

268 *Spray Volume and Temperature*

269 Volume of spray applied was 200 (3) mL in the control-spray and 200 (2) mL in the
270 menthol-spray conditions which were similar ($t = 0.110$, $p = 0.460$). The temperature
271 of the control-spray averaged 37.4 (1.2) °C and was 38.3 (1.6) °C in the menthol-spray
272 condition and were not different ($t = 1.766$, $p = .097$).

273

274 *TTE Performance*

275 TTE was 2.4 (1.55) minutes and 4.6 (1.74) minutes in the control-spray and menthol-
276 spray conditions respectively and was significantly greater after menthol-spraying
277 application ($t = -3.63$, $p = 0.004$; 95% CI 0.53 to 3.82 minutes).

278

279 *Perceptual Responses*

280 Participants' TS responses were similar in each condition before the first spray (i.e. at
281 20 minutes) and corresponded to the worded descriptor 'hot'. At 25-minutes, 5-
282 minutes after spraying, TS was significantly lower (main effect for condition: $F_{(1, 7)} =$
283 13.139, $p = 0.008$, $\eta^2 = .652$ & interaction effect: $F_{(8, 56)} = 12.843$, $p = 0.001$, $\eta^2 =$

284 .441) in the menthol-spray condition (11.0 (2.4) cm) compared to the control-spray
285 (15.7 (1.6) cm; $p = 0.02$). These ratings corresponded to the worded descriptors
286 'warm' to 'hot' in the control-spray and 'cold' in the menthol-spray condition. The
287 differences due to menthol-spraying remained until 40-minutes where TS was not
288 different ($p = .255$). Following the second administration of menthol-spray TS once
289 again declined (i.e. participants felt cooler) significantly ($p = .035$); see figure 1A.

290
291

292 ***Insert figure 1 near here***

293
294

295 The differences in TS only resulted in numerical changes in TC after spray
296 application (no condition effect: $F_{(1, 7)} = 2.297$, $p = .173$, $\eta^2 = .247$; no interaction
297 effect: $F_{(8, 56)} = 4.789$, $p = .270$, $\eta^2 = .155$) probably because of larger variation in
298 the TC response than TS. At 25-minutes, after first spray application, TC averaged 9.0
299 (3.9) cm and 11.8 (1.6) cm in the control-spray and in the menthol-spray conditions
300 respectively corresponding to the worded descriptors 'uncomfortable' and
301 'comfortable'; see figure 1B.

302

303 RPE did not differ between conditions (condition effect: $F_{(1, 7)} = .057$, $p = .819$, $\eta^2 =$
304 $.008$) or show any interaction effect ($F_{(6, 42)} = .782$, $p = .620$, $\eta^2 = .101$). RPE was
305 always within one RPE rating between condition; see figure 1C.

306

307 ***Thermal Responses (Including fc)***

308 One T_{rec} file was corrupted and consequently data from this participant were removed
309 (T_{rec} data $n = 7$). T_{rec} increased steadily throughout FI exercise and the TTE,
310 indicating that the exercise produced heat at a rate that was uncompensable (main
311 effect for time: $F_{(7, 42)} = 49.490$, $p = .001$, $\eta^2 = .892$); see figure 2A. There was no
312 difference between condition ($F_{(1, 6)} = .017$, $p = .899$, $\eta^2 = .003$) or interaction effect
313 for T_{rec} ($F_{(7, 42)} = 2.097$, $p = .182$, $\eta^2 = .259$). Terminal rectal temperature was 38.5
314 (0.26) and 38.4 (0.37) °C in the control-spray and menthol-spray conditions
315 respectively. The T_{skin} response was similar for the first 20-minutes of FI exercise
316 before spray application. Despite the changes in TS, there was no evident condition
317 effect for T_{skin} ($F_{(1, 7)} = .444$, $p = .527$, $\eta^2 = .105$) or any interaction effect ($F_{(7, 49)} =$
318 $.575$, $p = .389$, $\eta^2 = .147$) although T_{skin} did change numerically in the same direction
319 as the TS ratings. These data indicate an uncoupling of the T_{skin} and thermal
320 perceptual response; see figure 2B. Following the first menthol-spray application the
321 T_{skin} response had a tendency to be numerically lower until the commencement of the
322 TTE; see figure 2B. fc was similar throughout each condition and averaged 171 (14)
323 $b \cdot min^{-1}$ and 174 (7) $b \cdot min^{-1}$ in the control-spray and menthol-spray condition at test
324 cessation. There was no difference between condition ($F_{(1, 7)} = .053$, $p = .825$, $\eta^2 =$
325 $.008$) or interaction ($F_{(5, 35)} = .108$, $p = .990$, $\eta^2 = .015$).

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Insert figure 2 near here

329

330

331 ***Fluid Consumed, Sweat Produced, Blood lactate and Cardiac Frequency***

332 The volume of fluid consumed by each participant was relatively consistent between
333 conditions and averaged 630 (169) mL and 545 (187) in the control-spray and

334 menthol-spray conditions ($t = 1.12$, $p = .149$). These data combined with naked body
335 mass measurements generated an estimated sweat production of 1180 (380) mL and
336 994 (380) mL in the control-spray and menthol-spray conditions with production
337 being lower after menthol-spray ($t = 3.002$, $p = .020$). Due to the significantly longer
338 exercise duration in the menthol-spray condition the estimated sweat rate (827 (327)
339 mL·hr⁻¹) was reduced ($t = 2.392$, $p = .048$) versus the control-spray condition (941
340 (319) mL·hr⁻¹).

341

342 Terminal B_{lac} at the end of the FI period was 4.3 (2.1) mmol/L and 5.1 (3.1) mmol/L
343 in the control-spray and menthol-spray conditions and was not different ($t = 1.189$, p
344 = 0.273); further B_{lac} data not shown.

345

346 Discussion

347

348 The present study sought to examine whether *repeated* application on menthol-spray
349 to the torso enhanced exercise performance in trained cyclists in an exercise task
350 which was limited by tolerance rather than power output. Our data showed an
351 improvement in TTE performance of 133 (104) seconds after menthol-spraying in
352 contrast to a control-spray condition; H₁ is therefore accepted. We also suggested that
353 *repeated* menthol-spray application would provide a greater benefit to thermal
354 perception thereby driving behavioural thermoregulation. Our data suggest that only
355 thermal sensation was significantly improved although thermal comfort did alter
356 subjectively in the hypothesised direction. The performance change through
357 perceptual mechanisms did not manifest itself through lowered perceived exertion; we
358 therefore only provide partial support for H₂. An additional novel finding was the
359 change observed in sweat production and sweat rate following repeated menthol-
360 spray application which we have not seen previously with single application studies
361 using this menthol concentration.

362

363 To our knowledge, this is the first study to investigate the possibility of an ergogenic
364 effect of repeated menthol application using an ecologically valid protocol and a
365 menthol concentration similar to those commercially available. The fact that repeated
366 menthol application is required to produce an ergogenic effect provides a challenge to
367 performers of sports where a weight bearing component may be limiting to their
368 performance (e.g. running, tour cycling). A decision to carry and deploy menthol
369 must be balanced against any performance decrement induced by bearing the
370 additional weight. Moreover, our evidence that the perturbation in thermal perception
371 was lesser after the second menthol spray application also suggests that repeated
372 chemical stimulation of the skin may have limitations especially in a hot environment.
373 Indeed, we speculate that repeated menthol application is likely to have a lesser effect
374 because of acute habituation to the sensation¹⁸ or because of an increased contribution
375 of raised deep body temperature to thermoreception thereby reducing the contribution
376 T_{skin} makes to thermal perception¹⁹. Even in the scenario of hot skin and a
377 normothermic deep body temperature, menthol may evoke warm sensations if the
378 mean T_{skin} is over 37 °C which has been shown in isolated cells to activate warm
379 sensitive thermoreceptors TRPA and TRPV⁶. In the present study, activation of these
380 thermoreceptors by menthol may also contribute to the lessened perceptual effect with
381 repeated application. Consequently, a combination of peripheral and visceral
382 thermoreceptor stimulation may be a more viable target for performance enhancement
383 rather than visceral or peripheral alone. There is good evidence that menthol ingestion

384 is performance enhancing¹⁰ and we show here it is premature to conclude that topical
385 application is not. It is now also plausible that topical menthol application could be
386 ergogenic in other activities (e.g. strength and power-based activities) which could be
387 limited by hot environments or the perceptual mechanisms we describe here and
388 elsewhere in relation to RPE³.

389

390 The fact that repeated menthol-spray also altered sweating response by reducing it is
391 also a novel finding although others have reported delayed sweating and reduced
392 sweat production occurs after 4.6% menthol sediment application¹². The extent of the
393 reduction we see in the present study, albeit using different protocols and menthol
394 concentrations (i.e. 0.20% *cf* 4.6%), was far lower (i.e. 12% *cf* 63% of sweat response
395 seen in the control condition) than reported elsewhere¹² indicating a dose response
396 relationship for menthol application to the skin. Others have also reported that
397 menthol application activates different heat gain responses including vasoconstriction
398 with resultant increases in rectal temperature^{11,12}. Although we did not see the latter,
399 we also saw evidence that T_{skin} was lowered after menthol-spray application (see
400 figure 2B) indicating possible vasoconstriction. Any change in T_{skin} was also less
401 substantial on secondary application supporting the idea that visceral thermoreceptors
402 are applying a greater predominance of thermoregulatory input as deep body
403 temperature increases²⁰. Collectively across our study and those of others, we must be
404 cautious when titrating the concentration and frequency of menthol application during
405 exercise to avoid inducing heat gain responses which may increase heat illness risk,
406 especially during high intensity efforts where heat load would be high or when
407 performing in high ambient temperatures. This is especially prudent since an
408 uncoupling of thermal state from thermal perception is plausible with menthol
409 application thereby placing biophysical and behavioural thermoregulatory drivers in
410 conflict. Using a menthol-spray of lower concentration which still induces perceptual
411 benefits but does not alter thermoregulatory response (e.g. .05% concentration) may
412 be a safer option to safeguard health^{8,11,12}. Moreover, the addition of ethanol to the
413 spray mix, which was deliberately excluded in the present and previous studies to
414 maximise perceptual cooling through chemical stimulation and minimise
415 physiological cooling through evaporation, may ensure the perceptual and
416 thermoregulatory responses converge²⁰.

417

418 **Practical Applications**

419

420 Menthol-spray application triggers heat gain responses which could increase risk of
421 heat illness in some circumstances and care should be taken with the concentration
422 and frequency of application. The performance benefit of menthol-spray could be
423 extended to other population groups (i.e untrained persons) and activities where
424 perceptions are partially limiting. However, this must be balanced against the
425 logistical burden to carrying and deploying the spray.

426

427 **Conclusion**

428

429 Repeated menthol-spray application is ergogenic in trained participants during cycling
430 in hot conditions. The perceptual benefits of repeated menthol spraying are likely to
431 be dependent on thermal profile with a diminishing effect when there is an increasing
432 contribution of visceral thermoreceptors to thermoreception; i.e. when deep body
433 temperature is raised.

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References

1. Tucker R, Marle T, Lambert EV, Noakes TD. The rate of heat storage mediates an anticipatory reduction in exercise intensity during cycling at a fixed rate of perceived exertion. *J Physiol*, 2006, 574, 905–915.
2. Schlader ZJ, Simmons SE, Stannard SR, Mundel T. The independent roles of temperature and thermal perception in the control of human thermoregulatory behaviour. *Physiol Behav*, 2011, 103: 217-24.
3. Barwood MJ, Corbett J, Thomas K, Twentyman P. Relieving thermal discomfort: effects of sprayed L-Menthol on perception, performance and time trial cycling in the heat. *Scand J Med Sci in Sport*, 2015, S211-S218.
4. McKemy DD, Neuhausser WM, Julius D. Identification of a cold receptor reveals a general role for TRP channels in thermosensation. *Nature*, 2002, 416, 52-58.
5. Peier AM, Moqrich A, Hergarden AC, Reeve AJ, Andersson DA, Story GM *et al.* A TRP channel that senses cold stimuli and menthol. *Cell*, 2002, 108, 705-15.
6. Macpherson LJ, Hwang SW, Miyamoto T, Dublin AE, Patapoutian A, Story GM. More than cool: promiscuous relationships of menthol and other sensory compounds. *Mol Cell Neurosci*, 2006, 32, 335-343.
7. Green BG. Menthol modulates oral sensations of warmth and cold. *Physiol Behav*, 1985, 35, 427-434.
8. Barwood MJ, Corbett J, James J, White D. Early change in thermal perception is not a driver of anticipatory exercise pacing in the heat. *Br J Sports Med*, 2012, 46(13), 936-942.
9. Barwood MJ, Corbett J, White D. Spraying with 0.20% L-Menthol does not enhance 5k running performance in the heat in untrained runners. *J Sports Med Phys Fitness*, 2014, 54(5), 595-604.
10. Stevens CJ, Best R. Menthol: a fresh ergogenic aid for athletic performance. *Sports Med*, 2018, 47(6), 1035-1042.
11. Gillis DJ, House JR, Tipton MJ. The influence of menthol on thermoregulation and perception during exercise in warm, humid conditions. *Eur J Appl Physiol*, 2010, 110, 609-618.
12. Kounalakis SN, Botonis PG, Koskolou MD, Geladas ND. The effect of menthol application to the skin on sweating rate response during exercise in swimmers and controls. *Eur J Appl Physiol*, 2010, 109, 183-189.
13. DuBois D, DuBois EF (1915) The measurement of the surface area of man. *Int Arch Med*, 15, 868-81.
14. Bentley D J, McNaughton LR, Thompson D, Vleck VE, Batterham AM. Peak power output, the lactate threshold, and time trial performance in cyclists. *Med Sci Sport Exerc*, 2001, 33(12), 2077-2081.
15. Olesen BW. How many sites are necessary to estimate a mean skin temperature? In: Hales JRS, ed. *Thermal Physiology*. New York: Raven Press; 1980, 33-38.
16. Zhang H. *Human thermal sensation and comfort in transient and non-uniform thermal environments (published dissertation)*. Berkley: CA; University of California; 2003.
17. Borg GAV. Psychological bases of perceived exertion. *Med Sci Sport Exerc* 1982, 14, 377–381.

- 483 18. Gillis DJ, Weston N, House JR, Tipton MJ. Influence of daily menthol
484 exposure on human temperature regulation and perception. *Physiol Behav*,
485 2015, 139, 511-518.
- 486 19. Attia M, Engel P. Thermal pleasantness sensation: an indicator of
487 thermal stress. *Eur J Appl Physiol Occup Physiol*, 1982, 50(1), 55-70.
- 488 20. Nadel ER, Horvath SM, Dawson CA, Tucker A. Sensitivity to
489 peripheral and central thermal stimulation in man. *J Appl Physiol*, 1970, 29(5),
490 603-609.
- 491 21. Gillis DJ, Barwood MJ, Newton PS, House JR, Tipton MJ. The
492 influence of a menthol and ethanol soaked garment on human temperature
493 regulation and perception during exercise and rest in warm, humid conditions.
494 *J Therm Biol*, 2016, 58, 99-105.
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502 **Figure Legends**

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504 **Figure 1.** Mean (SD) TS (panel A), TC (panel B) and RPE (panel C) response at rest,
505 during FI exercise and at TTE end in the control-spray (circles) and menthol-spray
506 (squares) conditions; *indicates significant difference between conditions at a given
507 time point; --- indicates application of spray.

508

509 **Figure 2.** Mean (SD) T_{rec} (panel A) and T_{skin} (panel B) response at rest, during FI
510 exercise and at TTE end in the control-spray (circles) and menthol-spray (squares)
511 conditions; *indicates significant difference between conditions at a given time point;
512 --- indicates application of spray.

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