

Native Xanthomonas sp. α -1 \rightarrow (3,4) Fucosidase solution

Cat. No. NATE-0263 Lot. No. (See product label)

| Introduction | |
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| 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 spondylometaphyseoepiphyseal dysplasia in yet another form. |
| Synonyms | α -1→ (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 | > 2 units/mg protein |
| Concentration | > 0.5 unit/mL |
| Unit Definition | One unit will release 1.0 μ mole of fucose from Lewis X trisaccharide, 4-methylumbelliferyl glycoside per min at pH 5.0 at 37°C. |
| Usage and Packaging | |
| Package | vial of 0.02 unit |
| Storage and Shipping Information | |
| Storage | 2-8°C |
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