

CEIVD Mouse anti-Cytokeratin Broad Spectrum (AE1 & AE3) Cat. No.: BMS006 (16 ml Ready-to-use)

Instructions for use

Intended Use

This antibody is designed for the specific localization of cytokeratins in formalin-fixed, paraffin-embedded tissue sections.

Anti-Cytokeratin Broad Spectrum antibody is intended for in vitro diagnostic use.

Specifications	
Specificity:	Human Cytokeratin pan
Clone:	Cocktail composed of AE1 and AE3
Isotype:	Mouse IgG1/k / Mouse IgG1/k
Species reactivity:	Human +, others not tested

Summary and explanation

Cytokeratins (CK) are intermediate filaments that constitute the cytoskeletal structure of virtually all epithelial but also of some non-epithelial cells. According to R. Moll they are divided into Type I (acidic cytokeratins, CK9 to 20) and Type II (basic cytokeratins, CK1 to 8) cytokeratins. Each Type I cytokeratin is co-expressed with a Type II cytokeratin inside a single cell. Hence, it follows that all epithelial cells contain at least two different cytokeratins. Only CK19 is expressed unpaired.

The antibody of clone AE1 detects the acidic (Type I) cytokeratins 10, 15, 16 and 19. The antibody derived from clone AE3 detects all basic (Type II) cytokeratins, i.e. CK1 to 8. That means the antibody cocktail AE1/AE3 detects all cells containing cytokeratins and is therefore called "pan-Cytokeratin" antibody. Detection of cytokeratins with a broad spectrum ("pan"-) antibody allows for the staining of epithelial cells in normal

Detection of cytokeratins with a broad spectrum ("pan"-) antibody allows for the staining of epithelial cells in normal and abnormal tissues. It is especially useful in characterisation of metastases with unknown origin. Lerwill (2004) has shown that cocktail AE1/AE3 detects carcinomic metastases in lymph nodes more specific than other pan-CK antibodies.

Reagent provided

Cocktail of mouse monoclonal antibodies in TBS with carrier protein and preservative for stabilisation in the following format:

Ready-to-use: 16 ml (Cat. No. BMS006)

Dilution of primary antibody

None

Storage and handling

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

If necessary, dilutions of the antibody should be done with 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 stable 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. ProClin300 and sodium azide (NaN₃) are used for stabilisation. Reaction of sodium azide with lead or copper in drainage pipes can result in the formation of highly explosive metallic azides. Discard the antibody solution in a large volume of running water to avoid formation of deposits. A material safety data sheet (MSDS) for the pure substances is available upon request.

Staining procedure

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	Zytomed Systems recommendations
*Pre-treatment:	Heat Induced Epitope Retrieval (for example in Citrate buffer pH 6.0)
*Control tissue	Skin, tonsil
*Working dilution	None
*Incubation time	30 minutes

Quality control

The recommended positive control tissues for this antibody are skin or tonsil. 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 guality 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

The antibody stains positive in the cytoplasm of epithelial cells in formalin-fixed, paraffin-embedded tissue. The interpretation of the 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, pseudoperoxidase activity in erythrocytes 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 utilising a standard detection system. The product has been found to be sensitive and specific to the antigen of interest with minimal or no crossreactivity.

Bibliography

Moll R et al. Cell 31:11-24, 1982 Southgate J et al. Histopathol 14:657-664, 1999 Chu PG und Weiss LM. Histopathol 40:403-439, 2002 Lerwill MF. Am J Surg Pathol 28:1076-1091, 2004 Battifora H. Am J Surg Pathol. 1988; 12:24.

Cooper D, et al. Lab Invest. 1985; 52:243-56 Gown AM, et al. AM J Clin Pathol. 1985; 84:413 Kriho UK, et al. Virehows Arch. 1997; 431:139-47 Nadji M and Morales AR. Ann N.Y. Acad Sci 1983; 420:134-139 Omata M et al Am J Clin Pathol 1980; 73: 626-632



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Explanations of the symbols on the product

label

Symbols are used in accordance with ISO 15223-1. Further symbols on the product label might be:



GSH07: Warning / Attention

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