



Fluorescent Ion and pH Indicators

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OVERVIEW

Calcium Indicators

To study the role of the important intracellular messenger calcium in regulating various cellular events, it is essential to quantitatively monitor its concentration. The most widely used method of Ca^{2+} detection is by the use of fluorescent Ca^{2+} indicators, a technique pioneered by professor Roger Tsien and colleagues (1). Calcium indicators are fluorophores coupled to a BAPTA calcium chelator structure. They differ in their Ca^{2+} dissociation constants (K_d) or Ca^{2+} response range, excitation/emission wavelengths, spectral shift, and relative fluorescent quantum yields. Therefore, you should select a Ca^{2+} indicator that best suits your needs in consideration of your biological system, instrument settings and any other fluorescent probes that you may use at the same time. The K_d values can give you an estimate of the detectable Ca^{2+} concentration range, usually 0.1 K_d to 10 K_d . However, one should be cautious in using these *in vitro* determined K_d values as the values in cells could differ considerably due to differences in ionic strength, pH, viscosity and Ca^{2+} buffering by cellular lipids and proteins (2).

Other Ion and pH Indicators

Mag-fura-2 (Table 1) can be used as a magnesium indicator or low affinity calcium indicator. We also provide a selection of fluorescent indicators for chloride and zinc (page 5), as well as pH indicators (pages 6-7).

Cell Membrane-Permeable AM Esters

A wide selection of ion indicators are available in both the membrane-impermeant salt forms and the membrane-permeant AM ester forms. The salt forms of the indicators are water-soluble and can be loaded into cells via microinjection. The AM esters of the indicators themselves do not bind ions. However, once they have entered cells, they are readily hydrolyzed by intracellular esterases into the parent ion indicators. AM esters are membrane-permeant and thus can be loaded into cells by simple incubation of the cell or tissue preparation in a buffer containing the AM ester. Biotium also supplies Pluronic F-127, a mild non-ionic detergent that can facilitate cell loading of AM ester compounds (see page 7).

Accessory Products

Biotium also offers calcium ionophores (page 5), as well as calibration buffers, anhydrous solvents and loading agents for AM esters, and other accessory products for ion indicator studies (page 7).

References: 1) Methods in Cell Biology, Vol. 30, 1989. pp. 127-156. 2) Cell Calcium, 1997. 21, 233.

Table 1. Physical properties of calcium indicators

Indicator	MW ¹	Excitation ²	Emission ²	K_d ³	Page
Bis-fura-2 ⁴	779	363/335 nm	512/505 nm	370 nM	3
Fluo-3	770	506 nm	525 nm	390 nM	2
Fluo-4	737	494 nm	506 nm	335 nM	2
Fura-2	642	363/335 nm	512/505 nm	145 nM	3
Indo-1	650	349/331 nm	482/398 nm	230 nM	3
Furaptra (mag-fura-2)	435	369/330 nm	511 nm	1.9 mM (Mg^{2+}) 25 μM (Ca^{2+})	3
Rhod-2	755	556 nm	576 nm	1 μM	4
Rhod-590	912	595 nm	616 nm	610 nM	4

¹ Molecular weights for free acid form of indicators are listed.

² For non-ratiometric indicators, excitation/emission with calcium is shown; for ratiometric indicators, a pair of excitation/emission wavelengths for no calcium/high calcium are shown.

³ Calcium dissociation constant measured at 22 °C in pH 7.2 buffer.

⁴ Bis-fura-2 has similar calcium response as fura-2 but with a 75% larger extinction coefficient.

CALCIUM INDICATORS

Fluo-3

Fluo-3 absorbs at 506 nm and emits at 526 nm when bound to calcium (Figure 1). It is one of the most widely used fluorescent indicators because it can be optimally excited by the commonly-used 488 nm laser line. Fluo-3 is a non-ratiometric indicator; it is essentially non-fluorescent without Ca^{2+} present, but the fluorescence increases at least 40 times on Ca^{2+} binding. Because fluo-3 binds Ca^{2+} more weakly (higher K_d) than fura-2 and indo-1, it is more useful for measuring high transient Ca^{2+} concentration during Ca^{2+} spikes.

References: 1) J. Biol. Chem. 1989. 264, 8171. 2) J. Biol. Chem. 1989. 264, 8179.

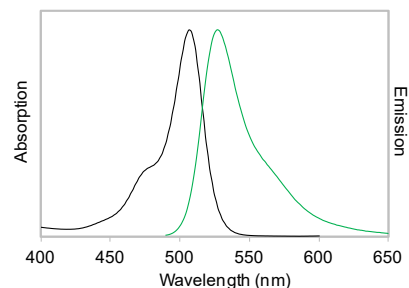
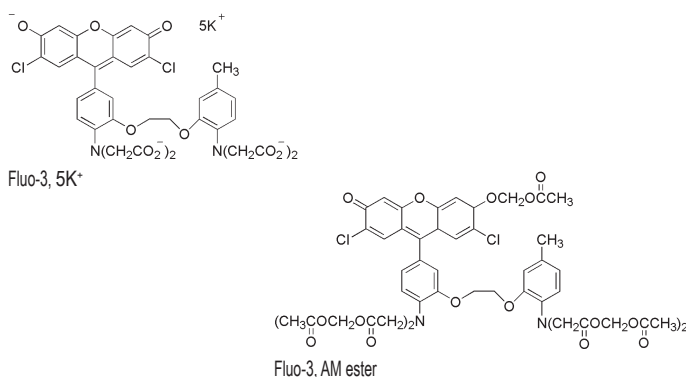


Figure 1. Absorption and emission spectra of fluo-3 saturated with Ca^{2+} .



Fluo-4

Fluo-4 is an analog of fluo-3 with the two chlorine substituents replaced by fluorines, which results in increased fluorescence excitation at 488 nm that gives higher fluorescence signal.

Reference: Cell Calcium. 2000. 27 (2), 97.

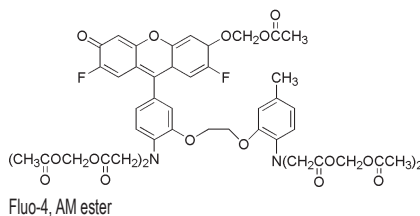


Table 2. Fluo-3 and Fluo-4 products

Cat. #	Product	Unit Size
50010	Fluo-3, 5NH_4^+	1 mg
50011	Fluo-3, 5K^+	1 mg
50012	Fluo-3, 5Na^+	1 mg
50013	Fluo-3, AM ester	10 x 100 μg
50014	Fluo-3, AM ester	1 mg
50015	Fluo-3, AM ester, 1 mM in anhydrous DMSO	1 mL
50016	Fluo-3, AM ester	20 x 50 μg
50018	Fluo-4, AM ester	10 x 50 μg

CALCIUM INDICATORS

Fura-2

Fura-2 is a widely used UV-excitable fluorescent calcium indicator (1-5). Upon calcium binding, the fluorescent excitation maximum of the indicator undergoes a blue shift from 363 nm (Ca²⁺-free) to 335 nm (Ca²⁺-saturated), while the fluorescence emission maximum is relatively unchanged at ~510 nm (Figure 2). The indicator is typically excited at 340 nm and 380 nm and the ratio of the fluorescence emission intensities is used to calculate the Ca²⁺ concentration. Measurement of calcium concentration using ratiometric methods avoids artifacts due to uneven dye distribution and photobleaching (6). Fura-2 has been used in many cellular systems and applications, particularly in microscopic imaging.

Bis-fura-2 has two fura-type fluorophores coupled to one BAPTA chelator. It has a slightly lower affinity for calcium and is membrane impermeable (7-11).

References: 1) J. Mol. Cell. Cardiol. 1992. 24, 937. 2) J. Biol. Chem. 1991. 266, 23739. 3) J. Cell Biol. 1990. 110, 1555. 4) Cell Calcium. 1990. 11, 385. 5) J. Biol. Chem. 1985. 260, 3340. 6) Methods in Cell Biology, Vol. 30. 1989. p. 157. 7) J Neurophysiol. 2000. 84, 2777. 8) Brain Res. 1999. 831, 113. 9) J Neurophysiol. 1999. 81, 2508. 10) J Neurosci Res. 1999. 57, 906. 11) J. Neurochem. 1999. Int 34, 391.

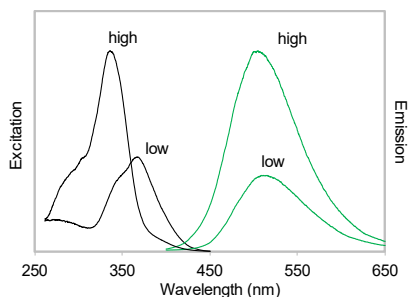
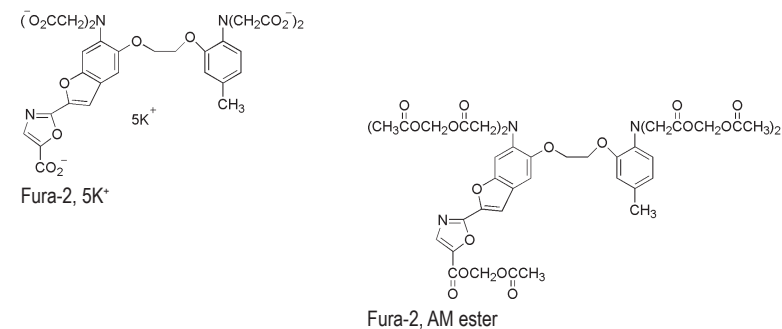


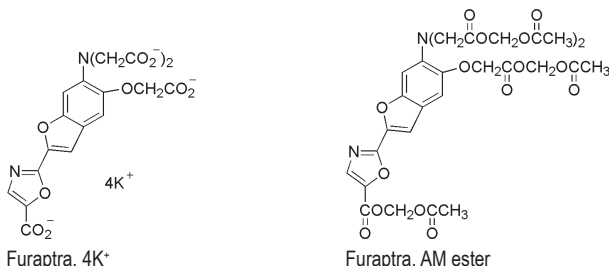
Figure 2. Absorption and emission spectra of fura-2 at high and low calcium concentrations.



Furaptra

Furaptra, which is also commonly called mag-fura-2, was originally developed as a fluorescent magnesium indicator, with a K_d of 1.9 mM for magnesium. However, it also is an excellent indicator for high calcium concentrations in the range of 1 μM to 100 μM (1-2). Fura-2 and furaptra share the same chromophore and as a result have similar spectral change in response to calcium concentration change.

References: 1) Pflügers Arch. 1995. 429, 587. 2) Neuron. 1993. 10, 21.



CALCIUM INDICATORS

Indo-1

Indo-1 is a UV-excitable fluorescent Ca²⁺ indicator that can be optimally excited by the 351-356 nm spectral lines of the argon-ion laser. The emission maximum of the indicator shifts from ~475 nm in Ca²⁺-free medium to ~400 nm in Ca²⁺-saturated medium (Figure 7). Ca²⁺ concentration can thus be determined by measuring the ratio of the fluorescence emission intensities at the two wavelengths. As with fura-2, this ratiometric technique avoids problems associated with uneven dye distribution and photobleaching. Indo-1 has been widely used in flow cytometry studies.

References: 1) J. Biol. Chem. 1985. 260, 3340. 2) Circulation Res. 1991. 69, 46. 3) Am. Heart J. 1990. 120, 590. 4) Cell Calcium. 1990. 11, 487. 5) Meth. Cell Biol. 1994. 41, 150.

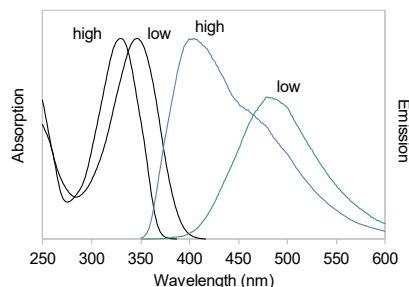
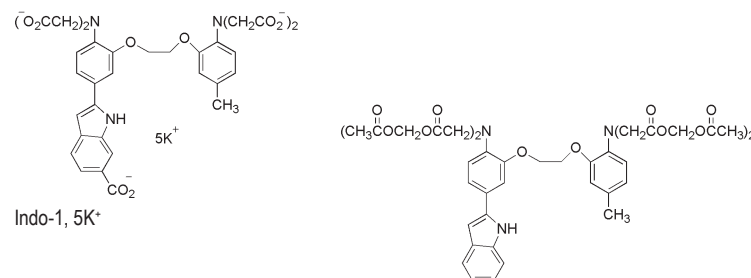


Figure 3. Absorption and emission spectra of indo-1 at high and low calcium concentrations.

Table 3. Fura-2, Furaptra, and Indo-1 products

Cat. #	Product	Unit Size
50030	Fura-2, 5NH ₄ ⁺	1 mg
50031	Fura-2, 5K ⁺	1 mg
50032	Fura-2, 5Na ⁺	1 mg
50033	Fura-2, AM ester	10 x 100 ug
50033-1	Fura-2, AM ester	20 x 50 ug
50034	Fura-2, AM ester	1 mg
50029	Fura-2, AM ester, 1 mM in anhydrous DMSO	1 mL
50045	Bis-Fura-2, 6K ⁺	1 mg
50035	Furaptra, 4K ⁺	1 mg
50036	Furaptra, 4Na ⁺	1 mg
50037	Furaptra, AM ester	10 x 100 ug
50038	Furaptra, AM ester	1 mg
50039	Furaptra, AM ester	20 x 50 ug
50040	Indo-1, 5NH ₄ ⁺	1 mg
50041	Indo-1, 5K ⁺	1 mg
50042	Indo-1, 5Na ⁺	1 mg
50043	Indo-1, AM ester	10 x 100 ug
50043-1	Indo-1, AM ester	20 x 50 ug
50044	Indo-1, AM ester	1 mg

CALCIUM INDICATORS

Rhod-2

Rhod-2 is a fluorescent calcium chelator bearing a rhodamine-like fluorophore. Similar to fluo-3, rhod-2 has excitation and emission spectra that do not undergo significant shift with changing calcium concentration (Figure 4). The indicator is essentially non-fluorescent before Ca^{2+} binding but becomes more fluorescent with increasing Ca^{2+} concentration. Compared to Fluo-3, Rhod-2 is much less fluorescent and has lower fluorescence enhancement with Ca^{2+} .

References: 1) J. Physiol. 1998. 507. 4051. 2) J. Biol. Chem. 1989. 264. 8179.

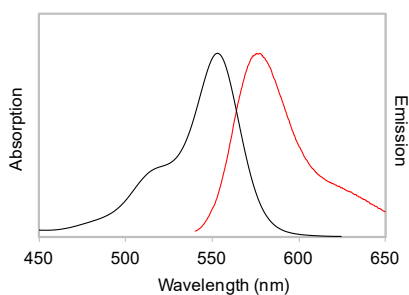
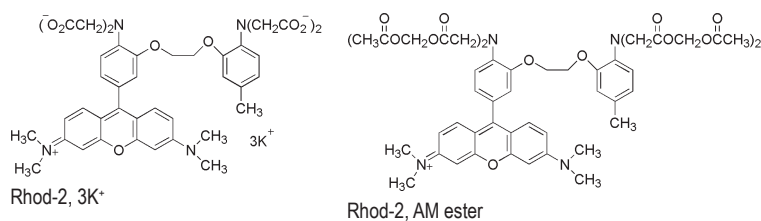


Figure 4. Absorption and emission spectra of rhod-2 saturated with Ca^{2+} .

Rhod-590

Rhod-590 has absorption (595 nm) and emission (616 nm) maxima that are longer than those of fluo-3. Compared to fluo-3, rhod-590 has a smaller fluorescent enhancement with Ca^{2+} .

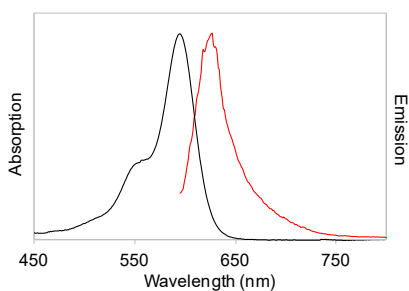
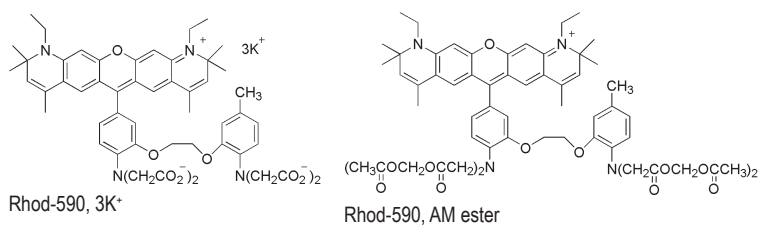


Figure 5. Absorption and emission spectra of rhod-590 saturated with Ca^{2+} .

Table 4. Indo-1, Rhod-2, and Rhod-590 products

Cat. #	Product	Unit Size
50020	Rhod-2, 3NH_4^+	1 mg
50021	Rhod-2, 3K^+	1 mg
50022	Rhod-2, 3Na^+	1 mg
50023	Rhod-2, AM ester	10 x 100 ug
50024	Rhod-2, AM ester	1 mg
50026	Rhod-590, 3K^+	500 ug
50025	Rhod-590, AM ester	10 x 50 ug

CALCIUM CHELATORS

BAPTA Chelators

BAPTA and its derivatives are calcium chelators that are commonly used to form calcium buffers with well-defined calcium concentrations. By injecting the chelators into cells or by incubating cells with the AM ester form of the chelators, one can control the cytosolic calcium concentration, an important means to study the roles of calcium.

Key advantages of these calcium chelators include relative insensitivity toward intracellular pH change and fast release of calcium. Biotium offers several BAPTA chelators with calcium dissociation constants covering the biologically significant range from 10^{-7} to 10^{-2} M.

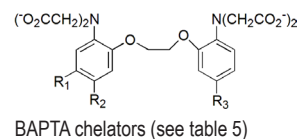


Table 5. Physical properties of BAPTA chelators

Chelator	R ₁	R ₂	R ₃	MW *	K _d (mM) **
BAPTA	H	H	H	477	No Mg^{2+} : 0.59 1 mM Mg^{2+} : 0.70
5,5'-Dibromo BAPTA	H	Br	Br	635	No Mg^{2+} : 3.6
5,5'-Difluoro BAPTA	H	F	F	513	No Mg^{2+} : 0.61 1 mM Mg^{2+} : 0.72
5,5'-Dimethyl BAPTA	H	CH_3	CH_3	505	No Mg^{2+} : 0.16 1 mM Mg^{2+} : 0.44
5-Methyl-5'-nitro BAPTA	H	NO_2	CH_3	536	No Mg^{2+} : 53
5-Mononitro BAPTA	H	NO_2	H	522	No Mg^{2+} : 94
4-Trifluoromethyl BAPTA	CF_3	H	H	545	No Mg^{2+} : 0.57

*Molecular weights for free acid forms are listed

**Cell Calcium 10, 491 (1989).

Table 6. List of BAPTA products

Cat. #	Product	Unit Size
50001	BAPTA, 4Cs^+	1 g
50002	BAPTA, 4K^+	1 g
50003	BAPTA, 4Na^+	1 g
50000	BAPTA, AM ester	25 mg
50000-1	BAPTA, AM ester	20 x 1 mg
50004	5,5'-Dibromo BAPTA, 4K^+	100 mg
50005	5,5'-Difluoro BAPTA, AM ester	25 mg
50006	5,5'-Difluoro BAPTA, 4K^+	100 mg
50007	5,5'-Dimethyl BAPTA, AM ester	25 mg
50008	5,5'-Dimethyl BAPTA, 4K^+	100 mg
50009	5-methyl-5'-nitro BAPTA, 4K^+	10 mg
50017	5-mononitro BAPTA, 4K^+	10 mg

IONOPHORES

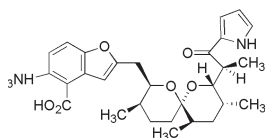
Calcium ionophores

Calcium ionophores are ion carriers that facilitate the equilibration of calcium across cellular membranes. They are useful tools for calibrating calcium indicators and manipulating intracellular calcium concentration. Biotium offers the ionophores listed below.

A-23187

A-23187 (calcimycin, calcium ionophore III) is a calcium ionophore that rapidly equilibrates intracellular and extracellular calcium concentrations. It is commonly used for in situ calibration of fluorescent calcium indicators. The ionophore allows Mn^{2+} to enter the cells and quench intracellular fluorescence of calcium indicators.

References: 1) Anal. Chem. 1989. 61, 382. 2) FASEB J. 1994. 8, 639. 3) Merck Index 12, 1678. 4) Science. 1982. 217, 943.

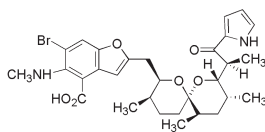


A-23187, free acid

4-Bromo-A23187

The calcium ionophore 4-bromo A-23187 is similar to A-23187. However, because 4-bromo A-23187 is non-fluorescent, it is preferred over the blue fluorescent A-23187 for calibrating UV-excited Ca^{2+} indicators such as indo-1 and fura-2 in order to keep background fluorescence to a minimum.

Reference: Anal. Chem. 1985. 146, 349.

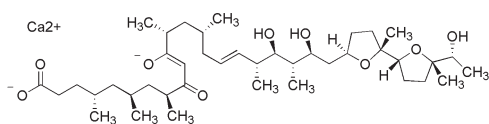


4-Bromo A-23187, free acid

Ionomycin

Ionomycin is commonly used to modify intracellular calcium concentrations and to calibrate fluorescent calcium indicators. Ionomycin is commonly used to stimulate cytokine production and cell proliferation in lymphocytes. In other cell types it can induce apoptosis and inhibit proliferation.

References: 1) J. Biol. Chem. 2000. 275, 7071. 2) J. Immunol. 1995. 155, 3297. 3) J. Urol. 1999. 162, 916.



Ionomycin, calcium salt

Table 7. Calcium ionophore products

Cat. #	Product	Unit Size
59001	A-23187, free acid	1 mg
59006	4-Bromo A-23187, free acid	1 mg
59007	Iononycin, calcium salt	1 mg

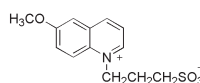
OTHER ION INDICATORS

Chloride indicators

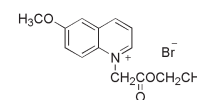
The fluorescence of SPQ 6(-methoxy-N-(3-sulfopropyl)quinolinium) is specifically quenched by chloride via collision. Therefore, chloride concentration is measured by monitoring the degree of fluorescence decrease. The dye can be loaded into cells by hypotonic shock. The dye has excitation/emission at 344/443 nm (1-3).

MQAE (N-(ethoxycarbonylmethyl)-6-methoxyquinolinium bromide) is an improved chloride indicator that has greater sensitivity to chloride ($K_{sv} = 200 M^{-1}$) than SPQ ($K_{sv} = 118 M^{-1}$) and higher fluorescence quantum yield. The ester group of MQAE may slowly hydrolyze inside cells, resulting in a change in its fluorescence response. The dye has excitation/emission at 350/460 nm (4-6).

References: 1) Neurosci. Lett. 1989. 104, 326. 2) Biophys. J. 1989. 56, 1071. 3) J. Biol. Chem. 1991. 266, 20590. 4) Am J Physiol. 1990. 259, C375. 5) Anal Biochem. 1989. 178, 355. 6) Anal Biochem. 1996. 241, 51.



SPQ

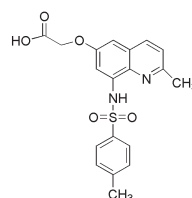


MQAE

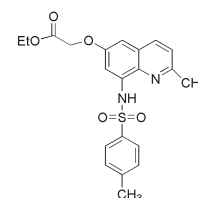
Zinc indicators

Zinc is believed to be involved in the suppression of apoptosis and play important roles in many neural activities. Zinquin is an UV-excitable, blue fluorescent zinc indicator with excitation/emission at 350/460 nm. Zinquin free acid is membrane-impermeant, while Zinquin ethyl ester is membrane-permeable and is hydrolyzed into Zinquin free acid after entering cells.

References: 1) Biochem. J. 1994. 303, 781. 2) Biochem. J. 1993. 296, 403.



Zinquin free acid



Zinquin ethyl ester

Table 8. Other ion indicators

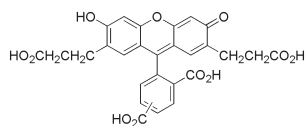
Cat. #	Product	Unit Size
52010	SPQ 6(-methoxy-N-(3-sulfopropyl)quinolinium)	50 mg
52011	MQAE (N-(ethoxycarbonylmethyl)-6-methoxyquinolinium bromide)	100 mg
52022	Zinquin free acid	5 mg
52020	Zinquin ethyl ester	5 mg

pH INDICATORS

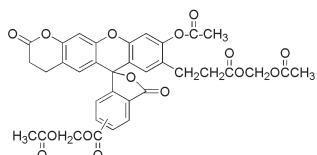
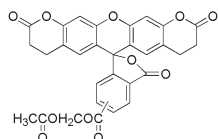
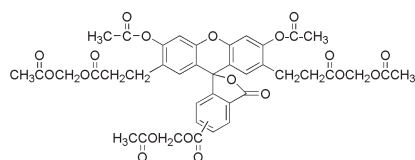
BCECF

BCECF is the most widely used fluorescent pH sensor. With a pK_a of 6.97, close to physiological pH, BCECF can detect changes in cytosolic pH with high sensitivity. At low pH, the dye is weakly fluorescent but becomes more fluorescent with increasing pH. The excitation spectrum of the dye undergoes a slight shift with pH change, while the wavelength of the emission maximum remains unchanged. The pH is determined ratiometrically by the relative fluorescent intensities at 535 nm when the dye is excited at 439 nm and 505 nm. Available in membrane-impermeant free acid and cell-permeable AM ester forms.

References: 1) J. Immunol. Meth. 1988. 108, 255. 2) Biochem. J. 1994. 304, 751. 3) J. Immunol. Meth. 1994. 172, 255.



BCECF



BCECF AM ester (mixture of 3 compounds)

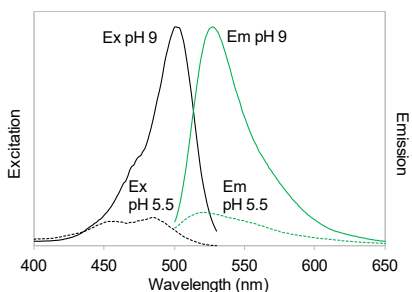


Figure 6. Excitation and emission spectra of BCECF at pH 5.5 and pH 9.0

Carboxyfluorescein derivatives

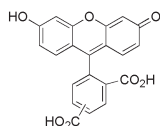
Carboxyfluorescein (FAM) has a pK_a of 6.5 and can be used as a pH indicator or cellular tracer. Its excitation spectrum and fluorescence response to pH are similar to those of BCECF (1). Carboxyfluorescein is also available in membrane permeable diacetate form (CFDA) (2-5). CFDA-SE is a membrane-permeable amine-reactive form that covalently reacts with cytoplasmic proteins for stable labeling (6-8).

5-(and-6)-Carboxy-2',7'-dichlorofluorescein is similar to carboxyfluorescein, but has a pK_a of 4.8, and therefore is useful for detecting pH in more acidic environment (9-11). It is also available in membrane-permeable diacetate, and membrane-permeable, amine-reactive diacetate succinimidyl ester forms.

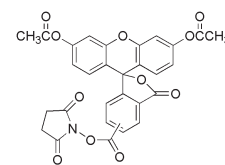
The pK_a of 5-carboxy-2',7'-dichlorosulfonefluorescein is near 4.0 and thus the dye is potentially an excellent pH indicator for acidic organelles.

Fluorescein derivatives are available as single isomer or mixed isomers; the isomers have the same spectral properties. For certain coupling reactions, single isomer may be preferable, but mixed isomers are suitable for most applications.

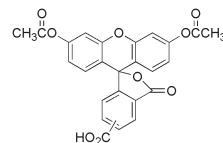
References: 1) Methods Enzymol. 1986. 128, 65. 2) J. Cell Sci. 1991. 98, 343. 3) Nature. 1982. 295, 524. 4) J. Immunol. Meth. 1987. 100, 261. 5) J. Immunol. Meth. 1980. 33, 33. 6) J. Cell Biol. 1985. 101, 610. 7) J. Cell Biol. 1986. 103, 2649. 8) Cell Transplantation. 1994. 3, 397. 9) Biochemistry. 1987. 26, 6330. 10) Biochim. Biophys. Acta. 1989. 985, 75. 11) Anal. Biochem. 1990. 187, 109.



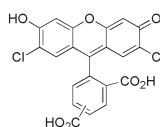
5(6)-FAM



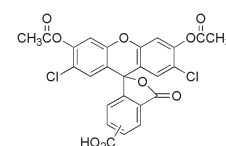
5(6)-CFDA-SE



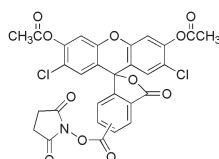
5(6)-CFDA



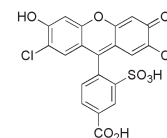
5-(and-6)-Carboxy-2',7'-dichlorofluorescein



5-(and-6)-Carboxy-2',7'-dichlorofluorescein diacetate



5(6)-Carboxy-2',7'-dichlorofluorescein diacetate, SE.



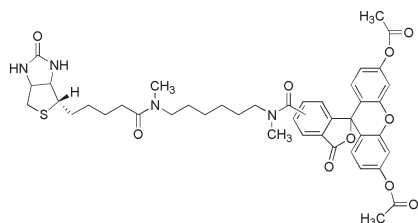
5-Carboxy-2',7'-dichlorosulfonefluorescein

pH INDICATORS

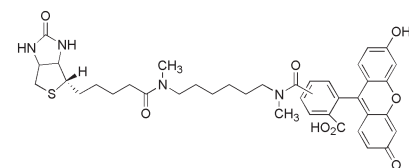
Flubida-2 and Flubi-2

Flubida-2 has been used to detect pH at a specific site in a cell such as cell organelles by directing the probe to where avidin-chimera proteins are located. The probe is a conjugate of biotin and fluorescein diacetate, which is non-fluorescent until the probe has entered the cells and is hydrolyzed by cellular esterases. Flubida-2 is membrane-permeable and can be delivered into cells via simple incubation. Flubi-2 is the membrane-impermeable hydrolyzed product of Flubida-2. Its excitation spectrum and fluorescence response to pH are similar to those of BCECF.

Reference: Chemistry & Biology. 2000. 7, 197.



Flubida-2



Flubi-2

Table 9. pH Indicators

Cat. #	Product	Unit Size
51010	BCECF, free acid	1 mg
51011	BCECF, AM ester	10 x 100 ug
51011-1	BCECF, AM ester	20 x 50 ug
51012	BCECF, AM ester	1 mg
51009	BCECF, AM ester, 1 mg/mL in anhydrous DMSO	1 mL
51019	5-Carboxyfluorescein (5-FAM, single isomer)	100 mg
51020	6-Carboxyfluorescein (6-FAM, single isomer)	100 mg
51013	5-(and-6)-Carboxyfluorescein (5(6)-FAM, mixed isomers)	100 mg
51018	5-CFDA (5-Carboxyfluorescein diacetate, single isomer)	100 mg
51021	6-CFDA (6-Carboxyfluorescein diacetate, single isomer)	100 mg
51014	5(6)-CFDA (5-(and-6)-Carboxyfluorescein diacetate, mixed isomers)	100 mg
90041	5(6)-CFDA, SE (5-(and-6)-Carboxyfluorescein diacetate, succinimidyl ester)	25 mg
51017	5-Carboxy-2',7'-dichlorofluorescein	100 mg
51015	5-(and-6)-Carboxy-2',7'-dichlorofluorescein	200 mg
51016	5-(and-6)-Carboxy-2',7'-dichlorofluorescein diacetate	100 mg
90040	5-(and-6)-Carboxy-2',7'-dichlorofluorescein diacetate, succinimidyl ester (SE)	25 mg
51023	5-Carboxy-2',7'-dichlorosulfonefluorescein	10 mg
51024	Flubi-2	5 mg
51022	Flubida-2	5 mg

ACCESSORY REAGENTS

Calcium Calibration Kit

The Calcium Calibration Kit is used to prepare buffers with a range of accurate calcium concentrations, and is useful for the calibration of fluorescent Ca²⁺ indicators (1-2). The kit contains 50 mL each of zero calcium buffer and 40 uM free Ca²⁺ buffer, with a detailed protocol for combining the two buffers to obtain a Ca²⁺ calibration curve from 0.017-39.8 uM free calcium.

Component A (zero free Ca²⁺):

Zero mM CaEGTA (10 mM K₂EGTA, 100 mM KCl and 10 mM MOPS; pH 7.20)

Component B (40 uM free Ca²⁺):

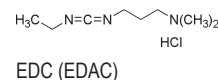
10 mM CaEGTA (10 mM CaEGTA, 100 mM KCl and 10 mM MOPS; pH 7.20)

References: 1) Cell Calcium. 1991. 12, 279. 2) Neuropharmacology. 1995. 34, 1423.

EDC for fixation of indicators

EDC (or EDAC) has recently been found to be very useful for fixing chelators in situ, including the fluorescent ion indicators in this catalog. The fixation of ion indicators makes it useful for post histological studies following the physiological experiments. EDC is also a widely used reagent to activate carboxy groups for amine coupling.

References: 1) Cell Calcium. 1997. 21(3), 175. 2) Tetrahedron Lett. 1993. 34, 7685.

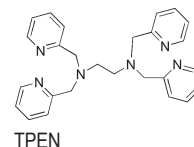


EDC (EDAC)

TPEN heavy metal chelator

TPEN is an excellent heavy metal chelator, but it does not affect Ca⁺, Mg⁺⁺, Na⁺ or K⁺ concentrations. Thus, TPEN is a useful tool to identify effects of heavy metals (Zn⁺⁺, Fe⁺⁺/Fe⁺⁺⁺, Cu⁺⁺, and Mn⁺⁺, etc.) on fluorescent indicators of Ca⁺⁺, Mg⁺⁺, Na⁺ and K⁺.

Reference: J. Biol. Chem. 1985. 260, 2719.



TPEN

Pluronic F-127 for AM ester loading

Pluronic F-127 is a non-ionic detergent that is useful for facilitating loading of AM ester compounds into cells.

References: 1) Science. 1986. 233, 886. 2) J. Biol. Chem. 1987. 262, 12801.

Anhydrous DMSO

Anhydrous Dimethylsulfoxide (DMSO) is recommended for preparing stock solutions of moisture-sensitive AM ester compounds.

Table 10. Accessory products

Cat. #	Product	Unit Size
59100	Calcium Calibration Kit	1 kit
59002	EDC (EDAC)	100 mg
59003	TPEN (tetrakis-(2-pyridylmethyl)ethylenediamine)	100 mg
59000	Pluronic F-127	1 g
59004	Pluronic F-127, 20% in DMSO	1 mL
59005	Pluronic F-127, 10% in H ₂ O	30 mL
90082	DMSO, anhydrous	10 mL



Biotium, Inc.

Toll Free: 800-304-5357

Phone: 510-265-1027

Fax: 510-265-1352

General Inquiries

btinfo@biotium.com

Quotes and Ordering

order@biotium.com

Technical Support

techsupport@biotium.com