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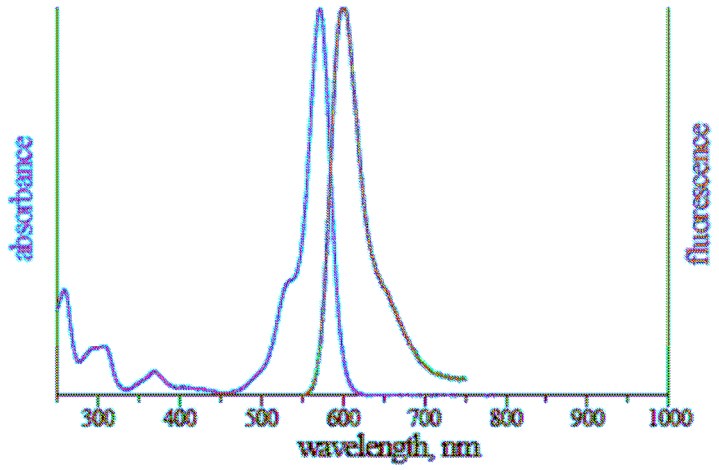
72763 Atto Rho12 maleimide

Application
Atto Rho12 is a new rhodamine dye for application in life sciences, e.g. labelling of DNA, RNA or proteins. The label shows strong absorption, high fluorescence quantum yield, and high photostability. After coupling to a substrate Atto Rho12 carries a net electrical charge of +1.
The dye is moderately hydrophilic. Atto Rho12 consists of a mixture of three isomers with practically identical absorption and fl uorescence properties.

Product Description

MW	872 g/mol
λ_{abs}	576 nm
ε_{max}	1.2 x 10^5 M^{-1} cm^{-1}
λ_{fl}	601 nm
η_{fl}	80 %
τ_{fl}	4.0 ns
CF_{260}	0.27
CF_{280}	0.09

Optical data of the carboxy derivative (in water):



Storage: store at ≤ -20°C. Protect from long-term exposure to moisture and light.

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General procedure for labelling proteins with maleimides

1) Dissolve the protein at 50–100 µM in a suitable buffer at pH 7.0–7.5 at room temperature. Common buffers include 10–100 mM phosphate, Tris, HEPES. Under those conditions, the protein thiol groups are sufficiently nucleophilic so that they react almost exclusively with the reagent. Other protein amines mostly remain protonated and relatively unreactive.
2) Redue disulfide bonds in the protein. A 10-fold molar excess of a reducing agent such as DTT (43817) or TCEP (93284) is usually sufficient. If DTT is used, then dialysis is required to remove the excess DTT prior to introducing the reactive dye. This is not necessary for TCEP.
3) As thiols can be oxidized to disulfides, It may be advisable to carry out thiol modifications in an oxygen-free environment. This is particularly important if the protein has been treated with a reagent such as dithiothreitol prior to thiol modification. In this case, all buffers should be deoxygenated and the reactions carried out under an inert atmosphere to prevent reformation of disulfides.
4) Prepare a 10–20 mM stock solution of the reactive dye in a suitable solvent immediately prior to use (DMSO is the most common choice). Protect all stock solutions from light as much as possible by wrapping containers in aluminum foil.
5) Add sufficient protein-modification reagent from a stock solution to achieve an 10–20 molar excess compared to protein. Add the reagent dropwise to the protein solution as it is stirring .
6) Let the reaction proceed for 2 hours at room temperature or overnight at 4°C. In both cases reaction should take place in the dark.
7) Upon completion of the reaction with the protein, an excess soluble low molecular weight thiol (e.g. glutathione, mercaptoethanol) can be added to consume excess thiol-reactive reagent, thus ensuring that no reactive species are present during the purification step.
8) Separate the conjugate on a gel filtration column, such as a Sephadex G-25 column or equivalent matrix, or by extensive dialysis at 4°C in an appropriate buffer.