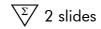


EGFR Control Slide Set

REF E-4009-2



For the detection of EGFR gene amplification by chromogenic *in situ* hybridization (CISH)

For research use only



Product Description

Content: <u>EGFR Control Slide</u> (**SC3**), consisting of two

different cell lines, affected by different levels of EGFR gene amplification, and one tissue (canine

myocardial muscle).

Product: E-4009-2 (2 slides)

Specificity: The EGFR Control Slide (**SC3**) is designed to be

used as a positive control for the detection of EGFR gene amplification by chromogenic *in situ*

hybridization (CISH).

Storage/Stability: The <u>EGFR Control Slide</u> (SC3) must be stored at

2...8°C and is stable through the expiry date

printed on the label.

Use: This product is designed for research purposes

only and not for use in diagnostic applications.

Safety Precautions: Read the operating instructions prior to use!

Although the fixation process renders the slides non-infectious, the user is advised to observe the same safety precautions as for handling/

disposing potentially infectious agents!

Avoid any direct contact with the slides. Take appropriate protective measures (use disposable gloves, protective glasses, and lab garments).

Slide Description

The <u>EGFR Control Slide</u> (**SC3**) consists of two different cell lines, affected by different levels of EGFR gene amplification, and one tissue (canine myocardial muscle).

The <u>EGFR Control Slide</u> (**SC3**) should be used for monitoring the correct performance of a CISH experiment for the detection of EGFR gene amplification in formalin-fixed, paraffin-embedded tissues. For use as an on-slide control simply mount a tissue sample of interest next to the cell lines of the <u>EGFR Control Slide</u> (**SC3**) before baking the slide.

Source: Mammalian cell lines containing human EGFR gene

sequences, and canine myocardial muscle tissue

Fixative: 10% neutrally buffered formalin (24 h, pH 7.0)

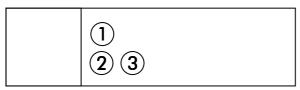
Embedding: Paraffin, red-colored

Thickness: $4 \mu m$

Mounting: Positively charged slides

Pretreatment: Pre-baked for 15 min at 58°C

Design:



1: no EGFR signals

2: no EGFR amplification, 1-2 gene copies per nucleus

3: high level EGFR amplification, large cluster per nucleus

Instructions

- 1. Remove label of the <u>EGFR Control Slide</u> (**SC3**) and label slide with a pencil
- 2. For on-slide control mount tissue sample of interest
- **3.** Bake slide at 60°C for a minimum of 2 h up to 16 h

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

For a particularly user-friendly performance, we recommend the use of a Zyto Dot CISH system by Zyto Vision.

Results

When using appropriate EGFR probes, a positive staining can be obtained on the different spots of the <u>EGFR Control Slide</u> (**SC3**), as for example two dot-shaped EGFR signals in a nucleus of a cell line not affected by an EGFR gene amplification. If the cells on the <u>EGFR Control Slide</u> (**SC3**) fail to show a positive staining, results of test samples should be considered as being invalid. In case of cross-reaction using an EGFR probe please alter washing steps and/or increase stringency of hybridization/washing.

Our experts are available to answer your questions.

As of: January 1, 2010 (4.5)		
Trademarks:		

 ${\it ZytoVision}^{\it @} \ {\it and} \ {\it Zyto} {\it Dot} \ {\it are} \ {\it trademarks} \ {\it of} \ {\it ZytoVision} \ {\it GmbH}.$

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