

## CAB24866

## **Product Information**

Product SKU:	CAB24866	Gene ID:	14257	Size:	20uL, 100uL
Clone No:	-	Host Species:	Rabbit	<b>Reactivity</b> :	Human,Rat

## **Additional Information**

Observed MW:	Refer to figures	Conjugate:	Unconjugated
Calculated MW:	153kDa	lsotype:	lgG

## **Immunogen Information**

Background:	Defects in FLT4 are the cause of lymphedema hereditary type 1A (LMPH1A) [MIM:153100]; also known as Nonne-Milroy lymphedema or Milroy disease. Hereditary lymphedema is a chronic disabling condition which results in swelling of the autremities due to altered lymphetic flow. Batiants with
	condition which results in swelling of the extremities due to altered lymphatic flow. Patients with lymphedema suffer from recurrent local infections and physical impairment. Note=Defects in FLT4 are found in juvenile hemangioma. Juvenile hemangiomas are the most common
	tumors of infancy, occurring as many as 10% of all births. These benign vascular lesions enlarge rapidly during the first year of life by hyperplasia of endothelial cells and attendant pericytes, and then
Recommended Dilution:	spontaneously involute over a period of years, leaving loose fibrofatty tissue. IF/ICC,1:50 - 1:200
Synonyms:	flt 4; FLT-4; VEGF R3; VEGFR 3; VEGFR-3; FLT4; LMPH1A; PCL; Vegfr3; VEGFR3/Flt-4
Purifcation Method:	Affinity purification
Immunogen:	Recombinant fusion protein containing a sequence corresponding to amino acids 25-229 of mouse VEGFR3/Flt-4(NP_032055.1).
Storage:	Store at -20°C. Avoid freeze / thaw cycles.Buffer: PBS with 0.05% proclin300,50% glycerol,pH7.3.