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Data Sheet Fluorogenic DPP8 Assay Kit

Catalog #: 80208

DESCRIPTION: Dipeptidyl peptidase-8 (DPP8) is a ubiquitously expressed serine exopeptidase that cleaves X-proline dipeptides from the N-terminus of polypeptides. DPP8 shares homology and substrate recognition patterns with DPP4 and like DPP4, is thought to play a role in T-cell activation and immune function. The *Fluorogenic DPP8 Assay Kit* is designed to measure DPP8 activity using purified DPP8 for screening and profiling applications. It comes in a convenient 96-well format, with purified DPP8 enzyme, DPP substrate, and DPP assay buffer for 100 enzyme reactions. The key to the *Fluorogenic DPP8 Assay Kit* is the fluorogenic substrate. Using this kit, only one simple step on a microtiter plate is required for DPP8 reactions. The fluorometric substrate is incubated with a sample containing DPP8 enzyme to produce a fluorophore that can then be measured using a fluorescence reader.

COMPONENTS:

Catalog #	Component	Amount	Storage				
80080	DPP8 human recombinant enzyme	5 μg	-80℃				
80300	DPP assay buffer	10 ml	-20℃	Augid			
80305	Fluorogenic DPP substrate 1 in DMSO	0 100 μl -80 °C Avoid		freeze/			
	(0.5 mM)	•		thaw			
	AMC Fluorescent standard (50 μM)	500 μl	-20℃	cycles!			
	black, low binding NUNC black	1 plate	Room	cycles:			
	microtiter plate	·	temp.				

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

REFERENCES:

1. Lee, H.J. et al., *J. Biol. Chem.* **281 (50):** 38653-38662 (2006)

2. Yu, D.M. et al., *FEBS J.* **273 (11):** 2447-2460 (2006)

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ASSAY PROTOCOL:

Immediately prior to assay:

- 1) Dilute DPP substrate 1 0.5 mM stock 5-fold with DPP assay buffer to make a 100 μM solution. (Make only sufficient quantity needed for the assay; store remaining 0.5 mM stock solution in aliquots at -20 °C.)
- 2) Dilute DPP8 in DPP assay buffer to 2 ng/µl (20 ng/reaction)*. Aliquot any remaining enzyme and store undiluted at -80 °C. Keep diluted enzyme on ice. Discard any remaining diluted enzyme after use. *Note: optimal enzyme concentration may vary with the specific activity of the enzyme.
- 3) Dilute 25 μ I of the Fluorescent AMC standard (50 μ M stock) 2-fold with DPP assay buffer to make a 25 μ M solution. Make serial 2-fold dilutions of the fluorescent AMC standard in DPP buffer as follows: 12.5 μ M, 6.25 μ M, 3.12 μ M, 1.56 μ M, 0.78 μ M, 0.39 μ M, 0.19 μ M, 0.10 μ M. Aliquot the remaining 50 μ M AMC standard and store undiluted at -20 °C.

Step 1:

In duplicate, add the reaction mixtures (below) to the microtiter black plate. Incubate at $22 \, ^{\circ}\text{C}$ for 10 min.

	Enzyme Positive Control	Test Inhibitor	AMC Standard Curve	Inhibitor Negative Control	"Blank" Negative Control
DPP8 (2 ng/μl)	10 μΙ	10 μΙ	ı	ı	ı
DPP substrate 1 (100 μM)	5 μΙ	5 μΙ	_	5 μΙ	-
AMC standard (0.1 μM – 50 μM)	-	-	5 μΙ	-	-
Inhibitor (in DPP assay buffer)	-	Xμl	-	ХμΙ	_
DPP assay buffer	85 μl	85 - X μl	95 μl	95 - X μl	100 μΙ
Total	100 μΙ	100 μΙ	100 μΙ	100 μΙ	100 μΙ

Step 2:

Read sample in a microtiter-plate fluorimeter that is capable of excitation at wavelengths ranging from 350-380 nm and detection of emitted light ranging from 440-460 nm.

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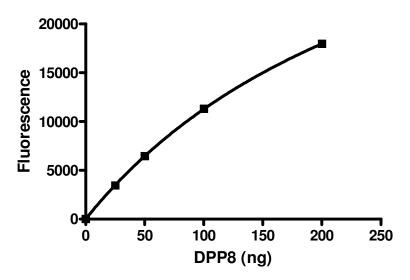


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Example of Assay Results:



DPP8 enzyme activity, measured using the *Fluorogenic DPP8 Assay Kit*, BPS Bioscience Cat.# 80208. *Note: Data shown is lot-specific. For lot-specific information, please contact BPS Bioscience, Inc. at* info@bpsbioscience.com.

RELATED PRODUCTS:

Product Name	Catalog #	<u>Size</u>
DPP8 enzyme	80080	10 μg
DPP assay buffer	80300	20 ml
Fluorogenic DPP substrate 1	80305	100 μl
DPP3 enzyme	80030	10 μg
DPP4 enzyme	80040	10 μg
DPP7 enzyme	80070	10 μg
DPP9 enzyme	80090	10 μg
FAP enzyme	80100	10 μg
POP enzyme	80105	20 μg
Fluorogenic DPP3 assay kit	80203	96 rxns
Fluorogenic DPP4 assay kit	80204	96 rxns
Fluorogenic DPP7 assay kit	80207	96 rxns
Fluorogenic DPP9 assay kit	80209	96 rxns
Fluorogenic FAP assay kit	80210	96 rxns
Fluorogenic POP assay kit	80106	96 rxns
Fluorogenic DPP substrate 2	80332	100 µl

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