



# Collagenase Activity Colorimetric Assay Kit

rev 07/20

(Catalog # K792-100; 100 assays; Store at -20°C)

## I. Introduction:

Collagenase (EC 3.4.24.3) is an enzyme in the matrix metalloproteinase family that breaks down collagen, assisting in degradation of the extracellular matrix, a key step in the pathogenesis of bacteria. Collagen is an abundant structural protein present in the connective tissue of animals. Collagenase has been used clinically for the treatment of Dupuytren's contracture, an affliction characterized by a thickening of connective tissue. **BioVision's Collagenase Activity Assay Kit** provides a quick and easy way to determine activity of Collagenase. The Kit measures collagenase activity using a synthetic peptide (FALGPA) that mimics collagen's structure. It is suitable for measuring activity of bacterial collagenases such as from *Clostridium histolyticum* type I-XI. In addition, it can also be used to screen/characterize collagenase inhibitors. The limit of detection for this assay is 0.02 mU Collagenase.

#### II. Applications:

- Measurement of collagenase activity
- For screening/studying/characterizing collagenase inhibitors

#### III. Sample Type:

Purified Collagenase, bacterial extract

#### IV. Kit Contents:

Components	K792-100	Cap Code	Part Number
Collagenase Assay Buffer	20 ml	WM	K792-100-1
Collagenase	1 ml	Red	K792-100-2
Collagenase Substrate (FALGPA)	4 ml	NM	K792-100-3
Inhibitor (1,10-Phenanthroline) [1 M]	50 µl	Yellow	K792-100-4

## V. User Supplied Reagents and Equipment:

- 96-well clear plate with flat bottom
- Multi-well spectrophotometer

## VI. Storage Conditions and Reagent Preparation:

Store kit at -20°C, protected from light. Briefly centrifuge small vials prior to opening. Read entire protocol before performing the assay.

- Collagenase Assay Buffer: Bring to room temperature before use. Store at -20°C or 4°C.
- Collagenase, Collagenase Substrate (FALGPA), and Inhibitor: Ready to use as supplied. Aliquot and store at -20°C. Avoid repeated freeze/thaw. Keep on ice during use. Stable for two months.

#### VII. Collagenase Assay Protocol:

1. Sample Preparation: Dissolve test collagenase in cold dH<sub>2</sub>O or Hank's Balanced Salt Solution (HBSS). Bacterial extract can be prepared in cold PBS. Suggested range of collagenase for measuring is 0.02-10 mU. Add 2-10 µl of test collagenase or bacterial extract into desired well(s) in 96-well plate and adjust the volume to 100 µl with Collagenase Assay Buffer. For positive control, add 10 µl of provided Collagenase. For Inhibitor Control, add 10 µl of provided Collagenase and 2 µl of Inhibitor (1,10-Phenanthroline) into desired well(s). Adjust the volume of positive control and inhibitor control wells to 100 µl with Collagenase Assay Buffer. For reagent background control, add 100 µl of Collagenase Assay Buffer.

## Notes:

a) For unknown collagenase activity, we suggest testing several amounts of collagenase or bacterial extract to ensure the activity is within the assay range.

b) High activity samples will consume substrate within 3 min. Dilute enzyme and remeasure if necessary.

2. Inhibitor Screening (optional): To test Collagenase inhibitors, dissolve test inhibitor to 100X in an appropriate solvent. Add 2 μl of test inhibitor and 10 μl of provided Collagenase into Test Inhibitor well(s). Prepare a parallel well as Enzyme Control (EC) by adding 10 μl of provided Collagenase. Adjust the volume of Test Inhibitor and Enzyme Control wells to 100 μl with Collagenase Assay Buffer. Incubate at room temperature for 10 min.

### Note:

Prepare a solvent control well by adding the same volume of solvent as the test inhibitor to test its effect on Collagenase activity. Provided Inhibitor (1,10-Phenanthroline) does not require a solvent control.

3. Reaction Mix: Prepare enough Reaction Mix for the number of wells (test collagenase, positive control, inhibitor control, reagent background, test inhibitor, and Enzyme Control) to be analyzed. For each reaction, prepare 100 µl reaction mix:

Collagenase Substrate (FALGPA)	40 µl
Collagenase Assay Buffer	60 µl

Mix and add 100  $\mu$ l of the reaction mix into each well, mix well.

4. Measurement: Immediately measure absorbance kinetically at OD 345 nm in a microplate reader at 37°C for 5-15 min. Low activity samples can be measured for 1-3 hr.





5. Calculation: Take the absorbance  $(A_{345nm1} \text{ and } A_{345nm2})$  at two time points  $(T_1 \text{ and } T_2)$  in the linear range. There should be at least two readings in between and at least 1 min. apart.

To determine Activity, use the following equation:

Sample Collagenase Activity = 
$$\frac{\left(\frac{-\Delta A_{345 \text{ nm}}}{\Delta T}\text{Test} - \frac{-\Delta A_{345 \text{ nm}}}{\Delta T}\text{Reagent Background}\right)X(0.2) \text{ X DF}}{(0.53) \text{ X V}} \text{ U/ml}$$

Where:  $\Delta A_{345nm}$  = Difference between  $A_{345 nm2}$  and  $A_{345 nm1}$   $\Delta T$  = Difference between  $T_2$  and  $T_1$ 0.2 = Reaction volume (mI) DF = Dilution Factor 0.53 = millimolar extinction coefficient of FALGPA V = Enzyme volume (mI)

For inhibitor screen, calculate percent inhibition using the following equation:



Figures: (a) Collagenase activity over 10 min. (b) Enzyme activity of collagenase and inhibition by 10 mM (1,10)-Phenanthroline. Assay was performed following the kit protocol.

### VIII. RELATED PRODUCTS:

Collagen-I, human recombinant (4796) Collagen-III, human recombinant (4797) MMP-1 Antibody (5781) MMP-9 Antibody (3529, 3969, 5565) MMP-11 Antibody (3531R) MMP-1 Inhibitor Screening Kit (K794) MMP-8 (rat) ELISA Kit (K4743) MMP-2 Antibody (5562) MMP-9, human recombinant (7789) MMP-17 Antibody (3537)

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