

Supplementary Material

A Surface Modifier for the Production of Selectively Activated Amino Surface Groups

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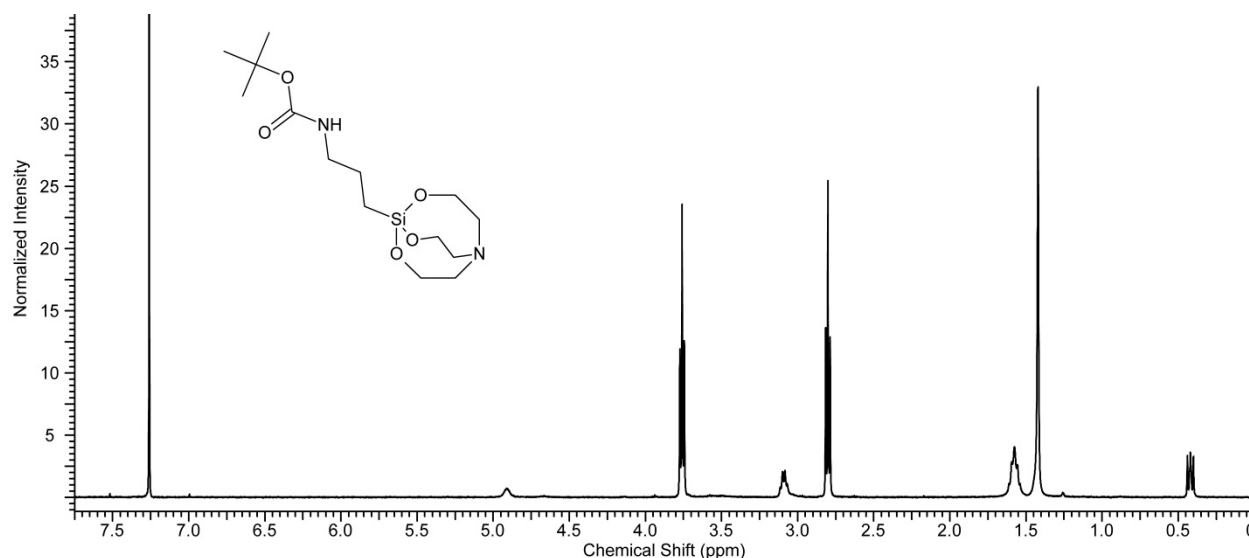


Figure S1. ^1H NMR spectra of BocAPS, CDCl_3 .

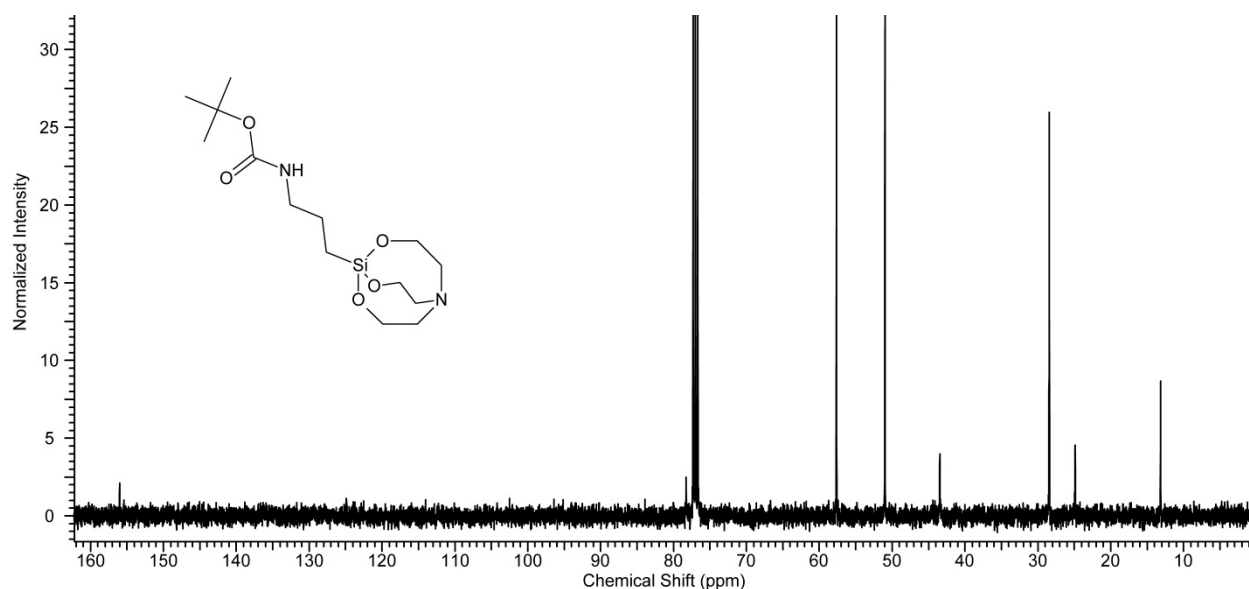
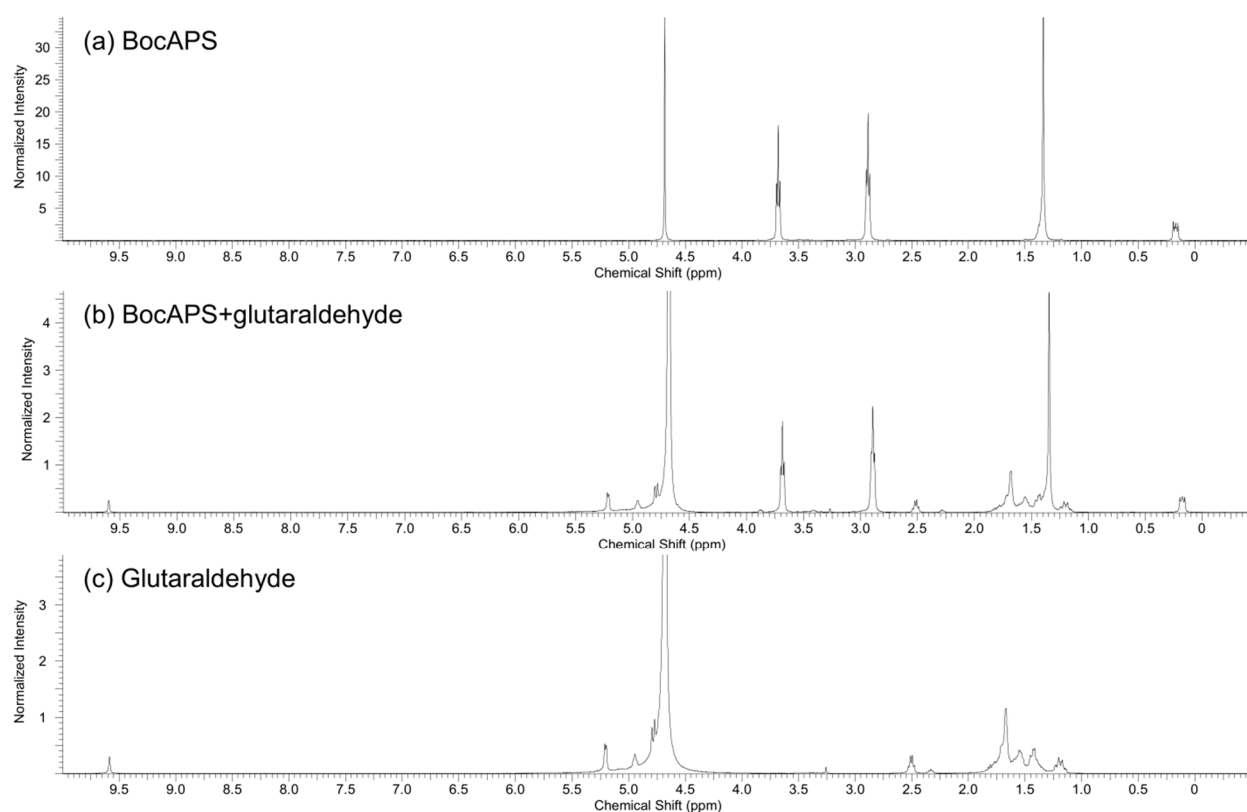


Figure S2. ^{13}C NMR spectra of BocAPS, CDCl_3 .

Table S1. Selected crystallographic data and details of refinement for *N-tert-butoxycarbonylamino*propyl silatrane.

Empirical Formula	C ₁₄ H ₂₈ N ₂ O ₅ Si
Formula weight	332.47
Crystal system	Triclinic
Space group	P-1
a (Å)	9.5590(10)
b (Å)	10.294(5)
c (Å)	10.763(4)
α(°)	67.14(4)
β(°)	84.85(3)
γ(°)	62.28(3)
V (Å ³)	858.5(5)
Z	2
D _{calcd} (Mg/m ³)	1.286
Radiation (CuKα)	λ = 1.54178 Å
T (K)	290(2)
θ range (°)	4.49–65.17
Total reflections	5713
Unique reflections	2920
Parameters / restraints	231 / 15
R ₁ ; wR ₂ (I > 2σ(I))	0.0404, 0.0880
Goodness-of-fit on F ²	1.029

**Figure S3.** ¹H NMR spectra of **BocAPS** (a), glutaraldehyde (c), mixture of **BocAPS** and glutaraldehyde after 30 min (b), D₂O.

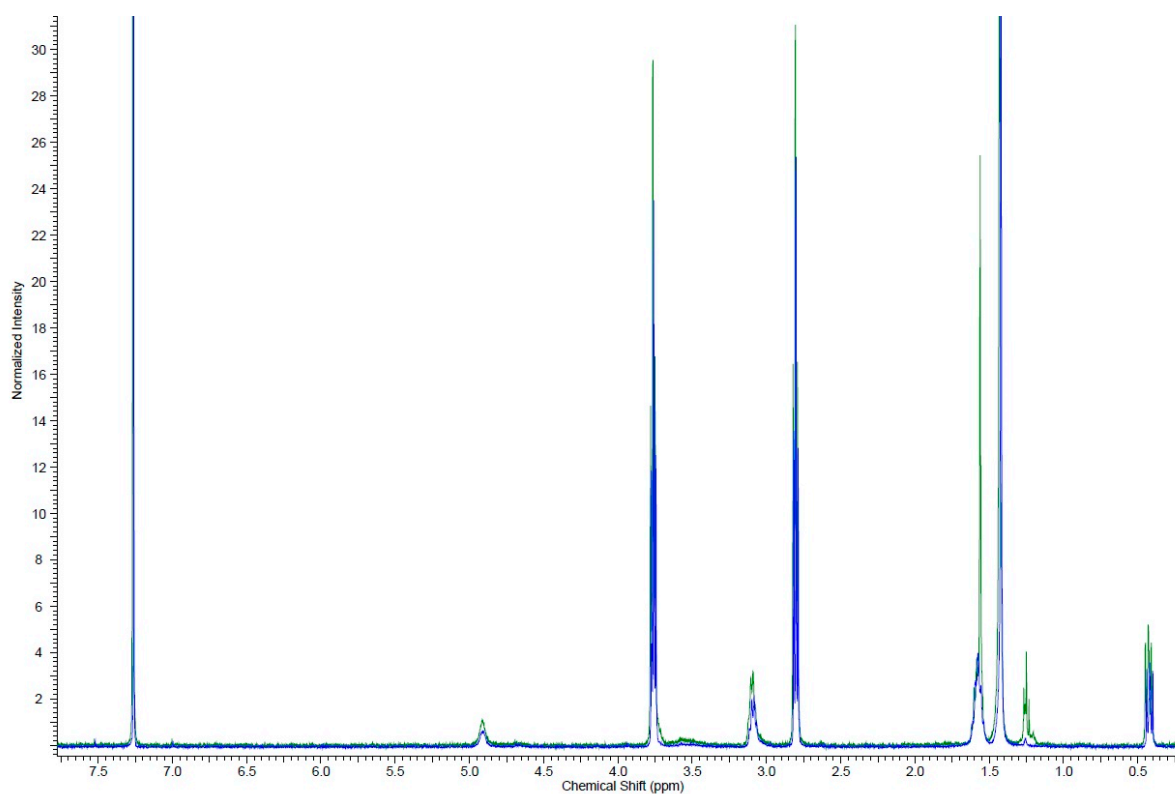


Figure S4. ^1H NMR spectra of **BocAPS** obtained recently (green, there are residues of DMF) and after two years of storage at room conditions (blue), CDCl_3 .

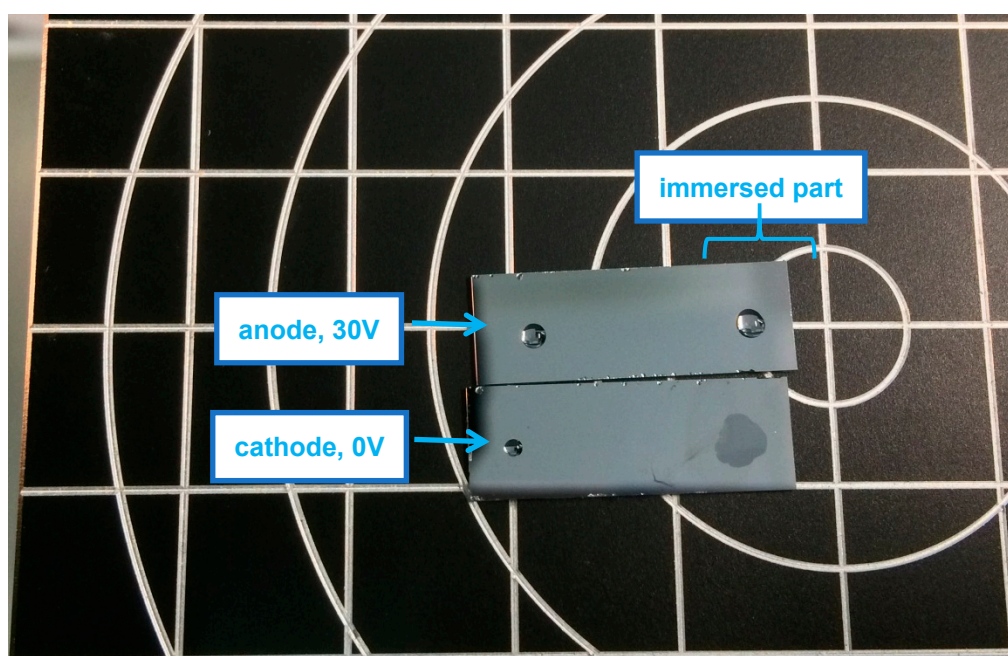


Figure S5. Wettability of parts of the anode and cathode after electrochemical removal of Boc-groups.

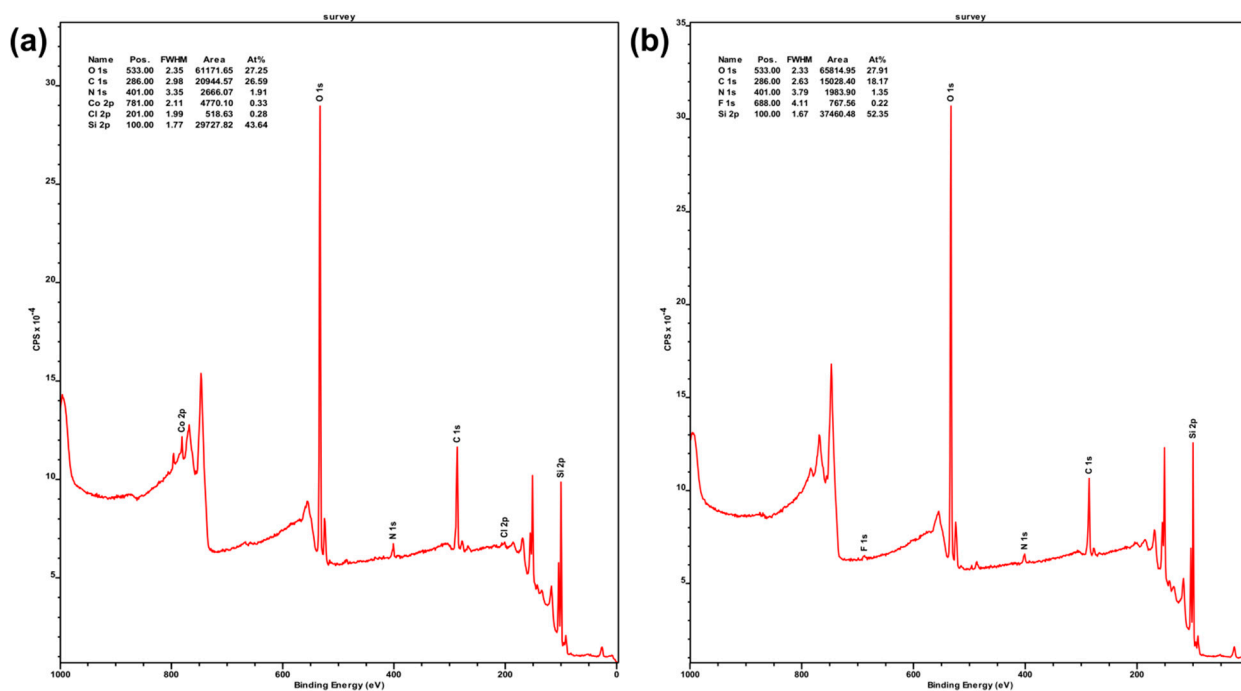


Figure S6. Survey XPS spectra of the BocAPS-modified SiO₂ surface (a) and BocAPS-modified SiO₂ surface with removed Boc-groups (b).

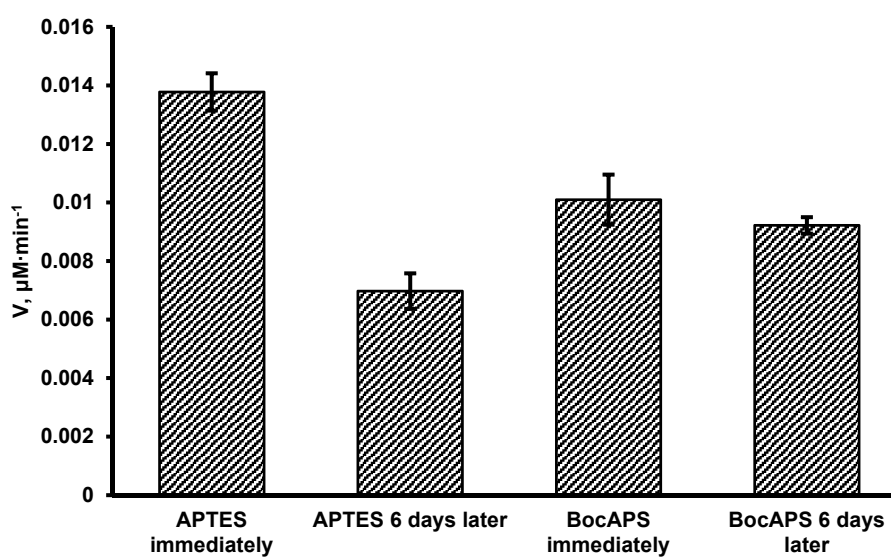


Figure S7. The activity of glucose oxidase immobilized on silicon slides, which were modified by APTES or BocAPS and had different holding times (at room conditions) between the formation of SAM and the immobilization of the enzyme.

