isotype control PE (open

histogram).

PE anti-human CD44 isoform 9 (CD44v9)

Catalog # / Size:	2572015 / 25 tests 2572020 / 100 tests	[]
Clone:	RV3	i 🏊
lsotype:	Rat IgG2a, к	e Cell Number
Immunogen:	Human CD44v8-10 transfected cells	
Reactivity:	Human	
Preparation:	The antibody was purified by affinity chromatography and conjugated with PE under optimal conditions. The solution is free of unconjugated PE and unconjugated antibody.	Log Fluorescence Intensity
Formulation:	Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide and 0.2% (w/v) BSA (origin USA).	PHA-stimulated (3 days) human peripheral blood lymphocytes were stained with CD44 isoform 9 (CD44v9) (clone RV3) PE (filled histogram) or Rat IgG2a, κ
Concentration:	Lot-specific	

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 μ l per million cells in 100 μ l staining volume or 5 μ l per 100 μ l of whole blood.	
Application Notes:	Additional reported applications (for the relevant formats) include: immunohistochemical staining of frozen tissue sections ³ .	
Application References:	 Kakehashi, A., et al. 2016. Cancer Sci. 107: 609. (IHC-P) Aso T, et al. 2015. PLoS One. 10: e0116596. (IHC-P) Yoshikawa M, et al. 2013. Cancer Res. 73: 1855. (IHC-F) Ishimoto T, et al. 2011. Cancer Cell. 387-400. (IHC-P) 	
Description:	CD44 isoform 9, also known as CD44v9, is a CD44 isofom consisting of exons 1-5, 13 (v9) and 15 to 19, product of alternative splicing. CD44v9 is expressed by cancer stem cells, epithelial-type carcinomas, subsets of monocytes and subsets of lymphocytes; have a role in tumor initiation, maintenance and metastasis of cancer cells. CD44v9 interacts with the glutamate-cystine transporter xCT, increasing reduced glutathione and contributing to the ROS resistance of cancer cells, and is a potential predictive marker for recurrence of some cancers and a potential therapeutic target.	
Antigen References:	 Hagiwara M, et al. 2018. BMC Cancer. 18:113 Matsumoto T, et al. 2017. Oncogenesis. 6:397 Bertaux-Skeirik N, et al. 2017. J. Pathol. 242:463 Kodama H, et al. 2017. Br. J. Cancer. 116:186 	

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