

# FitAmp™ General Tissue Section DNA Isolation Kit

Base Catalog # P-1003

## PLEASE READ THIS ENTIRE USER GUIDE BEFORE USE

The *FitAmp*™ kits are very suitable for isolating tiny amounts of DNA from microdissection samples, fresh tissue sections, formalin-fixed and paraffin-embedded tissues, plasma, serum, body fluids, etc. The quality of extracted DNA from formalin-fixed and paraffin-embedded tissues may be affected by the quality of the embedded tissue.

The *FitAmp*™ kits allow isolation of DNA size from 50 bp to 20 kb; DNA quantity from 1 ng to 2 µg, optimal at between 10 ng and 1 µg.

## KIT CONTENTS

Components	50 samples P-1003-1	100 samples P-1003-2
S1 (DNA Digestion Solution)	0.3 ml	0.6 ml
S2 (DNA Digestion Powder)	1 vial	1 vial
S3 (DNA Isolation Buffer)	6 ml	11 ml
S4 (DNA Binding Buffer)	12 ml	22 ml
S5 (DNA Elution Solution)	1 ml	2 ml
F-Spin Column	50	100
F-Collection Tube	50	100
User Guide	1	1

## SHIPPING & STORAGE

Upon receipt: (1) **S2** should be stored at  $-20^{\circ}\text{C}$ , or stored at  $4^{\circ}\text{C}$  as soon as it is dissolved in **S1** (up to 6 months); (2) Store **all other components** at room temperature. The kit can be stable for up to 6 months from the shipment date when stored properly.

## GENERAL PRODUCT INFORMATION

**Quality Control:** Epigenetek guarantees the performance of all products in the manner described in our product instructions.

**Product Updates:** Epigenetek reserves the right to change or modify any product to enhance its performance and design.

**Usage Limitation:** The *FitAmp*<sup>™</sup> kits are for research use only and are not intended for diagnostic or therapeutic application.

**Intellectual Property:** *FitAmp*<sup>™</sup> is a trademark of Epigenetek, Inc.

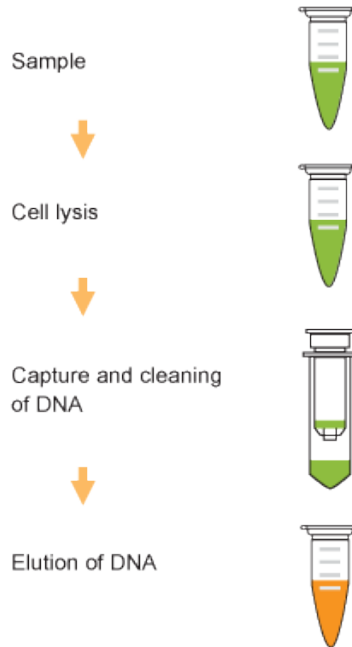
## A BRIEF OVERVIEW

The *FitAmp*<sup>™</sup> General Tissue Section DNA Isolation Kit is designed for isolating DNA from tissue sections. The kit uses a unique procedure and composition to efficiently isolate DNA in any targeted microscopic tissue area on a slide. The kit has the following features:

- The fastest procedure available, which can be finished within 2 hours, depending on sample types, with consistent isolation conditions.
- High efficiency of DNA isolation from tissue sections containing tiny amounts of DNA (as low as 1 ng).
- Use of non-toxic reagents and no phenol chloroform.

## PRINCIPLE & PROCEDURE

The *FitAmp*<sup>™</sup> General Tissue Section DNA Isolation Kit simply applies our proprietary DNA isolation buffer to a selected microscopic tissue area. The area is removed and transferred into a tube. After treatment with DNA digestion buffer, the DNA is easily recovered in 8-20  $\mu$ l by our specially designed Fast-Spin Column. DNA is ready for down-stream application.



Schematic Procedure for Using the *FitAmp*<sup>™</sup> General Tissue Section DNA Isolation Kit

## PROTOCOL

Note: Always cap spin columns before placing them in the microcentrifuge.

Before starting, prepare the following required solutions (not included): 90% ethanol, and 70% ethanol

1. Add 0.3 ml of **S1** to **S2** in order to create the **S1/S2 solution**. Vortex until solution is clear. Spin the solution down to the bottom.
2. Treat the tissue with the DNA Isolation Buffer:

For **microdissection samples**, directly collect the sample into a vial containing 100  $\mu$ l of **S3**, followed by adding 5  $\mu$ l of **S1/S2 solution**. Mix well and incubate at 65°C for 60-90 minutes.

For **tissues from fresh sections**, add 0.5  $\mu$ l of **S3** to 1 mm<sup>2</sup> (about 500-1000 cells) of tissue area and immediately remove the tissue area you need (1- 20 mm<sup>2</sup>) from the slide with a scalpel. Transfer it to a 1.5 ml vial containing 100  $\mu$ l of **S3**, followed by adding 5  $\mu$ l of **S1/S2 solution**. Mix

well and incubate this mixed solution at 65°C for 60-90 minutes or until the tissue is completely lysed (usually it is less than 2 hours). Vortex the sample for 5 seconds every 30 minutes. If tissue is not properly in this mixed solution after vortexing, spin it down into the solution.

**For paraffin samples**, remove the paraffin first with deparaffin reagents according to the manufacturer's instructions or according to the following procedures:

- 1) Drop the slide into 100% of *xylene* at room temperature for 10 min. Repeat once with new *xylene*.
- 2) Drop the slide in 100% of *ethanol*, 95% and 70% for 5 minutes each. Air dry the slide. Add 0.5  $\mu\text{l}$  of **S3** to 1 mm<sup>2</sup> of tissue area and immediately remove the tissue area you need (1-20 mm<sup>2</sup>) from the slide. Transfer it to a vial containing 100  $\mu\text{l}$  of **S3**, followed by adding 5  $\mu\text{l}$  of **S1/S2 solution**. Mix well and incubate this mixed solution at 65°C for 60-90 minutes, or until tissue is completely lysed (it is usually less than 2 hours). Vortex the sample for 5 seconds, every 30 minutes. If tissue is not properly in this mixed solution after vortexing, spin it down into the solution.
3. Place a spin column into a 2 ml collection tube. Vortex the mixture for 5 seconds after incubation. Add 200  $\mu\text{l}$  of **S4** to the mixture and transfer it to the column. Centrifuge at 12,000 rpm for 30 seconds. Discard the flowthrough. Replace the column to the collection tube (*Note: maximum volume of the column is 600  $\mu\text{l}$ .*)
4. Add 200  $\mu\text{l}$  of 70% *ethanol* to the column and centrifuge at 12,000 rpm for 20 seconds. Add 200  $\mu\text{l}$  of 90% *ethanol* to the column and centrifuge at 12,000 rpm for 20 seconds. Discard the flowthrough and replace the column to the collection tube.
5. Add an additional 200  $\mu\text{l}$  of 90% *ethanol* to the column and centrifuge at 12,000 rpm for 40 seconds.
6. Place the column in a new 1.5 ml vial. Add 8-18  $\mu\text{l}$  of **S5** directly to the column filter and centrifuge at 12,000 rpm for 20 seconds to elute DNA.

## RELATED PRODUCTS

P-1004	<i>FitAmp</i> <sup>™</sup> Plasma/Serum DNA Isolation Kit
P-1009	<i>FitAmp</i> <sup>™</sup> Paraffin Tissue Section DNA Isolation Kit

