# **Technical Data Sheet** V450 Mouse Anti-Human CD69

#### **Product Information**

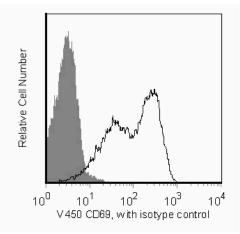
Material Number: Alternate Name: Size Vol. per Test: **Clone:** Isotype: **Reactivity:** Workshop: **Storage Buffer:** 

560740 Very Early Activation Antigen 50 tests 5 µl **FN50** Mouse IgG1, ĸ QC Testing: Human IV A091 Aqueous buffered solution containing protein stabilizer and ≤0.09% sodium azide.

## Description

Reacts with a 28/34 kDa dimeric glycoprotein expressed early during activation of lymphocytes and monocytes. FN50 monoclonal antibody labels NK cells and most lymphocytes of the follicular mantle and perifollicular/interfollicular zone as well as intragerminal center T cells of lymph nodes and tonsils.

The antibody is conjugated to BD Horizon™ V450, which has been developed for use in multicolor flow cytometry experiments and is available exclusively from BD Biosciences. It is excited by the Violet laser Ex max of 406 nm and has an Em Max at 450 nm. Conjugates with BD Horizon<sup>™</sup> V450 can be used in place of Pacific Blue<sup>™</sup> conjugates.



Flow cytometric analysis for CD69 in stimulated peripheral blood mononuclear cells (PBMC). Human PBMC were stimulated for at least 4-6 hours with 20 ng/mL PMA (Sigma-Aldrich Cat. No. P-8139) and 250 ng/mL calcium ionophore A23187 (Sigma-Aldrich Cat. No. C-9275). Cells were then stained with either a BD Horizon<sup>™</sup> V450 Mouse IgG1, κ isotype control (shaded) or with the BD Horizon™ V450 Mouse Anti-Human CD69 antibody (unshaded). Histograms were derived from gated events based on light scattering characteristics for lymphocytes. Flow cytometry was performed on a BD™ LSR II flow cytometry system.

## **Preparation and Storage**

Store undiluted at 4°C and protected from prolonged exposure to light. Do not freeze.

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

The antibody was conjugated with BD Horizon<sup>™</sup> V450 under optimum conditions, and unreacted BD Horizon<sup>™</sup> V450 was removed.

## **Application Notes**

Flow cytometry		Routinely Tested					
Suggested Comp	anion Products						
Catalog Number	Name	Name		Clone			
560373	V450 Mouse IgG1, κ Isotype Control		0.1 mg	MOPC-21			
555899	Lysing Buffer		100 ml	(none)			
sample (a test).	1	recommended Volume per Test. We typically use oncentration as the antibody of interest.	e 1 × 10^6 cells in a 100	)-μl experimental			
BD Biosciences bdbiosciences.com United States Canada	Europe Japan	Asia Pacific Latin America/Caribbean		A DI			

United States 877.232.8995	<b>Canada</b> 888.259.0187	Europe 32.53.720.550	<b>Japan</b> 0120.8555.90	Asia Pacific 65.6861.0633	Latin America/Caribbean 55.11.5185.9995			
For country-specific contact information, visit bdbiosciences.com/how_to_order/								
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- 3. BD Horizon<sup>TM</sup> V450 has a maximum absorption of 406 nm and maximum emission of 450 nm. Before staining with this reagent, please confirm that your flow cytometer is capable of exciting the fluorochrome and discriminating the resulting fluorescence.
- 4. 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.
- 5. Pacific Blue<sup>™</sup> is a trademark of Molecular Probes, Inc., Eugene, OR.
- 6. For fluorochrome spectra and suitable instrument settings, please refer to our Fluorochrome Web Page at www.bdbiosciences.com/colors.
- 7. Please refer to www.bdbiosciences.com/pharmingen/protocols for technical protocols.

#### References

Knapp W, Dorken B, Rieber EP, et al, ed. *Leucocyte Typing IV*. New York: Oxford University Press; 1989. (Biology) Schlossman S, Boumell L, et al, ed. *Leucocyte Typing V*. New York: Oxford University Press; 1995. (Biology)