

## Native *Xanthomonas* sp. $\alpha$ -1 $\rightarrow$ (2,3,4) Fucosidase solution

Cat. No. NATE-0262

Lot. No. (See product label)

### Introduction

**Description** Tissue alpha-L-fucosidase is an enzyme that in humans is encoded by the FUCA1 gene. 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**  $\alpha$ -1 $\rightarrow$  (2,3,4) Fucosidase; alpha-L-fucosidase; Alpha-Fucosidase; FUCA1; FUCA; EC 3.2.1.51; 9037-65-4

### Product Information

**Source** *Xanthomonas* sp.

**Form** buffered aqueous solution, Solution in 20 mM Tris-HCl, pH 7.5, 25 mM NaCl

**EC Number** EC 3.2.1.51

**CAS No.** 9037-65-4

**Activity** > 0.5 units/mg protein

**Unit Definition** One unit will hydrolyze 1  $\mu$ mole fucose from 3-fucosyllactose per min at pH 5.0 at 37°C.

### Usage and Packaging

**Package** vial of 0.004 unit

### Storage and Shipping Information

**Storage** 2-8°C