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Estrone Saliva LUM

REF IB57401

Effective: February 10, 2011

INTENDED USE For the determination of estrone in human saliva by a chemiluminescence immunoassay (LUM). For Research Use Only. Not for Use in Diagnostic Procedures.

PRINCIPLE OF THE TEST

The principle of the following chemiluminescence immunoassay (LUM) test follows a two-step competitive binding scenario. Competition occurs between an unlabeled antigen (present in calibrators, control and unknowns) and a biotin-labelled antigen for a limited number of anti-estrone antibody binding sites on the microwell plate. After washing the streptavidin-horseradish peroxidase conjugate is incubated and forms a complex with the streptavidin-norseradish peroxidase conjugate is incubated and forms a complex with the bound biotinylated estrone. The washing and decanting procedures remove unbound materials, and then luminescence substrate solution is added. The relative luminescence units (RLUs) are measured on a microtiter plate luminometer. The RLU values are inversely proportional to the concentration of estrone in the unknown. A set of calibrators is used to plot a calibration curve from which the amount of estrone in saliva unknowns and controls can be directly read.

PROCEDURAL CAUTIONS AND WARNINGS 1. Users should have a thorough understanding of this protocol for the successful use of this kit. Reliable performance will only be attained by strict and careful adherence to the instructions provided.

Control materials or saliva pools should be included in every run at a high and low level for assessing the reliability of results.
 When the use of water is specified for dilution or reconstitution, use deionized or

distilled water.
4. In order to reduce exposure to potentially harmful substances, gloves should be worn when handling kit reagents and human unknowns.
5. All kit reagents and unknowns should be brought to room temperature and mixed gently but thoroughly before use. Avoid repeated freezing and thawing of reagents and

unknowns

6. A calibrator curve must be established for every run.
7. The kit controls should be included in every run and fall within established confidence limits.

Improper procedural techniques, imprecise pipetting, incomplete washing as well as improper reagent storage may be indicated when assay values for the controls do not reflect established ranges.
 The luminescence substrate solutions (A and B) are sensitive to light and should be

stored in the original dark bottle away from direct sunlight.

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10. When dispensing the substrate, do not use pipettes in which these liquids will come into contact with any metal parts.
11. To prevent contamination of reagents, use a new disposable pipette tip for dispensing each reagent, unknown, calibrator and control.
12. Do not mix various lot numbers of kit components within a test and do not use any

14. Kit reagents must be regarded as hazardous waste and disposed of according to national regulations.

LIMITATIONS

All the reagents within the kit are calibrated for the determination of estrone in human saliva. The kit is not calibrated for the determination of estrone in serum, plasma or other unknowns of human or animal origin.
 Any unknowns or control sera containing azide or thimerosal are not compatible with this kit, as they may lead to false results.

this kit, as they may lead to false results. 3. Only calibrator A may be used to dilute any high saliva unknowns. The use of any other reagent may lead to false results. 4. The occurrence of heterophilic antibodies in subjects regularly exposed to animals or animal products has the potential of causing interferences in immunological tests. Consequently, all aspects of a subject's background, including the frequency of exposure to animals/products, should be considered if false results are suspected.

SAFETY CAUTIONS AND WARNINGS POTENTIAL BIOHAZARDOUS MATERIAL

Human fluids that may be used in the preparation of the calibrators and controls have been tested and found to be non-reactive for Hepatitis B surface antigen and have also been tested for the presence of antibodies to HCV and Human Immunodeficiency Virus (HIV) and found to be negative. However no test method can offer complete assurance that HIV, HCV and Hepatitis B virus or any infectious agents are absent. The reagents should be considered a potential biohazard and handled with the same precautions as applied to any human unknown.

CHEMICAL HAZARDS

Avoid direct contact with reagents. In case of contact, wash with plenty of water

COLLECTION AND HANDLING OF UNKNOWNS

Approximately 1 mL of saliva is required per duplicate determination. Collect 2-3 mL of saliva into a clean glass tube without force or inducement and before eating, drinking or brushing the teeth. Simply rinse the mouth with water before collection and wait a few minutes to start. Do not use blood-contaminated unknowns. Store unknowns at 4°C for up to 24 hours or at -10°C or lower if the analyses are to be done at a later date. Consider all human unknowns as possible biohazardous materials and take appropriate precautions when handling.

PRETREATMENT OF UNKNOWNS

1.Unknowns are to be centrifuged. The supernatants are to be transferred into

clean tubes. 2. The tubes containing the supernatant are to be placed in a water bath and heated

at 60-70°C for 1 hour, 3. Allow heated unknowns to reach room temperature before assaying.

REAGENTS AND EQUIPMENT NEEDED BUT NOT PROVIDED 1. Precision pipettes to dispense 100, 150 and 300 μL 2. Disposable pipette tips 3. Distilled or deionized water

4. Plate shaker

5. Microwell plate luminometer

REAGENTS PROVIDED AND PREPARATION

1. Rabbit Anti-Estrone Antibody Coated Microwell Plate-Break Apart Wells - Ready To Use.

Contents: One 96 well (12x8) polyclonal antibody-coated microwell plate in a resealable pouch with desiccant. Storage: Refrigerate at 2-8°C Stability: 12 months or as indicated on label.

2. Estrone-Biotin Conjugate Concentrate - X50

Contents: Estrone-biotin conjugate in a protein-based buffer with a non-mercury preservative

preservative. Volume: 0.3 mL/vial Storage: Refrigerate at 2-8°C Stability: 12 months or as indicated on label. Preparation of conjugate working solution: Dilute biotin conjugate concentrate 1:50 in biotin conjugate buffer before use (example: 20 µL of biotin conjugate concentrate in 1 mL of biotin conjugate buffer). If the whole plate is to be used dilute 240 µL of biotin conjugate concentrate in 12 mL of biotin conjugate buffer. Discard any that is left over.

3. Streptavidin-Horse Radish Peroxidase (HRP) Conjugate Concentrate - X50

Contents: Streptavidin-HRP conjugate in a protein-based buffer with a non-mercury

Contents: Streptavidin-HRP conjugate in a protein-based buffer with a non-mercury preservative. Volume: 0.4 mL/vial Storage: Refrigerate at 2-8°C Stability: 12 months or as indicated on label. Preparation of conjugate working solution: Dilute HRP conjugate concentrate 1:50 in HRP conjugate buffer before use (example: 20 µL of HRP conjugate concentrate in 1 mL of HRP conjugate buffer). If the whole plate is to be used dilute 320 µL of HRP conjugate concentrate in 16 mL of HRP conjugate concentrate in 16 mL of HRP conjugate buffer. Discard any that is left over.

4. Estrone Saliva Calibrators - Ready To Use. Contents: Six vials containing estrone in a protein-based buffer with a non-mercury preservative. Prepared by spiking buffer with a precise quantity of estrone. *Listed below are approximate concentrations, please refer to vial labels for exact concentrations.

Concentration	Volume/Vial
0 pg/mL	4.0 mL
2 pg/mL	1.0 mL
8 pg/mL	1.0 mL
25 pg/mL	1.0 mL
100 pg/mL	1.0 mL
300 pg/mL	1.0 mL
	0 pg/mL 2 pg/mL 8 pg/mL 25 pg/mL 100 pg/mL

Storage: Refrigerate at 2-8°C Stability: 12 months or as indicated on label.

5. Controls - Ready To Use.

Contents: Two vials containing estrone in a protein-based buffer with a non-mercury preservative. Refer to vial labels for acceptable range.

, Volume: 1 mL/vial

Storage: Refrigerate at 2-8°C Stability: 12 months or as indicated on label.

6. Biotin Conjugate Buffer - Ready To Use.

Volume: 13 mL/bottle Storage: Refrigerate at 2-8°C Stability: 12 months or as indicated on label.

7. HRP Conjugate Buffer - Ready To Use.

Contents: One bottle containing a protein-based buffer with a non-mercury preservative. Volume: 20 mL/bottle Storage: Refrigerate at 2-8°C Stability: 12 months or as indicated on label.

8. Wash Buffer Concentrate - X10 Contents: One bottle containing buffer with a non-ionic detergent and a non-mercury

preservative. Volume: 50 mL/bottle

Volume: 50 mL/bottle Storage: Refrigerate at 2-8°C Stability: 12 months or as indicated on label. Preparation of wash buffer working solution: Dilute wash buffer concentrate 1:10 in distilled or deionized water before use. If the whole plate is to be used dilute 50 mL of wash buffer concentrate in 450 mL of water.

9. Rinse Solution - Ready To Use.

Contents: Two bottles containing buffer with a non-ionic detergent and a non-mercury preservative.

Volume: 2 x 50 mL/bottle

Storage: Refrigerate at 2-8°C Stability: 12 months or as indicated on label.

10. LUM Substrate Reagent A - Requires Preparation.

Contents: One vial containing luminol enhancer. Volume: 0.8 mL/vial

Storage: Refrigerate at 2-8°C Stability: 12 months or as indicated on label. Preparation: See preparation of LUM working substrate solution.

11. LUM Substrate Reagent B - Requires Preparation.

Volume: 1.6 mL/vial Storage: Refrigerate at 2-8°C Stability: 12 months or as indicated on label. Preparation: See preparation of LUM working substrate solution.

12. LUM Substrate Reagent C - Requires Preparation.

Contents: One bottle containing buffer with a non-mercury preservative. Volume: 16 mL/bottle Storage: Refrigerate at 2-8°C Stability: 12 months or as indicated on label. Preparation: See preparation of LUM working substrate solution.

13. Preparation of LUM Working Substrate Solution In a clean container mix 1 part of LUM substrate reagent A with 2 parts of LUM substrate reagent B and 20 parts of LUM substrate reagent C. This gives the ready to use substrate solution. If the whole plate is to be used prepare working substrate solution as follows: Combine 0.6 mL of LUM substrate reagent A with 1.2 mL of LUM substrate reagent B and 12 mL of LUM substrate reagent C. It is suggested to wait at least 30 minutes prior to use after preparation of the working substrate solution. The working substrate solution is stable for up to 8 hours at room temperature. Discard the leftovers.

CE - European Conformity	- Consult Ins	structions For Use	- In Vitro Diagnostic Use	LOT - Lot Number	Temperature Limit See By
- Manufacturer	REF - Catalogue Number	X# - Dilute 1:# Before Use	+ # mL- Recor	stitute With # mL H_2O	EC REP - Authorized Representative

ASSAY PROCEDURE

Important Notes:

- 1. All reagents must reach room temperature before use.
- Once the procedure has been started, all steps should be 2. completed without interruption to ensure equal elapsed time for each pipetting step.
- 1. Prepare working solutions of both conjugates, wash buffer and LUM substrate (refer to reagents provided and preparation section).
- 2 Remove the required number of microwell strips. Reseal the bag and return any unused strips to the refrigerator.
- Pipette 100 µL of each calibrator, control and pretreated 3. unknown (refer to pretreatment of unknowns section) into correspondingly labelled wells in duplicate.
- 4. Pipette 100 µL of the estrone-biotin conjugate working solution into each well (We recommend using a multichannel pipette).
- Cover the plate and incubate for 1 hour on a plate shaker 5. (approximately 200 rpm) at room temperature.
- 6. Wash the wells 3 times with 300 µL of wash buffer working solution per well and tap the plate firmly against absorbent paper to ensure that it is dry (The use of a washer is recommended).
- 7. Pipette 150 µL of the streptavidin-HRP conjugate working solution into each well (We recommend using a multichannel pipette).
- 8. Cover the plate and incubate for 30 minutes on a plate shaker (approximately 200 rpm) at room temperature.
- 9. Wash the wells 3 times in the same manner as step 6.
- 10. Rinse the well 3 times with 300 µL/well of Rinse Solution and tap the plate firmly against absorbent paper to ensure that it is dry
- 10. Pipette 130 μL of LUM working substrate solution into each well (We recommend using a multichannel pipette).
- 11. Measure the RLU/second in each well on a microplate luminometer between 5-15 minutes after addition of the substrate.

RESULTS

- RESULTS
 Calculate the mean RLU of each calibrator duplicate.
 Draw a calibrator curve on semi-log paper with the mean RLUs on the Y-axis and the calibrator concentrations on the X-axis. If immunoassay software is being used, a 4-parameter curve is recommended.
 Calculate the mean RLU of each unknown duplicate.
 Read the values of the unknowns directly off the calibrator curve.
 If an unknown equation area to a 200 projet then different in with calibrator A at a dilution of the calibrator curve.

- 5. If an unknown reads more than 800 pg/mL then dilute it with calibrator A at a dilution of no more than 1:8. The result obtained should be multiplied by the dilution factor.

PERFORMANCE CHARACTERISTICS

SPECIFICITY (CROSS-REACTIVITY)

The following compounds were tested for cross-reactivity with the IBL-America estrone saliva LUM kit with estrone cross-reacting at 100%:

Steroid	%Cross-Reactivity
Estrone	100
Estrone-3-Sulfate	4.9
17β-Estradiol	2.2
Estrone-3-Glucuronide	1.2
178-Estradiol-3-Glucuronide	0.14

The following steroids were tested but cross-reacted at less than 0.1%: Androstenedione Cholesterol, Corticosterone, Cortisol, Cortisone, DHEAS, Diethylstilbesterone, Estriol, 17ß-Estradiol-3-Glucuronide, Estradiol-Sulfate, Progesterone, 17-OH Progesterone and Testosterone

RECOVERY

Spiked unknowns were prepared by adding accurate amounts of estrone to three saliva unknowns. The results (in pg/mL) are tabulated below:

Unknown	Obs.Result	Exp.Result	Recovery%
1 Unspiked	5.21	-	-
+ 50(5:1)	15.89	14.2	114
+200(5:1)	50.82	44.0	116
+800(5:1)	154.48	164.0	94
2 Unspiked	41.80	-	-
+ 50(5:1)	34.86	36.0	97
+200(5:1)	57.39	66.4	86
+800(5:1)	165.73	186.0	89
3 Unspiked	52.57	-	-
+ 50(5:1)	53.67	52.08	97
+200(5:1)	85.77	83.00	97
+800(5:1)	179.00	203.00	89

COMPARATIVE STUDY

The estrone assay results of 20 human saliva unknowns obtained with the IBL-America direct LUM method were compared with the same method but included an ether-extraction

Unknown	Direct	Extracted
1	12.4	8.6
2	6.5	6.6
3	17.5	12.2
4	22.1	17.7
5	36.9	28.4
6	44.3	31.1
7	2.8	5.4
8	4.5	6.3
9	12.3	8.3
10	11.5	8.7
11	9.3	6.4
12	66.2	45.6
13	22.1	18.3
14	13.5	10.2
15	4.7	4.1
16	7.5	5.5
17	8.7	5.7
18	11.4	9.8
19	10.5	8.5
20	8.9	12.5

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