

**S-2238™**

For Laboratory Use Only

For General Laboratory Use

**S-2238™**

S-2238 is a chromogenic substrate for thrombin.

**COMPOSITION**

Each vial contains chromogenic substrate S-2238 25 mg and mannitol 120 mg as a bulking agent.

**PRECAUTIONS AND WARNINGS:****Hazard class:** None**Hazard statements:** None**Precautionary statements:** None

**Supplemental Hazard Information:** Up to 2.5% of the mixture consists of component of unknown acute toxicity (oral, dermal, inhalation) for the human health and unknown hazard to the aquatic environment.

**CHEMISTRY**

**Chemical name:** H-D-Phenylalanyl-L-pipecolyl-L arginine-p-nitroaniline dihydrochloride.

**Formula:** H-D-Phe-Pip-Arg-pNA · 2 HCl

**Mol. wt:** 625.6

$\epsilon_{316 \text{ nm}^{-1}}$ :  $1.27 \cdot 10^4 \text{ mol}^{-1} \cdot \text{L} \cdot \text{cm}^{-1}$

**Solubility:** > 10 mmol/L in H<sub>2</sub>O

**Stability:** Substance: Stable until expiry date if stored at 2-8°C.

Avoid exposure to light.

The substance is hygroscopic and should be stored dry.

Solution: 1 mmol/L in H<sub>2</sub>O is stable for more than 6 months at 2-8°C. Contamination by microorganisms may cause hydrolysis.

**Suitable stock solution:**

1-2 mmol/L in H<sub>2</sub>O.

**PRINCIPLE**

## Enzyme

H-D-Phe-Pip-Arg-pNA  $\xrightarrow{\text{Enzyme}}$  H-D-Phe-Pip-Arg-OH+pNA

The method for the determination of activity is based on the difference in absorbance (optical density) between the pNA formed and the original substrate. The rate of pNA formation, i.e. the increase in absorbance per second at 405 nm, is proportional to the enzymatic activity and is conveniently determined with a photometer.

**CHROMOGENIX**

## KINETIC DATA

*Human thrombin:*  $K_m = 0.7 \cdot 10^{-5}$  mol/L  
 $V = 1.7 \cdot 10^{-7}$  mol/min · NIH-U

*Bovine thrombin:*  $K_m = 0.9 \cdot 10^{-5}$  mol/L  
 $V = 2.2 \cdot 10^{-7}$  mol/min · NIH-U

Both determined at 37°C in 2.5 mL 0.05 mol/L Tris buffer pH 8.3, I 0.15.

## STANDARDIZATION

An activity of  $\Delta A/\text{min} = 0.05$  (37°C) is obtained by using 0.1 mmol/L substrate and:

1. 0.03 NIH-U/mL of bovine thrombin (Roche or Parke-Davis)
2. 0.04 U/mL of human thrombin (MRC standard thrombin)

Aprotinin (Trasylol®) may be added to the buffer in a concentration of 75 KIU/L in order to inhibit other activities than that of thrombin.

**Note:** For thrombin standardization against the MRC-standard the natural substrate fibrinogen, is recommended as the primary substrate. The clotting and amidolytic activities of degraded thrombins do not always develop in parallel.

## APPLICATIONS

The substrate has been used for the determination of:

1. Prothrombin in plasma (1,2)
2. Antithrombin in plasma (3,4)
3. Platelet factor 3 in plasma (5,6)
4. Heparin in plasma (7)



1. AXELSSON G et al.: Prothrombin determination by means of chromogenic peptide Substrate. *Thromb Haemost* 36, 517 (1976).
2. BERGSTRÖM K & EGBERG N: Determination of plasma prothrombin using a synthetic chromogenic substrate. Paper read at the 2nd European Congress on Clinical Chemistry, October 1976.
3. ABILGAARD et al.: Antithrombin (heparin cofactor) assay with "new" chromogenic substrates. *Thromb Res* 11, 549-553 (1977).
4. Chromogenix AB: Determination of antithrombin-heparin cofactor in plasma. Laboratory Instruction.
5. SANDBERG H & ANDERSSON L O: A highly sensitive assay of platelet factor 3 using a chromogenic substrate. *Thromb Res* 14, 113-124 (1979).
6. SANDBERG H et al.: Determination of platelet factor 3 in whole blood by a chromogenic peptide substrate assay. *Thromb Res* 18, 871-882 (1980).
7. Chromogenix AB: Determination of heparin in plasma. Laboratory Instruction.

**CHROMOGENIX**

## ***S-2238***

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## ***LANGUAGES***

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ENGLISH

## ***TECHNICAL SPECS***

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PAPER: White paper,  
50-60 g/m<sup>2</sup> weight.

SIZE: 4.1 x 5.9" (104 x 150 mm.).

PRINT: Front/Back.

PRINT COLOR: All type in black.