

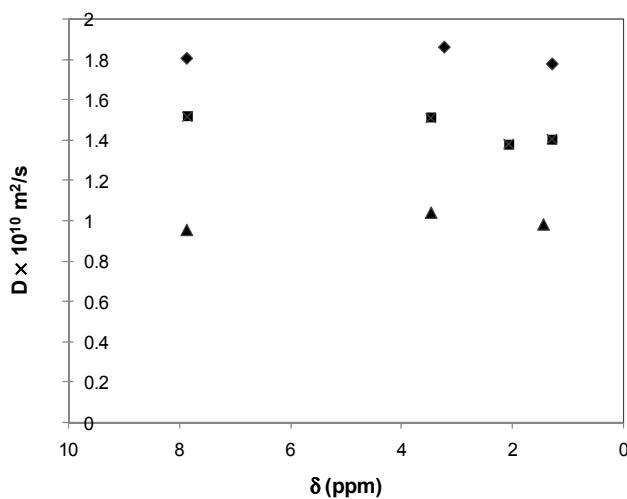
Synthesis and Biological Evaluation of Multivalent Carbohydrate Ligands Obtained By Click Assembly of Pseudo-Rotaxanes †

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Supporting Information Available: MS spectra and TLC profiles of rotaxane **13** and oligorotaxanes **14**, **15**, **16**. Figure displaying the diffusion coefficients of **13**, **14** and **16** calculated by NMR.

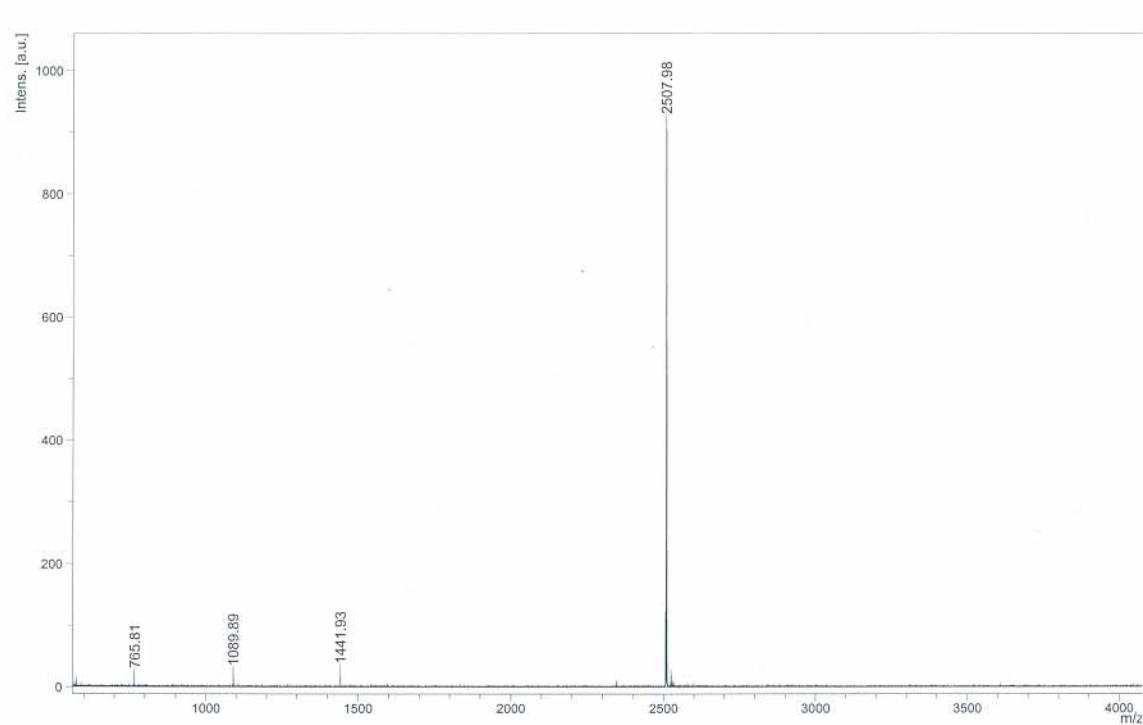
Diffusion coefficients of **13** (♦), **14** (■) and **16** (▲) calculated by NMR(500 MHz, 298K, D_2O) from proton signals of the cyclodextrin and the lactosyl group ($\delta=3.4\text{-}4$ ppm), of the triazole group ($\delta=7.3\text{-}8$ ppm) and of the decane chain ($\delta=0.9\text{-}1.6$ ppm).



TLC profiles of 1) Lactosyl-cyclodextrin **6**, 2) Rotaxane **13**, 3) Oligorotaxane **14**, 4) Oligorotaxane **15**, 5) Oligorotaxane **16**. Elution conditions: CH₃CN/H₂O (2% NaBr) 6:4.

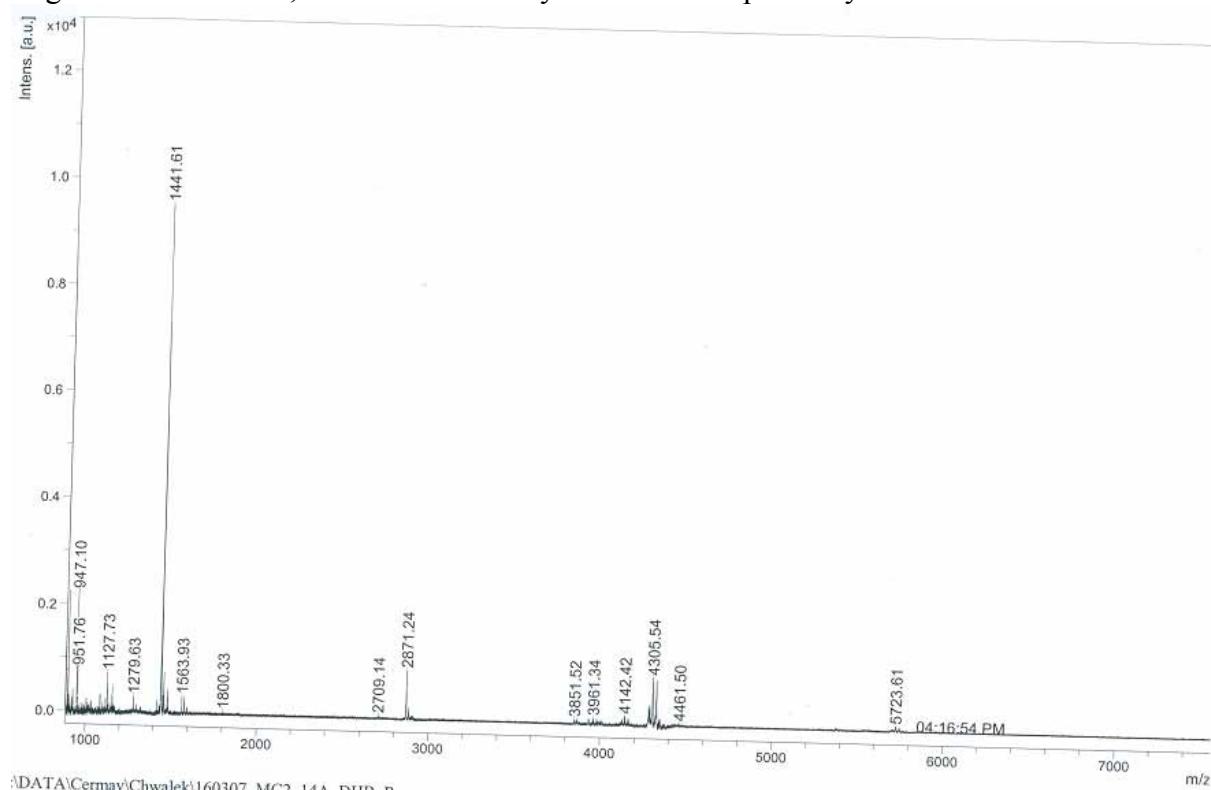


MALDI-MS spectrum of rotaxane **13**. The peak at 2508 was assigned to **13**

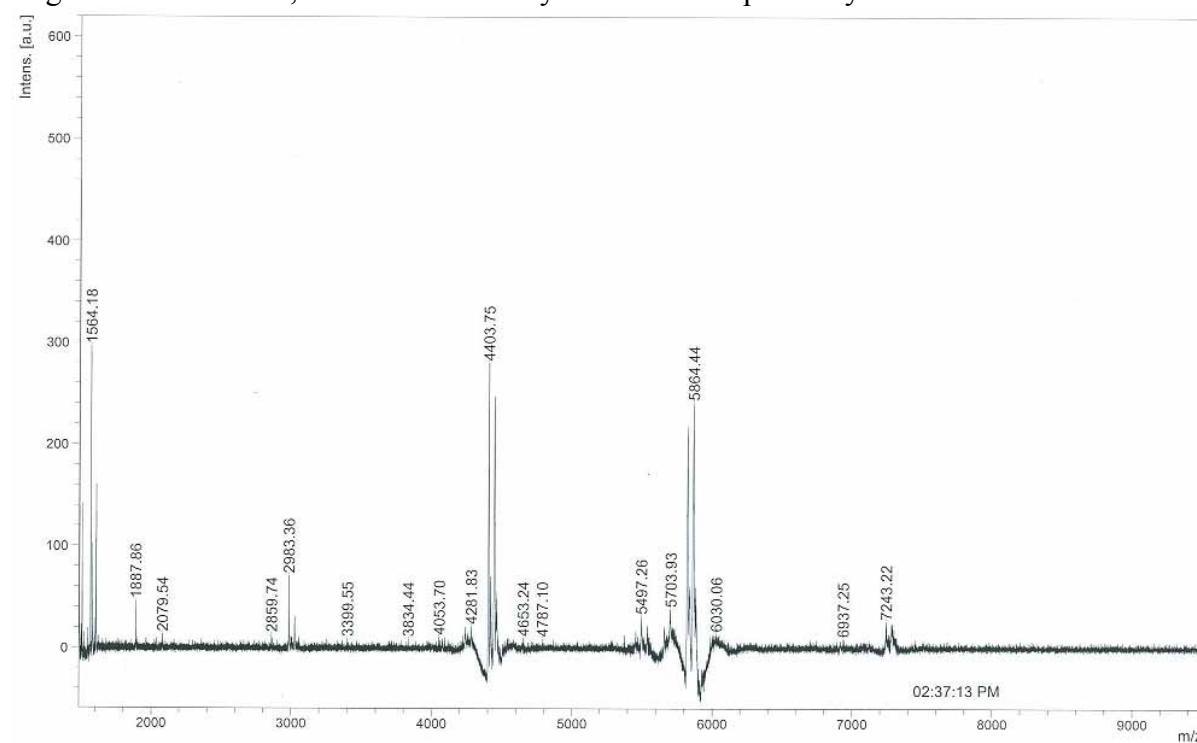


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MALDI-MS spectrum of oligorotaxane **14**. Peaks at 2871, 4305 and 5723 were assigned to oligorotaxanes with 1, 2 and 3 threaded cyclodextrins respectively.



MALDI-MS spectrum of oligorotaxane **15**. Peaks at 4403, 5864 and 7243 were assigned to oligorotaxanes with 2, 3 and 4 threaded cyclodextrins respectively.



MALDI-MS spectrum of oligorotaxane **16**. Peaks at 5819, 7237, 8653 and 10072 were assigned to oligorotaxanes with 3, 4, 5 and 6 threaded cyclodextrins respectively.

