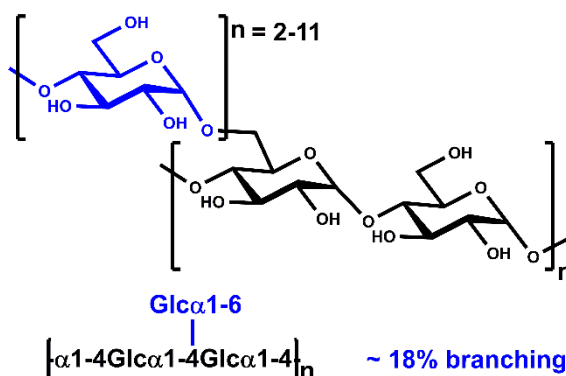


GLYCOGEN (ALGAE) (Lot 210101)

CAT. NO: P-GLYAL
CAS: 9005-79-2
Source: *Galdieria sulphuraria*

02/21

STRUCTURE



Schematic representation of algae glycogen subunit composed of α -(1,4) linked glucose monosaccharide backbone with α -(1,6) branches of α -(1,4) linked glucose monosaccharides.

DESCRIPTION

Algae glycogen is a branched polysaccharide containing a Glc- α -1,4-linked backbone. Branching points are connected via a α -1,6 linkage and side-chains are composed of Glc- α -1,4-linked oligosaccharides. Its structure is similar to that of glycogen from other sources (e.g. oyster), however, algae glycogen has higher degree of branching and shorter side chains.¹

PROPERTIES

| | |
|------------------------------|---|
| Purity: | > 77% (Enzyme incubation using amyloglucosidase, alpha-amylase and isoamylase followed by glucose quantitation) |
| Sugar Composition: | Glucose = 100 |
| Protein: | 1.5% |
| Ash: | 0.08% |
| Moisture: | 5.15% |
| Physical Description: | White/Off-white/Slightly off-white, odourless powder |
| Solubility: | > 150 mg/mL |

¹ Martinez-Garcia, M., Stuart, M. C. A. & van der Maarel, M. J. E. C. (2016). *Int. J. Biol. Macromol.*, **89**, 12-18.

STORAGE CONDITIONS

Store dry at room temperature in a well-sealed container. Under these conditions, the product is stable for several years.

HPAEC-PAD

Column: CarboPac PA200 guard and analytical columns (3 × 250 mm)

Temperature: 30°C

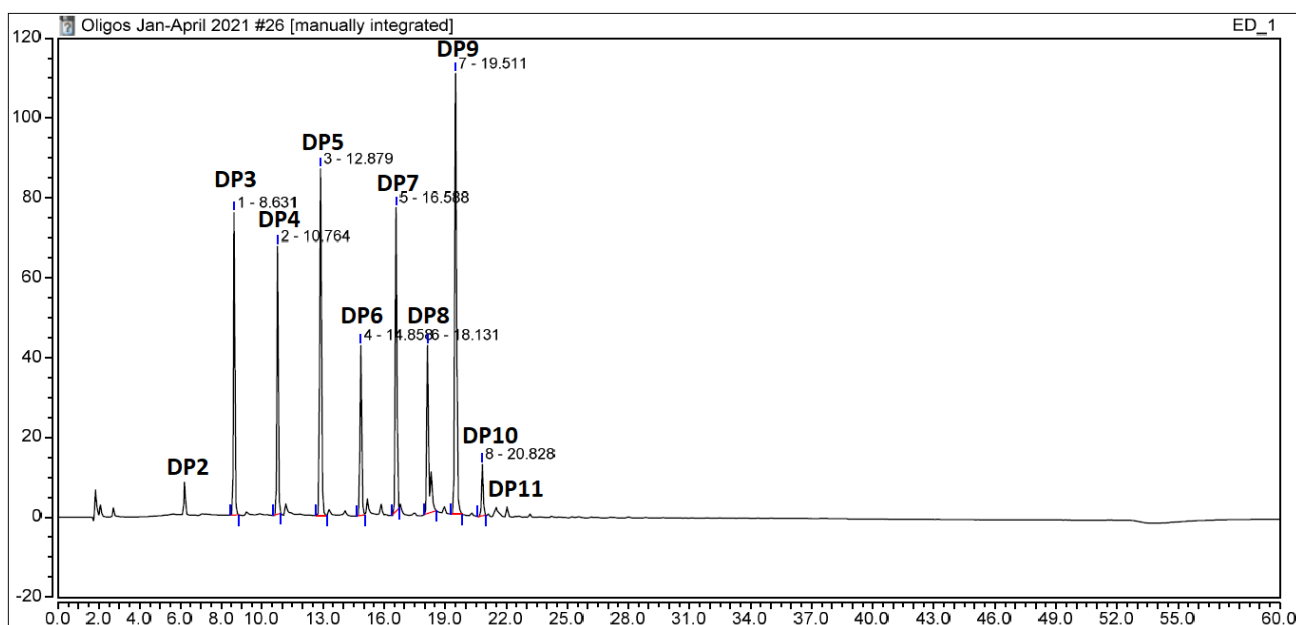
Detector: Au electrode; waveform Carbohydrate, standard quad

Flow rate: 0.5 mL/min

IC system: Dionex ICS5000 + DP system and Chromeleon 7 software

A stepwise linear gradient method was employed as shown.

| Time (min) | 100 mM NaOH (%) | 320 mM NaOAc (%) in 100 mM NaOH |
|------------|-----------------|---------------------------------|
| 0 | 100 | 0 |
| 40 | 0 | 100 |
| 50 | 0 | 100 |
| 50.5 | 100 | 0 |
| 60 | 100 | 0 |



HPAEC-PAD chromatogram showing algae glycogen side-chain composition after enzymatic hydrolysis with a purified debranching enzyme (i.e. isoamylase).

Enzymatic hydrolysis conditions:

10 mg of **P-GLYAL** was incubated with 6 µL of isoamylase **E-ISAMYHP** (Megazyme) in 1 mL of 0.1 M pH 4.0 Sodium Acetate buffer at 40°C for 90 min. Sample was then centrifuged and supernatant injected on the HPAEC-PAD system.