CLEANED/DERIVED VARIABLE METADATA TOP SHEET

| | 07/06/2019 |
|---|--|
| Date of submitting documentation | 07/06/2019 |
| Categories of variables ^{*:} (may be more than one) | UPSIT |
| Summary of work undertaken | See methods below All returned booklets were marked by a member of the study team and updated to XNAT Any participants with missing scores: TP searched for any unmarked booklets to ensure all returned booklets were marked Excel spreadsheet generated by TP |
| Source data file(s) | UPSIT booklets XNAT |
| Date source file(s) created: | 07/06/2019 |
| Names of source variables | Xnat (participant number) Upsit (score out of 40, missing data -99) change_smell (recent subjective change in sense of smell) reason data missing/test not done |
| Syntax provided | No |
| Location of syntax file | N/A |
| Date syntax file created: | N/A |
| Format of syntax | N/A |
| Output variables (please list names of new variables created) | Same as source variables |
| Output data file provided | Yes |
| Date output file created: | 07/06/2019 |
| Location of output file | N:\Test Data and Video Files\Phase 1\3_Cleaned Data\Insight46_upsit_cleaned_final_20190607 |
| Format of output file | excel |
| Documentation provided | N/A |
| List any papers in which cleaned/derived variables have been used | nil |

For Submission to the NSHD Scientific Support Team

Prior to olfactory testing participants were asked whether they had noticed a decline in their sense of smell over the past few years? In addition, participants were prospectively excluded and did not undertake smell testing if they provided a clear history one of the following: nasal bone fracture; nasal surgery; current diagnosis of rhinitis or nasal polyps; or another chronic nasal condition that could potentially affecting olfactory function.

The University of Pennsylvania Smell Identification Test (UPSIT) was used to asses olfactory function. The UPSIT is a commercially available, well-established, reliable, and standardized olfactory test that can be self-administered. The 'British' version was selected as the most culturally appropriate to the NSHD. Each test comprises four booklets with one odorant (embedded in 10–50-micron diameter microcapsules fixed in a proprietary binder and positioned on brown strips) at the bottom of each page (ten pages per booklet). Accompanying each strip is a multiple-choice question with four responses following an alternative forced choice paradigm. Packs were provided to participants to complete at home and returned using a stamped address envelope provided. All participants were advised if they had or developed an upper respiratory tract infection to complete the test after being symptom free for 2 weeks. If a participant only completed 38 or 39 items, a score of 0.25 was assigned for each missing item. If a participant completed 37 or less items, they were excluded from the analysis.