

Datasheet for ABIN6151499
anti-CD44 Standard antibody (CF®488A)



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Overview

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| Quantity: | 100 µL |
| Target: | CD44 Standard (CD44s) |
| Reactivity: | Human, Mouse |
| Host: | Mouse |
| Clonality: | Monoclonal |
| Conjugate: | This CD44 Standard antibody is conjugated to CF®488A |
| Application: | Flow Cytometry (FACS), Immunofluorescence (IF), Immunohistochemistry (Formalin-fixed Sections) (IHC (f)) |

Product Details

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| Purpose: | Mouse Monoclonal anti-CD44 Standard (DF1485), CF488A Conjugate |
| Immunogen: | Purified CD44 antigen (PGp-1) from lymphocyte membrane |
| Clone: | DF1485 |
| Isotype: | IgG1 kappa |
| Characteristics: | This antibody recognizes a cell surface glycoprotein of 80-95 kDa (CD44) on lymphocytes, monocytes, and granulocytes (Leucocyte Typing Workshop V). Its epitope is resistant to digestion by trypsin and chymotrypsin. The CD44 family of glycoproteins exists in a number of variant isoforms, the most common being the standard 85-95 kDa or hematopoietic variant (CD44s). Higher molecular weight isoforms are described in epithelial cells (CD44v), which are believed to function in intercellular adhesion and stromal binding. CD44 immunostaining is commonly used for the discrimination of urothelial transitional cell carcinoma in-situ from non- |

Product Details

neoplastic changes in the urothelium. Primary antibodies are available purified, or with a selection of fluorescent CF® dyes and other labels. CF® dyes offer exceptional brightness and photostability. Note: Conjugates of blue fluorescent dyes like CF®405S and CF®405M are not recommended for detecting low abundance targets, because blue dyes have lower fluorescence and can give higher non-specific background than other dye colors.

Target Details

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| Target: | CD44 Standard (CD44s) |
| Alternative Name: | CD44 Standard (CD44s Products) |
| Molecular Weight: | 80-95 kDa |
| Gene ID: | 960, 502328 |
| UniProt: | P16070 |

Application Details

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| Application Notes: | Immunohistology formalin-fixed 0.5-1 µg/mL <ul style="list-style-type: none">• Staining of formalin-fixed tissues requires boiling tissue sections in 10 mM citrate buffer, pH 6.0, for 10-20 min followed by cooling at RT for 20 minutes• Immunofluorescence 0.5-1 µg/mL• Flow Cytometry 0.5-1 µg/million cells/0.1 mL• Optimal dilution for a specific application should be determined by user |
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| Comment: | HeLa cells or paracortex in tonsil or lymph node. |
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| Restrictions: | For Research Use only |
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Handling

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| Format: | Liquid |
| Concentration: | 100 µg/mL |
| Buffer: | PBS/0.1 % BSA/0.05 % azide |
| Preservative: | Sodium azide |
| Precaution of Use: | This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only. |
| Handling Advice: | Protect from light |