

*ZytoLight*  
SPEC D13S319/13q34/CEN 12  
Triple Color Probe

REF Z-2160-200  $\nabla_{\Sigma}$  20 (0.2 ml)

REF Z-2160-50  $\nabla_{\Sigma}$  5 (0.05 ml)

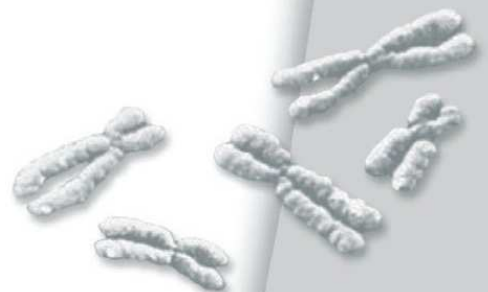
For the detection of the human D13S319 locus, human 13q34 specific sequences, and alpha-satellites of chromosome 12 by fluorescence *in situ* hybridization (FISH)



IVD

In vitro diagnostic medical device

according to EU directive 98/79/EC



Fluorescence-labeled polynucleotide probe for the detection of the human D13S319 locus, human 13q34 specific sequences, and alpha-satellites of chromosome 12 centromeres, ready to use

## Product Description

- Content:** ZytoLight SPEC D13S319/13q34/CEN 12 Triple Color Probe (PL116) in hybridization buffer. The probe contains orange-labeled polynucleotides (ZyOrange: excitation at 547 nm and emission at 572 nm, similar to rhodamine), which target the D13S319 region, blue-labeled polynucleotides (ZyBlue: excitation at 418 nm and emission at 467 nm, similar to DEAC), which target chromosome 13q34 specific sequences, and green-labeled polynucleotides (ZyGreen: excitation at 503 nm and emission at 528 nm, similar to FITC), which target alpha-satellite-sequences of the centromere of chromosome 12.
- Product:** Z-2160-200: 0.2 ml (20 reactions of 10 µl each)  
Z-2160-50: 0.05 ml (5 reactions of 10 µl each)
- Specificity:** The ZytoLight SPEC D13S319/13q34/CEN 12 Triple Color Probe (PL116) is designed to be used for the detection of the human D13S319 region as well as human 13q34 specific sequences and chromosome 12 alpha-satellites in formalin-fixed, paraffin-embedded tissue or cells by fluorescence *in situ* hybridization (FISH).
- Storage/Stability:** The ZytoLight SPEC D13S319/13q34/CEN 12 Triple Color Probe (PL116) must be stored at 2...8°C protected from light and is stable through the expiry date printed on the label.
- Use:** This product is designed for *in vitro* diagnostic use (according to EU directive 98/79/EC). Interpretation of results must be made within the con-

text of the patient's clinical history with respect to further clinical and pathologic data of the patient by a qualified pathologist!

**Safety Precautions:**

Read the operating instructions prior to use!

Do not use the reagents after the expiry date has been reached!

This product contains substances (in low concentrations and volumes) that are harmful to health. Avoid any direct contact with the reagents. Take appropriate protective measures (use disposable gloves, protective glasses, and lab garments)!

If reagents come into contact with skin, rinse skin immediately with copious quantities of water!

A material safety data sheet is available on request for the professional user!

## Principle of the Method

The presence of certain nucleic acid sequences in cells or tissue can be detected by *in situ* hybridization using labeled DNA probes. The hybridization results in duplex formation of sequences present in the test object with the labeled DNA probe.

Duplex formation (with sequences of the D13S319 and the 13q34 regions, as well as chromosome 12 alpha-satellites in the test material) is directly detected by using the tags of fluorescence-labeled polynucleotides.

# Instructions

Pretreatment (dewaxing, proteolysis, post-fixation) should be carried out according to the needs of the user.

Denaturation and hybridization of probe:

- 1.** Pipette 10 µl ZytoLight SPEC D13S319/13q34/CEN 12 Triple Color Probe (PL116) each onto individual samples

*A gentle warming of the probe, as well as using a pipette tip which has been cut off to increase the size of the opening, can make the pipetting process easier. Avoid long exposure of the probe to light.*

- 2.** Avoiding trapped bubbles, cover the samples with a coverslip (22 mm x 22 mm). Seal the coverslip, e.g. with a layer of hot glue from an adhesive pistol or with rubber cement

- 3.** Denature the slides at 75°C ( $\pm 2^\circ\text{C}$ ) for 10 min, e.g. on a hot plate

*Depending upon the age of the sample and variations in the fixation stage, it may be necessary to optimize the denaturing temperature (73°C-77°C).*

- 4.** Transfer the slide to a humidity chamber and hybridize overnight at 37°C (e.g. in a hybridization oven)

*It is essential that the tissue/cell samples do not dry out during the hybridization step.*

Further processing, such as washing and counter-staining, can be completed according to the user's needs. For a particularly user-friendly performance, we recommend the use of a *ZytoLight* FISH system by ZytoVision. These systems were also used for the confirmation of appropriateness of the ZytoLight SPEC D13S319/13q34/CEN 12 Triple Color Probe (PL116).

## Results

With the use of appropriate filter sets, the hybridization signals of the labeled D13S319 region appear orange; the hybridization signals of the labeled 13q34 specific sequences appear blue, and the hybridization signals of the labeled alpha-satellite-sequences of the centromere of chromosome 12 appear green. In interphases of normal cells or cells without aberrations of chromosome 12 and 13, two D13S319 signals, two 13q34 signals, and two chromosome 12 signals appear. In cells with a deletion of the D13S319 locus, a reduced number of orange signals is visible in interphases. In cells with a trisomy of chromosome 12 three green signals will be observed.

The polynucleotides contained in the ZytoLight SPEC D13S319/13q34/CEN 12 Triple Color Probe (PL116) which recognize the chromosomal region 13q34 function in themselves as an internal control that a successful hybridization has occurred, as well as proving the integrity of the cellular DNA.

In order to judge the specificity of the signals, every hybridization should be accompanied by controls. We recommend using at least one control sample in which the chromosome 12, D13S319, and 13q34 number is known.

Care should be taken not to evaluate overlapping cells, in order to avoid false results, e.g. an amplification of genes. Due to decondensed chromatin, single FISH signals can appear as small signal clusters. Thus, two or three signals of the same size, separated by a distance equal to or less than the diameter of one signal, should be counted as one signal.

**Our experts are available to answer your questions.**

# Literature

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