

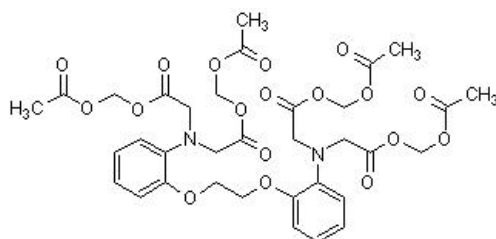
Product datasheet

BAPTA-AM, Ca²⁺ chelator ab120503

7 References 1 Image

Overview

Product name	BAPTA-AM, Ca ²⁺ chelator
Description	Selective Ca ²⁺ chelator. Analog of BAPTA.
Biological description	Selective Ca ²⁺ chelator. Cell-permeable analog of BAPTA (ab120449). Useful for manipulation of intracellular free Ca ²⁺ levels. Shows varied biological activity. Blocks hK _v 1.5, K _v 11.1 (hERG) and hK _v 1.3 channels (K _i values are 1.23, 1.30 and 1.45 μM, respectively).
Purity	> 97%
CAS Number	126150-97-8
Chemical structure	



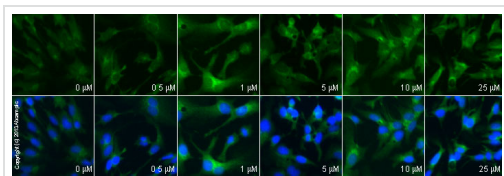
Properties

Chemical name	1,2-Bis(2-aminophenoxy)ethane- <i>N,N,N',N'</i> -tetraacetic acid tetrakis(acetoxymethyl ester)
Molecular weight	764.68
Molecular formula	C ₃₄ H ₄₀ N ₂ O ₁₈
Storage instructions	Store at -20°C. Store under desiccating conditions. The product can be stored for up to 12 months.
Solubility overview	Soluble in DMSO to 100 mM
Handling	<p>Wherever possible, you should prepare and use solutions on the same day. However, if you need to make up stock solutions in advance, we recommend that you store the solution as aliquots in tightly sealed vials at -20°C. Generally, these will be useable for up to one month. Before use, and prior to opening the vial we recommend that you allow your product to equilibrate to room temperature for at least 1 hour.</p> <p>Refer to SDS for further information.</p> <p>For more information on AM esters please visit our AM esters FAQ page.</p> <p>Need more advice on solubility, usage and handling? Please visit our frequently asked questions (FAQ) page for more details.</p>

Source

Synthetic

Images



Functional Studies - BAPTA-AM, Ca²⁺ chelator
(ab120503)

[ab66705](#) staining PAI1 in HepG2 cells treated with BAPTA-AM (ab120503), by ICC/IF. Increase in PAI1 expression correlates with increased concentration of BAPTA-AM, as described in literature. The cells were incubated at 37°C for 4 hours in media containing different concentrations of ab120503 (BAPTA-AM) in DMSO, fixed with 100% methanol for 5 minutes at -20°C and blocked with PBS containing 10% goat serum, 0.3 M glycine, 1% BSA and 0.1% tween for 2h at room temperature. Staining of the treated cells with [ab66705](#) (5 μg/ml) was performed overnight at 4°C in PBS containing 1% BSA and 0.1% tween. A DyLight 488 goat anti-rabbit polyclonal antibody ([ab96899](#)) at 1/250 dilution was used as the secondary antibody. Nuclei were counterstained with DAPI and are shown in blue.

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