

Electronic Supporting Information for

**Ice Recrystallisation Inhibition by Polyols: Comparison of
Molecular and Macromolecular Inhibitors and Role of
Hydrophobic Units**

Robert C. Deller,^{a, b} Thomas Congdon,^a Mohammed A. Sahid,^a Michael Morgan,^a Manu
Vatish,^c Daniel A. Mitchell,^c Rebecca Notman^a and Matthew I. Gibson^{a,*}

^a *Department of Chemistry, University of Warwick, Gibbet Hill Road, Coventry, UK, CV4*

7AL

^b *Molecular Organisation and Assembly in Cells (MOAC) Doctoral Training Centre,*

University of Warwick, CV4 7AL

^c *Clinical Sciences Research Laboratories, University of Warwick*

Clifford Bridge Road, Coventry, CV2 2DX

Corresponding Author Information

Fax: +44(0)2476 524112;

E-mail : m.i.gibson@warwick.ac.uk.

Ice Recrystallisation Inhibition: Additional Data

Concentration dependant activity of glucose derivates (main text, Figure 5) is shown in Figure S1 and the activity of mannose and O-ethylmannose is shown in Figure S2.

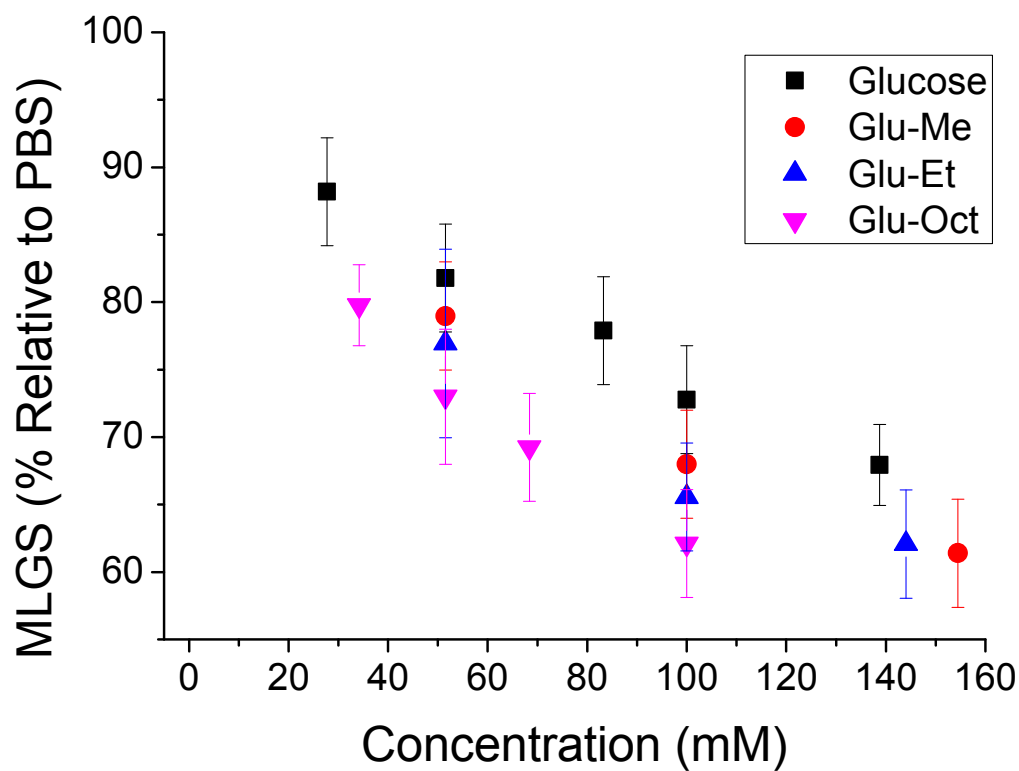


Figure S1. Ice recrystallisation inhibition activity of glucose and its derivatives.

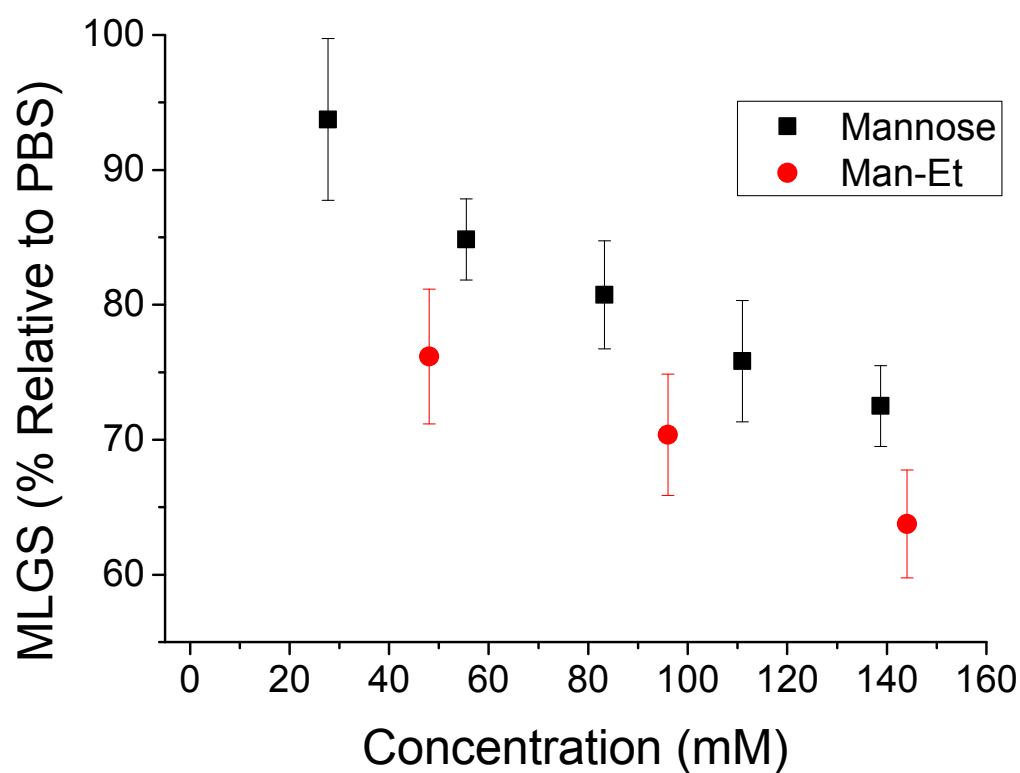


Figure S1. Ice recrystallisation inhibition activity of mannose o-ethylmannose.

Additional Experimental

Synthesis of 15 kDa poly(vinyl alcohol) used in DPH inclusion assay

Methyl(ethoxycarbonothioyl)sulfanyl benzene (0.21 g, 0.99 mmol), vinyl acetate (43.05 g, 0.500 mol) and ACVA (4,4'-azobis(4-cyanovaleric acid)) (0.013 g) were added to a stoppered vial. The solution was thoroughly degassed under a flow of N₂ for 20 mins and the reaction mixture was then allowed to polymerise at 68°C for 15 h. The yellow solution was then cooled to room temperature. Poly (vinyl acetate) was then recovered as a yellow sticky solid after precipitation into hexane. The hexane was then decanted and the poly (vinyl acetate) was re-dissolved in THF, which was then concentrated *in vacuo* and was then thoroughly dried under vacuum at 40°C for 24 h, forming a white crystalline solid. The PVAc precursor was characterised by ¹H NMR and SEC in THF. PVAc was then dissolved in a methanol (20 mL) and hydrazine hydrate solution (15 mL, 80% in water) in a round bottom flask. The reaction mixture was stirred at 30°C for 2 hours, then dialysed using distilled water. Poly(vinyl alcohol) was recovered as a spongy white solid by freeze drying the dialysis solution. Hydrolysis was confirmed by ¹H NMR.

PVAc Precursor: ¹H NMR (400 MHz, CDCl₃): δ=4.61 (-CHO-CH₂ br 1H), δ=1.74 (-CO-CH₃ br 3H), δ=1.53 (-CH₂- br 2H), M_n^{GPC}(THF) = 34400 Da, M_n/M_w = 1.22.

PVA: ¹H NMR (400 MHz, CDCl₃): δ=4.00 (-CHOH- br 1H), δ=1.68-1.60 (-CH₂- br 2H).