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# <u>Data Sheet</u> BRD3 (BD2) Inhibitor Screening Assay Kit Catalog # 32523

**DESCRIPTION:** The *BRD3* (*BD2*) *Inhibitor Screening Assay Kit* is designed to measure the inhibition of BRD3 bromodomain 2 (BD2) from binding to its substrate. The *BRD3* (BD2) *Inhibitor Screening Assay Kit* comes in a convenient AlphaLISA® format, with biotinylated histone peptide substrate, assay buffer, detection buffer and purified, GST-tagged BRD3 BD2 to perform a total of 384 enzyme reactions. The key to the *BRD3* (*BD2*) *Inhibitor Screening Assay Kit* is the highly specific binding of the BRD3 bromodomain 2 to the acetylated histone substrate. With this kit, only three simple steps on a microtiter plate are required. First, a sample containing BRD3 bromodomain 2 and an inhibitor of choice is incubated with the biotinylated substrate for thirty minutes. Next, acceptor beads are added, then donor beads, followed by reading the Alpha-counts.

### **COMPONENTS:**

| Catalog # | Component                         | Amount | Storage |          |
|-----------|-----------------------------------|--------|---------|----------|
| 31033     | BRD3 (306-417), BD2, GST-tag      | 20 µg  | -80°C   |          |
|           | BET Bromodomain Ligand            | 400 µl | -80°C   | (Avoid   |
|           | Non-acetylated Ligand             | 200 µl | -80°C   | freeze/  |
| 33001     | 3x BRD Homogeneous Assay Buffer 1 | 4 ml   | -20°C   | thaw     |
| 33002     | 3x BRD Homogeneous Detection      | 3 ml   | -20°C   | cycles!) |
|           | Buffer 1                          |        |         |          |

### MATERIALS OR INSTRUMENTS REQUIRED BUT NOT SUPPLIED:

AlphaLISA GSH acceptor beads, 5 mg/ml (PerkinElmer #AL109C)

AlphaScreen Streptavidin-conjugated donor beads, 5 mg/ml (PerkinElmer #6760002S)

Optiplate -384 (PerkinElmer #6007290)

AlphaScreen microplate reader

Adjustable micropipettor and sterile tips

**APPLICATIONS:** Useful for the study of bromodomain binding assays, screening inhibitors and selectivity profiling.

**CONTRAINDICATIONS:** Only limited amounts of DMSO can be included, as it has been shown to disrupt BRD-ligand interaction. Avoid green and blue dyes that absorb light in the AlphaScreen signal emission range (520-620 nm), such as Trypan Blue. Avoid the use of the potent singlet oxygen quenchers such as sodium azide (NaN<sub>3</sub>) or metal ions (Fe<sup>2+</sup>, Fe<sup>3+</sup>, Cu<sup>2+</sup>, Zn<sup>2+</sup> and Ni<sup>2+</sup>). The presence of >1% RPMI 1640 culture medium leads to a signal reduction due to the presence of excess biotin and iron in this medium. MEM, which lacks these components, does not affect AlphaScreen assays.

**STABILITY:** At least one year from date of receipt when stored as directed.

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**REFERENCE:** Muller, S., Filippakopoulos, P., Knapp, S., *Expert Rev. Mol. Med.* 2011 Sep 13:**13**:e29.

### **ASSAY PROTOCOL:**

All samples and controls should be tested in duplicate. Use slow shaking for all incubations.

# Step 1:

- 1) Prepare the master mixture: N wells x (2.5 μl **3x BRD Homogeneous Assay Buffer 1** + 1 μl **BET Bromodomain Ligand** + 1.5 μl **H<sub>2</sub>O**).
- 2) Thaw **BRD3** (**BD2**) on ice. Upon first thaw, briefly spin tube containing protein to recover full content of the tube. Aliquot both proteins into single use aliquots. Store remaining undiluted protein in aliquots at -80°C immediately. *Note:* **BRD3** is very sensitive to freeze/thaw cycles. Do not re-use thawed aliquots or diluted protein.
- 3) Dilute **BRD3 (BD2)** in **1x BRD Homogeneous Assay Buffer 1** at 16 ng/µl. Keep diluted proteins on ice until use. Discard any unused diluted protein after use.

Add 5  $\mu$ l of master mixture to each well designated for the "Positive Control", "Test Inhibitor", and "Blank". For the "Substrate Control", add 2.5  $\mu$ l **3x BRD Homogeneous Assay Buffer 1** + 1  $\mu$ l **Non-acetylated Ligand** + 1.5  $\mu$ l **H<sub>2</sub>O**.

|                                      | Blank  | Substrate<br>Control | Positive Control | Test<br>Inhibitor |
|--------------------------------------|--------|----------------------|------------------|-------------------|
| 3x BRD Homogeneous Assay<br>Buffer 1 | 2.5 µl | 2.5 µl               | 2.5 µl           | 2.5 µl            |
| BET Bromodomain Ligand               | 1 µl   | _                    | 1 µl             | 1 µl              |
| Non-acetylated Ligand                | -      | 1 µl                 | -                | -                 |
| H <sub>2</sub> O                     | 1.5 µl | 1.5 µl               | 1.5 µl           | 1.5 µl            |
| Test Inhibitor/Activator             | -      | _                    | -                | 2.5 µl            |
| Inhibitor buffer (no inhibitor)      | 2.5 µl | 2.5 µl               | 2.5 µl           | ı                 |
| 1x BRD Homogeneous Assay<br>Buffer 1 | 2.5 µl |                      |                  |                   |
| BRD3 (BD2) (16 ng/µl)*               | _      | 2.5 µl               | 2.5 µl           | 2.5 µl            |
| Total                                | 10 µl  | 10 µl                | 10 µl            | 10 µl             |

- 4) Add 2.5 µl of **inhibitor solution** to each well designated "Test Inhibitor". For the "Positive Control", "Substrate Control" and "Blank", add 2.5 µl of the same **solution without inhibitor** (inhibitor buffer). *Note: Keep DMSO concentration below 0.5 %*.
- 5) Add 2.5 µl of **1x BRD Homogeneous Assay Buffer 1** to the well designated "Blank".

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6) Initiate reaction by adding 2.5 µl of diluted **BRD3 (BD2)** prepared as described above. Incubate at room temperature for 30 minutes.

# Step 2:

### Note: Protect your samples from direct exposure to light!

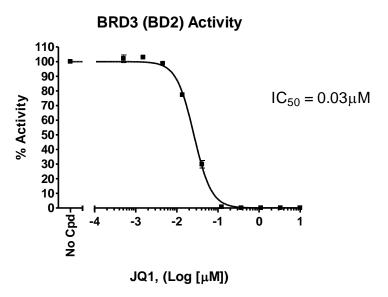
1) Dilute GSH Acceptor beads (PerkinElmer #AL109C) 250-fold with 1x BRD Homogeneous Detection Buffer 1. Add 10 µl per well. Shake plate briefly. Incubate at room temperature for 30 minutes.

# Step 3:

- 1) Dilute Streptavidin-conjugated donor beads (PE #6760002S) 250-fold with 1x Homogeneous Detection Buffer 1. Add 10 µl per well. Incubate at room temperature for 15 30 minutes.
- 2) Read Alpha-counts.

Due to lot to lot variability in AlphaScreen® bead performance, it may be necessary to optimize assay conditions. For example, slight adjustments to bromodomain or ligand concentrations may improve signal-to-noise ratio.

### **Example of Assay Results:**



BRD3 (BD2) binding activity, measured using the BRD3 (BD2) Inhibitor Screening Assay Kit, BPS Bioscience, Catalog #32523 and (+)-JQ1 Inhibitor, Catalog #27400. Data shown is lot-specific. For lot-specific information, please contact BPS Bioscience, Inc. at <a href="mailto:support@bpsbioscience.com">support@bpsbioscience.com</a>.

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# **RELATED PRODUCTS:**

| <b>Product Name</b>                | Catalog # | <u>Size</u> |
|------------------------------------|-----------|-------------|
| (+)-JQ1                            | 27400     | 10 mg       |
| BRD2 (339 – 459), His-tag          | 31020     | 100 µg      |
| BRD2 (65 – 187), GST-tag           | 31021     | 100 µg      |
| BRD3 (29 – 145), His-tag           | 31030     | 100 µg      |
| BRD3 (29 – 145), GST-tag           | 31032     | 100 µg      |
| BRD3 (306 – 417), His-tag          | 31031     | 100 µg      |
| BRD3 (306 – 417), GST-tag          | 31033     | 100 µg      |
| BRD4 (49 – 170), GST-tag           | 31040     | 100 µg      |
| BRD4 (342 – 460), GST-tag          | 31041     | 100 µg      |
| BRD4 (49 – 170), His-tag           | 31042     | 100 µg      |
| BRD4 (342 – 460), His-tag          | 31043     | 100 µg      |
| BRD9 (135 – 242), His-tag          | 31090     | 100 µg      |
| BRDT (22 – 138), GST-tag           | 31108     | 100 µg      |
| BRDT (22 – 138), His-tag           | 31101     | 100 µg      |
| BRDT (257 – 382), His-tag          | 31100     | 100 µg      |
| BET Bromodomain Ligand             | 33000     | 0.5 mL      |
| BRD3 (BD1) Inhibitor Screening Kit | 32513     | 384 rxns.   |
| BRD4 (BD1) Inhibitor Screening Kit | 32514     | 384 rxns.   |
| BRD4 (BD2) Inhibitor Screening Kit | 32524     | 384 rxns.   |

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