

## Supersensitive connection causes hatred of noises

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A supersensitised brain connection has been identified in people who suffer from misophonia, an extreme reaction to "trigger" sounds.

For the first time, researchers led by Newcastle University, have discovered increased connectivity in the brain between the <u>auditory cortex</u> and the motor control areas related to the face, mouth and throat.

Publishing today, in the *Journal of Neuroscience*, lead author Dr. Sukhbinder Kumar, Newcastle motor con University Research Fellow in the Biosciences trigger sou Institute said: "Our findings indicate that for people with misophonia there is abnormal communication between the auditory and motor brain regions—you condition. could describe it as a 'supersensitised connection'.

"This is the first time such a connection in the brain also found a similar pattern of communication between the visual and motor regions, which

Misophonia, which means literally 'hatred of sound', is a condition in which sufferers experience intense and involuntary reactions to certain sounds made by other people, referred to as 'trigger'

sounds. Trigger sounds are often the sound of someone chewing, breathing or speaking and for sufferers, usually related to mouth, throat or facial activity.

Their reaction is often extreme, and tends to consist of a combination of anger, disgust, fight-or-flight response, sometimes an urge to hurt the person making the sound or to leave the situation.

The condition is common affecting anywhere between 6% to 20% of people. Those with the more severe forms can find themselves unable to tolerate family, work, public or social situations.

Previously, misophonia had been considered a disorder of sound processing. This new research suggests that alongside this there is an abnormal type of communication between the brains hearing center, the auditory cortex, and the areas of the ventral pre-motor cortex that are responsible for movement of the face, mouth and throat.

In response to trigger or neutral sound, scans on people with misophonia showed that the brain's auditory cortex (hearing center) responded similarly to people without the condition, however, people with misophonia showed increased communication between the auditory cortex and the motor control areas related to the face, mouth and throat. These motor control regions were strongly activated by trigger sounds in people with misophonia in response only to their trigger sounds, but not to other sound types or in people without the condition.

Dr. Kumar adds: "What surprised us was that we also found a similar pattern of communication between the visual and motor regions, which reflects that misophonia can also occur when triggered by something visual.

"This lead us to believe that this communication activates something called the 'mirror system',



which helps us process movements made by other individuals by activating our own brain in a similar way—as if we were making that movement ourselves.

"We think that in people with misophonia involuntary overactivation of the mirror system leads to some kind of sense that sounds made by other people are intruding into their bodies, outside of their control.

"Interestingly, some people with misophonia can lessen their symptoms by mimicking the action generating the trigger sound, which might indicate restoring a sense of control. Using this knowledge may help us develop new therapies for people with the condition."

Tim Griffiths, Professor of Cognitive Neurology at Newcastle University, who is a senior author on the study and also a neurologist, added: "The study provides new ways to think about the treatment options for misophonia. Instead of focussing on sound centers in the brain, which many existing therapies do, effective therapies should consider motor areas of the brain as well."

The team will be further investigating whether this understanding can help develop more effective treatments for misophonia in future.

**More information:** Sukhbinder Kumar et al, The motor basis for misophonia, *The Journal of Neuroscience* (2021). <u>DOI:</u> 10.1523/JNEUROSCI.0261-21.2021

Provided by Newcastle University

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