

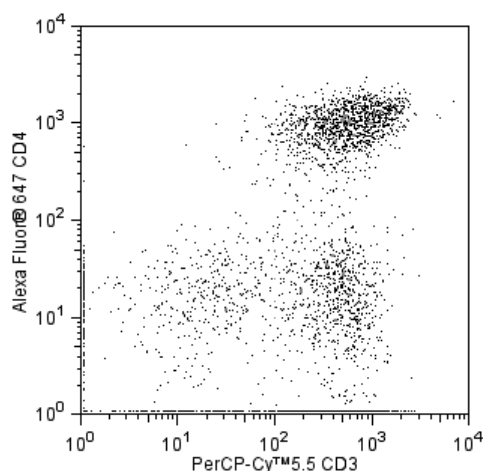
## Technical Data Sheet

**PerCP-Cy™5.5 Mouse Anti-Pig CD3ε****Product Information**

<b>Material Number:</b>	<b>561478</b>
<b>Alternate Name:</b>	CD3 epsilon subunit; CD3ε; T-cell surface glycoprotein CD3 epsilon chain
<b>Size:</b>	50 µg
<b>Concentration:</b>	0.2 mg/ml
<b>Clone:</b>	BB23-8E6-8C8
<b>Immunogen:</b>	Pig peripheral blood mononuclear cells
<b>Isotype:</b>	Mouse (BALB/c) IgG2a, κ
<b>Reactivity:</b>	QC Testing: Pig
<b>Storage Buffer:</b>	Aqueous buffered solution containing ≤0.09% sodium azide.

**Description**

The BB23-8E6-8C8 monoclonal antibody specifically binds to the 25-kDa ε chain of the T-cell receptor-associated CD3 complex. It recognizes all CD4+ and most CD8+ peripheral blood T lymphocytes, most thymocytes and phytohemagglutinin-stimulated blasts, and subsets of spleen and Peyer's patch lymphocytes. BB23-8E6-8C8 is a immunoglobulin isotype switch variant of the BB23-8E6 clone. This isotype-switch variant induces a proliferative response of peripheral blood mononuclear cells. The epitope recognized by BB23-8E6 mAb was designated CD3a by the Second International Swine CD Workshop.



**Multicolor flow cytometric analysis of CD3 expression on pig peripheral blood lymphocytes.** Pig whole blood was stained simultaneously with PerCP-Cy™5.5 Mouse Anti-Pig CD3ε antibody (Cat. No. 561478) and Alexa Fluor® 647 Mouse Anti-Pig CD4 antibody (Cat. No. 561472). The erythrocytes were lysed with BD PharmLyse™ Lysing Buffer (Cat. No. 555899). A two-color flow cytometric dot plot showing the correlated expression of CD3 versus CD4 was derived from gated events with the forward and side light-scatter characteristics of viable lymphocytes. Flow cytometry was performed using a BD™ LSR II Flow Cytometer System.

**Preparation and Storage**

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

The antibody was conjugated with PerCP-Cy5.5 under optimum conditions, and unconjugated antibody and free PerCP-Cy5.5 were removed. Storage of PerCP-Cy5.5 conjugates in unoptimized diluent is not recommended and may result in loss of signal intensity. Store undiluted at 4°C and protected from prolonged exposure to light. Do not freeze.

**Application Notes****Application**

Flow cytometry	Routinely Tested
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**Suggested Companion Products**

Catalog Number	Name	Size	Clone
555899	Lysing Buffer	100 ml	(none)
554656	Stain Buffer (FBS)	500 ml	(none)
561472	Alexa Fluor® 647 Mouse Anti-Pig CD4a	50 µg	74-12-4
550927	PerCP-Cy™5.5 Mouse IgG2a, κ Isotype Control	0.1 mg	G155-178

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## Product Notices

1. Since applications vary, each investigator should titrate the reagent to obtain optimal results.
2. Cy is a trademark of Amersham Biosciences Limited. This conjugated product is sold under license to the following patents: US Patent Nos. 5,486,616; 5,569,587; 5,569,766; 5,627,027.
3. Please observe the following precautions: Absorption of visible light can significantly alter the energy transfer occurring in any tandem fluorochrome conjugate; therefore, we recommend that special precautions be taken (such as wrapping vials, tubes, or racks in aluminum foil) to prevent exposure of conjugated reagents, including cells stained with those reagents, to room illumination.
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. This PerCP-conjugated product is sold under license to the following patent: US Patent No. 4,876,190.
6. PerCP-Cy5.5-labelled antibodies can be used with FITC- and R-PE-labelled reagents in single-laser flow cytometers with no significant spectral overlap of PerCP-Cy5.5, FITC, and R-PE fluorescence.
7. PerCP-Cy5.5 is optimized for use with a single argon ion laser emitting 488-nm light. Because of the broad absorption spectrum of the tandem fluorochrome, extra care must be taken when using dual-laser cytometers, which may directly excite both PerCP and Cy5.5™. We recommend the use of cross-beam compensation during data acquisition or software compensation during data analysis.
8. For fluorochrome spectra and suitable instrument settings, please refer to our Fluorochrome Web Page at [www.bdbiosciences.com/colors](http://www.bdbiosciences.com/colors).
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10. Please refer to [www.bdbiosciences.com/pharming/protocols](http://www.bdbiosciences.com/pharming/protocols) for technical protocols.
11. An isotype control should be used at the same concentration as the antibody of interest.

## References

Pescovitz MD, Book BK, Aasted B. Summary of workshop findings for antibodies reacting with porcine T-cells and activation antigens: results from the Second International Swine CD Workshop. *Vet Immunol Immunopathol.* 1998; 60(3-4):251-260. (Clone-specific)

Pescovitz MD, Book BK, Aasted B. Analyses of monoclonal antibodies reacting with porcine CD3: results from the Second International Swine CD Workshop. *Vet Immunol Immunopathol.* 1998; 60(3-4):261-268. (Clone-specific)