# **Technical Data Sheet**

# Purified Mouse Anti-Rat CD4

#### **Product Information**

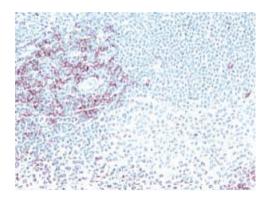
**Material Number:** 550297 Size: 1.0 ml  $125 \mu g/ml$ Concentration: OX-38 Clone:

Immunogen: Rat thymocyte glycoproteins Isotype: Mouse (BALB/c) IgG2a, κ Reactivity: QC Testing: Rat

Storage Buffer: Aqueous buffered solution containing BSA, goat serum, and ≤0.09% sodium

#### Description

The OX-38 antibody has been reported to react with the CD4 antigen on most thymocytes, a subpopulation of mature T lymphocytes (i.e., MHC class II-restricted T cells, including most T helper cells), monocytes, macrophages, and some dendritic cells. CD4 is an antigen coreceptor on the T-cell surface which interacts with MHC class II molecules on antigen-presenting cells. It participates in T-cell activation through its association with the T-cell receptor complex and protein tyrosine kinase lck. The OX-38 antibody has been reported to bind to the same epitope of CD4 as that recognized by W3/25 mAb, which is a different epitope than that recognized by OX-35 mAb (Cat. No. 554837). In vivo blocking of some cell-mediated immune responses by mAb OX-38 has been reported. Injection of OX-38 mAb induces allograft unresponsiveness in rats, with varying results depending on the rat strain used (high or low responder). Furthermore, in vivo depletion of CD4+ lymphocytes has been reported with this antibody.



Immunohistochemical staining of rat CD4 + T lymphocytes. Frozen sections of normal rat spleen were stained with Purified Mouse Anti-Rat CD4 antibody (Cat. No. 550297). CD4+ T lymphocytes can be identified by the intense brown labeling of their cell surface membranes. Amplification 20X.

## **Preparation and Storage**

Store undiluted at 4°C.

The monoclonal antibody was purified from tissue culture supernatant or ascites by affinity chromatography.

# **Application Notes**

Application

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Flow cytometry	Routinely Tested
Immunohistochemistry-frozen	Tested During Development
Immunohistochemistry-formalin (antigen retrieval required)	Not Recommended

### **Recommended Assay Procedure:**

Immunohistochemistry: Clone OX-38 is tested for immunohistochemical staining of acetone-fixed frozen sections of rat spleen or thymus. IHC of formalin-fixed paraffin embedded sections is not recommended. This antibody has been reported to stain the CD4 subset of T lymphocytes. The isotype control recommended for use with this antibody is purified mouse IgG2a (Cat. No. 550339). For optimal indirect immunohistochemical staining, the OX-38 antibody should be titrated (1:10 to 1:50 dilution) and visualized via a three-step staining procedure in combination with biotin conjugated rat anti-mouse IgG2a (Cat. No. 550332) as the secondary antibody and Streptavidin-HRP (Cat. No. 550946) together with the DAB detection system (Cat. No. 550880).

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## **Suggested Companion Products**

Catalog Number	<u>Name</u>	Size	Clone	_
550880	DAB Substrate Kit	500 tests	(none)	
550946	Streptavidin HRP	50 ml	(none)	
550339	Purified Mouse IgG2a κ Isotype Control	1.0 ml	C1.18.4	
550332	Biotin Rat Anti-Mouse IgG2a	1.0 ml	R19-15	
550296	Purified Mouse Anti-Rat CD4	1.0 ml	OX-35	
554835	Purified Mouse Anti-Rat CD4	0.5 mg	OX-35	

#### **Product Notices**

- Since applications vary, each investigator should titrate the reagent to obtain optimal results.
- Source of all serum proteins is from USDA inspected abattoirs located in the United States.
- 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.
- 4 An isotype control should be used at the same concentration as the antibody of interest.
- Sodium azide is a reversible inhibitor of oxidative metabolism; therefore, antibody preparations containing this preservative agent must not be used in cell cultures nor injected into animals. Sodium azide may be removed by washing stained cells or plate-bound antibody or dialyzing soluble antibody in sodium azide-free buffer. Since endotoxin may also affect the results of functional studies, we recommend the NA/LE (No Azide/Low Endotoxin) antibody format, if available, for in vitro and in vivo use.
- This antibody has been developed for the immunohistochemistry application. However, a routine immunohistochemistry test is not performed on every lot. Researchers are encouraged to titrate the reagent for optimal performance.
- Please refer to www.bdbiosciences.com/pharmingen/protocols for technical protocols.

#### References

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Liu L, Zhang M, Jenkins C, MacPherson GG. Dendritic cell heterogeneity in vivo: two functionally different dendritic cell populations in rat intestinal lymph can be distinguished by CD4 expression. J Immunol. 1998; 161(3):1146-1155. (Biology)

Stitz L, Sobbe M, Bilzer T. Preventive effects of early anti-CD4 or anti-CD8 treatment on Borna disease in rats. J Virol. 1992; 66(6):3316-3323. (Clone-specific: Blocking)

Suzuki H, Hara MH, Miyahara T, et al. Microchimerism and graft acceptance: IV. Cardiac allograft acceptance following anti-adhesion molecule antibody therapy. Transplant Proc. 1996; 28(4):2058-2060. (Clone-specific: Blocking)

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