

## **α(1-2) Fucosidase from Xanthomonas manihotis, Recombinant**

Cat. No. NATE-1258

Lot. No. (See product label)

### **Introduction**

#### **Description**

Alpha-Fucosidase is an enzyme that breaks down fucose. Fucosidosis is an autosomal recessive lysosomal storage disease caused by defective alpha-L-fucosidase with accumulation of fucose in the tissues. Different phenotypes include clinical features such as neurologic deterioration, growth retardation, visceromegaly, and seizures in a severe early form; coarse facial features, angiokeratoma corporis diffusum, spasticity and delayed psychomotor development in a longer surviving form; and an unusual spondylometaphyseal dysplasia in yet another form.

#### **Synonyms**

α-L-fucoside fucosidase; Alpha-Fucosidase; FUCA1; FUCA; EC 3.2.1.51

### **Product Information**

#### **Species**

Xanthomonas manihotis

#### **Source**

E. coli

#### **Molecular Weight**

70000 daltons

#### **Concentration**

20,000 units/ml

#### **Unit Definition**

One unit is defined as the amount of enzyme required to cleave > 95% of the α-L-fucose from 1 nmol of Fucα1-2Galβ1-4Glc-7-amino-4-methyl-coumarin (AMC), in 1 hour at 37°C in a total reaction volume of 10 μl.

### **Storage and Shipping Information**

#### **Storage**

4°C