

Collagen Assay Kit (Fluorometric)

LS-K259-100 (100 Tests) • See Storage Conditions Below



Introduction

COLLAGEN is the key structural protein of connective tissue and the most abundant protein in mammals. It occurs in many different types and forms with Types I -V being the most common. Aside from the crucial role it plays in the body, it has numerous medical applications such as its use in reconstructive surgery including bone and skin grafts. It is also commonly used in cosmetics due to its anti-aging and skin healing properties. Assay Methods available for quantifying collagen currently range from needing extensive hydrolysis procedures with acids and bases to using expensive antibodies and complicated protocols. This collagen assay kit delivers a very simple, safe, and sensitive Method to quantify collagen in samples. In the first step of this procedure, collagen in the sample is enzymatically digested into peptides. Subsequently, the N-terminal glycine containing peptides react with the dye reagent to form a fluorescent complex. The fluorescence intensity of this product, measured at $\lambda_{ex/em} = 375/465$ nm, is directly proportional to collagen concentration in the sample.

Key Features

- Safe. Non-radioactive assay.
- Fast and sensitive. Linear detection range (20 μ L sample): 2 μ g/mL to 50 μ g/mL collagen in 96-well plate assay.
- Convenient and high-throughput. "Add-mix-read" type assay. No wash and reagent transfer steps are involved. Can be readily automated for processing thousands of samples per day.

Applications

- Direct Assays: Collagen in biological samples. Collagen in cosmetic products.

Components

Component	K259-100
	100 Tests
Dye Reagent	5 mL
Buffer	5 mL
Digest Enzyme	70 μ L
Collagen Standard (3 mg/mL)	40 μ L
Developer	1 mL

Materials Not Supplied

Pipetting devices, centrifuge tubes, black flat-bottom 96-well plates, and plate reader.

Storage

The kit is shipped on ice. Store kit components (except Collagen Standard) at -20°C upon receiving. Collagen Standard is stored at 4°C. Shelf life: 6 months after receipt.

FOR RESEARCH USE ONLY! Not for use in humans.

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Assay Procedure

Sample Preparation

This assay is optimized for detection of very low levels of collagen. Due to the assay's sensitivity and the abundance of collagen, large dilution factors are often needed to place samples within the linear detection range of this kit. A serial dilution to determine the optimal dilution factor for a type of sample is highly recommended.

Biological fluid samples (e.g. serum, cell lysate, tissue lysate) first centrifuge to remove any particulates. A serial dilution may be needed to determine the optimal dilution factor for each type of sample in this assay.

Procedure Using 96-Well Plate

- Standards. Prepare 50 µg/mL Collagen Standard by mixing 5 µL 3 mg/mL Collagen Standard and 295 µL dH₂O. Next prepare standards in 1.5-mL centrifuge tubes as described in the Table.

No	50 µg/mL Standard + H ₂ O	Collagen (µg/mL)
1	100 µL + 0 µL	50
2	60 µL + 40 µL	30
3	30 µL + 70 µL	15
4	0 µL + 100 µL	0

- Transfer 20 µL Standards into separate wells of a black, flat-bottom 96-well plate. Transfer 20 µL of each sample in duplicate into separate wells (one well as "Sample" and one well as "Sample Blank").
- Prepare sufficient Digest Mix for the Standards and the "Sample" wells by mixing, for each well: 35 µL Buffer and 0.5 µL Digest Enzyme.
Add 30 µL Digest Mix to the Standards and the "Sample" wells.
Add 30 µL Buffer to the "Sample Blank" wells.
Tap plate to mix briefly and thoroughly. Cover plate and incubate 60 minutes at 37°C.
- Add 40 µL Dye Reagent to all wells. Incubate 10 minutes at 37°C.
- Add 8 µL Developer to all wells. Incubate 10 minutes at 37°C.
- Read Fluorescence at $\lambda_{\text{ex/em}} = 375/465$ nm.

Calculations

Subtract the blank value (#4) from the standard values and plot the ΔF against standard concentrations. Determine the slope and calculate the collagen concentration of Sample,

$$[\text{Collagen}] = \frac{F_s - F_B}{\text{Slope } (\mu\text{g/mL}^{-1})} \times n \quad (\mu\text{g/mL})$$

where F_s and F_B are fluorescence readings of the Sample and Sample Blank, respectively. n is the sample dilution factor.

Note: if the sample F value is higher than F for the 50 µg/mL Collagen Standard, dilute sample in water and repeat the assay. Multiply the results by the dilution factor.

Conversions: 50 µg/mL equals 5mg/dL, or 50 ppm.

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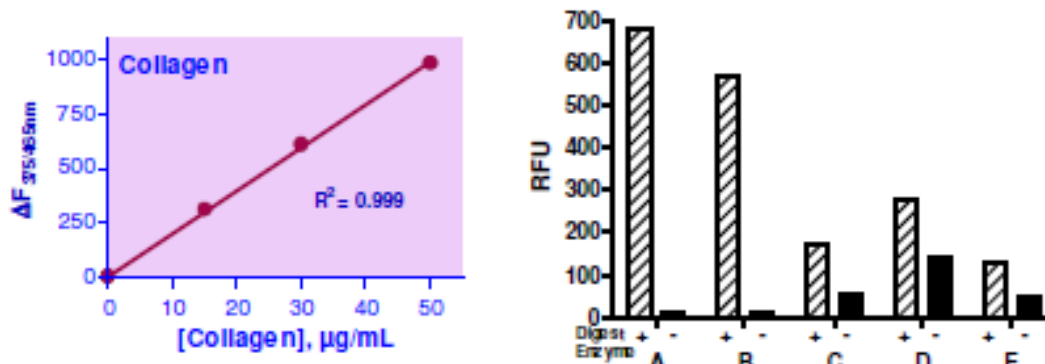
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Sample Data



Collagen Assay in 96-Well Plate

Left: Standard curve in 96-well plate assay

Right: Collagen Samples with (+) and without (-) Digest Enzyme. A) Collagen Standard 30 $\mu g/mL$. B) Essential Wholesale Marine Collagen, 250-fold diluted. C) Rat Serum, 20-fold diluted (2.7 mg/mL protein). D) Bovine Serum, 10-fold diluted (7.0 mg/mL protein). E) Human Serum, 20-fold diluted (3.14 mg/mL protein).

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