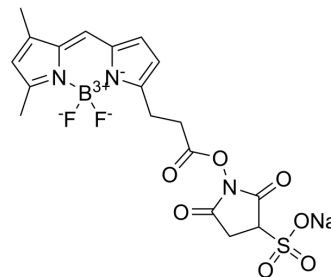


## BODIPY FL SSE

|                           |   |
|---------------------------|---|
| <b>Cat. No.:</b>          | HY-D1607  |
| <b>CAS No.:</b>           | 217190-17-5   |
| <b>Molecular Formula:</b> | C <sub>18</sub> H <sub>17</sub> BF <sub>2</sub> N <sub>3</sub> NaO <sub>7</sub> S         |
| <b>Molecular Weight:</b>  | 491.21  |
| <b>Target:</b>            | Fluorescent Dye   |
| <b>Pathway:</b>           | Others  |
| <b>Storage:</b>           | Please store the product under the recommended conditions in the Certificate of Analysis. |



## BIOLOGICAL ACTIVITY

|                    |  |
|--------------------|--|
| <b>Description</b> | BODIPY FL SSE is a potent fluorescent dye. BODIPY FL SSE is used to label the primary amines (R-NH <sub>2</sub> ) of proteins, amine-modified oligonucleotides, and other amine-containing molecules. BODIPY FL SSE can reactive with primary amines on biomolecules to emit green fluorescence. ( $\lambda_{ex}$ =502 nm, $\lambda_{em}$ =511 nm) <sup>[1][2]</sup> .   |
| <b>In Vitro</b>    | <p>Guidelines (Following is our recommended protocol. This protocol only provides a guideline, and should be modified according to your specific needs).</p> <p>BODIPY FL SSE assay (to quantify GAPDH)<sup>[1]</sup>:</p> <ol style="list-style-type: none"> <li>1. Make a 2 mg/ mL stock of GAPDH (MW 37 kDa) in PBS (pH 7.4).</li> <li>2. Add 54 nM of BODIPY-FL-SSE to 1 mL of the 2 mg/ mL (54 <math>\mu</math>M) GAPDH solution. Incubate the mixture for 4 h on ice.</li> <li>3. Dialyze the reaction mixture against 500 ml of PBS, pH 7.4, in a 10,000 MW-cutoff dialysis cassette for 1 h at 4 °C. Change the buffer and dialyze again overnight. Change the dialysis buffer once more the following morning and dialyze further for 1 h.</li> <li>4. Run the dialyzed sample through a Zeba desalting column at 1000 g for 2 min.</li> <li>5. Analyze the BD-GAPDH spectrophotometrically in a quartz cuvette. Set the wavelength scan to 250-700 nm. Blank the instrument against PBS and add the BD-GAPDH to the cuvette.</li> </ol> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> |

## REFERENCES

- [1]. Hill BG, et, al. Methods for the determination and quantification of the reactive thiol proteome. *Free Radic Biol Med.* 2009 Sep 15;47(6):675-83.
- [2]. Gite S, et, al. Ultrasensitive fluorescence-based detection of nascent proteins in gels. *Anal Biochem.* 2000 Mar 15;279(2):218-25.

**Caution: Product has not been fully validated for medical applications. For research use only.**

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