

Glycosylated nanomaterials: Detection and neutralisation of pathogenic bacteria and toxins

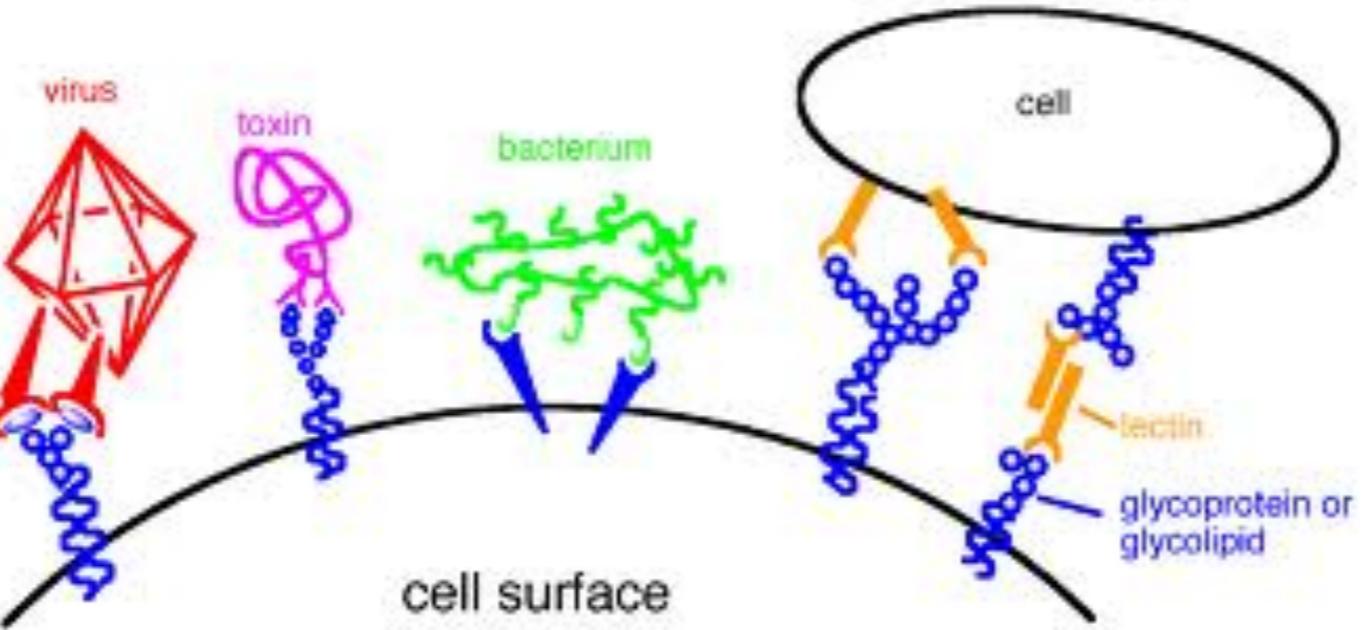
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Background

Protein-carbohydrate interactions mediate a multitude of critical biological recognition processes.¹ The proteins responsible for deciphering this information are termed lectins.²

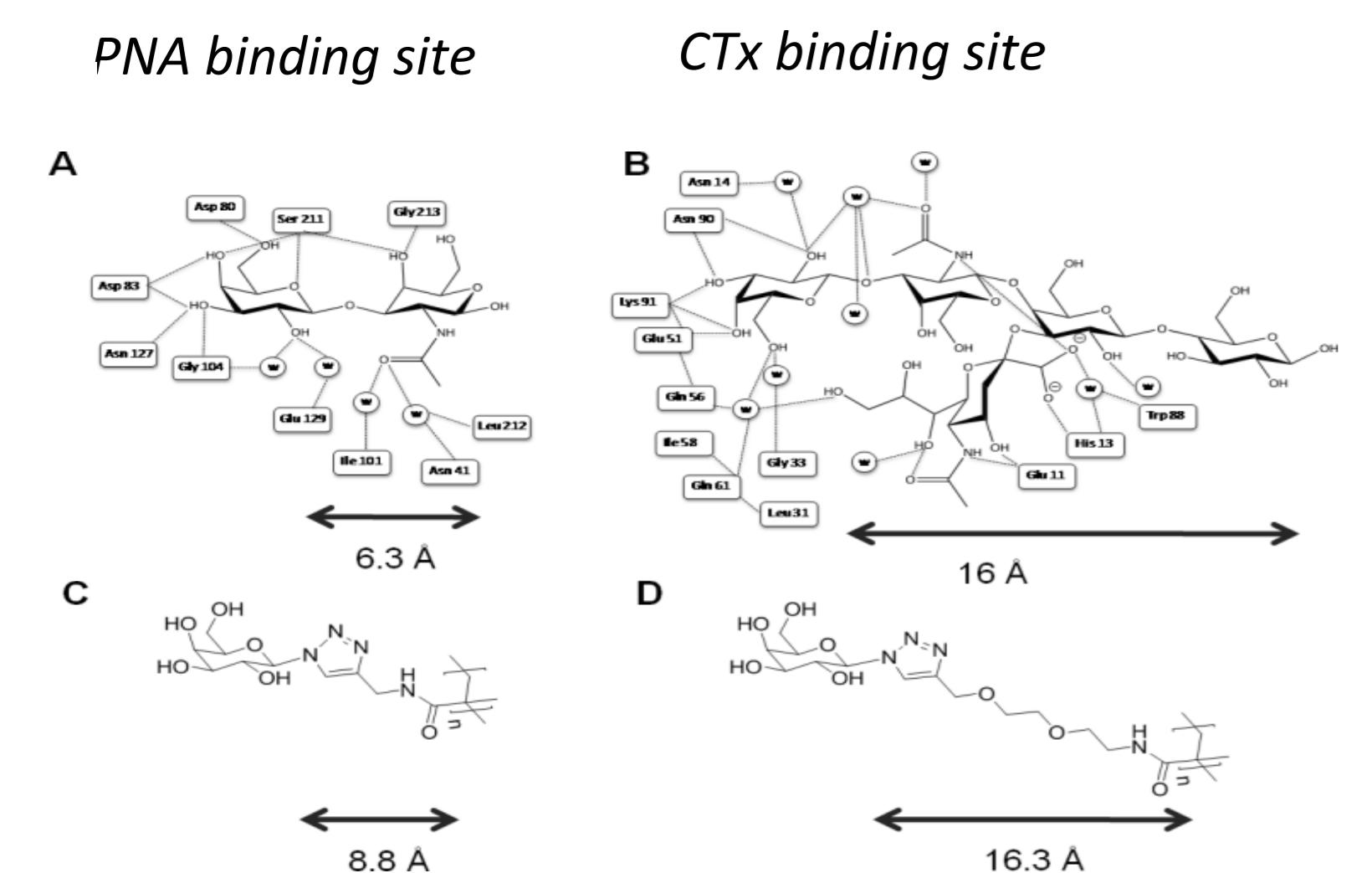


- Polymers with pendent carbohydrate moieties (glycopolymers) interact with lectins and have demonstrated binding affinities several orders of magnitude greater than a single carbohydrate.³
- The identification and treatment of bacterial infections remains a major healthcare challenge, especially to ensure appropriate application of a limited spectrum of antibiotics.

Inhibition of Bacterial Toxins

Bacterial Toxin Binding

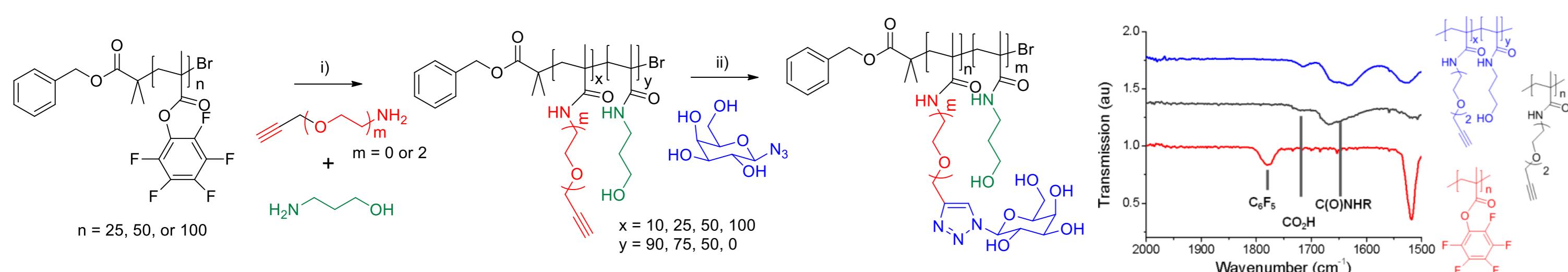
The cholera toxin (CTx) secreted by *Vibrio cholerae* binds glycosides expressed on the cell surface. Materials with high-affinity and selectivity for these lectins could find applications as anti-adhesive agents.



- Probe influence of chain length, carbohydrate density and linker length on binding inhibition.
- Glycopolymer library produced by tandem post - polymerisation modifications.
- Structural biology indicates CTx has deeper binding site than other galactose binding lectins such as Peanut agglutinin (PNA).

Tandem Post-Polymerisation Modification

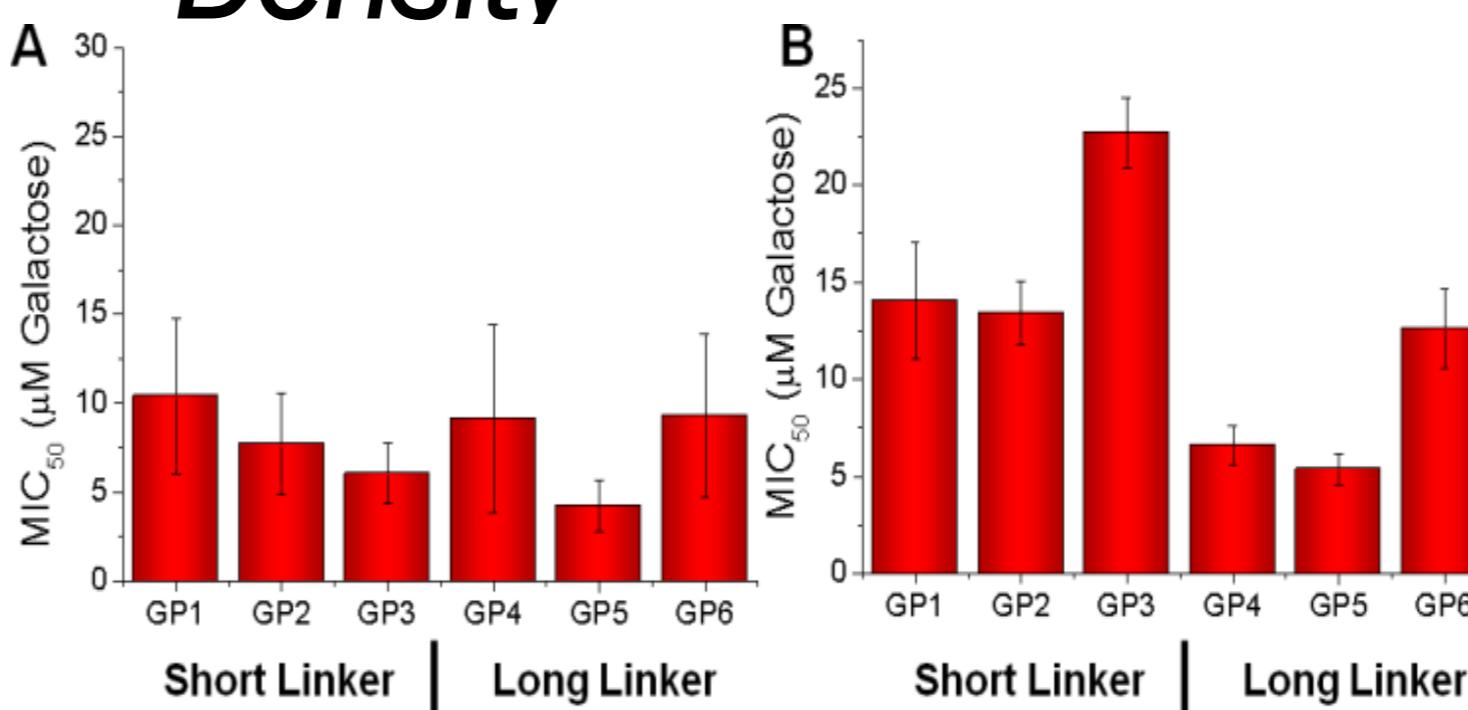
'Clickable' units are not compatible with controlled radical polymerisation. Instead, tandem-post polymerisation modification are performed.



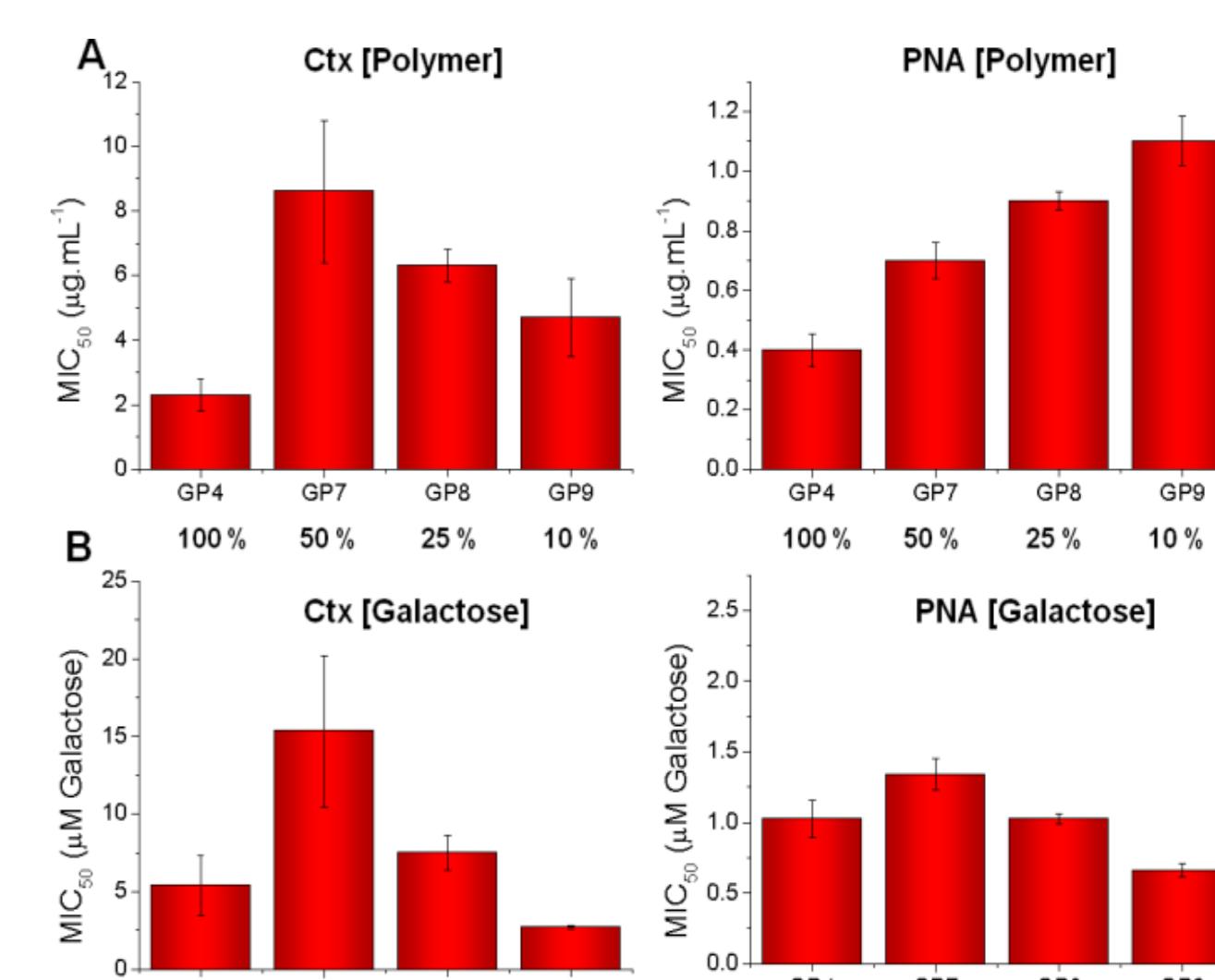
- Poly(pentafluorophenyl methacrylate) for easy modification.
- β -D-galactose was 'clicked-on' to pendent alkyne moieties.
- Results in biocompatible methacrylamide based (co)polymers.

Richards, S-J.; Jones, M. W.; Hunaban, M. I.; Haddleton, D. M.; Gibson M. I. *Angew. Chem Int. Ed.*, 2012

Role of Linker Length and Carbohydrate Density



- Polymers synthesised with 10, 25 and 50 % galactose.
- By mass (A), low galactose densities lead to a relative decrease in binding affinity/inhibitory activity.
- By mole of galactose (B), 10 % and 100 % functionalised polymers are the most active suggesting several features (e.g. sterics and site spanning) contribute to inhibitory activity.



Summary

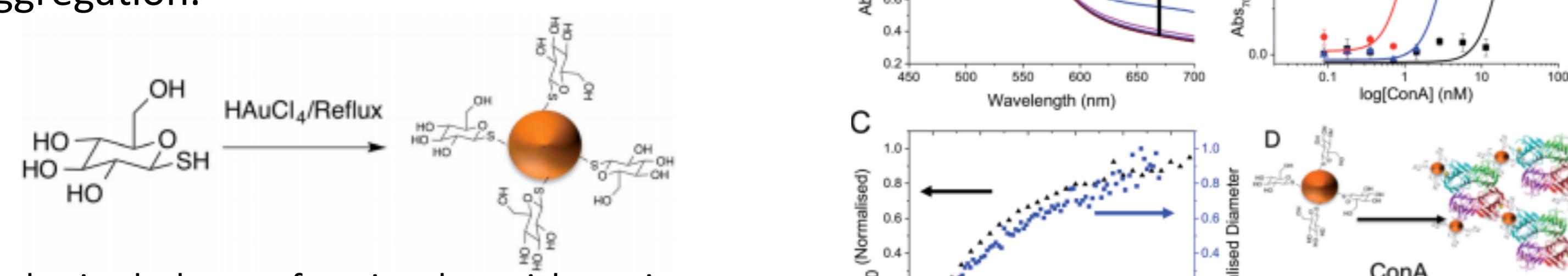
- Tandem post-polymerisation modification allows synthesis of polymers from same chain length distribution.
- Longer linker has better binding site accessibility.
- Inhibitors have to be developed for the binding site.

Jones, M.W. et. al., *Polym. Chem.*, 2013
Jones, M. W. et. al., *Chem. Sci.*, 2014

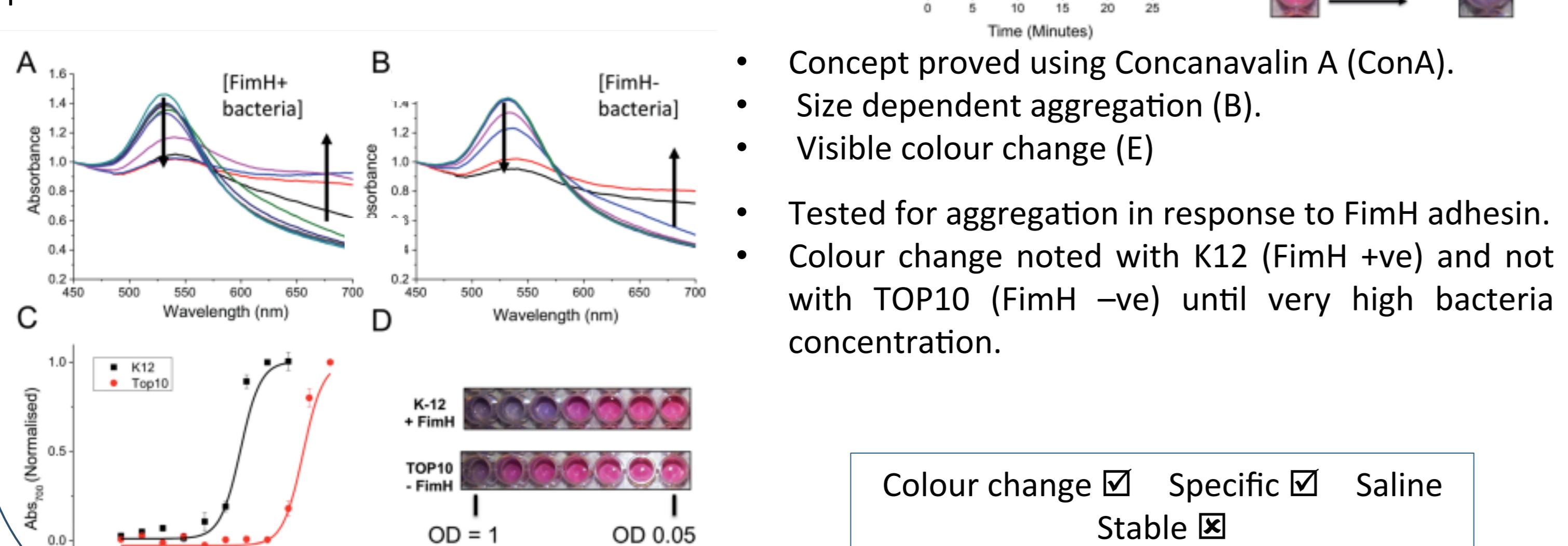
Detection of Lectins and Bacteria

Glycogoldnanoparticles

Goldnanoparticles have interesting optical properties. Red in solution → blue upon aggregation.



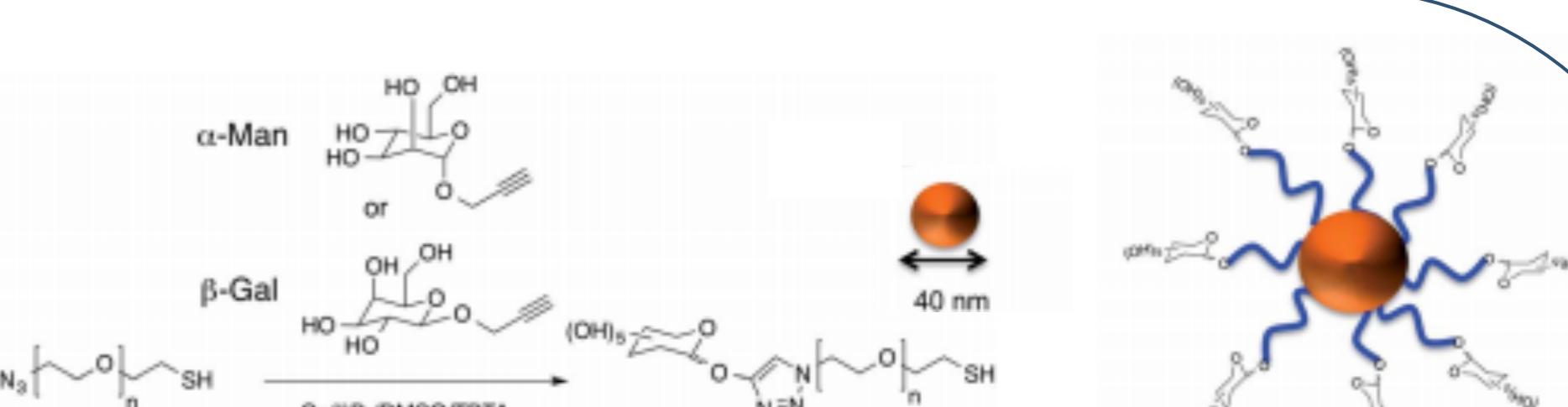
Synthesised glucose functional particles using a one-pot method.



Richards, S-J.; Fullam, L. C.; Besra, G. S.; Gibson M. I., *J. Mater. Chem.*, 2014

Increased saline stability by using a Poly(ethylene glycol) linker.

Carbohydrate introduced using a alkyne azide 'click' reaction.

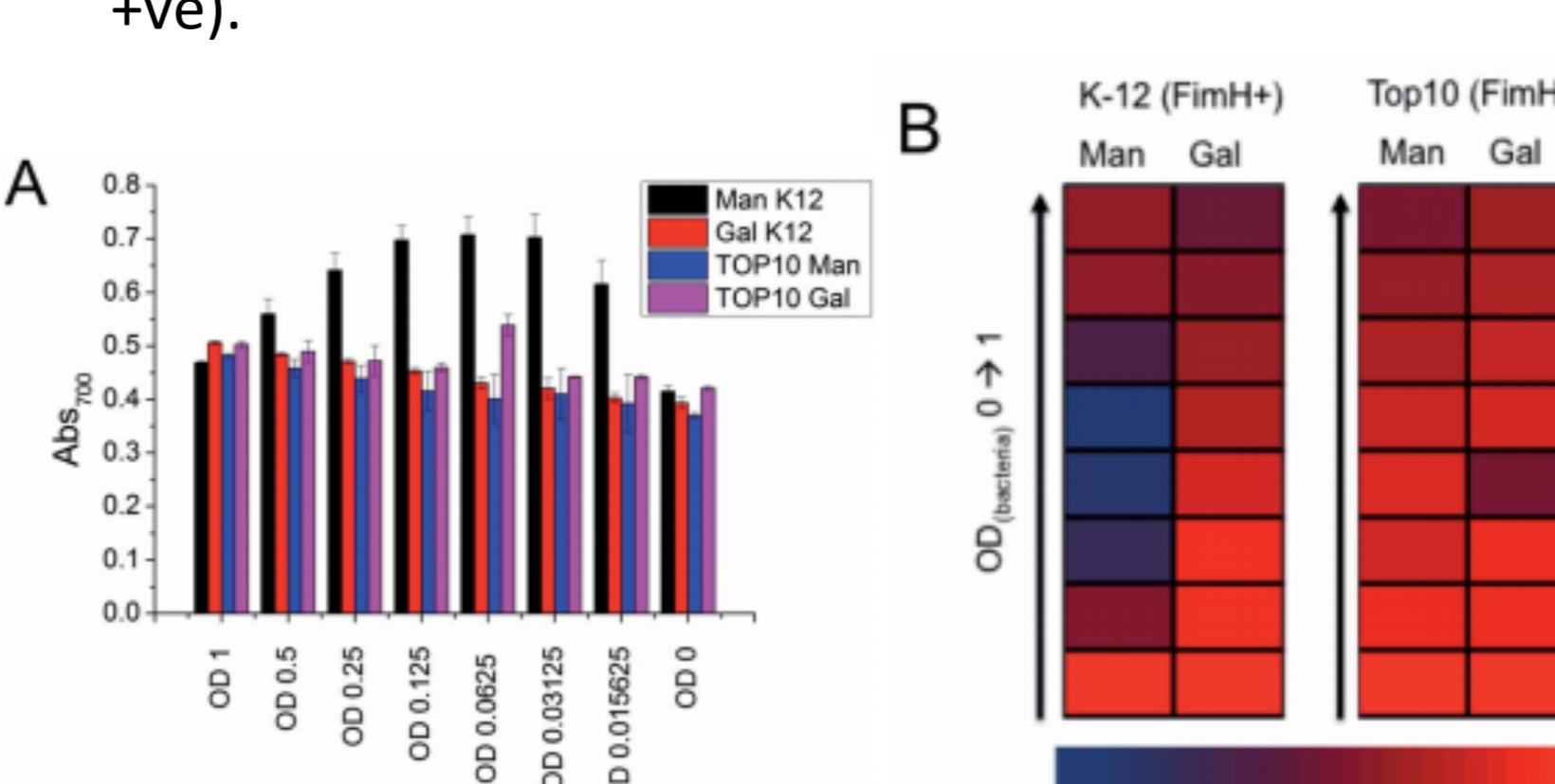


- Stable up to at least 1 M NaCl
- Lectin Specificity – aggregation with ConA (mannose specific) but not with PNA (galactose specific)

Colour change Specific Saline Stable

Summary

- Developed a sensitive, rapid colourimetric bioassay for the detection of lectins and Type I fimbriated bacteria.
- PEG layer increases saline stability (important for point of care diagnostics) but visual output is dramatically reduced.



References

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