

CERTIFICATE OF ANALYSIS

PRODUCT NAME: GLYKO® ASIALO, MONOGALACTOSYLATED, BIANTENNARY CORE-SUBSITUTED

WITH FUCOSE COMPLEX N-GLYCAN (NA2G1F or G1F)

PRODUCT CODE: GKC-014301

LOT NUMBER: DP19D1702a

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

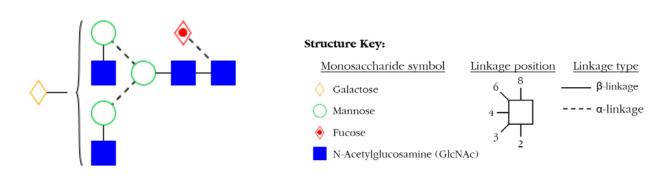
PURITY: ≥90% of glycan by UPLC®

FORM: Dry solid

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

EXPIRATION: May 2024, may be used for 1 year after reconstitution

STRUCTURE^{1,2,3}:



Quality Control:

Sample Preparation: NA2G1F was labeled with 2-aminobenzamide (2-AB) by reductive amination⁴ using the Signal[™] 2-AB Labeling Kit (product code GKK-404).

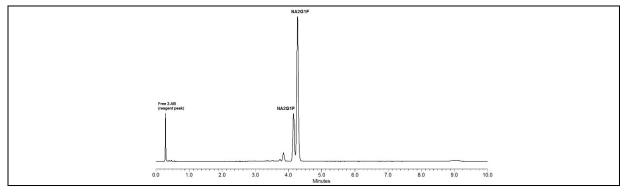


Figure 1 - **UPLC**[®] **Results**: 3 - 6 pmol (1 μl, aqueous) of the 2-AB-labeled⁴ glycan was injected on a Waters ACQUITY UPLC[®] H Class System utilizing a 10-minute method under the conditions below:

Time (min)	Flow (ml/min)	%ACN	%Buffer
0.00	1.0	75.0	25.0
8.0	1.0	60.0	40.0
8.1	0.5	40.0	60.0
8.5	0.5	40.0	60.0
8.6	1.0	40.0	60.0
8.8	1.0	75.0	25.0
10.0	1.0	75.0	25.0

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

ACN: Acetonitrile

Buffer: 100 mM ammonium formate, pH 4.4

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

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

Fluorescence Detection: λ_{ex} = 330 nm, λ_{em} = 420 nm

Average Mass⁵: 1625.5

Monoisotopic Mass⁵: 1624.5973

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.

Application:

- Qualitative standard for various analytical procedures
- Fluorescent-labeling or formation of a variety of oligosaccharide derivatives
- Substrate for glycosidase and glycosyl transferase assays

Handling & Reconstitution:

The oligosaccharide library is shipped as a dried solid. Use ultra-pure water or an aqueous buffer to dissolve the materials.

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 ultra-pure water or aqueous buffer, re-cap and mix thoroughly to redissolve all the material.

For maximal recovery, ensure that the cap lining is also rinsed. Centrifuge the reconstituted vial briefly before use.

Make sure that any glassware, plasticware, solvents or reagents used are free of glycosidases and carbohydrate contaminants.

Minimize exposure to elevated temperatures or extremes of pH. Store the reconstituted glycan library at -20° C. Allow the vial to equilibrate to ambient temperature before use.

REFERENCES

- Ceroni A, Maass K, Geyer H, Geyer R, Dell A, Haslam SM. GlycoWorkbench: a tool for the computer-assisted annotation of mass spectra of glycans. J Proteome Res. 2008 Apr; 7(4): 1650-9.
- Harvey DJ, Merry AH, Royle L, Campbell MP, Dwek RA, Rudd PM. Proposal for a standard system for drawing structural diagrams of N– and O-linked carbohydrates and related compounds. Proteomics 2009 Aug; 9(15): 3796-801.
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- Bigge JC, Patel T, Bruce JA, Goulding PN, Charles SM, Parekh RB. Nonselective and efficient fluorescent labeling of glycans using 2-amino benzamide and anthranilic acid. Anal Biochem 1995 Sep 20;230(2):229-238.
- 5. Average mass and monoisotopic mass of the glycan were calculated using the ExPASy GlycanMass calculator:

http://web.expasy.org/glycanmass/

- Ahn J, Bones J, Yu YQ, Rudd PM, Gilar M. Separation of 2aminobenzamide labeled glycans using hydrophilic interaction chromatography columns packed with 1.7 microm sorbent. J Chromatogr B Analyt Technol Biomed Life Sci. 2010 Feb 1; 878(3-4): 403-8.
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Authorized Signature