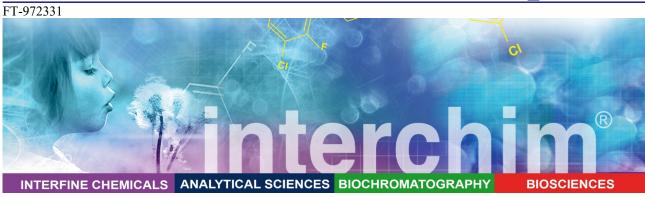
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## **Coelenterazines**

Bioluminescent substrates for bioassays (Ca<sup>2+</sup> measurements, gene reporter assays, ROS assays)

## **Products Information**

Product name cat.number *	MW	R1 £	R2 £	R3 £	λ max. Emission (nm)	Relative Luminescence capacity §	Relative Intensity §	Half-rise Time (s) § (ms) §§
Coelenterazine Native UP972333	423.50	OH	ОН	Phe	465	1.00	1.00	0.4-0.8 6-30ms
Coelenterazine cp UPR30793	415.48	OH	ОН	СР	442	0.95 0.63	20 28	0.15-0.3 5-5ms
Coelenterazine e T8677A	449.50	OH	OH#	Phe	405 and 465	0.5	4	0.15-0.3
Coelenterazine f 438762	425.45	F	ОН	Phe	473	0.80 0.80	18 20	0.4-0.8 6-30ms
e-Coelenterazine f 0C1061	451.46	F	ОН	Phe	413 and 475			
Coelenterazine fcp R47113	417.48	F	ОН	СР	452	0.57	135	0.4-0.8
Coelenterazine h R30783	407.50	Н	ОН	Phe	464	0.82 0.75	10 16	0.4-0.8 6-30ms
Coelenterazine hcp 083534	399.49	Н	ОН	СР	444	0.67 0.65	190 500	0.15-0.3 2-5ms
Coelenterazine i R30801	533.36	Ι	ОН	Phe	476	0.70	0.03	8
Coelenterazine ip R47122	389.45	Ι	ОН	2P	441	0.54	47	1
Coelenterazine n 398194	457.52	Naph	ОН	Phe	467	0.26 0.25	0.01 0.15	5 6-30ms
Coelenterazine 2-methyl T88892	331.37				N/A	N/A	N/A	N/A
Coelenterazine 400a BB8392	391.46				~400	N/A	N/A	N/A
Sample kit 421761	contains $25\mu$ g each of eight coelenterazine analogs: native coelenterazine, coelenterazine <i>cp</i> , coelenterazine <i>f</i> , coelenterazine <i>fcp</i> , coelenterazine <i>h</i> , coelenterazine <i>hcp</i> , coelenterazine <i>ip</i> , and coelenterazine <i>n</i> .							

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\*:other quantities on inquire (250 µg, bulk)

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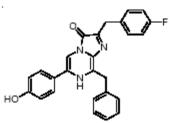


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£: Substituents groups R1, R2 and R3, in positions 2, 6 and 8, are: hydrogen (H), hydroxyl (OH), Phenyl (Phe), CycloPentyl (CP), 2-propionyl (2P), Napthyl (Naph), methyl (Met) <sup>#</sup> Coelenterazine e has a -CH<sub>2</sub>CH- bridge between the 6-phenyl-OH and position 2 of the imidazopyrazinone core.

§ Data from BioChem. J. 261, 913(1989) [normal characters]

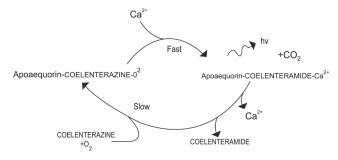
§§ Data from O.Shimoraura in Cell Calcium 14, 373 (1993) for calcium measurements [*smaller size and italic characters*]



### Introduction to Bioluminescence / Coelenterazines

**Bioluminescence** is generation of light by a biochemical reaction involving oxidation of a substrate via an enzyme. This phenomenon has been used extensively in different formats for life science research and drug discovery owing to its extremely high sensitivity, replacing advantagely hazardous methods as radioelement.

Coelenterazine (native form) is a bioluminescent substrate for enzymes apoaequorin and *Renilla* luciferase. Its structure has been investigated <u>ref</u>. Uptima<sup>®</sup> offer include a large selection of coelenterazine analogs, with each of them giving unique luminescent properties (table above). Careful selection of a coelenterazine derivative may be necessary to suit at best your application (see technical information below).



The involved enzymatic complex (~22Kda) contains apoaequorin protein (a protein from *Aequorea victoria* jelly fish and other marine organisms), molecular oxygen, and the luminogen coelenterazine. It releases carbon dioxide and blue light (465 nm) upon oxidation in presence of  $Ca^{2+}$ .

The **apoaequorin** / **Coelenterazine** system has shown privileged advantages for bioassays, notably for measurements of  $Ca^{2+}$  and of Reactive Oxygen Species (ROS).

Examples of bioluminescence applications include:

- Calcium detection in live cells or tissues  $\binom{1-7}{\pm}$
- Superoxide and peroxynitrite anion detection (ROS) (<sup>10-14</sup>)
- Reporter assays (<sup>8</sup>): aequorin has been largely used to tag (by plasmid or other engineering methods) recombinant proteins, and then monitor their expression (localization, regulation...). ±
- ELISA, bioluminescence resonance energy transfer (BRET) for protein interaction studies (9,23)
- Drug high throughput screening.

## **Directions for use**

#### Handling and Storage

All coelenterazines are soluble in MeOH or EtOH (not for Methoxy e-Coelenterazine); DO NOT DISSOLVE IN DMSO. Solution may be stored frozen, protected from light, preferably under dry nitrogen. Solubility in water in rather low. To avoid adverse effects from methanol solutions, aqueous solutions >1mM can be prepared pH 7 buffer containing 50 mM 2-hydroxypropyl-cyclodextrin.(see also ADVASEP and FluoCD<sup>TM</sup> technology).

#### Using coelenterazine / aequorin complex for calcium measurement

Coelenterazine analogs provide various affinities of  $Ca^{2+}$  complexation, and different spectral properties. Compared with fluorescent calcium indicators, aequaporin complex/Coelenterazine system has several advantages in detecting calcium:

• The Ca2+/aequorin complex can detect a broad range of calcium concentrations, from ~0.1 μM to >100 μM,

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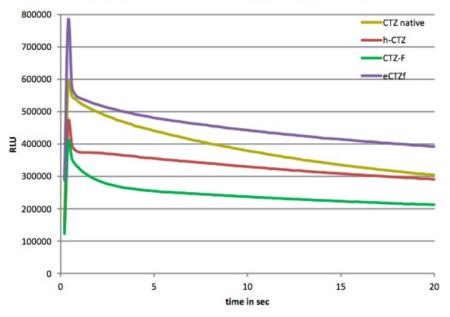
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- Background is lower than with fluorescence, and no autofluorescence of sample occurs,
- Although signal is lower, higher signal/noise ratios can be obtained with imaging equipment,
- The aequorin complex is not exported from cells, allowing to follow calcium concentration changes in cells for hours to days.

Coelenterazine cp, f, h, hcp and n have been used for these applications. Especially, Coelenterazine hcp shows a 500 fold enhancement of light intensity (and thus improved  $Ca^{2+}$  sensitivity) over the native Coelenterazine. Coelenterazine *e* has two emission peaks at 405 and 465 nm, respectively, making it possible to measure calcium concentration via the ratio of emission intensities <sup>17</sup>.

eCoelenterazine-F is a new luciferin utilized by *Renilla* luciferase with an additional ethyl group forming an additional ring system. In addition a Fluoride replace the hydroxyl group on the phenol ring. These changes in the structure result in unique properties in the emission spectrum and luminescent kinetics.



### comparison of kinetics (29.1ng RmLuc)

#### Using coelenterazine / aequorin complex reporter assays

The development of aequoporin vectors prompted many gene reporter assays, as well aequaporin tagged recombinant protein reporter assays. Recombinant proteins can even be targeted in definite cell compartments, for fine measurements.

Uptima<sup>®</sup> offer the largest choice of synthetic coelenterazine derivatives that allow users to choose the ones that suit at best their specific applications. All these coelenterazine analogs are highly purified (purity : >98%) to ensure optimal results in bioassays.

#### Using coelenterazine / BRET

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Standard BRET<sup>1</sup> uses coelenterazine H (UPR30783), that allows the transfer of energy to YFP acceptor (25).

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For BRET<sup>2</sup> application with Coelenterazine 400a (<u>UPBB8392</u>), the substrate for *Renilla luciferase* generates an emission peak centered around 400 nm. It is the preferred Rluc substrate for BRET<sup>2</sup> studies with the GFP acceptor.<sup>23</sup> Coelenterazine 400a is hardly soluble in water, stock solution can be made in ethanol at 1mM (or ~0.4mg/ml) concentration and stored at  $-20^{\circ}$ C in the dark. Avoid using DMSO to dissolve the material as it may cause oxidation. Prepare a 20-fold (50µM) dilution of the stock solution with the recommended buffer <sup>(3,9)</sup> we recommend Dulbecco's phosphate buffered saline containing CaCl<sub>2</sub> (0.1g/l), MgCl<sub>2</sub>.6H<sub>2</sub>O (0.1g/l) and D-glucose(1g/l) supplemented with aprotinin (2µg/ml). The working solution is susceptible to oxidation by air, thus should not be stored. See reference <sup>(Q)</sup>.

For BRET<sup>2</sup> application with Methoxy e-Coelenterazine (Me-O-e-CTZ), the substrate for *Renilla luciferase* generates an emission peak centered around 405 nm.

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Storage and Shelf Life: It is best stored as completely DRY powder under argon in air-tight O-ring plastic tubes at -20°C or for longer storage at -70°C, protected from light.

Usage: It is always best to make FRESH SOLUTIONS immediately before luminometer assays or experiments. 1. Dissolve lyophilized Me-O-e-CTZ in METHANOL as a 1 mg/ml solution (Ethanol won't dissolve Me-O-eCTZ). 2. Use this stock solution to make an aqueous solution in PBS or TBS (e.g. 50  $\mu$ M for luminometer assays equal to 231.8  $\mu$ l (1mg/ml) in 10 ml PBS).

3. Store methanol dissolved Methoxy e-Coelenterazine at -80°C, do not store the aqueous working solution (it will oxidize over time). Me-O-e-CTZ has the same stability in aqueous solutions like any other Coelenterazine analogue.

#### Using coelenterazine / aequorin complex for ROS species

Coelenterazine products are widely used to detect superoxide and peroxynitrite via chemiluminescence (11, 13).

Methyl coelenterazine has been reported to be a superior antioxidant for cells against reactive oxygen species (ROS) such as singlet oxygen and superoxide anion.<sup>14</sup> The coelenterazine derivative is membrane-permeant, nontoxic and highly reactive toward ROS. As oxidative stress is believed to be a mediator of apoptosis,<sup>12</sup> methyl coelenterazine should be another important tool for apoptosis study.

See related products: chromogenic probes (NBT, MTT), fluorescent and luminescent probes (MCLA,... cf FP-38544A).

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## **Technical information**

<b>Coelenterazine Native</b>	UP972331, 50 µg	The standard substrate widely used in many applications.				
C <sub>26</sub> H <sub>21</sub> N <sub>3</sub> O <sub>3</sub> ; MW: 423.5;	UP972333, 1 mg *	Coelenterazine is the luminophore of the native aequorin complex and also the substrate for Renilla luciferase. Bioluminescent detection of calcium concentration is highly sensitive in a broad concentration range $(0.1\mu M \text{ to }>100\mu M)^{1.4}$ . Monitoring of reporter genes (phot gene and luc gene) using coelenterazine is also a major application. Other uses of coelenterazine include bioluminescence resonance energy transfer (BRET) <sup>5,23</sup> and chemiluminescent detection of superoxide anion and peroxynitrite in cells or tissues <sup>6-9</sup> . Coelenterazine native is recommended when a fast regeneration is important.				
<b>Coelenterazine cp</b> C <sub>28</sub> H <sub>23</sub> N <sub>3</sub> O <sub>3</sub> ; MW: 415.48	UPR30791, 50 μg UPR30793, 1 mg *	Coelenterazine $cp$ acquorin complex generates luminescence intensity 15 times higher and has a faster response time to calcium than native coelenterazine does $\frac{20.21.22}{20.21.22}$ .				
<b>Coelenterazine e</b> C <sub>25</sub> H <sub>25</sub> N <sub>3</sub> O <sub>3</sub> ; MW: 449.5	UPT86770, 50 μg *	Coelenterazine <i>e</i> has the highest rate in the <i>in vitro</i> regeneration of aequorin, with two emission peaks at 405 and 465 nm, respectively. This make it possible to measure calcium concentration via the ratio of emission intensities in the range of pCa 5-7 <sup>17</sup> , improving measurement accuracy (correction for variation of coelenterazine concentration). However it do not suit intracellular applications because of poor permeability <sup>19</sup> . It was also found less stable in solution <sup>19</sup> .				
$\begin{array}{c} \hline \textbf{Coelenterazine f} \\ C_{26}H_{20}N_3O_2F; \ MW:457.45, \\ (M) \end{array}$	UP438761, 50 μg UP438763, 1 mg *	Coelenterazine $f$ with aequorin complex gives an almost 20 times higher luminescence intensity than that of native coelenterazine while its emission maximum is about 8 nm longer. Coelenterazine $f$ has been found the most cell permeant <sup>(19)</sup> . It is recommended when a high Ca <sup>2+</sup> sensitivity of regenerated aequorin is needed <sup>(20.21,22)</sup> .				
<b>Coelenterazine fcp</b> C <sub>25</sub> H <sub>24</sub> FN <sub>3</sub> O <sub>2</sub> ; MW:417.5	UPR47110, 50 μg UPR47111, 1 mg *	Coelenterazine <i>fcp</i> is a synthetic derivative of coelenterazine. Its luminescence intensity is 135 times higher than that of native coelenterazine $^{(20.21.22)}$ .				
$\begin{array}{c} \hline \textbf{Coelenterazine h} \\ C_{26}H_{21}N_3O_2; \ MW: \ 407.5; \\ \ (M) \end{array}$	UPR30782, 50 µg UPR30783, 1 mg *	Its luminescence intensity is more than 10 times higher than that of aequorin complex formed from native coelenterazine. Coelenterazine <i>h</i> is also more sensitive to $Ca^{2+}$ . It is also used for reporter assays (Blood, 94.6, 1999, 1899-1905, Blanpain C <u>Article</u> )				
Coelenterazine hcp C <sub>25</sub> H <sub>25</sub> N <sub>3</sub> O <sub>2</sub> ; MW:: 399.5.	UP083532, 50 µg UP083534, 1mg *	It's luminescence intensity is the highest (190 times than that of acquorin complex formed from native coelenterazine) with a fast response time to calcium <sup>(20,21,22)</sup> .				
Coelenterazine i C <sub>26</sub> H <sub>20</sub> IN <sub>3</sub> O <sub>2</sub> ; MW:533.4; 100	UPR30801, 50 μg UPR30803, 1 mg *	Its luminescence intensity is almost 50 times higher than that of native coelenterazine while its response time to calcium is much slower than the latter (20.21.22).				
Coelenterazine ip C <sub>23</sub> H <sub>23</sub> N <sub>3</sub> O <sub>3</sub> ; MW: 389.5	UPR47120, 50 μg UPR47122, 1 mg *	Its luminescence intensity is more than 10 times higher than that of aequorin complex formed from native coelenterazine.				
<b>Coelenterazine n</b> C <sub>30</sub> H <sub>23</sub> N <sub>3</sub> O <sub>3</sub> ; MW: 457.5	UP398192, 50 μg UP398193, 1 mg *	Its luminescence intensity is the weakest of all coelenterazine analogs and its response time to calcium is also much slower than that of native Coelenterazine. Coelenterazine $n$ is reported to be a very useful low-sensitivity coelenterazine.				
<b>2-methyl Coelenterazin</b> C <sub>20</sub> H17N <sub>3</sub> O <sub>2</sub> ; MW: 331.4	UPT88890, 50 μg UPT88891, 1 mg *	Methyl coelenterazine has been reported to be a superior antioxidant for cells against reactive oxygen species (ROS) such as singlet oxygen and superoxide anion 14. It is membrane-permeant, nontoxic and highly reactive toward ROS (potent antioxidant). As oxidative stress is believed to be a mediator o apoptosis, methyl coelenterazine should be important tool for apoptosis study. Biochem. Pharmacol. 60, 471 (2000) ;Immunol. Today 15, 7 (1994) ;Anal. Biochem. 206, 273 (1992) ; Circ. Res. 84, 1203 (1999)				
<b>Coelenterazine 400a</b> C <sub>26</sub> H <sub>21</sub> N <sub>3</sub> O; MW: 391.5, (M)	UPBB8391, 50 µg UPBB8392, 1 mg *	Coelenterazine 400a, also called DeepBlue C is a coelenterazine derivative that serves as a substrate for a Renilla luciferase (Rluc) and generates an emission peak centered around 400nm. It is the best Rluc substrate for BRET studies because it has minimal interference with the emission of GFP acceptor. $\lambda_{abs.}$ 424nm: 3.9 See guidelines for BRET application $\pm$ .				
<b>Methoxy e-</b> <b>Coelenterazine</b> C <sub>29</sub> H <sub>25</sub> N <sub>3</sub> O; MW: 463.54	1J4380, 250µg 1J4381, 500µg 1J4382, 1mg 1J4383, 10mg	13-fold higher luminescent than Coelenterazine-400a. New synthetic analogue of e-Coelenterazine with an additional methoxy group, that works with Renilla Luciferase (RLuc) and Renilla Luciferase 8 (RLuc8) to emit light at ~405 nm.				

\* other quantities on inquire (250  $\mu$ g , bulk)

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#### FT-972331 Related products

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**Pinterchim** 

- Coelenterazine Solvent (Coelenterazine concentration x10 / MeOH or EtOH), 1J4660
- BTA-2 (Dual luciferase assay), FP-QU636A
- Coenzyme A, <u>627374</u>
- D-Luciferin (also a luminescent substrate, used for gene reporter assays with Firefly luciferase) <u>FP-</u><u>M1224A</u>
- Luciferase assay kits with luminescent substrate in solution, <u>FP-BX0320</u>
- Coelenterazines *in vivo* grade (for animal *in-vivo* imaging) <u>FP-BV073A</u>
- UptiFectin-On Transfection Reagent, <u>CK5060</u>

## **Ordering information**

Catalog size quantites and prices may be found at <u>http://www.interchim.com</u> Please inquire for higher quantities (avaibility, shipment conditions). For any information, please ask : Uptima / Interchim; Hotline : +33(0)4 70 03 73 06

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