Direct solid phase synthesis and fluorescence labeling of large, monodisperse mannosylated dendrons in a peptide synthesizer.

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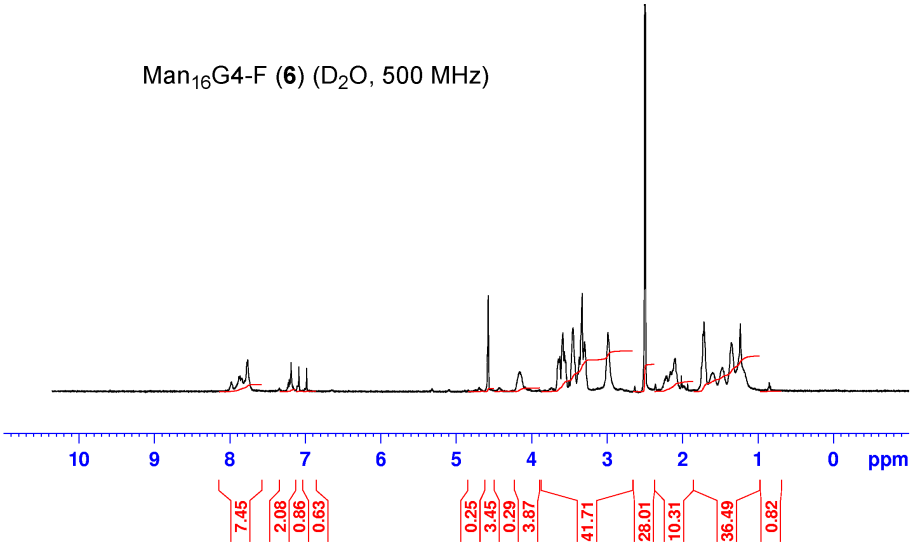
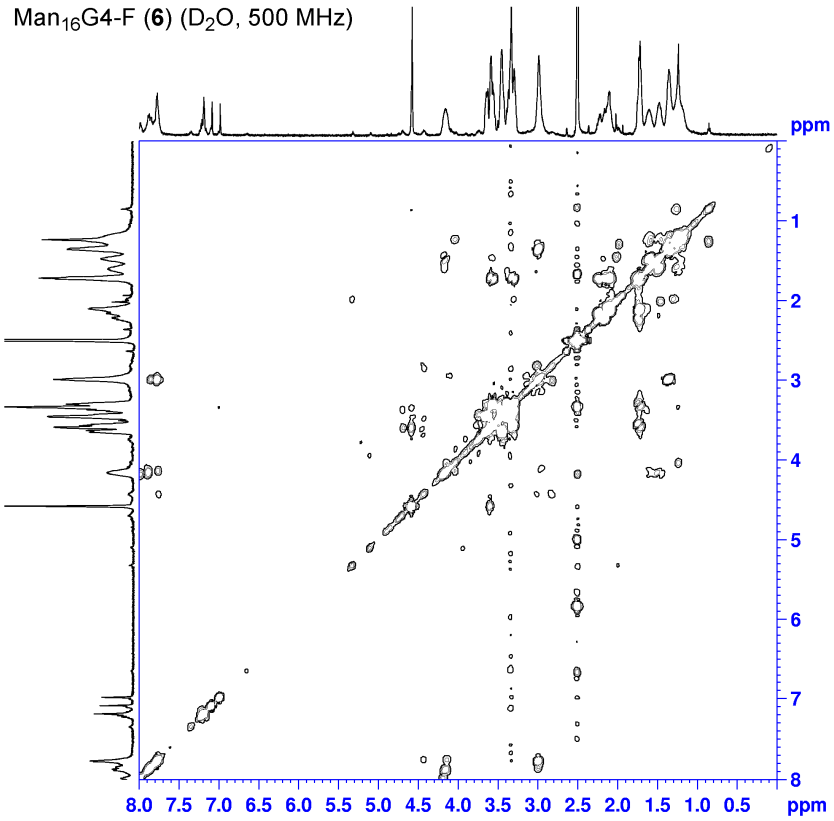
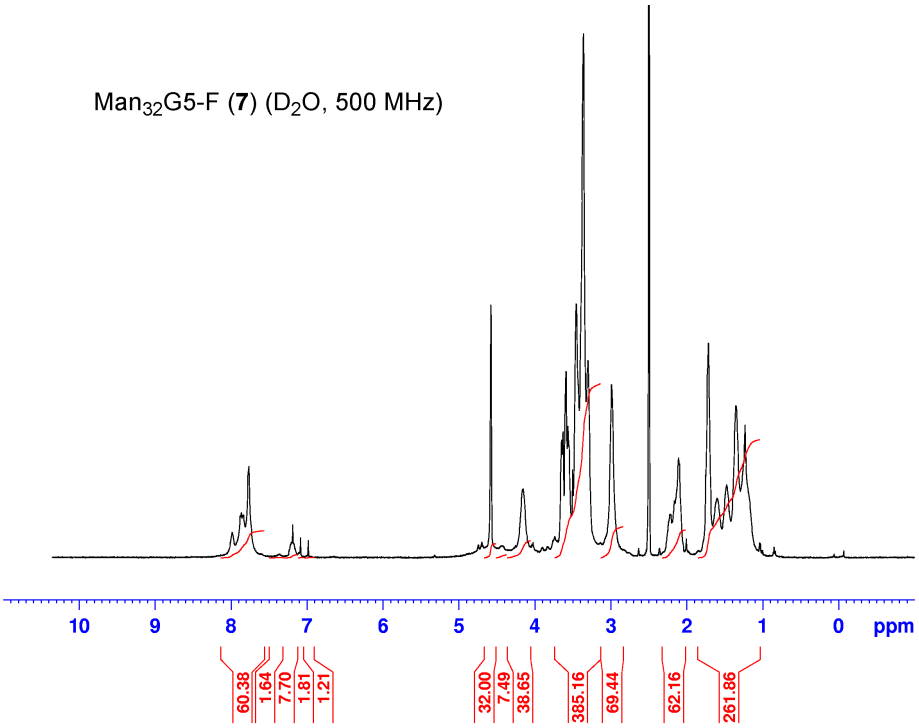
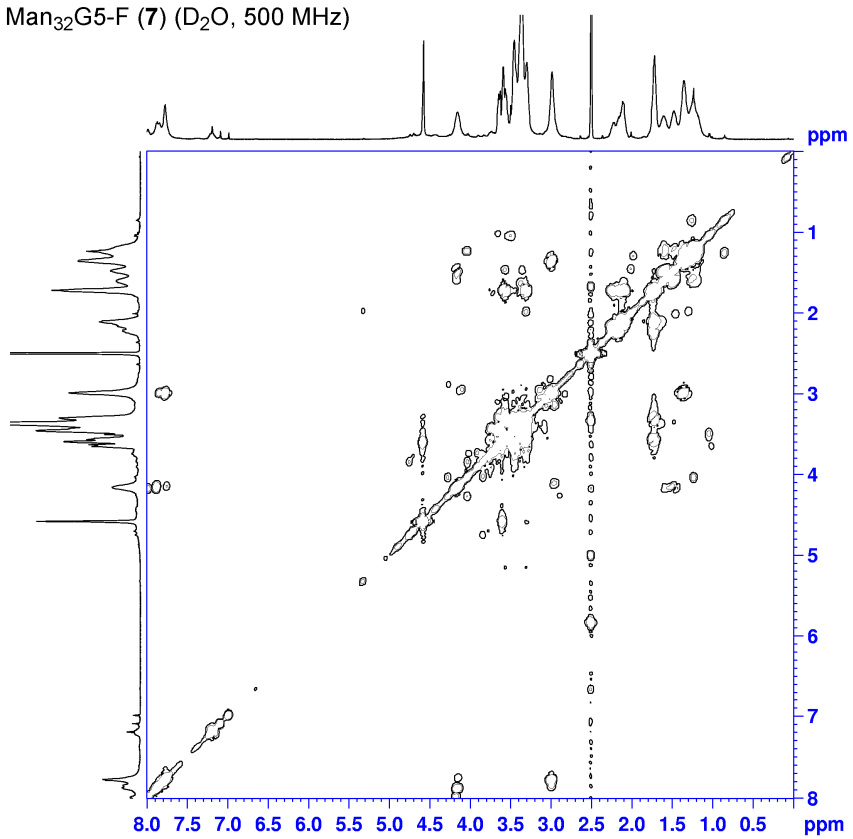
Supporting information.

NMR spectra


Man16G4-F (6) (D2O, 500 MHz)
10
9
8
7
6
5
4
3
2
1
0
ppm
7.45
2.08
0.86
0.63
0.25
3.45
0.29
3.87
41.71
28.01
10.31
36.49
0.82

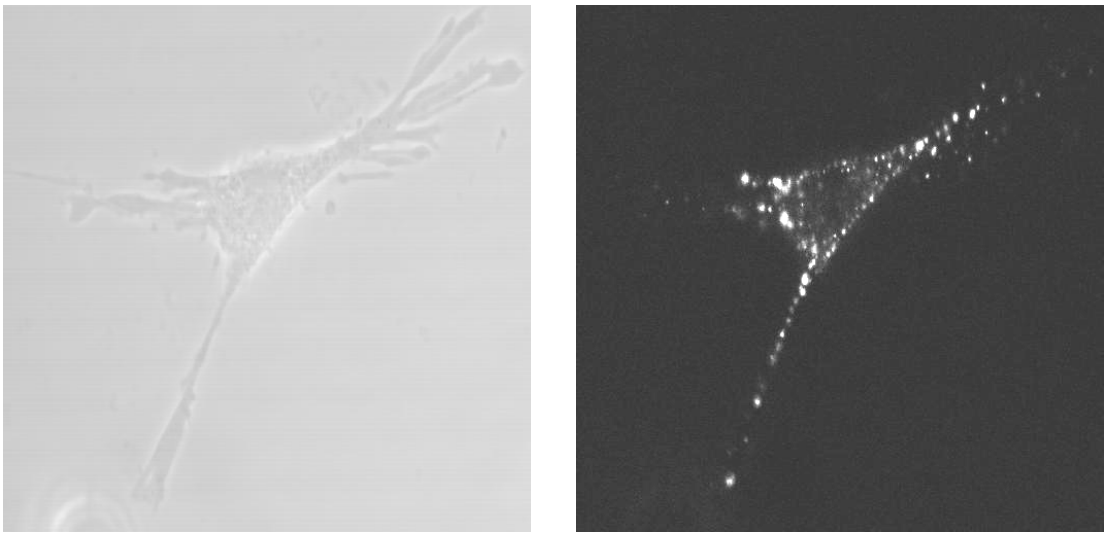

Man16G4-F (6) (D2O, 500 MHz)
ppm
1
2
3
4
5
6
7
8
8.0 7.5 7.0 6.5 6.0 5.5 5.0 4.5 4.0 3.5 3.0 2.5 2.0 1.5 1.0 0.5
ppm


Man32G5-F (7) (D2O, 500 MHz)
10
9
8
7
6
5
4
3
2
1
0
ppm
60.38
1.64
7.70
1.81
1.21
32.00
7.49
38.65
385.16
69.44
62.16
261.86

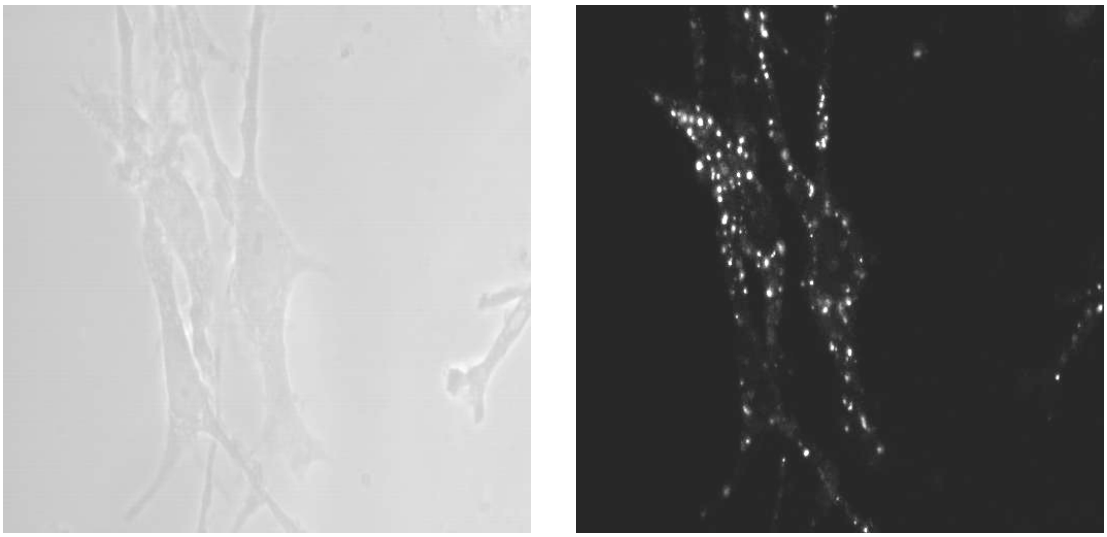

Man32G5-F (7) (D2O, 500 MHz)
ppm
1
2
3
4
5
6
7
8
8.0 7.5 7.0 6.5 6.0 5.5 5.0 4.5 4.0 3.5 3.0 2.5 2.0 1.5 1.0 0.5
ppm

Images from the confocal microscopy analysis of the entry of the FITC-labelled glycodendron Man_{16}-G4-K(FITC) (18) in dendritic cells

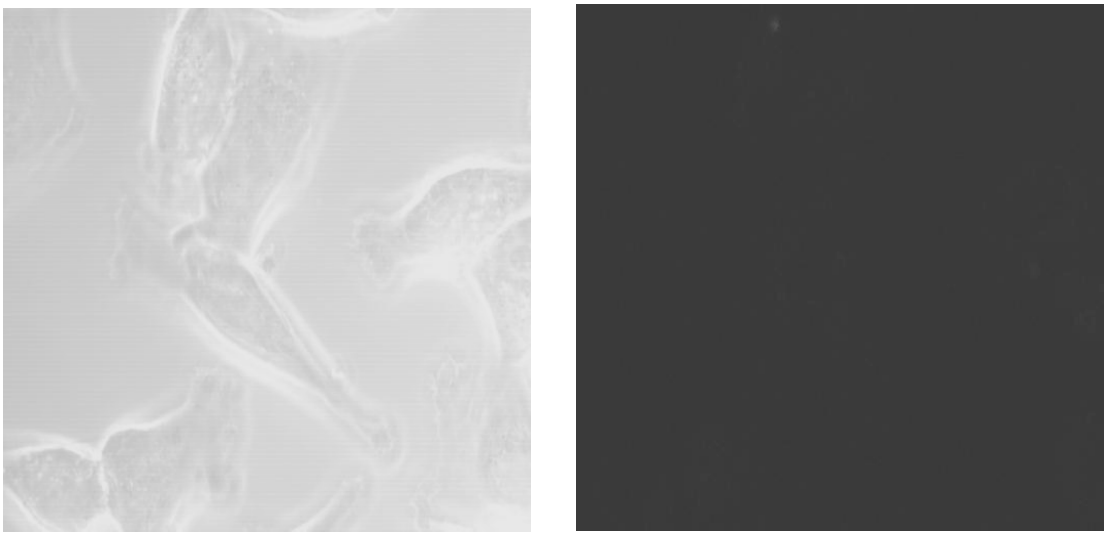
a)



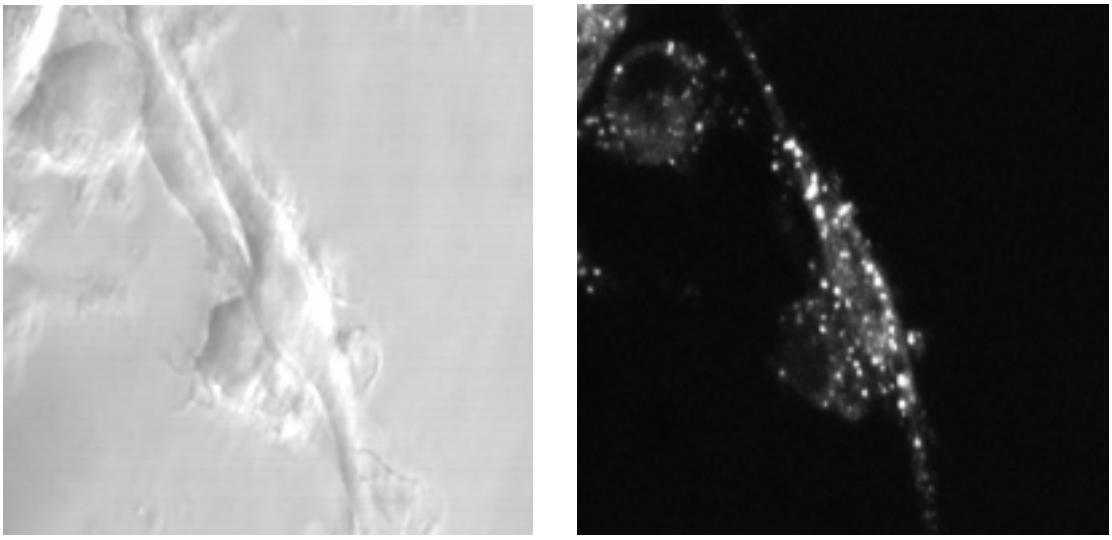
b)



c)



d)



The phase contrasts are shown on the left. (a) Incubation for 1 h at 4°C with 12 (1μM), washing, then 2 h chase at 37°C. (b) Incubation 2 h at 37°C with 12 (1 μM). Incubation for 1 h at 4°C with 12 (1μM), washing, then 2 h chase at 37°C. (c) Negative control: Incubation with HeLa cells under the same conditions as (b). (d) Incubation for 2 h at 37°C with 12 (10μM), washing, then 24 h chase at 37°C.