abcam

Product datasheet

Neuraminidase Activity Assay Kit (Fluorometric) ab 185436

3 Images

Overview

Product name Neuraminidase Activity Assay Kit (Fluorometric)

Detection method Fluorescent

Sample type Serum, Tissue, Adherent cells, Suspension cells

Assay type Enzyme activity (quantitative)

Sensitivity < 2 mU/ml

Species reactivity Reacts with: Mammals, Other species

Product overview Abcam's Neuraminidase Activity Assay Kit (Fluorometric) (ab185436) provides a simple and

sensitive method for measuring neuraminidase (NA) activity using fluorescence (Ex/Em = 530/590 nm). The assay utilizes NA Probe to detect the neuraminidase activity. This high-throughput adaptable assay kit can detect NA activity as low as 2.0 mU/mL in a variety of

samples.

NotesThis product is manufactured by BioVision, an Abcam company and was previously called K732

Neuraminidase Activity Fluorometric Assay Kit. K732-100 is the same size as the 100 test size of

ab185436.

Neuraminidase (NA) is a very common enzyme that hydrolyzes terminal sialic acid residues on polysaccharide chains; most often a galactose residue. NA activity plays a key role in the invasion

of target cells and the replication of influenza virus. NA activity also assists in the elution of progeny viruses from infected cells, and prevents self-aggregation of virus. Thus, NA is an

important target for drug development.

Platform Microplate reader

Properties

Storage instructions Store at -20°C. Please refer to protocols.

| Galactose Standard (100nmol/ul) 1 x 0.1ml | Components | 100 tests |
|---|---------------------------------|-----------|
| | Galactose Standard (100nmol/ul) | 1 x 0.1ml |
| NA Assay Buffer 1 x 30ml | NA Assay Buffer | 1 x 30ml |

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| Components | 100 tests |
|---------------------|-----------|
| NA Enzyme Mix I | 1 vial |
| NA Enzyme Mix II | 1 vial |
| NA Positive Control | 1 x 0.1ml |
| NA Probe | 1 x 0.2ml |
| NA Substrate | 1 vial |

Function Catalyzes the removal of sialic acid (N-acetylneuramic acid) moities from glycoproteins and

glycolipids. To be active, it is strictly dependent on its presence in the multienzyme complex.

Appears to have a preference for alpha 2-3 and alpha 2-6 sialyl linkage.

Tissue specificity Highly expressed in pancreas, followed by skeletal muscle, kidney, placenta, heart, lung and liver.

Weakly expressed in brain.

Involvement in disease Defects in NEU1 are the cause of sialidosis (SIALIDOSIS) [MIM:256550]. It is a lysosomal

storage disease occurring as two types with various manifestations. Type 1 sialidosis (cherry red spot-myoclonus syndrome or normosomatic type) is late-onset and it is characterized by the formation of cherry red macular spots in childhood, progressive debilitating myoclonus, insiduous visual loss and rarely ataxia. The diagnosis can be confirmed by the screening of the urine for sialyloligosaccharides. Type 2 sialidosis (also known as dysmorphic type) occurs as several variants of increasing severity with earlier age of onset. It is characterized by the presence of abnormal somatic features including coarse facies and dysostosis multiplex, vertebral

deformities, mental retardation, cherry-red spot/myoclonus, sialuria, cytoplasmic vacuolation of

peripheral lymphocytes, bone marrow cells and conjunctival epithelial cells.

Sequence similarities Belongs to the glycosyl hydrolase 33 family.

Contains 4 BNR repeats.

Domain A C-terminal internalization signal (YGTL) appears to allow the targeting of plasma membrane

proteins to endosomes.

Post-translational N-glycosylated.

modifications Phosphorylation of tyrosine within the internalization signal results in inhibition of sialidase

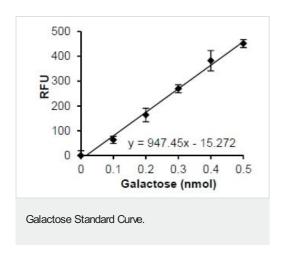
internalization and blockage on the plasma membrane.

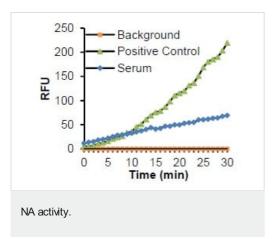
Cellular localization Lysosome membrane. Lysosome lumen. Cell membrane. Cytoplasmic vesicle. Localized not only

on the inner side of the lysosomal membrane and in the lysosomal lumen, but also on the plasma

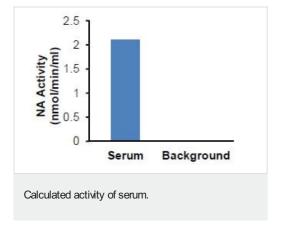
membrane and in intracellular vesicles.

Images





NA activity in normal Human serum (1 µl) and positive control (1 µl).



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