

Supporting Information

Effect of Zwitterionic Additive on Electrode Protection through Electrochemical Performances of Anatase TiO₂ Nanotube Array Electrode in Ionic Liquid Electrolyte

Aleksandra Roganović ¹, Milan Vraneš ¹, Nikola Cvjetićanin ², Xiaoping Chen ³ and Snežana Papović ^{1,*}

¹ Faculty of Sciences, Department of Chemistry, Biochemistry and Environmental Protection, University of Novi Sad, Trg Dositeja Obradovića 3, 21000 Novi Sad, Serbia

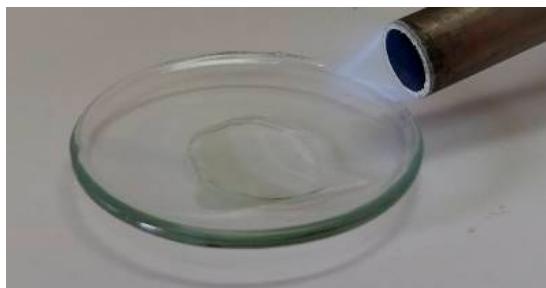
² Faculty of Physical Chemistry, University of Belgrade, Studentski trg 12-16, 11158 Belgrade, Serbia

³ Hangzhou Bay Automotive College/Mechanical College, Ningbo University of Technology, No.769, Bihai Road 2, 315336, Hangzhou Bay New Zone, 315000 Ningbo, China

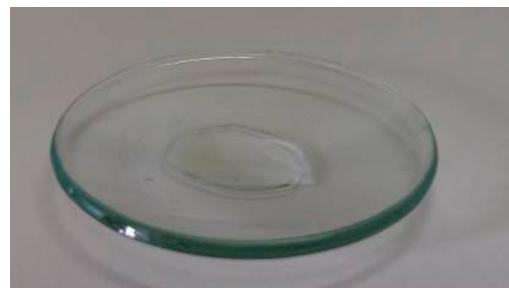
* Correspondence: snezana.papovic@dh.uns.ac.rs; Tel: +381-21-485-2751;
Fax: +381-21-454-065

Table S1. Summarized the identification results and normalized ion currents for investigated sample of synthetized zwitterionic compound C₁C₄imSO₃.

m/z fragment	Normalized ion current for compound C ₁ C ₄ imSO ₃
14	1·10 ⁻⁹
16	5·10 ⁻¹⁰
17	2·10 ⁻⁹
18	8·10 ⁻⁹
28	1·10 ⁻⁸
32	5·10 ⁻⁹
36	4·10 ⁻⁹
44	1·10 ⁻¹⁰
48	2·10 ⁻¹⁰
64	2·10 ⁻¹⁰



(a)



(b)

Figure S1. Flammability test pictures of LiTFSI/C₂C₂imTFSI/C₁C₄imSO₃ when is: **a**) exposed to flame and **b**) after exposition of 120 s to flame.

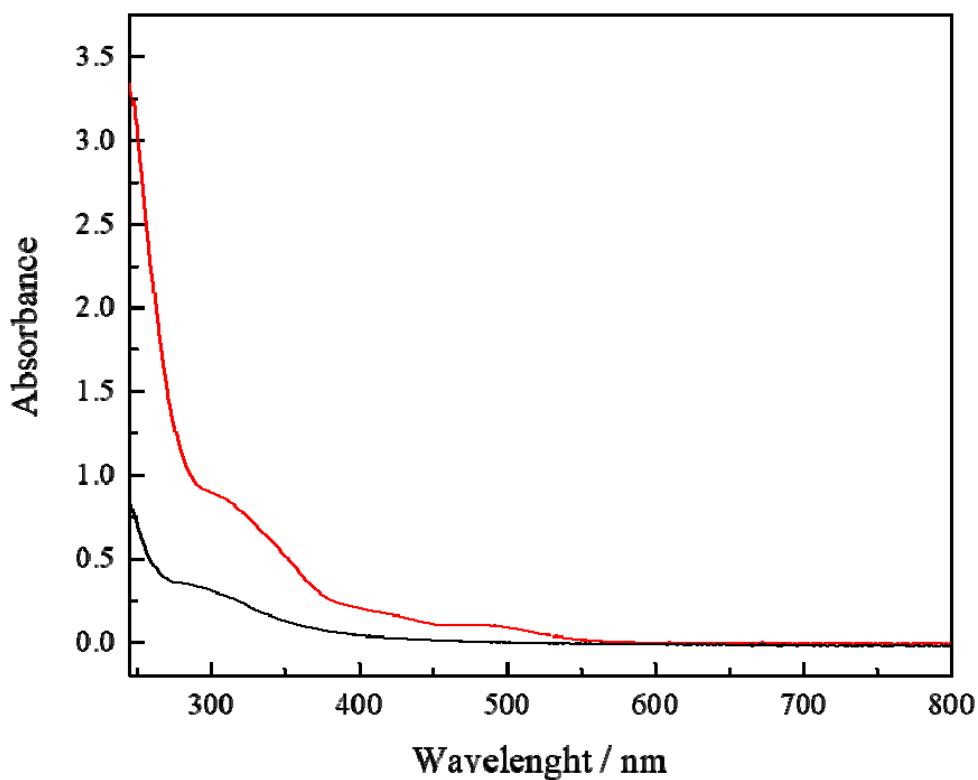


Figure S2. UV/Vis spectra of electrolyte recorded in the range from 250–800 nm at room temperature, where is red line related to the LiTFSI/C₂C₂imTFSI and the black line related to the LiTFSI/C₂C₂imTFSI/C₁C₄imSO₃ (spectra for electrolytes were recorded after galvanostatic cycling).

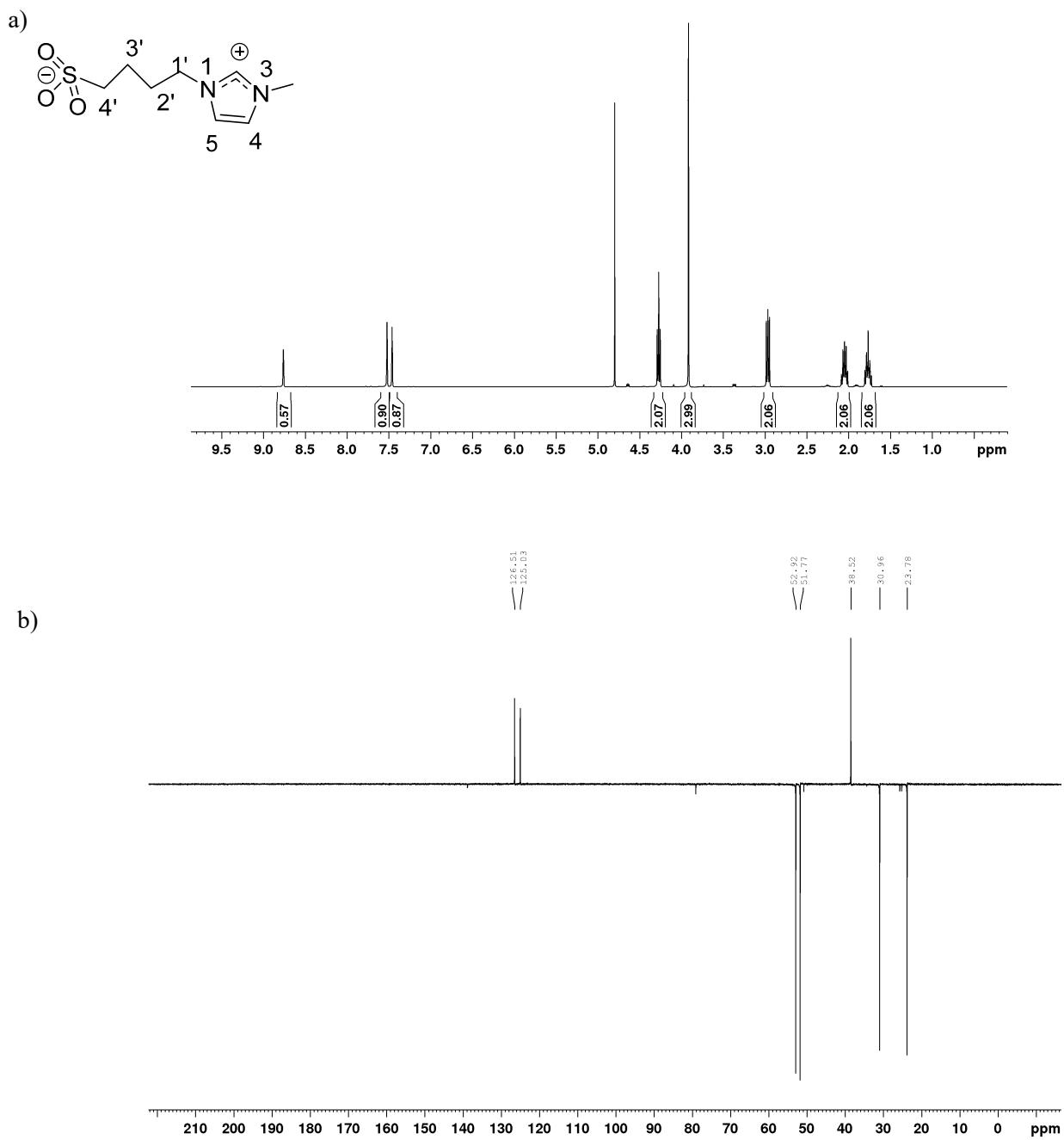


Figure S3. a) ^1H and b) ^{13}C NMR spectra for synthesized $\text{C}_1\text{C}_4\text{imSO}_3$

a) ^1H NMR: 1.76, m, 2H, $\text{NCH}_2\text{CH}_2\text{CH}_2\text{SO}_3^-$; 2.05, m, 2H, $\text{NCH}_2\text{CH}_2\text{CH}_2\text{CH}_2\text{SO}_3^-$; 2.96, m, 2H, $J=7.9$ Hz, $\text{NCH}_2\text{CH}_2\text{CH}_2\text{CH}_2\text{SO}_3^-$; 3.92, s, 3H, NCH_3 ; 4.27, t, 2H, $J=7.1$ Hz, $\text{NCH}_2\text{CH}_2\text{CH}_2\text{CH}_2\text{SO}_3^-$; 7.46, t, 1H, $J=1.9$ Hz, H-4; 7.52, t, 1H, $J=1.9$ Hz, H-5; 8.52, s, 0.6H, H-2.

b) ^{13}C NMR: 23.78 ($\text{NCH}_2\text{CH}_2\text{CH}_2\text{CH}_2\text{SO}_3^-$); 30.96 ($\text{NCH}_2\text{CH}_2\text{CH}_2\text{CH}_2\text{SO}_3^-$); 38.52 (NCH_3); 51.77 ($\text{NCH}_2\text{CH}_2\text{CH}_2\text{CH}_2\text{SO}_3^-$); 125.03 (C-5); 126.51 (C-4); 138.85 (C-2).

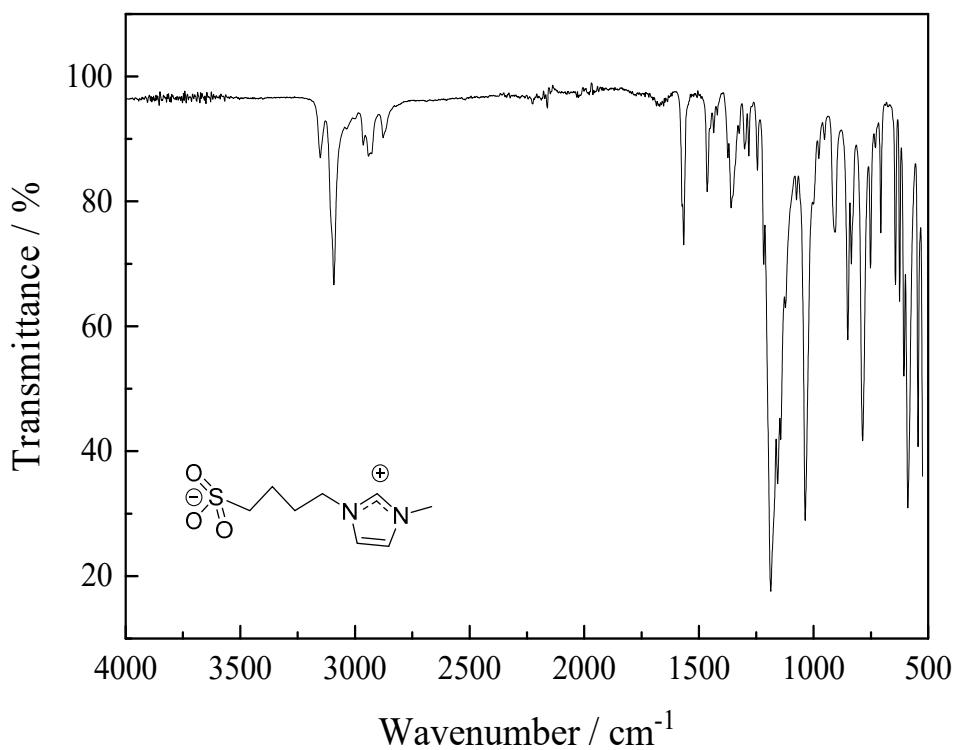


Figure S4. IR spectra of $\text{C}_1\text{C}_4\text{imSO}_3$

IR neat: 3158, 3114 (sym. stretching ν HC((4)C(5)H); 2959 (sym. stretching ν of imidazole ring ν CH_3 HCH); 2954, 2922 (asym. stretching ν C-H of $-\text{CH}_3$ and $=\text{CH}_2$); 2949 (asym. stretching ν CH_2); 1642 (stretching ν C=C); 1575 (in-plane ν of imidazolium ring); 1460 (C-N stretching of imidazole ring); 1450 (from methyl group, C-H bending); 1338 (SO_3^- assymmetric stretching); 1176 (skeletal ν of imidazolium ring); 1042 (out-of-plane C-H bending ν of imidazole ring); 974 ($-\text{S}=\text{O}-$ stretching); 651 (C2-N-C5 bending ν of imidazole ring); 622, 607 (CCC in-plane ring bending).

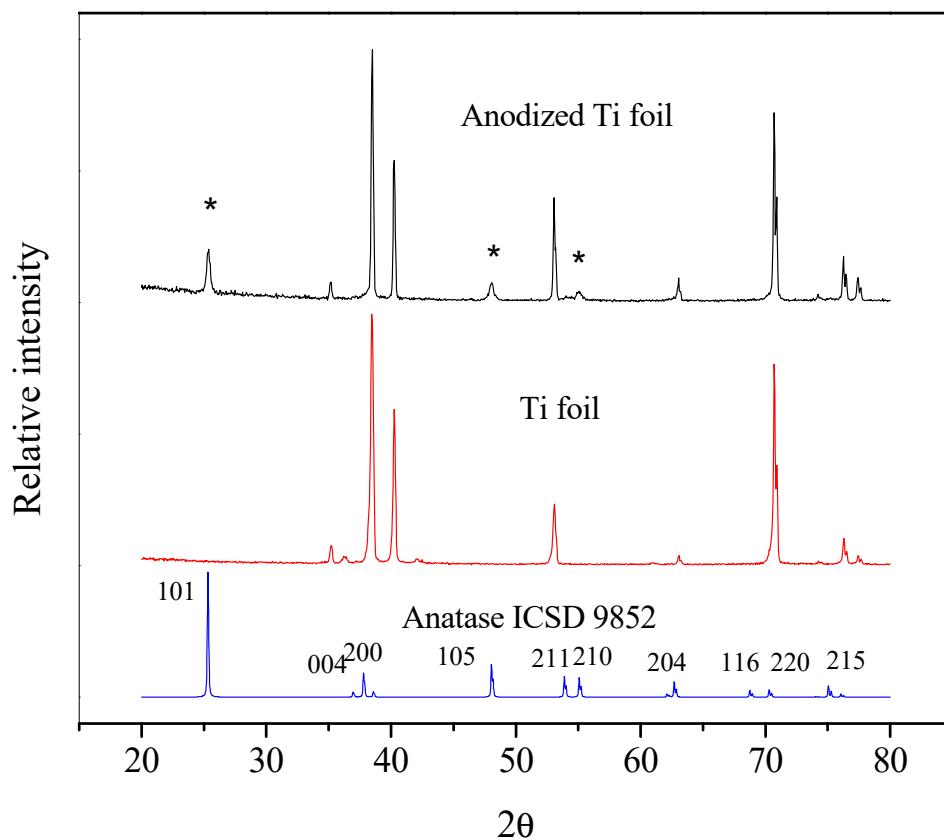
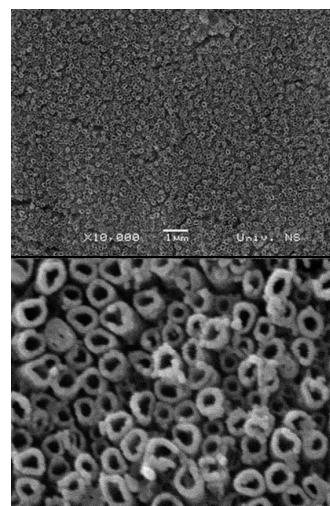


Figure S5. X-ray diffraction data for anodized Ti foil, Ti foil and anatase TiO_2 .



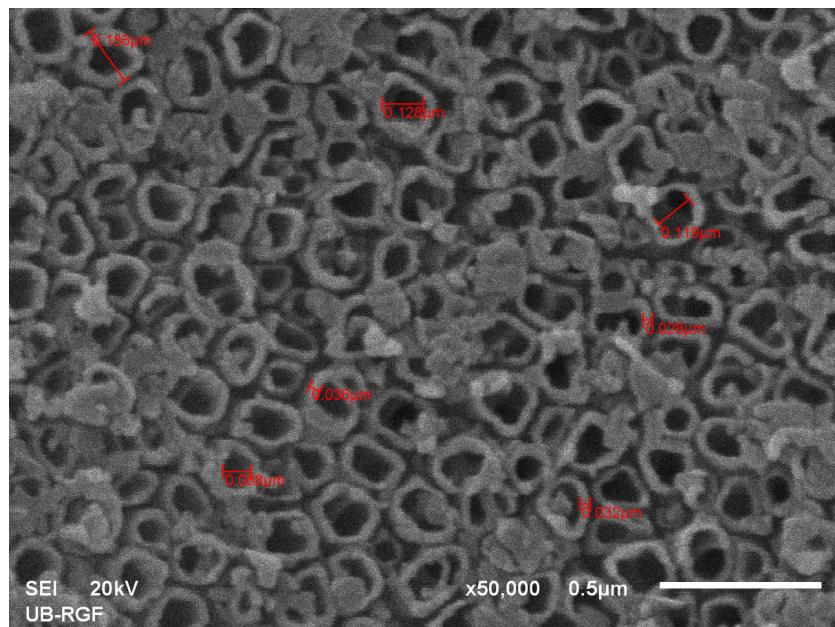


Figure S6. SEM images of TiO₂ nanotubes prepared by anodic oxidation (30 V, 6 hours) of Ti foil, with the thickness of the nanotube wall and the diameter of the nanotube opening.