



11 β -PROSTAGLANDIN F_{2 α} *ELISA Kit Instructions*

Please read all instructions carefully before beginning this assay

PRODUCT #404410
For Research Use Only

*****Store kit at 4°C at all times*****

Do not freeze kit components

DESCRIPTION

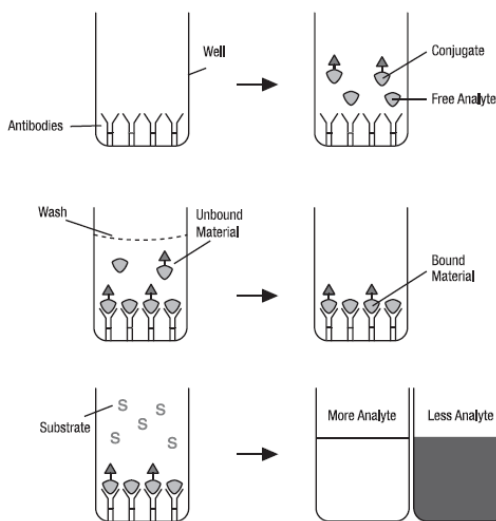
11 β -Prostaglandin F_{2 α} (11 β -PGF_{2 α}) is derived from PGD₂ by the action of 11-ketoreductase. It is a potent bronchoconstrictor agent. It also causes hypertension in animals. Plasma levels of 11 β -PGF_{2 α} are highly elevated in patients with systemic mastocytosis with an overproduction of PGD₂.

PRINCIPLE OF ASSAY

This is an ELISA (Enzyme-Linked ImmunoSorbent Assay) for the quantitative analysis of 11 β -Prostaglandin F_{2 α} levels in biological fluid. This test kit operates on the basis of competition between the enzyme conjugate and the 11 β -PGF_{2 α} in the sample for a limited number of binding sites on the antibody coated plate.

The sample or standard solution is first added to the microplate. Next, the diluted enzyme conjugate is added and the mixture is shaken and incubated at room temperature for one hour. During the incubation, competition for binding sites is taking place. The plate is then washed removing all the unbound material. The bound enzyme conjugate is detected by the addition of substrate which generates an optimal color after 30 minutes. Quantitative test results may be obtained by measuring and comparing the absorbance reading of the wells of the samples against the standards with a microplate reader at 650 nm. The extent of color development is inversely proportional to the amount of 11 β -PGF_{2 α} in the sample or standard. For example, the absence of 11 β -PGF_{2 α} in the sample will result in a bright blue color, whereas the presence of 11 β -PGF_{2 α} will result in decreased or no color development.

PRINCIPLE OF ASSAY (continued)



MATERIALS PROVIDED

1. **EIA BUFFER:** 30 mL. Provided to dilute enzyme conjugate and 11β -PGF_{2 α} standards.
2. **WASH BUFFER (10X):** 20 mL. Dilute 10 fold with deionized water. Diluted wash buffer is used to wash all unbound enzyme conjugate, samples and standards from the plate after the one hour incubation.
3. **K-BLUE SUBSTRATE:** 20 mL. Stabilized 3,3', 5,5' Tetramethylbenzidine (TMB) plus Hydrogen Peroxide (H₂O₂) in a single bottle. It is used to develop the color in the wells after they have been washed. Keep refrigerated. **LIGHT SENSITIVE.**
4. **EXTRACTION BUFFER (5X):** 30 mL. Dilute 5 fold with deionized water. This is used for diluting extracted and non-extracted samples.
5. **11β -PROSTAGLANDIN F_{2 α} ENZYME CONJUGATE:** 150 μ L. 11β -PGF_{2 α} horseradish peroxidase concentrate. Blue capped vial.
6. **11β -PROSTAGLANDIN F_{2 α} STANDARD:** 100 μ L. 11β -PGF_{2 α} standard at the concentration of 1 μ g/mL. Green capped vial.
7. **11β -PROSTAGLANDIN F_{2 α} ANTIBODY-COATED MICROPLATE:** A 96 well MaxiSorp™ Nunc microplate with anti- 11β -PGF_{2 α} rabbit antibody precoated on each well. The plate is ready for use as is. **DO NOT WASH!**

MATERIALS NEEDED BUT NOT PROVIDED

1. 300 mL deionized water for diluting wash buffer and extraction buffer.
 2. Precision pipettes that range from 10 μ L - 1000 μ L and disposable tips.
- NOTE: If all or several strips are to be used at one time, it is suggested that a multichannel pipette be used.**
3. Clean test tubes used to dilute the standards and conjugate.
 4. Graduated cylinders to dilute and mix wash buffer and extraction buffer.
 5. Microplate reader with 650 nm filter.
 6. Plate cover or plastic film to cover plate during incubation.

OPTIONAL MATERIALS

7. 1 N HCl or Neogen's Red Stop Solution.
8. Microplate shaker.
If performing an extraction on samples, the following will be required:
9. Methanol
10. Methyl formate
11. 0.1 M Sodium Phosphate buffer, pH 7.5
12. C₁₈ Sep-Pak® column (Waters® Corporation)
13. Petroleum ether
14. Nitrogen gas
15. Vortex
16. Centrifuge

WARNINGS AND PRECAUTIONS

1. **DO NOT** use components beyond expiration date.
2. **DO NOT** mix any reagents or components of this kit with any reagents or components of any other kit. This kit is designed to work properly as provided.
3. **DO NOT** pipette reagents by mouth.
4. Always pour substrate out of the bottle into a clean test tube - **DO NOT** pipette out of the bottle. If pipette tip is unclean this could contaminate the substrate.
5. All specimens should be considered potentially infectious. Exercise proper handling precautions.
6. **DO NOT** smoke, eat or drink in areas where specimens or reagents are being handled.
7. Use aseptic technique when opening and removing reagents from vials and bottles.
8. Keep plate covered except when adding reagents, washing or reading.
9. Kit components should be refrigerated at all times when not in use.

PROCEDURAL NOTES

1. It is not necessary to allow reagents to warm to room temperature before use.
2. Desiccant bag must remain in foil pouch with unused strips. Keep zip-lock pouch sealed when not in use to maintain a dry environment.
3. Always use new pipette tips for the buffer, enzyme conjugate, standards and samples.
4. Before pipetting a reagent, rinse the pipette tip three times with that reagent (i.e. fill the tip with the desired amount of reagent and dispense back into the same vial - repeat 2 times). Now the tip is properly rinsed and ready to dispense the reagent into your well or test tube.
5. When pipetting into the wells, **DO NOT** allow the pipette tip to touch the inside of the well, or any of the reagents already in the well - this can result in cross contamination.
6. Standards and samples should be assayed in duplicate.
7. To quantitate, always run samples alongside a standard curve. If testing a sample that is not extracted, standards should be diluted in the same type of medium being tested. This medium should be known to be negative.
8. Gently mix specimens and reagents before use. Avoid vigorous agitation.
9. When using only partial amounts of a kit, it is recommended to transfer the appropriate volume of each reagent to a clean vessel for repeated dispensing. This will reduce reagent contamination by repeated sampling from the original container.
10. The enzyme conjugate is most stable in its concentrated form. Dilute only the volume necessary for the amount of strips currently being used.
11. Before taking an absorbance reading, wipe the outside bottom of the wells with a lint-free wiper to remove dust and fingerprints.
12. Before opening the enzyme conjugate and standard vial, tap vial in an upright position to remove any liquid in the cap.

SAMPLE PREPARATION

This assay is non-species specific. Usually, urine and tissue culture supernatant can be assayed directly by diluting them with the diluted extraction buffer. Plasma and most other mediums will need to be extracted.

EXTRACTION OF 11 β -PGF_{2 α}

1. Add 0.2 mL of methanol to 1 mL of biological fluid and vortex.
2. For tissue, homogenize it in 15% methanol in 0.1 M sodium phosphate buffer, pH 7.5 (100 mg in 1 mL methanol-buffer). Centrifuge the homogenate for five (5) minutes. Collect the supernatant in a clean tube.
3. Precondition the C₁₈ Sep-Pak® column (Waters® Corporation) by washing the column with 2 mL of methanol followed by 2 mL of water.
4. Apply the above sample into the column and adjust the flow rate to 1 mL per minute.
5. Wash the column with 2 mL of 15% methanol in water followed by 2 mL of petroleum ether.
6. The Prostaglandin is eluted by 2 mL of methyl formate.
7. Evaporate methyl formate eluate with a stream of nitrogen gas.
8. Dissolve the residue with 1 mL of diluted extraction buffer and assay 50 μ L in duplicates.
9. If the concentration is higher than the high range of the standard curve, the samples in # 8 need to be further diluted and assayed.

NOTE: Extraction buffer must be diluted 5 fold with deionized water before use. Any precipitant present must be brought into solution before dilution.

TEST PROCEDURES

1. Prepare standards as follows:

Standard	Preparation
A	stock solution 1 μ g/mL (provided in green capped vial.)
B	take 20 μ L of A, add to 980 μ L of EIA buffer and mix=20 ng/mL
C	take 200 μ L of B, add to 1.8 mL of EIA buffer and mix=2 ng/mL
D	take 200 μ L of C, add to 1.8 mL of EIA buffer and mix=0.2 ng/mL

Continue standard preparation following Scheme I.

Scheme I

Standards	ng/mL	EIA buffer (μL added)	C standard μL	D standard μL
S ₀	0	as is	-	-
S ₁	0.01	950	-	50
S ₂	0.02	900	-	100
S ₃	0.04	800	-	200
S ₄	0.1	500	-	500
S ₅	0.2	-	-	as is
S ₆	0.4	800	200	-
S ₇	1	500	500	-

2. Determine the number of wells to be used.
3. Dilute the 11β -PGF_{2 α} enzyme conjugate. Add 1 μ L of enzyme conjugate into 50 μ L total volume of EIA buffer for each well assayed. For the whole plate, add 110 μ L of the enzyme conjugate into 5.5 mL total volume of EIA buffer. Mix the solution thoroughly.
4. Add 50 μ L or standards (S) or unknown (U) (some samples may require diluting) to the appropriate wells in duplicate.

See Scheme II for suggested template design.

5. Add 50 μ L of the diluted enzyme conjugate to each well. Use 8-channel pipette or 12-channel pipette for rapid addition.
6. Mix by shaking plate gently. A microplate shaker may be used.
7. Cover plate with plastic film or plate cover and incubate at room temperature for one hour.

NOTE: Keep plate away from drafts and temperature fluctuations.

8. Dilute concentrated wash buffer with deionized water (i.e. 20 mL of wash buffer plus 180 mL of deionized water). Mix thoroughly.
9. After incubation, dump out the contents of the plate. Tap out contents thoroughly on a clean lint-free towel.
10. Wash each well with 300 μ L of the diluted wash buffer. Repeat for a total of three washings. An automated plate washer can be used, however, increase wash cycles from three to five.
11. Add 150 μ L of substrate to each well. Use multichannel pipette for best results. Mix by shaking plate gently.
12. Incubate at room temperature for 30 minutes.
13. Gently shake plate before taking a reading to ensure uniform color throughout each well.
14. Plate is read in a microplate reader at 650 nm. If a dual wavelength is used, set W₁ at 650 nm and W₂ at 490 nm.
15. If accounting for substrate background, use 2 to 8 wells as blanks with only substrate in the wells (150 μ L/well). Subtract the average of these absorbance values from the absorbance values of the wells being assayed.

NOTE: Some microplate readers can be programmed to do these subtractions automatically when reading the plate. Consult your instrument manual.

OPTIONAL TEST PROCEDURES

16. Add 50-100 μ L of 1 N HCl or Neogen's Red Stop Solution to each well to stop enzyme reaction.
17. Read plate at 450 nm, if 1N HCl solution was used. Read plate at 650 nm, if Neogen's Red Stop Solution was used.
18. Plot the standard curve and estimate the concentrations of the samples from the curve.
See "CALCULATIONS."

NOTE: Absorbance readings will approximately double when stopped with acid. If absorbance readings are too high for measuring with your microplate reader, decrease the substrate incubation approximately 10 minutes but no more than 15 minutes.

Scheme II

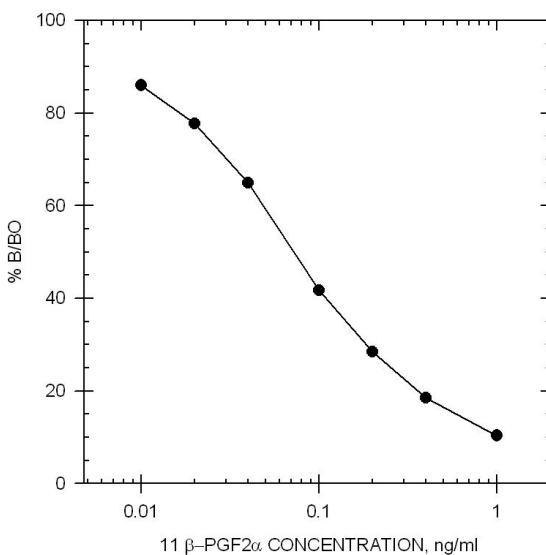
	1	2	3	4	5	6	7	8	9	10	11	12
A	^s 0	^s 0	^u 1	^u 1	^u 9	^u 9	^u 17	^u 17	^u 25	^u 25	^u 33	^u 33
B	^s 1	^s 1	^u 2	^u 2	^u 10	^u 10	^u 18	^u 18	^u 26	^u 26	^u 34	^u 34
C	^s 2	^s 2	^u 3	^u 3	^u 11	^u 11	^u 19	^u 19	^u 27	^u 27	^u 35	^u 35
D	^s 3	^s 3	^u 4	^u 4	^u 12	^u 12	^u 20	^u 20	^u 28	^u 28	^u 36	^u 36
E	^s 4	^s 4	^u 5	^u 5	^u 13	^u 13	^u 21	^u 21	^u 29	^u 29	^u 37	^u 37
F	^s 5	^s 5	^u 6	^u 6	^u 14	^u 14	^u 22	^u 22	^u 30	^u 30	^u 38	^u 38
G	^s 6	^s 6	^u 7	^u 7	^u 15	^u 15	^u 23	^u 23	^u 31	^u 31	^u 39	^u 39
H	^s 7	^s 7	^u 8	^u 8	^u 16	^u 16	^u 24	^u 24	^u 32	^u 32	^u 40	^u 40

CALCULATIONS

1. After the substrate background has been subtracted from all absorbance values, average all of your duplicate well absorbance values.
2. The average of your two S_0 values is now your B_0 value. (S_1 now becomes B_1 , etc.)
3. Next, find the percent of maximal binding ($\%B/B_0$ value). To do this, divide the averages of each standard absorbance value (now known as B_1 through B_n) by the B_0 absorbance value and multiply by 100 to achieve percentages.
4. Graph your standard curve by plotting the $\%B/B_0$ for each standard concentration on the ordinate (y) axis against concentration on the abscissa (x) axis. Draw a curve by using a curve-fitting routine (i.e. 4-parameter or linear regression).
5. Divide the averages of each sample absorbance value by the B_0 value and multiply by 100 to achieve percentages.
6. Using the standard curve, the concentration of each sample can be determined by comparing the $\%B/B_0$ of each sample to the corresponding concentration of $11\beta\text{-PGF}_{2\alpha}$ standard.
7. If the samples were diluted, the concentration determined from the standard curve must be multiplied by the dilution factor. If samples were concentrated, the concentration determined from the standard curve must be divided by the concentration factor.

TYPICAL STANDARD CURVE

$11\beta\text{-Prostaglandin F}_{2\alpha}$ in EIA Buffer



TYPICAL DATA

NOTE: "Typical data" is a representation. Variances in data will occur. Optical density readings may fluctuate during the shelf-life of the kit, but the %B/B₀ should remain comparable. Measuring wavelength: 650 nm.

Standard	Standard Concentration (ng/mL)	Optical Density (Absorbance Value)	%B/B ₀
S ₀ (B ₀)	0	1.450	100
S ₁ (B ₁)	0.01	1.257	87
S ₂ (B ₂)	0.02	1.044	72
S ₃ (B ₃)	0.04	0.900	62
S ₄ (B ₄)	0.1	0.544	38
S ₅ (B ₅)	0.2	0.319	22
S ₆ (B ₆)	0.4	0.245	17
S ₇ (B ₇)	1	0.131	9

CROSS REACTIVITY

11β-PROSTAGLANDIN F _{2α}	100.0%
PROSTAGLANDIN F _{2α}	0.24%
PROSTAGLANDIN E ₂	0.21%
THROMBOXANE B ₂	0.21%
PROSTAGLANDIN D ₂	0.01%
LEUKOTRIENE B ₁	<0.01%
PROSTAGLANDIN A ₁	<0.01%
PROSTAGLANDIN A ₂	<0.01%
PROSTAGLANDIN B ₁	<0.01%
PROSTAGLANDIN B ₂	<0.01%
PROSTAGLANDIN F _{1α}	<0.01%
6-KETO-PROSTAGLANDIN E ₁	<0.01%
6-KETO-PROSTAGLANDIN F _{1α}	<0.01%
13,14-DIHYDRO-15-KETO-PROSTAGLANDIN F _{2α}	<0.01%
15-KETO-PROSTAGLANDIN F _{2α}	<0.01%
11-DEHYDRO-THROMBOXANE B ₂	<0.01%

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TECHNICAL ASSISTANCE

Technical assistance is available Monday-Friday, between 8:00 a.m. and 6:00 p.m. EST.



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