

# SIGMA QUALITY CONTROL TEST

# **ProductInformation**

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

#### PRINCIPLE:

Abbreviation used:

BAEE = N□-Benzoyl-L-Arginine Ethyl Ester

**CONDITIONS:** T = 25°C, pH = 7.6,  $A_{253nm}$ , 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□-Benzoyl-L-Arginine Ethyl Ester Solution (BAEE) (Prepare 50 ml in Reagent A using Nα-Benzoyl-L-Arginine Ethyl Ester, Hydrochloride, Sigma Prod. No. B-4500.)
- C. 1 mM Hydrochloric Acid Solution (HCI) (Prepare 50 ml in deionized water using concentrated Hydrochloric Acid, Sigma Prod. No. H-7020.)
- 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.)
- 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>)

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## PROCEDURE:

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

### Part A:

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

Mix by inversion and equilibrate to  $25^{\circ}$ C. Monitor the A<sub>253nm</sub> 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  $\Delta A_{253nm}$ /minute using the maximum linear rate for the Tests, Blank, and Uninhibited Solution.

# **CALCULATIONS:**

Trypsin Activity in BAEE units/ml enzyme =

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

df = Dilution factor

0.001 = The change in A<sub>253nm</sub>/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)

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CALCULATIONS:	(continued)
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Units/mg solid = 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)

Mg Trypsin Inhibited by 1 mg Trypsin Inhibitor = 

mg Trypsin/RM (normalizing factor)

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  $\Delta A_{253nm}$  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 HCI, 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 ΔAbs<sub>253nm</sub>/minute of 0.0545 to 0.0835. With this rate and an inhibition of 20% to 80% the ΔAbs<sub>253nm</sub>/minute should be above the spectophotometric rate detection limit of 0.0020.

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**NOTES:** (continued)

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

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