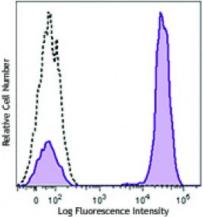
Product Data Sheet

Brilliant Violet 605[™] anti-human CD3

Catalog # / Size:	2324180 / 100 tests 2324175 / 25 tests	
Clone:	SK7	
Isotype:	Mouse lgG1, к	nber
Reactivity:	Human	Al Nu
Preparation:	The antibody was purified by affinity chromatography and conjugated with Brilliant Violet 605 [™] under optimal conditions. The solution is free of unconjugated Brilliant Violet 605 [™] and unconjugated antibody.	Relative Cell N
Formulation:	Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide and BSA (origin USA).	Hu Iyn
Concentration:	Lot-specific	(clo (fill Bri



Human peripheral blood lymphocytes were stained with CD3 (clone SK7) Brilliant Violet 605[™] (filled histogram) or mouse IgG1, κ Brilliant Violet 605[™] isotype control (open histogram).

Applications:

Applications:	Flow Cytometry
Recommended Usage:	Each lot of this antibody is quality control tested by immunofluorescent staining with flow cytometric analysis. For flow cytometric staining, the suggested use of this reagent is \leq 5 microL per million cells or 5 microL per 100 microL of whole blood. It is recommended that the reagent be titrated for optimal performance for each application.
	Brilliant Violet 605 [™] excites at 405 nm and emits at 603 nm. The bandpass filter 610/20 nm is recommended for detection, although filter optimization may be required depending on other fluorophores used. Be sure to verify that your cytometer configuration and software setup are appropriate for detecting this channel. Refer to your instrument manual or manufacturer for support. Brilliant Violet 605 [™] is a trademark of Sirigen Group Ltd.
	This product is subject to proprietary rights of Sirigen Inc. and is made and sold under license from Sirigen Inc. The purchase of this product conveys to the buyer a non-transferable right to use the purchased product for research purposes only. This product may not be resold or incorporated in any manner into another product for resale. Any use for therapeutics or diagnostics is strictly prohibited. This product is covered by U.S. Patent(s), pending patent applications and foreign equivalents.
Application Notes:	Additional reported application (for the relevant formats) include: immunohistochemical staining of frozen tissue sections ^{4,5,8} , immunofluorescent staining ⁶ , and Western blotting3.
Application References:	 Kan EA, <i>et al.</i> 1983. <i>J. Immunol.</i> 131:536. Wood GS, <i>et al.</i> 1985. <i>Am. J. Pathol.</i> 120:371. Van Dongen JJM, <i>et al.</i> 1988. <i>Blood</i> 71:603. (WB) Haringman JJ, <i>et al.</i> 2005. <i>Arthritis Res. Ther.</i> 7:R862. (IHC) Carbone A, <i>et al.</i> 1999. <i>Blood</i> 93:2319. (IHC) Goval JJ, <i>et al.</i> 2006. <i>J. Histochem. Cytochem.</i> 54:75. (IF)

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- Rutjens E, *et al.* 2007. *J. Immunol.* 178:1702.
 Kap Y, *et al.* 2009. *J. Histochem. Cytochem.* 57:1159. (IHC)
 Yoshino N, *et al.* 2000. *Exp. Anim. (Tokyo)* 49:97. (FC)

Description:	CD3 ϵ is a 20 kD chain of the CD3/T-cell receptor (TCR) complex, which is composed of two CD3 ϵ , one CD3 γ , one CD3 δ , one CD3 ζ (CD247), and a T-cell receptor (α/β or γ/δ) heterodimer. It is found on all mature T cells, NK T cells, and some thymocytes. CD3, also known as T3, is a member of the immunoglobulin superfamily that plays a role in antigen recognition, signal transduction, and T cell activation.
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Antigen	1. Barclay N, <i>et al.</i> 1993. The Leucocyte FactsBook. Academic Press. San Diego.
References:	2. Beverly P, et al. 1981. Eur. J. Immunol. 11:329.
	3. Lanier L, <i>et al.</i> 1986. <i>J. Immunol.</i> 137:2501.