MANUAL

PROGESTERONE KITS

Part # 6FPROPEG & 6FPROPEH

Test size#: 500 tests (6FPROPEG) and 10,000 tests (6FPROPEH) - assay volume: 20 µL

Revision: #05 of September 2023

Store at: -16°C or below (6FPROPEG); -16°C or below (6FPROPEH)

For research use only. Not for use in diagnostic procedures.

ASSAY PRINCIPLE

This kit is intended for the simple and rapid quantification of Progesterone directly from cell supernatants or purified solutions and offers a fast alternative to ELISA.

The detection principle of this kit is based on HTRF® technology (Homogeneous Time-Resolved Fluorescence). As shown in Figure 1, Progesterone is detected in a competitive assay by using anti Progesterone antibody labeled with Europium cryptate (donor), and Progesterone labeled with d2 (acceptor).

When the dyes are in close proximity, the excitation of the donor with a light source (laser or flash lamp) triggers a Fluorescence Resonance Energy Transfer (FRET) towards the acceptor, which in turn fluoresces at a specific wavelength (665 nm). The Progesterone present in the sample competes with the binding between the two HTRF detection solutions and thereby prevents FRET from occurring. The specific signal is inversely proportional to the Progesterone concentration.

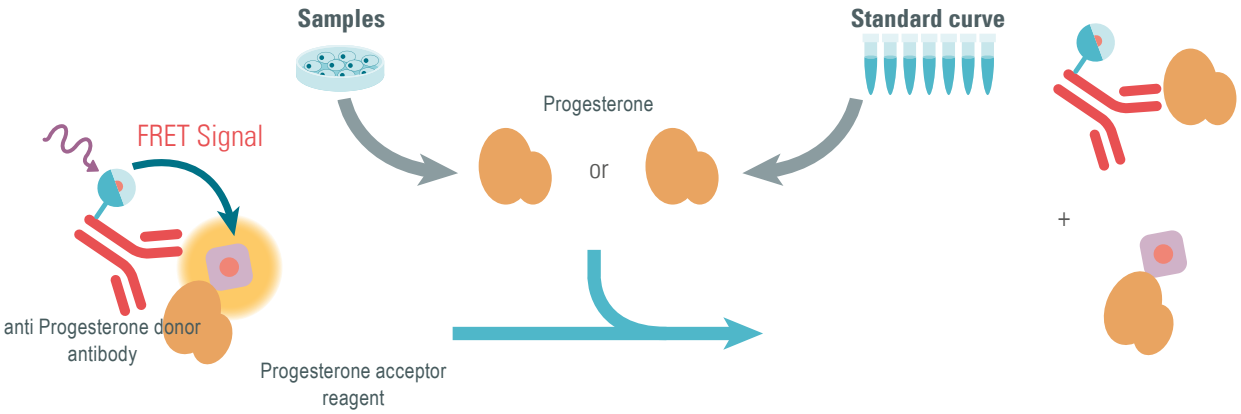
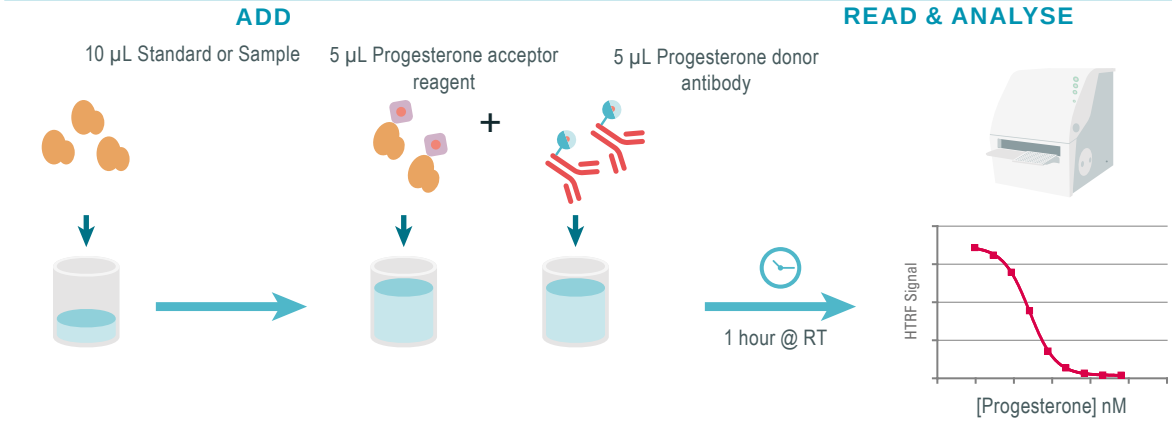


Figure 1: Principle of HTRF Progesterone competitive assay.

MNAUAL AT A GLANCE



Do not pre-mix the d2 and Cryptate solutions prior to dispensing.

Make sure to use the set-up for Eu Cryptate.

MATERIALS PROVIDED:

KIT COMPONENTS	500 TESTS * CAT # 6FPROPEG	10,000 TESTS * CAT # 6FPROPEH
Progesterone Standard Frozen - 100 X	1 vial - 10 µL 318 µM (100X)	1 vial - 10 µL 318 µM (100X)
anti Progesterone antibody Eu Cryptate antibody	1 vial - 50 µL Frozen - 50X	1 vial - 1 mL Frozen - 50X
Progesterone d2 reagent	1 vial - 50 µL Frozen - 50X	1 vial - 1 mL Frozen - 50X
Diluent ** ready-to-use	1 vial 20 mL	1 vial 20 mL
Detection buffer *** ready to use	1 vial 7 mL	1 vial 105 mL

* When used as advised, the two available kit sizes will provide sufficient reagents for 500 tests and 10,000 tests respectively in 20 µL final volume.. Assay volumes can be adjusted proportionally to run the assay in 96 or 1536 well microplates.
** Medium like cell culture medium can be an alternative to the diluent.
*** The Detection buffer is used to prepare working solutions of acceptor and donor reagents.

PURCHASE SEPARATELY:

- HTRF®-Certified Reader. Make sure the setup for Eu Cryptate is used.

For a list of HTRF-compatible readers and set-up recommendations, please visit www.revvity.com

- Small volume (SV) detection microplates - .

For more information about microplate recommendations, please visit our website at: www.revvity.com

STORAGE AND STABILITY

Store the kit at -16°C or below.

Under proper storage conditions, reagents are stable until the expiry date indicated on the label.Diluent and detection buffer are shipped frozen, but can be stored at 2-8°C in your premises.

If lyophilized, reconstituted reagents, antibodies, and standard stock solutions may be frozen and thawed only once. To avoid freeze/thaw cycles, it is recommended to dispense remaining stock solutions into disposable plastic vials for storage at -60°C or below .

REAGENT PREPARATION

BEFORE YOU BEGIN:

- It is very important to prepare reagents in the specified buffers. The use of an incorrect diluent may affect reagent stability and assay results.
- Before use, allow Diluent and Detection buffer to warm up at room temperature and homogenize them with a vortex.
- Thaw all reagents at room temperature, allow them to warm up.
- Antibody solutions must be prepared in individual vials and can be mixed prior to dispensing.
- Progesterone standards (for standard curve) must be prepared in diluent or in the same medium as the samples.

TAKE CARE TO PREPARE STOCK AND WORKING SOLUTIONS ACCORDING TO THE DIRECTIONS FOR THE KIT SIZE YOU HAVE PURCHASED.

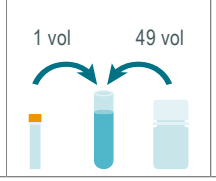
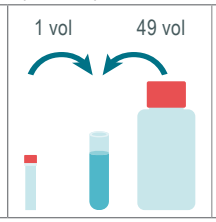
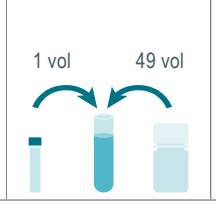
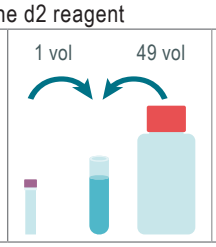
TO PREPARE REAGENT STOCK SOLUTIONS:

500 TESTS KIT - 6FPROPEG		10,000 TESTS KIT - 6FPROPEH	
anti Progesterone antibody Eu Cryptate antibody			
Thaw the anti Progesterone antibody Eu Cryptate antibody . Mix gently. This 50X stock solution can be frozen and stored at -60°C or below.			Thaw the anti Progesterone antibody Eu Cryptate antibody . Mix gently. This 50X stock solution can be frozen and stored at -60°C or below.
Progesterone d2 reagent			
Thaw the Progesterone d2 reagent . Mix gently. This 50X stock solution can be frozen and stored at -60°C or below.			Thaw the Progesterone d2 reagent . Mix gently. This 50X stock solution can be frozen and stored at -60°C or below.
Progesterone Standard			
Thaw the Progesterone standard solution in order to obtain a 318 µM (see vial label) stock solution. Mix gently this 100 X Progesterone standard stock solution. The progesterone standard stock solution is highly concentrated, in order to prepare the working standard solutions for the standard curve, we recommend to prepare a 1 X intermediate standard solution #A (3.18 µM). See section: TO PREPARE WORKING STANDARD SOLUTION p4.			Thaw the Progesterone standard solution in order to obtain a 318 µM (see vial label) stock solution. Mix gently this 100 X Progesterone standard stock solution. The progesterone standard stock solution is highly concentrated, in order to prepare the working standard solutions for the standard curve, we recommend to prepare a 1 X intermediate standard solution #A (3.18 µM). See section: TO PREPARE WORKING STANDARD SOLUTION p4.
Diluent			
The diluent is ready-to-use			The diluent is ready-to-use
Detection buffer			
The Detection buffer is ready-to-use.			The Detection buffer is ready-to-use.w

TO PREPARE ANTIBODY WORKING SOLUTIONS:

Each well requires 5 µL anti Progesterone antibody Eu Cryptate antibody and 5 µL Progesterone d2 reagent.

Prepare the two solutions in separate vials.

500 TESTS KIT - 6FPROPEG		10,000 TESTS KIT - 6FPROPEH	
anti Progesterone antibody Eu Cryptate antibody			
Dilute 50-fold the stock solution of Progesterone Eu Cryptate antibody with detection buffer#3 e.g. take 0.05 mL of Eu Cryptate antibody stock solution and add it to 2.45 mL of detection buffer #3.			Dilute 50-fold the stock solution of Progesterone Eu Cryptate antibody with detection buffer#3 e.g. take 1 mL of Eu Cryptate antibody stock solution and add it to 49 mL of detection buffer #3.
Progesterone d2 reagent			
Dilute 50-fold the stock solution of Progesterone d2 reagent with detection buffer: e.g. take 0.05 mL of d2 reagent stock solution and add it to 2.45 mL of detection buffer #3.			Dilute 50-fold the stock solution of Progesterone d2 reagent with detection buffer: e.g. take 1 mL of d2 reagent stock solution and add it to 49 mL of detection buffer #3.
Antibody mix			
Do not pre-mix the d2 and the Eu Cryptate solutions prior to dispensing.			

TO PREPARE STANDARD WORKING SOLUTIONS:

- Each well requires 10 µL of standard.
- Dilute the standard stock solution serially with diluent
- If the sample to test is a cell supernatant, replace the diluent by culture medium.
- In order to check for a potential interference effect from your own assay buffer when using the assay for the first time, we highly recommend the parallel preparation of a standard curve in your own supplemented cell culture medium and in diluent .
- In order to counteract any standard sticking, we recommend changing tips between each dilution.

A recommended standard dilution procedure is listed and illustrated below:

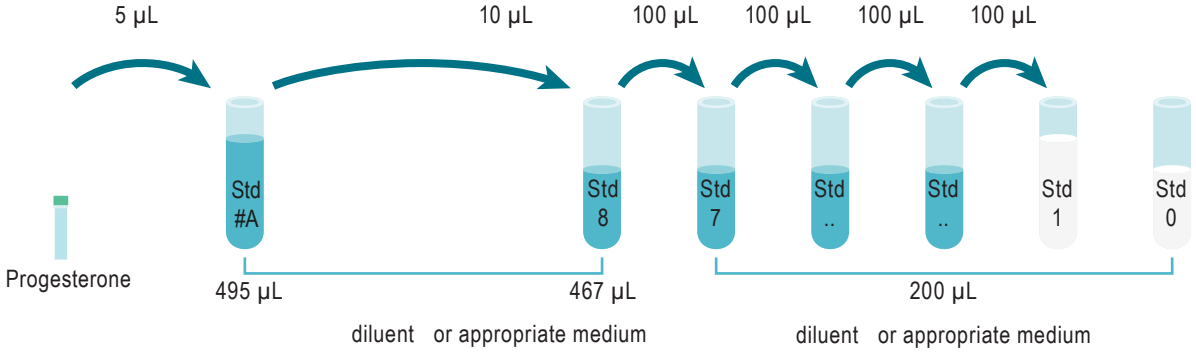
Dilute the standard stock solution 100-fold with diluent; this yields the Intermediate Standard Solution # A (3 180 nM). e.g: take 5 µL of standard stock solution and add it to 495 µL of diluent. Mix gently.

Dilute the intermediate Standard Solution #A 47.7-fold with diluent to prepare high standard (Std 8): e.g. take 10 µL of intermediate Standard Solution #A and add it to 467 µL of diluent . Mix gently.

Use the high standard (Std 8) to prepare the standard curve using 1/3 serial dilutions as follows:

- Dispense 200 µL of diluent in each vial from Std 7 to Std 0.
- Add 100 µL of standard to 200 µL of diluent , mix gently and repeat the 1/3 serial dilution to make standard solutions: std8, std7, std6, std5, std4, std3, std2, std1.

This will create 8 standards for the analyte. Std 0 (Positive control) is diluent or appropriate culture medium alone.



STANDARD	SERIAL DILUTIONS	PROGESTERONE WORKING SOLUTION (nM)
Standard Stock solution	Thawed stock solution	318,000
Intermediate standard solution #A	5µl Standard stock solution + 495µL Diluent	3,180
Standard 8	10 µL standard Stock solution + 467 µL Diluent	66.67
Standard 7	100 µL standard 8 + 200 µL Diluent	22.22
Standard 6	100 µL standard 7 + 200 µL Diluent	7.41
Standard 5	100 µL standard 6 + 200 µL Diluent	2.47
Standard 4	100 µL standard 5 + 200 µL Diluent	0.82
Standard 3	100 µL standard 4 + 200 µL Diluent	0.27
Standard 2	100 µL standard 3 + 200 µL Diluent	0.09
Standard 1	100 µL standard 2 + 200 µL Diluent	0.03
Standard 0	200µL Diluent	0

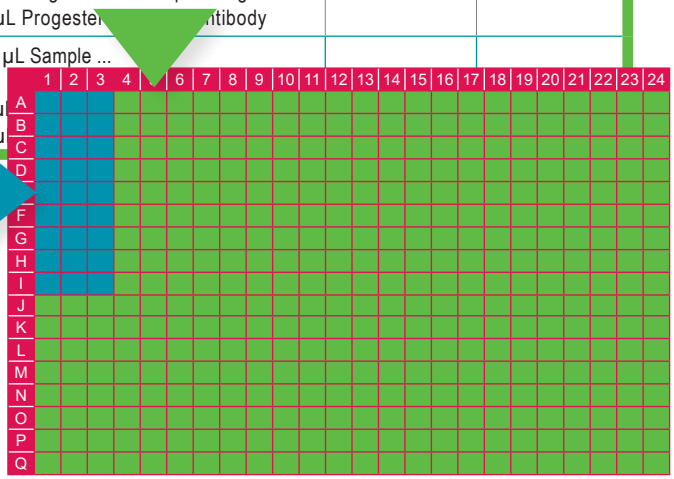
TO PREPARE SAMPLES:

- Each well requires 10 µL of sample.
- Just after their collection, put the samples at 4°C and test them immediately. For later use, samples should be dispensed into disposable plastic vials and stored at -60°C or below. Avoid multiple freeze/thaw cycles.
- Samples with a concentration above the highest standard (Std 8) must be diluted diluent or in your appropriate sample medium.

ASSAY MANUAL

	Negative control or Cryptate control	Standard (Std 0 - Std 8)	Samples
Step 1 	Dispense 10 µL of diluent into each negative control well	Dispense 10 µL of each Progesterone standard (Std 0 - Std 8) into each standard well	Dispense 10 µL of each sample into each sample well
Step 2 	Add 5 µL of detection buffer to all negative control wells	Add 5 µL Progesterone acceptor reagent working solution to all wells	
Step 3 	Add 5 µL Progesterone donor antibody working solution to all wells		
Step 4 	Seal the plate and incubate 1 hour @ RT		
Step 5	Remove the plate sealer and read on an HTRF® compatible reader		

	1	2	3	4	5	6
A	10 µL diluent (Negative control) 5 µL Detection Buffer# 3 5 µL Progesterone donor antibody	Repeat Well A1	Repeat Well A1	10 µL Sample 1 5 µL Progesterone acceptor reagent 5 µL Progesterone donor antibody	Repeat Well A4	Repeat Well A4
B	10 µL Std 0 (Positive control) 5 µL Progesterone acceptor reagent 5 µL Progesterone donor antibody	Repeat Well B1	Repeat Well B1	10 µL Sample 2 5 µL Progesterone acceptor reagent 5 µL Progesterone donor antibody	Repeat Well B4	Repeat Well B4
C	10 µL Std 1 5 µL Progesterone acceptor reagent 5 µL Progesterone donor antibody	Repeat Well C1	Repeat Well C1	10 µL Sample 3 5 µL Progesterone acceptor reagent 5 µL Progesterone donor antibody	Repeat Well C4	Repeat Well C4
D	10 µL Std 2 5 µL Progesterone acceptor reagent 5 µL Progesterone donor antibody	Repeat Well D1	Repeat Well D1	10 µL Sample ... 5 µL Progesterone acceptor reagent 5 µL Progesterone donor antibody	Repeat Well D4	Repeat Well D4
E	10 µLStd ... 5 µL Progesterone acceptor reagent 5 µL Progesterone donor antibody	Repeat Well E1	Repeat Well E1	10 µL Sample ... 5 µL Progesterone acceptor reagent 5 µL Progesterone donor antibody	Repeat Well E4	Repeat Well E4
F	10 µL Std ... 5 µL Progesterone acceptor reagent 5 µL Progesterone donor antibody	Repeat Well F1	Repeat Well F1	10 µL Sample ... 5 µL Progesterone acceptor reagent 5 µL Progesterone donor antibody	Repeat Well F4	Repeat Well F4
G	10 µL Std ... 5 µL Progesterone acceptor reagent 5 µL Progesterone donor antibody	Repeat Well G1	Repeat Well G1	10 µL Sample ... 5 µL Progesterone acceptor reagent 5 µL Progeste[illegible]ntibody	Repeat Well G4	Repeat Well G4
H	10 µL Std ... 5 µL Progesterone acceptor reagent 5 µL Progesterone donor antibody	Repeat Well H1	Repeat Well H1	10 µL Sample ... 5 µ 5 µ		



DATA REDUCTION

1. Calculate the ratio of the acceptor and donor emission signals for each individual well.

$$\text{Ratio} = \frac{\text{Signal 665 nm}}{\text{Signal 620 nm}} \times 10^4$$

2. Calculate the % CVs. The mean and standard deviation can then be worked out from ratio replicates.

$$\text{CV (\%)} = \frac{\text{Standard deviation}}{\text{Mean Ratio}} \times 100$$

3. Calculate the % delta F which reflects the signal to background of the assay. The negative control plays the role of an internal assay control. Delta F is used for the comparison of day to day runs of the same assay.

$$\text{delta F (\%)} = \frac{\text{Ratio Standard or sample - Ratio Negative Control}}{\text{Ratio Negative Control}} \times 100$$

For more information about data reduction, please visit www.revvity.com

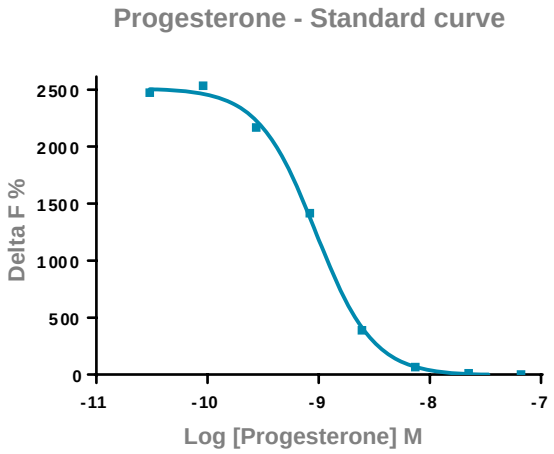
RESULTS

This data must not be substituted for the data obtained in the laboratory and should be considered only as an example.

Results may vary from one HTRF® compatible reader to another.

The assay standard curve is created by plotting delta F% versus the analyte concentration. To determine sample concentration, we recommend to use a log scale for the Progesterone concentrations and analyze the data with the sigmoidal dose response curve with variable slope.

	Ratio (1)	CV (2)	Delta F% (3)
Negative control	630	2.28%	
Std 0 – Positive control	17,593	4.04%	2,692%
Std 1 -0.03 nM	16,216	5.13%	2,473%
Std 2 - 0.09 nM	16,595	5.88%	2,533%
Std 3 - 0.27 nM	14,293	6.98%	2,168%
Std 4 - 0.82 nM	9,556	4.72%	1,416%
Std 5 - 2.47 nM	3,076	3.05%	388%
Std 6 - 7.41 nM	1,042	1.51%	65%
Std 7 - 22.22 nM	703	4.42%	12%
Std 8 - 66.67 nM	635	5.41%	1%



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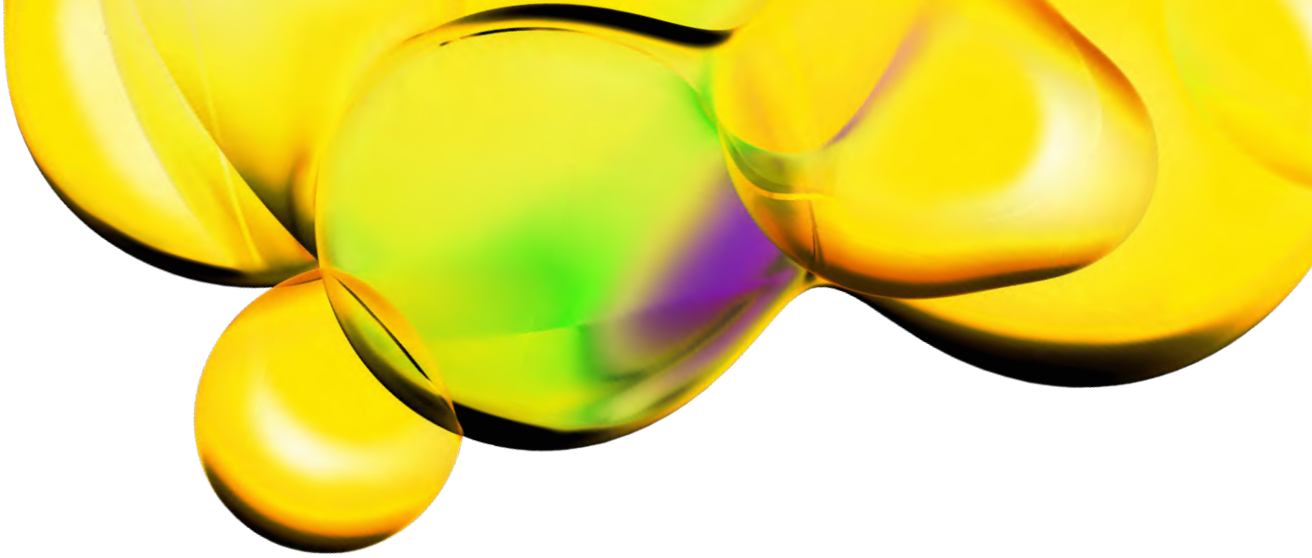

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