



## Mouse anti-ERG

Cat. No.: MSK090-05 (0.5 ml Concentrate); MSG090 (6 ml Ready-to-use)

### Instructions for use

#### Intended use

This antibody is designed for the specific localisation of ERG in formalin-fixed, paraffin-embedded tissue sections. Anti-ERG antibody is intended for in vitro diagnostic use.

#### Specifications

|                            |                            |
|----------------------------|----------------------------|
| <b>Specificity:</b>        | ERG                        |
| <b>Clone:</b>              | 9FY                        |
| <b>Isotype:</b>            | Mouse IgG1                 |
| <b>Species reactivity:</b> | Human +, others not tested |

#### Summary and Description

ERG protein expression in formalin-fixed paraffin-embedded (FFPE) tissues may be an extremely useful tool for the diagnosis of prostatic adenocarcinoma.

In terms of epithelial tumours, ERG expression is highly restricted to prostate carcinomas.

A recently developed mouse monoclonal anti-ERG antibody shows an extremely high specificity for detecting prostatic adenocarcinomas. Miettinen *et al.* found that ERG expression in high-grade PIN is correlated up to 90% with the existence of prostate carcinoma – even if the carcinoma itself cannot be found in the investigated biopsy. According to the findings of Yaskiv *et al.* ERG shows slightly lower sensitivity but much higher specificity than P504S.

Further utility for the mouse monoclonal anti-ERG antibody has been demonstrated recently in detecting endothelial malignancies, such as Kaposi sarcoma.

#### Reagent provided

Mouse monoclonal antibody in buffer with carrier protein and preservative for stabilisation in the following formats:

|                      |        |                      |
|----------------------|--------|----------------------|
| <b>Concentrate:</b>  | 0.5 ml | (Cat. No. MSK090-05) |
| <b>Ready-to-use:</b> | 6 ml   | (Cat. No. MSG090)    |

#### Dilution of primary antibody

Dilution of Zytomed Systems' concentrated antibody depends on the detection system used. The final working dilution must always be determined by the user. The elaboration of staining protocol should be done by an experienced specialist. For Zytomed Systems' recommendations see chapter 'Staining procedure'.

#### Storage and handling

The antibody should be stored at 2-8°C without further dilution.

Dilutions of the concentrated antibody should be done in a suitable antibody dilution buffer (e.g. ZUC025 from Zytomed Systems). The diluted antibody should be stored at 2-8°C after use. Stability of this working solution depends on various parameters and has to be confirmed by appropriate controls. The antibody provided is suitable for use until the expiry date indicated on the label, if stored at 2-8°C. Do not use product after the expiry date. Positive and negative controls should be run simultaneously with all specimens. If unexpected staining is observed which cannot be explained by variations in laboratory procedures and a problem with the antibody is suspected, contact Zytomed Systems' technical support or your local distributor.

#### Precautions

Use through qualified personnel only.

Wear protective clothing to avoid contact of reagents and specimens with eye, skin and mucous membranes. If reagents or specimens come in contact with sensitive area, wash with large amounts of water.

Microbial contamination of the reagent must be avoided, since otherwise non-specific staining may occur.

Sodium azide ( $\text{NaN}_3$ ), used for stabilisation, is not considered hazardous material in the concentration used.

Reaction of sodium azide with lead or copper in drainage pipes can result in the formation of highly explosive metallic azides. Sodium azide should be discarded in a large volume of running water to avoid formation of deposits. Material safety data sheets (MSDS) are available upon request.

#### Staining procedure for formalin-fixed paraffin sections

Refer to the following table for conditions specifically recommended for this antibody. Also refer to detection system data sheets for guidance on specific staining protocols or other requirements.

### Parameters

\*Pre-treatment  
 \*Control tissue  
 \*Working dilution  
 \*Incubation time

### Zytomed Systems recommendations

Heat Induced Epitope Retrieval (for example in Citrate Buffer pH 6.0 (ZUC028))  
 Prostate adenocarcinoma  
 1:50-1:100 (for concentrates)  
 30 - 60 minutes

### Quality control

The recommended positive control tissue for this antibody is prostate adenocarcinoma. We recommend carrying out a positive and a negative control with every staining run. Please refer to the instructions of the detection system for guidance on general quality control procedures.

### Troubleshooting

If you observe unusual staining or other deviations from the expected results please read these instructions carefully, refer to the instructions of the detection system for relevant information or contact your local distributor.

### Expected results

This antibody stains positive in nuclei of malignant and premalignant prostate gland cells in formalin-fixed, paraffin-embedded tissue sections. Further details about the expression pattern of ERG can be found in the chapter 'Summary and Description'. Interpretation of the staining results is solely the responsibility of the user. Any experimental result should be confirmed by a medically established diagnostic procedure.

### Limitations of the Procedure

Immunohistochemistry is a complex technique involving both histological and immunological detection methods. Tissue processing and handling prior to immunostaining, for example variations in fixation and embedding or the inherent nature of the tissue can cause inconsistent results (Nadji and Morales, 1983). Endogenous peroxidase, alkaline phosphatase or biotin may cause non-specific staining depending on the detection system used. Tissues containing Hepatitis B Surface Antigen (HBsAg) may give false positive results with HRP (horse radish peroxidase) detection systems (Omata *et al*, 1980). Inadequate counterstaining and mounting can influence the interpretation of the results.

Zytomed Systems warrants that the product will meet all requirements described from its shipping date until the expiry date is reached, if the product is stored and utilised as recommended. No additional guarantees can be given. Under no circumstances shall Zytomed System be liable for any damages arising out of the use of the reagent provided.

### Performance characteristics

Zytomed Systems has conducted studies to evaluate the performance of the antibody for use with a standard detection system. The product has been found to be sensitive and specific to the antigen of interest with minimal or no cross-reactivity.

### Bibliography

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Han B *et al*. 2009 Mod Pathol 22(8):1083-1093  
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 Mehra R *et al*. 2007 Cancer Res 67(17):7991-7995  
 Nadji M and Morales AR Ann N.Y. Acad Sci 420:134-9, 1983  
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Explanations of the symbols on the product label:

|            |  |   |   |   |   |
|------------|--|---|---|---|---|
| <b>REF</b> | Bestellnummer<br>Catalog Number<br>Référence du catalogue  |  | Verwendbar bis<br>Use By<br>Utiliser jusque                             |  | Gebrauchsanweisung beachten<br>Consult Instructions for use<br>Consulter les instructions d'utilisation   |
| <b>LOT</b> | Chargenbezeichnung<br>Batch Code<br>Code du lot  |  | Lagerungstemperatur<br>Temperature Limitation<br>Limites de température | <b>RUO</b>  | Nur für Forschungszwecke<br>For Research Use Only<br>Pour la recherche uniquement   |
| <b>IVD</b> | In vitro Diagnostikum<br>In Vitro Diagnostic Medical Device<br>Dispositif médical de diagnostic in vitro |  | Achtung<br>Warning<br>Attention   |  | Hersteller / Manufacturer / Fabricant<br>Zytomed Systems GmbH • Anhaltinerstraße 16<br>14163 Berlin, Germany • Tel: (+49) 30-804 984 990<br>www.zytomed-systems.com |