

For research use only

## Anti Human CD44 ICD (intracellular domain) Polyclonal Antibody

K0601 Code No. Cancer Category Target CD44

Type Polyclonal Antibody

Concentration 0.25 mg/ml

 $200 \,\mu\,\mathrm{g} \ (800 \,\mu\,\mathrm{L/vial})$ Contents (Volume)

Gene ID 960 **Primary Source** CD44

IN; LHR; MC56; MDU2; MDU3; MIC4; Pgp1; CDW44; CSPG8; HCELL; HUTCH-I; ECMR-III; MGC10468; CD44 Synonyms

Immunogen Partial peptide of Human CD44 (DQFMTADETRNLQNVDMKIGV)

Raised in Rabbit Purification ProteinG Source Rabbit Serum Cross Reactivity Not Tested Label Unlabeled

**Buffer** PBS [containing 2% Block Ace as a stabilizer, 0.1% Proclin as a bacteriostat]

Storage Store below -20°C. Once thawed, store at 4°C. Repeated freeze-thaw cycles should be avoic

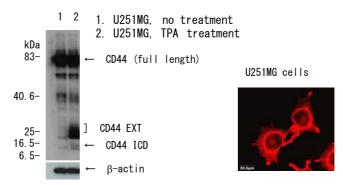
**Application** IHC, ICC, WB, IP, IF

## Recommended Antibody Dilutions

IIn: Diat Commons

ELISA	W B	HС	IC C
NotTested	15–20	100-150	100-150
P	FCM	F	N eutra lization
15-20	NotTested	100-150	NotTested

(μg/m L)



The data is provided from Division of Gene Regulation Institute for Advanced Medical Research School of Medicine, Keio Univ., Japan

Function Recentor for hyaluronic acid (HA) Mediates call-call and call-matrix interactions through its

UniPlot Summary	affinity for HA, and possibly also through its affinity for other ligands such as osteopontin, collagens, and matrix metalloproteinases (MMPs). Adhesion with HA plays an important role in cell migration, tumor growth and progression. Also involved in lymphocyte activation, recirculation and homing, and in hematopoiesis.  Altered expression or dysfunction causes numerous pathogenic phenotypes. Great protein heterogeneity due to numerous alternative splicing and post-translational modification events.  //Subcellular location Membrane: Single-pass type I membrane protein.  //Tissue specificity Isoform 10 (epithelial isoform) is expressed by cells of epithelium and highly expressed by carcinomas. Expression is repressed in neuroblastoma cells.  //Sequence similarities Contains 1 Link domain.
Reference 1)	Okamoto I, et al: Oncogene (1999) 18:1435-1446
2)	Okamoto I, et al: J Biol Chem (1999) 36: 25525-25534
3)	Kawano Y, et al: J Biol Chem (2000) 38: 29628-29635
4)	Okamoto I, et al: J Cell Biol (2001) 155: 755-762
5)	Murakami D, et al., Oncogene (2003) 22:1511-1516
6)	Nagano O, et al., J Cell Biol (2004) 165: 893-902
7)	Takahashi E, et al., J Biol Chem. (2010) 285: 4060-73.





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