revvity



Part # 62PROPEG & 62PROPEH

Test size#: 500 tests (62PROPEG) and 10,000 tests (62PROPEH) - assay volume: 20 μL Revision: #04 of September 2023 Store at: -60°C or below (62PROPEG); -60°C or below (62PROPEH) For research use only. Not for use in diagnostic procedures.

ASSAY PRINCIPLE

This kit is intended for the simple and rapid quantification of Progranulin in cell/tissue culture supernatants 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, Progranulin is detected in a sandwich assay by using anti Progranulin antibody labeled with Europium cryptate (donor), and anti Progranulin antibody 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). Signal intensity is proportional to the number of antigen-antibody complexes formed and therefore to the progranulin concentration.

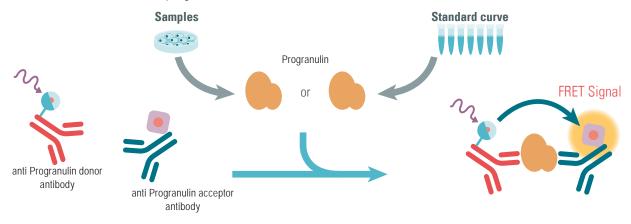
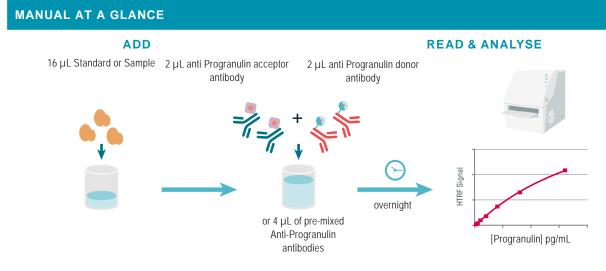


Figure 1: Principle of HTRF Progranulin sandwich assay.



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

MATERIALS PROVIDED:

KIT COMPONENTS	500 TESTS * CAT # 62PROPEG	10,000 TESTS * CAT # 62PROPEH
Progranulin Standard	1 vial - 50 µL	1 vial - 50 µL
Frozen	1 µg/mL	1 µg/mL
Progranulin Eu Cryptate Antibody	1 vial - 20 µL	1 vial - 0.4 mL
Frograndin Lu Cryptale Antibody	Frozen - 50X	Frozen - 50X
Progranulin d2 Antibody	1 vial - 20 µL	1 vial - 0.4 mL
Progranulin d2 Antibody	Frozen - 50X	Frozen - 50X
Diluent #5 **	1 vial	1 vial
5X	2 mL	10 mL
Detection buffer ***	2 vials	1 vial
	1.5 mL	50 mL
ready-to-use	Detection Buffer #3	Detection Buffer #3

* 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 - Use white plate only. For more information about microplate recommendations, please visit our website at: www.revvity.com

STORAGE AND STABILITY

Store the kit at -60°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.

Volume of Progranulin standard aliquots should not be under 10 µL.

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.
- Thaw the frozen reagents at room temperature, allow them to warm up to room temperature for at least 30 mins before use
- · Before use, allow Diluent and Detection buffer to warm up at room temperature and homogenize them with a vortex.
- · Antibody solutions must be prepared in individual vials and can be mixed prior to dispensing.
- · Progranulin 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 - 62PROPE	G	10,000 TESTS KIT - 62PROPEH		
	Anti-Progranulin E	u Cryptate antibody		
Thaw the Progranulin Eu Cryptate antibody. Mix gently. This 50X stock solution can be frozen and stored at -60°C or below.	Ī	I	Thaw the Progranulin Eu Cryptate antibody. Mix gently. This 50X stock solution can be frozen and stored at -60°C or below.	
	Anti-Progranu	lin d2 antibody		
Thaw the Progranulin d2 antibody. Mix gently. This 50X stock solution can be frozen and stored at -60° C or below.		Ī	Thaw the Progranulin d2 antibody. Mix gently. This 50X stock solution can be frozen and stored at -60° C or below.	
	Progranuli	n Standard		
Thaw the Progranulin standard solution in order to obtain a 1 μ g/mL stock solution. Mix gently.			Thaw the Progranulin standard solution in order to obtain a 1 μ g/mL stock solution. Mix gently.	
	Dilu	ient		
Dilute 5-fold the 5 X diluent #5 with distilled water: homogenize the 5 X diluent #5 with a vortex and add 1 volume of stock solution in 4 volumes of distilled water (e.g., 1 mL of diluent + 4 mL of distilled water). Mix gently after dilution. This 1X diluent can be frozen and stored at -60°C or below.	4 vol	1 vol	Dilute 5-fold the 5 X diluent #5 with distilled water: homogenize the 5 X diluent #5 with a vortex and add 1 volume of stock solution in 4 volumes of distilled water (e.g., 10 mL of diluent + 40 mL of distilled water). Mix gently after dilution. This 1X diluent can be frozen and stored at -60°C or below.	
	Detectio	n buffer		
The Detection buffer is ready-to-use.			The Detection buffer is ready-to-use.	

TO PREPARE ANTIBODY WORKING SOLUTIONS:

Each well requires 2 µL of Progranulin-Eu Cryptate Antibody and 2 µL of Progranulin-d2 Antibody.

Prepare the two antibody solutions in separate vials.

500 TESTS KIT - 62PROPEG		10,000 TESTS KIT - 62PROPEH				
Progranulin Eu Cryptate antibody						
Dilute 50-fold the 50X stock solution (thawed reagent) of Progranulin Eu Cryptate antibody stock solution with the Detection buffer #3 : add 1 volume of Eu Cryptate antibody stock solution in 49 volumes of Detection buffer #3 (e.g., 20 μ L of Eu Cryptate antibody stock solution + 980 μ L of Detection Buffer #3).	1 vol 49 vol	1 vol 49 vol	Dilute 50-fold the 50X stock solution (thawed reagent) of Progranulin Eu Cryptate antibody stock solution with the Detection buffer #3 : add 1 volume of Eu Cryptate antibody stock solution in 49 volumes of Detection buffer #3 (e.g., 0.4 mL of Eu Cryptate antibody stock solution + 19.6 mL of Detection Buffer #3).			
	Progranulin d2 antibody					
Dilute 50-fold the 50X stock solution (thawed reagent) of Progranulin d2 antibody stock solution with the Detection buffer #3 : add 1 volume of d2- antibody stock solution in 49 volumes of Detection buffer #3 (e.g., 20 μ L of d2-antibody stock solution + 980 μ L of Detection Buffer #3).	1 vol 49 vol	1 vol 49 vol	Dilute 50-fold the 50X stock solution (thawed reagent) of Progranulin d2 antibody stock solution with the Detection buffer #3 : add 1 volume of d2 antibody stock solution in 49 volumes of Detection buffer #3 (e.g., 0.4 mL of d2 antibody stock solution + 19.6 mL of Detection Buffer #3).			
Antibody mix						
It is possible to pre-mix the two ready-to-use antibody solutions just prior to dispensing the reagents by adding 1 volume of d2 antibody solution to 1 volume of Cryptate antibody solution (e.g. 1 mL of d2 antibody + 1 mL of Cryptate antibody).			It is possible to pre-mix the two ready-to-use antibody solutions just prior to dispensing the reagents by adding 1 volume of d2 antibody solution to 1 volume of Cryptate antibody solution (e.g. 1 mL of d2 antibody + 1 mL of Cryptate antibody).			

TO PREPARE STANDARD WORKING SOLUTIONS:

- Each well requires 16 µL of standard.
- Dilute the standard stock solution serially with diluent #5 (1X)
- 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 #5 (1X).
- · In order to counteract any standard sticking, we recommend changing tips between each dilution.

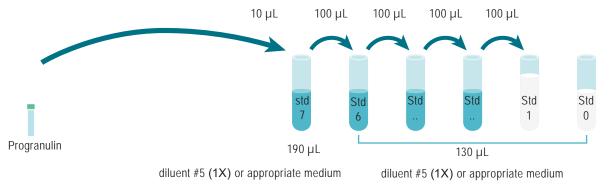
A recommended standard dilution procedure is listed and illustrated below:

Dilute the standard stock solution 20-fold with diluent #5 (1X) to prepare high standard (std 7): e.g. take 10 μ L of standard stock solution and add it to 190 μ L of diluent #5 (1X). Mix gently.

Use the high standard (std 7) to prepare the standard curve using 1/2.3 serial dilutions as follows:

- Dispense 130 μ L of diluent #5 (1X) in each vial from Std 6 to Std 0.
- Add 100 μL of standard to 130 μL of diluent #5 (1X), mix gently and repeat the 1/2.3 serial dilution to make standard solutions: std6, std5, std4, std3, std2, std1.

This will create 7 standards for the analyte. Std 0 (Negative control) is diluent #5 (1X) or appropriate culture medium alone.



STANDARD	SERIAL DILUTIONS	PROGRANULIN WORKING SOLUTIONS (PG/ML)
Standard Stock solution	Thawed stock solution	1,000,000
Standard 7	10 μL stock solution + 190 μL Diluent #5 (1X)	50,000
Standard 6	100 µL standard 7 + 130 µL Diluent #5 (1X)	21,739
Standard 5	100 µL standard 6 + 130 µL Diluent #5 (1X)	9,452
Standard 4	100 µL standard 5 + 130 µL Diluent #5 (1X)	4,109
Standard 3	100 µL standard 4 + 130 µL Diluent #5 (1X)	1,787
Standard 2	100 µL standard 3 + 130 µL Diluent #5 (1X)	777
Standard 1	100 µL standard 2 + 130 µL Diluent #5 (1X)	338
Standard 0	100 µL Diluent #5 (1X)	0

TO PREPARE SAMPLES:

- Each well requires 16 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 7) must be diluted diluent #5 (1X) or in your appropriate sample medium.
- In order to measure human Progranulin in cell lysates, cells must be lyzed with Lysis Buffer #4 (1X) for 30 min at RT under gentle shaking. Please note that the 4X stock solution of Lysis Buffer #4 must be ordered separately (Ref# 64KL4FDF, 130 mL) and 4-fold diluted with distilled water before use.

ASSAY MANUAL

		Standard (Std 0 - std 7)	Samples			
Step 1		Dispense 16 µL of each Progranulin standard (Std 0 - std 7) into each standard well	Dispense 16 μL of each sample into each sample well			
Step 2		Add 2 µL of Progranulin d2 antibody working solution to all wells				
Step 3		Add 2 μL of Progranulin Eu Cryptate antibody working solution to all wells				
Step 4	0	Seal the plate and incubate overnight				
Step 5	9	Remove the plate sealer and read on an HTRF® compatible reader				

	1	2	3	4	5	6
A	16 μL Std0 (Negative control) 2 μL Progranulin-d2	Repeat Well A1	Repeat Well A1	16 μL Sample 1 2 μL Progranulin-d2	Repeat Well A4	Repeat Well A4
	2 μL Progranulin-Eu Cryptate 16 μL Std 1			2 μL Progranulin-Eu Cryptate 16 μL Sample 2		
В	2 µL Progranulin-d2 2 µL Progranulin-Eu Cryptate	Repeat Well B1	Repeat Well B1	2 μL Progranulin-d2 2 μL Progranulin-Eu Cryptate	Repeat Well B4	Repeat Well B4
С	16 µL Std 2	Repeat Well C1	Repeat Well C1	16 µL Sample 3	Repeat Well C4	Repeat Well C4
	2 μL Progranulin-d2 2 μL Progranulin-Eu Cryptate			2 μL Progranulin-d2 2 μL Progranulin-Eu Cryptate		
D	16 μL Std 2 μL Progranulin-d2 2 μL Progranulin-Eu Cryptate	Repeat Well D1	Repeat Well D1	16 μL Sample Repeat Well D4 2 μL Progranulin-d2 Repeat Well D4 2 μL Progranulin-Eu Cryptate Repeat Well D4		Repeat Well D4
E	16 μLStd 2 μL Progranulin-d2 2 μL Progranulin-Eu Cryptate	Repeat Well E1	Repeat Well E1	16 μL Sample 2 μL Progranulin-d2 2 μL Progranulin-Eu Cryptate	Repeat Well E4	Repeat Well E4
F	16 μL Std 2 μL Progranulin-d2 2 μL Progranulin-Eu Cryptate	Repeat Well F1	Repeat Well F1	16 μL Sample 2 μL Progranulin-d2 2 μL Progranulin-Eu Cryptate		Repeat Well F4
G	16 μL Std 2 μL Progranulin-d2 2 μL Progranulin-Eu Cryptate	Repeat Well G1	Repeat Well G1	16 μL Sample Repeat Well G4 2 μL Progranuline Repeat Well G4		Repeat Well G4
н	16 μL Std 2 μL Progranulin-d2 2 μL Progranulin-Eu Cryptate	Repeat Well H1	Repeat Well H1	16 µ 1 2 3 4 6 7 8 9 10 11 A 2 µL B C I D	1 12 13 14 15 16 1	7 18 19 20 21 22 23

DATA REDUCTION

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.

3. Calculate the delta ratio of the acceptor and donor emission signals for each individual well. The Standard 0 (Negative control) plays the role of an internal assay control.

delta Ratio = Ratio Standard or sample - Ratio Standard 0

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

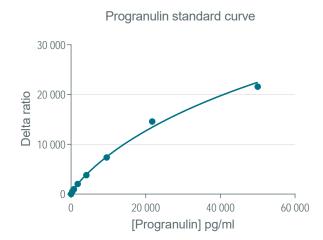
RESULTS

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.

Standard curve fitting with the 4 Parameter Logistic (4PL) model (with 1/Y² weighting):

	Ratio (1)	CV (2)	Delta Ratio
Standard 0 - Negative control	687	2%	0
Standard 1 - 338 pg/mL	1222	4%	535
Standard 2 - 777 pg/mL	1673	1%	986
Standard 3 - 1,787 pg/mL	2718	2%	2031
Standard 4 - 4,109 pg/mL	4489	1%	3802
Standard 5 - 9,452 pg/mL	8038	2%	7351
Standard 6 - 21,739 pg/mL	15262	1%	14575
Standard 7 - 50,000 pg/mL	22282	1%	21595



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