

Load N' Go™ *Quick-DNA/RNA*™ HT

Automation-ready: Rapid, high-throughput nucleic acid extraction from any biological sample

Highlights

- **Save Time and Focus on Discovery:** Pre-filled 96-well reagent plate technology that offers multi-platform compatibility and reduces hands-on time by 75%.
- **Precision Meets Versatility:** High-throughput, magnetic-bead based purification of DNA and RNA from plasma, serum, urine, cell culture media, blood, saliva, cellular suspensions, swab, fecal, and biopsy samples.

Catalog Numbers:
R2152



Scan with your smart-phone camera to
view the online protocol/video.



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Product Contents

Load N' Go™ Quick-DNA/RNA™ HT	Volume / Qty
DNA/RNA Shield™ (2x)	125 ml
Proteinase K (lyophilized)	60 mg
Proteinase K Storage Buffer	10 ml
Plate 1: HT MagBinding Beads	250 µl x 96
Plate 2: HT Binding Plate	600 µl x 96
Plate 3: MagBead DNA/RNA Wash 1	250 µl x 96
Plate 4: MagBead DNA/RNA Wash 2	250 µl x 96
Plate 5: E-Wash 1	250 µl x 96
Plate 6: E-Wash 2	250 µl x 96
Plate 7: Elution	30 µl x 96
96 Tip Combs ¹ (For V-Bottom Deep Well Plate)	2 pc
Instruction Manual	1 pc

- Materials/Equipment Needed (user provided)**

- ✓ Beta-mercaptoethanol (β-Me)
- ✓ Nuclease-free water
- ✓ Centrifuge with microplate carriers
- ✓ Vortex Mixer
- ✓ Liquid handler or bead mover laboratory automation

- Materials Available Separately (not provided)**

- ZR BashingBead™ Lysis Tubes (S6003-50; 2.0 mm beads), (S6012-50; 0.1 & 0.5 mm beads), (S6014-50; 0.1 & 2.0 mm beads)
- DNase I Set (E1011; 1500 U DNase I (lyophilized) supplied with DNA Digestion Buffer, 4 ml)
- DNA/RNA Prep Buffer (D7010-2-50, 50 ml)

¹ Compatible with platforms such as KingFisher™ Flex, KingFisher™ Apex, IsoPure™ 96, Auto-Pure 96 systems.

Specifications

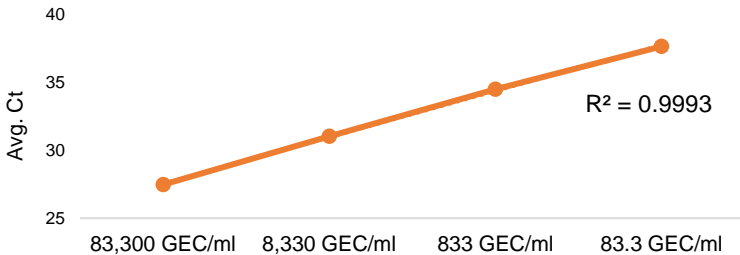
- **Sample Sources** – Any biological sample (e.g., swabs, liquids, cells, tissue, etc.), and samples collected/stored in media (e.g., UTM, VTM, saline, PBS, DNA/RNA Shield™, PAXgene®, RNA/ater™, RNAprotect®, etc.).
- **Binding Capacity** – 5 µg DNA/RNA per prep.
- **Elution Volume** – 30 µl **DNase/RNase-Free Water**.
- **Purity** – High quality, inhibitor-free DNA/RNA is eluted with DNase/RNase Free Water and is suitable for all downstream applications including PCR and Next-Generation Sequencing.
- **Storage Temperature and Stability** –
 - ✓ Store all components (i.e., buffers/reagents, columns) at room temperature (15-30°C).
 - ✓ Expiration dates for each of the unopened components are indicated on the individual component labels. These storage conditions apply to both opened and unopened components.
 - ✓ Eluted DNA/RNA can be used immediately or stored frozen (-20/-80°C).

Product Description

The **Load N' Go™ Quick-DNA/RNA™ HT** kit is intended for rapid, high-throughput nucleic acid extraction from any biological or clinical sample (e.g., swabs – nasal/nasopharyngeal, oropharyngeal, etc.; biological liquids – blood, plasma, serum, saliva, sputum, cells in suspension, etc.; tissue – needle biopsies, LCM, etc.) and/or samples stored in most collection matrices and devices (e.g., UTM, VTM, DNA/RNA Shield™, RNAlater™, RNAProtect®, etc.).

The kit is compatible with robotic-type sample processors (e.g., bead movers, liquid handlers) in combination with sensitive downstream molecular amplification assays. High-quality DNA/RNA extracted with the **Load N' Go™ Quick-DNA/RNA™ HT** kit can be used for Next-Gen sequencing, RT/qPCR and more.

Limit of Detection Preliminary Assay from Biological Specimens using Automated Extraction



Concentration in Dilution Tested (GEC/ml)

Concentration in Dilution	83,300 GEC/ml (5,000 GEC/rxn)	8,330 GEC/ml (500 GEC/rxn)	833 GEC/ml (50 GEC/rxn)	83.3 GEC/ml (5 GEC/rxn)	8.33 GEC/ml (0.5 GEC/rxn)
Avg. Ct	27.5	31.0	34.5	37.6	Not detected
Positive, n=5	5/5	5/5	5/5	5/5	0/5

Automated extraction of whole genome viral RNA (i.e., SARS-CoV-2) spiked in sputum/swab samples collected in DNA/RNA Shield™ was performed with the **Load N' Go™ Quick-DNA/RNA™ HT** kit format, followed by quantification by RT-qPCR. Preliminary limit of detection (LoD) assay determined the lowest concentration for which 5/5 independent replicates tested positive.

Protocol

The protocol consists of: (I) Loading Scripts, (II) Buffer Preparation, (III) Sample Preparation and (IV) DNA/RNA Purification.

(I) Loading Scripts

Contact automation@zymoresearch.com to obtain the script and other reference materials related to this **Load N' Go™ Quick-DNA/RNA™ HT** kit on your automation platform¹.

Examples of popular systems that are compatible include, but are not limited to:

- ✓ AllSheng Auto-Pure 96
- ✓ Accuris IsoPure™ 96
- ✓ Thermo Scientific™ KingFisher™ Flex
- ✓ Thermo Scientific™ KingFisher™ Apex
- ✓ Tecan Fluent®
- ✓ Hamilton Microlab® Star™
- ✓ Opentrons™ OT-2

If you are unsure about compatibility, reach out to automation@zymoresearch.com for verification.

(II) Buffer Preparation

- ✓ Reconstitute lyophilized **Proteinase K** at 20 mg/ml with **Proteinase K Storage Buffer** and mix by vortexing. Use immediately or store frozen aliquots:

For each **60 mg**, add 3.12 ml **Proteinase K Storage Buffer**

- ✓ Optional: add 3 µl beta-mercaptoethanol (user supplied) to each well for **Plate 2: HT Binding Plate**, (final 0.5% (v/v)).
- ✓ Optional: To prepare DNA/RNA Shield™ (1X)², dilute the 2X concentrate with an equal volume of nuclease-free water (user provided) (1:1) and mix well.

¹ If intended workflow requires a DNase I treatment, specify this in the outreach email and the appropriate scripts will be sent out.

² DNA/RNA Shield™ is a sample collection medium for storage and preservation of nucleic acids. It also inactivates pathogens and prevents viral infectivity. Specimens stored and transported in DNA/RNA Shield™ can be processed directly without reagent removal, using standard laboratory operating procedures, for the detection of nucleic acids with molecular amplification assays.

(III) Sample Preparation

- ✓ Perform all steps at room temperature (15-30°C)

Sample Input – Depending on the sample type, up to 200 µl can be processed per prep (see examples below).

Swabs		
Nasal/Nasopharyngeal	Oropharyngeal	Buccal/cheek
Vaginal	Fecal	
<p>Stored in UTM, VTM, saline, PBS, DNA/RNA Shield™, etc.</p> <p>Optional - To inactivate, store and preserve samples at room temperature, add 100 µl DNA/RNA Shield™ (2X concentrate)¹ to 100 µl sample (1:1). Mix well².</p>		

Liquids (I)		
Plasma	Serum	CSF
Cells in suspension		
<p>Optional - To inactivate, store and preserve samples at room temperature, add 100 µl DNA/RNA Shield™ (2X concentrate)¹ to 100 µl sample (1:1). Mix well².</p>		

Liquids (II)		
Whole blood	Saliva	Sputum
Urine		
<p>Recommended: To inactivate, lyse and preserve samples at room temperature, add 100 µl DNA/RNA Shield™ (2X concentrate)¹ to 100 µl sample (1:1). Mix well², centrifuge debris and process the cleared supernatant.</p>		

Tissue	
Tissue (< 5 mg)	Feces (< 20 mg)
<p>LCM, needle biopsy or samples stored in media/device.</p> <p>Recommended: To inactivate, lyse and preserve samples at room temperature, add 200 µl DNA/RNA Shield™ (1X)^{1,3} and homogenize⁴. Mix well, centrifuge debris and process the cleared supernatant.</p>	

¹ At this point, samples in DNA/RNA Shield™ can be stored at ambient temperature (4-30°C) for a month, 7 days at 35°C, or long-term (> 1 year) at -20°C or below.

² For all buffer additions and incubation steps, **mix well** for ≥1 minute by pipetting the beads up and down and/or by shaking (vortexing) at ~1,300 rpm. Optimization may be required.

³ To prepare DNA/RNA Shield™ (1X), dilute the **2X concentrate** with an equal volume of nuclease-free water (1:1) and mix well.

⁴ For efficient homogenization of tough-to-lyse tissue samples, bead-beat with ZR BashingBead™ Lysis Tubes (S6003, S6012, S6014), see Ordering Information on page 10 for more details.

(III) Sample Preparation (continued)

Proteinase K Treatment (optional)

- ✓ At this point, protein-rich samples (e.g., plasma, serum, saliva, sputum, tissue) can be treated by **Proteinase K**.
- 1. Add 1% **Proteinase K** (v/v) at 20 mg/ml¹ directly to a liquid sample. Mix well².

Note: Up to 5% Proteinase K (v/v) at 4 mg/ml can be added to protein-rich samples (e.g., tissue).
- 2. Incubate at room temperature for 15 minutes.

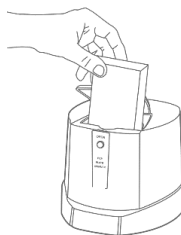
¹ For automation platforms with "dead volume" liquid handler dispensing, the lyophilized Proteinase K can be reconstituted to a working concentration of 4-20 mg/ml.

² For all buffer additions and incubation steps, **mix well** for ≥1 minute, by pipetting the beads up and down and/or by shaking (vortexing) at ~1,300 rpm. Optimization may be required.

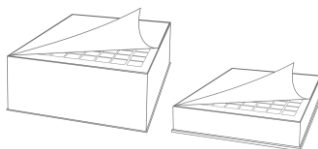
(IV) DNA/RNA Purification

- ✓ Perform all steps at room temperature (15-30°C)

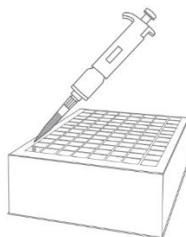
1. Spin down elution plate.



2. Remove foil seals from all plates.



3. Load 200 μ L of prepped sample¹ into each well of sample plate.



4. Run extraction script



¹ If using < 200 μ L sample, increase the total sample input volume to 200 μ L using DNA/RNA Shield (1x).

Appendices

Appendix A: FAQ

Question	Suggested Solutions
How do I avoid background contamination?	<ul style="list-style-type: none">- Clean workspace, centrifuge, and pipettes with 10% bleach to routinely to avoid contamination.- If use of kit is in an exposed environment without proper filtration. Check pipettes, pipette tips, microcentrifuge tubes, workspace, etc. for contamination.
How do I fix low DNA yield?	<p><u>Lysis Method</u></p> <ul style="list-style-type: none">- Bead beating devices that oscillate in a single dimension (only vertically or only horizontally) have been observed to inefficiently lyse very recalcitrant species. Devices that oscillate three-dimensionally or in a figure-8 motion often lyse microbes efficiently. <p><u>Input</u></p> <ul style="list-style-type: none">- Reference Section III on Page 5 for information on your input limit based on sample.
How do I know if my automation instrument is compatible?	<p>Examples of popular systems that are compatible include, <u>but are not limited to</u>:</p> <ul style="list-style-type: none">- AllSheng Auto-Pure 96- IsoPure™ 96- KingFisher™ Flex- KingFisher™ Apex- Tecan Fluent®- Hamilton Microlab® STAR™- Opentrons™ OT-2 <p>If you are unsure about compatibility, reach out to automation@zymoresearch.com for verification.</p>
How do I get scripts for this kit on my automation platform?	Please contact automation@zymoresearch.com to send a request for scripts and additional reference material.

For any other technical assistance, please email automation@zymoresearch.com

Appendix B: DNase I Treatment

- ✓ For DNA-free RNA, DNase I treatment can be performed using **DNase I Set** and **DNA/RNA Prep Buffer**, materials sold separately¹.
- ✓ To run the DNase I treatment on your automation platform, please contact automation@zymoresearch.com for additional support and scripts.

For each sample to be treated, prepare **DNase I Reaction Mix** in an RNase-free tube (not provided) and mix by gentle inversion:

DNase I Reaction Mix	
Nuclease-free water (user provided)	40 µl
DNA Digestion Buffer	5 µl
DNase I (reconstituted; 1 U/µl) ²	5 µl

Proceed with additional script and protocol supplied from automation@zymoresearch.com to complete DNase I treatment³.

¹ See Ordering Information on page 10 for more information.
² Prior to use, reconstitute lyophilized 250 U DNase I (E1009-A) to 1 U/µl (final concentration) with 275 µl nuclease-free water (user provided), mix by gentle inversion and store frozen aliquots.
³ Additional plastics will be required for the use of DNase I on an automation platform, see Ordering Information on page 10 for more details.

Ordering Information

Product Description	Catalog No.	Size
Load N' Go™ <i>Quick</i> -DNA/RNA™ HT	R2152	96 preps
DNase I Set	E1011	1500 U
DNA/RNA Shield™	R1100-250	250 ml
DNA/RNA Shield™ (2x)	R1200-125	125 ml
DNA/RNA Prep Buffer	D7010-2-50	50 ml
96 Deep Well Plate (V-Bottom 2.2 ml)	C2018-5	5 Plates
96 Deep Well Plate (V-Bottom 2.2 ml)	C2018-50	50 Plates
ZR BashingBead™ Lysis Tubes (2 mm)	S6003-50	50 Tubes
ZR BashingBead™ Lysis Tubes (0.1 & 0.5 mm)	S6012-50	50 Tubes
ZR BashingBead™ Lysis Tubes (0.1 & 2.0 mm)	S6014-50	50 Tubes

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