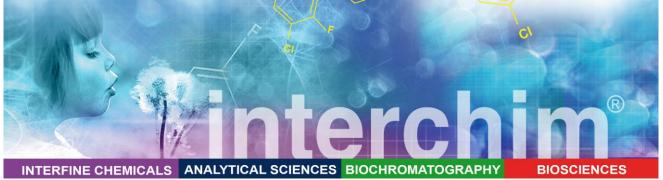


FT- 51254A



FluoProbes[®] Membrane markers FPMM[™] & SynapTracer[™]

Product Information

Product name	MW	$\lambda_{\text{exc}} \backslash \lambda_{\text{em}\bullet}$	mol. abs.	Comments
Cat. number	(g∙mol⁻¹)	max. MetOH (nm) membrane (nm)	(M-1cm-1)	
Green SynapTracer™ 1-1 FP-AM312A, 5mg	527.4	510 / 625 N/A		Shortest lipophilic tail and the most water soluble, thus expected to show the slowest "on-rate" and fastest "off-rate".
FP Membrane Marker 1-43 (Green Synaptracer 2-5) FP-51254A, 1 mg FP- 51254B, 5 x 1 mg FP-51254C, 5 mg	611.56	510 / 625 479 / 598	56 000	Used in synaptic functional studies and vesicle follow up. It is used to study vacuolar organelle morphology and dynamics, the endocytic pathway and vacuole fusion in yeast, endosomal marker and vital stain.Used with Fura-2 or Sulfo Rhodamine101, it has allowed to study membrane turn over and discriminate nonsynaptic labeling.
FP Membrane Marker 1-43 FX (Green SynapTracer [™] 1-4FX) FP-T2982A, 1 mg	560.1	510 / 625 479 / 598	50 000	Contains an amine group that renders it fixable with glutaraldehyde in situ. Ideal if subsequent immunochemistry is desired. Has been used for detection of yeast vacuole membrane staining with FPMM TM 4-64
FP Membrane Marker 3-25 FP-JW6950, 5 mg FP-JW6951, 5 x 1 mg	1226			Larger FPMM [™] 1-43 analog with lower brightness in cuvette studies
FP Membrane Marker 1-44 FX (Green SynapTracer [™] 1- 4BFX) FP-AN100A, 1 mg		N/A		Improved version of FPMM [™] 1-43 FX with better fixability and can be used as a general probe to monitor endocytosis.
FP Membrane Marker 2-10 (Green SynapTracer [™] 1-2) FP-77563A, 5mg	555.45	505 / 620 N/A	50 000	More hydrophilic than FPMM [™] 1-43, thus faster destaining rate. It may be preferred to FPMM [™] 1-43 for quantitative measurements.
FP Membrane Marker 2-10 FX (Green SynapTracer [™] 1-2FX) FP-AM307A, 1 mg	499 55	502 / 625 N/A		Analog of FPMM ^{TM2-10} , but contains an amine group that renders it fixable with glutaraldehyde.

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Product name Cat. number	MW (g·mol⁻¹)	λ _{exc} \λ _{em} . max. MetOH (nm) membrane (nm)	mol. abs. (M-1cm-1)	Comments
Green SynapTracer [™] 1-3 FP-80270A, 5 mg	542	510 / 625 480 / 598		Slightly more hydrophilic that FPMM [™] 1-43 and a lipophilic tail with one carbon shorter. The hydrophilic end is a trimethylammonium group.
RH 414 FP-47009A, 1 mg FP-47009B, 5 mg	581.48	533 / 717	50 500	A water soluble fluorescent probe for membrane labeling, widely used for functional imaging and tracing of neurons (monitoring membrane potential, synaptic activity and ion channel activity of neurons).
FP Membrane Marker 1-84 (Green SynapTracer [™] 1-5) FP-AM3229, 1 mg FP-AM322A, 5 mg FP-AM322B, 5 x1 mg	639.62	511 / 627 N/A		Fluorescent lipophilic tracer for staining and identifying actively firing neurons and investigating the mechanisms of activity-dependent vesicle cycling. A fluorescent nerve terminal dye suitable for monitoring synaptic activity at synapses and at neuromuscular junctions. Less hydrophilic than FPMM TM 1-43, thus faster staining rate but slower destaining rate.
Red SynapTracer™ 3-1 FP- AM323A, 5 mg	579.48	543 / weak N/A		More hydrophilic than FPMM [™] 1-43, thus faster destaining rate, thus a greater "off-rate".
FP Membrane Marker 4-64 (Red SynapTracer [™] 3-2) FP-41109A, 1 mg	607.53	543 / weak 558 / 737	48 000	Most popular red dye of FPMM TM series. It also used to study vacuolar organelle morphology and dynamics, the endocytic pathway and yeast endocytosis mutants. The green FPMM TM 1-43 and red dye FPMM TM 4-64 have become the most used, allowing dual color imaging, to follow synaptic activities at neuromuscular junctions or synapses, as well as in endocytosis vesicules and vacuoles. Fluorescence spectra are similar for all dyes and show 30-40 nm blue shift from polar environment to membrane one.
FP Membrane Marker 4-64 FX (SynaptoRed [™] C2) FP-BJ1011, 1 mg	872.85	558 / 734	46 000	N-CH=CH-CH=C
FP Membrane Marker 5-95 FP-R1422A, 1 mg FP-R1422B, 5 mg	565.43	543 / weak 558 / 737	43 000	Slightly more water soluble than FPMM [™] 4-64

(*) All FPMMTM and SynapTracerTM dyes are non-fluorescent in water.

Storage: Room temperature >1 year. (M)

Protect from light and moisture

Introduction

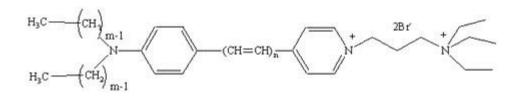
FPMMTM and SynapTracerTM dyes are a range of non toxic cationic styryl dyes, that are non-fluorescent in water but highly fluorescent upon membrane binding and internalization.

FPMMTM and SynapTracerTM dyes are typically formed by a highly hydrophilic, cationic charged, head group and of a lipophilic tail, separated by a linker that contains 1 double bond (giving green fluorescence), or 3 double bonds (giving red fluorescence). The dye name SynapTraceTM n-m indicates the number (n) of double bond and the number (m) carbons in the lipidic tails. Some are available derivatized with an amino group (SynapTracerTM « FX »), that makes the dye fixable in situ with glutaraldehyde. Others have modified chains.

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Applications :

- Terminal nerve and neuro-muscular junction
- Activity-dependent vesicle cycling
- Identifying cell membrane boundaries
- Labeling membranes of living cells
- Morphology or dynamics of endocytosis vesicules and vacuoles

Due to frequent problem of background fluorescence and when repeated washing is not sufficient, some quencher agents are available to remove background fluorescence :

The cyclodextrin ADVASEP-7 forms a water soluble inclusion complex with FPMM TM1-43, thus leaving the dye in the aqueous phase.

SCAS is used with FPMMTM and SynapTracerTM and highly reduce background when it is added to the preparation without the need for repeated washing.

SulfoRhodamine 101 can be used to reduce background staining via FRET

Directions for use

Handling and Storage

Powder can be stored 6 months at room temperature. Solutions of FX analogs, are quite unstable (2 weeks at -20°C). Soluble in water or

DMSO.

Protocol - Green FPMMTM1-43 FX staining protocol on fixated cells

- 1. Mix 40 μ l of FPMMTM1-43 FX (stock solution 400 μ M) with 4 ml of 50 mM K Tyride solution to obtain a final dye concentration of 4 μ M. Place the coverslip in this solution for 1 min at room temperature.
- 2. Transfer the coverslip to Tyrode + 0.5 μ M TTX solution for 1 min at room temperature.
- 3. Transfer the coverslip to quencher solution (SCAS in Tyrode + $0.5 \mu M$ TTX solution) for 4 min at room temperature. Typical concentration of SCAS working solution is 0.5 mM.
- 4. Transfer the coverslip to fixation solution (4% formaldehyde, 4% sucrose, 1µM TTX in PBS) for 20 min at room temperature.
- 5. Transfer the coverslip directly to pre-cooled 0.01% Triton solution for 12 min at $+4^{\circ}$ C.
- 6. Wash 3 times, 1 min with cold PBS.

Other protocol may found in the literature.

Related products

- ADVASEP-7, FP-AM305A
- SCAS, quencher to reduce background with FPMMTM dyes, <u>FP-AM308A</u>
- SulfoRhodamine 101, FP-46999A

- α-bungarotoxin, <u>FP-38034A</u>
- Fura-2 AM, <u>FP-42776C</u>
- Biotin-xx-a-bungarotoxin



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References

- Audrey C., *et al.*, « Using FM1-43 to study neuropeptide granule dynamics and exocytosis », *Methods* 33, 287 (2004) <u>Article</u>
- Beh C.T., *et al.*, « A role for yeast oxysterol-binding protein homologs in endocytosis and in the maintenance of intracellular sterol-lipid distribution », *J. Cell Sci.*, Jun 2004; 117: 2983 (2004) <u>Article</u>
- Brager D H., *et al.*, « Regulation of Exocytosis from Single Visualized GABAergic Boutons in Hippocampal Slices », *J. Neurosci.*, 23, 10475 (2003) <u>article</u>
- Dadsetan S., *et al.*, « Intracellular Ca2+ release triggers translocation of membrane marker FM1-43 from the extracellular leaflet of the plasma membrane into endoplasmic reticulum in T lymphocytes », *J. Biol. Chem*; 10.1074/jbc.M501202200. (2005) <u>Article</u>
- **Drew L.** *et al.*, FM1-43 is a permeant blocker of mechanosensitive ion channels in sensory neurons and inhibits behavioural responses to mechanical stimuli, *Molecular Pain* 3:1 (2007) <u>Article</u>
- Gale J. E., *et al.*, « FM1-43 Dye Behaves as a Permeant Blocker of the Hair-Cell Mechanotransducer Channel », *J. Neurosci.*, **21**, 7013 (2001) <u>Article</u>
- Griesinger C.B., *et al.*, « FM1-43 Reveals Membrane Recycling in Adult Inner Hair Cells of the Mammalian Cochlea », *J. Neurosci.*, **22**, 3939 (2002) <u>Article</u>
- Kilic G., « Exocytosis in Bovine Chromaffin Cells: Studies with Patch-Clamp Capacitance and FM1-43 Fluorescence », *Biophys. J.*, 83, 849(2002) <u>Article</u>
- Lee Jen-Yi, et al., « Mechanisms of cell positioning during *C. elegans* gastrulation », *Development*, 130, 307 (2003) <u>Article</u>
- **Meyers J.** et al., Lighting up the Senses: FM1-43 Loading of Sensory Cells through Nonselective Ion Channels, *The Journal of Neuroscience*, 23(10):4054-4065 (2003) <u>Article</u> (+ FM3-25, and FM 4-64)
- **Parton R. M.**, *et al.*, « Dynamics of the apical vesicle accumulation and the rate of growth are related in individual pollen tubes », *J. Cell Sci.*, **114**, 2685 (2001) <u>Article</u>
- Prévéral S. *et al.*, A common highly-conserved cadmium detoxification mechanism from bacteria to humans. Heavy metal tolerance conferred by the ABC transporter SpHMT1 requires glutathione but not metal-chelating phytochelatins peptides, *J. Biol. Chem.* (2008) <u>Article</u> (FM 4-64)
- **Pyle JL**, *et al.*, « Visualization of synaptic activity in hippocampal slices with FM1-43 enabled by fluorescence quenching. », *Neuron.*, **24**(4):803(1999) <u>Abstract</u>
- Samhan-Arias AK et al., Regionalization of plasma membrane-bound flavoproteins of cerebellar granule neurons in culture by fluorescence energy transfer imaging, *J Fluoresc.* 16(3):393-401 (2006) <u>Abstract (RH-414, FM 4-64)</u>
- Sato H. *et al.*, Differential Columnar Processing in Local Circuits of Barrel and Insular Cortices, *J. Neurosci.*, 28: 3076 3089 (2008) <u>Article</u> (RH-414)
- Walther A., *et al.*, « Polarized Hyphal Growth in *Candida albicans* Requires the Wiskott-Aldrich Syndrome Protein Homolog Wal1p », *Eukaryot. Cell*, **3**, 471 (2004) <u>Article</u>
- Wiederkehr A., *et al.*, « The F-box Protein Rcy1p Is Involved in Endocytic Membrane Traffic and Recycling Out of an Early Endosome in *Saccharomyces cerevisiae* », *J. Cell Biol.*, **149**, 397 (2000) <u>Article</u>

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