Technical Data Sheet

Biotin Rat Anti-Mouse CD94

Product Information

 Material Number:
 550773

 Size:
 0.1 mg

 Concentration:
 0.5 mg/ml

 Clone:
 18d3

 Immunogen:
 Transfected cell line

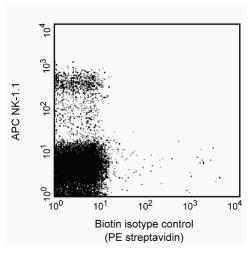
 Isotype:
 Rat (LEW) IgG2a, κ

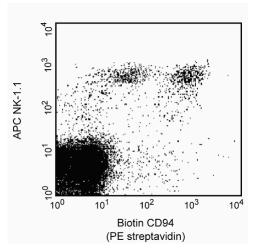
 Reactivity:
 QC Testing: Mouse

Storage Buffer: Aqueous buffered solution containing ≤0.09% sodium azide.

Description

The 18d3 antibody reacts with CD94 on all NK cells, NK1.1- or DX5-positive T lymphocytes (NK-T cells), and a subset of CD8-positive T lymphocytes in most strains tested (eg, A/J, AKR/J, BALB/c, C3H/He, C57BL/6, CBA/J, DBA/1, FVB/N, 129/Sv, NOD, SWR, and most DBA/2 substrains, but not DBA/2J). DBA/2J mice do not express CD94.3 CD94 is also expressed on CD8+ T lymphocytes activated in vivo. CD94 is a type-II transmembrane protein with an extracellular lectin-like domain and a short cytoplasmic tail which is not believed to have any signalling function. Heterodimers of CD94 with NKG2A, NKG2C, or NKG2E recognize Qa-1 (a non-classical MHC class I antigen) presenting the Qdm peptide. Studies on CD94/NKG2 heterodimers on human NK cells have demonstrated that the NKG2 components mediate signal transduction for the receptor, NKG2A being inhibitory and NKG2C being stimulatory. Similarly, the mouse NKG2A molecule contains two intracytoplasmic sequences which resemble the ITIM (Immunoreceptor Tyrosine-based Inhibitory Motif) consensus sequence. CD94/NKG2 receptors appear on fetal NK cells before the Ly- 49 MHC class I receptors, suggesting that CD94/NKG2 receptors and their ligand, Qa-1, may play a role in maintenance of self-tolerance in developing NK cells.





Expression of CD94 on mouse splenic NK cells. C57BL/6 splenocytes were preincubated with Mouse BD Fc Block^T purified anti-mouse CD16/CD32 mAb 2.4G2 (Cat. no. 553141/553142), then simultaneously stained with biotinconjugated rat IgG2a isotype control mAb R35-95 (Cat. no. 553928, Left panel) or biotinylated mAb 18d3 (Right panel) and APC-conjugated anti- mouse NK-1.1 mAb PK136 (Cat. no. 550627), followed by Streptavidin-PE (Cat. no. 554061). The CD94[dim] and CD94[bright] subpopulations of NK cells are observed in all strains tested, although the relative proportions of the two subsets may vary. Flow cytometry was performed on a BD FACSCalibur™ flow cytometry system.

Preparation and Storage

The monoclonal antibody was purified from tissue culture supernatant or ascites by affinity chromatography. The antibody was conjugated with biotin under optimum conditions, and unreacted biotin was removed. Store undiluted at 4° C and protected from prolonged exposure to light. Do not freeze.

Application Notes

Application

-	pheadon		
	Flow cytometry	Routinely Tested	
	Immunohistochemistry-frozen	Not Recommended	

Recommended Assay Procedure:

We have found that the use of Mouse BD Fc Block, purified anti-mouse CD16/CD32 mAb 2.4G2 (Cat. no. 553141/553142) reduces the non-specific staining of non-NK cells by this biotin conjugate. This antibody is not useful for immunohistochemical staining of frozen sections.

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550773 Rev. 5

Suggested Companion Products

Catalog Number	Name	Size	Clone	
553141	Purified Rat Anti-Mouse CD16/CD32 (Mouse BD Fc Block TM)	0.1 mg	2.4G2	
553928	Biotin Rat IgG2a κ Isotype Control	0.25 mg	R35-95	
550627	APC Mouse Anti-Mouse NK-1.1	0.1 mg	PK136	
554061	PE Streptavidin	0.5 mg	(none)	

Product Notices

- 1. Since applications vary, each investigator should titrate the reagent to obtain optimal results.
- Please refer to www.bdbiosciences.com/pharmingen/protocols for technical protocols.
- 3. Caution: Sodium azide yields highly toxic hydrazoic acid under acidic conditions. Dilute azide compounds in running water before discarding to avoid accumulation of potentially explosive deposits in plumbing.

References

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Sivakumar PV, Gunturi A, Salcedo M, et al. Cutting edge: expression of functional CD94/NKG2A inhibitory receptors on fetal NK1.1+Ly-49- cells: a possible mechanism of tolerance during NK cell development. *J Immunol.* 1999; 162(12):6976-6980.(Biology)

Toomey JA, Salcedo M, Cotterill LA. Stochastic acquisition of Qa1 receptors during the development of fetal NK cells in vitro accounts in part but not in whole for the ability of these cells to distinguish between class I-sufficient and class I-deficient targets. *J Immunol.* 1999; 163(6):3176-3184.(Biology)

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Vance RE, Jamieson AM, Raulet DH. Recognition of the class lb molecule Qa-1(b) by putative activating receptors CD94/NKG2C and CD94/NKG2E on mouse natural killer cells. *J Exp Med*. 1999; 190(12):1801-1812.(Immunogen)

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550773 Rev. 5