



For research use only. Not for use in diagnostic procedures.

Prepito truXTRAC DNA FFPE Kit

DNA purification from 10 μ m sections or equivalent of FFPE tissue

Kit Components

Magnetic Beads	Wash Buffer 6
Lysis Buffer	Elution Buffer
Binding Buffer	Deep Well Plates
Wash Buffer 3	0.75 mL Reaction Tubes
Wash Buffer 4	Disposable Tips
Wash Buffer 5	

Completion time: approx. 80 minutes

Storage Conditions and Safety Information

Expiry dates are stated on the box and on each single component of the kit. Do not use any component of the kit beyond the expiry date. All kit components can be stored at room temperature.

Any further questions?

chemagen Technology technical support: +49 (0) 2401 805-501 | support.chemagen@perkinelmer.com

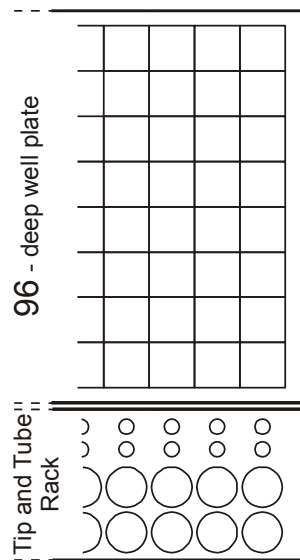




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Positioning Procedure

See **“Protocol Steps”** for detailed information



120 μ L lysate

Pos. 4 second row for Disposable Tips; **! not used in this protocol !**

Pos. 3 Disposable Tips

Pos. 2 0.75 mL reaction tubes with 150 μ L **Magnetic Beads**

Pos. 1 0.75 mL reaction tubes for 50 - 100 μ L **Elution Buffer**

Before You Start

- Check all kit components for integrity. In case of damages contact your supplier.
- Connect the tubes according to their numbering to the respective counterparts at the **chemagic 8-Pack**. Remove the lids from the individual buffer bottles in the **chemagic 8-Pack** and pierce the septum with the spike placed at the end of each tube. Place the **chemagic 8-Pack** upside down on the reagent holder and use the manual priming function for a complete filling of the dispensing system.

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Protocol Steps (chemagic Prepito serial numbers 1 – 99)

1. For FFPE tissue material pre-treatment please refer to the Covaris manual: “truXTRAC FFPE DNA microTUBE Kit for chemagen Technology”.
2. Switch the **chemagic Prepito** on and wait until the self test is finished.
3. Press [**change protocol**].
4. Select the **Prepito truXTRAC DNA FFPE Kit** protocol by pressing [**FFPE**].
5. Enter the access code [**4444**] for authorization and confirm by pressing [**enter**].
6. Confirm the selection of the correct protocol by pressing [**enter**].
7. Read the protocol information in the appearing information screen. Confirm by pressing [**continue**].
8. Select the sample positions and confirm by pressing [**continue**].
9. Enter the kit barcode with the barcode scanner and confirm by pressing [**ok**].
10. For the registration of the samples and storage tubes press [**yes**] and follow the instructions on the touch screen panel to enter the according barcodes.
11. Prepare the **chemagic Tip & Tube Rack** with the required materials. Place one 0.75 mL reaction tube filled with 50 – 100 μ L **Elution Buffer** (position 1), one 0.75 mL reaction tube filled with 150 μ L of **Magnetic Beads** (position 2) and one **Disposable Tip** (position 3) for each sample into positions according to the sample positions.

! *Shake the vessel with the Magnetic Beads vigorously until all Magnetic Beads are completely resuspended. An incomplete resuspension of the Magnetic Beads can cause a decreased yield of extracted nucleic acids.*

12. Transfer the lysate to each well of the Deep Well Plate (DWP, riplate SW) defined as sample well (Pos. “120 μ L tissue lysate”, see section above “Positioning Procedure”).
13. An information screen indicates the previously selected sample positions. Ensure that the sample positions in the DWP correspond to the selected positions. Place the DWP on its default position on the tracking system and press [**continue**].
14. Place the **chemagic Tip & Tube Rack** on its default position on the tracking system. Check for accurate fit of the DWP and **chemagic Tip & Tube Rack** and lock both by closing the safety latch.

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15. Close the front door and immediately start the automated isolation process by pressing [start] immediately.

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Protocol Steps (**chemagic Prepito** serial numbers **100 and later**)

1. For FFPE tissue material pre-treatment please refer to the Covaris manual: “truXTRAC FFPE DNA microTUBE Kit for chemagen Technology”.
2. Switch on the **chemagic Prepito** and wait for the self test to finish.
3. Press [**Change Protocol**].
4. Press [**Tissue**] in the Select Protocol Group window.
5. Select the **Prepito truXTRAC DNA FFPE Kit** protocol by pressing [**FFPE**] and confirm by pressing [**OK**].
6. Confirm the protocol selection in the Select Protocol Group window by pressing [**OK**].
7. Enter the 4 digit access code [**4444**] for authorization and confirm by pressing [**Enter**].
8. Press [**Start Process**].
9. Read the protocol information in the appearing information screen and confirm by pressing [**Continue**].
10. Select the sample positions and confirm by pressing [**OK**].
11. Enter the kit barcode with the barcode scanner and confirm by pressing [**OK**].
12. For the registration of the samples and the storage tubes press [**Yes**] and follow the instructions on the touch screen panel to enter the according barcodes.
13. Prepare the **chemagic Tip & Tube Rack** with the required materials. Place one 0.75 mL reaction tube filled with 50 – 100 µL **Elution Buffer** (position 1), one 0.75 mL reaction tube filled with 150 µL of **Magnetic Beads** (position 2) and one **Disposable Tip** (position 3) for each sample into positions according to the sample positions.

! *Shake the vessel with the Magnetic Beads vigorously until all Magnetic Beads are completely resuspended. An incomplete resuspension of the Magnetic Beads can cause a decreased yield of extracted nucleic acids.*

14. Transfer the lysate to each well of the Deep Well Plate (DWP, riplate SW) defined as sample well (Pos. “120 µL tissue lysate”, see section above “Positioning Procedure”).

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15. An information screen indicates the previously selected sample positions. Ensure that the sample positions in the DWP correspond to the selected positions. Place the DWP on its default position on the tracking system and press [**Continue**].
16. Place the **chemagic Tip & Tube Rack** on its default position on the tracking system. Check for accurate fit of the DWP and **chemagic Tip & Tube Rack** and lock both by closing the safety latch.
17. Close the front door and start the automated isolation process by pressing [**Start**] immediately.

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General Remarks

The **Elution Buffer** included in this kit is 10 mM Tris-HCl pH 8.0.

The **Magnetic Bead** suspension should be mixed vigorously before dispensing, otherwise the suspension is not homogenous and the DNA yield could be low.

UV Measurements

In some cases you may find traces of **Magnetic Beads** left in the eluate. Such particles will not interfere with PCR and most downstream applications but may increase the background in UV measurements. In such a case, prior to UV analysis, we recommend an additional separation step using a manual separator (e.g. **chemagic Stand 2x12**, art. No. CMG-300) in order to separate any traces of particles.

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