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# Produktinformation

- Forschungsprodukte & Biochemikalien
- Zellkultur & Verbrauchsmaterial
- Diagnostik & molekulare Diagnostik
- Laborgeräte & Service

Weitere Information auf den folgenden Seiten!
See the following pages for more information!

## Lieferung & Zahlungsart

siehe unsere Liefer- und Versandbedingungen

## Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand


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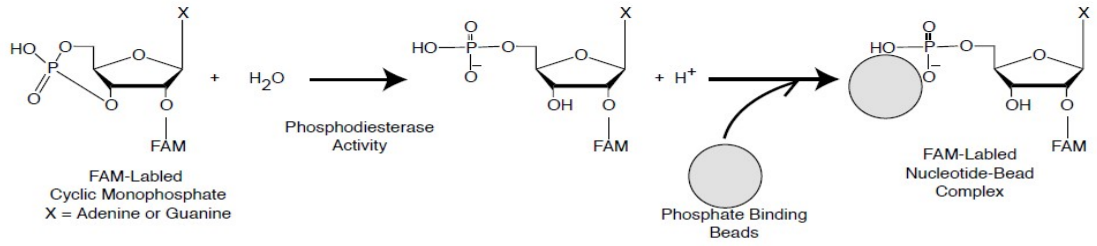
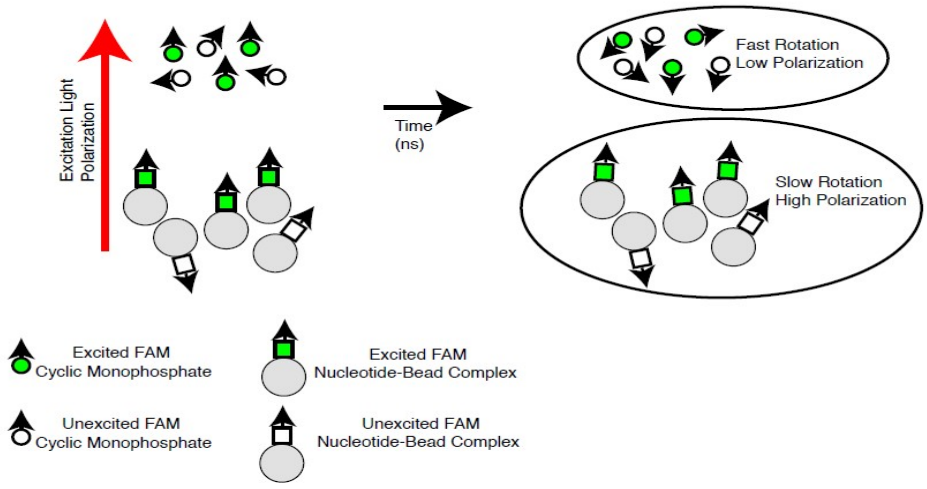
# Data Sheet
# *PDE4A1A Assay Kit*
**Catalog # 60340**

**DESCRIPTION:** Phosphodiesterases (PDEs) play an important role in dynamic regulation of cAMP and cGMP signaling. PDE4A1A is widely expressed in brain tumors and promotes their growth and its inhibitor Rolipram is identified as a potent anti-inflammatory agent. The *PDE4A1A Assay Kit* is designed for identification of PDE4A1A inhibitors using fluorescence polarization. The assay is based on the binding of a fluorescent nucleotide monophosphate generated by PDE4A1A to the binding agent.

Phosphodiesterases catalyze the hydrolysis of the phosphodiester bond in dye-labeled cyclic monophosphates. Beads selectively bind the phosphate group in the nucleotide product. This increases the size of the nucleotide relative to unreacted cyclic monophosphate. In the polarization assay, dye molecules with absorption transition vectors parallel to the linearly-polarized excitation light are selectively excited. Dyes attached to the rapidly-rotating cyclic monophosphates will obtain random orientations and emit light with low polarization. Dyes attached to the slowly-rotating nucleotide-bead complexes will not have time to reorient and therefore will emit highly polarized light.



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The PDE4A1A inhibitor screening assay kit comes in a convenient 96-well format, including purified PDE4A1A enzyme, fluorescently labeled PDE4A substrate (cAMP), binding agent, and PDE assay buffer for 100 enzyme reactions. The key to the *PDE4A1A Assay Kit* is the specific binding agent. Using this kit, only two simple steps on a microtiter plate are required for PDE4A1A reactions. First, the fluorescently labeled cAMP is incubated with a sample containing PDE4A1A for 1 hour. Second, a binding agent is added to the reaction mix to produce a change in fluorescent polarization that can then be measured using a fluorescence reader equipped for the measurement of fluorescence polarization.

**COMPONENTS:**

| Catalog # | Component | Amount | Storage | |
|---|---|---|---|---|
| 60040 | PDE4A1A recombinant enzyme | 1 µg | -80°C | ***(Avoid freeze/ thaw cycles!)*** |
| 60200 | FAM-Cyclic-3', 5'-AMP: 20 µM | 50 µl | -80°C | |
| 60393 | PDE assay buffer | 25 ml | -20°C | |
| 60390 | Binding Agent | 100 µl | +4°C | |
| 60391 | Binding Agent Diluent (cAMP) | 10 ml | +4°C | |
| 79685 | Black, low binding, microtiter plate | 1 | Room temp. | |

**MATERIALS OR INSTRUMENTS REQUIRED BUT NOT SUPPLIED:**
Fluorescent microplate reader capable to measure fluorescence polarization

**APPLICATIONS:** Great for studying enzyme kinetics and screening small molecular inhibitors for drug discovery and HTS applications.

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

**REFERENCE(S):** Goldhoff P., et al. *Clin Cancer Res*. 2008; **14(23):**7717-25.

**ASSAY PROTOCOL:**
All samples and controls should be tested in duplicate.

**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 sufficient quantity needed for the assay; store remaining stock solution in aliquots at -20°C.

2) Add 25 µl of FAM-Cyclic-3',5'-AMP (200 nM) to each well designated for the "Substrate Control", "Positive Control", and "Test Inhibitor". Add 25 µl of PDE assay buffer to each well designated for the "Blank".

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3) Add 5 µl of inhibitor solution of each well designated for “Test Inhibitor”. Add 5 µl of the same solution without inhibitor (Inhibitor buffer), to the “Positive Control”, “Substrate Control”, and “Blank”.

4) Add 20 µl of PDE assay buffer to the wells designed for the "Blank" and "Substrate Control”.

5) Thaw PDE4A1A on ice. Upon first thaw, briefly spin tube containing enzyme to recover full content of tube. Aliquot PDE4A1A enzyme into single use aliquots. Store remaining undiluted enzyme in aliquots at -70°C immediately. *Note: PDE4A1A is very sensitive to freeze/thaw cycles. Do not re-use thawed aliquots or diluted enzyme.*

6) Dilute PDE4A1A in PDE buffer to 4 pg/µl (80 pg/reaction) in PDE buffer*. Initiate reaction by adding 20 µl of diluted PDE4A1A to the well 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.*

7) Incubate at room temperature for 1 hour.

| | Blank | 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 | – | – | – | 5 µl |
| Inhibitor Buffer (no inhibitor) | 5 µl | 5 µl | 5 µl | – |
| PDE4A1A (4 pg/µl) | – | – | 20 µl | 20 µl |
| Total | **50 µl** | **50 µl** | **50 µl** | **50 µl** |

**Step 2:**

1) Shake the tube containing the binding agent to ensure it is thoroughly mixed. Mix **binding agent** thoroughly and dilute **binding agent** 1:100 with the cAMP binding agent diluent.

2) Add 100 µl diluted binding agent to each well. Incubate at room temperature for 30 minutes with slow shaking.

3) Read the fluorescent polarization of the sample in a microtiter-plate reader capable of excitation at wavelengths ranging from 485 ± 5 nm and detection of emitted light ranging from 528 ± 10 nm. Blank value is subtracted from all other values.

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**CALCULATING RESULTS:**

**Definition of Fluorescence Polarization:**

$$P = \frac{I_{II} - I_{\perp}}{I_{II} + I_{\perp}}$$

where $I_{\parallel}$ = Intensity with polarizers parallel and $I_{\perp}$= Intensity with polarizers perpendicular. Most instruments display fluorescence polarization in units of mP.

$$mP = \left(\frac{I_{II} - I_{\perp}}{I_{II} + I_{\perp}}\right) x\ 1000$$

The equation above assumes that light is transmitted equally well through both parallel and perpendicular oriented polarizers. In practice, this is generally not true and a correction must be made to measure the absolute polarization state of the molecule. This correction factor is called the "G Factor".

$$mP = \left(\frac{I_{II} - G(I_{\perp})}{I_{II} + G(I_{\perp})}\right) x\ 1000 \qquad \textbf{OR} \qquad mP = \left(\frac{G(I_{II}) - I_{\perp}}{G(I_{II}) + I_{\perp}}\right) x\ 1000$$

The G-factor is instrument-dependent and may vary slightly depending upon instrument and conditions. Please check the manual of your instrument to obtain the information about the establishment of the G-factor.

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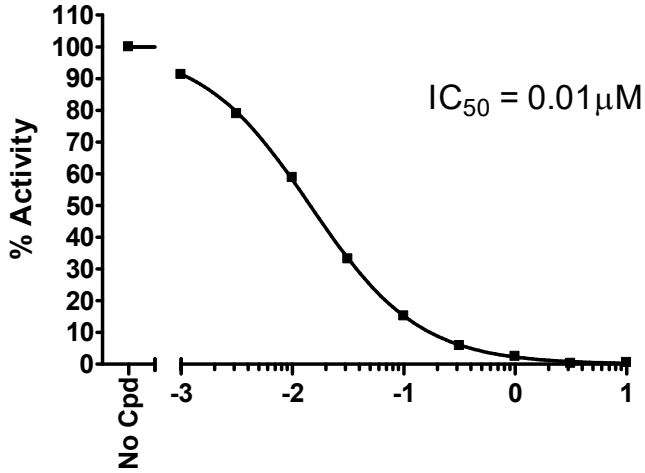
**EXAMPLE OF ASSAY RESULTS:**

**PDE4A1A Activity**

Substrate Conc. = 100nM (cAMP)




Inhibition of PDE4A by Apremilast, measured using the *PDE4A Assay Kit*, BPS Bioscience # 60340. Fluorescence polarization was measured at 528 nm using a Tecan M1000 fluorescent microplate reader. *Data shown is lot-specific. For lot-specific information, please contact BPS Bioscience, Inc. at info@bpsbioscience.com*

**RELATED PRODUCTS :**

| Product | Catalog # | Size |
|---|---|---|
| PDE4A1A | 60040 | 10 µg |
| PDE4A4B | 60039 | 10 µg |
| PDE4B1 | 60041 | 10 µg |
| PDE4B2 | 60042 | 5 µg |
| PDE Assay Kit | 60300 | 96 rxns. |
| PDE1B Assay Kit | 60311 | 96 rxns. |
| PDE2A Assay Kit | 60320 | 96 rxns. |
| PDE3A Assay Kit | 60330 | 96 rxns. |
| PDE3B Assay Kit | 60331 | 96 rxns. |
| PDE4D Assay Kit | 60345 | 96 rxns. |
| PDE4B2 Assay Kit | 60343 | 96 rxns. |
| PDE10A Assay Kit | 60400 | 96 rxns. |
| Apremilast | 27735 | 5 mg |