



ZYMO RESEARCH

The Beauty of Science is to Make Things Simple

INSTRUCTION MANUAL

Quick-gDNA™ MicroPrep

Catalog Nos. **D3020 & D3021**

Highlights

- Quick purification of high quality DNA from whole blood, plasma, serum, body fluids, buffy coat, lymphocytes, tissue, swabs or cultured cells in less than 15 minutes using innovative *Fast-Spin* column technology.
- Compatible with commonly used anticoagulants (i.e., EDTA, heparin, citrate).
- Unique extraction technology excludes the use of Proteinase K and organic denaturants.
- Isolated DNA is ideal for PCR, endonuclease digestion, bisulfite conversion/methylation detection, sequencing, genotyping, etc.

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Satisfaction of all Zymo Research products is guaranteed. If you are not satisfied with this product please call 1-888-882-9682.

Product Contents

Quick-gDNA™ MicroPrep (Kit Size)	D3020 (50 Preps.)	D3021 (200 Preps.)	Storage Temperature
Genomic Lysis Buffer*	50 ml	2 x 100 ml	Room Temp.
DNA Pre-Wash Buffer**	15 ml	50 ml	Room Temp.
g-DNA Wash Buffer	50 ml	100 ml	Room Temp.
DNA Elution Buffer	10 ml	2 x 10 ml	Room Temp.
Zymo-Spin™ IC Columns	50	200	Room Temp.
Collection Tubes	100	400	Room Temp.
Instruction Manual	1	1	-

Note - Integrity of kit components is guaranteed for up to one year from date of purchase. Reagents are routinely tested on a lot-to-lot basis to ensure they provide maximal performance and reliability.

* For optimal performance, add beta-mercaptoethanol to 0.5%(v/v) i.e., 250 µl per 50 ml or 500 µl per 100 ml.

**A precipitate may have formed in the DNA Pre-Wash Buffer during shipping. To completely resuspend the buffer, incubate the bottle at 30-37 °C for 30 minutes and mix by inversion. DO NOT MICROWAVE.

Specifications

- **Sample Sources** – Whole blood, plasma, or serum from humans, mice, rats, etc. Also, tissue, cells from culture, buccal cells, as well as a variety of biological liquids are effectively processed using this kit.
- **DNA Purity** – High-quality DNA is eluted with **DNA Elution Buffer** or water. DNA is especially well suited for PCR and other downstream applications. $A_{260}/A_{280} > 1.8$
- **DNA Size Limits** – Capable of recovering genomic DNA up to and above 40 kb. In most instances, mitochondrial DNA and viral DNA (if present) will also be recovered.
- **DNA Recovery** – Up to 5 µg total DNA is eluted into ≥ 10 µl (6 µl minimum) **DNA Elution Buffer** or water. Human whole blood will typically yield 1.5-3.5 µg DNA per 50 µl blood sampled. Mammalian tissues yield: 1-3 µg DNA per mg skeletal, heart, and brain tissues and 3-5 µg DNA per mg liver, kidney and lung tissues.
- **Product Detergent Tolerance** – $\leq 5\%$ Triton X-100, $\leq 5\%$ Tween-20, $\leq 5\%$ Sarkosyl, $\leq 0.1\%$ SDS.
- **Equipment** – microcentrifuge, vortex

Note - ™ Trademarks of Zymo Research Corporation. This product is for research use only and should only be used by trained professionals. It is not intended for use in diagnostic procedures. Some reagents included with this kit are irritants. Wear protective gloves and eye protection. Follow the safety guidelines and rules enacted by your research institution or facility.

For high-throughput purification (96-well, 5 µg DNA/well) use the **ZR-96 Quick-gDNA™** (D3010, D3011, D3012).

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Product Description

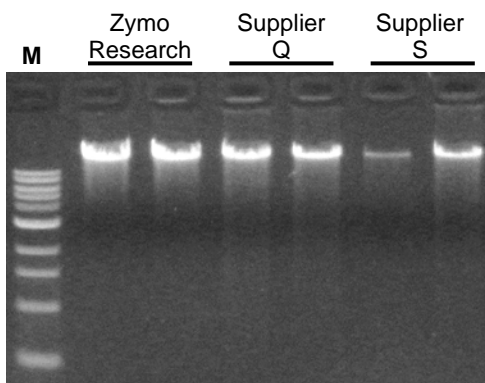
The **Quick-gDNA™ MicroPrep** is a simple procedure for the rapid isolation of total DNA (e.g., genomic, mitochondrial, viral) from a variety of biological sample sources. This product has been optimized for maximal recovery of ultra-pure DNA without RNA contamination and is compatible with whole blood (fresh or stored), serum, plasma, buffy coat, solid tissue, bone marrow and buccal cells, cells from culture, and many biological liquid samples.

For processing, simply add the specially formulated **Genomic Lysis Buffer** to a sample, vortex, and transfer the mixture to the supplied **Zymo-Spin™ Column**. There is no need for organic denaturants or Proteinase K digestion because of the unique chemistries featured in the kit. Instead, the product features *Fast-Spin* technology to yield high-quality, purified DNA in just minutes (see below). PCR inhibitors are effectively removed during the purification process. DNA purified using the **Quick-gDNA™ MicroPrep** is suitable for PCR, nucleotide blotting, DNA sequencing, restriction endonuclease digestion, bisulfite conversion/methylation analysis, and other downstream applications.



Ultra-pure DNA is ideal for...

- ✓ PCR
- ✓ Endonuclease Digestion
- ✓ Genotyping
- ✓ Bisulfite Conversion & Methylation Analysis



High yield/quality DNA is successfully isolated from porcine whole blood using the **Quick-gDNA™ MiniPrep** (D3024). Equivalent amounts (100 µl) of blood were processed without Proteinase K using the **Quick-gDNA™ MiniPrep** in half the time as compared to the kits from suppliers Q and S. Equal volumes of eluted DNA were then analyzed (in duplicate) in a 0.8% (w/v) TAE/agarose/ethidium bromide gel. The size marker "M" is a 1 kb ladder (Zymo Research).

The **ZymoBead™ Genomic DNA Kit (D3004, D3005)** and **ZR Serum DNA Kit™ (D3013)** are recommended for scaleable DNA yields >25 µg/prep. Both feature silica beads instead of a spin column.

Zymo Research offers the following for rapid, precise DNA methylation detection...

- 1.) **EZ DNA Methylation™ Kit** (D5001, D5002, D5003, D5004)
- 2.) **EZ DNA Methylation-Gold™ Kit** (D5005, D5006, D5007, D5008)
- 3.) **EZ DNA Methylation-Direct™ Kit** (D5020, D5021, D5022, D5023)

For **Technical Assistance**, please contact 1-888-882-9682 or E-mail tech@zymoresearch.com.

For the inclusion of small DNAs from serum, add 0.3 volumes isopropanol to the mixture. (For example, to a 1 ml mixture of serum and Genomic Lysis Buffer add 300 µl isopropanol.)

The column capacity is ~800 µl.

Elution of DNA from the column is dependent on pH and temperature. If water is used, ensure the pH is >6.0. Also, the total yield may be improved by eluting the DNA with Elution Buffer or water pre-equilibrated to 60-70°C.

Buffer Preparation

- ✓ For optimal performance, add beta-mercaptoethanol (user supplied) to the **Genomic Lysis Buffer** to a final dilution of 0.5%(v/v) i.e., 250 µl per 50 ml or 500 µl per 100 ml.

PROTOCOLS

Whole Blood, Serum, and Plasma Samples

The following is for the purification of DNA from whole blood, serum or plasma (the volumes can be adjusted depending on your requirements). Fresh, frozen, or preserved blood (in EDTA, citrate, or heparin) can be used. If material cannot be processed immediately, the sample can be “stabilized” for later processing (as noted below) although the immediate processing of blood samples is recommended.

1. In a 1.5 ml microcentrifuge tube, add up to 50 µl (max.) of blood, serum, or plasma to 200 µl of **Genomic Lysis Buffer**. Mix completely by vortexing 4-6 seconds, then let stand 5-10 minutes at room temperature.
2. Transfer the mixture to a **Zymo-Spin™ IC Column** in a **Collection Tube**. Centrifuge at 10,000 x g for one minute. Discard the **Collection Tube** with the flow through.
3. Transfer the **Zymo-Spin™ IC Column** to a new **Collection Tube**. Add 200 µl of **DNA Pre-Wash Buffer** to the spin column. Centrifuge at 10,000 x g for one minute.
4. Add 500 µl of **g-DNA Wash Buffer** to the spin column. Centrifuge at 10,000 x g for one minute.
5. Transfer the spin column to a clean microcentrifuge tube. Add ≥10 µl **DNA Elution Buffer** or water to the spin column. Incubate 2-5 minutes at room temperature and then centrifuge at top speed for 30 seconds to elute the DNA. The eluted DNA can be used immediately for molecular based applications or stored ≤-20°C for future use.

Delayed Processing (Stabilization) of Blood Samples: The immediate processing of blood with this kit is recommended. However, if blood cannot be processed immediately, samples can be “stabilized” in **Genomic Lysis Buffer** for processing at a later time. To do this, add *four* volumes of **Genomic Lysis Buffer** to *each* volume of whole blood (4:1), then vortex. Blood samples mixed with **Genomic Lysis Buffer** can be stored at room temperature for 1-2 weeks, 0-4°C for 1-2 months, -20°C for 6 months to a year, or <-70°C for many years. Samples stored at ≤4°C should reach room temperature prior to processing. Begin at Step 2 in the standard protocol (above) when purifying DNA from blood samples stabilized in **Genomic Lysis Buffer**.

Buccal Cells and Swabs

Buccal cells can be isolated using a rinse- or swab-based isolation method.

- A. **Rinse Method:** Vigorously rinse 10-20 ml of saline solution or mouthwash orally for 30 seconds. The more vigorous the rinsing action, the more cells that will be recovered. Spit the saline into a 50 ml tube and pellet the cells at 1,500 rpm for 5 minutes. Discard the supernatant without disturbing the cell pellet. Add 500 μ l of **Genomic Lysis Buffer** to the pellet then vortex 4-6 seconds, then let stand at room temperature for 5-10 minutes.
- B. **Swab Isolation Method:** Thoroughly rinse mouth out before isolating cells. Brush the inside of the cheek with a *buccal swab* for 15 seconds (approximately 20 brushes), making sure to cover the entire area of the inner cheek. Rinse the brush with 500 μ l of **Genomic Lysis Buffer** into a microcentrifuge tube, vortex 4-6 seconds, and then let stand at room temperature for 5-10 minutes.
 1. Transfer the mixture to a **Zymo-Spin™ IC Column** in a **Collection Tube**. Centrifuge at 10,000 \times *g* for one minute. Discard the **Collection Tube** with the flow through.
 2. Transfer the **Zymo-Spin™ IC Column** to a new **Collection Tube**. Add 200 μ l of **DNA Pre-Wash Buffer** to the spin column. Centrifuge at 10,000 \times *g* for one minute.
 3. Add 500 μ l of **g-DNA Wash Buffer** to the spin column. Centrifuge at 10,000 \times *g* for one minute.
 4. Transfer the spin column to a clean microcentrifuge tube. Add \geq 10 μ l **DNA Elution Buffer** or water to the spin column. Incubate 2-5 minutes at room temperature and then centrifuge at top speed for 30 seconds to elute the DNA. The eluted DNA can be used immediately for molecular based applications or stored \leq -20°C for future use.

The column capacity is ~800 μ l.

Solid Tissue Samples

Note: For Proteinase K digested materials (e.g., tailsnips) follow the protocol for **Cell Suspensions and Proteinase K Digested Samples** (pg. 6). Otherwise, mechanically homogenize up to 5 mg of fresh or frozen tissue in 500 μ l of **Genomic Lysis Buffer**.

1. Centrifuge the lysate at top speed (10,000 \times *g*) for 5 minutes. Making sure not to disturb the pelleted debris, transfer the supernatant to a **Zymo-Spin™ IC Column** in a **Collection Tube** and centrifuge at 10,000 \times *g* for one minute. Discard the **Collection Tube** with the flow through.
2. Transfer the **Zymo-Spin™ IC Column** to a new **Collection Tube**. Add 200 μ l of **DNA Pre-Wash Buffer** to the spin column. Centrifuge at 10,000 \times *g* for one minute.

Soft tissue samples are readily homogenized using our **Squisher™-Single**, **Squisher™-8**, and **Squisher™-96** products.

Typical yields are: 1-3 μ g DNA per mg skeletal, heart, and brain tissues and 3-5 μ g per mg liver, kidney, and lung tissues.

3. Add 500 µl of **g-DNA Wash Buffer** to the spin column. Centrifuge at 10,000 x g for one minute.
4. Transfer the spin column to a clean microcentrifuge tube. Add ≥10 µl **DNA Elution Buffer** or water to the spin column. Incubate 2-5 minutes at room temperature and then centrifuge at top speed for 30 seconds to elute the DNA. The eluted DNA can be used immediately for molecular based applications or stored ≤-20 °C for future use.

Cell Monolayer Samples

*The following procedure is designed for up to 1.0x10⁶ (max.) monolayer cells (roughly equal to one well of a 6-well plate). Although cell types and culture conditions may vary, the protocol will work with high-density growth cells (e.g., HeLa cells) as well as with low-density growth cells (e.g., neuronal cells). The procedure may be scaled up or down for increases or decreases in the amounts of monolayer cells sampled (see the **Guidelines for Monolayer Cell DNA Isolation** below).*

1. Trypsinize or manually scrape adherent cells from the growth surface of a culture flask or plate. Centrifuge the cell suspension at approximately 500 x g for 5 minutes. Remove the supernatant and add 400 µl¹ of **Genomic Lysis Buffer** directly to the cell pellet. Resuspend pellet by vortexing 4-6 seconds and let stand for 5-10 minutes at room temperature.

Alternatively: Cells can be lysed directly in the culture container by removing the medium and adding the Genomic Lysis Buffer directly to the monolayer surface.

2. Transfer the mixture to a **Zymo-Spin™ IC Column** in a **Collection Tube**. Centrifuge at 10,000 x g for one minute. Discard the **Collection Tube** with the flow through.
3. Transfer the **Zymo-Spin™ IC Column** to a new **Collection Tube**. Add 200 µl of **DNA Pre-Wash Buffer** to the spin column. Centrifuge at 10,000 x g for one minute.
4. Add 500 µl of **g-DNA Wash Buffer** to the spin column. Centrifuge at 10,000 x g for one minute.
5. Transfer the spin column to a clean microcentrifuge tube. Add ≥10 µl **DNA Elution Buffer** or water to the spin column. Incubate 2-5 minutes at room temperature and then centrifuge at top speed for 30 seconds to elute the DNA. The eluted DNA can be used immediately for molecular based applications or stored ≤-20 C for future use.

¹**Guidelines for Monolayer Cell DNA Isolation:** The above procedure is designed for the processing of 0.1-1.0x10⁶ cells. However, cell numbers (growth densities) can vary between different cell types. Table 1 (pg. 6) provides an approximation of what can be recovered from different culture containers for high-density growth cells like CV1 and HeLa cells. If processing more than 1.0x10⁵ cells, double the volume of **Genomic Lysis Buffer** added (i.e., 800 µl) to the sample.

Generally, no more than 1.0x10⁶ cells should be sampled, for larger samples will exceed the binding capacity of the spin column. See **Guidelines for Monolayer Cell Isolation** (below).

It may be necessary to centrifuge the sample mixture before transferring the supernatant to the **Zymo-Spin™ Column** to remove insoluble material that may clog the column.

The column capacity is ~800 µl.

Table 1: Culture Plate/Flask Growth Area (cm²) and Cell Number

Culture Container	Well /Flask Surface Area	Cell Number
96-well plate (each well)	0.32-0.6 cm ²	4-5x10 ⁴
24-well plate (each well)	2 cm ²	1-3x10 ⁵
12-well plate (each well)	4 cm ²	4-5x10 ⁵
6-well plate (each well)	9.5 cm ²	0.5-1x10 ⁶
T25 Culture Flask	25 cm ²	2-3x10 ⁶
T75 Culture Flask	75 cm ²	0.6-1x10 ⁷
T175 Culture Flask	175 cm ²	2-3x10 ⁷

Cell Suspensions and Proteinase K Digested Samples

The following protocol is designed for up to 200 µl of biological liquid sample including CSF, buffy coat, body fluids (semen), and cell suspensions containing less than 1.0x10⁶ cells as well as lysates derived from Proteinase K digested samples.

1. Add 4 volumes of **Genomic Lysis Buffer** to each volume of liquid sample (4:1). (e.g., add 400 µl of **Genomic Lysis Buffer** to 100 µl liquid sample). Mix briefly by vortexing, then let stand at room temperature for 5-10 minutes.

Note: For Proteinase K digested material, add 4 volumes of **Genomic Lysis Buffer** to each volume of lysate then mix briefly by vortexing. Centrifuge the mixture at 10,000 x g for 5 minutes. Transfer up to 800 µl supernatant to the Zymo-Spin™ IC Column in Step 2.

2. Transfer the mixture to a **Zymo-Spin™ IC Column** in a **Collection Tube**. Centrifuge at 10,000 x g for one minute. Discard the **Collection Tube** with the flow through.
3. Transfer the **Zymo-Spin™ IC Column** to a new **Collection Tube**. Add 200 µl of **DNA Pre-Wash Buffer** to the spin column. Centrifuge at 10,000 x g for one minute.
4. Add 500 µl of **g-DNA Wash Buffer** to the spin column. Centrifuge at 10,000 x g for one minute.
5. Transfer the spin column to a clean microcentrifuge tube. Add ≥10 µl **DNA Elution Buffer** or water to the spin column. Incubate 2-5 minutes at room temperature and then centrifuge at top speed for 30 seconds to elute the DNA. The eluted DNA can be used immediately for molecular based applications or stored ≤-20°C for future use.

Cells should be processed directly from biological fluids or from suspension in PBS, TE, or compatible buffers.

The column capacity is ~800 µl.

Typical yields from Proteinase K digested tissues are: 1-3 µg DNA per mg skeletal, heart, and brain tissues and 3-5 µg per mg liver, kidney, and lung tissues.

Troubleshooting:

1. **DNA degradation:** Check for DNase contamination. All reagents supplied with the **Quick-gDNA™ MicroPrep** are DNase-free. However, DNase contamination could result during the processing of some samples. Check pipets, pipet tips, microcentrifuge tubes, etc., and exercise the appropriate precautions during the DNA purification procedure.
2. **DNA is not performing well in subsequent experiments:** Ensure the correct volume of **Genomic Lysis Buffer** has been added to the sample. Also, make sure all centrifugation steps are completed for the indicated times and speeds (rcfs). Failure to do so may result in incomplete washing, which may cause salts to be eluted with the DNA affecting quantitation and subsequent experiments including enzymatic processes like PCR.
3. **RNA contamination:** The buffers in this kit are designed to efficiently hydrolyze and remove RNA during the DNA purification procedure.

Ordering Information

Product Description	Cat. No.	Kit Size
Quick-gDNA™ MicroPrep	D3020	50 preps.
	D3021	200 preps.
Quick-gDNA™ MiniPrep w/ uncapped columns	D3006	50 preps.
	D3007	200 preps.
Quick-gDNA™ MiniPrep w/ capped columns	D3024	50 preps.
	D3025	200 preps.
Quick-gDNA™ MidiPrep	D3100	25 preps.
	D3010	2x96 well
ZR-96 Quick-gDNA™	D3011	4x96 well
	D3012	10x96 well

For Individual Sale	Cat. No.	Amount
Genomic Lysis Buffer	D3004-1-50	50 ml
	D3004-1-100	100 ml
DNA Pre-Wash Buffer	D3004-5-15	15 ml
	D3004-5-30	30 ml
	D3004-5-50	50 ml
g-DNA Wash Buffer	D3004-2-50	50 ml
	D3004-2-100	100 ml
DNA Elution Buffer	D3004-4-10	10 ml
Zymo-Spin™ IC Columns	C1004-50	50
	C1004-250	250
Collection Tubes	C1001-50	50
	C1001-500	500
	C1001-1000	1,000

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Popular DNA Purification Products from Zymo Research

Product	Format	Kit Size	Cat No.
Fragment DNA Clean-up, Concentration & Recovery			
DNA Clean & Concentrator™-5	Spin Column Format (up to 5 µg/prep.)	50 preps. 200 preps.	D4003*, D4013 D4004*, D4014
DNA Clean & Concentrator™-25	Spin Column Format (up to 25 µg/prep.)	50 preps. 200 preps.	D4005*, D4033 D4006*, D4034
DNA Clean & Concentrator™-100	Spin Column Format (up to 100 µg/prep.)	25 preps. 50 preps.	D4029 D4030
DNA Clean & Concentrator™-500	Spin Column Format (up to 500 µg/prep.)	10 preps. 20 preps.	D4031 D4032
ZR-96 DNA Clean & Concentrator™-5	96-Well Format (up to 5 µg/well; deep well)	2x96 preps. 4x96 preps.	D4023 D4024
Genomic DNA Clean & Concentrator™	Spin Column Format (up to 10 µg/prep.)	25 preps. 100 preps.	D4010 D4011
ZR-96 DNA Clean-up Kit™	96-Well Format (up to 5 µg/well; shallow well)	2x96 preps. 4x96 preps.	D4017 D4018
ZR DNA Sequencing Clean-up Kit™	Spin Column Format (up to 5 µg/prep.)	50 preps. 200 preps.	D4050 D4051
ZR-96 DNA Sequencing Clean-up Kit™	96-Well Format (up to 5 µg/well)	2x96 preps. 4x96 preps.	D4052 D4053
OneStep™ PCR Inhibitor Removal Kit	Spin Column Format (up to 25 µg/prep.)	50 preps.	D6030
OneStep-96™ PCR Inhibitor Removal Kit	96-Well Format (up to 5 µg/well)	2x96 preps.	D6035
Zymoclean™ Gel DNA Recovery Kit	Spin Column Format (up to 5 µg/prep.)	50 preps. 200 preps.	D4001 D4002
ZR-96 Zymoclean™ Gel DNA Recovery Kit	96-Well Format (up to 5 µg/well)	2x96 preps. 4x96 preps.	D4021 D4022
Zymoclean™ Large Fragment DNA Recovery Kit	Spin Column Format (up to 10 µg/prep.)	25 preps. 100 preps.	D4045 D4046
Plasmid DNA Isolation			
Zyppy™ Plasmid Miniprep Kit	Pellet Free, Spin Column Format	50 preps. 100 preps. 400 preps. 800 preps.	D4036 D4019 D4020 D4037
Zyppy™ Plasmid Midiprep Kit	Pellet Free, Spin Column Format	25 preps. 50 preps.	D4025 D4026
Zyppy™ Plasmid Maxiprep Kit	Spin/Vacuum Column Format	10 preps. 20 preps.	D4027 D4028
ZR Plasmid Miniprep™-Classic	Spin Column Format	100 preps. 400 preps. 800 preps.	D4015 D4016 D4054
ZR BAC DNA Miniprep Kit	BAC/PAC plasmid DNA Isolation. Spin Column Format	25 preps. 100 preps.	D4048 D4049
Environmental DNA Isolation			
ZR Soil Microbe DNA MicroPrep™	Bead Bashing, Spin Column Format (up to 5 µg/prep.)	50 preps.	D6003
ZR Soil Microbe DNA MiniPrep™	Bead Bashing, Spin Column Format (up to 25 µg/prep.)	50 preps.	D6001
ZR Soil Microbe DNA MidiPrep™	Bead Bashing, Spin Column Format (up to 125 µg/prep.)	25 preps.	D6101
ZR-96 Soil Microbe DNA Kit™	Bead Bashing, 96-Well Format (up to 5 µg/well)	2x96 preps.	D6002
ZR Fungal/Bacterial DNA MicroPrep™	Bead Bashing, Spin Column Format (up to 5 µg/prep.)	50 preps.	D6007
ZR Fungal/Bacterial DNA MiniPrep™	Bead Bashing, Spin Column Format (up to 25 µg/prep.)	50 preps.	D6005
ZR Fungal/Bacterial DNA MidiPrep™	Bead Bashing, Spin Column Format (up to 125 µg/prep.)	25 preps.	D6105
ZR-96 Fungal/Bacterial DNA Kit™	Bead Bashing, 96-Well Format (up to 5 µg/well)	2x96 preps.	D6006
ZR Fecal DNA MicroPrep™	Bead Bashing, Spin Column Format (up to 5 µg/prep.)	50 preps.	D6012
ZR Fecal DNA MiniPrep™	Bead Bashing, Spin Column Format (up to 25 µg/prep.)	50 preps.	D6010
ZR Fecal DNA MidiPrep™	Bead Bashing, Spin Column Format (up to 125 µg/prep.)	25 preps.	D6110
ZR-96 Fecal DNA Kit™	Bead Bashing, 96-Well Format (up to 5 µg/well)	2x96 preps.	D6011
ZR Tissue & Insect DNA MicroPrep™	Bead Bashing, Spin Column Format (up to 5 µg/prep.)	50 preps.	D6015
ZR Tissue & Insect DNA MiniPrep™	Bead Bashing, Spin Column Format (up to 25 µg/prep.)	50 preps.	D6016
ZR Tissue & Insect DNA MidiPrep™	Bead Bashing, Spin Column Format (up to 125 µg/prep.)	25 preps.	D6115
ZR-96 Tissue & Insect DNA Kit™	Bead Bashing, 96-Well Format (up to 5 µg/well)	2x96 preps.	D6017
ZR Plant/Seed DNA MicroPrep™	Bead Bashing, Spin Column Format (up to 5 µg/prep.)	50 preps.	D6022
ZR Plant/Seed DNA MiniPrep™	Bead Bashing, Spin Column Format (up to 25 µg/prep.)	50 preps.	D6020
ZR Plant/Seed DNA MidiPrep™	Bead Bashing, Spin Column Format (up to 125 µg/prep.)	25 preps.	D6120
ZR-96 Plant/Seed DNA Kit™	Bead Bashing, 96-Well Format (up to 5 µg/well)	2x96 preps.	D6021

* Uncapped Spin Column Format

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Phone: (949) 679-1190 ▪ Toll Free: (888) 882-9682 ▪ Fax: (949) 266-9452 ▪ info@zymoresearch.com ▪ www.zymoresearch.com