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The Beauty of Science is to Make Things Simple

INSTRUCTION MANUAL

Quick-gDNA™ MidiPrep

Catalog No. **D3100**

Highlights

- For the purification of high quality DNA from up to 3 ml whole blood, plasma, serum, body fluids (e.g., semen) and buffy coat, lymphocytes, tissue, swabs and cultured cells in less than 20 minutes using innovative *Fast-Spin* column technology. Up to 125 µg/prep.
- Compatible with commonly used anticoagulants (i.e., EDTA, heparin, citrate).
- Eluted, inhibitor-free 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™ MidiPrep (Kit Size)	D3100 (25 preps.)	Storage Temperature
Genomic Lysis Buffer*	2 x 150 ml	Room Temp.
DNA Pre-Wash Buffer**	15 ml	Room Temp.
g-DNA Wash Buffer	50 ml	Room Temp.
DNA Elution Buffer	16 ml	Room Temp.
Zymo-Spin™ V-E Columns w/ Zymo-Midi Filters™	25	Room Temp.
Collection Tubes	50	Room Temp.
Instruction Manual	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., 750 µl per 150 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** – Up to 3 ml (see protocols) whole blood, plasma, or serum from humans, mice, rats, etc. Also, tissue, cells from culture, 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 125 µg total DNA is eluted into ≥ 150 µl DNA Elution Buffer or water. Human whole blood will typically yield 3-7 µg DNA per 100 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** – Centrifuge or vacuum source and manifold, 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 purification of up to 25 µg DNA/prep use the **Quick-gDNA™ MiniPrep** (D3006, D3007, D3024, D3025).

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™ MidiPrep** 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 cells from culture, and many biological liquid samples.

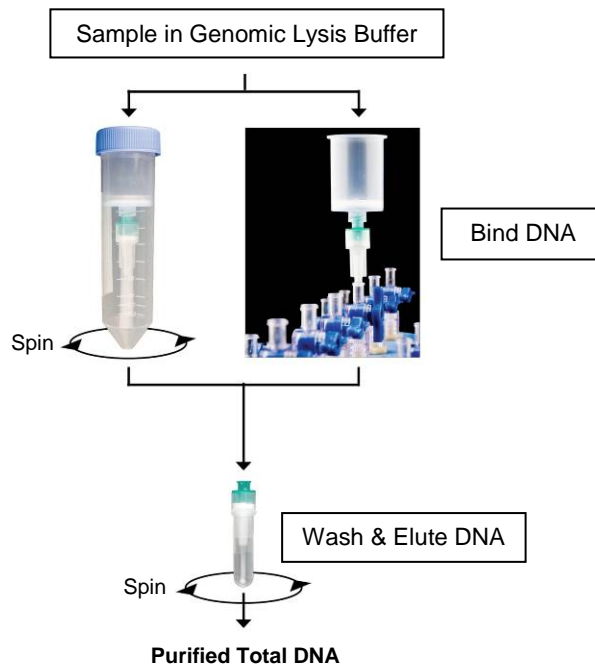
For processing, simply add the specially formulated **Genomic Lysis Buffer** to a sample, vortex, and transfer the mixture to the supplied **Zymo-Spin™ Column w/ Zymo-Midi Filter™**. There is no need for organic denaturants or Proteinase K digestion because of the unique chemistries featured in the kit. Instead, the product features *Fast-Spin* technology to yield high-quality, purified DNA in just 20 minutes (see below). PCR inhibitors are effectively removed during the purification process. DNA purified using the **Quick-gDNA™ MidiPrep** is suitable for PCR, nucleotide blotting, DNA sequencing, restriction endonuclease digestion, bisulfite conversion/methylation analysis, and other downstream applications.



Zymo-Spin™ V-E Column w/ Zymo-Midi Filter™



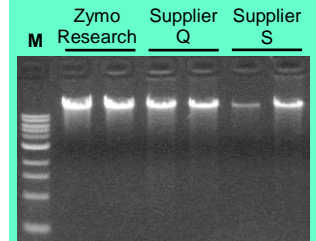
Zymo-Spin™ V-E Column



For FFPE tissues use the **ZR Genomic DNA™-Tissue MiniPrep** (D3050, D3051) or **MidiPrep** (D3110).

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)



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).

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

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., 750 µl per 150 ml.

PROTOCOLS

Whole Blood, Serum, and Plasma Samples

The following is for the purification of DNA from up to 3 ml¹ 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. Add 12 ml of **Genomic Lysis Buffer** to 3 ml¹ (4:1) of blood, serum, or plasma. Mix completely by vortexing 4-6 seconds, then let stand 5 minutes at room temperature.

Note: Add 12 ml Genomic Lysis Buffer to all samples < 3 ml.

2. Transfer the mixture to a **Zymo-Spin™ V-E Column/Zymo-Midi Filter™** assembly in a 50 ml tube². Centrifuge the tube at ≥1,000 x g (2,000 x g max.) for 5 minutes³.

Note: If using a vacuum manifold, the processing capacity is reduced to 2 ml of blood, serum, or plasma + 12 ml Genomic Lysis Buffer per prep. This filtration step may take up to twenty minutes when using vacuum.

3. Disconnect the **Zymo-Spin™ V-E Column/Zymo-Midi Filter™** assembly and transfer the **Zymo-Spin™ V-E Column** to a **Collection Tube**. Spin at 10,000 x g for 1 minute in a microcentrifuge⁴ to remove residue from the column.
4. Add 300 µl **DNA Pre-Wash Buffer** to the column and spin at 10,000 x g for 1 minute. Discard the flow through.
5. Add 400 µl of **g-DNA Wash Buffer** to the column and centrifuge at 10,000 x g for one minute. Discard flow through and repeat wash step.
6. Transfer the **Zymo-Spin™ V-E Column** to a 1.5 ml microcentrifuge tube and add 150 µl **DNA Elution Buffer** directly to the column matrix⁵ and allow column to stand for 1 minute at room temperature. Centrifuge at 10,000 x g for 1 minute to elute the DNA⁶. 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**.

For inclusion of small DNAs from serum, add 0.3 volumes isopropanol to the mixture. (For example, to a 15 ml mixture of serum and Genomic Lysis Buffer add 4.5 ml isopropanol.)

Notes:

¹Processing volumes are up to 3 ml for centrifugation and up to 2 ml for vacuum based manipulations, respectively.

²**Caution:** Make sure the connection between the column and filter is secure (finger tight) prior to centrifugation.

³Alternatively, the **Zymo-Spin™ V-E Column/Zymo-Midi Filter™** assembly can be mounted on a vacuum manifold with a vacuum source set at ≥500 mm Hg.

⁴ Leave the rotor cover off the microcentrifuge if clearance with the column tops is a problem.

⁵ 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 warmed to 60-70°C.

⁶ DNA yields can be increased by performing a second elution and pooling the eluates.

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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 100 mg of fresh or frozen tissue in 2.5 ml of **Genomic Lysis Buffer**. Increase the reagents proportionally if more than 100 mg of solid tissue is used.

1. Transfer the lysate to a **Zymo-Spin™ V-E Column/Zymo-Midi Filter™** assembly in a 50 ml tube¹. Centrifuge the tube at $\geq 1,000 \times g$ ($2,000 \times g$ max.) for 5 minutes².
2. Disconnect the **Zymo-Spin™ V-E Column/Zymo-Midi Filter™** assembly and transfer the **Zymo-Spin™ V-E Column** to a **Collection Tube**. Spin at $10,000 \times g$ for 1 minute in a microcentrifuge³ to remove residue from the column.
3. Add 300 μl **DNA Pre-Wash Buffer** to the column and spin at $10,000 \times g$ for 1 minute. Discard the flow through.
4. Add 400 μl of **g-DNA Wash Buffer** to the column and centrifuge at $10,000 \times g$ for 1 minute. Discard flow through and repeat wash step.
5. Transfer the **Zymo-Spin™ V-E Column** to a 1.5 ml microcentrifuge tube and add 150 μl **DNA Elution Buffer** directly to the column matrix⁴ and allow column to stand for 1 minute at room temperature. Centrifuge at $10,000 \times g$ for 1 minute to elute the DNA⁵. The eluted DNA can be used immediately for molecular based applications or stored $\leq -20^{\circ}\text{C}$ for future use.

For some solid tissues, proteinase K treatment may be required: use conventional methods or the **ZR Genomic DNA™-Tissue MidiPrep** (D3110).

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

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

Notes:

¹**Caution:** Make sure the connection between the column and filter is secure (finger tight) prior to centrifugation.

²Alternatively, the **Zymo-Spin™ V-E Column/Zymo-Midi Filter™** assembly can be mounted on a vacuum manifold with a vacuum source set at ≥ 500 mm Hg.

³ Leave the rotor cover off the microcentrifuge if clearance with the column tops is a problem.

⁴ 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 $60-70^{\circ}\text{C}$.

⁵ DNA yields can be increased by performing a second elution and pooling the eluates.

Generally, no more than 25×10^6 cells should be sampled, for larger samples will exceed the binding capacity of the spin column.

Cell Monolayer Samples

The following procedure is designed for up to 25×10^6 (max.) monolayer cells. 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 table (below) is provided for estimating cell numbers.

1. Trypsinize or manually scrape adherent cells from the growth surface of a culture flask or plate. Centrifuge the cell suspension at approximately $500 \times g$ for 5 minutes. Remove the supernatant and add 2.5 ml of **Genomic Lysis Buffer** directly to the pellet. Resuspend pellet by vortexing 4-6 seconds and let stand for 5 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™ V-E Column/Zymo-Midi Filter™** assembly in a 50 ml tube¹. Centrifuge the tube at $\geq 1,000 \times g$ ($2,000 \times g$ max.) for 5 minutes².
7. Disconnect the **Zymo-Spin™ V-E Column/Zymo-Midi Filter™** assembly and transfer the **Zymo-Spin™ V-E Column** to a **Collection Tube**. Spin at $10,000 \times g$ for 1 minute in a microcentrifuge³ to remove residue from the column.
8. Add 300 μ l **DNA Pre-Wash Buffer** to the column and spin at $10,000 \times g$ for 1 minute. Discard the flow through.
9. Add 400 μ l of **g-DNA Wash Buffer** to the column and centrifuge at $10,000 \times g$ for one minute. Discard flow through and repeat wash step.
3. Transfer the **Zymo-Spin™ V-E Column** to a 1.5 ml microcentrifuge tube and add 150 μ l **DNA Elution Buffer** directly to the column matrix⁴ and allow column to stand for 1 minute at room temperature. Centrifuge at $10,000 \times g$ for 1 minute to elute the DNA⁵. The eluted DNA can be used immediately for molecular based applications or stored $\leq -20^\circ\text{C}$ for future use.

Notes:

¹ **Caution:** Make sure the connection between the column and filter is **secure** (finger tight) prior to centrifugation.

² Alternatively, the **Zymo-Spin™ V-E Column/Zymo-Midi Filter™** assembly can be mounted on a vacuum manifold with a vacuum source set at ≥ 500 mm Hg.

³ Leave the rotor cover off the microcentrifuge if clearance with the column tops is a problem.

⁴ 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 $60-70^\circ\text{C}$.

⁵ DNA yields can be increased by performing a second elution and pooling the eluates.

Guidelines for Monolayer Cell DNA Isolation: The above procedure is designed for the processing of up to 25×10^6 cells. However, cell numbers (growth densities) can vary between different cell types. The table below provides a reference for the approximation of what can be recovered from different culture containers for high-density growth cells like CV1 and HeLa cells.

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

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Cell Suspensions and Proteinase K Digested Samples

The following protocol is designed for up to 1 ml of biological liquid sample including CSF, buffy coat, body fluids (semen), and cell suspensions containing less than 25×10^6 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. (For example, for 1 ml of sample, add 4 ml of **Genomic Lysis Buffer**). Mix briefly by vortexing, then let stand at room temperature for 5 minutes.

Note: For Proteinase K digested material, add 4 volumes of **Genomic Lysis Buffer** to each volume of lysate then mix briefly by vortexing. Transfer up to 5.0 ml supernatant to the Zymo-Spin™ Column in Step 2.

2. Transfer the mixture to a **Zymo-Spin™ V-E Column/Zymo-Midi Filter™** assembly in a 50 ml tube¹. Centrifuge the tube at $\geq 1,000 \times g$ ($2,000 \times g$ max.) for 5 minutes².
6. Disconnect the **Zymo-Spin™ V-E Column/Zymo-Midi Filter™** assembly and transfer the **Zymo-Spin™ V-E Column** to a **Collection Tube**. Spin at $10,000 \times g$ for 1 minute in a microcentrifuge³ to remove residue from the column.
7. Add 300 μ l **DNA Pre-Wash Buffer** to the column and spin at $10,000 \times g$ for 1 minute. Discard the flow through.
8. Add 400 μ l of **g-DNA Wash Buffer** to the column and centrifuge at $10,000 \times g$ for one minute. Discard flow through and repeat wash step.
3. Transfer the **Zymo-Spin™ V-E Column** to a 1.5 ml microcentrifuge tube and add 150 μ l **DNA Elution Buffer** directly to the column matrix⁴ and allow column to stand for 1 minute at room temperature. Centrifuge at $10,000 \times g$ for 1 minute to elute the DNA⁵. The eluted DNA can be used immediately for molecular based applications or stored $\leq -20^\circ\text{C}$ for future use.

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

Typical yields from Proteinase K digested tissues 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.

Notes:

¹**Caution:** Make sure the connection between the column and filter is secure (finger tight) prior to centrifugation.

²Alternatively, the **Zymo-Spin™ V-E Column/Zymo-Midi Filter™** assembly can be mounted on a vacuum manifold with a vacuum source set at ≥ 500 mm Hg.

³Leave the rotor cover off the microcentrifuge if clearance with the column tops is a problem.

⁴ 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 $60-70^\circ\text{C}$.

⁵ DNA yields can be increased by performing a second elution and pooling the eluates.

Troubleshooting:

1. **DNA degradation:** Check for DNase contamination. All reagents supplied with the **Quick-gDNA™ MidiPrep** 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	Catalog 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	Catalog No.	Amount
Genomic Lysis Buffer	D3004-1-150	150 ml
DNA Pre-Wash Buffer	D3004-5-15	15 ml
g-DNA Wash Buffer	D3004-2-50	50 ml
DNA Elution Buffer	D3004-4-16	16 ml
Zymo-Spin™ V-E Columns w/ Zymo-Midi Filters™	C1021-25	25 columns/filters
	C1001-50	50 tubes
Collection Tubes	C1001-500	500 tubes
	C1001-1000	1,000 tubes

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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™ ZR Soil Microbe DNA MiniPrep™ ZR Soil Microbe DNA MidiPrep™ ZR-96 Soil Microbe DNA Kit™	Bead Bashing, Spin Column Format (up to 5 µg/prep.) Bead Bashing, Spin Column Format (up to 25 µg/prep.) Bead Bashing, Spin Column Format (up to 125 µg/prep.) Bead Bashing, 96-Well Format (up to 5 µg/well)	50 preps. 50 preps. 25 preps. 2x96 preps.	D6003 D6001 D6101 D6002
ZR Fungal/Bacterial DNA MicroPrep™ ZR Fungal/Bacterial DNA MiniPrep™ ZR Fungal/Bacterial DNA MidiPrep™ ZR-96 Fungal/Bacterial DNA Kit™	Bead Bashing, Spin Column Format (up to 5 µg/prep.) Bead Bashing, Spin Column Format (up to 25 µg/prep.) Bead Bashing, Spin Column Format (up to 125 µg/prep.) Bead Bashing, 96-Well Format (up to 5 µg/well)	50 preps. 50 preps. 25 preps. 2x96 preps.	D6007 D6005 D6105 D6006
ZR Fecal DNA MicroPrep™ ZR Fecal DNA MiniPrep™ ZR Fecal DNA MidiPrep™ ZR-96 Fecal DNA Kit™	Bead Bashing, Spin Column Format (up to 5 µg/prep.) Bead Bashing, Spin Column Format (up to 25 µg/prep.) Bead Bashing, Spin Column Format (up to 125 µg/prep.) Bead Bashing, 96-Well Format (up to 5 µg/well)	50 preps. 50 preps. 25 preps. 2x96 preps.	D6012 D6010 D6110 D6011
ZR Tissue & Insect DNA MicroPrep™ ZR Tissue & Insect DNA MiniPrep™ ZR Tissue & Insect DNA MidiPrep™ ZR-96 Tissue & Insect DNA Kit™	Bead Bashing, Spin Column Format (up to 5 µg/prep.) Bead Bashing, Spin Column Format (up to 25 µg/prep.) Bead Bashing, Spin Column Format (up to 125 µg/prep.) Bead Bashing, 96-Well Format (up to 5 µg/well)	50 preps. 50 preps. 25 preps. 2x96 preps.	D6015 D6016 D6115 D6017
ZR Plant/Seed DNA MicroPrep™ ZR Plant/Seed DNA MiniPrep™ ZR Plant/Seed DNA MidiPrep™ ZR-96 Plant/Seed DNA Kit™	Bead Bashing, Spin Column Format (up to 5 µg/prep.) Bead Bashing, Spin Column Format (up to 25 µg/prep.) Bead Bashing, Spin Column Format (up to 125 µg/prep.) Bead Bashing, 96-Well Format (up to 5 µg/well)	50 preps. 50 preps. 25 preps. 2x96 preps.	D6022 D6020 D6120 D6021

* Uncapped Spin Column Format

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