

Non-invasive brain stimulation relieves phantom limb pain in amputees; an fMRI study

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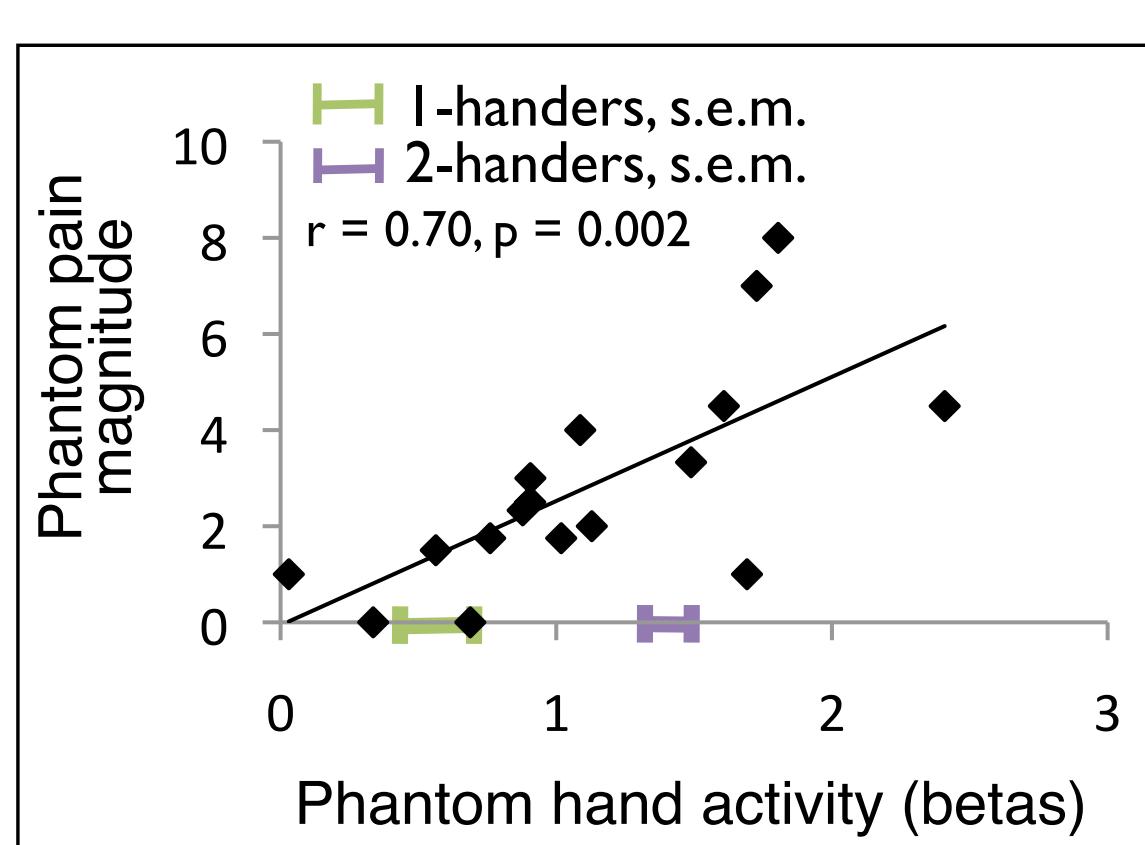
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Background

- Up to 80% of upper-limb amputees suffer from chronic phantom limb pain.
- Current analgesic treatments are ineffective.
- Neuropathic pain may be relieved using non-invasive brain stimulation (tDCS).
- We previously showed that chronic phantom limb pain associated with stronger sensorimotor phantom hand representation (see figure).



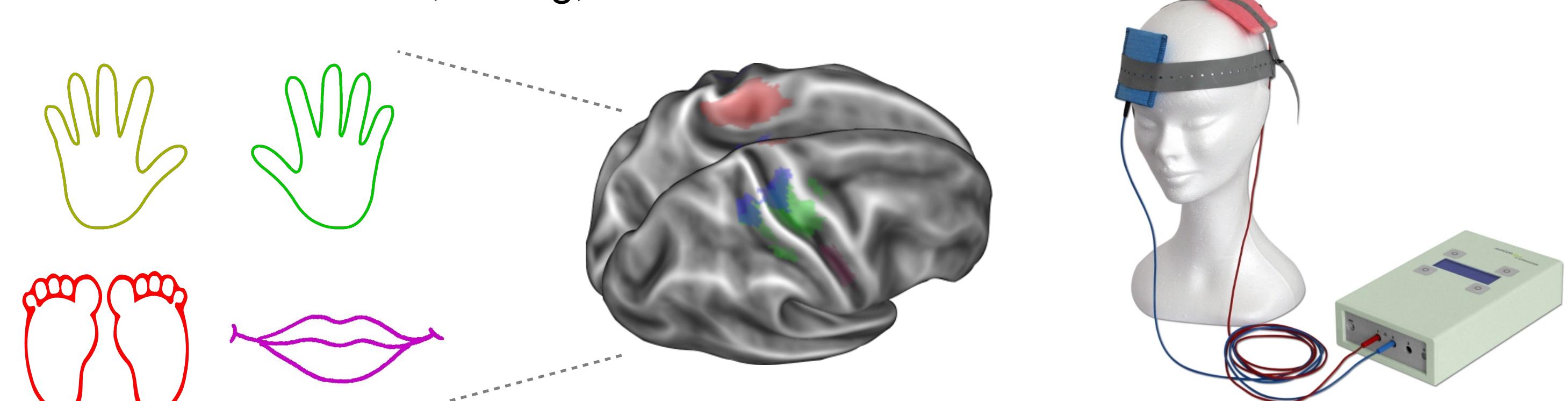
Makin et al., (2013), Nature comm.
Replicated in: Kikkert et al., (Under revisions).

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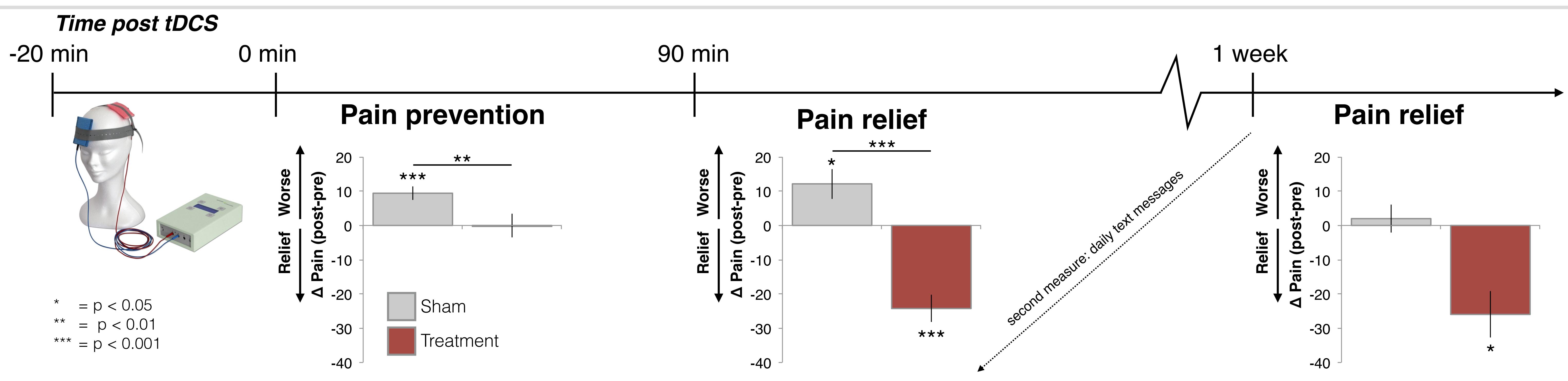
- Can we relieve phantom limb pain using non-invasive stimulation?
- What are the neural mechanisms underlying phantom limb pain relief?

Experimental set-up

- 15 unilateral upper limb amputees suffering from chronic phantom limb pain.
- Sham-controlled, counterbalanced and double-blind design.
- 20 minutes of excitatory non-invasive brain stimulation (anodal tDCS; 1mA) over deprived primary sensorimotor cortex, during phantom hand movements.
- Subjective pain ratings before and after brain stimulation.
- 3 tesla fMRI before, during, and after stimulation.

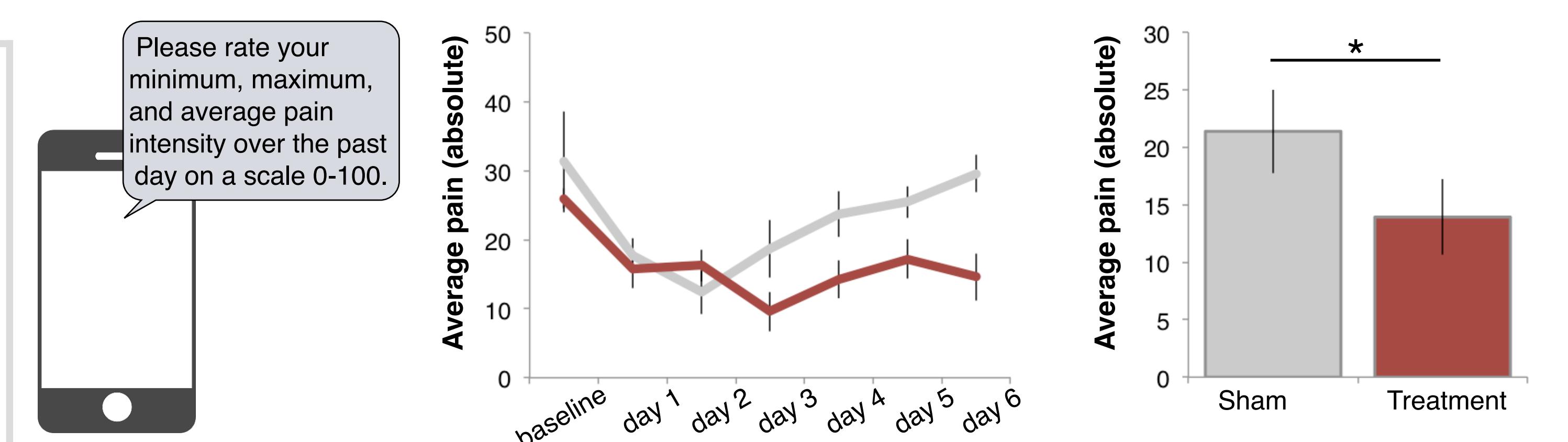
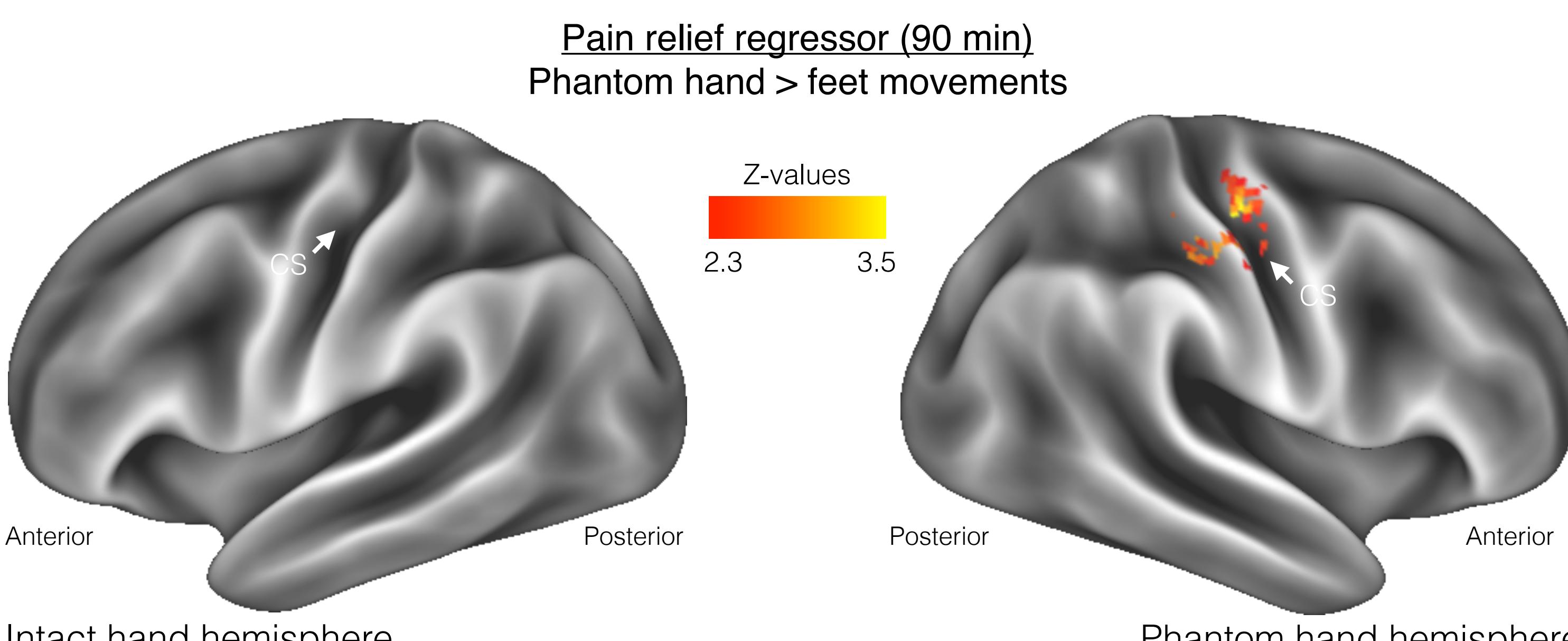


Non-invasive brain stimulation causes lasting phantom limb pain relief



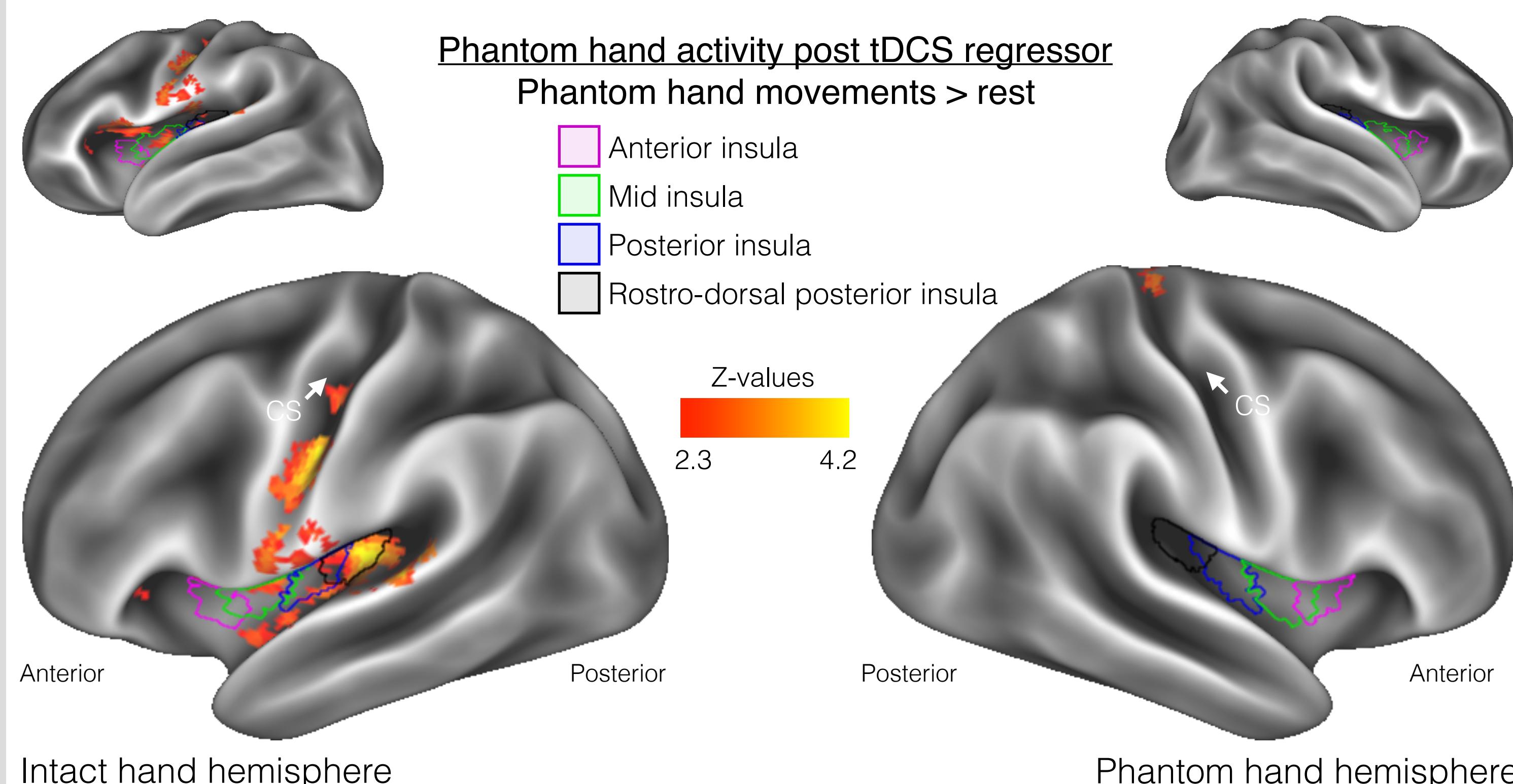
What are the neural correlates of pain relief post tDCS?

Amputees experiencing less phantom limb pain post treatment, activate the primary sensorimotor (S1/M1) phantom hand area less post treatment.



What happens during tDCS to modulate S1/M1 activity?

Amputees who activate the phantom hand area less after tDCS, engage the mid and posterior insula more during tDCS.



Conclusions

- tDCS is a promising tool for managing phantom pain: A single 20 min treatment of non-invasive brain stimulation can relieve phantom pain for at least a week.
- Reduced activity in the cortical phantom hand area reflects reduced phantom limb pain after treatment.
- Both reduced activity in the cortical phantom hand area and phantom pain relief rely on increased insula activity during treatment.

Our findings highlight the role of pain-related networks in mediating phantom pain.