A Clinical Study of Smell Disorders in COVID-19 Patients in a Tertiary Care Hospital in Pondicherry: A Cross Sectional Study

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Abstract Anosmia and ageusia are the first and maybe the only symptom in patients affected with COVID-19 especially if the patient is paucisymptomatic. This aim of this study was to determine the demographic details of patients with anosmia, prevalence of anosmia and the time taken for it to resolve in patients who are positive for COVID-19 and took treatment in our hospital. Cross Sectional Study. Patients with real time polymerase chain reaction (RT-PCR) positive nasopharyngeal and oropharyngeal swabs, who met the inclusion and exclusion criteria were included in the study. The study group was interviewed through telephonic calls and a questionnaire filled to see the development and regression of their symptoms. Of the study population of 1000, 742 patients had some sort of a smell disturbance. There was a positive correlation between the severity of the disease and history of smoking. The prevalence of smell disturbances among COVID-19 patients in our study was 74.2%. One important finding that we found out was that majority of the smokers had moderate disease. Most of the patients had complete recovery form smell disturbance in the due course of time. The mean time for resolution of smell disturbance was found to be 9.89 days. Anosmia and ageusia can represent the only symptomatology present in patients with COVID-19 and they are completely reversible and hence they can be used as early predictors of infection.

Keywords COVID-19 · Coronavirus · Anosmia · Otorhinolaryngology

Introduction

Several patients with pneumonia of unidentified causes were seen in the Wuhan city of China by the end of 2019, which was later on named as COVID-19 by the World Health Organization. From then on, there was a steady and very heavy spill over to various parts of the world. Coronavirus disease is an acute respiratory illness caused by a novel human coronavirus (SARS-CoV-2, called COVID-19 virus), which causes higher mortality in people aged ≥ 60 years and also in people with underlying co-morbid conditions such as cardiovascular disease, chronic respiratory disease, diabetes, and cancer. The COVID-19 outbreak was announced as a public health emergency of international concern on January 30, 2020 [1].

The initial symptoms of the disease are fatigue, fever, dry or productive cough, shortness of breath, muscle pain, diarrhea, vomiting, anorexia, headache, sore throat, dizziness, palpitations and chest pain. As more and more cases were reported, the spectrum of symptoms also got expanded and included few symptoms of smell disturbances and taste disturbances also. As the number of cases reported in medical literature increased, there has been an increment in the number of cases which presents with anosmia or ageusia alone as the presenting complaint. A retrospective study by Gilani et al., in April 2020 reported 8 cases of anosmia of which 5 patients were confirmed cases of COVID-19 [2]. Corona virus is known to cause the common cold like symptoms which is associated with olfactory loss [3]. The initial reports from China did not include anosmia as a presenting symptom of COVID-19.
A possible association between COVID-19 and anosmia was first reported in a short communication from Italy in the month of July, 2020 and it was a warning symptom especially for the otorhinolaryngologists regarding this possible new manifestation of COVID-19. They reported the presence of anosmia and ageusia as the first and maybe the only symptom in patients affected with COVID-19 especially if the patient is paucisymptomatic [4].

We aimed to determine the demographic details of patients with anosmia, prevalence of anosmia and the time taken for it to resolve in patients who are positive for COVID-19 and took treatment in our hospital.

Materials and Methods

This was a cross sectional study done at Mahatma Gandhi Medical College and Research Institute, Pondicherry, India from a period of August 2020 to December 2020 after obtaining clearance from the Institutional Research Committee of our University. Patients with real time polymerase chain reaction (RT-PCR) positive nasopharyngeal and oropharyngeal swabs, (positive for COVID-19) were the study population. The first reported case of COVID-19 in Pondicherry was in March 2020, however in-patient care of positive patients in our hospital was started only in August 2020. The real time PCR of the nasopharyngeal and oropharyngeal swabs for the detection of SARS-CoV-19 was done in our centre and worldwide, and they were subjected to only telephonic interviews and therefore no ethical consideration is involved in this study.

Inclusion criteria were patients who are laboratory proven SARS-CoV-2 infection with a mild or moderate presentation of COVID-19 according to the severity classification of the Massachusetts General Hospital COVID-19 treatment guidance [5] above 18 years who give informed consent to participate in the study. Exclusion criteria were, patients with a severe form of COVID-19, who did not give consent to take part, subjects with incomplete data, previous anosmia and those who were lost to follow-up (not responding to 2 telephone calls) were excluded.

The study included a convenience sample of SARS-CoV-2 polymerase chain reaction (PCR) confirmed cases being admitted for care or assessed and discharged during the study period and was optimized to 1000. Patient underwent a telephonic interview on the development of symptoms and then they were serially interviewed at one week intervals for up to 5 weeks or when the symptoms regress, on the progression and regression of symptoms and the time taken for the smell disturbance to resolve was noted. Interview data were anonymously collected and no reward was offered for participation.

Data regarding the demography, history of recent travel, presenting symptoms (fever, myalgia, cough, dyspnea, fatigueability, sore throat, anosmia, ageusia, diarrhea, etc.), the severity of the disease, past nasal and paranasal diseases (NPND) and history of smoking were taken from every patient who were included in the study (Fig. 1).

Results

The age of the patients ranged from 18 to 68 years with a mean age of 42.35 years. There was a male preponderance 580 patient (58%), with females of 388 (38.8%) and 32 (3.2%) patients were transgender population. 865 (86.5%) patients had mild disease as per severity classification of the Massachusetts General Hospital COVID-19 treatment guidance and 135 (13.5%) had moderate symptoms (Tables 1 and 2; Fig. 2).

Most of the cases were without any history of smoking 798 (79.8%) and majority of the patients with history of smoking were having moderate degree of symptoms (99%—200 patients) and 2 patients (1%). Majority of our patients did not have a history of travel, 948 patients (94.8%). 625 patients had a previous history of diagnosed nose or paranasal sinus disease like deviated nasal septum, allergic rhinitis, or sinusitis and 204 patients gave a previous history of nasal surgeries in the past (Tables 1 and 2; Fig. 3).

We had a total study population of 1000 subjects of which 742 patients had some sort of a smell disturbance, 258 patients did not have any smell disturbance. 163 patients had reduced smell sensation, 123 had altered smell sensation and 456 patients had complete loss of smell sensation. Out of the 742 patients with smell disturbances, 652 (87.8%) had complete recovery, 84 (11%) had partial recovery which means symptoms were better than before and 6 patients (0.8%) had no recovery.

Of the study population, 562 patients (56.2%) had altered smell sensation as the only presenting complaint, 438 (43.8%) patients had smell disturbance along with taste abnormalities or other complaints. There was an average resolution time of 7–28 days with a mean of 9.89 days. No death was reported in our study during the study period.

Discussion

Various theories have been postulated as the causative factor for the development of smell and taste abnormalities in COVID-19. A study by Zhou et al. have reported that COVID-19 uses the same receptor [cellular angiotensin-converting enzyme 2 (ACE 2)] as SARS-CoV [6]. The enzyme found in the tongue. Therefore, it is possible that the COVID-19 causes taste dysfunction in the same way as
the ACE2 inhibitors [7]. Secondly, the taste disturbance in the presence of smell abnormality is that both chemosensory senses are closely associated [8]. A study by Brann et al. [9] suggested that the involvement of non-neuronal olfactory epithelium by COVID-19 virus is what causes anosmia and the associated taste dysfunctions. This is the most acceptable mechanism because most of the patients with COVID-19 infection are of the mild severity, outpatient cases, and most of the Smell and Taste abnormalities (STA) resolve within short period of less than 2 weeks [10]. Damaged olfactory cells can regenerate, but they may not regenerate normally and this faulty regeneration is what results in parosmia experienced by most of the patients. However, a larger number of a cohort studies and other meta analyses with long term follow-up and better methods of assessment are needed to assess the exact time and rate of recovery of the STA in confirmed cases of this novel coronavirus infection.

**Table 1** Incidence of smoking in patients with COVID-19

| History of smoking | Number of patients |
|--------------------|--------------------|
| Yes                | 202 (200 had moderate symptoms and 2 had mild symptoms) |
| No                 | 798                |

**Fig. 1** Questionnaire used for the study
Disorders of smell and taste are newly found problems during the pandemic of COVID-19. Smell disturbance alone, smell disturbance with or without taste abnormality or smell disturbance with other COVID symptoms are the usual presenting features noted. Smell and taste abnormalities were the first and the only complaints in 10% of subjects, 19% experienced STA before other classical symptoms like fever and cough, as well as 25% of the children had only STA at the time of the presentation in a study reported by Qiu et al. [11]. Therefore, these disorders are early indicators of COVID-19 disease and can be helpful for screening and infection control. Our study design included a questionnaire for patients about the development of anosmia alone and the time taken for it to resolve. In a prospective, cross-sectional telephone questionnaire study by Speth et al., who reported that the prevalence rate of olfactory dysfunction was 61.2% [12]. Another study by Paderno et al., showed that the olfactory dysfunction was seen in 83% and gustatory dysfunctions in 89% of the patients [13]. A systematic review of literature by Tong et al. reported the prevalence rate of olfactory dysfunction in COVID-19 patients as 52.73% [14]. In a large prospective multicenter case series study by Chary et al., it was found that the rate of STA without nasal complaints was 70% [15]. This wide variability among various studies might be due to the variation in the ethnicity, differences in sample size, and type of the vast sample (children or adults or both, hospitalized, non-hospitalized or both, with or without nasal symptoms, subjective, objective or both assessment, and severity of the disease).

Our study had a male preponderance of 58% with an incidence in transgender population in 3.2%. Incidence of COVID-19 has not been reported in literature and the incidence in our study can be because our hospital is a centre for sex reassignment surgery in south India and we have a perennial transgender clinic running in our hospital. This difference between our study and the previous studies might be attributed to the differences in the social and cultural behaviors and geographical locations. All the patients in our study had mild symptoms (86.5%) to moderate symptoms (13.5%) only. This is because our samples were taken from the COVID ward of our hospital and not from the ICU patients or from patients who needed oxygen support.

In the early period of COVID-19 pandemic, recent travel was considered as an important factor for the transmission of the disease especially to China or International travel. But as the time progressed and the infection progressed to become a worldwide disease, with community spread, history of recent travel became less and less important. Moreover, the contact with suspicious, asymptomatic or proven COVID-19 cases is an important way of the transmission of the infection to healthy people. Our findings did not reveal any significance between recent history of travel and the development of disease.

Disorders of smell and taste are newly found problems during the pandemic of COVID-19.
rather than nonsmokers in the MERS-CoV outbreak of 2012 [16]. Smoking has a negative impact on the severity and clinical outcome in patients with the COVID-19 disease [16]. In our study, majority of the patients were non-smokers (79.8%). However, 99% of the patients with moderate disease were smokers which strengthens the well-known fact that smoking causes irreversible lung injury and make them prone to infectious diseases. These findings were contradictory to two similar studies by Speth et al. [12] and by Al-Ani who reported that 8.8% of their patients were current smokers and 20% smokers respectively [17].

Olfactory dysfunction presenting as anosmia or hyposmia are generally one of the first symptoms in COVID-19 disease [12, 18]. A study by Yan et al. [19] reported that a statistically significant association was seen between STA and mild disease and severe disease type was usually not associated with the STA. Therefore, the STA can act as a major positive factor for the prognosis of COVID-19 disease and to segregate patients as mild and moderate disease. Of the study population, 562 patients (56.2%) had anosmia as the only presenting complaint, 438 (43.8%) patients had anosmia along with taste abnormalities or other symptoms like throat pain and dry cough.

In our study, we found that 742 patients in the total study population had smell abnormalities. The prevalence of smell abnormality was found to be 74.2%. Complete recovery was observed in 88% of the cases, partial recovery was noted in 11% cases, and lack of recovery in 6 patients (1%) after 5 weeks of follow-up. We noted a resolution time range of 7–28 days with a mean of 9.89 days. A study by Vaira et al. noted a spontaneous resolution time range of 7–28 days with a mean of 9.89 days. Anosmia and ageusia can represent the only symptomatology present in patients with COVID-19 and they are completely reversible and hence they can be used as early predictors of infection especially for Otorhinolaryngologists.

**Conclusion**

The COVID-19 pandemic caused by novel human coronavirus has made knowledge regarding respiratory and personal hygiene very important [21]. The prevalence of smell disturbances among COVID-19 patients in our study was 74.2%. There was a male preponderance with incidence of disease in transgender population also and the mean age group was 42.35 years. Majority of the patients in our study group did not have any travel history and only 20% of the patients had history of smoking. One important finding that we found out was that majority of the smokers had moderate disease. Most of the patients had complete recovery form smell disturbance in the due course of time. The mean time for resolution of smell disturbance was found to be 9.89 days.

Anosmia and ageusia can represent the only symptomatology present in patients with COVID-19 and they are completely reversible and hence they can be used as early predictors of infection especially for Otorhinolaryngologists.

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**Declarations**

**Conflict of interest** The authors declare that they have no conflict of interest.

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