

# MagMAX™ CORE Nucleic Acid Purification Kit

## USER GUIDE

Automated purification of high-quality DNA and RNA from  
veterinary samples

for use with:

KingFisher™ Flex Purification System

MagMAX™ Express-96 Deep Well Magnetic Particle Processor

KingFisher™ Duo Prime Purification System

KingFisher™ mL Purification System

Catalog Numbers A32700, A32702

Publication Number MAN0015944

Revision C.0



**Manufacturer:** Life Technologies Corporation | 2130 Woodward Street | Austin, TX 78744

The information in this guide is subject to change without notice.

**DISCLAIMER:** TO THE EXTENT ALLOWED BY LAW, LIFE TECHNOLOGIES AND/OR ITS AFFILIATE(S) WILL NOT BE LIABLE FOR SPECIAL, INCIDENTAL, INDIRECT, PUNITIVE, MULTIPLE, OR CONSEQUENTIAL DAMAGES IN CONNECTION WITH OR ARISING FROM THIS DOCUMENT, INCLUDING YOUR USE OF IT.

**Revision history:** Pub. No. MAN0015944 (English)

Revision	Date	Description
C.0	13 December 2017	<ul style="list-style-type: none"><li>• Updated the usage statement on the front cover.</li><li>• Minor reorganization of the required materials section for style and clarity.</li><li>• Minor corrections to product names.</li><li>• Minor edits for style and consistency.</li></ul>
B.0	30 June 2017	<ul style="list-style-type: none"><li>• Combined and renamed workflows:<ul style="list-style-type: none"><li>– Simple: formerly Workflows A and C</li><li>– Complex: formerly Workflow B</li><li>– Digestion: formerly Workflow D</li></ul></li><li>• Added new workflow: Lysis Incubation.</li><li>• Added plate processing of samples in the Digestion workflow.</li><li>• Added list of instrument scripts.</li><li>• Reorganized into chapters for better navigation and clarity.</li></ul>
A.0	22 December 2016	New document.

**Important Licensing Information:** These products may be covered by one or more Limited Use Label Licenses. By use of these products, you accept the terms and conditions of all applicable Limited Use Label Licenses.

**Limited Use Label License No. 569: Veterinary and Research Use:** Notice to Purchaser : The purchase of this product conveys to the purchaser the limited, non-transferable right to use the purchased amount of the product for research and veterinary use only. No other right is hereby granted expressly, by implication, or by estoppel. The purchase of this product does not grant the purchaser any additional rights, including (without limitation) the right to transfer or resell the product in any form or the right to use the product as a therapeutic agent or human diagnostics test component. For information on obtaining additional rights, please contact [outlicensing@thermofisher.com](mailto:outlicensing@thermofisher.com) or Licensing and Commercial Supply, Thermo Fisher Scientific, 5823 Newton Drive, Carlsbad, CA, 92008, United States.

**TRADEMARKS:** All trademarks are the property of Thermo Fisher Scientific and its subsidiaries unless otherwise specified.

©2017 Thermo Fisher Scientific Inc. All rights reserved.

# Contents

■	<b>CHAPTER 1</b>	<b>Product information</b>	5
		Product description	5
		Contents and storage	5
		Required materials not supplied	6
		Recommended workflows	8
■	<b>CHAPTER 2</b>	<b>Before you begin</b>	9
		Procedural guidelines	9
		Before first use of the kit	9
		Determine the maximum plate shaker setting	9
		Download and install the script	10
■	<b>CHAPTER 3</b>	<b>Simple Workflow</b>	11
		Workflow: Simple	11
		Set up the processing plates	12
		Prepare Bead/PK Mix	12
		Prepare Lysis/Binding Solution	13
		Prepare the sample	14
		Combine the sample with Bead/PK Mix and Lysis/Binding Solution	15
		Process samples on the instrument	15
■	<b>CHAPTER 4</b>	<b>Complex Workflow</b>	16
		Workflow: Complex	16
		Set up the processing plates	17
		Prepare Bead/PK Mix	17
		Prepare the Lysis Solution	18
		Prepare the clarified lysate	19
		Combine the clarified lysate with Bead/PK Mix and MagMAX™ CORE Binding Solution	20
		Process samples on the instrument	21

■	<b>CHAPTER 5</b>	<b>Digestion Workflow</b>	22
		Workflow: Digestion	22
		Set up the processing plates	23
		Prepare the Lysis/Binding Solution	23
		Prepare PK Solution	24
		Treat the samples with PK Solution	25
		Treat the samples with PK Solution (tube processing)	25
		Treat the samples with PK Solution (plate processing)	27
		Combine Proteinase K-treated samples with beads and Lysis/Binding Solution	28
		Process samples on the instrument	29
■	<b>CHAPTER 6</b>	<b>Lysis Incubation Workflow</b>	30
		Workflow: Lysis Incubation	30
		Set up the processing plates	31
		Prepare Lysis/Binding/Bead Mix	31
		Prepare the ear punch lysate	31
		Combine the lysate with Proteinase K and Lysis/Binding/Bead Mix	32
		Process samples on the instrument	33
■	<b>APPENDIX A</b>	<b>Troubleshooting</b>	34
■	<b>APPENDIX B</b>	<b>Purification with the KingFisher™ Duo Prime or KingFisher™ mL instrument</b>	36
		Required materials not supplied	36
		Purification procedure	37
■	<b>APPENDIX C</b>	<b>Safety</b>	38
		Chemical safety	39
		Biological hazard safety	40
		<b>Documentation and support</b>	41
		Related documentation	41
		Customer and technical support	41
		Limited product warranty	41



# Product information

---

**IMPORTANT!** Before using this product, read and understand the information in the “Safety” appendix in this document.

---

## Product description

The MagMAX™ CORE Nucleic Acid Purification Kit is designed for rapid purification of high-quality DNA and RNA for downstream molecular analysis. The kit uses magnetic bead-based separation, and it is compatible with the following instruments:

- KingFisher™ Flex Magnetic Particle Processor
- MagMAX™ Express-96 Deep Well Magnetic Particle Processor
- KingFisher™ Duo Prime Magnetic Particle Processor
- KingFisher™ mL Magnetic Particle Processor

The kit is optimized for a wide range of sample types. See “Recommended workflows” on page 8.

## Contents and storage

Table 1 MagMAX™ CORE Nucleic Acid Purification Kit

Contents	Cat. No. A32700 (100 reactions)	Cat. No. A32702 (500 reactions)	Storage
MagMAX™ CORE Lysis Solution <sup>[1]</sup>	50 mL	275 mL	15–30°C (room temperature)
MagMAX™ CORE Binding Solution	45 mL	220 mL	
MagMAX™ CORE Wash Solution 1	60 mL	300 mL	
MagMAX™ CORE Wash Solution 2	60 mL	300 mL	
MagMAX™ CORE Elution Buffer	12 mL	55 mL	
MagMAX™ CORE Magnetic Beads	2.2 mL	11 mL	
MagMAX™ CORE Proteinase K (20 mg/mL)	1.25 mL	5 mL	

<sup>[1]</sup> Available for purchase separately (Cat. No. A32837).

## Required materials not supplied

Unless otherwise indicated, all materials are available through **thermofisher.com**.  
MLS: Fisher Scientific (**fisherscientific.com**) or other major laboratory supplier.

Table 2 Materials required for all workflows<sup>[1]</sup>

Item	Source
<b>Instrument and equipment</b>	
One of the following instruments: <ul style="list-style-type: none"> <li>KingFisher™ Flex Purification System</li> <li>MagMAX™ Express-96 Deep Well Magnetic Particle Processor</li> </ul> See page 36 for other compatible instruments.	Contact your local sales office.
Benchtop microcentrifuge capable of 15,000 × g	MLS
Laboratory mixer, Vortex or equivalent	MLS
<b>Reagents</b>	
PBS, pH 7.4 <sup>[2]</sup>	10010023
<b>(Optional) Internal positive control (IPC), one of the following:</b>	
VetMAX™ Xeno™ Internal Positive Control DNA	A29764
VetMAX™ Xeno™ Internal Positive Control RNA	A29763
IPC supplied with your VetMAX™ PCR Kit	<b>thermofisher.com</b>
<b>Tubes, plates, and other consumables</b>	
Adhesive PCR Plate Foils, or equivalent	AB0626
KingFisher™ Flex Microtiter Deepwell 96 plates, 50 plates	95040460
KingFisher™ 96 KF microplates (200 µL), 48 plates	97002540
KingFisher™ 96 tip comb for DW magnets, 100 combs	97002534

<sup>[1]</sup> See Table 4 and Table 5 for additional materials required for the Simple and Digestion workflows.

<sup>[2]</sup> Not required for the Lysis Incubation workflow.

Table 3 Optional equipment

Item	Source
Biotang Inc Microplate Shaker, or equivalent titer plate shaker (for mixing beads with samples; all workflows)	Fisher Scientific™ 50-751-4965
Benchtop centrifuge with plate adaptors (for lysate preparation in plates; Complex and Digestion workflows)	MLS

**Table 4** Additional materials required for the Simple workflow (tissue samples only)

Item	Source
Fisher Scientific™ Bead Mill 24 Homogenizer	Fisher Scientific™ 15-340-163
PYREX™ Solid Glass Beads for Distillation Columns (3 mm)	Fisher Scientific™ 11-312-10A

**Table 5** Additional materials required for the Digestion workflow

Item	Source
PK Buffer for MagMAX™ -96 DNA Multi-Sample Kit	4489111

## Recommended workflows

**Note:** For tough-to-lyse bacteria, for example, *M. paratuberculosis* (MAP), use the MagMAX™ CORE Mechanical Lysis Module (Cat. Nos. A32836, A37487).

Sample matrix	Nucleic acid	Recommended workflow
<ul style="list-style-type: none"> <li>Ear punch (circular shape, 2- to 3-mm diameter) in Lysis Solution</li> </ul>	Viral nucleic acid	Lysis Incubation <sup>[1]</sup> (page 30)
<ul style="list-style-type: none"> <li>Ear punch (circular shape, 2- to 3-mm diameter) in PBS</li> <li>Ear notch (triangular shape, approximately 1-cm width)</li> <li>Milk</li> <li>Plasma</li> <li>Serum</li> </ul>	<ul style="list-style-type: none"> <li>Viral nucleic acid</li> <li>Bacterial DNA</li> </ul>	Simple (page 11)
<ul style="list-style-type: none"> <li>Biomed Diagnostics InPouch™ TF (<i>Tritrichomonas foetus</i>) culture</li> </ul>	<i>Tritrichomonas foetus</i> DNA	
<ul style="list-style-type: none"> <li>Semen</li> </ul>	Viral nucleic acid	
<ul style="list-style-type: none"> <li>Swabs—animal</li> <li>Whole blood</li> </ul>	<ul style="list-style-type: none"> <li>Viral nucleic acid</li> <li>Genomic DNA</li> </ul>	
<ul style="list-style-type: none"> <li>Tissue or organ</li> </ul>	<ul style="list-style-type: none"> <li>Viral nucleic acid</li> <li>Bacterial DNA<sup>[2]</sup></li> </ul>	
<ul style="list-style-type: none"> <li>Hair follicles</li> </ul>	Genomic DNA	Digestion (page 22)
<ul style="list-style-type: none"> <li>Environmental samples</li> <li>Feces</li> <li>Swabs—environmental or fecal</li> </ul>	Bacterial DNA	
<ul style="list-style-type: none"> <li>Oral fluid</li> </ul>	<ul style="list-style-type: none"> <li>Viral nucleic acid</li> <li>Bacterial DNA</li> </ul>	Complex (page 16)

<sup>[1]</sup> Recommended if overnight incubation is required.

<sup>[2]</sup> If concurrent isolation of viral nucleic acid and bacterial DNA is not required, use the Digestion workflow.





# Before you begin

## Procedural guidelines

- Before use, invert bottles of solutions and buffers to ensure thorough mixing.
- Mix samples with reagents using a plate shaker or by pipetting up and down.  
**Note:** Do not use a plate shaker with the large tube strips required by the KingFisher™ mL instrument.
- To prevent cross-contamination:
  - Cover the plate or tube strip during the incubation and shaking steps, to prevent spill-over.
  - Carefully pipet reagents and samples, to avoid splashing.
- To prevent nuclease contamination:
  - Wear laboratory gloves during the procedures. Gloves protect you from the reagents, and they protect the nucleic acid from nucleases that are present on skin.
  - Use nucleic acid-free pipette tips to handle the reagents, and avoid putting used tips into the reagent containers.
  - Decontaminate lab benches and pipettes before you begin.

## Before first use of the kit

### Determine the maximum plate shaker setting

If a plate shaker is used, determine the maximum setting:

1. Verify that the plate fits securely on your shaker.
2. Add 1 mL of water to each well of the plate, then cover with sealing foil.
3. Determine the maximum setting that you can use on your shaker without any of the water splashing onto the sealing foil.

## Download and install the script

The appropriate script for the MagMAX™ CORE Nucleic Acid Purification Kit must be installed on the instrument before first use.

1. On the MagMAX™ CORE Nucleic Acid Purification Kit product web page (at [thermofisher.com](http://thermofisher.com), search by catalogue number), scroll to the **Product Literature** section.
2. Right-click the appropriate file to download the latest version of the MagMAX\_CORE script for your instrument.

**Table 6** Recommended scripts

Instrument	Script name
KingFisher™ Flex	MagMAX_CORE_Flex.bdz
KingFisher™ 96 MagMAX™ Express-96	MagMAX_CORE_KF-96.bdz
KingFisher™ Duo Prime	MagMAX_CORE_DUO.bdz
KingFisher™ mL	MagMAX_CORE_mL_no_heat.bdz

If required by your laboratory, use one of the following scripts, which do not heat the samples during the elution step.

**Table 7** Alternate scripts without heated elution step

Instrument	Script name
KingFisher™ Flex	MagMAX_CORE_Flex_no_heat.bdz
KingFisher™ 96 MagMAX™ Express-96	MagMAX_CORE_KF-96_no_heat.bdz
KingFisher™ Duo Prime	MagMAX_CORE_DUO_no_heat.bdz
KingFisher™ mL	MagMAX_CORE_mL_no_heat.bdz

3. See your instrument user guide or contact Technical Support for instructions for installing the script.

# 3

## Simple Workflow

The Simple workflow is recommended for the following sample types. The nucleic acid that this workflow is optimized for varies by sample type; see “Recommended workflows” on page 8 for details.

- Biomed Diagnostics InPouch™ TF (*Tritrichomonas foetus*) culture
- Ear notch (triangular shape, approximately 1-cm width)
- Ear punch (circular shape, 2- to 3-mm diameter; PBS incubation)
- Milk
- Plasma
- Semen
- Serum
- Swabs—animal
- Tissue or organ
- Whole blood

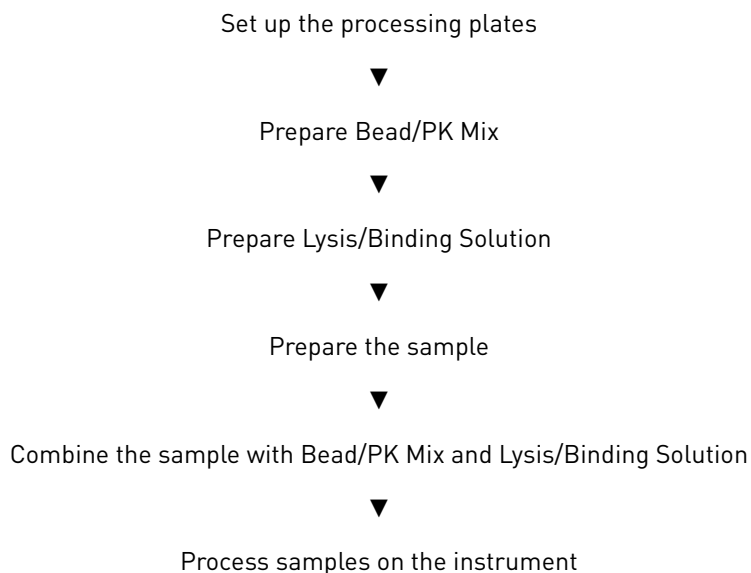
Follow this procedure if you are using these instruments:

- KingFisher™ Flex
- MagMAX™ Express-96

Follow Appendix B, “Purification with the KingFisher™ Duo Prime or KingFisher™ mL instrument” if you are using these instruments:

- KingFisher™ Duo Prime
- KingFisher™ mL

### Workflow: Simple



## Set up the processing plates

1. Set up the processing plates.

**Table 8** Plate setup: KingFisher™ Flex or MagMAX™ Express-96 instrument

Plate ID	Plate position <sup>[1]</sup>	Plate type	Reagent	Volume per well
Wash Plate 1	2	Deep Well	MagMAX™ CORE Wash Solution 1	500 µL
Wash Plate 2	3	Deep Well	MagMAX™ CORE Wash Solution 2	500 µL
Elution	4	Standard	MagMAX™ CORE Elution Buffer	90 µL
Tip Comb	5	Standard	Place a tip comb in the plate.	

<sup>[1]</sup> Position on the instrument.

2. (Optional) To prevent evaporation and contamination, cover the prepared processing plates with sealing foil until they are loaded into the instrument.

## Prepare Bead/PK Mix

We recommend that you prepare new Bead/PK Mix for each processing run. If necessary, you can store Bead/PK Mix at 4°C for up to 1 week.

1. Vortex the MagMAX™ CORE Magnetic Beads thoroughly to ensure that the beads are fully resuspended.
2. Combine the following components for the required number of samples plus 10% overage.

Component	Volume per sample
MagMAX™ CORE Magnetic Beads	20 µL
MagMAX™ CORE Proteinase K	10 µL
<b>Total Bead/PK Mix</b>	<b>30 µL</b>

## Prepare Lysis/Binding Solution

1. Combine the following components for the required number of samples plus 10% overage.

Component	Volume per sample
MagMAX™ CORE Lysis Solution	350 µL
MagMAX™ CORE Binding Solution	350 µL
<b>Total Lysis/Binding Solution (-IPC)</b>	<b>700 µL</b>
<i>(Optional)</i> Internal positive control (IPC), one of the following:	
VetMAX™ Xeno™ Internal Positive Control DNA	2 µL
VetMAX™ Xeno™ Internal Positive Control RNA	2 µL
Internal positive control (IPC) supplied with your VetMAX™ PCR Kit	As indicated in the instructions for the kit
<b>Total Lysis/Binding Solution (+IPC)</b>	<b>700 µL + volume of IPC</b>

2. Mix by inverting the tube or bottle at least 10 times.

*(Optional)* Store Lysis/Binding Solution at room temperature for up to 24 hours.

## Prepare the sample

Prepare samples according to sample type.

For...	Do this...
Biomed Diagnostics InPouch™ TF culture	Proceed with 300 µL of previously enriched culture media.
Ear notch (triangular shape, approximately 1-cm width)	<ol style="list-style-type: none"> <li>1. Add one ear notch to a 5-mL specimen tube.</li> <li>2. Add 2 mL of PBS, pH 7.4 to each sample.</li> <li>3. Incubate at room temperature with or without shaking: <ul style="list-style-type: none"> <li>• Without shaking—15 minutes</li> <li>• With moderate shaking—10 minutes</li> </ul> </li> <li>4. Proceed with 200 µL of supernatant.</li> </ol>
Ear punch (circular shape, 2- to 3-mm diameter)	<ol style="list-style-type: none"> <li>1. Add one ear punch to a 2-mL tube.</li> <li>2. Add 200 µL of PBS, pH 7.4 to each sample.</li> <li>3. Incubate at room temperature with or without shaking: <ul style="list-style-type: none"> <li>• Without shaking—15 minutes</li> <li>• With moderate shaking—10 minutes</li> </ul> </li> <li>4. Proceed with 50–200 µL of supernatant.</li> </ol>
Milk, plasma, serum, or whole blood	Proceed with 200 µL of sample.
Semen	<ol style="list-style-type: none"> <li>1. Add 500 µL of semen to a fresh tube.</li> <li>2. Centrifuge at 15,000 × <i>g</i> for 2 minutes.</li> <li>3. Proceed with 200 µL of supernatant.</li> </ol>
Swabs—animal	<p>Follow the manufacturer's recommended protocol, or follow this procedure:</p> <ol style="list-style-type: none"> <li>1. Break off the tip of the swab and add to a 2-mL tube.</li> <li>2. Add 1 mL of PBS, pH 7.4 to each sample.</li> <li>3. Vortex for 3 minutes.</li> <li>4. Proceed with 200 µL of supernatant.</li> </ol>
Tissue or organ	<ol style="list-style-type: none"> <li>1. Add the following components to a 2-mL tube: <ul style="list-style-type: none"> <li>• Tissue—20 to 30 mg</li> <li>• PBS, pH 7.4—1 mL</li> <li>• PYREX™ Solid Glass Beads for Distillation Columns (3 mm)—2 beads</li> </ul> </li> <li>2. Disrupt (bead-beat) the samples in a Fisher Scientific™ Bead Mill 24 Homogenizer at 6 m/s for 45 seconds.</li> <li>3. Centrifuge at 1,000 × <i>g</i> for 1 minute.</li> <li>4. Proceed with 100 µL of supernatant.</li> </ol>

## Combine the sample with Bead/PK Mix and Lysis/Binding Solution

1. Invert the tube of Bead/PK Mix several times to resuspend the beads, then add 30  $\mu$ L of the Bead/PK Mix to the required wells in the plate or tube strip.
2. Transfer the appropriate volume of each prepared sample to a well with Bead/PK Mix.

For...	Use...
Biomed Diagnostics InPouch™ TF culture	300 $\mu$ L of supernatant
Ear notch (triangular shape, approximately 1-cm width) Semen Swabs—animal	200 $\mu$ L of supernatant
Ear punch (circular shape, 2- to 3-mm diameter)	50–200 $\mu$ L of supernatant
Milk, plasma, serum, or whole blood	200 $\mu$ L of sample
Tissue or organ	100 $\mu$ L of supernatant

3. Mix the sample with Bead/PK Mix for 2 minutes at room temperature according to your mixing method.
  - **Using a plate shaker**—Shake vigorously for 2 minutes (see “Determine the maximum plate shaker setting” on page 9).
  - **By pipetting**—Pipet up and down several times, then incubate for 2 minutes at room temperature. (For downstream processing on the KingFisher™ mL Magnetic Particle Processor, you must mix by pipetting.)
4. Add 700  $\mu$ L of Lysis/Binding Solution to each sample-containing well or tube.
5. Immediately proceed to process samples on the instrument (next section).

## Process samples on the instrument

1. Select the appropriate script on the instrument (see “Download and install the script” on page 10).
2. Start the run, then load the prepared plates or tube strips in the appropriate positions when prompted by the instrument.

Store purified nucleic acid on ice for immediate use, at  $-20^{\circ}\text{C}$  for up to 1 month, or at  $-80^{\circ}\text{C}$  for long-term storage.

# 4

## Complex Workflow

The Complex workflow is recommended for the following sample types. The nucleic acid that this workflow is optimized for varies by sample type; see “Recommended workflows” on page 8 for details.

- Environmental samples
- Feces
- Oral fluid
- Swabs—environmental or fecal

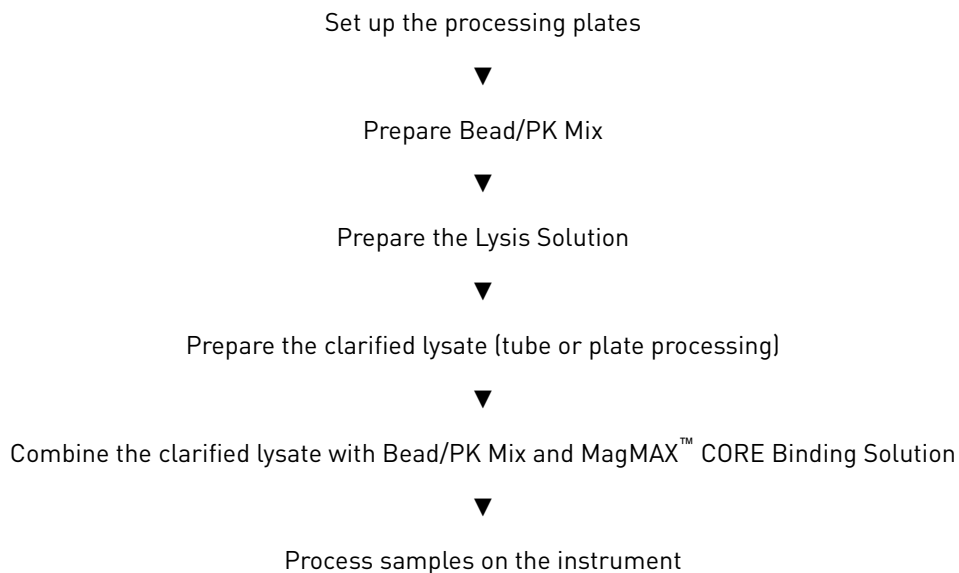
Follow this procedure if you are using these instruments:

- KingFisher™ Flex
- MagMAX™ Express-96

Follow Appendix B, “Purification with the KingFisher™ Duo Prime or KingFisher™ mL instrument” if you are using these instruments:

- KingFisher™ Duo Prime
- KingFisher™ mL

### Workflow: Complex





## Set up the processing plates

1. Set up the processing plates.

**Table 9** Plate setup: KingFisher™ Flex or MagMAX™ Express-96 instrument

Plate ID	Plate position <sup>[1]</sup>	Plate type	Reagent	Volume per well
Wash Plate 1	2	Deep Well	MagMAX™ CORE Wash Solution 1	500 µL
Wash Plate 2	3	Deep Well	MagMAX™ CORE Wash Solution 2	500 µL
Elution	4	Standard	MagMAX™ CORE Elution Buffer	90 µL
Tip Comb	5	Standard	Place a tip comb in the plate.	

<sup>[1]</sup> Position on the instrument.

2. (Optional) To prevent evaporation and contamination, cover the prepared processing plates with sealing foil until they are loaded into the instrument.

## Prepare Bead/PK Mix

We recommend that you prepare new Bead/PK Mix for each processing run. If necessary, you can store Bead/PK Mix at 4°C for up to 1 week.

1. Vortex the MagMAX™ CORE Magnetic Beads thoroughly to ensure that the beads are fully resuspended.
2. Combine the following components for the required number of samples plus 10% overage.

Component	Volume per sample
MagMAX™ CORE Magnetic Beads	20 µL
MagMAX™ CORE Proteinase K	10 µL
<b>Total Bead/PK Mix</b>	<b>30 µL</b>

## Prepare the Lysis Solution

1. Combine the following components for the required number of samples plus 10% overage.

Component	Volume per sample
MagMAX™ CORE Lysis Solution	450 µL
[Optional] Internal positive control (IPC), one of the following:	
VetMAX™ Xeno™ Internal Positive Control DNA	2 µL
VetMAX™ Xeno™ Internal Positive Control RNA	2 µL
Internal positive control (IPC) supplied with your VetMAX™ PCR Kit	As indicated in the instructions for the kit
<b>Total Lysis Solution (+IPC)</b>	<b>450 µL + volume of IPC</b>

2. Mix by inverting the tube or bottle at least 10 times.

(Optional) Store Lysis Solution at room temperature for up to 24 hours.

## Prepare the clarified lysate

1. Prepare samples according to sample type.

For...	Do this...
Environmental samples Feces	<ol style="list-style-type: none"> <li>Transfer 0.2-0.3 g of sample to a 2-mL tube.</li> <li>Add 1 mL of PBS, pH 7.4, then vortex vigorously for 3 minutes.</li> <li>Centrifuge as indicated. <ul style="list-style-type: none"> <li><b>For viral nucleic acid purification</b>—centrifuge at 15,000 × <i>g</i> for 1 minute.</li> <li><b>For bacterial DNA purification or concurrent purification of bacterial and viral nucleic acids</b>—centrifuge at 100 × <i>g</i> for 1 minute.</li> </ul> </li> <li>Proceed with 200 µL of supernatant.</li> </ol>
Oral fluid	<ol style="list-style-type: none"> <li>Briefly mix the oral fluid sample.</li> <li>Proceed with 300 µL of sample.</li> </ol>
Swabs— environmental or fecal	<ol style="list-style-type: none"> <li><b>Fecal samples</b>—swirl a clinical swab in a fecal sample. <b>Environmental swabs</b>—proceed with an environmental swab.</li> <li>Add 1 mL of PBS, pH 7.4 to a 2-mL tube.</li> <li>Swirl the swab in 1 mL of PBS, pH 7.4 for 5–10 seconds, removing as much sample material as possible, then discard the swab. Alternatively, break off the swab tip and leave the swab in the PBS, pH 7.4.</li> <li>Vortex vigorously for 3 minutes, or until the sample is suspended.</li> <li>Centrifuge as indicated. <ul style="list-style-type: none"> <li><b>For viral nucleic acid purification</b>—centrifuge at 15,000 × <i>g</i> for 1 minute.</li> <li><b>For bacterial DNA purification or concurrent purification of bacterial and viral nucleic acids</b>—centrifuge at 100 × <i>g</i> for 1 minute.</li> </ul> </li> <li>Proceed with 200 µL of supernatant.</li> </ol>

## 2. Add Lysis Solution, then clarify the lysate.

For...	Do this...
Processing in tubes	<ol style="list-style-type: none"> <li>1. For each sample, add 450 µL of Lysis Solution to a new 2-mL tube.</li> <li>2. Add the indicated volume of sample from step 1 on page 19 to the Lysis Solution.</li> <li>3. Vortex vigorously for 3 minutes.</li> <li>4. Centrifuge at 15,000 × <i>g</i> for 2 minutes.</li> <li>5. Remove the supernatant (clarified lysate) without disturbing the pellet.</li> </ol>
Processing in plates	<ol style="list-style-type: none"> <li>1. For each sample, add 450 µL of Lysis Solution to the appropriate wells of a deep-well plate.</li> <li>2. Add the indicated volume of sample from step 1 on page 19 to the Lysis Solution.</li> <li>3. Seal the plate with sealing foil.</li> <li>4. Shake the plate at moderate speed for 5 minutes.</li> <li>5. Centrifuge at 3,000 × <i>g</i> for 5 minutes.</li> <li>6. Remove the supernatant (clarified lysate) without disturbing the pellet.</li> </ol>

## Combine the clarified lysate with Bead/PK Mix and MagMAX™ CORE Binding Solution

1. Invert the tube of Bead/PK Mix several times to resuspend the beads, then add 30 µL of the Bead/PK Mix to the required wells in the plate or tube strip.
2. Transfer the appropriate volume of each clarified lysate (see “Prepare the clarified lysate” on page 19) to a well with the Bead/PK Mix.

For...	Use...
Oral fluid	600 µL
Environmental samples, fecal samples, and swabs	500 µL

3. Mix the sample with Bead/PK Mix for 2 minutes at room temperature according to your mixing method.
  - **Using a plate shaker**—Shake vigorously for 2 minutes (see “Determine the maximum plate shaker setting” on page 9).
  - **By pipetting**—Pipet up and down several times, then incubate for 2 minutes at room temperature. (For downstream processing on the KingFisher™ mL Magnetic Particle Processor, you must mix by pipetting.)
4. Add 350 µL of MagMAX™ CORE Binding Solution.
5. Immediately proceed to process samples on the instrument (next section).

## Process samples on the instrument

1. Select the appropriate script on the instrument (see “Download and install the script” on page 10).
2. Start the run, then load the prepared plates or tube strips in the appropriate positions when prompted by the instrument.

Store purified nucleic acid on ice for immediate use, at  $-20^{\circ}\text{C}$  for up to 1 month, or at  $-80^{\circ}\text{C}$  for long-term storage.

# Digestion Workflow

The Digestion workflow is recommended for the following sample types. The nucleic acid that this workflow is optimized for varies by sample type; see “Recommended workflows” on page 8 for details. The Digestion workflow is not recommended for purification of RNA.

- Environmental samples
- Feces
- Hair follicles
- Swabs—environmental or fecal
- Tissue or organ

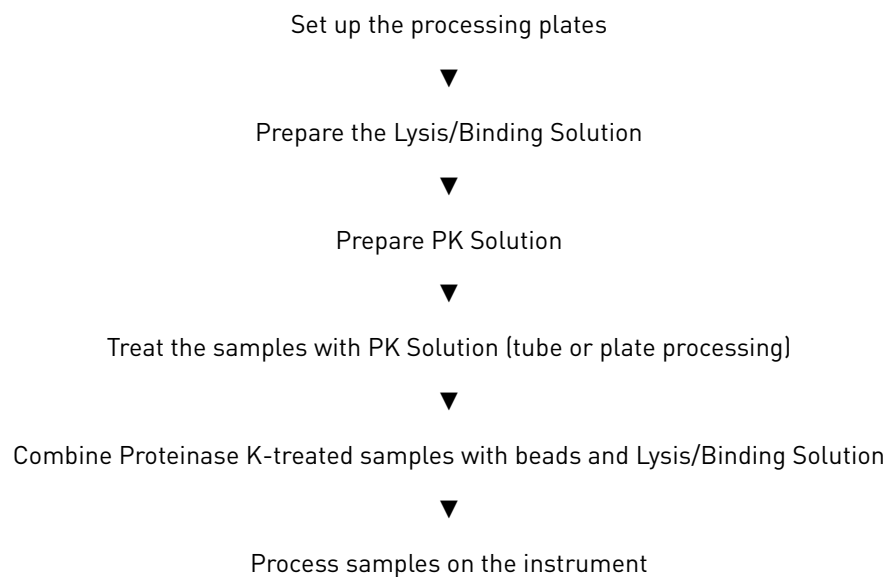
Follow this procedure if you are using these instruments:

- KingFisher™ Flex
- MagMAX™ Express-96

Follow Appendix B, “Purification with the KingFisher™ Duo Prime or KingFisher™ mL instrument” if you are using these instruments:

- KingFisher™ Duo Prime
- KingFisher™ mL

## Workflow: Digestion



## Set up the processing plates

1. Set up the processing plates.

Table 10 Plate setup: KingFisher™ Flex or MagMAX™ Express-96 instrument

Plate ID	Plate position <sup>[1]</sup>	Plate type	Reagent	Volume per well
Wash Plate 1	2	Deep Well	MagMAX™ CORE Wash Solution 1	500 µL
Wash Plate 2	3	Deep Well	MagMAX™ CORE Wash Solution 2	500 µL
Elution	4	Standard	MagMAX™ CORE Elution Buffer	90 µL
Tip Comb	5	Standard	Place a tip comb in the plate.	

<sup>[1]</sup> Position on the instrument.

2. (Optional) To prevent evaporation and contamination, cover the prepared processing plates with sealing foil until they are loaded into the instrument.

## Prepare the Lysis/Binding Solution

1. Combine the following components for the required number of samples plus 10% overage.

Component	Volume per sample
MagMAX™ CORE Lysis Solution	350 µL
MagMAX™ CORE Binding Solution	350 µL
<b>Total Lysis/Binding Solution (-IPC)</b>	<b>700 µL</b>
<i>(Optional)</i> Internal positive control (IPC), one of the following:	
VetMAX™ Xeno™ Internal Positive Control DNA	2 µL
VetMAX™ Xeno™ Internal Positive Control RNA	2 µL
Internal positive control (IPC) supplied with your VetMAX™ PCR Kit	As indicated in the instructions for the kit
<b>Total Lysis/Binding Solution (+IPC)</b>	<b>700 µL + volume of IPC</b>

2. Mix by inverting the tube or bottle at least 10 times.

*(Optional)* Store Lysis/Binding Solution at room temperature for up to 24 hours.

## Prepare PK Solution

Prepare PK Solution immediately before use.

1. Combine the following components for the required number of samples plus 10% overage.

Component	Volume per sample
PK Buffer for MagMAX™ -96 DNA Multi-Sample Kit	90 µL
MagMAX™ CORE Proteinase K	10 µL
<b>Total PK Solution</b>	<b>100 µL</b>

2. Invert the tube several times to mix, then centrifuge briefly to collect contents at the bottom of the tube.
3. Proceed immediately to the next step:
  - **For tube processing**—proceed to “Treat the samples with PK Solution (tube processing)” on page 25.
  - **For plate processing**—proceed to “Treat the samples with PK Solution (plate processing)” on page 27.



## Treat the samples with PK Solution

### Treat the samples with PK Solution (tube processing)

Treat samples with PK Solution according to the sample type.

Sample type	Procedure
Environmental samples Feces	<ol style="list-style-type: none"> <li>1. Transfer 0.2–0.3 g of sample to a 2-mL tube.</li> <li>2. Add 1 mL of PBS, pH 7.4, then vortex vigorously for 3 minutes.</li> <li>3. Centrifuge at <math>100 \times g</math> for 1 minute.</li> <li>4. Transfer 200 <math>\mu</math>L of the supernatant to a new tube.</li> <li>5. Add 100 <math>\mu</math>L of PK Solution to the transferred supernatant, then vortex briefly to mix.</li> <li>6. Incubate for 30 minutes at 55°C.</li> <li>7. Centrifuge at <math>15,000 \times g</math> for 2 minutes.</li> <li>8. Proceed with 200 <math>\mu</math>L of digested sample.</li> </ol>
Hair follicles	<ol style="list-style-type: none"> <li>1. Place 10–15 hair follicles in a 2-mL tube.</li> <li>2. Add 100 <math>\mu</math>L of PK Solution to the sample.</li> <li>3. Incubate for 30 minutes at 55°C.</li> <li>4. Centrifuge briefly to collect the contents to the bottom of the tube.</li> <li>5. Proceed with the volume of digested sample that is available to pipet. The available volume will be less than 100 <math>\mu</math>L.</li> </ol>
Swabs— environmental or fecal	<ol style="list-style-type: none"> <li>1. <b>Fecal samples</b>—swirl a clinical swab in a fecal sample. <b>Environmental swabs</b>—proceed with an environmental swab.</li> <li>2. Add 1 mL of PBS, pH 7.4 to a 2-mL tube.</li> <li>3. Swirl the swab in the PBS, pH 7.4 for 5–10 seconds, removing as much sample material as possible, then discard the swab. Alternatively, break off the swab tip and leave the swab in the PBS, pH 7.4.</li> <li>4. Vortex vigorously for 3 minutes, or until the sample is suspended.</li> <li>5. Centrifuge at <math>100 \times g</math> for 1 minute.</li> <li>6. Transfer 200 <math>\mu</math>L of the supernatant to a new tube.</li> <li>7. Add 100 <math>\mu</math>L of PK Solution to the transferred supernatant, then vortex briefly to mix.</li> <li>8. Incubate for 30 minutes at 55°C.</li> <li>9. Centrifuge at <math>15,000 \times g</math> for 2 minutes.</li> <li>10. Proceed with 200 <math>\mu</math>L of digested sample.</li> </ol>
Tissue or organ	<ol style="list-style-type: none"> <li>1. Transfer 20–30 mg of tissue to a 2-mL tube.</li> <li>2. Add 100 <math>\mu</math>L of PK Solution to the sample.</li> <li>3. Incubate for 2 hours at 55°C.</li> <li>4. Centrifuge briefly to collect the contents to the bottom of the tube.</li> <li>5. Proceed with the volume of digested sample that is available to pipet. The available volume will be less than 100 <math>\mu</math>L.</li> </ol>

Sample type	Procedure
	Use a P1000 pipette tip to transfer the viscous sample.

Treat the samples with PK Solution (plate processing)

Treat samples with PK Solution according to the sample type.

Sample type	Procedure
Environmental samples Feces	<ol style="list-style-type: none"> <li>1. Transfer 0.2–0.3 g of sample to a well of a 2-mL tube.</li> <li>2. Add 1 mL of PBS, pH 7.4 to each sample, then vortex vigorously for 3 minutes.</li> <li>3. Centrifuge at <math>100 \times g</math> for 1 minute.</li> <li>4. Transfer 200 <math>\mu</math>L of each supernatant to a deep-well plate.</li> <li>5. Add 100 <math>\mu</math>L of PK Solution to each transferred supernatant, then pipet up and down to mix.</li> <li>6. Seal the plate with sealing foil.</li> <li>7. Incubate for 30 minutes at 55°C.</li> <li>8. Centrifuge at <math>3,000 \times g</math> for 5 minutes.</li> <li>9. Proceed with 200 <math>\mu</math>L of digested sample.</li> </ol>
Hair follicles	<ol style="list-style-type: none"> <li>1. Place 10–15 hair follicles in a well of a deep-well plate.</li> <li>2. Add 100 <math>\mu</math>L of PK Solution to each sample.</li> <li>3. Seal the plate with sealing foil.</li> <li>4. Incubate for 30 minutes at 55°C.</li> <li>5. Centrifuge briefly to collect the contents to the bottom of the plate.</li> <li>6. Proceed with the volume of digested sample that is available to pipet. The available volume will be less than 100 <math>\mu</math>L.</li> </ol>
Swabs—environmental or fecal	<ol style="list-style-type: none"> <li>1. <b>Fecal samples</b>—swirl a clinical swab in a fecal sample. <b>Environmental swabs</b>—proceed with an environmental swab.</li> <li>2. Add 1 mL of PBS, pH 7.4 to a 2-mL tube.</li> <li>3. Swirl the swab in the PBS, pH 7.4 for 5–10 seconds, removing as much sample material as possible, then discard the swab. Alternatively, break off the swab tip and leave the swab in the PBS, pH 7.4.</li> <li>4. Vortex vigorously for 3 minutes, or until the samples are suspended.</li> <li>5. Centrifuge at <math>100 \times g</math> for 1 minute.</li> <li>6. Transfer 200 <math>\mu</math>L of each supernatant to a deep-well plate.</li> <li>7. Add 100 <math>\mu</math>L of PK Solution to each transferred supernatant, then pipet up and down to mix.</li> <li>8. Seal the plate with sealing foil.</li> <li>9. Incubate for 30 minutes at 55°C.</li> <li>10. Centrifuge at <math>3,000 \times g</math> for 2 minutes.</li> <li>11. Proceed with 200 <math>\mu</math>L of digested sample.</li> </ol>
Tissue or organ samples	<ol style="list-style-type: none"> <li>1. Transfer 20–30 mg of tissue to a well of a deep-well plate.</li> <li>2. Add 100 <math>\mu</math>L of PK Solution to each sample.</li> <li>3. Seal the plate with sealing foil.</li> <li>4. Incubate for 2 hours at 55°C.</li> </ol>

Sample type	Procedure
	<ol style="list-style-type: none"> <li>Centrifuge briefly to collect the contents to the bottom of the plate.</li> <li>Proceed with the volume of digested sample that is available to pipet. The available volume will be less than 100 <math>\mu\text{L}</math>. Use a P1000 pipette tip to transfer the viscous sample.</li> </ol>

## Combine Proteinase K-treated samples with beads and Lysis/Binding Solution

- Vortex the tube of MagMAX™ CORE Magnetic Beads several times to resuspend the beads, then add 20  $\mu\text{L}$  of the beads to the required wells in the plate or tube strip.

**Note:** Do not use Bead/PK Mix.

- Add the appropriate volume of each Proteinase K-treated sample to a well with beads.

For...	Use...
Environmental samples, feces Swabs	200 $\mu\text{L}$
Hair follicles Tissue or organ samples	Up to 100 $\mu\text{L}$

- Mix the sample with beads for 2 minutes at room temperature according to your mixing method.
  - Using a plate shaker**—Shake vigorously for 2 minutes (see “Determine the maximum plate shaker setting” on page 9).
  - By pipetting**—Pipet up and down several times, then incubate for 2 minutes at room temperature. (For downstream processing on the KingFisher™ mL Magnetic Particle Processor, you must mix by pipetting.)
- Add 700  $\mu\text{L}$  of Lysis/Binding Solution to each sample.
- Immediately proceed to process samples on the instrument (next section).

## Process samples on the instrument

1. Select the appropriate script on the instrument (see “Download and install the script” on page 10).
2. Start the run, then load the prepared plates or tube strips in the appropriate positions when prompted by the instrument.

Store purified nucleic acid on ice for immediate use, at  $-20^{\circ}\text{C}$  for up to 1 month, or at  $-80^{\circ}\text{C}$  for long-term storage.

# 6

## Lysis Incubation Workflow

The Lysis Incubation workflow is recommended for ear punches that require processing with:

- An extended lysis step before nucleic acid isolation.
- Addition of punches directly to a lysis solution.

Follow this procedure if you are using these instruments:

- KingFisher™ Flex
- MagMAX™ Express-96

Follow Appendix B, “Purification with the KingFisher™ Duo Prime or KingFisher™ mL instrument” if you are using these instruments:

- KingFisher™ Duo Prime
- KingFisher™ mL

### Workflow: Lysis Incubation

The Lysis Incubation workflow can be performed with a 15-minute or an overnight incubation in Lysis Solution. If samples are incubated overnight, set up the processing plates and prepare Lysis/Binding/Bead Mix after the incubation is complete.

#### 15-min incubation

Set up the processing plates



Prepare Lysis/Binding/Bead Mix



Prepare the ear punch lysate  
(15-min incubation)



Combine the lysate with Proteinase K and Lysis/  
Binding/Bead Mix



Process samples on the instrument

#### Overnight (16–18 hours) incubation

Prepare the ear punch lysate  
(overnight incubation; 16–18 hours)



Set up the processing plates



Prepare Lysis/Binding/Bead Mix



Combine the lysate with Proteinase K and Lysis/  
Binding/Bead Mix



Process samples on the instrument

## Set up the processing plates

1. Set up the processing plates.

Table 11 Plate setup: KingFisher™ Flex or MagMAX™ Express-96 instrument

Plate ID	Plate position <sup>[1]</sup>	Plate type	Reagent	Volume per well
Wash Plate 1	2	Deep Well	MagMAX™ CORE Wash Solution 1	500 µL
Wash Plate 2	3	Deep Well	MagMAX™ CORE Wash Solution 2	500 µL
Elution	4	Standard	MagMAX™ CORE Elution Buffer	90 µL
Tip Comb	5	Standard	Place a tip comb in the plate.	

<sup>[1]</sup> Position on the instrument.

2. (Optional) To prevent evaporation and contamination, cover the prepared processing plates with sealing foil until they are loaded into the instrument.

## Prepare Lysis/Binding/Bead Mix

1. Combine the following components, in the order indicated, for the required number of samples plus 10% overage.

Component	Volume per sample
MagMAX™ CORE Lysis Solution	350 µL
MagMAX™ CORE Binding Solution	350 µL
MagMAX™ CORE Magnetic Beads	20 µL
<b>Total Lysis/Binding/Bead Mix (-IPC)</b>	<b>720 µL</b>
<i>(Optional)</i> Internal positive control (IPC), one of the following:	
VetMAX™ Xeno™ Internal Positive Control DNA	2 µL
VetMAX™ Xeno™ Internal Positive Control RNA	2 µL
Internal positive control (IPC) supplied with your VetMAX™ PCR Kit	As indicated in the instructions for the kit
<b>Total Lysis/Binding/Bead Mix (+IPC)</b>	<b>720 µL + volume of IPC</b>

2. Mix by inverting the tube or bottle at least 10 times.

## Prepare the ear punch lysate

1. Add 300 µL of MagMAX™ CORE Lysis Solution to each ear punch.
2. Incubate without shaking at room temperature for the desired time.
  - 15 minutes
  - Overnight (16–18 hours)

3. Proceed with individual or pooled supernatants.

For...	Do this...
Individual samples	Proceed with 250 $\mu$ L of supernatant.
Pooled samples	<ol style="list-style-type: none"> <li>1. Combine 50 <math>\mu</math>L of individual supernatants in a 2-mL microcentrifuge tube.</li> <li>2. If the volume of pooled supernatants is less than 250 <math>\mu</math>L, add MagMAX™ CORE Lysis Solution to a total of 250 <math>\mu</math>L.</li> <li>3. Vortex briefly to mix the pooled samples.</li> <li>4. Proceed with 250 <math>\mu</math>L of pooled supernatant.</li> </ol> <p>For example:</p> <ul style="list-style-type: none"> <li>• For a pool of 10 samples, the combined volume is 500 <math>\mu</math>L (10 <math>\times</math> 50 <math>\mu</math>L). Proceed to the next step with 250 <math>\mu</math>L of the pool.</li> <li>• For a pool of 4 samples, the combined volume is 200 <math>\mu</math>L (4 <math>\times</math> 50 <math>\mu</math>L). Add 50 <math>\mu</math>L of MagMAX™ CORE Lysis Solution and proceed to the next step with the 250-<math>\mu</math>L pool.</li> </ul>
Individual analysis of a positive pool	Proceed with the remaining supernatant of each individual sample in the positive pool. The volume may be less than 250 $\mu$ L.

Store individual and pooled lysates for retesting: up to 48 hours at room temperature, or longer term below  $-16^{\circ}\text{C}$ .

## Combine the lysate with Proteinase K and Lysis/Binding/Bead Mix

1. Add 10  $\mu$ L of MagMAX™ CORE Proteinase K to the required wells in the plate or tube strip.
2. Add 250  $\mu$ L of individual or pooled supernatant.
3. Mix the supernatant with Proteinase K by pipetting up and down several times, then incubate for 2 minutes at room temperature.
4. Invert the tube of Lysis/Binding/Bead Mix several times to resuspend the beads, then add 720  $\mu$ L of Lysis/Binding/Bead Mix to each sample.
5. Immediately proceed to process samples on the instrument (next section).



## Process samples on the instrument

1. Select the appropriate script on the instrument (see “Download and install the script” on page 10).
2. Start the run, then load the prepared plates or tube strips in the appropriate positions when prompted by the instrument.

Store purified nucleic acid on ice for immediate use, at  $-20^{\circ}\text{C}$  for up to 1 month, or at  $-80^{\circ}\text{C}$  for long-term storage.



# Troubleshooting

Observation	Possible cause	Recommended action
The eluate is light brown in color	Magnetic beads were carried over into the eluate.	<p>A small quantity of beads in the sample does not inhibit RT-PCR or PCR reactions.</p> <p>Remove the beads from the eluted nucleic acid by placing the plate or tube strip on a magnetic stand (~1 minute), then transfer the nucleic acid solution to a new nuclease-free plate or tube strip.</p>
<p>Poor or no RNA or DNA signal (that is, the <math>C_t</math> value is higher than expected)</p> <p>In test samples, the <math>C_t</math> value of the IPC target is outside of the validated value range (non-compliant IPC <math>C_t</math> value; invalid sample).</p>	<p>Inhibitors are present in the recovered nucleic acid.</p> <p>These workflows yield high-quality nucleic acid for most samples. However, samples that contain exceptionally high amounts of inhibitors can carry over inhibitors at levels sufficient to affect RT-PCR or PCR.</p>	<ol style="list-style-type: none"> <li>1. Dilute the invalid nucleic acid sample 1:10 in 1X TE buffer.</li> <li>2. Perform a new PCR analysis with the diluted nucleic acid.</li> </ol> <ul style="list-style-type: none"> <li>• If the diluted nucleic acid is positive for the target, or if it is negative for the target with a compliant IPC <math>C_t</math> value, the result is validated.</li> <li>• If the diluted nucleic acid is negative for the target with a non-compliant IPC <math>C_t</math> value, the result is not validated.</li> </ul> <p>In this case, dilute the original biological sample 1:10 in 1X PBS, then repeat the purification and PCR.</p> <p>If the result is still not validated, then repeat the purification and PCR on a new biological sample.</p>
	<p>Samples with high amounts of nucleic acid, such as tissue, avian blood, and bacterial cultures, can saturate the magnetic beads. Bead saturation reduces nucleic acid recovery.</p>	<p>For the samples that show reduced recovery of the IPC RNA or DNA, dilute samples 1:2, 1:4, 1:8, and 1:16 in 1X PBS. Use the dilution that shows the best IPC recovery.</p>
	<p>The IPC DNA or RNA did not bind efficiently to the magnetic beads, due to extracellular material in the sample.</p>	<p>Add MagMAX™ CORE Magnetic Beads to the Lysis/Binding Solution, instead of preparing Bead/PK Mix or adding beads directly to the sample.</p>
	<p>Repeat the purification using the Complex workflow.</p>	
Poor yield of viral RNA from tissue, fecal or environmental samples, or swabs	The Digestion workflow was used for viral nucleic acid purification.	Follow the appropriate workflow. See “Recommended workflows” on page 8.

Observation	Possible cause	Recommended action
Well-to-well variation in RNA/DNA yield from replicate samples	The magnetic beads were not fully resuspended/dispersed.	<p>In general, the magnetic beads disperse more easily when the temperature of the mixture is &gt; 20°C. Be sure that you:</p> <ul style="list-style-type: none"> <li>• Vortex the magnetic beads thoroughly before preparing a bead mix.</li> <li>• Fully resuspend the bead mix before adding it to the samples.</li> </ul>
Positive samples are clustered in the PCR plate	High-titer samples (exhibiting a low or early C <sub>t</sub> ) have contaminated nearby wells.	Repeat the nucleic acid purification of the positive or suspect samples without the high-titer samples.
	If the same plate layout is used from nucleic acid purification through PCR, it can be difficult to determine if contamination occurred during nucleic acid purification or during PCR.	Avoid splashing when pipetting the reagents or samples.



# Purification with the KingFisher™ Duo Prime or KingFisher™ mL instrument

## Required materials not supplied

Unless otherwise indicated, all materials are available through [thermofisher.com](http://thermofisher.com).  
MLS: Fisher Scientific ([fisherscientific.com](http://fisherscientific.com)) or other major laboratory supplier.

**Table 12** Materials required for processing on the KingFisher™ Duo Prime instrument

Item	Source
KingFisher™ Duo Prime Purification System	5400110
KingFisher™ Duo Combi pack for Microtiter 96 Deepwell plate (tip combs, plates and elution strips for 96 samples)	97003530
KingFisher™ Duo Elution Strip, 40 pieces <sup>[1]</sup>	97003520
KingFisher™ Duo 12-tip comb, for Microtiter 96 Deepwell plate, 50 pieces <sup>[1]</sup>	97003500
KingFisher™ Flex Microtiter Deepwell 96 plates <sup>[1]</sup>	95040460

<sup>[1]</sup> Included in the KingFisher™ Duo Combi pack (Cat. No. 97003530).

**Table 13** Materials required for processing on the KingFisher™ mL instrument

Item	Source
KingFisher™ mL Purification System	5400050
KingFisher™ mL Tubes and tip combs for 240 samples	97002141
KingFisher™ mL Tip comb, 800 pieces	97002111
KingFisher™ mL Tube, 20 x 45 pieces	97002121

## Purification procedure

**Note:** When performing this procedure for processing on the KingFisher™ mL instrument, mix samples by pipetting up and down. Do not use a plate shaker with the large tube strips required by this instrument.

1. Follow the workflow for your sample type, starting with sample lysate preparation through combining the samples with beads and lysis solution.

**Note:** Do not set up processing plates or tubes before preparing samples.

2. Add MagMAX™ CORE Wash Solutions and MagMAX™ CORE Elution Buffer to the indicated positions, according to your instrument.

Load the Tip Comb and all of the plates or tube strips at the same time. The instrument does not prompt you to load items individually.

**Table 14** Plate setup: KingFisher™ Duo Prime instrument

Row ID	Row in the plate	Plate type	Reagent	Volume per well
Sample	A	Deep Well	Sample lysate/bead mix	Varies by sample
Wash 1	B		MagMAX™ CORE Wash Solution 1	500 µL
Wash 2	C		MagMAX™ CORE Wash Solution 2	500 µL
Elution <sup>[1]</sup>	Separate tube strip <sup>[2]</sup>	Elution strip	MagMAX™ CORE Elution Buffer	90 µL
Tip Comb	H	Deep Well	Place a tip comb in the plate.	

<sup>[1]</sup> Ensure that the elution strip is placed in the correct direction in the elution block.

<sup>[2]</sup> Placed on the heating element.

**Table 15** Tube strip setup: KingFisher™ mL instrument

Position ID	Tube strip position	Tube	Reagent	Volume per well
Sample	1	Standard	Sample lysate/bead mix	Varies by sample
Wash 1	2		MagMAX™ CORE Wash Solution 1	500 µL
Wash 2	3		MagMAX™ CORE Wash Solution 2	500 µL
Elution	4		MagMAX™ CORE Elution Buffer	90 µL
Tip Comb	N/A	N/A	Slide the tip comb into the tip comb holder.	

3. Follow “Process samples on the instrument” on page 15.



# Safety



**WARNING! GENERAL SAFETY.** Using this product in a manner not specified in the user documentation may result in personal injury or damage to the instrument or device. Ensure that anyone using this product has received instructions in general safety practices for laboratories and the safety information provided in this document.

- Before using an instrument or device, read and understand the safety information provided in the user documentation provided by the manufacturer of the instrument or device.
  - Before handling chemicals, read and understand all applicable Safety Data Sheets (SDSs) and use appropriate personal protective equipment (gloves, gowns, eye protection, etc). To obtain SDSs, see the “Documentation and Support” section in this document.
-



## Chemical safety



---

**WARNING! GENERAL CHEMICAL HANDLING.** To minimize hazards, ensure laboratory personnel read and practice the general safety guidelines for chemical usage, storage, and waste provided below. Consult the relevant SDS for specific precautions and instructions:

- Read and understand the Safety Data Sheets (SDSs) provided by the chemical manufacturer before you store, handle, or work with any chemicals or hazardous materials. To obtain SDSs, see the “Documentation and Support” section in this document.
  - Minimize contact with chemicals. Wear appropriate personal protective equipment when handling chemicals (for example, safety glasses, gloves, or protective clothing).
  - Minimize the inhalation of chemicals. Do not leave chemical containers open. Use only with adequate ventilation (for example, fume hood).
  - Check regularly for chemical leaks or spills. If a leak or spill occurs, follow the manufacturer's cleanup procedures as recommended in the SDS.
  - Handle chemical wastes in a fume hood.
  - Ensure use of primary and secondary waste containers. (A primary waste container holds the immediate waste. A secondary container contains spills or leaks from the primary container. Both containers must be compatible with the waste material and meet federal, state, and local requirements for container storage.)
  - After emptying a waste container, seal it with the cap provided.
  - Characterize (by analysis if necessary) the waste generated by the particular applications, reagents, and substrates used in your laboratory.
  - Ensure that the waste is stored, transferred, transported, and disposed of according to all local, state/provincial, and/or national regulations.
  - **IMPORTANT!** Radioactive or biohazardous materials may require special handling, and disposal limitations may apply.
-

## Biological hazard safety



**WARNING! BIOHAZARD.** Biological samples such as tissues, body fluids, infectious agents, and blood of humans and other animals have the potential to transmit infectious diseases. Conduct all work in properly equipped facilities with the appropriate safety equipment (for example, physical containment devices). Safety equipment can also include items for personal protection, such as gloves, coats, gowns, shoe covers, boots, respirators, face shields, safety glasses, or goggles. Individuals should be trained according to applicable regulatory and company/ institution requirements before working with potentially biohazardous materials. Follow all applicable local, state/provincial, and/or national regulations. The following references provide general guidelines when handling biological samples in laboratory environment.

- U.S. Department of Health and Human Services, *Biosafety in Microbiological and Biomedical Laboratories (BMBL)*, 5th Edition, HHS Publication No. (CDC) 21-1112, Revised December 2009; found at:  
[www.cdc.gov/biosafety/publications/bmb15/BMBL.pdf](http://www.cdc.gov/biosafety/publications/bmb15/BMBL.pdf)
  - World Health Organization, *Laboratory Biosafety Manual*, 3rd Edition, WHO/CDS/CSR/LYO/2004.11; found at:  
[www.who.int/csr/resources/publications/biosafety/Biosafety7.pdf](http://www.who.int/csr/resources/publications/biosafety/Biosafety7.pdf)
-



# Documentation and support

## Related documentation

Document	Publication Number
<i>Thermo Scientific™ KingFisher™ Flex User Manual</i>	N07669
<i>Thermo Scientific™ KingFisher™ Duo Prime Technical Manual</i>	N16621
<i>Thermo Scientific™ KingFisher™ mL User Manual</i>	1508260
<i>Applied Biosystems™ MagMAX™ Express 96 User Manual</i>	N07849
<i>MagMAX™ CORE Mechanical Lysis Module User Guide</i>	MAN0015945

## Customer and technical support

Visit [thermofisher.com/support](http://thermofisher.com/support) for the latest in services and support, including:

- Worldwide contact telephone numbers
- Product support, including:
  - Product FAQs
  - Software, patches, and updates
  - Training for many applications and instruments
- Order and web support
- Product documentation, including:
  - User guides, manuals, and protocols
  - Certificates of Analysis
  - Safety Data Sheets (SDSs; also known as MSDSs)

**Note:** For SDSs for reagents and chemicals from other manufacturers, contact the manufacturer.

## Limited product warranty

Life Technologies Corporation and/or its affiliate(s) warrant their products as set forth in the Life Technologies' General Terms and Conditions of Sale found on Life Technologies' website at [www.thermofisher.com/us/en/home/global/terms-and-conditions.html](http://www.thermofisher.com/us/en/home/global/terms-and-conditions.html). If you have any questions, please contact Life Technologies at [www.thermofisher.com/support](http://www.thermofisher.com/support).

