

### Datasheet for ABIN6151487

# anti-CD44 Standard antibody (CF®555)



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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®555	
Application:	Flow Cytometry (FACS), Immunofluorescence (IF), Immunohistochemistry (Formalin-fixed Sections) (IHC (f))	

### **Product Details**

Purpose:	Mouse Monoclonal anti-CD44 Standard (DF1485), CF555 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,	

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-

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**

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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Immunohistology formalin-fixed 0.5-1 µg/mL

- 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

#### Comment:

HeLa cells or paracortex in tonsil or lymph node.

Restrictions:

For Research Use only

# Handling

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