## Human Immune Cell Phenotyping IHC **Antibody Sampler Kit**



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For Research Use Only. Not for Use in Diagnostic Procedures.

1 Kit (9 x 20 microliters)

Product #	Quantity	Mol. Wt	Isotype/Source
85061	20 µl	23 kDa	Rabbit IgG
70306	20 µl		Mouse IgG1
98377	20 µl	45 kDa	Rabbit IgG
49420	20 µl	170 kDa	Rabbit IgG
76437	20 µl		Rabbit IgG
45581	20 µl	145 kDa	Rabbit IgG
90176	20 µl	95 kDa	Rabbit IgG
4545	20 µl	46-58 kDa	Mouse IgG1
99746	20 µl	120 to 220 kDa	Rabbit IgG
	85061 70306 98377 49420 76437 45581 90176 4545	85061     20 μl       70306     20 μl       98377     20 μl       49420     20 μl       76437     20 μl       45581     20 μl       90176     20 μl       4545     20 μl	85061     20 μl     23 kDa       70306     20 μl     45 kDa       98377     20 μl     45 kDa       49420     20 μl     170 kDa       76437     20 μl     145 kDa       90176     20 μl     95 kDa       4545     20 μl     46-58 kDa

Please visit cellsignal.com for individual component applications, species cross-reactivity, dilutions, protocols, and additional product information.

Description The Human Immune Cell Phenotyping IHC Antibody Sampler Kit provides an economical means of detecting the accumulation of immune cell types in formalin-fixed, paraffin-embedded tissue samples. Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 µg/ml BSA, 50% glycerol and less than Storage 0.02% sodium azide. Store at -20°C. Do not aliguot the antibody. Background Cluster of Differentiation 3 (CD3) is a multiunit protein complex expressed on the surface of T cells that directly associates with the T cell receptor (TCR). CD3 is composed of four polypeptides:  $\zeta$ ,  $\gamma$ ,  $\varepsilon$  and  $\delta$ . Engagement of the TCR complex with antigens presented in Major Histocompatibility Complexes (MHC) induces tyrosine phosphorylation in the immunoreceptor tyrosine-based activation motif (ITAM) of CD3 proteins. CD3 phosphorylation is required for downstream signaling through ZAP-70 and p85 subunit of PI-3 kinase, leading to T cell activation, proliferation, and effector functions (1). CD8 is a transmembrane glycoprotein expressed primarily on cytotoxic T cells, but has also been described on a subset of dendritic cells in mice (2,3). On T cells, CD8 is a co-receptor for the TCR, and these two distinct structures are required to recognize antigen bound to MHC Class I. CD8 ensures specificity of the TCRantigen interaction, prolongs the contact between the T cell and the antigen presenting cell, and recruits the tyrosine kinase Lck, which is essential for T cell activation (2). Forkhead box P3 (FoxP3) is crucial for the development of T cells with immunosuppressive regulatory properties and is a wellestablished marker for CD4<sup>+</sup> T regulatory cells (Tregs) (4). Cluster of differentiation molecule 11b (CD11b)/Integrin alpha M (ITGAM) is a transmembrane protein forming heterodimers that are composed of  $\alpha$  and  $\beta$  subunits (5). CD11b is expressed by, and commonly used as a marker for, myeloid lineage cells, including neutrophils, monocytes, macrophages, and microglia (6). CD68 (macrosialin) is a heavily glycosylated transmembrane protein that is expressed by and commonly used as a marker for monocytes and macrophages (7,8). It is found on the plasma membrane, as well as endosomal and lysosomal membranes (7-9). CD11c (integrin αΧ, ITGAX) is a transmembrane glycoprotein highly expressed by dendritic cells, and has also been observed on activated NK cells, subsets of B and T cells, monocytes, granulocytes, and some B cell malignancies including hairy cell leukemia (10,11). CD19 is a co-receptor expressed on B cells that amplifies the signaling cascade initiated by the B cell receptor (BCR) to induce activation. It is a biomarker of B lymphocyte development, lymphoma diagnosis, and can be utilized as a target for leukemia immunotherapies (12,13). NCAM (neural cell adhesion molecule, CD56) is an adhesion glycoprotein with five extracellular immunoglobulin-like domains followed by two fibronectin type III repeats (14). CD56 and CD16 are commonly used to identify NK cells although some cells with the T cell markers CD3 and CD4 also express CD56 (15). Keratins (cytokeratins) are intermediate filament proteins that are mainly expressed in epithelial cells. Keratin heterodimers composed of an acidic keratin (or type I keratin, keratins 9 to 23) and a basic keratin (or type II keratin, keratins 1 to 8) assemble to form filaments (16,17). Keratin isoforms demonstrate tissue- and differentiation-specific profiles that make them useful as research biomarkers (16). **Background References** 1. Kuhns, M.S. et al. (2006) Immunity 24, 133-9. 2. Zamoyska, R. (1994) Immunity 1, 243-6.

	<ol> <li>Shortman, K. and Heath, W.R. (2010) <i>Immunol Rev</i> 234, 18-31.</li> <li>Ochs, H.D. et al. (2007) <i>Immunol Res</i> 38, 112-21.</li> <li>Solovjov, D.A. et al. (2005) <i>J Biol Chem</i> 280, 1336-45.</li> <li>Murray, P.J. and Wynn, T.A. (2011) <i>Nat Rev Immunol</i> 11, 723-37.</li> <li>Rabinowitz, S.S. and Gordon, S. (1991) <i>J Exp Med</i> 174, 827-36.</li> <li>Holness, C.L. and Simmons, D.L. (1993) <i>Blood</i> 81, 1607-13.</li> <li>Ramprasad, M.P. et al. (1995) <i>Proc Natl Acad Sci U S A</i> 92, 9580-4.</li> <li>Kohrgruber, N. et al. (1999) <i>J Immunol</i> 163, 3250-9.</li> <li>Qualai, J. et al. (2016) <i>PLoS One</i> 11, e0154253.</li> <li>Tedder, T.F. et al. (1997) <i>Immunity</i> 6, 107-18.</li> <li>Scheuermann, R.H. and Racila, E. (1995) <i>Leuk Lymphoma</i> 18, 385-97.</li> <li>Cunningham, B.A. et al. (1987) <i>Science</i> 236, 799-806.</li> <li>Robertson, M.J. and Ritz, J. (1990) <i>Blood</i> 76, 2421-38.</li> <li>Moll, R. et al. (1982) <i>Cell</i> 31, 11-24.</li> <li>Chang, L. and Goldman, R.D. (2004) <i>Nat Rev Mol Cell Biol</i> 5, 601-13.</li> </ol>
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