

CERTIFICATE OF ANALYSIS

PRODUCT NAME: GLYKO® ASIALO-, AGALACTO- TETRAANTENNARY COMPLEX

N-GLYCAN (NGA4)

PRODUCT CODE: GKC-006300

LOT NUMBER: DP15H0703a

PACK SIZE: 10 µg (qualitative standard for glycan identification)

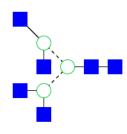
PURITY: ≥90% of glycan by UPLC®

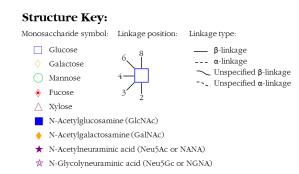
FORM: Dry solid

STORAGE: Store at -20°C before and after reconstitution

EXPIRATION: January 2022, may be used for 1 year after reconstitution

STRUCTURE 1,2,3 :





Quality Control:

Sample Preparation: NGA4 was labeled with 2-aminobenzamide (2-AB) by reductive amination⁴ using the Signal[™] 2-AB Labeling Kit (product code GKK-404).

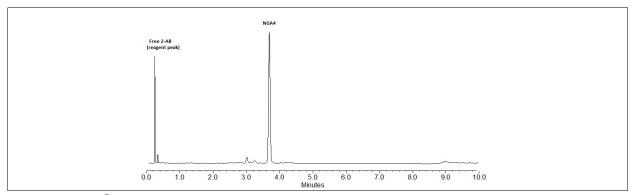


Figure 1 - UPLC® Results: 3-6 pmol (1 μ l) of the 2-AB-labeled glycan was injected on a Waters ACQUITY UPLC® H Class System utilizing a 10-minute method under the conditions below:

Time (min)	Flow (ml/min)	% ACN	% Buffer
0	1.0	75	25
8.0	1.0	60	40
8.1	0.5	40	60
8.5	0.5	40	60
8.6	1.0	40	60
8.8	1.0	75	25
10.0	1.0	75	25

Waters ACQUITY UPLC BEH Glycan Column (1.7 μ m, 2.1 x 100 mm) Column:

ACN: Acetonitrile

Buffer: 100 mM ammonium formate, pH 4.4

Flow rate: As stated in table, in ml/min

60°C Temperature: 15,000 psi Max Pressure:

Fluorescence Detection: $\lambda_{ex} = 330 \text{ nm}$ $\lambda_{em} = 420 \text{ nm}$

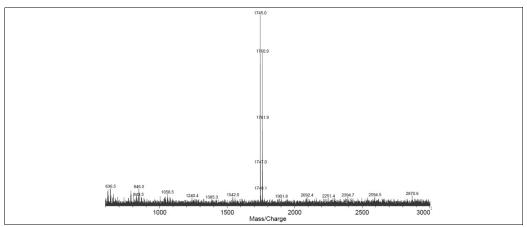


Figure 2 - Mass Spectrum of NGA4 [M + Na]⁺

Average Mass⁵: 1723.6

Monoisotopic Mass⁵: 1722.6454

Structural Analysis: The purity and structural integrity of the glycan was assessed by UPLC⁶ (as described above) and MALDI-TOF^{7,8} or LC-MS. Agreement was found between the results from mass spectrometry and UPLC.

Applications:

- qualitative standard for various analytical procedures
- fluorescent-labeling or formation of a variety of oligosaccharide derivatives
- substrate for glycosidase and glycosyl transferase assays

Reconstitution: Use HPLC-grade water or an aqueous buffer to dissolve the glycan. Store the reconstituted glycan at -20°C in working aliquots. Avoid multiple freeze/thaw cycles.

Handling: The oligosaccharide is shipped as a dried solid. Allow the unopened vial to reach ambient temperature and tap on a solid surface to ensure that most of the material is at the bottom of the vial. Gently remove the cap, add the desired volume of water or buffer, re-cap and mix thoroughly to redissolve all the oligosaccharide. For maximal recovery, ensure that the cap lining is also rinsed, and centrifuge the reconstituted vial briefly before use.

Make sure that any glassware, plasticware, solvents or reagents which come into contact with the glycan are free of glycosidases and carbohydrate contaminants.

Minimize exposure to elevated temperatures or extremes of pH.

REFERENCES

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- Average mass and monoisotopic mass were calculated using the ExPASy GlycanMass calculator: http://web.expasy.org/glycanmass/
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- James DC and Jenkins N. Analysis of N-glycans by matrix-assisted laser desorption/ionization mass spectrometry. In: Jackson P, Gallagher JT, editors. A laboratory guide to glycoconjugate analysis, BioMethods Vol. 9. Basel: Birkhäuser; 1997. p. 91-112.
- 8. Papac DI, Wong A and Jones AJS. Analysis of acidic oligosaccharides by matrix-assisted laser desorption/ionization time of flight mass spectrometry. Anal Chem 1996 Sep 15;68(18):3215-3223.