The efficacy of transcranial random noise stimulation (tRNS) on mood may depend on individual differences including age and trait mood

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HIGHLIGHTS
- At a group level, transcranial random noise stimulation (tRNS) over bilateral dorsolateral prefrontal cortex did not significantly improve mood across participants.
- Individual differences in age and trait mood affect the direction of mood change in response to tRNS.
- Mood change was comparable in older adults with and without cardiovascular risk factors.

ABSTRACT
Objectives: To assess whether changes in brain microstructures associated with ageing and presence of cardiovascular risk factors (CVRF) reduce the efficacy of transcranial electrical stimulation (tES) improving mood in euthymic older adults.

Methods: Using excitatory high-frequency transcranial random noise stimulation (tRNS) over bilateral dorsolateral prefrontal cortex, the effect on mood was assessed in euthymic young adults (YA), older adults (HOA) and older adults with CVRF (OVR). Active-tRNS or sham was applied over two sessions. Positive and Negative Affect Schedule and Warwick Edinburgh Mental Well-being Scale measured self-reported state mood before and after stimulation. Trait mood was also measured using the Geriatric Depression Scale.

Results: Response to tRNS seemed dependent on individual differences in age and trait mood. In HOA, more negative trait mood was associated with more positive mood change after tRNS. OVR showed a similar but reduced pattern of mood change to HOA. In YA, more positive trait mood was associated with greater positive mood change after tRNS.

Conclusions: Age and trait mood may be important factors when examining the efficacy of tES as an alternative treatment for depression.

Significance: Future studies should consider how response to tES is affected by individual differences.

1. Introduction

Major depressive disorder (MDD) is common among otherwise healthy older adults, occurring in more than 13% of individuals over 60 years old (Beekman et al., 1999). Despite this, less than a quarter of older adults with depression receive treatment (Roose and Schatzberg, 2005). Older adults diagnosed with MDD may be reluctant to start drug therapies and have residual cognitive difficulties even when depression symptoms abate (Alexopoulos, 2005; Barch et al., 2012). Beyond MDD, 15% of older adults are estimated to have sub-clinical levels of depression and persistent low mood (dysthymia) that impairs quality of life and is associated with poor cognitive performance (Barch et al., 2012). Given the ageing population and prevalence of dysthymia, alternate therapies for MDD and dysthymia are vital. Transcranial electrical stimulation (tES), a non-invasive method where a weak electrical current is applied to the head to excite or inhibit neural activity (Jorge and Robinson, 2011) may be an alternative to standard treatments. Preliminary evidence suggests that tES may improve mood in young adulthood but its efficacy among older adults has not been examined. This study will examine the effects of tES on mood in euthymic younger and older adults.

Studies exploring the effects of tES have predominantly used transcranial direct current stimulation (tDCS; tES where a direct
current is applied) and have demonstrated that tES applied to the dorsolateral prefrontal cortex (DLPFC) can improve mood in young adults with and without MDD (Boggio et al., 2008; Brunoni et al., 2014; Ferrucci et al., 2009). The left DLPFC is often targeted for active tES as it may be hypoactive in MDD (Grimm et al., 2008; Koenigs and Grafman, 2009). In young adults with MDD, tDCS of the left DLPFC resulted in reduced depression ratings (Boggio et al., 2008). Among adults with MDD (18–65 years), Brunoni and colleagues (Brunoni et al., 2014) demonstrated a reduction in depressive symptoms (immediately after treatment and at two-week follow-up) when bilateral DLPFC tDCS was applied while a working memory task was performed over 10 sessions. Positive effects of left DLPFC stimulation have been reported in healthy participants, where stimulation facilitated cognitive reappraisal (Feeser et al., 2014) and reduced self-reported emotional distress when viewing aversive stimuli (Boggio et al., 2009). TES has not yet been applied to elevate mood in older adults either with or without MDD.

Whilst tES shows promise, research suggests its effects on mood or cognition may vary depending on the individual. These findings warrant the need to assess whether the effects of tES may differ in late-life compared to young adulthood. Although tES has not yet been applied in a systematic fashion to effect mood in ageing, one case report has been identified. In a 92-year-old-patient with MDD, improved mood was demonstrated after 10 sessions of tDCS without MDD.

Cardiovascular risk factors (CVRF) such as high-blood pressure or diabetes which lead to cardiovascular damage in the brain, are common in ageing (Kennedy and Raz, 2009; Knopman et al., 2011; Raz et al., 2007) and may also influence the efficacy of tES. CVRF have been shown to disrupt white matter microstructure that support neural networks involved in cognition and mood regulation (Alexopoulos, 2005; Taylor et al., 2013; Kennedy and Raz, 2009; Verdelho et al., 2007). Age-related disruption in white matter microstructure has been demonstrated throughout the brain but particularly in anterior regions including the DLPFC where degradation is associated with late-life depression (Bae et al., 2006; O’Sullivan et al., 2001). It is not yet clear whether cardiovascular damage to white matter may further influence the efficacy of tES or the spread of stimulation across the affected neural networks in older adults. Therefore the current study will also examine CVRF and its impact on response to tES.

In this study we examine the efficacy of tRNS applied bilaterally to the DLPFC (implicated in mood regulation by previous research) to improve mood in older euthymic adults with and without CVRF, and young adults. Participants received both active and sham tRNS using a double-blind design; presence of CVRF among older adults was measured. We hypothesised that: (1) tRNS may be as effective in older compared to younger euthymic adults and (2) that efficacy of tRNS among older adults would be associated with fewer CVRF. We also sought to explore whether individual differences in trait mood levels (i.e. long-term mood characteristics) influences response to tRNS.

2. Methods

2.1. Participants and procedure

Participants: recruitment was conducted through community outreach and local newspaper advertising. Exclusion criteria included any contraindications to tRNS (Rossi et al., 2009) and current or recent history of depression, presence of psychiatric or neurological conditions. These criteria minimised the variance in baseline cortical excitability introduced by psychological or psychiatric problems, which can moderate and potentially reverse the anticipated excitatory effect of tES (Krause and Cohen Kadosh, 2014). Furthermore, since the efficacy of tES has not yet been established, an initial study on euthymic individuals was deemed a necessary step prior to interventions with patients, which may have negative effects. Ten people were excluded from the study at initial screening (current/recent history of depression
n = 8; contraindication to tRNS n = 1; reason not given for ineligibility n = 1). Sixty older adults aged over 60 years (Mage = 67.33 ± 6.7, 25 male) and 30 younger adults (YA) aged 20–40 years (M_age = 26.37 ± 5.3, 14 male) participated. Older adults (OA) were divided into two groups, healthy older adults (HOA) and older adults with CVRF (OVR). OVR had a diagnosis of hypertension, diabetes, or had a high average blood pressure reading (>140 systolic and/or >90 diastolic) at the time of testing. A modified version of the Framingham Stroke Risk Profile excluding age as a risk factor (FSRP; Wolf et al., 1991) was recorded to quantify CVRF. See Table 1 for demographic information. All participants gave written informed consent and were given an honorarium of up to £40 for taking part. The study was approved by Goldsmiths University of London Ethics committee.

2.2. Assessment

**Background Measures:** depressive symptoms were measured using the Geriatric Depression Scale (GDS; Yesavage, 1988) in both sessions of the study; GDS scores quantified trait mood whereby a higher score indicated lower trait mood. The scale was administered to all participant groups as it has been shown to be appropriate for use with young adults (Weintraub et al., 2007). Whilst trait mood was expected to be stable, the GDS was measured in both sessions to rule out any anomalous changes in mood. The Mini-mental state exam (MMSE; Molloy et al., 1997) was used as a screen for cognitive impairment across participants (at risk <24); all participants scored above this cut-off (see Table 1). The Wechsler Test of Adult Reading (WTAR; Wechsler, 2001) was used to estimate full-scale IQ (FSIQ).

**Experimental Measures:** in both sessions, participants completed the Positive and Negative Affect Schedule (PANAS; Watson et al., 1988) and Warwick-Edinburgh Mental-Well-being Scale (WEMWBS; Tennant et al., 2007) before and after stimulation (active-tRNS/sham). Participants completed each questionnaire in relation to how they felt at that moment in time to quantify cur-

2.3. Transcranial random noise stimulation (tRNS)

After completing the mood questionnaires, stimulation was applied using a DC-Stimulator Plus device (neuroConn, Germany). During active-tRNS, 1 mA of high-frequency stimulation was applied for 10 min. The duration of after effects was unknown, yet based on previous research were anticipated to be shorter than the 60 min observed when targeting the motor cortex (Nitsche and Paulus, 2011). Two 5 x 5 cm (25 cm²) rubber electrodes were encased in saline-soaked sponges and positioned bilaterally over the DLPFC (identified using F3 and F4 based on the 10-20 system as per previous studies (Boggio et al., 2008; Brunoni et al., 2014; Shiozawa et al., 2014b). Stimulation was ramped up and down for 15 s at the beginning and end of stimulation. During sham, the current was applied for 30 s after ramping up before switching off. Such a short duration of tRNS has been shown to have no effect on cortical excitability (Ambrus et al., 2010; Kadosh, 2013) and to be indistinguishable from active-tRNS (Ambrus et al., 2010; Kadosh, 2013).

2.4. Procedure

All participants attended two sessions where they received active-tRNS or sham stimulation using a double-blind procedure. Sessions were between 2 and 14 days apart (M = 5.9 days ± 4.2). Stimulation type (active-tRNS/sham) was randomised and counterbalanced across participants. In the first session participants were given a brief interview to assess the presence of any CVRF and completed the cognitive tests. Participants also identified two positive memories, one of which would be recalled and described to the researcher in each session. In both sessions, participants completed the GDS to describe trait mood followed by the pre-stimulation PANAS and WEMWBS describing state mood. During the first five minutes of stimulation participants sat quietly and for the remaining five minutes participants described one of the identified positive memories in order to induce a more positive mood. Immediately after stimulation, participants completed the post-stimulation PANAS and WEMWBS.

2.5. Data analysis

As presence of depression was an exclusion criterion for the study, GDS scores were reviewed. One YA was removed from the analyses having scored 23 (severe depression) on the GDS. Among the remaining participants a maximum of 15/30 was recorded on the GDS (criteria for mild depression score >10) with an average score of 4.13 (SD = 3.87). Grubbs’ test (Grubbs, 1969) for outliers on Mood Change identified three participants (two HOA; one YA) as outliers who were removed from the analyses. As GDS scores were not normally distributed (Shapiro-Wilk test = .864, p < .001) they were converted into z scores for use in the analyses. This was expected due to our exclusion criteria. Group differences in demographic variables were analysed using ANOVA or Chi-Square as appropriate.

The effect of stimulation (active-tRNS/Sham) on mood state was assessed using a Mood Change difference score calculated by deducting the total score for positive affect/well-being prior to stimulation from the total score after stimulation: Mood Change = Mood score After Stimulation – Mood score Before Stimulation. A positive score indicated an improvement in mood state after stimulation.

<table>
<thead>
<tr>
<th>Table 1</th>
<th>Demographics for each participant group, mean (standard deviation).</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>YA (n = 28)</td>
</tr>
<tr>
<td>Sex (M:F)</td>
<td>13:15</td>
</tr>
<tr>
<td>Age</td>
<td>26.68 (5.4)</td>
</tr>
<tr>
<td>Education level</td>
<td>4.50 (1.2)</td>
</tr>
<tr>
<td>Trait Mood (GDS)</td>
<td>5.32 (3.5)</td>
</tr>
<tr>
<td>MMSE</td>
<td>29.57 (0.7)</td>
</tr>
<tr>
<td>FSFP % Age</td>
<td>1.79 (1.3)</td>
</tr>
<tr>
<td>WTAR FSIQ</td>
<td>109.89 (6.6)</td>
</tr>
</tbody>
</table>

* p < .05.
** p < .001. –: comparable between groups; <: significant difference between adjacent groups.

Education level coding: 0 = no qualification; 1 = GCSE/NVQ (16 yrs); 2 = City & Guilds or other post 16 qualification; 3 = A Level/BTEC/Access course; 4 = Diploma/HND; 5 = Degree; 6 = Masters; 7 = PhD.
and a negative score indicated a reduction in mood state after stimulation. A repeated-measures ANOVA assessed Mood Change across the whole participant sample across Stimulation conditions (active-tRNS/Sham). Mixed model ANOVAs explored whether Mood Change differed depending on Stimulation and Age (YA/OA), and depending on Stimulation and Age/CVRF (YA/HAO/OVR). A Bonferroni correction was applied to post hoc analyses to correct for multiple comparisons.

To assess the effect of individual differences on response to stimulation, separate multiple regression analyses were conducted for PANAS-Positive and WEMWBS Mood Change scores. These were performed for each sham and active-tRNS sessions using the enter method to examine the relationship between Mood Change and four potential predictors: Trait Mood (GDS), Age, FSRP (%), and the interaction term Group × GDS score (YA × GDS; HAO × GDS; OVR × GDS).

3. Results

3.1. Demographics & GDS scores between sessions

3.1.1. Group differences

Participant groups were comparable in terms of sex, education level, and cognition (MMSE, FSIQ). As expected, YA were significantly younger than the HOA and OVR. Similarly, OVR demonstrated significantly higher CVRF than YA and HOA when quantified with the FSRP. YA had significantly lower trait mood compared to HOA and OVR participants, see Table 1. GDS scores measuring trait mood did not differ significantly between sessions for the whole sample (t(85) = -1.057, p = .293) or each group: HOA (t(27) = -9.57, p = .347), OVR (t(28) = -1.092, p = .284), or YA (t(29) = -4.29, p = .671). The relationship between GDS score and baseline mood (positive affect and well-being scores pre-stimulation) was also consistent across all groups in both active-tRNS and sham sessions; HOA, OVR, and YAs showed a significant negative correlation between GDS and baseline well-being, and non-significant negative correlation between GDS and baseline positive affect (data not shown).

3.2. Effect of active-tRNS/sham on state mood

ANOVA assessed whether tRNS improved positive affect/well-being across the whole sample, and whether the effect of tRNS on state mood differed depending on age (YA/OA) or presence of cardiovascular risk factors (YA/HAO/OVR).

3.2.1. PANAS-positive affect ANOVAs

Whole sample: repeated-measures ANOVA confirmed a non-significant main effect of Stimulation (active-tRNS/sham) indicating that change in positive affect was consistent between active-tRNS (M = 1.29 ± 4.35) and sham (M = .85 ± 4.06) conditions (F(1,85) = .611, p = .437, ηp² = .007). YA and OA: mixed model ANOVA demonstrated non-significant main effects for Stimulation (F(1,84) = .380, p = .539, ηp² = .005), Age (F(1,84) = .443, p = .507, ηp² = .005), and interaction Stimulation × Age (F(1,84) = .104, p = .748, ηp² = .001) indicating that Mood Change was comparable for both YA and OA in both active-tRNS and sham conditions.

YA, HAO, and OVR: non-significant main effects for Stimulation (F(1,35) = .567, p = .453, ηp² = .007), Age/CVRF group (F(2,83) = .234, p = .792, ηp² = .006), and interaction Stimulation × Age/CVRF (F(2,83) = .199, p = .820, ηp² = .005) confirmed Mood Change did not differ depending on stimulation condition, age, or presence of cardiovascular risk factors.

3.2.2. WEMWBS ANOVAs

Whole sample: change in well-being was comparable between active-tRNS (M = 1.40 ± 4.08) and sham (M = 1.44 ± 3.52) conditions (F(1,85) = .008, p = .927, ηp² < .001) across all participants.

YA and OA: non-significant main effects of Stimulation (F(1,84) = .017, p = .896, ηp² < .001), Age (F(1,84) = .446, p = .506, ηp² = .005), and interaction Stimulation × Age (F(1,84) = .017, p = .896, ηp² < .001) confirmed that well-being did not significantly differ depending on stimulation condition or age.

YA, HAO, and OVR: Mood Change did not differ depending on the age of the participants or presence of cardiovascular risk factors as demonstrated by non-significant main effects of Stimulation (F(1,83) = .005, p = .943, ηp² < .001), Age/CVRF (F(2,83) = .237, p = .790, ηp² = .006), and interaction Stimulation × Age/CVRF (F(2,83) = .309, p = .735, ηp² = .007).

3.3. Effect of individual differences on mood change

Although non-significant effects of tRNS on Mood Change was observed at a group level, regression analyses assessed whether individual differences influenced participants response to tRNS. These were performed for each sham and active-tRNS sessions using the predictors age, trait mood (GDS), cardiovascular risk (FSRP%), and interaction terms YA × GDS, HOA × GDS, and OVR × GDS.

3.3.1. PANAS-positive affect regressions

Active-tRNS: the model significantly explained 14.4% of the variance in Mood Change (F(5,80) = 2.702, p = .026, R² = .144, R²Adjusted = .091) with the interaction terms YA × GDS (β = -2.450, t(80) = -2.596, p = .011) and HOA × GDS (β = 2.129, t(80) = 2.184, p = .032) contributing significantly to the model. In YA, mood reduced by 2.45 points for each point on the GDS; the lower trait mood – the greater negative change in mood state after active-tRNS. In HOA, mood increased by 2.18 with each point on the GDS; the lower trait mood – the more positive change in mood state after active-tRNS. Age, GDS, FSRP, and OVR × GDS did not contribute significantly to the model (p ≥ .219). Fig. 1 indicates that both HOA and OVR demonstrate a similar association between Mood Change after tRNS and GDS.

Sham: the model did not significantly explain any variance in Mood Change after sham (F(5,80) = .738, p = .597, R² = .044, R²Adjusted = -.016).

Including baseline state mood: to confirm that baseline state mood was not driving the results, positive affect score pre-stimulation was included as an additional predictor. Results confirmed that baseline state mood did not significantly contribute to either model. During active-tRNS, the model significantly explained 14.4% of the variance in Mood Change (F(6,79) = 2.275, p = .045, R² = .147, R²Adjusted = .083). During sham, the model did not significantly explain variance in Mood Change (F(6,79) = .644, p = .695, R² = .047, R²Adjusted = -.026).

3.3.2. WEMWBS regressions

Active-tRNS: the model significantly explained 16.8% of the variance in well-being change (F(5,80) = 3.241, p = .010, R² = .168, R²Adjusted = .116); the interaction term YA × GDS (β = -2.941, t(80) = -3.370, p = .001) contributed to the model. In YA, mood reduced by 2.94 points for each point on the GDS. The lower trait mood scores, the greater the negative change in mood state after tRNS. Age, GDS, FSRP, HOA × GDS, and OVR × GDS did not contribute significantly to the model (p ≥ .194).

Sham: Variance in Mood Change after sham stimulation was not significantly explained by the model (F(5,80) = 1.434, p = .221, R² = .082, R²Adjusted = .025).
Including baseline state mood: the inclusion of well-being score pre-stimulation as an additional predictor confirmed that baseline state mood did not significantly contribute to either model. During active-tRNS, the model significantly explained 17.0% of the variance in Mood Change ($F_{(6,79)} = 2.687, p = .020, R^2 = .170, R^2_{Adjusted} = .103$), whereas during sham, the model did not significantly explain variance in Mood Change ($F_{(6,79)} = 1.183, p = .324, R^2 = .082, R^2_{Adjusted} = .013$).

4. Discussion

This study examined the efficacy of tRNS to improve mood and whether this differs depending on individual differences in euthymic younger and older adults with and without CVRF. Results from two measures (PANAS-positive affect & WEMWBS) indicated that active-tRNS did not improve positive affect or well-being across all participants or within each participant group (YA/HOA/OVR), suggesting that tRNS was not sufficient in facilitating greater positive mood than a positive memory mood induction alone. Notably however, the patterns of response to active-tRNS appeared to differ depending on individual differences in age and trait mood. A small but significant proportion of the variance in positive affect after active-tRNS was explained by interaction terms for trait mood in the YA and HOA group. This trend was also observed for well-being, where the interaction term of trait mood and YA group reached significance. These results suggest that age and trait mood may be important variables affecting the efficacy of tES and are in line with a growing body of research emphasising the role of individual differences on the effect of stimulation.

Among older adults, individuals with lower trait mood reported the most positive change in mood after active-tRNS stimulation. This association was significant when mood was measured in terms of positive affect but did not reach significance for well-being. The relationship between trait mood and mood change after active-tRNS is positive and in keeping with some of the tDCS...
results in YA with MDD (Boggio et al., 2008; Brunoni et al., 2014). This relationship, whereby HOA at most risk may show the greatest benefit, also follows a similar pattern to previous studies of tDCS applied to improve cognitive performance, where a relationship between baseline performance and response to stimulation has been observed (Berryhill and Jones, 2012; Learmonth et al., 2015). One explanation for these results is that those with the lowest mood have the greatest capacity for positive change. However, as the scores on all measures were not at ceiling for any group, all participants could have reported positive change in mood. Although the overall positive trait mood ratings were higher in the HOA group compared to YA, GDS scores alone did not significantly contribute to explaining the variance in tRNS response, suggesting that additional factors including age are influencing the effects of tRNS on mood. Coupled with the non-significant effect of tRNS on mood observed at a group level, the influence of age and trait mood on response to tRNS emphasises the importance of examining how individual differences may impact the efficacy of tES as a treatment for depression.

Whilst HOA with poorer trait mood showed the most positive change after tRNS, YA displayed the opposite pattern. In the current euthymic YA sample, applying tRNS bilaterally to DLPFC seemed to reduce mood state in those with lowest trait mood for both mood measures, positive affect and well-being. Yet, YA reporting more positive trait mood showed greater positive change. This pattern of results differs to previous studies that have shown reduced depressive symptoms in young adults with MDD when applying 1–2 mA of tDCS to the left DLPFC for approximately 20 min (Boggio et al., 2008; Ferrucci et al., 2009; Pérez et al., 2016).

It is possible that the relationship between tRNS and mood change in this euthymic adult sample is due to the use of bilateral DLPFC tRNS, which contrasts previous studies of young adults and those described here. For YA bilateral DLPFC tRNS could intensify the trait mood of the individual, therefore stimulation could be positive or negative depending on current mood (Feerer et al., 2014; Möbius et al., 2017). Due to reduced task-specific hemispheric asymmetry observed in older adults (Cabeza, 2002), bilateral stimulation was selected in an attempt to optimise the protocol for older adults. The relationship between trait mood and positive effects of tRNS observed in HOA in the current study support this theory. However it is important to note that it may be detrimental to young adults who do not typically rely on bilateral activation (Herrington et al., 2005). Bilateral tRNS to the DLPFC may have a different mechanism and effects compared to anodal tDCS over the left DLPFC (the most commonly used stimulation protocol). Only one previous case study in a 35-year-old woman with depression has demonstrated improvements in mood using bilateral DLPFC tRNS, although the cathode electrode was placed over F8 rather than F4 (Chan et al., 2012). The optimum stimulation protocol may therefore differ between younger and older adults.

As well as bilateral stimulation potentially having differing effects on younger and older adult populations, the effects of tRNS in this euthymic sample may be due to differences in the severity of depressive symptoms. However the regression model suggests that it is the interaction between depressive symptoms and age that is important. It is also worth noting that all groups described here reported normal mood levels and the effects of tRNS may differ in MDD or late-life depression. High-frequency tRNS may be more effective for improving mood in severe low mood where criteria for MDD are met. Research by Nitsche et al. (2012) suggest the effect of tDCS may be limited to emotion processing and not self-reported mood in healthy participants. Nevertheless, facilitatory effects have been noted elsewhere in the literature when using tDCS in healthy populations. tDCS has been shown to facilitate cognitive reappraisal both positively and negatively depending on the regulatory goal (Feerer et al., 2014). Although not a study of mood per se, Boggio and colleagues (Boggio et al., 2009) also demonstrated that healthy young adults reported less “unpleasantness” when viewing aversive stimuli, suggesting that lower mood is not necessary for tES to have an elevatory effect. It is worth noting that studies in MDD have reported mixed results, with some studies not demonstrating improvements in mood after tDCS (Nitsche et al., 2012; Palm et al., 2012), so other factors apart from depressive symptoms may influence response to stimulation (Loo et al., 2012).

One major issue raised by these results is the suggestion that individual differences including age and the level of depressive symptoms as well as interactions between such factors may influence response to tES for the elevation of mood. This is an important consideration for future studies using tES to modulate mood, as it highlights the importance of individual differences which may need careful consideration to optimise stimulation outcomes (Fertonani et al., 2016). Inter-individual differences in neurophysiology, anatomy, and psychology have been reported to result in substantial within- and between-group variances in response to tDCS (for reviews see Horvath et al., 2014; Krause and Cohen Kadosh, 2014). Though speculative, results suggest that individual differences may explain response to tES even in the absence of differences at the group level; when inter-individual variability can result in a different or even opposing response to tES, regressing to the mean of the sample may collapse any effect of tES that is occurring at an individual level. Another issue raised by the discrepancy in mood change across YA and HOA is the need to explore the anatomical specificity of these effects by comparing mood change across different electrode montages including stimulation of unilateral or bilateral DLPFC, and a control site. This was not explored in the current study as the focus was on individual differences and the efficacy of tRNS.

Another purpose of this study was to examine the effects of cardiovascular risk on response to tRNS in ageing. There were no significant differences between response to tRNS in the HOA compared to the OVR group, although the magnitude of the effect were lower in OVR. Although preliminary, the comparable relationship between age, trait mood, and response to tRNS suggest that tRNS may be as efficacious in OVR as in HOA. It is worth noting however that the OVR group described here does not include severe cardiovascular disease such as stroke, transient ischaemic attacks, or chronic obstructive pulmonary disease. Only one individual had diabetes (an additional participant was borderline), and 13 had diagnosed hypertension, with most individuals controlling their CVRF through diet and medication. Thus it remains to be established if the pattern reported here in individuals with relatively mild CVRF is comparable with more severe or uncontrolled cardiovascular disease. TNS efficacy may reduce with severity of cardiovascular disease due to severity of damage in brain and disruption of neural networks (Alexopoulos, 2005). A limitation of this study was the lack of information about cardiovascular damage in the brain among older adults. Although one commonly observes strong associations between CVRF and cardiovascular damage in brain, individual differences do occur (Charlton et al., 2014). Future studies should aim to quantify cardiovascular damage in brain to fully explore the effect of cardiovascular disease on tES. This is particularly important if treatment controlled cardiovascular disease reduces the commonly associated cardiovascular damage.

It is important to acknowledge a number of study limitations that may also help understand the results observed in the current study. In the tRNS protocol, individuals received stimulation in a single session, which has been argued to be insufficient to affect mood in euthymic participants (for review see Remue et al., 2016). Mood was measured using self-report measures, which may be relatively insensitive to small changes in mood; using a
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