

# Synthesis of Glycoconjugate Vaccines against *Candida albicans* Using the Novel Linker Methodology

Xiangyang Wu, J. James Bailey, Eugenia Paszkiewicz and David R. Bundle\*

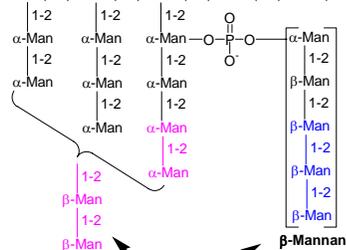
Department of Chemistry, University of Alberta, Edmonton, Alberta, Canada, T6G 2G2



## Introduction

The cell wall phosphomannan of *Candida* species is a glycoprotein containing predominantly  $\alpha$ -linked mannose residues. However, it is the minor  $\beta$ -mannan component of the phosphomannan of clinically important *Candida* strains that provides immunological protection in animal models of fungal disease and hence holds promise as a component of conjugate vaccines. This important antigen occurs in different forms linked to the  $\alpha$ -mannan backbone via a phosphodiester bond (acid labile  $\beta$ -mannan) or directly via a glycosidic bond.<sup>1</sup> Evidence from immunochemistry and solution properties of this antigen implied that (1 $\rightarrow$ 2)- $\beta$ -mannan oligomers have potential as the key epitope of conjugate vaccines.<sup>2</sup>

$\alpha$ -Man(1-6) $\alpha$ -Man(1-6) $\alpha$ -Man(1-6) $\alpha$ -Man(1-6) $\alpha$ -Man(1-6)-inner core-Asn



At least two forms of the  $\beta$ -mannan

Here, we report gram scale syntheses of the both forms of the  $\beta$ -mannan antigen.

## References

- Shibata, N.; Fukasawa, S.; Kobayashi, H.; Tojo, M.; Yonezu, T.; Ambo, A.; Ohkubo, Y.; Suzuki, S. *Carbohydr. Res.* **1989**, 187, 239.
- Nitz, M.; Ling, C. C.; Otter, A.; Cutler, J. E.; Bundle, D. R. *J. Biol. Chem.* **2002**, 277, 3440.

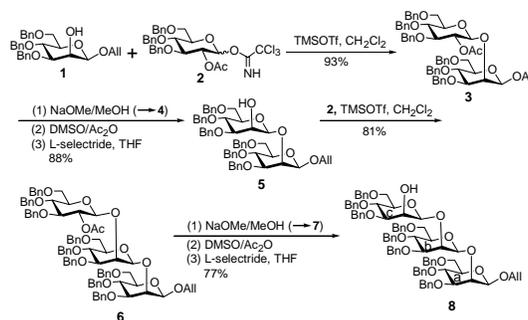
## Summary

Compounds that are derivatives of the unique (1 $\rightarrow$ 2)- $\beta$ -D-mannopyranan found in the cell wall of *C. albicans* have been synthesized on a multi-gram scale employing the C2 inversion methodology. Coupling larger oligosaccharides to BSA or TT using the linear homobifunctional linker with high efficiency under very mild conditions was confirmed. Glycoconjugates **26** and **27** are being evaluated for their efficiency as anti-*C. albicans* vaccine in rabbit and mice. (see poster # 44 by Tomasz Lipinski *et al.*)

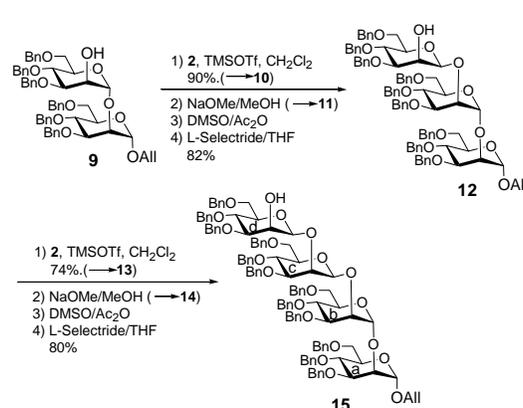
## Preparation of 5, 8 and 15.

The glucosyl trichloroacetimidate donor **2** was employed to establish a  $\beta$ -glucopyranosyl linkage. Subsequent *Swern* oxidation and selective reduction facilitated an efficient approach to the  $\beta$ -mannopyranosides **5**, **8** and **15** on a multi-gram scale.

### Synthesis of acid labile epitope



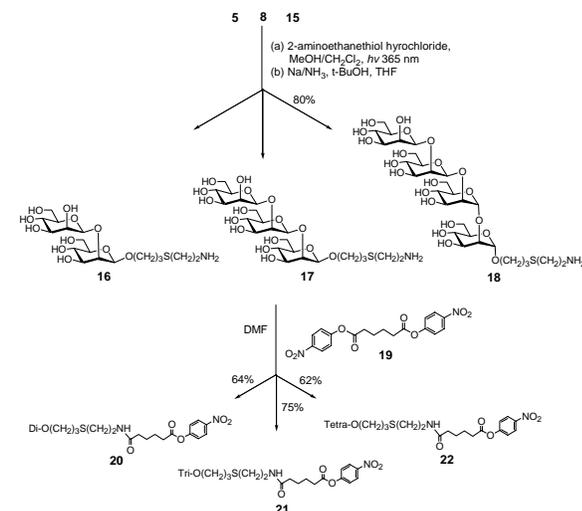
### Synthesis of acid stable epitope



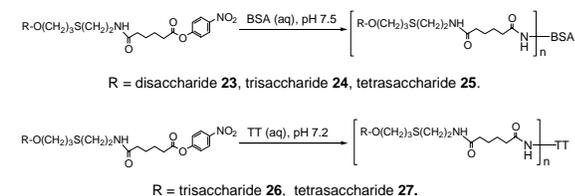
## Conjugation Chemistry

Reaction of glycosides **16**, **17** and **18** with homobifunctional adipic acid p-nitrophenyl diesters in dry DMF gave the corresponding half esters in good yields, and of sufficient stability to permit chromatographic purification.

### Synthesis of half esters 20, 21 and 22



## Synthesis of neoglycoproteins



## Acknowledgements

Financial support for this work was provided by research grants from the Alberta Ingenuity Centre for Carbohydrate Science and the Canadian Institutes of Health Research (CIHR).

