

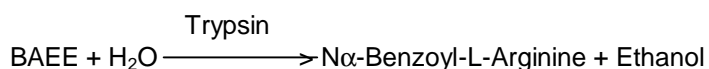


SIGMA QUALITY CONTROL TEST

## Product Information

### Enzymatic Assay of TRYPSIN INHIBITOR<sup>1</sup>

#### PRINCIPLE:



Abbreviation used:

BAEE = N $\square$ -Benzoyl-L-Arginine Ethyl Ester

**CONDITIONS:** T = 25°C, pH = 7.6, A<sub>253nm</sub>, Light path = 1 cm

**METHOD:** Continuous Spectrophotometric Rate Determination

#### REAGENTS:

- A. 67 mM Sodium Phosphate buffer, pH 7.6 at 25°C  
(Prepare 100 ml in deionized water using Sodium Phosphate, Monobasic, Anhydrous, Sigma Prod. No. S-0751. Adjust to pH 7.6 at 25°C with 1 M NaOH.)
- B. 0.25 mM N $\square$ -Benzoyl-L-Arginine Ethyl Ester Solution (BAEE)  
(Prepare 50 ml in Reagent A using N $\alpha$ -Benzoyl-L-Arginine Ethyl Ester, Hydrochloride, Sigma Prod. No. B-4500.)
- C. 1 mM Hydrochloric Acid Solution (HCl)  
(Prepare 50 ml in deionized water using concentrated Hydrochloric Acid, Sigma Prod. No. H-7020.)
- D. Trypsin Enzyme Solution (Trypsin)  
(Immediately before use, prepare a solution containing 1 mg protein/ml of Trypsin, Prod. No. T-8003, in cold Reagent C.)
- E. Trypsin Inhibitor Solution (Inhib)  
(Immediately before use, prepare a solution containing 1.0 mg/ml of Trypsin Inhibitor in cold Reagent A.<sup>2,3</sup>)

## Enzymatic Assay of TRYPsin INHIBITOR<sup>1</sup>

### PROCEDURE:

Pipette (in milliliters) the following reagents into suitable containers:

Part A:

	<u>Uninh</u>	<u>Test1</u>	<u>Test2</u>	<u>Test3</u>	<u>Test4</u>	<u>Test5</u>
Reagent E (Inhib)	---	0.10	0.15	0.20	0.25	0.30
Reagent D (Trypsin)	0.50	0.50	0.50	0.50	0.50	0.50
Reagent C (HCl)	9.50	9.40	9.35	9.30	9.25	9.20

Allow to stand at 25°C for a minimum of five minutes and no longer than six minutes.

Mix by inversion and pipette (in milliliters) the following reagents into suitable cuvettes:

Part B:

	<u>Uninh</u>	<u>Test1</u>	<u>Test2</u>	<u>Test3</u>	<u>Test4</u>	<u>Test5</u>	<u>Blank</u>
Reagent B (BAEE)	3.00	3.00	3.00	3.00	3.00	3.00	3.00
Reagent C (HCl)	0.10	0.10	0.10	0.10	0.10	0.10	0.20

Mix by inversion and equilibrate to 25°C. Monitor the  $A_{253nm}$  until constant, using a suitably thermostatted spectrophotometer. Then add:

Uninh (Part A)	0.10	---	---	---	---	---	---
Test 1 (Part A)	---	0.10	---	---	---	---	---
Test 2 (Part A)	---	---	0.10	---	---	---	---
Test 3 (Part A)	---	---	---	0.10	---	---	---
Test 4 (Part A)	---	---	---	---	0.10	---	---
Test 5 (Part A)	---	---	---	---	---	0.10	---

Immediately mix by inversion and record the increase in  $A_{253nm}$  for approximately 5 minutes. Obtain the  $\Delta A_{253nm}/\text{minute}$  using the maximum linear rate for the Tests, Blank, and Uninhibited Solution.

### CALCULATIONS:

Trypsin Activity in BAEE units/ml enzyme =

$$\frac{(\Delta A_{253nm}/\text{min Test} - \Delta A_{253nm}/\text{min Blank})(df)(10.0)}{(0.001)(0.10)(0.5)}$$

df = Dilution factor

0.001 = The change in  $A_{253nm}/\text{minute}$  per unit of Trypsin at pH 7.6 at 25°C in a 3.2 ml reaction mix

0.10 = Volume (in milliliters) enzyme used (Part B)

10.0 = Total volume in milliliters of assay (Part A)

0.5 = Volume (in milliliters) of enzyme used (Part A)

## Enzymatic Assay of TRYPSIN INHIBITOR<sup>1</sup>

### CALCULATIONS: (continued)

$$\text{Units/mg solid} = \frac{\text{units/ml enzyme}}{\text{mg solid/ml enzyme}}$$

Plot the Trypsin activity (BAEE units/mg protein) vs ml of Trypsin Inhibitor/RM

Mg Trypsin Inhibitor = (ml of Trypsin Inhibitor)(Conc. of Trypsin Inhibitor, mg/ml)

$$\text{Mg Trypsin Inhibited by 1 mg Trypsin Inhibitor} = \frac{\text{mg Trypsin/RM (normalizing factor)}}{\text{mg Trypsin Inhibitor (from plot)}}$$

Normalizing Factor = (BAEE Units of Uninhibited Trypsin per mg solid/10,000 BAEE units of Trypsin per specification.)

### UNIT DEFINITION:

One trypsin unit will produce a  $\Delta A_{253\text{nm}}$  of 0.001 per minute with BAEE as substrate at pH 7.6 at 25°C in a reaction volume of 3.2 ml.

### FINAL ASSAY CONCENTRATION:

In a 3.20 ml reaction mix, the final concentrations are 63 mM sodium phosphate, 0.23 mM BAEE, 0.002 mM HCl, 0.005 mg trypsin, and 0.0003 - 0.001 mg trypsin inhibitor.

### NOTES:

1. This enzyme assay is used to assay Sigma Prod. Nos. T-9003, T-9008, T-9128, T-9253, T-2011, T-1886, T-4385, T-9378, and T-0256.
2. When assaying Trypsin Inhibitor, Ovoinhibitor, Prod. No. T-1886, the diluent used is 200 mM sodium phosphate, monobasic, pH 7.6 at 25°C.
3. When assaying Trypsin Inhibitor, Type II-S, Sigma Prod. No. T-9128, prepare a solution containing 0.60 mg/ml of Trypsin Inhibitor in cold Reagent A.
4. The uninhibited Trypsin activity should be within 85% of the release value for activity. With a 11,700 to 13,005 Trypsin units/mg solid per label, the acceptable range for activity of the uninhibited Trypsin reaction should be 10,000 to 15,300 Trypsin units/mg solid. This range should also correspond to a corrected  $\Delta \text{Abs}_{253\text{nm}}$ /minute of 0.0545 to 0.0835. With this rate and an inhibition of 20% to 80% the  $\Delta \text{Abs}_{253\text{nm}}$ /minute should be above the spectrophotometric rate detection limit of 0.0020.

## Enzymatic Assay of TRYPSIN INHIBITOR<sup>1</sup>

**NOTES:** (continued)

4. Where Sigma Product or Stock numbers are specified, equivalent reagents may be substituted.

**Sigma warrants that the above procedure information is currently utilized at Sigma and that all Sigma-Aldrich, Inc. products conform to the information in this and other Sigma-Aldrich, Inc. publications. Purchaser must determine the suitability of the information and product(s) for their particular use. Additional terms and conditions may apply. Please see reverse side of the invoice or packaging slip.**