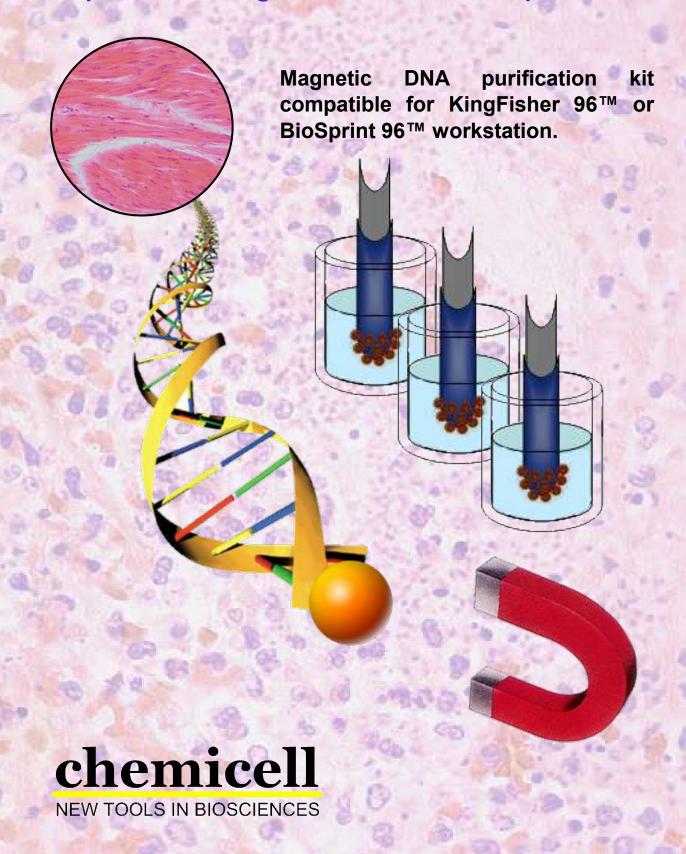
geneMAG-DNA 96 / Tissue

compatible for KingFisher™ 96 and BioSprint™ 96

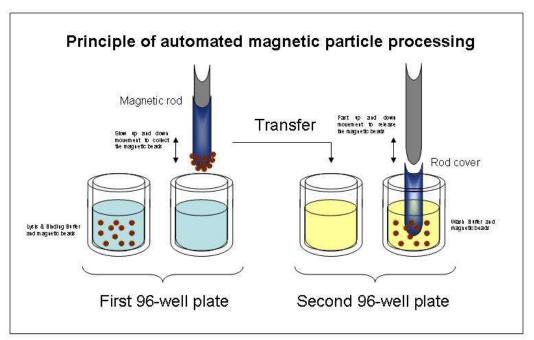


Technology

The **geneMAG-DNA 96** / **Tissue** kit is a novel, simple and highly efficient tool for the isolation of genomic DNA from tissue with magnetic silica beads using the KingFisher™96 or BioSprint™ 96 workstations.

The lysis of cells and binding of DNA is carried out under non-chaotropic conditions with the Lysis & Binding Buffer. The wash steps with Wash Buffer I and II guarantee a clean DNA which is suitable for PCR reactions or other biochemical applications.

geneMAG-DNA 96 / *Tissue* is highly suitable for a variety of automatization platforms since it requires no centrifugation or vacuum filtration procedures.



The magnetic bead processing of KingFisher™ 96 / BioSprint™ 96 workstation

The KingFisher™ 96 workstation is a trademark of Thermo Fisher Scientific.

The BioSprint™ 96 workstation is a trademark of Qiagen.

Products

Kits	Contents	Number of isolations	Price Euro/US\$
geneMAG-DNA 96 / Tissue (Cat. No.: KF3801-96)	100 ml Lysis & Binding Buffer200 ml Wash Buffer I10 ml SiMAG/KF-DNA Beads	1 x 96 preps per 100 mg tissue	220 / 286
geneMAG-DNA 480 / Tissue (Cat. No.: KF3801- 480)	 500 ml Lysis & Binding Buffer 1000 ml Wash Buffer I 50 ml SiMAG/KF-DNA Beads 	5 x 96 preps per 100 mg tissue	900 / 1170

Reagents and Equipment to be Supplied by the User

• Wash Buffer II: 70% ethanol or 70% 2-propanol

• Elution Buffer: ddH2O

• KingFisher™ 96 / BioSprint™ 96 workstation

• Deep well 96-well plates (2,2 ml) squared well

KingFisher™ 96 plate (0,3 ml)

Magnet Head for deep well 96-well plates

Storage

The kit compounds are stable at room temperature. If there are salt precipitates in the Lysis/Binding Buffer or Wash Buffer I dissolve these precipitates by warming in a water bath.

Safety Note

Wash Buffer I contain chaotropic salts, which are irritant. Take appropriate laboratory safety measures and wear gloves when handling. **Avoid skin and eye contact**

Protocol for KingFisher™ 96 or BioSprint 96™

This protocol describes the isolation of genomic-DNA from 100 mg tissueper each well of 96-well plate with the geneMAG-DNA 96 / *Tissue* kit using KingFisher™ 96 or BioSprint™ 96 workstation.

Homogenization of sample:

Homogenize approx. **100 mg tissue** to reduce to small pieces with a scalpel or better with a commercial homogenizer.

Preparation of the deep well 96-well plates (2,2 ml)

First 96-well plate:

- 1. Add 100 mg sample in each well
- 2. Add 1000 µl Lysis & Binding Buffer and 100 µl SiMAG/KF-DNA.

Second 96-well plate:

1. Add 1000 µl Wash Buffer I in each well

Third 96-well plate:

1. Add 1000 µl Wash Buffer I in each well

Fourth 96-well plate:

1. Add 1000 µl Wash Buffer II (70% 2-propanol) in each well

Fifth 96-well plate:

1. Add 1000 µl Wash Buffer II (70% 2-propanol) in each well

Sixth 96-well plate: Use 96-well plate with max. volume of 0,3 ml

1. Add 100 µl Elution Buffer (dH2O) in each well

Seventh 96-well plate: Parking station!!!!

Protocol for KingFisher™ 96 or BioSprint 96™

Settings of the processing times for Lysis- & Binding-, Wash- and Elution steps:

Start the KingFisher Software and set the following parameters:

- 1. Lysis & Binding process (first deep-well plate):6 minutes with low stirring
- Wash process with Wash Buffer I (second deep-well plate):1 minutes with medium stirring
- 3. Wash process with Wash Buffer I (third deep-well plate):1 minutes with medium stirring
- 4. Wash process with Wash Buffer II (fourth deep-well plate):1 minutes with medium stirring
- 5. Wash process with Wash Buffer II (fifth deep-well plate):1 minutes with medium stirring
- **6.** Elution process with Elution Buffer (dH2O):

 Heat time: 10 minutes with high stirring

Temperature: 80°C

Contact

chemicell GmbH

Eresburgstrasse 22-23 12103 Berlin Germany



info@chemicell.com

Tel.: +49-30-2141481 Fax.: +49-30-21913737 e-mail: info@chemicell.com Internet: www.chemicell.com