Elabscience®

ALPL Polyclonal Antibody

catalog number: E-AB-70384

Note: Centrifuge before opening to ensure complete recovery of vial contents.

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Description			
Reactivity	Human;Mouse;Rat		
Immunogen	Recombinant protein corresponding to Mouse Alkaline Phosphatase		
Host	Rabbit		
Isotype	IgG		
Purification	Affinity purification		
Conjugation	Unconjugated		
Buffer	Phosphate buffered solution, pH 7.4, containing 0.05% stabilizer, 1% protein		
	protectant and 50% glycerol.		
Applications	Recommended Dilution		
WB	1:500-1:2000		
Data			
72KDa - 55KDa - 72KDa - 72KDa - 43KDa - 34KDa - 34KDa -		72KDa- 55KDa- 43KDa- 34KDa-	
Western Blot analysis of various samples using ALPL		Western Blot analysis of various samples using ALPL	
Polyclonal Antibody at dilution of 1:1000.		Polyclonal Antibody at dilution of 1:1000.	
Observed-MW:55 kDa		Observed-MW:55 kDa	
Calculated-MW:57 kDa		Calculated-MW:57 kDa	
Preparation & Storage			
64	Stars at 2000 Mal 10 12		

Storage	Store at -20°C Valid for 12 months. Avoid freeze / thaw cycles.	
Shipping	The product is shipped with ice pack,upon receipt, store it immediately at the	
	temperature recommended.	

Background

There are at least four distinct but related alkaline phosphatases: intestinal, placental, placental-like, and liver/bone/ kidney (tissue non-specific). The first three are located together on chromosome 2, while the tissue non-specific form is located on chromosome 1. The product of this gene is a membrane bound glycosylated enzyme that is not expressed in any particular tissue and is, therefore, referred to as the tissue-nonspecific form of the enzyme. The exact physiological function of the alkaline phosphatases is not known. A proposed function of this form of the enzyme is matrix mineralization; however, mice that lack a functional form of this enzyme show normal skeletal development. This enzyme has been linked directly to hypophosphatasia, a disorder that is characterized by hypercalcemia and includes skeletal defects. The character of this disorder can vary, however, depending on the specific mutation since this determines age of onset and severity of symptoms. Alternatively spliced transcript variants have been described.