

1 COVID-19 Anosmia Reporting Tool: Initial Findings

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20 Abstract

21 There is accumulating anecdotal evidence that anosmia and dysgeusia are associated
22 with the COVID-19 pandemic. In order to investigate their relationship to SARS-CoV2
23 infection, the American Academy of Otolaryngology–Head and Neck Surgery (AAO-
24 HNS) developed the COVID-19 Anosmia Reporting Tool for Clinicians **for the basis of**
25 **this pilot study**. This tool allows healthcare providers to confidentially submit cases of
26 anosmia and dysgeusia related to COVID-19. We analyzed the first 237 entries **which**
27 **revealed that** anosmia was noted in 73% of subjects prior to COVID-19 diagnosis and
28 was the initial symptom in 26.6%. Some improvement was noted in 27% of patients,
29 with mean time to improvement of 7.2 days in this group (85% of this group improved
30 within 10 days). Our findings suggest that anosmia can be a presenting symptom of
31 COVID-19, consistent with other emerging international reports. Anosmia may be critical
32 in timely identification of individuals infected with SARS-CoV2 **who may unwittingly be**
33 **transmitting the virus**.

34

35 Anosmia (loss of sense of smell) and dysgeusia (alteration of sense of taste) have been
36 reported in association with the COVID-19 pandemic. Dysgeusia may be related to
37 alteration in the perception of taste due to loss of the sense of olfaction. There have
38 been published and non-published anecdotal reports of anosmia related to COVID-19
39 emanating from around the world, including South Korea, Germany, Italy, United
40 Kingdom, Iran and the United States. In South Korea, the Daegu City Council's informal
41 phone survey found 15.3% of 3191 confirmed SARS-CoV-2 cases had anosmia or
42 dysgeusia.¹ Hendrick Streeck, a German virologist, reported a loss of smell and taste in
43 over two-thirds of 100 COVID-19 positive people interviewed with mild symptoms.²
44 Massimo Galli, an Italian infectious disease specialist at the University of Milan, noted
45 that anosmia and dysgeusia seems to be observed in patients even with modest
46 symptoms or limited severity, however, they appear to present later in the course of
47 infection.³ A non-peer reviewed Iranian study⁴ on 10,069 subjects with anosmia or
48 hyposmia (unknown COVID-19 status) noted sudden symptom onset in 76.2%
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50
51 In an effort to establish a platform allowing healthcare providers of all specialties
52 worldwide to submit data to confidentially report on anosmia symptoms related to
53 COVID-19, the American Academy of Otolaryngology—Head and Neck Surgery (AAO-
54 HNS) established the COVID-19 Anosmia Reporting Tool for Clinicians (Supplement
55 Data 1).⁵ The survey was developed by expert panelists and stakeholders from the
56 AAO-HNS Infectious Disease and Patient Safety Quality Improvement Committees.
57 After multiple iterations, consensus was reached, satisfying adequate face validity. The
58 content of the tool, especially the data elements, was based on review of multiple

59 COVID-19 reports related to anosmia. As this is a pilot study conducted in an expedited
60 manner to address the rapidly changing situation, additional validation will be
61 forthcoming. Data collection is hosted on a platform similarly used for the AAO-HNS
62 Patient Safety Event Reporting Tool,⁶ with digital safeguards built to ensure anonymity.
63 No identifiable data about the user was solicited. Computer's IP address was not
64 captured from the submitting provider to preserve the confidentiality of the
65 report/reporter. If identifiable information was inadvertently provided by the reporter in
66 the free-text entry boxes, this information was immediately discarded. This preliminary
67 analysis is based on data collected from the opening of the survey March 25, 2020 to
68 April 3, 2020. Descriptive statistics were used to describe submissions to the database.
69
70 In 10 days of data accrual, 240 entries were made. Three were removed due to clinical
71 inconsistencies and thus 237 entries were analyzed. While otolaryngologists contributed
72 47% of entries, the majority came from a variety of other specialists (Figure 1).
73 Demographics are shown in Table 1. Age distribution histogram is found in Appendix 1.
74 Over one-third of the cases were of healthcare workers.
75
76 One of the most relevant findings is the timing of anosmia in relationship to diagnosis
77 and the presence—or absence—of other symptoms (Table 2). Anosmia was noted in
78 73% prior to diagnosis. Anosmia contributed to recommending testing in 40%. More
79 critically, anosmia was the initial symptom in more than a quarter of patients. The
80 remainder demonstrated more common symptoms of COVID-19, with myalgias and
81 sore throat highly noted as free-text entries in the “Other” category. Some improvement
82 in anosmia was noted in 27% of patients, with mean time to improvement of 7.2 days in

83 this group (85% of this group improved within 10 days). This data is subject to
84 significant interpretation as many entries were submitted before long-term follow up was
85 achieved.

86
87 Although the exact pathophysiology of how SARS-CoV-2 could produce olfactory
88 dysfunction has not been firmly established, direct extension through the nasal mucosa
89 (via angiotensin-converting enzyme 2 receptor on the basal layer of the nasal
90 epithelium) and/or extension to the olfactory bulb are potential hypotheses. Post-viral
91 olfactory dysfunction is a common cause of olfactory dysfunction and is thought to be
92 caused by neuroepithelial dysfunction.^{7,8} Deems et al⁶ reported 26% of patients had
93 anosmia as a result of an upper respiratory infection or cold, with a preponderance of
94 females (63%) being affected more than males. More recently, Fornazieri et al⁹ reported
95 a 13% incidence of post-viral loss of smell.

96
97 Although coronaviruses are a known etiology for post-viral olfactory dysfunction,¹⁰ there
98 is only one case report of SARS-CoV producing anosmia.¹¹ SARS-CoV-2 appears to
99 differ in this regard, as the mounting anecdotal evidence caused ENT UK and the AAO-
100 HNS to announce the potential association.^{12,13} For the current COVID-19 pandemic,
101 scientific data is emerging. A European multi-center study released recently had 417
102 COVID-19 patients with mild-to-moderate symptoms (mean age = 36.9 years; 263
103 (63%) female).¹⁴ Olfactory dysfunction was reported in 11.8% of cases before any other
104 symptom.

105

106 Currently, neither the WHO nor the CDC recognize anosmia as a screening symptom.
107 As we continue to treat this pandemic, it is vital to identify additional symptoms outside
108 of the classic triad of fever, cough, and dyspnea in an effort to promote timely
109 identification of infected individuals who may unknowingly transmit the virus.
110 Characteristic symptoms inclusive of anosmia can be utilized to direct early and
111 widespread testing to mitigate this disease.

112
113 With this survey, some caveats should be considered. Entries are provider initiated and
114 limited by the awareness of the tool. Because of limitations to testing availability,
115 diagnosis of some patients is presumed and not confirmed. By the nature of the capture
116 strategy, analysis has been made of a subset of COVID-19 patients only: those that
117 have anosmia. It is difficult to determine the prevalence of this symptom among all
118 COVID-19 patients.

119
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123

124 **Table 1. Demographics, n=237**

125 Age, y

126 Mean (\pm SD) 39.6 (\pm 14.6)

127 Median 36

128 Range 2-89

129 Sex, M/F (%) 108/129 (46/54)

130 Patient location (%)

131 United States 158 (67)

132 Mexico 11 (5)

133 Italy 9 (4)

134 UK 7 (3)

135 Other 52 (22)

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Table 2. Timing of Anosmia

Anosmia onset (%)	
Before diagnosis	172 (73)
Anosmia contributed to	
testing for COVID-19	94 (40)
After diagnosis	65 (27)
Symptoms before anosmia (%)	
None	64 (27)
Fever	90 (38)
Chills	63 (27)
Malaise	93 (39)
Cough	98 (41)
Headache	88 (37)
Nasal congestion	60 (25)
Rhinorrhea	42 (18)
GI distress	24 (10)
Other	28 (13)
Resolution of Anosmia (%)	
Complete resolution	30 (13)
Partial resolution	32 (14)
None, not yet	175 (74)
Mean time to improvement, d (\pm SD)	7.2 (\pm 3.1)

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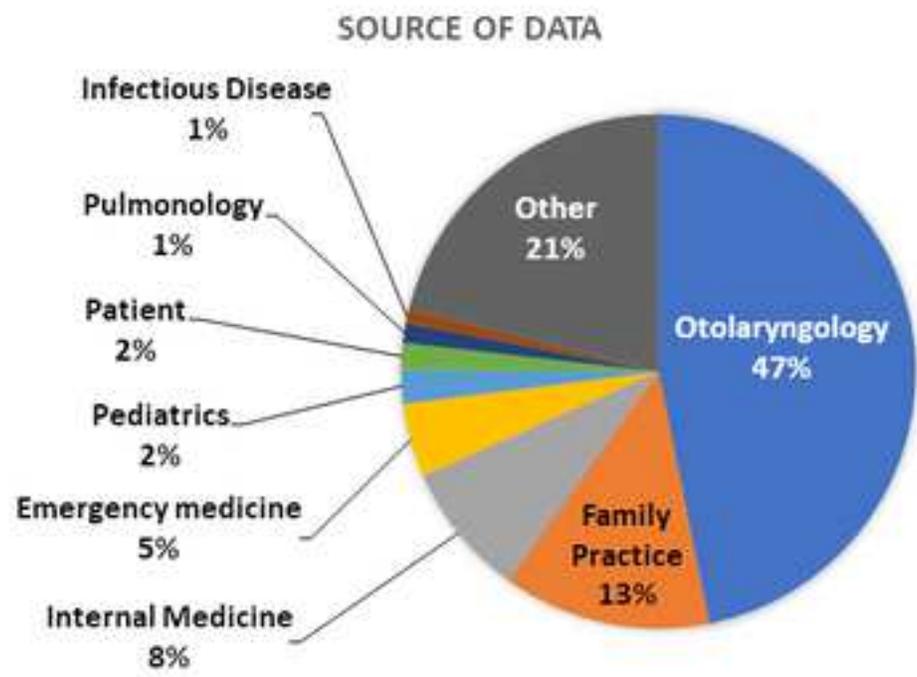
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207 Figure 1. Sources of Data.

208 Supplement Data 1. The COVID-19 Anosmia Reporting Tool.

209 Supplement Data 2. Age Distribution of Patients

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COVID-19 Anosmia Reporting Tool

There is rapidly accumulating anecdotal evidence that anosmia with resultant dysgeusia are frequently reported symptoms associated with the COVID-19 pandemic. Similar reports are surfacing from multiple countries around the world including the United States. In an effort to establish the importance of these symptoms in diagnosis and progression of COVID-19, the American Academy of Otolaryngology-Head and Neck Surgery (AAO-HNS) has established an **Anosmia Reporting Tool**. This tool was developed by the AAO-HNS Infectious Disease and Patient Safety Quality Improvement Committees to allow healthcare providers of all specialties worldwide to submit data to confidentially report on anosmia and dysgeusia related to COVID-19.

There are safeguards to ensure the confidentiality of reporting. No identifiable data about the user is submitted with the report and the computer's IP address is not captured from the submitting provider to preserve the confidentiality of the report/reporter. Further, any report containing identifiable information (hospital name, location, practice name, etc.) is immediately discarded.

To maintain confidentiality, this report needs to be completed online. However, for your convenience, you may view the questions and download a pdf of the Anosmia Reporting Tool below:

Instructions:

- Please respond to as many questions as possible
- Do not report any information that may identify the physician, patient or institution in the questions with the free text boxes
- No identifiable information is collected from the submitting provider
- Once your data is submitted, an email will be sent to AAO-HNS staff notifying them of the new submission

You may contact the AAO-HNS research business unit regarding the Anosmia Reporting Tool at Anomsia@entnet.org

COVID-19 ANOSMIA REPORTING TOOL

1. Medical Specialty of Submitting Provider: (free text)_____
2. Patient age: ____ (Only enter a number)
3. Patient Gender: ____ Male ____ Female
4. Patient location at time of diagnosis
 ____ United States
 ____ Other
 4a. If Other, please specify country: (free text)_____
5. Is the source of the COVID-19 infection identifiable? ____ Yes ____ No

_____ Other

9a. If "Other," please provide detail: (free text)_____

10. What symptoms did the patient have **AT THE TIME** of anosmia/dysgeusia?

- _____ None
- _____ Fever
- _____ Chills
- _____ Malaise
- _____ Cough
- _____ Headache
- _____ Nasal Congestion
- _____ Rhinorrhea
- _____ Gastrointestinal Distress
- _____ Other

10a. If "Other," please provide detail: (free text)_____

11. What was the condition of the COVID-19 infection at the time the anosmia/dysgeusia was observed?

- _____ Inpatient /hospitalized
- _____ Outpatient

12. Did the patient's condition worsen or improve after the anosmia/dysgeusia was observed?

- _____ Worsen
- _____ Improve

13. What is the patient's current COVID-19 infection status

- _____ Active
- _____ Recovered
- _____ Deceased

14. Did the anosmia/dysgeusia resolve?

- _____ Yes
- _____ No

If "Yes" to above:

14a. How long after the anosmia/dysgeusia was observed: (free text)_____

14b. Was it:

- _____ Complete resolution
- _____ Partial Resolution

15. Did the patient receive treatments for the anosmia?

- _____ Yes
- _____ No

If "Yes" to above:

- 15a. What treatments if any did the patient receive?
- 15b. How soon after did the anosmia/dysgeusia resolve:
- 17. Comments/Additional information: (free text)_____

