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Relationship between interoception and stress in patients with Functional

Neurological Symptom Disorder

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Objectives / aims

Self-report studies of alexithymic traits in individuals with Functional Neurological Symptom

Disorder (FND), suggest that emotion dysregulation in this population is characterised by an

impaired ability to detect and identify their own emotions (identification impairments) (1). This

regulatory deficit might be particularly problematic for a patient group with an increased

incidence of stressful life events relative to healthy controls (2), for whom the ability to

regulate emotions might therefore be more crucial. Examining sensitivity to changes in

physiological cues associated with emotional experience (interoception) is a way of assessing

one aspect of participants' capacity to identify their own emotions. However, no studies have

yet experimentally investigated how stress might interact with interoception in this population.

Therefore, the aim of this study was to investigate patients' interoceptive sensitivity both at

baseline and under stress.

Methods

Twenty-six patients with FND and twenty-seven healthy controls performed the Heartbeat

Detection Task (HBDT) pre- and post- stress-induction with the Cold Pressor Test. The HBDT

is a behavioural paradigm, measuring participants' sensitivity to a physiological cue associated

with emotional experience - the heartbeat. Participants also completed a self-report measure of

emotion dysregulation (The Emotional Processing Scale-25) which includes a subscale

capturing 'a detached experience of one's emotions due to poor emotional insight', and a

measure of Major Depressive symptomology (The PHQ-9).

Results

Relative to healthy controls, patients with FND performed more poorly on the HBDT both at

baseline and following stress-induction (p = .032). Patients also reported greater impairments

across all domains of the EPS-25 and higher scores on the PHQ-9 than healthy controls (both

p <.001). Group differences on HBDT performance were not explained by group differences

in age or depressive symptomology.

Conclusions

Impaired HBDT performance suggests that patients with FND lack sensitivity to their heartbeat, both under 'normal' conditions and following stress-induction. Physiological cues (like the heartbeat) are an important source of interoceptive information for emotional experience, for example during stress. Our findings therefore represent a form of identification impairment that may contribute to stress-vulnerability in this population. Raised levels of self-reported 'impoverished emotional experience' corroborate the suggestion that patients with FND have difficulty identifying and understanding their emotions. These findings have direct implications for understanding and treating emotion dysregulation in FND.

References

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