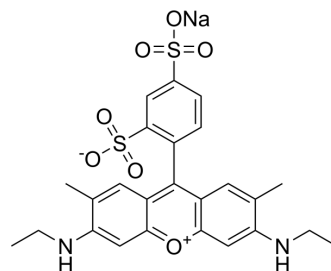


## Sulforhodamine G

<b>Cat. No.:</b>	HY-D1674
<b>CAS No.:</b>	5873-16-5
<b>Molecular Formula:</b>	C <sub>25</sub> H <sub>25</sub> N <sub>2</sub> Na <sub>2</sub> O <sub>7</sub> S <sub>2</sub> <sup>+</sup>
<b>Molecular Weight:</b>	575.58
<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>	Sulforhodamine G is a fluorescent stain with broad dynamic ranges. Sulforhodamine G can be used for the research of protein stains <sup>[1]</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>Labeling of fluorescent internal protein:</p> <p>A. Prepared of protein samples:</p> <ol style="list-style-type: none"> <li>1. Protein samples to be analyzed are spiked with 0.1% of the total protein load of ALIS647 (ALIS internal standard) prior.</li> <li>2. Protein sample is separated with 2-DE in the dark.</li> </ol> <p>B. Purify Sulforhodamine G:</p> <ol style="list-style-type: none"> <li>1. Sulforhodamine G (60% purity, 10 mg) dissolved in 100 mL of 1% v/v acetic acid to purify by RP chromatography.</li> <li>2. Collect the pool with an absorbance maximum at 528 nm, lyophilized to dryness and stored as a dry powder at 47°.</li> </ol> <p>C. Staining:</p> <ol style="list-style-type: none"> <li>1. staining was performed in polypropylene staining dishes wrapped in aluminum foil to prevent photobleaching of the stains.</li> <li>2. Sulforhodamine G staining is performed overnight in 35% methanol with a four-fold molar excess of dye: protein based on an average protein molecular weight of 50 kDa.</li> <li>3. Following 4×15 min washes in 35% methanol and 2×15 min equilibrations in water.</li> <li>4. Total protein and ALIS are visualized with a laser scanner using different channels.</li> <li>5. Protein spots visualized using total protein stain (λ<sub>ex</sub>=532 nm) and ALIS (λ<sub>ex</sub>=633 nm) are quantified separately using 2-DE software, and statistical analysis.</li> </ol> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p>

### REFERENCES

[1]. Asa M Wheelock, et al. Use of a fluorescent internal protein standard to achieve quantitative two-dimensional gel electrophoresis. *Proteomics*. 2006 Mar;6(5):1385-98.

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

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