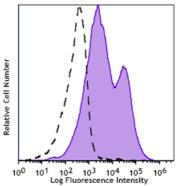
## PE/Fire<sup>™</sup> 640 anti-human CD95 (Fas)

DX2	
Mouse IgG1, к	5
CD95 transfected L cells	qumN
Human, Non-human primate	Relative Cell Number
The antibody was purified by affinity chromatography and conjugated with PE/Fire™ 640 under optimal conditions.	Selati 100
Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide and 0.2% (w/v) BSA (origin USA)	Human p
VI C-64	lymphoc anti-hun DX2) PE
Lot-specific	histogra
	DX2 Mouse IgG1, κ CD95 transfected L cells Human, Non-human primate The antibody was purified by affinity chromatography and conjugated with PE/Fire™ 640 under optimal conditions. Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide and 0.2% (w/v) BSA (origin USA) VI C-64



Human peripheral blood lymphocytes were stained with anti-human CD95 (Fas) (clone DX2) PE/Fire™ 640 (filled histogram), or mouse lgG1, ĸ PE/Fire™ 640 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 5  $\mu$ L per million cells in 100  $\mu$ L staining volume or 5  $\mu$ L per 100  $\mu$ L of whole blood. It is recommended that the reagent be titrated for optimal performance for each application.

\* PE/Fire<sup>™</sup> 640 has a maximum excitation of 566 nm and a maximum emission of 639 nm.

Application Notes: The DX2 antibody is useful for inducing apoptosis of Fas-positive cells. Additional reported applications (for the relevant formats) include: *in vitro* induction of apoptosis<sup>3</sup> (DX2 antibody is required to be cross-linked for effective induction of apoptosis) and immunohistochemical staining<sup>4,5</sup> of acetone-fixed frozen tissue sections and formalin-fixed paraffin-embedded tissue sections. The Ultra-LEAF™ purified antibody (Endotoxin < 0.01 EU/µg, Azide-Free, 0.2 µm filtered) is recommended for functional assays (Cat. No. 305655 and 305656).

**Note:** EOS9.1 antibody can induce apoptosis without cross-linking.

Application References:	<ol> <li>Schlossman S, et al. Eds. 1995. Leucocyte Typing V. Oxford University Press. New York.</li> <li>Kishimoto T, et al. Eds. 1997. Leucocyte Typing VI. Garland Publishing Inc. New York.</li> <li>Cifone M, et al. 1994. J. Exp. Med. 180:1547. (Apop)</li> <li>Zietz C, et al. 2001. Am. J. Pathol. 159:963. (IHC)</li> <li>Sergi C, et al. 2000. Am. J. Pathol. 156:1589. (IHC)</li> <li>Xie S, et al. 2010. J. Immunol. 184:2289. (FC) PubMed</li> <li>Yoshino N, et al. 2000. Exp. Anim. (Tokyo) 49:97. (FC)</li> <li>Sestak K, et al. 2010. PLoS One 5:e9787. (FC)</li> <li>Dixit N, et al. 2012. J. Immunol. 189:5954. PubMed</li> </ol>
Description:	CD95 is a 45 kD single chain type I glycoprotein also known as Fas, APO-1, and TNFRSF6. It is a member of the TNF receptor superfamily. CD95 is expressed on T and B lymphocytes, monocytes, neutrophils, and fibroblasts. CD95 expression is upregulated by activation. The extracellular region of CD95 binds to CD178 (Fas ligand). CD178 binding to CD95 induces apoptosis and has been shown to play a role in the maintenance of peripheral tolerance.

Antigen 1. Krammer P, *et al.* 1994. *Immunol Rev.* 142:175.

References: 2. Nagata S, et al. 1995. Science. 267:1449.