



Cornell University
Office of
Research Integrity and Assurance

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Institutional Review Board for Human Participants

Notice of Exemption

To: Hannah Kelson
From: Andrew Willford,
IRB Chairperson

A handwritten signature in blue ink, appearing to read 'Andrew Willford'.

Protocol ID#: 1912009302
Protocol Title: Measuring Smell and Emotion Using the Sniff Olfactometer
Approval Date: March 05, 2020
Expiration Date: None

Your protocol has been granted exemption from IRB review according to Cornell IRB policy and under paragraph(s) 3 of the Department of Health and Human Services Code of Federal Regulations 45CFR 46.104(d).

• Paragraph 3 allows to be exempted from IRB review research activities in which the only involvement of human subjects will be in the following category: Benign Behavioral Interventions - Research involving benign behavioral interventions in conjunction with the collection of information from an adult subject through verbal or written responses (including data entry) or audiovisual recording if the subject prospectively agrees to the intervention and information collection, and if: i) the information obtained is recorded in such a manner that the human subjects cannot be identified, directly or through identifiers linked to the subjects; or ii) any disclosure of the human subjects' responses outside the research would not reasonably place the subjects at risk of criminal or civil liability or be damaging to the subjects' financial standing, employability, educational advancement, or reputation..

Please note the following:

- Investigators are responsible for ensuring that the welfare of research subjects is protected and that methods used and information provided to gain participant consent are appropriate to the activity. Please familiarize yourself with and conduct the research in accordance with the ethical standards of the Belmont Report (<https://www.hhs.gov/ohrp/regulations-and-policy/belmont-report/index.html>).
- Investigators are responsible for notifying the IRB office of change or amendments to the protocol and acquiring approval or concurrence **BEFORE** their implementation.
- Progress reports, requests for personnel or other administrative changes, or requests for continuation of approval are not required for the study. However, upon conclusion of the study, please submit a Project Closure form: <http://www.irb.cornell.edu/forms>.

For questions related to this application or for IRB review procedures, please contact the IRB office at irbhp@cornell.edu or 607-254-5162. Visit the IRB website at www.irb.cornell.edu for policies, procedures, FAQs, forms, and other helpful information about Cornell's Human Participant Research Program. Please download the latest forms from the IRB website www.irb.cornell.edu/forms/ for each submission.

Cc: Terry Acree