



SZABO SCANDIC

Part of Europa Biosite

Produktinformation



Forschungsprodukte & Biochemikalien



Zellkultur & Verbrauchsmaterial



Diagnostik & molekulare Diagnostik



Laborgeräte & Service

Weitere Information auf den folgenden Seiten!
See the following pages for more information!



Lieferung & Zahlungsart

siehe unsere [Liefer- und Versandbedingungen](#)

Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

SZABO-SCANDIC HandelsgmbH

Quellenstraße 110, A-1100 Wien

T. +43(0)1 489 3961-0

F. +43(0)1 489 3961-7

mail@szabo-scandic.com

www.szabo-scandic.com

[linkedin.com/company/szaboscandic](https://www.linkedin.com/company/szaboscandic) 



ZYMO RESEARCH

The Beauty of Science is to Make Things Simple

INSTRUCTION MANUAL

Quick-DNA™ Urine Kit

Catalog No. **D3061**

Highlights

- Purify cellular and/or cell-free DNA easily and reliably from up to 40 ml of urine.
- Uniquely formulated urine conditioning reagent allows stabilization of DNA in urine for up to 1 month at ambient temperature.
- Clean-Spin™ column technology ensures DNA is ready for all sensitive downstream applications including qPCR, DNA sequencing, arrays, and DNA methylation analysis.

Contents

Product Contents.....	1
Product Specifications.....	1
Product Description.....	2
Buffer Preparation.....	3
Protocol.....	3-4
Protocol at a Glance.....	5
Appendix A: Isolating Cellular or Cell-free DNA Only.....	6
Appendix B: Tough-to-Lyse Samples.....	7
Quick Protocol.....	8-9
Troubleshooting.....	10-11
Ordering Information.....	12
Other Products.....	13-16

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-DNA™ Urine Kit (Kit Size)	D3061 (50 Preps.)	Storage Temperature
Urine Conditioning Buffer	140 ml	Room Temp.
Clearing Beads	1 ml	Room Temp.
Urine Pellet Digestion Buffer	20 ml	Room Temp.
Proteinase K*	2 x 20 mg	-20°C (after mixing)
Genomic Lysis Buffer**	50 ml	Room Temp.
Urine DNA Prep Buffer	10 ml	Room Temp.
Urine DNA Wash Buffer*** (concentrate)	12 ml	Room Temp.
DNA Elution Buffer	4 ml	Room Temp.
Zymo-Spin™ IC-S Columns	50	Room Temp.
Collection Tubes	100	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.

* Add 1,040 µl of **Proteinase K Storage Buffer** to reconstitute the lyophilized 20 mg **Proteinase K** at 20 mg/ml prior to use and store at -20°C after use.

** Add beta-mercaptoethanol to a final dilution of 0.5% (v/v) i.e., 250 µl per 50 ml.

*** Add 48 ml 100% ethanol (52 ml 95% ethanol) to the 12 ml **Urine DNA Wash Buffer** (concentrate).

Product Specifications

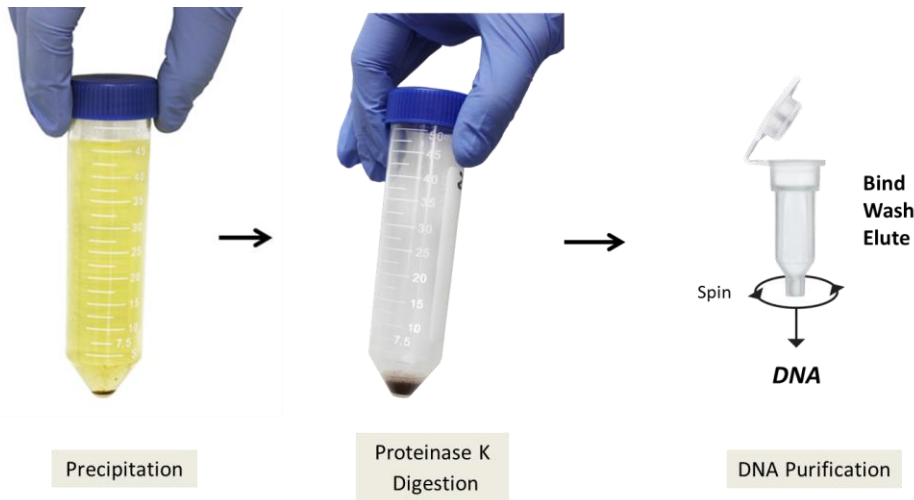
- **Sample Source** – Urine.
- **Sample Volume** – Up to 40 ml of urine per standard preparation. The sample volume can be scaled up or down if required.
- **DNA Quality**– High quality DNA can be used for downstream application, such as PCR, bisulfite treatment/methylation detection, array, etc.
- **DNA Yield** – The DNA binding capacity of the column is 5 µg. Note that DNA yield may vary depending on the urine itself. Female urine typically yields more DNA than male urine. Urine DNA from healthy female individual ranges on average from 6 - 1000 ng/ml. DNA from healthy male individual ranges on average from 2 - 20 ng/ml.
- **DNA Size** – Capable of recovering DNA fragments from 100 bp to 23 kb.
- **Equipment** – A centrifuge, a microcentrifuge, and a heat block / water bath.

Note - ™ Trademarks of Zymo Research Corporation. This product is for research use only and should 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 safety guidelines and rules enacted by your research institution or facility.

ZYMO RESEARCH CORP.

Product Description

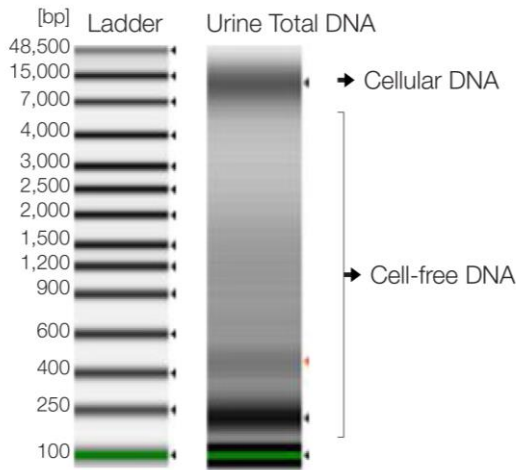
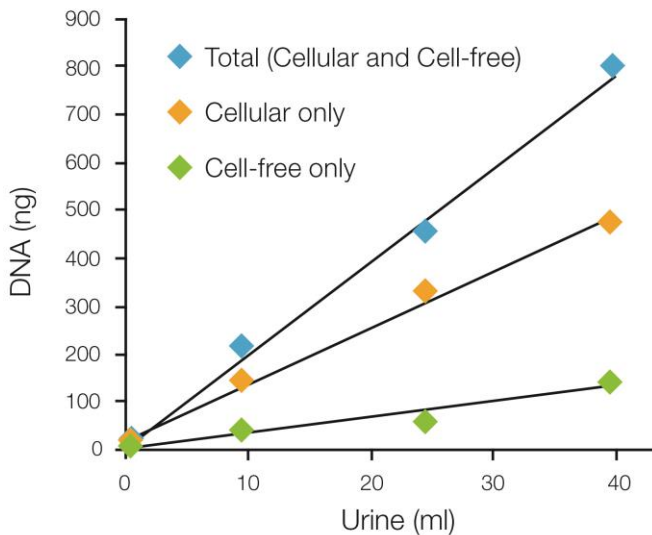
The **Quick-DNA™ Urine Kit** is an innovative product designed for easy, reliable, and rapid isolation of cellular and/or cell-free DNA from up to 40 ml of urine. The product features a uniquely formulated urine DNA stabilization reagent that also functions as a precipitation reagent. After collection, total (cellular and cell-free) or cell-free urine can be stored at ambient temperature for up to one month by adding the **Urine Conditioning Buffer**. When ready to extract the urine DNA, just add the **Clearing Beads**, vortex, and centrifuge to collect the precipitate. Following precipitation, chemical lysis and enzymatic digestion are used to extract DNA from the precipitate. The DNA is subsequently purified and concentrated using **Zymo-Spin™ IC-S Columns**. Urine DNA isolated with the **Quick-DNA™ Urine Kit** is ideal for qPCR, array, methylation analysis¹, and other downstream applications.



Notes:

¹ Zymo Research offers the following for rapid and precise DNA methylation detection.

- 1.) **EZ DNA Methylation-Lightning™ Kit** (D5030, D5031, D5032, D5033, D5046, D5047)
- 2.) **EZ DNA Methylation-Direct™ Kit** (D5020, D5021, D5022, D5023, D5044, D5045)
- 3.) **EZ DNA Methylation-Gold™ Kit** (D5005, D5006, D5007, D5008, D5042, D5043)
- 4.) **5-mC DNA ELISA Kit** (D5325, D5326)



DNA yields increase linearly with increasing volumes of urine from healthy subjects extracted using the Quick-DNA™ Urine Kit. DNA was isolated from 1 ml, 10 ml, 25 ml, and 40 ml urine. DNA concentration was quantified using the **Femto™ Human DNA Quantification Kit** (Zymo Research, E2005).

Both cellular and cell-free DNA can be effectively purified from urine using the Quick-DNA™ Urine Kit. 5 ml of urine from a healthy female donor was processed and DNA was eluted in 20 µl final volume. 1 µl of the sample was analyzed using a 2200 TapeStation.

ZYMO RESEARCH CORP.

Buffer Preparation

- ✓ 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.
- ✓ Add 1,040 µl of **Proteinase K Storage Buffer** to reconstitute the lyophilized 20 mg **Proteinase K** at 20 mg/ml prior to use and store at -20°C after use.
- ✓ Add 48 ml 100% ethanol (52 ml 95% ethanol) to the 12 ml **Urine DNA Wash Buffer** (concentrate).

Notes:

¹ The volume of urine sample can be scaled up or down if required.

² Warm up the urine at 37 °C for 5 - 10 minutes if the urine is visually cloudy. This may be caused by salt precipitation. If the urine does not turn clear, it may be caused by bacterial contamination.

³ Mix **Clearing Beads** well before use by vortexing.

Protocol

The following protocol is designed for the isolation of DNA from up to 40 ml urine sample¹. All steps should be performed at room temperature (20 - 30 °C).

The protocol consists of three sections:

- (A)** Total (cellular and cell-free) DNA Precipitation,
- (B)** Protein Digestion, and
- (C)** DNA Purification.

(A) Total (cellular and cell-free) DNA Precipitation

*(If only cellular or cell-free DNA is desired, refer to **Appendix A** on page 6)*

1. Transfer up to 40 ml urine² into a microcentrifuge tube or a conical tube.

Note: Use a microcentrifuge tube if processing ≤ 1 ml urine.
Use a 15 ml conical tube if processing > 1 ml to 14 ml urine.
Use a 50 ml conical tube if processing > 14 ml to 40 ml urine.

2. Add 70 µl **Urine Conditioning Buffer** for every 1 ml of urine (e.g. Add 350 µl **Urine Conditioning Buffer** to 5 ml urine). Mix the urine mixture well by vortexing.

After adding and mixing urine with **Urine Conditioning Buffer**, urine can be stored up to 1 month at ambient temperature. At time of processing, mix the urine mixture well by vortexing and continue to the next step.

3. Add 10 µl **Clearing Beads**³ if processing ≤ 14 ml urine and add 20 µl **Clearing Beads**³ if processing >14 to 40 ml urine. Mix the urine mixture well by vortexing.
4. Centrifuge at 3,000 x g for 15 minutes.
*(If isolating DNA from gram (+) bacteria, fungal, yeast, or other tough-to-lyse samples in urine, continue to **Appendix B** on page 7)*

(B) Protein Digestion

- Without disturbing the pellet, slowly decant or pipette out the urine supernatant leaving behind 100 - 400 μ l of pellet

Recommended: leave at least 200 μ l of pellet if processing 15 - 40 ml urine.

Note: you could adjust the volume to 100 - 400 μ l by adding DNase/RNase-Free Water to the pellet.

- Add an equal volume of **Urine Pellet Digestion Buffer** to the pellet (e.g. Add 100 μ l Urine Pellet Digestion Buffer to 100 μ l pellet). Resuspend the pellet well by pipetting or vortexing.
- Add 5% (v/v) of **Proteinase K** to the resuspended pellet (e.g. Add 10 μ l Proteinase K to 200 μ l mixture), mix well by vortexing, and incubate the pellet mixture at 55 °C for 30 minutes.

(C) DNA Purification

All centrifugation steps should be performed $\geq 16,000 \times g$, unless stated otherwise.

- Add 1 volume of **Genomic Lysis Buffer** to the digestion mixture (e.g. Add 210 μ l Genomic Lysis Buffer to 210 μ l digestion mixture) and mix well by vortexing.
 - Transfer the sample into a **Zymo-Spin™ IC-S Column** in a **Collection Tube**. Centrifuge for 1 minute. Discard flow-through. Repeat with the remaining mixture¹.
- Note:** The column may need to be loaded multiple times. Transfer 800 μ l maximum for each load of mixture.
- Transfer the **Zymo-Spin™ IC-S Column** to a new **Collection Tube**.
 - Add 200 μ l of **Urine DNA Prep Buffer** to the spin column. Centrifuge for 1 minute. Discard flow-through.
 - Add 700 μ l **Urine DNA Wash Buffer** to the column and centrifuge for 1 minute. Discard flow-through. Repeat this step with 200 μ l **Urine DNA Wash Buffer**.
 - Transfer the spin column to a DNase/RNase-free microcentrifuge tube. Add $\geq 10 \mu$ l² **DNA Elution Buffer**³ or water directly on the column matrix and let stand for 3 - 5 minutes at room temperature. Centrifuge at full speed for 1 minute. The eluted DNA can be used immediately or stored at ≤ -20 °C for future use.

Note: If the concentration of the DNA is > 100 ng/ μ l in the first elution. Most of the DNA will be recovered on the first elution however the loading eluate from the first elution back onto the column, incubating for 3 - 5 minutes at room temperature, and centrifuging again can increase total yield

Notes:

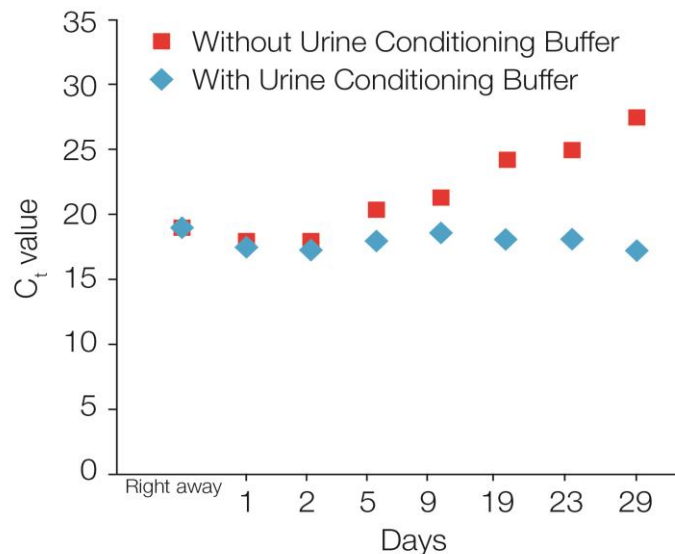
¹ **Clearing Beads** will be retained in the column.

² DNA yield may be increased by warming the **DNA Elution Buffer** to 65 °C before eluting and by performing a re-elution. Do not worry about the Clearing Beads that are retained at the ridge of the column. The DNA binding to the Clearing Beads are negligible. It is important to load the DNA Elution Buffer directly on the column matrix.

³ **DNA Elution Buffer:** 10 mM Tris-HCl, pH 8.5, 0.1 mM EDTA. If water is used, make sure the pH is > 6.0 .

Protocol at a Glance

DNA Precipitation	Urine volume	≤ 1 ml	> 1 - 14 ml	> 14 - 40 ml
	Tube to be used	microcentrifuge	15 ml conical	50 ml conical
	Urine Conditioning Buffer to be added	70 µl / ml urine		
	Clearing Beads to be added	10 µl	10 µl	20 µl
Centrifuge at 3000 x g for 15 minutes				
Protein Digestion <i>(minimum pellet volume is listed)</i>	Pellet volume to be left	100 µl	100 µl	200 µl
	Urine Pellet Digestion Buffer to be added	100 µl	100 µl	200 µl
	Reconstituted Proteinase K to be added	10 µl	10 µl	20 µl
Incubate at 55 °C for 30 minutes				
DNA Purification	BIND: Genomic Lysis Buffer to be added	210 µl	210 µl	420 µl
	Load to Column IC-S			
	WASH 1: Urine DNA Prep Buffer to wash	200 µl		
	WASH 2: Urine DNA Wash Buffer to wash	700 µl, 200 µl		
	ELUTE: DNA Elution Buffer to be added	≥ 10 µl		



DNA in urine can be preserved for up to 1 month at ambient temperature. Total (cellular and cell-free) urine DNA was incubated with and without **Urine Conditioning Buffer** for 29 days. For each sample, 5 ml urine from a healthy female donor was processed using the **Quick-DNA™ Urine Kit** and the C_t values were determined by qPCR using the **Femto™ Human DNA Quantification Kit** (Zymo Research, E2005).

ZYMO RESEARCH CORP.

Appendix A

Isolating Cellular or Cell-free DNA Only

1. Cellular DNA only

- i. Centrifuge up to 40 ml urine in a microcentrifuge tube or a conical tube at 3,000 x g for 15 minutes.
- ii. Proceed to **Protocol** section **(B) Protein Digestion** on **page 4**,

OR

Proceed to **Appendix B** on **page 7** if isolating DNA from gram (+) bacteria, fungal, yeast, or other tough-to-lyse samples in urine.

2. Cell-free DNA only

- i. Centrifuge up to 40 ml urine in a microcentrifuge tube or a conical tube at 3,000 x g for 15 minutes.
- ii. Without disturbing the pellet, carefully transfer urine supernatant to a new microcentrifuge tube or a conical tube.

Note: Use a microcentrifuge tube if processing ≤ 1 ml cell-free urine.
Use a 15 ml conical tube if processing > 1 ml to 14 ml cell-free urine.
Use a 50 ml conical tube if processing > 14 ml to 40 ml cell-free urine.

- iii. Add 70 μ l **Urine Conditioning Buffer** for every 1 ml of urine supernatant (e.g. Add 350 μ l **Urine Conditioning Buffer** to 5 ml urine) and mix well by vortexing.

After adding and mixing urine with **Urine Conditioning Buffer**, urine can be stored up to 1 month at ambient temperature. At time of processing, mix the urine mixture well by vortexing and continue to the next step.

- iv. Add 10 μ l **Clearing Beads**¹ if processing ≤ 14 ml urine and add 20 μ l **Clearing Beads**¹ if processing >14 to 40 ml urine. Mix the urine mixture well by vortexing.
- v. Centrifuge at 3000 x g for 15 minutes.
- vi. Proceed to **Protocol** section **(B) Protein Digestion** on **page 4**,

Notes:

¹ Mix **Clearing Beads** well before use by vortexing.

Notes:

¹The following additional components are sold separately:

ZR BashingBead Lysis Tube (0.1 & 0.5 mm) -
50 pack: S6012-50

BashingBead Buffer -
40 ml: D6001-3-40
150 ml: D6001-3-150

Zymo-Spin III-F Filters -
50 pack: C1057-50

Collection Tubes -
50 pack: C1001-50
500 pack: C1001-500

² DNA yield may be increased by warming the **DNA Elution Buffer** to 65 °C before eluting and by performing a re-elution.

Appendix B

Tough-to-Lyse Samples

Order the **ZR BashingBead Lysis Tubes, BashingBead Buffer, Zymo-Spin III-F Filters,** and additional **Collection Tubes** before proceeding¹. The following protocol is an adapted protocol for lysing tough-to-lyse organisms in urine, such as gram (+) bacteria, yeast, and other fungi.

Sample Lysis

1. Without disturbing the pellet, slowly decant or pipette out the urine supernatant leaving behind 100 - 400 µl of pellet. Transfer the pellet to a **ZR BashingBead™ Lysis Tube (0.1 & 0.5 mm)** and then add **BashingBead Buffer** to a final volume of 800 µl (e.g. *Add 700 µl of BashingBead Buffer to a 100 µl pellet*).

2. Secure in a bead beater fitted with a 2 ml tube holder assembly (e.g., Disruptor Genie™) and process at maximum speed for 5 minutes.

Note: Processing times may be as little as 40 seconds when using high-speed cell disrupters (e.g. the portable Xpedition™ Sample Processor, FastPrep®-24 or similar). See manufacturer's literature for operating information.

3. Centrifuge the ZR BashingBead Lysis Tube at 10,000 x g for 1 minute.

4. Transfer 400 µl of the supernatant to a **Zymo-Spin™ III-F Filter** in a **Collection Tube**. Centrifuge at 10,000 x g for 1 minute. **Keep** flow-through and discard the filter.

DNA Purification

All centrifugation steps should be performed $\geq 16,000$ x g, unless stated otherwise.

5. Add 1,200 µl **Genomic Lysis Buffer** to the flow-through. Mix well by pipetting up and down several times. Transfer 800 µl of the mixture to the **Zymo-Spin™ IC-S Column** in a clean **Collection Tube**. Centrifuge for 1 minute. Discard the flow-through. Repeat with the remaining mixture.

6. Transfer the **Zymo-Spin™ IC-S Column** to a new **Collection Tube**.

Note: The column may need to be loaded multiple times. Transfer 800 µl maximum for each load of mixture.

7. Add 200 µl **Urine DNA Prep Buffer** to the column. Centrifuge for 1 minute. Discard the flow-through.

8. Add 700 µl **Urine DNA Wash Buffer** to the column. Centrifuge for 1 minute. Discard the flow-through. Repeat this step with 200 µl **Urine DNA Wash Buffer**.

9. Transfer the column to a DNase/RNase-free microcentrifuge tube. Add ≥ 10 µl **DNA Elution Buffer**² directly to the column matrix and let stand for 3 - 5 minutes at room temperature. Centrifuge at full speed for 1 minute. The eluted DNA can be used immediately or stored at ≤ -20 °C for future use.

Note: If the concentration of the DNA is > 100 ng/µl in the first elution. Most of the DNA will be recovered on the first elution however the loading eluate from the first elution back onto the column, incubating for 3 - 5 minutes at room temperature, and centrifuging again can increase total yield

ZYMO RESEARCH CORP.

Quick-DNA™ Urine Kit

Catalog Nos. D3061



ZYMO RESEARCH

The Beauty of Science is to Make Things Simple

Quick Protocol

The quick protocol is designed for the isolation of total DNA from urine (both cellular and cell-free DNA).

See the Instruction Manual page 6 for selective isolation of cellular or cell-free DNA from urine.

See the Instruction Manual page 7 if processing tough-to-lyse samples such as yeast and bacteria from urine.

Buffer Preparation:

- ✓ 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.
 - ✓ Add 1,040 µl of **Proteinase K Storage Buffer** to reconstitute the lyophilized 20 mg **Proteinase K** at 20 mg/ml prior to use and store at -20°C after use.
 - ✓ Add 48 ml 100% ethanol (52 ml 95% ethanol) to the 12 ml **Urine DNA Wash Buffer** (concentrate).
-

The following procedure should be performed at room temperature (15 - 30°C).

1. Transfer up to 40 ml urine into a microcentrifuge tube or a conical tube.

Note: Use a microcentrifuge tube if processing ≤ 1 ml urine.
Use a 15 ml conical tube if processing > 1 ml to 14 ml urine.
Use a 50 ml conical tube if processing > 14 ml to 40 ml urine.

2. Add 70 µl **Urine Conditioning Buffer** for every 1 ml of urine (e.g. Add 350 µl *Urine Conditioning Buffer* to 5 ml urine). Mix the urine mixture well by vortexing.

After adding and mixing urine with **Urine Conditioning Buffer**, urine can be stored up to 1 month at ambient temperature. At time of processing, mix the urine mixture well by vortexing and continue to step 3.

3. Add 10 µl **Clearing Beads** if processing ≤ 14 ml urine and add 20 µl **Clearing Beads** if processing >14 to 40 ml urine. Mix the urine mixture well by vortexing.

4. Centrifuge at 3,000 x g for 15 minutes.
-

ZYMO RESEARCH CORP.

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

Quick-DNA™ Urine Kit

Catalog Nos. D3061



ZYMO RESEARCH

The Beauty of Science is to Make Things Simple

-
- Without disturbing the pellet, slowly decant or pipette out the urine supernatant leaving behind 100 - 400 µl of pellet.

Recommended: leave at least 200 µl of pellet if processing 15 - 40 ml urine.

Note: you could adjust the volume to 100 - 400 µl by adding DNase/Rnase-Free Water to the pellet.

- Add an equal volume of **Urine Pellet Digestion Buffer** to the pellet (e.g. Add 100 µl Urine Pellet Digestion Buffer to 100 µl pellet). Resuspend the pellet well by pipetting or vortexing.
- Add 5% (v/v) of **Proteinase K** to the resuspended pellet (e.g. Add 10 µl Proteinase K to 200 µl mixture), mix well by vortexing, and incubate the pellet mixture at 55 °C for 30 minutes.
- Add 1 volume of **Genomic Lysis Buffer** to the digestion mixture (e.g. Add 210 µl Genomic Lysis Buffer to 210 µl digestion mixture) and mix well by vortexing.
- Transfer the sample into a **Zymo-Spin™ IC-S Column** in a **Collection Tube**. Centrifuge for 1 minute. Discard flow-through. Repeat with the remaining mixture.

Note: The column may need to be loaded multiple times. Transfer 800 µl maximum for each load of mixture.

- Transfer the **Zymo-Spin™ IC-S Column** to a new **Collection Tube**.
- Add 200 µl of **Urine DNA Prep Buffer** to the spin column. Centrifuge for 1 minute. Discard flow-through.
- Add 700 µl **Urine DNA Wash Buffer** to the column and centrifuge for 1 minute. Discard flow-through. Repeat this step with 200 µl **Urine DNA Wash Buffer**.
- Transfer the spin column to a DNase/RNase-free microcentrifuge tube. Add ≥ 10 µl **DNA Elution Buffer** or water directly on the column matrix and let stand for 3 - 5 minutes at room temperature. Centrifuge at full speed for 1 minute.

Ver 1.0

For the full Instruction Manual, visit
<http://www.zymoresearch.com/m/D3061>

ZYMO RESEARCH CORP.

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

Troubleshooting

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

Problem	Possible Causes and Suggested Solution
----------------	---

Volume Input

- For low DNA containing urine, you can increase the input volume to as high as 40 ml.

Increasing DNA Yields

- The total yield may be improved by eluting the DNA with DNA Elution Buffer pre-heated to 60 - 70°C.
- Loading the eluate from the first elution back onto the column, incubate for 3 - 5 minutes at room temperature, and centrifuge at full speed can maximize recovery.

Low DNA Yield

Incomplete Lysis/Digestion

- Ensure Proteinase K digestions are performed at 55 °C as indicated. It is possible to extend digestion times to 1 hour if samples are high in protein.
- Vortex samples longer after the addition of Genomic Lysis Buffer to ensure that the lysate is homogenous.

Variety in Urine Samples

- Urine DNA from healthy female individual ranges on average from 6 - 1000 ng/ml.
- DNA from healthy male individual ranges on average from 2 - 20 ng/ml.

Urine Storage

Short Term Storage

- Urine samples could be stored up to 1 month at ambient temperature by adding Urine Conditioning Buffer. Add 70 µl Urine Conditioning Buffer for every 1 ml of urine and mix thoroughly by vortexing. At time of processing, mix the urine well by vortexing and continue by adding the Clearing Beads.

Long Term Storage

- If you want to store the urine samples for more than 1 month, we recommend pelleting after adding the Urine Conditioning Buffer. Add 70 µl Urine Conditioning Buffer for every 1 ml of urine and mix thoroughly by vortexing. Centrifuge at 3,000 x g for 15 minutes. Discard urine supernatant and store the pellet at -20 °C or -80 °C.

Keep in mind that after adding Urine Conditioning Buffer, the cellular and cell-free fractions of the urine DNA cannot be separated. If required, separate the cellular and cell-free urine DNA fractions of the urine by following **Appendix A page 6**.

ZYMO RESEARCH CORP.

Sample Condition

Cloudy urine

- This may be caused by salt precipitation. Warm up the urine at 37 °C for 5 - 10 minutes if the urine is visually cloudy. If the urine does not turn clear, it may be caused by bacterial contamination.

Clearing Beads in the flow through or eluate

- Urine with high amount of cellular content may cause clearing beads to pass through the spin column.
- If the Clearing Beads are present in the eluate, centrifuge the eluate at full speed for 1 minute and transfer the supernatant to a DNase/RNase-free microcentrifuge tube for storage or analysis.

The presence of oil-like solution after the 15 minutes 3,000 x g centrifugation

- We recommend removing the oil-like solution before adding the Urine Pellet Digestion Buffer and proceed with Protein Digestion Section on page 4.
-

Ordering Information

Product Description	Catalog No.	Kit Size
Quick-DNA™ Urine Kit	D3061	50 Preps.

For Individual Sale	Catalog No.	Amount
Proteinase K	D3001-2-5	5 mg
	D3001-2-20	20 mg
Genomic Lysis Buffer	D3004-1-50	50 ml
	D3004-1-100	100 ml
DNA Elution Buffer	D3004-4-1	1 ml
	D3004-4-4	4 ml
	D3004-4-10	10 ml
	D3004-4-50	50 ml
Urine Conditioning Buffer	D3061-1-140	140 ml
Clearing Beads	D3061-2-1	1 ml
Urine Pellet Digestion Buffer	D3061-3-20	20 ml
Urine DNA Prep Buffer	D3061-4-10	10 ml
Urine DNA Wash Buffer (concentrate)	D3061-5-12	12 ml
Zymo-Spin™ IC-S Columns	C1015-50	50 columns
Collection Tubes	C1001-50	50 tubes
	C1001-500	500 tubes
	C1001-1000	1,000 tubes

Supplementary Components (for tough-to-lyse samples)	Catalog No.	Amount
ZR BashingBead Lysis Tube (0.1 & 0.5 mm)	S6012-50	50 tubes
BashingBead Buffer	D6001-3-40	40 ml
	D6001-3-150	150 ml
Zymo-Spin III-F Filters	C1057-57	50 pack
Collection Tubes	C1001-50	50 tubes
	C1001-500	500 tubes

ZYMO RESEARCH CORP.

Related Products from Zymo Research

Product	Description	Kit Size (Preps)	Catalog No. (Format)
DNA Methylation Detection			
EZ DNA Methylation-Lightning™ Kit	Complete bisulfite conversion in about an hour using a unique liquid format conversion reagent that requires no preparation. Fast-Spin technology ensures ultra-pure, converted DNA for subsequent DNA methylation analysis. Magnetic bead format for adaptation to automated liquid handling platforms.	50 Rxns. 200 Rxns. 2x96 Rxns. 2x96 Rxns. 4x96 Rxns. 8x96 Rxns.	D5030 (spin column) D5031 (spin column) D5032 (shallow-well plate) D5033 (deep-well plate) D5046 (magnetic bead) D5047 (magnetic bead)
EZ DNA Methylation-Direct™ Kit	Features simple and reliable DNA bisulfite conversion directly from blood, tissue (FFPE/LCM), and cells without the prerequisite for DNA purification in as little as 4-6 hrs. The increased sensitivity of this kit makes it possible to amplify bisulfite converted DNA from as few as 10 cells or 50 pg DNA. Magnetic bead format for adaptation to automated liquid handling platforms.	50 Rxns. 200 Rxns. 2x96 Rxns. 2x96 Rxns. 4x96 Rxns. 8x96 Rxns.	D5020 (spin column) D5021 (spin column) D5022 (shallow-well plate) D5023 (deep-well plate) D5044 (magnetic bead) D5045 (magnetic bead)
EZ DNA Methylation-Gold™ Kit	For the fast (3 hr.) conversion of unmethylated cytosines in DNA to uracil via heat/chemical-denaturation of DNA and a specially designed CT Conversion Reagent. Fast-Spin technology ensures ultra-pure, converted DNA for subsequent DNA methylation analysis. Magnetic bead format for adaptation to automated liquid handling platforms.	50 Rxns. 200 Rxns. 2x96 Rxns. 2x96 Rxns. 4x96 Rxns. 8x96 Rxns.	D5005 (spin column) D5006 (spin column) D5007 (shallow-well plate) D5008 (deep-well plate) D5042 (magnetic bead) D5043 (magnetic bead)
5-mC DNA ELISA Kit	A convenient and powerful tool that allows the researcher to accurately quantitate 5-mC in any DNA sample in less than 3 hours. The kit features a unique anti-5-Methylcytosine monoclonal antibody that is both sensitive and specific for 5-mC.	1x96 Rxns. 2x96 Rxns.	D5325 D5326
Methylated-DNA IP Kit	IP with a highly specific anti-5-methylcytosine monoclonal antibody. Designed for the enrichment of 5-methylcytosine-containing DNA from any pool of fragmented genomic DNA for use in genome-wide methylation analysis.	10 Rxns.	D5101
Other...			
Universal Methylated Human DNA Standards	Human (male) genomic DNA having all CpG sites methylated. To be used for the evaluation of bisulfite-mediated conversion of DNA. Supplied with a control primer set.	1 set	D5011
ZymoTaq™ DNA Polymerase	ZymoTaq™ "hot start" DNA Polymerase is specifically designed for the amplification of "difficult" DNA templates including: bisulfite-treated DNA for methylation detection. The product generates specific amplicons with little or no by-product formation. Available either as a single buffer premix or as a polymerase system with components provided separately.	50 Rxns. 200 Rxns. 50 Rxns. 200 Rxns.	E2001 (system) E2002 (system) E2003 (premix) E2004 (premix)
Femto™ Human DNA Quantification Kit	Highly specific detection and quantification of human DNA with increased accuracy, precision, and sensitivity.	100 Rxns.	E2005
Femto™ Bacterial DNA Quantification Kit	Highly specific detection and quantification of bacterial DNA with increased accuracy, precision, and sensitivity.	100 Rxns.	E2006
Femto™ Fungal DNA Quantification Kit	Highly specific detection and quantification of fungal DNA with increased accuracy, precision, and sensitivity.	100 Rxns.	E2007

Please visit our website to see our complete line-up of products.

ZYMO RESEARCH CORP.

DNA PURIFICATION



What is Clean-Spin™ Technology?

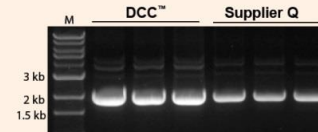
The spin columns from Zymo Research have been designed to ensure complete elution with no binding/wash buffer carryover. The result is ultra-pure inhibitor-free DNA and RNA.

Purify DNA from PCR & other sources

DNA Clean & Concentrator™ (DCC™)

- ✓ Recovery of ultra-pure DNA that is free of salts and contaminants.
- ✓ Small ($\geq 6 \mu\text{l}$) elution volume.
- ✓ DNA is ideal for ligation, PCR, Next-Gen sequencing, etc.

Product	Size (Cat. No.)
DNA Clean & Concentrator™-5	50 Preps. (D4013) 200 Preps. (D4014)
ZR-96 DNA Clean & Concentrator™-5	2 x 96 Preps. (D4023) 4 x 96 Preps. (D4024)
Genomic DNA Clean & Concentrator™	25 Preps. (D4010) 100 Preps. (D4011)



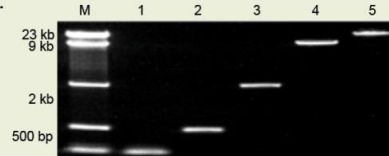
High efficiency DNA recovery with the DCC™-5 compared to Supplier Q.

Boost DNA recoveries from agarose gels to >80%

Zymoclean™ Gel DNA Recovery

- ✓ Rapid (15 min.) recovery of ultra-pure DNA from agarose gels in $\geq 6 \mu\text{l}$.
- ✓ Ultra-pure DNA ideal for DNA ligation, sequencing, etc.
- ✓ Format also available for large DNA >20 kb.

Product	Size (Cat. No.)
Zymoclean™ Gel DNA Recovery Kit	50 Preps. (D4001) 200 Preps. (D4002)
Zymoclean™ Large Fragment DNA Recovery Kit	25 Preps. (D4045) 100 Preps. (D4046)



DNA fragments recovered from an agarose gel using the Zymoclean™ Gel DNA Recovery Kit. Lanes: M: DNA Ladder; 1-5: individual ladder DNA fragments.

Recover transfection-quality plasmid DNA directly from culture

Zyppy™ Plasmid Prep Kits

- ✓ The fastest, simplest method available for purifying high quality plasmid DNA from *E. coli*.
- ✓ Pellet-Free™ procedure omits conventional cell-pelleting and resuspension steps.
- ✓ Transfection quality plasmid DNA directly from culture in under 15 minutes.

Easy, Pellet-free Procedure: Add Lysis Buffer **Directly** to Bacterial Culture



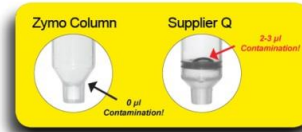
Product	Size (Cat. No.)
Zyppy™ Plasmid Miniprep Kit	50 Preps. (D4036) 100 Preps. (D4019) 400 Preps. (D4020) 800 Preps. (D4037)

ZYMO RESEARCH CORP.



RNA PURIFICATION

Get RNA *directly* from TRIzol® without phase separation

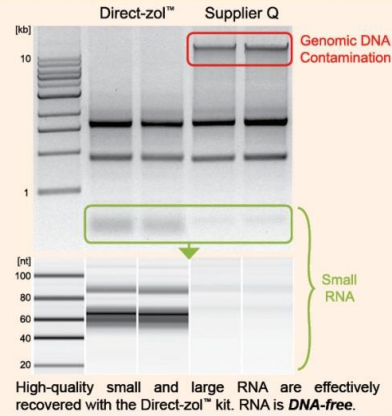


Direct-zol™ RNA

- ✓ For purification of high-quality small and large RNA *directly* from TRIzol®, TRI Reagent®, or similar.
- ✓ Bypasses phase separation and precipitation procedures allowing for unbiased recovery of miRNA

Product	Size (Cat. No.)
Direct-zol™ RNA MiniPrep	50 Preps. (R2050)
	50 Preps. (R2051)*
	200 Preps. (R2052)
	200 Preps. (R2053)*
96-well and MagBead formats also available!	

DNase I included in all kits.
* Supplied with TRI-Reagent®

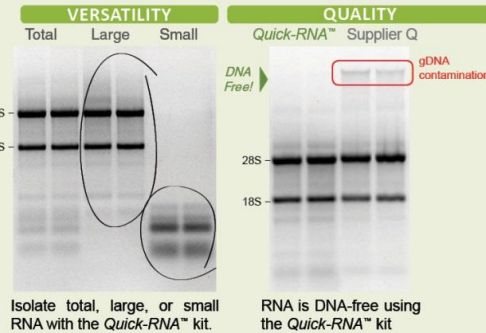


Isolate DNA-free RNA from 1 to 10⁷ cells in minutes

Quick-RNA™

- ✓ Isolation of total, large, or small RNA – *You decide!*
- ✓ Ultra clean, high-quality RNA from a single cell to 10⁷ cells.
- ✓ DNA-free RNA ideal for any downstream application – *DNase I included.*

Product	Size (Cat. No.)
Quick-RNA™ MicroPrep	50 Preps. (R1050)
	200 Preps. (R1051)
Quick-RNA™ MiniPrep	50 Preps. (R1054)
	200 Preps. (R1055)
ZR-96 Quick-RNA™	2 x 96 Preps. (R1052)
	4 x 96 Preps. (R1053)



Purify RNA from enzymatic and labeling reactions in 5 minutes

RNA Clean & Concentrator™

- ✓ Recover ultra-pure RNA in small (≥6 µl) elution volumes.
- ✓ Compatible with TRIzol®, phenol, chloroform, and RNase inhibitors (RNAlater®).
- ✓ RNA is ideal for RT-PCR, q-PCR, hybridization, arrays, RNA interference, etc.

Product	Size (Cat. No.)
RNA Clean & Concentrator™-5	50 Preps. (R1015)
	200 Preps. (R1016)
RNA Clean & Concentrator™-25	50 Preps. (R1017)
	100 Preps. (R1018)
ZR-96 RNA Clean & Concentrator™	2x96 well plates (R1080)
DNA-Free RNA Kit™	50 Preps. (R1013)
	200 Preps. (R1014)

The following are trademarks of other companies: pGEM®, Promega Corp.; TRIzol® and TRI Reagent®, Molecular Research Center, Inc.; DH5® and DH10B™, Life Technologies, Inc.

ZYMO RESEARCH CORP.

OTHER INNOVATIVE PRODUCTS FROM ZYMO RESEARCH...

Competent cells for transformations *without* heat shock!

Mix & Go! Pre-made Competent *E. Coli*

- ✓ High efficiency: 10⁸-10⁹ transformants/μg plasmid DNA
- ✓ Just Mix & Go! Simply add DNA then spread. Transformation in as little as 20 seconds!

Product	Size (Cat. No.)
Zymo 5α (Same as DH5α)	10 x 100 μl aliquots (T3007) 96 x 50 μl aliquots (T3009) 96 x 50 μl aliquots PCR-plate (T3010)
Zymo 10B (Same as DH10B)	10 x 100 μl aliquots (T3019) 96 x 50 μl aliquots (T3020)
JM109	10 x 100 μl aliquots (T3003) 96 x 50 μl aliquots (T3005)
HB101	10 x 100 μl aliquots (T3011) 96 x 50 μl aliquots (T3013)
C600	10 x 100 μl aliquots (T3015)
TG1	10 x 100 μl aliquots (T3017)

- ✓ No heat shock
- ✓ No incubations
- ✓ No outgrowth
- ✓ No wait!!!

Mix & Go
Competent cells
E. coli + DNA

Mix & Go!
for 20 second*
transformations!



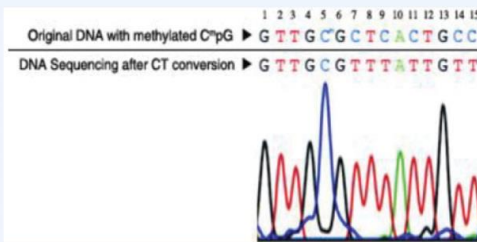
Others
E. coli + DNA
Ice 45 min.
42°C 2 min. (Heatshock)
Place on Ice
Add SOC
Spin to Concentrate Cells
Remove Supernatant

* For Ampicillin selection only.

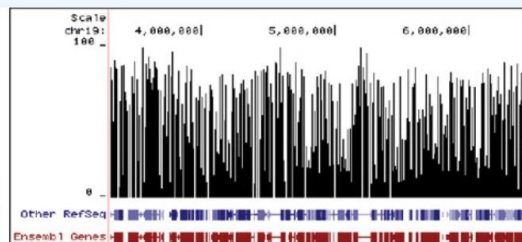
The fastest method for complete bisulfite conversion of DNA

EZ DNA Methylation-Lightning™ Kits

- ✓ The next generation of bisulfite conversion technology by the most cited provider in the industry
- ✓ Guarantees high conversion efficiencies of cytosine (>99.5%)
- ✓ Maintains the highest template integrity following bisulfite conversion
- ✓ Recovered DNA is ideal for PCR, MSP, array, bisulfite, and next-generation sequencing.



DNA Sequencing Results Following Bisulfite Treatment



Methylation Plot From Reduced Representation Bisulfite Sequencing (RRBS)

Product	Size (Cat. No.)
EZ DNA Methylation-Lightning™ Kit	50 rxns. (D5030) 200 rxns. (D5031)
EZ-96 DNA Methylation-Lightning™ Kit	Shallow-Well 2 x 96 rxns. (D5032) Deep-Well 2 x 96 rxns. (D5033)
EZ-96 DNA Methylation-Lightning™ MagPrep	4 x 96 rxns. (D5046) 8 x 96 rxns. (D5047)

ZYMO RESEARCH CORP.