

KI-60005 FISH Reagent Kit

For freshly prepared samples

RUO - Research Use Only Not for use in diagnostic procedures

Instructions for use for KI-60005 FISH Reagent Kit in combination with Kreatech™ fluorescent labeled DNA probes

Fluorescent in situ hybridization (FISH) identifies or labels target genomic sequences so that their location can be studied. DNA sequences from appropriate chromosome specific probes are first labeled with reporter molecules. The labeled FISH probe is then hybridized to the metaphase chromosomes or interphase nuclei on a slide. After washing, the specimen is screened for the reporter molecules by fluorescence microscopy.

Kreatech™ probes do not contain Cot-1 DNA. Hybridization efficiency is therefore increased and background, due to unspecific binding, is highly reduced.

For use on metaphase and interphase cells from peripheral blood cultures or direct preparations prepared by standard cytogenetic methods, see: The ACT cytogenetics laboratory manual. 3rd ed. New York: Raven Press; 1996.

This pretreatment kit is specifically developed to obtain optimal results on FRESHLY prepared cytological samples.

Note: background or difficult samples

It is advised to use the $\underline{\text{KI-60006 FISH Digestion kit}}$ in case older samples, slides with cytoplasmic background or difficult samples are used or expected.

For paraffin embedded tissues it is recommended to use pretreatment kits KI-60004 or KI-60007.

For more info consult our website: www.LeicaBiosystems.com.

Pretreatment:

- 1. Pretreat freshly prepared sample slides in 2 x SSC / 0.5% Igepal (LK-105B) at 37 °C for 15 minutes (min).
- 2. Dehydrate in 70%, 85% and 100% ethanol for 1 minute each. Airdry.

Proceed with Probe preparation.

Probe preparation:

Kreatech™ FISH probes are supplied Ready to Use (RtU). SE, ST, and WC Kreatech™ FISH probes are provided 5 x concentrated and must be diluted as follows: 2 µl 5 x conc. Probe in 8 µl FISH Hybridization

Buffer (FHB or WCB, supplied with probes). To combine several 5 x conc. probes, replace FISH Hybridization Buffer (FHB or WCB) with $2 \mu l$ for each probe added.

Co-denaturation:

Apply 10 μ l of probe or probe-mix per 22 x 22 mm field. Cover with glass cover slip and seal with Fixogum or rubber cement. Denature sample and probe on a Thermo*brite* (TS-01/02) or on a hotplate at 75 \pm 1 °C for 5-10 min. Continue with hybridization.

Hybridization:

Incubate overnight at 37 ± 1 °C in a Thermo*brite* (TS-01/02) in a humidified chamber.

Post-Hybridization Wash:

- 1. Pre-warm Wash Buffer I (0.4 x SSC / 0.3% Igepal) (LK-102) to 72 °C
- 2. Remove rubber cement.
- Place up to 14 slides in 200 ml of Wash Buffer II (2 x SSC / 0.1% Igepal) (LK-103), incubate for 2 min at RT. Slide off coverslips.
 Re-use only once for a total of 28 slides.
- Place up to 14 slides in 200 ml of pre-warmed Wash Buffer I (0.4 x SSC / 0.3% Igepal) (LK-102), incubate for 2 min at 72 °C (± 1 °C) without agitation. Re-use only once for a total of 28 slides.
- Place up to 14 slides in 200 ml of fresh Wash Buffer II (2 x SSC / 0.1% Igepal) (LK-103), incubate for 1 min at RT without agitation. Re-use only once for a total of 28 slides.
- Dehydrate in fresh 70%, 85% and 100% ethanol, incubate for 1 min each at RT. Air dry at RT and proceed to counterstaining.

Counterstaining:

Apply 15 μ I DAPI counterstain (DAPI/Antifade 0.1 μ g/ml) (LK-095B) and apply glass cover slip. Proceed with microscopy.

Procedural recommendations:

Temperature and buffer concentration (stringency) of hybridization and washing are important, as lower stringency can result in non-specific binding of the probe to other sequences, and higher stringency can result in a lack of signal. Incomplete denaturation of target DNA can result in lack of signal.

LeicaBiosystems.com



Material provided:

| LK-105B | 2 x SSC / 0.5% Igepal |
|---------|-------------------------------|
| LK-102B | Wash buffer I |
| LK-103B | Wash buffer II |
| LK-095B | DAPI counterstain (0.1 μg/ml) |
| LK-097B | Counterstain diluent |

Recommendations for Fluorescence Microscopy:

For optimal visualization use a well maintained and regularly calibrated microscope equipped with a 100 W mercury lamp and a 63x or 100x fluorescent objective. Triple band-pass filters (DAPI/FITC/Cy3 or DAPI/FITC/TRITC) are used to view multiple colors, single band-pass filters are used for individual color visualization.

Suitable excitation and emission range for KreatechTM fluorophores:

Platinum $Bright^{\text{TM}}$ 415 Ex 429 ±20 nm Em 470 ±30 nm Platinum $Bright^{\text{TM}}$ 495 Ex 495 ±20 nm Em 525 ±30 nm Platinum $Bright^{\text{TM}}$ 500 Ex 546 ±12 nm Em 580 ±30 nm

Material required, but not supplied:

- Ethanol 100%, 85% and 70%
- Fixogum (LK-071A) or rubber cement
- Hot plate with accurate temperature control up to 75 °C or Thermo Brite™ (TS01/TS02)
- Incubator with accurate temperature control at 37 °C
- Waterbath with accurate temperature range of 37-75 °C
- Plastic or glass coplin jars
- Variable micropipettes (1 μl 200 μl)
- Fluorescence microscope equipped with suitable filters (see recommendations for Fluorescence Microscopy).

Warnings and Precautions:

- 1. For in vitro use only. For professional use only. In case of emergencies check SDS sheets for safety information.
- 2. FISH probes and hybridization buffers contain formamide which is a teratogen; do not inhale or allow skin contact. Wear gloves and a lab coat when handling FISH probes and DAPI counterstain.
- All materials should be disposed of according to your institution's guidelines for hospital waste disposal.

OSHA HazCom Standard (2012)

| Code # | Description | Signal Word | Pictogram | Hazard Statements | | |
|---------|--|----------------|------------|----------------------|--|--|
| LK-095B | DAPI/Antifade 0.1 µg/ml (DAPI Counterstain) | Warning | (1) | H315, H319, H335 | | |
| LK-097B | Antifade (Counterstain diluent) | Warning | (1) | H315, H319, H335 | | |
| LK-102B | 0.4 x SSC / 0.3% Igepal (Wash Buffer I) | N.A. | N.A. | N.A. | | |
| LK-103B | 2 x SSC / 0.1% Igepal (Wash Buffer II) | N.A. | N.A. | N.A. | | |
| LK-104B | 2 x SSC | N.A. | N.A. | N.A. | | |

(*Please note that LK-102 and LK-103 products containing the same number but different letters are identical except for the volume. A= 100 ml, B=250 ml)

Technical support

Technical support is available at

<u>www.LeicaBiosystems.com</u> or toll free at 800-248-0123 or via e-mail: <u>kreatech-support@leicabiosystems.com</u>.

Customer service

Kreatech probes may be ordered through Leica Customer Service toll free at 800-248-0123

or order via e-mail: purchase.orders@leica-microsystems.com.



Kreatech Biotechnology B.V. Vlierweg 20 1032 LG Amsterdam The Netherlands

Patents & Tradenames:

Kreatech™ is a trade name of Kreatech Biotechnology B.V., ULS™ is a trademark of Kreatech Biotechnology B.V. Repeat-Free™ is a trademark of Veridex LLC. Thermo-brite™ is a trademark of Statspin.

Nothing disclosed herein is to be construed as recommendation to use any of Kreatech products in violation of any patents. Furthermore, Kreatech does not warrant that the product, either alone, or in combination with other products, is immune from charges of patent infringement. Kreatech will not be held responsible for patent infringements or other violations that may occur from the use of any of its products.