

X-ray Diffraction Analysis of the O(H) Blood-group-specific *Ulex europaeus* Lectin I

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Introduction

The first lectin from *Ulex europaeus* (UE-I) has been shown to bind the human H-type 2 blood group determinant [1]. The H-type 2 blood group determinant is the trisaccharide [α -L-Fuc α (1 \rightarrow 2)- β -D-Gal β (1 \rightarrow 4)- β -D-GlcNAc α -], and it is the antigenic determinant present on O-type erythrocytes [2].

Methods and Materials

As part of the examination into the structural nature of the recognition of the H-type 2 blood group determinant by UE-I, crystals of UE-I in complex with the methylglycoside of the H-type 2 trisaccharide (H-type 2-OMe) were subjected to x-ray diffraction analysis. Diffraction data were collected on beamline BM-14-C of the Bio Consortium for Advanced Radiation Sources (BioCARS) facility at sector 14 of the APS. The UE-I:H-type 2-OMe complex crystallizes in the orthorhombic space group C222₁, with unit cell dimensions $a = 88.80 \text{ \AA}$, $b = 164.75 \text{ \AA}$, and $c = 77.42 \text{ \AA}$, and a single UE-I dimer is present within the asymmetric unit. Diffraction data were collected to 2.3- \AA resolution; however, the data were truncated to 3.0- \AA resolution.

Results and Discussion

The preliminary structure of the UE-I:H-type 2-OMe complex at 3.0- \AA resolution has contributed to the understanding of the critical protein:carbohydrate

interactions that occur between UE-I and the H-type 2 blood group determinant during O-type erythrocyte recognition and binding.

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References

- [1] W. C. Boyd and E. Shapleigh, *Blood* **9**, 1195-1198 (1954).
- [2] W. M. Watkins, in *Glycoproteins. Their Composition, Structure and Function*, edited by A. Gottschalk (Elsevier, Amsterdam, 1972), pp. 830-891.
- [3] G. F. Audette, Ph. D. thesis, University of Saskatchewan, Saskatoon, Canada, 2002.
- [4] G. F. Audette, D. J. H. Olson, A. R. S. Ross, J. W. Quail, and L. T. J. Delbaere, *Can. J Chem.* (submitted).