

## Milk Bacterial DNA Isolation Kit

Product #21500

## Product Insert

Norgen's Milk Bacterial DNA Isolation Kit is designed for the rapid preparation of genomic DNA from the various bacterial species found within milk. The kit allows for the isolation of genomic DNA from both Gram negative and Gram positive bacteria found in milk samples. The genomic DNA is preferentially purified from other cellular proteinaceous components. Typical yields of genomic DNA will vary depending on the bacterial density of the milk sample, as well as the bacterial species present. The purified genomic DNA is fully digestible with all restriction enzymes tested, and is completely compatible with PCR, quantitative PCR and Southern Blot analysis.

### Norgen's Purification Technology

Purification is based on spin column chromatography using Norgen's proprietary resin as the separation matrix. Norgen's resin binds DNA under high salt concentrations and releases the bound DNA under low salt and slightly alkali conditions. The process for the isolation of genomic DNA from the bacteria found in milk involves first centrifuging the milk sample in order to pellet any bacteria which may be present. If DNA is to be isolated from Gram positive or unknown strains of bacteria, the pellet is then resuspended in Digestion Buffer in order to break down the cell wall of the bacteria. For all types of bacteria, the next step involves lysing the bacteria cells using a combination of the provided Lysis Buffer and Proteinase K. After a 30 minute incubation, Binding Solution and ethanol are added to the lysate, and the solution is loaded onto a spin column. Norgen's resin binds DNA in a manner that depends on ionic concentrations, thus the DNA will bind to the column while most of the RNA and the digested proteins will flowthrough or be retained on the top of the resin. The bound DNA is then washed twice using the provided wash buffer in order to remove any remaining impurities, and the purified bacterial genomic DNA is eluted with the elution buffer.

### Specifications

Kit Specifications	
Maximum Milk Input	1 mL
Time to Complete 10 purifications	45 minutes
Bacteria Species Processed	Gram positive and Gram negative
Minimum Detection Limit	10 bacteria in 1 mL of milk

### Advantages

- Fast and easy processing using a rapid spin-column format
- Isolate genomic DNA from both Gram positive and Gram negative bacteria found in milk
- DNA can be isolated and detected from milk samples with very low bacterial densities (10 bacteria in 1 mL of milk)
- Isolate high quality genomic DNA

## Kit Components

Component	Product #21500 (25 samples)
Digestion Buffer	3 mL
Lysis Solution	12 mL
Binding Solution	1.2 mL
Wash Solution I	3.75 mL
Wash Solution II	15 mL
Elution Buffer	6 mL
Proteinase K (lyophilized)	6 mg
Lysozyme (powder)	60 mg
Micro Spin Columns	25
Collection Tubes	25
Elution tubes (1.7 mL)	25
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## Storage Conditions and Product Stability

All solutions should be kept tightly sealed and stored at room temperature. The Lysozyme should be stored at -20°C upon arrival, and the Digestion Buffer should be stored at -20°C after addition of the lysozyme. The lyophilized Proteinase K should be stored at -20°C upon arrival and after reconstitution. These reagents should remain stable for at least 1 year in their unopened containers.

## Precautions and Disclaimers

This kit is designed for research purposes only. It is not intended for human or diagnostic use.

Ensure that a suitable lab coat, disposable gloves and protective goggles are worn when working with chemicals. For more information, please consult the appropriate Material Safety Data Sheets (MSDSs). These are available as convenient PDF files online at [www.norgenbiotek.com](http://www.norgenbiotek.com).

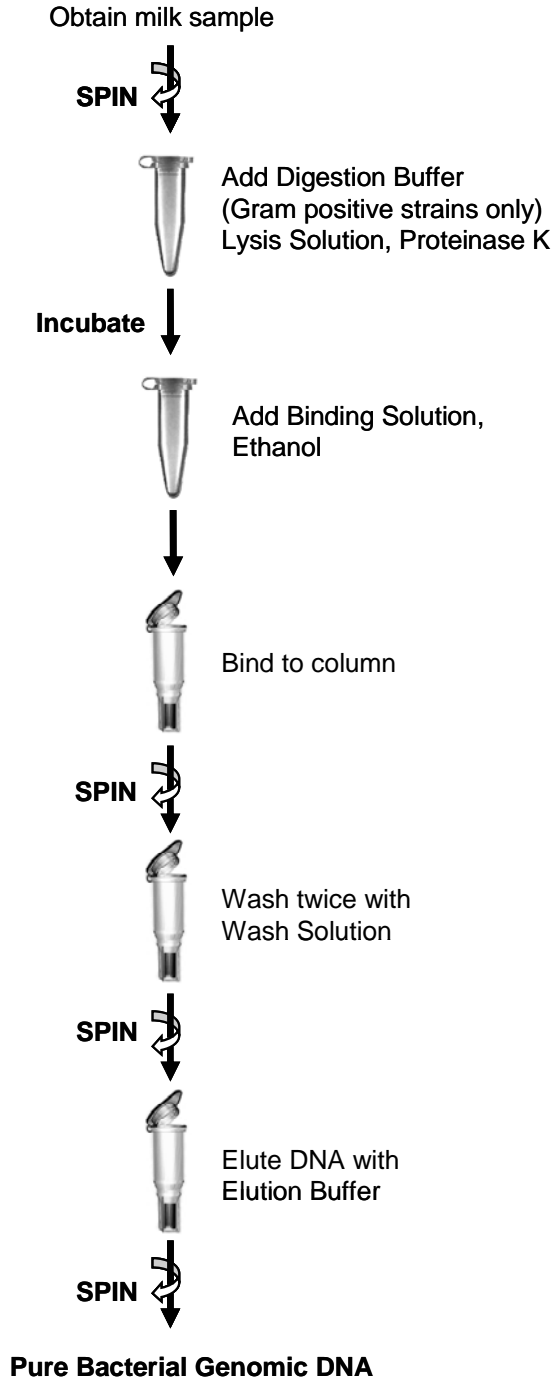
The **Binding Solution**, **Wash Solution I** and **Wash Solution II** contain guanidine hydrochloride, and should be handled with care. Guanidine hydrochloride forms highly reactive compounds when combined with bleach, thus care must be taken to properly dispose of any of these solutions.

## Customer-Supplied Reagents and Equipment

- Microcentrifuge tubes
- Benchtop microcentrifuge
- Micropipettors
- 55°C incubator
- 37°C incubator (for Gram positive strains only)
- 96 – 100% ethanol
- Cotton swab

## Flow Chart

Procedure for Purifying Bacterial Genomic DNA from Milk Samples using Norgen's Milk Bacterial DNA Isolation Kit



## Procedure

All centrifugation steps are carried out in a benchtop microcentrifuge. Various speeds are required for different steps, so please check your microcentrifuge specifications to ensure that it is capable of the proper speeds. All centrifugation steps are performed at room temperature. The correct rpm can be calculated using the formula:

$$\text{RPM} = \sqrt{\frac{\text{RCF}}{(1.118 \times 10^{-5}) (r)}}$$

where *RCF* = required gravitational acceleration (relative centrifugal force in units of g); *r* = radius of the rotor in cm; and *RPM* = the number of revolutions per minute required to achieve the necessary *g*-force.

### Notes prior to use:

- Preheat an incubator or heating block to 55°C (and 37°C for Gram positive strains).
- Prepare a working concentration of **Wash Solution I** by adding 11.25 mL of 96 - 100% ethanol (to be provided by the user) to the supplied bottle containing concentrated **Wash Solution I**. This will give a final volume of 15 mL. The label on the bottle has a box that can be checked to indicate that ethanol has been added.
- Reconstitute the **Proteinase K** in 300 µL of molecular biology grade water, aliquot into small fractions and store the unused portions at -20°C until needed.
- Add the provided amount of **Digestion Buffer** to the tube containing the **Lysozyme**, and mix well. Aliquot the **Digestion Buffer** into small fractions and store the unused portions at -20°C until needed.

### 1A. Lysate Preparation (Gram Negative Bacteria)

- a. Aliquot a maximum of 1 mL of milk into a microcentrifuge tube.
- b. Centrifuge at 14,000 x g (~14,000 RPM) for 2 minutes.
- c. Pour off the supernatant by quickly inverting the tube and gently tapping it against the wall of the waste container. This tapping is to ensure that the creamy layer present on the top of the milk sample after centrifugation is removed. Clean any remaining white solid from the microcentrifuge tube wall by using either a clean 200 µL pipette tip or a cotton swab. Ensure that the pellet is not dislodged.
- d. Add 400 µL of **Lysis Solution** and 10 µL of reconstituted **Proteinase K** to the cell pellet and mix well by vortexing.
- e. Incubate the lysate at 55°C for 30 minutes.

## 1B. Lysate Preparation (Gram Positive Bacteria)

- a. Aliquot a maximum of 1 mL of milk into a microcentrifuge tube.
- b. Centrifuge at 14,000 x g (~14,000 RPM) for 2 minutes.
- c. Pour off the supernatant by quickly inverting the tube and gently tapping it against the wall of the waste container. This tapping is to ensure that the creamy layer present on the top of the milk sample after centrifugation is removed. Clean any remaining white solid from the microcentrifuge tube wall by using either a clean 200  $\mu$ L pipette tip or a cotton swab. Ensure that the pellet is not dislodged.
- d. Resuspend the pellet in 100  $\mu$ L of **Digestion Buffer**. Incubate at 37°C for 30 minutes.

**Note:** Ensure that the provided lysozyme has been added to the Digestion Buffer.

- e. After incubation, add 300  $\mu$ L of **Lysis Solution** and 10  $\mu$ L of reconstituted **Proteinase K** to the digestion mixture and mix well by vortexing.
- f. Incubate the lysate at 55°C for 30 minutes.

## 2. Sample Binding to Column

- a. After incubation, add 40  $\mu$ L of **Binding Solution** and 180  $\mu$ L of 96-100% ethanol to the lysis mixture, and mix by vortexing.
- b. Spin the sample for 10 seconds at 14,000 x g (~14,000 RPM). A thin layer of lipid may form on the top of the aqueous phase. Using a pipette, carefully transfer the clear aqueous phase only to a spin column that has been attached to a collection tube.
- c. Centrifuge the column assembly for 4 minutes at 5,200 x g (~8,000 RPM) to bind the bacterial DNA. If all the liquid does not pass through the column, spin for an additional 1 minute at 14,000 x g (~14,000 RPM).

## 3. Column Wash

- a. Apply 500  $\mu$ L of **Wash Solution I** to the column and centrifuge for 3 minutes at 14,000 x g (~14,000 RPM).

**Note:** Ensure the appropriate amount of ethanol has been added to **Wash Solution I**.

- b. Discard the flowthrough and reassemble the column and the collection tube.
- c. Apply 500  $\mu$ L of **Wash Solution II** to the column and centrifuge again for 3 minutes at 14,000 x g (~14,000 RPM).
- d. Ensure that all the **Wash Solution II** has passed through the column and that the column is dry. Spin for an additional minute if necessary.
- e. Discard the collection tube with the flowthrough.

## 4. DNA Elution

- a. Transfer the spin column to a provided 1.7 mL Elution tube.
- b. Apply 100  $\mu$ L of **Elution Buffer** to the column and centrifuge at 2,600 x g (~6,000 RPM) for 2 minutes.
- c. Spin for an additional 2 minutes at 14,000 x g (~14,000 RPM) to complete the DNA elution.

## Troubleshooting Guide

Problem	Possible Cause	Solution and Explanation
The micro spin column is clogged	The sample is too large	Ensure that no more than 1 mL of milk be used for the procedure in order to prevent clogging of the column. If the milk sample is known to have a high bacterial content, it is recommend that less than 1 mL be used for the input. Clogging can be alleviated by increasing the g-force and/or centrifuging for a longer period of time until the lysate passes through the column.
	White, creamy layer was not removed after initial spin	After the first 2 minute spin, ensure that the white, creamy layer floating on top of the milk sample is removed. This layer should be removed in order to prevent clogging of the column.
The lysate is very gelatinous prior to loading onto the column	White, creamy layer was not removed after initial spin	After the first 2 minute spin, ensure that the white, creamy layer floating on top of the milk sample is removed. This layer should be removed in order to minimize the thickness of the lysate.
	The sample is too large	Too many cells are in the lysate preparation. Ensure that no more than 1 mL of milk is used for the procedure. If the milk sample is known to have a high bacterial content, it is recommended that less than 1 mL is used for the input.
The yield of genomic DNA is low	The milk sample may not contain any bacterial species	If the milk sample does not contain any bacteria then no genomic DNA will be detected.
	Incomplete lysis of cells	Extend the incubation time of Proteinase K digestion or reduce the amount of milk used for the input. Increase the lysozyme incubation time for Gram positive strains.
	The DNA elution is incomplete	Ensure that centrifugation at 16,000 x g is performed after the 3,000 x g centrifugation cycle, to ensure that all the DNA is eluted.
The genomic DNA is sheared	The genomic DNA was handled improperly	Pipetting steps should be handled as gently as possible. Reduce vortexing times during mixing steps (no more than 10-15 seconds).

<b>Related Products</b>	<b>Product #</b>
Bacterial Genomic DNA Isolation Kit	17900
Yeast Genomic DNA Isolation Kit	18600
Blood Genomic DNA Isolation Kit	18200

### **Technical Support**

Contact our Technical Support Team between the hours of 8:30 and 5:30 (Eastern Standard Time) at (905) 227-8848 or Toll Free at 1-866-667-4362.

Technical support can also be obtained from our website ([www.norgenbiotek.com](http://www.norgenbiotek.com)) or through email at [techsupport@norgenbiotek.com](mailto:techsupport@norgenbiotek.com).

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