

## Phalloidin-TRITC

<b>Cat. No.:</b>	HY-P2270
<b>CAS No.:</b>	915013-10-4
<b>Molecular Formula:</b>	C <sub>60</sub> H <sub>70</sub> N <sub>12</sub> O <sub>13</sub> S <sub>2</sub>
<b>Molecular Weight:</b>	1231.4
<b>Target:</b>	Arp2/3 Complex; Fluorescent Dye
<b>Pathway:</b>	Cytoskeleton; Others
<b>Storage:</b>	Sealed storage, away from moisture and light, under nitrogen Powder    -80°C    2 years -20°C    1 year * In solvent : -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture and light, under nitrogen)

### BIOLOGICAL ACTIVITY

<b>Description</b>	Phalloidin-TRITC is a fluorescein derivative of Phalloidin, which can specifically label myofilin and display red fluorescence when labeled and can be observed using Tetrad channels <sup>[1]</sup> .
<b>In Vitro</b>	<ol style="list-style-type: none"> <li>Preparation of Phalloidin-TRITC working solution           <ol style="list-style-type: none"> <li>Preparation of the stock solution Dissolve Phalloidin-TRITC in Methanol to obtain 10 mM of stock solution. Note: It is recommended to store the stock solution at -20 °C or -80 °C away from light and avoid repetitive freeze-thaw cycles.</li> <li>Preparation of Phalloidin-TRITC working solution Dilute the stock solution in serum-free cell culture medium to obtain 1-10 μM of working solution. Note: Please adjust the concentration of Phalloidin-TRITC working solution according to the actual situation.</li> </ol> </li> <li>Cell staining           <ol style="list-style-type: none"> <li>Suspension cells (6-well plate)               <ol style="list-style-type: none"> <li>Centrifuge at 1000 g at 4°C for 3-5 minutes and then discard the supernatant. Wash twice with PBS, 5 minutes each time. The cell density is 1×10<sup>6</sup>/mL.</li> <li>Add 1 mL of working solution, and then incubate at room temperature for 30-60 minutes.</li> <li>Centrifuge at 400 g at 4°C for 3-4 minutes and then discard the supernatant.</li> <li>Wash twice with PBS, 5 minutes each time.</li> <li>Resuspend cells with serum-free cell culture medium or PBS.</li> </ol>               Observation by fluorescence microscopy or flow cytometry.             </li> <li>Adherent cells               <ol style="list-style-type: none"> <li>Culture adherent cells on sterile coverslips.</li> <li>Remove the coverslip from the medium and aspirate excess medium.</li> <li>Add 100 μL of working solution, gently shake it to completely cover the cells, and then incubate at room temperature for 30-60 minutes.</li> <li>Wash twice with medium, 5 minutes each time. Observation by fluorescence microscopy.</li> </ol> </li> </ol> </li> </ol> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p>

### CUSTOMER VALIDATION

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- Small. 2022 Jun 9;e2201147.

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## REFERENCES

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[1]. J A Cooper, et al. Effects of cytochalasin and phalloidin on actin. J Cell Biol. 1987 Oct;105(4):1473-8.

[2]. J Wehland, et al. Phalloidin-induced actin polymerization in the cytoplasm of cultured cells interferes with cell locomotion and growth. Proc Natl Acad Sci U S A. 1977 Dec;74(12):5613-7.

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

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