Audiological and Other Factors Predicting the Presence of Misophonia Symptoms Among a Clinical Population Seeking Help for Tinnitus and/or Hyperacusis

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This paper evaluates the proportion and the audiological and other characteristics of patients with symptoms of misophonia among a population seeking help for tinnitus and/or hyperacusis at an audiology clinic ($n = 257$). To assess such symptoms, patients were asked “over the last 2 weeks, how often have you been bothered by any of the following problems? Feeling angry or anxious when hearing certain sounds related to eating noises, lip-smacking, sniffing, breathing, clicking sounds, tapping?”. The results of routine audiological tests and self-report questionnaires were gathered retrospectively from the records of the patients. Measures included: pure tone audiometry, uncomfortable loudness levels (ULLs), and responses to the tinnitus impact questionnaire (TIQ), the hyperacusis impact questionnaire (HIQ), and the screening for anxiety and depression in tinnitus (SAD-T) questionnaire. The mean age of the patients was 53 years (SD = 16) (age range 17 to 97 years). Fifty four percent were female. Twenty-three percent of patients were classified as having misophonia. The presence and frequency of reporting misophonia symptoms were not related to audiometric thresholds, except that a steeply sloping audiogram reduced the likelihood of frequent misophonia symptoms. Those with more frequent misophonia symptoms had lower values of ULLmin (the across-frequency average of ULLs for the ear with lower average ULLs) than those with less frequent or no reported symptoms. The reported frequency of experiencing misophonia symptoms increased with increasing impact of tinnitus (TIQ score $\geq 9$), increasing impact of hyperacusis (HIQ score $> 11$), and symptoms of anxiety and depression (SAD-T score $\geq 4$). It is concluded that, when assessing individuals with tinnitus and hyperacusis, it is important to screen for misophonia, particularly when ULLmin is abnormally low or the TIQ, HIQ or SAD-T score is high. This will help clinicians to distinguish patients with misophonia, guiding the choice of therapeutic strategies.

Keywords: misophonia, hyperacusis, hearing loss, tinnitus, uncomfortable loudness levels
INTRODUCTION

Tinnitus is the perception of sound without an acoustical source external to the body. Hyperacusis is intolerance of certain everyday sounds, which are perceived as too loud or uncomfortable and cause significant distress and impairment in the individual’s day-to-day activities (Aazh et al., 2016, 2022a). Misophonia is characterized by a decreased tolerance for specific sounds (Jastreboff and Jastreboff, 2002; Brout et al., 2018; Swedo et al., 2022). These sounds are known as “triggers,” and they are usually man or animal-made sounds, and often orofacial sounds (generated by the mouth and nose), such as sniffing and chewing. In addition, there is evidence to suggest that, regardless of the source of the triggers, they share similar properties, including repetition (Brout et al., 2018; Erfanian et al., 2019; Enzler et al., 2021b; Hansen et al., 2021). People with misophonia may also be intolerant of certain visual and tactile stimuli (Kumar et al., 2017, 2021; Rouw and Erfanian, 2018; Schroder et al., 2019; Eijsker et al., 2021b). It may be the case that the action of the trigger-producing person is what causes the reaction, rather than the sound itself (Kumar et al., 2021).

The reported prevalence of misophonia varies from 6 to 19%, although a prevalence as high as 37% has been found (Wu et al., 2014; Zhou et al., 2017; Naylor et al., 2021). The prevalence depends on the population studied and on the way that misophonia is diagnosed; the prevalence differs markedly across populations with and without co-morbid disorders. A growing body of literature shows co-morbidity of misophonia with a range of affective disorders as diagnosed in mental health settings, such as major depressive disorder (MDD), obsessive-compulsive personality disorder (OCPD), and post-traumatic stress disorder (PTSD) (Schroder et al., 2013; Rouw and Erfanian, 2018; Erfanian et al., 2019; Jager et al., 2020b), and developmental disorders like attention-deficit hyperactivity disorder (ADHD) and autism spectrum disorder (ASD) (Jager et al., 2020a; McKay and Acevedo, 2020; Haq et al., 2021). The overlapping symptomology of misophonia and psychiatric, developmental, and audiological disorders makes the diagnosis and treatment complicated.

Although auditory disorders, including tinnitus and hyperacusis, often co-occur with misophonia (Jastreboff and Jastreboff, 2014; Danesh and Aazh, 2020), studies focused on misophonia in the field of audiology are scarce (Porcaro et al., 2019). Nevertheless, audiologists play a key role in providing therapy and support for this patient population. Often, audiologists who are specialized in the management of tinnitus and hyperacusis also provide counseling and sound therapy (Jastreboff and Jastreboff, 2014) and/or audiologist-delivered cognitive behavioral therapy for the management of misophonia (Aazh et al., 2014, 2019b). Although the term misophonia was suggested based on studies related to therapy for tinnitus and hyperacusis (Jastreboff and Jastreboff, 2002), most of the research literature on misophonia comes from the fields of psychiatry, psychology and neuroscience, with little or no attention paid to the audiological profile of the population studied. A possible reason for this is that in the field of audiology misophonia is often considered as a subtype of hyperacusis rather than a distinct disorder (Tyler et al., 2014). Therefore, most research studies in the field of audiology have not distinguished misophonia from hyperacusis (Fackrell et al., 2015; Sheldrake et al., 2015; Zaugg et al., 2016; Aazh et al., 2017).

Most studies of misophonia performed in mental health settings have not conducted full audiological evaluations, but some have performed pure tone audiometry on a sub-group of patients (Sztuka et al., 2010; Schroder et al., 2013, 2014; Jager et al., 2020a,b; Siepsiak et al., 2022). Generally, no hearing loss was found, although some cases of tinnitus and/or hyperacusis were reported. However, Enzler et al. (2021b) conducted a study on the development of a psychoacoustic test for assessment of misophonia and reported that among 78 patients with misophonia diagnosed via the MisoQuest questionnaire (Siepsiak et al., 2020), 17 reported hearing problems, 14 had tinnitus, and 55 had hyperacusis. These results suggest that hearing loss, tinnitus and hyperacusis may not be uncommon among patients with misophonia.

Published studies have not assessed the relationship between hearing-related variables and misophonia. In theory, hearing loss could affect the experience of misophonia. The trigger sounds for misophonia often have a spectrum that is dominated by high-frequency components (Dacremont, 1995; Enzler et al., 2021b). A steeply sloping audiogram, with the greatest loss at high frequencies, would reduce the audibility of such sounds, perhaps making it less likely for an individual to have misophonia or reducing the severity of misophonia. On the other hand, people with hearing loss also often experience loudness recruitment, a more rapid than normal growth of loudness with increasing sound level once the sound becomes audible (Moore and Glasberg, 2004). Hence a sound that is only just above the detection threshold may be of moderate loudness and may be annoying. Analysis of the audiometric characteristics of people with misophonia can indicate if hearing loss influences the likelihood or severity of misophonia.

An audiological measure that is often used in the assessment and diagnosis of hyperacusis is the uncomfortable loudness level (ULL) (Aazh and Moore, 2017b). People with hyperacusis often have lower ULLs than people without hyperacusis (Blaesing and Kroener-Herwig, 2012; Fornby et al., 2015). In addition, the difference between ULLs at 1 and 8 kHz, a measure of the variation of ULLs across frequency, may be an indicator of a dislike of specific sounds, especially high-frequency sounds. Aazh and Moore (2017b) reported that among patients seeking help for tinnitus and/or hyperacusis the difference between ULLs at 1 and 8 kHz was ≥20 dB for about 10%, perhaps indicating misophonia. Siepsiak et al. (2022) compared ULLs for 62 patients with misophonia and 51 individuals with no sound sensitivity symptoms. The average ULL across ears was about 85 dB HL (standard deviation, SD = 16 dB) for the misophonia group and 90 dB HL (SD = 14 dB) for the control group, but the difference was not statistically significant, perhaps because of the large SD within each group.

Another audiological factor that may be relevant to misophonia is asymmetrical hearing threshold levels (HTLs) or ULLs (i.e., between-ear differences). A large between-ear difference in ULLs might indicate some specific abnormality in monaural pathways. For example, a disorder of the olivo-cochlear efferent system, which reduces the gain of the cochlea in response to high-level sounds, might increase sound sensitivity.
The study was registered, reviewed and approved as a clinical Ethical Approval. MATERIALS AND METHODS

Information could be used to guide the choice of therapy. This would be associated with a greater frequency of experiencing of misophonia symptoms. In the general population (Wu et al., 2014; Zhou et al., 2017; Rouw and Moore, 2017a), making the individual more likely to develop a strong reaction to trigger sounds, i.e., misophonia. Alternatively, tinnitus may distract the individual, preventing them from attending to potentially annoying trigger sounds. Past studies have not assessed the relationship between the impact of tinnitus and/or hyperacusis and symptoms of misophonia.

The first aim of the current study was to assess the proportion of patients with symptoms of misophonia among a clinical population of patients seeking help for tinnitus and hyperacusis. We predicted that this proportion would be higher than for the general population (Wu et al., 2014; Zhou et al., 2017; Rouw and Erfanian, 2018).

The second aim was to compare the audiological characteristics and severity of tinnitus, hyperacusis, anxiety and depression among patients who reported different frequencies of experiencing symptoms of misophonia in a two-week period (i.e., 0-1 days, 2-6 days, 7-10 days, and 11-14 days). We predicted that a sloping audiogram, with greater hearing loss at high frequencies would be associated with a smaller number of days of experiencing misophonia symptoms and that lower ULLs and more severe tinnitus, hyperacusis, anxiety and depression would be associated with a greater frequency of experiencing of misophonia symptoms.

The results were intended to inform those working in audiology clinics of the likelihood of misophonia among their patients and of factors that are related to it, i.e., factors that increase the probability of misophonia being present. This information could be used to guide the choice of therapy.

MATERIALS AND METHODS

Ethical Approval

The study was registered, reviewed and approved as a clinical audit by the Quality Governance Department at RSFT. The need for patient consent was waived as this was a retrospective analysis of available clinical data. Analysis of the data was approved by the South West-Cornwall and Plymouth Research Ethics Committee and the Research and Development department at the RSFT (Project ID: 182924).

Study Design and Patients

This was a retrospective cross-sectional study conducted at the Tinnitus and Hyperacusis Therapy Specialist Clinic (THTSC), RSFT, Guildford, United Kingdom. Data were included for all patients who attended the THTSC seeking help for tinnitus and/or hyperacusis in 2019-2020 and who answered a question assessing symptoms of misophonia (n = 257). Administration of the self-report questionnaires (including the question about misophonia) and audiological measurements included in this study were part of the routine care for patients at THTSC. This routine care did not include the administration of validated questionnaires for assessing misophonia; this issue is addressed in the Discussion section.

Demographic data for the patients, results of their audiological investigations and the outcomes of their self-report questionnaires were imported from their records held at the Audiology Department. All questionnaires were completed prior to the start of any treatment, at each patient’s first visit to the clinic. Patients completed the questionnaires in the clinic waiting area without involvement of their audiologist. The mean age of the patients was 53 years (SD = 16 years) (age range = 17 - 97 years). Fifty four percent (139/257) were female.

Audiological Measures

Audiological measures were:

1. Pure tone audiogram measured using the procedure recommended by the British Society of Audiology (BSA, 2011a), but with some modifications proposed by Aazh and Moore (2017c) to limit discomfort. The starting presentation level at 0.25, 0.5, 2, 3, 4, 6, and 8 kHz was equal to the HTL at the adjacent frequency (e.g., if the HTL at 1 kHz was 20 dB HL, the starting level for measuring the HTL at 2 kHz was 20 dB HL, instead of 50 dB HL as recommended by the BSA). The severity of hearing loss was categorized based on the values of the PTA across the frequencies 0.25, 0.5, 1, 2, and 4 kHz, as recommended by the British Society of Audiology (BSA, 2011a): Mild (20 – 40 dB HL), Moderate (41 – 70 dB HL), Severe (71 – 95 dB HL) and Profound (over 95 dB HL). To explore asymmetries in HTLs across the ears, patients were classified into five groups based on the between-ear difference in PTA: <5 dB, ≥5 and <10 dB, ≥10 and <20, ≥20 and <30, and ≥30. The absolute values of the differences in HTLs between 8 and 1 kHz, referred to here as HTL slope were calculated separately for the right and left ears.

2. ULLs measured following the BSA recommended procedure (BSA, 2011b), but with the modifications proposed by Aazh and Moore (2017c), to limit discomfort. The instructions were “I will gradually make the sound
louder in your ear, and you must press the button (or raise your hand) as soon as the sound becomes uncomfortable (uncomfortably loud). This is not a test to find the loudest sound you can tolerate; it is a test to find what level of sound you find uncomfortable. You should press the button (or raise your hand) only when the sound becomes uncomfortable; but make sure you press (raise) it as soon as the sound reaches that level.” The starting presentation level was equal to the measured HTL at the test frequency. In addition, levels above 80 dB HL were not used. If the ULL was not reached at 80 dB HL, the ULL at the test frequency was recorded as 85 dB HL. The across-frequency average (0.25, 0.5, 1, 2, 3, 4, 6, and 8 kHz) ULL for the ear with lower average ULL is denoted ULLmin. When ULLmin was ≤ 77 dB HL, hyperacusis was deemed to be present (Aazh and Moore, 2017b). Patients were diagnosed with severe hyperacusis if the ULL for any frequency for either ear was 30 dB HL or less (Aazh and Moore, 2018). To explore asymmetries in ULLs across the ears, patients were classified into three groups based on the between-ears difference in average ULLs (across 0.25, 0.5, 1, 2, 3, 4, 6, and 8 kHz): symmetrical (between-ear difference <5 dB), mildly asymmetrical (between-ear difference ≥5 dB and <10 dB), asymmetrical (between-ear difference ≥10 dB). The absolute values of the differences in ULLs between 8 and 1 kHz (i.e., ULL at 8 kHz minus ULL at 1 kHz), referred to here as ULL slope, were calculated separately for the right and left ears.

**Questionnaires**

**Assessment of Misophonia Symptoms**

Item 4 of the Sound Sensitivity Symptoms Questionnaire (SSSQ) (Aazh et al., 2022a) was used to identify patients with symptoms of misophonia. This item asks, “Over the last 2 weeks, how often have you been feeling angry or anxious when hearing certain sounds related to eating noises, lip smacking, sniffing, breathing, clicking sounds, tapping?” The response choices are: 0-1 days, 2-6 days, 7-10 days and 11-14 days. Scores of 0 were assigned for 0-1 days, 1 for 2-6 days, 2 for 7-10 days and 3 for 11-14 days. Cronbach’s alpha for the SAD-T, based on responses from a tinnitus and hyperacusis clinical population, is 0.91 (Aazh et al., 2022a). The overall score for the SAD-T ranges from 0 to 12. Scores of 4 or more indicate symptoms of anxiety and/or depression. This was calculated but not reported during a study on the acceptability and relevance of psychological questionnaires in the assessment of patients with tinnitus and/or hyperacusis (Aazh and Moore, 2017d).

**Questions About History of Mental Health**

Given the high prevalence of mental illness among patients seeking help for tinnitus and/or hyperacusis, the patients were asked several questions about mental health as part of routine history taking (Aazh and Moore, 2017d; Aazh et al., 2018). The questions were: (1) Do you have any history of mental illness? (2) Have you seen mental health professionals? (3) While you were growing up during the first 18 years of life did your parent(s) have depression or mental illness? The responses for these questions were “yes” or “no.” The third question is taken from the questionnaire for Adverse Childhood Experiences (Felitti et al., 1998).

**Hyperacusis Impact Questionnaire**

The hyperacusis impact questionnaire (HIQ) has eight items assessing the impact of hyperacusis on the patient’s life. The HIQ asks respondents how often (in number of days in the last 14 days) each of several situations occurred because of certain environmental sounds that seemed too loud to them, but that other people could tolerate well. Responses choices and the score for each choice were the same as for the SAD-T, as described above. Cronbach’s alpha for the HIQ is 0.93. The overall score ranges from 0 to 24. Scores above 11 indicate a clinically significant impact of hyperacusis (Aazh et al., 2022a).

**Tinnitus Impact Questionnaire**

This 7-item questionnaire assesses how often respondents experience a number of problems because of hearing a sound in their ears or head with no external source (e.g., buzzing, high-pitched whistle, hissing), over a two-week period. Responses choices and the score for each choice were the same as for the SAD-T, as described above. Cronbach’s alpha for the TIIQ is 0.89 (Aazh et al., 2022b). The overall score ranges from 0 to 21. A score below 5 indicates no impact of tinnitus, a score of 5 or 6 indicates mild impact, a score of 7 or 8 indicates moderate impact, and a score of 9 or more indicates a severe impact (Aazh et al., 2022b).

**Data Analyses**

The data were anonymized prior to statistical analysis. Descriptive statistics for the demographic variables, hearing thresholds and ULLs, and the scores for the self-report questionnaires were calculated. Welch’s t-tests (Delacre et al., 2017) and chi-squared ($\chi^2$) tests were used to compare audiological variables across frequencies and to assess the differences in the scores for the questionnaires between Miso Cat 1 and Miso Cat 0. Cohen’s $d$ was calculated.
to assess effect sizes (ES) based on mean comparison for unequal variances (Lakens, 2013; Delacre et al., 2017).

One-way analyses of variance (ANOVA) were used to assess the differences in scores for the HIQ, TIQ, SAD-T, ULLmin and PTA across ears among patients with scores of 0, 1, 2, and 3 for item 4 of the SSSQ (SSSQ4 score). The Šidák method was used for post hoc tests (Kirk, 2012). ES values following ANOVA were assessed using the $\xi$ measure (Smithson, 2001).

Spearman correlation was used to assess the relationships between SSSQ4 scores and scores for the HIQ, TIQ, SAD-T, ULLmin, PTA across ears, HTL and ULL slopes and age. The strength of the correlation coefficient ($\rho$) was considered as weak if $\rho < 0.2$, moderate if $\rho$ was between 0.2 and 0.5, and strong if $\rho > 0.5$ (Cohen, 1988; Hemphill, 2003). Variables that were significantly correlated with SSSQ4 scores were included in a logistic regression model to assess whether the SSSQ4 score (dependent variable) was related to ULLmin, scores for the HIQ, TIQ, SAD-T, and ULL and HTL slopes (independent variables). Odds ratios (ORs) and their 95% confidence intervals were obtained, both unadjusted and adjusted for (a) age and gender (b) categories of tinnitus impact as measured via the TIQ, (c) hyperacusis impact as measured via the HIQ, (d) anxiety and depression as measured via the SAD-T, (e) hyperacusis as measured via ULLmin, (f) ULL slope, and (g) HTL slope. Hearing loss categories and between-ear differences in ULLs and PTA were not included in the model as they were not correlated with SSSQ4 scores. The $p$ value required for statistical significance was $p < 0.05$.

The analyses were restricted to patients with complete data for all variables required for a particular analysis. The number of patients included in each analysis ($n$) is reported. The STATA program (version 13) (StataCorp, 2013) and MATLAB 2020a (The MathWorks, 2020) were used for statistical analyses.

**RESULTS**

**Characteristics of the Study Population**

The means and SDs of the HTLs and ULLs for each ear and each frequency are shown in Table 1. The grand mean PTA across ears was 22 dB HL (SD = 15 dB) ($n = 244$). The grand mean PTA for the better ear was 18 dB HL (SD = 13 dB). The grand mean PTA for the worse ear was 26 dB HL (SD = 19 dB). Based on the PTA for the better ear, 65% of the patients had no hearing loss, 28% had mild hearing loss, and 7% had moderate hearing loss. Based on the PTA for the worse ear, 49% of the patients had no hearing loss, 34% had mild hearing loss, 13.5% had moderate hearing loss, 2.9% had severe hearing loss and 0.8% had profound hearing loss.

For 64% of the patients (156/244), there was less than a 5-dB difference in PTA between the two ears. The difference in PTA between ears was $\geq 5$ and $< 10$ dB for 20% of cases, $\geq 10$ and $< 20$ dB for 5% of cases, $\geq 20$ and $< 30$ dB for 4.5% of cases, and $\geq 30$ dB for 6.2% of cases. The mean HTL slope was 22.7 dB (SD = 19.5 dB) for the left ears and 20 dB (SD = 19 dB) for the right ears. The HTL slope was $\geq 20$ dB, for at least one ear, for 58% of the patients (143/248).

The grand average ULL across 0.25, 0.5, 1, 2, 4, 6, and 8 kHz and across ears was 78.5 dB HL (SD = 8.3) ($n = 169$). The average value of ULLmin was 77.7 dB HL (SD = 9) ($n = 191$). ULLmin values were 77 dB HL or below, suggesting hyperacusis for 30% of patients. About 1.5% of the patients were diagnosed with severe hyperacusis, based on them having a ULL of 30 dB HL or less for any frequency for either ear (Aazh and Moore, 2018). ULLs were symmetrical for 83% of patients, mildly asymmetrical for 14% and asymmetrical for 2.4%. The mean ULL slope was 5 dB (SD = 8 dB) for both ears. About 11.5% of patients had a ULL slope $\geq 20$ dB for at least one ear.

For the study population, the mean scores for the HIQ, TIQ and SAD-T were 8 (SD = 7.5, $n = 224$), 8.4 (SD = 6, $n = 170$), and 4 (SD = 4, $n = 253$), respectively. Based on scores for the HIQ, 30% of patients had hyperacusis. Based on scores for the TIQ, 28% of patients had no tinnitus handicap, 20.5% had a mild tinnitus handicap, 10.5% had a moderate tinnitus handicap, and 41% (70/170) had a severe tinnitus handicap. Based on scores for the SAD-T, 44.5% of patients had symptoms of anxiety and/or depression. About 47% of the patients (113/241) reported a history of mental illness, 39% (94/240) reported seeing mental health professionals, and 31.5% reported that when they were under 18 years of age at least one of their parents had mental illness.

**Comparison of Miso Cat 0 and Miso Cat 1**

Overall, 23% of patients (59/257) were classified as Miso Cat 1. Patients in Miso Cat 1 were younger on average than those in Miso Cat 0 (Table 2). The percentage of females was 61% for Miso Cat 1 and 52% for Miso Cat 0, and the difference was not significant ($\chi^2 = 1.5, p = 0.22$). There was no significant difference in PTA between those in Miso Cat 0 and those in Miso Cat 1, as shown in Figures 1, 2.

Based on the PTA for the better ear, among the 58 patients in Miso Cat 1, 59% had no hearing loss, 38% had mild hearing loss, and 3.5% had moderate hearing loss. Among the 190 patients in Miso Cat 0, 67% had no hearing loss, 24% had mild hearing loss, and 8% had moderate hearing loss. Based on the PTA for the worse ear, among the patients in Miso Cat 1, 45% had no hearing loss, 38% had mild hearing loss, 14% had moderate hearing loss, and 3.5% had severe hearing loss. Among the patients in Miso Cat 0, 50% had no hearing loss, 33% had mild hearing loss, 13% had moderate hearing loss, 3% had severe hearing loss, and 1% had profound hearing loss. The differences in distributions of hearing loss categories for the better and worse ears between Miso Cat 1 and Miso Cat 0 were not statistically significant ($\chi^2 = 5.08, p = 0.079$ and $\chi^2 = 0.22$). The percentage of females was 61% for Miso Cat 1 and 52% for Miso Cat 0, and the difference was not significant ($\chi^2 = 0.86, p = 0.37$).

The HTL slope averaged across ears was $\geq 20$ dB for 52% of those in Miso Cat 1 and for 59.5% of those in Miso Cat 0 ($\chi^2 = 1.1, p = 0.3$). The PTA differed across ears by less than 5 dB for 65.5% of those in Miso Cat 1 and 63.4% of those in Miso Cat 0. The difference in PTA across ears was $\geq 5$ and $< 10$ dB for 17% of cases in Miso Cat 1 and 21% of Miso Cat 0, $\geq 10$ and $< 20$ dB for 3% of those in Miso Cat 1 and 5.9% of Miso Cat 0, $\geq 20$ and $< 30$ dB for 5.3% of those in Miso Cat 1 and 4.3% of those in Miso Cat 0,
TABLE 1 | Means (SDs) of hearing threshold levels (HTLs) and uncomfortable loudness levels (ULLs) in dB HL for each ear of the study population across different frequencies.

| Frequency, kHz | 0.25 | 0.5 | 1 | 2 | 3 | 4 | 6 | 8 |
|---------------|------|-----|---|---|---|---|---|---|
| **HTL right** |      |     |   |   |   |   |   |   |
| n = 247       | 18   | 18  | 19 | 21 | 26 | 30 | 38 | 36 |
| **HTL left**  |      |     |   |   |   |   |   |   |
| n = 247       | 18   | 19  | 19 | 23 | 29 | 34 | 39 | 40 |
| **ULL right** |      |     |   |   |   |   |   |   |
| n = 198       | 78   | 79  | 79 | 79 | 79 | 79 | 77 | 77 |
| **ULL left**  |      |     |   |   |   |   |   |   |
| n = 198       | 78   | 79  | 80 | 79 | 80 | 79 | 77 | 77 |

The number of patients included in each analysis is indicated by n.

TABLE 2 | Results of independent-samples Welch’s t-tests comparing the PTA (pure tone average) averaged across ears, between-ears difference in PTA, ULLmin (across-frequency average uncomfortable loudness level for the ear with lower average ULL), between-ears difference in average ULL, ULL slope (the value of the difference in ULLs between 8 and 1 kHz) for each ear and averaged across ears, HTL slope (absolute values of the differences in hearing threshold levels between 8 and 1 kHz) for each ear and averaged across ears, scores for the TIIQ (Tinnitus Impact Questionnaire), HIQ (Hyperacusis Impact Questionnaire), SAD-T (Screening for Anxiety and Depression-Tinnitus), and age for groups Miso Cat 0 and Miso Cat 1. Significant p values are indicated in bold font.

|               | Miso Cat 0 Mean (SD) | Miso Cat 1 Mean (SD) | Difference: mean and 95% confidence intervals (CI) | P-value | ES and 95% CI |
|---------------|----------------------|----------------------|-----------------------------------------------------|---------|---------------|
| PTA across ears | 22 (15.5)            | 22.5 (14)            | −0.61 (−4.9 to 3.7)                                  | 0.78    | −0.04         |
| Between-ears difference in PTA (dB) | 6.8 (11)            | 9.0 (15.5)           | −2.2 (−6.6 to 2.2)                                  | 0.31    | −0.17         |
| ULLmin (dB HL) | 79 (8)               | 74 (11)              | 5.0 (1.5 to 8.5)                                    | 0.006   | 0.56          |
| Between-ears difference in average ULL (dB) | 1.9 (2.9)        | 2.7 (3.4)            | −0.8 (−2.0 to 0.4)                                  | 0.18    | −0.27         |
| ULL slope for right ears (dB) | 3.8 (6.5)           | 9 (10.8)             | −5.1 (−8.7 to −1.6)                                 | 0.006   | −0.67         |
| ULL slope for left ears (dB) | 4.1 (6.7)           | 9.0 (10.4)           | −4.8 (−8.2 to −1.5)                                 | 0.005   | −0.62         |
| ULL slope averaged across ears (dB) | 4.2 (6.1)           | 8.7 (9.4)            | −4.5 (−7.8 to −1.3)                                 | 0.007   | −0.64         |
| HTL slope for right ears (dB) | 21.7 (20)           | 15.4 (16)            | 6.2 (1.1 to 11.3)                                   | 0.017   | 0.33          |
| HTL slope for left ears (dB) | 24.2 (20)           | 18.1 (15)            | 6.1 (1.1 to 11.0)                                   | 0.016   | 0.31          |
| HTL slope averaged across ears (dB) | 23.0 (18)           | 16.8 (14)            | 6.2 (1.8 to 10.6)                                   | 0.007   | 0.36          |
| TiQ score (0-21) | 6.8 (4.9)           | 13.7 (6.6)           | −7.0 (−9.2 to −4.7)                                 | < 0.0001| −1.3          |
| HIQ score (0-24) | 5.7 (5.9)           | 16.0 (6.8)           | −10.3 (−12.4 to −8.2)                               | < 0.0001| −1.1          |
| SAD-T score (0-12) | 3 (3.4)             | 7.5 (4.1)            | −4.5 (−5.7 to −3.3)                                 | < 0.0001| −1.3          |
| Age (years)   | 54.5 (17)           | 49.5 (12)            | 5.0 (1.0 to 8.9)                                    | 0.014   | 0.31          |

The sixth column shows ES values based on Cohen’s d with 95% CIs.

and ≥30 dB for 8.6% of those in Miso Cat 1 and 5.4% of those in Miso Cat 0. The proportions of patients falling in each asymmetry category did not differ significantly for Miso Cat 1 and Miso Cat 0 ($\chi^2 = 1.7, p = 0.79$).
As shown in Figures 1, 2, those in Miso Cat 1 had significantly lower (worse) mean ULLs than those in Miso Cat 0 for all frequencies and both ears (all \( p < 0.01 \)). The ULL slope averaged across ears was \( \geq 20 \text{ dB} \) for 26% of patients \((n = 12/46)\) in Miso Cat 1 compared to 7% \((n = 10/145)\) in Miso Cat 0, and this difference in proportions was significant \((\chi^2 = 12.6, p < 0.001)\).

Among patients in Miso Cat 1 \((n = 38)\), ULLs were symmetrical for 82%, mildly asymmetrical for 16% and asymmetrical for 3%. Corresponding values for those in Miso Cat 0 were 84, 14, and 2%. The proportions of patients falling in each asymmetry category did not differ significantly for Miso Cat 1 and Miso Cat 0 \((\chi^2 = 0.12, p = 0.94)\).

Among patients in Miso Cat 1, based on TIQ scores there was no impact of tinnitus for 8% \((3/39)\), a mild impact for 13% \((5/39)\), a moderate impact for 5% \((2/39)\), and a severe impact for 74% \((29/39)\). Among patients in Miso Cat 0, corresponding values were 34% \((44/131)\), 23% \((30/131)\), 12% \((16/131)\), and 31% \((41/131)\). The proportions falling in the different tinnitus impact categories differed significantly between Miso Cat 1 and 0 \((\chi^2 = 23.7, p < 0.001)\).
significant ($\chi^2 = 5.56, p < 0.001$).

Based on the SAD-T score, 81% (47/58) of patients in Miso Cat 1 had symptoms of anxiety and depression compared to 33.5% (65/194) of patients in Miso Cat 0, and this difference in proportions was significant ($\chi^2 = 6.4, p = 0.011$). Among patients with a history of parental mental illness in their childhood, 59% (43/73) had an abnormal SAD-T score compared to 39.5% (64/162) of those with no history of parental mental illness, and this difference in proportions was significant ($\chi^2 = 7.6, p = 0.006$).

Table 2 shows the results of Welch’s $t$-tests comparing various measures for those in Miso Cat 0 and Miso Cat 1. The mean ULLmin values were significantly lower and total scores for the SAD-T, TIQ, HIQ were significantly worse for Miso Cat 1 than for Miso Cat 0. The ULL slope was higher for Miso Cat 1 than for Miso Cat 0, but the HTL slope was lower for Miso Cat 1 than for Miso Cat 0. There were no significant differences between Miso Cat 1 and 0 in terms of the between-ear differences in ULL or PTA.

Audiological and Psychological Factors Related to Misophonia Symptoms

Of 257 patients, SSSQ4 scores of 0, 1, 2, and 3 were obtained for 149 (58%), 49 (19%), 21 (8%) and 38 (15%), respectively. As shown in Table 3, there were significant differences in ULLmin, HIQ, TIQ and SAD-T scores among patients with different SSSQ4 scores. Post hoc pairwise comparisons indicated that ULLmin was significantly lower only for patients who scored 3 compared to 0 for SSSQ4 ($p = 0.013$). The other pairwise comparisons were not significant ($p > 0.05$). HIQ scores were significantly worse for patients whose SSSQ4 scores were 3 vs. 0 ($p < 0.0001$), 2 vs. 0 ($p = 0.009$), 3 vs. 1 ($p < 0.0001$) and 3 vs. 0 ($p < 0.0001$). TIQ scores were significantly worse for patients whose SSSQ4 scores were 3 vs. 0 ($p < 0.0001$), 3 vs. 1 ($p < 0.0001$) and 3 vs. 2 ($p = 0.005$). SAD-T scores were significantly worse for patients whose SSSQ4 scores were 3 vs. 0 ($p < 0.0001$), 2 vs. 0 ($p = 0.001$), 3 vs. 1 ($p < 0.0001$) and 3 vs. 2 ($p = 0.041$). The other pairwise comparisons were not significant ($p > 0.05$).

There was no significant difference in the average PTA across ears for patients with different SSSQ4 scores.

As shown in Table 4, there was no significant correlation between the SSSQ4 scores and the average PTA values across ears. There were moderate to strong correlations between SSSQ4 scores and the HTL slopes; and (f) HTL slope < 20 dB versus ≥ 20 dB; and (f) HTL slope < 20 dB versus ≥ 20 dB. When all measures were treated as independent (columns 2 and 3 of Table 5), the resulting non-adjusted ORs differed significantly from 1 for all predictors, with the largest effects for tinnitus impact category being moderate or severe, significant impact of hyperacusis, and presence of symptoms of anxiety and depression.

DISCUSSION

Our results showed that 42% of patients seeking help for tinnitus and/or hyperacusis presented with some symptoms of misophonia. Twenty three percent of patients reported being bothered by certain sounds on 7-14 days in the last 14 days (Miso Cat 1). There was no difference in the prevalence of different degrees of hearing loss among patients in Miso Cat 1 and Miso Cat 0, but a significant proportion of patients in both groups (more than 33%) had some degree of hearing loss, indicating that misophonia is not restricted to those with normal hearing. The percentage of patients with hearing loss among those with misophonia symptoms reported here is higher than reported in previous studies. For example, Enzler et al. (2021b) reported that 22% of individuals with misophonia as measured via the MisoQuest had self-reported hearing issues and Siepsia
TABLE 3 | Means (SD) of ULLmin (across-frequency average uncomfortable loudness level for the ear with lower average ULL), PTA (pure tone average) across ears, and scores for the HIQ (Hyperacusis Impact Questionnaire), TIQ (Tinnitus Impact Questionnaire), and SAD-T (Screening for Anxiety and Depression-Tinnitus) for patients giving each SSSQ4 score, indicating the number of days that they were bothered by certain sounds in the last 2 weeks.

| Number of days bothered in the last 14 days | 0-1 days | 2-6 days | 7-10 days | 11-14 days | F-value degrees of freedom | p-value | ES (95%CI) |
|-------------------------------------------|---------|----------|----------|-----------|---------------------------|--------|-----------|
| ULLmin (dB HL)                            | 80 (8.2)| 77 (7.3)| 74 (7.8)| 74 (13)   | 4.45                      | 3, 187 | 0.07 (0.007 to 0.13) |
|                                          | n = 105 | n = 40  | n = 16  | n = 30    |                           |        |            |
| PTA across ears (dB HL)                   | 23 (16) | 20 (14) | 20 (13) | 24 (15)   | 0.59                      | 3, 240 | 0.007 (0 to 0.03) |
|                                          | n = 137 | n = 49  | n = 21  | n = 37    |                           |        |            |
| HIQ score (0-24)                          | 5.1 (5.8)| 7.3 (5.9)| 10.2 (6.5)| 18.4 (5.4)| 50.5                     | 3, 220 | 0.0001 (0.3 to 0.48) |
|                                          | n = 129 | n = 44  | n = 15  | n = 36    |                           |        |            |
| TIQ score (0-21)                          | 6.0 (4.8)| 8.8 (4.8)| 9.8 (5.4)| 15.7 (6.3)| 25.5                     | 3, 166 | 0.0001 (0.19 to 0.41) |
|                                          | n = 94  | n = 37  | n = 13  | n = 26    |                           |        |            |
| SAD-T score (0-12)                        | 2.7 (3.3)| 4.1 (3.4)| 5.9 (3.6)| 8.5 (4.1) | 29.6                     | 3, 248 | 0.0001 (0.16 to 0.34) |
|                                          | n = 146 | n = 49  | n = 21  | n = 37    |                           |        |            |

The right-most column shows the outcomes of one-way ANOVAs with factor SSSQ4 score. The number of patients included in each analysis is indicated by n. Significant p values are indicated in bold font. The seventh column shows ES values based on η² with 95% CIs.

TABLE 4 | Spearman correlations (ρ) and corresponding p values between the number of days out of 14 when bothered by certain sounds (based on SSSQ4 score) with: PTA (pure tone average) across ears, TIQ (Tinnitus Impact Questionnaire) scores, HIQ (Hyperacusis Impact Questionnaire) scores, SAD-T (Screening for Anxiety and Depression-Tinnitus) scores, ULL slope (the value of the difference in ULLs between 8 and 1 kHz) for each ear and averaged across ears, ULLmin (across-frequency average uncomfortable loudness level for the ear with lower average ULL), HTL slope (value of the differences in hearing threshold levels between 8 and 1 kHz) for each ear and averaged across ears, and age.

| PTA across ears | TIQ score | HIQ score | SAD-T score | ULL slope | ULLmin | HTL slope | Age |
|-----------------|-----------|-----------|-------------|-----------|---------|-----------|-----|
| p = 0.014       | p = 0.49  | p = 0.53  | p = 0.47    | p = 0.28  | p = −0.29| p = −0.16 | p = −0.15 |
| p = 0.82        | p < 0.0001| p < 0.0001| p < 0.0001  | p < 0.0001| p < 0.0001| p < 0.0001| p = 0.015 |
| n = 244         | n = 170   | n = 224   | n = 253     | n = 170   | n = 191  | n = 246   | n = 257 |

Each cell also shows the number of patients (n). Significant p values are indicated in bold font.

eet al. (2022) reported hearing loss in 16% of participants with misophonia, as diagnosed using the criteria of Schroder et al. (2013). Most of the patients in those studies were recruited via social media, so their study population was different from that for our study.

Although the presence or absence of hearing loss did not seem to be related to the presence of misophonia symptoms, a steep slope of the audiogram, with greater loss at high frequencies, was associated with a reduced risk of misophonia. This probably occurs because some of the triggers for misophonia are sounds whose spectrum is dominated by high frequencies, such as the sound of crispy foods (Dacremont, 1995). Hearing loss at high frequencies reduces the likelihood that such trigger sounds will be audible.

The presence of symptoms of misophonia was not significantly related to between-ear differences in HTL or ULL. This indicates that the underlying mechanism of misophonia is unlikely to be related to asymmetric pathologies of the peripheral auditory pathway; rather, a more central mechanism is involved. This is consistent with imaging studies reporting altered non-auditory areas in the brain among patients with misophonia compared with healthy controls (Kumar et al., 2017; Lin et al., 2020).

In this paper, one of the criteria for indicating the presence of hyperacusis was a ULLmin value ≤77 dB HL (the other criterion was HIQ score). The use of ULLs for diagnosing hyperacusis has been challenged by several authors; some studies have reported that ULLs averaged across frequency were not significantly correlated with self-report measures of hyperacusis (Khalifa et al., 2002; Meeus et al., 2010). In addition, there are differences in the criteria for diagnosing hyperacusis based on ULLs (Goldstein and Shulman, 1996; Anari et al., 1999; Jastreboff and Jastreboff,
TABLE 5 | Results of a logistic regression model showing the odds ratio (OR) of the SSSQ4 score (dependent variable) relative to a baseline.

| Hyperacusis impact category | Non-adjusted OR (95% CI) | P-value | Adjusted OR (95% CI) | P-value |
|-----------------------------|--------------------------|---------|----------------------|---------|
| No (ULLmin > 77 dB HL)      | 1.0                      | 0.005   | 1.0                  | 0.86    |
| Yes (ULLmin ≤ 77 dB HL)     | 2.45 (1.3 to 4.6)        |         | 1.1 (0.34 to 3.56)   |         |
|                             | n = 191                  |         |                      |         |

| Tinnitus impact category    | Non-adjusted OR (95% CI) | P-value | Adjusted OR (95% CI) | P-value |
|-----------------------------|--------------------------|---------|----------------------|---------|
| No impact (TIQ score < 5)   | 1.0                      | 0.076   | 1.0                  | 0.06    |
| Mild (TIQ score 5 or 6)     | 2.5 (0.9 to 7.14)        |         | 3.54 (0.93 to 13.5)  |         |
| Moderate (TIQ score 7 or 8) | 6.1 (1.83 to 20.25)      |         | 4.39 (0.85 to 22.8)  |         |
| Severe (TIQ score ≥ 9)      | 9.3 (3.8 to 23.1)        |         | 5.42 (1.3 to 20.17)  |         |
|                             | n = 170                  |         |                      |         |

| Hyperacusis impact category | Non-adjusted OR (95% CI) | P-value | Adjusted OR (95% CI) | P-value |
|-----------------------------|--------------------------|---------|----------------------|---------|
| No impact (HIQ score ≤ 11)  | 1.0                      | < 0.0001| 1.0                  | 0.032   |
| Significant impact (HIQ score > 11) |           |         |                      |         |

| Anxiety and depression      | Non-adjusted OR (95% CI) | P-value | Adjusted OR (95% CI) | P-value |
|-----------------------------|--------------------------|---------|----------------------|---------|
| No (SAD-T score < 4)        | 1.0                      | < 0.0001| 1.0                  | 0.044   |
| Yes (SAD-T score ≥ 4)       | 5.4 (3.1 to 9.3)         |         | 2.8 (1.03 to 7.4)    |         |
|                             | n = 252                  |         |                      |         |

| Across-frequency difference in ULLs | Non-adjusted OR (95% CI) | P-value | Adjusted OR (95% CI) | P-value |
|------------------------------------|--------------------------|---------|----------------------|---------|
| No (across ears ULL slope < 20 dB) | 2.96 (1.15 to 7.63)      | < 0.0125| 1.0                  | 0.13    |
| Yes (across ears ULL slope ≥ 20 dB)|                         |         |                      |         |
|                             | n = 191                  |         |                      |         |

| Across-frequency difference in HTLs | Non-adjusted OR (95% CI) | P-value | Adjusted OR (95% CI) | P-value |
|------------------------------------|--------------------------|---------|----------------------|---------|
| No (across ears HT slope < 20 dB)  | 0.59 (0.36 to 0.99)      | < 0.006  | 1.0                  | 0.047   |
| Yes (across ears HT slope ≥ 20 dB) |                         |         |                      |         |
|                             | n = 191                  |         |                      |         |

Variables included in the model were the presence or absence of hyperacusis based on ULLmin (across-frequency average uncomfortable loudness level for the ear with lower average ULL), tinnitus impact category based on scores for the TIQ (Tinnitus Impact Questionnaire), hyperacusis impact category based on scores for the HIQ (Hyperacusis Impact Questionnaire), presence of anxiety and depression symptoms as measured via the SAD-T (Screening for Anxiety and Depression-Tinnitus), presence or absence of across-frequency difference in ULLs based on the average ULL slope (the values of the difference in hearing threshold levels between 8 and 1 kHz) across ears, and presence or absence of across-frequency difference in HTLs based on the HTL slope (the values of the differences in hearing threshold levels between 8 and 1 kHz) across ears. Unadjusted and adjusted OR values and their 95% confidence intervals (CIs) are shown. The adjusted OR takes into account the effects of age and gender in addition to the effects of other variables in the model. Significant p values are indicated in bold font.

For example, Goldstein and Shulman (1996) suggested that ULLs between 80 and 90 dB HL at two or more frequencies indicate mild hyperacusis, ULLs between 65 and 75 dB HL indicate moderate hyperacusis and ULLs below 60 dB HL indicate severe hyperacusis. Anari et al. (1999) suggested 70 dB HL as the cutoff value indicating significant hyperacusis. Jastreboff and Jastreboff (2000) suggested that "threshold of significant hyperacusis is defined as average LDLs below 100 dB HL" (LDL stands for loudness discomfort level, which is another term for ULL). Sheldrake et al. (2015) reported that if a criterion value of 100 dB HL for ULLs averaged across 0.5, 1, 2, and 4 kHz (denoted ULL0.5–4) is used, this results in a positive diagnosis for 90% of those with hyperacusis, but results in a high false positive rate of 60% (and a corresponding specificity of only 40%).

Sherlock and Formby (2005) reported that among individuals with no complaint of hyperacusis the average value of ULL0.5–4 was 102 dB HL (SD = 12 dB). They showed that 50% of people with no hyperacusis had average ULL0.5–4 values less than 105 dB HL, 25% had ULL0.5–4 values less than 94 dB HL, and 5% had ULL0.5–4 values less than 80 dB HL. To avoid excessive false positives when diagnosing hyperacusis based on ULL0.5–4, the lower 95% bound of the global mean for people without hyperacusis can be used as the cutoff; this is obtained by subtracting from that mean 1.96 times the square root of the variance of the mean, giving a value of 80 dB HL based on the data of Sherlock and Formby (2005).

Aazh and Moore (2017b) took a different approach. As noted earlier, they based their analyses on the average ULL across 0.25, 0.5, 1, 2, 3, 4, 6, and 8 kHz for the ear with lower average ULLs, denoted ULLmin. They chose a cutoff value for ULLmin corresponding to the 95% upper bound of the ULLmin values for people with hyperacusis as diagnosed via the score for Hyperacusis Questionnaire (HQ) (Khalfa et al., 2002); this was obtained by adding 1.96 times the
indicating that patients with misophonia symptoms are likely
slope values were moderately correlated with SSSQ4 scores,
been even larger if the cap of 80 dB HL had not been imposed.
values between those in Miso Cat 0 and Miso Cat 1 would have
in average ULLmin values for those diagnosed as having versus
having hyperacusis based on ULLmin. For patients who were
frequencies. Thus, the cap of 80 dB HL had very little influence
ULLmin who did not press the button indicating the onset of
the left and right ears, respectively. Therefore, the cap of 80 dB
level used by Siepsiak et al. (2022) ranged from 90 to 120 dB
HL depending on the test frequency. In our study, the level was
limited to 80 dB HL regardless of frequency in order to avoid
discomfort, as recommended by Aazh and Moore (2017c).
One concern is the extent to which the cap of 80 dB HL
influenced the values of ULLmin. There were very few cases of
patients who were classified as having hyperacusis based on
ULLmin who did not press the button indicating the onset of
discomfort at a level of 80 dB HL or below, so the artificial value of
85 dB was used only rarely, and then usually only for one or two
frequencies. Thus, the cap of 80 dB HL had very little influence
on the values of ULLmin among those who were classified as
having hyperacusis based on ULLmin. For patients who were
not classified as having hyperacusis based on ULLmin, 59 and 70
(out of 133) did not press the button at 80 dB HL or below for
the left and right ears, respectively. Therefore, the cap of 80 dB
HL would have reduced the mean ULLmin value among patients
who were not classified as having hyperacusis based on ULLmin
values. The overall effect of the cap was to reduce the difference
in average ULLmin values for those diagnosed as having versus
not having hyperacusis. It is likely that the differences in ULLmin
values between those in Miso Cat 0 and Miso Cat 1 would have
been even larger if the cap of 80 dB HL had not been imposed.
The ULL slope was significantly higher (steeper) for patients
in Miso Cat 1 than for patients in Miso Cat 0. The ULL
slope values were moderately correlated with SSSQ4 scores,
indicating that patients with misophonia symptoms are likely
to be more bothered by high-frequency sounds than by low-
frequency sounds. This is consistent with the finding that
sounds with strong concentrations of energy in the range 2.5 to
5.5 kHz are associated with auditory perceptual unpleasantness
for normal subjects (Halpern et al., 1986; Kumar et al., 2008). The
auditory system is maximally sensitive over this frequency range,
in that absolute thresholds are lowest, and for a given sound level
loudness is greatest (Moore, 1997). This sensitivity may be
magnified in patients with misophonia, as has been observed for
individuals with noise sensitivity (Kliuchko et al., 2016). High
sensitivity to high-frequency sounds has also been reported for
cases of severe hyperacusis (Aazh and Moore, 2018) and many
of the patients in our sample with higher SSSQ4 scores also had
hyperacusis as measured via the HIQ and ULLmin.
The proportion of patients who had seen mental health
professionals was significantly higher for Miso Cat 1 than for
Miso Cat 0. This is consistent with the finding of Kulc et al. (2021)
that contact with mental health services for any psychological
problem was more common among those with misophonia than
among those without (48 vs. 29%). The present study showed
that SAD-T scores were moderately correlated with SSSQ4
scores. This is consistent with other reports of a relationship
between misophonia and mental illness (Guetta et al., 2022;
Siepsiak et al., 2022). More in-depth investigation is needed
to shed light on the directionality of the association between
misophonia and psychiatric disorders/symptoms. Specifically, it
would be useful to assess whether the chance of being affected
by psychiatric disorders is higher when misophonia already exists
(Erfanian et al., 2019).
A new finding of our study was that the proportion of patients
with a childhood history of parental mental illness was higher for
Miso Cat 1 (45%) than for Miso Cat 0 (27%). This is consistent
with reports of a higher impact of tinnitus, hyperacusis-induced
anxiety, and depression symptoms among patients who reported
that during their first 18 years of life their parent(s) suffered
from a mental illness (Aazh et al., 2018, 2019a,c, 2020). Mounting
evidence suggests that adverse childhood experiences play a
major lifelong role in mental and physical problems (Anda
et al., 2006, 2010; Erfanian, 2018). Parental mental illness is an
important form of adverse childhood experiences (Anda et al.,
2006). Future studies should explore the history of exposure to
various childhood adverse experiences, ranging from different
forms of abuse (physical, emotional, or sexual), neglect (physical
and emotional) and various aspects of household dysfunction
(substance abuse in the family, parental mental illness, mother
treated violently, imprisoned household member, or parental
separation) among patients with misophonia (Felitti et al.,
1998; Felitti, 2009). This is important because, if a significant
relationship exists, the presence of more severe misophonia
symptoms in patients could be an indicator of childhood adverse
experiences. Patients with a history of childhood adversities often
need more complex and in-depth psychological treatments for
their mental health should they develop emotional problems
(Pigeon et al., 2009; Kajeepeta et al., 2015).
Scores for the TIQ and HIQ were significantly worse for
patients in Miso Cat 1 than for those in Miso Cat 0. Also,
TIQ scores were moderately correlated with SSSQ4 scores
and HIQ scores were strongly correlated with SSSQ4 scores.
The adjusted logistic regression model showed that patients
with a severe impact of tinnitus and a significant impact of hyperacusis were more likely to have a higher SSSQ4 score. To the best of our knowledge, these are novel findings. This is consistent with the similarity of the neuropathology of misophonia, hyperacusis and tinnitus, as indicated by altered auditory-limbic system connections (Kumar et al., 2017; Lin et al., 2020), micro-structural alternations of white matter in non-auditory regions (Chen et al., 2020; Eijsker et al., 2021a), and functional connectivity among auditory cortex, cerebellum and the limbic system (Kumar et al., 2017; Cai et al., 2019; Eijsker et al., 2021b). A similar relationship has been reported between tinnitus and hyperacusis: patients with a more severe impact of tinnitus also tend to have more severe symptoms of hyperacusis (Aazh and Moore, 2017d; Cederroth et al., 2020; Aazh et al., 2021).

The adjusted logistic regression model also showed that a score of 4 or more for the SAD-T significantly increased the odds of having a higher SSSQ4 score, consistent with the idea that misophonia is associated with anxiety and depression. Given that misophonia leads to significant emotional distress, interpersonal and social difficulties, disability, and interference with daily life, it is not surprising that it contributes to the development of anxiety and depression. Sufferers may also experience functional impairments, such as difficulty in performing their job and concentration difficulties (Swedo et al., 2022). Finally, the adjusted model showed that a slope of the audiogram of 20 dB or more significantly decreased the odds of having a higher SSSQ4 score, consistent with the idea that reduced audibility of high-frequency sounds decreases the chances of misophonia trigger sounds being audible. There is a gap in our understanding of the function of auditory system in this patient population, and future studies should explore other characteristics of the auditory system among patients with misophonia using psycho-acoustic and electrophysiological measures.

This study was based on a retrospective analysis of the available clinical data for patients seen during the years 2019 and 2020. Therefore, we were limited to the measures that were obtained as a part of routine clinical practice at the THTSC during that time. Misophonia was assessed based on only one question (item 4 of the SSSQ). This is not unusual for clinical services, since misophonia questionnaires have not yet been widely adopted by audiologists in day-to-day clinical practice. However, we recognize that using only one question to assess misophonia is not ideal, although it has been done by other researchers for assessing misophonia, hyperacusis severity, hearing impairment and tinnitus severity (Schecklmann et al., 2014; Greenberg and Carlos, 2018; Cederroth et al., 2020; Jaswal et al., 2021). Also, the validity and reliability of using SSSQ4 to assess the frequency of reported misophonia symptoms have not been evaluated. Therefore, the results of our correlational and regression modeling need to be interpreted with caution. To address this limitation, future studies should use validated measures to assess the relationship between misophonia and measures of the impact of tinnitus and hyperacusis, measures of anxiety and depression, and hearing-related variables. Examples of these measures are MisoQuest (Siepsiak et al., 2020), the Amsterdam Misophonia Scale (Schroder et al., 2013; Naylor et al., 2021), the Misophonia Response Scale (Dibb et al., 2021), the Core Discriminant Sounds of Misophonia (Enzler et al., 2021b), the Duke Misophonia Questionnaire (Rosenthal et al., 2021) and the Misophonia Questionnaire (MQ) (Wu et al., 2014).

Another limitation is that all patients were referred to an audiology clinic for tinnitus and/or hyperacusis management. Therefore, our results are probably not representative of the general population or of patients referred to mental health services.

**CONCLUSION**

Among a population seeking help from an audiology clinic for tinnitus and/or hyperacusis, 23% were classified as having misophonia. The presence and frequency of reported symptoms of misophonia were not related to audiometric thresholds, or to the asymmetry of audiometric thresholds across ears, except that a steeply sloping audiogram reduced the likelihood of more frequently reported misophonia symptoms in a 2-week period. The latter effect may reflect the finding that the sounds that trigger misophonia often contain significant energy at high frequencies, and high-frequency hearing loss reduces the likelihood of such sounds being audible. Those with higher SSSQ4 scores had lower values of ULLmin (the across-frequency average of ULLs for the ear with lower average ULLs) than those with lower SSSQ4 scores. The frequency of reported misophonia symptoms as measured via SSSQ4 increased with increasing impact of tinnitus. Using a logistic regression model adjusted for the effects of age and gender, it was found that a TIQ score ≥9 increased the odds of reporting misophonia symptoms by a factor of 5.4. Using the same adjusted model, it was found that an HIQ score >11 (indicating a significant impact of hyperacusis) increased the odds of reporting misophonia symptoms by a factor of 3.9. Using the same adjusted model, it was found that a SAD-T score ≥4 (indicating symptoms of anxiety and depression) increased the odds of reporting misophonia symptoms by a factor of 2.8. We conclude that, when assessing individuals with tinnitus and hyperacusis, it is important to screen for misophonia, particularly when ULLmin is abnormally low or the TIQ, HIQ or SAD-T score is abnormally high. This will help clinicians to distinguish misophonia from similar disorders, guiding the choice of therapeutic strategies.

**DATA AVAILABILITY STATEMENT**

The original contributions presented in this study are included in the article/supplementary material. Further inquiries can be directed to the corresponding author.

**ETHICS STATEMENT**

The study was registered, reviewed and approved as a clinical audit by the Quality Governance Department at RSFT. The need for patient consent was waived as this was a retrospective analysis.
of available clinical data. Analysis of the data was approved by the South West-Cornwall and Plymouth Research Ethics Committee and the Research and Development department at the RSFT (Project ID: 182924).

AUTHOR CONTRIBUTIONS

HA collected the data. HA, ME, AD, and BM collaborated on analysis of the data, interpretations of the results and preparing the manuscript. All authors contributed to the article and approved the submitted version.

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