Research Article

The Supraclavicular Skin Temperature Response to Mild Cold Stimulation is Dependent on Ambient Temperature

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Purpose: This study investigated the basal activity, and cold-induced thermogenic response, of supraclavicular brown adipose tissue (BAT) under warm (23°C) and cool (18°C) ambient conditions using supraclavicular skin temperature as a measure of BAT activity. As a highly metabolic, heat-producing tissue, it has been hypothesised that under-active/dysfunctional BAT may underlie a pathological energy imbalance leading to obesity.

Methods: Five lean, healthy participants underwent infrared thermography (IRT) of supraclavicular BAT before, and during, mild cold exposure (single-hand immersion in cool water at 20°C), once at 18°C and once at 23°C. Energy expenditure (EE) was measured simultaneously using indirect calorimetry, and mean skin temperature (Tmsk) was calculated at 1-minute intervals in parallel to IRT using wireless data loggers.

Results: Following 30 minutes of hand cooling, supraclavicular skin temperature (TscB) rose significantly from baseline at an ambient temperature of 23°C (ΔTscB: 0.17 ± 0.03°C, P < 0.01), and EE rose by 0.22 ± 0.02 kJ/min, P < 0.001. At an ambient room temperature of 18°C, Tmsk after hand cooling was similar to baseline, and EE remained unchanged. The Tmsk response was indicative of a systemic vasoconstrictive response of similar magnitude in both warm and cool ambient temperatures.

Conclusions: At 18°C in light clothing, BAT may already be maximally stimulated at baseline, and respond minimally to additional cold exposure. Ambient temperature is recognised as a determinant of glucose uptake in BAT. In this study, we show, that it also modulates the TscB response to further localised cold-stimulation, indicating an effect on BAT thermogenesis.

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Introduction

Following the recent discovery of functional, metabolically active brown adipose tissue (BAT) in humans, interest lies in elucidating the mechanisms underlying the BAT mediated non-shivering thermogenic (NST) component of the physiological response to acute cold [9, 10, 28, 29, 30]. It is well established that cold-induced activation of uncoupling protein (UCP)1 on the inner mitochondrial membrane of thermogenic brown adipocytes uncouples oxidative phosphorylation from the generation of ATP, and that the excess chemical energy is dissipated as heat [6]. As a result, fat depots containing an abundance of thermogenic UCP1 containing adipocytes generate heat and expend energy [3, 11]. It is estimated that cold-induced BAT activation could increase resting energy expenditure by at least 2.5-5% [27]. The most superficial thermogenic BAT depot in humans is found in the neck and upper thorax [9], and supraclavicular skin temperature has been used in conjunction with varying cold stimuli (e.g.: localised chest cooling, personalised whole body cooling protocols designed to achieve maximal NST) as a proxy measure of thermogenesis [15, 14, 5, 16]. As a highly metabolic tissue, BAT has a high glucose requirement, and radio-labelled glucose

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uptake is often used as an indicator of BAT activity. 18F-fluorodeoxyglucose (FDG) positron emission tomography (PET) is frequently used for clinical purposes, and warming patients prior to scanning is recommended to reduce FDG uptake in BAT (which may obscure the region of clinical interest [26]). Retrospective studies of clinically indicated FDG-PET scans have identified that BAT is less frequently observed in individuals who have been subject to a range of pre-warming procedures and, prospective BAT dedicated studies confirm an acute BAT response to both warming and cooling [8, 25, 17, 31].

Furthermore, the primary substrates of BAT are lipids, and UCP1 activation is fatty acid-dependent [4]. Glucose uptake is, therefore, a crude measure of BAT activity, and when quantified as standard uptake values (SUV) on static FDG-PET-CT (e.g. SUVmean, SUVmax) may be considered semi-quantitative at best [20]. Importantly, these measures do not quantify thermogenic output. How exposure to warm ambient conditions before exposure to a cold stimulus affects the subsequent heat production from BAT, has not been explored. In this study, we used direct measurements of supraclavicular skin temperature to investigate the basal activity and cold-induced thermogenic response of supraclavicular brown adipose tissue (BAT) under warm (23˚C) and cool (18˚C) ambient conditions. We hypothesised that the basal activity of BAT would be lowest under warm conditions and that the response to mild cold exposure would, therefore, be greater.

Methods

The study was undertaken following the University of Nottingham School of Medicine Ethics Committee approval (Reference no: D052011) during the months of October and November. All participants gave written informed consent to take part, and the study conformed to the standards set by the Declaration of Helsinki 2008, in place at the time.

I Participants

Five healthy non-obese male volunteers aged 18-36 years with a mean body mass index of 24 ± 0.75 kg/m², fat mass of 17.1 ± 2.2 kg and fat free mass of 67.4 ± 4.2 kg took part. Each attended twice after an overnight fast (from midnight). Visits were on separate days and were allocated in random order. Experiments were conducted in a quiet, temperature-controlled laboratory with participants in a semi-reclined supine position. Participants wore standardised clothing consisting of a cotton short sleeved shirt, long cotton trousers and socks (0.35 clo). Participants were asked to avoid alcohol and strenuous exercise in the preceding 24 hours. Participants with any condition, disease or drug known to affect metabolic rate or BAT activity were excluded. Height was measured using a stadiometer to the nearest 0.1cm (Leicester height measure; Child Growth Foundation, Sutton Coldfield, United Kingdom) weight was measured to the nearest 0.1 kg using calibrated portable digital scales, and body composition was determined using bioelectrical impedance using a hand to foot, single-frequency (50 kHz) battery-operated bioimpedance analyser (BIM4; Impedimed P/L, Capalaba, Austria).

II Cooling protocol

Participants attended the laboratory on two mornings, and the cooling protocol was undertaken at room temperatures of 18˚C and 23˚C. Measurements commenced after a minimum of 45 minutes’ stabilisation to the room temperature. After a 15 minute basal period, the left hand was immersed in a 9L bucket of cool water (20˚C) to the level of the ulnar styloid process. Water temperature was checked every 5 minutes, ice water added if it had risen by more than 0.2˚C until temperature was restored to 20˚C, and equal volumes of water removed to maintain the water depth to the level of the ulnar styloid process.

III Energy expenditure

Indirect calorimetry was performed during baseline and single-hand immersion in cool water. Continuous recordings of oxygen consumption and carbon dioxide production were made using a mask collection method (Oro-nasal reusable face mask, V7450 series, Hans Rudolph Inc., Shawnee, USA) with the Europa gas exchange monitor (GEM; Europa Scientific Ltd., Crewe, UK). Resting energy expenditure (REE) was recorded for 30 minutes before, and during, the 30 minutes hand cooling period. Average values for REE during baseline and the 30 minute hand cooling period were calculated from the last 15 minutes of each period.

IV Infrared thermography

Supraclavicular skin temperature was measured using infrared thermography (FLIR B425, thermal resolution 320x240 pixels; FLIR Systems, Danderyd, Sweden) as described previously and used as an indicator of BAT activity [21, 22, 24]. In brief, the camera was positioned so that the lens was perpendicular to the larynx and the field of view included, as a minimum, the full width of the shoulders laterally, the manubriosternal joint inferiorly and angle of the mandible superiorly. The distance from the camera required to achieve this was measured and entered alongside ambient and reflective temperatures into the thermal camera during setup as per the manufacturer’s instructions. Images were taken at 1 minute intervals during baseline and cooling. During thermographic image analysis, a region of interest (ROI) was defined as that bounded by the left sternocleidomastoid muscle, clavicle and lateral contour of the neck using ThermaCAM Researcher Pro 2.10 (FLIR systems AB, Taby, Sweden) as described previously [22, 21]. ROIs were exported into Excel (Microsoft, Redmond, WA, USA) and a custom written script in R (A Language and Environment for Statistical Computing, version 3.4.3 (R Core Team)) was used to calculate the 87.5th percentile temperature value (T roc).

V Mean skin temperature

Mean skin temperature (Tmsk) was evaluated using measurements of skin temperature obtained from wireless data loggers (iButton, model no. DS1219H-F50, resolution 0.125˚C; Maxim, Sunnyvale, CA, USA) placed at seven body sites (i.e. forehead, trunk, arm, hand, lower leg, thigh and foot) and calculated using the Hardy Du Bois formula [13]. Measurements were taken at 1 minute intervals throughout, in parallel to the acquisition of thermograms as described above.

VI Statistical analysis

All analyses were performed using GraphPad Prism for Windows
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Version 7 (GraphPad Software, La Jolla California USA). Data are reported as means ± SEM unless otherwise stated. All data were normally distributed as indicated by the Kolmogorov-Smirnov normality test. Comparison between baseline and 30 minute energy expenditure and skin temperature of the thigh was made using a two-tailed paired t-test. A two-way repeated measures ANOVA was used to determine whether any interaction was present between cold stimulation and ambient room temperature. Where ANOVA analysis revealed a significant F-ratio for the interaction, a post-hoc paired two-tailed t-test was employed to define the simple effect of cold stimulation at each ambient room temperature. A P value < 0.05 was considered to be statistically significant; where comparisons were made at both 18˚C and 23˚C, this threshold was adjusted (using a simple Bonferroni correction) to < 0.025.

**Figure 1:** Effect of single-hand immersion in cool water on supraclavicular temperature at 18˚C and 23˚C ambient room temperature

A significant interaction was observed between ambient temperature and thermogenic response to hand cooling (F (1, 4) = 12.22, P = 0.025), **P < 0.01 following 30 minutes of hand cooling (open circle) compared with baseline (circle); n = 5. TSCR – supraclavicular skin temperature

**Results and Discussion**

TSCR increased following single-hand immersion in cool water at an ambient room temperature of 23˚C (Figure 1). This was accompanied by a small, statistically significant increase in REE from baseline (baseline EE: 4.76 ± 0.30 kJ/min, ∆EE: 0.22 ± 0.02 kJ/min, P = 0.0008) (Figure 2). Neither change was observed in the same group of participants examined at a room temperature of 18˚C. In contrast, skin temperature measurements taken over a central location not overlying BAT (i.e. thigh) remained static at 23˚C (Figure 3), suggesting that this rise in temperature was localised to BAT, rather than secondary to a wider systemic thermogenic response.

**Figure 3:** Skin temperature of the anterior thigh and supraclavicular region prior to and following 30 minutes’ hand cooling at 23˚C **P < 0.01 when compared to baseline, n = 5. Closed circles = baseline, open circles = 30 minutes cooling

Consistent with the well-described insulative response to acute cold exposure, TMSK fell from baseline throughout the duration of cooling [7]. The degree and pattern of this fall was remarkably similar at both ambient room temperatures (Figure 4), suggesting that at both 18˚C and 23˚C hand immersion in cool water acts as an effective cold stimulus of similar magnitude. Skin temperature was, as expected lower at all sites when measured at 18˚C. Moreover, the skin temperature of the non-immersed hand was well above ambient temperature during baseline at 18˚C (mean: 28.01˚C, 95% CI 23.33 to 32.70˚C) and dropped significantly (-0.61˚C, 95% CI -0.93 to -0.29˚C) following immersion of the left hand in water at 20˚C. This is consistent with the recognised vasoconstrictor response of the contralateral hand to indirect cooling, further supporting the efficacy of the cold stimulus at a cooler room temperature [19]. However, the extent to which vasoconstriction within the hand and fingers at 18˚C prior to immersion in water may have attenuated subsequent heat extraction is not certain.

The supraclavicular BAT response to acute localised cold exposure under varying ambient conditions has not been reported before. However, the outcomes of a recent study in a similar study cohort of
non-obese males at 22-23°C support our findings [1]. In that study, acute mild cold exposure prior to the onset of significant involuntary superficial muscle activity (i.e. shivering) resulted in a similar increase in energy expenditure and a similar pattern of reduction in mean skin temperature. However, supraclavicular temperature measured using a single iButton placed within the “supraclavicular zone” rather than with thermography was unchanged throughout [1]. Gashi et al (2018) also reported an isolated increase in TMSK during cooling in contrast to a fall in skin temperature at 7 other sites, in addition they identified a significant positive linear relationship between ΔTMSK and cold induced thermogenesis [12].

Ambient temperature is a determinant of FDG-detected BAT prevalence, and acute cold exposure has been shown to increase BAT activity on PET-CT [23, 25, 31]. Our findings are in line with this but are the first to show a differential effect of ambient temperature on supraclavicular heat production using thermography in response to cold stimulation, which suggests that supraclavicular BAT can be activated with minimal cold exposure. The REE and TMSK response observed at 23°C may have been even greater had we also examined our subjects under confirmed thermoneutral conditions. One explanation for our findings is that BAT at a room temperature of 18°C is already close to its maximal activity. However, REE at baseline was similar at 18°C and 23°C (4.76 ± 0.30 kJ/min and 4.88 ± 0.35 kJ/min respectively), indicating that the overall increase in REE may not be maintained long term. This may reflect a compensatory reduction in energy expenditure from another component of REE or may indicate that cold-induced BAT activation is transient.

Maximising BAT activity by increasing time spent outside of thermoneutrality presents an attractive mechanism for enhancing lipid/glucose metabolism, particularly in the context of diabetes, obesity and their metabolic sequelae. Although the changes in energy expenditure seen following cold stimulation in our study group were small (c. 3-5.5% of baseline REE), if these were sustained over a long period they could contribute significantly to energy balance. For example, assuming that 1 kg of body fat contains 37,000 kJ and a prior neutral energy balance, an increase in EE of 0.22 kJ/min for just 50% of each day would increase daily energy expenditure by 160 kJ, which, if accumulated as WAT over a single year, would equate to around 1.5 kg.

As a pilot study, our main limitation was small sample size. Nonetheless, the magnitude of the response to the mild cool stimulus at 23°C was sufficient to reach statistical significance. How these findings may relate to a larger, more diverse population is unclear and will only be identified by further, more comprehensive studies. Although we did not observe shivering, nor did our subjects report it, EMG measurements would have provided input into the final interpretation. The manuscript was drafted by LJR, MES, IAM and HB. All authors revised the manuscript, approved the final version and agree to be accountable for all aspects of the work. All persons designated as authors qualify for authorship, and all those who qualify for authorship are listed.

Disclosures

The authors have no competing interests.

Abbreviations

BAT – brown adipose tissue
CT – computed tomography
FDG – 18F-fluorodeoxyglucose
NST – nonshivering thermogenesis
PET – positron emission tomography
REE – resting energy expenditure
TMSK – mean skin temperature
TSCR – supraclavicular skin temperature
UCP1 – uncoupling protein 1

Supraclavicular thermography measures infrared radiation from the skin surface, this is determined not only by the heat produced from the deeper structures such as BAT but also by an overlying “insulative layer” consisting of the skin, subcutaneous adipose tissue and also a dynamic cutaneous vasculature network. At present there are no methods to control for this, however, given that cold induced changes in supraclavicular temperature closely approximate to FDG uptake on PET-CT, and also to changes in energy expenditure we speculate that these effects are minimal [12, 14, 15,16]. As chemical-shift water-fat MRI becomes more available for the assessment of BAT, large scale studies in conjunction with perfusion assessment may enable the thermographic evaluation of BAT to be refined.

Conclusion

Our initial findings show that supraclavicular BAT can be activated under warm conditions by a very mild cold stimulus, generating heat in association with an increase in energy expenditure, and that supraclavicular BAT may be maximally stimulated at an ambient room temperature of c.18°C. Whether sustained activation of BAT occurs at lower ambient temperatures under free-living conditions, and whether this can directly impact on overall energy balance, is unknown but presents an exciting avenue for further study.

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Contributions

MES, HB and LJR conceived and designed the study. IAM advised on the final version of the protocol. LJR was responsible for data collection and image analysis, and MES, HB and LJR interpreted the data and IAM provided input into the final interpretation. The manuscript was drafted by LJR, MES, IAM and HB. All authors revised the manuscript, approved the final version and agree to be accountable for all aspects of the work. All persons designated as authors qualify for authorship, and all those who qualify for authorship are listed.
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