



Sheep anti Human Fibrinopeptide A polyclonal antibody (CABT-L400)

This product is for research use only and is not intended for diagnostic use.

PRODUCT INFORMATION

Specificity	This antibody is specific for fibrinopeptide A as demonstrated by immunoelectrophoresis and ELISA.
Target	Fibrinopeptide A
Immunogen	Synthetic fibrinopeptide Aα 1-16 conjugated to carrier.
Isotype	IgG
Source/Host	Sheep
Species Reactivity	Human
Purification	Affinity purified
Conjugate	Unconjugated
Applications	Suitable for use in IEP, ELISA. Each laboratory should determine an optimum working titer for use in its particular application. Other applications have not been tested but use in such assays should not necessarily be excluded.
Format	Liquid
Size	0.5 mg
Buffer	10 mM HEPES, pH 7.4, 150 mM NaCl, 50% (v/v) glycerol.
Preservative	None
Storage	Store between -10 and -20°C. Product will become viscous but will not freeze. Avoid storage in

frost-free freezers. Keep vial tightly capped. Allow product to warm to room temperature and gently mix before use.

BACKGROUND

Introduction

Human fibrinogen is a 340 kDa plasma protein produced in the liver. Plasma concentrations are typically 1.7-3.5 g/L (5-12 μ M). The intact fibrinogen molecule consists of two identical subunits, each consisting of three non-identical polypeptide chains ded as A α , B β and γ . The letters A and B in the A α and B β chains designate, respectively, fibrinopeptide A (FpA, residues 1-16), and fibrinopeptide B (FpB, residues 1-14), which are cleaved by thrombin upon conversion of fibrinogen to fibrin. The fibrin monomers polymerize in a half-overlap fashion to form insoluble fibrin fibrils. The polymerised fibrin is subsequently stabilized by activated Factor XIII that forms amide linkages between γ chains and, to a lesser extent, α chains of the fibrin molecules. Proteolysis of fibrinogen by plasmin initially liberates C-terminal residues from the A α chain to produce fragment X (intact D-E-D, which is still clottable). Fragment X is further degraded to nonclottable fragments Y (D-E) and D. Fragment Y can be digested into its constituent D and E fragments. Proteolysis of crosslinked fibrin by plasmin results in fragment DD (D-Dimer consisting of the D domains of 2 fibrin molecules crosslinked via the γ chains), fragment E (central E domain) as well as DDE in which fragment E is noncovalently associated with DD. The molecular weights of the cleavage fragments produced from human crosslinked fibrin are: 184 kDa for fragment DD, 92 kDa for D, 50 kDa for E, 1.54 kDa for FpA and 1.57 kDa for FpB. Most of the fibrinogen in the circulation consists of 2 copies of each chain (A α 2, B β 2, γ A2), but in normal plasma approximately 10% of the fibrinogen molecules contain one γ A chain and one variant γ chain (termed γ'), in which the c-terminal AGDV residues are replaced with the amino acid sequence VRPEHPAETEDSLYPEDDL. This variant fibrinogen is commonly referred to as fibrinogen gamma prime (γ A/ γ') but has also been called fibrinogen 2 or peak 2 fibrinogen because it elutes separately from fibrinogen 1 (γ A2) by ion exchange chromatography. Residues 414-427 of the γ' chain of fibrin gamma prime (contain a high-affinity binding site for exosite II of thrombin, which allows the active site of bound thrombin to remain available to interact with substrates while demonstrating resistance to heparin mediated inhibition by antithrombin.

Keywords

fibrinogen alpha chain;FGA;Fib2;fibrinogen, A alpha polypeptide;

GENE INFORMATION

Entrez Gene ID

[2243](#)

UniProt ID

[P02671](#)