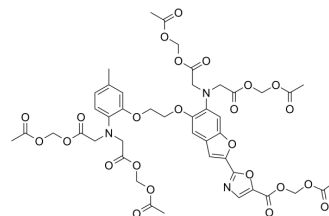


## Fura-2 AM

<b>Cat. No.:</b>	HY-101897
<b>CAS No.:</b>	108964-32-5
<b>Molecular Formula:</b>	C <sub>44</sub> H <sub>47</sub> N <sub>3</sub> O <sub>24</sub>
<b>Molecular Weight:</b>	1001.85
<b>Target:</b>	Others
<b>Pathway:</b>	Others
<b>Storage:</b>	-20°C, protect from light

\* The compound is unstable in solutions, freshly prepared is recommended.



### BIOLOGICAL ACTIVITY

<b>Description</b>	Fura-2 AM is a high affinity, intracellular, UV light-excitabile and ratiometric fluorescent Ca <sup>2+</sup> indicator.
<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>Fura-2 AM diffuses across the cell membrane and is de-esterified by cellular esterases to yield Fura-2 free acid.</p> <ol style="list-style-type: none"> <li>1. First, prepare the 1 mM Fura-2 AM stock by adding 50 µL of DMSO to a 50 µg vial. It is important to use dry DMSO packed under nitrogen and it is necessary to remove the DMSO with a needle by puncturing the septum to prevent hydration of the DMSO. After preparing the Fura-2 AM solution keep it in a dark dry place. Fura-2 AM in DMSO is stable at RT for 24 hours and is stable at -20 degrees in a dry container for several months.</li> <li>2. Aliquot 2 mL of culture media into a 15 mL conical tube, warm to 37 deg. and add 2 µL of Fura-2 AM stock to generate a 1µM Fura-2 AM solution. Vortex the solution vigorously for 1 min.</li> <li>3. Transfer the loading solution to a 35 mm tissue culture dish and transfer the coverslip with the cells into the dish.</li> <li>4. Incubate the neurons at 37 degrees for 30 minutes in a dark incubator. Time the incubation precisely.</li> <li>5. Prepare a 35 mm dish containing 2 mL of tissue culture media without Fura-2 AM. Remove the coverslip from the loading solution and place in the new dish.</li> <li>6. Mount the coverslip on the imaging chamber.</li> </ol> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p>

### CUSTOMER VALIDATION

- Elife. 2021 Mar 24;10:e64830.
- Biochem Biophys Res Commun. 2020 Dec 17;533(4):1276-1282.
- bioRxiv. 2021 Jan 13.

See more customer validations on [www.MedChemExpress.com](http://www.MedChemExpress.com)

### REFERENCES

- [1]. Odmara L Barreto-Chang, et al. Calcium imaging of cortical neurons using Fura-2 AM. J Vis Exp. 2009 Jan 19;(23):1067.

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

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