



Sheep anti Human Fibrinogen polyclonal antibody (CABT-L391)

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

PRODUCT INFORMATION

| Specificity | This antibody is specific for Fibrinogen as demonstrated by immunoelectrophoresis and ELISA. |
|--------------------|--|
| Target | Fibrinogen |
| Immunogen | Human fibrinogen purified from plasma. |
| Isotype | IgG |
| Source/Host | Sheep |
| Species Reactivity | Human |
| Purification | Affinity purified |
| Conjugate | Unconjugated |
| Applications | IEP, ELISA |
| 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

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Introduction

Fibrinogen is an abundant plasma protein (5-10 uM) produced in the liver. The intact protein has a molecular weight of 340 kDa and is composed of 3 pairs of disulphide-bound polypeptide chains named $A\alpha$, $B\beta$ and γ . Fibrinogen is a triglobular protein consisting of a central E domain and terminal D domains. Proteolysis by thrombin results in release of Fibrinopeptide A (FPA, A α1-16) followed by Fibrinopeptide B (FPB, Bβ1-14) and the fibrin monomers that result polymerize in a half-overlap fashion to form insoluble fibrin fibrils. The chains of fibrin are referred to as α , β and γ , due to the removal of FPA and FPB. The polymerised fibrin is subsequently stabilized by the transglutaminase 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 Aα 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. Digestion of non-crosslinked fibrin with plasmin is very similar to the digestion of fibrinogen, which results in production of fragments D and E. Degradation 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 non-covalently associated with DD. For human crosslinked fibrin, the relative weights of the cleavage fragments produced 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.

Keywords

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

GENE INFORMATION

Entrez Gene ID 2243

UniProt ID P02671

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