### Description

Covalent conjugation to ubiquitin (Ub) is one of the major post-translational modifications that regulates protein stability, function, and localization. Ubiquitination is the concerted action of three enzymes: a Ub-activating enzyme (E1), a Ub-conjugating enzyme (E2), and a Ub ligase (E3). The specificity and efficiency of ubiquitination are largely determined by the E3 enzyme, which directs the last step of the Ub-conjugating cascade by binding to both an E2~Ub conjugate and a substrate protein. This step ensures the transfer of Ub from E2~Ub to the substrate, leading to its mono- or poly-ubiquitination.

The Cereblon (CRBN) protein via its interaction with the DNA damage-binding protein-1 (DDB1), Cullin 4 (Cul4A or Cul4B), and regulator of Cullins 1 (RoC1) forms the functional E3 Ub ligase complex. In this complex, Cereblon functions as a substrate receptor that mediates the ubiquitination and subsequent proteasomal degradation of target proteins. Cereblon complex is involved in many biological processes including cell proliferation and apoptosis. That is why it has been employed for targeted protein degradation in the treatment of cancer. Like most E3 ligases, Cereblon complex ubiquitinates itself and this auto-ubiquitination promotes its Ub ligase activity. The Cereblon intrachain TR-FRET Assay Kit is a sensitive high-throughput screening (HTS) TR-FRET Assay Kit, designed to measure Cereblon auto-ubiquitination activity in a homogeneous 384 reaction format. It utilizes a Europium cryptate-labeled Ub (donor) as well as Cy5-labeled Ub (acceptor) to complete the TR-FRET pairing. Since both the TR-FRET donor and acceptor are incorporated into poly-ubiquitin chains formed on Cereblon, this FRET-based assay requires no time-consuming washing steps, making it especially suitable for HTS applications as well as real-time kinetics analyses of polyubiquitination.

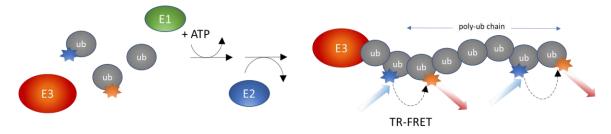


Figure 1. Cereblon intrachain TR-FRET Assay Kit schematic

#### **Applications**

Great for screening molecules that inhibit Cereblon Ub ligase activity in drug discovery HTS applications, for determination of compound IC<sub>50</sub>, and for Cereblon real-time kinetics analyses.

#### **Supplied Materials**

Catalog #	Name	Amount	Storage	
80301	UBE1 (E1)*	50 μg	-80°C	
80314	UBCH5b (E2)*	300 μg	-80°C	Avoid
100329	Cereblon/DDB1/Cul4A/Rbx1 Complex (E3)*	60 μg	-80°C	multiple
78307	TRF Ubiquitin Mix (200x)	50 μΙ	-80°C	freeze/ thaw
	ATP (4 mM)	150 μΙ	-80°C	cycles
78269	CBL assay buffer 2	2x10ml	-80°C	
	White, nonbinding, low volume microtiter plate		Room Temp	

<sup>\*</sup> The initial concentration of enzyme is lot-specific and will be indicated on the tube containing the protein.



#### **Materials Required but Not Supplied**

- Fluorescent microplate reader capable of measuring Time Resolved Fluorescence Resonance Energy Transfer (TR-FRET)
- Adjustable micropipettor and sterile tips
- Rotating or rocker platform

### **Storage Conditions**



This assay kit will perform optimally for up to 6 months from date of receipt when the materials are stored as directed. **Avoid multiple freeze/ thaw cycles!** 

#### Safety



This product is for research purposes only and not for human or therapeutic use. This product should be considered hazardous and is harmful by inhalation, in contact with skin, eyes, clothing, and if swallowed. If contact occurs, wash thoroughly.

#### **Contraindications**

The Cereblon intrachain TR-FRET Assay Kit is compatible with up to 1% final DMSO concentration. We recommend preparing the inhibitor in no higher than 5% DMSO solution in buffer and using 4  $\mu$ l per well.

#### **Assay Protocol**

- All samples and controls should be performed in triplicates
- The assay should include a "Blank", a "Positive control", and a "Negative control"
- 1) Thaw UBE1, UBCH5b, Cereblon complex, TRF Ubiquitin Mix, CBL assay buffer 2, and ATP on ice. Aliquot each protein, assay buffer, and ATP into single-use aliquots and immediately store at -80°C. Note: UBE1, UBCH5b, Cereblon complex, TRF Ubiquitin Mix, and CBL assay buffer 2 are sensitive to freeze/thaw cycles. Avoid multiple freeze-thaw cycles.
- 2) Carefully calculate the amount of each protein needed and prepare appropriate amounts of diluted proteins:

Prepare 5x TRF Ubiquitin Mix in CBL assay buffer 2 (40-fold dilution of the 200x TRF Ubiquitin Mix); Dilute the UBE1 in CBL assay buffer 2 at 800 nM (96 ng/ $\mu$ l) (final concentration in reaction 40 nM); Dilute the UBCH5b in CBL assay buffer 2 at 10  $\mu$ M (720 ng/ $\mu$ l) (final concentration in reaction 500 nM); Dilute the Cereblon complex in CBL assay buffer 2 at 100 nM (25 ng/ $\mu$ l) (final concentration in reaction 25 nM);

Keep all diluted proteins on ice until use.



3) Prepare the compound solution.

If the compound is dissolved in DMSO, make a 100-fold higher concentration of the compound in DMSO than the highest concentration you want to test in the assay. Then dilute 20-fold in CBL assay buffer 2 (at this step the compound concentration is 5-fold higher than the desired final concentration). If you want to run an  $IC_{50}$  or test lower concentrations of the compound, prepare serial dilutions using 1X assay buffer containing 5% DMSO, so the final concentration of DMSO will be 1% in all samples.

If the compound is soluble in water, prepare a solution of the compound in CBL assay buffer 2 that is 5-fold higher than the final assay concentration.

4) To the wells designated as "Blank", add 4  $\mu$ l of **5x TRF Ubiquitin Mix** + 1  $\mu$ l of **UBE1** + 1  $\mu$ l of **UBCH5b** + 4  $\mu$ l of **diluent solution** (for example DMSO 5%) + 5  $\mu$ l of **CBL assay buffer 2**.

	Blank
TRF Ubiquitin Mix (5x)	4 μΙ
UBE1	1 μΙ
UBCH5b	1 μl
Cereblon complex	-
Test Compound	-
Diluent solution* (no inhibitor)	4 μΙ
CBL assay buffer 2	5 μΙ
ATP (4 mM)	5 μΙ
Total	20 μΙ

<sup>\*</sup>The diluent solution contains the assay buffer with the same concentration of solvent (e.g. DMSO) as the test compound solution.

5) Make the master mixture using diluted reagents:

N wells  $\times$  (4  $\mu$ l 5x TRF Ubiquitin Mix + 1  $\mu$ l UBE1 + 1  $\mu$ l UBCH5b + 5  $\mu$ l Cereblon complex).

- 6) Add 11  $\mu$ l of master mixture to each well designated for the "Negative Control", "Positive Control", "Test Sample".
- 7) Add 4 µl of inhibitor solution to each well designated "Test Inhibitor". For all other wells: "Positive Control", "Negative Control", add 4 µl of the diluent solution without inhibitor.
- 8) Initiate the reaction by adding 5  $\mu$ l of **ATP** to the wells labeled "Positive Control," "Test Inhibitor," and "Blank." Add 5  $\mu$ l of **CBL** assay buffer 2 to the well designated "Negative Control." Cover the plate with a plate sealer. Incubate the reaction at room temperature for two hours or at 30°C for one hour.



	Test Sample	Negative Control	Positive Control
Master Mix	11 μΙ	11 μl	11 μΙ
Test compound	4 μΙ	_	_
Diluent solution* (no inhibitor)	_	4 μΙ	4 μΙ
CBL assay buffer 2	-	5 μΙ	_
ATP (4 mM)	5 μΙ	_	5 μΙ
Total	20 μΙ	20 μΙ	20 μΙ

<sup>\*</sup>The diluent solution contains the assay buffer with the same concentration of solvent (e.g. DMSO) as the test compound solution.

9) Read the fluorescent intensity in a microtiter-plate reader capable of measuring TR-FRET. "Blank" value is subtracted from all other values.

#### **Instrument Settings**

Reading Mode	Time Resolved	
Excitation Wavelength	340±20 nm	
Emission Wavelength	620±10 nm	
Lag Time	60 μs	
Integration Time	500 μs	
Excitation Wavelength	340±20 nm	
Emission Wavelength	665±10 nm	
Lag Time	60 μs	
Integration Time	500 μs	

#### **CALCULATING RESULTS:**

Two sequential measurements should be conducted. Tb-donor emission should be measured at 620 nm followed by dye-acceptor emission at 665 nm. Data analysis is performed using the TR-FRET ratio (665 nm emission/620 nm emission).

When percentage activity is calculated, the FRET value from the Blank (it is expected that Blank and Negative Control represent similar value) can be set as zero percent activity and the FRET value from the positive control can be set as one hundred percent activity.

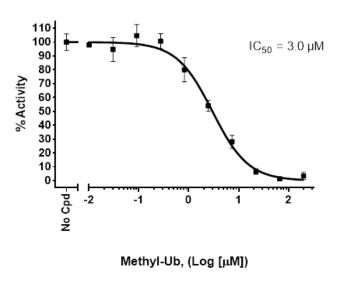
% Activity = 
$$\frac{FRET_s - FRET_{blank}}{FRET_p - FRET_{blank}} \times 100\%$$

Where FRETs = Sample FRET, FRET<sub>blank</sub> = Blank FRET, and FRET<sub>P</sub> = Positive control FRET.



## **Example Results**

# Cerebion TR-FRET Activity



**Figure 1:** Inhibition of Cereblon complex auto-ubiquitination by Methylated Ubiquitin, measured using the Cereblon intrachain TR-FRET Assay Kit, BPS Bioscience #78301 Data shown is representative. For lot-specific information, please contact BPS Bioscience, Inc. at support@bpsbioscience.com.

## **Troubleshooting Guide**

Visit bpsbioscience.com/assay-kits-faq for detailed troubleshooting instructions. For all further questions, please email support@bpsbioscience.com



#### **Related Products**

Products	Catalog #	Size
MDM2 intrachain TR-FRET Assay Kit	78302	384 rxns.
SMURF1 intrachain TR-FRET Assay Kit	78303	384 rxns.
SMURF2 intrachain TR-FRET Assay Kit	78304	384 rxns.
VHL intrachain TR-FRET Assay Kit	78305	384 rxns.
XIAP intrachain TR-FRET Assay Kit	78306	384 rxns.
MDM2 TR-FRET Assay Kit	79773	384 rxns.
CBL-B TR-FRET Assay Kit	79575	384 rxns.
c-CBL TR-FRET Assay Kit	79786	384 rxns.
Cereblon Ubiquitination Homogenous Assay Kit	79881	384 rxns.
UBCH13 TR-FRET Assay Kit	79741	384 rxns.
UBCH5a TR-FRET Assay Kit	79900	384 rxns.
UBCH5c TR-FRET Assay Kit	79901	384 rxns.
UBCH5b TR-FRET Assay Kit	79896	384 rxns.
MDM2, GST-Tag (Human)	80751	20 μg
UBE1 (UBA1), FLAG-tag	80301	100 μg
UBE1, GST-Tag	100402	100 μg
UBE2A, His-Tag	79368	20 μg
UBE2C, His-Tag	79369	20 μg
UBE2D2, His-Tag	79370	20 μg
UBE2E3 (UBCH9), His-Tag	79371	20 μg
UBE2G1 (UBC7), His-Tag	79372	20 μg
UBE2K (UBC1), His-Tag	79373	20 μg
UBE2O, GST-Tag	79374	20 μg
UbcH5a (UBE2D1), His-tag	80315	100 μg
UbcH5b, His-Tag (Human)	80314	100 μg
UbcH6 (UBE2E1), His-tag	80316	100 μg
UbcH7, His-tag (E. coli-derived)	80317	100 μg
UbcH7, His-tag (Sf9-derived)	80318	50 μg
UbcH13 (UBE2N), His-tag	80323	100 μg
CBL-B, GST-Tag (Human)	80415	100 μg
c-CBL, GST-Tag (Human)	100370	100 μg
XIAP, FLAG-tag	80401	20 μg
SMURF1, FLAG-tag	80402	20 μg
SMURF2, FLAG-tag	80403	20 μg
Cereblon/DDB1/Cul4A/Rbx1 Complex	100329	10 μg
VHL/CUL2/ELOB/ELOC/RBX1 Complex	100373	10 μg
Ubiquitin, His-Tag	79293	2 mg
Ubiquitin, His-Avi-Tag, Biotin Labeled	11236	50 μg

