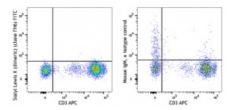
Product Data Sheet

FITC anti-human Sialyl Lewis X (dimeric)

Catalog # / Size:	2440560 / 100 tests 2440555 / 25 tests
Clone:	FH6
Isotype:	Mouse IgM, κ
Immunogen:	Purified 6B fucoganglioside absorbed to Salmonella minnesota.
Reactivity:	Human
Preparation:	The antibody was purified by affinity chromatography and conjugated with FITC under optimal conditions. The solution is free of unconjugated FITC and unconjugated antibody.
Formulation:	Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide and 0.2% (w/v) BSA (origin USA).
Concentration:	Lot-specific



Human peripheral blood Lymphocytes were stained with sialyl lewis X (dimeric) (clone FH6) FITC and UCHT1 APC (left) or mouse IgM

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Applications:

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Applications:	Flow Cytometry	
Recommended Usage:	Each lot of this antibody is quality control tested by immunofluorescent staining with flow cytometric analysis. For flow cytometric staining, the suggested use of this reagent is 5 µl per million cells or 5 µl per 100 µl of whole blood. It is recommended that the reagent be titrated for optimal performance for each application.	Human peripheral blood granulocytes were stained with sialyl lewis X (dimeric) (clone FH6) FITC (filled histogram) or mouse IgM
Application References:	1. Fukushi Y, <i>et al.</i> 1984. <i>J. Biol. Chem.</i> 259:10511. 2. Kannagi R, <i>et al.</i> 1986. <i>Cancer Research</i> 5:2619. 3. Nakasaki H, <i>et al.</i> 1989. <i>Cancer Research</i> 49:3662. 4. Dohi T, <i>et al.</i>	
Description:	The FH6 antibody recognizes Sialyl Lewis X (demeric) on glycolipids or glycoproteins. It also recognizes Sialyl Lewis X with long carbohydrate attachments (Sialyl Lewis X-i). These antigens are expressed on human granulocytes, monocytes, small subsets of lymphocytes, some fetal tissues such as the fetal stomach, fetal colon, and fetal intestine, and a variety of cancer tissues. It is believed that these antigens are involved in cell adhesion.	
Antigen References:	1. Fukushi Y, <i>et al.</i> 1984. <i>J. Biol. Chem.</i> 259:10511. 2. Kannagi R, <i>et al.</i> 1986. <i>Cancer Research</i> 5:2619. 3. Nakasaki H, <i>et al.</i> 1989. <i>Cancer Research</i> 49:3662.	

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