



**HAL**  
open science

## **The best COVID-19 predictor is recent smell loss: a cross-sectional study**

Richard Gerkin, Kathrin Ohla, Maria Geraldine Veldhuizen, Paule Joseph, Christine Kelly, Alyssa Bakke, Kimberley Steele, Michael Farruggia, Robert Pellegrino, Marta Pepino, et al.

### ► **To cite this version:**

Richard Gerkin, Kathrin Ohla, Maria Geraldine Veldhuizen, Paule Joseph, Christine Kelly, et al.. The best COVID-19 predictor is recent smell loss: a cross-sectional study. 2020. hal-02979656v1

**HAL Id: hal-02979656**

**<https://inserm.hal.science/hal-02979656v1>**

Preprint submitted on 16 Nov 2020 (v1), last revised 27 Oct 2020 (v2)

**HAL** is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

# The best COVID-19 predictor is recent smell loss: a cross-sectional study

## Byline authors:

Richard C. Gerkin,<sup>1\*</sup> Kathrin Ohla,<sup>2\*</sup> Maria G. Veldhuizen,<sup>3</sup> Paule V. Joseph,<sup>4,5,6</sup> Christine E. Kelly,<sup>7</sup> Alyssa J. Bakke,<sup>8</sup> Kimberley E. Steele,<sup>6,9</sup> Michael C. Farruggia,<sup>10</sup> Robert Pellegrino,<sup>11</sup> Marta Y. Pepino,<sup>12</sup> Cédric Bouysset,<sup>13</sup> Graciela M. Soler,<sup>14,15</sup> Veronica Pereda-Loth,<sup>16</sup> Michele Dibattista,<sup>17</sup> Keiland W. Cooper,<sup>18</sup> Ilja Croijmans,<sup>19</sup> Antonella Di Pizio,<sup>20</sup> M. Hakan Ozdener,<sup>21</sup> Alexander W. Fjaeldstad,<sup>22</sup> Cailu Lin,<sup>21</sup> Mari A. Sandell,<sup>23</sup> Preet B. Singh,<sup>24</sup> V. Evelyn Brindha,<sup>25</sup> Shannon B. Olsson,<sup>26</sup> Luis R. Saraiva,<sup>27</sup> Gaurav Ahuja,<sup>28</sup> Mohammed K. Alwashahi,<sup>29</sup> Surabhi Bhutani,<sup>30</sup> Anna D'Errico,<sup>31</sup> Marco A. Fornazieri,<sup>32</sup> Jérôme Golebiowski,<sup>13</sup> Liang-Dar Hwang,<sup>33</sup> Lina Öztürk,<sup>3</sup> Eugeni Roura,<sup>34</sup> Sara Spinelli,<sup>35</sup> Katherine L. Whitcroft,<sup>36</sup> Farhoud Faraji,<sup>37</sup> Florian Ph.S Fischmeister,<sup>38</sup> Thomas Heinbockel,<sup>39</sup> Julien W. Hsieh,<sup>40</sup> Caroline Huart,<sup>41</sup> Iordanis Konstantinidis,<sup>42</sup> Anna Menini,<sup>43</sup> Gabriella Morini,<sup>44</sup> Jonas K. Olofsson,<sup>45</sup> Carl M. Philpott,<sup>46</sup> Denis Pierron,<sup>16</sup> Vonnie D.C. Shields,<sup>47</sup> Vera V. Voznessenskaya,<sup>48</sup> Javier Albayay,<sup>49</sup> Aytug Altundag,<sup>50</sup> Moustafa Bensafi,<sup>51</sup> María Adelaida Bock,<sup>52</sup> Orietta Calcinoni,<sup>53</sup> William Fredborg,<sup>45</sup> Christophe Laudamiel,<sup>54</sup> Juyun Lim,<sup>55</sup> Johan N. Lundström,<sup>56</sup> Alberto Macchi,<sup>57,58</sup> Pablo Meyer,<sup>59</sup> Shima T. Moein,<sup>60</sup> Enrique Santamaría,<sup>61</sup> Debarka Sengupta,<sup>28</sup> Paloma Rohlf's Dominguez,<sup>62</sup> Hüseyin Yanik,<sup>63</sup> GCCR Group Author, Thomas Hummel,<sup>90</sup> John E. Hayes,<sup>8</sup> Danielle R. Reed,<sup>21</sup> Masha Y. Niv,<sup>108</sup> Steven D. Munger,<sup>85,109†</sup> Valentina Parma<sup>110†c</sup>

<sup>1</sup>School of Life Sciences, Arizona State University, <sup>2</sup>Institute of Neuroscience and Medicine (INM3), Forschungszentrum Jülich, <sup>3</sup>Department of Anatomy, Mersin University, <sup>4</sup>National Institutes of Nursing Research, <sup>5</sup>National Institute of Alcohol Abuse and Alcoholism, <sup>6</sup>National Institutes of Health, <sup>7</sup>AbScent, <sup>8</sup>Food Science, The Pennsylvania State University, <sup>9</sup>National Institute of Diabetes and Digestive and Kidney Diseases, <sup>10</sup>Interdepartmental Neuroscience Program, Yale University, <sup>11</sup>Food Science, University of Tennessee, <sup>12</sup>Department of Food Science and Human Nutrition, University of Illinois at Urbana Champaign, <sup>13</sup>Institut de Chimie de Nice, UMR CNRS 7272, Université Côte d'Azur, <sup>14</sup>Grupo de Estudio de Olfato y Gusto (GEOG), Buenos Aires, <sup>15</sup>Department of Otorhinolaryngology University of Buenos Aires, <sup>16</sup>Medecine Evolutive UMR5288, University of Toulouse, <sup>17</sup>Department of Basic Medical Science, Neuroscience and Sense Organs, Università degli Studi di Bari A. Moro, <sup>18</sup>Department of Neurobiology and Behavior, University of California, Irvine, <sup>19</sup>Department of Psychology, Utrecht University, <sup>20</sup>Leibniz-Institute for Food Systems Biology at the Technical University of Munich, <sup>21</sup>Monell Chemical Senses Center, <sup>22</sup>Flavour Clinic, Department of Otorhinolaryngology, Regional Hospital West Jutland, <sup>23</sup>Department of Food and Nutrition, University of Helsinki, <sup>24</sup>Department of Oral Surgery and Oral Medicine, Faculty of Dentistry, University of Oslo, <sup>25</sup>Department of Electrical and Electronics Engineering, Karunya Institute of Technology and Sciences, <sup>26</sup>National Centre for Biological Sciences, Tata Institute of Fundamental Research, <sup>27</sup>Research Branch, Sidra Medicine, <sup>28</sup>Computational Biology, Indraprastha Institute of Information Technology, <sup>29</sup>ENT Division, Surgery Department, Sultan Qaboos University, <sup>30</sup>School of Exercise and Nutritional Sciences, San Diego State University, <sup>31</sup>Cellular and Molecular Neurobiology, Goethe University Frankfurt, <sup>32</sup>Clinical Surgery, Universidade Estadual de Londrina, <sup>33</sup>The University of Queensland Diamantina Institute, The University of Queensland, <sup>34</sup>Centre for Nutrition and Food Sciences, Queensland Alliance for Agriculture and Food Innovation, The University of Queensland, <sup>35</sup>Department of Agriculture, Food, Environment and Forestry (DAGRI), University of Florence, <sup>36</sup>Ear Institute, University College London, <sup>37</sup>Department of Surgery, Division of Otolaryngology-Head and Neck Surgery, UC San Diego Health, <sup>38</sup>Department of Psychology, University of

Graz,<sup>39</sup> Department of Anatomy, Howard University College of Medicine,<sup>40</sup> Department of Otorhinolaryngology, Rhinology-Olfactology Unit, Geneva University Hospitals,<sup>41</sup> ENT Department, Cliniques universitaires Saint-Luc,<sup>42</sup> ORL Department, Papageorgiou Hospital, Aristotle University,<sup>43</sup> Neuroscience Area, SISSA, International School for Advanced Studies,<sup>44</sup> University of Gastronomic Sciences,<sup>45</sup> Department of Psychology, Stockholm University,<sup>46</sup> Norwich Medical School, The Norfolk Smell & Taste Clinic, University of East Anglia,<sup>47</sup> Biological Sciences Department, Fisher College of Science and Mathematics, Towson University,<sup>48</sup> Severtsov Institute of Ecology and Evolution RAS,<sup>49</sup> Department of General Psychology, University of Padova,<sup>50</sup> Otorhinolaryngology Department, Biruni University,<sup>51</sup> Lyon Neuroscience Research Center, CNRS,<sup>52</sup> Departamento de Salud Pública ORL, Hospital General Barrio Obrero,<sup>53</sup> Private practice, Milan,<sup>54</sup> Scent Engineering, DreamAir LLC,<sup>55</sup> Food Science and Technology, Oregon State University,<sup>56</sup> Department of Clinical Neuroscience, Karolinska Institutet,<sup>57</sup> ENT Department University of Insubria Varese, Assi-Sette Laghi,<sup>58</sup> Italian Academy of Rhinology,<sup>59</sup> Health Care and Life Sciences, IBM TJ. Watson Research Center,<sup>60</sup> School of Biological Sciences, Institute for Research in Fundamental Sciences,<sup>61</sup> Clinical Neuroproteomics Unit, Navarrabiomed-IdiSNA,<sup>62</sup> Psychology and Anthropology, University of Extremadura,<sup>63</sup> Department of Electrical and Electronics Engineering, Mersin University,<sup>90</sup> Smell & Taste Clinic, Department of Otorhinolaryngology, TU Dresden,<sup>108</sup> The Faculty of Agriculture, Food and Environment, The Hebrew University of Jerusalem,<sup>109</sup> Department of Pharmacology and Therapeutics, University of Florida, Department of Psychology,<sup>1110</sup> Temple University

**Non-byline authors (to be listed as collaborators in PubMed under the GCCR Group Author):**

Sanne Boesveldt,<sup>64</sup> Jasper H.B. de Groot,<sup>65</sup> Caterina Dinnella,<sup>35</sup> Jessica Freiherr,<sup>66</sup> Tatiana Laktionova,<sup>48</sup> Sajidxa Mariño,<sup>67</sup> Erminio Monteleone,<sup>35</sup> Alexia Nunez-Parra,<sup>68</sup> Olagunju Abdulrahman,<sup>69</sup> Marina Ritchie,<sup>18</sup> Thierry Thomas-Danguin,<sup>70</sup> Julie Walsh-Messinger,<sup>71</sup> Rashid Al Abri,<sup>29</sup> Rafieh Alizadeh,<sup>72</sup> Emmanuelle Bignon,<sup>13</sup> Elena Cantone,<sup>73</sup> Maria Paola Cecchini,<sup>74</sup> Jingguo Chen,<sup>75</sup> Maria Dolores Guàrdia,<sup>76</sup> Kara C. Hoover,<sup>77</sup> Noam Karni,<sup>78</sup> Marta Navarro,<sup>34</sup> Alissa A. Nolden,<sup>79</sup> Patricia Portillo Mazal,<sup>80</sup> Nicholas R. Rowan,<sup>81</sup> Atiye Sarabi-Jamab,<sup>82</sup> Nicholas S. Archer,<sup>83</sup> Ben Chen,<sup>84</sup> Elizabeth A. Di Valerio,<sup>85</sup> Emma L. Feeney,<sup>86</sup> Johannes Frasnelli,<sup>87</sup> Mackenzie E. Hannum,<sup>21</sup> Claire Hopkins,<sup>88</sup> Hadar Klein,<sup>89</sup> Coralie Mignot,<sup>90</sup> Carla Mucignat,<sup>91</sup> Yuping Ning,<sup>92</sup> Elif E. Ozturk,<sup>93</sup> Mei Peng,<sup>94</sup> Ozlem Saatci,<sup>95</sup> Elizabeth A. Sell,<sup>96</sup> Carol H. Yan,<sup>37</sup> Raul Alfaro,<sup>12</sup> Cinzia Cecchetto,<sup>49</sup> Gérard Coureaud,<sup>97</sup> Riley D. Herriman,<sup>21</sup> Jeb M. Justice,<sup>85,98</sup> Pavan Kumar Kaushik,<sup>26</sup> Sachiko Koyama,<sup>99</sup> Jonathan B. Overdeest,<sup>100</sup> Nicola Pirastu,<sup>101</sup> Vicente A. Ramirez,<sup>102</sup> S. Craig Roberts,<sup>103</sup> Barry C. Smith,<sup>104</sup> Hongyuan Cao,<sup>105</sup> Hong Wang,<sup>21</sup> Patrick Balungwe Birindwa,<sup>106</sup> Marius Baguma<sup>106</sup>

<sup>64</sup>Division of Human Nutrition and Health, Wageningen University, <sup>65</sup>Behavioral Science Institute, Radboud University, <sup>66</sup>Department of Psychiatry and Psychotherapy, Friedrich-Alexander- Universität Erlangen-Nürnberg, <sup>67</sup>Centro de Otorrinolaringología Respira Libre, <sup>68</sup>Department of Biology, University of Chile, <sup>69</sup>Department of Physiology, The Federal University of Technology, Akure, Nigeria, <sup>70</sup>CSGA-Centre for Taste and Feeding Behavior, INRAE, CNRS, AgroSup Dijon, University Bourgogne Franche- Comté, <sup>71</sup>Department of Psychology, University of Dayton, <sup>72</sup>ENT and Head and Neck Research Center and Department, the Five Senses Institute, Iran University of Medical Sciences, <sup>73</sup>Neuroscience Department, Federico II University, <sup>74</sup>Department of Neuroscience, Biomedicine and Movement Sciences, Anatomy and Histology Section, University of Verona, <sup>75</sup>Department of Otolaryngology-Head and Neck Surgery, Second Affiliated Hospital of Xi'an Jiaotong University, <sup>76</sup>Food Technology, IRTA, <sup>77</sup>Department of Anthropology, University of Alaska Fairbanks, <sup>78</sup>Department of Medicine, Hadassah - Hebrew University Medical Center, Mt Scopus, <sup>79</sup>University of Massachusetts, <sup>80</sup>Servicio de Otorrinolaringología, Hospital Italiano de Buenos Aires, <sup>81</sup>Department of Otolaryngology-Head and Neck Surgery, Johns Hopkins University School of Medicine, <sup>82</sup>School of Cognitive Sciences, Institute for Research in Fundamental Sciences (IPM), <sup>83</sup>Agriculture and Food, Commonwealth Scientific and Industrial Research Organisation (CSIRO), <sup>84</sup>Department of Psychiatry, The Affiliated Brain Hospital of Guangzhou Medical University, <sup>85</sup>Center for Smell and Taste, University of Florida, <sup>86</sup>Institute of

Food and Health, University College Dublin, <sup>87</sup>Department of Anatomy, Université du Québec à Trois-Rivières, <sup>88</sup>ENT Department, Guy's and St Thomas' Hospitals, <sup>89</sup>Department of Biochemistry, Food Science and Nutrition, The Hebrew University of Jerusalem, <sup>90</sup>Department of Otorhinolaryngology, Smell and Taste Clinic, TU Dresden, <sup>91</sup>Department of Molecular Medicine, University of Padova, <sup>92</sup>Department of Neurology, The Affiliated Brain Hospital of Guangzhou Medical University, <sup>93</sup>Nutrition and Dietetics, Kilis 7 Aralik University, <sup>94</sup>Sensory Neuroscience Laboratory, University of Otago, <sup>95</sup>Department of Otorhinolaryngology, Sancaktepe Education and Research Hospital, <sup>96</sup>University of Pennsylvania, Perelman School of Medicine, <sup>97</sup>CMOENES groups, Centre de Recherche en Neurosciences de Lyon, <sup>98</sup>Department of Otolaryngology, University of Florida, <sup>99</sup>Department of Chemistry, Indiana University, <sup>100</sup>Department of Otolaryngology - Head & Neck Surgery, Columbia University Medical Center, <sup>101</sup>Centre for Global Health, Usher Institute, University of Edinburgh, <sup>102</sup>Department of Public Health, University of California Merced, <sup>103</sup>Division of Psychology, University of Stirling, <sup>104</sup>Centre for the Study of the Senses, University of London, <sup>105</sup>Department of Statistics, Florida State University, <sup>106</sup>First Department of Specialties, Hôpital Provincial Général de Bukavu, Faculty of Medicine, Université Catholique de Bukavu

\* Co-first authors

† Co-last authors

<sup>c</sup> Corresponding author:

Valentina Parma, PhD  
Department of Psychology  
Temple University  
1701 N 13th St  
Philadelphia (PA), 19122  
USA  
Email: [valentina.parma@temple.edu](mailto:valentina.parma@temple.edu)

# Abstract

**Background:** COVID-19 has heterogeneous manifestations, though one of the most common symptoms is a sudden loss of smell (anosmia or hyposmia). We investigated whether olfactory loss is a reliable predictor of COVID-19.

**Methods:** This preregistered, cross-sectional study used a crowdsourced questionnaire in 23 languages to assess symptoms in individuals self-reporting recent respiratory illness. We quantified changes in chemosensory abilities during the course of the respiratory illness using 0-100 visual analog scales (VAS) for participants reporting a positive (C19+; n=4148) or negative (C19-; n=546) COVID-19 laboratory test outcome. Logistic regression models identified singular and cumulative predictors of COVID-19 status and post-COVID-19 olfactory recovery.

**Results:** Both C19+ and C19- groups exhibited smell loss, but it was significantly larger in C19+ participants (mean±SD, C19+: -82.5±27.2 points; C19-: -59.8±37.7). Smell loss during illness was the best predictor of COVID-19 in both single and cumulative feature models (ROC AUC=0.72), with additional features providing negligible model improvement. VAS ratings of smell loss were more predictive than binary chemosensory yes/no-questions or other cardinal symptoms, such as fever or cough. Olfactory recovery within 40 days was reported for ~50% of participants and was best predicted by time since illness onset.

**Conclusions:** As smell loss is the best predictor of COVID-19, we developed the ODoR-19 tool, a 0-10 scale to screen for recent olfactory loss. Numeric ratings  $\leq 2$  indicate high odds of symptomatic COVID-19 ( $4 < OR < 10$ ), which can be deployed when viral lab tests are impractical or unavailable.

# Introduction

The novel coronavirus SARS-CoV-2 responsible for the global COVID-19 pandemic has left a staggering level of morbidity, mortality, and societal and economic disruption in its wake.<sup>1</sup> Early publications<sup>2-8</sup> indicate that sudden smell and taste loss are cardinal, early and potentially specific symptoms of COVID-19,<sup>9</sup> including in otherwise asymptomatic individuals.<sup>10-13</sup> While fever and cough are common symptoms of diverse viral infections, the potential specificity of early chemosensory loss to COVID-19 could make it valuable in screening and diagnosis.

Anosmia and other chemosensory disorders have serious health and quality-of-life consequences for patients. However, the general lack of awareness of anosmia and other chemosensory disorders by clinicians and the public, including their association with upper respiratory infections,<sup>14</sup> contributed to an underappreciated role of chemosensory symptoms in the diagnosis of COVID-19. Additionally, the impact of smell loss as a clinical consequence of COVID-19 has not been adequately addressed. Thus, there is an urgent need to better define the chemosensory dysfunctions associated with COVID-19 and to determine their relevance as predictors of this disease. It is critical to develop rapid clinical tools to efficiently and effectively integrate chemosensory assessments into COVID-19 screening and treatment protocols. Information on the duration and reversibility of post-COVID-19 chemosensory impairment is also lacking.

We used binary, categorical and continuous self-report measures to determine the chemosensory phenotype, along with other symptoms and characteristics, of COVID-19-positive (C19+) and COVID-19-negative (C19-) individuals who had reported recent symptoms of respiratory illness. Using those results in logistic regression models, we identified predictors of COVID-19 and recovery from smell loss. Finally, we propose the **Olfactory Determination Rating** scale for COVID-19 (ODO<sub>R</sub>-19), a quick, simple-to-use, telemedicine-friendly tool to improve the utility of current COVID-19 screening protocols, particularly when access to rapid testing for SARS-CoV-2 is limited.

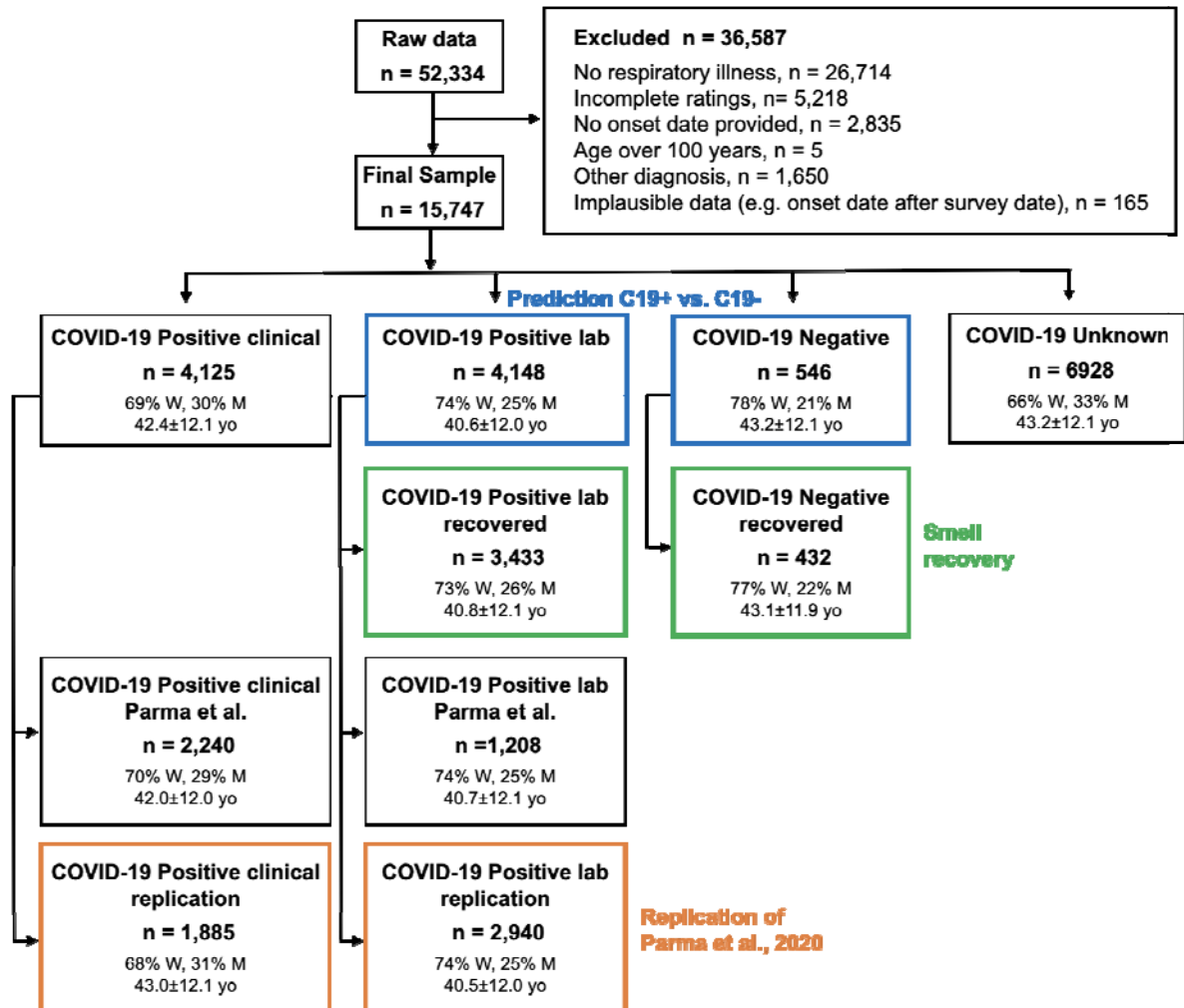
## Methods

### Study design

This preregistered,<sup>15</sup> cross-sectional online study was approved by the Office of Research Protections of The Pennsylvania State University (STUDY00014904); it is in accordance with the revised Declaration of Helsinki, and compliant with privacy laws in the U.S.A. and European Union. Data reported here were collected between April 7 and July 3, 2020 from the Global Consortium for Chemosensory Research (GCCR) core questionnaire (**Appendix 1** and <https://gcchemosensr.org>),<sup>8</sup> an online crowdsourced survey deployed in 32 languages that used binary response and categorical questions (e.g. **Appendix 1**, Questions 6,9) and visual analog scales (e.g., **Appendix 1**, Question 13) to measure self-reported chemosensory ability and other symptoms in adults with current or recent respiratory illness. Data reported here include responses in Arabic, Bengali, Chinese (Simplified and Traditional), Danish, Dutch, English, Farsi, Finnish, French, German, Greek, Hebrew, Hindi, Italian, Japanese, Korean, Norwegian, Portuguese, Russian, Spanish, Swedish, Turkish, and Urdu.

## Participants (272)

A convenience sample of 52,334 volunteers accessed the GCCR questionnaire; 25,620 met eligibility criteria ( $\geq 19$  years old, respiratory illness or suspicion thereof within the past two weeks). After applying pre-registered exclusion criteria, 15,747 participants were included in reported analyses (**Figure 1**).



**Figure 1.** Flow diagram showing the demographics of participants included and excluded in the present analyses. Participants included in the prediction of COVID-19 status are highlighted in blue. Participants included in the smell recovery models are highlighted in green. Participants included in the replication of our prior work are highlighted in orange. N = number of participants; yo = age in years; W = women; M = men. Gender percentages do not include <1% of participants who answered “other” or “preferred not to say”. Participants described in the green boxes are a subset of those described in the boxes listed above them.

Based on the self-reported outcome of a COVID-19 lab test, participants were labeled as either C19+ (positive result) or C19- (negative result). The specific collider bias characterizing this sample (high fraction of C19+ participants and high prevalence of chemosensory disorders in both groups) underestimates the positive correlation between smell loss and COVID-19 (**Figure S1**). Thus, it represents a conservative scenario to test the hypothesis that smell loss reliably predicts COVID-19 status. We benchmarked the GCCR dataset to the representative samples collected with the Imperial College London YouGov Covid 19 Behaviour Tracker (henceforth, YouGov; countries shared across datasets: Brazil, Canada, Denmark, Finland, France, Germany, Italy, Mexico, Netherlands, Norway, Spain, Sweden, UK, USA; YouGov: N=8,674, GCCR: N=3,962; data publicly available at <https://github.com/YouGov-Data/covid-19-tracker>). Benchmarking shows the GCCR

sample underestimates the positive association between smell loss and C19+ (**Figure S1, Table S1**). The country-wise fraction of C19+ participants is correlated ( $r \sim 0.45$ ) when responses from the same calendar week are aligned (**Figure S2**). These findings are in line with other comparisons between crowdsourced versus representative health data,<sup>16</sup> confirming that trends identified in crowdsourced data reasonably approximate population data. Because the GCCR cohort is not demographically balanced, it should not be used to estimate prevalence. However, the representative YouGov cohort indicates globally  $\sim 33\%$  of C19+ individuals report smell loss (**Table S1**).

## Statistical analyses

Statistical analyses were performed in Python 3.7.6 using the pandas,<sup>17</sup> scikit-learn,<sup>18</sup> and statsmodels<sup>19</sup> packages. The data and annotated code is included as supplemental material and will be publicly available on GitHub (<http://github.com/GCCR/GCCR002>) upon publication. Missing values for covariates used in prediction models of COVID-19 status and smell recovery were imputed as follows: binary features = 0.5, numeric features = median, categorical variables = "Missing". Prediction targets themselves were never imputed. Responses incompatible with model generalization (e.g., open ended questions) were excluded. A one-hot encoding was applied to all categorical variables to produce binary indicators of category membership. L1-regularized logistic regression (penalty  $\alpha=1$ ) consistently produced sparse models with comparable cross-validation accuracy and were therefore the prediction test of choice. Model quality was measured using receiver operating characteristic (ROC) area under the curve (AUC). Cross-validation was performed in 100 random splits of 80% training set and 20% test set, and ROC curves are concatenated over each test set. ROC curves are computed on predicted probabilities from each model, circumventing the high-cardinality bias of AUC. For single feature models, AUC is independent of most modeling details, including all rank-invariant decisions. To correctly compute p-values for model coefficients, the normalized data were standardized (mean 0, variance 1) and then coefficients back-transformed to normalized form after fitting.

## Results

### Chemosensory loss associates with COVID-19

A preregistered replication of our prior study<sup>8</sup> confirmed that reported smell, taste, and chemesthesis abilities drop significantly in both lab-tested C19+ participants and those diagnosed by clinical assessment (**Figure S3, Table S2**).

Next, we compared lab-tested C19+ and C19- participants. C19+ participants reported a greater loss of smell (C19+:  $-82.5 \pm 27.2$  points; C19-:  $-59.8 \pm 37.7$  points;  $p=2.2e-46$ , extreme evidence of difference:  $BF_{10}=8.97e+61$ ; **Figure 2A,B; Table S3**), taste (C19+:  $-71.6 \pm 31.8$  points; C19-:  $-55.2 \pm 37.5$  points;  $p=6.7e-26$ , extreme evidence of difference:  $BF_{10}=6.67e+24$ ; **Figure 2C,D; Table S3**) and chemesthesis ability (C19+:  $-36.8 \pm 37.1$  points; C19-:  $-28.7 \pm 37.1$  points;  $p=1.6e-07$ , extreme evidence of difference:  $BF_{10}=3182$ ; **Figure 2E,F; Table S3**). However, both groups reported a similar degree of nasal obstruction (**Figure 2G,H; Table S3**). Self-reported changes in smell, taste, and chemesthesis were highly correlated within both groups (C19+:  $0.71 < r < 0.83$ ; C19-:  $0.76 < r < 0.87$ ) and orthogonal to nasal obstruction changes (C19+:  $r=-0.20$ ; C19-:  $r=-0.13$ ).

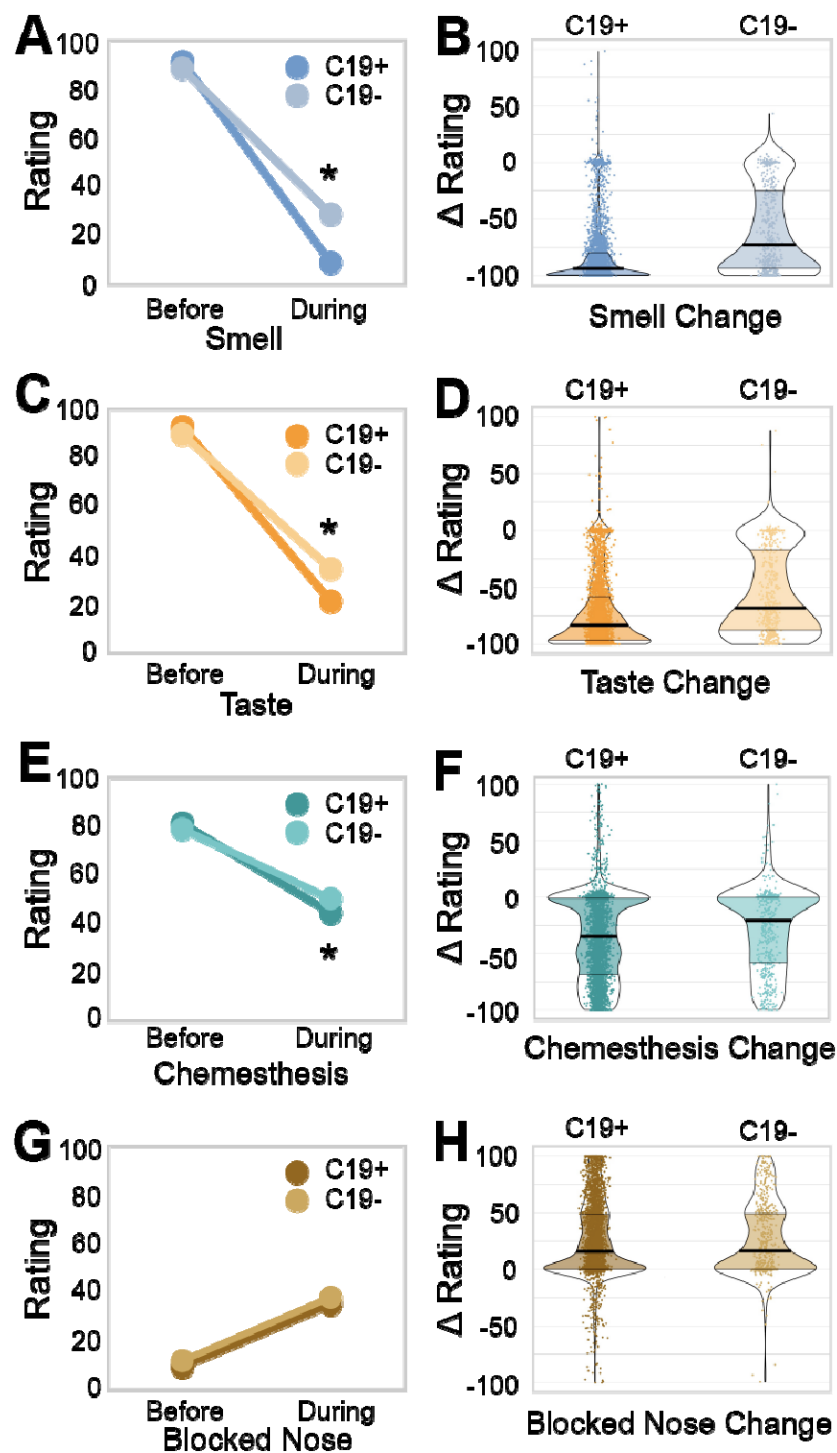


## Prediction of COVID-19 status from survey responses

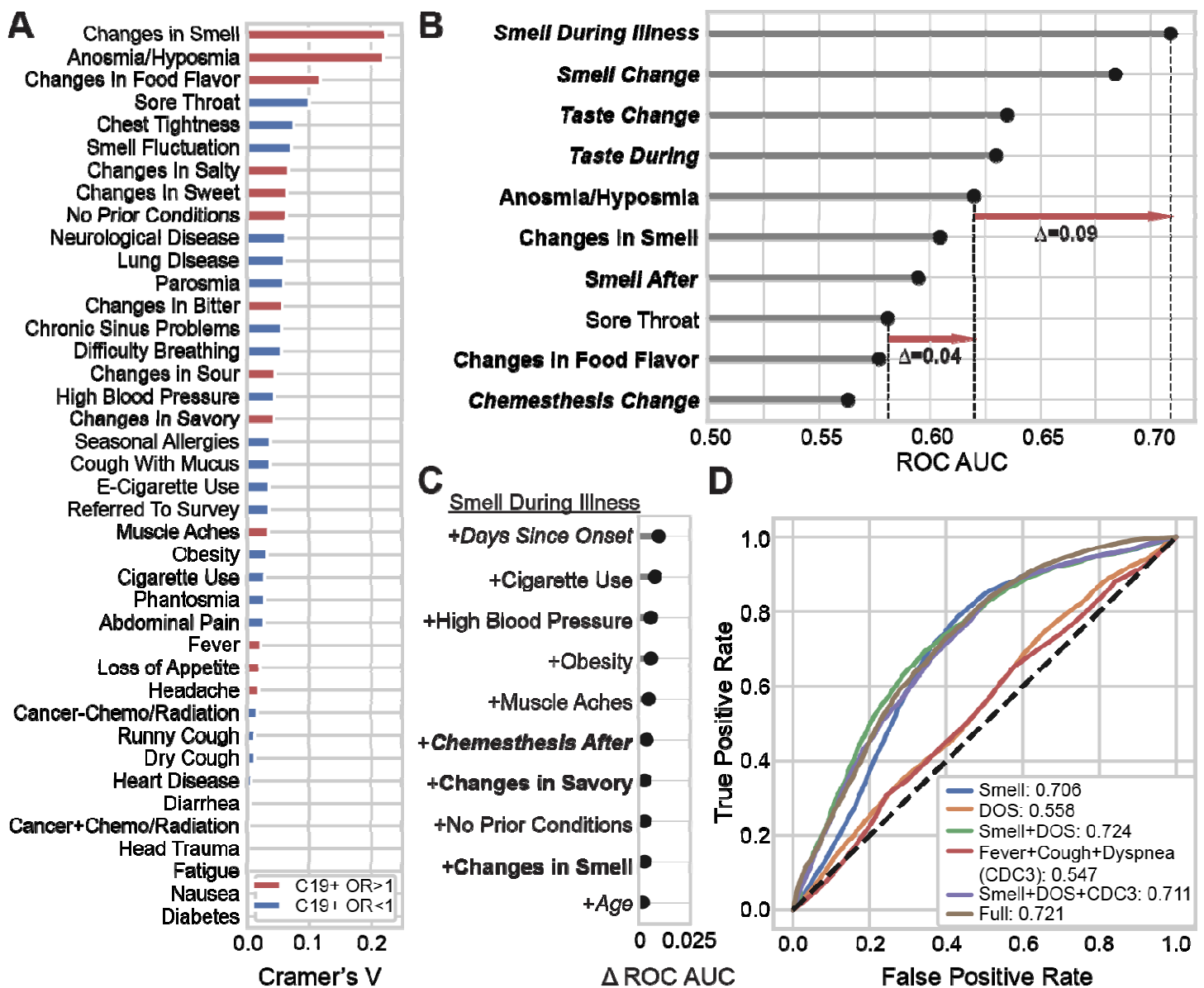
Binary (yes/no) and categorical reported symptoms suggested that COVID-19 is more strongly associated with chemosensory than with non-chemosensory symptoms, including fever, cough, and shortness of breath, cardinal symptoms currently highlighted by the US Centers for Disease Control and Prevention (CDC) (**Figure 3A**). Using AUC to assess prediction quality (**Figure 3B**), we found that self-reported smell ability during illness, reported on a continuous scale, was the most predictive survey question for COVID-19 status (AUC=0.71). Changes in smell as a result of illness (the difference between smell ability during and before illness) was similarly predictive (AUC=0.69). Changes in taste ability (assessed via rating) were the next most predictive features (AUC=0.64-0.65) (**Figure 3B**). Models fit to the same data but with shuffled COVID-19 status consistently produced AUC~0.5 for all features. The most predictive non-chemosensory symptom, sore throat (which was negatively associated with COVID-19) was substantially less predictive (AUC=0.58) than the top chemosensory symptoms. Nasal obstruction was not predictive (AUC=0.52). Responses given on a continuous scale were more predictive (AUC=0.71) than binary responses to parallel questions (e.g., **Appendix 1**, Question 13 versus 14, **Figure S5**) (AUC=0.60-0.62), likely because a continuous scale contains a greater amount of diagnostic information (**Figure S4**).

Next, we examined which simple multi-feature model would best predict COVID-19 status. As some questions have highly correlated responses, the question most complementary to “Smell during illness” is unlikely to be one that carries redundant information. Adding “Days since onset of respiratory symptoms” (DOS) to “Smell during illness” (Smell Only) produced the largest incremental gain in predictive performance (AUC=0.72, +0.01 versus the Smell Only model) (**Figure 3C**).

We directly compared the Smell Only+DOS model to other candidate models. The Smell Only+DOS model (**Figure 3D**) yielded an equal or higher AUC than the model including the three cardinal CDC symptoms (AUC=0.55) or the full model using 70 features (AUC=0.72). Because the Smell Only+DOS model exhibits the same AUC as the full model it strikes a good balance between model parsimony and predictive accuracy for C19+. However, the Smell Only model also offers reasonable sensitivity of 0.85 (at specificity=0.51, cutoff=13 on the 100-point VAS) and/or specificity of 0.75 (at sensitivity=0.51, cutoff=1) as desired. By sharp contrast, fever has a sensitivity of only 0.54 with specificity of 0.49 and dry cough has sensitivity of 0.52 and specificity of 0.46.



**Figure 2.** Chemosensory ability and nasal obstruction in C19+ and C19- participants. Self-reported smell (A,B), taste (C,D), chemesthesis (E,F), and nasal obstruction (G,H; formulated as “How blocked was your nose?”) before and during respiratory illness in C19+ (darker shades) and C19- (lighter shades) participants. Ratings were given on 0-100 visual analog scales. Left panels (A,C,E,G) show mean values. Right panels (B,D,F,H) show distributions of the change scores (during minus before). Thicker sections indicate relatively more subjects (higher density of responses). The thick black horizontal bar indicates the median, the shaded area within each violin indicates the interquartile range. Each dot represents the rating of a single participant.



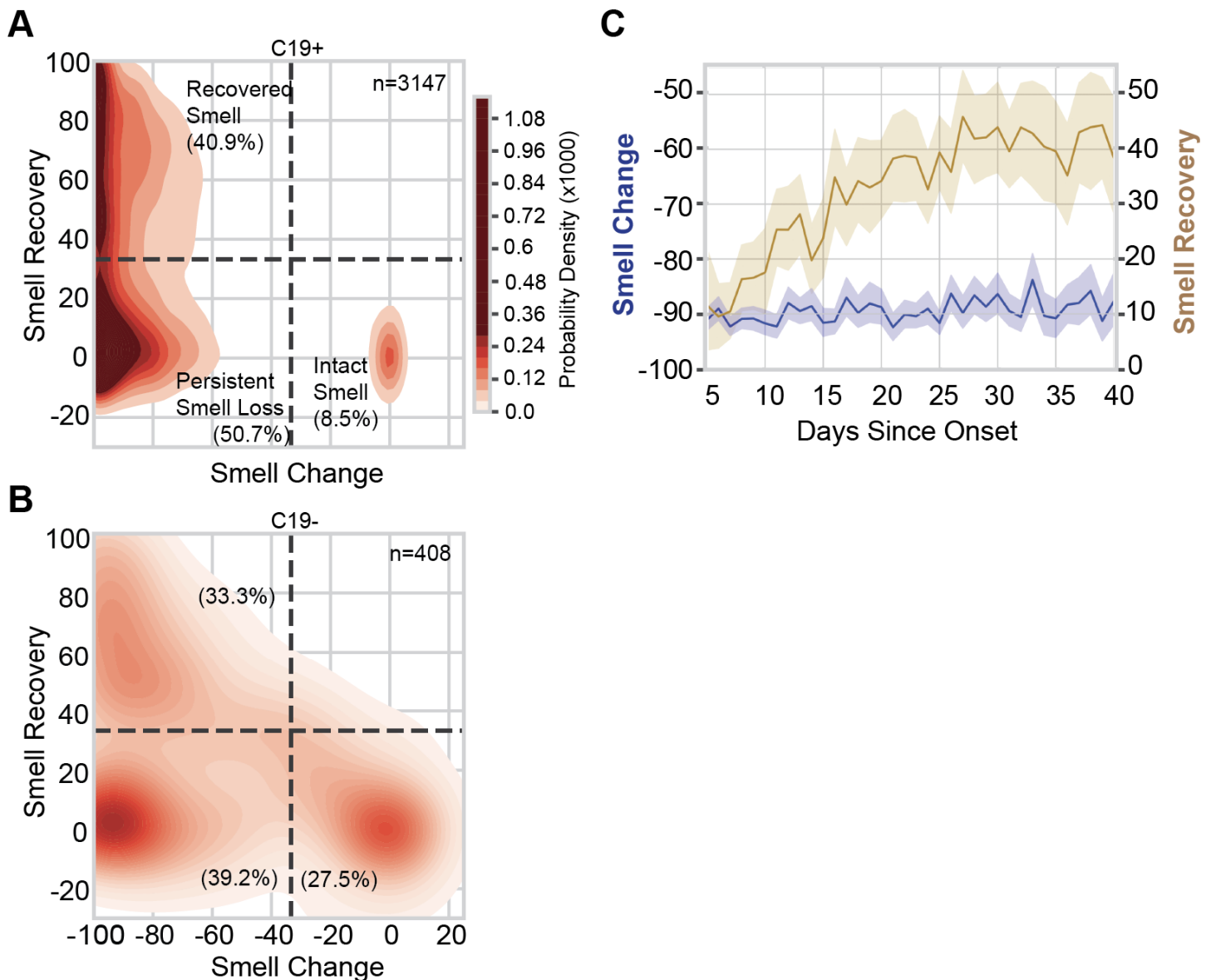
**Figure 3.** Smell loss is the strongest predictor of COVID-19 status. **(A)** A normalized measure of association (Cramer's V) between binary or categorical responses on COVID-19 status.  $V=0$  reflects no association between the response and COVID-19 status;  $V=1$  reflects a perfect association;  $V>0.1$  is considered a meaningful association. Features in red are positively associated with C19+ (odds ratio  $> 1$ ); features in blue are negatively associated with C19+ (odds ratio  $< 1$ ). **(B)** Logistic regression is used to predict COVID-19 status from individual features. Top-10 single features are ranked by performance (cross-validated area under the ROC curve, AUC). Chemosensory-related features (bold) show greater predictive accuracy than non-chemosensory features (non-bold). Responses provided on the numeric scale (italic) were more informative than binary responses (non-italic). Red arrows indicate differences in prediction quality (in AUC) between features. **(C)** Adding features to "Smell During Illness" results in little improvement to the model; only Days Since Onset of Respiratory Symptoms (DOS) yields meaningful improvement. **(D)** ROC curves for several models. A model using "Smell during illness" (Smell Only, abbreviated "Smell" in figure) is compared against models containing this feature along with DOS, as well as models including the three cardinal CDC features (fever, dry cough, difficulty breathing). "Full" indicates a regularized model fit using 70 dozen survey features, which achieves prediction accuracy similar to the parsimonious model "Smell Only+DOS".

## Recovery from smell loss

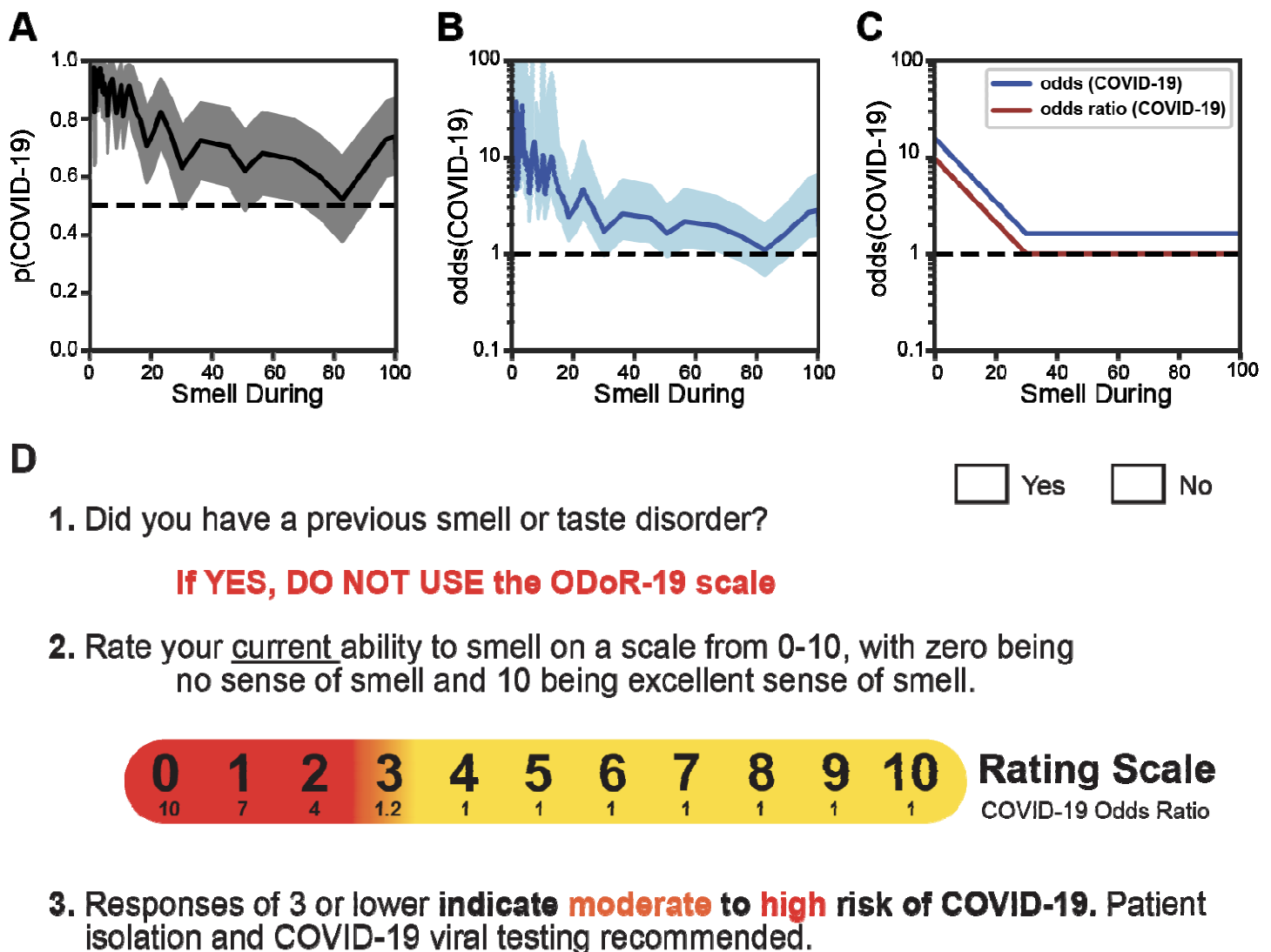
Recovery from smell loss was modest (approximately half the initial average loss) in C19+ participants with full or partial resolution of respiratory symptoms. Overall, self-reported, post-illness olfactory ability was still lower for C19+ ( $39.9 \pm 34.7$ ) than C19- ( $52.2 \pm 35.2$ ,  $p=2.8e-11$ , **Figure S6A**). However, the mean recovery of smell (after illness relative to during illness) was greater for C19+ ( $30.5 \pm 35.7$ ) than C19- ( $24.6 \pm 31.9$ ,  $p=0.0002$ , **Figure S6B**). A similar but smaller effect of COVID-19 status on recovery was observed for taste (**Figure S6C, D**), while little to no association with COVID-19 was observed for recovery of chemesthesis (**Figure S6E,F**) or nasal obstruction (**Figure S6G,H**). When illness-induced change in olfactory function (during minus before illness) and recovery of olfactory function (after minus during illness) were evaluated, we identified three respondent clusters: those self-reporting no loss of smell (Intact Smell), those reporting recovery from smell loss (Recovered Smell), and those reporting smell loss without recovery by up to 40 days (Persistent Smell Loss, **Figure 4, Table S3**). Intact smell was reported by only 8.5% of the participants in the C19+ group but by 27.5% in the C19- group ( $p=3.8e-31$ ). A greater proportion of C19+ participants were included in both the Recovered Smell group (C19+: 40.9%, C19-: 33.3%;  $p=4.9e-10$ ) and the Persistent Smell Loss group (C19+: 50.7%, C19-: 39.2%;  $p=5e-5$ ; **Figure 4A, B**). C19+ participants in both the Recovered Smell and Persistent Smell Loss clusters reported a similar extent of olfactory loss, irrespective of time since respiratory symptom onset. By contrast, the rate of self-reported smell recovery increased over time, with a plateau at 30 days (**Figure 4C**). Finally, DOS was the best predictor (AUC=0.62) between the Persistent Smell Loss and the Recovered Smell groups (**Figure S6A, Table S3**).

## Simple screening for COVID-19: the Olfactory Determination Rating scale in COVID-19 (ODoR-19) (136)

Our results indicate that a continuous rating of current olfactory function is the single best predictor of COVID-19 and improves the discrimination between C19+ and C19- over a binary question on smell loss. For example, the Smell Only model can reach a specificity of 0.83 at the low end of the VAS (sensitivity=0.36, cutoff=0). We propose here a numeric variation of the rating scale (0-10), the ODoR-19, that can be administered in person or via telemedicine to improve early COVID-19 screening for individuals without preexisting smell and/or taste disorders. Responses to the ODoR-19 scale  $\leq 2$  indicate high odds of COVID-19 positivity ( $4 < OR < 10$ , **Figure 5D**). An ODoR-19 response of 3 indicates a borderline risk ( $OR=1.2$ ).



**Figure 4:** Smell loss, recovery, and time course. **(A, B)** Joint distribution of smell loss (during minus before illness ratings) and smell recovery (after minus during illness ratings) for C19+ (A) and C19- (B) participants. Darker color indicates a higher probability density; the color map is shared between (A) and (B); dashed lines are placed at a third of the way across the rating scale to aid visualization of the clusters. Severe smell loss that is either persistent (lower left) or recovered (upper left) was more common in C19+ than C19-. *n* indicates the number of participants in each panel. % indicates the percentage of participants of the given COVID status in each quadrant. **(C)** In C19+ participants who lost their sense of smell (Recovered Smell + Persistent Smell Loss), the degree of smell recovery (right y axis) increased over ~30 days since onset of respiratory symptoms before plateauing; the degree of reported smell change (left y axis) did not vary in that window of observation. Solid lines indicate the mean of the measure, the shaded region indicates the 95% confidence interval.



**Figure 5.** The odds of a COVID-19 diagnosis as a function of olfactory loss. **(A)** The solid line indicates the probability of a COVID-19 diagnosis as a function of “Smell during illness” ratings. The shaded region indicates the 95% confidence interval. **(B)** The solid line expresses the probability of a COVID-19+ diagnosis as a function of “Smell during illness” in odds ( $p/(1-p)$ ); it is shown on a logarithmic scale. The shaded region indicates the 95% confidence interval. **(C)** Stylized depiction of change in the odds of a COVID-19 diagnosis and of the odds ratio. **(D)** The ODoR-19 screening tool. After healthcare providers or contact tracers have excluded previous smell and/or taste disorders such as those resulting from head trauma, chronic rhinosinusitis, or previous viral illness, the patient can be asked to rate their current ability to smell on a scale from 0-10, with 0 being no sense of smell and 10 being excellent sense of smell. If the patient reports a value below or equal to 3, there is a high (red) or moderate (orange) probability that the patient has COVID-19. Values in yellow (ratings above 3) cannot rule out COVID-19.

## Discussion

Self-reported smell loss was more common in C19+ than C19- participants, but present in both groups. The use of a VAS to assess olfactory loss better predicted COVID-19 status than using a binary question. We found that the best predictor of COVID-19-associated smell recovery, within the time frame captured by the survey (~40 days), was days since onset of COVID-19.

The SARS-CoV-2 pandemic requires healthcare providers and contact tracers to quickly and reliably assess an individual's COVID-19 risk, often remotely. Thus, reliable screening tools are critical to assess a person's likelihood of having COVID-19 and to justify self-quarantine and/or testing recommendations. Indeed, some reports suggest that COVID-19-associated smell loss might be an indicator of disease severity.<sup>2,20</sup> Current symptom criteria (e.g., fever, dry cough) are less specific than severe olfactory loss. Indeed, the value of our ODoR-19 tool lies in the high specificity of values  $\leq 2$  for indicating COVID-19 positivity, therefore representing a valuable addition to the current repertoire of COVID-19 screening tools. Those who receive a negative outcome from a COVID-19 viral test, yet report significant idiopathic smell loss, should be considered as high-priority candidates for COVID-19 re-testing.

Our online survey and sampling methodology likely selected participants with a heightened interest in smell and taste and/or their disturbances. This self-selection bias could be viewed as a limitation since the C19- group also showed chemosensory loss. However, finding difference between groups in a sample with a higher barrier for discriminating between C19+ and C19- supports the robustness of this tool when used in a typical clinical population; our collider bias analysis also suggests that our findings are likely conservative estimates (**Figure S1, Table S1**).

Our results suggest that chemosensory impairment has strong COVID-19 predictive value and is useful when access to viral testing is limited or absent. As with any self-report measure, veracity of self-reports cannot be guaranteed. However, the ability to screen individuals in real-time should outweigh this potential confound.<sup>21</sup> While objective smell tests are the gold standard for assessing olfactory function,<sup>22,23</sup> they are costly, time-consuming to administer, and can require in-person interactions with potentially infectious patients.<sup>23,24</sup> By contrast, the ODoR-19 is free, quick, and can be administered in person or remotely. We cannot exclude that our C19- sample contains COVID-19 false negatives.<sup>25</sup> However, self-reported smell during illness distinguishes between C19+ and C19-, but not between randomly shuffled cases, suggesting that the difference between C19+ and C19-, even in a sample with over-represented chemosensory dysfunction, is substantial and can be captured via self-report.

Approximately half of the participants in the C19+ group recovered their sense of smell within 40 days from the onset of respiratory symptoms. This suggests the presence of at least two subgroups of patients: one that recovers quickly (<40 days, 40.9%) and another that may present a more variable time course of recovery (50.7%). Since these data are collected before the full recovery of all symptoms, we cannot offer a complete picture of recovery from olfactory loss in COVID-19-positive individuals, but they align with other early reports.<sup>26</sup> The COVID-19 pandemic will greatly increase the number of patients suffering from anosmia and other chemosensory disorders,<sup>27</sup> conditions that significantly affect quality-of-life,<sup>28,29</sup> dietary behavior,<sup>30</sup> cardiovascular health,<sup>31</sup> and mental health.<sup>32,33</sup> Thus, it is necessary to prepare healthcare providers to address the long-term needs of these patients.

Based on our results, we propose the use of the ODoR-19 tool, a quick, free, and effective smell-based screening method for COVID-19. This 0-10 rating scale accurately predicts COVID-19 in individuals without pre-existing smell and taste disorders (e.g., from head trauma, chronic rhinosinusitis<sup>34</sup>). ODoR-19 combines the utility of a continuous scale with the ease and speed needed for a screening tool. ODoR-19 is safe for remote administration during an illness with high viral spread and can precede and complement viral testing. This tool will improve screening for patients with limited or no access to medical care around the globe.

## Acknowledgements

The authors wish to thank all study participants, patients, and patient advocates that have contributed to this project, including members of the AbScent Facebook group. The authors wish to thank Micaela Hayes, MD for her input on the clinical relevance of this project, Shannon Alshouse and Olivia Christman for their help in implementing the survey, Sara Lipson for her support, and the international online survey research firm YouGov for providing data gathered with the Imperial College London YouGov Covid 19 Behaviour Tracker.

## Funding

Deployment of the GCCR survey was supported by an unrestricted gift from James and Helen Zallie to support sensory science research at Penn State. Richard C. Gerkin is supported by NINDS (U19NS112953) and NIDCD (R01DC018455). Paule V. Joseph is supported by the National Institute of Nursing Research under award number 1ZIANR000035-01. PVJ is also supported by the Office of Workforce Diversity, National Institutes of Health and the Rockefeller University Heilbrunn Nurse Scholar Award. Vera V. Voznessenskaya is supported by IEE RAS basic project 0109-2018-0079. Mackenzie Hannum is supported by NIH T32 funding (DC000014). Masha Niv is supported by Israel Science Foundation grant #1129/19.

## Conflict of interest

Richard C. Gerkin is an advisor for Climax Foods, Equity Compensation (RCG); John E. Hayes has consulted for for-profit food/consumer product corporations in the last 3 years on projects wholly unrelated to this study; also, he is Director of the Sensory Evaluation Center at Penn State, which routinely conducts product tests for industrial clients to facilitate experiential learning for students. Since 2018 Thomas Hummel collaborates with and received funding from Sony, Stuttgart, Germany; Smell and Taste Lab, Geneva, Switzerland; Takasago, Paris, France: aspuraclip, Berlin, Germany. Christine E. Kelly is the founder of AbScent, a charity registered in England and Wales, No. 1183468. Christophe Laudamiel has received fundings from scent related institutions and corporations, however for work totally unrelated to the field of the present study.

## References

1. Coronavirus. World Health Organ. 2020 (<https://www.who.int/westernpacific/health-topics/coronavirus>). ([https://www.nlm.nih.gov/dr2/COVID-19\\_BSSR\\_Research\\_Tools.pdf](https://www.nlm.nih.gov/dr2/COVID-19_BSSR_Research_Tools.pdf))
2. Paderno A, Schreiber A, Grammatica A, et al. Smell and taste alterations in Covid-19: a cross-sectional analysis of different cohorts. *Int Forum Allergy Rhinol* 2020;(Epub ahead of print).
3. Giacomelli A, Pezzati L, Conti F, et al. Self-reported Olfactory and Taste Disorders in Patients With Severe Acute Respiratory Coronavirus 2 Infection: A Cross-sectional Study. *Clin Infect Dis* ciaa330.
4. Yan CH, Faraji F, Prajapati DP, et al. Association of chemosensory dysfunction and COVID-19 in patients presenting with influenza-like symptoms. *Int Forum Allergy Rhinol* 2020;10(7):806–13.
5. Lechien JR, Chiesa-Estomba CM, De Siaty DR, et al. Olfactory and gustatory dysfunctions as a clinical presentation of mild-to-moderate forms of the coronavirus disease (COVID-19): a multicenter European study. *Eur Arch Otorhinolaryngol* 2020;277:2251–61.
6. Moein ST, Hashemian SMR, Mansourafshar B, et al. Smell dysfunction: a biomarker for COVID-19. *Int Forum Allergy Rhinol* 2020;(Epub ahead of print).
7. Hornuss D, Lange B, Schröter N, et al. Anosmia in COVID-19 patients. *Clin Microbiol Infect* 2020;(Epub ahead of print).
8. Parma V, Ohla K, Veldhuizen MG, et al. More than smell – COVID-19 is associated with severe impairment of smell, taste, and chemesthesis. *Chem Senses* 2020;(Epub ahead of print).



9. Menni C, Valdes AM, Freidin MB, et al. Real-time tracking of self-reported symptoms to predict potential COVID-19. *Nat Med* 2020;26:1037–1040.
10. Menni C, Sudre CH, Steves CJ, et al. Quantifying additional COVID-19 symptoms will save lives. *The Lancet* 2020;395(10241):e107–e108.
11. Kim G -u, Kim M-J, Ra SH, et al. Clinical characteristics of asymptomatic and symptomatic patients with mild COVID-19. *Clin Microbiol Infect* 2020;26(7):948.e1-948.e3.
12. Walsh-Messinger J, Kaouk S, Manis H, et al. Standardized Testing Demonstrates Altered Odor Detection Sensitivity and Hedonics in Asymptomatic College Students as SARS-CoV-2 Emerged Locally. June 1, 2020 (<https://www.medrxiv.org/content/10.1101/2020.06.17.20106302v1>). preprint.
13. Hopkins C, Surda P, Kumar N. Presentation of new onset anosmia during the COVID-19 pandemic. *Rhinology* 2020;50:1–4.
14. Soler ZM, Patel ZM, Turner JH, et al. A primer on viral-associated olfactory loss in the era of COVID-19. *Int Forum Allergy Rhinol* 2020;10(7):814–20.
15. Parma V, Veldhuizen M, Ohla K, et al. Is olfactory loss a sensitive symptomatic predictor of COVID-19? A preregistered, crowdsourced study. 2020 (<https://osf.io/gxu7e>).
16. Kraemer JD, Strasser AA, Lindblom EN, et al. Crowdsourced data collection for public health: A comparison with nationally representative, population tobacco use data. *Prev Med* 2017;102:93–9.
17. Reback J, McKinney W, Jbrockmendel, et al. pandas-dev/pandas: Pandas 1.0.5. Zenodo; 2020 [cited 2020 Jul 10].
18. Pedregosa F, Varoquaux G, Gramfort A, et al. Scikit-learn: Machine Learning in Python. *J Mach Learn Res* 2011;12(85):2825–30.
19. Seabold S, Perktold J. Statsmodels: Econometric and statistical modeling with python. In: *Proceedings of the 9th Python in Science Conference*. Austin, TX; 2010. p. 61.
20. Yan CH, Faraji F, Prajapati DP, et al. Self-reported olfactory loss associates with outpatient clinical course in COVID-19. *Int Forum Allergy Rhinol* 2020;10(7):821–31.
21. Mermelstein R, Hedeker D, Flay B, et al. *The Science of Real-Time Data Capture: Self-Reports in Health Research*. 2007.
22. Hummel T, Sekinger B, Wolf SR, et al. ‘Sniffin’ Sticks’: Olfactory Performance Assessed by the Combined Testing of Odor Identification, Odor Discrimination and Olfactory Threshold. *Chem Senses* 1997;22(1):39–52.
23. Doty RL, Shaman P, Kimmelman CP, et al. University of pennsylvania smell identification test: A rapid quantitative olfactory function test for the clinic. *The Laryngoscope* 1984;94(2):176–8.
24. Oleszkiewicz A, Schriever VA, Croy I, et al. Updated Sniffin’ Sticks normative data based on an extended sample of 9139 subjects. *Eur Arch Otorhinolaryngol* 2019;276(3):719–28.
25. Kucirka LM, Lauer SA, Laeyendecker O, et al. Variation in False-Negative Rate of Reverse Transcriptase Polymerase Chain Reaction–Based SARS-CoV-2 Tests by Time Since Exposure. *Ann Intern Med* 2020;(Epub ahead of print).
26. Chiesa Estomba CM, Lechien JR, Radulesco T, et al. Patterns of smell recovery in 751 patients affected by the COVID-19 outbreak. *Eur J Neurol* (Epub ahead of print).
27. Rawal S, Hoffman HJ, Bainbridge KE, et al. Prevalence and Risk Factors of Self-Reported Smell and Taste Alterations: Results from the 2011–2012 US National Health and Nutrition Examination Survey (NHANES). *Chem Senses* 2016;41(1):69–76.
28. Smeets MAM, Veldhuizen MG, Galle S, et al. Sense of smell disorder and health-related quality of life. *Rehabil Psychol* 2009;54(4):404–12.
29. Croy I, Nordin S, Hummel T. Olfactory Disorders and Quality of Life--An Updated Review. *Chem Senses* 2014;39(3):185–94.
30. Kershaw JC, Mattes RD. Nutrition and taste and smell dysfunction. *World J Otorhinolaryngol - Head Neck Surg* 2018;4(1):3–10.
31. Gallo S, Byham-Gray L, Duffy VB, et al. Associations of olfactory dysfunction with anthropometric and cardiometabolic measures: Findings from the 2013–2014 national health and nutrition examination survey (NHANES). *Physiol Behav* 2020;215:112702.
32. Croy I, Symmank A, Schellong J, et al. Olfaction as a marker for depression in humans. *J Affect Disord* 2014;160:80–6.
33. Malaty J, Malaty IAC. Smell and taste disorders in primary care. *Am Fam Physician* 2013;88(12):852–9.
34. Hummel T, Landis BN, Hüttenbrink K-B. Smell and taste disorders. *GMS Curr Top Otorhinolaryngol Head*

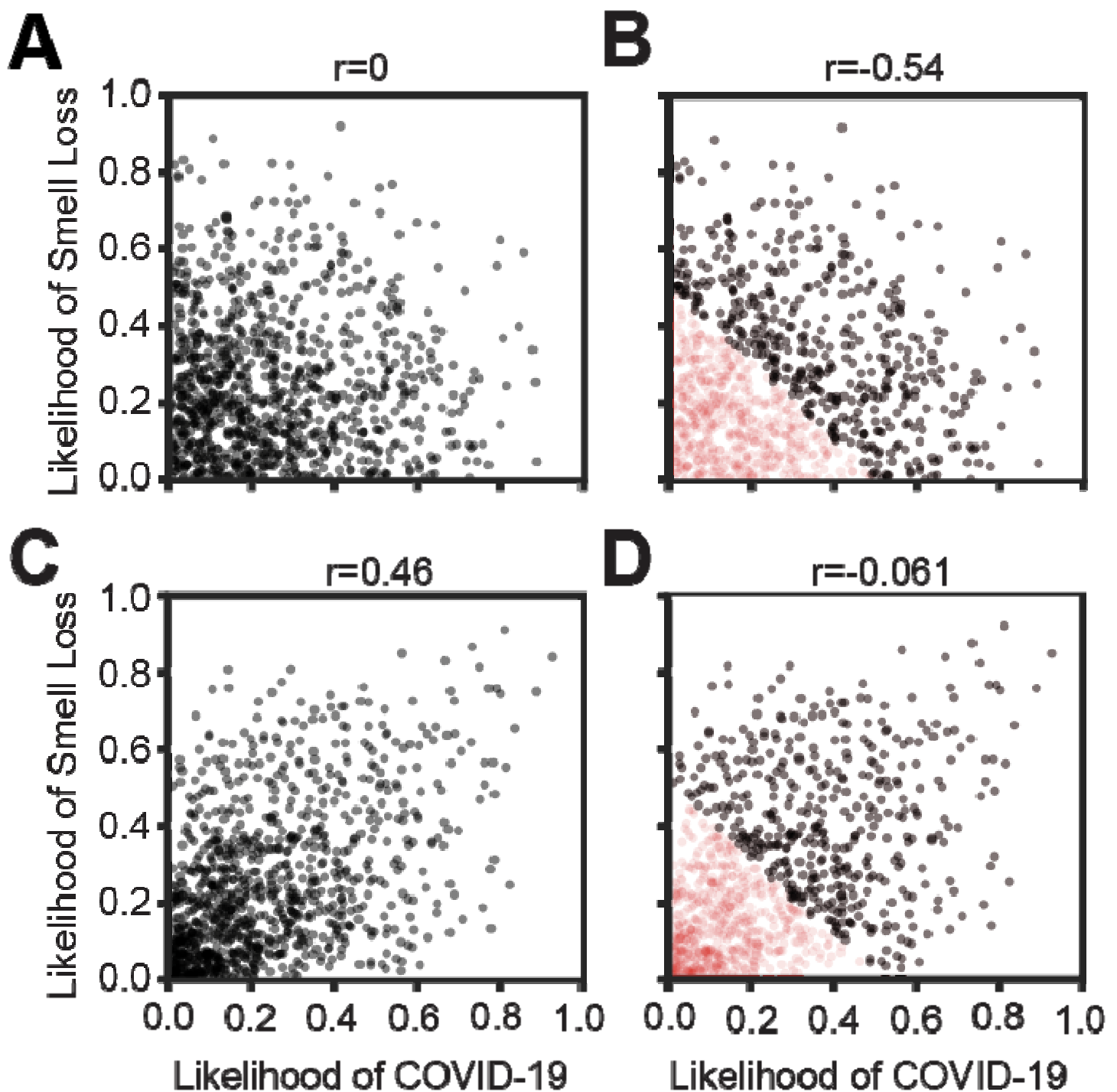
Neck Surg 2011;10:Doc04.

35. Griffith G, Morris TT, Tudball M, et al. Collider bias undermines our understanding of COVID-19 disease risk and severity. June 20, 2020 (<https://www.medrxiv.org/content/10.1101/2020.05.04.20090506v3>). preprint.
36. YouGov-Data/covid-19-tracker. YouGov Data; 2020(<https://github.com/YouGov-Data/covid-19-tracker>)
37. Lee MD, Wagenmakers E-J. Bayesian Cognitive Modeling: A Practical Course. Cambridge: Cambridge University Press; 2013 [cited 2020 Jul 19].
38. Jeffreys H. The Theory of Probability. OUP Oxford; 1998.
39. COVID-19 BSSR Research Tool. 2020 ([https://www.nlm.nih.gov/dr2/COVID-19\\_BSSR\\_Research\\_Tools.pdf](https://www.nlm.nih.gov/dr2/COVID-19_BSSR_Research_Tools.pdf))

## Supplementary Material

### Collider bias results in underestimation of an association between smell loss and COVID-19

As others have noted,<sup>35</sup> collider bias, resulting from selection or conditioning on variables involved in the analysis, may result in the distorted association between COVID-19 and candidate symptoms or patient attributes. In the present sample, it is likely that we have selected for both a higher probability of COVID-19 and a higher probability of smell and taste disorders than the population at large. However, rather than leading to an overestimation of the positive correlation between smell loss and COVID-19, collider bias is expected to lead to an underestimation of this correlation (**Figure S1**). If we consider the hypothetical scenario in which there is no association between smell loss and COVID-19 status in the general population, we would expect a distribution similar to that depicted in **Figure S1A**, where the correlation between the likelihood of smell change and likelihood of COVID-19 is  $r = 0$ . Based on our recruitment method, we expect that the participants who elected to complete the GCCR core questionnaire were likely to have COVID-19, smell loss, or both. We can simulate participant selection to reflect this hypothesis by censoring subjects which do not meet a fixed sum of smell loss and COVID-19 probabilities (i.e., the red dots are excluded from the calculation of the correlation; **Figure S1B**). As a result, the estimated correlation between smell loss and COVID-19 status originating from a population with  $r = 0$  would be negative (**Figure S1B**). A similar scenario would manifest if the association between smell loss and COVID-19 status in the general population is positive (**Figure S1C**). Again, simulating the removal of participants with low likelihood of having COVID-19 and/or reporting smell loss would result in a bias of the estimated correlation towards more negative values (**Figure S1D**). This collider bias indicates that the positive correlation between smell loss and C19+ is underestimated in the present sample. Indeed, a direct comparison of the binary (y/n) smell loss questions in the two empirical samples yields an C19 odds ratio of 5.96 in the YouGov sample (**Table S1**) but only 4.89 for GCCR. Therefore, our analyses represent a conservative scenario for the prediction of C19+ and C19- based on chemosensory alterations.



**Figure S1.** Collider bias leads to underestimation of the positive correlation between smell loss and COVID-19-positive status. **(A)** Hypothetical scenario depicting no relationship between smell change and likelihood of COVID-19 positive status. Black dots indicate individual potential subjects, each of whom has a latent likelihood of COVID-19 and of smell loss. **(B)** Hypothetical scenario depicting the emergence of a negative correlation between smell change and likelihood of COVID-19 positive status following a baseline lack of correlation, if participants with greater smell loss and/or COVID-19 positive are preferentially included in the sample. Red dots indicate subjects not observed due to this selection bias; subjects observed remain in black. **(C)** Hypothetical scenario depicting a positive relationship between smell change and likelihood of COVID-19 positive status. **(D)** Hypothetical scenario depicting the emergence of a negative correlation between smell change and likelihood of COVID-19 positive status following a positive baseline correlation, if participants with greater smell loss and/or COVID-19 positive are preferentially included in the sample.

**Table S1.** Comparison with the representative YouGov database shows that the GCCR sample underestimates the positive correlation between smell loss and COVID-19 positive status.

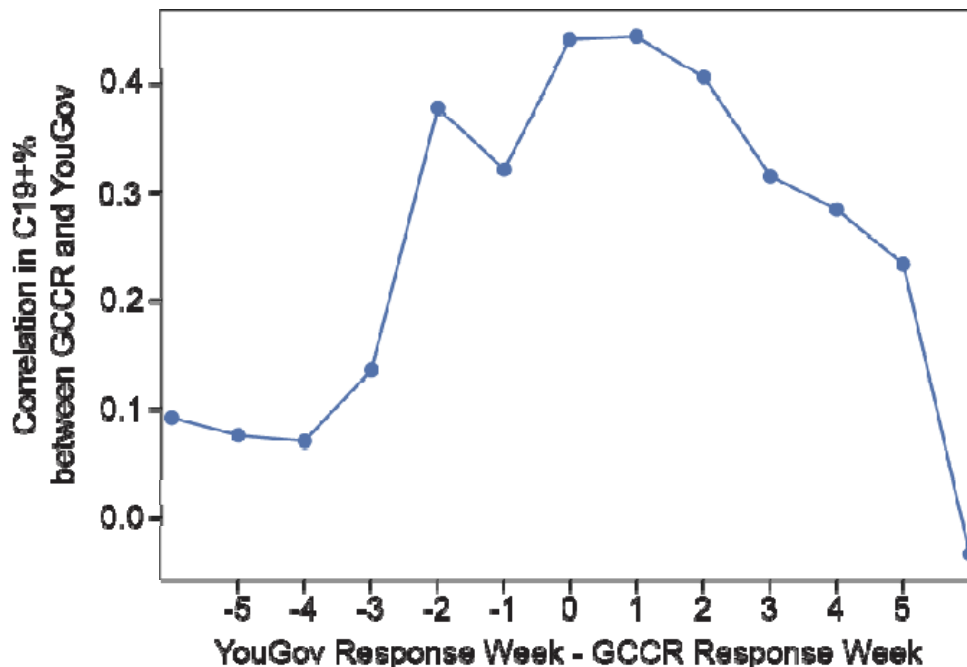
	N	% C19+	% C19-	OR (C19+ vs C19-)	OR (C19+ vs Not Tested)	OR (C19- vs Not Tested)	p(Smell Loss   C19+)
<b>Global</b>	109395	0.51	3	5.96	49.7	8.33	0.33
<b>Brazil</b>	4167	0.91	5.4	8.39	33.9	4.04	0.47
<b>Canada</b>	5524	0.47	3.4	7.76	71.6	9.22	0.35
<b>Denmark</b>	5839	0.36	5.7	9.35	77.8	8.32	0.33
<b>Finland</b>	5927	0.32	1.9	1.85	27.8	15.1	0.21
<b>France</b>	9820	0.46	1.7	7	57.9	8.27	0.29
<b>Germany</b>	9468	0.49	2.1	4.56	94.2	20.7	0.37
<b>Italy</b>	9790	0.33	2.8	9.17	50.8	5.54	0.22
<b>Mexico</b>	5840	0.24	3.5	8.91	27.6	3.1	0.21
<b>Netherlands</b>	3822	1.2	2.7	2.73	16.1	5.92	0.3
<b>Norway</b>	5794	0.86	4.4	5.82	51.2	8.8	0.32
<b>Spain</b>	9789	0.37	3.5	5.81	43.5	7.48	0.33
<b>Sweden</b>	9741	0.44	2	3.69	38	10.3	0.35
<b>United Kingdom</b>	13565	0.15	1.2	23.1	108	4.66	0.5
<b>United States</b>	10309	1.2	4.8	4.61	54.7	11.9	0.34

Column 1 includes the list of countries available in the YouGov database. Column 2 indicates the number of participants over the history of their survey up to July 3, 2020. Minimum N=1000 per time point. Column 3 reports the percentage of participants who reported C19+. Column 4 reports the percentage of participants who reported C19-. Columns 5-7 report the odds ratios using Smell Loss for either C19+ vs. C19- (Column 5, and as reported in Figure 5 based on the data of the GCCR survey), for C19+ vs. untested individuals (Column 6) and C19- vs. untested individuals (Column 7). Untested individuals in YouGov's survey are those who did not report to be sick but were contacted as representative participants of a country. Column 8 reports the probability of smell loss in the C19+ group. The first row indicates the Global average (across countries) weighted by sample size. The global odds ratio for Smell loss calculated from a binary question for the group C19+ vs. C19- is 6.72, which is greater than what we identify in the GCCR survey (OR for changes in smell (binary question) = 4.89). The OR for C19+ vs. untested individuals is 58 and lowers to 11 for C19-. This confirms that smell loss is also associated with other etiologies, but is not nearly as prevalent as in participants with C19+

## How representative is the GCCR sample?

As with most COVID-19 studies,<sup>19</sup> the sample studied here is not representative of the general population. To better understand the extent to which this is the case, we computed a cross-correlation between GCCR and YouGov data.<sup>36</sup> These data were aligned by weighting YouGov samples to achieve an identical survey date distribution to the GCCR samples. Specifically, GCCR survey dates were converted to a YouGov “week number” because YouGov surveys only weekly. The distribution of week numbers was computed for each country in the GCCR data. The YouGov data for the same country was then weighted by week number to match the corresponding GCCR distribution for that country. So, for example, if a country had 10 GCCR survey responses in week 1 of the YouGov survey period, and 30 in week 2 of that period, then the YouGov data in week 1 would be weighted at 25% and in the YouGov data in week 2 at 75%. This procedure was applied independently for each country, and the weights were used to compute a weighted mean COVID-19-positive rate for each country from the YouGov data. This was then directly compared against the raw COVID-19-

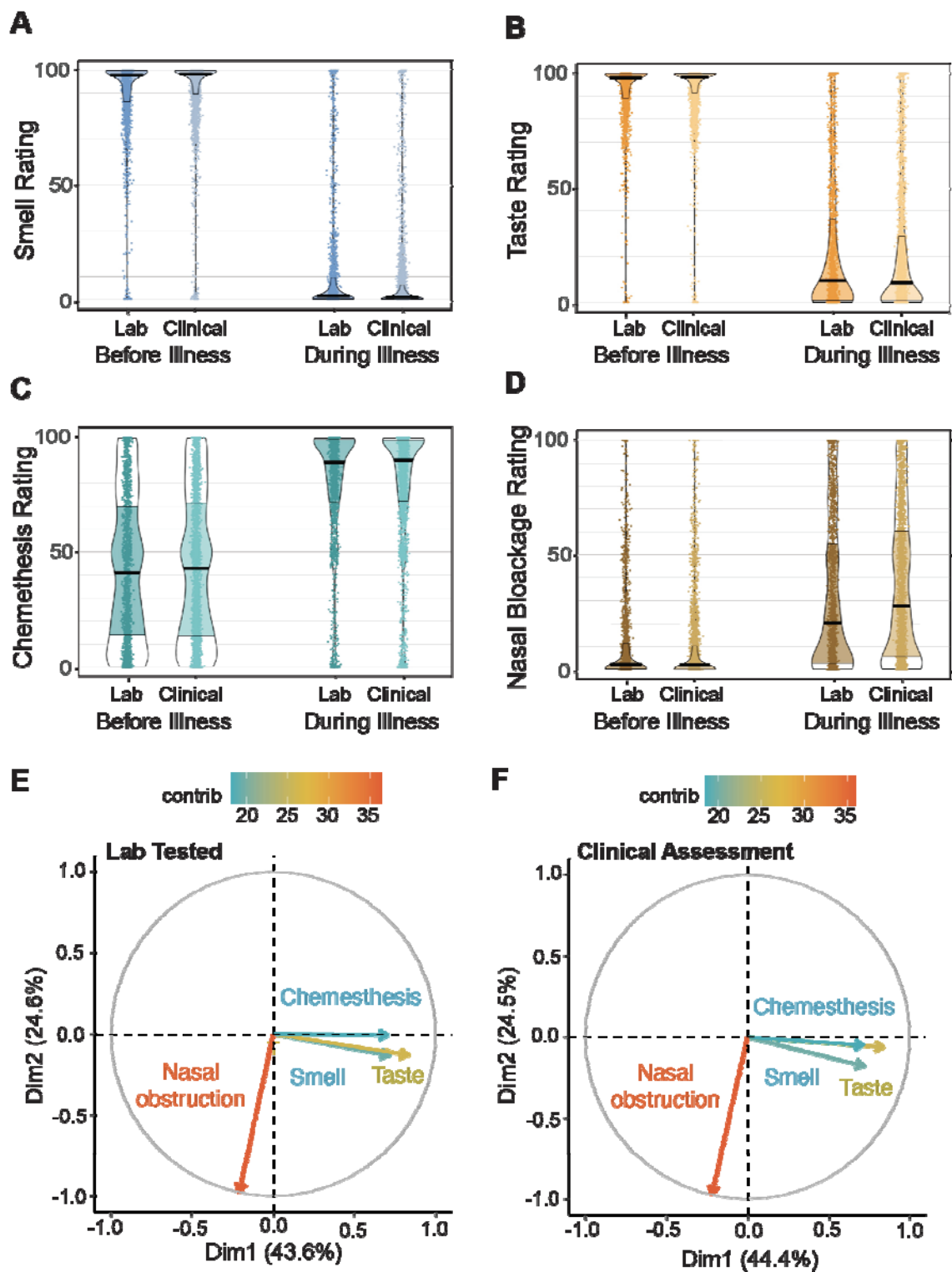
positive rate for each country in the GCCR data. A lag (x-axis value in **Figure S2**) of 0 exactly reflects the above description. Other values of the lag indicate that the alignment was shifted: for example, a lag of one week means that the hypothetical GCCR responses above would be weighted 25/75 towards weeks 2 and 3, instead of weeks 1 and 2. Under the hypothesis that the COVID-19-positive rates in the two surveys are related, but may have different temporal dynamics, changing the lag allows these dynamics to be estimated. **Figure S2** depicts the country-wise correlation in participants with a positive COVID-19 test results (C19+) fraction between the two datasets, as a function of the lag between GCCR survey date and YouGov survey date. The country-wise C19+ fraction is correlated ( $r \sim 0.45$ ) when responses from the same calendar week are aligned, but diminishes outside of that window, showing both surveys capture a similar within-country temporal component of the epidemic.



**Figure S2.** COVID-19 status in the GCCR cohort is correlated with a representative YouGov sample.

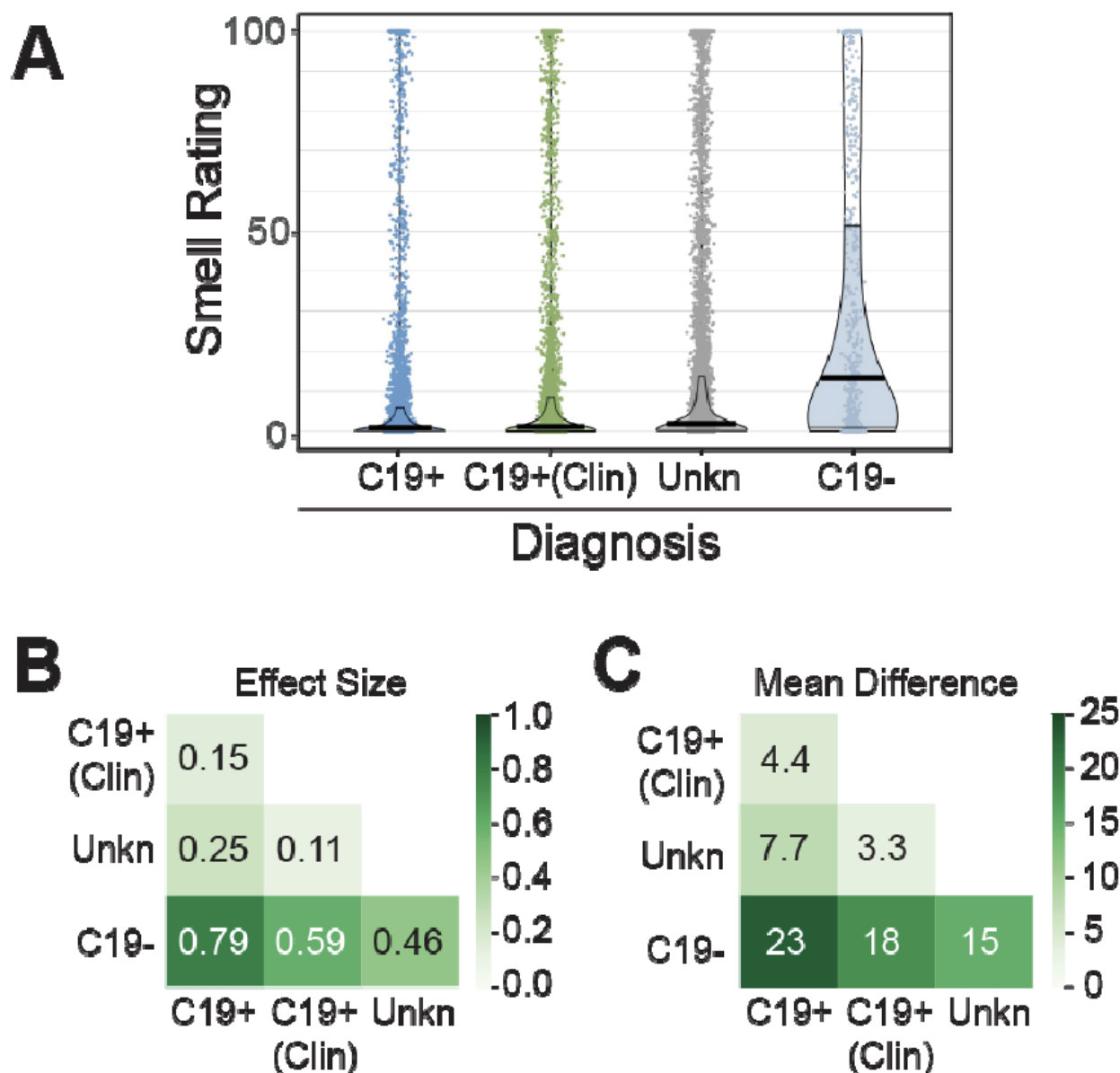
## Sample description

Based on responses to question 7 of the GCCR survey (“Have you been diagnosed with COVID-19?”, **Appendix 1**), participants can be split into six groups (see **Figure 1**). Participants who responded with Option 2 (“Yes – diagnosed with viral swab”) or 3 (“Yes – diagnosed with another lab test”) were classified as C19+; participants who responded with option 5 (“No – I had a negative test, but I have symptoms”) were classified as symptomatic C19-; participants who responded with option 4 (“No – I was not diagnosed, but I have symptoms”) were classified as C19 Unknown; participants who responded with option 6 (“No – I do not have any symptoms”), with option 7 (“Don’t know”), or with option 8 (“Other”) were classified as undefinable and excluded from the final analyses. To replicate our previous findings,<sup>8</sup> we first compared individuals newly included in the GCCR dataset (responses from 14 May to 2 July, 2020, replication sample in **Figure 1**) with COVID-19 who were lab tested and those who were diagnosed by a clinician based on the self-reported quantitative changes in smell, taste, chemesthesis, and nasal obstruction (**Figure S3**). Participants with lab-test confirmed C19+ did show slightly greater chemosensory deficits than did those diagnosed with C19+ clinically, but the difference was not clinically meaningful (smell:  $4.4 \pm 28.6$ ,  $p=2.7e-13$ ) (**Figure S3, Table S2**). We then focused our descriptive and predictive analyses of participants who received a positive (C19+) or a negative (C19-) lab test for COVID-19. We also computed descriptive and predictive analysis for the C19+ subsample who reported partial or full signs of recovery from their recent respiratory illness. Lastly, the unknown group was originally hypothesized as similar to the C19- group. Yet the ratings of smell ability during illness suggest that the majority of these participants has a smell profile closer to C19+ than C19- (**Figure S4**). To maximize the validity of the COVID-19 diagnosis in our sample, we therefore excluded the C19 Unknown group from further analyses.



**Figure S3.** This figure describes a pre-registered replication of Parma et al, 2020 and includes only new data collected between May 14th and July 3rd 2020 via the GCCR survey. (A-D) Changes in smell (A), taste (B), chemesthesis (C) and nasal blockage (D) during versus before in COVID-19-positive individuals (Groups 1, 2 and 3, see Figure 1). All subjects had a COVID-19-positive status either via lab test (darker shades) or via clinical assessment (lighter shades). (E-F) Principal component analysis shows that smell, taste, and chemesthesis changes in both the lab test (E) and clinical assessment (F) groups) were orthogonal to blocked nose changes, i.e., the three chemosensory changes were highly correlated across subjects whereas blocked nose changes were mostly uncorrelated.





**Figure S4.** (A) Self-reported smell change and comparison of smell change between four diagnosis groups: Positive COVID-19 lab-test (C19+), positive COVID-19 clinical assessment (C19+ (Clin)), COVID-19 Unknown (Unkn; lack of clinical and lab test diagnosis, but reported symptoms), and negative COVID-19 lab test (C19-). Solid horizontal lines reflect the median; dashed lines reflect the quartiles. (B, C) Differences between groups, in terms of (B) effect size (Cohen's D) and (C) means (on a 0-100 scale).

## Replication of previous analyses

The replication of Parma et al.<sup>8</sup> used the same Bayesian linear regression approach with Cauchy prior [ $r = \sqrt{2}/2$ ]. This approach is appropriate for estimating the strength of the evidence in support of the alternative hypothesis: the clinical assessment and the lab test C19+ groups show similar smell, taste, chemesthesis and nasal obstruction changes before vs. during the illness. The interpretation of the Bayes factors BF follows the classification scheme proposed by Lee and Wagenmakers<sup>37</sup> and adjusted from Jeffreys<sup>38</sup>, which considers BF

> 3 as moderate evidence, BF > 10 as strong evidence, BF > 30 as very strong evidence and BF > 100 as extreme evidence for H<sub>0</sub> or H<sub>1</sub>.

**Table S2.** Differences between lab-tested and clinically-assessed COVID-19-positive participants on changes in smell, taste, chemesthesis and nasal blockage.

	Smell Change	Taste Change	Chemesthesis Change	Change in Nasal Blockage
$\Delta$	-4.4	-3.4	0.37	3.9
$\sigma$	0	0	37	33
se $\Delta$	-0.048	-0.037	0.0041	0.043
$D$	-0.15	-0.1	0.01	0.12
$p$	2.70E-13	2.00E-06	0.38	4.00E-09

Change means the rating "before" illness minus the rating "during" illness on the 0-100 visual-analog scale.  $\Delta$  indicates the mean difference in change between lab-test and clinically-assessed COVID-19-positive subjects, while  $\sigma$  indicates the standard deviation.  $D$  indicates effect size (Cohen's D).  $p$  indicates p-value from a Mann-Whitney U-test. In contrast to the prediction of the pre-registration, we found statistically significant differences between groups. However, the effect sizes are small and thus unlikely to be of practical importance.

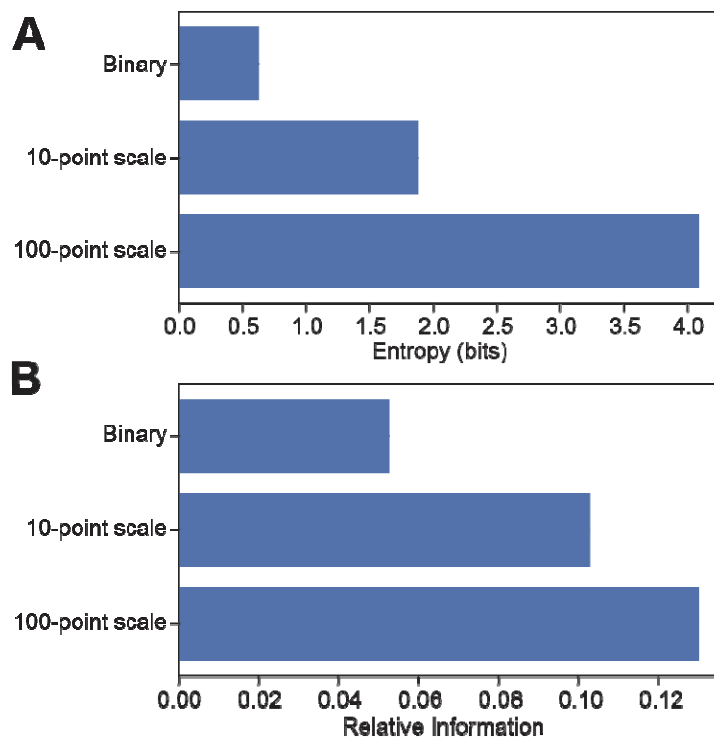
## Chemosensory characterization of C19+ and C19-

We asked how accurately COVID-19 status could be predicted from the survey responses. The data matrix had strictly non-negative values and was normalized (column-wise min=0, max=1) to apply regularization in an equitable fashion across features and give regression coefficients the same interpretation for each feature. Compared with the main text, models with similar AUC values (but with non-zero coefficients for additional, likely spurious features) were obtained for smaller values of  $\alpha$ , and inferior results for larger ones (which contained fewer or no non-zero coefficients). Quantitatively similar AUC values were obtained for other models predicting COVID-19 status using multiple features including ridge regression and random forest, but L1-regularized logistic regression consistently produced sparser models with comparable cross-validation accuracy. Each logistic regression model included an intercept term and one or more normalized features. Each model attempted to predict, using the value of the response to a single question (and an additive constant), whether a subject reported a C19+ or C19- status. Coefficients in a logistic regression model can be interpreted as changes in odds, or as odds ratios when two values are compared. Each ROC curve -- constructed using predictions on holdout test sets and concatenated over these test sets -- summarizes the tradeoff between sensitivity (fraction of C19+ cases correctly identified) and specificity (fraction of C19- cases correctly identified) as the threshold value for the predictor is varied.

## Value of using a scale rather than a binary response to detect C19+

We quantified the information entropy for each survey question used the following standard equation:  $I = -\sum_i^n (p_i * \log_2(p_i))$  evaluated over the  $n$  response options. Re-binning to mimic new scales was achieved by dividing response values by a constant and rounding to the nearest integer. Relative mutual information was calculated by computing the mutual information between survey response and COVID status based on the following standard equation:  $MI = \sum_i^n \sum_j^2 (p_{ij} * \log_2(\frac{p_{ij}}{p_i p_j}))$  where survey response options are indexed with  $i$  and the C19+/C19- status (two possible values) are indexed with  $j$ , and then dividing by the entropy available from that same C19 status distribution, calculated using the first equation. Results indicate that soliciting responses

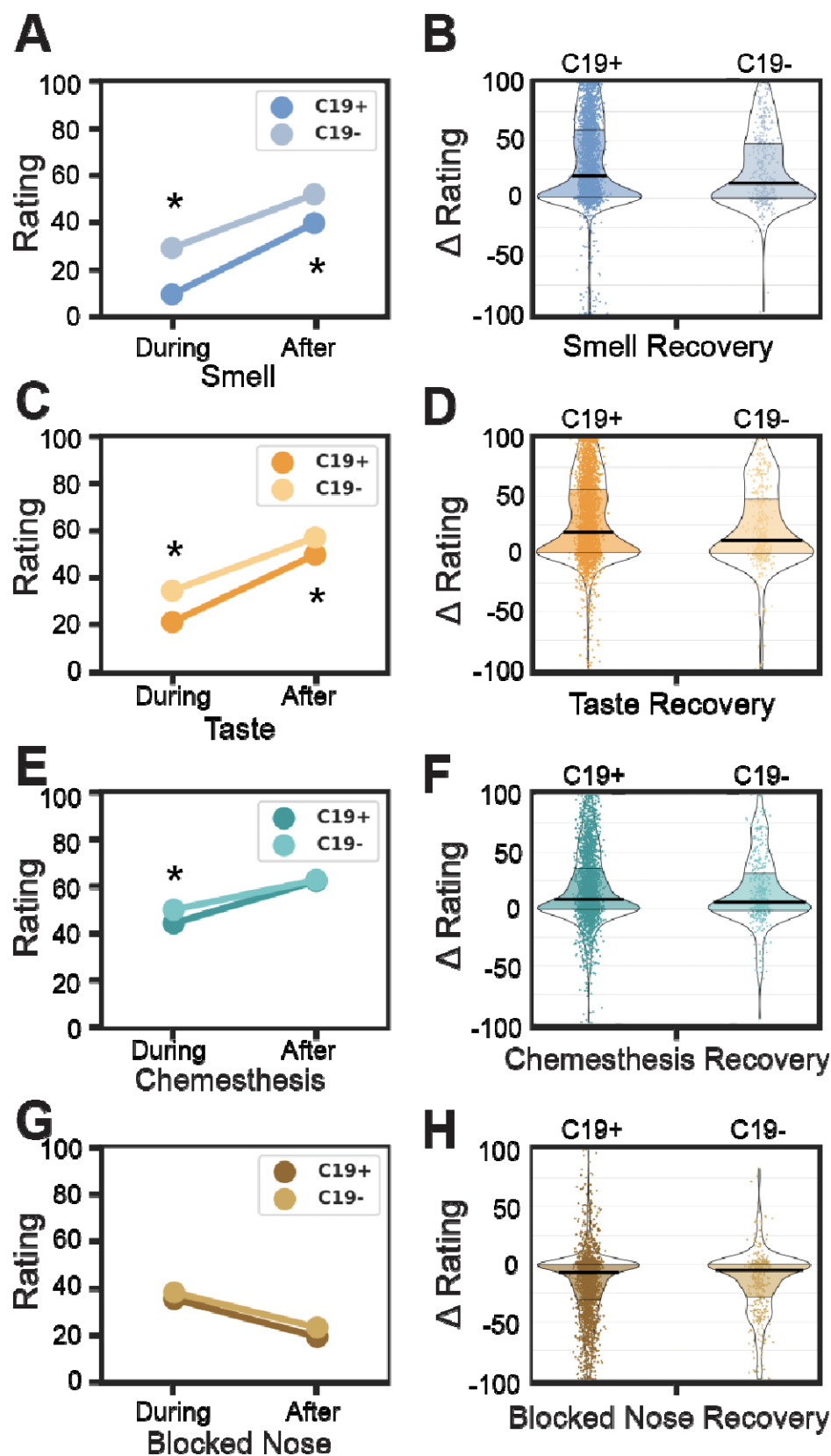
on either a continuous 100-point scale or a downsampled 10-point numeric version of the scale is more informative about symptoms themselves and about COVID-19 status (given the symptoms) than soliciting binary responses (**Figure S5**).



**Figure S5. (A)** Relative information available from the distribution of responses to the two primary “Smell” survey questions. *Binary* refers to the yes/no question about symptomatic smell loss. A relative information of 1 would correspond to a question whose response is perfectly informative about COVID-19 status. By contrast, a similar question asked on a numeric scale (0-100, the original scale; or a hypothetical 10-point scale obtained by rounding responses) contains substantially more information due to the resolution of the scale. A 10-point scale may be familiar from clinical self-reports of pain. **(B)** The relative mutual information about COVID status contained in the survey response is also higher for the full numeric scale or the hypothetical 10-point scale than for the binary question.

## Prediction of recovery from COVID-19-associated smell loss

We applied the same predictive modeling framework used in **Figure 4** to try to predict smell recovery in C19+ participants. In other words, we asked which survey responses predicted that a subject would fall into the Recovered Smell rather than the Persistent Smell Loss cluster, given both smell loss during the disease and C19+ status. The only predictive feature of any practical significance was “Days Since Onset” of respiratory symptoms (AUC=0.62), indicating that those who experienced their first respiratory symptoms less recently are more likely to have Recovered Smell (**Figure S6A**). Adding additional features to the model provided modest improvement (AUC=0.65 for the optimal model), but overall it was difficult to predict whether a C19+ participant would exhibit Recovered Smell or Persistent Smell Loss based on the data available (**Figure S6B**). **Table S3** includes the means and SD by recovery group for C19+ and C19-participants.



**Figure S6.** COVID-19 recovery. Similar to Figure 1, but self-reported smell (A,B), taste (C,D), chemesthesis (E,F), and nasal blockage(G,H) during and after respiratory illness in C19+ (darker) versus C19- (lighter). (A,C,E,G) mean values during and after respiratory illness, respectively. (B,D,F,H) Change (after minus during) as a distribution over subjects.

**Table S3.** Main chemosensory and relevant demographic features in the three clusters of recovering C19+ and C19- participants.

	Intact Smell			Persistent Smell Loss			Recovered Smell		
	C19+	C19-	p	C19+	C19-	p	C19+	C19-	p
<b>Taste Change</b>	-34 ± 39	-21 ± 30	0.011	-75 ± 28	-68 ± 32	0.0018	-75 ± 29	-69 ± 31	0.0043
<b>Chemesthesis Change</b>	-17 ± 32	-5.9 ± 25	0.016	-40 ± 37	-42 ± 38	0.2	-36 ± 37	-37 ± 39	0.41
<b>Changes In Sweet</b>	0.31 ± 0.46	0.19 ± 0.39	0.0067	0.46 ± 0.5	0.4 ± 0.49	0.067	0.47 ± 0.5	0.45 ± 0.5	0.36
<b>Changes In Sour</b>	0.27 ± 0.45	0.14 ± 0.35	0.0037	0.43 ± 0.5	0.42 ± 0.5	0.42	0.4 ± 0.49	0.42 ± 0.5	0.29
<b>Smell Change</b>	-3.6 ± 14	-8.1 ± 12	0.022	-89 ± 13	-82 ± 19	1.50E-07	-89 ± 13	-83 ± 17	4.20E-05
<b>Phantosmia</b>	0.09 ± 0.29	0.11 ± 0.31	0.35	0.09 ± 0.28	0.11 ± 0.31	0.21	0.08 ± 0.28	0.10 ± 0.3	0.32
<b>Days Since Onset</b>	28 ± 18	27 ± 17	0.35	27 ± 20	33 ± 26	0.0039	32 ± 17	33 ± 20	0.32
<b>Parosmia</b>	0.15 ± 0.36	0.15 ± 0.36	0.48	0.09 ± 0.28	0.12 ± 0.33	0.063	0.12 ± 0.33	0.15 ± 0.36	0.19
<b>Changes In Savory</b>	0.18 ± 0.39	0.16 ± 0.37	0.32	0.4 ± 0.49	0.36 ± 0.48	0.17	0.32 ± 0.47	0.35 ± 0.48	0.26
<b>Onset Day</b>	95 ± 24	91 ± 23	0.078	1e+02 ± 29	95 ± 30	0.00036	90 ± 21	91 ± 27	0.47
<b>Taste Recovery</b>	15 ± 33	13 ± 28	0.43	9.2 ± 23	6 ± 29	0.07	55 ± 30	50 ± 31	0.043
<b>Changes In Bitter</b>	0.27 ± 0.45	0.17 ± 0.38	0.018	0.43 ± 0.5	0.4 ± 0.49	0.2	0.41 ± 0.49	0.4 ± 0.49	0.41
<b>Changes In Salty</b>	0.36 ± 0.48	0.21 ± 0.41	0.0026	0.47 ± 0.5	0.42 ± 0.49	0.12	0.47 ± 0.5	0.44 ± 0.5	0.23
<b>Fraction Women</b>	0.63 ± 0.48	0.79 ± 0.41	0.0017	0.76 ± 0.43	0.77 ± 0.42	0.38	0.73 ± 0.44	0.75 ± 0.43	0.32
<b>Anosmia/Hyposmia</b>	0.43 ± 0.5	0.14 ± 0.35	3.50E-08	0.94 ± 0.23	0.86 ± 0.35	1.40E-05	0.88 ± 0.32	0.86 ± 0.35	0.23
<b>Chemesthesis Recovery</b>	8.3 ± 27	8.3 ± 24	0.24	9.5 ± 26	10 ± 27	0.41	29 ± 33	25 ± 38	0.25
<b>Smell Recovery</b>	-3.8 ± 39	4.4 ± 22	0.14	6.3 ± 10	6.3 ± 11	0.22	67 ± 19	62 ± 18	0.007
<b>Age</b>	44 ± 13	44 ± 13	0.33	41 ± 12	42 ± 12	0.092	40 ± 12	43 ± 12	0.0015

# APPENDIX 1

## GCCR core questionnaire

The core questionnaire of the Global Consortium for Chemosensory Research (GCCR) has been deployed in Compusense Cloud in 32 languages. The questionnaire was published previously<sup>8</sup> and also appears in the NIH Office of Behavioral and Social Sciences Research (OBSSR) research tools for COVID-19.<sup>39</sup> Responses to the GCCR core questionnaire in 23 languages were collected between April 7 and July 2, 2020 and included in the final dataset, on which we conducted the analyses reported in this paper.

## Global Consortium for Chemosensory Research (GCCR) Questionnaire



Global Consortium for  
Chemosensory Research

### Welcome and Consent

1. I consent to participate.  [1] Yes  
 [0] No

Question Type: Choose only 1  
Branching logic: if [0]=checked, then go to Section 4) End of test

### About your illness

- 
2. In which year (YYYY) were you born?

Question Type: Numeric  
Branching logic: if year of birth greater than 2001 then go to Section 4) End of test

- 
3. What is your current country of residence?

Question Type: Comment

- 
4. **Optional:** What city, town, or region do you currently live in?

Question Type: Comment

- 
5. Which gender do you most identify with?

- [1] Female  
 [2] Male  
 [3] Another not listed here  
 [4] Prefer not to say

Question Type: Choose only 1

- 
6. Within the past two weeks, have you been diagnosed with or suspect that you have a respiratory illness?  [1] Yes  
 [0] No

Question Type: Choose only 1  
Branching: if [0]=checked, then go to Section 4) Re-contact

- 
7. What **date** did you **first notice symptoms** of your recent respiratory illness? Provide your best guess or leave blank if you do not remember. Click the box below to display a calendar.

Question Type: Numeric

- 
8. Have you been diagnosed with **COVID-19**?

- [1] Yes-diagnosed based on symptoms only  
 [2] Yes-diagnosed with viral swab  
 [3] Yes-diagnosed with another lab test  
 [4] No-I was not diagnosed, but I have symptoms  
 [5] No-I had a negative test, but I have symptoms  
 [6] No-I do not have any symptoms  
 [7] Don't Know  
 [8] Other

Question Type: Choose only 1

- 
9. Were you diagnosed with any **other respiratory illnesses (not COVID-19)** in the **last two weeks**?

(Select all that apply)

- [1] Strep throat (Streptococcal bacteria)  
 [2] Another bacterial illness  
 [3] Flu (influenza)  
 [4] Another viral illness  
 [5] Other  
 [6] None

Question Type: Choose n

---

10. Have you had any of the following symptoms with **your recent respiratory illness or diagnosis**? (Select all that apply)

- [1] Fever
- [2] Dry cough
- [3] Cough with mucus
- [4] Difficulty breathing/shortness of breath
- [5] Chest tightness
- [6] Runny nose
- [7] Sore throat
- [8] Changes in food flavor
- [9] Changes in smell
- [10] Loss of appetite
- [11] Headache
- [12] Muscle aches
- [13] Fatigue
- [14] Diarrhea
- [15] Abdominal pain
- [16] Nausea
- [17] No symptoms

Question Type: Choose n

---

11. **Optional:** Please describe the progression or order you noticed your symptoms

Question Type: Comment

---

12. **Optional:** What treatment(s) or medication(s) have you received for your recent **respiratory illness or diagnosis**?

Question Type: Comment

The next section of this survey is focused on your experience of smell, taste, and food flavor during your recent respiratory illness or diagnosis.

**These questions relate to your sense of smell (for example, sniffing flowers or soap, or smelling garbage) but not the flavor of food in your mouth.**

---

13. Rate your ability to **smell** BEFORE your **recent respiratory illness or diagnosis**

No sense of smell Excellent sense of smell



(Place a mark on the scale above)

Question Type: Line Scale

---

14. Rate your ability to **smell** DURING your **recent respiratory illness or diagnosis**

No sense of smell Excellent sense of smell



(Place a mark on the scale above)

Question Type: Line Scale

---

15. Have you experienced any of the following changes in **smell** with your **recent respiratory illness or diagnosis**? (Select all that apply)

- [1] I cannot smell at all / Smells smell less strong than they did before
- [2] Smells smell different than they did before (the quality of smell has changed)
- [3] I can smell things that aren't there (e.g. I smell burning when nothing is on fire)
- [4] Sense of smell fluctuates (e.g. comes and goes)

Question Type: Choose n

---

16. **Optional:** Please describe any changes in smell

\_\_\_\_\_



Question Type: Comment

---

17. How **blocked** was your nose BEFORE your recent respiratory illness or diagnosis?

Not at all blocked Completely blocked

=====

*(Place a mark on the scale above)*

Question Type: Line Scale

---

18. How **blocked** was your nose DURING your recent respiratory illness or diagnosis?

Not at all blocked Completely blocked

=====

*(Place a mark on the scale above)*

Question Type: Line Scale

---

**The following questions are related to your sense of taste. For example sweetness, sourness, saltiness, bitterness experienced in the mouth**

---

19. Rate your ability to **taste** BEFORE your recent respiratory illness or diagnosis

No sense of taste Excellent sense of taste

=====

*(Place a mark on the scale above)*

Question Type: Line Scale

---

20. Rate your ability to **taste** DURING your recent respiratory illness or diagnosis

No sense of taste Excellent sense of taste

=====

*(Place a mark on the scale above)*

Question Type: Line Scale

---

21. **OPTIONAL:** Have you experienced changes to **specific tastes** with your recent respiratory illness or diagnosis? (Select all that apply)

[1]Sweet  
 [2]Salty  
 [3]Sour  
 [4]Bitter  
 [5]Savory/Umami

Question Type: Choose n

---

22. **Optional:** Describe any **changes in taste** during your recent respiratory illness or diagnosis.

\_\_\_\_\_

Question Type: Comment

---

**The following questions are related to other sensations in your mouth, like burning, cooling, or tingling. For example chili peppers, mint gum or candy, or carbonation.**

---

23. Rate your **ability to feel these other sensations** BEFORE your recent respiratory illness or diagnosis.

Not sensitive at all Very sensitive

=====

*(Place a mark on the scale above)*

Question Type: Line Scale

---

24. Rate your **ability to feel these other sensations** DURING your recent respiratory illness or diagnosis

Not sensitive at all Very sensitive

=====

*(Place a mark on the scale above)*

Question Type: Line Scale

**25. Optional:** Describe any changes in these other sensations during your **recent respiratory illness or diagnosis**.

Question Type: Comment

**26. Optional:** Think about a food or beverage you consume regularly - for example, your morning coffee or tea or a piece of fruit you have each day. Has the taste, smell, or flavor changed with your recent respiratory illness or diagnosis? If so, **please describe how and be sure to indicate which food or beverage you are describing**.

Question Type: Comment

**27. Optional:** Is there anything else you would like to tell us about how your recent respiratory illness or diagnosis has affected your sense of smell, taste, and flavor?

Question Type: Comment

**28.** Have you recovered from your **recent respiratory illness or diagnosis**? (For example you no longer have a cough, fever, or shortness of breath.)

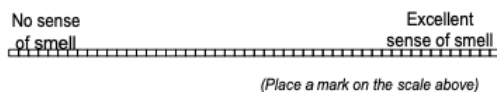
- No
- Yes - partly
- Yes - fully
- Don't know

Question Type: Choose only 1

Branching: if [0]=checked, then go to Section 3) General Health Information

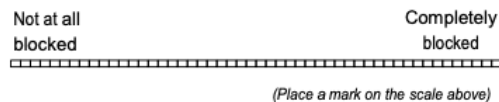
The next section of this survey is focused on your experiences of smell, taste, and food flavor after **your** recovery from your recent respiratory illness or **diagnosis**.

**29.** Rate your ability to **smell** AFTER your recovery



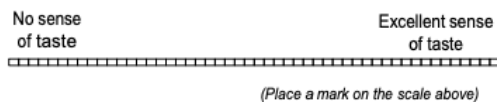
Question Type: Line Scale

**30.** How **blocked** was your nose AFTER your recovery



Question Type: Line Scale

**31.** Rate your ability to **taste** AFTER your recovery



Question Type: Line Scale

---

32. Rate your **ability to feel these other sensations like burning, cooling, and tingling** AFTER your recovery

Not sensitive at all Very sensitive  
-----

(Place a mark on the scale above)

Question Type: Line Scale

---

33. How were you directed to this survey?

- [1] Clinician or healthcare professional
- [2] Media (social media, print, radio, tv, etc)
- [3] Word of mouth
- [4] Other

Question Type: Choose only 1

The next section of this survey will ask some optional questions about your habits and general health.

## General Health Information

34. **Optional:** Have you smoked at least 100 combustible cigarettes or cigars in your entire life?

- [1] No
- [2] Yes
- [3] Prefer not to say
- [4] Don't know

Question Type: Choose only 1

---

35. **Optional:** During the past 30 days, on how many days did you smoke combustible cigarettes or cigars?

Question Type: Numeric

---

36. **Optional:** Have you ever used an e-cigarette ('vaped'/'Juuled') even one time? (E-cigarettes are battery-powered devices that usually contain liquid nicotine, and do not produce smoke.)

- [1] No
- [2] Yes
- [3] Prefer not to say
- [4] Don't know

Question Type: Choose only 1

---

37. **Optional:** During the past 30 days, on how many days did you use an e-cigarette?

Question Type: Numeric (range 0-30; integer)

---

38. Did you have any of the following **in the 6 months prior** to your recent respiratory illness or diagnosis? (Select all that apply)

- [1] High blood pressure
- [2] Heart disease (heart attack or stroke)
- [3] Diabetes (high blood sugar)
- [4] Obesity
- [5] Lung disease (asthma/COPD)
- [6] Head trauma
- [7] Neurological disease
- [8] Cancer that required chemotherapy or radiation
- [9] Cancer that did NOT require chemotherapy or radiation
- [10] Chronic sinus problems
- [11] Seasonal allergies/hay fever
- [12] None

Question Type: Choose n

---

39. **Optional:** Any other medical conditions that you would like to mention?

Question Type: Comment

---

40. **Optional:** Which medication(s) do you take regularly? For example, medications for pain, blood pressure, thyroid function, anti-viral, etc.

Question Type: Comment

---

41. **Optional:** Is there anything we didn't ask about that you would like to share with us?

---

Question Type: Comment

## Re-contact

---

42. We may want to re-contact you for follow up research on this topic. **Is it okay if our team or other researchers re-contact you to participate in future research?** By saying yes, you agree that we can share your email address with other researchers for this purpose.

[1] Yes  
 [0] No

Question Type: Choose only 1

Branching logic: if [0]=checked, then go to: Section 5)End of test

---

43. Please provide your full email address, so you can be contacted for future studies by our team or other researchers.

---

Question Type: Comment

## End of Test

**You have now completed the survey and may close your browser**

Thank you for your time!

## Notes

"In which year (YYYY) were you born?"

-- value must be 1900 or greater

"What **date** did you **first notice symptoms** of your recent respiratory illness? Provide your best guess or leave blank if you do not remember.

Click the box below to display a calendar"

-- format (mm/dd/yyyy)

"Have you been diagnosed with COVID-19?"

-- if [8] Other was selected, a comment is required

"Were you diagnosed with any other respiratory illnesses (not COVID-19) in the last two weeks? (Select all that apply)"

-- if [6] None was selected, no other options can be selected.

"Have you had any of the following symptoms with your **recent respiratory illness or diagnosis**? (Select all that apply)"

-- if [17] No symptoms was selected, no other options can be selected.

"Rate your ability to smell BEFORE your **recent respiratory illness or diagnosis**"

-- Line Scale Range 0-100, intervals of 1. All following line scales formatted similarly

All line scales had anchors indented at ~10% and 90% of scale range.

'**OPTIONAL:** During the past 30 days, on how many days did you smoke combustible cigarettes or cigars?' and "**OPTIONAL:** During the past

-- value must be between 0-30

Did you have any of the following in the 6 months prior to your **recent respiratory illness or diagnosis**? (Select all that apply)

-- if [12] None was selected, no other options can be selected.