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ProductInformation

IMMUNOFILE

Monoclonal Anti Protein Kinase Cβ₂ Mouse Ascites Fluid Clone PK-B26

Product Number P 2584

TECHNICAL BULLETIN

Product Description

Monoclonal Anti Protein Kinase $C\beta_2$ (mouse IgG_1 isotype) is derived from the PK-B26 hybridoma produced by the fusion of mouse myeloma cells and splenocytes from BALB/c mice immunized with a synthetic peptide (Cys)-Ser-Phe-Val-Asn-Ser-Glu-Phe-Leu-Lys-Pro-Glu-Val-Lys-Ser, corresponding to the C-terminal variable (V5) region (amino acids 660-673) of PKC β_2 , conjugated to maleimido activated KLH (mKLH) as carrier protein. The isotype is determined using Sigma ImmunoType Kit (Product Code ISO-1) and by a double diffusion immunoassay using Mouse Monoclonal Antibody Isotyping Reagents (Product Code ISO-2). The product is provided as ascites fluid with 0.1% sodium azide as a preservative.

Protein kinase C (PKC, 77-90 kDa), is a family of homologous serine-threonine protein kinases, which are key regulatory enzymes in signal transduction, cellular regulation, tumor promotion and oncogenesis. PKC is a calcium-dependent and phospholipid-dependent enzyme that is activated *in vivo* by the lipid diacylglycerol, produced in response to a variety of hormones and growth factors. PKC consists of a single polypeptide chain, containing four conserved regions and five variable regions. Sequence information defined a putative domain structure for the enzyme which can be divided into an amino-terminal regulator and a carboxyterminal catalytic domain joined by a hinge region.

Proteolysis of purified native PKC by trypsin yields two major fragments, representing the regulatory and the kinase domains of the enzyme, due to cleavage in a proposed hinge region between residue 292 and residue 317.4 There is evidence that in vivo, agonist-induced generation of a catalytic fragment of the enzyme may occur as well. The PKC family of isoenzymes can be subdivided into two major classes; conventional (c) isoforms (α , β_1 , β_2 and γ), which are Ca²⁺ and phospholipid-dependent kinases, and novel (n) isoforms $(\delta, \epsilon, \zeta, \eta \text{ and } \Theta)$ that are Ca²⁺-independent, phospholipidstimulated kinases.⁵ PKC is widely distributed in all tissues and cells. The majority of cells coexpress multiple PKC isoforms, indicating a specific function for each isoform in the cell. PKCβ₁ and PKCβ₂ isoforms are encoded by the same gene but diverge at the C-terminal (V5) region as a result of differential mRNA splicing. The $PKC\beta_1/\beta_2$ isoenzymes appear to be widely expressed, in the brain, lung, liver, spleen, thymus, skeletal muscle and skin but not in kidney, rat and mouse fibroblasts.6 PKCβ₂ is also reported to be expressed in a wider variety of tissues and cell lines and in higher levels than PKCβ₁.6 Antibodies that react specifically with PKC isoenzymes are useful for the study of the differential tissue expression, intracellular and subcellular distribution, of these isoenzymes. Furthermore, they also allow the detection and localization of PKC in normal and malignant tissues. The monoclonal nature of the product guarantees the continuous production of a constant titer of Anti Protein Kinase Cβ₂ antibody with the same specificity and chemical identity.

Monoclonal Anti Protein Kinase $C\beta_2$ is a homogenous population of antibody molecules which may be used for the localization of Protein Kinase $C\beta_2$ using various immunochemical assays such as ELISA, immunoblot and dot blot.

Storage/Stability

For continuous use, store at 0-5 °C. For extended storage freeze in working aliquots. Repeated freezing and thawing is not recommended. Storage in "frost-free" freezers is not recommended. If slight turbidity occurs upon prolonged storage, clarify the solution by centrifugation before use.

Specificity

Monoclonal Anti Protein Kinase $C\beta_2$ recognizes an epitope located within the amino acid residues 660-673 at the C-terminal variable (V5) region of PKC β_2 . It reacts with the 80 kDa polypeptide of PKC, applying the immunoblotting technique, using rat brain extract. The product reacts in dot-blot immunobinding and in ELISA with the PKC β_2 peptide conjugated to BSA with 1-ethyl-3-(3-dimethylaminopropyl)-carbodiimide (EDCI). The antibody shows no cross-reactivity with PKC peptides corresponding to C-terminal sequences from PKC β_1 (658-671) and PKC γ (684-697) conjugated to BSA with EDCI.

References

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