



CD34

Monoclonal Mouse Anti-CD34 Clone QBEnd/10

REF 014-1053

Ready-To-Use ■ 100 Tests/50 Tests

Concentrate ■ 1mL

INTENDED USE

IVD For in vitro diagnostic use.

Celerus monoclonal mouse anti-CD34, clone QBEnd/10, is intended for laboratory use in identifying the human endothelial cell marker molecule, CD34, in normal and neoplastic tissue using light microscopy. It may be used with frozen tissue or with formalin-fixed paraffin-embedded tissue.

Positive results aid in the classification of normal and abnormal cells/tissues and serve as an adjunct to conventional histopathology. The clinical interpretation of any positive staining or its absence should be complemented by morphological and histological studies with proper controls. Evaluations should be made by a qualified individual in conjunction with the patient's clinical history and other diagnostic test results.

Refer to the Wave Instrument Operator's Manual for additional information concerning Materials Required but Not Provided; Storage; Staining Procedure; Troubleshooting; Interpretation of Staining; and General Limitations.

SUMMARY AND EXPLANATION

The CD34 molecule is a cell membrane glycoprotein found on certain lymphoid and myeloid hematopoietic progenitor cells and vascular endothelium. The CD34 antibody can be used to identify CD34 expression in a variety of neoplasias, including vascular tumors and acute lymphoblastic leukemias.

PRINCIPLE OF PROCEDURE

Immunohistochemistry is a multi-step process to identify specific cell markers within tissue biopsies or tumor specimens. The sequential steps include antigen retrieval (optional), antibody application, and antibody visualization followed by optional counterstaining. Specimens are then coverslipped and observed under light microscopy by trained personnel. Normally, multiple antibodies are tested to determine lineage and cell cycle markers. The Celerus Wave is an automated instrument that performs immunohistochemistry stains. For further information on the staining procedure, refer to the Celerus Wave Operator's Manual.

MATERIALS AND METHODS

Reagent Provided

Clone
QBEnd/10

Ig Class
IgG, kappa

Immunogen

Vascular suspension prepared from perfusate of human term placenta.

Ready-To-Use in Primary Antibody Cartridge

Celerus anti-CD34 is provided with ProClin 300 as a preservative, in a Primary Antibody Cartridge (PAC), a self-contained dispenser of reagents. Each PAC contains sufficient reagent to complete 100 stained slides. PACs must remain upright to avoid spilling. PAC must be primed before first use. See Celerus Wave Operator's Manual for details.

Concentrated Antibody

Liquid

Liquid concentrated antibody is provided containing 15 mM sodium azide as a preservative and 1% bovine serum albumin as a carrier protein.

Lyophilized

Lyophilized antibody is provided containing 15mM sodium azide as a preservative. Reconstitute vial with 1.0 ml distilled water.

Dilution

The suggested dilution is 1:35- 1:70. This is a guide only and users should determine their own optimal working dilutions.

SPECIFICITY

Human CD34 molecule.

MATERIALS REQUIRED BUT NOT PROVIDED

- Wave instrument
- Wave slide rack
- Positively-charged microscope slides, appropriately labeled
- Timer
- Celerus Riptide for antigen retrieval (or equivalent)
- Celerus Target Retrieval Solution (or equivalent)
- Slide drying chamber
- Xylene or xylene substitute
- Reagent alcohol or ethyl alcohol
- Distilled or deionized water
- TBS wash buffer, pH 7.6
- Positive and negative tissue controls
- Celerus Negative Control Reagent (or equivalent)
- Mounting Medium
- Cover slips

STORAGE AND HANDLING

Ready-to-Use PAC, Liquid Concentrated and Lyophilized Antibody

Store reagent at 2-8 °C. Do not freeze. The reagent is stable until the expiration date on the container. Do not use reagent after the expiration date, as the activity cannot be ensured.

Reconstituted Antibody

For reconstituted antibody, the reagent is stable for at least two months when stored at 4 °C. For long-term storage it is recommended that aliquots of the antibody be stored at -20 °C. Repeated freezing and thawing of the antibody should be avoided.

There are no signs to indicate instability of this reagent. To ensure a valid staining assay, the use of positive and negative tissue controls is recommended. Contact your Celerus representative if there are stability concerns prior to the expiration date.

PRECAUTIONS

- For professional users.
- Minimize microbial contamination of reagents or an increase in nonspecific staining may occur.
- As with any product derived from biological sources, proper handling procedures should be used.
- A Material Safety Data Sheet is available for professional users on request.

- ProClin 300 is classified per applicable European Community (EC) Directives as: Irritant (Xi). The following are the appropriate Risk (R) and Safety (S) phrases.



- R36** Irritating to eyes
R43 May cause sensitization by skin contact
S24 Avoid contact with skin
S26 In case of contact with eyes, rinse immediately with plenty of water and seek medical advice
S35 This material and its container must be disposed of in a safe way
S37/39 Wear suitable gloves and eye/face protection
S46 If swallowed, seek medical advice immediately and show this container or label.

WASTE DISPOSAL

Adhere to all local laws when disposing of the PAC.

PACKAGING DAMAGE

DO NOT USE a PAC if it is leaking, has leaked, has spilled, cannot be primed, or has visually apparent physical damage.

SPECIMEN COLLECTION AND HANDLING

Formalin-fixed paraffin embedded (FFPE) tissues, frozen tissues, or smears are suitable for use. Wave detection kits have been optimized for tissues fixed with 10% formalin. Ideally, each 4-6µ tissue section should be placed on charged slides on the lower 2/3 of the slide. Very large sections should be placed 1/4 inch below the lower end of the slide label.

Slides should be baked overnight at 37 °C, or at 60 °C for one hour.

Use standard histochemical techniques to deparaffinize processed slides. For uniformity of staining results, it is recommended that target retrieval be performed using the Celerus Riptide and Celerus Target Retrieval Solution (or equivalent) at 112 °C for 5 minutes. Avoid drying of the tissue specimen during this process. After all slides to be stained have been inserted and reagents mounted on the instrument, start the staining run.

When the slides have completed the staining run, remove them from the instrument, coverslip, and view under light microscopy.

PRODUCT-SPECIFIC LIMITATIONS

CD34 antibody, when used on the Wave instrument, detects antigens that survive routine formalin fixation, tissue processing, and sectioning. Users who deviate from recommended test procedures are responsible for interpretation and validation of patient results.

RESULTS EXPECTED/ PERFORMANCE CHARACTERISTICS

Normal Tissues

Clone QBEnd/10 detected the endothelial cell marker, CD34, in the cytoplasm of vascular endothelium in a variety of normal tissues.

Abnormal Tissues

Clone QBEnd/10 stained the endothelium of a variety of neoplastic tissues.

REFERENCES

- Nishio H, Tada J, Hashiyama M, Hirn J, Ingles-Esteve J, Suda T. MC7. CD34 workshop panel report. In: Kishimoto T, Kikutani H, von dem Borne AEG, Goyert SM, Mason DY, Miyasaka M, et al., editors. Leucocyte typing VI. White cell differentiation antigens. Proceedings of the 6th International Workshop and Conference; 1996 Nov 10-14; Kobe, Japan. New York, London: Garland Publishing Inc.; 1997. p. 974-76.
- Civin CI, Trischmann TM, Fackler MJ, Bernstein ID, Bühring HJ, Campos L, et al. M7.1. Report on the CD34 cluster workshop. In: Knapp W, Dörken B, Gilks WR, Rieber EP, Schmidt RE, Stein H, et al., editors. Leucocyte typing IV. White cell differentiation antigens. Proceedings of the 4th International Workshop and Conference; 1989 Feb 21-25; Vienna, Austria, Oxford, New York, Tokyo; Oxford University Press; 1989. p. 818-25.
- Civin CI, Strauss LC, Fackler MJ, Trischmann TM, Wiley JM, Loken MR. Positive stem cell selection-basic science. Prog Clin Biol Res 1990;333:387-402.
- Krause DS, Fackler MJ, Civin CI, May WS. CD34: structure, biology, and clinical utility (review). Blood 1996;87:1-13.
- Campos L, Guyotat D, Archimbaud E, Devaux Y, Trelle D, Larese A, et al. Surface marker expression in adult acute myeloid leukaemia; correlations with initial characteristics, morphology and response to therapy. Dr J Haematol 1989;72:161-6.
- Ramani P, Bradley NJ, Fletcher CDM. QBEND/10, a new monoclonal antibody to endothelium; assessment of its diagnostics utility in paraffin sections. Histopathology 1990;17:237-242.
- Dercksen MW, Daams GM, de Haas M, von dem Borne AEG, dan der Schoot CE. M10.2 Characterization of the CD34 cluster. In: Schlossman SF, Boumsell L, Gilks W, Harlan JM, Kishimoto T, Morimoto C, et al., editors. Leucocyte typing V. White cell differentiation antigens. Proceedings of the 5th International Workshop and Conference; 1993 Nov 3-7; Boston, USA, Oxford, New York, Tokyo: Oxford University Press; 1995. p. 850-3.
- Fina L, Molgaard HV, Robertson D, Bradley NJ, Monaghan P, Delia D, et al. Expression of the CD34 gene in vascular endothelial cells. Blood 1990;75(12):2417-2426.
- Serke S, van Lessen A., Huhn D. Quantitative fluorescence flow cytometry; a comparison of the three techniques for direct and indirect immunofluorescence. Cytometry 1998;33:179-87.



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