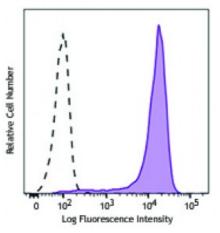
## **Product Data Sheet**

## PE/Dazzle<sup>™</sup> 594 anti-mouse CD326 (Ep-CAM)

Catalog # / Size:	1191175 / 25 μg 1191180 / 100 μg
Clone:	G8.8
Isotype:	Rat IgG2a, к
Immunogen:	TE-71 thymic epithelial cell line
<b>Reactivity:</b>	Mouse
Preparation:	The antibody was purified by affinity chromatography and conjugated with PE/Dazzle <sup>™</sup> 594 under optimal conditions. The solution is free of unconjugated PE/Dazzle <sup>™</sup> 594 and unconjugated antibody.
Formulation:	Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide.
<b>Concentration:</b>	0.2



TE-71 (mouse thymic epithelial stromal cell line) cells were stained with CD326 (clone G8.8) PE/Dazzle<sup>m</sup> 594 (filled histogram) or rat IgG2a,  $\kappa$  PE/Dazzle<sup>m</sup> 594 isotype control (open histogram).

## **Applications:**

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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 $\leq 0.125$ microg per million cells in 100 microL volume. It is recommended that the reagent be titrated for optimal performance for each application.
	* PE/Dazzle™ 594 has a maximum excitation of 566 nm and a maximum emission of 610 nm.
Application Notes:	Additional reported applications for clone G8.8 (for the relevant formats) include: immunohistochemistry of frozen sections: acetone fixed1, with or without OCT embedding <sup>2,4</sup> .
Application References:	<ol> <li>Farr A, <i>et al.</i> 1991. <i>J. Histochem. Cytochem.</i> 39:645. (FC, IHC)</li> <li>Dooley J, <i>et al.</i> 2005. <i>J. Immunol.</i> 175:4331. (FC, IHC)</li> <li>Hinterberger M, <i>et. al.</i> 2010. <i>Nat. Immunol.</i> 11:512. (FC) PubMed</li> <li>Gracz AD, <i>et al.</i> 2010. <i>Am J. Physiol Gastrointest Liver Physiol.</i> 298:590. (IHC)</li> <li>PubMed</li> <li>Nudel I, <i>et al.</i> 2011. <i>J. Immunol.</i> 186:891. PubMed</li> <li>Morimoto H, <i>et al.</i> 2012. <i>Biol Reprod.</i> 86:148. PubMed</li> <li>Takehashi M, <i>et al.</i> 2012. <i>Biol Reprod.</i> 86:178. PubMed</li> <li>Takehashi M, <i>et al.</i> 2013. <i>PLoS One.</i> 8:73270. PubMed</li> <li>Taguchi K, <i>et al.</i> 2014. <i>Mol Cell Biol.</i> 34:900. PubMed</li> <li>Hirokawa Y, <i>et al.</i> 2014. <i>Am J Physiol Gastrointerest Liver Physiol.</i> 306:547. PubMed</li> <li>Ding X, <i>et al.</i> 2015. <i>Cancer Res.</i> 75:330. PubMed</li> </ol>

**Description:** EpCAM (CD326) mediates calcium-independent homophilic cell to cell adhesion. It may also function as a growth factor receptor. It is thought to be involved in

For research use only. Not for diagnostic use. Not for resale. Sony Biotechnology Inc. will not be held responsible for patent infringement or other violations that may occur with the use of our products. Sony Biotechnology Inc. 1730 North First Street, San Jose, CA 95112 www.sonybiotechnology.com maintaining cells in position during proliferation. Expression of EpCAM seems to correlate inversely with the level of E-cadherin (CD324). EpCAM is considered important in tumor biology.

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 1. Borkowski TA, et al. 1996. Eur. J. Immunol. 26:110.

 References:
 2. Bergsagel PL, et al. 1992. J. Immunol. 148:590.