Supplementary Material

Design, Modeling and Synthesis of 1,2,3-Triazole-linked Nucleoside-Amino Acid Conjugates as Potential Antibacterial Agents

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Trial	Reagents	Reaction Conditions	Workup and purification	Result
			Added MeOH, Conc.	
1	5, TsCl, pyr	0° C to rt, 16 h	Column 1: CH2Cl2:MeOH	12%
			Column 2: Hex:EtOAc	
			Diluted with CH ₂ Cl ₂	
2	5, TsCl, pyr	0° C to rt, 17.25 h	Wash: HCl, brine	9%
			Column: CH2Cl2:MeOH	
			Diluted with CH ₂ Cl ₂	
3	5 , Ts ₂ O, pyr	0° C to 35° C, 4.5 h	Wash: HCl, brine, NaHCO ₃	23%
			Column: CH2Cl2:MeOH	
4	5 , Ts ₂ O, pyr	0° C to rt, 16.5 h	Diluted with CH ₂ Cl ₂	0%
			Wash: HCl, brine, NaHCO ₃	
			Conc.	56%,
5	5, TsCl, pyr	0° C to rt, 14.75 h	Column 1: CH2Cl2:MeOH	· ·
			Column 2: CH2Cl2:MeOH	impure

Table S1. Reagents and results for the formation of the acyclic tosylate, 6

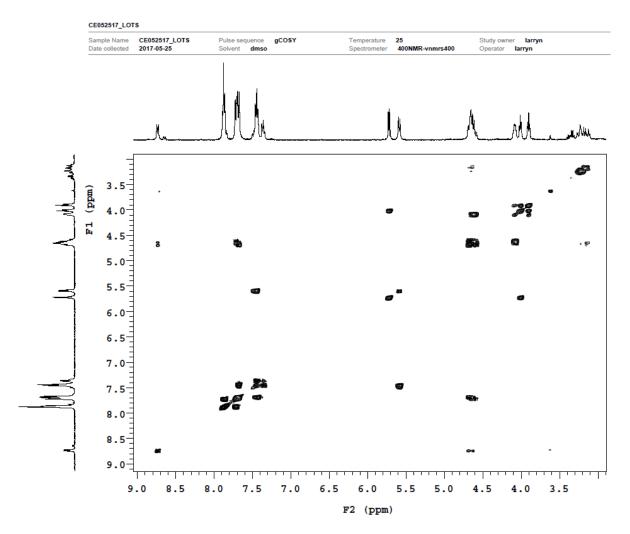


Figure S1. ¹H-¹H COSY of 4.

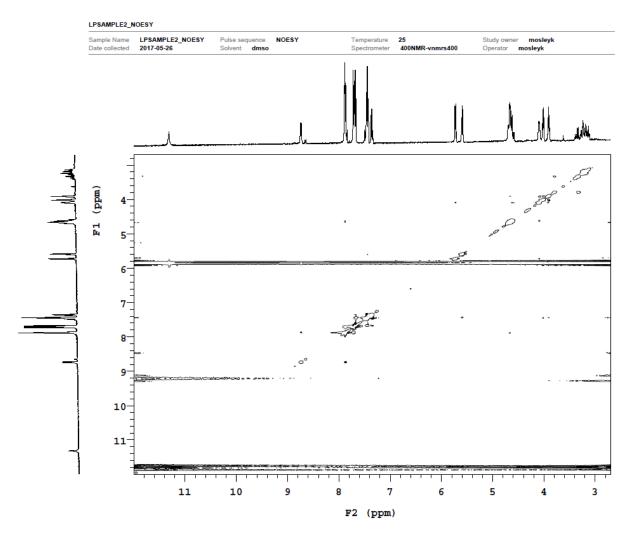


Figure S2. NOESY spectrum of **4**.