

## Anti-CD158e1 Antibody-PE labeled

Catalog #	Source	Reactivity	Applications
CFF8965	Mouse	H	IF, FC
<b>Description</b>	Mouse monoclonal antibody PE labeled to CD158e1		
<b>Immunogen</b>	Human NK cell clone VL186-1.6		
<b>Purification</b>	The antibody was purified by affinity chromatography.		
<b>Specificity</b>	Recognizes human CD158e1		
<b>Clonality</b>	Monoclonal (clone: DX9)		
<b>Conjugation</b>	PE		
<b>Form</b>	Mouse IgG1 kappa. Liquid in PBS, pH 7.3, and 0.02% sodium azide.		
<b>Dilution</b>	10 µl / assay		
<b>Gene Symbol</b>	KIR3DL1		
<b>Alternative Names</b>	CD158E; NKAT3; NKB1; Killer cell immunoglobulin-like receptor 3DL1; CD158 antigen-like family member E; HLA-BW4-specific inhibitory NK cell receptor; MHC class I NK cell receptor; Natural killer-associated transcript 3; NKAT-3; p70 natural killer cell receptor clones CL-2/CL-11; p70 NK receptor CL-2/CL-11; CD158e		
<b>Entrez Gene</b>	3811 (Human)		
<b>SwissProt</b>	P43629 (Human)		
<b>Directions for Use</b>	<ol style="list-style-type: none"> <li>1. Take 100 µl peripheral blood anticoagulated by EDTA and add to the bottom of 5 ml tube.</li> <li>2. Add 10 µl labeled antibody to the bottom of flow tube mixing with the whole blood, incubate for 20 minutes at room temperature away from light.</li> <li>3. Add 2 ml RBC lysis buffer, incubate for 10 minutes away from light after mixing, dissolve red blood cells.</li> </ol>		

**Application key:** E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, RNAi- RNA interference

**Species reactivity key:** H- Human, M- Mouse, R- Rat, B- Bovine, C- Chicken, D- Dog, G- Goat, Mk- Monkey, P- Pig, Rb- Rabbit, S- Sheep, Z- Zebrafish

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## Product Data Sheet

4. Sample tube is set to 1000 rpm centrifugation for 5 minutes, discard the supernatant.
5. Add 2 ml PBS wash buffer to resuspend the cells, then 1000 rpm centrifugation for 5 minutes, discard the supernatant.
6. Add 0.5 ml PBS wash buffer to resuspend the cells and detect by flow cytometry (sample should be determined on the day on the machine and can also be added fixation overnight at 4 °C then measured).

### Storage/Stability

Shipped and store at 4°C for one year. Do not freeze.

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