Cotinine
PT Event ID: $\quad 201702 \mathrm{SCOH}$
Participant:
Analyst:
Reviewer:

Units of Result: \begin{tabular}{l}

| Form Approved |
| :--- |
| OMB No. |
| Exp. Date $\mathrm{xx} / \mathrm{mx} / \mathrm{xxxx}$ | \\

\hline
\end{tabular}

| Sample ID | Analyte | Reported Value |
| :---: | :--- | :--- |
| SCOH-QP3 | cotinine |  |


| Sample ID | Analyte | Reported Value |
| :--- | :--- | :--- |
| SCOH-909 | cotinine |  |


| Sample ID | Analyte | Reported Value |
| :---: | :--- | :--- |
| SCOH-806 | cotinine |  |

By submitting this form, we attest that the results reported were produced in this laboratory from the analysis of proficiency testing samples that were introduced into the routine workflow of the laboratory and analyzed using protocols and procedures with the same frequency routinely applied to patient specimens.

We further attest that the laboratory did not discuss or engage in any communications with anyone outside of our laboratory regarding the proficiency test or the results obtained.

CDC estimates the average public reporting burden for this collection of information as 45 minutes per response, including the time for reviewing instructions, searching existing data/information sources, gathering and maintaining the data/information needed, and completing and reviewing the collection of information. An agency may not conduct or sponsor, and a person is not required to respond to a collection of information unless it displays a currently valid OMB Control Number. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden to CDC/ATSDR Information Collection Review Office, 1600 Clifton Road NE, MS D-74, Atlanta, Georgia 30333; ATTN: PRA 0920-xxxx).

