

HIGH SENSITIVITY SALIVARY COTININE QUANTITATIVE ENZYME IMMUNOASSAY KIT

Catalog No. 1-2002/1-2012, 96-Well Kit

For Research Use

Intended Use

This assay kit is a competitive immunoassay designed for the quantitative measurement of the presence of cotinine in saliva samples. Please read the complete kit insert before performing this assay. For further information about this kit, its application, or the procedures in this insert, please contact the technical service team at Salimetrics or your local sales representative.

Introduction

Since primitive times, tobacco leaves have been processed and used by humans to deliver nicotine to the central nervous system. Historically, the preferred route of nicotine administration has varied from snuff, chew, and inhalation of smoke from burned tobacco leaves to contemporary methods such as transdermal patches, chewing gums, and smokeless inhalers. Regardless of delivery route, nicotine has addictive properties that cause the user to continue use despite efforts to quit. When tobacco leaves are smoked in cigarettes, nicotine is absorbed and distributed in the body within seconds. Metabolism is mainly by oxidation to cotinine and nicotine-N-oxide. Volumes of literature document the negative economic impact and health consequences of tobacco use. The costs to individuals and societies associated with smoking (and secondhand exposure to tobacco smoke) have led to widespread public health interventions to curb smoking behavior.

The detection of exposure to tobacco smoke by measurement of cotinine is the preferred method. Cotinine can be measured noninvasively (in saliva) following nicotine administration. Nicotine is not considered a valid marker of smoking status due to its relatively short half-life (approximately 2 hours). By contrast, cotinine in saliva has a half-life of more than 10 hours and the literature has documented it to be a specific and sensitive marker for determining exposure to tobacco/nicotine.

The majority of commercially available assays for salivary cotinine are qualitative. They return "positive" or "negative" determinations with respect to tobacco/nicotine exposure. However, many studies show that levels of cotinine in saliva show large inter-individual differences. The sources of these differences include factors related to intrinsic and extrinsic predispositions that affect the physiology of nicotine metabolism, the dose of nicotine present in the cigarette (or alternative source), and health behaviors relevant to how cigarettes are smoked (e.g., vent blocking, duration and frequency of puffs). There is a clear need for inexpensive, accurate, quantitative, and noninvasive means of validating smoking status, measuring the immediate physiological consequences of individual differences and intra-individual change in smoking behaviors, and determining secondhand tobacco smoke exposure. Salimetrics has designed this research tool to provide biomedical researchers with a highly sensitive and reliable means to do so.

ELISA methods have been shown to be 100% effective in distinguishing non-smokers from smokers (1). EIA results have shown near perfect agreement with GC/MS confirmation of smoking status (2). Levels, however, may be higher with EIA since 3-OH-cotinine, a metabolite of nicotine is also measured. Cotinine levels in saliva have been found to be independent of salivary flow rate (3). Due to the ease of collection and accuracy of measurement, saliva has been declared the matrix of choice for the measurement of environmental tobacco smoke (1).

Test Principle

Standards and unknowns are added to a 96-well microtiter plate along with rabbit antibodies to cotinine and cotinine linked to horseradish peroxidase (conjugate). The cotinine in standards and unknowns and the conjugate compete for the antibody binding sites. After incubation, unbound components are washed away. Bound conjugate is measured by the reaction of the peroxidase enzyme on the substrate tetramethylbenzidine (TMB). This reaction produces a blue color. A yellow color is formed after stopping the reaction with 2-molar sulfuric acid. Optical density is read on a standard plate reader at 450 nm. The amount of cotinine peroxidase detected is inversely proportional to the amount of cotinine present (4).

pH Indicator

A pH indicator in the assay diluent alerts the user to samples with high or low pH values. Acidic samples will turn the diluent yellow. Alkaline samples will turn the diluent purple. Dark yellow or purple wells indicate that a pH value for that sample should be obtained using pH strips. Cotinine values from samples with a pH ≤ 3.5 or ≥ 9.0 may be artificially inflated or lowered (5).

Precautions

1. Stop solution is a 2-molar solution of sulfuric acid. This solution is caustic; use with care.
2. This kit uses break-apart microtitre strips. Unused wells must be stored at 2 - 8°C in the sealed foil pouch and used in the frame provided.
3. Do not mix components from different lots of kits.
4. When using a multichannel pipette, reagents should be added to duplicate wells at the same time. Follow the same sequence when adding additional reagents so that incubation time with reagents is the same for all wells.
5. See 'Material Safety Data' at the end of procedure.
6. Routine calibration of pipettes is critical for the best possible assay performance.
7. Pipetting of samples and reagents must be done as quickly as possible (without interruption) across the plate.
8. When running multiple plates, or multiple sets of strips, a standard curve should be run with each individual plate and/or set of strips.
9. The temperature of the laboratory may affect assays. Salimetrics' kits have been validated at 68 - 74°F (20 - 23.3°C). Higher or lower temperatures will cause an increase or decrease in OD values, respectively. Salimetrics cannot guarantee test results outside of this temperature range.

Storage

All components of this kit are stable at 2 - 8°C until the kit's expiration date.

Reagents and Reagent Preparation

1. **Microtitre plate:** A 96-well high affinity binding plate.
2. **Antiserum:** 15 mL of a solution of rabbit anti-cotinine antibody.
3. **Cotinine Standard:** 0.5 mL of cotinine in a saliva-like matrix with a non-mercury preservative, at a concentration of 200 ng/mL.
4. **Wash Buffer:** 100 mL of a 10X phosphate buffered solution containing detergents and a non-mercury preservative. Dilute only the amount needed for current day's use. Discard any leftover reagent. Dilute the wash buffer concentrate 10-fold with room temperature deionized water (100 mL of 10X wash buffer to 900 mL of deionized H₂O). (*Note: If precipitate has formed in the concentrated wash buffer, it may be heated to 60°C for 15 minutes. Cool to room temperature before use in assay.*)
5. **Assay Diluent:** 63 mL of a phosphate buffered solution containing a pH indicator and a non-mercury preservative.
6. **Enzyme Conjugate:** 75 μ L of a solution of cotinine labeled with horseradish peroxidase. Dilute prior to use with assay diluent.
7. **Tetramethylbenzidine (TMB):** 25 mL of a non-toxic ready-to-use solution.
8. **Stop Solution:** 12.5 mL of a 2-molar solution of sulfuric acid (USA customers only). Stop solution is provided in powdered form to customers outside the USA. Reconstitute the powdered stop solution with 12.5 mL of deionized water. Let sit for 10 minutes before use.
9. **Cotinine Controls:** Two controls representing high and low levels of cotinine in a saliva-like matrix with a non-mercury preservative. Each vial contains 0.25 mL. See vials for target ranges.

Materials Needed But Not Supplied

- Precision pipette to deliver 20 μ L, 50 μ L, and 100 μ L
- Precision multichannel pipette to deliver 50 μ L, 100 μ L, and 200 μ L
- Vortex
- Microplate incubator/shaker
- Plate reader with a 450 nm filter
- Software for data reduction
- Deionized water
- Reagent reservoirs
- One 15 mL disposable tube
- Small disposable tubes
- Pipette tips
- 10 mL serological pipette

Specimen Collection

The preferred saliva collection method (5) is to use plain (non-citric acid) cotton Salivettes, P/N 5001. Samples may also be collected using Sorbettes, P/N 5029 (for infants) or cotton ropes, P/N 5016 or by passive drool. For accurate results collection devices should be completely saturated before removal. **Do not** add sodium azide to saliva samples as a preservative. Samples visibly contaminated with blood should be recollected. Cotinine in saliva has been reported in the literature to be stable at room temperature for up to 12 days (6). However, room temperature exposure should be avoided as much as possible to avoid bacterial growth in the specimen. Freeze at -20°C or lower for long-term storage. Contact the technical service team at Salimetrics for more detailed information on specimen collection.

Saliva samples should be frozen prior to assay to precipitate the mucins. On day of assay, thaw completely, vortex, and centrifuge at 1500 x g (@3000 rpm) for 15 minutes. Pipette clear sample into appropriate wells. Particulate matter may interfere with antibody binding, leading to falsely elevated results.

Sample Preparation

Known smokers: Dilute saliva samples x 10 (10 µL saliva into 90 µL assay diluent).

Non-smokers: Run saliva sample straight.

Procedure

Bring all reagents to room temperature.

Step 1: Determine your plate layout. Here is a suggested layout.

	1	2	3	4	5	6	7	8	9	10	11	12
A	200 Std	200 Std	C-H	C-H								
B	66.7 Std	66.7 Std	C-L	C-L								
C	22.2 Std	22.2 Std	Unk-1	Unk-1								
D	7.4 Std	7.4 Std	Unk-2	Unk-2								
E	2.5 Std	2.5 Std	Unk-3	Unk-3								
F	0.8 Std	0.8 Std	Unk-4	Unk-4								
G	Zero	Zero	Unk-5	Unk-5								
H	NSB	NSB	Unk-6	Unk-6								

Step 2: Keep the desired number of wells in the strip holder and return any remaining strips to the foil pouch. Reseal the zip-lock and refrigerate at 2 - 8°C.

Step 3:

- Label five small tubes, such as microcentrifuge tubes, with numbers 2 through 6.
- Pipette 100 µL of assay diluent in tubes 2 through 6. Serially dilute the standard 3X by adding 50 µL of the 200 ng/mL standard (tube 1) to tube 2. Mix well. After changing pipette tips, remove 50 µL from tube 2 to tube 3. Mix well. Continue for tubes 4, 5 and 6. The final concentrations of standards for tubes 1 through 6 are respectively, 200 ng/mL, 66.7 ng/mL, 22.2 ng/mL, 7.4 ng/mL, 2.5 ng/mL and 0.8 ng/mL.
- Pipette 15 mL of assay diluent into a disposable tube. Set aside for Step 5.

Step 4:

- Pipette 20 µL of standards, controls and unknowns into appropriate wells. Standards, controls and unknowns should be assayed in duplicate.
- Pipette 20 µL of assay diluent into 2 wells to serve as the zero.
- Pipette 120 µL of assay diluent into 2 wells to serve as the non-specific binding.

Step 5: Dilute the enzyme conjugate 1:300 by adding 50 µL of the conjugate to the 15 mL of assay diluent prepared in Step 3 (full plate only). Immediately mix the diluted conjugate solution and add 100 µL to each well using a multichannel pipette.

Step 6: Pipette 100 µL of antiserum into all wells, except the nonspecific binding wells, using a multichannel pipette.

Step 7: Cover the plate with a plate cover. Incubate the plate on a microplate incubator/shaker for 1.5 hours at 37°C with constant mixing at 500-600 rpm.

Step 8: Wash the plate 4 times with 1X wash buffer. A plate washer is recommended. However, washing may be done by gently squirting wash buffer into each well with a squirt bottle or by pipetting 300 µL of wash buffer into each well and then flipping the liquid into a sink. After each wash, the plate should be thoroughly blotted on paper towels before turning upright. If using a plate washer, blotting is still recommended after the last wash.

Step 9: Add 200 µL of TMB solution to each well using a multichannel pipette.

Step 10: Mix at 500 rpm for 5 minutes (or tap to mix) and incubate in the dark for an additional 25 minutes.

Step 11:

- Add 50 µL of stop solution using a multichannel pipette.
- Mix on a plate rotator at room temperature for 3 minutes at 500 rpm (or tap to mix). Be sure all wells have turned yellow. If green color remains, continue mixing until green color turns to yellow.
- Wipe off bottom of plate with a water-moistened lint-free cloth and wipe dry.
- Read in a plate reader at 450 nm. Read plate within 10 minutes of adding stop solution (correction at 620 is desirable).

Calculations

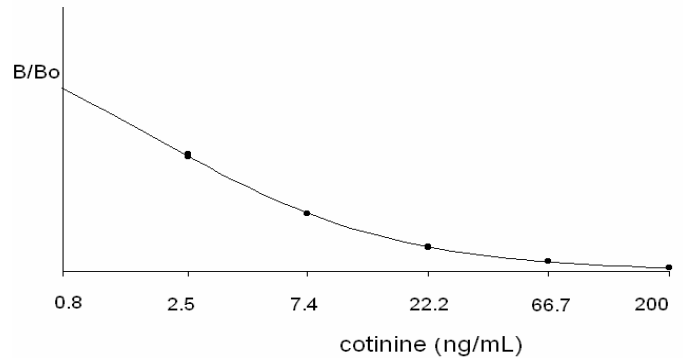
- Compute the average optical density (OD) for all duplicate wells.
- Subtract the average OD for the NSB wells from the average OD of the zero, standards, controls and unknowns.
- Calculate the percent bound (B/Bo) for each standard, control and unknown by dividing the average OD (B) by the average OD for the zero (Bo).
- Determine the concentrations of the controls and unknowns by interpolation using software capable of logistics. We recommend a 4-parameter curve fit.

Typical Results

The results shown below are for illustration only and should not be used to calculate results from another assay.

Well	Sample	Average OD	B	B/Bo	Cotinine
A1,A2	S1	0.045	0.037	0.022	200
B1,B2	S2	0.096	0.088	0.051	66.7
C1,C2	S3	0.222	0.214	0.125	22.2
D1,D2	S4	0.489	0.481	0.280	7.4
E1,E2	S5	0.893	0.885	0.516	2.5
F1,F2	S6	1.278	1.270	0.740	0.8
G1,G2	Bo	1.724	1.716	NA	NA
H1,H2	NSB	0.008	NA	NA	NA

Cotinine 4-Parameter Curve Fit



Material Safety Data*

Hazardous Ingredients

Stop solution is a 2-molar solution of sulfuric acid. This solution is caustic; use with care.

We recommend the procedures listed below for all kit reagents.

Handling

Follow good laboratory procedures when handling kit reagents. Laboratory coats, gloves, and safety goggles are recommended. Wipe up spills using standard absorbent materials while wearing protective clothing. Follow local regulations for disposal.

Emergency Exposure Measures

In case of contact, immediately wash skin or flush eyes with water for 15 minutes. Remove contaminated clothing. If inhaled, remove individual to fresh air. If individual experiences difficulty breathing, give oxygen and call a physician.

*The above information is believed to be accurate but is not all-inclusive. This information should only be used as a guide. Salimetrics shall not be liable for accidents or damage resulting from contact with reagents.

Performance Characteristics

A. Recovery:

Eight saliva samples were spiked with known quantities of cotinine and assayed.

Sample	Endogenous (ng/mL)	Added (ng/mL)	Expected (ng/mL)	Observed (ng/mL)	Recovery (%)
1	0	100	100	87.61	87.6
1	0	10	10	10.64	106.4
2	0	30	30	28.26	94.2
2	0	3.75	3.75	3.47	92.5
3	594.90	30	624.90	602.70	96.4
4	627.95	3.75	631.70	661.95	104.8
5	267.39	30	297.39	290.40	97.6
6	68.46	1.00	69.46	60.11	86.5
7	188.03	500	688.03	612.17	89.0
8	100.44	500	600.44	538.24	89.6

B. Precision:

The intra-assay precision was determined from 28 samples of high and 12 samples of low levels of cotinine.

Sample	N	Mean (ng/mL)	Std Dev (ng/mL)	Coefficient of Variation (%)
Low	12	5.91	0.13	2.3
High	28	92.67	5.41	5.8

The inter-assay precision was determined from the mean of average duplicates for 12 separate runs.

Sample	N	Mean (ng/mL)	Std Dev (ng/mL)	Coefficient of Variation (%)
Low	12	5.09	0.49	8.2
High	12	101.96	5.06	5.0

C. Sensitivity:

The lower limit of sensitivity was determined by interpolating the mean minus 2 SD's for 20 zero values. The minimal concentration of cotinine that can be distinguished from zero is 0.05 ng/mL.

D. Linearity of Dilution:

Two saliva samples were diluted with assay diluent and assayed.

Sample	Dilution Factor	Expected (ng/mL)	Observed (ng/mL)	Recovery (%)
Sample 1			87.61	
	1:2	43.81	43.13	98.4
	1:4	21.90	21.99	100.4
	1:8	10.95	12.30	112.3
	1:16	5.48	6.10	111.3
	1:32	2.73	2.99	109.5
	1:64	1.37	1.50	109.5
Sample 2			586.39	
	1:2	293.20	291.84	99.5
	1:4	146.60	148.91	101.6
	1:8	73.30	73.63	100.5
	1:16	36.65	38.56	105.2
	1:32	18.32	19.14	104.5
	1:64	9.16	10.29	112.3

E. Cross-Reactivity

Nicotine	1.4%	Nicotinamide	0.015%
Nicotinic acid	ND	3-OH-cotinine	54.86 %

F. Measurement of Salivary Cotinine in Smokers and Non-smokers Using the Salimetrics EIA

Group	N	Mean (ng/mL)	Std Dev (ng/mL)	Range (ng/mL)
Adult Smokers	21	206.33	123.47	47.87 - 586.39
Non-smokers	10	0	0	NA

The Salimetrics EIA is able to distinguish smokers from non-smokers with 100% accuracy.

G. Salimetrics quantitative enzyme immunoassay for salivary cotinine discriminates smokers from non-smokers, and differentiates primary from secondary smoke exposure (7)

Table 1: Salivary cotinine levels (ng/mL) in smoking and non-smoking mothers and their 6-month old infants

Mother's Self-Reported Status: Smokers (n=27)			
Group	Mean	Std Dev	Ranges
Number of cigarettes smoked in prior 48 hours by mother	14.26	14.67	3 – 60
Mother's salivary cotinine (ng/mL)	241.27	176.67	0 – 565.36
Infant salivary cotinine (ng/mL)	9.59	8.58	0.91 – 33.19

Mother's Self-Reported Status: Non-smokers (n=20)			
Group	Mean	Std Dev	Ranges
Number of cigarettes smoked in prior 48 hours by mother	0.0	NA	NA
Mother's salivary cotinine (ng/mL)	2.15	5.45	0 - 23.9
Infant salivary cotinine (ng/mL)	2.12	3.54	0 - 12.34

Notes:

- Smoking status determined by number of cigarettes smoked in the past 48 hours, "0" = non-smoker, "> 3" = smoker.
- Independent sample t-test comparing smoking and non-smoking groups, $p < 0.001$.
- Only 1 infant of a smoking mother had received breastmilk in prior 7 days.

Figure 1. Positive association between mother and infant salivary cotinine

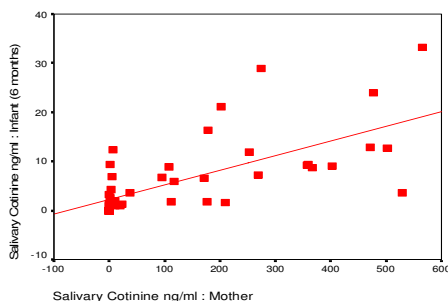
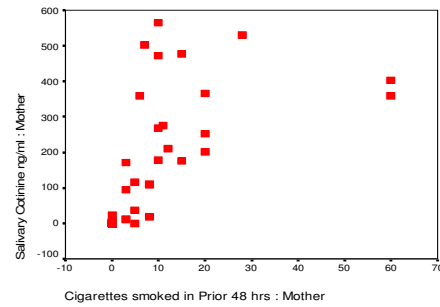


Figure 2. Positive association between number of cigarettes smoked in the prior 48 hours and salivary cotinine in mothers



G. Comparison of Cotinine Measurement by LC-ES/MS/MS to EIA

(n=40)	Cotinine Results by LC-ES/MS/MS*	Cotinine Results by Salimetrics EIA
LC-ES/MS/MS		
Pearson Correlation (p-value)	--	0.901 (0.000)
Salimetrics Cotinine EIA		
Pearson Correlation (p-value)	0.901 (0.000)	--
Total Hrs. Exposed[^]		
Pearson Correlation (p-value)	0.394 (0.012)	0.484 (0.002)

*Liquid Chromatography Electrospray Ionization Tandem Mass Spectroscopy

[^] Self-reported hours of exposure to secondhand smoke

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Seller's Limited Warranty

"Seller warrants that all goods sold hereunder will be free from defects in material and workmanship. Upon prompt notice by Buyer of any claimed defect, which notice must be sent within thirty (30) days from date such defect is first discovered and within three months from the date of shipment, Seller shall, at its option, either repair or replace the product that is proved to Seller's satisfaction to be defective. All claims should be submitted in written form. This warranty does not cover any damage due to accident, misuse, negligence, or abnormal use. Liability, in all cases, will be limited to the purchased cost of the kit.

It is expressly agreed that this limited warranty shall be in lieu of all warranties of fitness and in lieu of the warranty of merchantability. Seller shall not be liable for any incidental or consequential damages that arise out of the installation, use or operation of Seller's product or out of the breach of any express or implied warranties."