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<u>Data Sheet</u> PDE10A2 TR-FRET Assay Kit Catalog # 60710

DESCRIPTION: Phosphodiesterases (PDEs) play an important role in dynamic regulation of cAMP and cGMP signaling. PDE10A is a dual substrate PDE highly expressed in striatal medium spiny neurons. PDE10A inhibitors can improve the cognitive symptoms of schizophrenia, and exhibit potential therapeutic value for Huntington's disease. PDE10A2 is located in cytosol, whereas PDE10A2 is a membrane-associated protein.

The *PDE10A2 TR-FRET Assay Kit* is designed for identification of inhibitors of PDE10A2 using TR-FRET (Time Resolved Fluorescence Resonance Energy Transfer) technology. The assay is based on the generation of FAM-labeled nucleotide monophosphates by the phosphodiesterase. These phosphate groups bind to terbium-labeled nanoparticles, resulting in energy transfer from the terbium to the FAM, which emits a fluorescent signal at 520 nm. The change in fluorescent intensity can be easily measured using a fluorescence plate reader.

The PDE10A2 TR-FRET Assay Kit comes in a convenient 96-well format, with purified PDE10A2 enzyme, fluorescently labeled PDE substrate (cAMP), binding agent, and PDE assay buffer for 100 enzyme reactions. Using this kit, only two simple steps on a microtiter plate are required for the PDE10A2 activity assay. First, the fluorescent-labeled cAMP is incubated with a sample containing PDE10A2 for 1 hour. Second, a binding agent and a terbium donor are added to the reaction mix and incubated for 1 hour. Then, fluorescence intensity can be measured using a fluorescence reader.

APPLICATIONS: Great for screening small molecular inhibitors for drug discovery and HTS applications.

STABILITY: 6 months from date of receipt when stored as directed.



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COMPONENTS:

Catalog #	Component	Amount	Ç	Storage
60100	PDE10A2 recombinant enzyme	1 µg	-80°C	
60200	FAM-Cyclic-3', 5'-AMP: 20 µM	50 µl	-80°C	
60393	PDE assay buffer	25 ml	-20°C	
	Tb donor	30 µl	-80°C	(Avoid
60390	Binding Agent	200 µl	+4°C	freeze/ thaw
	Binding Buffer A	20 ml	+4°C	cycles!)
	Binding Buffer B	20 ml	+4°C	
79685	Black, low binding NUNC microtiter	1	Room	
	plate		temp.	

MATERIALS OR INSTRUMENTS REQUIRED BUT NOT SUPPLIED:

Fluorescent microplate reader capable of measuring Time Resolved Fluorescence Resonance Energy Transfer (TR-FRET)

REFERENCE(S):

- 1) Akinori Nishi and Gretchen L. Snyder. *J Pharmacol Sci.*, 2010, **114**: 6 16.
- 2) Kenji Omori and Jun Kotera. Circulation Research. 2007; 100: 309-327.

ASSAY PROTOCOL:

All samples and controls should be tested in duplicate.

Protocol for PDE10A2 assay

Step 1:

- 1) Dilute 20 µM FAM-Cyclic-3',5'-AMP substrate stock solution 100-fold with PDE buffer to make a 200 nM solution. Make only a sufficient quantity needed for the assay; store remaining stock solution in aliquots at -20°C.
- Add 25 μI of FAM-Cyclic-3',5'-AMP (200 nM) to each well designated "Substrate Control", "Positive Control", and "Test Inhibitor". Add 25 μI of PDE assay buffer to each well designated "Tb-only Control".
- 3) Add 5 µl of inhibitor solution to each well designated "Test Inhibitor". Add 5 µl of the same solution without inhibitor (inhibitor buffer) to the "Tb-only Control", "Substrate Control" and "Positive Control".
- 4) Thaw PDE10A2 on ice. Upon first thaw, briefly spin tube containing enzyme to recover the full contents of the tube. Aliquot PDE10A2 enzyme into single-use aliquots. Store remaining undiluted enzyme in aliquots at -80 °C immediately. Note: PDE10A2 is very sensitive to freeze/thaw cycles. Do not re-use thawed aliquots or diluted enzyme.



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5) Dilute PDE10A2 in PDE buffer to 0.6 pg/μl (12 pg/reaction) in PDE buffer*. Add 20 μl of PDE assay buffer to the wells designated as the "Tb-only Control" and "Substrate Control". Initiate reaction by adding 20 μl of PDE10A2 (0.6 pg/μl) to the wells designated for the "Positive Control" and "Test Inhibitor". Discard any remaining diluted enzyme after use. *Note: optimal enzyme concentration may vary with the specific activity of the enzyme.

	Tb only Control	Substrate Control	Positive Control	Test Inhibitor
FAM-Cyclic-3',5'-AMP (200 nM)	_	25 µl	25 µl	25 µl
PDE assay buffer	45 µl	20 µl	_	_
Test Inhibitor	_	I	_	5 µl
Inhibitor Buffer (no inhibitor)	5 µl	5 µl	5 µl	_
PDE10A2 (0.6 pg/µl)	_		20 µl	20 µl
Total	50 µl	50 μl	50 µl	50 μl

6) Incubate at room temperature for 1 hour.

Step 2:

- 1) Make binding dilution buffer by mixing equal volumes of Binding buffer A and Binding buffer B. For example, mix 1 ml Binding buffer A with 1 ml Binding buffer B.
- 2) **Mix binding agent** thoroughly and dilute **binding agent** 1:50 with binding dilution buffer made in Step 1.
- 3) Add Tb donor (1:1,000 dilution) to the mixture in Step 2.
- 4) Add 100 µl to each well. Incubate at room temperature for 1 hour with slow shaking.
- 5) Read the fluorescent intensity in a microtiter-plate reader capable of TR-FRET.

Instrument Settings

Reading Mode	Time Resolved	
Excitation Wavelength	330±20	
Emission Wavelength	490±10	
Lag Time	50 μs	
Integration Time	50 μs	
Excitation Wavelength	330±20	
Emission Wavelength	520±10	
Lag Time	50 μs	
Integration Time	50 μs	

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CALCULATING RESULTS:

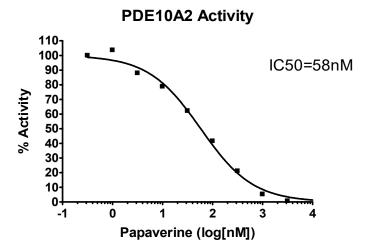
$$FRET = \frac{S_{520} - \left(\frac{Tb_{520}}{Tb_{490}} \times S_{490}\right)}{S_{490}} \times 1000$$

Where S_{520} = Sample 520 nm reading, S_{490} = Sample 490 nm reading, Tb_{520} = Tb only 520 nm reading, Tb_{490} = Tb only 490 nm reading. When percentage activity is calculated, the FRET value from substrate only control can be set as zero percent activity and the FRET value from positive control can be set as one hundred percent activity.

$$\% Activity = \frac{FRET_S - FRET_{Sub}}{FRET_P - FRET_{Sub}} \times 100\%$$

Where $FRET_s = Sample FRET$, $FRET_{Sub} = Substrate only control FRET$, and $FRET_P = Positive control FRET$.

EXAMPLE OF ASSAY RESULTS:



Inhibition of PDE10A2 by Papaverine, measured using the *PDE10A2 TR-FRET Assay Kit*, BPS Bioscience # 60710. *Data shown is lot-specific. For lot-specific information, please contact BPS Bioscience, Inc. at info@bpsbioscience.com*



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RELATED PRODUCTS:

Cat.#	Size
60099	10 µg
60100	10 µg
60101	10 µg
60102	5 µg
60110	10 µg
60011	10 µg
60012	10 µg
	5 µg
	96 rxns.
60705	96 rxns.
60701	96 rxns.
	60099 60100 60101 60102 60110 60011 60012 60046 60300 60400 60411 60311 60312 60345 60704 60705